IMÁGENES EN DERMATOLOGÍA

Síndrome de CLAPO

CLAPO Syndrome

M.Á. Flores-Terry, P. Zamberk-Majlis, M.P. Cortina-de la Calle, M. García-Arpa

Servicio de Dermatología Médico-Quirúrgica y Venereología, Hospital General Universitario de Ciudad Real, Ciudad Real, Spain

We evaluated a 1-month-old boy with no personal or family history of interest for congenital lesions on the lip and right leg. The examination revealed an erythematous-violaceous plaque on the central part of the lower lip that extended beyond the vermillion border, asymmetric violaceous macular lesions on the right leg, and a barely perceptible increase in the diameter of the leg (Fig. 1). No lesions were detected on the tongue, oral cavity, neck, or face. The results of the neurologic examination were normal, and the extension study did not reveal vascular abnormalities in other organs. At 2 years of follow-up, the lesion on the lip persisted, although it was less intense, and the diameter of the right leg had increased slightly, with the same macular lesions. No complications were observed. Overgrowth syndromes with complex vascular abnormalities comprise a heterogeneous group of disorders characterized by vascular malformations and local or generalized hypertrophy. CLAPO syndrome is an example of overgrowth syndromes. It was described by López-Gutiérrez and is characterized by the presence of a capillary malformation on the lower lip, lymphatic malformation on the face and neck, asymmetry, and partial or generalized overgrowth. It is important to recognize this condition and follow patients in order to detect major abnormalities and prevent complications.

Figure 1


* Corresponding author.
E-mail address: miguelterry85@hotmail.com (M.Á. Flores-Terry).

1578-2190/© 2017 Elsevier España, S.L.U. and AEDV. All rights reserved.