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***Retrospective Study***

**Efficacy of intragastric balloon on weight reduction: Saudi perspective**

Almeghaiseeb ES *et al*. Intragastric balloon on weight reduction

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**Abstract**

### *AIM*

### To evaluate the safety and efficacy of intragastric balloon (IGB) in weight reduction in obese patients referred to a tertiary hospital in the Kingdom of Saudi Arabia.

***METHODS***

Three hundred and one consecutive obese individuals, who underwent IGB placement during January 2009 to May 2015, were analyzed. The subjects aged 18 to 60 years and had a minimum BMI of 27 kg/m2. The IGB was placed under conscious sedation and kept for 6 mo. Anthropometric measurements were recorded during and after 6 mo of IGB removal.

***RESULTS***

The body weight, excess body weight, and BMI were significantly reduced at the time of IGB removal and 6 mo later. Body weight loss >10% was achieved in 224 subjects at removal of IGB. End of treatment success (ETS) and long-term success (LTS) were both significantly observed in women (70 *vs* 11) (71 *vs* 12.5) respectively. Excess BMI loss (EBMIL%) was significantly higher in subjects retaining the IGB for over 6 mo both at the removal [43.44 ± 19.46 (*n* = 221) *vs* 55.60 ± 28.69 (*n* = 80); *t* = 4.19, *P* = 0.0001] as well as at the end of 6 mo’ follow-up [46.57 ± 24.89 (*n* = 221) *vs* 63.52 ± 31.08 (*n* = 80); *t* = 4.87, *P* = 0.0001]. Within 3 d of IGB placement, two subjects developed pancreatitis and one subject developed cardiac arrhythmia. Intestinal obstruction due to displacement of IGB occurred in two subjects. All these subjects recovered uneventfully after immediate removal of the IGB.

***CONCLUSION***

IGB was effective in our cohorts. The observed weight reduction was maintained for at least 6 mo post IGB removal. IGB placement was safe with a satisfactory tolerance rate.

**Key words:** Intragastric Balloon; Weight Reduction; Saudi

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**Core tip:** Intragastric balloon (IGB) is a minimally invasive option for weight reduction. Several studies have demonstrated its superiority to lifestyle changes in reducing the morbidity and mortality associated with morbid obesity. This study evaluated the safety and efficacy of Medsil IGB in weight reduction of patients referred for weight reduction to a tertiary center in the Kingdom of Saudi Arabia.Endoscopic placement and keeping the Medsil IGB in situ for six months was proven to be safe, well tolerated and very effective for short and long term weight loss.

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## INTRODUCTION

Obesity, a medical condition in which body fat is accumulated in excess leading to severe negative health effects including reduced lifespan, affects an estimated 700 million people worldwide[1]. This pandemic health problem is much more serious threat to public health even when compared to alcohol consumption or tobacco smoking. Obesity decreases life expectancy by 6 to 7 years[2],while a BMI of 30–35 kg/m2 and > 40 kg/m2 reduces life expectancy by 2 to 4 years and by 10 years respectively[3]. Some estimates have even predicted that a BMI of > 40 reduces life expectancy by almost 20 years[4]. In addition to decline in the lifespan, this preventable cause of death also leads to quality of life deterioration owing to severe cardiologic, respiratory, dermatological, gastrointestinal, urinary, reproductive and psychiatric complications[5]. A recent estimate published in 2013 has put the overall prevalence of obesity at 28.7% (body mass index ≥ 30 kg/m2) in The Kingdom of Saudi Arabia, with women much more prone than men (33.5% *vs* 24.1%)[6].

Advanced cases of obesity require surgical interventions with drastic lifestyle modifications. Alternatively, mild to moderately obese subjects can achieve 5%-10% weight loss through exercise and dietary changes[7]. However, the weight gain recurs at high rates on cessation of these weight loss programs[8].Further, pharmacological agents have not been found to be any better than dietary and exercise programs. The therapeutic or lifestyle management of obesity is a long-term and arduous undertaking. Literature shows that long-term treatments, both dietary regimens and weight-loss programs following pharmacotherapy remain largely ineffective[9]. Further, conservative treatment is clearly ineffective in morbid obesity (BMI ≥ 40 kg/m2)[10], while bariatric surgery remains the only option with promising long-term results. However, subjects who are unwilling to consent for or do not qualify for the bariatric surgery end up having intragastric balloon as the best possible alternative[11].

These factors have fostered a spurt in interest in the utility of intragastric balloon (IGB) to achieve weight loss in excess of 10%. While earlier studies have documented the utility of various intragastric balloons in Saudi subjects[12,13], data on newer variants of balloons is lacking. Hence, this study is an attempt to evaluate the end of treatment success rates (ETS) and long-term treatment success rates (LTS) for recently introduced, GOST R certified intragastric balloon MEDSIL® in obese patients referred to a tertiary health clinic in Saudi Arabia.

**MATERIALS AND METHODS**

### *Subjects*

Three hundred and one subjects, consecutively opting for IGB therapy for weight loss at The Prince Sultan Military Medical City Hospital, Kingdom of Saudi Arabia between January 2009 and May 2015 were included in the study. Both men and women aged 14 to 65 years with a minimum BMI of 27 kg/m2 and medically free from or with one or two of the comorbidities namely diabetes mellitus, hypertension, bronchial asthma, back pain, or Knee joint complains were included. In general, patients rated only up to ASA class II were preferred. Further, subjects without active endocrine diseases and ability to tolerate the procedure were selected for the procedure. Patients classified into ≥ ASA category were excluded from the procedure. Baseline characteristics of the study subjects are presented in Table 1.

All the subjects underwent a routine clinical examination where information on anthropometrics and medical history was collected. Weight and height was measured with patients wearing no shoes and light clothing. Fasting blood sample was collected in the morning for the estimation of blood glucose concentration. Subsequent to the clinical evaluation of these results and after obtaining an informed written consent, MEDSIL® IGB was placed in the stomach through endoscopy.

***Intragastric Balloon Implantation and Removal***

All the subjects were treated with MEDSIL® IGB, silicon based saline filled bioenetric intragastric balloon (BIB) with a maximal volume of 700 mL (CSC MEDSIL, Russia). Patients were explained all the risks including perforation, bleeding, infection and adverse effects to the medicine, as well as the benefits and alternatives to the procedure prior to obtaining the informed written consent. All the patients understood the details and stated so.

For insertion of balloon, the patient was connected to monitoring devices and placed in left lateral position. In 297 patients, the device was implanted under procedural sedation and analgesia, with midazolam and fentanyl or pethidine. In 4 patients, the device implantation was done by inducing anesthesia using midazolam and intravenous propofol. Oxygen was provided continuously through a nasal cannula. Intravenous medications were administered through an indwelling cannula. After adequate conscious sedation was achieved, the patient was intubated and the endoscope was advanced under direct visualization to the duodenum.

Endoscope was withdrawn after complete examination for the presence of grossly anatomical contraindications. Balloon was inserted into the oral cavity and pushed into the stomach with a trocar followed by re-introduction of the endoscope. Under direct visualization, balloon was adjusted for proper placement followed by retraction of the push trocar wire. Balloon was then inflated with 400 mL to 700 mL of saline mixed with methylene blue. On achieving the desired inflation, the balloon catheter was gently pulled out leaving the balloon in stomach. The scope was gently retracted with careful examination of the colour, texture, anatomy, and integrity of the mucosa on the way out. The patient was subsequently transferred to the recovery area for observation.

The IGB was removed on completion of 6 months, with the duration extending by one or more months in some subjects owing to various reasons. Anthropometric measurements were recorded during and after 6 mo of IGB removal. Overnight fasting blood specimen was collected at removal for the estimation of blood concentration. Patients were also provided with walk in follow ups/clinic appointments on quarterly basis, but two third of the patients were seen on the fixed appointment only.

Similar protocol was followed for the removal of the balloon. After achieving adequate conscious sedation, the patient was intubated and the scope was advanced under direct visualization to the Stomach. After confirming absence of food or organic particles, an aspiration needle was inserted into the balloon followed by complete withdrawal of the fluid. The fully deflated balloon was withdrawn using a toothed forceps along the scope followed by routine follow up procedures.

Patients remained on their regular food without prescription of a hypocaloric diet. Post-insertion fasting blood glucose level estimation were scheduled and followed up to ensure that it was done as soon as treatment duration is completed.

***Measurements and statistics***

Weight loss variables like body weight (kg), BMI, body weight loss (BWL %) and excess BMI loss (EBL %) were measured at baseline, and during and after 6 months of IGB removal. A BWL value of > 10% at the time of IGB removal and after 6 months of IGB removal was considered an end of treatment success (ETS) and long-term success (LTS) respectively. EBL% was calculated using the formula [(Baseline BMI-Current BMI)/(Baseline BMI-25)] × 100. All the descriptive data are expressed as mean ± SD. Paired t-test was used to compare baseline and outcome variables for individuals, whereas unpaired *t*-test was used for gender and age based comparisons. Fisher’s exact test was used to evaluate the occurrence of number of patients with BWL% >10 between groups. The association of initial BMI and age with BWL% and EBL% was measured through Pearson correlation coefficient. A two-tailed *P* value of < 0.05 was considered statistically significant. Statistical Package for the Social Sciences (SPSS v. 18) was used for all the statistical tests.

## RESULTS

Apart from the expected post procedure symptoms like nausea, vomiting and upper abdominal discomfort, and no serious complications were observed during recovery from IGB placement. Balloon was removed a day to week earlier than 6 mo in 20 subjects, at the completion of 6 mo in 201 subjects and after a week to few months over 6 months in 80 subjects. In addition to the removal of IGB in 221 subjects owing to completion of the treatment duration, balloons were removed due to numerous other reasons as listed in the Table 2.

At the end of treatment, body weight, excess body weight, and BMI were significantly lowered as compared to initial measurements (Table 3; Figures 1, 2 and 3). The ETS rate, represented by number of patients with BWL% of > 10 was 74% (224 of 302 subjects). The fasting blood glucose remained statistically similar to the initial measurements. At the end of 6 mo after IGB removal, the body weight, excess body weight and BMI still remained significantly lower (Table 3). The body weight loss and BMI loss continued during post IGB removal phase, resulting in significantly higher measurements after 6 mo of removal as compared to the measurements taken during IGB removal. The LTS rates remained similar to ETS with just 2 more subjects added to the > 10% weight loss by the end of 6 mo IGB post-removal.

Statistical sub-analysis revealed different outcome when patients where compared according to gender and exercise habits (Table 4). The BWL%, BMI loss, and EBMIL% was significantly lesser in women at the end of treatment as well as after 6 mo of removal. However, significantly higher proportion of women achieved ETS (70 *vs* 11) and LTS (71 *vs* 12.5) rates. As expected, BWL%, BMI loss and EBMIL% was significantly higher in exercising cohort at the end of treatment. Fasting blood glucose level changes remained statistically similar with gender and exercise habit.

Age was not correlated with initial BMI or EMBIL% at initial or later phases of the study (Table 5). However, initial BMI was strongly correlated with BMI as well as EBMIL% measured at IGB removal as well as 6 months after removal.

The duration of IGB removal was also important in determining the EBMIL% (Figure 4). The EBMIL% was significantly higher in subjects retaining the IGB for over 6 months both at the removal [43.44 ± 19.46 (*n* = 221) *vs* 55.60 ± 28.69 (*n* = 80); *t* = 4.19, *P* = 0.0001] as well as at the end of 6 mo follow-up [46.57 ± 24.89 (*n* = 221) *vs* 63.52 ± 31.08 (*n* = 80); *t* = 4.87, *P* = 0.0001]. Duration of the IGB was also important in determining adverse complications in some of the individuals as outlined below.

In addition to the routine adverse events associated with IGB therapy, some of the patients experience unusual complication of spontaneous deflation and passage out of the digestive system. It must be noted that most of these patients had the balloon beyond the treatment duration. Out of five such cases, 2 women who had undergone IGB insertion at other institute, consulted us for removal after 1 year of insertion and three women operated at our clinic approached for removal at 8 mo. These five women underwent gastroscopy using X-ray and CT scan, which failed to reveal any traces of the IGB in the GI tract. The only plausible explanation for this is spontaneous rupture and excretion of the IGB without any knowledge of these patients.

One woman and one man developed clinical and biochemical pancreatitis on third day post IGB insertion. These symptoms subsided completely after immediate removal of IGB. One man developed arrhythmia 2 d post IGB insertion and recovered fully following immediate removal of IGB.

A woman with IGB got pregnant before the due date of removal and approached us at the end of first trimester with symptoms correlating intestinal obstruction. Esophagogastroduodenoscopy failed to detect balloon in the stomach. Imaging showed that balloon was lodged in middle of the jejunum. This IGB was surgically removed. The woman completed pregnancy without complications and gave birth to a normal offspring. Another man developed intestinal obstruction due to IGB dislodgement 7 mo post insertion that was detected through abdominal CT scan. This man recovered completely and uneventfully after laparoscopic surgery.

**DISCUSSION**

Subsequent to their introduction in 1982[14], numerous studies showed that IGBs are an effective and low cost method to achieve temporary weight loss in morbidly obese individuals, leading to significant decrease in morbidity and mortality rates[15,16].The promising outcomes have fuelled the development of numerous fluid or air filled IGBs over the years. The newer variants are becoming much less invasive compared to surgical interventions in morbid obesity, though latter options remain primary approach in super-obese patients with a BMI over 50[17].IGBs are also employed as a preoperative tool in bariatric surgery as its weight reducing effects significantly reduces the mortality, morbidity and risks associated with this invasive surgery[18].

Variety of intragastric balloons have been studied in numerous studies for safety and efficacy in Saudi subjects[12,19]. However, the newer variant of intragastric balloon Medsil remains to be tested in this population. In this study, we tested the end of the treatment and long term success rates for this device in a large Saudi cohort. Similar to a 2014 report on Czech subjects[20],these balloons were well tolerated in Saudi subjects analysed in this study.

This study demonstrated a clear benefit of Medsil balloon on body composition, as six mo placement of the balloon lead to significant reduction in body weight. The mean BMI loss (4.75 and 5.20 kg/m2) and body weight loss (13.08 and 14.30 kg) at IGB removal and after six months of removal were comparable to both the results of using other balloons or Medsil balloons. These studies have reported a BMI loss of 5.7-6.7 kg/m2 and weight loss of 14.7-17.8 kg[16,21,22],with one study using the same device reporting a BMI loss of 5.5 kg/m2 and weight loss of 18.4 kg[20].Table 6 presents a comparison showing the similarities of outcomes in body composition reported by earlier studies. It must be noted that the BMI loss which corresponds to weight-loss was on the higher end as compared to other studies after one year of completion of the treatment.

Two reviews have extensively evaluated the weight loss due to IGBs. Dumonceau *et al*[21] 2008 analysed 4877 patients from 30 studies and recorded a mean weight loss of 17.8 kg (or a BMI loss of 4-9 kg/m2). Another systematic review reported similar outcomes and revealed that, combined with lifestyle changes, IGBs provide an effective means for achieving a significant temporary weight loss, though the long term outcomes remain yet to be understood[16].Our results of a significant weight loss and a large number of patients achieving and maintaining > 10% BWL from the IGB removal to follow-up after 6 months, clearly suggests that long term results can be achieved through this method. In addition to the balloon, initial BMI, adherence to the lifestyle changes and patients level of motivation are highly likely to play an important role in achieving long term results.

Bioenetric intragastric balloons, which are now known as Orbera Intragastric Balloon (Apollo Endosurgery, Austin, TX, United States) are the most commonly used balloons. A comparison with the existing literature showed that body weight loss was less than expected, however not too less (Table 7).

The number of individuals achieving > 10% BWL increased from 224 to 226 from balloon removal to at 6 mo follow-up. This amounts to an increase in the number of patients achieving > 10% BWL during follow-up. These results are highly impressive as two of the earlier studies have showed that only 48%[23] or 55%[24] of the patients went on to continue losing weight from balloon removal to follow-up at 1 year. Our study results are very promising in this aspect.

The fasting blood glucose level remained statistically similar both during balloon removal as well as after 6 mo of follow-up. Bugza *et al*[20] 2014 also reported a similar result, though they demonstrated a positive effect of the balloon on glucose tolerance. On the contrary, earlier studies by Mathus-Vliegen and Konopko-Zubrycka have demonstrated a statistically significant reduction in fasting blood glucose levels through intragastric balloons[11,24].These contradictory findings remain to be evaluated by meta-analysis to reveal the actual association.

Our study, though of higher strength due to large sample size, had a few limitations. A follow-up period of more than six months (at least 1 year) including tracking of comorbidities along with body conversion parameters would have been more insightful. The evaluation of fasting blood glucose levels could have been more meaningful if glucose tolerance and glycated hemoglobin levels were also included. Compared to earlier report of maintenance of > 10% weight loss in about 25% of patients for almost 30 mo[25],75% of the subjects who achieved this result after 6 mo follow-up in our study seem to be responding much better. Looking at the similarities of BIB and Medsil balloons, it is highly likely that our subjects will be able to maintain weight loss for long term. However, it must be noted that our study provides the first report on the follow-up parameters for Medsil balloons.

In conclusion, it could be concluded that Medsil intragastric balloons are safe and effective for Saudi subjects and more than three fourth of the subjects can be expected to achieve long term weight loss.

**COMMENTS**

***Background***

Obesity is a major pan-endemic health problem in the Kingdom of Saudi Arabia affecting about 30% of the population. Literature shows that dietary regimens and weight-loss programs following pharmacotherapy remain largely ineffective. Bariatric surgery is the most effective long term option, however the majority are either reluctant to undergo surgery or do not qualify for medical reasons.

***Research frontiers***

Intragastric balloons are of proven benefit as an alternative or a bridge to surgery, however the evidence for its utility particularly the newer version such as the intragastric balloon (IGB) MEDSIL® in Saudi Arabia is lacking.

***Innovations and breakthroughs***

This study is an attempt to evaluate its long-term treatment success rate in obese patients referred to a tertiary health clinic in Saudi Arabia. Endoscopic placement and keeping the Medsil IGB in situ for six months was proven to be safe, well tolerated and very effective for short and long term weight loss.

***Applications***

The intragastric balloons are well tolerated and are effective in weight reduction.

***Peer-review***

It is a retrospective study but of a very big cohort and the IGB is a new commercialized one. It is a well done paper and the language is good too.

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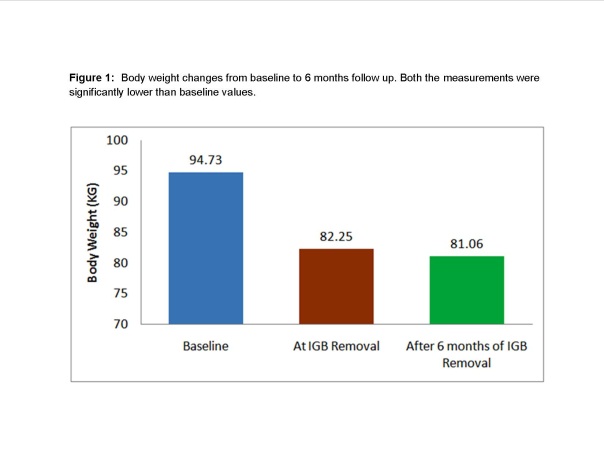
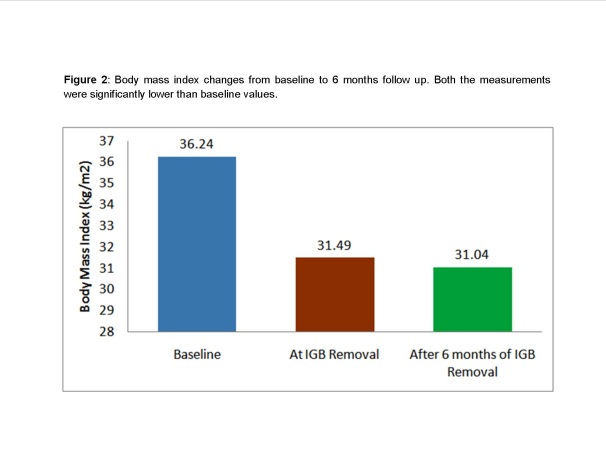
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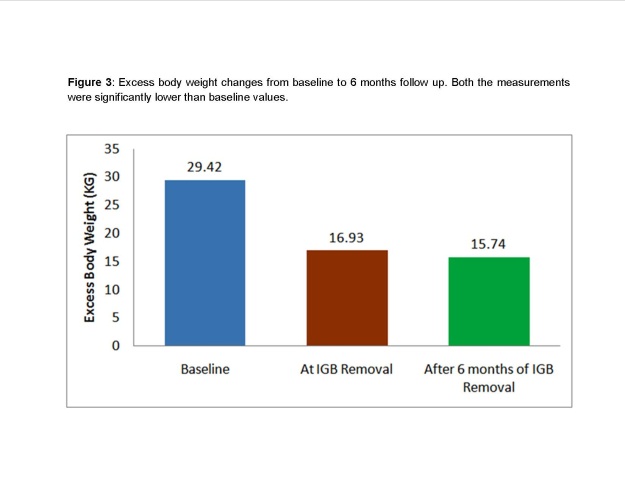
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**P-Reviewer:** de Castro ML, Doldi SB **S-Editor:** Qi Y **L-Editor: E-Editor:**

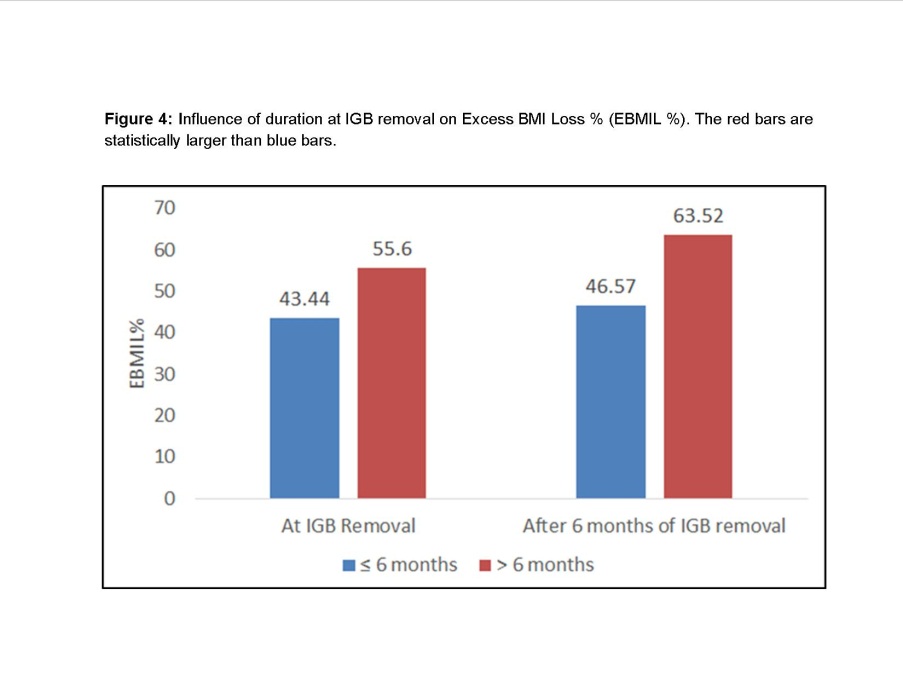
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A B

****

**C**

**Figure 1 from baseline to 6 mo follow up.** A: Body weight changes. Both the measurements were significantly lower than baseline values; B: Body mass index changes. Both the measurements were lower than baseline values; C: Excess body weight changes. Both the measurements were significantly than baseline values.

****

**Figure 4 Influence of duration at intragastric balloon removal excess BMI loss% (EBMIL %).** The red bars are statistically larger than bars.

**Table 1** **Baseline characteristics of the subjects (*n* = 301) included for intragastric balloon therapy**

|  |  |  |
| --- | --- | --- |
| **Characteristics** | **Average (mean ± SD)1** | **Range** |
| Age (yr) | 34.34 ± 10.38 | 14 – 65 |
| Female (%) | 67 | |
| Body Weight (Kg) | 94.73 ± 16.38 | 67 – 203 |
| Height (cm) | 161.52 ± 6.11 | 150 – 179 |
| Body Mass Index (BMI; kg/m2) | 36.24 ± 5.24 | 27.10 – 70.24 |
| Excess Body Weight (ideal BMI 25) | 29.42 ± 14.61 | 5.6 – 130.8 |
| Fasting Blood Glucose (mg/dL) | 100.15 ± 29.26 | 69 – 240 |

1Applicable to all the values except % females.

**Table 2 Reasons for removal of the intragastric balloon**

|  |  |  |  |
| --- | --- | --- | --- |
| **Reason for removal** | **Number** | **Reason for removal** | **Number** |
| Treatment duration complete (TDC) | 221 | Intolerance | 4 |
| Abdominal pain | 1 | Vomition | 1 |
| Miscellaneous | 2 | TDC and intolerance | 1 |
| TDC and abdominal pain | 19 | TDC and vomition | 10 |
| TDC and discomfort | 24 | TDC and other reasons | 2 |
| Intolerance and abdominal pain | 4 | Intolerance and vomition | 2 |
| Intolerance and other reason | 1 | Abdominal pain and vomition | 3 |
| Abdominal pain and other reason | 1 | Discomfort and other reason | 1 |
| TDC, abdominal pain and vomition | 1 | Intolerance, abdominal pain and vomition | 2 |
| TDC, abdominal pain and discomfort | 1 |  |  |

*Abdominal discomfort* was a state of tolerable uneasiness without pain; *Abdominal pain* involved a state of colic; *Vomition* was a state of uncontrolled expulsion of gastric contents; *Intolerance* was a state wherein subjects experienced a mix of side effects and were unable to tolerate. *Other reasons* were a variety of situations that did not show a consistent pattern. TDC: Treatment duration complete.

**Table 3 Weight related measurements at intragastric balloon removal and after 6 mo of removal**

|  |  |  |
| --- | --- | --- |
| **Characteristics** | **At removal**  **(mean ± SD)a** | **After 6 mo of**  **removal** |
| Body weight (kg)  (Min - Max) | 82.25 ± 14.73b  (55 - 181) | 81.06 ± 14.84b  (53 - 181) |
| Body Weight Loss (kg)  (Min - Max) | 12.48 ± 5.16  (0 - 30) | 13.67 ± 6.65b  (-1 - 42) |
| Body Weight Loss (%)  (Min - Max) | 13.08 ± 4.81  (0 – 35.29) | 14.30 ± 6.12b  (-0.85 – 32.14) |
| Number of Patients with BWL% > 10 | 224 | 226 |
| Excess Body Weight (ideal BMI 25)  (Min - Max) | 16.93 ± 13.44b  (-10.61 – 108.75) | 15.74 ± 13.73b  (-12.72 – 107.04) |
| Body Mass Index (BMI; kg/m2)  (Min - Max) | 31.49 ± 4.88b  (20.96 – 62.63) | 31.04 ± 5.01b  (20.44 – 63.14) |
| Body Mass Index Loss (BMI; kg/m2)  (Min - Max) | 4.75 ± 1.87  (0 – 11.43) | 5.20 ± 2.40b  (-0.36 – 15.43) |
| Excess BMI Loss (EBMIL %)  (Min - Max) | 46.67 ± 22.88  (0 – 161.09) | 51.07 ± 27.66b  (-2.12 – 195.53) |
| Fasting Blood Glucose (mg/dL)  (Min - Max) | 98.67 ± 20.28  (71 - 187) | -- |

b*P* < 0.001, a*P* < 0.05.

**Table 4** **Association of gender and exercise on end of treatment success and long term success**

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| **Characteristics** | **At removal (mean ± SD)a** | | | | **After 6 mo of removal** | |
| **Gender** | | **Exercise** | | **Gender** | |
| **Male**  **(*n* = 72)** | **Female**  **(*n* = 229)** | **Yes (*n* = 131)** | **No**  **(*n* = 170)** | **Male**  **(*n* = 72)** | **Female**  **(*n* = 229)** |
| Body Weight Loss (%) | 14.99 ± 4.72 | 12.48 ± 4.68b | 15.22 ± 4.81 | 11.43 ± 4.11b | 16.71 ± 6.72 | 13.53 ± 5.72b |
| Number of Patients with BWL% > 10 | 8 | 160b | 113 | 111b | 9 | 163b |
| Body Mass Index Loss (BMI; kg/m2) | 5.56 ± 1.92 | 4.49 ± 1.78b | 5.47 ± 1.85 | 4.19 ± 1.68b | 6.19 ± 2.72 | 4.89 ± 2.20b |
| Excess BMI Loss (EBMIL %) | 51.61 ± 24.75 | 45.12 ± 22.09a | 55.66 ± 25.12 | 39.74 ± 18.24b | 57.48 ± 30.27 | 49.06 ± 26.53a |
| Fasting Blood Glucose Reduction (%) | -0.59 ± 13.29 | -0.92 ± 13.14 | -0.37 ± 13.10 | -1.25 ± 13.22 | -- | -- |

b*P* < 0.001, a*P* < 0.05.ETS: End of treatment success; LTS: Long term success.

**Table 5** **Correlation of age and initial BMI with weight-linked parameters**

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
|  | **Age** | **Initial**  **BMI** | **BMI**  **At removal** | **EBMIL%**  **At removal** | **BMI**  **After 6m of removal** | **EBMIL%**  **After 6m of removal** |
| Age | 1 | 0.074 | 0.082 | -0.108 | 0.082 | -0.108 |
| Initial  BMI | 0.074 | 1 | 0.934b | -0.413 b | 0.891b | -0.376b |

Values are Pearson Correlation Coefficient (r); b*P* < 0.01.

**Table 6** **Studies reporting outcomes in body composition after placement of intragastric balloon**

|  |  |  |
| --- | --- | --- |
| **Study** | **Weight loss at removal (Duration of placement in months)** | **Weight loss at follow-up (Duration of follow-up in months)** |
| Mathus-Vliegen *et al*[26], 2005 | 21.3 (12) | 12.6 (12) |
| Herve *et al*[27], 2005 | 12.0 (6) | 8.6 (12) |
| Doldi *et al*[28], 2004 | 15.5 (6) | -1.3 (14) |
| Melissas *et al*[29], 2006 | 41.6% EWL (6) | 23.9% EWL (6-30) |
| Angrisani *et al*[30], 2006 | 32.9% EWL (6) | 27.1% EWL |
| Ganesh *et al*[31], 2007 | 4.4 (6) | 1.5 (6-12) |
| Ohta *et al*[32], 2009 | 12 (6) | 6.4 (12) |
| Gumurdulu *et al*[33], 2013 | 12.4 (6) | 9.7 (6) |
| Buzga *et al*[20], 2014 | 18.4 (6) | - |
| Present study | 13.08 (6) | 14.30 (6) |

Weight in kgs; EWL: Excess weight loss.

**Table 7 Comparison of weight-loss with other types of intragastric balloons**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Balloon** | **Type (volume)** | **Material** | **Weight loss**  **(EOT in months)** | **Ref.** |
| Medsil BIB | Fluid-supplied  (400-700 ml saline) | Silicone | 12.48 ± 5.16 Kg (6 mo) | This study |
| Orbera (Apollo Endosurgery) | Fluid-supplied  (400-700 ml saline) | Silicone | 16.9 ± 0.9 Kg (6 mo) | Gaur *et al*[34], 2015 |
| The Elipse™ (Allurion Technologies) | Fluid-supplied  (450-550 mL filling fluid) | NA | 2.4 Kg (6 wk) | Machytka *et al*[35], 2016 |
| ReShape Duo® Integrated DualBalloon System (ReShape medical) | Fluid-supplied  (900 mL; 450 mLX2 saline) | Silicone | 25.1 ± 1.6% EWL (6 mo) | Ponce *et al*[36], 2015 |
| Spatz Adjustable Balloon system (Spatz FGIA) | Fluid-supplied  (400-600 mL saline) | Silicone | 24 kg (at 12 mo) | Brooks *et al*[37], 2014 |
| Heliosphere BAG® (Helioscopie) | Air-supplied  (950 mL air) | Polyurethane and silicone | 16 ± 7 kg (6 mo) | Giardiello *et al*[38], 2012 |
| Obalon® Gastric Balloon (Obalon Therapeutics) | Air-supplied  (250 mL air, nitrogen) | NA | 5 kg (12 wk) | Mion *et al*[39], 2013 |

EOT: End of treatment.