

Case report

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Primitive Neuroectodermal Tumor (PNET) of the kidney: a case report

Giorgio Pomara*¹, Francesco Cappello², Maria G Cuttano¹,
Francesca Rappa², Girolamo Morelli¹, Pierantonio Mancini¹ and Cesare Selli¹

Address: ¹Department of Surgery – Urology Section – S. Chiara Hospital – University of Pisa, Pisa – Italy and ²Department of Experimental Medicine – Human Anatomy Section – University of Palermo, Palermo – Italy

Email: Giorgio Pomara* - g.pomara@libero.it; Francesco Cappello - francapp@hotmail.com; Maria G Cuttano - mg.cuttano@med.unipi.it; Francesca Rappa - francapp@hotmail.com; Girolamo Morelli - g.pomara@libero.it; Pierantonio Mancini - g.pomara@libero.it; Cesare Selli - c.selli@dc.med.unipi.it

* Corresponding author

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Abstract

Background: A case of Primitive Neuroectodermal Tumor (PNET) of the kidney in a 27-year-old woman is presented. Few cases are reported in the literature with a variable, nonspecific presentation and an aggressive behaviour. In our case, a radical nephrectomy with lymphadenectomy was performed and there was no residual or recurrent tumour at 24-month follow-up.

Methods: The surgical specimens were formalin-fixed and paraffin embedded. The sections were stained with routinary H&E. Immunohistochemistry was performed.

Results: The immunohistochemical evaluation revealed a diffuse CD99 positivity in the cytoplasm of the neoplastic cells. Pankeratin, cytokeratin AE1/AE3, vimentin, desmin, S100, cromogranin were negative. The clinical presentation and the macroscopic aspect, together with the histological pattern, the cytological characteristic and the cellular immunophenotype addressed the diagnosis towards primary PNET of kidney.

Conclusions: Since sometimes it is difficult to discriminate between PNET and Ewing's tumour, we reviewed the difficulties in differential diagnosis. These tumors have a common precursor but the stage of differentiation in which it is blocked is probably different. This could also explain their different biological behaviour and prognosis.

Background

The peripheral Primitive Neuroectodermal Tumor (PNET), firstly recognized by Arthur Purdy Stout in 1918, is a member of the family of "small round-cell tumors". Primitive renal localization is very rare. There are almost 50 cases reported in the literature, although it is difficult

to estimate the exact number since often it has not been differentiated from Ewing's Sarcoma [1-13]. Renal PNET is more aggressive than in the other sites. It frequently arises during childhood or adolescence, having an aggressive clinical course towards metastatic disease and death. It often recurs locally and metastasises early to regional

lymph nodes, lungs, liver, bone and bone marrow, resulting in a poor prognosis. The 5-year disease-free survival rate, for patients presenting well confined extra-skeletal PNET, is around 45–55% and cases with advanced disease at presentation have a median relapse-free survival of only 2 years [1].

Case presentation

A 27-year-old woman was referred because of a mild left flank pain and haematuria. Ultrasonography identified a left renal mass homogeneously hyperechogenic in comparison with renal parenchyma. CT scan showed a 11 mm × 8 mm × 6 mm tumor replacing the upper half of the left kidney with extension into the renal vein. Chest x-ray was

negative. Pathological stage after radical nephrectomy was T3aN0Mx.

The surgical specimens were formalin-fixed and paraffin embedded. The sections were stained with routine H&E. Immunohistochemistry was performed using avidin biotin complex technique and diaminobenzidine as chromogen. The antibodies used included CD99 (Dako, M3601), pankeratin (Dako, M0821), cytokeratin AE1/AE3 (Dako, M3515), vimentin (Dako, M7010), desmin (Dako, M0760), S100 (Dako, Z0311), and chromogranin A (Dako, M0869), at suggested dilution. We performed also appropriate routinely positive and negative controls.

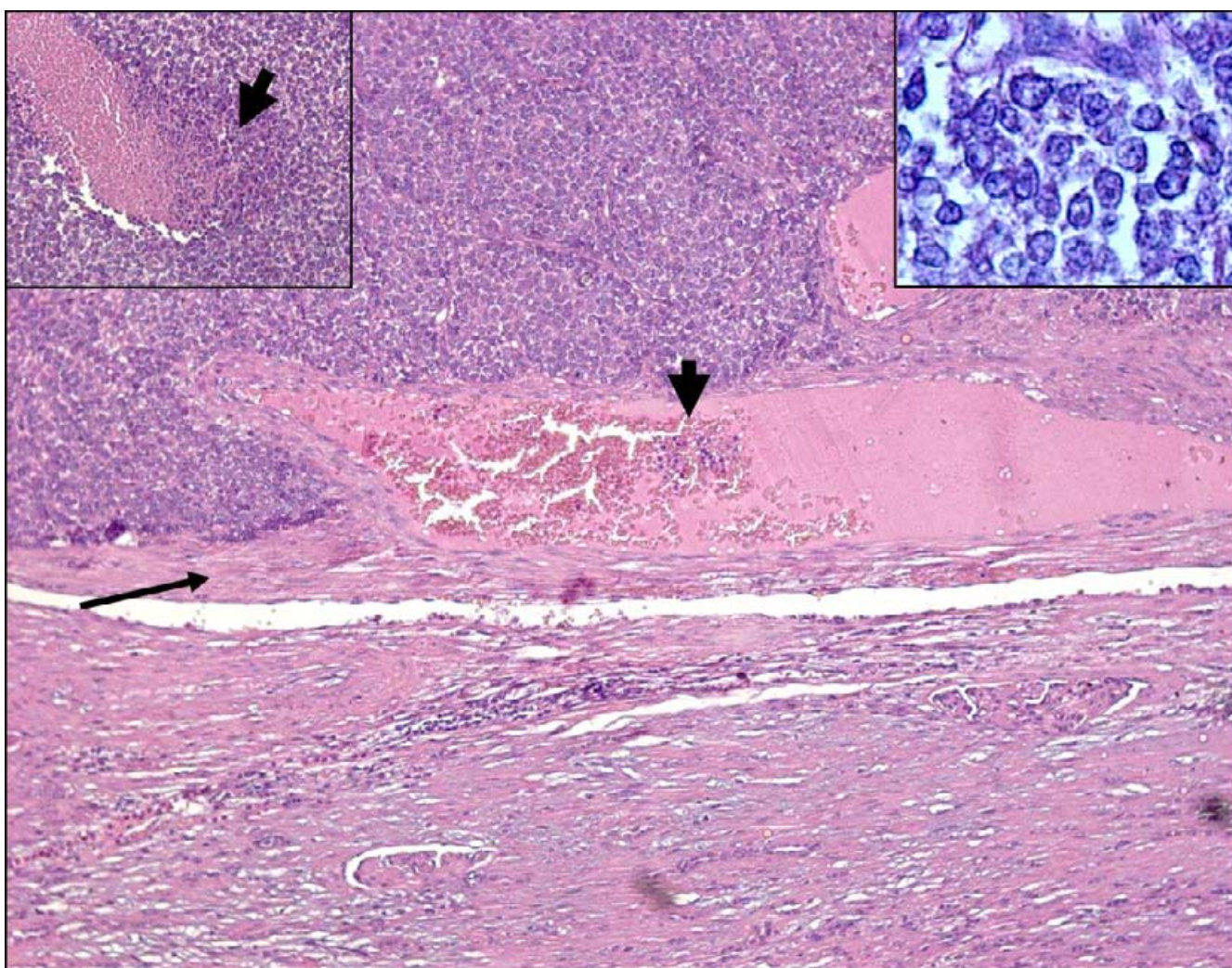


Figure 1

H&E features of the tumour (original magnification 10×; inset 40 ×): sheets of monotonous cells infiltrating vessels (small arrow) even near the capsule (long arrow). Homer-Wright rosettes were common (inset).

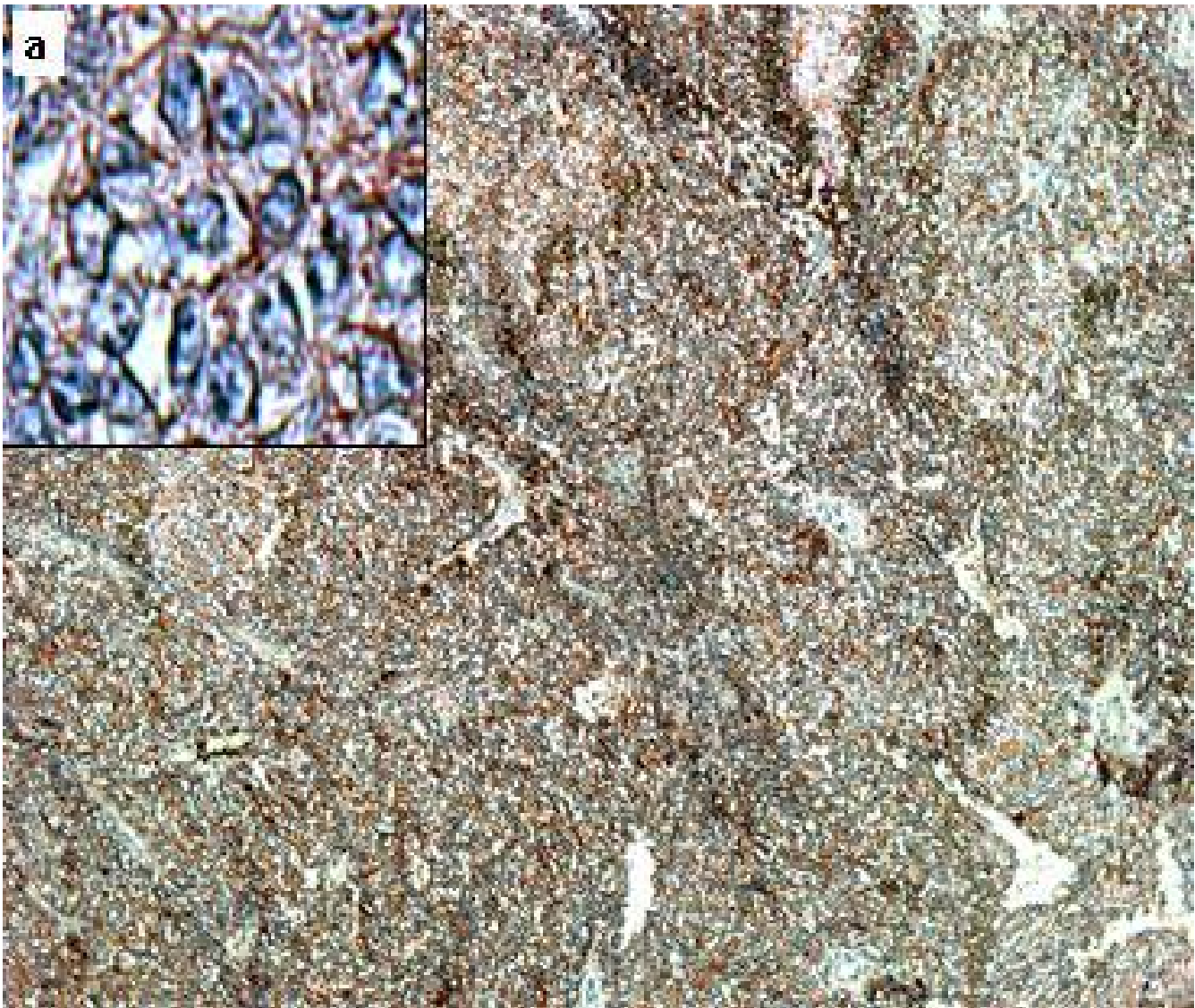


Figure 2

Immunoperoxidase for CD99 (10 \times) showed the tumor composed of small round cells with round nuclei and scant cytoplasm arranged in cohesive lobules. There are spindle cellular elements with diffuse CD99 cytoplasmic staining. Inset a (40 \times) shows Homer-Wright type rosettes positive for CD99.

The tumor was multilobular, grey, glistening, focally hemorrhagic, surrounded by a capsule and with a sharp demarcation from the uninvolved kidney. Histologically, the tumor consisted of small round cells with round nuclei and scant cytoplasm. It presented different patterns, with cohesive lobules or rosettes and perivascular pseudo-rosettes or, in some areas, spindle cellular elements (fig. 1).

The immunohistochemical evaluation revealed a diffuse CD99 positivity in the cytoplasm of the neoplastic cells (fig. 2); tumoral cells were also visible in the vascular lumens (fig. 3). By contrast, pankeratin, cytokeratin AE1/AE3, vimentin, desmin, S100, cromogranin were negative.

The clinical presentation and the macroscopic aspect, together with the histological pattern, the cytological characteristic and the cellular immunophenotype addressed the diagnosis towards primary PNET of kidney.

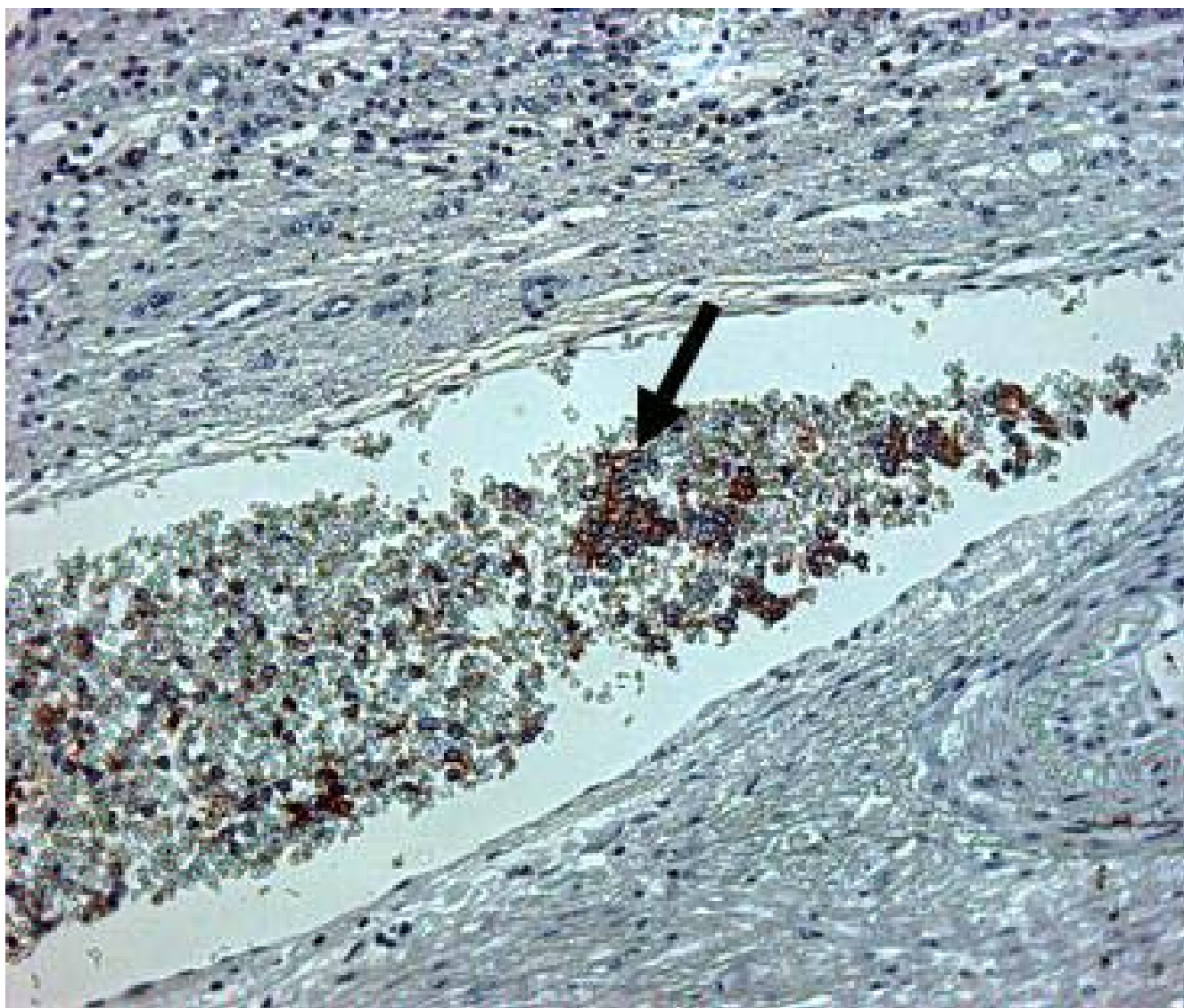


Figure 3
Immunoperoxidase for CD99 (X25) revealed positive tumoral cells in the vascular lumens (arrow).

A bone scan did not reveal positive areas. Eight cycles of chemotherapy with Vincristine, Ifosfamide and Adriamycin, four cycles of Ifosfamide and VP16 and eight sittings of local radiotherapy were sequentially performed. Follow-up examinations with CT and bone scan failed to show residual or recurrent tumor after 24 months.

Conclusions

Primitive Neuroectodermal Tumor of the kidney is a rare entity. The few cases reported revealed a variable presentation and an aggressive behaviour. The distinction from other primary malignancies of the kidneys is crucial for

prognosis. The differential diagnosis includes extra-osseous Ewing's sarcoma, rhabdomyosarcoma, Wilm's tumor, carcinoid, neuroblastoma, clear cell sarcoma of the kidney, lymphoma, the small cell variant of osteosarcoma, desmoplastic small round cell tumor and nephroblastoma [5].

The *Homer-Wright type* rosettes, commonly scarce of number or less defined in extra skeletal Ewing's sarcoma (ES), are a typical histological feature for PNET and can address the diagnosis although they can be found also in neuroblastoma [5]. To better address the diagnosis, an

immunohistochemical analysis is necessary. In our case the presence of MIC-2 gene products, known also as CD99, 12E7, E2, 013 and HBA71, suggested a PNET diagnosis. Primitive neuroectodermal tumors only immunoreactive to CD99, even if uncommon, are reported in the literature [13]. The reactivity to vimentin, NSE and S-100 may facilitate the diagnosis but is not pathognomonic, while CD 99 positivity is nowadays a clue for the diagnosis. Moreover cytogenetic studies (not performed in our case) demonstrated that PNET and Ewing's sarcoma can both be associated to a translocation of the long arms of chromosome 11 and 22, t(11;22)(q22;q12) [5]. Despite their genetic and antigenic similarity, many authors currently recognize PNET and extra-skeletal Ewing's sarcoma of the kidney as separate entities. It is also important to keep separate renal PNET and malignant rhabdomyosarcoma tumor (MRT). Weeks et al reported 8 cases suggestive for PNET but mimicking MRT [14]. Although renal PNET and MRT show similar clinico-pathological features, the latter usually occurs in very young children, having a more aggressive prognosis.

Rodriguez et al postulated that these two renal neoplasms share a common undifferentiated precursor to explain their similarity and we agree with these Authors [12]. Indeed, the hypothesis that tumors arise from stem cells (SCs) as a consequence of a maturative arrest is now growing [15]. SCs are present in almost all tissues and may originate different cellular lineages by the multi-step process named "differentiation". The role of SCs in tumorigenesis was clearly demonstrated in a number of carcinogenic models showing that solid and haematopoietic cancers could arise from tissue-specific SCs [16-19]. In agreement with Sell and Pierce, we retain that the degree of malignancy of a carcinoma depends by the stage in which SCs differentiation stopped during carcinogenesis [19]. In particular, since PNET, Ewing's tumour and MRT have a similar morphology, our hypothesis is that the mesenchymal stem precursor of these tumors is the same, but the stage of differentiation in which it is blocked is different. This could explain why sometimes it is difficult to discriminate between these tumors, notwithstanding they present a different biological behaviour.

Competing interests

None declared.

Authors' contribution

All authors contributed.

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Written consent was obtained from the patient for publication of the patient's details

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