

Images of Interest / Imagens de Interesse

Bladder Rhabdomyosarcoma in an Infant with Schimmelpenning Syndrome*Rabdomiossarcoma de Bexiga em Recém-Nascido com Síndrome de Schimmelpenning*Bruna Suda Rodrigues¹, Mayara Oliveira da Silva², Márcio Luís Duarte³, Élcio Duarte¹¹Irmandade da Santa Casa de Misericórdia de Santos, Santos, Brasil²Clínica MegalImagem, Santos, Brasil³WEBIMAGEM Telerradiologia, São Paulo, Brasil**Address**Márcio Luís Duarte
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Abstract

Bladder tumors occur more frequently between the fifth and seventh decades of life - they are rare before 40 years of age. Rhabdomyosarcoma is a rare embryonic tumor originating from skeletal muscle tissue. The tumor can appear anywhere in the body, even where there is no skeletal muscle. When located in the genitourinary tract, it can present with hematuria and anuria. The nevus sebaceous of Jadassohn (NSJ) is a congenital hamartoma of cutaneous structures, with epithelial and adnexal origin, present in approximately 0.3% of newborns, and may contain any component of the skin, including sebaceous and apocrine glands or hair. When associated with syndromic features such as mental retardation, central nervous system abnormalities, cardiovascular defects such as the cardiac arrhythmia reported in this case, ocular or skeletal abnormalities, it is called linear sebaceous nevus syndrome, or Schimmelpenning syndrome.

Keywords

Infant newborn; Nevus sebaceous of jadassohn; X-ray computed tomography; Magnetic resonance imaging.

Resumo

Os tumores de bexiga ocorrem com mais frequência entre a quinta e a sétima décadas de vida, sendo raros antes dos 40 anos de idade. O rabdomiossarcoma é um tumor embrionário raro originado do tecido muscular esquelético que pode aparecer em qualquer parte do corpo, mesmo onde não há músculo esquelético. Quando localizado no trato geniturinário, pode apresentar-se com hematuria e anúria. O nevo sebáceo de Jadassohn (NSJ) é um hamartoma congênito de estruturas cutâneas, de origem epitelial e anexial, presente em aproximadamente 0,3% dos recém-nascidos, podendo conter qualquer componente da pele, incluindo glândulas sebáceas e apócrinas ou cabelos. Quando está associado a características síndromicas, como atraso mental, anomalias do sistema nervoso central, defeitos cardiovasculares (como a arritmia cardíaca relatada neste caso), anomalias oculares ou esqueléticas, é denominado síndrome do nevo sebáceo linear ou síndrome de Schimmelpenning.

Palavras-chave

Recém-nascido; Nevo sebáceo de jadassohn; Tomografia computadorizada por raios X; Imagem por ressonância magnética.

Bladder tumors occur more frequently between the fifth and seventh decades of life - they are rare before 40 years of age. Rhabdomyosarcoma is a rare embryonic tumor originating from skeletal muscle tissue.¹ When located in the genitourinary tract, it can present with hematuria and anuria.¹ Being a tumor of embryonic origin, it is more common in children and very rare in adults: 85% occur in children and adolescents.¹

Although most musculoskeletal tumors occur in the chest, arms, and legs, rhabdomyosarcomas often appear elsewhere in the body. The most common site of involvement is the head and neck (30% to 40%), followed by the urinary and reproductive systems (20% to 25%). However, the tumor can appear anywhere in the body, even where there is no skeletal muscle.¹ When located in the genitourinary tract, it can present with hematuria and anuria.¹

In most cases, bladder tumors are asymptomatic. The clinical condition, when apparent, progresses with hematuria, intermittent or persistent without apparent causes, with an approximate average of 25% of patients presenting macroscopic and 9% microscopic hematuria.^{1,2} Other manifestations include flank pain, irritative and obstructive symptoms, such as increased voiding frequency, voiding urgency, retention, dysuria, nocturia, hydronephrosis,

and changes in renal function. Signs and symptoms vary according to the location and extent of the tumor mass.^{1,2} Diagnosis consists of vesical visualization through cystoscopy and abdominal and pelvic evaluation through imaging exams, followed by a biopsy or resection.² It is a very aggressive tumor, which usually requires surgical treatment, with partial resections and even complete removal of a large part of the lower genitourinary tract.²

The presentation of the tumor on ultrasound demonstrates a heterogeneous lesion with sonolucent areas, corresponding to sites of hemorrhage or necrosis. On CT, we should look for an image consistent with a heterogeneous, infiltrative attenuation mass, usually with imprecise borders. On the other hand, MRI reveals a low signal on T1-weighted images and a high signal on T2-weighted images, allowing for a better assessment than ultrasound and CT of the local extension of the tumor and the involvement of adjacent structures.

The nevus sebaceous of Jadassohn (NSJ) is a congenital hamartoma of cutaneous structures, with the epithelial and adnexal origin, present in approximately 0.3% of newborns, and may contain any component of the skin, including sebaceous and apocrine glands or hair.¹ Most often seen in the head and neck region, it presents as a well-demarcated yellowish alopecia plaque.¹ In our case, the nevus was

confirmed with a biopsy and pathological examination of the cephalic lesion. The nature of secondary tumors that arise in sebaceous nevus and the risk of malignant neoplasm are controversial issues.¹

When a Jadassohn sebaceous nevus is associated with syndromic features such as mental retardation, central nervous system abnormalities, cardiovascular defects such as the cardiac arrhythmia reported in this case, ocular or skeletal abnormalities, it is called linear sebaceous nevus syndrome, or Schimmelpenning syndrome.¹ The etiology is still unknown, but recent studies point to a possible link with the mother's positivity for the human papillomavirus or mutations in the patched gene (PTCH).² Luo et al. noticed a novel mutation in the HRAS gene (c.38G>T; p.Gly13Val) was noticed in a mosaic state in NS and rhabdomyosarcoma, which could be a connection between these entities.³

We report a male infant aged four months and 13 days old, weighing 4.3 kg, with a history of diarrhea for ten days, low formula acceptance, and prostration. He had not urinated for four days and had started incoercible vomiting. He was born at 34 weeks, remaining in the neonatal intensive care unit for 48 days due to a previous diagnosis of fetal cardiac arrhythmia.

On physical examination, the baby was hypoactive/reactive, pale 2+/4+ and dehydrated 1+/4+. The abdomen was globular with reduced air-fluid sounds, distended, painful on diffuse palpation, with a thickened spermatic cord, there was a bilateral hydrocele.

He also had large, hairless, yellowish-brown plaques on the scalp (Figure 1), which were already present at birth according to his mother. Histopathological examination of these lesions on the scalp revealed hyperkeratosis, acanthosis, and hyperplasia of the sebaceous glands, compatible with nevus sebaceous of Jadassohn. Karyotype evaluation showed: karyotype 46, XY, presence of variant15CENH+ (increase in centromeric heterochromatin in one of chromosomes 15). Renal function deteriorated – urea (323 mg/ml) and creatinine (7.20 mg/ml).



Figure 1 – Hairless, yellowish-brown plaques on the scalp of a four-month-old infant. Histopathological examination revealed hyperkeratosis, acanthosis, and hyperplasia of the sebaceous glands, compatible with Jadassohn sebaceous nevus.

Urethrocytography was performed (Figure 2), which demonstrated a defect in the bladder contrast. Ultrasonography and computed tomography (CT) showed a bladder mass measuring approximately 6.5 cm (Figure 2), later confirmed by magnetic resonance imaging (MRI) (Figure 3).

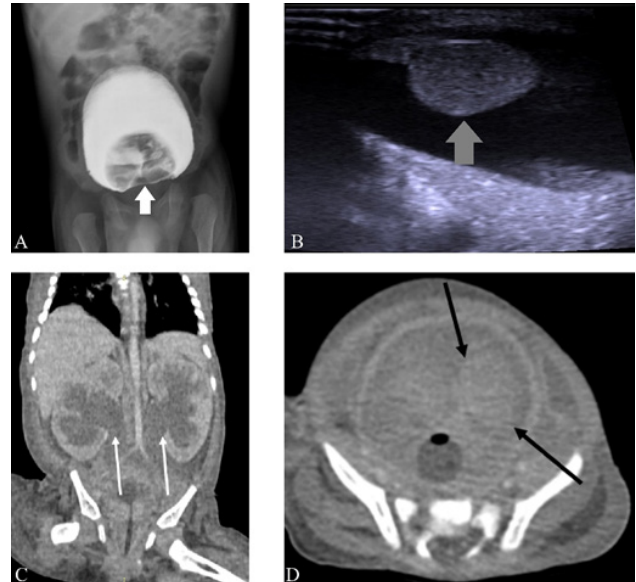


Figure 2 – **A:** urethrocytography showing a defect in the bladder contrast (white arrow). **B:** Ultrasonography showing expansive and infiltrative lesion affecting the bladder wall (grey arrow). **C:** contrast-enhanced CT in the coronal section demonstrating bilateral ureterohydronephrosis (white arrows). **D:** contrast-enhanced CT in the axial section showing an expansive and infiltrative lesion affecting the lateral wall and the vesical floor, invading the ureteral meatus (black arrows), causing bilateral ureterohydronephrosis.

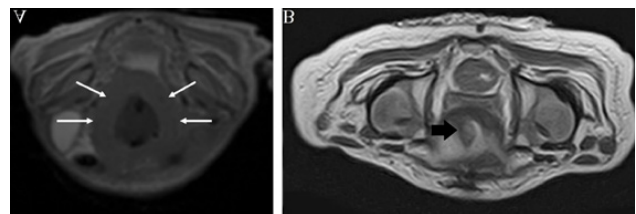


Figure 3 – **A:** MRI in the axial section of the sequence T1 showing diffuse thickening of the bladder walls (white arrows). **B:** MRI in the axial section of the sequence T2 showing an expansive lesion affecting the bladder wall (black arrow).

The bladder tumor was excised – the anatomopathological analysis diagnosed botryoid-type embryonal rhabdomyosarcoma with compromised margins – and the patient is currently undergoing chemotherapy. Bladder ultrasonography and CT scan were normal after the surgical procedure.

Ethical disclosures / Divulgações Éticas

Conflicts of interest: The authors have no conflicts of interest to declare.

Conflitos de interesse: Os autores declaram não possuir conflitos de interesse.

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Confidentiality of data: The authors declare that they have followed the protocols of their work center on the publication of data from patients.

Confidencialidade dos dados: Os autores declaram ter seguido os protocolos do seu centro de trabalho acerca da publicação dos dados de doentes.

Protection of human and animal subjects: The authors declare that the procedures followed were in accordance with the regulations of the relevant clinical research ethics committee and with those of the Code of Ethics of the World Medical Association (Declaration of Helsinki).

Proteção de pessoas e animais: Os autores declaram que os procedimentos seguidos estavam de acordo com os regulamentos estabelecidos pelos responsáveis da Comissão de Investigação Clínica e Ética e de acordo com a Declaração de Helsínquia da Associação Médica Mundial.

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