Should parafibromin staining replace *HRTP2* gene analysis as an additional tool for histologic diagnosis of parathyroid carcinoma?

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Abstract

**Objective:** *HRPT2* gene mutations are associated with parathyroid carcinomas, and absence of parafibromin immunoreactivity has been suggested as a diagnostic marker of malignancy. The aim of our study was to extend parafibromin studies in a series of benign and malignant parathyroid tumors and cross-validate the results of immunohistochemistry with those of *HRPT2* analysis.

**Design and patients:** We performed parafibromin and cyclin D1 immunostaining and *HRPT2* gene analysis using loss of heterozygosity studies and sequencing analysis in parathyroid specimens from 11 patients with carcinoma (eleven primary tumors, one skin, and four lung metastases), 22 with sporadic adenomas, and 4 with atypical adenomas.

**Results:** Ten out of eleven parathyroid cancers were negative for parafibromin staining and showed *HRPT2* gene abnormalities. The remaining sample was negative for immunostaining and genetic analyses. All but one sporadic adenomas showed parafibromin immunoreactivity and no *HRPT2* gene abnormalities. The sample with negative immunostaining carried an *HRPT2* mutation. Two atypical adenomas were positive and two negative with parafibromin staining. No *HRPT2* abnormalities were found in these samples. Cyclin D1 expression was heterogeneous and there was no relationship between expression/expression level of cyclin D1 and parafibromin expression.

**Conclusions:** We have shown that negative parafibromin staining is almost invariably associated with *HRPT2* mutations and confirm that loss of parafibromin staining strongly predicts parathyroid malignancy. In clinical practice, these tests could be particularly useful in the subset of parathyroid tumors with equivocal histological examination. However, their diagnostic value in this setting remains to be proven.

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Introduction

Primary hyperparathyroidism (PHPT) is the most common cause of hypercalcemia in the general population and is caused in approximately 85% of cases by a single hyperfunctioning adenoma or, more rarely, by a carcinoma (1).

The course of parathyroid carcinoma is rather indolent and it is common that the diagnosis of malignancy is made retrospectively following recurrence of the disease (2). Indeed, at initial parathyroidectomy, the diagnosis of malignancy can be established with certainty only in those tumors showing evidence of vascular invasion, perineural space invasion and growth into adjacent tissues or distant metastases (3). Other histological features characteristic of carcinoma, such as trabecular pattern, mitoses, thick fibrous bands and capsular invasion, may also be present in the so-called atypical adenoma and therefore cannot be considered diagnostic (4–6).

In recent years, several attempts have been made to identify a diagnostic marker of parathyroid malignancy (7–10). Loss of Rb1 protein immunostaining was shown to be a potentially useful diagnostic tool (7) but, because of subsequent contradictory results, this test is not routinely used in clinical practice.

The recent demonstration that the *HRPT2* gene is involved in the pathogenesis of sporadic parathyroid cancer has raised the possibility that analysis of this gene might have some diagnostic utility (11–14). Parafibromin, the 531 amino acid protein encoded by the *HRPT2* gene (15), is part of the human Paf1 complex and involved in transcription regulation, histone modification and cell proliferation (16–21). It has been shown that parafibromin down-regulates the expression of cyclin D1 (16, 20). The protein is mostly located in the
nucleus and the nuclear localization signal (NLS) was originally identified at residues 125–139 (19) and most recently at residues 136–139 (21). Inactivating mutants of parafibromin were reported in 77% of sporadic parathyroid carcinomas (11–13) and loss of immunoreactivity for parafibromin was shown in the majority of parathyroid carcinomas (22, 23). Thus, parafibromin immunostaining, because of its lower cost and greater feasibility when compared with genetic analysis, could become a promising molecular marker for the diagnosis of parathyroid carcinoma.

The aim of the present study was to extend parafibromin immunostaining studies in a series of benign and malignant parathyroid tumors, and cross-validate their results with those of HRPT2 gene analysis.

Materials and methods

PHPT patients
The study was approved by our Internal Review Board. All patients gave their informed consent for genetic studies.

The diagnosis of PHPT was based on increased total or ionized serum calcium (>2.55 and >1.32 mmol/l respectively) and inappropriately elevated serum parathyroid hormone (PTH) levels (normal range 10–75 ng/l).

Sporadic parathyroid carcinomas All patients had either local recurrence and/or distant metastases after initial parathyroid surgery. The clinical and genetic data of seven patients (nos 1–7) were previously reported (13). The remaining four patients (nos 8–11) are described in more detail herein (Table 1). Parathyroid cancer was diagnosed at the age of 45, 51, 56, and 52 years respectively. Patients underwent parathyroidectomy and later had evidence of either local recurrence (n = 2), cervical lymph node metastasis (n = 1), or multiple lung metastases (n = 3). The interval between initial surgery and the first recurrence ranged between 7 months and 4 years (mean 2 years). Various degrees of hypercalcemia developed during follow-up (maximum mean serum calcium 3.5 mmol/l). No patient was treated with chemotherapy; two (nos 1, 10) received external neck irradiation and one (no. 8) a short treatment with cinacalcet. Intravenous pamidronate or zoledronate were used in most cases in an attempt to reduce the degree of hypercalcemia.

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>Age at diagnosis (year/sex)</th>
<th>Cystic features</th>
<th>HRPT2 mutational status</th>
<th>Effect on parafibromin</th>
<th>HRPT2 mutation</th>
<th>LOH\textsuperscript{c} at HRPT2</th>
<th>Parafibromin immunoreactivity\textsuperscript{d}</th>
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</thead>
<tbody>
<tr>
<td>1 Carcinoma</td>
<td>43/F</td>
<td>No</td>
<td>Wild type</td>
<td>–</td>
<td>–</td>
<td>Yes</td>
<td>Negative</td>
</tr>
<tr>
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<td>32/F</td>
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<td>Wild type</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Skin metastasis</td>
<td>46/M</td>
<td>No</td>
<td>Wild type</td>
<td>–</td>
<td>–</td>
<td>Yes</td>
<td>Negative</td>
</tr>
<tr>
<td>2 Carcinoma</td>
<td>63/M</td>
<td>No</td>
<td>700C&gt;T in exon 7\textsuperscript{d}</td>
<td>Stop codon R234X</td>
<td>Germ line</td>
<td>No</td>
<td>Negative</td>
</tr>
<tr>
<td>Lung metastasis</td>
<td>53/M</td>
<td>No</td>
<td>195insT in exon 3</td>
<td>Stop codon at 104</td>
<td>Somatic</td>
<td>Yes</td>
<td>Negative</td>
</tr>
<tr>
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<td>No</td>
<td>25C&gt;T in exon 1</td>
<td>Stop codon R9X</td>
<td>Germ line</td>
<td>No</td>
<td>Negative</td>
</tr>
<tr>
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<td>56/F</td>
<td>No</td>
<td>195insA in exon 2</td>
<td>Stop codon at 104</td>
<td>Somatic</td>
<td>Yes</td>
<td>Negative</td>
</tr>
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</tr>
<tr>
<td>6 Carcinoma</td>
<td>45/M</td>
<td>No</td>
<td>700 C&gt;T in exon 7\textsuperscript{d}</td>
<td>Stop codon R234X</td>
<td>Germ line</td>
<td>Yes</td>
<td>Negative</td>
</tr>
<tr>
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<td>Negative</td>
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<td>Wild type</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>10 Carcinoma</td>
<td>52/M</td>
<td>No</td>
<td>Wild type</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>11 Carcinoma</td>
<td>44/M</td>
<td>No</td>
<td>Wild type</td>
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<td>–</td>
<td>–</td>
<td>–</td>
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<td>Wild type</td>
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<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>2.2 Atypical adenoma</td>
<td>52/M</td>
<td>No</td>
<td>Wild type</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
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<tr>
<td>2.3 Atypical adenoma</td>
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<td>Wild type</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>2.4 Atypical adenoma</td>
<td>48/M</td>
<td>No</td>
<td>Wild type</td>
<td>–</td>
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</tbody>
</table>

\textsuperscript{a}Clinical data and HRPT2 gene analyses of patients nos 1–7 have previously been reported (Ref. (13)).

\textsuperscript{b}Number in parentheses indicate the percentage of positive cells.

\textsuperscript{c}Loss of heterozygosity.

\textsuperscript{d}These two patients, showing the same HRPT2 mutation, were unrelated.

Table 1 Clinical data, HRPT2 gene analyses, and parafibromin immunostaining in parathyroid carcinomas and atypical adenomas\textsuperscript{a}.
fibrosis, mitoses and fibrous bands, but lack of unequivocal signs of capsular or vascular invasion (5).

Follow-up after surgery ranged from 8 months to 3.2 years (mean 2 years). One patient (no. 2.5) was cured, one (no. 2.1) had permanent hypoparathyroidism treated with calcium and calcitriol, one (no. 2.2) had persistent PHPT, and one (no. 2.3) had moderately elevated PTH levels (126 pg/ml) with normocalcemia. None of the latter two showed abnormalities in the imaging studies (neck ultrasound, neck and lung CT scan, and sestamibi scan).

**Tissue samples**

Twenty-two parathyroid adenomas, four atypical adenomas, and sixteen cancer specimens (eleven primary tumors, one skin metastasis, and four lung metastases) were studied.

Tissues were obtained at the time of surgery, immediately snap frozen in liquid nitrogen, and stored at −80 °C until use.

**Pathological findings**

All tumors were histologically defined according to the recent World Health Organization classification (4, 5).

All parathyroid specimens were carefully reviewed by two pathologists (PV and GF) in order to confirm the histological diagnosis by the use of homogeneous criteria, eventually describe the presence of cystic features, and check that all samples had a sufficient proportion of tumor cells for DNA analyses (>70%).

**HRPT2 gene analyses**

Genomic DNA was isolated from peripheral blood leukocytes and parathyroid tissue by the standard proteinase K-SDS digestion and the phenol/chloroform method. Allelic deletions and direct sequencing of HRPT2 gene were assessed as described previously (13).

**Immunohistochemistry**

Archival sections were deparaffinized in xylene and rehydrated in alcohol. Endogenous peroxide activity was blocked by incubating the slides in 1% hydrogen peroxide in methanol for 10 min. In order to unmask the antigen, the slides were microwave treated in 10 mM citrate buffer (pH 6.0) for 10 min. After blocking nonspecific staining with normal serum, the sections were incubated for 1 h with the primary monoclonal antibody (MAb) used at the dilution ratio of 1:50. The parafibromin antibody (kindly donated by Bin Tean Teh) is directed against the portion of the protein corresponding to amino acid positions 87–100 (22). The cyclin D1 antibody (clone SP4) was from Lab Vision (Fremont, CA, USA). The sections were then incubated with biotin-labeled secondary antibody (dilution 1:500) and avidin–biotin complex (Vector Burlingame, Burlingame, CA, USA) for 30 min each. Sites of binding were visualized using 3,3-diaminobenzidine as the chromogen. Finally, sections were counterstained with hematoxylin, dehydrated, and mounted. The positive control was normal parathyroid tissue for parafibromin and a breast carcinoma overexpressing cyclin D1 for cyclin D1; negative controls included experiments omitting primary antibody or using primary antibody preabsorbed with a 20-fold excess of the immunizing peptide. For each case, six different sections were analyzed.

Tumors were scored as positive if specific nuclear staining was detected, and the staining was quantified according to the percentage of positive cells, independent of the intensity of staining. Tumors were scored as negative when no tumor cells showed a specific nuclear staining.

Each section was evaluated by two independent observers (PV and GF) without knowledge of the diagnosis or outcome. In cases where the assessment of percentage positivity differed between the two observers, the disagreements were resolved by reaching a consensus after joint review using a conference microscope.

**Statistical analysis**

Sensitivity, specificity, predictive values and 95% confidence interval (CI) for proportions were calculated using standard methods for binomial distribution. The χ² test, Fisher’s exact test and the Mann–Whitney test were used as indicated.

**Results**

**HRPT2 gene analyses**

Results of loss of heterozygosity (LOH) and mutational analyses of the 22 adenomas and carcinomas specimens from patients nos 1–7 have previously been reported (13). We report herein the results observed in additional samples (four atypical adenomas and four carcinomas) (Table 1). All tissues were informative at least for one polymorphic site at LOH analysis and, with the exception of a cancer specimen, showed retention of heterozygosity. HRPT2 mutations resulting in a premature stop codon were found in three out of four carcinomas (Table 1); all atypical adenomas were negative. The HRPT2 mutations described here were not found in 50 unrelated normal individuals of Italian origin.

**Parafibromin immunohistochemistry**

In each experiment normal parathyroid gland tissue was used as the positive control and showed a diffuse nuclear staining. Immunoreactivity was highly specific for...
parathyroid chief cells, without staining of adipose or connective tissues or blood vessels (Fig. 1A). Controls with antibody, pre-absorbed with a 20-fold excess of specific peptide, did not show any immunoreactivity.

All parathyroid carcinomas and metastases were negative for parafibromin (Table 1; Fig. 1C and D). In one tumor a weak cytoplasmic signal was observed. In contrast, all but one parathyroid adenomas were parafibromin positive (Fig. 1B). Parafibromin expression was heterogeneous with a range of positive cells between 10 and 80%. Immunoreactivity was not detected in the adenoma carrying an \( \text{HRPT2} \) mutation (Fig. 1E).

Two atypical adenomas specimens were positive (patients nos 2.3 and 2.5; 20 and 40% of positive cells respectively) and two (nos 2.1 and 2.2) were negative (Table 1). A representative case is shown in Fig. 1F.

**Correlation between parafibromin immunostaining and HRPT2 gene alterations**

Status of parafibromin expression was compared with \( \text{HRPT2} \) gene alterations (Tables 1 and 2 and Ref. [13]). Ten out of eleven parathyroid carcinomas, all negative for parafibromin immunostaining, showed \( \text{HRPT2} \) gene

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**Figure 1** Immunohistochemical analyses of parafibromin expression. A: Normal parathyroid gland. The parathyroid cells showed a diffuse nuclear immunoreactivity (×200). B: Parathyroid adenoma. The majority of cells showed a nuclear immunoreactivity for parafibromin. Some nuclei were negative. C: Parathyroid carcinoma. The neoplastic cells were completely negative for parafibromin (×200). D: Lung metastasis of parathyroid carcinoma. The tumor showed a diffuse loss of staining. Adjacent lung parenchyma was also negative (×200). E: Parathyroid adenoma carrying \( \text{HRPT2} \) mutation. The tumor cells were completely negative for parafibromin (×400). F: Atypical parathyroid adenoma (no. 2.1). The tumor showed sheets of neoplastic cells separated by fibrous bands. No invasion of the capsule was evident (hematoxylin–eosin, ×100). Inset shows that the cells were completely negative for parafibromin (×200).

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abnormalities (mutations in nine cases, associated \((n = 6)\) or not \((n = 3)\) with LOH, and LOH alone in one case). The remaining sample was negative at immunostaining and genetic analyses.

All but one adenoma showed parafibromin immunoreactivity and no \(HRPT2\) gene abnormalities were detected in these samples. The tumor with negative immunostaining carried an \(HRPT2\) mutation.

As mentioned before, all mutations generated premature stop codons which are predicted to cause truncated proteins of different size. Figure 2 depicts a schematic representation of parafibromin, including the binding site of the MAb used in the present study, the NLS of the protein (19), and the truncated proteins resulting from the \(HRPT2\) mutations detected in our tumor series. Six mutations (R9X, E24X, IVS1+1G>A, L61X, 195insA, and 195insT) were located in the N-terminal part of the protein before the antibody-binding site. Thus, the resulting truncated proteins do not contain either the amino acid sequence recognized by the antibody or the NLS. The R139X mutation was located just at the end of the C-terminal arm of the NLS, and the R234X far downstream with respect to either the antibody-binding site or the NLS. Therefore, in both cases, the altered proteins included the antibody-binding site and the NLS.

Two atypical adenomas were positive and two negative at immunohistochemistry. No LOH or \(HRPT2\) mutations were found in these.

In summary, 11 out of 14 cases (ten carcinomas and one adenoma) showed an association between the loss of parafibromin staining and the \(HRPT2\) gene abnormalities, whereas no gene alteration was detected in three cases (one carcinoma and two atypical adenomas). On the other hand, parafibromin expression was associated in all cases with normal \(HRPT2\) gene analysis (21 sporadic and 2 atypical adenomas \((P < 0.0001\) by Fisher’s exact test)).

### Diagnostic values of \(HRPT2\) gene abnormalities

The diagnostic values of \(HRPT2\) gene abnormalities for differentiating parathyroid carcinoma from adenoma are shown in Table 2. Loss of parafibromin staining had a sensitivity of 100%, greater than that of \(HRPT2\) mutation and LOH; on the other hand, the latter two tests had a greater specificity when compared with immunostaining. It is commonly estimated that parathyroid cancer accounts for <1% of cases of sporadic PHPT. At our institution, the prevalence of parathyroid cancer is 0.5% (3 out of 555 consecutive patients with sporadic PHPT submitted to parathyroidectomy over the last 8 years). A higher prevalence (5%) has been reported by another Italian group in 290 PHPT patients submitted to surgery between 1980 and 1996 (24). We have no explanation for such a discrepancy. The positive

<table>
<thead>
<tr>
<th></th>
<th>Loss of parafibromin immunostaining</th>
<th>(HRPT2) mutation</th>
<th>Loss of heterozygosity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensitivity (95% CI)</td>
<td>100 (68–100)</td>
<td>82 (48–97)</td>
<td>64 (32–88)</td>
</tr>
<tr>
<td>Specificity (95% CI)</td>
<td>88 (69–97)</td>
<td>96 (78–100)</td>
<td>96 (78–100)</td>
</tr>
<tr>
<td>Positive predictive value at 0.5% prevalence (95% CI)</td>
<td>4 (0–11)</td>
<td>10 (1–19)</td>
<td>8 (0–16)</td>
</tr>
<tr>
<td>Negative predictive value at 0.5% prevalence (95% CI)</td>
<td>100 (100–100)</td>
<td>100 (99–100)</td>
<td>100 (98–100)</td>
</tr>
<tr>
<td>Positive predictive value at 5% prevalence (95% CI)</td>
<td>31 (16–46)</td>
<td>53 (37–69)</td>
<td>46 (30–63)</td>
</tr>
<tr>
<td>Negative predictive value at 5% prevalence (95% CI)</td>
<td>100 (100–100)</td>
<td>100 (96–100)</td>
<td>98 (94–100)</td>
</tr>
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</table>
predictive values (PPV) and negative predictive values (NPV), with 95% CI. for an estimated prevalence of 0.5 and 5% of cancer among patients with PHPT, are also reported in Table 2.

**Relationship between cyclin D1 expression and parafibromin expression**

Cyclin D1 expression was heterogeneous with a range of positive cells between 1 and 80% and was graded on the following scale: mild (1–10% positive nuclei), moderate (11–30% positive nuclei), and strong (> 30% positive nuclei). There was no relationship between expression/level of expression of cyclin D1 and parafibromin expression (Table 3) or HRPT2 gene alterations (data not shown).

**Discussion**

The aim of the present study was to further validate the use of parafibromin staining in the diagnosis of parathyroid malignancy, by comparing immunohistochemistry results and HRPT2 genetic analysis in a series of benign and malignant parathyroid tumors. Twenty-one out of twenty-two sporadic adenomas and two out of four atypical adenomas showed parafibromin immunoreactivity. All carcinomas, one adenoma and the remaining two atypical adenomas were negative at immunostaining. Thus, we confirm that loss of parafibromin staining strongly predicts malignancy.

With the exception of three cases (one carcinoma and two atypical adenomas), all parafibromin negative samples showed HRPT2 abnormalities, responsible for the synthesis of differently truncated proteins. Lack of the antibody-binding site could account for the negative immunostaining results in five cases. In the remaining six, the HRPT2 abnormalities were located downstream to the antibody-binding sequence and therefore the negative staining might be due to a less antigenic or a less stable mutated protein.

The combined evaluation of parafibromin expression and HRPT2 gene analysis was previously carried out in a 56 parathyroid tumors (41 sporadic adenomas (25), 2 sporadic carcinomas (23), and 13 hyperparathyroidism-jaw tumor adenomas (22, 23)). Parafibromin staining was negative in 16 out of 19 samples carrying HRPT2 mutations (all carcinomas and HPT-JT adenomas, and four adenomas). All the remaining samples (37 adenomas) showed positive parafibromin staining and absence of HRPT2 gene alterations. This data together with our results indicate that negative parafibromin staining is almost invariably associated with HRPT2 mutations (27 out of 30 cases, 90%) and positive staining with no HRPT2 alterations (60 out of 63 cases, 95%). Only a minority of cases (6 of a total of 93 studied) showed discrepant results (parafibromin staining in the presence of HRPT2 alterations or vice versa). In three cases (present study), the finding of negative parafibromin staining in the absence of abnormalities in the HRPT2 coding and splice site regions could be explained by the occurrence of still unidentified mutations in the promoter or other regulatory regions of the gene, or alterations in the post-transcriptional processing of parafibromin with loss of immunoreactivity. On the other hand, missense mutations could explain retained parafibromin staining in two out of the three tumors reported by other authors.

As discussed by Gill (23), missense mutations result in the production of a full-length parafibromin, which is still recognized by the antibody but is presumably less biologically active.

It could be argued that the two atypical parathyroid adenomas with negative parafibromin immunostaining and no HRPT2 gene abnormalities might have been misdiagnosed and be early stage parathyroid cancers. Indeed, loss of parafibromin not associated with HRPT2 gene abnormalities was observed in one of our carcinomas. This hypothesis is not supported by the follow-up, showing no evidence of local recurrence or distant metastasis after 33 and 44 months.

As mentioned before, all but one of the adenomas we studied were negative at HRPT2 gene analysis, thus confirming the low frequency of HRPT2 gene abnormalities in sporadic parathyroid adenomas (12–13, 15, 25–27). The association in this adenoma of a negative parafibromin staining prompted us to re-evaluate

<table>
<thead>
<tr>
<th>Cyclin D1 staining</th>
<th>Adenomas (n=22)</th>
<th>Atypical adenomas (n=4)</th>
<th>Carcinomas (n=11)</th>
<th>Total (n=37)</th>
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<tr>
<td></td>
<td>Parafibromin staining</td>
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</tr>
<tr>
<td></td>
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</tr>
<tr>
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<td>13</td>
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<td>Mild (1–10)</td>
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<td>Moderate (11–30)</td>
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<td>Strong (&gt;30)</td>
<td>2</td>
<td>–</td>
<td>1</td>
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</table>

*aNumber in parentheses indicates the percentage of positive cells.

*bStatistical analysis was performed in the whole series of parathyroid tumors: (i) P=0.244 (by χ² test). (ii) P=0.183 (by Fisher’s exact test after grouping together all samples with cyclin D1 positive cells); (iii) P=0.23 (by Mann–Whitney test using individual percentages of cells positive for cyclin D1 expression).
the histology. No features commonly observed in malignant lesions were found. The benign nature of this adenoma was also suggested by the long-term remission of PHPT.

In vitro studies have shown that parafibromin down regulates cyclin D1 expression. Our data show that both high and low level of cyclin D1 expression were found in malignant and benign parathyroid tumors, independent of the HRPT2 mutational status, indicating that other mechanisms are involved in the control of cyclin D1 expression in parathyroid cells. Similar results have recently been reported by Juhlin et al. in parathyroid adenomas (25).

The clinical utility of a diagnostic test depends on the prevalence of a disease in a given population, as well as the sensitivity and specificity. Given the rarity of parathyroid carcinoma (0.5% of all cases of PHPT in our series), loss of parafibromin staining, as well as HRPT2 gene abnormalities, has a rather low PPV and therefore these both tests are not recommended for analysis of all parathyroid tumors, but would be most applicable to a subset of tumors with atypical features. Limiting these diagnostic tests to this subset will increase the PPV. However, because of the limited number of atypical adenomas no definite indication can be provided, even though the genetic analysis seems to have a better diagnostic accuracy. All tests showed an NPV of 100% but, based on our results, only the presence of parafibromin staining excludes malignancy (95% CI, 100–100). This conclusion cannot be generalized since three carcinomas described by Gill showed a positive staining (23).

Parafibromin antibodies are commercially available and therefore parafibromin immunostaining, being cheaper and less time consuming when compared with HRPT2 genetic analysis, is expected to become widely used in cases which pose diagnostic challenges. The question of whether it should be preferred to genetic analysis can only be answered by studies including a greater number of equivocal parathyroid tumors.

Acknowledgements

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