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Case Report

Nephrogenicadenoma of the Urinary bladder Faketumor or Witchhunt?

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Abstract

The optimal management of the nephrogenicadenoma of the bladder remains one of the concerns of the scientific community. The diagnosis is often misleading with other malignant lesions of the bladder and its lack of knowledge can lead to disastrous medical mistakes.

We report the clinical case of a 65-year-old man hospitalized for cripplingpollakiuriawithhematuria. Ultrasonography found a left lateral intravesical mass with ipsilateral kidney a large pyelocalicielle dilatation and laminated parenchyma Trans-urethral resection of the intra-vesical tumorbud found a nephrogenicadenoma of the bladder. Supra-trigonal enlargement enterocystoplasty with left nephrectomy was recommended. The patient did not have local recurrence at cystoscopic control, with a 20-month follow-up.

Through this clinical case and a literature review, we analyze the epidemiological, aetiology diagnostic and therapeutic aspects of this rareentity.

Introduction

The nephrogenicadenoma of the bladder (NAUB) is a rare lesion [1]. Easily diagnosed as a malignant lesion, mis understanding of this entity could lead to many medical faults. A number of irritative factors have been in criminated in ethiopathogeny: recurrent infections, relapsing stones, intra-vesical instillations, foreign bodies, radiotherapy ... but none is un animous [2]

Through this case of NAUB we will analyze the different etiopathogenic and therapeutic aspects of this often unknown entity.

Case Report

We report the case of a 65-year-old patient, followed in our training for disabling pneumakiuria with an episode of hematuria. A renovesical ultra sound demonstrated a suspicious budding lesion of the bladder wall, with reduced bladder capacity and bilateral pyelocalicar dilatation laminating the left renal parenchyma. Biologically, he had s ever e renal impairment with a glomerular filtration rate of 22 ml / min / 1.73m³. Trans-urethral resection of the intra-vesical tumorbud seen on ultrasound re turned to a NAUB (Figure 2-3). The thoracoabdominopelvic CT, performed as part of the extension assessment, did not find lymphadenopathy or secondary locations (Figure 1). Renal scintigraphy with DMSA showed a functional value reduced to 5% of the left renal unit.

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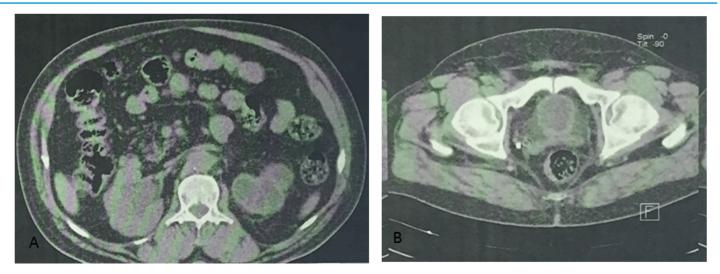


Figure 1. CT cross section.

(A): Bilateralpyelocalicielle dilation and laminating the leftrenalparenchyma.

(B): Bladderwiththickwall and lowcapacity.

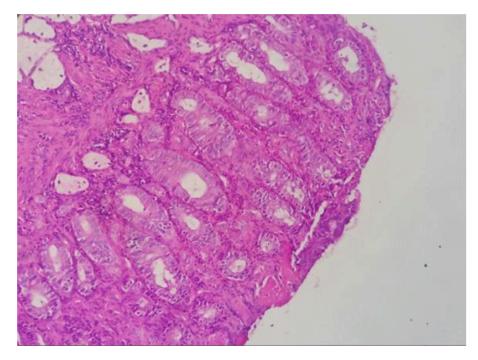


Figure 2. Histological section at medium magnification showing tubular structures surrounded by epitheloid cells (HE.200).

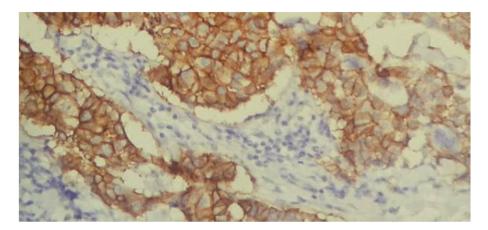


Figure 3. Immunohistochemical study: Positive labeling of tubular structures by anti-CK 7 antibody.

After multidisciplinary consultation, it was decided to perform a trigonal enlargementent erocystoplasty with left nephrectomy.

Nephrectomy was performed laparoscopically, and median umbilical laparotomy was recommended for entericcystoplasty with supratrigonal enlargement.

The postoperative course was simple. The evolution was marked by the improvement of the of lower urinary tractsymptoms and renal function.

The patient did not have local recurrence at cystoscopic control, with a 20-month follow-up.

Discussion

Although the nephrogenicadenoma of the bladder (NAUB) was first described by David in 1949, it was only a year later that Friedman and Kuhlenbeck were named "nephrogenicadenoma" because of the similarity to tubular cells. kidney [3, 4]. It is a metaplasia of the bladder with papillary or cryptic structures composed of small hollow tubules resembling the mesonephric tubules. There is often a layer of cuboid or studded cells surrounding the tubular cells [5]. Other alternative terminologies have been used to describethisentity: mesonephricmetaplasia, adenomatoid tumor and adenomatoidmetaplasia [5].

NAUB is more common in men than in women (2/3 in men) [6]. It is the preserve of the adult even if rare cases in children have been reported [7].

The NAUB is ubiquitous throughout the urinary shaft often identified in the bladder, but it can be ubiquitous throughout the urinary shaft. In the litterature, It is most often located in the bladder neck and the adjacent urethra [8].

Although now a days there are many hypotheses for etiopathogeny, the first one was developed in 1954 by Most of iwho reported that squamous and glandular metaplasia of the urothelium was frequently associated with chronic infection [2].

Thus, the literature incriminates intravesical instillations, stones, chronic catheterization, exstrophy, interstitial cystitis and surgery. At the same time, thereis an increased incidence of nephrogenic adenomas in transplant patients [9].

Our patient was followed for chronic cystitis for 3 years and was probably the implantation bed of nephrogenicadenoma.

In addition, recent studies have shown a link between nephrogenicadenoma and certain drugs, such as libuprofen and phenacetin [10]. Our patient was also on diclofenac NSAIDs of 150 mg daily, but intermittently and limited in time.

Clinical Presentation of the NAUB is not specific. It tends to present with lower urinary tract irritative symptoms such as: increased frequency of urination, increased urgency of urination, urge incontinence and excessive passage of urine at nigh. It rarely presents with haematuria.

Cystoscopy, found so litary or multiple lesions, often papillary or polypoid. However flat lesions have also been reported. This appearancemimics a malignant tumor and can lead to medical errors. In addition, flat lesions may also be confused with urothelial carcinoma in situ [11]. It is important to note that the literature reports rare cases of malignant transformation that should not be ignored and that completely change the prognosis [12].

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Histologically, the nephrogenicadenoma has several different histological profiles: tubular, tubulocystic, polypoid, papillary, fibromyxoid, and flat [13]. This morphological variability, it can mimicurothelial carcinoma, metastatic tumors or clear cell adenocarcinoma. However, unlike malignant tumors, nephrogenic adenomas are inactive at the mitotic level and have no atypia or risk of invasion of the muscularis. There is no evidence of necrosis and no desmoplasia.

Finally, they are often positive for PAX2, PAX8, cytokeratin 7, high molecular weight cytokeratins and a-methyl-acyl-coenzyme A racemase (AMACR), CD10 et MUC. (14-17).

Cytogenetic s studies found that monosomy 9 and trisomy 7 have been associated with this entity [18].

A conservative attitude with trans-urethral resection of the NAUB is the optimal strategy. Partial cystectomy can be preeconised for large tumors. Some authors propose the use of hyaluronic acid instilled in the bladder [19]. In our case, the trans-urethral resection of the bladder allowed us to specify the histological nature of the tumor. Enterocystoplasty on trigonal enlargement was indicated in front of reduced bladder capacity and repercussion on the upper urinary tract.

Because of the rare reported case of malignant transformation, a long-term follow-up with cystoscopy should be recommended.

Conclusion

Nephrogenicadenoma is a rare un common benign lesion of the urinary system that occurs in patients with chronic urinary tract infection or in transplant patients. We report a case of nephrogenic adenoma of the bladder in a 65-year-old man with chronic cystitis treated with ibuprofen. Through this case we wish to remind the existence of this benignentity to avoid medical errors that could be disastrous.

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