Large uterine leiomyoma presenting as pseudo-Meigs’ syndrome with an elevated CA 125: Case report and literature review

Jake Alan Gibbons

ABSTRACT

Introduction: Uterine leiomyomas presenting as pseudo-Meigs’ syndrome with an elevated CA 125 are rare and pose a diagnostic challenge for physicians. Case Report: A 35-year-old G0P0 presented to the emergency department with a chief complaint of abdominal swelling and weight gain beginning one month ago. Laboratory work was significant for an elevated CA 125. She was found to have a large mass originating from her uterus. Following resection of the mass, her symptoms resolved. Pathology of the mass came back as a uterine leiomyoma. Conclusion: Generally, findings of ascites, a pelvic mass, and an elevated CA 125 in a female suggests a diagnosis of ovarian cancer. However, as demonstrated by our case and others, benign causes such as Meigs’ syndrome and pseudo-Meigs’ syndrome should also be considered. Surgery is the mainstay of treatment in pseudo-Meigs’ syndrome. Resolution of the ascites and hydrothorax occurs spontaneously following resection of the tumor.

Keywords: Ascites, CA 125, Gynecology, Leiomyoma, Pseudo-Meigs’ syndrome

INTRODUCTION

Meigs’ syndrome is the triad of a benign ovarian fibroma or thecoma combined with ascites and hydrothorax [1]. A clinically similar yet distinct pathological entity is pseudo-Meigs’ syndrome. Like Meigs’ syndrome, pseudo-Meigs’ syndrome presents with ascites and hydrothorax. However, unlike Meigs’ syndrome, pseudo-Meigs’ syndrome is associated with non-thecoma, non-fibroma ovarian tumors or uterine leiomyomas [1]. Ovarian tumors reported in pseudo-Meigs’ syndrome include: strumaovarii tumors, germ cell tumors, ovarian metastases from gastric or colon cancer, and serous or mucinous cystadenomas [2]. The uterine leiomyoma, however, is the most commonly reported tumor in cases of pseudo-Meigs’ syndrome [3]. Leiomyomas have an incidence of 20-50% in females over the age of 30, but only rarely do they present with ascites and hydrothorax [1]. On its own, pseudo-Meigs’ syndrome is a rare diagnosis. A concomitant elevation in cancer antigen 125 (CA 125) in a patient presenting with pseudo-Meigs’ syndrome is an even more uncommon finding [4-7]. We report a case of pseudo-Meigs’ syndrome in a 35-year-old female who was also found to have a markedly elevated CA 125. Additionally, a literature review was
performed and past cases of pseudo-Meigs’ syndrome were identified. Data from those cases is presented in the discussion section of this paper following the case report.

CASE REPORT

A 35-year-old GoPo who presented to the emergency department with a chief complaint of abdominal swelling and weight gain beginning one month ago. She reported increasing abdominal pain, abdominal swelling, nausea, and vomiting. She denied chest pain, shortness of breath, heat or cold intolerance, palpitations, and any motor or sensory deficits. She had no history of cancer. Her past surgical history was significant for an exploratory laparotomy sixteen years ago due to an abdominal stab wound.

On physical examination, her abdomen was distended and mildly tender to palpation in all four quadrants. A fluid wave was present. A large mass was palpated in the left lower quadrant. No other abnormalities were observed on physical examination. Her blood work was significant for a leukocyte count of 12.8 \(10^3/\mu L\) and a markedly elevated CA 125 level of 272.9 U/ml. Given the elevated CA 125, combined with the ascites and mass on physical examination, a presumptive diagnosis of ovarian carcinoma was made.

A CT scan was ordered, which demonstrated a large intra abdominal mass measuring 28.8 cm and significant ascites. The mass contained both cystic and solid components, as well as internal vascularity. Omental carcinomatosis was also present.

The patient was taken to the operating room for a planned exploratory laparotomy, lysis of adhesions, omentectomy, excision of intra abdominal mass, hysterectomy, and bilateral salpingo-oophorectomy. An incision was made from the diploid process to the symphysis pubis. The wound was deepened using electrocautery. Upon entry into the abdomen, two ventral hernias were identified. The hernia sacs were divided and sent to pathology. Two liters of intra abdominal fluid was aspirated. Peritoneal washings were sent to pathology.

The mass was identified. It was closely adherent to the anterior abdominal wall, and also involved the omentum, sigmoid colon, and two large areas of small bowel. It was dissected from the anterior abdominal wall and the two segments of small bowel. Both of the segments of small bowel were resected. Two side-to-side functional end-to-end anastomoses were performed to close the common enterotomy.

A portion of the sigmoid mesentery involving the mass was removed. The tumor, which appeared to originate from the uterine fundus, was completely resected and sent to pathology. A total abdominal hysterectomy and bilateral salpingo-oophorectomy were performed. The abdomen was examined for additional sites of metastasis, but none were identified. The abdomen was closed using #1 looped Polydiaxanone sutures. The wound was irrigated with saline and subsequently closed with a skin stapler. The patient tolerated the procedure well and was extubated without complications. Pathology demonstrated that the mass was an aggressive-behaving uterine leiomyoma (Figure 1). Additionally, ascitic cytology was negative for malignant cells.

DISCUSSION

Findings of ascites, a pelvic mass, and an elevated CA 125 in a female carry a grim prognosis, most likely that of an ovarian carcinoma [8, 9]. However, as demonstrated by this case and others, benign etiologies must also be considered on the differential diagnosis [9]. Cases of Meigs’ syndrome and pseudo-Meigs’ syndrome with elevations in CA 125 have been reported, which should prompt physicians to consider these diagnoses as well [9–11]. Furthermore, an elevated CA 125 is not 100% specific for ovarian carcinoma [12]. Pelvic inflammatory disease, pregnancy, endometriosis, benign ovarian tumors, and uterine leiomyomas can cause CA 125 to become elevated [13]. In cases of Meigs’ syndrome, it has been conjectured that mesothelial expression of CA 125, as opposed to expression by the fibroma, is the cause of the elevation [10, 11]. In pseudo-Meigs’ syndrome, peritoneal inflammation has been implicated as the cause of the elevation in CA 125 [9].

In both Meigs’ syndrome and pseudo-Meigs’ syndrome, spontaneous resolution of the pleural effusion and ascites occurs following resection of the underlying tumor [14]. The cause of the ascites and hydrothorax in both Meigs’ syndrome and pseudo-Meigs’ syndrome is not entirely known, but likely multifactorial. It has been reported that, intraoperatively, cutting a uterine leiomyoma caused fluid leakage from myometrial cysts within the tumor [9]. Thus, it is possible that the fluid production is the result of cystic degeneration of the tumor [15]. In addition to fluid from the tumor itself, peritoneal fluid production is also a likely factor in the production of the ascites. Mechanical irritation of the peritoneum from the tumor in addition to pressure from pre-existing ascites causes peritoneal inflammation, which results in further fluid production [16]. The development of pleural effusion in these syndromes is thought to be a result of transdiaphragmatic transport of the ascitic fluid into the thorax through intercellular gaps [9].

A comprehensive review of the English literature using PubMed and Medline was performed, which resulted in 21 cases describing uterine leiomyomas presenting as pseudo-Meigs’ syndrome with an elevated CA 125 (Table 1). Review of the literature indicates that the mean age of presentation was 40 years (22–66 years). The majority of patients, 15 of 21 cases, presented with abdominal distention. Additionally, in 8 of 21 cases, patients reported dyspnea. Other symptoms less commonly reported include: nausea, vomiting, dysuria, and urinary frequency. The average CA 125 level of the 21 cases was
Table 1: Literature review of uterine leiomyomas presenting as pseudo-Meigs' syndrome with an elevated CA 125

<table>
<thead>
<tr>
<th>Author</th>
<th>Year</th>
<th>Age</th>
<th>Symptoms</th>
<th>CA 125 (U/mL)</th>
<th>Tumor size</th>
<th>Presence of ascites</th>
<th>Presence of pleural effusion</th>
<th>Type of surgery</th>
<th>Recurrence</th>
</tr>
</thead>
<tbody>
<tr>
<td>Frank et al.</td>
<td>1973</td>
<td>66</td>
<td>Dyspnea, fatigue, palpitations, orthopnea, and anorexia</td>
<td>NS</td>
<td>18x16x13 cm</td>
<td>Y</td>
<td>Y, right side</td>
<td>EX LAP and removal of tumor</td>
<td>NS</td>
</tr>
<tr>
<td>Rush</td>
<td>1976</td>
<td>47</td>
<td>8 months of weakness with exertion, intermittent painless abdominal swelling, constipation, lethargy, blood streaked stools</td>
<td>NS</td>
<td>20 cm in diameter</td>
<td>Y</td>
<td>Y, left side</td>
<td>EX LAP, TAH, BSO, colon polypectomy</td>
<td>NS</td>
</tr>
<tr>
<td>Handler et al.</td>
<td>1982</td>
<td>35</td>
<td>2 months of progressively increasing dyspnea, dysmenorrhea, deep dyspareunia</td>
<td>NS</td>
<td>9x9 cm</td>
<td>Y</td>
<td>Y, right side</td>
<td>EX LAP and excision of the mass</td>
<td>No recurrence after 1 year</td>
</tr>
<tr>
<td>Buckshee et al.</td>
<td>1990</td>
<td>46</td>
<td>Progressive abdominal distention and loss of appetite for 1.5 months</td>
<td>NS</td>
<td>20x20 cm</td>
<td>Y</td>
<td>Y, bilateral</td>
<td>TAH with myomectomy and RSO</td>
<td>No recurrence, but time not specified</td>
</tr>
<tr>
<td>Ollendorf et al.</td>
<td>1997</td>
<td>31</td>
<td>2 year history of increasing abdominal girth and six months of progressive shortness of breath</td>
<td>301</td>
<td>27x18x13 cm</td>
<td>Y</td>
<td>Y, right side</td>
<td>EX LAP and myomectomy</td>
<td>No recurrence after 8 months</td>
</tr>
<tr>
<td>Brown et al.</td>
<td>1998</td>
<td>31</td>
<td>Dyspnea, abdominal swelling, intermittent difficulty in passing urine, 6kg weight loss</td>
<td>83</td>
<td>17x11.5x8.5 cm</td>
<td>Y</td>
<td>Y, right side</td>
<td>NS</td>
<td>No recurrence after 18 months</td>
</tr>
<tr>
<td>Dunn et al.</td>
<td>1998</td>
<td>46</td>
<td>Nausea, vomiting, diarrhea, tachypnea</td>
<td>254</td>
<td>15x30x18 cm</td>
<td>Y</td>
<td>Y, right side</td>
<td>EX LAP, subtotal abdominal hysterectomy, BSO, omentectomy, and lymph node sampling</td>
<td>NS</td>
</tr>
<tr>
<td>Domingo et al.</td>
<td>1998</td>
<td>46</td>
<td>Menorrhagia for several years, 2 months of fatigue and malaise, 3 weeks of pedal edema, abdominal swelling, shortness of breath</td>
<td>317</td>
<td>20 cm in diameter</td>
<td>Y</td>
<td>Y, bilateral</td>
<td>TAH with BSO</td>
<td>NS</td>
</tr>
<tr>
<td>Migishima et al.</td>
<td>2000</td>
<td>51</td>
<td>Gradual abdominal distention for 3 years, progressive dyspnea for 5 months</td>
<td>820</td>
<td>12.3x24.3x20.5 cm</td>
<td>Y</td>
<td>Y, left side</td>
<td>EX LAP, TAH, BSO</td>
<td>No recurrence after 4 months</td>
</tr>
<tr>
<td>Amant et al.</td>
<td>2001</td>
<td>39</td>
<td>Abdominal swelling</td>
<td>785</td>
<td>30x30x15 cm</td>
<td>Y</td>
<td>Y, left side</td>
<td>TAH</td>
<td>NS</td>
</tr>
<tr>
<td>Author et al.</td>
<td>Year</td>
<td>Age</td>
<td>Symptoms</td>
<td>Mass Size</td>
<td>Side</td>
<td>Procedure</td>
<td>Recurrence</td>
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<tr>
<td>Kebapci et al. [22]</td>
<td>2002</td>
<td>38</td>
<td>1 week of low back pain, abdominal distention, weakness, and loss of appetite</td>
<td>281 x 10 x 10.5 cm</td>
<td>Y, left side</td>
<td>EX LAP and excision of the mass, aspiration of the ascitic fluid, omentectomy, and appendectomy</td>
<td>NS</td>
<td></td>
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<tr>
<td>Weise et al. [23]</td>
<td>2002</td>
<td>27</td>
<td>Increasing abdominal girth for 2 months</td>
<td>1854 x 7 x 8 x 6 cm</td>
<td>Y, right side</td>
<td>EX LAP and myomectomy</td>
<td>No recurrence after 3 months</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Weinrach et al. [24]</td>
<td>2004</td>
<td>40</td>
<td>2 year history of increasing abdominal distention and several months of progressive shortness of breath</td>
<td>734 x 19 x 11 x 10 cm</td>
<td>Y, right side</td>
<td>TAH with BSO</td>
<td>No recurrence after 6 months</td>
<td></td>
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<tr>
<td>Landrum et al. [25]</td>
<td>2008</td>
<td>47</td>
<td>Symptoms of upper respiratory infection, worsening anasarca and profound increase in abdominal girth</td>
<td>475 x 20 x 22 cm</td>
<td>Y, bilateral</td>
<td>EX LAP, TAH, BSO, pelvic and para-aortic lymphadenectomy</td>
<td>NS</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ricci et al. [26]</td>
<td>2009</td>
<td>35</td>
<td>Abdominal distention 2 days after a spontaneous vaginal delivery</td>
<td>231.4 x 15 x 10 x 8.5 cm</td>
<td>Y</td>
<td>EX LAP and removal of pedunculated mass</td>
<td>No recurrence after 3 years</td>
<td></td>
<td></td>
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<tr>
<td>Chourmouzi et al. [3]</td>
<td>2010</td>
<td>41</td>
<td>12 month history of abdominal swelling, discomfort, urinary frequency and incontinence</td>
<td>436.7 x 13 x 16 cm</td>
<td>Y, bilateral</td>
<td>EX LAP and excision of the mass</td>
<td>NS</td>
<td></td>
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</tr>
<tr>
<td>Makris et al. [27]</td>
<td>2012</td>
<td>26</td>
<td>Noticed pelvic mass</td>
<td>93.9 x 11 x 8.3 x 10.2 cm</td>
<td>Y</td>
<td>EX LAP and open myomectomy</td>
<td>No recurrence after 2 months</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yip et al. [13]</td>
<td>2013</td>
<td>41</td>
<td>Abdominal fullness and prolonged menstrual periods for 3 years</td>
<td>939.7 x 12 x 11 x 7.8 cm</td>
<td>Y</td>
<td>EX LAP and removal of mass</td>
<td>NS</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Oguma et al. [28]</td>
<td>2014</td>
<td>50</td>
<td>3 month history of shortness of breath</td>
<td>218 x 14 x 8 x 7 cm</td>
<td>Y, right side</td>
<td>TAH with BSO</td>
<td>Not specified</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Seo et al. [29]</td>
<td>2014</td>
<td>22</td>
<td>10 days of painless abdominal distention</td>
<td>450 x 8.9 x 5.2 cm</td>
<td>Y</td>
<td>EX LAP and removal of mass</td>
<td>No recurrence after 5 years</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dong et al. [12]</td>
<td>2015</td>
<td>37</td>
<td>Right lower abdominal dull pain for 2 years, abdominal distention for 6 days, nausea</td>
<td>920.4 x 20 x 18 x 10 cm</td>
<td>Y, bilateral</td>
<td>EX LAP and TAH</td>
<td>No recurrence after 82 months</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Y- Yes  
N- No  
NS- Not specified  
EX LAP- Exploratory laparotomy  
TAH- Total abdominal hysterectomy  
BSO- Bilateral salpingo-oophorectomy
541 U/mL (83 U/mL-1854 U/mL). Ascites was present in all 21 cases, and hydrothorax was reported in 16 of the 21 cases. All patients were treated surgically, with no reported cases of tumor recurrence.

CONCLUSION

Uterine leiomyomas presenting as pseudo-Meigs’ syndrome with an elevated CA 125 are exceedingly rare. In a patient with a clinical presentation consistent ovarian cancer with the added finding of an elevated CA 125, benign etiologies such as Meigs’ syndrome and pseudo-Meigs’ syndrome should also be considered in the differential diagnosis. Surgery is the mainstay of treatment in pseudo-Meigs’ syndrome. Resolution of the ascites and hydrothorax occurs spontaneously following resection of the tumor. No cases of recurrence have been reported in the literature.

REFERENCES


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Author Contributions
Jake Alan Gibbons – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Guarantor of Submission
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Conflict of Interest
Authors declare no conflict of interest.

Data Availability
All relevant data are within the paper and its Supporting Information files.

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