## LETTER TO THE EDITOR

## Papillary thyroid microcarcinoma in struma ovarii

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A 62-year-old woman presented abdominal pain and a rightsided pelvic mass. Ultrasonography showed a large, thickwalled, cystic ovarian mass with solid components. The patient underwent laparotomic monolateral annessectomy. Histology showed an endometriosic cyst, with a 17-cm struma ovarii containing a 3-mm single papillary thyroid carcinoma. Pelvic ultrasonography showed a normal left ovary, abdomino-pelvic CT scan showed only two renal cysts. The patient had no previous history of thyroid disease. Thyroid evaluation showed a 12 × 17 mm hypoechoic nodule with benign cytological findings in the right thyroid lobe. Serum TSH was 4.8 mIU/l, thyroglobulin concentration was 126 ng/nl, and both thyroglobulin and TPO antibodies were undetectable. Because of persisting slightly increased TSH levels, L-thyroxine (75 µg/die) administration was started 2 months after pelvic surgery. Serum TSH and thyroglobulin levels decreased to 0.5 mUI/l and 1.5 ng/dl, respectively.

The definition of *struma ovarii* applies to ovarian teratomas consisting of 50% or more thyroid tissue [1]. Its presentation usually resembles a solid tumor and the patients are readily addressed to surgery. Less than 5% of *struma ovarii* cases may also show hyperthyroidism. Occasionally, the diagnosis is made during the follow up of differentiated thyroid carcinoma, due to increased serum thyroglobulin levels and/or focal pelvic uptake at wholebody scan. Histological features of thyroid carcinoma are

found in 5–37% of *struma ovarii* cases (malignant *struma ovarii*). Most reported cases are classical or follicular variant of papillary carcinomas [1–4]. Individual cases of follicular and mixed (follicular and papillary) carcinoma have been described [3]. Ovarian metastasis of primary thyroid carcinoma may rarely occur [2].

The clinical behavior of malignant *struma ovarii* is based on single case reports. Metastases, mainly to the adjacent pelvic occur in up to 23% cases. Distant metastases (lungs, bone, liver, and brain) have occasionally been reported [2].

There is no consensus on the surgical approach and postoperative management of malignant struma ovarii patients. A total abdominal hysterectomy is usually performed with mono/bilateral salpingo-oophorectomy. When extra-ovarian invasion is present, omentectomy, peritoneal washings and lymph node sampling can be associated. If histology shows a malignant struma ovarii, a risk stratification similar to that used in thyroid carcinoma should be performed. PTCs larger than 20 mm, disease extension outside the struma ovarii, or aggressive histological variants should be considered high risk. In such cases total thyroidectomy and radioactive iodine ablation are required [4]. This would allow both serum thyroglobulin monitoring and radioactive iodine treatment of recurrent disease [5]. The case patient was 62 years old and presented a unique papillary microcarcinoma in her struma ovarii, with no evidence of either carcinoma in the thyroid gland or metastases at pelvic imaging studies. For these reasons, we excluded thyroidectomy. L-thyroxine therapy was administered to normalize serum TSH levels, so minimizing serum thyroglobulin fluctuations. We also recommended periodic pelvic imaging and measurements of serum thyroglobulin. After 1 year follow up, the thyroid nodule remained unchanged and both pelvic ultrasonography and

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abdomino-pelvic CT scan did not show any evidence of tumor relapse. Serum TSH and thyroglobulin levels were stable.

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