A healthy primigravida with a 36-week gestation presented with fetal right ventriculomegaly. Magnetic resonance imaging (MRI) showed an encephaloclastic porencephaly and residual hemorrhagic components. Trauma, hypertension, TORCH infection and alloimmune thrombocytopenia were excluded. A caesarean section was planned and a term boy with normal platelet count and coagulation tests was delivered. Brain MRI confirmed the previous findings (Fig. 1). A heterozygous pathogenic variant (c.518G>A, p.Gly173Asp), maternally inherited, was identified in the COL4A1 gene. Abdominal and cardiac ultrasound were normal. At three years old the child presented left hemiparesis, inaugural epilepsy, and strabismus with no retinal arterial tortuosity or cataract.

Collagen type IV alpha 1 (COL4A1) is a structural component of the vascular basement membrane of many tissues. Its fragility can trigger rupture and intrauterine intraventricular hemorrhage, leading to porencephaly and neurological symptoms, like hemiparesis and epilepsy, which were previously unexplained. Early diagnosis allowed an appropriate follow-up and genetic counseling.

Keywords: Cerebral Hemorrhage/genetics; Collagen Type IV/genetics; Fetal Diseases/genetics; Infant, Newborn

PALAVRAS-CHAVE: Colagénio Tipo IV/genética; Doenças Fetais/genética; Hemorragia Cerebral/genética; Recém-Nascido

Figure 1 – Brain MRI. Coronal T2 weighted-image (WI) (A) and axial T2*-WI (B), depicting right frontal porencephalic cavity (star) with peripheral hemorrhagic residues (arrow)

Laura LEITE DE ALMEIDA1,2, Ana GRANGEIA2,3, Luísa SAMPAIO2,4, Ana VILANT."5
Acta Med Port (In Press) • https://doi.org/10.20344/amp.20515
AUTHOR CONTRIBUTIONS
LLA: Data collection, literature search and drafting of the manuscript.
AG: Critical review of the manuscript.
LS: Data collection and analysis.
AV: Data collection, literature search and critical review of the manuscript.

PROTECTION OF HUMANS AND ANIMALS
The authors declare that the procedures were followed according to the regulations established by the Clinical Research and Ethics Committee and to the Helsinki Declaration of the World Medical Association updated in 2013.

DATA CONFIDENTIALITY
The authors declare having followed the protocols in use at their working center regarding patients’ data publication.

PATIENT CONSENT
Obtained.

COMPETING INTERESTS
The authors have declared that no competing interests exist.

FUNDING SOURCES
This research received no specific grant from any funding agency in the public, commercial, or not-for-profit sectors.

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