Uterus-Like Ovarian Mass with Elevated Serum Levels of CA 19-9 and CA 125: A Case Report and Review of the Literature

**ARTICLE TYPE**
Case Report

**ABSTRACT** Uterus-like ovarian mass is a rare benign entity with controversial etiopathogenesis. A 37-year-old woman, gravida 2, para 2, presented with the complaint of pain in right lower abdominal quadrant. Physical evaluation revealed tenderness in this area and the ultrasonographic examination showed a right adnexal mass. Biochemical blood tests showed minimal leucocytosis and elevated serum levels of both CA 19-9: 117.79 mIU/mL (N: 0-37) and CA 125: 41.4 u/mL (N: 0-35). To exclude a malignancy, the specimen was examined with frozen sections and was subsequently diagnosed as endometriosis. Therefore, the surgical procedure was limited to right salpingo-oophorectomy. The right ovary was 6 x 4 x 2.5 cm in size and the dissected surface of ovary resembled a miniature uterus. Histologic sections as well as immunohistochemistry (smooth muscle actin, caldesmon and CD10) were consistent with the endometrial mucosal tissue that surrounded the bundles of smooth muscle.

**Key Words:** Endometriosis; CA 19-9 antigen; adenomyoma

**TURKISH ABSTRACT** Uterus-benzeri ovaryan kitle etyopatogenezi tartışmalı, nadir görülen, benign bir lezyondur. 37 yaşında kadın hasta, gravida 2, para 2, sağ alt abdominal kadran ağrısı ile başvurdu. Fizik muayene ile hassasiyet gözlenen aynı bölge, ultrason ile incelemede sağ adneksiyel kitle satıldı. Biyokimyasal incelemelerde, minimal lökositoz yansırsı serum Ca19.9: 117.79 mIU/mL (N: 0-37) ve Ca 125: 41.4 u/mL (N: 0-35) değerleri yüksekliği izlendi. Malignite dışlanmasını için olgu frozen ile inceledi ve kitle endometriozis olarak değerlendirildi. Bunun üzerine cerrahi uygulama sağ salpingoöferektomi ile sınırlandırıldı. Sağ over 6 x 4 x 2.5 cm boyutlarında olup kesi yüzü minyatür bir uterus görülmüştü. Histolojik kesitleri yansırsı düz kas aktin, kaldesmon ve CD10’u içeren immünhisstokimyasal incelemeleri, düz kas demetleri ile çevrelenmiş endometriyal mukoza ile uyumlu idi.

**Anahtar Kelimeler:** Endometriyozis; CA 19-9; adenomyoma

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**Uterus-like ovarian mass (UOM) was first described by Cozzutto in 1981.1 ‘Endomyometrioma’, ‘adenomyoma’ and ‘tumor of mullerian type’ are representative of some of the terms, reported for UOM since 1981. This lesion shows a central cavity lined by endometrial mucosa surrounded by thick muscle wall. To our knowledge, 11 cases of UOM, aged 11-54 years old, were reported.1-7 It was documented that five out of 11 cases were followed because of infertility, three had renal anomalies, two had breast cancers and three had elevated serum CA 125 levels (Table 1). UOM was also found in some other locations such as cervix uteri, broad
ligament, peritoneum, pelvic lymph nodes, scrotum, small bowel, bladder and conus medullaris.8-11 Two hypotheses have been proposed for the origin of UOM: 1- In the presence of endometriosis, through the metaplasia of endometrial stromal cells to the smooth muscle cells, 2- In the presence of the urogenital anomaly, the probability of being a part of congenital malformation.1-5,12,13

CASE REPORT

A 37-year-old woman presented with abdominal pain. The patient had allergic asthma. Physical evaluation revealed tenderness in the right lower abdominal quadrant. The ultrasonographic examination showed a right adnexial mass (6 cm) and no other gynecologic or urinary abnormalities. On abdominal computed tomography (CT), liver, spleen, gallbladder and bilateral kidneys were normal in size and location. There were no suspicious lymphadenopathies in the abdomen or retroperitoneal area. During biochemical laboratory evaluation, minimal leucocytosis (12.1 K/ul), elevated serum CA 19-9: 117.79 mIU/mL (N: 0-37) and slightly elevated serum CA 125: 41.4 u/mL (N: 0-35) levels were detected. Blood count, TSH, fT3, fT4, CEA and AFP values were within the normal ranges. To exclude a malignancy, the specimen was examined with frozen sections and diagnosed as endometriosis. Therefore, the surgical procedure was limited to right salpingo-oophorectomy. Intraoperatively, the right ovary was observed in Douglas sac adhered to the left ovary and rectal serosa. The uterus was of normal size and appearance. No foci suspicious for endometriosis were found in the pelvic or abdominal region or in another organ of genital tract. The right ovary was 6 x 4 x 2.5 cm in size and composed of two adherent nodules. One of the nodules resembled a miniature uterus since the cut section had a cavity with hemorrhagic appearance, which was surrounded with the bundles of smooth muscle (Figure 1). Microscopic evaluation revealed a cavity lined with columnar epithelium associated with endometrial stroma and surrounded by a thick muscular wall (Figure 2). Cells with atypia such as pleomorphic nuclei, hyperchromasia, nuclear vesiculation, prominent nucleolus and eosinophilic cytoplasm were observed in the wide areas of surface columnar epithelium (Figure 3). Hemosiderin-laden macrophages were present in endometrial stroma, which were surrounded by thick walls of smooth muscle bundles. From the block of this tissue, immunohistochemical stainings were applied for CD 10 (56C6 clone, 1:60 dilution, Neomarker Labvision), smooth muscle actin (1A4 clone, 1:100 dilution, Biogenex) and caldesmon (h-CD clone, 1/100 dilution, Biogenex)

<table>
<thead>
<tr>
<th>Case</th>
<th>Year, Author</th>
<th>Age</th>
<th>Urogenital System Anomaly</th>
<th>Breast Cancer</th>
<th>CA 125</th>
<th>Other</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1981, Cozutto C</td>
<td>32</td>
<td>Unilateral renal agenesis</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>2</td>
<td>1985, Peublitz-Peredo et al.</td>
<td>18</td>
<td>Double excretory system, double ureter, bifid pelvis</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>3</td>
<td>1991, Rahilly et al.</td>
<td>38</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>Endometrioid cancer in both ovary and endometrium</td>
</tr>
<tr>
<td>4</td>
<td>1994, Noel et al.</td>
<td>49</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>Cyto genetic anomaly (+), congenital anomaly (-)</td>
</tr>
<tr>
<td>5</td>
<td>1997, Mitra et al.</td>
<td>34</td>
<td>Not known</td>
<td>-</td>
<td>-</td>
<td>-</td>
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<tr>
<td>6</td>
<td>1998, Pai-Sanjay et al.</td>
<td>39</td>
<td>-</td>
<td>+</td>
<td>+</td>
<td>-</td>
</tr>
<tr>
<td>7</td>
<td>1998, Pai-Sanjay et al.</td>
<td>43</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>-</td>
</tr>
<tr>
<td>8</td>
<td>1998, Pai-Sanjay et al.</td>
<td>38</td>
<td>-</td>
<td>+</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>9</td>
<td>2005, Shutter</td>
<td>11</td>
<td>Resected right pelvic kidney</td>
<td>-</td>
<td>-</td>
<td>-</td>
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<tr>
<td>10</td>
<td>2007, Gurel et al.</td>
<td>54</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>Uterus unicornis</td>
</tr>
<tr>
<td>11</td>
<td>2007, Zennoni et al.</td>
<td>33</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>Histologic atypia</td>
</tr>
<tr>
<td>12</td>
<td>2008, Present</td>
<td>37</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>Elevated CA 19-9 and histologic atypia</td>
</tr>
</tbody>
</table>

TABLE 1: Clinicopathologic features in patients with uterus-like ovarian masses.
with use of a standard labeled streptavidin biotin peroxidase method. Positive staining with CD10 (consistent with endometrial stroma), smooth muscle actin and caldesmon (consisting with smooth muscle tissue) confirmed the histological diagnosis. In one pole of this nodule, the bundles of smooth muscles were intermingling with the spindle cells of the ovarian stroma of the other nodule, composed of ovarian stroma with follicular cysts and hemorrhagic corpus luteum. No abnormality was found in the sections of right tubal tissues. The patient had been followed for two years due to infertility. Meanwhile, the only pregnancy achieved through in vitro fertilization was diagnosed as ‘blighted ovum’, thus finally curettage was applied to the patient. There has been no pregnancy since then.

**DISCUSSION**

The etiopathogenesis of UOM is controversial. There are two main hypotheses. Cozzutto suggests that metaplasia of ovarian stroma in the background of endometriosis may explain the cause of the lesion.\(^1\) This hypothesis has been supported by several other cases.\(^3,4,13\) On electron microscopic evaluation, Cozzutto observed myofibroblasts as well as smooth muscle cells, and consequently, he suggested that myofibroblasts could be a transitional cell during the metaplastic changes. Supporting this view, Scully suggested that pressure of an expanding endometriotic cyst could trigger a metaplasia through smooth muscle cells in the cells surrounding the cyst.\(^14\) He also indicated that smooth muscle fibers could be seen in some of other ovarian lesions such as leiomyoma, mucinous cystadenoma and ovarian stromal hyperplasia. In cases with absence of demonstrable endometrial mucosa due to bleeding or pressure, differentiation from ovarian stromal hyperplasia and leiomyoma should be considered. Although the period which triggers the metaplasia cannot be clearly determined, there is some evidence to support the role of ovarian hormones. Breast cancers were determined in two out...
of three cases reported by Pai et al.\textsuperscript{4} Endometrioid carcinoma was identified in both ovary and uterus as well as endometriosis in the case of Rahilly et al.\textsuperscript{13} Peterson et al. reported a case with precocious puberty and this points to the role of estrogen in formation of UOM.\textsuperscript{9}

The other hypothesis proposed by Rosen is that this lesion may be a part of urogenital congenital malformation as a fusion defect of Mullerian duct or a real partial duplication of Mullerian system.\textsuperscript{12} The detection of renal agenesis in the case of Cozzutto retrospectively and the presence of urinary anomaly in two other cases also support this hypothesis.\textsuperscript{2,5} The hypothesis of metaplasia seems more likely in the cases without any uterus and/or renal congenital anomaly.\textsuperscript{3,4,13}

It is known that the cases with endometriosis should be evaluated carefully by the pathologists in terms of the presence of cytologic atypia. Epithelial abnormalities (hyperplasia or atypia) have been considered as a step in the progression to malignancy.\textsuperscript{15,16} Endometrioid carcinomas and clear cell carcinomas are the two most common tumors developed in ovarian endometriosis.\textsuperscript{15,16} Rarely, various tumors in the background of ovarian endometriosis have been reported such as ovarian serous cystadenoma of low malignant potential, benign and malignant mucinous tumors, squamous cell carcinoma, granulosa cell tumor, endometrioid stromal sarcoma, malignant mesodermal mixed tumor and adenosarcoma.\textsuperscript{16,17} Atypical cells display eosinophilic cytoplasm, large hyperchromatic or pale pleomorphic nuclei, increased nuclear to cytoplasmic ratio, cellular crowding, stratification or tufting. Zannoni et al. reported the first case of ovarian endomyometrioma with histologic atypia (hyperchromasia, pleomorphic nuclei, eosinophilic cytoplasm) and foci of mucinous metaplasia on cavity lined by columnar epithelium.\textsuperscript{18} In this study, authors have mentioned that the simultaneous presence of endometrial epithelium with histologic atypia and metaplasia supported the metaplastic theory of UOM. They have suggested the term ‘endomyometrioma with atypias’ for these cases. Similarly, our case showed epithelial tufting with pleomorphic nuclei, hyperchromasia, nuclear vesiculation, prominent nucleolus and eosinophilic cytoplasm.

High serum levels of both CA 19-9 and CA 125 have been reported in some benign lesions such as leiomyoma, endometriosis, teratoma of the fallopian tube and tuba-ovarian abscesses.\textsuperscript{19-22} In endometriosis, serum level of CA 125 and CA 19-9 is expected to be no more than 100 IU/ml and 1000 IU/ml, respectively. However, very high serum levels of CA 125 and CA 19-9 (CA 125: 9537 IU/ml, CA 19-9: 15.653 IU/ml) was recorded in a patient with ruptured ovarian endometrioma.\textsuperscript{20} The mechanisms that may elevate serum levels of CA 125 and CA 19-9 in endometriosis are not yet clearly understood; however, the diffusion of cyst fluid through the peritoneal surface into the circulation is considered in some cases.\textsuperscript{20}

We believe that UOM was developed due to the metaplasia of ovarian endometrioma in our patient. First, no urogenital abnormalities were found on radiologic evaluations. Second, high serum levels of CA 19-9 and CA 125 were associated with the presence of ovarian endometriosis. Third, the presence of histologic atypia in columnar epithelium of endometrial mucosa supports the metaplastic theory.

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REFERENCES