INTRODUCTION

Endoscopic radiofrequency therapy for gastro-esophageal reflux disease (GERD) has been proven to be successful. STRETTA is the only radiofrequency device licensed for this indication in the UK and is approved by The National Institute for Health and Care Excellence (NICE) for cases refractory to medical therapy.  

STRETTA improves the quality of life and reduces the need for anti-reflux medication in select patients, especially those with uncomplicated gastro-esophageal reflux disease (GERD). We aimed to review the outcomes of STRETTA in patients with medially refractory GERD, who had undergone previous gastric surgery.

Methods: This was a review of a prospective database in a British center. Since 2016, all GERD patients who underwent STRETTA and had a history of previous gastric surgery were studied (n = 11). Anti-reflux medication pre- and post-STRETTA was evaluated. The outcomes were assessed objectively by the change in anti-reflux medication and subjectively through a pre- and post-procedure GERD-health-related quality of life (HRQL) questionnaire.

Results: The median length of follow-up was 23 months. Nine patients demonstrated improved GERD-HRQL scores following STRETTA (82%). Of the 7 patients who underwent fundoplication, all reported improved symptoms, with 3 patients discontinuing the medication and 3 patients on a reduced dose of proton pump inhibitor. Four patients underwent surgery other than fundoplication, of which 2 reported improvement and discontinued the proton pump inhibitor. Two patients reported no improvement.

Conclusions: This study demonstrates that STRETTA is successful in reducing refractory GERD in patients with previous gastric surgery. The outcomes were comparable to published outcomes in patients with uncomplicated GERD with no previous history of gastric surgery.

Key Words: Endoscopic management; Gastro-esophageal reflux disease; Revisional anti-reflux procedure; STRETTA
Two prospective studies with a small sample size that reviewed STRETTA following anti-reflux surgery demonstrated good outcomes in patients with treatment-refractory GERD; 15/18 patients had improved outcomes, and 6/7 patients were satisfied with the results.

We aimed to review the outcomes of STRETTA at a specialist center in patients who had undergone previous gastric surgery and had GERD refractory to medication. This case series will add to the limited evidence for the use of STRETTA in patients with a history of gastric surgery, having GERD refractory to medication.

MATERIALS AND METHODS

This was a review of a prospectively maintained database at a tertiary referral center in the UK. In all, 196 patients underwent STRETTA since 2016. Of these, 11 patients who had undergone previous gastric surgery were identified. All patients were assessed by pre-operative endoscopy, 24-hour pH and manometry, and barium studies. Demographic data such as age, sex, and history of previous surgery were recorded in a prospectively maintained database. Anti-reflux medication before and after STRETTA were reviewed. The reported outcome of STRETTA was assessed objectively by the change in anti-reflux medication post-procedure and subjectively through a pre- and post-procedure questionnaire. The questionnaire used was the GERD-health-related quality of life (GERD-HRQL), which has been validated for assessing the severity of GERD. The pre-procedure GERD-HRQL was administered at the pre-procedure clinic appointment.

Due to the waiting period in the National Health Service, the time from this appointment to the actual procedure was variable. However, all patients were screened on the day of the procedure to confirm that the ongoing symptoms persisted. The post-procedure GERD-HRQL was administered at the post-procedure follow-up appointment, between 4 and 12 weeks after the procedure. The overall length of follow-up was determined by the time since the procedure as the medications post-procedure were reviewed on the same date, April 29, 2020 (follow-up range, 2–42 months). A local ethics committee approved the maintenance of the prospective database, and patients provided written consent for the treatment and study.

The indications for STRETTA in our center are in conformity with the NICE guideline according to which, if proton pump inhibitor (PPI) medication has failed to resolve the symptoms, endoscopic radiofrequency therapy may be used before or after the surgery.

RESULTS

Eleven patients were included in the case series. The median follow-up duration was 23 months (range, 2–42 months). Nine patients demonstrated improved GERD-HRQL scores following STRETTA (82%) (Fig. 1). Of these, 3 patients required no further anti-reflux medication. Before STRETTA, all 11 patients were taking a PPI. After the procedure, 5 of them were able to discontinue the PPI altogether, and 3 patients had a reduced dose of PPI.

Fundoplication was the most common intervention (Table 1). Of the 7 patients who had undergone a previous fundoplication...

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**Fig. 1.** Flowchart of the outcomes following STRETTA. GIST, gastrointestinal stromal tumor; LNF, laparoscopic Nissen fundoplication.
plication, all reported an improvement in symptoms (100%), with 3 patients discontinuing the medication completely (43%) and 3 patients on a reduced dose of PPI (43%). Only 1 patient with a history of fundoplication did not discontinue or reduce the PPI usage following STRETTA; however, the H2-receptor antagonist (H2RA) was no longer required. Endoscopic images before and after the procedure of a patient with a history of previous fundoplication are demonstrated in Fig. 2A, B.

Four patients underwent gastric interventions other than fundoplication (Table 2). Two of these patients reported an improvement (50%). Of these, 1 patient with a history of vagotomy and pyloroplasty discontinued the PPI altogether, and is now only on an antacid. The other patient, with a history of resection for a fundal gastrointestinal stromal tumor (GIST), reported an excellent outcome, with a significant improvement in symptoms. The endoscopic images of these patients are shown in Fig. 2C, D.

One patient, with a history of an Esophyx or transoral incisionless fundoplication (TIF), was only on PPI before STRETTA; however, following the procedure is on PPI and an antacid. The endoscopic images of this patient are shown in Fig. 2E, F. This patient underwent 2 additional revisional anti-reflux surgeries after EsophyX. Biopsies showed benign esophageal inflammatory changes without Barrett’s esophagus.

A patient with a history of gastric bypass, who showed no improvement on the subjective scoring system, now requires only a PPI on a PRN (pro re nata) basis, although he required regular PPI and antacids before the procedure.

The time from original gastric surgery to STRETTA varied from 2 years to 40 years. There was no link between the time since surgery and outcomes following STRETTA.

<table>
<thead>
<tr>
<th>Patient</th>
<th>Date of STRETTA</th>
<th>Age</th>
<th>Sex</th>
<th>Follow-up (mo)</th>
<th>Pre STRETTA meds</th>
<th>Post STRETTA meds</th>
<th>Reported outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>2016</td>
<td>57</td>
<td>M</td>
<td>42</td>
<td>PPI + Antacid</td>
<td>Reduced PPI dose, antacid stopped</td>
<td>Improved</td>
</tr>
<tr>
<td>2</td>
<td>2016</td>
<td>66</td>
<td>F</td>
<td>40</td>
<td>PPI</td>
<td>No medication dependence</td>
<td>Improved</td>
</tr>
<tr>
<td>3</td>
<td>2017</td>
<td>66</td>
<td>M</td>
<td>39</td>
<td>PPI</td>
<td>Reduced PPI dose</td>
<td>Improved</td>
</tr>
<tr>
<td>4</td>
<td>2018</td>
<td>44</td>
<td>M</td>
<td>23</td>
<td>PPI + Antacid + H2RA</td>
<td>Reduced PPI dose, antacid and H2RA stopped</td>
<td>Improved</td>
</tr>
<tr>
<td>5</td>
<td>2018</td>
<td>66</td>
<td>F</td>
<td>16</td>
<td>PPI + Antacid</td>
<td>No medication dependence</td>
<td>Improved</td>
</tr>
<tr>
<td>6</td>
<td>2019</td>
<td>62</td>
<td>F</td>
<td>7</td>
<td>PPI + Antacid + H2RA</td>
<td>No medication dependence</td>
<td>Improved</td>
</tr>
<tr>
<td>7</td>
<td>2020</td>
<td>54</td>
<td>F</td>
<td>4</td>
<td>PPI + Antacid + H2RA</td>
<td>PPI and Antacid the same, H2RA stopped</td>
<td>Slightly improved</td>
</tr>
</tbody>
</table>

H2RA, H2-receptor antagonist; PPI, proton pump inhibitor.

<table>
<thead>
<tr>
<th>Table 1. Patients Who Had Undergone Previous Fundoplication and Their Results following STRETTA</th>
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<tbody>
<tr>
<td>Patient</td>
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<tr>
<td>---------</td>
</tr>
<tr>
<td>1</td>
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<td>3</td>
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<td>4</td>
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GIST, gastrointestinal stromal tumor; H2RA, H2-receptor antagonist; PPI, proton pump inhibitor.
**Fig. 2.** Endoscopic images pre- and post-STRETTA. GEJ, gastro-esophageal junction; GIST, gastrointestinal stromal tumor.

<table>
<thead>
<tr>
<th>Procedure</th>
<th>Pre-STRETTA endoscopic appearance</th>
<th>Post-STRETTA endoscopic appearance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Previous fundoplication</td>
<td><img src="image1" alt="A" /> Loose wrap with the preserved angle of His, and a lax GEJ, with refluxing gastric contents (arrow)</td>
<td><img src="image2" alt="B" /> Spherical thermal lesions following STRETTA (arrow)</td>
</tr>
<tr>
<td>Resection of Fundal GIST with fundus scarring</td>
<td><img src="image3" alt="C" /> Fundal scar (arrow)</td>
<td><img src="image4" alt="D" /> Spherical thermal lesions following STRETTA (arrow)</td>
</tr>
<tr>
<td>Esophyx and two further revisional antireflux surgeries</td>
<td><img src="image5" alt="E" /> Esophyx fastener at GEJ (arrow), maintained angle of His, with laxity of GEJ, and prolapse of the gastric mucosa</td>
<td><img src="image6" alt="F" /> Ongoing prolapse of gastric mucosa, which may explain the patient's failure to improve. Spherical thermal lesions (arrow)</td>
</tr>
</tbody>
</table>
DISCUSSION

Symptoms of GERD are common in patients who have undergone previous gastric surgery. This might be due to non-resolution of the symptoms following an attempt at anti-reflux surgery or as a direct result of the procedure itself, such as in Roux-en-Y gastric bypass. In patients with no history of previous gastric surgery, STRETTA has shown a reduction in the dependence on PPI. However, in most patients, PPI cannot be completely discontinued. Considering PPI usage as a measure of success of STRETTA, only 73% of the patients in this series could discontinue or reduce PPI. A previous study that evaluated STRETTA in patients with no previous gastric surgery reported that 78% of patients could discontinue or reduce PPI. This suggests that patients who have undergone previous gastric interventions might have outcomes comparable to those who did not.

Laparoscopic Nissen fundoplication (LNF) has been considered as the gold standard intervention for GERD resistant to optimal medical therapy. However, it has been reported that 21% of the patients after LNF will have ongoing GERD symptoms requiring medication, and 17% will require revisional surgery. These patients face the prospect of lifelong dependence on medication or a second surgery. Revisional Nissen fundoplication is associated with a lower probability of success and a greater risk of morbidity. Alternatively, long-term anti-reflux medication usage is associated with a high cost and long-term risks such as infectious complications, cardiac complications, and nutritional problems. STRETTA is proven to be more cost-effective than surgical management. A study involving 7 patients with persistent reflux after anti-reflux surgery, showing that 85.7% of the patients were satisfied with the result following STRETTA. In the present case series, similar results were achieved. 100% of the patients who underwent fundoplication could discontinue or reduce the PPI medication following STRETTA. Overall, patients with failed previous Nissen fundoplication are likely to have a significant benefit from STRETTA as they can potentially reduce or discontinue PPI medications and can avoid a revisional surgery, which has a higher risk. We have demonstrated good outcomes in patients who have undergone previous fundoplication, and therefore STRETTA might augment the neo-GEJ following fundoplication.

Clinicians should remember changes in the gastric anatomy, and therefore physiology following gastric surgery might be the reason for the failure of STRETTA in these patients. For example, in patients who have undergone Roux-en-Y gastric bypass or pyloroplasty, GERD symptoms are likely to be due to bile reflux following changes in the anatomy of the distal stomach. Furthermore, the formation of the gastric tube during a gastric bypass obliterates the angle of His, which normally prevents reflux of bile into the esophagus. Although STRETTA is postulated to stimulate muscular hypertrophy, increasing the tone of the lower esophageal sphincter, it does not affect the angle of His. In the present study, we had a single patient with persistent symptoms following STRETTA after Roux-en-Y gastric bypass, although the medication requirements had decreased. However, in a study on outcomes of STRETTA following laparoscopic Roux-en-Y gastric bypass, 83% of the patients had complete resolution of symptoms. The sample size was small (n = 6); however, the patients were followed up for a mean of 20 months, by which time 5 patients were asymptomatic. Further research on STRETTA following gastric bypass surgery is necessary with larger sample size.

Only one patient in our data set had undergone a previous TIF or EsophyX, a novel alternative anti-reflux endoscopic procedure. The patient reported no improvement and required additional anti-reflux medication post-procedure. EsophyX is effective in managing GERD. In a study that followed patients for 6 years following EsophyX, 86% of the patients required a reduced dose of PPI, and around 50% of them had discontinued the PPI completely. In TIF, the stomach, and distal esophagus are pulled together into a roll to form a valve at the lower esophageal sphincter, held in place by multiple polypropylene fasteners. These fasteners might disrupt the delivery of energy during STRETTA, which might be responsible for the failure of STRETTA in this patient.

The small sample size is a limitation of this case series. Nevertheless, though STRETTA is not routinely used for recurrent or primary GERD after gastric surgery, we believe this is the second-largest cohort of patients on this topic. An additional limitation is the lack of post-operative pH or manometry studies to help establish reasons for the success or failure of the STRETTA procedure. These tests are not routinely performed postoperatively at our center. This is an initial study that demonstrates that STRETTA is safe and likely to be beneficial in this cohort; however, further research is necessary to predict treatment success. This includes correlation with additional measures for GERD-HRQL, such as symptom index or symptom association probability.

In summary, this case series demonstrates that STRETTA is successful in reducing refractory GERD in patients with a history of gastric surgery. The outcomes in this cohort of patients were comparable to published outcomes in patients with uncomplicated GERD who have not previously undergone a gastric intervention.

Conflicts of Interest

The authors have no potential conflicts of interest.
Funding

None.

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