Biodegradable stents for the treatment of bowel strictures in Crohn's disease: technical results and challenges

Authors
John Gásdal Karstensen1,2, Katrine Risager Christensen1, Jørn Brynskov1, Claus Ronholt1, Peter Vilmann1, Jakob Hendel1

Institutions
1 Gastro Unit, Division of Endoscopy, Copenhagen University Hospital Herlev, Denmark
2 Gastro Unit, Division of Surgery, Copenhagen University Hospital Hvidovre, Denmark
3 Gastro Unit, Division of Surgery, Copenhagen University Hospital Herlev, Denmark

Background and study aims: In patients with Crohn's disease, the idea of biodegradable stents for treatment of bowel strictures with limited effect of endoscopic balloon dilation is tempting and initial results have been promising. The aim of this study was to evaluate the technical and clinical success of biodegradable stents for treatment of inflamed Crohn’s strictures refractory to endoscopic balloon dilatation.

Patients and methods: Consecutive patients treated with biodegradable stents due to Crohn’s disease and inflamed bowel strictures refractory to endoscopic balloon dilatation were included. Technical and clinical success were evaluated.

Results: Six patients were included in the study. Technical success was obtained in five patients (83%); clinical success was limited to one patient (20%); failure was observed due to mucosal overgrowth (n=2), stent migration (n= 1), and stent collapse (n =1).

Conclusions: In Crohn’s disease, it is technically feasible to treat bowel strictures with biodegradable stents. However, we have stopped using biodegradable stents due to lack of clinical success and side effects such as mucosal overgrowth and stent collapse.

Introduction

Bowel obstruction due to intestinal stricture formation is a well-known complication of Crohn’s disease (CD). The recommended treatment of short strictures before surgical resection is endoscopic balloon dilation (EBD) combined with optimized medical therapy in case of significant active luminal disease [1]. However, 50% to 75% of patients experience recurrence of the stricture within 1 year after EBD [2–4]. In these patients, self-expanding metal stents (SEMS) could be a minimally invasive alternative to repeat EBD or surgery. Both covered and partially covered metal stents have been used clinically, but rather high rates of adverse events have been reported and removal of the stent within the first month after deployment is required to avoid intestinal impaction [5–7]. Biodegradable stents have recently been introduced in an attempt to overcome these drawbacks, and initial case reports and series have shown promising results with acceptable clinical results and no mucosal overgrowth [8–11]. However, the results from some of these studies have been criticized because they included patients naïve to EBD [12]. The aim of this study was to evaluate the feasibility and clinical success of biodegradable stents to treat inflamed CD associated strictures refractory to standard EBD.

Case reports

Patients
Between December 2011 and February 2015, six patients with known CD and inflamed bowel strictures were included in the study. The median age was 51 years (range 25 – 60) and four were female. Three out of six strictures were located at an ileocolic anastomosis, while the remaining were located in the duodenal bulb, the sigmoid colon, and the ascending limb of a J-pouch, respectively. The patients had been treated by a median of 5.5 (range 4 – 7) EBDs prior to stent placement. Table 1 summarizes patient demographics. The decision to treat the patients with biodegradable stents was made in multidisciplinary meetings. The reason for this decision was that the stents are CE approved, and this retrospective case series was not subject to approval by the local ethics committee.
Biodegradable stents
The biodegradable stents consisted of polydioxanone monofilament, which is expected to secure the integrity and maintenance of radial force for 6 to 8 weeks before degradation (SX-ELLA BD biodegradable stent; ELLA-CS, Hradec Králové, Czech Republic) (Fig. 1). The biodegradable stents were initially designed to treat esophageal strictures and intestinal use has been limited to the left side of the colon using the same delivery device. However, a custom-made system is now available to facilitate stent deployment in the proximal colon, ileum, or duodenum. In a few previous cases the stents were placed during fluoroscopic control via an overtube (TS13140, Fujifilm, Tokyo, Japan). The length of the stents was 6 cm with a diameter of 18 mm and flares of 25 mm. In one case a stent was custom-made for a long, narrow stricture [10].

Endoscopy
Standard gastroscopes (EG 29-i10, Pentax Medical, Tokyo, Japan) or colonoscopes (EC 38-i10L, Pentax Medical, Tokyo, Japan) were used for stent placement in this study. Endoscopic follow-up was performed after 1 and 3 weeks. In case of suspected complications or lack of clinical success, a follow-up examination was performed within 48 hours. Technical success was defined as deployment of the stent at the desired anatomic location, while clinical success was defined as six-months relief of obstructive symptoms.

Results

Technical success
Technical success was achieved in five patients (83%) (Table 2). No adverse events were registered in relation to stent deployment. In one patient with an anastomotic ileocolic stricture (Case 2), stent deployment failed because a sharp angulation of the colon made it impossible to deliver the stent via the overtube. Since this attempt, the patient has been managed with repetitive EBDs.

Clinical outcome
Clinical success was achieved in only one of five patients (20%) and even in that individual (Case 4) the stricture did not resolve and 8 months later, EBDs were needed to avoid bowel obstruction (Table 2). The stent migrated in one patient (Case 1), who
<table>
<thead>
<tr>
<th>No</th>
<th>Etiology</th>
<th>Site</th>
<th>Length of stricture (mm)</th>
<th>Route</th>
<th>Design of stent</th>
<th>Stent length (mm)</th>
<th>Stent diameter – body (mm)</th>
<th>Stent diameter – flares (mm)</th>
<th>Dilation before stent delivery</th>
<th>Technical success</th>
<th>Procedure time (minutes)</th>
<th>Stent migration</th>
<th>Follow-up period (months)</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>de novo</td>
<td>Duodenal bulb</td>
<td>4</td>
<td>Oral</td>
<td>Standard</td>
<td>60</td>
<td>20</td>
<td>25</td>
<td>No</td>
<td>Yes</td>
<td>35</td>
<td>Yes</td>
<td>42</td>
<td>Migration. Two additional EBDs were needed.</td>
</tr>
<tr>
<td>2</td>
<td>Anastomotic</td>
<td>Ileocolic</td>
<td>2</td>
<td>Anal</td>
<td>Standard, but custom-made delivery device</td>
<td>60</td>
<td>20</td>
<td>25</td>
<td>No</td>
<td>No</td>
<td>80</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>Anastomotic</td>
<td>Ileorectal</td>
<td>4</td>
<td>Anal</td>
<td>Standard</td>
<td>60</td>
<td>20</td>
<td>25</td>
<td>Yes</td>
<td>Yes</td>
<td>59</td>
<td>No</td>
<td>16</td>
<td>The stent collapsed. After four subsequent EBDs, the stricture was surgically resected.</td>
</tr>
<tr>
<td>4</td>
<td>de novo</td>
<td>Ascending limp from a J-pouch</td>
<td>10</td>
<td>Anal</td>
<td>Custom-made</td>
<td>150</td>
<td>15</td>
<td>18</td>
<td>No</td>
<td>Yes</td>
<td>34</td>
<td>No</td>
<td>15</td>
<td>Success, but after stent degradation three subsequent EBDs were needed.</td>
</tr>
<tr>
<td>6</td>
<td>de novo</td>
<td>Sigmoid colon</td>
<td>4</td>
<td>Anal</td>
<td>Standard</td>
<td>60</td>
<td>20</td>
<td>25</td>
<td>No</td>
<td>Yes</td>
<td>45</td>
<td>No</td>
<td>4</td>
<td>Severe hypergranulation. Sigmoid resection.</td>
</tr>
</tbody>
</table>

EBD, endoscopic balloon dilation
had to be managed by EBDs; however, despite the location of the stricture in the duodenal bulb, retrieval of the migrated stent was unnecessary because it degraded. In one patient (Case 3) with an inflamed anastomotic ileorectal stricture, the stent collapsed (Fig. 2a). Afterward, that patient needed repetitive EBDs before finally having surgical bowel resection (Fig. 2b and Fig. 2c). In two patients (Case 5 and Case 6) the stents had to be removed either endoscopically (Fig. 3a and Fig. 3b) or by bowel resection because of severe hypergranulation (Video 1).
stents in CD strictures were promising, with clinical success rates ranging from 70% to 83% with no cases of mucosal overgrowth. Unfortunately, we were unable to reproduce these results and beside one technical failure in relation to stent deployment, we experienced two patients with severe hypergranulation, one stent migration, and one stent that collapsed [8, 9]. A possible explanation could be that the patients included in our case study had received multiple EBDs due to refractory stricture formation and all of them had symptoms of bowel obstruction. This differs from the largest case series, where 36% were naive to EBD and only 45% had obstructive symptoms. Based on our experience, we have now stopped using this treatment modality and if endoscopic stenting is required, we will use SEMSs. Furthermore, because case series are often influenced by publication bias, we find that it is relevant to report our negative results.

In conclusion, the idea of biodegradable stents is tempting and deployment is technically feasible, but due to limited clinical success, we have stopped using this treatment modality until the stents can be modified to address the problems described in this case series.

Competing interests: None

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References