Primary Female Urethral Carcinoma A Case Report. A Review of the Diagnosis and Treatment

Abstract

Primary urethral carcinoma is a very rare pathology that makes up less than 1% of all genitourinary tumors [1,2]. There are several cases report published in the medical literature, although there is no consensus for the management. The prognosis is poor when diagnosed in advanced stages. Treatment includes local or radical surgery with pelvic lymph node dissection, radiotherapy, brachytherapy, and neo or adjuvant chemotherapy. Many authors have suggested a combination of surgery, chemosensibilization and radiotherapy [1,3].

Keywords: Urethral carcinoma; Diagnosis; Tumors

Case

A 54-year old white woman, without significant past medical history other than menopause since 2 years ago presented with post coital hematuria and also she had urinary tract infections (UTIs), documented by urine cultures (Proteus Mirabilis). No other associated symptoms. She received antibiotic therapy with quinolones. The urine cytology was negative. The assessment by the Gynecology service was normal. The renal ultrasound (US) Each kidney was normal, preserved cortico medullary differentiation and without dilated collecting systems. There was irregularity zone in the mucosa close to bladder neck, measuring 1 x 0.5 cm. Cystoscopy revealed an edematous and erythematosus lesion of 1 x 1.5 cm in the posterior wall of the bladder neck that extended into the urethra. In pelvic magnetic resonance imaging (MRI) a pelvic mass was evident measuring 35x 38x 47mm, widening the bladder neck at this level, nonspecific inguinal adenopathies and a left iliac node with malignancy aspect.

Initial US

The pelvic examination during the transurethral resection of bladder (TURB) revealed a palpable nodule on the anterior wall of the vagina and there was not others remarkable findings. The pathology of transurethral resection of bladder confirmed urethra adenocarcinoma muscle-invasive with invasion of bladder neck. She was taken to an extension study (toracoabdominopelvic scan) in which the metastatic lesions were discarded (Figure 1-3).

Radical cystourethrectomy was performed with bilateral pelvic lymph node (extense iliac region) dissection (Figure 4) and anterior vaginal wall excision (vaginoplasty) followed by ileal urinary diversion (Bricker type). Ureteral end edges were sent to frozen biopsy and the results were negative. The specimen’s pathology was “adenocarcinoma with clear cell areas and focus of papillary, located in the trigone region, affecting the peri vesical tissues without infiltrating the muscular layer of the vagina (pT4a). Bilateral pelvic lymph nodes were free of tumor (12 nodes in the left side and 14 in the right side), like the rest of organs. Neoplasia stage: pT4a pN0 pM0.”

Figure 1: Initial pelvic ultrasound.

Figure 2: Initial pelvic ultrasound.

Figure 3: Initial pelvic ultrasound.
After a month postoperative follow-up she was asymptomatic, no findings in the physical examination. So along with Oncologists, she started adjuvant chemotherapy with cisplatine/Taxol/Gemcitabine (3 cycles) during one month. No other complementary treatment. During one year she had oncologic remission with toracoabdominopelvic scans and physical examinations every 6 months without changes. Twenty months after the surgery in the control scan showed a 42 x 30 mm mass in the right pelvic lymphadenectomy area with ilion infiltration. She also had anemia secondary digestive bleeding. General surgery dismissed curative treatment, for the lesion’s localization and the infiltration of piri for muscle (right catic nerve compression). She started chemotherapy CDDP/Gem (6 cycles) and right hemi pelvic radiotherapy (18 sessions, total doses 4500cGy) for pain control. Nowadays, she is finishing the oncology treatment, does not need blood transfusion and also the right leg pain improved.

Discussion

Urethra carcinomas are a very rare pathology (less than 0.02% in female tumors); most of them are transitional cell carcinoma (55%), followed by squamous cell carcinoma (21%) and adenocarcinomas (16.4%) [4]. The age of onset is over the 50s and the most frequent symptoms are bleeding, infections, urethral diverticulum. This pathology is 4 times more common in women compared to men [5,6].

There have been a few reports of urethral carcinomas (Figure 5) diagnosed within dilated paraurethral Skene glands (diverticul), half of them being adenocarcinomas [7]. Some authors have proposed that the urethra adenocarcinomas may arise from urethritis glandularis, Mullein ducts (paramesonephric ducts), Skene’s glands (paraurethral glands), or mixed origins. The most frequent histological subtype is columnar/mucinous adenocarcinoma and there are around 15% of dear cell of them [1,3].

Tumors origins in distal paraurethral duct come from female Skene’s glands (homogeneity of male prostate gland) are matched by positive immunostains of PSA and prostatic acid phosphatase. PSA elevation is objective and declines after surgery. The tumors origins in proximal paraurethral ducts can produce carcinoembryonic antigen (CEA). The immuno histo chemical study can be helpful identifying the type of adenocarcinoma [1,8].
The diagnosis of this condition can be delayed because of vague and nonspecific symptoms such as hematuria, genital mass feeling, increased urinary frequency, dysuria and urethral obstruction. Urine tract infection with no answer to antibiotic treatment and urine incontinence are less frequently seen as clinical manifestations [1]. The diagnosis made with urethrocystoscopy and lesion biopsy. The Scan and MRI are made for local and distal extension.

The most important factor for the prognosis/ survival is disease stage, in the early stages the survival in 5 years is 50%, in advance disease the survival in 3 years 56% and there is no survival in 5 years [5,6]. In the Netherlands, an epidemiological study based on data from the National Cancer Registry during the years 1989 to 1998, concluded that survival is determined by factors such as the TNM staging and histological type of tumor. The prognosis is poor in advanced stages, when most of them are diagnosed. It has not been possible to establish the most effective treatment for the type of tumor (better prognosis for squamous type and worse if is adenocarcinomas), so the current guidelines are extrapolated from other pathologies [4]. Other survival factor is the lesion’s size, less 2 cm survival is 60% 2-4 cm is 46% and when lesion is more than 4 cm the survival is only 13% [5,6].

The surgical treatment includes a range from local excisions to anterior pelvic extirpation [9,10]. The local excisions or urethrectomies resulted in postoperative complications, local recurrence or distal metastasis and aggressive treatment as anterior extirpation (specially the posterior urethral lesion), radical cystectomy with urinary diversion and pelvic/groin lymphadenectomy showed promising results with 87% of recurrence disease-free at 6 months to 2 years [3,11,6]. In anterior urethral lesions preferred wide extirpation (total or parcialurethrectomy) and if a superficial lesion is, can be treated with local extirpation [5,12].

For tumors less than 2cm curative radiation or surgery with or without adjuvant radiotherapy have been proposed as treatment options, whereas; bulky injury is being treated with brachy (BR) / external beam radiotherapy (RE)(45-50 Gy) as a conservative organ therapeutic and if there are nodal disease the doses are 60-65 Gy [2,5,13]. Moinuddin Ali and Cols [14], treated three patients in early state with RE (40-45 Gy) and extra doses with BR (28-30Gy) presented recurrence disease-free after 30 months. RT as monotherapy could be optimal treatment in early state of distal urethral tumors [10]. For locally advanced tumors, combined treatment with neo or adjuvant radiotherapy, surgery and chemotherapy, has been associated to cyto reduction [4,2,10]. There are some series with urethra carcinoma advance state with unfavorable results with surgery or RT as monotherapy [10].

Dalbagniand Cols, proposed adjuvant RT as part of the surgical treatment with the mean goal to reduce the local recurrence, which improve the disease- specific survival. In the same way, the authors published improvement in the recurrence disease-free after 5 year 37% with neoadjuvant [12]. A retrospective cohort study has shown that treatment with chemotherapy is beneficial, especially when is given prior to surgery in patients with positive nodes, as it increases disease-free survival [15]. There are some reports that shown combination therapy with several chemotherapeutic agents in locally advanced female urethral adenocarcinomas like cisplatin- based or cisplatin with 5-fluorouracil or with Carboplatin, plaquitaxel and irinotecan with adjuvant radiotherapy with good results [1,4,11,16,17].

It has proven the Douillard regimen (irinotecan, fluorouracil and leucovorin) in urethral adenocarcinomas and is also used in colon carcinoma and has had evidence in uraco carcinoma as well [1]. There is a series of 72 women treated at Memorial Sloan-Kettering in a period of 36 years, the multivariate analyse showed as in independent risk factors for survival the initial stage of tumor, the presence or absence of regional lymph nodes and the site of the disease (proximal versus distal urethra). Being a better prognosis small masses (less than 2 cm) in distal urethra, this kind of tumors could be treated with single modality treatment, on the other hand, multimodal treatments has been recommended for larger and proximal tumors [7,18-20].

Conclusion

Does the treatment of urethra carcinoma early states T1 or T2 is surgery with pelvic lymph node dissection and adjuvant RT? Does the treatment of the primary urethral cancer is RT? And the early local recurrence should be removed? Does the chemotherapy is the first line treatment in the primary urethra cancer or should be used with RT or adjuvant treatment in advance state?

The treatment of urethral carcinoma is still controversial, it is a rare condition and there are not enough published retrospective studies, non-prospective studies to clarify. Despite there is not guidelines or protocols for the urethra adenocarcinoma, the combination therapy seems to be the most effective. Besides the histology tumor, the size, location and disease stage are prognostic factors in female urethra carcinoma.

References


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