Case Report

Metastasizing Malignant Granular Cell Tumor (Abrikossoff Tumor) of the Anterior Abdominal Wall, with Prolonged Survival

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Malignant granular cell tumor (MGCT) is a rare high-grade mesenchymal tumor of Schwann cell origin. MGCTs commonly affect thigh, extremity, and trunk; however, involvement of the abdominal wall is exceedingly rare [12]. Herein, we report a case of MGCT arising in the anterior abdominal wall, who underwent surgical excision with wide resection margin and received chemotherapy, but developed massive metastases in the lungs and in the right inguinal lymph nodes, with prolonged survival for 11 years, with a review of the relevant literature.

1. Introduction

Granular cell tumors (GCTs), previously known as granular cell myoblastoma, are uncommon mesenchymal neoplasms of Schwann cell origin, composed of cells with granular eosinophilic cytoplasm, usually presenting as asymptomatic nodule, affecting adults, with women affected twice as men [1]. They occur in the dermis or subcutaneous tissues [2], oral mucosa including tongue [2, 3], gum (congenital epulis) [4], breast [5], vulva [6], and gastrointestinal and biliary tracts [7, 8]. In 5% of the reported cases, the tumor was multiple [2].

On the other hand, malignant granular cell tumors (MGCTs) are exceedingly rare high-grade malignant mesenchymal neoplasms, representing only 1%-2% of all GCTs [9]. In a study of 6,412 soft tissue sarcomas, 26 cases (0.4%) were diagnosed as MGCT [10], with 157 cases in various anatomical sites on record, documented in a review in 2018 [11]. When the tumor has malignant cytological features or when a benign-appearing tumor causes metastasis or death, it is considered malignant.

Most MGCTs commonly affect lower extremities, especially thighs and the trunk; however, involvement of the subcutaneous tissue of the abdominal wall is exceedingly rare [12].

Herein, we report a case of MGCT arising in the anterior abdominal wall, who underwent surgical excision with wide resection margin and received chemotherapy, but developed massive metastases in the lungs and in the right inguinal lymph nodes, with prolonged survival for 11 years, with a review of the relevant literature.

2. Case Report

A 50-year-old female presented in July 2007 to a peripheral general hospital with progressively enlarging mass in the anterior abdominal wall of one-year duration. On physical examination, she was well built with normal vital signs. A 7 x 6 cm firm irregular subcutaneous mass was felt in the right lower para-umbilical area of the anterior abdominal wall. On palpation, the abdomen was soft and lax with no intra-abdominal masses. There were no palpable lymph nodes. Her cardiac and respiratory examinations were unremarkable. Computerized Tomography (CT) scan of the abdomen revealed a mass within the abdominal wall. Her laboratory
investigations were within normal limits. A plain chest X-ray was unremarkable.

Wide local excision with safety margins of the tumor was done. The postoperative period was uneventful, and she was discharged on the second postoperative day in a satisfactory condition.

Gross examination of the specimen showed an ellipse of skin that measured 13 x 10 cm with subcutaneous fatty tissue measuring 11 cm in thickness. On cut section, a well-defined nonencapsulated grayish yellow mass measuring 7 x 6 cm with foci of necrosis was seen. No skin infiltration or abdominal skeletal muscle involvement was noted.

Microscopic examination revealed well-defined but nonencapsulated subcutaneous mesenchymal neoplasm composed of irregular islands of large cells with mostly round to oval pleomorphic vesicular nuclei, with markedly granular cytoplasm, and several large eosinophilic globules (Figure 1(a)). The nuclear cytoplasmic ratio was variable, with several cells having large nuclei. The tumor nests were surrounded by bands of fibrous connective tissue (Figure 1(b)).

Many foci of tumor necrosis and scattered mitotic figures, 4 per 10 HPF at a magnification of 200, were identified (Figures 1(c) and 1(d)). Margins of resection were free of tumor.

Immunohistochemical stains show the tumor cells to be strongly positive for vimentin, S100 and CD68 (Figures 2(a), 2(b), and 2(c)). Stains for cytokeratin (CK), smooth muscle actin (SMA), desmin, and myogenin were all negative. Ki-67 proliferative index was 3-5% (Figure 2(d)). PAS stain highlighted the granules and the eosinophilic globules within the cytoplasm of tumor cells.

The patient was referred in September 2007 to the Oncology Clinic at our hospital for regular follow-up. A chest CT done on May 2008 showed small right lung nodules measuring 0.48 cm of uncertain significance. In April 2010, a follow-up of her chest CT showed bilateral multiple lung nodules; the largest was in the right upper lobe measuring 1.5 x 1 cm. Pelvic CT on February 2011 revealed a large right inguinal mass measuring 3.2 x 2.1 cm. Palliative chemotherapy was planned for the patient. She was started on doxorubicin and ifosfamide for 6 cycles; the last cycle was in October 2011 and then she refused to continue treatment.

In September 2012, FNAC of the inguinal lymph node showed numerous clusters and single cells with abundant granular cytoplasm and large pleomorphic nuclei.

In December of 2015 she was started on docetaxel single agent for 6 cycles; the last cycle was completed by April 2016. In January 2018, her last follow-up chest CT revealed an increase in the size of all metastatic nodules. The largest nodule was in the right upper lobe and measured up to 5.2 x 4.9 cm. It had increased in size since July 2017 when it measured at that time 4.4 x 4.3 cm (Figure 3(a)).
(a) Tumor cells and intervening stromal cells in connective tissue between tumor nests are positive for vimentin; x20

(b) Tumor cells are strongly positive for S100 protein; x20

(c) Tumor cells are strongly positive for CD68; x20

(d) Ki-67 stain featuring scattered positive nuclear staining in 4% of tumor cells; x40

Figure 2

(a) Coronal CT-scan of thorax featuring bilateral lung metastases (Cannon balls) with calcification

(b) Coronal CT-scan of pelvic region showing markedly enlarged right inguinal lymph node

Figure 3

the inguinal lymph node increased in size up to 8.24 x 7.89 cm (Figure 3(b)). The patient is still alive 11 years after her initial surgery and is followed up at the Oncology Clinic.

3. Discussion

Alexei I Abrikossoff, a Russian pathologist, described the first case of benign GCT in the skeletal muscle of the tongue in 1926. He thought that these tumors were of skeletal muscle cells origin and called them “myoblastoma” [35]. Although the origin of these tumors is still uncertain, further research has suggested that it is mostly of Schwann cell origin [36, 37].

There are several large series of GCTs. In a study of 95 patients from Memorial Hospital in New York, with their age ranging from 11 months to 68 years, an average of 38.1 years, there were 92 benign GCTs and three MGCTs. There
were 31 males and 64 females with a male to female ratio of 1:2. The average size of the tumors was 1.85 cm. Multiple lesions were encountered in 8 patients. The tumor distribution was ubiquitous in many anatomic locations, including two in the abdominal wall [38]. A study of 52 GCTs in 42 patients from the Medical College of Virginia revealed an average age of 37 years, with females constituting 67% of the patients, and a dominance of the tumor in African American patients (74%). There was a high rate of multicentricity (14%). The commonest sites of involvement were the tongue (13 cases), the breast (7 cases), anogenital region (7 cases), the upper extremity (5 cases), and the abdominal wall (5 cases). These cases are reported without data on the age and sex of the patients or the size of the tumor [39]. Another clinicopathologic study of 50 cases of GCTs revealed a mean age of 38.6 years and a mean size of 2.1 cm. Most patients (64%) were females. There was a predilection for the upper extremities and the upper trunk [40].

On the other hand, MGCT is a rare and aggressive tumor, representing less than 2% of all GCT and 0.2% of all soft tissue sarcomas [3, 9]. Ordonez reviewed 43 cases of MGCTs published by 1998. There were 14 males and 29 females with male to female ratio of 1:2. The age ranged from 23 to 82 years, with a mean of 50 years. The size of the tumor ranged from 2 to 17 cm, with a mean of 7 cm. Three patients had multiple tumors. The commonest site of involvement was the lower limb (11 cases) [41]. In a review of 113 cases of MGCTs from Surveillance, Epidemiology, and End Results (SEERS) database, the median size of the tumor was 4 cm [42], unlike other benign GCTs with median size of less than 3 cm [1]. Most MGCTs were located within the soft tissues and the skin. The median age was 54 years, and 77.0% were female [42]. In a recent review of the literature of 157 reported cases of MGCT, the median tumor size was 6 cm. The median age was 51 years. There were 99 females (63% of total cases) and 58 males (37%), with female to male ratio of 1.7:1. The commonest sites of the tumor were the trunk and the thighs [11]. Fanburg-Smith et al. reviewed 46 cases of MGCTs from the Armed Forces Institute of Pathology, with a mean age of 40 years. There were 32 females, 12 males, and 2 patients with unknown age. Male to female ratio was 1:2.7. The upper extremity was the commonest site affected (40%), followed by trunk (38%) and the lower extremity (22%) [9].

Histologically, sheets of polygonal to spindly cells were shown, with an abundant granular eosinophilic cytoplasm with scattered globules; both are positive for Periodic Acid-Schiff (PAS) stain [10]. Immunohistochemical stains of all GCTs are essentially the same, which are positive for S100, vimentin, NSE, and to lesser extent for CD68 and CD57 [1].

The differential diagnosis of MGCT includes pleomorphic rhabdomyosarcoma, granular leiomyosarcoma, and alveolar soft part sarcoma (ASPS) [46]. MGCTs are negative for muscle markers, and rhabdomyosarcomas are almost always negative for S100 protein. In a review of nine smooth muscle tumors with granular cell changes, including four malignant ones, all tumors were S100 protein negative and were positive for anti-muscle actin antibody (HHF-35) [47]. Macareno et al. reported a case of leiomyosarcoma of the saphenous vein with granular cell changes. They stated that other neoplasms can exhibit granular cell change including dermatofibrosarcoma protuberans, angiosarcoma, atypical fibroxanthoma, dermatofibroma, basal cell carcinoma, and melanocytic nevi. The granular cells in their cases stained negative for S100 protein, but positive for CD8. The spindle cell component of the tumor stained positive for smooth
Table 1: Reported cases of benign, atypical, and malignant granular cell tumor of the abdominal wall.

<table>
<thead>
<tr>
<th>No.</th>
<th>Authors Year (Reference)</th>
<th>Age</th>
<th>Gender</th>
<th>Size Cm.</th>
<th>Type</th>
<th>Location in Abdominal Wall</th>
<th>Metastasis, outcome and remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Cave et al Case 3 1955 [13]</td>
<td>6.5</td>
<td>Female</td>
<td>2</td>
<td>Benign</td>
<td>Dermis and subcutaneous</td>
<td>Not applicable. Patient had one tumor in right submental area of head and one tumor in the abdominal wall.</td>
</tr>
<tr>
<td>2</td>
<td>Baraf and Bender Case 1 1964 [14]</td>
<td>29</td>
<td>Male</td>
<td>0.5-2.5</td>
<td>Benign</td>
<td>Dermis and subcutaneous</td>
<td>Not applicable. Patient had multiple cutaneous GCTs (21 tumors) at various sites including abdominal wall</td>
</tr>
<tr>
<td>3</td>
<td>Baraf and Bender Case 2 1964 [14]</td>
<td>18</td>
<td>Male</td>
<td>0.5-3</td>
<td>Benign</td>
<td>Dermis and subcutaneous</td>
<td>Not applicable. Patient had multiple cutaneous GCTs (15 tumors) at various sites, including abdominal wall</td>
</tr>
<tr>
<td>4</td>
<td>Baraf and Bender Case 3 1964 [14]</td>
<td>33</td>
<td>Male</td>
<td>0.5-1.5</td>
<td>Benign</td>
<td>Dermis and subcutaneous</td>
<td>Not applicable. Patient had multiple cutaneous GCTs (about 40 tumors) within ten years at various sites</td>
</tr>
<tr>
<td>5</td>
<td>Gorelkin et al. 1978 [15]</td>
<td>58</td>
<td>Female</td>
<td>8</td>
<td>Benign</td>
<td>Intramuscular</td>
<td>Not applicable</td>
</tr>
<tr>
<td>6</td>
<td>Apisarnthanarax Case 6 1981 [16]</td>
<td>43</td>
<td>Female</td>
<td>3</td>
<td>Benign</td>
<td>Subcutaneous</td>
<td>Not applicable</td>
</tr>
<tr>
<td>7</td>
<td>Apisarnthanarax Case 7 1981 [16]</td>
<td>36</td>
<td>Female</td>
<td>2.5</td>
<td>Benign</td>
<td>Subcutaneous</td>
<td>Not applicable</td>
</tr>
<tr>
<td>8</td>
<td>Kucan et al 1982 [17]</td>
<td>8</td>
<td>Male</td>
<td>Small, Size not stated</td>
<td>Benign</td>
<td>Dermis and Subcutaneous</td>
<td>Metastases to lungs and lymph nodes. Dead of disease at 96 months post diagnosis.</td>
</tr>
<tr>
<td>9</td>
<td>Geisinger et al. 1985 [19]</td>
<td>69</td>
<td>Female</td>
<td>Not stated</td>
<td>Malignant</td>
<td>Not stated</td>
<td>Not applicable. Patient had total of 57 cutaneous GCTs excised during 13-year period from age 10 to 23 years at various sites including abdominal wall</td>
</tr>
<tr>
<td>10</td>
<td>Rifkin et al. Case 1 Mother 1986 [20]</td>
<td>23</td>
<td>Female</td>
<td>Not stated</td>
<td>Benign</td>
<td>Subcutaneous</td>
<td>Not applicable. Patient had tracheal GCT with a recurrence, two abdominal wall tumors and a perianal tumor</td>
</tr>
<tr>
<td>11</td>
<td>Rifkin et al. Case 2 Son 1986 [20]</td>
<td>6</td>
<td>Male</td>
<td>Not stated</td>
<td>Benign</td>
<td>Subcutaneous</td>
<td>Not applicable. Patient had multiple subcutaneous GCTs in fingers, arms, neck, buttock, abdominal wall, and the largest involved the clitoris (4.5x5.5 cm)</td>
</tr>
<tr>
<td>12</td>
<td>Rubenstein et al Case 1 1987 [21]</td>
<td>7</td>
<td>Female</td>
<td>3</td>
<td>Benign</td>
<td>Subcutaneous</td>
<td>Not applicable. Patient had one tumor in right submental area of head and one tumor in the abdominal wall.</td>
</tr>
</tbody>
</table>
Table 1: Continued.

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<th>Type</th>
<th>Location in Abdominal Wall</th>
<th>Metastasis, outcome and remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>13</td>
<td>Khansur et al 1987 [22]</td>
<td></td>
<td>Male</td>
<td></td>
<td>Malignant</td>
<td>Not stated</td>
<td>Presented with systemic metastasis in the liver and lung which rapidly progressed and caused death in five months. Son had MGCT of chest wall.</td>
</tr>
<tr>
<td>14</td>
<td>Vamsy et al. 1992 [23]</td>
<td>30</td>
<td>Female</td>
<td>10x8x5</td>
<td>Malignant</td>
<td>Intramuscular</td>
<td>None. Alive at 24 months postop.</td>
</tr>
<tr>
<td>15</td>
<td>Menaker and Sanger 1997 [24]</td>
<td>50</td>
<td>Female</td>
<td>4x2x2</td>
<td>Atypical (Uncertain Malignant Potential)</td>
<td>Subcutaneous</td>
<td>Underwent wide local excision. No recurrence or metastasis at 16 months postop.</td>
</tr>
<tr>
<td>18</td>
<td>Joshi and Aqel 2003 [25]</td>
<td>37</td>
<td>Male</td>
<td>2x1.6</td>
<td>Benign</td>
<td>Intramuscular</td>
<td>Not applicable</td>
</tr>
<tr>
<td>19</td>
<td>Chelly et al. 2005 [18]</td>
<td>67</td>
<td>Female</td>
<td>6x4x3</td>
<td>Malignant</td>
<td>Subcutaneous and intra-muscular</td>
<td>None. Patient died due to pulmonary embolism three months postop.</td>
</tr>
<tr>
<td>20</td>
<td>An et al. 2007 [26]</td>
<td>44</td>
<td>Female</td>
<td>4</td>
<td>Benign</td>
<td>Intramuscular</td>
<td>Not applicable</td>
</tr>
<tr>
<td>21</td>
<td>Chaudhry et al. 2008 [27]</td>
<td>70</td>
<td>Female</td>
<td>10</td>
<td>Benign</td>
<td>Intramuscular</td>
<td>Not applicable. Alive and well 5 months postop.</td>
</tr>
<tr>
<td>22</td>
<td>Panunzi et al. 2012 [28]</td>
<td>29</td>
<td>Female</td>
<td>1.5</td>
<td>Benign</td>
<td>Adherent to muscle</td>
<td>Not applicable</td>
</tr>
<tr>
<td>24</td>
<td>Toelen et al. 2013 [30]</td>
<td>68</td>
<td>Female</td>
<td>3</td>
<td>Benign</td>
<td>Subcutaneous</td>
<td>Not Applicable</td>
</tr>
<tr>
<td>25</td>
<td>Porta et al. 2015 [31]</td>
<td>45</td>
<td>Female</td>
<td>3</td>
<td>Benign</td>
<td>Intramuscular</td>
<td>Not Applicable</td>
</tr>
<tr>
<td>26</td>
<td>Liu et al. 2015 [32]</td>
<td>66</td>
<td>Male</td>
<td>3x2.8</td>
<td>Malignant</td>
<td>Dermis and Subcutaneous</td>
<td>No recurrence or metastasis at 12 months postop.</td>
</tr>
<tr>
<td>27</td>
<td>Yoon et al. 2016 [33]</td>
<td>66</td>
<td>Male</td>
<td>4.5x3.4x3</td>
<td>Malignant</td>
<td>Intramuscular</td>
<td>No metastasis or recurrence at 30 months.</td>
</tr>
<tr>
<td>28</td>
<td>Imanishi et al. Case 13 2016 [34]</td>
<td>48</td>
<td>Male</td>
<td>8</td>
<td>Malignant</td>
<td>Not stated</td>
<td>Local recurrence at 27 months. Mets to lung and bone at 26 months. Died at 87 months postop.</td>
</tr>
</tbody>
</table>
muscle actin [48]. Alveolar soft part sarcoma has a distinctive organoid pattern and lacks S100 protein positivity [46].

MGCTs have a poor prognosis with 32% local recurrence and 50% metastatic rate [9]. It can metastasize several years following the initial surgical excision, as in our case, and can recur before that. However, benign and atypical GCTs have favorable outcome with no potential for metastasis [1, 9]. Common metastatic sites for MGCT are lymph nodes, lungs, liver, and bones. It has 39% mortality rate in 3-year interval [9]. Our patient had prolonged survival although she had significant metastatic disease. It has been established that older age group, larger tumor size, local recurrence, metastasis, Ki67 >10%, and p53 immunoreactivity are all adverse prognostic factors [9].

The treatment of choice is complete surgical resection with safe margins and regional lymph node dissection. Marked resistance of the tumor to radiotherapy and chemotherapy made them of low value [8, 9]. Some authors stated that even when negative margins are not obtained, the prognosis is still favorable [9]. Follow-up guidelines for MGCTs are needed, although annual follow-up is advised to rule out local recurrence or metastatic spread [7].

4. Conclusion

Our patient had prolonged survival in spite of the presence of metastatic tumor deposits in both lungs and right inguinal lymph node. She received chemotherapy but it is not clear whether this contributed to her long survival or not. The presence of metastases is currently considered as the only unequivocal sign of true malignancy. This opinion is supported by some degree of overlap in clinicopathologic data between benign and malignant GCTs [1, 2]. Only a large tumor size (> 5 cm) is considered a clinical sign of potential malignancy.

Disclosure

This case was presented as a poster form at the 32nd Congress of the International Academy of Pathology held at King Hussein Bin Talal Convention Center by the Dead Sea in Jordan, on October 14-18, 2018.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this article.

Table 1: Continued.

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<th>Authors</th>
<th>Year (Reference)</th>
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<th>Gender</th>
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<th>Type</th>
<th>Location in Abdominal Wall</th>
<th>Metastasis, outcome and remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>29</td>
<td>Alnashwan et al.</td>
<td>Current case 2019</td>
<td>50</td>
<td>Female</td>
<td>7x6</td>
<td>Malignant</td>
<td>Subcutaneous</td>
<td>Metastases to both lungs and right inguinal lymph node. Alive at 132 months postop.</td>
</tr>
</tbody>
</table>

References


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