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Renal Cell Carcinoma: A Case Report and Literature Review

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Abstract

To publish the various clinical presentations that may manifest this type of pathology through the case of a 58-year-old male who comes to the clinic for weight loss of 10 kilos in 1 month and pain in the left renal fossa which radiates to the left iliac fossa and left testicle. Imaging studies were performed, and he found a large left kidney tumour which in turn was causing a varicocele confirmed by testicular ultrasound. The treatment of choice was open radical nephrectomy. It is extremely important to determine and prevent risk factors for patients who may develop this pathology, as well as to identify the paraneoplastic symptoms that can guide us to a more accurate diagnosis, as well as rule out differential diagnoses.

Keywords: Renal Cancer; Nephrectomy; Risk factors; Clear cell carcinoma; Renal cell carcinoma; Varicocele

1. Introduction

Renal cell carcinoma is a common entity in the general population and is the mostlethal of all urological cancers [1]. They account for 2% to 3% of them. The age of predominance of this pathology ranges between 55 and 75 years of age and they are almost twice as common in men as in women [1-3]. The most common type of these tumours is adenocarcinoma or also known as renal cell tumour, 9out of 10 tumours belong to this type. Of this type, the most prevalent type are clear cell tumours [4].

This clinical entity is of utmost relevance because its risk factors are increasingly present in our society, the most common of which are age, race (more common in African people), smoking (doubles the chance of developing it), obesity, hypertension and lately diabetes mellitus 1 and 2 have been associated with an increased risk of it [5,6].

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The triad of renal fossa pain, macroscopic haematuria and palpable abdominal mass is becoming less frequent nowadays [7]. The main reason for patient consultation is increasingly due to paraneoplastic symptoms, such as high blood pressure, cachexia, weight loss and anaemia in some cases [7,8].

The diagnosis is made clinically by correlating it with the findings of various imaging studies such as CT and MRI scans [9,10]. The treatment for almost any renal tumour of considerable size is nephrectomy, either partial or total [7,8,11].

2. Clinical Case

Male patient aged 58 years, type 2 diabetes mellitus for 25 years treated with metformin, systemic arterial hypertension for 8 months treated with losartan, positive smoking, smoking 15 cigarettes a day without mentioning how long it has been going on, alcoholism denied.

He attended the urology department due to stabbing pain in the left renal fossa radiating to the left iliac fossa and to the left testicle, in addition to a weight loss of 10 kilos in the last month.

Physical examination corroborated the pain radiating to the left iliac fossa and to the left testicle, with a score of 7 on the visual analogue scale.

Laboratory studies with blood biometry, liver function tests and serum electrolytes within normal. General urine examination revealed proteinuria and glycosuria, renal function tests reported urea 72.2, blood urea nitrogen 33.7, creatinine 2.07 and a glomerular filtration rate of 34.3 ml/min/1.72 m² which places him in stage 3B of the Glomerular Filtration Rate interpretation criteria. Testicular ultrasonography showed the presence of a cyst in the head of the epididymis in the right testicle and a varicocele in the left testicle (FIG. 1), the latter due to continuous pressure from the left renal vein.



FIG. 1. Ultrasonography of the testicles. The right testicle with an occupying lesion in the right testicle with cystic echostructure in the epididymis and the left testicle with varicocele.

In the abdominopelvic tomography the left kidney was observed with a significant increase inits dimensions, at the time of the study of $14.79 \times 8.88 \times 9.85$ in its longitudinal, anteroposterior and transverse axes respectively. As well as significant striation of the perirenal fat and renal hilum fat. Retroperitoneal and mesenteric nodes were normal (FIG. 2).

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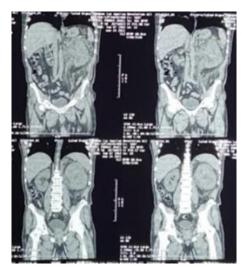


FIG. 2. Coronal abdominopelvic CT scan showing marked enlargement of the left kidney.

In the MRI of the abdomen, the left kidney showed an apparent tumour mass in the upper poleand infiltration in the lower pole, with areas of cystic degeneration. It also showed hypointenseADC maps, which is suggestive of a diffuse tumour infiltration of the kidney. Right kidney with involvement of the perirenal fat due to change of a right pyelonephritis.

The diagnosis was a left renal tumour which underwent surgery for a left radical nephrectomyfinding a tumour of approximately 15 cm.

During the surgical procedure, the left kidney was removed without any problems, and the kidney was weighed, giving a total weight of 1.305 kg (FIG. 3 and 4).



FIG. 3. Total weight of tumour removed.



FIG. 4. Tumour removed.

The nephrectomy specimen was sent to pathology, and it was reported that the tumour invades calyxes, renal pelvis, ureter and renal vein with a medium increase, and there is also contact with multiple areas of invasion to the same extent towards the perihilar adipose tissue, anterior fascia of Gerota and retroperitoneal adipose tissue.

Microscopic examination showed large, round, polygonal cells with well-defined cytoplasmic borders, in some areas with abundant, clear cytoplasm, central nucleus with variable nuclear atypia and prominent nucleoli, in some cells the nuclei are small (8-10 microns) with inconspicuous nucleoli (GradeII), in others the nuclei are pleomorphic, large and irregular with prominent reddish nucleoli (Grade IV).

In the areas adjacent to the necrotic areas, discohesive, pleomorphic cells, some spindle-shaped, others epithelioid with sparse dense eosinophilic cytoplasm, some intracytoplasmic inclusions are observed. Hematoxylin and eosin stain was used in this pathological study (FIG. 5).

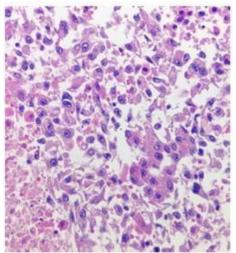


FIG. 5. Cut at ×100 magnification.

3. Discussion

Renal cell tumours are an extremely frequent entity in urological consultations [1,2]. Clear cell tumours are the most common, accounting for 90% of these tumours [4]. Early identification of risk factors in our patients as well as early diagnosis makes patient survival even higher [3,4].

Furthermore, it should not only be limited to exploring the renal system, as the clinical manifestations can range from the genitals (varicocele) to distant metastases (lung, bone, liver). It is therefore important to always look for organ involvement that may occur alongside the main disease with the help of imaging studies. The importance of a multidisciplinary team forboth diagnosis and treatment is key to a better prognosis for patients [9,10].

4. Conclusion

The selection of patients who will undergo this type of procedure must be careful, multidisciplinary management is essential to maintain the optimal conditions possible for each case, since the risks are usually high, it is a fact that the diameter of the distal bile duct it is the most important predictive factor when selecting the diameter of the balloon to use, since excessive dilation increases the risk of perforation.

5. Declarations

Conflict of interest: There is no conflict of interest.

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