

Neonatal Subgaleal Hemorrhage

Hemorragia Subgaleal Neonatal

Mariana Portela^{1*}, Maria Ventura Nogueira¹, Marta Alves², Almerinda Pereira²

*Corresponding Author/Autor Correspondente:

Mariana Portela [mariana.fpo@hotmail.com]

Hospital de Braga, Sete Fontes - São Victor, 4710-243 Braga, Portugal

ORCID iD: 0000-0002-3297-9431

KEYWORDS: Hemorrhage; Infant, Newborn; Scalp

PALAVRAS-CHAVE: Couro Cabeludo; Hemorragia; Recém-Nascido

A full-term female newborn was born by eutocic deliver, after an uneventful pregnancy. Apgar score 10/10. Right after birth, an occipital swelling was noted, without other findings. Two hours later, the dimensions of the scalp collection had increased significantly, with the edema extending to temporal and periorbital regions (Fig. 1). At admission in intensive care, she was pale, hypotonic, however vital signs and capillary refill time were normal. Head circumference (HC) was 37.5 cm (99th percentile). At admission she presented: hemoglobin 14.4 g/dL, normal platelet count, mild metabolic acidosis (pH 7.313, HCO₃⁻ 21.3 mmol/L, base excess of (-)5 and lactate 7.9 mmol/L) and normal coagulation parameters (prothrombin time was 24.9 sec, international normalized

ratio of 1.8 and activated partial thromboplastin clotting time was 45.9 sec). A 10 mL/kg bolus of 0.9% saline was administered. At day 2, hemoglobin dropped to 10.1 g/dL, without clinical repercussion, and a transfusion of red blood cells was administered. Maximum HC was 39 cm. Cranial ultrasound excluded intraventricular hemorrhage and cranial tomography confirmed an extensive subgaleal hemorrhage (SGH) (Fig. 2). Phototherapy was required from day 3 to 5 (maximum bilirubin level was 17.76 mg/dL). The newborn was discharged at day 6, with a stable hemoglobin level (17.9 g/dL) and decreasing HC (37.5 cm). Coagulopathy was excluded in follow-up (normal levels of von Willebrand factor and factor VIII procoagulant activity).

1. Pediatrics Department, Hospital de Braga, Braga, Portugal. 2. Neonatology Department, Hospital de Braga, Braga, Portugal.

Received/Recebido: 23/03/2022 - Accepted/Aceite: 18/07/2022 - Published online/Publicado online: 01/09/2022 - Published/Publicado: 30/09/2022

© Author(s) (or their employer(s)) and Gazeta Médica 2022. Re-use permitted under CC BY-NC. No commercial re-use. © Autor (es) (ou seu (s) empregador (es)) e Gazeta Médica 2022. Reutilização permitida de acordo com CC BY-NC. Nenhuma reutilização comercial.



FIGURE 1. Extensive subgaleal hemorrhage.

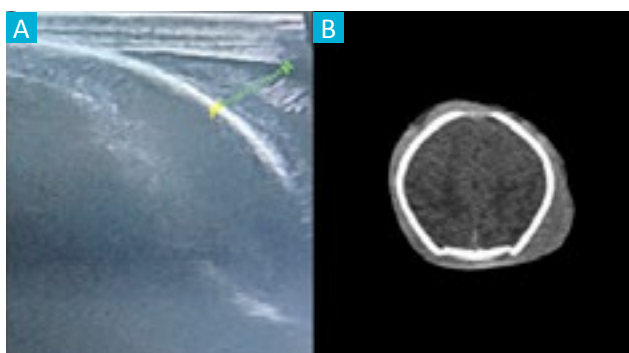


FIGURE 2. Ultrasound demonstrating the left side of the subgaleal hemorrhage, with 11.10 mm (A). Cranial tomography performed at 24 hours of life, showing an extensive hemorrhage (B).

SGH is a rare and potentially lethal condition in newborns, due to the rupture of emissary veins, causing accumulation of blood between the epicranial aponeurosis and the periosteum.¹⁻³ It should be considered in the presence of diffuse fluctuant swelling, that extends widely and crosses suture lines.²⁻⁴ SGH is more frequently associated with vacuum/forceps assisted delivery, but may also occur spontaneously, with an incidence of approximately 0.4/1000 spontaneous births.¹⁻³ Massive blood loss into that space may lead to acute anemia, hypotension, acidosis and eventually, hypovolemic shock.^{1,3} Early detection and close monitoring, with prompt administration of fluids and/or blood products are mandatory.³ Imaging study corroborates the diagnosis. After stabilization, the etiological investigation should proceed to discard the possibility of coagulopathy.²

AUTHORS CONTRIBUTION/ CONTRIBUIÇÃO AUTORAL

MP: Involved in patient care and drafted the manuscript
MVN, MA, AP: Involved in patient care and contributed to the revision of the manuscript

MP: Envolvimento no seguimento do paciente e redação do manuscrito

MVN, MA, AP: Envolvimento no seguimento do paciente e contribuição para a revisão do manuscrito

RESPONSABILIDADES ÉTICAS

CONFLITOS DE INTERESSE: Os autores declaram a inexistência de conflitos de interesse na realização do presente trabalho.

FONTES DE FINANCIAMENTO: Não existiram fontes externas de financiamento para a realização deste artigo.

CONFIDENCIALIDADE DOS DADOS: Os autores declaram ter seguido os protocolos da sua instituição acerca da publicação dos dados de doentes.

CONSENTIMENTO: Consentimento do doente para publicação obtido.

PROVENIÊNCIA E REVISÃO POR PARES: Não comissionado; revisão externa por pares.

ETHICAL DISCLOSURES

CONFLICTS OF INTEREST: The authors have no conflicts of interest to declare.

FINANCING SUPPORT: This work has not received any contribution, grant or scholarship.

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

PATIENT CONSENT: Consent for publication was obtained.

PROVENANCE AND PEER REVIEW: Not commissioned; externally peer reviewed.

REFERENCES

1. Acuña J, Adhikari S. Point-of-care Ultrasound to Distinguish Subgaleal and Cephalohematoma: Case Report. *Clin Pract Cases Emerg Med.* 2021;5:198-201. doi: 10.5811/cpcem.2021.3.51375.
2. Wetzel E, Kingma A. Subgaleal hemorrhage in a neonate with factor X deficiency following a non-traumatic cesarean section. *J Perinatol.* 2012;32:304-5. doi: 10.1038/jp.2011.122.
3. Davis DJ. Neonatal subgaleal hemorrhage: diagnosis and management. *CMAJ.* 2001;164:1452-3.
4. Lee SJ, Kim JK, Kim SJ. The clinical characteristics and prognosis of subgaleal hemorrhage in newborn. *Korean J Pediatr.* 2018;61:387-91. doi: 10.3345/kjp.2018.06800.