Adjuvant treatment with thyrotropin alpha for remnant ablation in thyroid cancer

Abstract: Various studies have demonstrated the safety and efficacy of recombinant human thyroid-stimulating hormone (rhTSH) for radioiodine remnant ablation. On this basis, rhTSH was approved in Europe for the radioiodine ablation of low-risk differentiated thyroid cancer (DTC) during thyroid hormone therapy with L-thyroxine (L-T4). Moreover, in December 2007, the US Federal Drug Administration approved the use of rhTSH for adjuvant treatment with radioiodine in patients with DTC without evidence of metastatic thyroid cancer. Quality of life was found to be better with rhTSH preparation than with L-thyroxine withdrawal, thereby resulting in benefits for society as a whole. Furthermore, rhTSH for radioiodine remnant ablation results in a longer effective radioiodine half-life within remnant thyroid tissue and a lower specific absorbed dose in the blood and exposure of bone marrow to X-rays. More studies are required to establish the amount of radioiodine to be administered especially in high-risk patients.

Keywords: thyroid cancer, thyrotropin, radioiodine (131I) remnant ablation (RRA), quality of life, ray exposure

There is general agreement that total thyroidectomy is the initial treatment-of-choice for patients with differentiated thyroid cancer (DTC). Radioiodine (131I) remnant ablation is recommended after thyroidectomy to destroy post-surgical residual thyroid tissue especially in patients at high-risk of recurrence and mortality. 131I ablation has two advantages: 1) it destroys any remaining microscopic tumoral foci; and 2) it eliminates all normal thyroid cells that would continue to produce thyroglobulin and confound interpretation of measurement of serum thyroglobulin (Tg), which is a specific marker of recurrent or persistent disease. Consequently, this procedure improves the follow-up and treatment of patients with DTC by increasing the specificity and sensitivity of Tg monitoring and 131I treatment. Moreover, 131I administration decreases the frequency of recurrences and mortality.

Elevated serum thyroid-stimulating hormone (TSH) levels (above 30 mU/L) are necessary to ensure sufficient trapping and retention of 131I by functioning thyroid tissue. Traditionally, the endogenous increase of TSH was achieved by withdrawal of thyroid hormone therapy (L-thyroxine; LT4) for 4 to 5 weeks, which induces clinical hypothyroidism. However, this short-term hypothyroid condition is associated with cognitive and physical impairment and alteration of quality of life in young and middle-aged patients. Elevated serum TSH levels (above 30 mU/L) are necessary to ensure sufficient trapping and retention of 131I by functioning thyroid tissue. Traditionally, the endogenous increase of TSH was achieved by withdrawal of thyroid hormone therapy (L-thyroxine; LT4) for 4 to 5 weeks, which induces clinical hypothyroidism. However, this short-term hypothyroid condition is associated with cognitive and physical impairment and alteration of quality of life in young and middle-aged patients. Moreover, withdrawal of LT4 can impair cardiac, cognitive and neurological function with consequent health risks especially for elderly people.

Lastly, it may not increase TSH levels in cases of persistent thyroid hormone production by large thyroid remnants or functional metastases, in elderly patients, and in the presence of hypothalamic or pituitary disease or long-term steroid therapy.

Recombinant human TSH (rhTSH) is a heterodimeric glycoprotein produced by recombinant DNA technology for the purpose of producing increased TSH levels without...
LT4 withdrawal and the consequent hypothyroidism. The use of rhTSH was initially limited to the field of DTC follow-up and was approved by the US Food and Drug Administration (FDA) in December 1998 for diagnostic use. Subsequently, rhTSH was found to be effective for ^131I remnant ablation,^13,14 and in February 2005, rhTSH was approved in Europe for the ^131I ablation of low-risk DTC during thyroid hormone therapy with LT4. In December 2007 the FDA approved the use of rhTSH for adjuvant treatment with ^131I in patients with DTC without evidence of metastatic thyroid cancer.

Here we review the studies on rhTSH-aided ablation with the aim of addressing such open questions as the exact protocol of rhTSH administration and the dose of ^131I to obtain maximum effectiveness.

**RhTSH-aided ablation: literature analysis**

Table 1 lists the studies that evaluated the effectiveness of rhTSH in the adjuvant treatment for ^131I remnant ablation in DTC patients. The criteria used to define successful thyroid ablation differed among studies from no visible uptake at whole body scan after rhTSH or undetectable basal and rhTSH-stimulated serum thyroglobulin. Despite these differences, there is general agreement that rhTSH for thyroid ablation gives results similar to those found after LT4 withdrawal.

The study by Perros et al was the first report on the use of rhTSH to increase ^131I uptake for remnant ablation.15 Subsequently, the effect of rhTSH-aided ablation was evaluated in a prospective non-randomized trial of 10 patients with papillary cancer.16 The dose of ^131I administrated varied between 30 and 250 mCi. The ablation rate was 100% when judged by the absence of visible uptake in the thyroid bed after diagnostic whole body scan 3 months after ablation.16 Another randomized study confirmed complete ablation after high doses of ^131I (approximately 108 mCi) by using TSH to stimulate ^131I uptake for the ablation of remnant thyroid tissue.17

A subsequent retrospective study from the Memorial Sloan-Kettering Cancer Center confirmed that a high dose of ^131I increased the rate of rhTSH-aided ablation in DTC patients.13 In this study, the rates of complete ablation did not differ significantly between a group of patients who were prepared by thyroid hormone withdrawal (THW) and a group of patients prepared by rhTSH when treated with 100 mCi (84% in 45 euthyroid patients after rhTSH vs 81% in 42 hypothyroid patients).13

However, a prospective study by Pacini et al did not confirm these results.18 In this prospective randomized study in which 1.1 GBq (30 mCi) was used as standard ablative activity, 162 DTC patients were randomized in three treatment arms: in the first arm, patients (n = 50) were treated by LT4 withdrawal (HYPO); in the second arm, patients

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**Table 1** Studies evaluating the efficacy of rhTSH for remnant ablation

| Authors               | Patients (n) | Stage of disease | Dose of ^131I (mCi) | Outcome                      |
|-----------------------|--------------|------------------|---------------------|------------------------------|
|                       | rhTSH | LT4W | rhTSH | LT4W | rhTSH | LT4W | rhTSH | LT4W |
| Robbins et al 2001    | 10    | n.p. | T1–T4 | n.p. | 30–250 | n.p. | 100% dWBS negative, 60% Tg < 1.0 |
| Prospective randomized study |
| Pacini et al 2002    | 70    | 50   | T1–T4 | T1–T4 | 30    | 30   | 54% dWBS negative, 86.8% Tg < 1.0 |
| Prospective randomized study |
| Robbins et al 2002   | 42 rhTSH + LT4WI | T1–T4 | T1–T4 | N0–N1 | N0–N1 | 30 | 78.5% dWBS negative, 84.8% Tg < 1.0 |
| Barbaro et al 2003   | 45 rhTSH | T1–T4 | T1–T4 | N0–N1 | N0–N1 | 110 ± 65, 128.9 ± 74 | 81% dWBS negative, 84% dWBS negative |
| Non-randomized randomized study |
| Pacini et al 2006    | 33    | 30   | T1–T4 | T1–T4 | 100   | 100  | 77% dWBS negative, 86.5% Tg < 1.0 |
| Prospective randomized study |
| Pilli et al 2007     | 36    | n.p. | T1–T4 | T1–T4 | 50    | n.p. | 88.9% dWBS negative, 78.9% Tg < 1.0 |
| Prospective randomized study |

n.p. = not performed.

**Abbreviations:** rhTSH, recombinant human thyroid-stimulating hormone; dWBS, diagnostic whole body scanner; LT4W, levo-thyroxine withdrawal; Tg, serum thyroglobulin.
(n = 42) were treated by LT4 withdrawal combined with rhTSH (HYPO + rhTSH); in the third arm, patients (n = 70) were stimulated with rhTSH in euthyroidism (EU + rhTSH). The follow-up was performed 6 to 10 months post ablation. When the criterion for successful ablation was no uptake on the thyroid bed on diagnostic whole body scan, the rate of successful ablation was similar in the HYPO and HYPO + rhTSH groups (84% and 78.5%, respectively) but significantly lower (54% p < 0.01) in the EU + rhTSH group.18 On the contrary when successful ablation was defined as no visible thyroid bed uptake on diagnostic whole body scan or undetectable serum Tg after rhTSH, the success rates were similar (95% vs 74%). However, the reduced rate of ablation in the EU group may be explained by the protocol of 131I administration used by Pacini et al.18 Indeed, ablative 131I administration was delayed by 24 h and it was delivered 48 h after the second injection of rhTSH. Therefore, the authors suggested that the dose of 131I be increased or that different protocols of rhTSH administration be used to obtain a satisfactory rate of rhTSH-aided thyroid ablation.

An international randomized controlled trial showed that the efficacy of rhTSH for ablation was similar to that of LT4 withdrawal with 100% ablation after 3.7 GBq (100 mCi).14 The predefined primary criterion for successful ablation was “no visible uptake in the thyroid bed, or a visible uptake less than 0.1%” on neck scans performed 8 months after therapy, and was satisfied in 100% of patients in both groups. A secondary criterion for ablation, a rhTSH-stimulated serum thyroglobulin concentration less than 2 ng/mL, was fulfilled by 23 of 24 (96%) euthyroid rhTSH patients and 18 of 21 (86%) hypothyroid patients (p = 0.2341). In this randomized prospective ablation trial, all rhTSH patients had an iodine excretion below 200 µL, indicating the absence of overt iodine excess.

Only two studies have evaluated the efficacy of rhTSH for remnant ablation with lower 131I doses.19,20 A recent study by Pilli et al showed that 1850 MBq (50 mCi) 131I had a similar success rate to 3700 MBq (100 mCi) in 72 patients prepared with rhTSH for thyroid ablation.19 This prospective, randomized study showed that 3700 MBq 131I is associated with high rates of successful thyroid ablation after rhTSH preparation and that similar ablation rates (88.9%) were obtained with lower 131I activity (1850 MBq). These results were obtained when the criterion of successful ablation was defined as no visible uptake at the 6- to 8-month control diagnostic 131I whole body scan after rhTSH stimulation, and also when the criterion of successful ablation was undetectable (1 ng/mL) rhTSH-stimulated serum Tg. Furthermore, successful ablation was also obtained in patients with nodal metastases. Lastly, the dosimetric study showed that thyroid uptake was similar in patients treated with 1850 or 3700 MBq.

Since thyroid hormones are an important source of iodine and may interfere with 131I uptake during thyroid ablation, Barbaro et al suggested LT4 therapy be discontinued before rhTSH injection.20 They compared ablation obtained with doses of 30 mCi in 2 groups of DTC patients: one group was prepared by hypothyroidism and the other group was prepared by rhTSH stimulation. In the rhTSH group, LT4 therapy was interrupted for 4 days starting the day before the first injection. In the rhTSH group, urinary iodine excretion was significantly lower than in a control group of euthyroid subjects who received rhTSH stimulation. One year later, patients underwent a whole body scan with a tracer dose of 131I and serum Tg was measured using rhTSH with the same protocol in both groups. The percentage of ablation (undetectable Tg and a negative whole body scan) was 81.2% in patients treated with rhTSH and 76% in patients treated by L-T4 withdrawal.

Similarly, Pitoia et al suggested replacing LT4 with LT3 therapy to maintain the euthyroid state and to minimize the iodine pool during rhTSH preparation.21 Indeed, LT3 has an iodine content 5-fold less than LT4.21

RhTSH-aided ablation: 131I dosimetry, safety and cost

Because 131I activity is associated with such important risks as bone marrow depression and pulmonary fibrosis, several dosimetric studies have been performed to evaluate the absorbed dose in the blood (a surrogate for bone marrow) and 131I activity in the lung to determine the minimum effective dose to reduce the these risks. It has been reported that a dose of 2 Gy of radiation in the blood is dose-limiting,22 whereas 3 GBq in the lung in 24 h is the safety limit to avoid pulmonary fibrosis.23

An international, prospective, randomized study compared the iodine biokinetics, dosimetry and the effectiveness of ablation therapy with 100 mCi in DTC after rhTSH stimulation or LT4 withdrawal.24 Iodine biokinetics differed between the two groups of patients.24 In fact, in the euthyroid state, renal clearance of iodine was 50% faster than in hypothyroidism.21 Indeed, fractional 131I uptake into thyroid remnants was lower after rhTSH stimulation than after LT4 withdrawal.24 However, this reduction was partially compensated for by an increased half-life of 131I in thyroid cells after rhTSH stimulation.24 rhTSH-treated patients showed a longer effective 131I half-life within remnant thyroid tissue, and the residence times of the radioisotope were comparable in the two groups.24 Moreover, the specific absorbed dose in the blood was significantly lower (one-third) after rhTSH preparation, suggesting that
higher $^{131}$I activities might be safely administered after rhTSH stimulation. Finally, another study confirmed that the bone marrow absorbed dose remained under 2 Gy after rhTSH-aided administration of high activities of $^{131}$I. Moreover, patients prepared with rhTSH had a better quality of life than hypothyroid patients. RhTSH-aided ablation was well tolerated with no important side effects, and it can be useful in elderly patients and in patients with associated comorbidities without increasing the risk of cardiac, cerebrovascular, pulmonary or neurological complications.

Finally, a recent study compared the cost-effectiveness of ablation after rhTSH stimulation or LT4 withdrawal. The additional cost of rhTSH procurement and administration was considered justified in relation to the clinical benefits and cost offsets such as avoidance of hypothyroidism, increased work productivity and quality life, reduced discharge from radioprotection and period of sick leave. These observations were recently confirmed by Borget et al who found that rhTSH can decrease the duration of sick leave, and that its high cost is compensated for by benefits to patients and society with a modest net cost.

**RhTSH-aided ablation: advantages and limits**

There is general agreement that rhTSH-aided ablation is effective and safe. Various studies have confirmed the efficacy of rhTSH in aiding ablation and show that rhTSH preparation is more beneficial than LT4 withdrawal in terms of quality of life and well-being and avoids the important side effects of short-term hypothyroidism. Moreover, rhTSH for remnant ablation decreases exposure of bone marrow to X-rays.

Several questions are still open, namely, the amount of $^{131}$I to be administered and the effect of iodine intake. More studies are required to evaluate whether rhTSH can be used effectively for remnant ablation in high risk patients with outcomes at least comparable to those seen with ablation after thyroxine withdrawal.

**Disclosures**

The authors have no conflicts of interest to disclose.

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