Terminal Ileum Perforation as a Consequence of a Migrated and Fractured Oesophageal Stent

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Abstract
Covered self-expanding metallic stents are commonly employed to relieve malignant oesophageal obstruction. We report a case of a patient with oesophageal cancer treated by stent insertion and curative chemoradiotherapy with subsequent stent migration to the stomach. The stent fractured with the distal fragment migrating as far as the terminal ileum where it caused perforation of the bowel.

Introduction

Oesophageal carcinoma is often a terminal diagnosis with most cases only presenting with symptoms when the disease is at an advanced stage, by which time curative therapy is improbable [1]. The three-year survival rate is only 20% and the rate of incidence in the West has increased significantly in recent decades [1]. The most common presenting complaint is progressive dysphagia, which is addressed by relieving the oesophageal obstruction to allow normal feeding. Treatment can involve surgery, radiotherapy, chemotherapy, laser and endoscopic stent insertion, either exclusively or in combination [2].

Radiologically guided endoscopic stent insertion can be employed for symptom relief in palliative cases suffering from malignant oesophageal obstruction or as an adjunct to other treatment. The ease of use and the relative speed with which lumen patency is restored makes stent use appealing in such cases, however complications can occur, including migration, oesophageal perforation, haemorrhage, pain, occlusion and recurrence of dysphagia.

In patients with early or low-stage disease, surgery in combination with chemotherapy and/or radiotherapy offers potential for cure. At present, surgery is the ‘gold standard’ for curative therapy, however it is clear from cohort studies of patients receiving...
chemoradiotherapy as definitive treatment that there are long-term survivors: of patients receiving definitive chemoradiotherapy, 5-year survival rates of 6.5–49.2% have been demonstrated, as compared to 23–64.9% in patients having chemoradiotherapy and complete resection [3–7]. It is yet to be established how chemoradiotherapy alone compares to surgery in terms of curative potential because as yet there are no completed randomised controlled trials [3].

In patients undergoing definitive chemoradiotherapy and having a stent, migration is highly likely as the tumour bulk recedes. In our case, the patient had a covered metallic oesophageal stent inserted soon after diagnosis and definitive chemoradiotherapy achieving remission. The stent migrated to the stomach, then fractured. The distal fragment progressed throughout the small bowel and perforated the terminal ileum. This is a rare complication of oesophageal stenting in oesophageal cancer patients having definitive chemoradiotherapy. In those who achieve remission, prophylactic removal of a migrated and fractured stent is worthwhile to avoid potential complications as highlighted by this case.

Case Report

A 79-year-old female with squamous cell carcinoma of the lower oesophagus (T3 N0 M0) diagnosed twelve months previously had a covered self-expanding metallic oesophageal stent (Choostent; Solco Intermed, Seoul, Korea) inserted to relieve her obstruction. She refused surgery and therefore underwent a course of chemoradiotherapy. This successfully reduced the tumour bulk such that six months after insertion, the stent had become loose and migrated to the stomach. At follow-up chest x-ray stent fracture was noted but it was decided to leave it in situ. She subsequently presented three times with dysphagia secondary to oesophageal strictures, which were treated by balloon dilatation on each occasion.

Six months after the migration and fracture had been noted, she presented to the surgical admissions unit complaining of a two-day history of progressive lower abdominal pain and persistent nausea. Examination demonstrated tenderness and fullness of the right iliac fossa. She was tachycardic, pyrexial and had a raised white cell count (12.5 × 10^9/l) and C-reactive protein level (60 mg/l). A plain abdominal radiograph was performed which demonstrated the stent within the stomach, a fragment of stent in the right lower quadrant and loops of dilated small bowel suggestive of small bowel obstruction (fig. 1). A plain erect chest radiograph was unremarkable with no air under the diaphragm. A contrast-enhanced CT scan of the abdomen and pelvis demonstrated additional findings of free air in the peritoneal cavity, sub-acute small bowel obstruction proximal to the right lower quadrant stent fragment with a local perforation and free fluid in the pelvis (fig. 2, fig. 3). A midline laparotomy was performed and perforation of the terminal ileum with faeces and pus in the pelvis was demonstrated. A right hemicolectomy was performed and the remaining stent was retrieved through an anterior gastrotomy. Postoperatively the patient recovered and was medically fit for discharge 15 days later.

Discussion

Self-expanding metallic stents are widely employed to relieve malignant oesophageal obstruction because they are easy to use, relatively safe and effective in allowing patients to consume a normal diet [8]. Stenting allows rapid establishment of patency relative to the alternative therapies, which involve longer periods of treatment, recovery and morbidity. Although covered stents are associated with an increased risk of migration in comparison to non-covered stents, they are believed to be advantageous in patients with malignant obstruction as they stem the inward growth of tumour, thereby slowing re-obstruction [9]. Metallic stents expand to have larger diameters that are more permissive to solid foods and are slower to occlude. Plastic stents are more likely to migrate, as illustrated by Bastos et al., who described a case in whom a plastic stent came
to be expelled per rectum [10]. Kato et al. reported a technique of endoscopic clipping of the upper branch of the stent to the oesophageal mucosa in 9 patients in whom stent migration was not encountered [11].

Perforation secondary to stent migration is rare and we only identified two reports of this scenario. Henne et al. described a case in whom two stents used to relieve a recurrent malignant oesophageal stenosis migrated and caused perforation at the site of the side-to-side jejunoojejunostomy of an earlier Bilroth II [12]. Kim et al. reported a duodenal perforation secondary to a migrated stent [13]. Stent fracture is also a very rare complication. Dogan and Egilmez reported a case in whom the fractured distal segment migrated only as far as the stomach and remained there uneventfully until the patient died [14]. The authors of this report suggest that such a fragment should only be removed if symptomatic. Altiparmak et al. reported a fractured stent that migrated to the stomach and caused a gastrocolic fistula to develop. Endoscopic retrieval methods (using grasping forceps, balloon dilatation catheters and polypectomy snare) were unsuccessful, necessitating gastrotomy [15].

In our case, stent insertion was required to relieve malignant obstruction early prior to a course of definitive chemoradiotherapy. Patients whose disease responds well to chemoradiotherapy are prone to stent migration as the cancer bulk shrinks. Non-fractured migrated stents can usually be left safely within the stomach. Endoscopic methods of removing migrated stents have been reported, but they carry a considerable risk of complications, such as perforation, and can be technically very difficult and prolonged [16].

**Conclusion**

We would advocate the elective surgical removal of fractured oesophageal stents in patients who are in remission having had definitive chemoradiotherapy. This approach would prevent the occurrence of potentially fatal scenarios as illustrated by this case.
Fig. 1. Plain abdominal radiograph.

Fig. 2. Contrast-enhanced CT axial view at level T12 demonstrating the stent in the stomach.
Fig. 3. Contrast-enhanced CT axial view at level S3 demonstrating the stent fragment in the terminal ileum, with localised collection, dilated loops of small bowel, and free air in the peritoneal cavity.
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