Case Report

Ovarian cystic tumor composed of Brenner tumor and struma ovarii

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Abstract: Ovarian tumor composed only of Brenner tumor and struma ovarii is very rare; only 6 cases have been reported in the English literature, to the best of the author’s knowledge. A 66-year-old woman underwent right oophorectomy because of torsion of right ovarian cyst. Macroscopically, the ovarian cyst was hemorrhagic and red. Cystic content was hemorrhagic fluid. Microscopically, the cyst walls were composed only of Brenner tumor (50% in area) and struma ovarii (50% in area). Hemorrhage and ischemic changes were seen. Other elements were not recognized. No malignant transformation was noted. These two elements were separately present, and no mergers between them were recognized. Immunohistochemically, the Brenner tumor element was positive for cytokeratins (AE1/3 and CAM5.2) and Ki67 (labeling=3%), but negative for thyroglobulin, TTF-1, p53, CA125, and vimentin. The struma ovarii element was positive for cytokeratins (AE1/3 and CAM5.2), thyroglobulin, TTF-1 and Ki67 (labeling=5%), but negative for p53, CA125 and vimentin. The findings suggest that there were cases of ovarian cyst composed only of Brenner tumor and struma ovarii, that such a case may be monodermal mature cystic teratoma or the Brenner tumor element was derived from surface epithelium in the preexisting struma ovarii, and that such a tumor manifest as cystic torsion.

Keywords: Brenner tumor, struma ovarii, ovarian cyst, torsion of ovarian cyst, histopathology, immunohistochemistry

Introduction

Ovarian tumor composed only of Brenner tumor and struma ovarii is very rare; only 6 cases have been reported in the English literature, to the best of the author’s knowledge [1-6]. Herein reported is a 66-year-old woman with Brunner tumor and struma ovarii of the right ovary. This patient presented with acute abdomen due to torsion of the right ovarian cyst.

Case report

A 65-year-old woman was found to have right ovarian cyst by gynecologic routine check. She was followed up. One year later, when she was 66-year-old, she suddenly complained of right abdominal pain. Physical examination and imaging techniques are suggestive of the torsion of the right ovarian cyst. Therefore, emergency right oophorectomy was performed.

Macroscopically, the ovarian cyst was hemorrhagic and red, and measured 7 x 6 x 6 cm (Figure 1). Cystic content was hemorrhagic fluid. Microscopically, the cyst walls were composed only of Brenner tumor (50% in area) and struma ovarii (50% in area) (Figure 2A). Hemorrhage and ischemic changes were seen (Figure 2A).

Figure 1. Macroscopic features of the right ovarian cyst. The cyst shows hemorrhagic infarction due to torsion. The cyst was red and measured 7 x 6 x 6 cm.
Coexistence of Brenner tumor and struma ovarii

The Brenner tumor element was composed of cellular islands of urothelial-like cells with coffee bean appearances embedded in fibrous stroma (Figure 2B). No atypia was recognized. The struma ovarii element was composed of normal follicular and macrofollicular thyroid tissue (Figure 2C). No atypia was recognized. No tumor formation was seen. No ground glass appearances, nuclear inclusions, or nuclear groove was seen. Other elements were not recognized. No malignant transformation was noted in these elements. These two elements were separately present, and no mergers between them were recognized (Figure 2A).

An immunohistochemical study was performed with the use of Dako’s EnVision method, as previously described [7, 8]. The immunohistochemical results are shown in Table 1. Immunohistochemically, the Brenner tumor element was positive for cytokeratins (AE1/3 and CAM5.2) (Figure 3A) and Ki-67 (labeling=3%), but negative for thyroglobulin, TTF-1, p53, CA125, and vimentin. The struma ovarii element was positive for cytokeratins (AE1/3 and CAM5.2), thyroglobulin (Figure 3B), TTF-1 (Figure 3C) and Ki-67 (labeling=5%), but negative for p53, CA125 and vimentin. A right ovarian tumor composed only of Brenner tumor and struma ovarii was pathologically diagnosed. The patient is now healthy 8 months after the operation.

Table 1. Immunohistochemical findings

<table>
<thead>
<tr>
<th>Antigen</th>
<th>Brenner tumor</th>
<th>Struma ovarii</th>
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</thead>
<tbody>
<tr>
<td>CKAE1/3</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>CKCAM5.2</td>
<td>+++</td>
<td>+++</td>
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<tr>
<td>Thyroglobulin</td>
<td>-</td>
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<tr>
<td>TTF-1</td>
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</tr>
<tr>
<td>Ki-67</td>
<td>3%</td>
<td>5%</td>
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<tr>
<td>P53</td>
<td>-</td>
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<tr>
<td>CA125</td>
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<td>Vimentin</td>
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</table>

Discussion

The present ovarian cyst was composed only of Brenner tumor and struma ovarii. Brenner tumor is relatively rare tumor, accounting 5% of all ovarian tumors [9]. Brenner tumor is classified into benign, malignant and borderline ones. The Brenner tumor element of the present tumor showed typical histologic features of this tumor. No significant atypia was seen. Immunohistochemically, p53 was negative and Ki-67 labeling...
Coexistence of Brenner tumor and struma ovarii

was very low, suggesting that the Brenner tumor component was benign. In contrast, struma ovarii belongs to monodermal teratoma [10]. Struma ovarii is rare ovarian tumor, accounting for 2.7% of all ovarian teratomas. Struma ovarii usually shows normal thyroid tissue, but may represent thyroid adenoma and carcinomas. In the present case, the thyroid tissue was normal or mildly hyperplastic, and no thyroid adenoma and carcinoma were seen. The negative p53 and low Ki-67 labeling indicates that the struma ovarii element of the current case is benign. The positive reaction of thyroglobulin and TTF-1 shows that the struma ovarii of the present tumor is true struma ovarii.

Most of Brenner tumor is solid tumor, while struma ovarii is a cystic tumor. The present tumor is entirely cystic, thus resembling struma ovarii macroscopically. The present tumor manifested with torsion of the ovarian cyst. Torsion of ovarian cyst is not rare. However, torsion of the cystic ovarian tumor composed of Brenner tumor and struma ovarii has not been reported [1-6].

The pathogenesis of this type of ovarian tumor composed of Brenner’s tumor and struma ovarii is uncertain. Brenner tumor is generally thought to be derived from surface epithelial cells [9], while struma ovarii is monodermal teratoma (germ cell tumor). Most of the literature [1-6] favors the germ cell origin of this ovarian tumor composed of Brenner tumor and struma ovarii. For example, Yoshida et al [5] reported a case of coexisting Brenner tumor and struma ovarii of the ovary. They found that both Brenner tumor and struma ovarii were immunoreactive for thyroglobulin, and suggested that the Brenner tumor element was metaplastic one and the ovarian tumor is germ cell origin. Similar case was reported by Takeuchi et al [6] who found thyroglobulin immunoreactivity in both the Brenner tumor and struma ovarii. They also speculated that this peculiar combination tumor may be derived from germ cells. In the present case, however, thyroglobulin and TTF-1 immunoreactivities were found in struma ovarii element but not in the Brenner tumor element. In addition, both the elements were histologically separated and no merges between the both elements were found. These findings may imply that both the elements were separate tumor. It is possible that the Brenner tumor is derived from surface epithelial cells in pre-existing struma ovarii in the present case. However, germ cell origin hypothesis is also possible in the present tumor. In any way, accumulation of cases of this kind of ovarian tumor is mandatory.

In conclusion, the author presented a very rare
Coexistence of Brenner tumor and struma ovarii

A case of ovarian cystic tumor composed only of Brenner tumor and struma ovarii.

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References


