Non-functional parathyroid cyst – diagnostic pitfall: A case-report

Nefunkcionalna cista paratiroidne žlezde – dijagnostička zamka

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Abstract

Introduction. Parathyroid cysts are relatively rare and they may be misdiagnosed with thyroid nodules. Parathyroid cysts are characterized by elevated level of parathyroid hormone (PTH) in cystic fluid. Case report. We reported about middle-aged woman with palpable node in the left thyroid lobe. Ultrasound showed anechoic 40 × 25 mm lesion in the left thyroid lobe. Fine needle aspiration (FNA) obtained 13 mL colorless, watery cystic fluid. PTH value in cystic fluid was ten fold more in comparison with serum PTH. Serum PTH was slightly elevated, D vitamin was under the reference range, serum calcium and phosphorus were normal as well as thyroid hormones. Thyroglobulin antibodies (TgAb) and thyroid peroxidase antibodies (TPOAb) were not detected. Radionuclide parathyroid scintigraphy indicated at physiological metaiodbenzylguanidine (MIBG) distribution. After six months of vitamin D supplementation, serum calcium, phosphorus, vitamin D and PTH were normal. This finding was indicative that was a nonfunctional parathyroid cyst.

Conclusion. This case report points out that thyroid cystic lesions with thin walls, and reverberation in ultrasound, must be observed as a potential parathyroid cyst. These cysts require caution during diagnostic aspiration because of danger of hypercalcemic crises due to FNA, which can be a life-threatening condition.

Key words: parathyroid diseases; cysts; biopsy, fine needle; hypercalcemia.

Introduction

Parathyroid cysts have been relatively rare described in clinical practice, and till now, less than 300 cases have been reported in the literature. They are often confused with thyroid nodules. Parathyroid cysts could be diagnosed by ultrasound and verified by elevated parathyroid hormone (PTH) in aspirated cystic fluid. First description of macroscopic characteristic of parathyroid cysts was described in 1880. About 40% to 80% parathyroid lesions have been diagnosed by finding parathyroid cells in aspirated cystic fluid, about 8% to 30% are misdiagnosed as a thyroid lesion, and 8% to 16% have no enough cell elements for cytological diagnose. Cystic fluid is watery, colorless and looks as

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lymph fluid, called "water from rocks". Usually, there is no relapse after fine needle aspiration (FNA) and evacuation of cystic fluid after one-year follow-up.

Case report

We examined a 40-year-old female with discomfort in the lower part of the anterior cervical region. In her left thyroid lobe there was elastic, palpable node, approximately 40 mm. An ultrasound examination revealed anechoic and oval-shaped lesion measuring $24 \times 40$ mm in the left thyroid lobe, with reverberation and without register of blood flow (Figure 1). In the right thyroid lobe there was a 5 mm hypoechoic, colloid node.

Fine-needle aspiration was performed in order to evacuate the cystic fluid and to make cytology analyses. Fluid obtained by aspiration was watery, colorless, and it suspected to parathyroid cyst. In 13 mL of cystic fluid, the level of PTH was 766.22 pg/mL (serum reference range 15–65 pg/mL). Serum PTH was slightly elevated: 88 pg/mL [chemiluminescent microparticle immunoassay (CMIA), ARCHITECT ci8200, Abbot]. Serum calcium was 2.24 mmol/L, phosphorus 0.91 mmol/L and vitamin D was under the reference range: 12.6 ng/mL. Thyroid tests were normal, thyroid peroxidase antibodies (TPOAb) and thyroglobulin antibodies (TgAb) were not detected (Table 1).

Table 1

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Baseline (reference range)</th>
<th>After 6 months</th>
</tr>
</thead>
<tbody>
<tr>
<td>PTH (serum), pg/mL</td>
<td>88 (15–65 pg/mL)</td>
<td>53.6</td>
</tr>
<tr>
<td>PTH (cystic fluid), pg/mL</td>
<td>766.22</td>
<td></td>
</tr>
<tr>
<td>Calcium, mmol/L</td>
<td>2.24 (2.1–2.55)</td>
<td>2.13</td>
</tr>
<tr>
<td>Phosphorus, mmol/L</td>
<td>0.91 (0.79–1.42)</td>
<td>0.94</td>
</tr>
<tr>
<td>25 OHD, ng/mL</td>
<td>12.6 (30–100)</td>
<td>33.2</td>
</tr>
<tr>
<td>TSH, µU/mL</td>
<td>1.75 (0.35–4.94)</td>
<td>1.3</td>
</tr>
<tr>
<td>FT4, pmol/L</td>
<td>12.2 (9.0–19.1)</td>
<td></td>
</tr>
<tr>
<td>TPOAb, IU/mL</td>
<td>2.4 (&lt; 5.61)</td>
<td>1.4</td>
</tr>
</tbody>
</table>

PTH – parathyroid hormone; TSH – thyroid stimulation hormone; FT4 – free thyroxine; TPOAb – thyroid peroxidase antibodies.

After one week ultrasound showed a residue of 2 mL of cystic fluid (Figure 2).

Cystic content was completely evacuated with repeated FNA. After the second aspiration ultrasound showed two hypoechoic solid structure lesions, 4 and 5 mm in diameter, without cystic content (Figure 3).

Radionuclide parathyroid scintigraphy 666 Tc99m methoxyisobutylisonitrile (MIBI) in 20, 60, and 120 min after applications indicated at physiological MIBI distribution (Figure 4).

After a 6-month vitamin D supplementation, serum calcium, phosphorus and PTH were normal, as well as serum vitamin D (Table 1). This finding was indicative that was a nonfunctional parathyroid cyst. Initially, the elevated PTH was a consequence of the low level of serum vitamin D.
Discussion

Parathyroid cysts are often misdiagnosed as a solitary thyroid adenoma or nodular goiter. They could be nonfunctional or functional. In both of them, the level of PTH in cystic fluid is high. Nonfunctional cysts are often located in the inferior parathyroid glands, particularly in the left inferior gland, while localization of functional cysts is unpredictable. They could be small, from some millimeters to more than 50 mm. A true ontogenetic parathyroid cyst, from the remnants of the third or fourth branchial clefts, has thin wall, lined by secretory epithelium and contains watery cystic fluid. A cyst formed by the coalescence of micro-cysts may contain hemorrhagic fluid. Cystic degeneration of parathyroid adenomas could be presented as pseudocysts. Their content is a brown fluid and they are rather functional. Just a few cases of intrathyroidal parathyroid cyst reported till now. The latest was a case of primary hyperparathyroidism due to intrathyroidal parathyroid cyst which was confirmed by founding of elevated PTH in needle aspirate. Other etiologies of parathyroid cysts have been proposed without any supporting evidence. Case report from 1965 was described that administration of high doses of calcium acetate, vitamin D2, vitamin D3 or extract of calcium acetate may cause small, from some millimeters to more than 50 mm cysts. These cysts require caution during diagnostic aspiration because of danger of hypercalcemic crises due to fine needle aspiration, which could be a life-threatening condition.

Conclusion

This case report elucidates the diagnostic pitfalls in fine needle aspiration of thyroid lesions. Also, some of thyroid cystic lesions with thin walls, and reverberation in ultrasound, have to be observed as a potential parathyroid cyst. These cysts require caution during diagnostic aspiration because of danger of hypercalcemic crises due to fine needle aspiration, which would be a life-threatening condition.

References