

Endoscopic intragastric balloon: a gimmick or a viable option for obesity?

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Abstract: Worldwide, the prevalence of obesity has doubled since 1980 in 70 countries. More than one in three adults now suffer from overweight or obesity. Health problems related to obesity include orthopedic problems, psychiatric conditions, metabolic and cardiovascular diseases, and of increasing concern, cancer. Thus, obesity has an enormous impact on the individual's wellbeing as well as on society's workforce and health care expenses. Medical efforts are ongoing to find safe and effective treatment options for obesity and its metabolic implications. At present, available treatment options include lifestyle interventions, pharmacotherapy, endoscopic applications, and bariatric surgery. Within the range of endoscopic treatment options, the intragastric balloon is the most widely used device. The idea is simple: the gastric volume is reduced by a balloon that is in most cases implanted by an endoscopic procedure similar to a gastroscopy. During the past decades, different models have been developed, which we will briefly introduce in this review. We aim at reviewing the pathophysiology underlying the effect of endoscopic intragastric balloon on weight loss and metabolic changes. We will assess expected short-term and long-term benefits for the patient, and we will discuss common side effects as well as rare complications. We will compare endoscopic intragastric balloon to conservative treatment options with or without pharmacological support on the one hand and to the spectrum of bariatric surgery on the other hand. In most patients, obesity must be considered a chronic disease that requires a lifelong treatment concept. In view of current treatment options for obesity, we will discuss whether endoscopic intragastric balloon is a viable treatment option, and who may be the right patient to benefit from it.

Keywords: Bariatric surgery; metabolic endoscopy; gastric balloon; obesity

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Introduction

The intragastric balloon (IGB) probably set the starting point of endoscopic bariatric therapy. Depending on the system, an empty balloon is introduced into the stomach by means of a gastroscopy or by just swallowing a capsule. The balloon is then inflated with air or saline to varying volumes. This reduces the stomach volume, resulting in a feeling of satiety which shall then lead to weight loss. The treatment is limited to 6 months for most systems. Ideally the procedure is imbedded in a continuous weight loss program, and the patient is supported by an interdisciplinary team of healthcare professionals.

The idea of today's IGB systems may date back to as early as 1939, when Michael DeBakey analyzed cases of gastric bezoars and concretions to find that 38% of the patients experienced weight loss. However, the vast majority of the patients suffered from nausea and vomiting even

Table 1 Overview of FI	DA approved and/or CE certif	ied intragastric balloon	s in alphabetic order

Balloon type	Implant/explant	FDA approved	CE mark	Time in situ (months)	Filling	Volume (mL)	TBWL (%)	EWL (%)	Ref.
Elipse [®]	Swallow/excretion	No*	Yes	4	Liquid	450–550	10.0	n/a	(4)
End-Ball [®]	Endo/endo	No	Yes	6	Liquid/gas	700	17.1	36.5	(5)
Heliosphere [®] BAG	Endo/endo	No	Yes	6	Air	700	13.4	33.2	(6)
Lexbal®	Endo/endo	No	Yes	6	Liquid	500-800	14.9	26.8	(7)
MedSil [®]	Endo/endo	No	Yes	6	Liquid	400–700	13.1	n/a	(8)
Obalon®	Swallow/endo	Yes	Yes	6	Gas	3×250	7.1	n/a	(9)
Orbera®	Endo/endo	Yes	Yes	6	Liquid	400–700	10.2	26.5	(10)
Reshape®	Endo/endo	Yes	Yes	6	Liquid	750–900	6.8	25.1	(11)
Spatz3 [®]	Endo/endo	No	Yes	12	Liquid	Adjustable	20.1	45.8	(12)

*, in progress. FDA, Food and Drug Administration; CE, Communauté Européenne; TBWL, total body weight loss; EWL, excess weight loss; endo, endoscopically; n/a, not applicable.

without necessarily losing weight (1). The first IGB, the "Garren-Edwards Gastric Bubble" was introduced in 1985 and experienced a similar problem. The FDA approved the system, but it was withdrawn after seven years because of major side effects, including gastric erosions, gastric ulcers, small bowel obstruction, and Mallory-Weiss tears, as well as insufficient efficacy on weight loss (2). Despite these first disappointing experiences, other systems have been developed and successfully introduced on the European and US American market: The Orbera® Intragastric Balloon System (Apollo Endosurgery Inc, Austin, TX, United States, formerly BIB), was approved for use in Europe in 1997 and culminated in an FDA approval in 2015. It is the system, among all balloon systems, for which most scientific data on efficacy and safety are available; thus, it may be the most widely implanted system worldwide. Since 2015, another two balloon systems have gained FDA approval: The ReShape[®] Integrated Dual BalloonSystem (ReShape Medical, Inc., San Clemente, CA, United States), and the Obalon[®] (Obalon Therapeutics, Inc.), the first swallowable balloon system that allows up to three balloons to be put in place without endoscopy. Removal of the Obalon® balloon system, however, also requires endoscopy. Finally, there is a number of Communauté Européenne (CE) approved but not yet U.S. food and drug administration (FDA) approved balloon systems under clinical investigation. Within these systems, the Spatz^{3®} intragastric balloon (Spatz FGIA Inc.) differs from the other balloons as it is adjustable and can stay in place for 12 months. The Elipse® Balloon (Allurion Technologies) is the only swallowable CE approved system that is removed by natural excretion and thus, ideally does not require any endoscopic procedure (3). An overview of FDA and CE approved IGBs is given in *Table 1*.

The idea of the IGB keeps being attractive: in contrast to bariatric surgery, the therapy is little or non-invasive, reversible and limited to a specified timeframe. However, the effect on weight loss is by far not comparable to what can be expected from operative procedures. As with any medical therapy decision, advantages and disadvantages must be weighed up to best support the patient.

Pathophysiology

It appears common-sense that a stomach that is continuously "pre-filled" by a rather large balloon will lead to an increased sensation of satiety and to reduced appetite. But which pathophysiological mechanisms may be underlying, and can the IGB indeed take advantage of these mechanisms?

Satiety and bodyweight are regulated by a complex neuro-humoral network which is, in part, represented in the gut brain axis. The current and highly accepted concept describes humoral and nerval afferences from adipose tissue and gut that are integrated in brain stem circuits. These brain stem circuits in turn provide hunger, saturation and satiety signals. The system is fed by mechanical stimuli such as stomach distension, and it is modulated by hormonal factors as well as by psychological factors, known for example as the reward system (13-15). Distension of the stomach in response to a meal signals saturation via parasympathetic afferents. Conversely, brain stem nuclei control the pre-prandial tonus and relaxation after a meal (13-15). Interestingly, functional neuro imaging studies

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have shown that there is a difference between balloon distension of the stomach and the physiological distension by nutrients: the pure mechanic distension by an inflated balloon addresses pain-processing brain region. This socalled "visceral pain neuromatrix" has been shown to be downregulated during nutrient infusion, which may constitute a necessity to ensure adequate food intake despite gastric distension (16). Considering this mechanism, pure gastric distension by an IGB without simultaneous food intake explains side effects in the first days after implantation and satiety after food reconstitution. However, there are limitations to this effect: functional MRI studies have shown that sweet beverages led to no significant effect on cortical brain circuits associated with food craving (17). Thus, Patients with an increased intake of liquid calories may not benefit from an IGB system.

On the humoral side, the satiety hormone, ghrelin, a 28 amino acid peptide expressed in the gastric fundus, is a major player in weight regulation (18). Its role in ballooninduced weight loss, however, is not clear (19,20). Ly *et al.* show that the mechanical effects of balloon inflation alone did not reduce plasma ghrelin levels, so that caloric intake seems a mandatory signal for ghrelin induced sensation of satiety (21). Still, ghrelin levels decreased during IGB treatment and positively correlated with weight loss, suggesting that food-induced ghrelin is active in regulating satiety and possibly weight loss in the presence of IGB (22).

Besides gastric distension and humoral factors, gastric emptying represents another important mechanism to induce weight loss following IGB insertion. Thus, three months after balloon placement, solid and liquid scintigraphic gastric emptying time significantly slowed down, which correlated positively with weight loss (23). Despite these findings, there are data questioning whether the initial effects of IGB persist during the time of treatment. While subjective rating of hunger and food desire decreases shortly after IGB placement, these alterations disappear in the same subjects two months after IGB placement (24).

To sum up, gastric distension, ghrelin signaling, and gastric emptying are probably the major players involved in weight loss during IGB treatment. The extent to which each factor is involved can be modulated by the amount and form of caloric intake and by the time since the placement of the gastric balloon.

In patients who had no previous bariatric interventions

and in whom IGB was the primary weight-loss agent, short-term weight reduction as reported after six months treatment averaged at 11.5 kg and was more pronounced in subjects with higher BMI (25).

A recent meta-analysis performed by the American Society for Gastrointestinal Endoscopy Bariatric (ASGE) Endoscopy Task Force indicates that the Orbera[®] IGB leads to an overall pooled total body weight loss (TBWL) of 13.16% after 6 months. Furthermore, in three randomized controlled trials, excess weight loss (%EWL) over controls was 26.9%. Thus Orbera[®] meets the "preservation and incorporation of valuable endoscopic innovations" (PIVI) thresholds for obesity therapy which are set at 5% TBWL and 15% EWL over control, respectively (26,27).

Considering that intensive lifestyle intervention programs report a weight loss of 7–10 kg after 6 months, the effect of IGB therapy has to be considered "modest". While behavioral therapy still results in long-term weight loss of approximately 5 kg, few long-term data are available after IGB therapy (28). Long-term follow up after IGB therapy is extremely heterogenic: a number of patients benefit from repeated use of the same endoscopic therapy, others seek medical therapy, and others decide on bariatric surgery during follow up. In one study, 100 patients who initially lost a mean of 12.6 kg with IGB therapy regained 6.5 kg during 2 years of follow-up, which compares well to other forms of weight loss strategies that are timely restricted (29). Tate et al. review controlled studies of the three FDA approved balloon systems, ReShape[®], Orbera[®] and Obalon[®] to conclude that all systems have comparable effects on weight loss. In summary, the effects were greater than control, smaller than bariatric surgery, but also smaller than the FDA approved weight loss drug, Qsymia (30). As mentioned before, gastric emptying time has been reported as a possible predictor for IGB success. Therefore, the combination of IGB with a medication that reduces gastric motility may potentiate the effect on weight loss (31,32). Another predictor for successful treatment might be the extent of overweight. In non-obese but overweight patients, IGB successfully reversed diabetes and hypertension in a significant amount. EWL was up to 59% at 6 months and 29% at 3 years (33). Similarly, Jamal et al. report a better performance in overweight and obese patients with a BMI of 27.5-35 kg/m² compared to patients with higher BMIs (34).

Adverse events

Adverse events after IGB placement are common and

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similarly seen with all systems that are currently used. The rate of adverse events reported in single studies varies extremely depending on the focus of the study. It is clear, however, that the majority of patients carrying a gastric balloon suffer from nausea and vomiting. Metaanalyses report that 23% to 72% of the patients experience nausea and 20% to 50% of gastric balloon carriers report abdominal pain (25,35,36). It is notable that even severe adverse events occurred frequently: the FDA pivotal trial for the most frequently used gastric balloon, Orbera[®], report gastric ulceration in 10% of the participants. Patients carrying ReShape[®] experienced gastric ulceration even in 35% of the cases probably due to high filling volumes of up to 900 mL (11,27). However, a large systematic review including 26 studies, mostly case series, reports gastric ulceration in only 0.3%. Other rare severe complications following gastric balloon include intestinal obstruction (0.8%), gastric perforation (0.1%) and death (0.05%) (25). Many patients tend to tolerate gastrointestinal side effects to the benefit of weight loss. Early removal of IGBs was reported in 3.5% to 7% of the patients (25,26). Interestingly, the tolerability but also the effectiveness of IGB depends upon localization with more side effects and more weight loss with an antral position (37). The Spatz3[®] Adjustable Balloon was developed to overcome tolerability issues by allowing to adjust the balloon's volume according to side effects and weight loss goals. However, a casecontrol study found no differences in side effects and weight loss outcome compared to Orbera[®] (38), and another study raised concerns regarding long-term safety (39). Single studies suggest, that air-filled balloons like the Heliosphere® BAG (Heliscopie, Vienne, France) or Obalon® will prevent nausea and vomiting while being also efficient in terms of weight loss (6,9). This is remarkable since early airfilled balloons were removed from the market because of ineffectiveness and safety issues. Whether the newer generations have overcome these problems still needs to be answered by head-to-head studies with fluid-filled balloons. In order to prevent or overcome nausea and vomiting, patients receive antiemetic treatment after IGB placement on a regular basis. Prospective studies have shown serotonin receptor antagonists to be effective, and even more so, if combined with midazolam (40,41).

Comparison to surgery and drugs

Very obese patients are often recommended to undergo bariatric surgery as the most effective way to achieve significant weight loss. The IGB and laparoscopic vertical sleeve gastrectomy are based on the same idea of reducing the stomach volume to induce weight loss. This raises the question in how far these procedures are comparable mechanistically. The average gastric volume is about 1,200 mL. In obesity, this volume can reach up to the threefold size. A successful weight reduction after sleeve gastrectomy does not correlate with the extent of the absolute volume reduction, but with the residual volume, whereby a residual volume of more than 100 mL is associated with a poorer outcome (42,43). The IGB, at a filling volume of 600 mL, would leave a residual volume of 600 mL in a normal weight person and much more in an obese or very obese person. Thus, the restrictive characteristics of the surgical and the endoscopic procedures differ tremendously, which explains the much weaker effect of the IGB. The great difference in residual gastric volume may also explain that humoral effects with reduced ghrelin and increased GLP-1 release are seen in sleeve gastrectomy but not in IGB (44,45). When translated into weight loss, the endoscopic therapy as a single treatment stands far behind surgical methods. Bariatric surgery initially achieves an exceptional weight loss of up to 31% depending on the type of surgery. After 10 years of follow up, patients remain a 29% weight reduction compared to baseline (46). Available data on long term weight loss after gastric balloon therapy show modest results with substantial regain of weight after balloon removal if no other therapies follow. Profound metabolic improvements are routinely seen following bariatric surgery. Metabolic data following IGB placement report only limited success that does not go beyond what could be achieved with a conservative therapy for weight reduction (47). In view of these facts, it seems plausible that some medical treatment options are superior to the IGB in respect of metabolic improvements. The GLP1 agonist, liraglutide, leads to weight loss and can stop the progression of pre-diabetes into diabetes over a relatively long period of 3 years (48). Semaglutide, which recently received FDA approval as an antidiabetic drug, seems even more potent reaching a TBWL of 13.8% after 1 year (49), and a novel combination drug of GLP and GIP receptor agonists proved still more effective. The combination drug has been tested in a phase 2 trial, where 73% of the patients reached the weight loss goal of 5% compared to 22% in the dulaglutide group and 0% in the placebo group (50). In conclusion, when compared to surgery or medical therapy, bariatric surgery will lead to a much more profound weight loss than IGB treatment, combined with clear metabolic

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benefits. Medical treatment may be similarly effective as IGB in inducing weight loss; however, differences will be seen depending on patient characteristics and choice of drug. In respect of metabolic effects, antidiabetic drugs based on GLP-1 and GIP related pathways promise to be more effective than the IGB, while the side effects nausea and vomiting are probably equally frequent.

Special indications

In view of the wide spectrum of available weight reduction interventions, the question arises, for which patients the IGB may represent the optimal solution. There are no comprehensive studies available on this topic. Reviewing the literature, however, brings up groups of patients who may specifically benefit from IGB treatment. As described before, some data suggest that the greatest effects on weight loss are achieved in overweight and moderately obese patients (34). This patient group, especially if metabolically healthy, will not qualify for bariatric surgery. The IGB can support weight loss and give a "kick-start" to successful weight loss, especially in situations where life style interventions were unsuccessful (51). In line with this theory, follow-up studies have shown successful weight maintenance after balloon removal mainly in the lower BMI groups (5,34,52).

On the other side of the obesity spectrum, IGB therapy has been discussed in a preoperative setting: in superobese it is reasonable to reduce body weight before bariatric surgery in order to improve associated comorbidities, to facilitate surgery by reducing liver volume and intraabdominal fat and to finally reduce complications. In fact, a 14-day very lowcalorie diet was shown to reduce postoperative complication rate (53). Similar results could be presumed following IGB treatment. However, Coffin et al. show in a multicenter prospective study that while IGB induced preoperative weight loss, neither complication rate nor postoperative weight loss was influenced (54). Thus, pretreatment with IGB before bariatric surgery does reduce body weight but has so far not been proven to be beneficial for surgery outcomes (54-56). Another subgroup which has been studied for pre-surgical IGB treatment are obese patients awaiting kidney transplantation, as BMI >35 kg/m² can preclude renal insufficient patients from being waitlisted (57) or put them at greater risk for complications after transplantation (58). IGB therapy resulted in weight loss in these patients and proved to be save. However, the study was too small to conclude on benefits regarding kidney transplantation after

IGB treatment (58).

While transplant patients surely are a minority within the obese population, children and adolescents represent a growing proportion of the obesity community. It is important to understand whether the IGB may be a worthy treatment option for younger patients. Numerous studies have shown that the benefits of conservative therapy for adolescents are very limited (59). Although more and more adolescents undergo surgical measures, they are considered to be the last resort. Long-term data on success and side effects of bariatric surgery in children and adolescents do not exist, and it is known that children and adolescents often times fail to take their supplement on a regular basis. Consequently, these patients are prone for deficiency symptoms after a few years. In face of this problem, the effects of gastric banding as a potentially reversible intervention has recently been studied in adolescents (60). Reversibility can also be argued as an advantage of the IGB. While there is little data on the use of a gastric balloon in adolescents, the available studies show that the procedure can be used safely and effectively (61-63). However, it needs to be emphasized that none of the balloons has to date been approved for use in minors. Their use is therefore currently only possible within the framework of studies.

In search of the "ideal patient" for IGB treatment one can sum up that lower BMI patients $(25-35 \text{ kg/m}^2)$ may experience more profound and longer lasting benefits on weight loss than more obese patients; however, data are diverse on this issue (26,34). In very obese patients, IGB treatment has been proven effective and safe in preoperative settings. However, available data fail to prove benefits as to the post-operative outcomes in these patients compared to patients who have not received IGB treatment.

Conclusions

After 35 years of experience with IGB systems in the spectrum of anti-obesity therapies, the concept has not found its position as a standard therapy for weight reduction. A number of meta-analyses have summed up weight benefits and side effect of IGB therapy, coming to the conclusion, that weight reduction is moderate and side effects are common, including severe but seldom complications such as gastric ulceration and intestinal obstruction. In addition, metabolic benefits are only seen after sufficient weight loss, clearly putting the IGB behind bariatric surgery but also behind drug interventions that involve the GLP-1 or GIP system. As a complement to life style interventions the IGB

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may have its place as it facilitates initial weight loss and thus may increase the patient's motivation to continue life style efforts. In this setting, IGB therapy shows the best results in overweight and moderately obese patients, which is in our eyes the main indication for this therapy. When we talk about "moderate weight effects" which are comparable to conservative treatment effects, we need to consider that in most studies mean TBWL% and standard deviation (SD) is given rather than ranges. This kind of data does not transparently show the large range of success and does not reveal what is maximally achievable given the right balloon and patient. For example, a 30 kg TBWL is a realistic goal with IGB treatment in selected subjects and could turn a 120 kg obese individual into an overweight or even normalweight 90 kg person. The answer to the question, whether the IGB is a useful tool, will finally reflect perspectives and expectations. IGB therapy is not a magic bullet that leads to long-term remissions from obesity in most cases. Such expectations would certainly be presumptuous. IGB is a tool for selected patients that can be part of a lifelong obesity therapy. In the ideal case, the IGB helps patients so that an initial weight reduction positively influences lifestyle changes that will then allow weight stabilization in the long term. Even failure of the IGB therapy may have a benefit if it strengthens the patient's insight that conservative methods are not the key to success and thus pave the way for metabolic surgery. It is accepted that effectiveness of IGB therapy will never reach that of bariatric surgery and in clinical practice IGB therapy will not be considered a true alternative to surgical procedures. The success of IGB therapy must be measured against conservative therapy programs and drugs. The individual preference will depend on multiple factors, including the patient's choice, local conditions and funding agencies.

Looking at alternative therapies, other endoscopic procedures are evolving that appear to be more effective in the metabolic respect: the duodenal-jejunal bypass sleeve mimics the situation after Y-roux bypass surgery by bypassing the duodenal mucosa. Despite beneficial effects on glycemic control and body weight (64,65), the CE Certificate of conformity was withdrawn in 2017 due to a deficient quality management system. The results of a new pivotal trial will decide on a comeback of the device (47). An innovative procedure that pursues reprogramming of neuroendocrine cells of the duodenum is the duodenal mucosal resurfacing with promising preliminary data (66,67). Finally, Endoscopic sleeve gastroplasty, as a minimally invasive procedure to reduce gastric volume, also seems to be more effective than IGB therapy (68). Thus, further endoscopic procedures may be promising future options for weight loss and metabolic improvement in obesity.

Now we come back to the initial question, whether or not endoscopic IGB therapy is a gimmick. "Gimmick" is defined as "something that is not serious or of real value that is used to attract people's attention or interest temporarily, especially to make them buy something" (69). If you take a significant reduction of body weight as a "real value", the benefit of the IGB is more than zero. IGB therapy consistently led to a reduction of the body weight of approx. 5% when compared to a control as shown in meta-analyses, thus meeting the criteria for a successful weight loss therapy (26). Therefore, IGB therapy has a value, albeit moderate and depending on the right patient choice. A certain advertising effect, however, can probably not be denied for IGB systems. The concept of restricting the gastric volume is easy to understand and might lead to expectations that will not be met in many cases. On the other hand, the low invasive procedure may suggest lower rates of side effects than we see in reality. Thus, it is the physician's responsibility to choose the right patient and to well explain benefits and side effects that can be expected. If this is done, IGB treatment ends up not to be a gimmick but a viable option for the selected patient and under the right conditions.

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