Association of antepartum and intrapartum SARS-CoV-2 infection on pregnancy outcomes in South African women, two observational studies

Marta Nunes¹, Stephanie Jones¹, Renate Strehlau¹, Vuyelwa Baba¹, Zanele Ditse¹, Kelly Da Silva¹, Lane Bothma¹, Natali Serafin¹, Vicky Baillie¹, Gaurav Kwatra¹, Megan Burke¹, Amy Wise¹, Mary Adam¹, Philiswa Mlandu², Mpolokeng Melamu², Juliette Phelp², Wendy Fraser³, Colleen Wright³, Elizabeth Zell⁴, Yasmin Adam⁵, and Shabir Madhi¹

¹University of the Witwatersrand Faculty of Health Sciences
²Chris Hani Baragwaneth Academic Hospital, University of the Witwatersrand, Johannesburg
³Lancet Laboratories
⁴Stat-Epi Associates, Inc.
⁵Chris Hani Baragwanath Academic Hospital and the University of the Witwatersrand

November 18, 2021

Abstract

Objective: Evaluate the impact of the timing of SARS-CoV-2 infection on pregnancy outcomes in a low-middle income setting.

Design: two parallel, observational studies. Setting and population: pregnant women or women presenting for labour, enrolled between April-September 2020, in South Africa. Methods: i) longitudinal follow-up study of symptomatic or asymptomatic pregnant women investigated for SARS-CoV-2 infection antenatally, ii) cross-sectional study of SARS-CoV-2 infection at time of labour. SARS-CoV-2 infection was investigated by nucleic acid amplification test (NAAT). Main Outcome Measures: association of SARS-CoV-2 infection on nasal swab and birth outcomes. Results: Antenatally, 793 women were tested for SARS-CoV-2. Overall SARS-CoV-2 infection was confirmed in 138 women, including 119/275 with symptomatic illness (COVID-19) and 19/518 asymptomatic women; 493 women were asymptomatic and SARS-CoV-2 non-reactive. Women with COVID-19 were 1.66-times (95%CI: 1.02, 1.71) more likely to have a low-birthweight newborn (30%) compared to asymptomatic women without SARS-CoV-2 (21%). Overall, 3117 women were tested for SARS-CoV-2 infection at delivery, including 1560 healthy women with an uncomplicated term delivery. Adverse birth outcomes or pregnancy-related complications were not associated with infection at delivery. Among women with SARS-CoV-2 infection at delivery, NAAT was reactive on 6/98 of maternal blood samples, 8/93 of cord-blood, 14/54 of placentas and 3/22 of nasopharyngeal swabs from newborns collected within 72-hours of birth. Conclusions: Antenatal, but not intrapartum, SARS-CoV-2 infection was associated with low-birthweight delivery. Maternal infection at the time of labour was associated with in utero foetal and placental infection, and possible vertical and/or horizontal viral transfer to the newborn.