

Images in Infectious Diseases

Congenital cytomegalovirus infection-related thrombocytopenia

Melis Deniz^[1], Mehmet Fatih Deveci^[2] and Nazlı Gülsüm Akyel^[3]

[1]. Sanliurfa Training and Research Hospital, Department of Pediatric Infectious Diseases, Sanliurfa, Turkey.

[2]. Sanliurfa Training and Research Hospital, Department of Neonatology, Sanliurfa, Turkey.

[3]. Sanliurfa Training and Research Hospital, Department of Pediatric Radiology, Sanliurfa, Turkey.

A neonate with respiratory distress and no petechial rash was admitted to the neonatal intensive care unit. The initial blood count revealed thrombocytopenia. Results from the peripheral blood culture and additional *biochemical* tests were negative. Due to persistent thrombocytopenia, investigations were conducted for congenital cytomegalovirus (cCMV) infection. Positive results for CMV-specific immunoglobulins G and M were found on two separate occasions. Elevated blood levels of CMV DNA (4,285 copies/mL) 5 days after birth and high urinary levels of CMV DNA (1.96 million copies/mL) within 2 weeks of birth were detected. A computed tomography scan of the head detected periventricular calcifications (**Figure 1**). Magnetic resonance imaging of the brain confirmed bilateral periventricular white matter disease and cystic abnormalities in the right lateral ventricle (**Figure 2**). While the ophthalmologic evaluation was unremarkable, the hearing test revealed a moderate deficit in both ears. After initiating treatment with intravenous (IV) ganciclovir, a gradual increase in the platelet count was observed. After the platelet count stabilized, IV ganciclovir was replaced with oral valganciclovir.

Newborns with cCMV infection may have nonspecific findings such as petechiae, decreased platelet count, hepatosplenomegaly, jaundice at birth, chorioretinitis, sensorineural hearing loss, and seizures^{1,2}. Intracranial calcifications, white matter disease, cystic abnormalities, periventricular leukomalacia, and ventriculomegaly may be observed on neuroimaging². Since early antiviral therapy reduces the long-term adverse outcomes (mostly hearing impairment) and neurologic sequelae, cCMV infection should be

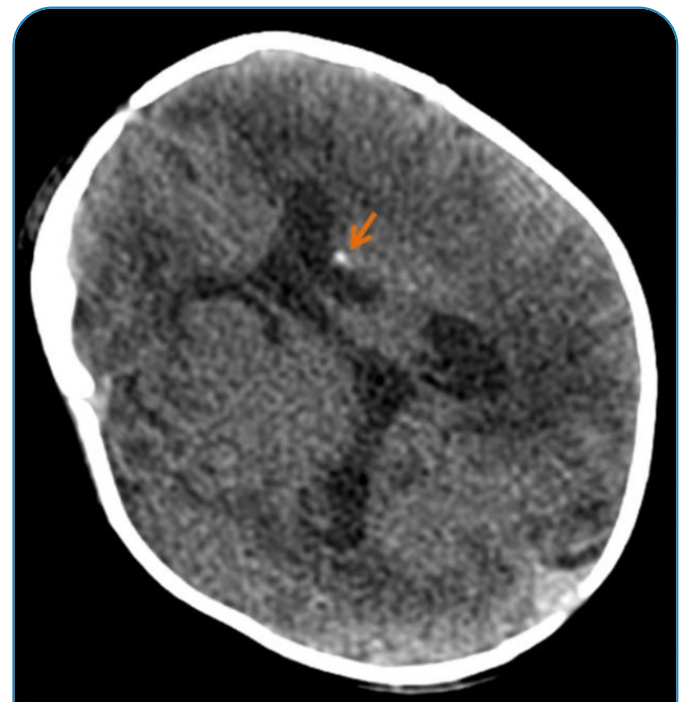


FIGURE 1: Unenhanced axial computed tomography image shows periventricular calcification (arrow) and hypodense white matter involvement.

Corresponding author: Dr. Melis Deniz. **e-mail:** mlsdnz@gmail.com

Authors' contribution: MD: Conception and design of the study, acquisition of data, drafting the article, final approval of the version to be submitted. MFD: Proofreading the article, final approval of the version to be submitted. NGA: Evaluating images, final approval of the version to be submitted.

Conflict of Interest: The authors report no conflict of interest.

Financial Support: The authors declare that there was not financial support.

Received 28 September 2022 | **Accepted** 22 November 2022

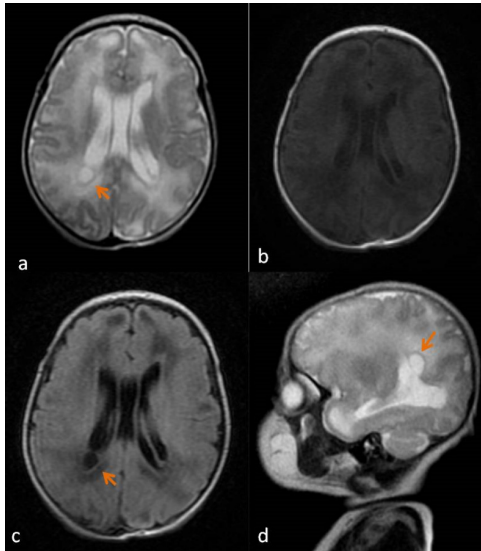


FIGURE 2: Axial T2-weighted (a), T1-weighted (b), FLAIR (c), and sagittal T2-weighted (d) images show diffuse white matter signal abnormality, a finding indicative of delayed myelination. Subependymal cyst (arrows) at the right posterior periventricular region is also demonstrated.

considered as one of the causes of thrombocytopenia in neonates and antiviral therapy should be started in the first month of life³.

ACKNOWLEDGEMENTS

We thank the staff and Sanliurfa Research and Training Hospital for their assistance.

REFERENCES

1. Dollard SC, Dreon M, Hernandez-Alvarado N, Amin MM, Wong P, Lanzieri TM, et al. Sensitivity of Dried Blood Spot Testing for Detection of Congenital Cytomegalovirus Infection. *JAMA Pediatr.* 2021;175(3):e205441. Available from: <https://doi.org/10.1001/jamapediatrics.2020.5441>
2. Kylat RI, Kelly EN, Ford-Jones EL. Clinical findings and adverse outcome in neonates with symptomatic congenital cytomegalovirus (SCCMV) infection. *Eur J Pediatr.* 2006;165(11):773-8.
3. Kimberlin DW, Jester PM, Sánchez PJ, Ahmed A, Arav-Boger R, Michaels MG, et al. Valganciclovir for symptomatic congenital cytomegalovirus disease. *N Engl J Med.* 2015;372(10):933-43.