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Case Report

Primary malignant melanoma of the ovary arising in a cystic teratoma; case report and review of the literature

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Abstract

Primary malignant melanoma of the ovary arising in a cystic teratoma is extremely rare. There is no melanocytic component in the normal ovarian tissue. However, melanocytes may be present in a cystic teratoma of the ovary, especially at the basal layer of squamous epithelium. We report a case of primary malignant melanoma of the ovary arising in a cystic teratoma. We demonstrate junctional activity of melanocytes showing malignant transformation with both radial and vertical growth phases, a feature diagnostic for primary malignant melanoma but not always demonstrated in such cases. We compiled and reviewed twelve cases that are also able to demonstrate junctional activity.

Keywords: Primary Melanoma, teratoma, ovarian cancer, junctional activity

Introduction

An ovarian malignant melanoma is rare with unfavorable prognosis, usually owing to metastases [1]. Primary malignant melanomas are even more rare in the ovary [2,3] and its diagnosis requires fulfillment of established criteria. Boughton et al. [4] and Cronje and Woodruff [5] proposed the following criteria for the diagnosis of primary ovarian melanoma: 1) no other possible sites of a primary tumor; 2) unilateral tumor within the ovarian teratoma; 3) good correlation of the patient’s age and symptoms with those of well-documented cases in the literature; and 4) demonstration of junctional activity (desirable but not necessary for diagnosis). A point of controversy for diagnosing the primary nature of this malignancy is whether or not it is necessary to fulfill the last criteria—demonstration of junctional activity. Indeed, establishing junctional activity goes further toward providing evidence that the malignant cells originated from this junctional activity and many cases reported as primary malignancies without junctional activity may actually be metastatic disease, thus portending higher mortality risk for the patient. In this report, we report a case of primary malignant melanoma of the ovary arising in a cystic teratoma. We also compare and contrast the metastatic and mortality rate of these cases showing junctional activity with those of cases without junctional activity in specimens reported as primary malignant melanoma of ovary.

Case synopsis

We report a case of a 38-year-old female who presented with a 2 month history of lower abdominal pain. A 12 x 13 cm left adnexal mass was discovered on CT for which the patient was treated with left salpingo-oophorectomy. The specimen was received in one container fixed in formalin labeled “left tube and ovary” and consisted of an ovary, tan to tan-yellow, with multiloculated
cysts that had been previously opened. It measured 12 x 7 x 3.5 cm. There was an overlying segment of fallopian tube that measured 6.5 cm in length and 1.1 cm in average diameter. The ovary showed some intact cystic structures; some were filled with keratinosebaceous material and others with clear watery fluid. Solid areas, on serial sectioning, revealed a smooth tan-brown cut surface. Some cystic structures were lined by flattened squamous epithelium with pilosebaceous structures, respiratory epithelium with underlying mucinous glands, and a solid area containing cartilage and thyroid tissue (Figure 1). The findings were diagnostic for a mature cystic teratoma.

Figure 1. Mature cystic teratoma consisting of cysts lined by flattened squamous epithelium and respiratory epithelium and containing cartilage and thyroid tissue (H&E x 217)

Junctional activity of melanocytes was observed within the squamous epithelium. There was a 2 cm tumor nodule in the cyst wall (Figure 2A). The cytomorphical features of the tumor cells included a moderate amount of eosinophilic cytoplasm with round to ovoid pleomorphic nuclei, which had a vesicular chromatin pattern and prominent nucleoli (Figure 2B). The tumor cells stained positively for S100, Melan-A, HMB 45, and vimentin, which supported a diagnosis of malignant melanoma. The tumor appeared continuous with overlying squamous epithelium, which contained melanocytes in the junctional areas. Junctional melanocytes showed evidence of malignant transformation with radial and vertical growth phases, which were highlighted by immunohistochemical stains for Melan-A and HMB-45 (Figure 3). Vertical grown phase melanoma with formation of a tumor nodule in the cyst wall strongly supported the primary nature of this malignant melanoma of ovary.

Figure 2. (A) A 2 cm tumor nodule in the cyst wall (H&E stain x 217). (B) Higher magnification showing epithelioid tumor cells and junctional activity of melanocytes in the overlying squamous epithelium (H&E stain x 650)
Other possible primary sites of melanoma, such as the skin, intestine, and eye were examined but no pigmented lesions were found in this patient. The malignant melanoma arose in a unilateral ovarian teratoma, thus fulfilling the main criteria as established by Boughton et al. [4] and Cronje and Woodruff et al [5] for diagnosing primary malignant melanoma of the ovary occurring in association with a dermoid cyst.

**Discussion**

Lee et al. [6] recently reviewed and compiled 19 cases of primary malignant melanoma of the ovary that matched some of the criteria established by Boughton et al. [4] and Cronje and Woodruff [5]. Amongst these cases, only five cases were identified as establishing junctional activity and thus fulfilling all the established criteria. We additionally found the remaining 5 more cases in the literature whereby junctional activity was also established (Table I), thus completing a comprehensive review of available cases of this entity. The unique histology we show for our case thoroughly establishes this junctional activity as being the origin of malignant cells in that we demonstrate radial and vertical growth phases of junctional melanocytes with progression into malignant melanoma in the cyst wall of cystic teratoma of ovary.

Compared to the cases whereby junctional activity could not be observed (see table II), cases exhibiting junctional activity (see Table I in this report) generally had less occurrence of distant metastases; 4 of the 12 cases with junctional activity had distant metastases, fewer than the 8 of the 13 cases without junctional activity previously reported. Likewise, 4 of the 12 (33%) with junctional activity were dead of disease at follow-up, again a lower frequency than the 8 from the 13 cases (62%) without junctional activity. The improved mortality in cases with junctional activity may suggest that the observation of junctional activity means that the disease was caught at an earlier stage when junctional activity in the primary ovarian malignant melanomas had not yet been destroyed by tumor invasion. An alternative explanation is that several of the cases whereby junctional activity could not be established are not indeed primary malignancies. However, owing to the limited number of cases reported thus far, it remains to be seen if lower metastatic and mortality rate in cases showing junctional activity continues to be observed as more cases become available.

**Table I. Clinical-pathologic features of previous cases of primary malignant melanoma of the ovary with established junctional activity.**

<table>
<thead>
<tr>
<th>Author</th>
<th>Age (yr)</th>
<th>Distant metastasis</th>
<th>Follow-up time</th>
<th>Patient outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nanbu et al.⁷</td>
<td>47</td>
<td>No</td>
<td>ND*</td>
<td>NED⁺</td>
</tr>
<tr>
<td>Parekh et al.⁸</td>
<td>26</td>
<td>Yes</td>
<td>8 months</td>
<td>DOD⁷⁺</td>
</tr>
<tr>
<td>Marcial-Rojas et al.⁹</td>
<td>77</td>
<td>Lost to followup</td>
<td>Lost to followup</td>
<td>NED</td>
</tr>
<tr>
<td>Author</td>
<td>Age (yr)</td>
<td>Distant metastasis</td>
<td>Follow-up time</td>
<td>Patient outcome</td>
</tr>
<tr>
<td>--------------------------------</td>
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<td>----------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>Leo et al.</td>
<td>72</td>
<td>No</td>
<td>8 mo</td>
<td>NED</td>
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<tr>
<td>Park et al.</td>
<td>49</td>
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<td>Not mentioned</td>
<td>Not mentioned</td>
</tr>
<tr>
<td>O’Gorman and Olaitan.</td>
<td>49</td>
<td>No</td>
<td>ND</td>
<td>NED</td>
</tr>
<tr>
<td>Vigliani et al.</td>
<td>67</td>
<td>Yes</td>
<td>3 mo</td>
<td>DOD</td>
</tr>
<tr>
<td>Takubo et al.</td>
<td>65</td>
<td>No</td>
<td>24 mo</td>
<td>NED</td>
</tr>
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<td>Cronje and Woodruff.</td>
<td>74</td>
<td>Yes</td>
<td>18 mo</td>
<td>DOD</td>
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<tr>
<td>Boughton et al.</td>
<td>27</td>
<td>No</td>
<td>2 yr</td>
<td>NED</td>
</tr>
<tr>
<td>Liberati et al.,</td>
<td>74</td>
<td>No</td>
<td>7 months</td>
<td>NED 7 months</td>
</tr>
<tr>
<td>Gregg et al.</td>
<td>56</td>
<td>Yes</td>
<td>4 months</td>
<td>DOD</td>
</tr>
</tbody>
</table>

*ND-(Not determined)  
+NED-(No evidence of disease)  
++DOD-(Dead of Disease)

**Table II. Clinical-pathologic features of previous cases of primary malignant melanoma of the ovary where junctional activity could not be determined or established (Table and information adapted from Lee et al.)**
<table>
<thead>
<tr>
<th>Gök et al.</th>
<th>67</th>
<th>No</th>
<th>7 mo</th>
<th>DOD</th>
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<tr>
<td>Tsukamoto et al.</td>
<td>46</td>
<td>Yes</td>
<td>12 mo</td>
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<td>Gao et al.</td>
<td>53</td>
<td>Yes</td>
<td>6 mo</td>
<td>DOD</td>
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<tr>
<td>(A) Lee et al.</td>
<td>46</td>
<td>Yes</td>
<td>2 mo</td>
<td>DOD</td>
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<tr>
<td>Choi et al.</td>
<td>45</td>
<td>Yes</td>
<td>37 mo</td>
<td>NED</td>
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<td>(B) Lee et al.</td>
<td>71</td>
<td>Yes</td>
<td>5 month</td>
<td>DOD</td>
</tr>
</tbody>
</table>

*ND-(Not determined)
+NED-(No evidence of disease)
++DOD-(Dead of Disease)

References


