Duodenal mucosal resurfacing: Multicenter experience implementing a minimally invasive endoscopic procedure for treatment of type 2 diabetes mellitus

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submitted 14.7.2020
accepted after revision 30.7.2020

ABSTRACT
Background and study aims Duodenal mucosal resurfacing (DMR) is an endoscopic procedure which improves insulin resistant metabolic disease, including type 2 diabetes mellitus (T2DM). The aim of this report was to evaluate the feasibility and procedural aspects of DMR and to provide more specific DMR procedural guidance for endoscopists.

Patients and methods In this international multicenter, prospective, open-label study, patients on oral anti-diabetic agents for treating T2DM underwent single DMR. DMR entails circumferential submucosal lifting followed by circumferential mucosal hydrothermal ablation using an over-the-guidewire balloon catheter for lifting and ablation. For the first 28 patients a dual catheter system was used. During the study, a new integrated catheter was developed which was used for the latter 18 patients. During DMR, procedure success (complete DMR: duodenal ablation length ≥9 cm) and procedure duration were captured.

Results Forty-six patients underwent DMR. Using the dual catheter system, a complete DMR was performed in 22 of 28 patients (79 %). In the next eighteen patients who underwent DMR with the integrated catheter, a complete DMR was performed in 15 of 18 patients (83 %). The integrated catheter facilitated the DMR procedure and resulted in a reduction in procedure time. A detailed table and video are provided for future endoscopists.

Conclusions In our multicenter study, DMR was found to be feasible in the hands of experienced endoscopists. The integrated DMR catheter was a welcome modification during the study, allowing for easier ablation administration. Further optimization of the technique would be valuable prior to widespread dissemination.

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Introduction

Worldwide, 387 million people are diagnosed with type 2 diabetes mellitus (T2DM), and the number continues to rise at an alarming rate in every country [1]. Despite lifestyle interventions and the increasing array of pharmacological therapies, most patients with T2DM fail to achieve established treatment goals to prevent complications [2]. Recently, minimally invasive endoscopic treatment options for T2DM have emerged. Evidence from bariatric surgery and the duodeno-jejunal bypass highlights the importance of the small bowel, especially the duodenum, in glucose homeostasis and metabolic regulation [3]. If contact between the duodenal mucosa, bile, and nutrients is prevented or altered, then insulin sensitivity improves and β-cell function increases [4–6]. These improvements are quickly reversed when the bypassed duodenum is re-exposed to nutrients, underscoring the gluoregulatory role of the duodenum [4, 7, 8].

Duodenal mucosal resurfacing (DMR) is a minimally invasive endoscopic procedure involving catheter-based hydrothermal ablation of duodenal mucosa [9]. DMR may offer a new minimally invasive treatment approach for insulin-resistant metabolic diseases, including T2DM, nonalcoholic fatty liver disease (NAFLD), and nonalcoholic steatohepatitis (NASH) [10–12]. The first multicenter study reported durable and substantial glycemic improvement in patients with T2DM as well as a decrease in liver transaminase levels [10] at 24 weeks and 12 months [11]. Here we report on and evaluate the procedural feasibility of DMR. More specifically, we elaborate on DMR catheter development and improvements and endoscopic procedure optimization. In addition, we provide tips and tricks for endoscopists in order to conduct a successful DMR procedure.

Patients and methods

Study design

We evaluated endoscopic feasibility data from the first open-label, single-arm, multicenter study (NCT02413567) conducted at seven clinical sites in the EU and Chile (Academic Medical Center of Amsterdam, Netherlands; Erasmus University Hospital Brussels, Belgium; Policlinico Gemelli, Catholic University of Rome, Italy; University College Hospital London, United Kingdom; CCO Clinical Center for Diabetes, Obesity and Reflux, Santiago, Chile; King’s College Hospital, London, United Kingdom; and University Hospital Leuven, Leuven, Belgium) [11]. At each site, a single endoscopist performed the DMR procedure after completing initial didactic and hands-on DMR training in a porcine model. The study protocol was approved by the independent ethics committees at each center. The study was conducted in accordance with Good Clinical Practice Guidelines and the Declaration of Helsinki.

Patients

Eligible patients had T2DM and used oral glucose-lowering medication, were aged 28 to 75 years, had a body mass index between 24 and 40 kg/m² and hemoglobin A1c (HbA1c) between 59 and 86 mmol/mol (7.5%–10.0%). Main exclusion criteria were Type 1 diabetes (including positive GAD antibodies), use of injectable glucose-lowering medication, previous gastrointestinal surgery that could affect the ability to treat the duodenum (eg, Billroth 2 or Roux-en-Y gastric bypass), history of pancreatitis, and upper gastrointestinal bleeding. The complete eligibility list is available in the recently published efficacy study [11]. Written informed consent was obtained from all patients at screening.

Endoscopic setting

The DMR procedure was performed under general anesthesia or deep sedation with propofol, according to local guidelines and the endoscopists’ preference. First, a screening gastroendoscopy was conducted to ensure there were no conditions that would impede the DMR procedure (eg, erosive esophagitis ≥grade C, gastroduodenal ulcers, varices, strictures, or telangiectasia). Next, the papilla of Vater was marked on the contralateral duodenal wall using argon plasma coagulation (APC) or a hemostasis clip, to mark the proximal margins of the intended ablation zone. Then, a guidewire (recommended: Jagwire 0.035” Stiff Shaft [Boston Scientific, Marlborough, Massachusetts, United States]) was placed past the ligament of Treitz to assist delivery of the DMR catheters. Fluoroscopy was used to verify the delivery and positioning of the guidewire.

Initial modifications of the DMR procedure

Prior to this multicenter study, three patients from the first-in-human study developed a duodenal stenosis within 6 weeks after the procedure [12]. The stenoses were resolved by endoscopic balloon dilatation without further sequelae. Root cause analysis revealed that overlapping ablations and ablation of non-lifted duodenal mucosa were the likely causes for development of these duodenal stenoses after DMR. Therefore, the following modifications were made to the DMR procedure prior to initiation of the multicenter study: 1) ablation was performed from proximal to distal (instead of distal to proximal as was done in the first-in-human study) to optimize endoscopic visualization and avoid overlapping ablation zones and minimize the length of time elapsed between submucosal injection and ablation; and 2) mucosal lifting was performed more extensively to avoid ablation of non-lifted mucosa. All participating endoscopists reached a consensus on the optimal DMR procedure during a joint meeting with hands-on training in a porcine model.

DMR procedure with double catheter system

Initially, DMR was performed with two catheters (Fractyl Laboratories, Inc., Lexington, Massachusetts, United States): a submucosal expansion (SE) catheter and a hot fluid (HF) catheter. The SE catheter was tracked over the guidewire into the horizontal part of the duodenum just distal to the clip/APC mark (papilla) to measure the inner diameter of the post-papillary duodenal lumen. The endoscope (pediatric colonoscope) was positioned just proximal to the balloon at the tip of the catheter. Then three vacuum-assisted needles around the tip of the SE catheter were used to inject saline into the submucosal space. Six to nine consecutive submucosal lifting cycles were
performed to ensure submucosal lifting of the postpapillary duodenal area targeted for ablation. Between each lifting cycle, the catheter and endoscope was positioned 1 to 2 cm distally. Then, the SE catheter was replaced with a HF catheter (with fixed balloon diameter of 19, 21, 23 or 25 mm based on the SE catheter measurements). Three consecutive non-overlapping hydrothermal ablations of the lifted area were performed with a 3-cm HF balloon under endoscopic visualization. Between each ablation cycle, the catheter and endoscope were positioned 3 cm distally. A complete DMR procedure was defined as three ablations corresponding with a 9-cm circumferential ablation of post-papillary duodenal mucosa.

DMR procedure with integrated catheter

During the course of this study, a single catheter (▶ Fig. 1) became available that integrated submucosal lift and hydrothermal ablation functions. The novel catheter eliminated the need for catheter exchanges during the procedure and helped ensure that ablation was performed immediately after submucosal lift of the same segment of duodenum (▶ Video 1). The DMR catheter was introduced over the guidewire and positioned just distally of the marking of the papilla in the duodenum. The tip of the endoscope (pediatric colonoscope) was positioned just proximal to the balloon guided by the reference markers on the catheter. The single integrated catheter has three vacuum-assisted submucosal injectors around a 2-cm ablation balloon at its distal end to enable lifting of the duodenal mucosa. To ensure optimal lifting prior to mucosal ablation, every second lifting cycle was followed by a circumferential mucosal ablation. The catheter was advanced proximally 1 cm distally after each lifting or lifting-ablation cycle. Five ablations were performed to achieve a total duodenal ablation length of 10 cm, defining a complete DMR procedure.

Post-procedural care

Patients were discharged within 24 hours after the procedure, depending on the local center’s guidelines. Prior to discharge, patients were prescribed a 2-week post-procedure diet that progressed from clear liquids to solid foods to facilitate mucosal healing. The follow-up protocol and efficacy outcomes can be appreciated from our recently published study [11].

Outcomes

Feasibility endpoints included successful completion of submucosal expansion and mucosal ablation, defined as three (double catheter system with ablation balloon of 3 cm) or five (single catheter system with ablation balloon of 2 cm) ablations and procedure time (calculated from the time of insertion of the endoscope until removal of the guidewire and catheter). Data are presented as mean (standard deviation [SD]) or percentage where applicable. We created a table with tips and tricks for future endoscopists to facilitate an optimal procedure.

Results

Procedure time

In total, 46 patients underwent DMR. The first 28 patients who received double-catheter DMR had a mean (SD) procedure time of 96 (34) minutes. Once available, the single catheter was used for all subsequent 18 patients, which reduced the mean DMR procedure time significantly to 71 (23) minutes (P = .031).

Successful completion

The DMR procedure was completed (i.e., 3 ablation zones totaling 9 cm) in 22 of 28 patients (79%) in the double-catheter group. The full DMR treatment was not performed in six patients, due to catheter failure (n = 4), tortuous anatomy (n = 1), and difficulty with catheter positioning (n = 1). In the single integrated catheter group, the DMR procedure was completed (i.e., 5 ablation zones totaling 10 cm) in 15 of 18 patients (83%). Full DMR treatment was not possible in three patients due to difficulty with catheter handling and positioning.

A stepwise explanation of the DMR procedure, including challenges and advice for how to overcome them, is provided in ▶ Table 1.
Table 1  Tips and tricks for a successful DMR procedure.

<table>
<thead>
<tr>
<th>Step</th>
<th>Challenges</th>
<th>Don’ts</th>
<th>Photo</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>The preferred patient position is left lateral. In centers where general anesthesia is used, patient position can be supine.</td>
<td>In case proximal looping is observed in the stomach or during passage into the duodenum, turn the patient left lateral/probe.</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>Introduction of the catheter</td>
<td>In case the guidewire loops in the cardia, retract the guidewire slightly without losing length in the jejunum, before the tip of the catheter turns to the fundus.</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>Lubricate the tip of the catheter with PAM or other lubrication agent.</td>
<td>Avoid looping of the guidewire at the cardia during catheter advancement over this guidewire.</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>Advance catheter along the greater curvature of the stomach until passage into the duodenum D1/D2.</td>
<td>Advance catheter along the greater curvature of the stomach until passage into the duodenum, then straighten the catheter and introduce the endoscope into the stomach to assist advancement of the catheter through the pylorus. Deflate the stomach as much as possible.</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>Advance endoscope by following the catheter along the greater curvature of the stomach and enter the pylorus.</td>
<td>Try not to disconnect the loop of the endoscope and the catheter during the procedure.</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>Position scope in duodenum just proximally of the single marker band on the catheter.</td>
<td>Do not advance the catheter past the clip, the endoscope will push the catheter even further to a too distal position in the duodenum. Withdrawal of the catheter will result in disconnecting the loop of the endoscope and catheter.</td>
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</tbody>
</table>

Guidewire loops in stomach

Guidewire retracted

Catheter enters duodenum with endoscope introduced

Follow catheter in greater curvature stomach

Black marking on ablation catheter aligned with clip
<table>
<thead>
<tr>
<th></th>
<th>Challenges</th>
<th>Don’ts</th>
<th>Photo</th>
</tr>
</thead>
<tbody>
<tr>
<td>7</td>
<td>Oriente and optimize X-ray so that the triangular tip of catheter is projected horizontal in X-ray image.</td>
<td>Suboptimal fluoroscopic support</td>
<td><img src="image1.png" alt="Suboptimal fluoroscopic support" /></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Optimal fluoroscopic support due to turn fluoroscopy unit</td>
<td><img src="image2.png" alt="Optimal fluoroscopic support due to turn fluoroscopy unit" /></td>
</tr>
<tr>
<td>8</td>
<td>Store first X-ray image.</td>
<td></td>
<td><img src="image3.png" alt="Balloon inflated just past clip for first lift" /></td>
</tr>
<tr>
<td></td>
<td>Start of DMR lifting and ablation cycles</td>
<td></td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>Lift 1</td>
<td></td>
<td><img src="image4.png" alt="First lift, just distal of clip (marking papilla)" /></td>
</tr>
<tr>
<td>10</td>
<td>Move scope and catheter for 2 duodenal folds distally (± 1 cm). In case the catheter/endoscope does not advance in the duodenum or it meets with too much resistance, undo the loop of endoscope and catheter in the stomach. Use the endoscope stiffener (if available) and gentle traction on the guidewire while advancing the scope and catheter. If this fails, change the patient’s position.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>11</td>
<td>Check with X-ray and compare with the previous stored image, and store this image.</td>
<td></td>
<td><img src="image5.png" alt="Catheter 1 cm distally of first lift to perform second lift" /></td>
</tr>
<tr>
<td></td>
<td>Challenges</td>
<td>Don’ts</td>
<td>Photo</td>
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<tr>
<td>12</td>
<td>Lift 2</td>
<td></td>
<td><img src="Second.lift.png" alt="" /></td>
</tr>
<tr>
<td>13</td>
<td>Check with X-ray for stable position after second lift and compare with previous stored image, and store this image.</td>
<td>When you need to reposition the scope or catheter, withdraw scope or catheter and advance until again forward movement occurs and both loops are restored in parallel.</td>
<td>Avoid withdrawal of endoscope or catheter separately during whole DMR session. This will uncouple the parallel loops in the stomach and scope and catheter will no longer move one on one.</td>
</tr>
<tr>
<td>14</td>
<td>Perform ablation 1.</td>
<td></td>
<td><img src="First.ablation.mucosa.gets.a.white.aspect.png" alt="" /></td>
</tr>
<tr>
<td>15</td>
<td>Move scope and catheter for 2 duodenal folds distally.</td>
<td></td>
<td><img src="Ablated.mucosa.after.deflation.balloon.png" alt="" /></td>
</tr>
<tr>
<td>16</td>
<td>Repeat steps 9 to 15 until 5 ablations are completed.</td>
<td>If a needle port disconnects from the balloon and kinks (visible on X-ray), further lifting cannot be performed adequately. Replace the dysfunctional catheter with a new catheter.</td>
<td><img src="Two.ports.slightly.deviating.(sign.of.disconnection).png" alt="" /></td>
</tr>
</tbody>
</table>

DMR, duodenal mucosal resurfacing.
Discussion
DMR is a promising minimally invasive endoscopic procedure that appears to elicit robust and sustained improvements in metabolic parameters after a single treatment in patients with T2DM. Data from the first international, multicenter study suggest that DMR is feasible, safe, and effective in patients with T2DM that is suboptimally controlled with oral glucagon-lowering medication with a considerable glucagon lowering effect and a reduction in hepatic transaminase levels [10, 11]. Reported AEs are reassuringly mild and consistent with what is observed with other general upper gastrointestinal therapeutic endoscopic approaches. No cases of duodenal stenosis were reported after the initial DMR procedure modifications prior to this study.

Conclusions
The goal of the multicenter study was to deliver a uniform DMR to the T2DM study population. During the study, the introduction of a new, single DMR catheter resulted in an overall reduction in procedure time and increased ability to perform a complete DMR procedure. DMR appears to be a safe endoscopic procedure which is feasible in the hands of experienced endoscopists. With the introduction of the integrated DMR catheter during this study, the system matured and is ready for more extensive studies and possibly also for the first clinical application. This study also raises the question of whether DMR could be performed during ERCP. Further studies are ongoing to determine the potential of this procedure in an endoscopic setting.

Competing interests
Dr. Rodriguez-Grunert has received honorarium for consultancy from Fractyl Laboratories. Dr. Galvao Neto reports honorarium from Fractyl Laboratories (proctor), GI Dynamics (proctor, scientific advisory board member), GI Windows (consultant), Apollo EndoSurgery (faculty in training courses, consultant, proctor), USGI (consultant, proctor), Colubris Mx (consultant, proctor), Keyron (scientific advisory board member), Ethicon EndoSurgery (faculty in training courses, consultant), Meditronics (faculty in training courses, consultant), Olympus (faculty in training courses, consultant). Dr. Hayee has received funding/grant support from Olympus, Fujifilm, Fractyl Laboratories and honorarium for consultancy from Takeda Pharmaceuticals UK, Fujifilm, Atlantic Pharmaceuticals, Endotools. Dr. Deviere has received funding/grant support from Fractyl Laboratories Inc. for IRB approved studies. Dr. Bergman received research support from Fractyl Laboratories Inc. for IRB approved studies and received a consultancy fee for a single advisory board meeting of Fractyl in September 2019. This study was funded by Fractyl Laboratories.

References