

Boosting Weight Loss after Revisional RYGB: A Double-Blind- RCT of Liraglutide and Placebo use

Mohamed Hany MD^{a,b}, Bart Torensma MSc, PhD^c, Mohamed Ibrahim MD^a, Ahmed Zidan MD^a, Ann Samy Shafiq Agayabya MD^a, Mohamed Hisham BSc^d, Iman El Sayed MSc, PhD^e*

^aDepartment of Surgery, Medical Research Institute, Alexandria University, Egypt

^bMadina Women's Hospital, Alexandria University, Egypt

^cClinical Epidemiologist, Leiden University Medical Center (LUMC), Leiden, The Netherlands

^dDepartment of Pharmacology, Alexandria University, Egypt

^eBiomedical Informatics and Medical Statistics Department, Medical Research Institute, Alexandria University, Alexandria, Egypt



In accordance with «EACCME criteria for the Accreditation of Live Educational Events», please disclose whether you have or you have not any conflict of interest with the companies:

We have no potential conflict of interest to report



Background

Laparoscopic sleeve gastrectomy (LSG) has high reported revision rates (**10-22%**)

Weight regain and insufficient weight loss (**±70%**)

Revisional bariatric surgery → RYGB is the most popular performed.

But **revisional bariatric surgery** has worse outcomes than
primary bariatric surgery

Study endpoints

Additional weight reduction with **liraglutide** after RRYGB compared to the RRYGB with **placebo** use.

Weight loss @1 months || 6 weeks || 6 months

&

Changes in metabolic biomarkers measured at the end of treatment



Liraglutide vs Placebo pens

Liraglutide (Saxenda)

Post-operative week-6
with
0.6 mg daily injections

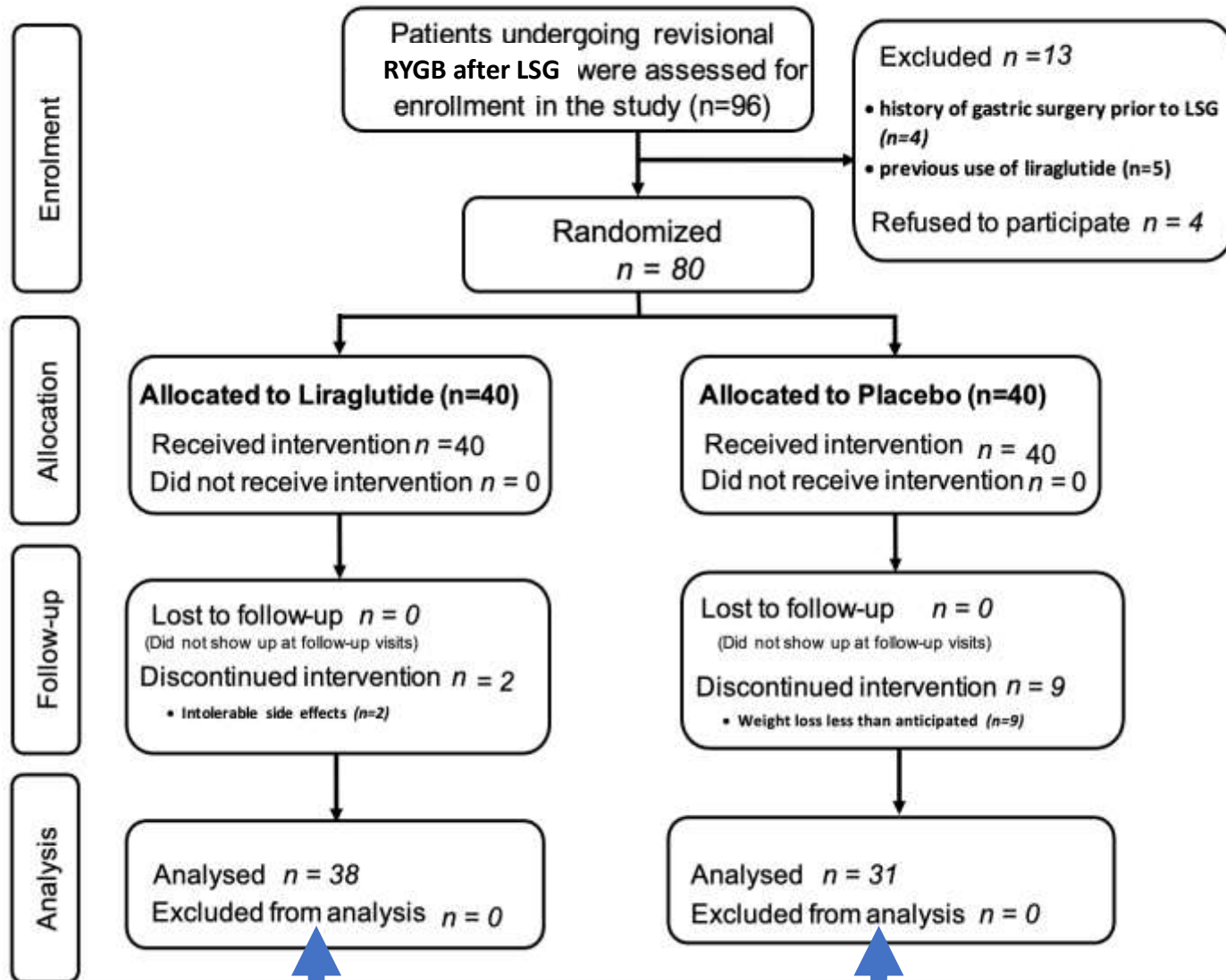
**Max of 3 mg daily after 5
weeks, continued for 24
weeks.**

Placebo (NALC 0.9%)

Post-operative week-6
with
0.6 mg daily injections

**Max of 3 mg daily after 5
weeks, continued for 24
weeks.**

CONSORT Flow Diagram



Results

Weight loss

**Liraglutide
%TWL**

6 weeks 12.66±1.69

6 months 18.17±2.17

**Delta
%TWL**

+1.41

+2.64

**Placebo
%TWL**

6 weeks 11.25±1.62

6 months 15.53±1.64

**Liraglutide
%EWL**

6 weeks 41.89±5.97

6 months 60.21±8.40

**Delta
%EWL**

+5.42

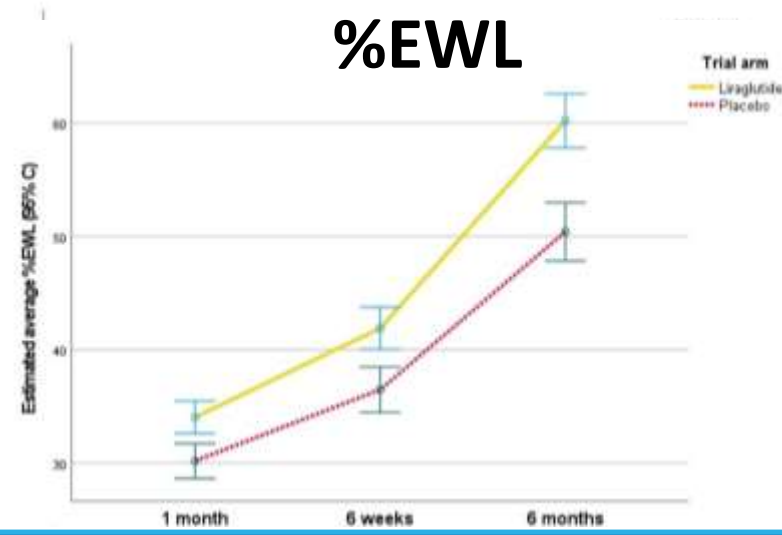
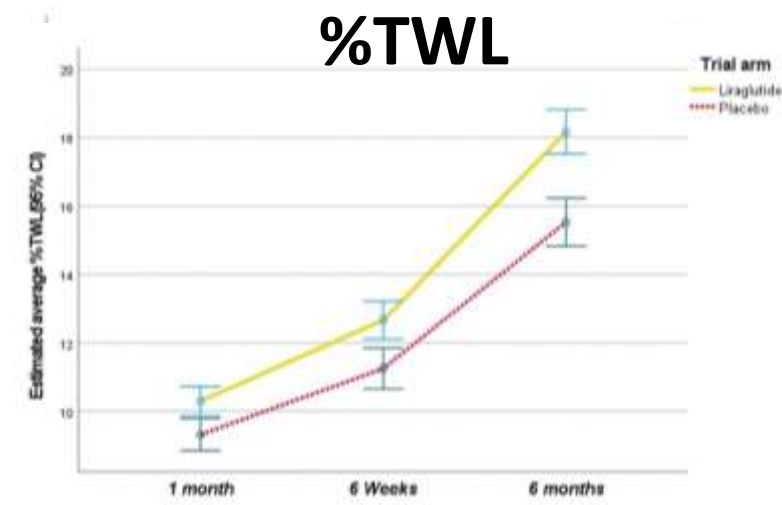
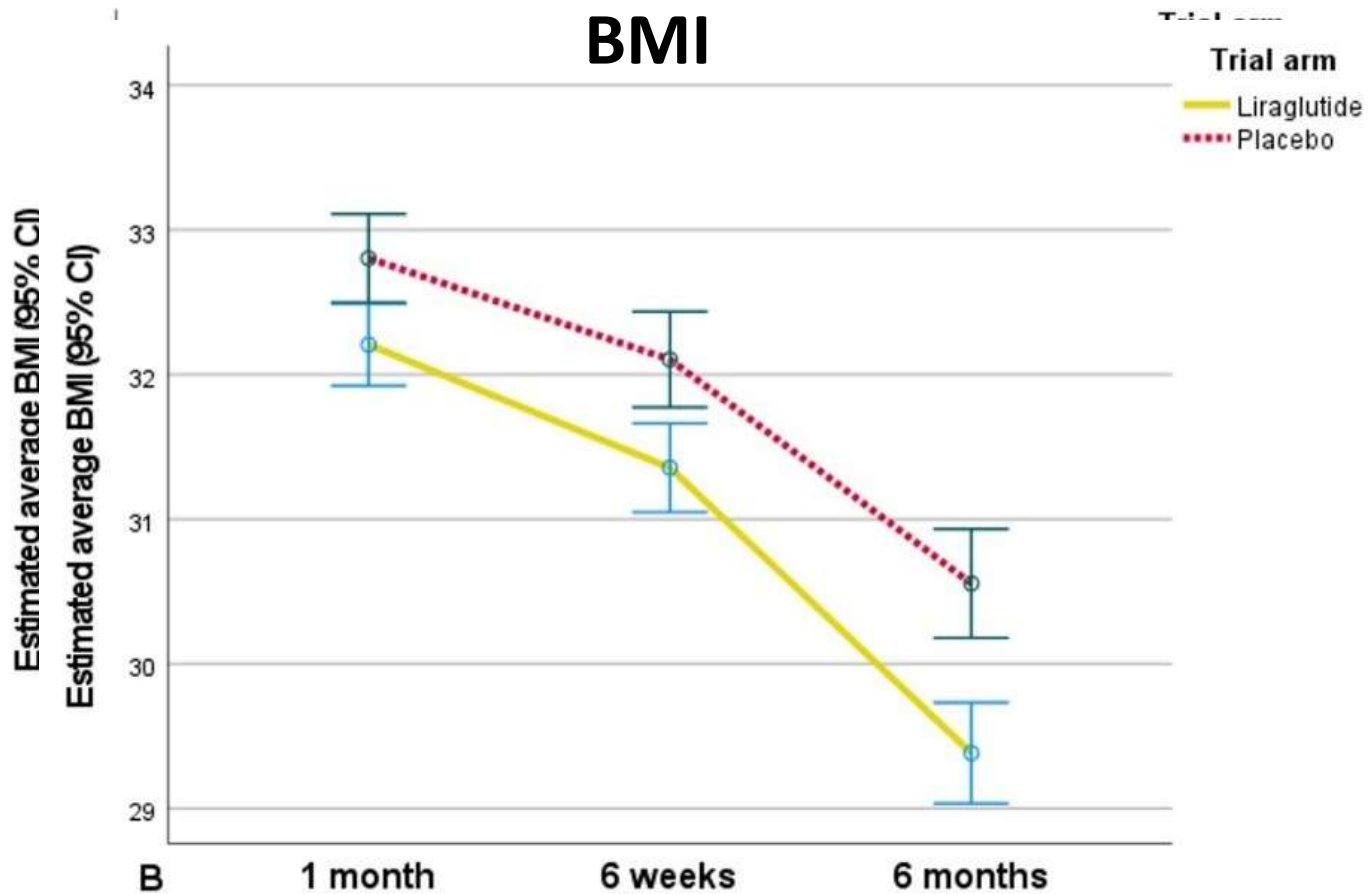
+9.8

**Placebo
%EWL**

6 weeks 36.47±5.06

6 months 50.41±5.37





Results

metabolic biomarkers

- Lipid profile, fasting blood sugar, HbA1c, fasting insulin levels, leptin, ghrelin, and PYY levels

NO Significant differences

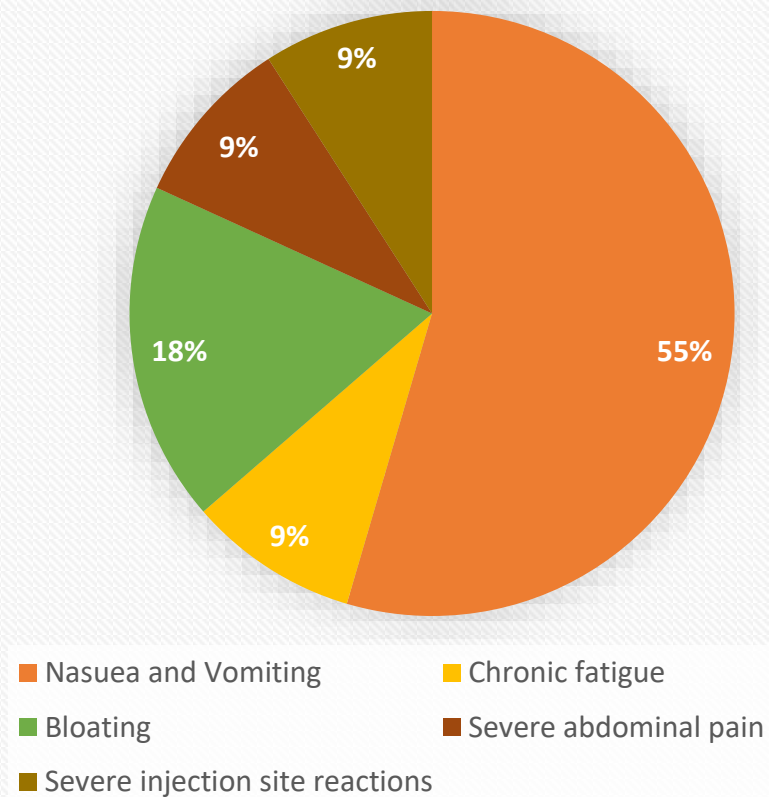
- GLP-1 **decreased** significantly in the **liraglutide** cohort ($p < 0.001$), (*GLP-1 increased in placebo*)
- GIP increased in both cohorts, with a **significantly more increase** in the **liraglutide** cohort ($p = .014$)

Severe and intolerable side effects

- 5.26% in the liraglutide group after 3 and 4 weeks.

- 81.6% could reach and tolerate the 3mg/day dose,
- 18.4% continued lower doses (range 2.4-1.2 mg/day).

From the 5.26%



Limitations

- A longer follow-up is needed to assess the durability of the weight loss outcomes.
- The resolution of diabetes mellitus should be confirmed when the patients are no longer on medications.

Conclusion

- Adjunctive use of liraglutide has significantly added weight loss and effect on associated medical problems with good tolerability.
- The added value from liraglutide use can help overcome the inherently lower weight loss outcomes of RBS.



mohamed.ashour@alexu.edu.eg

ORCID: 0000-0001-6650-8112

+20 100 2600970

Egypt



Thank you



bart@torensmaresearch.nl

ORCID: 0000-0003-0274-9608

+316 41 38 90 70

The
Netherlands

