

A Rare Non-Functional Bladder Paraganglioma Managed by Transurethral Resection

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Abstract

Background: Paraganglioma of the urinary bladder is rarely encountered. Its clinical signs and symptoms are usually non-specific.

Case Presentation: We report a case of a 48-year-old woman who presented with recurrent hematuria for last 2 months with a recently detected hypertension. Computed tomography (CT) revealed a lobulated mass arising from the posterior wall of the urinary bladder and protruding into the vesical lumen measuring 3cm with inhomogeneous postcontrast enhancement. Transurethral resection of the mass has instantly produced a hypertensive peak (200/110 mmHg) leading to an interruption of operation. Pathology of the specimen found a pheochromocytoma. Our patient denied a history of dizziness, sweating, and palpitations during micturition. Hormonal evaluation showed a normal level of both plasmatic and urinary metanephrine and normetanephrine. Metastatic workup was negative. Both adrenal glands were normal at CT. MIBG scan showed an increased localized activity in the bladder. After preoperative stabilization of hypertension and medical preparation, a complete transurethral resection of the tumor was performed. This case was discussed at a tumor board meeting and surveillance was decided. The patient has been disease-free for 9 months.

Conclusion: In patients with unusual, striking symptoms and an increase in blood pressure with micturition, diagnosis of pheochromocytoma of the urinary bladder must be considered. Imaging will help in confirming the diagnosis and localizing the tumor. Surgical removal is the treatment of choice.

Keywords: Pheochromocytoma; Bladder

Introduction

Paraganglioma of the urinary bladder (PUB) rarely encountered. Pheochromocytoma that arise outside the adrenal gland is called paraganglioma [1]. It is a neuroendocrine neoplasm arising from chromaffin cells located in the muscle of the bladder wall muscle [2]. Its clinical expression is paroxysmal associating sweating, headache and tachycardia [3]. It can be non-functional or functional by hormonal screening. Establishing the diagnosis may be difficult. Histological examination is often the only key leading to diagnosis [2]. Its treatment is surgical.

We report the clinical features and pathological characteristics of a patient diagnosed as paraganglioma of the urinary bladder managed by monobloc transurethral resection.

Case Presentation

We report a case of a 48-year-old woman who presented with recurrent hematuria for last 2 months with a recently detected hypertension. Computed tomography (CT) revealed a mass arising from the posterior wall of the urinary bladder measuring 3cm with inhomogeneous contrast enhancement (Figure 1). Cystoscopy showed a mass covered by smooth vesical mucosa, (Figure 2). Transurethral resection of the mass has

instantly produced a hypertensive peak (200/110 mmHg) leading to an interruption of operation. Pathology of the specimen has confirmed the diagnosis of a bladder paraganglioma (Figure 3).

Our patient denied a history of dizziness, sweating, and palpitations during micturition. Hormonal evaluation showed normal levels of plasmatic and urinary metanephrine and normetanephrine (VMA, HVA). Metastatic workup was negative. Adrenal glands were normal at CT. MIBG scan showed an increased isolated activity in the bladder. After preoperative stabilization of hypertension and medical preparation using alpha-blockers, a complete transurethral resection of the tumor has been performed (Figure 4).

The patient was discharged two days after the operation without any postoperative complications. This case was discussed at a tumor board meeting and surveillance was decided. The patient has been disease-free for 9 months.

During the postoperative follow-up, our patient was completely asymptomatic. It presented no more hematuria and neither high blood pressure. Post-operative MIBG was normal as well as the metanephrines.



Figure 1: Transverse CT-scan image showing a 3cm mass in the posterior urinary bladder wall

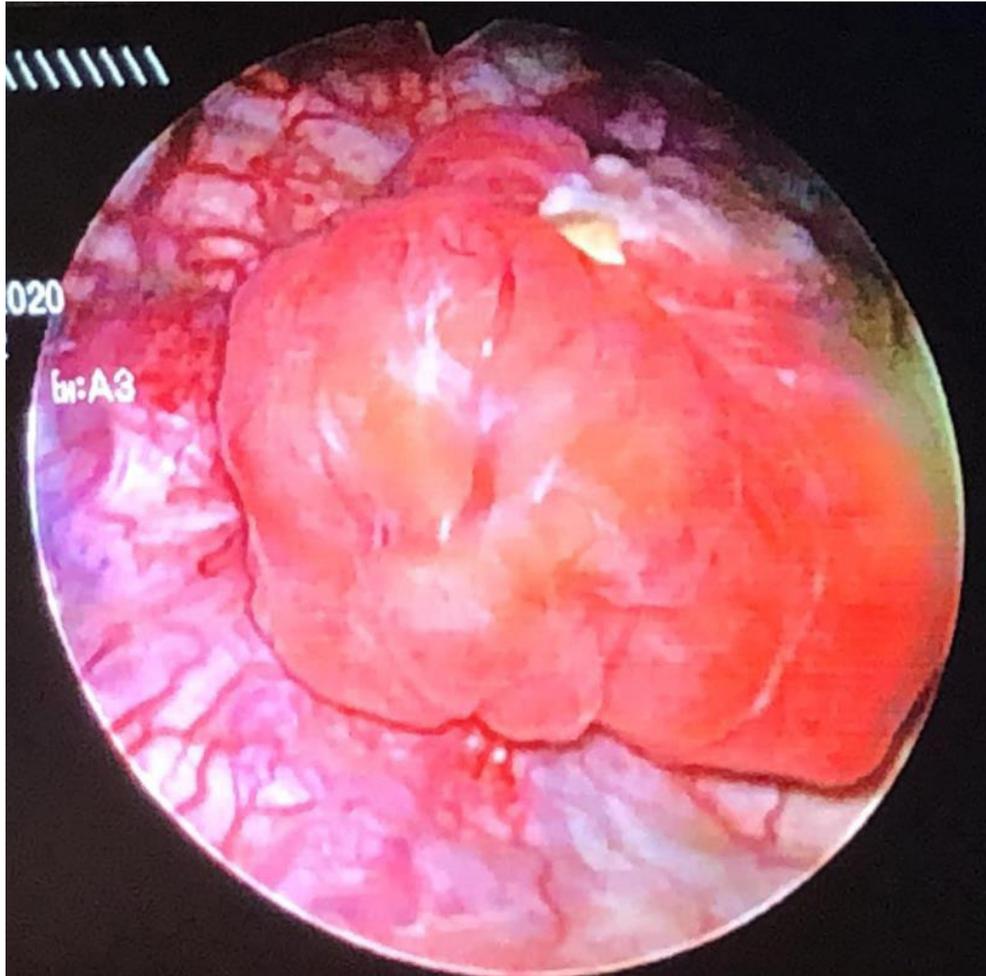


Figure 2: Endoscopic image of PUB showing a mass covered by smooth vesical mucosa

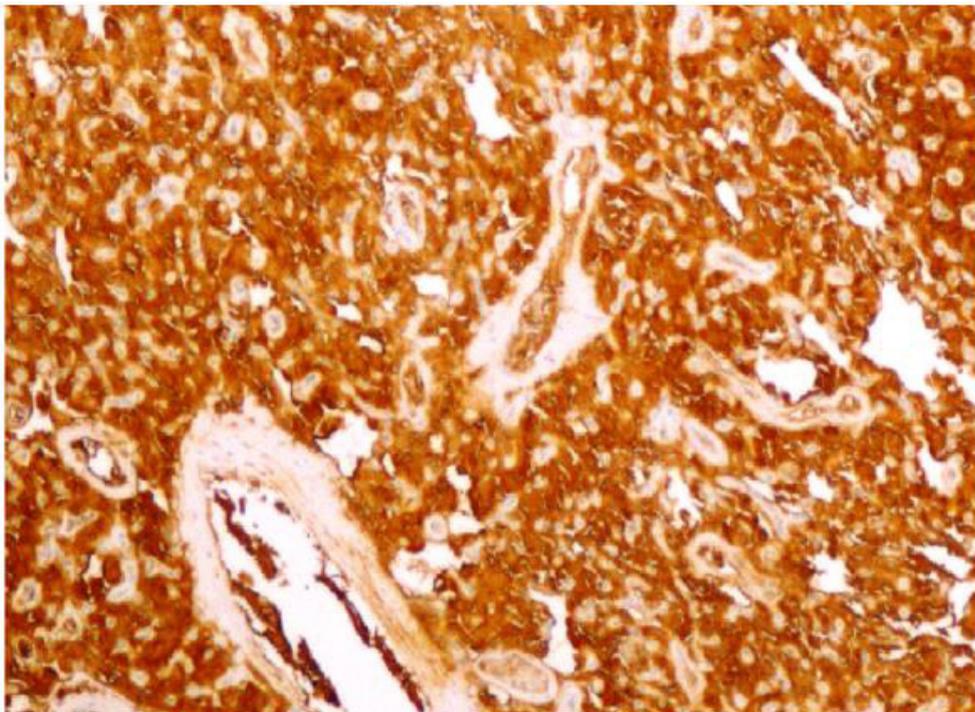


Figure 3: Tumor cells are strongly stained with synaptophysin ($\times 200$)



Figure 4: Macroscopic view of the resected bladder paraganglioma. the tumor is red-brown in color, lobulated and well-circumscribed

Discussion

Extra-adrenal localization of pheochromocytoma is rarely encountered [1,4], it is found in 10–15% of cases [5]. The first case with a bladder localization was described by Zimmerman in 1953 [6]. In the literature, fewer than 200 cases that has been described [7].

The average age of predilection is 40 years with extremes ranging from 10 to 74 years [1]. Women are more concerned than men [8]. Functional paragangliomas secrete catecholamines, which may lead to symptoms such as palpitations, dizziness, and sweating. These symptoms are mainly precipitated by micturition or overdistention of the bladder, defecation, sexual activity or any bladder instrumentation such as cystoscopy or transurethral resection [8,9].

Imaging findings are indistinguishable from other tumor types. The Meta-iodo-benzyl-guanidine (MIBG) scan is regarded as high specific for paraganglioma. However, its use is limited because of its expense and restricted availability [10]. In our patient, MIBG scan showed an increased localized activity in the bladder with no other localisations.

In some cases, such as in our case, urinary catecholamine

levels may be normal if the tumors are small at the time of diagnosis and because catecholamine secretion may increase for only brief periods during contraction of a distended urinary bladder [11].

In our case, if the PUB was suspected when transurethral resection of the mass has instantly produced a hypertensive peak (200/110 mmHg). Metanephrines levels were normal, metastatic workup was negative and both adrenal glands were normal.

There is no standard treatment for PUB. The benefit of chemotherapy and radiotherapy is not well established [3]. Surgical removal is the treatment of choice [3]. Various surgical options are available including transurethral resection, partial cystectomy and radical cystectomy. Some specialists indicated that partial cystectomy is the mainstream treatment [12]. To avoid hypertensive spikes during the mass manipulation, preoperative treatment with alpha-blockers is required, started 1 to 3 weeks before surgery [13]. Katiyar et al. [14] reported two non-functional PUB patients who underwent transurethral bladder resection tumor in which no recurrence after follow-up for 6 and 10 months, indicating that TURBT is an effective treatment for non-functional PUB. In contrast, it has been suggested that resection rarely excises all the tumor as the neoplasms often invade the submucosa and muscularis of the urinary bladder

leading to potential tumor recurrence [15].

We successfully treated our patient with a trans-urethral resection. No recurrence or metastasis were found at 9-months follow-up. Therefore, the complete resection of the tumors through the transurethral approach seems to be a curative option and may represent the mainstream future treatment [14].

In general, PUB is a rare tumor with a good prognosis and slow development [15]. However, paraganglioma has a tendency for recurrence and metastasis.

Conclusion

In patients with unusual, striking symptoms and an increase in blood pressure with micturition, diagnosis of paraganglioma of the urinary bladder must be considered. Imaging will help in confirming the diagnosis and localizing the tumor. Surgical removal is the treatment of choice. Complete transurethral resection is a safe and efficient treatment that serves both diagnostic and therapeutic purposes. Long-term follow-up is recommended.

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