CASE REPORT

Gastric antral vascular ectasia: maintenance treatment with oestrogen-progesterone

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Abstract
Gastric antral vascular ectasia (‘watermelon stomach’) is a rare cause of chronic gastrointestinal bleeding and various medical and surgical treatments have been described. We report a patient in whom an oestrogen-progesterone preparation successfully controlled recurrent blood loss.

Case report
A 73 year old woman was referred for investigation of recurrent occult gastrointestinal bleeding. She was first found to have a microcytic anaemia when aged 67 years and was treated with iron replacement without further investigation. She remained on iron and was well until 1989, when she again presented with symptomatic anaemia (haemoglobin 5·4 g/dl) and was admitted to her local hospital.

Investigations confirmed iron deficiency and faeces were persistently positive for occult blood. Coagulation studies were normal and there was no evidence of haemolysis. Urea, electrolytes, and liver function tests were all within normal limits, as were serum vitamin B12 and red cell folate. Upper gastrointestinal endoscopy on three occasions showed appreciable antral erythema; endoscopic biopsy specimens revealed no abnormality. Colonoscopy, a small bowel barium enema, and jejunal mucosal biopsy specimens were all normal. What was thought to be antral gastritis was treated with a variety of drugs including ranitidine, omeprazole, antacids, sucralfate, and tripotassium dicitratabismuthate (DeNol). The patient’s gastrointestinal blood loss continued, however, and she became dependent on transfusions, receiving 60 units of blood over the course of a year.

On assessment at Hammersmith Hospital in February 1989 her only symptom was recurrent fatigue when anaemic. She also had thyrotoxicosis and non-insulin dependent diabetes mellitus for which she took carbimazole and tolbutamide respectively. She did not drink alcohol or take non-steroidal anti-inflammatory drugs. On examination she had a goitre but no other abnormal findings. Investigation confirmed iron deficiency with occult blood loss, and showed antibodies to gastric parietal cells, with a titre of 1:160. At gastroscopy a series of longitudinally arrayed red streaks were seen radiating to the pylorus – the typical appearances of antral vascular ectasia (‘watermelon stomach’). The diagnosis was confirmed histologically: biopsy specimens from the gastric antrum showed dilated blood vessels within the lamina propria (some containing fibrin thrombi) together with prominent vertically orientated muscle fibres (Fig 1). Immunocytochemical staining for gastrin and somatostatin cells and with chromogranin showed normal antral neuroendocrine cell populations, biopsy specimens from the gastric body showed atrophic gastritis.

A technetium labelled red cell scan did not show a source of gastrointestinal blood loss. Inferior and superior mesenteric and coeliac axis angiography was performed. The latter showed considerable hypervascularity of the gastric antrum with early venous drainage, also seen on a selective common hepatic arteriogram (Fig 2). No other abnormalities were seen. Secretory studies disclosed hypochlorhydria with a basal acid output of 0·2 mEq/hour and a peak acid output after pentagastrin stimulation of 3·0 mEq/hour (off therapy). The fasting plasma gastrin concentration was correspondingly high (293 pmol/ml, normal <40). Prednisolone
therapy, initially at a dose of 30 mg, successfully stopped the bleeding (Fig 3) and other drugs were withdrawn apart from carbimazole and tolbutamide. Prednisolone also restored the gastric acid secretion to normal (basal acid output 2-7 mEq/hour, peak acid output 14 mEq/hour) with a corresponding fall in gastrin to 70 pg/ml.

However, prednisolone caused hyperglycaemia, even when the dose was reduced to 10 mg per day. We therefore changed to a standard oestrogen-progesterone pill (Loestrin 30, Parke-Davis, Hampshire) containing ethinyloestradiol 30 µg and norethisterone 1-5 mg daily taken for three weeks each month. She has now been taking this for over one year. Apart from one blood transfusion (4 units) after an endoscopic antral biopsy she has maintained her haemoglobin concentration on iron alone over this period, showing a considerable reduction in gastrointestinal bleeding (Fig 3).

Discussion
We have described a patient with gastric antral vascular ectasia causing chronic severe gastrointestinal blood loss which required 60 units of transfused blood over one year. Therapy, initially with prednisolone and then with an oestrogen/progesterone pill, has successfully controlled the bleeding.

Gastric antral vascular ectasia (watermelon stomach) was first described in 1953 by Rider et al. and about 50 further cases have been subsequently reported. The term ‘watermelon stomach’ was coined by Jabbari in 1984 to describe the endoscopic appearances. In contrast to the telangiectasia of portal hypertension and Osler-Weber-Rendu there are no lesions in other parts of the stomach. Most of the patients are elderly women in whom a source of occult bleeding is eventually traced to the gastric antrum. The significance of the endoscopic appearances is often overlooked initially. Consequently, these patients tend to be extensively investigated before the diagnosis is reached. Angiography is usually normal but endoscopic biopsy specimens may be diagnostic.

The aetiology of the condition is obscure although the pathological changes found are similar to those of solitary rectal ulcer. The condition is associated with autoimmune conditions, especially atrophic gastritis, demonstrated by achlorhydria, hypochlorhydria, and antiparietal cell antibodies in about a quarter of cases. An increased number of antral neuroendocrine cells has been reported in one case but there was no obvious increase in the number of cells in our patient.

It may be that mucosal atrophy, commonly present in the elderly, exposes the telangiectatic blood vessels to the gastric lumen, making them prone to bleed. Arteriolar changes in the gastric mucosa have been observed in atrophic gastritis and may be of aetiological importance.

A number of treatments have been described for gastric antral vascular ectasia including antrectomy and therapeutic endoscopy using either laser or diathermy. In view of the extensive vascular abnormalities in our patient we did not feel that coagulation treatment was appropriate, owing to the tendency to scar formation. Follow up of patients treated in this way is limited and circumferential scarring leading to stenosis is a risk. As our patient was keen to avoid surgery we opted to treat her with corticosteroids. Three out of four patients with gastric antral vascular ectasia and atrophic gastritis have improved with prednisolone. The rationale for using prednisolone is based upon treating the associated autoimmune atrophic gastritis. Prednisolone has improved gastric mucosal histology and restored gastric acid secretion in some patients with pernicious anaemia, though it is possible that other actions of the corticosteroid may have reduced the bleeding – for example via vasoconstriction or by an anti-angiogenic effect.

Because of the side-effects of prednisolone we elected to use oestrogen-progesterone as a maintenance therapy to control the bleeding. Van Cutsem et al have recently reported the successful treatment of chronic bleeding from gastrointestinal vascular malformations with an oestrogen-progesterone preparation. In a double blind controlled trial blood requirements in patients with hereditary haemorrhagic
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