Severe Brain Damage in a Moderate Preterm Infant as Complication of Post-COVID-19 Response during Pregnancy

Viktoria Engert\textsuperscript{a} Celine Siauw\textsuperscript{a} Annika Stock\textsuperscript{b} Monika Rehn\textsuperscript{c} Achim Wöckel\textsuperscript{c} Christoph Härtel\textsuperscript{a} Johannes Wirbelauer\textsuperscript{a}

\textsuperscript{a}Department of Pediatrics, University Hospital of Würzburg, Würzburg, Germany; \textsuperscript{b}Department of Neuroradiology, University Hospital of Würzburg, Würzburg, Germany; \textsuperscript{c}Department of Obstetrics and Women’s Health, University Hospital of Würzburg, Würzburg, Germany

Established Facts

- SARS-CoV-2 infections may be transmitted in utero and in the perinatal period.
- Multisystem inflammatory syndromes can occur after SARS-CoV-2 infections, particularly in the first 2 decades of life.

Novel Insights

- SARS-CoV-2 infection during pregnancy may cause neurological damage to the fetus.
- Etiological workup of unusual neonatal presentations should include serological screening for SARS-CoV-2 antibodies, in particular during COVID-19 pandemic.

Keywords

Intracranial hemorrhage · Periventricular leukomalacia · COVID-19 · Newborn

Abstract

Current evidence from the COVID-19 pandemic suggests that neonatal SARS-coronavirus-2 infections usually have a mild course. Data on how maternal infection during pregnancy affects fetal development are scarce. We present the unique case of a moderate preterm infant with intracranial bleeding and periventricular leukomalacia as a potential consequence of post-COVID-19 hyperinflammation during pregnancy.

Introduction

It is well acknowledged that infections during pregnancy with specific viruses, for example, cytomegalovirus, rubella virus, and zika virus, may represent major...
causes of permanent neurological disability in children. Current experience of the COVID-19 pandemic suggests that SARS-CoV-2 infection in pregnant women is associated with a higher risk for preterm delivery but usually mild or uneventful neonatal courses. Potential neurological manifestations of SARS-CoV-2 infection in the newborn are yet unclear [1–5]. Here, we report the complication of intracranial hemorrhage and periventricular leukomalacia in a moderate preterm infant born after maternal hyperinflammatory response following SARS-CoV-2 infection during the 2nd trimester of pregnancy.

Case Report

Perinatal History

The 25-year-old mother presented at 33 weeks of gestation with preterm labor but no other clinical abnormalities. Apart from mild cold symptoms at 16 weeks of gestation, maternal history including immune-mediated and coagulation disorders was unremarkable. Laboratory evaluation revealed increased markers of inflammation, for example, C-reactive protein 14.5 mg/dL and a total white blood cell count (WBC) of 183/mL with normal platelet counts and coagulation studies. There was no focus for any infection. Due to pathological cardiotocography and maternal signs of inflammation, caesarean section was performed at 33 5/7 weeks. The histological examination of the placenta revealed no signs of funisitis, chorioamnionitis, or placental infarction. Microbiological evaluation of swabs (cervix and placenta) was negative for any pathogen.

Neonatal Course

The preterm infant presented with Apgar score 4/7/7 after 1, 5, and 10 min (pH 7.28, base excess −5.0 mmol/L) and transitory tachypnea which was well controlled with respiratory support using continuous positive airway pressure. Ubiquitous petechial bleedings and pale hematomas were noticed on the skin but no other signs of clinical infection (shown in Fig. 1). Birth weight (2,100 g, 43rd percentile), length (45 cm, 48th percentile), and head circumference (30 cm, 21st percentile) were appropriate for gestational age. Day 1 cerebral ultrasound demonstrated cortical hyperechogenicity and cystic periventricular lesions (shown in Fig. 2). Laboratory studies revealed increased numbers of WBC (48.6/mL, nucleated red blood cells 27/100 WBCs), slightly increased procalcitonin (2.1 ng/mL, ref <0.5 ng/mL), but unremarkable C-reactive protein (<0.1 mg/dL). The initial hemoglobin value (17.9 g/dL, ref. 14.3–19 g/dL) was normal and the platelet count increased procalcitonin (2.1 ng/mL, ref <0.5 ng/mL), but unremarkable. Laboratory evaluation revealed increased markers of inflammation, caesarean section was performed at 33 5/7 weeks. The histological examination of the placenta revealed no signs of funisitis, chorioamnionitis, or placental infarction. Microbiological evaluation of swabs (cervix and placenta) was negative for any pathogen.

Maternal COVID-19 Infection

Maternal serology testing for SARS-CoV-2 (ELISA) revealed IgG antibodies against S1-protein (1.2 ratio; pos >1.0; Euroimmun, Germany) and against N-protein (anti-N-protein IgG/IgM/IgA index 15.2, ref. positive >1.0, Roche, Germany). Anti-N-protein IgG (34 U/mL, ref. positive >24; anti-N-protein IgG/IgM/IgA index 4.42, ref. positive >1.0) was also prevalent in the newborn serum, while specific anti-N-protein IgM and anti-S1/S2 IgM and IgG were negative. PCR test of SARS-CoV-2 after birth was negative in the mother (swab) and infant (swab and CSF). The infant recovered well and was discharged at 36 days of life without major clinical findings for further close follow-up.

Discussion

Our case report generates the hypothesis that a post-COVID systemic inflammatory response during pregnancy affected the fetal circulation followed by a coagulopathy and preterm labor. One proposed underlying mechanism might be the placental overexpression of angiotensin-converting enzyme 2 – the SARS-CoV-2 receptor – which can critically modulate hemodynamics within the uteroplacental unit [6]. Second, the proinflammatory cytokine storm invoked by SARS-CoV-2 may induce even more severe inflammation with deleterious consequences on the fetal brain during a critical time frame (e.g., development of periventricular leukomalacia). Further mechanistic links exist between hyperinflammation, endothelial activation, and dysregulated complement...
function which are known pathways to mediate microvascular injury, vasculitis, and coagulopathy [7–9].

Systemic inflammatory responses and coagulation disturbances are known complications of acute SARS-CoV-2 infection in intensive care patients [10, 11]. Hyperinflammatory responses also occur with temporal delay of several months after SARS-CoV-2 infection, for example, multisystem inflammatory syndrome in children. With a predominance in the first decades of life, post-COVID hyperinflammation is rather related to antibody-dependent enhancement of acquired immune responses to virus than direct virus replication. In the case of a 16-year-old boy, vascular inflammation was reported including stenosis of cerebral vessels leading to stroke [8]. In the specific context of immunotolerance during pregnancy, neonatal consequences of post-COVID hyperinflammation have not been described yet. In our unique case, diagnostic signs of inflammation and coagulopathy were intracranial and pettechial bleeding (without abnormal platelet count) as well as highly elevated D-dimers and WBC count.

Two other case reports describe neurological symptoms in newborns in relation to a SARS-CoV-2 infection.
due to perinatal transmission. The newborns presented with encephalitic symptoms such as lethargy, irritability, axial hypertonia, and high-pitched crying but unremarkable CSF results. Cerebral imaging was normal in 1 affected infant, while white matter injury was diagnosed in the other case being suggestive for cerebrovascular inflammation [12, 13].

To our knowledge, we present the first report of severe brain damage as a potential consequence of SARS-CoV-2-mediated hyperinflammation during pregnancy. We propose that the current pandemic situation requires serological screening of mother-infant dyads for SARS-CoV-2 antibodies when unusual clinical appearances of vasculitis, inflammation, or coagulopathy are obvious. Furthermore, all infants of mothers with known SARS-CoV-2 infection during pregnancy should be monitored for adverse neurological findings.

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**References**