Case report

Management and survival of patients with Mullerian adenosarcoma of the cervix without sarcomatous overgrowth desiring fertility preservation, a case report and review of the literature

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1. Introduction

Mullerian adenosarcomas are a rare malignancy, first described in 1974 (Roth et al., 1976). They are a form of mesenchymal tumor that most often arise in the uterus in postmenopausal women, however, there may be extrauterine manifestations including the cervix (Kanayama et al., 2017). They most commonly present in a low-grade fashion with a benign epithelial component and an internal malignant mesenchymal component, although high grade manifestations have been reported (Bagga et al., 2010). On evaluation of mesenchymal tumors, Mullerian adenosarcomas are often described as a stepping point between benign adenofibromas and malignant carcinomas (Andrade et al., 1992). Cervical Mullerian adenosarcomas primarily present in adolescents (Fleming et al., 2009).

There is limited information regarding primary cervical Mullerian adenosarcomas. Prognosis has historically been favorable, with the exception of deeply invasive tumors or those with high-grade sarcomatous overgrowth (Fleming et al., 2009). Twenty-nine cases of women with low grade disease have been collected for review. Twenty cases were primarily managed with hysterectomy. The remaining nine cases underwent local excision, a cold knife conization, or trachelectomy with five known cases of disease recurrence requiring additional intervention. Mortality has been identified in one case. We present a case of a 14-year-old woman treated with a cervical wedge resection and no disease recurrence 4 years after diagnosis (see Table 1).

2. Case

The patient presented at age 14 with intermenstrual spotting with a vaginal mass. The patient was otherwise healthy, not sexually active, with menarche at age 12 with intermittent dysmenorrhea. A pap smear was performed showing ASCUS, High Risk HPV negative. Pelvic ultrasound was noted to be significant for an antverted uterus measuring 6.7 × 4.0 × 5.2 cm and an endometrial stripe measuring 7 mm. A small 5 mm hypoechoic avascular structure was noted in the cervix and an ovoid hypoechoic structure was seen in the vaginal canal without vascularity, measuring 1.2 × 0.7 × 1.0 cm. Both ovaries were normal. An exam under anesthesia was performed, and a 2 × 3 cm polypoid cervical mass was removed using torsion. Electrocautery was used to remove the stump. Pathology was significant for adenosarcoma with low grade morphology and unclear margins. The lesion was described as a polypoid mass with phylloid type growth pattern with areas of immature spindled stroma condensing around irregular and dilated cystic spaces lined by benign endocervical type columnar epithelium (Fig. 1). No pleomorphic cells were noted in the stroma. As malignancy was not expected at the time of the procedure, the specimen was not marked for orientation, and thus no comment on margins were able to be made.

The patient was referred to gynecologic oncology. MRI pelvis showed a questionable lesion in the mesentery but no pelvic findings. PET CT was concerning for hypermetabolic activity in the subcarinal and right hilar lymph nodes and a possible hypermetabolic mesenteric implant. CT scan confirmed mildly enlarged mesenteric lymph node of unclear clinical significance. The patient underwent subcarinal endoscopic lymph node dissection that showed no carcinomatous or sarcomatous elements identified. A second dilation and curettage was performed. An endometrial polyp was noted and extracted. An additional small cervical polyp was removed and a 1.5 × 1.5 cm cervical wedge resection of the posterior cervical lip was performed at the site of the original lesion using a scalpel. Final pathology showed no evidence of malignancy. Management options were reviewed with the patient and family including hysterectomy or conservative management with observation. Conservative management was chosen. PET CT after 6 months showed no evidence of recurrence. Patient underwent repeat dilation and curettage and MRI pelvis secondary to intermenstrual vaginal bleeding 7 months after initial diagnosis. No evidence of recurrence was identified. The patient has remained disease free for four years to date with recent MRI pelvis with no evidence of disease. The patient and her mother have consented to participation in this case report (see Fig. 2).

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https://doi.org/10.1016/j.gore.2019.100525
Received 20 August 2019; Received in revised form 23 November 2019; Accepted 25 November 2019
Available online 07 December 2019
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| Author | Age (yo) | Diagnosis or Presentation | Available pathology | Treatment | Follow Up | Recurrence | Surveillance (if available) |
|--------|----------|---------------------------|---------------------|-----------|-----------|------------|---------------------------|
| Roth et al., 1976 | 14 | Cervical mass biopsy | Superficial invasion, heterologous elements | RH, BSO, LND, upper vaginectomy, CT, RT | 41 months | NED | n/a NED after RH |
| Martinelli et al., 1988 | 55 | Polypoid mass biopsy | Superficial invasion | TAH, BSO, RT | 4.5 years | NED | n/a NED after TAH |
| Ostor and Fortune, 1980 | 47 | Cervical polyp | Unavailable | Polyp excision | 14 years | Case A: Endometrium after 11 years | Case A: NED after hysterectomy |
| Zaloudek and Norris, 1981 | 15yo | Cervical mass biopsy | Unavailable | Trachelectomy | 6 yrs | Case A: NED for 11.5 yrs | Case A: NED after hysterectomy |
| Edinger et al., 1982 | 17 | ECC, polypectomy | Heterologous elements, no invasion | RT, TAH, BSO | 56 months | NED | n/a NED after hysterectomy/BSO and RT |
| Chen, 1985 | 11 | Cervical mass biopsy | Heterologous elements, no invasion | RH, partial vaginectomy, LND, TAH, BSO | 4.5 years | NED | n/a NED after hysterectomy |
| Gal et al., 1988 | 14 | D&C | Unavailable | CT, RT, TAH, BSO | 5 months | Recurrence in pelvis, vagina | Case D: Died of disease, 14 months after recurrence, CT, RT, TD, TAH, BSO, and hysterectomy |
| Gast et al., 1987 | 15 | D&C, polypectomy | Heterologous elements, no invasion | RAH | 4 years | NED | n/a NED after hysterectomy |
| Clement and Scully, 1990 | 19 | Cervical mass biopsy | Unavailable | Chemotherapy | 30 months | NED | n/a NED after hysterectomy/BSO and CT |
| Kerner, 1993 | 30 | D&C, cervical polyp | No invasion | RH | 3.5 years | NED | n/a NED after hysterectomy |
| Feroze et al., 1997 | 18 | Abdominal mass | Biopsy | RH | 3.5 years | NED | n/a NED after hysterectomy |
| Ramos et al., 2002 | 25 | D&C | Heterologous elements, no invasion | TAH, BSO | 2 years | NED | n/a NED after hysterectomy |
| Manoharan et al., 2007 | 26 | Abdominal mass biopsy | No invasion | RH | 10 months | NED | n/a NED after simple excision |
| Kanayama et al., 2017 | 15 | Cervical mass biopsy | Heterologous elements | TAH, BSO, LND, omentectomy | 10 months | NED | n/a NED after hysterectomy |
| Buyukyurt et al., 2010 | 14 | Exam under anesthesia, simple excision | Heterologous elements, no invasion | Simple excision | 15 months | NED | n/a NED after hysterectomy/BSO and CT |
## Table 1 (continued)

| Author | Age (yo) | Diagnosis or Presentation | Available pathology | Treatment | Follow Up | Recurrence | Surveillance (if available) and outcome |
|--------|----------|---------------------------|---------------------|-----------|-----------|------------|---------------------------------------|
| Fleming et al., 2009 | 10 | Cervical mass biopsy, Superficial invasion | Rh, LND, upper abdominal MRI, physical exam | 18 months NED | NED | n/a | Abdominal MRI, physical exam, NED after hysterectomy |
| Kanayama et al., 2017 | 28 | Cervical mass biopsy | Heterologous elements | CKC | 32 months | Cervix and vagina, Transcervical resection | NED after transcervical resection, pregnancy |
| Shinnick et al., 2017 | 14 | Bulging vaginal mass, No invasion | CKC | 6 months NED | n/a | MRI, US, physical exam | MRI, vaginal hysterectomy, NED: no evidence of disease, BSO: bilateral salpingo-oophorectomy, RH: radical hysterectomy, CT: chemotherapy, RT: radiation therapy, TVUS: transvaginal ultrasound, n/a: not applicable, VH: vaginal hysterectomy, CKC: cold knife cone, HSC: hysteroscopy, D&C: dilation and curettage, ECC: endocervical curettage. |

3. Discussion

Cervical Mullerian adenosarcomas are a rare form of malignancy with limited information regarding their management, with historic precedent primarily involving hysterectomy for reproductively aged women. This presents a management dilemma as low grade cervical adenosarcomas typically present in adolescents who may desire fertility preservation.

A literature review of all known cases with low grade cervical Mullerian adenosarcoma was performed and summarized. A total of 29 cases were collected and reviewed.

The patients identified ranged from ages 10 to 67 with the median age of 24.5. Of the total 29 cases reviewed, twenty were noted to have been treated with hysterectomy, with only one having recurrence of disease (Martinelli et al., 1988; Edinger et al., 1982; Chen, 1985).

The patient with recurrent disease was a 14-year-old with a large vaginal mass $12 \times 8 \times 8$ cm occluding her urethra diagnosed as Mullerian adenosarcoma of the cervix with heterologous elements. The patient received 3 cycles of preoperative vincristine, Adriamycin, cyclophosphamide, and Actinomycin D and then underwent pelvic and abdominal radiation. She underwent a hysterectomy and bilateral salpingo-oophorectomy, with negative resection margins. Vaginal recurrence occurred five months later with diagnosis of a 10 cm mass and subsequent demise (Gal et al., 1988). This case is unique in its aggressive and extensive nature of disease at the time of diagnosis and management, despite low grade pathology. This may identify this case as an outlier.

A total of nine patients underwent initial conservative management. Six cases were treated with local resection (Ostor and Fortune, 1980; Clement and Scully, 1990; Zaloudek and Norris, 1981; Buyykurt et al., 2010), one case underwent a trachelectomy (Zaloudek and Norris, 1981), and two cases underwent a cold knife cone as initial treatment (Kanayama et al., 2017; Shinnick et al., 2017).

Of those undergoing local resection, three had documented recurrences. One of these subjects underwent a hysterectomy (Ostor and Fortune, 1980), another was treated with a repeat dilation and curettage (Clement and Scully, 1990), and another underwent a cold knife cone (Clement and Scully, 1990). The patient who initially underwent a trachelectomy had no recurrence (Zaloudek and Norris, 1981). One of the patients who initially underwent a cold knife cone recurred, and was treated with a subsequent resection and has had a successful pregnancy (Kanayama et al., 2017). The other patient that underwent a cold knife cone initially is currently undergoing surveillance with biannual MRI, transvaginal US, and serial clinical exams (Shinnick et al., 2017). These cases summarize how varied the treatment has been for this diagnosis, the majority of which have been hysterectomy for reproductively aged women.

We present the case of a patient with low grade primary cervical adenosarcoma who underwent a cervical wedge resection and polypectomy, and who has remained disease free for four years since diagnosis. This is the first case to have had no evidence of recurrence with the most minimally invasive resection approach for this time frame. We have recommended the patient undergo continued surveillance visits, with specific counseling regarding menstrual disturbances as abnormal bleeding is often the initial symptom of recurrence. There are no formal recommendations by any governing body regarding surveillance timing or interventions. Recurrence has been documented as late as 11 years after diagnosis (Ostor and Fortune, 1980). Surveillance for that minimum of time can be considered. An initial assessment with dilation and curettage or local excision for a visible lesion would be reasonable to evaluate for recurrence, as many have been documented as masses. Although more aggressive surgical management has been undertaken in the past, recurrences appear to be local. Resection is a reasonable option, particularly in cases like ours where the presentation is a small polypoid mass amenable to torsion or local resection. Cold knife cones have also been identified to be successful, however, we present an even
more minimally invasive resection approach with similar results, which may lead to a better reproductive outcome for patients desiring fertility preservation.

Routine imaging does not appear critical to surveillance with some false positive results documented in our case, but may be a useful tool. Pathology expertise seems essential to ensure correct diagnosis and treatment options. Overall, this case report aims to discuss the possibility of offering more minimally invasive approaches for young women affected with a low grade rare tumor during their reproductive years.

**Author contribution**

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Assisted in drafting case report, performed literature review

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John Groth

Reviewed case report, performed and provided pathology images and captions

Nuzhath Hussain

Reviewed case report, performed initial procedure that led to diagnosis

Rajul Kothari

Edited and assisted in drafting case report, performed literature review, provided oncologic care to patient

**Conflict of Interest**

The authors declare that there are no conflicts of interest.
Acknowledgements

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