ORIGINAL CONTRIBUTIONS



TSH Normalization in Bariatric Surgery Patients After the Switch from L-Thyroxine in Tablet to an Oral Liquid Formulation

Poupak Fallahi¹ • Silvia Martina Ferrari¹ • Stefania Camastra¹ • Ugo Politti¹ • Ilaria Ruffilli¹ • Roberto Vita² • Giuseppe Navarra³ • Salvatore Benvenga² • Alessandro Antonelli¹

Published online: 7 June 2016 © Springer Science+Business Media New York 2016

Abstract

Objective Drug malabsorption is one of the potential troubles after bariatric surgery. Evidence for diminished levothyroxine (L-T4) absorption has been reported in patients after bariatric surgery.

Methods This study reports 17 cases of hypothyroid patients [who were well replaced with thyroxine tablets (for >1 year) to euthyroid thyrotropin (TSH) levels before surgery (13 Rouxen-Y gastric bypasses (RYGB); 4 biliary pancreatic diversions (BPD))]. From 3 to 8 months after surgery, these patients had elevated TSH levels. Patients were then switched from oral tablets to a liquid L-T4 formulation (with the same dosage, 30 min before breakfast).

Results Two-three months after the switch, TSH was significantly reduced both in patients treated with RYGB, as in those treated with BPD, while FT4 and FT3 levels were not significantly changed (RYGB group, TSH μ IU/mL: 7.58 ± 3.07 vs 3.808 ± 1.83, *P* < 0.001; BPD group, TSH μ IU/mL: 8.82 ± 2.76 vs 3.12 ± 1.33, *P* < 0.01).

Conclusions These results first show that liquid L-T4 could prevent the problem of malabsorption in patients with BPD and confirm those of previous studies in patients submitted to RYGB, suggesting that the L-T4 oral liquid formulation could circumvent malabsorption after bariatric surgery.

Alessandro Antonelli alessandro.antonelli@med.unipi.it

- ¹ Department of Clinical and Experimental Medicine, University of Pisa, Via Savi 10, 56126 Pisa, Italy
- ² Department of Clinical and Experimental Medicine, Section of Endocrinology, University of Messina, Messina, Italy
- ³ Department of Human Pathology, University Hospital of Messina, Messina, Italy

Keywords Bariatric surgery · Roux-en-Y gastric bypass · Biliary pancreatic diversion · Levothyroxine malabsorption · Liquid L-T4 · Hypothyroidism

Introduction

In the past decades, demand for bariatric surgery has globally increased, and about 101,000–180,000 surgeries are performed annually in the USA alone [1–4].

Surgical procedures include different types of operations: a—purely restrictive (gastric banding, gastroplasty); b—restrictive with limitation of digestive capacity (sleeve gastrectomy); c—restrictive/malabsorptive (gastric bypass); d purely malabsorptive (biliopancreatic diversion, jejunoileal bypass). Malabsorptive procedures can lead to nutritional deficiencies, [5] or drug malabsorption [6].

People with severe obesity receiving bariatric surgery often have multiple medical comorbidities requiring multidrug treatments. After diversionary procedures, drug malabsorption is a potential concern. In fact, nearly all oral agents are absorbed mainly in the small intestine, which is bypassed in several bariatric procedures. Other factors that impair drug absorption are the following: diminished opportunity for mucosal exposure and changes in drug solubility and dissolution resulting from alterations in intestinal pH [6]. The most consistent evidence for diminished absorption has been found for cyclosporine, phenytoin, rifampin, and thyroxine [7].

The prevalence of hypothyroidism in patients with morbid obesity range from 12 to 25 % [8, 9].

Moreover, in obese males, insulin resistance is significantly related with thyroid function impairment [10].

Levothyroxine (L-T4) is the gold standard and effective replacement therapy for patients with hypothyroidism for autoimmune thyroiditis or after radioablative therapies, or suppressive therapy after surgical removal of thyroid cancer [11–13].

To obtain an effective therapy, some requisites are necessary, as L-T4 products are of optimal quality, [14] patients' compliance, [15] dissolution of the hormone in the stomach, [16] adequate absorption in the intestines, [17, 18], and normal metabolism [19]. Several diseases and drugs affect the absorption and metabolism of L-T4 [20]. Gastritis causes L-T4 malabsorption, by altering the gastric juice pH, thereby affecting L-T4 dissolution [21–23].

Evidence for diminished L-T4 absorption has been reported in patients after bariatric surgery [24, 25].

Recently, four cases of hypothyroid patients who were well replaced with thyroxine tablets to euthyroid thyrotropin (TSH) levels prior to Roux-en-Y gastric bypass (RYGB) surgery, and developed elevated TSH levels after the surgery, have been reported [26]. In these patients, TSH responded reversibly to switching from L-T4 treatment with oral tablets to a liquid formulation.

Here, we report our experience with the use of liquid L-T4 in patients with bariatric surgery who had developed elevated TSH levels after the surgery.

Methods

We have evaluated 17 out-patients with morbid obesity, recruited from the end of 2013 to the first months of 2015 (ten females, seven males; age 31–59 years), who were diagnosed with hypothyroidism (11 affected by autoimmune thyroiditis and six treated with total thyroidectomy for multinodular goiter) before the bariatric surgery, and were well replaced with thyroxine tablets (for >1 year) to euthyroid TSH, free thyroxine (FT4), and free triiodothyronine (FT3) levels. Thirteen patients were treated with RYGB, [27] while four with biliary pancreatic diversions (BPD) [28].

The RYGB group included nine women and four men, mean age of 45 ± 9 years; body mass index (BMI) (before surgery) in this group was $\geq 40 \text{ kg/m}^2$ (mean, $42.9 \pm 3.5 \text{ kg/m}^2$). The hypothyroidism in this group was due in nine cases to autoimmune thyroiditis and in four to total thyroidectomy for multinodular goiter.

The BPD group included one woman and three men, mean age of 42 ± 7 years; BMI (before surgery) in this group was $\geq 40 \text{ kg/m}^2$ (mean, $44.1 \pm 4.4 \text{ kg/m}^2$). The hypothyroidism in this group was due in two cases to autoimmune thyroiditis and in two to total thyroidectomy for multinodular goiter.

From 3 to 8 months after surgery, these patients had elevated TSH levels.

We decided to treat these patients with oral liquid L-T4. Patients were switched from oral tablets to a liquid formulation of L-T4 (Tirosint® fiala monouso, IBSA Farmaceutici Italia) (with the same dosage, 30 min before breakfast). Two-three months after the switch, circulating TSH, FT4, and FT3 were re-evaluated. Clinical data and medication information were collected before recruitment from medical records. Patients with other possible causes of altered L-T4 absorption (such as atrophic gastritis and use of medications associated with impaired L-T4 absorption) were not evaluated.

All patients signed an informed consent. The study was approved by the Ethical Committee.

Serum FT4 (normal range, 0.7–1.7 ng/dL), FT3 (normal range, 2.7–4.7 pg/mL), and serum TSH (normal range, 0.4–4 μ IU/mL) were determined in all samples by electrochemiluminescence immunoassay (Roche Corporation, Indianapolis, IN, USA). The concentration of each hormone at baseline, and after the switch, was calculated as a mean of the two samples collected before the L-T4 dose.

Data Analysis

Values are given as mean \pm SD for normally distributed variables, otherwise as median and interquartile range. Mean group values were compared by using one-way analysis of variance (ANOVA) for normally distributed variables (age and BMI). Post hoc comparisons on normally distributed variables were carried out using the Bonferroni-Dunn test. Proportions were compared by the χ^2 test. Simple regression was used to evaluate the correlation among changes of TSH (after the switch—baseline), vs changes of FT4, or FT3.

Results

After bariatric surgery (at the time of TSH re-evaluation, 3 to 8 months after surgery) in the RYGB group, BMI was reduced from 42.9 ± 3.5 to 37.9 ± 2.7 kg/m² (P < 0.01), while in the BPD group, BMI was reduced from 44.1 ± 4.4 to 38.6 ± 3.7 kg/m² (P < 0.01).

Although the reduction of body weight, with the same L-T4 dosage (mean $192 \pm 32 \ \mu g/day$) in tablets, 30 min before breakfast, after bariatric surgery, TSH was increased in both groups (Table 1), while FT4 and FT3 levels were not significantly changed. The only symptom observed in patients with high TSH was fatigue in 11/17 (65 %) of patients. Then patients were switched from oral tablets to a liquid formulation of L-T4 (Tirosint® fiala monouso, IBSA Farmaceutici Italia) (with the same dosage, 30 min before breakfast). Two–three months after the switch, when circulating TSH, FT4, and FT3 were re-evaluated, in the RYGB BMI was reduced from 37.9 ± 2.7 to $36.3 \pm 2.5 \ \text{kg/m}^2$ (P < 0.01), while in the BPD group, BMI was reduced from 38.6 ± 3.7 to $37.9 \pm 3.4 \ \text{kg/m}^2$ (P > 0.05). Table 1TSH, FT4, and FT3levels before and after bariatricsurgery

RYGB		BPD		P value (by ANOVA)	
Before surgery	After surgery	Before surgery	After surgery	RYGB	BPD
2.21 ± 0.91	7.58 ± 3.07	2.65 ± 0.86	8.82 ± 2.76	<0.001	0.005
1.27 ± 0.31	1.18 ± 0.37	1.31 ± 0.28	1.20 ± 0.36	ns	ns
3.12 ± 0.91	2.98 ± 0.98	2.99 ± 0.92	2.87 ± 0.96	ns	ns

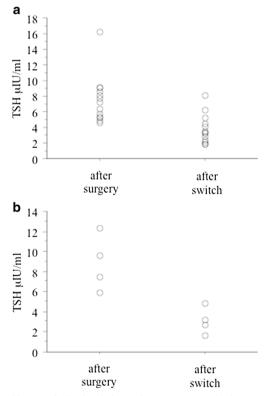
Serum TSH, normal range 0.4–4 μ IU/mL; serum FT4, normal range 0.7–1.7 ng/dL; serum FT3, normal range 2.7–4.7 pg/mL

RYGB Roux-en-Y gastric bypass, BPD biliary pancreatic diversions, ns no significance

After the switch from oral tablets to a liquid formulation of L-T4, circulating TSH levels were significantly reduced in both groups, while FT4 and FT3 levels were not significantly changed (RYGB group, TSH μ IU/mL: 7.58 ± 3.07 vs 3.808 ± 1.83, *P* < 0.001; BPD group, TSH μ IU/mL: 8.82 ± 2.76 vs 3.12 ± 1.33, *P* < 0.01) (Fig. 1).

TSH FT4 FT3

The comparison of TSH values before surgery, 3 to 8 months after surgery, and 2–3 months after the switch are shown in Fig. 2, and data show that the TSH values after the switch are higher, even if not significantly (P = 0.064), with respect to the TSH values before surgery.



A negative correlation between decrease of TSH (after the switch—baseline), vs the increase of FT4 (after the switch—baseline) was observed by simple regression (r = 0.754, P = 0.007) (Fig. 3). While no significant association was observed between changes of TSH (after the switch—baseline) and changes of FT3 (after the switch—baseline).

There was no significant difference between gender in changes of TSH values after surgery, or in the response to the switch to the liquid L-T4 formulation.

No patient before and after bariatric surgery, or after the switch had euthyroid sick syndrome [29].

Albumin levels were in the normal range (3.5 to 5.5 g/dL) in all patients before and after bariatric surgery, or after the switch.

Discussion

The abovementioned data suggest that the L-T4 oral liquid formulation could avoid the problem of the malabsorption after bariatric surgery. The obtained results first show that liquid L-T4 could prevent the problem of the malabsorption in patients with BPD and confirm those obtained in a previous study in patients submitted to RYGB. The serum TSH levels

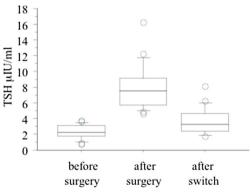


Fig. 1 Change of TSH levels in RYGB group (**a**), or in the BPD group (**b**), after surgery, and 2–3 months after the switch from oral tablets to a liquid formulation of levothyroxine (with the same dosage, 30 min before breakfast). There was no significant difference in the decrease of TSH in RYGB group vs the BPD group

Fig. 2 Comparison of TSH values before surgery, 3 to 8 months after surgery, and 2–3 months after the switch in all bariatric patients, are shown. The TSH values (box-plot) after the switch are higher, even if not significantly (P = 0.064), with respect to the TSH values before surgery

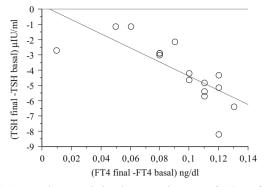


Fig. 3 A negative correlation between decrease of TSH (after the switch—baseline) (TSH final—TSH basal), vs the increase of FT4 (after the switch—baseline) (FT4 final—FT4 basal), was observed by simple regression (r = 0.754, P = 0.007)

in our patients (according to the selection criteria) were increased after bypass surgery, suggesting a malabsorption of L-T4, [24–26] in agreement with Azizi et al., Bevan et al., and Pirola et al.

RYGB reduces the size of the stomach to a small pouch about the size of an egg. It does this by stapling off a section of it. This reduces the amount of food patients can take in at meals. This pouch is then attached directly to the small intestine, bypassing most of the rest of the stomach and the upper part of the small intestine, and reducing the amount of fat and calories that are absorbed from the foods [30]. A recent review demonstrated that RYGB is associated with diminished absorption of lypophilic drugs such as cyclosporine, phenytoin, rifampin, and thyroxine [7].

BPD is a bariatric surgery for patients with severe obesity. The primary mechanism of weight loss with the BPD is malabsorption. BPD removes approximately three fourths of the stomach to produce both restriction of food intake and reduction of acid output. Leaving enough upper stomach is important to maintain proper nutrition. The small intestine is then divided with one end attached to the stomach pouch to create what is called an "alimentary limb." All the food moves through this segment, however, not much is absorbed. The bile and pancreatic juices move through the "biliopancreatic limb," which is connected to the side of the intestine close to the end. This supplies digestive juice in the section of the intestine now called the "common limb." The surgeon varies the length of the common limb to regulate the amount of absorption of protein, fat, and fat-soluble vitamins [31, 32]. A recent review demonstrated that also BPD is associated with diminished absorption of lypophilic drugs and thyroxine [31].

The mechanisms by which liquid L-T4 circumvent malabsorption in bariatric surgery remain to be studied. It has been suggested that absorption of thyroxine is greater with oral liquid formulations in patients after bariatric surgery [26]. In fact, normal gastric acid secretion is necessary for effective absorption of L-T4 [18] by dissolution of tablets, and drug dissolution and solubility may be altered by restrictive procedures that increase gastric pH in the newly created stomach pouch; this may occur in gastric bypass [7]. Since it has been shown that the liquid formulation of L-T4 is extremely effective to circumvent the problem of incomplete absorption of L-T4 caused by induced proton pump inhibitor, [33] this formulation could also circumvent the pH alteration resulting from gastric bypass [34].

Furthermore, the presence of alcohol in the L-T4 liquid formulation could also play a key role in thyroxine absorption. Indeed, oral mucosal is highly vascularized, and drugs that are absorbed through the oral mucosal directly enter the systemic circulation, bypassing the gastrointestinal tract [35]. Further studies are needed to clarify these intriguing points.

The comparison of TSH values before surgery, after surgery, and 2–3 months after the switch show that the TSH values after the switch are higher (near to the statistical significance, P = 0.064), with respect to the TSH values before surgery. These results suggest that in patients with bariatric surgery a slight malabsorption persist even when the liquid L-T4 formulation is given. However, there was no significant difference in the decrease of TSH in RYGB group vs the BPD group, nor in females vs males.

In conclusion, the abovementioned data suggest that the L-T4 oral liquid formulation could circumvent L-T4 malabsorption both in patients submitted to BPD, as in those submitted to RYGB; further studies are needed to enlarge the number of participants and to clarify the implicated mechanisms.

Compliance with Ethical Standards

Conflict of Interest The authors declare that they have no conflict of interest.

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

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

References

- Buchwald H, Oien DM. Metabolic/bariatric surgery worldwide 2008. Obes Surg. 2009;19(12):1605–11. PMID: 19885707.
- Buchwald H, Oien DM. Metabolic/bariatric surgery worldwide 2011. Obes Surg. 2013;23(4):427–36. PMID: 23338049.
- Frühbeck G. Bariatric and metabolic surgery: a shift in eligibility and success criteria. Nat Rev Endocrinol. 2015;11(8):465–77. PMID: 26055046.
- Bray GA, Frühbeck G, Ryan DH, Wilding JP. Management of obesity. Lancet. 2016. PMID: 26868660.
- Fujioka K. Follow-up of nutritional and metabolic problems after bariatric surgery. Diabetes Care. 2005;28(2):481–4. PMID: 15677821.

- Miller AD, Smith KM. Medication and nutrient administration considerations after bariatric surgery. Am J Health Syst Pharm. 2006;63(19):1852–7. PMID: 16990631.
- Padwal R, Brocks D, Sharma AM. A systematic review of drug absorption following bariatric surgery and its theoretical implications. Obes Rev. 2010;11(1):41–50. PMID: 19493300.
- Michalaki MA, Vagenakis AG, Leonardou AS, et al. Thyroid function in humans with morbid obesity. Thyroid. 2006;16(1):73–8. PMID: 16487017.
- Moulin de Moraes CM, Mancini MC, de Melo ME, et al. Prevalence of subclinical hypothyroidism in a morbidly obese population and improvement after weight loss induced by Roux-en-Y gastric bypass. Obes Surg. 2005;15(9):1287–91. PMID: 16259889.
- Galofré JC, Pujante P, Abreu C, et al. Relationship between thyroidstimulating hormone and insulin in euthyroid obese men. Ann Nutr Metab. 2008;53(3–4):188–94. PMID: 19011282.
- Pacini F, Castagna MG. Approach to and treatment of differentiated thyroid carcinoma. Med Clin North Am. 2012;96(2):369–83. PMID: 22443981.
- Almandoz JP, Gharib H. Hypothyroidism: etiology, diagnosis, and management. Med Clin North Am. 2012;96(2):203–21. PMID: 22443971.
- Antonelli A, Ferrari SM, Corrado A, et al. Autoimmune thyroid disorders. Autoimmun Rev. 2015;14(2):174–80. PMID: 25461470.
- Fish LH, Schwartz HL, Cavanaugh J, et al. Replacement dose, metabolism, and bioavailability of levothyroxine in the treatment of hypothyroidism. Role of triiodothyronine in pituitary feedback in humans. N Engl J Med. 1987;316(13):764–70. PMID: 3821822.
- Eledrisi MS, Szymajda A, Alshanti M, et al. Noncompliance with medical treatment: pseudomalabsorption of levothyroxine. South Med J. 2001;94(8):833–6. PMID: 11549198.
- Pabla D, Akhlaghi F, Zia H. A comparative pH-dissolution profile study of selected commercial levothyroxine products using inductively coupled plasma mass spectrometry. Eur J Pharm Biopharm. 2009;72(1):105–10. PMID: 18996189, Epub 2008 Nov 1.
- Sherman SI, Malecha SE. Absorption and malabsorption of levothyroxine sodium. Am J Ther. 1995;2(10):814–8. PMID: 11854792.
- Centanni M, Gargano L, Canettieri G, et al. Thyroxine in goiter, Helicobacter pylori infection, and chronic gastritis. N Engl J Med. 2006;354(17):1787–95. PMID: 16641395.
- Mandel SJ, Brent GA, Larsen PR. Levothyroxine therapy in patients with thyroid disease. Ann Intern Med. 1993;119(6):492– 502. PMID: 8357116.
- Liwanpo L, Hershman JM. Conditions and drugs interfering with thyroxine absorption. Best Pract Res Clin Endocrinol Metab. 2009;23(6):781–92. PMID: 19942153.
- Schubert ML, Peura DA. Control of gastric acid secretion in health and disease. Gastroenterology. 2008;134(7):1842–60. PMID: 18474247.

- Checchi S, Montanaro A, Pasqui L, et al. L-thyroxine requirement in patients with autoimmune hypothyroidism and parietal cell antibodies. J Clin Endocrinol Metab. 2008;93(2):465–9. PMID: 18042648.
- Sachmechi I, Reich DM, Aninyei M, et al. Effect of proton pump inhibitors on serum thyroid-stimulating hormone level in euthyroid patients treated with levothyroxine for hypothyroidism. Endocr Pract. 2007;13(4):345–9. PMID: 17669709.
- Azizi F, Belur R, Albano J. Malabsorption of thyroid hormones after jejunoileal bypass for obesity. Ann Intern Med. 1979;90(6): 941–2. PMID: 443690.
- Bevan JS, Munro JF. Thyroxine malabsorption following intestinal bypass surgery. Int J Obes. 1986;10(3):245–6. PMID: 3759332.
- Pirola I, Formenti AM, Gandossi E, et al. Oral liquid L-thyroxine (L-t4) may be better absorbed compared to L-T4 tablets following bariatric surgery. Obes Surg. 2013;23(9):1493–6. PMID: 23824980.
- Camastra S, Gastaldelli A, Mari A, et al. Early and longer term effects of gastric bypass surgery on tissue-specific insulin sensitivity and beta cell function in morbidly obese patients with and without type 2 diabetes. Diabetologia. 2011;54(8):2093–102. PMID: 21614570.
- Scopinaro N, Gianetta E, Civalleri D, et al. Bilio-pancreatic bypass for obesity: II. Initial experience in man. Br J Surg. 1979;66(9): 618–20. PMID: 497645.
- 29. Lee S, Farwell AP. Euthyroid sick syndrome. Compr Physiol. 2016;6(2):1071–80. PMID: 27065175.
- Carswell KA, Belgaumkar AP, Amiel SA, Patel AG. A systematic review and meta-analysis of the effect of gastric bypass surgery on plasma lipid levels. Obes Surg. 2015. PMID: 26210195
- Padwal RS, Lewanczuk RZ. Trends in bariatric surgery in Canada, 1993–2003. CMAJ. 2005;172(6):735. PMID: 15767602.
- 32. Pontiroli AE, Laneri M, Veronelli A, et al. Biliary pancreatic diversion and laparoscopic adjustable gastric banding in morbid obesity: their long-term effects on metabolic syndrome and on cardiovascular parameters. Cardiovasc Diabetol. 2009;8:37. PMID: 19619292.
- 33. Saraceno G, Vita R, Trimarchi F, et al. A liquid formulation of Lthyroxine (L-T4) solves problems of incomplete normalization/ suppression of serum TSH caused by proton pump inhibitors (PPI) on conventional tablet formulations of L-T4. Presented at European Society of Endocrinology ICE/ECE 2012 May Florence, Italy. Endocr Abstr. 2012;29:P1626.
- Vita R, Fallahi P, Antonelli A, et al. The administration of Lthyroxine as soft gel capsule or liquid solution. Expert Opin Drug Deliv. 2014;11(7):1103–11. PMID: 24896369.
- Zhang H, Zhang J, Streisand JB. Oral mucosal drug delivery: clinical pharmacokinetics and therapeutic applications. Clin Pharmacokinet. 2002;41(9):661–80. PMID: 12126458.