Effect of laparoscopic sleeve gastrectomy vs laparoscopic sleeve + Rossetti fundoplication on weight loss and de novo GERD in patients affected by morbid obesity: a randomized clinical study

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Abstract
Purpose To compare sleeve gastrectomy (SG) to SG associated with Rossetti fundoplication (SG + RF) in terms of de novo gastro-esophageal reflux disease (GERD) after surgery, weight loss, and postoperative complications.

Materials and methods Patients affected by morbid obesity, without symptoms of GERD, who were never in therapy with proton pump inhibitors (PPIs), were randomized into two groups. One group underwent SG and the other SG + RF. The study was stopped on February 2020 due to the COVID pandemic.

Results A total of 278 patients of the programmed number of 404 patients were enrolled (68.8%). De novo esophagitis was considered in those patients who had both pre- and postoperative gastroscopy (97/278, 34.9%). Two hundred fifty-one patients (90.3%) had completed clinical follow-up at 12 months. SG + RF resulted in an adequate weight loss, similar to classic SG at 12-month follow-up (%TWL = 35.4 ± 7.2%) with a significantly better outcome in terms of GERD development. One year after surgery, PPIs were necessary in 4.3% SG + RF patients compared to 17.1% SG patients (p = 0.001). Esophagitis was present in 2.0% of SG + RF patients versus 23.4% SG patients (p = 0.002). The main complication after SG + RF was wrap perforation (4.3%), which improved with the surgeon’s learning curve.

Conclusion SG + RF seemed to be an effective alternative to classic SG in preventing de novo GERD. More studies are needed to establish that an adequate learning curve decreases the higher percentage of short-term complications in the SG + RF group.

Keywords Rossetti sleeve · Nissen sleeve · GERD · Obesity

Introduction
Sleeve gastrectomy (SG) is the most performed bariatric intervention in the world [1]. A recent systematic review and meta-analysis reported an increase of 19% of gastro-esophageal reflux disease (GERD) after SG and a 23% rate of new onset GERD, with erosive esophagitis (EE) in 30% of patients, and Barrett’s esophagus (BE) in 6% [2]. The use of proton pump inhibitors (PPIs) increased in 38% of patients. Other authors pointed out a development of EE after SG in 15.5 to 66.7% of patients [3–9].

The cause of an increase in GERD after SG may be related to alimentary behavior and surgery itself. Changes in the anatomy of the esophago-gastric (EG) junction area, dissection of the angle of His and potential damage to the sling fibers [10], increased intragastric pressure [7] reduced compliance [11], and eventual vagus nerve damage [12] are all factors that may alter the physiological anti-reflux systems [2].

On the other hand, some authors reported an improvement of GERD after SG, explained by weight loss, which reduces the gastro-esophageal pressure, and by decreased acid production due to the reduced volume of the stomach [13, 14].
We began performing a combined SG + Rossetti anti-reflux fundoplication (RF) to improve both obesity and GERD [15] in our center in 2015. Sleeve associated with fundoplication has been described by other authors with some variations, mostly regarding the type of fundoplication [16–21]. In previous manuscripts, we described the feasibility and the efficacy of SG + RF [22, 23].

The purpose of this study was to compare the outcomes of standard SG and SG + RF, in patients without preoperative GERD, evaluated in terms of de novo GERD after surgery, weight loss, and postoperative complications.

### Materials and methods

#### Study design

The study was designed as a monocentric, two-arm (1:1), randomized, controlled clinical trial. All the surgical procedures were performed in a high-volume bariatric surgery center (Policlinico San Marco, GSD, Zingonia (BG), Italy), from May 2017 to February 2020. Patients were randomized, and data were collected in a prospectively held database and analyzed by another autonomous Research Center (Research Centre on Public Health (CESP) of the University of Milan-Bicocca, Monza (MB), Italy). The study was approved by the Ethics Committee and was conducted according to the Declaration of Helsinki.

Inclusion criteria were morbid obesity suitable for surgery according to the Italian Bariatric Society lines [24]; no preoperative typical or atypical reflux symptoms; no preoperative therapy with PPIs; and no esophagitis at the esophago-gastro-duodenoscopy (EGDS) when it was performed before surgery. The original protocol followed the 2016 Italian guidelines for bariatric surgery [24]; hence, the need for EGDS was assessed by clinical judgment. During the course of the study, we decided to perform EGDS preoperatively in all patients because it could add more objective information about GERD.

Exclusion criteria were clinical dysphagia; previous surgery for obesity or procedures on the EG junction; and any contraindication to laparoscopic surgery.

Eligible subjects were randomly assigned to one of the two arms according to the randomization code that was generated at CESP. In the experimental arm, patients underwent SG + RF. In the control arm, patients underwent standard SG.

Considering COVID-19 pandemic status, Italian Bariatric Society recommended performing only emergent and urgent surgery [25]. Therefore, in agreement with the statistician, the study was stopped in January 2020.

#### Surgical technique

Standard SG and SG + RF techniques were largely described in previous manuscripts [15, 22, 23]. In both interventions, four trocars were used: 10 mm left sub-costal (optic view), 5 mm epigastric (liver retraction), 5 mm right hypochondrium (left hand), and 15 mm mesogastrium (right hand). The dissection of the gastro-colic ligament began at 3–4 cm from the pylorus and reached the left diaphragmatic pillar. At this point, the two interventions differed. In the patients undergoing SG + RF, the phreno-esophageal membrane was dissected. A minimal opening of the posterior crura was made to allow the retro-esophageal passage of the fundus. A 1.5–2 cm long, floppy, 360° fundoplication was created over a 38Fr oro-gastric boogie, with two interrupted, non-absorbable, extracorporeal Roeder knots. No sutures on the esophagus were performed. After the formation of the fundoplication, the two procedures were continued and concluded in the same way. The stomach was sectioned over a 38Fr oro-gastric boogie with a linear articulable stapler (Tristaple Signia™ stapling system, Medtronic, Dublin, Ireland). The choice of the type of cartridge depended on the tissue thickness, as usual. The main difference between the two interventions was the fact that at the level of the fundoplication, a black cartridge was used. An indocyanine green (ICG) test was performed at the end of the SG + RF intervention, to check the correct vascularization of the wrap, using IMAGE, STM RUBINA™ system (KARL STORZ SE & CO. KG, Tuttlingen, Germany).

#### Postoperative management and follow-up

On postoperative day two, an upper gastro-intestinal (UGI) series with oral water-soluble contrast was performed. If negative, the patient started a liquid diet on the same day. If there were any signs of leakage after the UGI series, a computed tomography (CT) scan with oral water-soluble contrast was performed. If the postoperative course was regular, the patient was discharged on the third or fourth postoperative day. Patients had clinical follow-up at 1, 3, 6, and 12 months after surgery; some patients who had preoperative EGDS repeated the exam 12 months after surgery, but this was not specified in the initial protocol.

#### Statistical analysis

Descriptive statistics were used to summarize patient baseline characteristics and clinical data. Continuous variables were compared using non-parametric tests, Mann–Whitney–Wilcoxon test, or Wilcoxon signed-rank test for matched samples. The differences between groups were compared.
with Fisher’s exact test. A $p$ value $< 0.05$ was considered statistically significant. The analyses were performed with the software SAS version 9.4 for Windows.

**Results**

**Patient characteristics**

A total of 278 patients of the programmed number of 404 patients were enrolled (68.8%). The study was early stopped for COVID pandemic. One hundred thirty-eight patients were randomized in the experimental group and underwent SG + RF; the remaining 140 patients were randomized in the control group and underwent normal SG. All the patients enrolled in the study respect the inclusion criteria shown in Table 1. All the interventions were performed in laparoscopy. The mean operative time was not significant different between SG + RF and SG (47.4 ± 17.4 vs 48.4 ± 15.1 min, $p = 0.585$). 62.8% of the SG + RF were performed by an expert surgeon, who had performed this procedure more than 100 times, and the remaining 37.2% of the SG + RF were performed by the rest of the surgical equipe. Considering only the expert surgeon, the mean operative time for normal SG was 32.1 ± 4.9 min and 38.5 ± 13.1 min for SG + RF (+ 6.4 min in SG + RF, $p = 0.015$). Considering the rest of the surgeons of the equipe, the mean operative time for normal SG was 49.3 ± 14.9 min and 62.3 ± 12.9 min for SG + RF (+ 12.9 min in SG + RF, $p = 0.006 − 3$). The relatively small difference in global mean operative time for SG + RF and SG was related to the fact that most of the SG + RF were performed by the expert surgeon and most of the normal SG were performed by the rest of the equipe.

**New onset of GERD after surgery**

None of the patients had preoperative symptoms of GERD and none was on PPIs. None of the patients who had preoperative EGDS (224/278, 80.6%) presented with EE. 30/224 (13.4%) had a small sliding hiatal hernia (<2 cm). 97/278 (34.9%) patients had EGDS both pre- and postsurgery at a mean follow-up of 14.7 ± 5.4 months in SG group and of 16.9 ± 7.3 in SG + RF group. De novo esophagitis was considered in those patients who had both pre- and postoperative gastroscopy (33.6% in SG group and 36.2% in SG + RF group).

New onset GERD was higher in the SG group than in SG + RF group (Table 2), with a significantly higher rate of patients using PPIs 1 year after surgery (17.1% vs 4.3%, $p = 0.001$). The incidence of EE after SG was higher than after SG + RF (23.4% vs 2.0%, $p = 0.002$). In the SG group, the patients had grade A and grade B EE (54.4% and 45.5%). In the SG + RF group, only 1 patient, who did not respect the dietary indications, had grade A EE. GERD symptoms did not necessarily follow the finding of EE. In the SG group, only 5 out of the 11 patients with esophagitis referred GERD symptoms (45.5%). The only patient with EE in the SG + RF group did not report symptoms. On the contrary, 13 of the total of 84 patients without esophagitis reported reflux symptoms 1 year after surgery, 8 in the SG group and 5 in the SG + RF group.

The finding of postoperative EE was not related to the preoperative endoscopic finding of small sliding HH (< 2 cm). Of the 12 patients with HH in the SG group, only one had postoperative grade B esophagitis, and among the 18 patients with HH in the SG + RF group, none had postoperative esophagitis. HH was not surgically repaired because it was < 2 cm and asymptomatic.

**BMI variations and comorbidity improvements**

Two hundred fifty-one patients out of 278 (90.3%) had completed clinical follow-up at 12 months. One year after surgery, both groups reached a mean BMI < 30 kg/m² (Fig. 1), with a %TWL > 20%, which is considered adequate [26]. The SG group had a %TWL of 35.4 ± 7.2%, with a mean BMI at 12 months of 29.1 ± 5.8 kg/m². The SG + RF group had a %TWL of 32.2 ± 7.6%, with a mean BMI = 29.4 ± 5.0 kg/m². Obstructive sleep apnea syndrome (OSAS), type 2 Diabetes (T2D), and hypertension (HTN) improved significantly 1 year after surgery (Fig. 2).

**Complications**

Patients in the SG + RF group had a longer hospital stay than those in the SG group (Table 3). The percentage of re-intervention for early and late complications was not different in the two groups. Both groups had a 0.7% of bleeding that needed laparoscopy and hemostasis on day 2 after surgery. Leakage, a well-known complication after sleeve, appeared in 1 out of 140 SG patients (0.7%), and never appeared in those with SG + RF. However, wrap perforation complicated 6 out of 138 patients who underwent SG + RF (4.3%). It did not occur in the patients who had had SG ($p = 0.013$).

Leakage after SG was treated with an endogastric self-expandable metal stent (SEMS) placed endoscopically on day 5. On day 12, the SEMS was replaced, and a laparoscopic abdominal toilette and drainage was performed. The patient was discharged on day 32 after primary surgery. The SEMS was removed on day 39 with leakage resolution.

Wrap perforation was treated by laparoscopic valve resection. The mean time of presentation of wrap perforation was 2.4 ± 5.5 days. The diagnosis was performed by an abdominal CT scan with oral water-soluble contrast. The mean hospitalization time after revision surgery was 11.2 ± 2.6 days for 5 out of 6 patients (83.3%). One of the six patients had
a longer hospital stay because he developed leakage after valve resection and needed SEMS placement on day 5. The SEMS was replaced on day 20 and was definitively removed on day 38 with leakage resolution.

Regarding late complications, one patient in the SG group needed a laparoscopic conversion to Roux-en-Y gastric bypass (RYGB) 28 months after primary surgery because of weight regain and GERD. In the SG + RF group, one patient needed a surgical revision 24 months after primary surgery because of valve disruption and weight regain, and underwent conversion to RYGB. Another patient had weight regain and underwent laparoscopic valve resection 28 months after primary surgery.

**Conclusions**

After a year of follow-up, SG + RF was effective both in terms of weight loss (Fig. 1) and prevention of de novo GERD (Table 2). A previous manuscript reported an improvement of preoperative GERD and esophagitis in patients who underwent SG + RF [22].

![Figure 1](image-url)
SG + RF had a longer operative time than normal SG: a mean of +6.4 min when SG + RF was performed by an expert surgeon in both bariatric and GERD surgery (with already more than 100 SG + RF procedures), and a mean of +12.9 min when SG + RF was performed by the rest of the surgical equipe.

The group of patients who underwent SG + RF had a longer hospitalization than those who underwent SG (Table 3): respectively 3.9 ± 4.0 days vs 3.1 ± 0.5 days (p = 0.018). This was related to a higher frequency of wrap perforation after fundoplication (4.3%) than the classic leakage following normal SG (0.7%). Wrap perforation differed from leakage after SG and never occurred along the staple line. It seemed to be related to traumatism of the gastric serosa due to excessive manipulation of the fundus during fundoplication, and usually appeared within the first 2–3 days after surgery. The increased pressure in the wrap during feeding might overwhelm the serosa resistance. Perforation could also be associated to ischemic issues. For this reason, at the end of every fundoplication, the wrap was checked with ICG test. Presentation was usually pain irradiated to left shoulder, fever, and leukocytosis. CT scan with oral water-soluble contrast was diagnostic for perforation and peri-gastric collection. The therapeutic approach was laparoscopic resection of the valve associated with a toilette of the abdominal cavity. The rate of wrap perforation (4.3%), 6 times higher than classic leakage (0.7%), could be alarming, but it decreased with the learning curve of the surgeon (Fig. 3). In fact, during the last year of the study, from January 2019, no valve perforations were reported. In this period, 38 SG + RF were performed (27.5% of the total). Moreover, the hospital stay after wrap perforation and fundus resection was shorter (11.2 ± 2.6 days) than the postoperative course after leakage in SG (39 days).

No classic leakages were seen after SG + RF. This could be explained by the valve protecting the higher part of the stomach. After revision surgery for wrap perforation, one leakage was reported.

| Summary statistics | SG (n 140) | SG + RF (n 138) | p-value |
|-------------------|------------|----------------|---------|
| Hospital stay (days) | Mean±SD   | 3.1±0.5 | 3.9±4.0 | 0.0181* |
| Early reintervention (< 1 month) | %, n/Pts | 1.4%, 2/140 | 5.2%, 7/138 | 0.0768 |
| Bleeding | %, n/Pts | 0.7%, 1/140 | 0.7%, 1/138 | 1.0000 |
| Leakage | %, n/Pts | 0.7%, 1/140 | 0.0%, 0/138 | 0.3310 |
| Wrap perforation | %, n/Pts | 0.0%, 0/140 | 4.3%, 6/138 | 0.0133* |
| Late reintervention (> 1 month) | %, n/Pts | 0.7%, 1/140 | 1.4%, 2/138 | 0.5687 |
| Weight regain | %, n/Pts | 0.0%, 0/140 | 1.4%, 2/138 | 0.1608 |
| GERD | %, n/Pts | 0.7%, 1/140 | 0.0%, 0/138 | 0.3310 |

SG sleeve gastrectomy; RF Rossetti fundoplication; GERD gastro-esophageal reflux disease; n/Pts number of patients out of the total; *statistically significant

Fig. 2 Variations in co-morbidities measured as the necessity to assume at least 1 drug for pathology. The follow up was completed for 126/140 (90%) in the SG group and 125/138 in the SG + RF group (90.6%). T2D = type 2 diabetes, HTN = hypertension, CPAP = continuous positive airways pressure.
While conducting this study, other than the efficacy of the surgical procedure itself, we noted the importance of the preoperative EGDS, even if it is not considered mandatory in the principal guidelines for bariatric surgery [24, 28, 29]. We found it essential for establishing the presence and the gravity of esophagitis, according to the LA classification. Moreover, from the literature, we know that more than 25% of patients without GERD symptoms may have unknown esophagitis [8]. The preoperative data about the condition of the EG junction mucosa may lead to the surgical choice of adding a fundoplication to SG, improving the postoperative condition of the patient.

Esophageal manometry associated with EGDS may be the best way to select the best treatment for each patient, since 76.6% of the patients who underwent SG did not have GERD after 1 year. For this reason, it is important to select that 23.4% of patients affected by obesity without preoperative GERD who may benefit from the association between SG and fundoplication, to prevent de novo GERD. SG + RF may be the correct intervention for patients with preoperative EE (> B grade), and, to avoid de novo GERD, for those without GERD who have a hypotonic lesser esophageal sphincter (LES) at the manometry. Patients without GERD and with a normal LES may be the best candidate for normal SG.

The limits of this study are the lack of manometric and 24 h-PH metry studies in describing GERD. The role of 24 h-PH metry in bariatric patients is still debated [30]. In fact, it is more complicated to perform and require a higher level of patient compliance.

Further studies to associate preoperative EGDS with manometric analysis are ongoing. These could be an important diagnostic aid for correctly deciding the best intervention for each patient, in order to provide a tailored approach. Further studies with longer follow-up are needed to evaluate long-term maintenance of weight loss and GERD improvement in patients undergoing SG + RF.

Declarations

Ethics approval  All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Informed consent  Informed consent was obtained from all individual participants included in the study.

Conflict of interest  All the authors declare no competing interests.

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