Pyloric stenosis associated Crohn's disease responding to adalimumab therapy

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Abstract
Gastroduodenal Crohn's disease (CD) is rare and the response to standard medical therapy is often poor. Anti-tumor necrosis factor therapy has revolutionised the treatment of CD. We present a patient with pyloric stenosis associated with CD which improved with Adalimumab therapy. We recommend considering anti-tumor necrosis factor therapy in symptomatic gastroduodenal CD.

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Key words: Pyloric stenosis; Crohn's disease; Anti-tumor necrosis factor therapy; Adalimumab

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INTRODUCTION
CD is a chronic inflammatory condition and can involve the gut from the lips to the anus. Gastroduodenal involvement is seen in 0.5%-4% of all patients with CD[1-4]. While most patients with gastroduodenal disease are asymptomatic[5], some of them do develop features of gastric outlet obstruction[6]. Historically, strictureing gastroduodenal CD has been treated with resection[6,7], gastrojejunostomy bypass[7] and strictureplasty[8]. These have been associated with recurrence and morbidity. Corticosteroids and parenteral nutrition have been reported to be effective in acute gastric outlet obstruction secondary to CD while both thiopurines and methotrexate have been effective in maintaining remission[9-11]. Endoscopic balloon dilatation as adjunct to aggressive medical intervention has been described but has high recurrence rates[12,13]. There have been case reports describing the efficacy of anti-TNF therapy in strictureing gastroduodenal CD[14,15]. The patients in these case reports went through the combined treatment with mesalazine, corticosteroids, proton pump inhibitors and total parenteral nutrition before being considered for infliximab infusions.

CASE REPORT
A 23-year-old woman presented to the Gastroenterology Clinic in September 2010 with a 11 mo history of worsening bloating, malodorous belching, vomiting, reflux symptoms, 3 kg weight loss and borborygmi. She was diagnosed to have Crohn's disease (CD) in 2005 and had undergone a subtotal colectomy with ileorectal anastomosis in 2006. The pathology of the resected specimen was in keeping with CD. She later developed perianal fistulating disease in 2007 which was treated with initial Seton insertion and later marsupialisation. She was intolerant of azathioprine, developing abnormal liver function tests and was on methotrexate 20 mg once a week orally since June 2008. She had also been treated for suspected small
intestinal bacterial overgrowth with intermittent antibiotics between October 2008 and August 2010, with no symptomatic improvement. During this time, her lower gastrointestinal and perianal disease was well controlled on methotrexate.

In August 2010, her C reactive protein started rising and reached 78 mg/L (normal range: 0-5 mg/L). Her faecal calprotectin was high at 1840 ng/g of faeces (normal: < 60 ng/g) in September 2010. However, a flexible sigmoidoscopy in September 2010 showed a normal looking mucosa up to 35 cm from the anal verge. Random biopsies from the rectum and remaining colon showed minimally active, mild colitis with granuloma formation consistent with CD. A gastroscopy done at the same time was inconclusive due to food residue suggesting delayed gastric emptying.

She had an magnetic resonance imaging (MRI) of her small bowel in October 2010 which showed gross gastric dilatation with possible duodenal stricture (Figure 1A). The majority of the oral contrast had not passed into her small intestine. The rest of the small bowel was slightly dilated in keeping with post colectomy appearance. A repeat gastroscopy, after a prolonged fast, in December 2010 showed reflux oesophagitis, a normal stomach, and significant pyloric stenosis. The pylorus was slightly dilated in keeping with post colectomy appearance. Random biopsies from the rectum and remaining colon showed minimal activity, mild colitis with granuloma formation consistent with CD. A gastroscopy done at the same time was inconclusive due to food residue suggesting delayed gastric emptying.

The pyloric inflammation was likely due to her CD. The pylorus was negotiated and the first and second part of duodenum appeared normal. The biopsies from the pylorus showed focally erosive active chronic antral gastritis with intestinal metaplasia with no evidence of helicobacter infection. The pyloric inflammation was likely due to her CD. She was discussed at a Multi Disciplinary Team meeting. The surgical consensus was that she would likely require a subtotal gastrectomy. She was offered a trial of an anti-tumor necrosis factor (TNF) alpha antagonist therapy; B: Significantly reduced stomach size but with some focal small intestinal dilatation.

Her repeat MRI scan showed that the gastric dilatation was significantly less (Figure 1B). The oral contrast was seen to be passing into the small intestine and colon. Though there were focal areas of small bowel dilatation, they were likely to be secondary to adhesions or fibrotic strictures. The patient continues to take Adalimumab fortnightly.

**DISCUSSION**

There has been one case report of a patient with isolated gastric CD with gastric outlet obstruction initially who went into remission with infliximab infusions. When her symptoms recurred, she was restarted on infliximab. However, the patient developed mild arthralgias and later acute infusion reaction. This led to the patient being started on Adalimumab in addition to azathioprine and prednisolone. However, there was no significant clinical improvement after 3 mo. Topical corticosteroid therapy in form of fluticasone inhaler was introduced. The patient improved subsequently and went into clinical remission. Duodenal CD has been treated successfully with a combination of azathioprine, adalimumab and pantoprazole. The patient had isolated non-stenotic duodenal CD.

This is the first reported case in which the patient with symptomatic pyloric CD has responded to Adalimumab therapy. This patient was on an apparently adequate doses of oral methotrexate, although drug absorption may have been suboptimal, her lower gastrointestinal and perianal disease were well controlled. However, this was apparently not enough to control disease activity in the upper gut. By opting for Adalimumab directly, we avoided the side effects of long term corticosteroids and the morbidity associated with total parenteral nutrition, frequent endoscopic dilatations, surgery and repeated admissions to hospital for infliximab infusions.

In conclusion, we report a novel and safe way of treating stricturing gastroduodenal CD with Adalimumab. Treatment can be imparted with minimal disturbance to patient’s routine life. Adalimumab merits consideration in symptomatic gastroduodenal CD.

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