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CLINICAL REVIEW

Endo-bariatrics: A New Paradigm for Weight Loss

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Introduction

Obesity is a well-recognized disease that has reached pandemic proportions. Approximately 30% of the world population is over-weight or obese, and every year, almost 3 million patients die from complications of obesity. In the United States, a significant percentage of adults are obese, but also 17% of children suffer from obesity. Obesity and its complications represent a substantial burden on healthcare, estimated at 14.3% of the US healthcare spending, corresponding to \$427.8 billion and an incremental cost of \$1,429/obese person per year.^{1,2}

Only recently, in June 2013, was obesity considered a chronic disease by the American Medical Association.³ Multiple high-quality studies have shown associations between increased body-mass index (BMI) and multi-system conditions including coronary artery disease, type II diabetes, dyslipidemia, hypertension, stroke, atrial fibrillation, and osteoporosis.⁴⁻⁸ Non-alcoholic fatty liver diseases (NALFD) is often considered the hepatic manifestation of obesity and metabolic syndrome.⁹ Obese individuals who smoke have a markedly reduced life expectancy compared to non-obese smokers.¹⁰ Nearly 3.6% of all cancer cases in 2012 were attributable to obesity. These cancers include colorectal, esophageal, gallbladder, pancreas, liver, endometrial, postmenopausal breast, and thyroid cancer.¹¹⁻¹³

Management of obesity has traditionally revolved around two approaches: noninvasive weight-loss strategies such as dietary and lifestyle changes, and invasive options represented in bariatric surgery. Despite the superiority of bariatric surgery in achieving weight loss, only 2% of eligible patients undergo surgery due to patients' preference, increased cost, and limited access.¹⁴

As a result, a considerable gap exists in the treatment of obesity and its complications, and effective, less-invasive approaches are critically needed.

Diet and Lifestyle Changes

Although considered the first line of therapy, both diet and lifestyle modification have low efficacy in achieving significant weight loss. Approximately 3% of patients reach their target weight with dietary changes. The well-known limitation to any dietary therapy is modest weight loss and challenges of long term adherence to the diet. Furthermore, the optimal diet to achieve weight loss remains unknown.

Very low-calorie ketogenic diet (VLCKD) for weight loss has been studied for many years. In a meta-analysis of twelve studies, 10.0 Kg of total body weight loss (TBWL) was achieved with VLCKD after up to 4 weeks and was sustainable on follow up after two years. This diet was also associated with improvement in waist circumference (-12.6 cm), hemoglobin (Hb)A1C (-0.7%), total cholesterol (-28 mg/dl), triglycerides (-30 mg/dl), AST (-7 U/l), ALT (-8 U/l), GGT (-8 U/l), systolic and diastolic blood pressure (-8 and -7 mmHg, respectively).¹⁵

High-intensity interval training (HIIT), a type of exercise that involves short bursts of high-intensity exercise interrupted by light exercise or recovery periods, has been shown to lower triglycerides and fasting glucose, systolic and diastolic blood pressure, decrease oxidative stress and inflammatory markers and improve cardiac function.¹⁶ However, no difference was found in BMI or weight loss with HIIT when compared to moderate-intensity continuous training.^{17,18}

Intermittent fasting (IF) can be defined as alternating periods of eating and fasting, although no single quantitative description exists. Several different forms of IF have been described: alternate-day fasting, alternate-day modified fasting (ADMF), 5:2 diet, fasting-mimicking diet, and time-restricted feeding. IF has shown benefits in rodent models, independent of calorie intake.¹⁹ Multiple studies have focused on ADMF, described as alternating "fast" days (25% of daily calorie needs) with "feast" days (125% of daily calorie needs). Initial reports found that ADMF induces weight loss and improves glucose and insulin levels, blood pressure, lipids, and high-sensitivity C-reactive protein (hs-CRP) in humans.^{20,21} ADMF also reduces binge eating and depression in humans.²² However, the largest ADMF trial in humans showed ADMF is not superior to conventional diet for improving blood glucose, insulin, blood pressure, heart rate, lipids, visceral fat, hs-CRP, and homocysteine. Notably, the dropout rate for ADMF was higher than conditional diet (38% vs. 29%, respectively).²³

Pharmacotherapy

Patients are considered candidates for drug therapy after failure to achieve at least 5% of TBWL with diet and lifestyle interventions alone. Suitable candidates generally have BMI > 30 kg/m² or BMI > 27 kg/m² with related comorbidity. Lorcaserin was recommended first-line pharmacotherapy in non-diabetic

patients by some experts. It reduces appetite as a selective serotonin 2C receptor agonist.²⁴ The drug was initially approved by the US Food and Drugs Administration (FDA) in 2012, but was recently voluntarily withdrawn from the market due to concerns about long term adverse effects. Adverse effects (AEs) include headache, nausea, upper respiratory tract infection, pharyngitis, and dizziness. Caution should be taken with type 2 diabetes due to risks of hypoglycemia.²⁵

Phentermine is a noradrenergic, sympathomimetic amine that decreases appetite through its central nervous system (CNS) effects and the stimulation of the hypothalamus. It is one of the most commonly prescribed and least expensive drug therapies for weight loss.²⁶ Common AEs include tachycardia, hypertension, tremor, overstimulation of CNS, dry mouth, and constipation. A combination of extended-release topiramate and phentermine was approved by the FDA in 2012 for weight loss.^{27,28} Phentermine-topiramate can lead to 8-10% TBWL.²⁹ However, this drug combination should be avoided in patients with hypertension or coronary artery disease and is contraindicated in pregnancy.

Liraglutide is preferred in patients with type 2 diabetes. It is a chemically modified glucagon-like peptide-1 (GLP-1) agonist, stimulates insulin release from pancreatic islets,³⁰ and inhibits gastric emptying and reduces appetite through action on central satiety centers.³¹ When used at a higher dose (3mg daily injections) than usual doses in diabetes (1.8mg injection maximum daily dose), liraglutide was associated with a weight reduction of 2-4 kg compared to placebo.^{32,33} Saxenda is the trade name for the higher dose of liraglutide prescribed for weight loss. Liraglutide causes thyroid C-cell tumors in both rats and mice of both genders. No evidence of these tumors was found in human studies, and liraglutide is contraindicated in patients with a personal or family history of medullary thyroid carcinoma or multiple endocrine neoplasia syndrome types 2.^{32,34,35}

Bupropion-Naltrexone combination can be considered in obese smokers. Although the exact mechanism of action for weight loss is unclear, the combination is thought to work in the hypothalamus to promote satiety and inhibit the “reward system” that various food induce.^{36,37} Bupropion-Naltrexone leads to approximately 5% TBWL at 56 weeks.³⁸ After initial rejection by the FDA in 2011 for concerns of cardiovascular AEs,³⁹ the combination drug was approved in 2014 following additional clinical trials addressing its safety.⁴⁰

Orlistat is one of the earliest pharmacotherapies approved for weight loss (FDA approval in 1999 for obesity management).^{41,42} It works by inhibiting gastrointestinal lipase and reducing dietary fat absorption by approximately 30%. Compared to placebo, orlistat results in 3 kg average additional weight loss when combined with behavioral interventions, according to a meta-analysis of 12 trials.⁴³ Use is limited by side effects. Approximately 15-30% of patients taking orlistat experience abdominal cramps, flatus, and fecal incontinence.⁴⁴ Low levels of fat-soluble vitamins, especially vitamin

D, are common with Orlistat use. Acute kidney injury secondary to oxalate has been reported, although the incidence is rare.⁴⁵

In general, the efficacy of the current pharmacotherapy is limited to 5-10% TBWL in the majority of responding patients. It is important to recognize that individual responses to drugs vary widely. Weight regain is expected after medication discontinuation. Therefore, medications are usually used as an adjunct to lifestyle changes and sometimes adjunct to surgery or endoscopic interventions.

Endoscopic Interventions for Weight Loss

There are three main types of bariatric surgeries currently performed for weight loss. Roux-en Y gastric bypass (RYGB) is the most effective surgery, usually leading to 45%-55% TBWL. Laparoscopic sleeve gastrectomy (LSG) has been gaining more popularity over the last decade and has recently emerged as the most commonly performed bariatric surgery. It leads to approximately 25% TBWL. The adjustable gastric band is now less commonly used.

However, all current bariatric surgeries present multiple challenges. The post-surgical morbidity rate can be high at 3-20%. Mortality rates are approximately 0.1-0.5%. The need for re-operation is around 8%, and 15% of patients become malnourished. Additionally, 25% of patients regain weight following initial weight loss.³⁵

In a survey of 284 patients with a BMI >40 or a BMI >35 with obesity-related comorbidity, only 2% of eligible patients undergo surgery. Further questioning showed that half of the patients expressed a fear of operation, and 32% had a fear of dying.³⁶

Therefore, endoscopic bariatric (endobariatric) treatment is an attractive option. This approach is minimally invasive, scarless, can be performed in an outpatient procedure room, and is likely to be less costly than surgery. It is almost always reversible and repeatable. Most importantly, this approach may fill a large unmet need in the management of obesity. (Figure 1)

Endoscopic intervention for weight loss can be divided into stomach-focused interventions, and small bowel-focused interventions.

A. Stomach Focused Approaches

1. Endoscopic Sleeve Gastroplasty

Endoscopic sleeve gastroplasty (ESG) is a novel, incision-less, minimally-invasive procedure developed as a non-surgical alternative to sleeve gastrectomy for the management of obesity. The procedure incorporates applying intraluminal full-thickness sutures to plicate the stomach, resulting in significant shortening and reduction in gastric volume.

Evolution of the Procedure

Initial attempts of this procedure were reported in 2008, under the term endoluminal vertical gastropasty (EVG). This was performed using the EndoCinch® suturing device (C.R. Bard, Inc). Vogel et al. reported 58.1% ± 19.9% excessive weight loss (EWL) among the 64 patients who underwent the procedure, with no serious AEs. The suturing included the anterior wall and the posterior wall, but not the greater curve, which was later found to be an essential step for the success of the intervention.³⁷ Two years later, an updated version of the EndoCinch device was introduced under the name RESTORE® system (C.R. Bard, Inc). This was assessed in a pilot study (TRIM trial) of 18 patients with 27.7% ± 21.9% EWL at 12 months. The downside to the procedure was the loosening of plications in 13 out of 18 patients.^{38,39}

In 2012, the Apollo OverStitch® device (Apollo Endosurgery, Austin TX, US) was developed to help overcome the shortcoming of gastric plication mentioned above. This device allowed for full-thickness suturing that included the muscle layers. It was made possible with the use of a screw-like instrument (the tissue helix). (Figure 2) With the full-thickness sutures, an increase in tissue approximation durability was possible, allowing for better long-term results. Multiple patterns of suturing can be made, and this is the topic of ongoing debate. In general, running stitches, with 6–12 tissue purchase sites, are placed in a "U" shape or "N" shape at the anterior wall, greater curvature, and posterior wall. (Figure 3) Clinical experience has shown that incorporating full-thickness sutures in each site is critical to ensure the durability of the plication.⁴⁰

Safety of ESG

Mild AEs following ESG range from 5-30% and are nausea, vomiting, and abdominal discomfort, which are anticipated and can be well-managed or prevented with pharmacologic prophylaxis. The rate of serious events requiring intervention or hospitalization remains <2%. In a 2016 study, 1 of 91 (1.1%) patients developed post-procedural peri-gastric leak following dietary indiscretion and required antibiotics treatment.⁴¹ When using a different suturing system, there were no major AEs after a two year follow up in one study.⁴² However another study reported two patients (<2%) developed upper gastrointestinal bleeding, one of whom was on anticoagulation.⁴³ Overall, ESG is considered safe with reduced risk of major AEs as compared to other endobariatric options.

Efficacy of ESG

There are consistent outcomes across multiple studies showing that ESG leads to TBWL in the range of 15-20.9% at six months, 12 months and 24 months.⁴¹⁻⁴³ In a case-matched study, ESG resulted in significantly more TBWL at 12 months compared to high-intensity diet and lifestyle therapy (20.6% versus 14.3%, respectively, $P < 0.001$).⁴⁴ More recently, five-year outcome data report TBWL of 18.1%, 17.3%, 20.8%, and 18.7% at 1-year, 2-years, 3-years, and 5-years, respectively.⁴⁵

Importantly, ESG resulted in a significant reduction in metabolic comorbidities. In prediabetic and diabetic patients, HbA1C levels were reduced from an average of 6.6% to 5.6% one year following ESG. Some patients were able to stop insulin therapy. Systolic blood pressures decreased significantly by an average of 7mmHg and triglyceride levels by an average of 40mmol/dL after one year.⁴¹ Non-alcoholic fatty liver disease surrogates also improved following ESG.⁴⁶ These alterations in metabolic profiles with ESG are similar to previously published surgical series and suggest potential alternative benefits beyond weight loss.

Learning Curve for ESG

To achieve competency in performing ESG, endoscopists need experience in endoscopic suturing and plication. To date, there is one study involving one operator examining the learning curve to competently perform the procedure. Efficiency was reached at 38 cases and mastery at 55 cases.⁴⁷ Given the obesity epidemic, mastery of ESG with sufficient experience is possible and has the potential for wide adaptation in the care of bariatric patients.

PIVI Criteria

The Preservation and Incorporation of Valuable endoscopic Innovation (PIVI) thresholds, set jointly by the American Society of Gastrointestinal Endoscopy (ASGE) and the American Society for Metabolic & Bariatric Surgery (ASMBS), recommend efficacy targets of > 25% EWL at 12 months, and a safety threshold of < 5% risk of major complication for endoscopic bariatric treatments. ESG appears to meet these criteria.

2. Intra-gastric Balloon

Intra-gastric balloons (IGB) have been the major space-occupying stomach focused therapy for weight loss. Multiple balloons are currently available for commercial use and we discuss the commonly used IGBs:

ReShape Balloon: The ReShape Dual Integrated Balloon System (ReShape Medical) consists of two connected saline-filled spheres that are endoscopically placed and removed six months later. The device was FDA approved in 2015 following the REDUCE trial, a randomized sham-controlled trial comparing ReShape with diet and exercise against lifestyle modification alone.⁴⁸ ReShape subjects had 6.8% TBWL compared to 3.3% in the sham controls. However, AEs were seen in 7.5-75% of patients, largely due to accommodative symptoms. Gastric ulcerations were seen in 10.3% of patients with some improvement when decreasing balloon size. In 2018, ReShape Medical was purchased with plans for phasing out the balloon in favor of an alternative, the Orbera.

Orbera: The Orbera Intra-gastric Balloon System (Apollo Endosurgery) consists of a single saline-filled sphere that,

similar to ReShape, is endoscopically placed and removed six months later. It received FDA approval in 2015. In a multicenter randomized trial published in 2017, the Orbera arm achieved 10.2% TBWL compared to 3.3% in the lifestyle arm in 6 months.⁴⁹ Over 50% of patients had accommodative symptoms of nausea, vomiting, and abdominal pain. Due to device intolerance or per patient request, 18.8% of patients had the Orbera removed early. The Orbera balloon is currently being investigated as bridging therapy to bariatric surgery in superobese patients.

Obalon: The Obalon Balloon System (Obalon Therapeutics) consists of three gas-filled balloons that are swallowed as deflated capsules. Placement of the capsules is confirmed under fluoroscopy and then inflated with a gaseous mixture with eventual removal after six months. The Obalon was FDA approved in 2016. The SMART trial, a randomized sham-controlled trial, showed an average 6.6% TBWL in the treatment arm after two years versus 3.4% TBWL in the control arm.^{50,51} Patients who completed at least twenty weeks of the balloons in place achieved a mean 10.0% TBWL.

Other intragastric balloons are under clinical investigation. The Elipse balloon (Allurion Technologies) differs in design compared to the other balloons. It is a swallowed saline-filled balloon that self-deflates and eventually excretes through the GI tract in roughly 16 weeks. The Sptaz3 Adjustable Balloon System (Spatz FGIA) is a fluid-filled balloon placed endoscopically, and its size can be further adjusted endoscopically in response to intolerance or weight loss.⁵²

3. *Aspire Assist Device*

The Aspire Assist device ® (Aspire Bariatrics) consists of the placement of a large percutaneous gastrostomy tube, subsequently connected with a skin port to the external part of the device. Patients use this system to siphon off a portion of the ingested meal, typically one-third of the volume around twenty minutes after ingesting the food. The mechanism is an alternative to space-occupying therapies through partial disposal of ingested food to decrease the caloric burden following a meal. Unlike the standard PEG tubes, the Aspire requires both endoscopic placement and removal (Figure 4). The pilot study of aspiration therapy showed a 49% EWL after one year without any AE on eating behavior or compensatory eating.⁵³ A 52-week clinical trial showed a mean 31.5% TBWL with Aspire Assist compared to lifestyle counseling in patients with BMI of 35.0-55.0 kg/m².⁵⁴ They also reported clinically significant improvement in co-morbid metabolic parameters such as HbA1C and cholesterol, and a moderate improvement in blood pressure and low-density lipoprotein. Complications were mostly associated with gastrostomy tube placement and managed conservatively. The biggest concern is whether the Aspire Assist promotes bulimic tendencies however, multiple studies have demonstrated improved cognitive food restraint and increased satiety.^{53,54} For these reasons, this approach may be a good candidate for bridge therapy in high BMI patients to bariatric surgery.

4. *Transoral Outlet Reduction (TORe)*

One in four patients undergoing bariatric surgery regain weight. Enlarged gastric pouch and gastrojejunal anastomosis (GJA) is an independent predictor of weight regain following RYGB.⁵⁵ Studies report a linear correlation between weight regain and dilation in the GJA. Surgical revision is associated with increased morbidity and limited efficacy and can be technically challenging.^{56,57} Endoscopic transoral outlet reduction (TORe) is a minimally-invasive endoscopic approach that can be used to restore the ideal pouch size and reduce the GJA outlet. Full-thickness sutures are placed using an endoscopic suturing device in a purse-string or interrupted fashion. A large multicenter international trial of TORe reported a mean weight loss of 9.31 ± 6.7 kg at six months and 8 ± 8.8 kg at 18 months following the procedure.⁵⁸ A validation meta-analysis of 330 patients confirmed the efficacy and safety of TORe. The procedure duration is usually under 60 minutes, and it can be performed in an outpatient procedure room.

B. *Small Bowels-Focused Approaches*

1. *Trans-pyloric shuttle*

The trans-pyloric shuttle (TPS) (BAROnova) is a non-balloon space-occupying device that is inserted and removed endoscopically. It consists of a ball connected to a tether and passed through the pylorus into the duodenum to cause intermittent obstruction to stomach emptying (Figure 5). A small feasibility study of twenty patients, reported 25.1% and 41.0% EWL at 3 and 6 months, respectively.⁵⁹ The TPS system was FDA approved in 2016 for use in obese adults with a BMI of 35-40 kg/m² or those with BMI 30-35 kg/m² with co-morbidities.

2. *Dual-path Enteral Bypass / Incision-less Anastomosis System*

The paradigm of creating a permanent anatomic alteration using mechanical compression was first described by Kanshin et al. in 1978, to create a sutureless side-to-side anastomosis.⁶⁰ Multiple IAS have been developed in the last decade, with the latest version by Ryou et al. using modified nitinol exoskeleton magnets to innately endoscopically recreate a Roux-en-Y gastrojejunal anastomosis.⁶¹ The self-forming octagonal magnets are placed endoscopically and aligned within the gastrointestinal lumen to create an anchoring window. From a technical standpoint, two endoscopes are required to access the small bowel – one magnet is placed 50-100 cm distal to the ligament of Treitz, and the other magnet placed 50-100cm proximal to the ileocecal valve. Placement is confirmed either fluoroscopically or surgically in earlier trials, prior to deployment.⁶² Through compression and focal ischemia, a large caliber fistula is formed creating a partial jejunal diversion. The magnets eventually disengage within a few weeks of placement. The goal of the diversion is for digested food to circumvent a large portion of the small intestine to reduce nutritional absorption, resulting in increased secretions of gut hormones.

The first pilot study of 10 patients showed a mean TBWL of 14.6% and EWL of 40.2% after one year.⁶² There was also a reduction in HbA1C of 1.9% from baseline. One surgical AEs was reported, with trochar disruption of the gastric serosa, repaired with sutures. Common AEs were nausea and vomiting, as well as recurrent diarrhea in 40% of patients, which resolved with dietary intervention. Long term data are still pending.

3. Duodenal Mucosal Resurfacing

Endoscopic duodenal mucosal resurfacing (DMR) is a minimally invasive procedure that incorporates circumferential thermal ablation of the duodenal mucosa, using catheters (Fractyl Laboratories). This is hypothesised to alter nutrient interactions with the duodenal mucosa to improve metabolic homeostasis. In a study of 36 obese patients with type 2 diabetes, HbA1C and fasting blood glucose improved significantly following DMR.⁴⁶ However, only modest weight loss (-2.5 ± 0.6 kg) was reported and one patient developed serious AEs of fever, malaise, and elevated CRP.

Who is the ideal candidate for endoscopic interventions?

It is unknown which patients would be ideal for endoscopic interventions, and further studies are needed to address this question. However, the available procedures can be offered to patients who fail to reach their target weight with diet and lifestyle changes. The following criteria can be used when choosing endoscopic therapy:

- BMI 30-44 kg/m².
- Poor surgical candidate.
- Bridge to surgery (such as knee replacement or transplant surgery).
- Failed treatment of post-surgery complications.
- Early intervention.

Conclusions

Weight-loss approaches with the highest chance to succeed should be individualized and can include different therapeutic modalities, and should focus on the long-term outcomes. Endobariatric treatment is emerging as a safe, minimally-invasive, cost-effective approach. Endoscopic sleeve gastropasty has been endorsed for broader use by experts, but further research is needed to optimize patient selection before greater adoption of the new techniques.

Figures

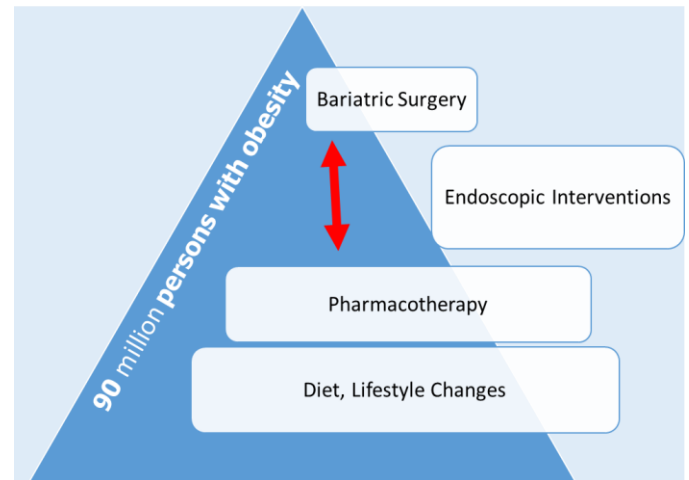
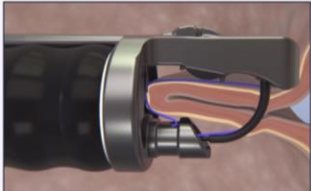
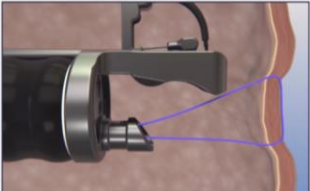


Figure 1: Endoscopic interventions for weight loss can fill in the gap in the management of obesity



OverStitch™

Endoscopic Suturing System

The OverStitch™ Endoscopic Suturing System enables advanced endoluminal surgery by allowing physicians to place **full-thickness sutures** through a flexible endoscope. With the capability to **customize suture patterns**, OverStitch™ facilitates a **broad range** of bariatric and gastrointestinal procedures.

Full thickness sutures incorporate muscle layer to increase tissue approximation durability.

Customize suture patterns to address simple or complex gastrointestinal and bariatric procedures.

Figure 2: Apollo OverStitch device ®



Figure 3: Body of stomach in an obese patient: (A): Prior to ESG. (B): Reduced stomach lumen size at the end of ESG.

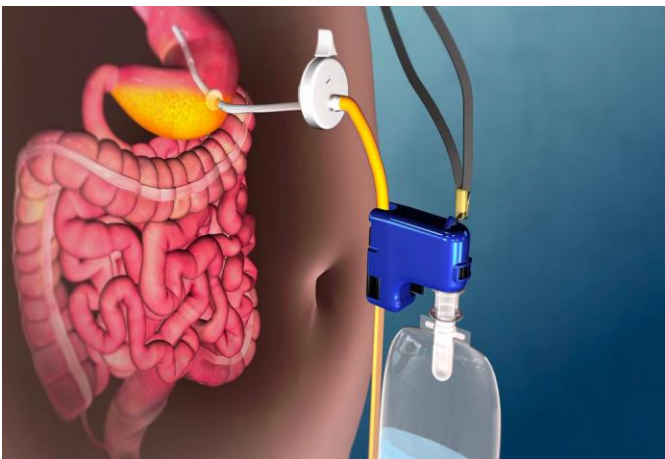


Figure 4: Aspire Assist device ® (Aspire Bariatrics)

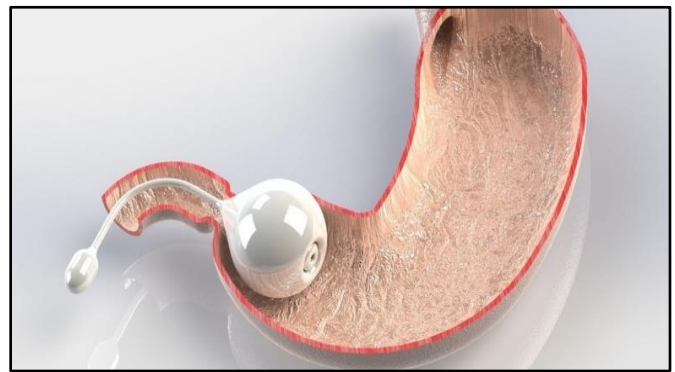


Figure 5: Trans-pyloric shuttle ®

REFERENCES

1. **Waters H, DeVol R.** Weighing down America: The health and economic impact of obesity. Milken Institute 2016. Available at: <https://milkeninstitute.org/reports/weighing-down-america-health-and-economic-impact-obesity>.
2. National Health Expenditures 2014 Highlights. Available at: <https://ccf.georgetown.edu/wp-content/uploads/2017/03/highlights.pdf>.
3. A. M. A . recognizes obesity as a disease. Available at: <https://www.nytimes.com/2013/06/19/business/ama-recognizes-obesity-as-a-disease.html>.
4. **Castellana M, Conte E, Cignarelli A, Perrini S, Giustina A, Giovannella L, Giorgino F, Trimboli P.** Efficacy and safety of very low calorie ketogenic diet (VLCKD) in patients with overweight and obesity: A systematic review and meta-analysis. *Rev Endocr Metab Disord.* 2019 Nov 9. doi: 10.1007/s11154-019-09514-y. [Epub ahead of print] Review. PubMed PMID: 31705259.
5. **Weston KS, Wisløff U, Coombes JS.** High-intensity interval training in patients with lifestyle-induced cardiometabolic disease: a systematic review and meta-analysis. *Br J Sports Med.* 2014 Aug;48(16):1227-34. doi: 10.1136/bjsports-2013-092576. Epub 2013 Oct 21. Review. PubMed PMID: 24144531.
6. **Iellamo F, Manzi V, Caminiti G, Vitale C, Castagna C, Massaro M, Franchini A, Rosano G, Volterrani M.** Matched dose interval and continuous exercise training induce similar cardiorespiratory and metabolic adaptations in patients with heart failure. *Int J Cardiol.* 2013 Sep 10;167(6):2561-5. doi: 10.1016/j.ijcard.2012.06.057. Epub 2012 Jul 4. PubMed PMID: 22769574.
7. **Tjønnå AE, Lee SJ, Rognmo Ø, Stølen TO, Bye A, Haram PM, Loennechen JP, Al-Share QY, Skogvoll E, Slørdahl SA, Kemi OJ, Najjar SM, Wisløff U.** Aerobic interval training versus continuous moderate exercise as a treatment for the metabolic syndrome: a pilot study. *Circulation.* 2008 Jul 22;118(4):346-54. doi: 10.1161/CIRCULATIONAHA.108.772822. Epub 2008 Jul 7. PubMed PMID: 18606913; PubMed Central PMCID: PMC2777731.
8. **Anson RM, Guo Z, de Cabo R, Iyun T, Rios M, Hagepanos A, Ingram DK, Lane MA, Mattson MP.**

- Intermittent fasting dissociates beneficial effects of dietary Restriction on glucose metabolism and neuronal resistance to injury from calorie intake. *Proc Natl Acad Sci U S A*. 2003 May 13;100(10):6216-20. Epub 2003 Apr 30. PubMed PMID: 12724520; PubMed Central PMCID: PMC156352.
9. **Harvie M, Wright C, Pegington M, McMullan D, Mitchell E, Martin B, Cutler RG, Evans G, Whiteside S, Maudsley S, Camandola S, Wang R, Carlson OD, Egan JM, Mattson MP, Howell A.** The effect of intermittent energy and carbohydrate restriction v. daily energy restriction on weight loss and metabolic disease risk markers in overweight women. *Br J Nutr*. 2013 Oct;110(8):1534-47. doi: 10.1017/S0007114513000792. Epub 2013 Apr 16. PubMed PMID: 23591120; PubMed Central PMCID: PMC5857384.
 10. **Ahmet I, Wan R, Mattson MP, Lakatta EG, Talan M.** Cardioprotection by intermittent fasting in rats. *Circulation*. 2005 Nov 15;112(20):3115-21. Epub 2005 Nov 7. PubMed PMID: 16275865.
 11. **Hoddy KK, Kroeger CM, Trepanowski JF, Barnosky AR, Bhutani S, Varady KA.** Safety of alternate day fasting and effect on disordered eating behaviors. *Nutr J*. 2015 May 6;14:44. doi: 10.1186/s12937-015-0029-9. PubMed PMID: 25943396; PubMed Central PMCID: PMC4424827.
 12. **Trepanowski JF, Kroeger CM, Barnosky A, Klempel MC, Bhutani S, Hoddy KK, Gabel K, Freels S, Rigdon J, Rood J, Ravussin E, Varady KA.** Effect of Alternate-Day Fasting on Weight Loss, Weight Maintenance, and Cardioprotection Among Metabolically Healthy Obese Adults: A Randomized Clinical Trial. *JAMA Intern Med*. 2017 Jul 1;177(7):930-938. doi: 10.1001/jamainternmed.2017.0936. PubMed PMID: 28459931; PubMed Central PMCID: PMC5680777.
 13. FDA approves Belviq to treat some overweight or obese adults. *Home Healthc Nurse*. 2012 Sep;30(8):443-4. PubMed PMID: 23097775.
 14. **Smith SR, Prosser WA, Donahue DJ, Morgan ME, Anderson CM, Shanahan WR; APD356-004 Study Group.** Lorcaserin (APD356), a selective 5-HT(2C) agonist, reduces body weight in obese men and women. *Obesity (Silver Spring)*. 2009 Mar;17(3):494-503. doi: 10.1038/oby.2008.537. Epub 2008 Dec 4. PubMed PMID: 19057523.
 15. **Fidler MC, Sanchez M, Raether B, Weissman NJ, Smith SR, Shanahan WR, Anderson CM; BLOSSOM Clinical Trial Group.** A one-year randomized trial of lorcaserin for weight loss in obese and overweight adults: the BLOSSOM trial. *J Clin Endocrinol Metab*. 2011 Oct;96(10):3067-77. doi: 10.1210/jc.2011-1256. Epub 2011 Jul 27. PubMed PMID: 21795446.
 16. **Apovian CM, Aronne LJ, Bessesen DH, McDonnell ME, Murad MH, Pagotto U, Ryan DH, Still CD; Endocrine Society.** Pharmacological management of obesity: an endocrine Society clinical practice guideline. *J Clin Endocrinol Metab*. 2015 Feb;100(2):342-62. doi: 10.1210/jc.2014-3415. Epub 2015 Jan 15. Erratum in: *J Clin Endocrinol Metab*. 2015 May;100(5):2135-6. PubMed PMID: 25590212.
 17. Prescribing Insert for Qysmia. Administration FaD. http://www.accessdata.fda.gov/drugsatfda_docs/label/2012/022580s000lbl.pdf. 2012.
 18. **Allison DB, Gadde KM, Garvey WT, Peterson CA, Schwiers ML, Najarian T, Tam PY, Troupin B, Day WW.** Controlled-release phentermine/topiramate in severely obese adults: a randomized controlled trial (EQUIP). *Obesity (Silver Spring)*. 2012 Feb;20(2):330-42. doi: 10.1038/oby.2011.330. Epub 2011 Nov 3. PubMed PMID: 22051941; PubMed Central PMCID: PMC3270297.
 19. **Gadde KM, Allison DB, Ryan DH, Peterson CA, Troupin B, Schwiers ML, Day WW.** Effects of low-dose, controlled-release, phentermine plus topiramate combination on weight and associated comorbidities in overweight and obese adults (CONQUER): a randomised, placebo-controlled, phase 3 trial. *Lancet*. 2011 Apr 16;377(9774):1341-52. doi: 10.1016/S0140-6736(11)60205-5. Epub 2011 Apr 8. Erratum in: *Lancet*. 2011 Apr 30;377(9776):1494. PubMed PMID: 21481449.
 0. **Astrup A, Rössner S, Van Gaal L, Rissanen A, Niskanen L, Al Hakim M, Madsen J, Rasmussen MF, Lean ME; NN8022-1807 Study Group.** Effects of liraglutide in the treatment of obesity: a randomised, double-blind, placebo-controlled study. *Lancet*. 2009 Nov 7;374(9701):1606-16. doi: 10.1016/S0140-6736(09)61375-1. Epub 2009 Oct 23. Erratum in: *Lancet*. 2010 Mar 20;375(9719):984. PubMed PMID: 19853906.
 21. **de Mello AH, Prá M, Cardoso LC, de Bona Schraiber R, Rezin GT.** Incretin-based therapies for obesity treatment. *Metabolism*. 2015 Sep;64(9):967-81. doi: 10.1016/j.metabol.2015.05.012. Epub 2015 May 23. Review. PubMed PMID: 26072135.
 22. **Pi-Sunyer X, Astrup A, Fujioka K, Greenway F, Halpern A, Krempf M, Lau DC, le Roux CW, Violante Ortiz R, Jensen CB, Wilding JP; SCALE Obesity and Prediabetes NN8022-1839 Study Group.** A Randomized, Controlled Trial of 3.0 mg of Liraglutide in Weight Management. *N Engl J Med*. 2015 Jul 2;373(1):11-22. doi: 10.1056/NEJMoa1411892. PubMed PMID: 26132939.
 23. **Wadden TA, Hollander P, Klein S, Niswender K, Woo V, Hale PM, Aronne L; NN8022-1923 Investigators.** Weight maintenance and additional weight loss with liraglutide after low-calorie-diet-induced weight loss: the SCALE Maintenance randomized study. *Int J Obes (Lond)*. 2013 Nov;37(11):1443-51. doi: 10.1038/ijo.2013.120. Epub 2013 Jul 1. Erratum in: *Int J Obes (Lond)*. 2013 Nov;37(11):1514. *Int J Obes (Lond)*. 2015 Jan;39(1):187. PubMed PMID: 23812094.
 24. **Greenway FL, Whitehouse MJ, Guttadauria M, Anderson JW, Atkinson RL, Fujioka K, Gadde KM, Gupta AK, O'Neil P, Schumacher D, Smith D, Dunayevich E, Tollefson GD, Weber E, Cowley MA.** Rational design of a combination medication for the treatment of obesity. *Obesity (Silver Spring)*. 2009

- Jan;17(1):30-9. doi: 10.1038/oby.2008.461. Epub 2008 Nov 6. PubMed PMID: 18997675.
25. **Billes SK, Greenway FL.** Combination therapy with naltrexone and bupropion for obesity. *Expert Opin Pharmacother.* 2011 Aug;12(11):1813-26. doi: 10.1517/14656566.2011.591382. Epub 2011 Jun 21. Review. PubMed PMID: 21689063.
 26. **Greenway FL, Fujioka K, Plodkowski RA, Mudaliar S, Guttadauria M, Erickson J, Kim DD, Dunayevich E; COR-I Study Group.** Effect of naltrexone plus bupropion on weight loss in overweight and obese adults (COR-I): a multicentre, randomised, double-blind, placebo-controlled, phase 3 trial. *Lancet.* 2010 Aug 21;376(9741):595-605. doi: 10.1016/S0140-6736(10)60888-4. Epub 2010 Jul 29. Erratum in: *Lancet.* 2010 Aug 21;376(9741):594. *Lancet.* 2010 Oct 23;376(9750):1392. PubMed PMID: 20673995.
 27. F.D.A. fails to approve diet drug. Available at: <https://www.nytimes.com/2011/02/02/business/02drug.ht ml>.
 28. Administration FaD. FDA approves weight-management drug Contrave. https://www.accessdata.fda.gov/drugsatfda_docs/label/2014/200063s000lbl.pdf
 29. **Torgerson JS, Hauptman J, Boldrin MN, Sjöström L.** XENical in the prevention of diabetes in obese subjects (XENDOS) study: a randomized study of orlistat as an adjunct to lifestyle changes for the prevention of type 2 diabetes in obese patients. *Diabetes Care.* 2004 Jan;27(1):155-61. Erratum in: *Diabetes Care.* 2004 Mar;27(3):856. PubMed PMID: 14693982.
 30. **Sjöström L, Rissanen A, Andersen T, Boldrin M, Golay A, Koppeschaar HP, Krempf M.** Randomised placebo-controlled trial of orlistat for weight loss and prevention of weight regain in obese patients. European Multicentre Orlistat Study Group. *Lancet.* 1998 Jul 18;352(9123):167-72. PubMed PMID: 9683204.
 31. **Leblanc ES, O'Connor E, Whitlock EP, Patnode CD, Kapka T.** Effectiveness of primary care-relevant treatments for obesity in adults: a systematic evidence review for the U.S. Preventive Services Task Force. *Ann Intern Med.* 2011 Oct 4;155(7):434-47. doi: 10.7326/0003-4819-155-7-201110040-00006. Review. PubMed PMID: 21969342.
 32. **Rucker D, Padwal R, Li SK, Curioni C, Lau DC.** Long term pharmacotherapy for obesity and overweight: updated meta-analysis. *BMJ.* 2007 Dec 8;335(7631):1194-9. Epub 2007 Nov 15. Erratum in: *BMJ.* 2007 Nov 24;335(7629). doi: 10.1136/bmj.39406.519132.AD. PubMed PMID: 18006966; PubMed Central PMCID: PMC2128668.
 33. **Singh A, Sarkar SR, Gaber LW, Perazella MA.** Acute oxalate nephropathy associated with orlistat, a gastrointestinal lipase inhibitor. *Am J Kidney Dis.* 2007 Jan;49(1):153-7. PubMed PMID: 17185156.
 34. **Heymisfield SB, Wadden TA.** Mechanisms, Pathophysiology, and Management of Obesity. *N Engl J Med.* 2017 Apr 13;376(15):1492. doi: 10.1056/NEJMc1701944. PubMed PMID: 28402780.
 35. **SAXENDA®** safely and effectively. 2018. (Package insert).
 36. **Ponce J, Nguyen NT, Hutter M, Sudan R, Morton JM.** American Society for Metabolic and Bariatric Surgery estimation of bariatric surgery procedures in the United States, 2011-2014. *Surg Obes Relat Dis.* 2015 Nov-Dec;11(6):1199-200. doi: 10.1016/j.soard.2015.08.496. Epub 2015 Aug 12. Review. PubMed PMID: 26476493.
 37. **Fogel R, De Fogel J, Bonilla Y, De La Fuente R.** Clinical experience of transoral suturing for an endoluminal vertical gastroplasty: 1-year follow-up in 64 patients. *Gastrointest Endosc.* 2008 Jul;68(1):51-8. doi: 10.1016/j.gie.2007.10.061. Epub 2008 Mar 19. PubMed PMID: 18355825.
 38. **Brethauer SA, Chand B, Schauer PR, Thompson CC.** Transoral gastric volume reduction as intervention for weight management: 12-month follow-up of TRIM trial. *Surg Obes Relat Dis.* 2012 May-Jun;8(3):296-303. doi: 10.1016/j.soard.2011.10.016. Epub 2011 Nov 9. PubMed PMID: 22178565.
 39. **Brethauer SA, Chand B, Schauer PR, Thompson CC.** Transoral gastric volume reduction for weight management: technique and feasibility in 18 patients. *Surg Obes Relat Dis.* 2010 Nov-Dec;6(6):689-94. doi: 10.1016/j.soard.2010.07.012. Epub 2010 Aug 6. PubMed PMID: 20947451.
 40. **Kumar N, Abu Dayyeh BK, Lopez-Nava Breviere G, Galvao Neto MP, Saldala NP, Shaikh SN, Hawes RH, Gostout CJ, Goenka MK, Orillac JR, Alvarado A, Jirapinyo P, Zundel N, Thompson CC.** Endoscopic sutured gastroplasty: procedure evolution from first-in-man cases through current technique. *Surg Endosc.* 2018 Apr;32(4):2159-2164. doi: 10.1007/s00464-017-5869-2. Epub 2017 Oct 26. PubMed PMID: 29075966; PubMed Central PMCID: PMC5845469.
 41. **Sharaiha RZ, Kumta NA, Saumoy M, Desai AP, Sarkisian AM, Benevenuto A, Tyberg A, Kumar R, Igel L, Verna EC, Schwartz R, Frissora C, Shukla A, Aronne LJ, Kahaleh M.** Endoscopic Sleeve Gastroplasty Significantly Reduces Body Mass Index and Metabolic Complications in Obese Patients. *Clin Gastroenterol Hepatol.* 2017 Apr;15(4):504-510. doi: 10.1016/j.cgh.2016.12.012. Epub 2016 Dec 23. PubMed PMID: 28017845.
 42. **Lopez-Nava G, Galvão MP, Bautista-Castaño I, Fernandez-Corbelle JP, Trelle M, Lopez N.** Endoscopic sleeve gastroplasty for obesity treatment: Two years of experience. *Arq Bras Cir Dig.* 2017 Jan-Mar;30(1):18-20. doi: 10.1590/0102-6720201700010006. English, Portuguese. PubMed PMID: 28489162; PubMed Central PMCID: PMC5424680.
 43. **Sartoretto A, Sui Z, Hill C, Dunlap M, Rivera AR, Khashab MA, Kalloo AN, Fayad L, Cheskin LJ, Marinos G, Wilson E, Kumbhari V.** Endoscopic Sleeve Gastroplasty (ESG) Is a Reproducible and Effective Endoscopic Bariatric Therapy Suitable for Widespread Clinical Adoption: a Large, International Multicenter Study. *Obes Surg.* 2018 Jul;28(7):1812-1821. doi: 10.1007/s11695-018-3135-x. PubMed PMID: 29450845.
 44. **Cheskin LJ, Hill C, Adam A, Fayad L, Dunlap M, Badurdeen D, Koller K, Bunyard L, Frutchey R, Al-**

- Grain H, Kahan S, Hedjoudje A, Khashab MA, Kalloo AN, Kumbhari V.** Endoscopic sleeve gastropasty versus high-intensity diet and lifestyle therapy: a case-matched study. *Gastrointest Endosc.* 2020 Feb;91(2):342-349.e1. doi: 10.1016/j.gie.2019.09.029. Epub 2019 Sep 27. PubMed PMID: 31568769.
45. **Hajifathalian K, Ang B, Dawod QM, Shah SL, Dawod E, Mehta A, Mukewar S, Mahadev S, Sampath K, Carr-Locke DL, Issa D, Aronne LJ, Kumar R, Shukla A, Sharaiha RZ.** 175 Long-term follow up and outcomes after endoscopic sleeve gastropasty for treatment of obesity (5 year data). *Gastrointest Endosc.* 2019; 89(6): AB60-AB61.
46. **Hajifathalian K, Ang B, Dawod QM, Shah SL, Dawod E, Mehta A, Mahadev S, Mukewar S, Sampath K, Carr-Locke DL, Shukla A, Aronne LJ, Issa D, Kumar R, Sharaiha RZ.** 179 improvement in non-alcoholic fatty liver disease after endoscopic sleeve gastropasty. *Gastrointest Endosc.* 2019; 89(6): AB58.
47. **Saumoy M, Schneider Y, Zhou XK, Shukla A, Kahaleh M, Aronne L, Sharaiha RZ.** A single-operator learning curve analysis for the endoscopic sleeve gastropasty. *Gastrointest Endosc.* 2018 Feb;87(2):442-447. doi: 10.1016/j.gie.2017.08.014. Epub 2017 Aug 24. PubMed PMID: 28843586.
48. **Ponce J, Woodman G, Swain J, Wilson E, English W, Ikramuddin S, Bour E, Edmundowicz S, Snyder B, Soto F, Sullivan S, Holcomb R, Lehmann J; REDUCE Pivotal Trial Investigators.** The REDUCE pivotal trial: a prospective, randomized controlled pivotal trial of a dual intragastric balloon for the treatment of obesity. *Surg Obes Relat Dis.* 2015 Jul-Aug;11(4):874-81. doi: 10.1016/j.soard.2014.12.006. Epub 2014 Dec 16. PubMed PMID: 25868829.
49. **Courcoulas A, Abu Dayyeh BK, Eaton L, Robinson J, Woodman G, Fusco M, Shayani V, Billy H, Pambianco D, Gostout C.** Intragastric balloon as an adjunct to lifestyle intervention: a randomized controlled trial. *Int J Obes (Lond).* 2017 Mar;41(3):427-433. doi: 10.1038/ijo.2016.229. Epub 2016 Dec 23. PubMed PMID: 28017964.
50. **Sullivan S, Swain J, Woodman G, Edmundowicz S, Hassanein T, Shayani V, Fang JC, Noar M, Eid G, English WJ, Tariq N, Larsen M, Jonnalagadda SS, Riff DS, Ponce J, Early D, Volckmann E, Ibele AR, Spann MD, Krishnan K, Bucobo JC, Pryor A.** Randomized sham-controlled trial of the 6-month swallowable gas-filled intragastric balloon system for weight loss. *Surg Obes Relat Dis.* 2018 Dec;14(12):1876-1889. doi: 10.1016/j.soard.2018.09.486. Epub 2018 Sep 29. PubMed PMID: 30545596.
51. **Trang J, Lee SS, Miller A, Cruz Pico CX, Postoev A, Ibikunle I, Ibikunle CA.** Incidence of nausea and vomiting after intragastric balloon placement in bariatric patients - A systematic review and meta-analysis. *Int J Surg.* 2018 Sep;57:22-29. doi: 10.1016/j.ijso.2018.06.038. Epub 2018 Jul 20. Review. PubMed PMID: 30031839.
52. **ASGE Bariatric Endoscopy Task Force and ASGE Technology Committee, Abu Dayyeh BK, Kumar N, Edmundowicz SA, Jonnalagadda S, Larsen M, Sullivan S, Thompson CC, Banerjee S.** ASGE Bariatric Endoscopy Task Force systematic review and meta-analysis assessing the ASGE PIVI thresholds for adopting endoscopic bariatric therapies. *Gastrointest Endosc.* 2015 Sep;82(3):425-38.e5. doi: 10.1016/j.gie.2015.03.1964. Epub 2015 Jul 29. Review. PubMed PMID: 26232362.
53. **Sullivan S, Stein R, Jonnalagadda S, Mullady D, Edmundowicz S.** Aspiration therapy leads to weight loss in obese subjects: a pilot study. *Gastroenterology.* 2013 Dec;145(6):1245-52.e1-5. doi: 10.1053/j.gastro.2013.08.056. Epub 2013 Sep 6. PubMed PMID: 24012983; PubMed Central PMCID: PMC4025911.
54. **Thompson CC, Abu Dayyeh BK, Kushner R, Sullivan S, Schorr AB, Amaro A, Apovian CM, Fullum T, Zarrinpar A, Jensen MD, Stein AC, Edmundowicz S, Kahaleh M, Ryou M, Bohning JM, Ginsberg G, Huang C, Tran DD, Glaser JP, Martin JA, Jaffe DL, Farraye FA, Ho SB, Kumar N, Harakal D, Young M, Thomas CE, Shukla AP, Ryan MB, Haas M, Goldsmith H, McCrea J, Aronne LJ.** Percutaneous Gastrostomy Device for the Treatment of Class II and Class III Obesity: Results of a Randomized Controlled Trial. *Am J Gastroenterol.* 2017 Mar;112(3):447-457. doi: 10.1038/ajg.2016.500. Epub 2016 Dec 6. PubMed PMID: 27922026; PubMed Central PMCID: PMC5350543.
55. **Heneghan HM, Yimcharoen P, Brethauer SA, Kroh M, Chand B.** Influence of pouch and stoma size on weight loss after gastric bypass. *Surg Obes Relat Dis.* 2012 Jul-Aug;8(4):408-15. doi: 10.1016/j.soard.2011.09.010. Epub 2011 Sep 23. PubMed PMID: 22055390.
56. **Parikh M, Heacock L, Gagner M.** Laparoscopic "gastrojejunal sleeve reduction" as a revision procedure for weight loss failure after roux-en-y gastric bypass. *Obes Surg.* 2011 May;21(5):650-4. doi: 10.1007/s11695-010-0274-0. PubMed PMID: 20835779.
57. **Coakley BA, Deveney CW, Spight DH, Thompson SK, Le D, Jobe BA, Wolfe BM, McConnell DB, O'Rourke RW.** Revisional bariatric surgery for failed restrictive procedures. *Surg Obes Relat Dis.* 2008 Sep-Oct;4(5):581-6. Epub 2007 Dec 11. PubMed PMID: 18065290.
58. **Vargas EJ, Bazerbachi F, Rizk M, Rustagi T, Acosta A, Wilson EB, Wilson T, Neto MG, Zundel N, Mundi MS, Collazo-Clavell ML, Meera S, Abu-Lebdeh HS, Lorentz PA, Grothe KB, Clark MM, Kellogg TA, McKenzie TJ, Kendrick ML, Topazian MD, Gostout CJ, Abu Dayyeh BK.** Transoral outlet reduction with full thickness endoscopic suturing for weight regain after gastric bypass: a large multicenter international experience and meta-analysis. *Surg Endosc.* 2018 Jan;32(1):252-259. doi: 10.1007/s00464-017-5671-1. Epub 2017 Jun 29. PubMed PMID: 28664438.
59. **Marinos G, Eliades C, Raman Muthusamy V, Greenway F.** Weight loss and improved quality of life with a nonsurgical endoscopic treatment for obesity: clinical results from a 3- and 6-month study. *Surg Obes Relat Dis.* 2014 Sep-Oct;10(5):929-34. doi: 10.1016/

j.soard.2014.03.005. Epub 2014 Mar 12. PubMed PMID: 25066439.

60. **Kanshin NN, Permiakov NK, Dzhagoniia RA, Nikulin BI, Kuznetsov AA.** [Sutureless anastomoses in gastrointestinal surgery with and without steady magnetic field (experimental study)]. *Arkh Patol.* 1978;40(8):56-61. Russian. PubMed PMID: 365148.
61. **Ryou M, Ryan MB, Thompson CC.** Current status of endoluminal bariatric procedures for primary and revision indications. *Gastrointest Endosc Clin N Am.* 2011 Apr;21(2):315-33. doi: 10.1016/j.giec.2011.02.004. Review. PubMed PMID: 21569983; PubMed Central PMCID: PMC3460649.
62. **Machytka E, Bužga M, Zonca P, Lautz DB, Ryou M, Simonson DC, Thompson CC.** Partial jejunal diversion using an incisionless magnetic anastomosis system: 1-year interim results in patients with obesity and diabetes. *Gastrointest Endosc.* 2017 Nov;86(5):904-912. doi: 10.1016/j.gie.2017.07.009. Epub 2017 Jul 14. PubMed PMID: 28716404.