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Case Report

Cervical Mesonephric Adenocarcinoma Treated with Neoadjuvant Chemotherapy. A Case Report and a Literature Review

Hiroaki Ishida ^{1,*}, Megumi Manrai ¹, Hiroki Egashira ¹, Mizuki Nonaka ¹, Nobuyuki Hiruta ², Reiko Watanabe ³ and Akiko Takashima ¹

¹ Department of Obstetrics and Gynecology, Toho University Medical Center Sakura

² Department of Pathology, Toho University Medical Center Sakura

³ Department of Pathology, St. Marianna University of Medicine

* Correspondence: ishida-04@sakura.med.toho-u.ac.jp; Tel.: +043-462-8811

Abstract: Cervical mesonephric ductal adenocarcinoma (MA) is a HPV-independent adenocarcinoma that originates from remnants of the wolffian duct in the male reproductive system during fetal development and occurs in middle-aged women. MA is a rare disease that accounts for less than 1% of all cervical adenocarcinomas. We report a case of MA in which abdominal radical hysterectomy (ARH) was performed after neoadjuvant chemotherapy (NAC). The patient was a 66-year-old woman with abnormal genital bleeding. A colposcopy examination revealed macroscopic invasive cancer. A pelvic MRI scan revealed a 53 × 26 mm tumor in the cervix, and the histological diagnosis of the cervix was endometrioid carcinoma, with the diagnosis being cervical adenocarcinoma cT1b3N0M0. One course of NAC with paclitaxel-carboplatin (PC) was administered to shrink the tumor and stop the bleeding, and ARH was performed. Postoperative histopathological diagnosis was MA. The surgical margins of the resected specimen were negative, and NAC had been effective, so the patient underwent five courses of PC therapy after surgery. There has been no recurrence 12 months after surgery. There is no established standard treatment, but there are reports that PC therapy is effective. It is necessary to search for effective treatments by following up and accumulating further cases.

Keywords: Cervical mesonephric adenocarcinoma; Neoadjuvant chemotherapy; Adjuvant chemotherapy

1. Introduction

Mesonephric adenocarcinoma (MA) of the cervix is HPV-independent adenocarcinoma [1]. It is currently believed that MA originates from remnants of the Wolffian duct in the male reproductive system during fetal development and occurs in middle-aged women. The average age of patients is 53 years, and it often originates from the lateral wall of the cervix [1]. MA is a rare form of cervical adenocarcinoma, accounting for less than 1% of all cervical adenocarcinomas. In a multicenter collaborative study by Pors J et al. [2], 30 cases of MA and their clinicopathological characteristics were analyzed. Follow-up data revealed that the prognosis of cervical MA was worse than that of HPV-related adenocarcinoma. No standard treatment has been established for cervical MA. We report a case of cervical MA in which radical hysterectomy was performed after neoadjuvant chemotherapy, based on a literature review.

2. Clinical Case

66 years old, G2P2, menopause at 58 years old. Medical history: diabetes.

She had been experiencing abnormal genital bleeding for the past six months, and one month ago she began experiencing daily bleeding, so she visited a local doctor. A cytological examination at

a local hospital revealed cervical cytology, AGC, and endometrial cytology class IIIb, and a tumor was found in the cervix. A cervical biopsy was performed, and the patient was diagnosed with endometrioid carcinoma, and referred to our hospital. Colposcopy of the cervix at our hospital revealed a bulky tumor occupying the vagina, and the tumor was diagnosed as macroscopic invasive carcinoma (Figure 1). A transvaginal ultrasound scan revealed a cervical tumor measuring 47×22 mm (Figure 2), and a pelvic MRI scan revealed a cervical tumor measuring 53×26 mm (Figure 3A). CT scan showed no distant metastasis. Tumor markers were CA125: 13.3 U/ml, CEA: 2.4 U/ml, CA19-9: 6.5 U/L, SCC: 0.5 ng/ml, and no abnormal values were found. A histological diagnosis of the cervix at our hospital also revealed a diagnosis of endometrioid carcinoma. Based on the above, the patient was diagnosed with cervical adenocarcinoma, clinical stage IB3 (cT1b3N0M0). Surgery was scheduled for 40 days later. As the tumor was bulky and bleeding continued, one course of neoadjuvant chemotherapy with paclitaxel-carboplatin (PC) was administered with the aim of reducing the tumor and stopping the bleeding. After one course of PC, the cervical tumor was reduced in size (Figure 3B), and the patient was diagnosed with partial response (PR).

40 days after the initial consultation, an abdominal radical hysterectomy + bilateral salpingo-oophorectomy + pelvic lymphadenectomy (ARH+BSO+PLA) was performed. The operation time was 6 hours and 22 minutes, the blood loss was 905ml, and the postoperative course was uneventful, and the patient was discharged on the 9th day after the operation. Macroscopic findings of the resected specimen showed that the resection margins between the tumor and the parametria were negative, and the resection was diagnosed as complete (Figure 4). HE staining revealed that atypical cells proliferated forming a lumen, and the lumen contained eosinophilic protein substances (Figure 5A). Morphologically, the tumor resembled endometrioid carcinoma, but remnants of the mesonephric duct are observed (Figure 5B). Immunohistochemistry showed that it was transacting T-cell-specific transcription factor GATA-3 (GATA3) (Figure 6A), clusters of differentiation 10 (CD10) (Figure 6B) and thyroid transcription factor 1 (TTF-1) positive (Figure 6C) and estrogen receptor (ER) negative, (Figure 6D) and the tumor was diagnosed as HPV-independent mesonephric adenocarcinoma. The surgical margins of the resected specimen were negative, and the tumor was sufficiently separated from the lesion. As preoperative chemotherapy was effective, the patient underwent five courses of PC therapy after surgery. There has been no recurrence as of 12 months after surgery.



Figure 3. A Pelvic MRI T2 weighted image : A cervical tumor measuring 53×26 mm. Before chemotherapy. **Figure 3B** Pelvic MRI T2 weighted image : After one course of neoadjuvant chemotherapy with paclitaxel-carboplatin (PC) . A cervical tumor measuring 43×23 mm. The tumor shrank and PR was diagnosed.

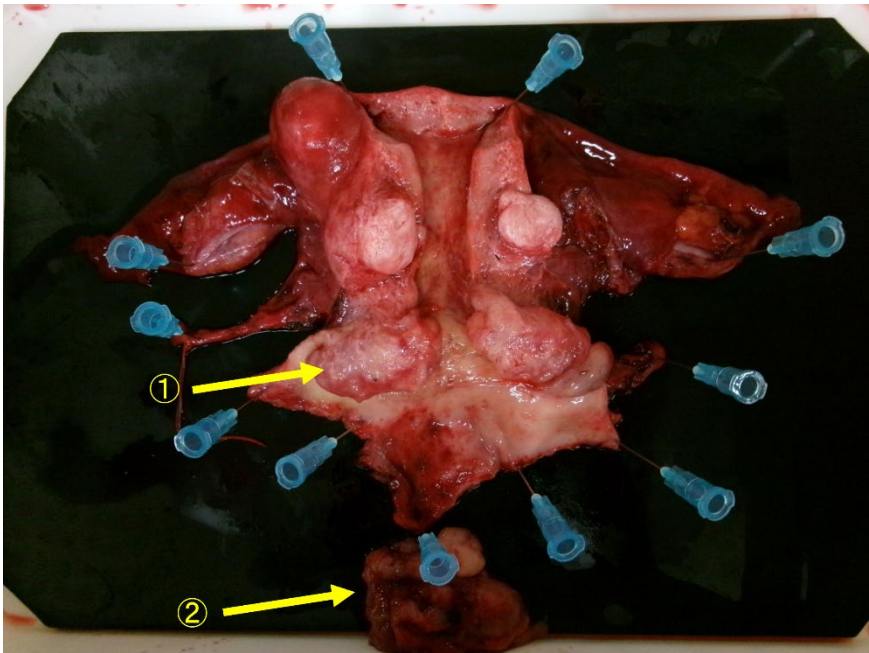


Figure 4. Macroscopic findings of the resected specimen showed that the resection margins between the tumor and the parametria were negative, and the resection was diagnosed as complete. ① → Cervical tumor ② → Tumor that has fallen off from the cervical tumor.

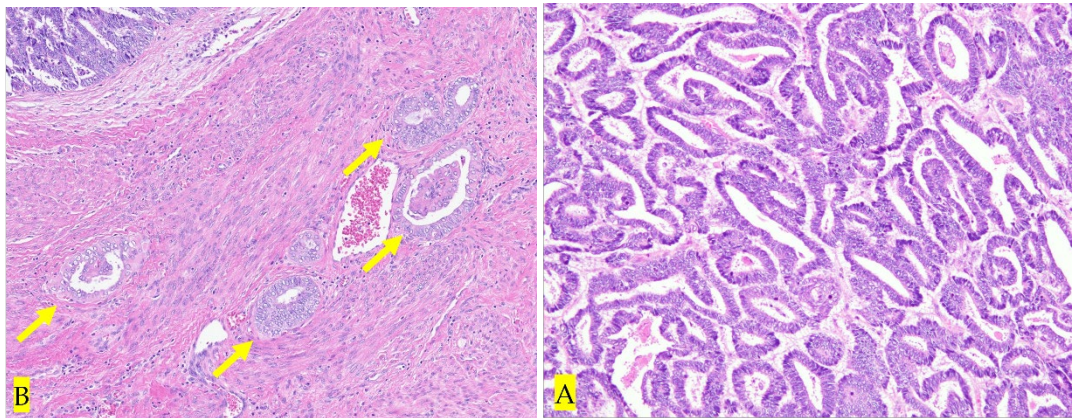
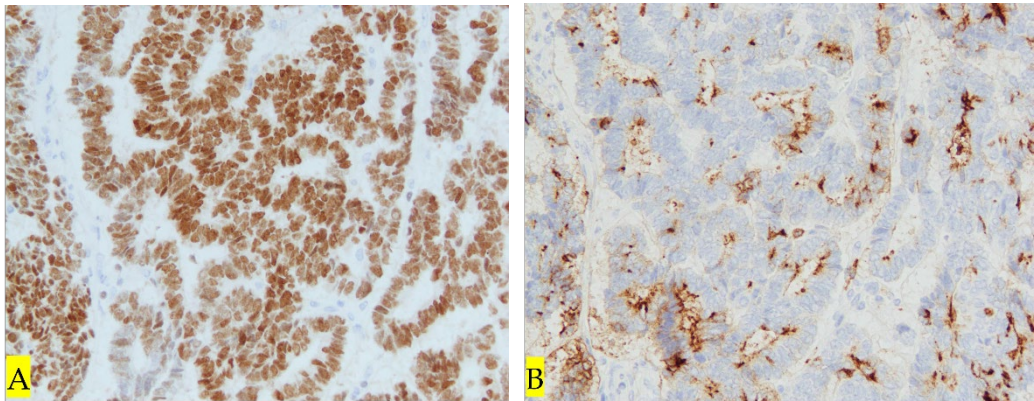


Figure 5. A HE staining 100×. Atypical cells proliferate and form a lumen that contains eosinophilic protein material. **Figure5B** HE staining 100×. Remnants of the mesonephric duct are observed (→).



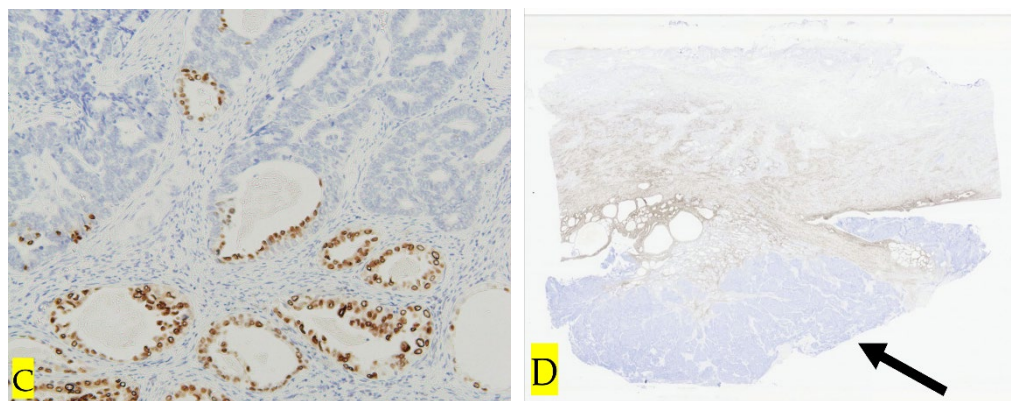


Figure 6. A Immunostaining: GATA 200×. GATA positive in tumor tissue. Figure6B Immunostaining: CD10 100×. CD10 positivity found in the stroma. Figure6C Immunostaining: TTF-1 200×. TTF-1 positive in some tumor tissues. Figure6D Immunostaining: ER 1×. ER negative in tumor tissue (→).

3. Discussion

The mesonephric duct, also known as the Wolffian duct, runs parallel to the paramesonephric duct (Mullerian duct) during early pregnancy, and in men it develops and differentiates into the male internal genital organs (epididymis and vas deferens) under the action of testosterone.

In women, unlike men, the paramesonephric ducts (Mullerian ducts) develop and differentiate to form the female internal genital organs (uterus, fallopian tubes, and upper one-third of the vagina [3]. In women, the mesonephric duct regresses, but residual tissue may be found around the ovary, within the broad ligament, uterus, and vagina [4]. Cervical mesonephric adenocarcinoma (MA) has been reported to arise mostly from the mesonephric duct located deep within the lateral wall of the cervix, but it can also arise from remnants of the mesonephric duct in the uterine body [5].

The initial symptom of cervical MA is often abnormal genital bleeding [5], and in this case, the diagnosis was made based on abnormal genital bleeding. It has been reported that the detection rate of abnormal cells in cytology is as low as 10% [5], but in this case, a solid tumor was already occupying the vagina at the time of initial diagnosis, and cervical cytology showed that AGC was diagnosed. Cervical MA has a wide variety of histological findings, and the preoperative diagnostic rate in biopsy tissue is reported to be 20% [6]. Therefore, it has been reported that it may be confused with other more common adenocarcinomas, such as serous, clear cell, or endometrial adenocarcinoma [7]. The classic histopathological pattern is the proliferation of columnar cells forming lumina containing acidophilic hyaline secretions [1]. In this case, preoperative pathological diagnosis showed that atypical cells had a glandular structure, and the case was diagnosed as cervical endometrioid carcinoma. In addition to morphological features, immunostaining is effective in diagnosing cervical MA. In tumor tissue, GATA-3, paired box 8 (PAX8), and CD10 were positive, and ER was negative. TTF-1 is rarely positive, p16 is not distributed, and HPV is usually not detected [1]. In this case, columnar cells proliferated and formed a lumen containing acidophilic hyaline-like secretions, and immunohistochemistry was positive for GATA3, PAX8, CD10, TTF-1, and negative for ER. The preoperative diagnosis was endometrioid carcinoma, but the final pathological diagnosis was MA..

There are few comprehensive reports on the prognosis of cervical MA, but Pors et al. reported the clinical characteristics in a multi-institutional study [2]. Of 30 cases of cervical MA, 25 were available for analysis; 10 cases (20%) were FIGO Stage I, and 15 cases (60%) were FIGO Stage II to IV. Lymph node metastasis was observed in 28% (4/14), and the recurrence rate was 50% (12/24), with the most common site of metastasis being the lungs 2). The 5-year progression-free survival rate was 60%, and the overall survival rate was 74%. Multivariate analysis revealed that the disease progression-free survival rate was associated with advanced stage (P value 0.04), but no association was observed with age, tumor size, or lymph node metastasis [2]. In a two-year follow-up study of cervical MA, HPV-dependent adenocarcinoma, and HPV-independent adenocarcinoma, the PFS of

HPV-dependent adenocarcinoma was favorable (P value <0.01), while no significant difference in PFS was observed between cervical MA and HPV-independent adenocarcinoma [2]. Based on these facts, cervical MA has a poorer prognosis than HPV-dependent cervical cancer, and is a pathological condition that is prone to distant metastasis, especially lung metastasis, but it is a rare disease and there is no evidence regarding treatment. There is a lack of standard treatment, and no standard treatment has been established.

We reviewed literature reporting on the treatment of cervical MA over the past 10 years from 2014 to 2024 (Table1). Neoadjuvant chemotherapy (NAC) was administered to 3 patients, including this case, and all were partial responded, and radical hysterectomy was subsequently performed. The initial treatment for all 12 patients was total hysterectomy, and as postoperative adjuvant therapy., 4 patients underwent CCRT and 1 patient received radiation therapy alone. Chemotherapy was administered in 3 patients, and no post-treatment was administered in 4 patients. The effect of NAC in improving prognosis in stage I-II cervical cancer is unclear [17]. However, NAC is being considered as an option for patients undergoing elective treatment for cervical cancer during the COVID-19 pandemic, according to the International Gynecologic Cancer Society (IGCS) and the European Society of Gynecological Oncology (ESGO) [18]. In this case, the preoperative pathological diagnosis was endometrioid carcinoma, which is HPV-independent, and radiation therapy may have been ineffective. In addition, the bulky tumor occupied the vagina and there was a lot of bleeding, so early treatment was necessary. Therefore, PC therapy was administered as NAC, and since the tumor were partial responded and bleeding decreased, a radical hysterectomy was performed after one course of chemotherapy was completed. Although the number of cases of NAC is limited, all three reported cases were successful, and therefore we believe that NAC should also be considered for cervical MA. Concerning postoperative adjuvant therapy for cervical cancer, CCRT is recommended for cases at risk of recurrence [19]. On the other hand, in a clinical trial investigating the effectiveness of radiation alone, CCRT, and chemotherapy as postoperative therapy for stage IB-II cervical adenocarcinoma, it was reported that there was no difference in survival rate among the three groups, and chemotherapy may also be effective as a postoperative adjuvant therapy [20]. Although no clear standard chemotherapy has been established for cervical MA, there have been reported cases in which PC therapy was effective in recurrent cases [21]. Considering these circumstances, in this case, because NAC had been effective and the tumor resection margins were adequately secured, chemotherapy with PC therapy was selected as postoperative adjuvant therapy.

Table 1. Literature reporting on the treatment of cervical MA over the past 10 years from 2014 to 2024.

Case	Age	Stage	NAC	operation	Adjuvant therapy	PFS months	Out come
Nili F 8)	46	I B3	—	ATH tubectomy	CCRT with CDDP	9m	Peritoneal dissemination
Reis-de-Carvalho C 9)	60	I B1	—	ARHBSO PLND	CCRT with CDDP	60m	No recurrence
Marani C 10)	58	I B2	—	ATHBSO	—	72m	Vaginal recurrence
Jiang LL 11)	48	IVB	—	ATHBSOPLND Omentectomy appendectomy	Chemotherapy	32m	Lung recurrence
Montalvo N 12)	48	I B1	—	ATH	—	36m	Lung recurrence
Papoutsis D 13)	67	II B	—	ARH BSO PLND	CCRT and brachytherapy	12m	No recurrence
Abdul-Ghafar J 14)	48	I B2	—	Vaginal hysterectomy	CCRT	24m	No recurrence
Xie C Case1 4)	50	I B1	—	ARHBSO	—	64m	No recurrence
Xie C Case2 4)	49	I B1	—	ARHBSO LND	—	70m	No recurrence
Ditto A 15)	51	II B	CDP	ARH BSO PLND	RT	6m	No recurrence
Kuratsune K 16)	30	II A2	PC 2kur	ARHBSO PLND	PC 4kur	13m	No recurrence
This Case	66	I B3	PC 1kur	ARH BSO PLND	PC 5kur	9m	No recurrence

ATH : Abdominal total hysterectomy, CCRT : Concurrent chemoradiotherapy, CDDP : Cisplatin, ARH : Abdominal radical hysterectomy, BSO: Bilateral salpingo-oophorectomy, PLND : Pelvic lymphadenectomy, LND : Lymph node dissection, CDP: Cisplatin Doxorubicin Paclitaxel, PC : Paclitaxel Carboplatin.

4. Conclusions

We report a case of cervical mesonephric adenocarcinoma in which preoperative chemotherapy was effective. Cervical MA is very rare, and preoperative diagnosis is difficult. Because it is a rare disease, there is currently no established standard treatment, but it has been suggested that PC therapy may be effective. This disease is characterized by a high incidence of distant metastasis to the lungs and other organs, and it is necessary to conduct long-term follow-up and accumulate more cases to find effective treatments.

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References

1. WHO Classification of Tumours Female Genital Tumours[M] (5th ed.), Lyon: IARC Press (2020), pp. 378-379
2. Pors J, Segura S, Chiu DS, Almadani N, Ren H, Fix DJ, Howitt BE, Kolin D, McCluggage WG, Mirkovic J, Gilks B, Park KJ, Hoang L. Clinicopathologic Characteristics of Mesonephric Adenocarcinomas and Mesonephric-like Adenocarcinomas in the Gynecologic Tract: A Multi-institutional Study. *Am J Surg Pathol*. 2021 Apr 1;45(4):pp498-506. doi: 10.1097/PAS.0000000000001612.
3. Howitt BE, Nucci MR. Mesonephric proliferations of the female genital tract. *Pathology*. 2018 Feb;50(2):pp141-150. doi: 10.1016/j.pathol.2017.11.084. Epub 2017 Dec 19.
4. Wu H, Zhang L, Cao W, Hu Y, Liu Y. Mesonephric adenocarcinoma of the uterine corpus. *Int J Clin Exp Pathol*. 2014 Sep 15;7(10):pp7012-7019.
5. Xie C, Chen Q, Shen Y. Mesonephric adenocarcinomas in female genital tract: A case series. *Medicine (Baltimore)*. 2021 Sep 3;100(35):e27174. doi: 10.1097/MD.00000000000027174.
6. McFarland M, Quick CM, McCluggage WG. Hormone receptor-negative, thyroid transcription factor 1-positive uterine and ovarian adenocarcinomas: report of a series of mesonephric-like adenocarcinomas. *Histopathology*. 2016 Jun;68(7):pp1013-1020. doi: 10.1111/his.12895. Epub 2016 Jan 4.
7. Secosan C, Balint O, Ilian A, Balan L, Balulescu L, Motoc A, Zahoi D, Grigoras D, Pirtea L. New Insights in the Diagnosis of Rare Adenocarcinoma Variants of the Cervix-Case Report and Review of Literature. *Healthcare (Basel)*. 2022 Jul 28;10(8):pp1410. doi: 10.3390/healthcare10081410.
8. Nili F, Salarvand S, Saffar H, Kalaghchi B, Ghalehtaki R. Mesonephric Adenocarcinoma of Uterine Cervix: A Case Report and Review of the Literature. *Iran J Pathol*. 2021 Spring;16(2):pp227-231. doi: 10.30699/IJP.2020.125459.2375. Epub 2020 Dec 21.
9. Reis-de-Carvalho C, Vaz-de-Macedo C, Ortiz S, Colaço A, Calhaz-Jorge C. Cervical Mesonephric Adenocarcinoma: A Case Report of a Rare Gynecological Tumor from Embryological Remains of the Female Genital Tract. *Rev Bras Ginecol Obstet*. 2021 Apr;43(4):pp329-333. doi: 10.1055/s-0041-1725051. Epub 2021 Mar 30.
10. Marani C, Akaev I, Yeoh CC, Walsh E, Rahimi S. Cervical malignant mixed mesonephric tumour: A case report with local recurrence after six-years and next-generation sequencing analysis with particular reference to the ataxia telangiectasia mutated gene. *Exp Ther Med*. 2021 Apr;21(4):pp394. doi: 10.3892/etm.2021.9825. Epub 2021 Feb 24.
11. Jiang LL, Tong DM, Feng ZY, Liu KR. Mesonephric adenocarcinoma of the uterine cervix with rare lung metastases: A case report and review of the literature. *World J Clin Cases*. 2020 May 6;8(9):pp1735-1744. doi: 10.12998/wjcc.v8.i9.1735.

12. Montalvo N, Redrobán L, Galarza D. Mesonephric adenocarcinoma of the cervix: a case report with a three-year follow-up, lung metastases, and next-generation sequencing analysis. *Diagn Pathol.* 2019 Jul 3;14(1):pp71. doi: 10.1186/s13000-019-0847-8.
13. Papoutsis D, Sahu B, Kelly J, Antonakou A. Perivascular epithelioid cell tumour and mesonephric adenocarcinoma of the uterine cervix: an unknown co-existence. *Oxf Med Case Reports.* 2019 Jan 24;2019(1):omy115. doi: 10.1093/omcr/omy115.
14. Abdul-Ghafar J, Chong Y, Han HD, Cha DS, Eom M. Mesonephric Adenocarcinoma of the Uterine Cervix Associated with Florid Mesonephric Hyperplasia: A Case Report. *J Lifestyle Med.* 2013 Sep;3(2):pp117-120. Epub 2013 Sep 30.
15. Ditto A, Martinelli F, Bogani G, Gasparri ML, Donato VD, Paolini B, Carcangiu ML, Lorusso D, Raspagliesi F. Bulky mesonephric adenocarcinoma of the uterine cervix treated with neoadjuvant chemotherapy and radical surgery: report of the first case. *Tumori.* 2016 Nov 11;102(Suppl. 2). doi: 10.5301/tj.5000386.
16. Kuratsune K, Ueda T, Tajiri R, Tohyama A, Hoshino K, Harada H, Kurita T, Kubo C, Komatsu K, Shiba E, Matsuura Y, Yoshino K. [A Case of Adenocarcinoma, HPV-independent, Mesonephric Type with Significant Response to Neoadjuvant Chemotherapy]. *J UOEH.* 2024;46(1):pp45-51. Japanese. doi: 10.7888/juoeh.46.45.
17. Zhao H, He Y, Yang SL, Zhao Q & Wu YM (2019):Neoadjuvant chemotherapy with radical surgery vs radical surgery alone for cervical cancer: a systematic review and meta-analysis. *Onco Targets Ther* 12:pp1881-1891.
18. Ramirez PT, Chiva L, Eriksson AGZ, Frumovitz M, Fagotti A, Gonzalez Martin A, Jhingran A, Pareja R. COVID-19 Global Pandemic: Options for Management of Gynecologic Cancers. *Int J Gynecol Cancer.* 2020 May;30(5):pp561-563. doi: 10.1136/ijgc-2020-001419. Epub 2020 Mar 27.
19. Rosa DD, Medeiros LR, Edelweiss ML, Pohlmann PR, Stein AT. Adjuvant platinum-based chemotherapy for early stage cervical cancer. *Cochrane Database Syst Rev.* 2012 Jun 13;6(6):CD005342. doi: 10.1002/14651858.CD005342.pub3. Update in: *Cochrane Database Syst Rev.* 2016 Nov 22;11:CD005342.
20. Shimada M, Nishimura R, Hatae M, Hiura M, Takehara K, Tase T, Yamada H, Kurachis H, Sugiyama T, Kigawa J. Comparison of adjuvant chemotherapy and radiotherapy in patients with cervical adenocarcinoma of the uterus after radical hysterectomy: SGSG/TGCU Intergroup surveillance. *Eur J Gynaecol Oncol.* 2013;34(5):pp425-428.
21. Montagut C, Mármol M, Rey V, Ordi J, Pahissa J, Rovirosa A, Gascón P, Mellado B. Activity of chemotherapy with carboplatin plus paclitaxel in a recurrent mesonephric adenocarcinoma of the uterine corpus. *Gynecol Oncol.* 2003 Aug;90(2):pp458-461. doi: 10.1016/s0090-8258(03)00228-2.

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