Case Report

Uterine adenosarcoma: a case report and review of the literature

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Abstract: Uterine adenosarcoma is a rare gynecological malignancy with no specific symptoms, and the optimal management is still inconclusive. Herein we present a case of uterine adenosarcoma in a 38-year-old woman with a good prognosis and review of literatures. The patient presented with abnormal vaginal bleeding with no special medical history. Sonographic scan revealed a heterogeneous echoic mass in the cavity, indicating a polypus or a submucous myoma. The pathology based on the specimen after the hysteroscopic tumor excision suggested diagnosis of uterine adenosarcoma. Subsequently, the patient received pelvic MRI scan before surgery. MRI identified a patchy lesion at the cervix-lower endometrial cavity with low signal in T1WI and a mixed high T2 signal in T2WI, with no sign of metastasis. Then total abdominal hysterectomy with bilateral salpingo-oopherectomy plus pelvic lymph node dissection was performed and 6 cycles of chemotherapy were administered. The patient remains disease-free on follow-up to date, more than 15 months after chemotherapy.

Keywords: Müllerian adenosarcoma, sarcoma, uterine cancers, gynecological malignancy, Müllerian malignancy

Introduction

Uterine adenosarcoma is a rare malignancy, accounts for 5%-10% of all uterine sarcomas [1, 2]. It was first reported by Clement and Scully in 1974, characterized by components of both sarcoma and benign epithelial glands [3]. The morbidity was hard to estimate worldwide but it only has an age-adjusted incidence of 2-3 per 1,000,000 in the US population [4]. Due to the rarity and the inconsistent of clinical management, here we present a case of uterine adenosarcoma in a 38-year-old woman with a good prognosis.

Case

A 38-year-old woman presented with abnormal vaginal bleeding for 1 year and a prolonged 10-day menstruation compared to the original 3-5 days with no other complaints such as menorrhagia, watery discharge, or abdominal fullness etc. She had no other medical history but caesarean section for one childbirth and she took no specific medications. The patient did not go to the clinic in the past year until her last menstrual period lasted for 15 days. In the outpatient department, no significant signs were found during the pelvic exam while transvaginal 3D ultrasound scan revealed a heterogeneous echoic mass, 3.8*2.2*2.1 cm in size with a cystic region of approximately 1.6*1.5*1.3 cm, started from the uterine cavity and descended to the cervical canal.

An endometrial polypus or a submucous myoma was suspected, the patient was admitted and hysteroscopy was performed. During the hysteroscopy, the endocervical canal was visually normal, but from the middle of endometrial cavity to the internal os, a soft, partly cystic, sessile mass about 4*2*2 cm in size was seen. It was difficult to specify whether the mass was a submucous myoma with cystic degeneration or an endometrial polypus intraoperatively, so a hysteroscopic tumor excision was done. The pathological examination found that the tumor was composed of benign glandular elements and malignant stromal components. The epithelial glands were papillary or slit-like while the stro-
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Figure 1. The tumor was composed of benign glandular elements and malignant stromal components. A. The slit-like epithelial glands were widely distributed in the stroma. B. The stromal cells forming intraglandular protrusions (blue arrow) and peri-glandular cuffs around the glands (black arrow).

mal cells were spindle-shaped, with increased mitotic activity. The stromal cells formed intraglandular protrusions and in some areas, formed peri-glandular cuffs around the glands, which was the typical microscopic characteristic of adenosarcoma (Figure 1). The final pathological diagnosis was uterine adenosarcoma, with the malignant stromal component was low grade endometrial stromal sarcoma.

In order to understand the characteristics of the tumor and whether there was myometrial invasion or distant metastasis, pelvic Magnetic Resonance Image (MRI) with contrast enhanced was performed before reoperation. It can be observed a patchy lesion at the cervix-lower endometrial cavity with low signal in T1 Weighted Image (T1WI) and a mixed high T2 signal in T2 Weighted Image (T2WI). No obvious diffusion limitation was observed in Diffusion Weighted Image (DWI), and uneven tissue enhancement was shown in T1WI with contrast-enhanced (Figure 2). There were no obvious enlarged lymph nodes in pelvic cavity and no sign of myometrial invasion on MRI.

A subsequent laparoscopy was scheduled. There were no visible tumor and no enlarged lymph nodes detected in the surgery, then total abdominal hysterectomy with bilateral salpingo-oophorectomy (TAH with BSO) and pelvic lymphadenectomy was performed.

Macroscopic examination found a protruding adenosarcoma sized 1.8*1.2*0.8 cm at the cervical-lower uterus. The tumor is mainly exogenous, with focal invasion to the superficial myometrium (<½ myometrial thickness), and no lymphovascular space invasion (LVSI), no involvement on surgical margins, bilateral parametria, bilateral uterine horns, bilateral fallopian tubes and ovaries, and 24 resected lymph nodes were negative. Given that the risk of recurrence increased while the presence of myometrial invasion, we conducted 6 cycles of chemotherapy, with regimen of pegylated liposomal doxorubicin and carboplatin. The patient has been followed up for more than 15 months after chemotherapy, and has no signs of recurrence so far.

Discussion

Uterine adenosarcoma is a biphasic neoplasm consisting of benign or occasionally atypical glandular component and malignant stromal component. It accounts for about 5%-10% of uterine sarcomas, while uterine sarcomas are only 1% of all gynecological malignancies and 3% of all uterine cancers [1, 2, 4, 5].

Uterine adenosarcoma occurs more frequently in postmenopausal women. According to three retrospective analyses, the median ages of patients diagnosed with uterine adenosarcoma were 54 [6], 56 [7] and 58 [8] respectively, ranging from 14 to 89 years old. Of 544 cases from the SEERs database, 51.5% of the patients were between 40 and 65 years old, and less than 10% were younger than 40 years old.
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Figure 2. MRI revealed a patchy lesion (arrow) of 17*9 mm at the cervix-lower endometrial cavity. A. A mixed high T2 signal in T2WI. B. Uneven tissue enhancement in T1WI with contrast-enhanced.

The patient in this study was a 38-year-old woman of reproductive age.

The most common symptom of uterine adenosarcoma is abnormal vaginal bleeding, but a small number of patients present with uterine enlargement, pelvic mass, pelvic pain, uterine prolapse, vaginal discharge, or some rare symptoms include dysmenorrhea, dysuria, infertility, abdominal distension caused by ascites [2, 4, 6, 8, 9]. During gynecological examination, tumors can be seen protruding from the external cervical orifice or even fill the vagina, which may be misdiagnosed as “cervical polyps”, especially in patients with histories of recurrent endometrial polyps or cervical canal polyps [2, 6, 8, 10-13].

Uterine adenosarcomas occur typically in the endometrium (87%) and may also be confined to the internal os (9%) or myometrium (4%) [6, 8-10, 14]. Similar tumors can originate in the cervix [12, 13, 15-17], ovaries [6, 15, 18-24], pelvis [25-28], and vagina [29-31]. The diameter of adenosarcoma varied from 0.1 cm to >20 cm, with an average of 5 cm [8, 10, 32]. Most tumors are lobulated and granular, grown as solitary polypoid, but multiple polypoid masses within the uterine cavity have also been seen [2, 6, 8-10, 32].

Due to well performance on soft tissue resolution, MRI can display the internal structure and changes such as hemorrhage or necrosis of the tumor, and can accurately determine the relationship between the lesion and the endometrium or myometrium. Typical MRI manifestations of uterine adenosarcoma are huge polypoid masses arising from the uterine cavity, protruding to the vagina through the cervical canal, with clear or blurred borders, and multiple small cystic areas inside [33-36]. Metastasis and enlarged lymph nodes could also be revealed on MRI in advanced patients. Compared with the signal of myometrium, the masses of uterine adenosarcoma often showed slightly high signal on T2WI and low or high signal on T1WI, which was quite similar to that of endometrial polyps or submucosal fibroids [34, 36]. Usually, there is heterogeneous enhancement in T1WI with contrast-enhanced [34, 35]. However, with regard to DWI, the results are inconsistent. Some researchers reported that adenosarcomas usually presented relatively low DWI signal intensity, especially low-grade adenosarcomas [35-38], while some other researchers found that uterine adenosarcomas showed slightly high signal or high signal on DWI [39-41]. The reasons for the different DWI signals are still unclear but could be explained by the different proportions of benign epithelial and malignant mesenchymal components to some extent. Other imaging modalities can also be used, such as sonography, CT or Positron emission tomography-computed tomography.
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(PET-CT). Sonography and CT are nonspecific but most reachable, usually revealed enlarged uterine with a heterogeneous mass inside the cavity [34, 42]. PET/CT can be used to identify distant metastases or relapse locations [43].

Without specific immunohistochemical markers, the diagnosis of uterine adenosarcomas mainly relies on morphological features. The typical microscopic characteristic of adenosarcoma is peri-glandular cuffs composed of benign epithelial glands and malignant stromal components [2, 6, 8, 44, 45]. The mitotic activity of the stroma cells varies greatly, ranging from 0 to 24 MF/10 high-power fields (hpf) [8, 10, 45, 46]. About 30% of cases were accompanied by endometrial dysplasia or endometrial adenocarcinoma [8]. Most uterine adenosarcomas were confined to the endometrium [6, 44, 45, 47], but Zaloudek et al. reported 7/25 cases of myometrial invasion [10], Clement et al. found that 15/100 cases had superficial myometrial invasion, while 4 cases had deep invasion [8]. The immunophenotype of most uterine adenosarcomas is similar to that of endometrial stromal sarcomas (positive ER, PR, WT1, and CD10), while patients with sarcoma overgrowth have loss of ER, PR, and CD10 expression [48-50].

There is no consensus on the optimal approach to treatment of uterine adenosarcoma. According to the published data, the recommended management for adenosarcoma is TAH with BSO [9, 51], but for women with fertility requirements, fertility sparing surgery (FSS) can be considered for appropriate selected patients [52-54]. ZiZolfi et al. reported one 23-year-old woman with Stage IA uterine adenosarcoma underwent hysteroscopic tumor resection followed by 160 mg/d oral megestrol acetate, and got full-term infant four years after diagnosis [54]. Goh et al. reported a 21-year-old patient with Stage I low-grade uterine adenosarcoma, underwent dilatation and curettage (D&C) and polypectomy, and got spontaneous conception and delivery four years later, but she received TAH+BSO+ lymphadenectomy when recurrence was confirmed 96 months after initial diagnosis [55]. L’Heveder et al. reported an 18-year-old girl who diagnosed as low-grade adenosarcoma after polypectomy, she conceived and delivered twins at 28 weeks, and underwent hysterectomy 20 years postoperatively with no recurrence [53]. Lee et al. reported seven cases, the largest case series to date, of Stage I uterine adenosarcoma managed with FSS, but two of them suffered recurrence while only one woman got livebirth [56]. Zaloudek et al. reported three cases of 14 to 15-year-old patients, one underwent TAH with ovary preservation, the second only received cervixectomy, and the third underwent only tumor excision. All three patients received radiotherapy after surgery, and no recurrence was observed during follow-up, but only the third woman got a normal pregnancy [10]. Michener et al. reported a 25-year-old female diagnosed with Stage IB uterine adenosarcoma who underwent hysterectomy but preserved ovaries, and was suggested to in vitro fertilization using a surrogate [52]. Due to the extremely rarity, it cannot to conclude a standard procedure nor unified criteria of FSS, but it seems it can be performed in young patients with low-grade, stage IA disease without sarcomatous overgrowth, and close long-term follow-up is needed. However, fertility-sparing surgery is still not the recommended approach due to the risk of recurrence, and hysterectomy should be considered after completion of childbearing.

Lymphadenectomy is still controversial. It has been reported that the incidence of lymph node metastasis in uterine adenosarcoma is very low, ranging from 0 to 6.5% [2, 6, 7, 9, 57]. Furthermore, some researchers found that lymphadenectomy did not show overall or progression-free survival benefit [6, 7]. But meanwhile, some studies found patients underwent lymphadenectomy showed a better survival [57, 58]. In a retrospective study of 994 cases and a systematic review of 230 surgically treated adenosarcomas, lymph node involvement revealed the greatest impact on progression-free survival compared to other factors such as deep myometrial invasion or sarcomatous overgrowth [57]. So, the question of whether lymph nodes should be removed during the primary surgery remained unanswered. But in any case, enlarged lymph nodes should be excised to rule out or confirm metastasis [57, 59]. Although the mainstream guidelines like the NCCN guideline suggested TH+BSO as standard surgery for uterine adenosarcoma, lymphadenectomy is practiced widely. Of 1027 cases from the SEERs database, 53.1% of patients underwent lymph node dissection [7]. In this case, consid-
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The involvement of lymph node metastasis in tumor staging and it may be an indicator of poor prognosis, we performed lymphadenectomy during the surgery.

The general 5-year overall survival of uterine adenosarcoma is about 60-87%, and the most important prognostic factors are tumor stage, presence of sarcomatous overgrowth and myometrial invasion [2, 4-7, 9, 46, 47, 51, 57, 60, 61]. Survival analyses from the SEERs database indicated that the 5-year survival for patients with stage I was 79% (95% CI, 75-84%) and for patients with stage III was 48% (95% CI, 29-65%) [9]. However, the 2-year PFS and OS were only 20%-50% with sarcomatous overgrowth while both 100% without sarcomatous overgrowth [6, 45, 60]. Myometrial invasion is related with a more aggressive behavior and a high rate of recurrence, the overall survival for early stage patients with myometrial invasion was only 63% [9]. The case reported here was stage IB with myometrial invasion, so the risk of recurrence cannot be ignored. Therefore, we administered 6 cycles of adjuvant chemotherapy. The patient was disease-free to date.

Conclusion

Most patients with uterine adenosarcoma are in the early stage and with relatively good prognosis. The recommended management is TAH with BSO, lymphadenectomy may not be routinely performed but enlarged lymph nodes should be resected. Sarcoma-specified chemotherapy regimens (eg, doxorubicin or ifosfamide) may be considered in patients at high risk of recurrence or death, such as myometrial invasion, sarcomatous overgrowth and advanced stage.

Disclosure of conflict of interest

None.

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