Conflict of Interest

The authors declare that they have no conflicts of interest.

References


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Classic Ehlers-Danlos Syndrome:
Clinical and Ultrasound Findings

Síndrome de Ehlers-Danlos clásico: hallazgos clínicos y ecográficos

To the Editor:

Ehlers-Danlos syndrome (EDS) is a heterogeneous group of congenital connective tissue diseases caused by mutations in genes involved in the synthesis or processing of collagen fibers.1 The phenotypic manifestations of EDS vary greatly, and mild cases can go unnoticed until late in life. We present the case of a girl who was diagnosed with classic EDS in our hospital based on clinical and ultrasound findings.

An 8-year-old girl was referred from the emergency department for evaluation of a painful lesion on the left leg that had appeared several weeks earlier after a fall from a ladder. Physical examination revealed a subcutaneous bulge of about 4 cm in diameter in the left pretibial region with yellowish-purpuric overlying skin and a strikingly gummy consistency. The patient had reticulated erythematous-violaceous lesions on the right leg and dehiscent an atrophic scars on the right leg and left knee (Fig. 1). High-frequency ultrasound (18 MHz) of the left pretibial lesion revealed an anechoic collection delimited by a thin pseudocapsule, compatible with an organized hematoma (Fig. 2A). Septal edema compatible with traumatic panniculitis was evident in the surrounding hyper-echogenic subcutaneous tissue. Doppler signal was absent. In the directed anamnesis the patient’s mother reported that the girl had been born preterm due to premature rupture of membranes and had muscular hypotonia during the neonatal period. The patient was undergoing tests in the endocrinology department of another hospital for short stature and disproportion between the trunk and limbs. The family history provided by the mother included joint hyperlaxity, abnormal scarring, and early osteoarthritis. The patient also presented with skin hyperextensibility, joint hypermobility, and Gorlin sign (ability to reach the nose with the tip of the tongue) (Fig. 3). Based on these data a suspected diagnosis of classic EDS was established. A cardiological examination, including electrocardiogram and echocardiography, revealed no findings of interest, and the

Figure 1 Extensive purpuric lesion on the anterior aspect of the middle and distal third of the left leg, and reticulated erythematous-violaceous lesions on the contralateral leg. Note the presence of 2 atrophic scars on the distal third of the right leg and on the left knee.
Figure 2 Ultrasound images of the left pretibial lesion (B-mode, 13-MHz probe). A, At the level of the ecchymotic area is an anechoic subcutaneous mass delimited by a thin pseudocapsule. B, Follow-up ultrasound 1 month later reveals a marked reduction in lesion thickness.

Figure 3 Characteristic clinical signs of classic Ehlers-Danlos syndrome. A, Joint hypermobility with hyperextension of the fingers. B, Atrophic scars with a cigarette-paper-like appearance. C, Skin hyperlaxity. D, Hyperextension of the tongue (Gorlin sign).

results of a laboratory workup, including a complete blood count and coagulation tests, were normal apart from slightly elevated D-dimer levels. A second ultrasound examination performed 1 month later revealed a reduction in the hematoma of approximately 50% (Fig. 2B).

Classic EDS is inherited in an autosomal dominant manner and is caused by mutations in COL5A1 or COL5A2, which encode the alpha-1 and alpha-2 chains, respectively, of collagen type V. The disease is characterized by skin hyperextensibility, joint hypermobility and associated complications (luxations, pain, early osteoarthritis), and other clinical characteristics that are reviewed in a recent international consensus paper. Dermatologic manifestations of classic EDS include skin hyperextensibility and abnormal scarring, which results in the formation of atrophic scars with a cigarette-paper-like appearance. Patients have thin, velvety skin that bruises in response to minimal trauma. Other potential lesions include nodular molluscoid pseudotumors secondary to calcification and fibrosis of hematomas; spheroids (hard spherical nodules on the forearms and pretibial areas); and piezogenic pedal papules. Our patient fulfilled the diagnostic criteria (Table 1). No genetic study was performed as this was not essential for confirmation of the diagnosis. Classic EDS is usually diagnosed when affected individuals begin to stand and walk, as this period coincides with the appearance of lacerations and bruising that tend to alarm parents. The main clinical differential diagnosis in childhood is bruising caused by child abuse, potentially distinguishing features of which include lesions located in areas not exposed to accidental
In conclusion, classic EDS should be suspected in children with joint and skin hyperlaxity and easy bruising. High-resolution ultrasound is very useful for the diagnosis and follow-up of hematomas associated with EDS.

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References


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<tr>
<th>Table 1 Diagnostic Criteria for Classic Ehlers-Danlos Syndrome</th>
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<tr>
<td><strong>Major Criteria</strong></td>
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<tr>
<td>1. Marked skin hyperextensibility and atrophic scarring</td>
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<td>2. Generalized joint hypermobility</td>
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<td>3. Skin fragility</td>
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<td>4. Molluscoid pseudotumors</td>
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<td>5. Subcutaneous spheroids</td>
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<td>6. Hernia (or history thereof)</td>
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For diagnosis it is necessary to fulfil major criterion 1 + major criterion 2, or major criterion 1 + 3 of the minor criteria. Source: Malfait et al., 2017.