

**Commentary**

# Towards larger volume reduction of nodular goitres by radioiodine therapy: a role for pretreatment with recombinant human thyrotropin?

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In the last two decades it has been demonstrated that radioiodine ( $^{131}\text{I}$ ) is an effective therapy for thyroid volume reduction in patients with toxic and nontoxic nodular goitre. In these patients,  $^{131}\text{I}$  treatment leads to a significant decrease in goitre size. In most patients compressive symptoms improve as well. The decrease in compressive symptoms is accompanied by significant tracheal widening and improvement of respiratory function (for review see Hegedüs *et al.*, 2003).

$^{131}\text{I}$  treatment is especially attractive in elderly patients who have a high operative risk and in those who refuse surgery. However, in patients with nontoxic, nodular goitre thyroid radioactive iodide uptake (RAIU) is usually rather low, especially in areas with a high iodine intake. As a result, high doses of  $^{131}\text{I}$  are often needed for thyroid volume reduction, causing a relatively high radiation burden to extrathyroidal organs (Huysmans *et al.*, 1996). Therefore, it is of interest to explore strategies to enhance RAIU in these patients. In the last decade recombinant human thyrotropin (rhTSH) has become available for diagnostic use in patients with differentiated thyroid cancer. It has been shown that rhTSH stimulates RAIU in thyroid remnants and thyroid cancer tissue (Ladenson *et al.*, 1997; Haugen *et al.*, 1999). Recently, we reported that rhTSH stimulates RAIU also in patients with nodular goitre: the administration of a single, low dose of 0.01 or 0.03 mg rhTSH doubled 24-h RAIU in these patients (Huysmans *et al.*, 2000). Pretreatment with rhTSH also caused a more homogeneous distribution of radioiodine on the thyroid scintigrams of nodular goitres by stimulating radioiodide uptake in relatively cold areas more than in relatively hot areas, especially in patients with a low baseline serum TSH level (Nieuwlaaf *et al.*, 2001). These observations suggest that administration of rhTSH before  $^{131}\text{I}$  therapy for volume reduction of nodular goitre may allow treatment with lower doses of  $^{131}\text{I}$  without diminishing the radiation-absorbed dose in the thyroid and the efficacy of this mode of therapy. Indeed, pretreatment with a single, low dose of 0.01 or

0.03 mg rhTSH allowed approximately 50–60% reduction of the therapeutic dose of radioiodine without compromising the efficacy of thyroid volume reduction (Nieuwlaaf *et al.*, 2003). As the radiation burden of radioiodine therapy to extrathyroidal organs is directly related to the administered  $^{131}\text{I}$  dose (Huysmans *et al.*, 1996), such a dose reduction may render radioiodine therapy more attractive for younger patients and may allow for more patients to be treated on an out-patient basis.

A major drawback of radioiodine therapy for nodular goitre (with  $^{131}\text{I}$  doses aimed at approximately 100–150  $\mu\text{Ci}$  retained per gram of thyroid tissue at 24 h) is that mean thyroid volume reduction is not greater than approximately 40% after 1 year, and 50–60% after 3–5 years (for review see Hegedüs *et al.*, 2003). Moreover, not all patients respond. Preliminary data by Le Moli *et al.* (1999) suggest that goitre reduction can be augmented by increasing the radiation-absorbed dose in the thyroid. Such an increase can be achieved by simply enlarging the administered dose of radioiodine, but this will further increase the radiation burden to extrathyroidal organs. Alternatively, it should be possible to increase the radiation-absorbed dose in the thyroid without increasing the administered dose of radioiodine, by stimulating RAIU using pretreatment with rhTSH.

In this issue of *Clinical Endocrinology*, Silva *et al.* (2004) report a study in which 34 patients with large, nodular goitres (22 were subclinically hyperthyroid and seven overtly hyperthyroid) were randomized to radioiodine therapy alone or to radioiodine therapy in comparable doses but with pretreatment with a relatively high dose of rhTSH (0.45 mg), given 24 h before radioiodine administration. Patients pretreated with rhTSH had a significantly larger thyroid volume reduction after 1 year ( $58 \pm 13\%$ ; mean  $\pm$  SD) than patients given radioiodine without rhTSH pretreatment ( $40 \pm 12\%$ ).

From data provided in Tables 1 and 2 of their report, we calculated the dose retained in the thyroid at 24 h for individual patients in both groups. This parameter was considerably higher in the rhTSH-pretreated patients (191  $\pm$  75  $\mu\text{Ci/g}$  of thyroid tissue) than in the patients not pretreated with rhTSH (73  $\pm$  22  $\mu\text{Ci/g}$ ). It seems likely that the higher thyroid volume reduction in the rhTSH-pretreated group can be explained by the higher retention of radioiodine in the thyroid. An alternative explanation is that pretreatment with rhTSH improves thyroid volume reduction by causing a more homogeneous distribution of radioiodine within the thyroid, especially increasing the uptake of radioiodine in relatively cold areas (Nieuwlaaf *et al.*, 2001).

It may be anticipated that a higher dose of radioiodine retained in the thyroid will be accompanied by more severe early adverse effects due to radiation-induced thyroiditis and oesophagitis. Indeed, in comparison with patients not pretreated with rhTSH, rhTSH-pretreated patients had a higher incidence of pain in the thyroid region (52% vs. 23%), complaints due to oesophagitis (17% vs. 11%) and weight loss (65% vs. 52%).

Another consequence of thyroiditis is acute enlargement of the thyroid gland with (further) compression of the trachea. For radioiodine doses aimed at approximately 100–150 µCi retained per gram of thyroid tissue at 24 h, increases in thyroid volume 1 week after radioiodine therapy up to 25% have been found (Nygaard *et al.*, 1994; Bonnema *et al.*, 1999; Nieuwlaat *et al.*, 2003). Silva *et al.* (2004) did not quantify thyroid volume changes in the first period after radioiodine therapy. Therefore, it remains to be investigated whether the larger <sup>131</sup>I doses retained in the thyroid as used by Silva *et al.* (2003) cause larger increases in thyroid volume in the first period after radioiodine treatment.

Mild increases in serum thyroid hormone levels due to radiation-induced thyroiditis are commonly seen in the first weeks after radioiodine treatment of nodular goitre, with maximum levels reached at approximately 2 weeks after therapy (Nygaard *et al.*, 1994). It has also been demonstrated that administration of low doses of rhTSH (0.01 or 0.03 mg) in patients with nodular goitre results in mild increases in serum thyroid hormone levels, with maximum levels reached at 1–4 days after administration of rhTSH (Huysmans *et al.*, 2000). It may be anticipated that pretreatment with a relatively high dose of rhTSH followed by a full dose of radioiodine, leading to a high radiation-absorbed dose in the thyroid, will result in larger increases in serum thyroid hormone levels. Indeed, peak FT4 levels reached in the rhTSH-pretreated patients studied by Silva *et al.* (2004) were much higher than those in the patients treated with radioiodine alone (59 ± 22 vs. 24 ± 7 pmol/l), despite a low-iodine diet in all patients and methimazole pretreatment in the seven overtly hyperthyroid patients. The highest FT4 levels were reached already at 1–3 days after radioiodine therapy, suggesting that rhTSH administration was the most important cause of the rise. Fortunately, no hyperthyroid symptoms or worsening of heart diseases were observed. The authors give three possible explanations: the rise in thyroid hormone levels was of short duration, most patients had cardiac medications, and all patients were confined to their beds for 5–9 days after radioiodine therapy.

A late adverse effect of radioiodine treatment of nodular goitre is development of hypothyroidism. Probably related to the higher radioiodine dose retained in the thyroid, Silva *et al.* (2004) found that hypothyroidism 1 year after radioiodine therapy was more frequent in the rhTSH-pretreated patients than in the patients not pretreated with rhTSH (65% vs. 21%).

The study of Silva *et al.* (2004) is the first to show that the efficacy of radioiodine therapy of nodular goitres can be

improved by pretreatment with rhTSH. However, conclusions are based on small numbers of patients and both toxic (four of them had used amiodarone) and nontoxic patients were included. Moreover, no attempt was made to calculate precisely the radioiodine doses to be administered (radioiodine doses were based on a rather simple algorithm with estimated thyroid volume as the only parameter).

Before rhTSH can be advised as an adjunct to improve the efficacy of radioiodine therapy in nodular goitre, further studies are needed. First, it has to be determined in a formal dose–response study which dose of rhTSH is optimal for this purpose. Such a dose should stimulate RAIU considerably, but should not cause unacceptable rises in serum thyroid hormone levels. Then, radioiodine therapy with and without pretreatment with that particular dose of rhTSH should be investigated in randomized studies. Such studies should look carefully at dose–response relationships with respect to efficacy and adverse effects. Given the heterogeneity of nodular goitres, large groups of patients will be needed.

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