complete resection is uncommon, having been reported in only approximately 10% of cases⁽⁵⁾.

REFERENCES

- Saab ST, McClain CM, Coffin CM. Fibrous hamartoma of infancy: a clinicopathologic analysis of 60 cases. Am J Surg Pathol. 2014;38:394– 401.
- Reye RD. A consideration of certain subdermal fibromatous tumours of infancy. J Pathol Bacteriol. 1956;72:149–54.
- Dickey GE, Sotelo-Avila C. Fibrous hamartoma of infancy: current review. Pediatr Dev Pathol. 1999;2:236–43.
- Eich GF, Hoeffel JC, Tschäppeler H, et al. Fibrous tumours in children: imaging features of a heterogeneous group of disorders. Pediatr Radiol. 1998;28:500–9.
- 5. Jesus LE, Gameiro VS, Novelli RJ, et al. Hamartoma fibroso infantil:

- lesão volumosa com envolvimento de plexo braquial. Acta Ortop Bras. 2006;14:229–30.
- Laffan EE, Ngan BY, Navarro OM. Pediatric soft-tissue tumors and pseudotumors: MR imaging features with pathologic correlation: part 2. Tumors of fibroblastic/myofibroblastic, so-called fibrohistiocytic, muscular, lymphomatous, neurogenic, hair matrix, and uncertain origin. Radiographics. 2009;29:e36.

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Left renal vein thrombosis secondary to compression by the uncinate process of the pancreas, mimicking the nutcracker syndrome

Dear Editor,

A 40-year-old woman presented with a five-month history of pain in a portion of the mesogastrium and in the left side. She reported recent weight loss (of 5 kg) after having had dengue fever. She also reported no comorbidities and stated that she was not using contraceptives. The physical examination revealed Giordano's sign on the left side. The results of the blood count and urinalysis were normal. Computerized tomography of the abdomen showed compression of the left renal vein (LRV) caused by the uncinate process of the pancreas pressing against the aorta,

leading to dilation of the proximal segment, with an intraluminal thrombus (Figures 1A and 1B), dilation of the collateral perirenal venous system, dilation of the gonadal veins, and ipsilateral pelvic varices (Figures 1C and 1D). The patient was treated with oral anticoagulants for four months and declined to have a stent placed in the LBV.

Vascular compressive syndromes occur in less than 1% of the cases and represent vascular trapping between rigid surfaces, which lead to manifestations caused by hypertension, venous congestion, thrombosis, and arterial ischemia⁽¹⁻⁴⁾.

The causes of compression of the LRV include expansive retroperitoneal formations, anatomical variations, and nutcracker syndrome (NCS)⁽²⁾. NCS is usually caused by the trapping of the LRV between the superior mesenteric artery and the abdominal

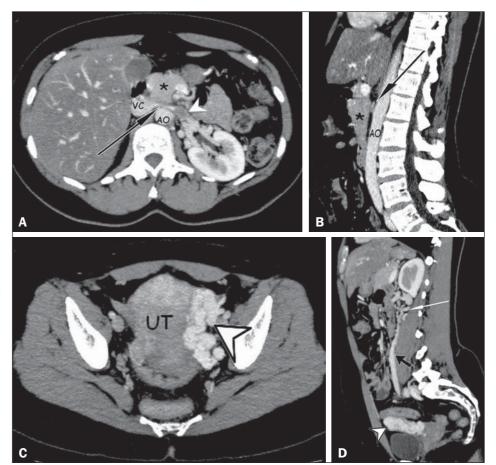


Figure 1. A,B: Contrast-enhanced arterial-phase CT of the abdomen, in the axial plane and in sagittal reconstruction, respectively, characterizing clots within the proximal portion of the LRV (white arrowhead), showing a reduction in its caliber at the aortomesenteric compression (black arrows), close to its junction with the inferior vena cava (VC), due to the extrinsic compression exerted by the uncinate process of the pancreas (asterisks) against the aorta (AO). Additional finding: diffuse hypointense signal in the hepatic parenchyma, suggesting fatty infiltration. C,D: Contrast-enhanced arterial-phase CT of the abdomen, in the axial plane and in sagittal reconstruction, respectively, showing dilation of the pelvic vessels (arrowheads) near the left lateral aspect of the uterus and the left gonadal vein (black arrow), with dilated collateral veins in the ipsilateral perirenal space (white arrow).

aorta (aortomesenteric compression) $^{(1-5)}$. In rare cases, the LRV is retroaortic. In such cases, compression occurring between the aorta and the spine is known as posterior NCS $^{(2-4)}$. The nutcracker phenomenon corresponds to these findings without clinical correlation $^{(2-4)}$. The prevalence of NCS is unknown, although it is known that it occurs predominantly in healthy, thin individuals between 20 and 40 years of age and in women $^{(1-4)}$. Clinically, hematuria is the most common finding, followed by pain on the left side, dyspareunia, dysmenorrhea, dysuria, varicoceles, and pelvic varices $^{(1-5)}$. In exceptionally rare cases, anatomic variations in the pancreas compress nearby vessels, including the LRV $^{(2,6,7)}$.

Renal vein thrombosis (RVT) is common in nephrotic syndrome and in severely hypotensive neonates. Other causes: traumas, surgery, infections, neoplasias, vasculitis, venous compressions, contraceptives and myeloproliferative diseases. It's infrequent in healthy adults, predominantly unilaterally $^{(8,9)}$. The clinical presentation of RVT is much like that of NCS, with the added features of an acute increase in renal volume, late atrophy, and progressive deterioration of renal function, as well as the complication of pulmonary thromboembolism in up to 50% of cases $^{(5,8,9)}$.

The pathophysiology of thromboses encompasses Virchow's triad: endothelial lesions, stasis, and hypercoagulability. Generally, thrombotic events involve at least two factors, although one may be sufficient (5,8,9).

One of the principal methods employed in the diagnosis of NCS is Doppler ultrasound, which is noninvasive and can be used in determining venous caliber and flow, the latter being suggestive of NCS when it exceeds 100 cm/s, with a sensitivity and specificity of 78% and 100%, respectively, for the diagnosis (1-4). It shows high sensitivity in the investigation of RVT⁽⁸⁾. Ultrasound, however, is operator-dependent and may not detect small thromboses^(8,9). For the diagnosis of NCS and RVT, angiography has a sensitivity of 66.7-100% and a specificity of $55.6-100\%^{(8)}$. It is able to evaluate the aortomesenteric angle (compression); possible compression and dilation of the LRV; filling defects; endoluminal blood clots; and signs of chronic thrombosis, such as thickening of the vessel walls and calcifications (1-4,9). However, it uses radiation and potentially nephrotoxic contrast agents^(8,9). Retrograde venography is the gold standard examination in NCS⁽¹⁻⁴⁾ and RVT⁽⁸⁾; it shows pressure gradients greater than 3 mmHg in the LRV, in

addition to the filling defects that represent thrombi $^{(1-4,8)}$. However, it is invasive, potentially triggering thrombosis, and uses intravenous iodine $^{(8)}$.

The therapeutic options are conservative treatment, reimplantation/transposition of the LRV, the use of an external or internal stent, renal autotransplantation, gonadocaval bypass, and nephrectomy $^{(1-5)}$. If RVT occurs, anticoagulation and thrombolysis can also be employed $^{(8-10)}$.

REFERENCES

- Eliahou R, Sosna J, Bloom AI. Between a rock and a hard place: clinical and imaging features of vascular compression syndromes. Radiographics. 2012;32:E33–49.
- Lamba R, Tanner D, Sekhon S, et al. Multidetector CT of vascular compression syndromes in the abdomen and pelvis. Radiographics. 2014;34:93-115.
- Butros SR, Liu R, Oliveira GR, et al. Venous compression syndromes: clinical features, imaging findings and management. Br J Radiol. 2013; 86:20130284.
- Fong JK, Poh AC, Tan AG, et al. Imaging findings and clinical features of abdominal vascular compression syndromes. AJR Am J Roentgenol. 2014;203:29–36.
- Mallat F, Hmida W, Jaidane M, et al. Nutcracker syndrome complicated by left renal vein thrombosis. Case Rep Urol. 2013;2013:168057.
- Yun SJ, Nam DH, Ryu JK, et al. The roles of the liver and pancreas in renal nutcracker syndrome. Eur J Radiol. 2014;83:1765–70.
- Chauhan R, Roy TS, Chaudhury A, et al. Variant human pancreas: aberrant uncinate process or an extended mesenteric process. Pancreas. 2003;27:267–9.
- Asghar M, Ahmed K, Shah SS, et al. Renal vein thrombosis. Eur J Vasc Endovasc Surg. 2007;34:217–23.
- Wang Y, Chen S, Wang W, et al. Renal vein thrombosis mimicking urinary calculus: a dilemma of diagnosis. BMC Urol. 2015;15:61.
- Yoshida RA, Yoshida WB, Costa RF, et al. Nutcracker syndrome and deep venous thrombosis in a patient with duplicated inferior vena cava. J Vasc Surg Venous Lymphat Disord. 2016;4:231–5.

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Atypical presentation of mature cystic teratoma ("floating balls")

Dear Editor,

A 43-year-old female patient with no known diseases sought medical attention complaining of increased abdominal volume. The patient underwent ultrasound and subsequent magnetic resonance imaging (MRI) of the pelvis (Figure 1), which showed an expansile cystic lesion, with heterogeneous content, measuring $16.0 \times 16.0 \times 10.0$ cm and containing numerous oval formations of various sizes. The lesion was hyperechoic on ultrasound and mobile upon a change in patient position. The oval formations showed intermediate signal intensity on T1- and T2-weighted MRI scans, with no evidence of signal loss in fat-saturated sequences or signal drop on an out-of-phase T1-weighted gradientecho sequence. These imaging findings, although uncommon, are pathognomonic of mature cystic teratoma (MCT). The patient underwent surgery, and the diagnosis was confirmed by histopathological analysis of the surgical specimen.

Also known as a dermoid cyst, MCT is the most common benign ovarian tumor, accounting for 10-25% of cases in adult

patients and 50% of those in pediatric patients^(1–3). MCTs are typically asymptomatic and slow-growing^(1,3). They are usually seen in women of reproductive age and are rarely diagnosed before puberty. Its growth ceases at menopause^(4–7). An MCT typically contains well-differentiated tissues of the three germ layers^(1,5): the ectoderm, (derived from the skin and neural tissues); the mesoderm (osteomuscular and adipose tissues); and the endoderm (ciliated and mucinous epithelium). The diversity of tissues in teratomas results in a wide variety of characteristics in imaging studies.

In most cases, pelvic tumors do not present imaging features that are considered diagnostic $^{(8-12)}$. However, MCTs often present typical imaging features, which facilitate the diagnosis. Among such features, one of the most common is that of a fatty tumor $^{(3)}$. In such cases, the most common ultrasound finding is that of a cystic mass with an echogenic tubercle (a Rokitansky nodule), presenting posterior acoustic shadowing secondary to calcifications, strands of hair, or foci of $fat^{(3,5,7)}$.

Characteristic findings on computed tomography include areas of fat attenuation, with or without foci of calcification. On