Importance of gender-specific calcitonin thresholds in screening for occult sporadic medullary thyroid cancer

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Abstract

Men and women differ in thyroidal C-cell mass and calcitonin secretion. This difference may have implications for the definition of calcitonin thresholds to distinguish sporadic C-cell hyperplasia from occult medullary thyroid cancer. This retrospective study examined the hypothesis that gender-specific calcitonin thresholds predict occult medullary thyroid cancer more accurately among patients with increased basal calcitonin levels than unisex thresholds. A total of 100 consecutive patients were evaluated with occult sporadic C-cell disease no larger than 10 mm who were referred for increased basal calcitonin levels and underwent pentagastrin stimulation preoperatively at this institution. Altogether, gender-specific calcitonin thresholds predicted medullary thyroid cancer better than unisex thresholds. At lower (< 50 pg/ml basally; < 500 pg/ml after stimulation), but not higher, calcitonin serum levels, women revealed medullary thyroid cancer four to eight times more often than men. Most discriminatory between C-cell hyperplasia and medullary thyroid cancer was a basal calcitonin threshold of 15 pg/ml (corrected 20 pg/ml) for women and 80 pg/ml (corrected 100 pg/ml) for men, based on the greatest accuracy at the lowest possible calcitonin level. The respective gender-specific stimulated peak calcitonin thresholds were 80 pg/ml (corrected 100 pg/ml) and 500 pg/ml. Corresponding positive predictive values for medullary thyroid cancer at these calcitonin thresholds were 89 and 90% for women, as opposed to 100% for men. To increase the positive predictive value for women to 100%, the respective calcitonin thresholds would have to be raised to 40 pg/ml (corrected 50 pg/ml) and 250 pg/ml. These findings indicate that gender-specific calcitonin thresholds predict sporadic occult medullary thyroid cancer better than unisex thresholds.

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Introduction

Originating from parafollicular C-cells, medullary thyroid cancer cells retain the ability of synthesizing, storing, and releasing calcitonin into the bloodstream. Because they broadly reflect overall C-cell mass (Iacobone et al. 2002, Machens et al. 2005), serum calcitonin levels have prognostic ramifications in medullary thyroid cancer patients regarding a) the extent of locoregional disease and neck dissection preoperatively, and b) locoregional recurrence and cancer-specific survival postoperatively (Machens et al. 2007, Oskam et al. 2008).

With the propagation of calcitonin screening, the prevalence of medullary thyroid cancer has surged from 1.1 to 3.2 cases per 1000 patients with nodular thyroid disease (Vierhapper et al. 2005). This increase was mainly due to the increasing detection of occult medullary thyroid cancer no larger than 10 mm. In the presence of multinodular goiter, these occult cancers are barely amenable to conventional diagnostic strategies, such as ultrasonographic visualization and fine-needle aspiration. Because occult medullary thyroid cancers are usually still confined to the thyroid gland without having spread to lymph nodes, total thyroidectomy usually will be curative at this early
stage (Elisei et al. 2004). In contrast, surgical cure is exceptional with larger tumors when 10 or more lymph nodes are involved (Machens et al. 2000a, Scollo et al. 2003).

As in any other screening program, the cost-effectiveness of calcitonin screening for medullary thyroid cancer hinges on a) the prevalence of the target disease (3.2 cases per 1000 patients with thyroid nodular disease), b) the predictive value of the screening test (which depends on the calcitonin threshold selected), and c) the availability of effective treatment strategies (e.g. thyroidectomy that will cure all cancers limited to the thyroid gland). For these reasons, calcitonin screening holds great promise as a cost-effective screening tool.

The most common alternative source of calcitonin secretion, other than medullary thyroid cancer, is C-cell hyperplasia, which also arises from the parafollicular C-cells of the thyroid gland. It comes in a reactive (sporadic) and a neoplastic (hereditary) variety. Unlike its reactive counterpart, hereditary neoplastic C-cell hyperplasia represents a preneoplastic condition for carriers of rearranged during transfection (RET) germline mutations, giving rise to hereditary medullary thyroid cancer in an age-dependent fashion (Machens et al. 2003). Because increased calcitonin serum levels can originate from either condition, it is often challenging, if not impossible, to distinguish C-cell hyperplasia from medullary thyroid cancer, especially at lower levels.

Strictly speaking, both C-cell hyperplasia patients and medullary thyroid cancer patients, revealing elevated calcitonin serum levels, yield ‘positive’ calcitonin test results. Yet, the ramifications of a positive calcitonin test differ greatly depending on the underlying condition: from expectant observation, or total thyroidectomy without any form of lymph node dissection, for sporadic C-cell hyperplasia to immediate total thyroidectomy, usually coupled with systematic lymph node dissection of at least the central neck compartment, for medullary thyroid cancer. To make this important distinction, biochemical screening programs for medullary thyroid cancer commonly have interpreted increased calcitonin serum levels as true positive in the presence of medullary thyroid cancer, and as false positive in the absence thereof. Indeed, such false-positive test results, mostly in connection with lower calcitonin levels, have tarnished the performance of calcitonin screening, triggering unnecessary thyroid operations for suspected but non-existent medullary thyroid cancer. For instance, the failure to define gender-specific calcitonin thresholds may have diminished the assay’s performance by covering up gender-specific differences in thyroid C-cell mass and calcitonin secretion (Heath & Sizemore 1977).

False-negative findings may have resulted from the inclusion of patients with medullary thyroid cancer >10 mm. This inclusion may have raised calcitonin thresholds so much that some patients with smaller medullary thyroid cancer became misclassified as tumor free. These gross tumors are often detectable using conventional methods, rendering the case for biochemical detection less compelling. Moreover, the unwillingness of some patients with increased calcitonin levels to undergo thyroidectomy may have caused underestimation of medullary thyroid cancer prevalence (Heath & Sizemore 1977, Vierhapper et al. 2005).

The current investigation aimed to examine the hypothesis that gender-specific calcitonin thresholds predict occult medullary thyroid cancer better than unisex thresholds in patients with sporadic C-cell disease.

Patients and methods

Inclusion and exclusion criteria

To be considered for this retrospective study, patients needed to have undergone total thyroidectomy with or without systematic lymph node dissection at this institution, in addition to meeting the following criteria:

(a) an increased basal calcitonin level stimulated preoperatively by i.v. bolus injection of 0.5 μg pentagastrin (Peptavlon, Laboratoires SERB, Paris, France) per kilogram body weight, measured at this institution between July 2004 and December 2008 with the Immulite 2000 calcitonin assay (Diagnostic Products Corporation, Los Angeles, CA, USA; normal range <5 pg/ml for women and <8.4 pg/ml for men; or <5 and <8.4 ng/l respectively in Système International Units). The Immulite 2000 automated calcitonin assay, instituted in July 2004 to enable determination of calcitonin levels immediately before surgery, is linearly related to other calcitonin assays, including the Nichols Advantage assay, but with a lower calibration relative to other calcitonin assays (Bieglmayer et al. 2007). This is most relevant for serum calcitonin levels <100 pg/ml. Bieglmayer et al. (2007) proposed to use a conversion factor of 0.8 for the transformation of calcitonin levels measured.
with other calcitonin assays into calcitonin levels obtained with the Immulite 2000 system. For the reverse calculation, the suggested conversion factor was 1.25 (or 1/0.8), which henceforth is referred to as the Bieglmayer correction;

(b) a histopathologic diagnosis of C-cell hyperplasia based on at least >50 C-cells per low-power field (×100) or medullary thyroid cancer (breach of the basement membrane) confined to the thyroid gland no larger than 10 mm in greatest dimension;

(c) negative personal histories of bronchial cancer (Machens et al. 2000b), neuroendocrine cancer (Machens et al. 2000b), renal failure, hyperparathyroidism, follicular cell-derived thyroid cancer, proton pump inhibitors, or any other drug known to increase serum calcitonin levels (Karges et al. 2004, Borget et al. 2007);

(d) a negative RET gene test for exons 10, 11, 13, 14, 15, and 16 (all medullary thyroid cancer patients); exon 8 was not tested in all these patients;

(e) negative family histories of medullary thyroid cancer, pheochromocytoma, or hyperparathyroidism.

Thyroidectomy and cervical lymph node dissection
Fulfilling all these criteria, 100 consecutive patients (74 men and 26 women) qualified for this analysis. In addition to thyroidectomy, 51 (51%) of these patients underwent central lymph node dissections (24 of 26 medullary thyroid cancer and 27 out of 74 C-cell hyperplasia patients), and 23 (23%) patients underwent lateral lymph node dissections for medullary thyroid cancer at the operating surgeon’s discretion. Surgical procedures were conducted using optical magnification and bipolar coagulation, as described elsewhere (Dralle 2002). Informed consent was obtained before each surgical procedure that represented standard practice of care.

Pathological examination and tumor staging
A total of 100 entire thyroid glands were available for histopathologic examination. After gross evaluation by the pathologist, the entire thyroid gland was divided vertically to separate the left and right lobes. The thyroid halves were then sectioned horizontally from the superior to the inferior pole, as previously described (Hinze et al. 1998). After fixation in formalin, the whole thyroid gland was embedded in paraffin. Soft tissue and lymph nodes were processed separately. Conventional staining (hematoxylin and eosin) and calcitonin immunohistochemistry were performed on every surgical specimen, using the standard avidin–biotin complex peroxidase approach and a commercial polyclonal antibody (Immunotech, Marseilles, France). A diagnosis of medullary thyroid cancer was made on evidence of tumor extension beyond the basement membrane, demonstration of lymphatic or vascular invasion on histopathology, or any combination thereof. Primary tumor diameter was ascertained by direct measurements on the surgical thyroid specimens. Histopathologic confirmation was required for diagnosis of lymph node metastasis.

Statistical analysis
Categorical and continuous data were tested with the two-tailed Fisher exact test and Mann–Whitney–Wilcoxon rank sum test respectively. Multiple testing was corrected with the Bonferroni method. Suitability for prediction of medullary thyroid cancer was evaluated using receiver operating characteristics (ROC) curves. The level of significance was set at <0.05.

Results
Study population
As depicted in Table 1, 26 patients harbored occult solitary medullary thyroid cancer, one of which was node positive (one positive among 104 removed nodes). These 26 medullary thyroid cancer patients differed significantly (P < 0.001) from the 74 C-cell hyperplasia patients regarding the prevalence of concomitant goiter (35 vs 76%), basal and stimulated peak calcitonin levels (respective means of 85 vs 19, and 1519 vs 116 pg/ml), the ratio of stimulated peak calcitonin to basal calcitonin serum levels (respective means of 22-fold versus sixfold), and underrepresentation of men (31 vs 89%). No significant differences were found among the two groups with regard to age, histopathologic presence of thyroiditis, or postoperative normalization of calcitonin levels (biochemical cure).

ROC curves
ROC plot analyses (not shown) confirmed the superiority of stimulated over basal calcitonin levels for prediction of medullary thyroid cancer, with mean areas under the curve totaling 0.89 and 0.74 respectively. The fit was better for women (0.99 and 0.85) than men (0.71 and 0.53).
| Table 1 Clinicopathologic composition of the study population with sporadic C-cell disease |
|-----------------------------------------------|----------------|----------------|----------------|----------------|----------------|
| Gender | CCH (n=74) | MTC (n=26) | CCH (n=74) | MTC (n=26) | CCH (n=74) | MTC (n=26) |
|        | Male, n (%) | Male, n (%) | Female (n=8) | Male (n=66) | Female (n=18) | Male (n=8) |
| Age, years, mean (95% CI) | 56 (53; 59) | 61 (57; 66) | 0.07 | 49 (40; 59) | 57 (54; 60) | 0.09 | 60 (55; 65) | 65 (57; 72) | 0.31 |
| Goiter, n (%) | 56 (76) | 9 (35) | <0.001* | 6 (75) | 50 (76) | >0.99 | 6 (33) | 3 (38) | >0.99 |
| Thyroiditis, a n (%) | 6 (8) | 3 (12) | 0.70 | 2 (25) | 4 (6) | 0.12 | 2 (11) | 1 (13) | >0.99 |
| Otherwise normal thyroid, n (%) | 16 (22) | 14 (54) | 0.005* | 1 (13) | 15 (23) | 0.68 | 10 (56) | 4 (50) | >0.99 |
| Basal calcitonin, b pg/ml, mean (95% CI) | 19 (16; 22) | 85 (30; 139) | <0.001* | 14 (7; 21) | 20 (17; 22) | 0.05 | 105 (29; 181) | 39 (0; 92) | 0.04 |
| Peak calcitonin, b, c pg/ml, mean (95% CI) | 116 (95; 137) | 1519 (379; 2659) | <0.001* | 84 (26; 143) | 119 (97; 142) | 0.13 | 2037 (409; 3665) | 352 (7; 698) | 0.005* |
| Peak-to-basal calcitonin ratio, d mean (95% CI) | 6 (5; 7) | 22 (13; 31) | <0.001* | 6 (3; 8) | 6 (5; 7) | 0.89 | 25 (15; 36) | 16 (0; 35) | 0.035 |
| Primary tumor size, mm, mean (95% CI) | 6 (5; 7) | 6 (5; 7) | | 4 (2; 6) | | 0.01 |
| Lymph node metastases, n (%) | 1 (4) | 0 | | 1 (6) | 0 | >0.99 |
| Biochemical cure, e n (%) | 74 (100) | 26 (100) | 1.00 | 8 (100) | 66 (100) | 1.00 | 18 (100) | 8 (100) | 1.00 |

CCH, C-cell hyperplasia; MTC, medullary thyroid cancer. *Significant after Bonferroni correction for multiple testing within each group of comparisons; †P<0.001.

aIncluding Hashimoto’s and Graves’ disease (based on histopathologic criteria).

bThe upper normal limit of the assay being <5 pg/ml for females and <8.4 pg/ml for males respectively.

cStimulated calcitonin level taken 2 or 5 min after i.v. bolus injection of pentagastrin, whichever was higher.

dStimulated peak calcitonin level divided by the corresponding basal calcitonin level.

ePostoperative normalization of increased calcitonin levels at discharge.

fOne positive of 104 removed lymph nodes (basal calcitonin 71 pg/ml; stimulated calcitonin 1569 pg/ml).
Prevalence of medullary thyroid cancer by preoperative calcitonin level

The prevalence of sporadic medullary thyroid cancer in women and men increased across the various segments of calcitonin levels (Table 2). Altogether, gender-specific calcitonin thresholds predicted sporadic medullary thyroid cancer better than unisex thresholds. Intriguingly, women revealed occult medullary thyroid cancer four to eight times more often than men in the lower calcitonin segments (Table 2): 36 vs 10% (increased basal levels of <20 pg/ml); 80 vs 11% (basal levels of 21–50 pg/ml) and 75 vs 9% (stimulated levels of 101–500 pg/ml). In the higher segments, this difference disappeared after levels exceeded 100 pg/ml (basal calcitonin) and 500 pg/ml (stimulated calcitonin) respectively. Because tumors larger than 10 mm were excluded, fewer occult medullary thyroid cancers were seen with higher calcitonin levels.

<table>
<thead>
<tr>
<th>Serum calcitonina pg/ml</th>
<th>Female (n=26)</th>
<th>Male (n=74)</th>
<th>Total (n=100)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Total, n MTC, n (%)</td>
<td>Size, mm, median (range)</td>
<td>Total, n MTC, n (%)</td>
</tr>
<tr>
<td>Basal</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤20</td>
<td>11 (436) 4 (10)</td>
<td>7 (2–8)</td>
<td>52 (11) 4 (2–5)</td>
</tr>
<tr>
<td>21–50</td>
<td>5 (80) 2 (11)</td>
<td>6 (4–6)</td>
<td>19 (11) 3 (4–4)</td>
</tr>
<tr>
<td>51–100</td>
<td>6 (100) 2 (100)</td>
<td>6 (2–9)</td>
<td>2 (0)</td>
</tr>
<tr>
<td>&gt;100</td>
<td>4 (100) 1 (100)</td>
<td>10 (9–10)</td>
<td>1 (100) 8</td>
</tr>
<tr>
<td>Peakc</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤100</td>
<td>6 (0) 3 (8)</td>
<td>2 (2–4)</td>
<td>39 (8)</td>
</tr>
<tr>
<td>101–500</td>
<td>8 (75) 3 (9)</td>
<td>6 (2–8)</td>
<td>33 (9)</td>
</tr>
<tr>
<td>501–1000</td>
<td>3 (100) 1 (100)</td>
<td>5 (4–6)</td>
<td>1 (100) 4</td>
</tr>
<tr>
<td>&gt;1000</td>
<td>9 (100) 1 (100)</td>
<td>9 (5–10)</td>
<td>1 (100) 8</td>
</tr>
</tbody>
</table>

MTC, medullary thyroid cancer.
aThe upper normal limit of the assay being <5 pg/ml for females and <8.4 pg/ml for males respectively.
bOne of which was node positive (one positive of 104 removed lymph nodes; basal calcitonin 71 pg/ml; stimulated calcitonin 1569 pg/ml).
cStimulated calcitonin level taken 2 or 5 min after i.v. bolus injection of pentagastrin, whichever was higher.

Sensitivity, specificity, positive and negative predictive value, and accuracy for medullary thyroid cancer

Sensitivity, specificity, positive and negative predictive value, and accuracy of various calcitonin thresholds were calculated to explore the impact of these gender-specific disparities on the prediction of medullary thyroid cancer. Most discriminatory between C-cell hyperplasia and medullary thyroid cancer was a basal calcitonin threshold of 15 pg/ml (20 pg/ml after Bieglmayer correction) for women and 80 pg/ml (corrected 100 pg/ml) for men, based on the greatest accuracy at the lowest possible calcitonin level (Table 3). The corresponding stimulated peak calcitonin thresholds were 80 pg/ml (corrected 100) for women and 500 pg/ml for men (Table 4).

These basal and stimulated peak calcitonin thresholds, however, were not ideal for women because

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Table 3 Sensitivity, specificity, positive and negative predictive value, and accuracy of basal calcitonin levels for medullary thyroid cancer

<table>
<thead>
<tr>
<th>Basal calcitonin threshold (pg/ml)</th>
<th>Female (n=26)</th>
<th>Male (n=74)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Immulite 2000 assay</td>
<td>10 15 20 40 80</td>
<td>20 40 60 80 100</td>
</tr>
<tr>
<td>Bieglmayer corrected (×1.25)a</td>
<td>10 20 25 50 100</td>
<td>25 50 75 100 (125)</td>
</tr>
<tr>
<td>Sensitivity (%)</td>
<td>89 89 83 67 33</td>
<td>50 13 13 13 13</td>
</tr>
<tr>
<td>Specificity (%)</td>
<td>25 75 75 100 100</td>
<td>65 96 99 100 100</td>
</tr>
<tr>
<td>Positive predictive value (%)</td>
<td>73 89 88 100 100</td>
<td>15 25 50 100 100</td>
</tr>
<tr>
<td>Negative predictive value (%)</td>
<td>50 75 67 57 40</td>
<td>92 90 90 90 90</td>
</tr>
<tr>
<td>Accuracy (%)</td>
<td>69 85 81 77 54</td>
<td>64 87 89 91 91</td>
</tr>
</tbody>
</table>

aConversion factor proposed by Bieglmayer et al. (2007), which may be less relevant for calcitonin concentrations >100 pg/ml. Converted calcitonin levels >100 pg/ml are parenthesized to indicate this fact.
Table 4 Sensitivity, specificity, positive and negative predictive value, and accuracy of stimulated peak calcitonin levels for medullary thyroid cancer

<table>
<thead>
<tr>
<th></th>
<th>Female (n=26)</th>
<th>Male (n=74)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Stimulated peak calcitonin threshold (pg/ml)</td>
<td>Stimulated peak calcitonin threshold (pg/ml)</td>
</tr>
<tr>
<td>Immulite 2000 assay</td>
<td>60 to 200</td>
<td>100 to 200</td>
</tr>
<tr>
<td>Bieglmayer corrected (×1.25)</td>
<td>75 to 250</td>
<td>100 to 250</td>
</tr>
<tr>
<td>Sensitivity (%)</td>
<td>100 to 94</td>
<td>38 to 25</td>
</tr>
<tr>
<td>Specificity (%)</td>
<td>63 to 88</td>
<td>53 to 97</td>
</tr>
<tr>
<td>Positive predictive value (%)</td>
<td>86 to 100</td>
<td>14 to 23</td>
</tr>
<tr>
<td>Negative predictive value (%)</td>
<td>100 to 80</td>
<td>92 to 92</td>
</tr>
<tr>
<td>Accuracy (%)</td>
<td>89 to 92</td>
<td>80 to 92</td>
</tr>
</tbody>
</table>

Table 4: Sensitivity, specificity, positive and negative predictive value, and accuracy of stimulated peak calcitonin levels for medullary thyroid cancer.

Discussion

This is the first series exploring gender-specific thresholds in patients with increased basal and stimulated calcitonin levels to differentiate between sporadic C-cell hyperplasia and occult sporadic medullary thyroid cancer. In these elusive conditions, calcitonin levels overlap widely so that the incremental diagnostic value of calcitonin screening is most evident in this setting (Borget et al. 2007).

Gender-specific disparities in calcitonin secretion

The most striking finding of this series was the four- to eightfold lower yield of medullary thyroid cancer for men in the lower calcitonin segment (≤50 pg/ml). Because male thyroids harbor more C-cells than female thyroids, serum calcitonin levels are more often higher in men than women (Heath & Sizemore 1977, Hahm et al. 2001, Vierhapper et al. 2005, Scheuba et al. 2009). This observation explains why men outnumbered women in the sporadic C-cell hyperplasia group by more than eightfold (66 men versus 8 women; P < 0.001), whereas women predominated twofold in the occult medullary thyroid cancer group (18 women versus 8 men; Table 1). Similar gender-specific disparities have been reported elsewhere (Rink et al. 2009, Scheuba et al. 2009).

In this study, increased basal calcitonin levels of ≤50 pg/ml for men were associated with much poorer positive predictive values for occult medullary thyroid cancer. It seems that occult medullary thyroid cancers have greater difficulties in men than women to release those quantities of calcitonin into the bloodstream necessary to overcome the background noise of basal calcitonin secretion. As a corollary, all calcitonin thresholds were consistently higher for men than women. Stimulation with pentagastrin afforded better discrimination between C-cell hyperplasia and medullary thyroid cancer in both women and men, especially when calcitonin levels were lower, as noted previously (Iacobone et al. 2002, Vierhapper et al. 2005, Costante et al. 2007, Scheuba et al. 2009).

By implication, pentagastrin stimulation should ideally be performed in all patients with increased basal calcitonin levels unless these levels are high. Of note, pentagastrin has become restricted in some countries.

Calcitonin thresholds in the literature

Despite the use of diverse selection criteria and different calcitonin assays in heterogeneous populations with variable degrees of histopathologic...
confirmation, there is universal agreement regarding the following statements:

(a) a basal calcitonin level within normal limits of the assay practically excludes medullary thyroid cancer (Elisei et al. 2004, Rink et al. 2009, Scheuba et al. 2009)

(b) the likelihood of medullary thyroid cancer is remote when stimulated calcitonin levels are <100 pg/ml (Hahm et al. 2001, Karges et al. 2004, Vierhapper et al. 2005, Rink et al. 2009, Scheuba et al. 2009)


Traditionally, calcitonin thresholds, most of which differ slightly between institutions, have been conceived as less-than-perfect static means of balancing the consequences of overtreatment and undertreatment. A more dynamic concept consisting of gender-specific relative (20 and 100 pg/ml for women) and absolute (50 and 250 pg/ml for women; 100 and 500 pg/ml for men) basal and stimulated peak calcitonin thresholds may yield greater accuracy and flexibility: close biochemical surveillance for the crossing of the relative threshold; immediate thyroidectomy for the breach of the absolute threshold.

Clinical work-up for sporadic C-cell disease

Calcitonin serum levels, representing a biologic continuum like any other biomarker levels, should always be interpreted within the clinical context. This involves consideration of thyroid nodules and/or suspect cervical lymph nodes visualizing on high-resolution ultrasonography and appreciation of suspicious or positive fine-needle aspiration cytology results using immunohistochemistry for calcitonin and determination of calcitonin in the washout of the needle. For occult medullary thyroid cancer, however, there is a dearth of information on the accuracy and utility of these diagnostic methods, which have been shown to work well in more advanced disease. An association between C-cell hyperplasia and thyroiditis or thyroid autoantibodies, which were not measured in this study, is controversial. As a matter of fact, no statistically significant correlations were found a) between the presence of C-cell hyperplasia and chronic lymphocytic thyroiditis on thyroid surgical specimens from 112 patients (Guyetant et al. 1994), and b) basal calcitonin serum levels and anti-thyroid peroxidase or anti-thyroglobulin antibodies among 298 age-matched hypothyroid or euthyroid goitrous women (Pantazi & Papapetrou 1998). In keeping, the prevalence of histopathologically diagnosed thyroiditidis in this series was broadly comparable among patients with C-cell hyperplasia and lower calcitonin serum levels, and patients with medullary thyroid cancer and higher calcitonin serum levels (Table 1).

Clinical implications

Because it represents an innocuous intervention with relatively low resource consumption, calcitonin measurement is a powerful tool for detecting C cell disease at a stage when it is still curable. Although the ultimate proof of a screening test’s value lies in the degree to which it extends the survival of those who submit to it, it may take decades for such an extension of life to show statistically. To augment the cost-effectiveness of calcitonin screening, its methodology must be refined. Although recent European and American cost-benefit studies have hailed calcitonin screening as cost-effective for patients with nodular goiter (Borget et al. 2007, Cheung et al. 2008), there is still room for improvement. A dynamic concept of gender-specific calcitonin thresholds attuned to the inherent risk of medullary thyroid cancer would help to reduce the unavoidable harms of screening by minimizing the numbers of false positives and unnecessary operations. These false positives have enormous clinical and economic potential for damage: if allowed to spiral out of control, they expand human suffering and inevitably drive up costs, damaging and ultimately destroying the cost-effectiveness of any screening program.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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