

Acknowledgment

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About the Author

Dr. Safdar is an infectious diseases physician, scientist, and hospital epidemiologist. Her research focuses on patient and healthcare worker safety in healthcare settings.

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Address for correspondence: Nasia Safdar, University of Wisconsin School of Medicine and Public Health, Department of Population Health Sciences, 1685 Highland Ave, Madison, WI 53705, USA; email: ns2@medicine.wisc.edu

Disappearance of SARS-CoV-2 Antibodies in Infants Born to Women with COVID-19, Wuhan, China

Jinzhi Gao,¹ Wei Li,¹ Xiaolin Hu, Ying Wei, Jianli Wu, Xiaoping Luo, Suhua Chen, Ling Chen

Author affiliation: Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, China

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¹These first authors contributed equally to this article.

We report the detection and decline over time of severe acute respiratory syndrome coronavirus 2 antibodies in infants born to women with coronavirus disease. Among 11 infants tested at birth, all had detectable IgG and 5 had detectable IgM. IgG titers with positive IgM declined more slowly than those without.

Although the diagnosis of coronavirus disease (COVID-19) by reverse transcription PCR (RT-PCR) is efficient and specific, IgM and IgG production and decay are useful to assess past or recent infection, especially for patients with negative nucleic acid tests (1). Evidence of IgM and IgG in adults with COVID-19 appeared around 13 days after illness onset (2). Plateau IgM levels lasted for 4 weeks and gradually declined (3). Although IgG lasted for a longer time, only 19.5% patients had a ≥ 4 -fold increase in titers during convalescence, a finding that was helpful for diagnosis of existing or acute infection (2,3).

However, to our knowledge, antibody persistence in infants born to women with COVID-19 has not yet been reported. IgM is the antibody isotype produced initially in the immune response and the first immunoglobulin class to be synthesized by a fetus or infant. Maternal IgM does not cross the placental barrier intact; therefore, positive IgM in early infants is potential evidence of intrauterine vertical transmission (1). Although IgG is transferred passively from mother to fetus through the placenta, the duration of passive immunity from maternal IgG is still unclear.

We implemented assays for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2)-specific antibodies and SARS-CoV-2 nucleic acid tests in 64 infants admitted to the neonatal section of Tongji Hospital (Wuhan, China) during January 19–April 12, 2020. Among these, 24 infants (ranging in gestational age from 31 weeks to 41 weeks, 2 days) were born to women with PCR-confirmed COVID-19 (Table) and 40 infants (ranging in gestational age from 35 weeks, 3 days, to 41 weeks, 3 days) were born to women without COVID-19. Because antibody testing was implemented in early March, the timing of antibody testing in infants was inconsistent. We conducted SARS-CoV-2 nucleic acid tests by using a qualitative SARS-CoV-2 RT-PCR (DAAn GENE Biotech, <http://www.daangene.com>). We performed quantitative assessment of IgG and IgM by using the IFlash3000 Chemiluminescence Immunoassay Analyzer (YHLO Biotech, <http://en.szyhlo.com>), which has been proven to be a highly accurate method to detect SARS-CoV-2 antibodies (4). We considered IgM or IgG titers >10 AU/mL to be positive.

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Among the 40 infants born to women without COVID-19, results of nucleic acid tests from throat and anal swab specimens and results of antibody assays were negative. Among the 24 infants born to women with COVID-19, 15 (62.5%) had detectable IgG and 6 (25.0%) had detectable IgM; nucleic acid test results were all negative. None of the 24 infants had complications related to pneumonia, a finding that is consistent with a previous report (5). Among

11 infants with antibody titers detected at birth, all had detectable IgG (100%) and 5 (45.5%) had detectable IgM, 1 of whom had high IgM levels (infant described in case 19 in the Table). Although the IgG titers in all 15 infants with positive IgG decreased gradually, the IgG levels declined more slowly in infants with positive IgM compared with those without (Figure). The infant described in case 19 was born 33 days after the mother had COVID-19

Table. Sequential severe acute respiratory syndrome coronavirus 2–specific antibodies assay in 24 infants born to mothers with PCR-confirmed coronavirus disease, Wuhan, China, January 19–April 12, 2020*

Case no.	IgM/IgG titers, AU/mL, by age of infant							GA, wk+d	
	0 d	7 d	14 d	28 d	45 d	60 d	75 d	At symptom onset	At birth
1	Mother						7.5/116.9*	33+4	36+3
	Infant						0.8/0.3		
2	Mother						4.4/118.3	38+5	38+5
	Infant						0.5/2.5		
3	Mother					5.0/83.3		37+5	38+2
	Infant					0.4/0.2			
4	Mother				14.5/100.8			39+1	39+5
	Infant				0.3/0.3				
5	Mother							34	35+2
	Infant	0.7/2.8	0.4/3.5	0.3/0.9					
6	Mother					7.0/93.0		39+3	40
	Infant					0.4/0.3			
7	Mother					51.1/94.6		39+4	41+2
	Infant					0.3/0.9			
8	Mother						67.0/92.6	37+4	38+4
	Infant						0.3/0.1		
9	Mother					17.8/104.3		38+2	39+5
	Infant					0.4/8.2			
10	Mother				15.3/95.4		9.9/108.7	38+1	39+5
	Infant				2.2/32.7		0.7/19.5		
11	Mother				11.3/112.5		5.3/105.6	35+5	38+2
	Infant	3.6/166.0	5.3/120.8	1.0/36.2			0.4/16.2		
12	Mother					+/+†	37.9/94.6	29+4	31
	Infant	5.7/142.0	2.6/113.9	1.8/75.3	0.3/22.0		0.4/11.3		
13	Mother				+/+			34+3	38
	Infant	3.5/134.1	2.4/125.0	2.8/119.0	0.7/24.6				
14	Mother						1.9/59.5	34+6	39
	Infant	2.2/78.4	1.5/76.9	3.0/80.9	7.0/61.1	0.8/28.6	0.5/13.0		
15	Mother	0.7/58.9						NA	37+2
	Infant	8.5/174.5			7.8/122.7				
16	Mother	+/+						NA	38+3
	Infant	0.7/101.0							
17	Mother	+/+				1.5/104.2		NA	38+3
	Infant	1.2/110.4	10.4/108.9		2.1/68.2				
18	Mother	-/+						NA	39+5
	Infant	6.6/96.2	11.3/86.0	11.9/88.3	4.8/54.5	2.5/60.9			
19	Mother	2,581.6/281			40.2/89.8	13.9/99.6	6.9/94.2	32+1	38
	Infant	184.3/147.2	42.6/161.6	21.2/83.8	5.5/79.2	1.6/57.0	0.3/25.9		
20	Mother						6.0/81.3	31+1	39+5
	Infant			6.9/82.8			0.4/47.4		
21	Mother	+/+					5.1/139.0	30+3	38
	Infant	26.0/145.8	41.8/120.1	42.9/110.1	38.0/89.5		14.2/105.3		
22	Mother	+/+						30	30
	Infant	22.8/116.5	11.8/104.4	8.7/99.8	1.9/75.2				
23	Mother	+/+					4.3/115.6	32+1	32+1
	Infant	15.8/222.1	33.8/182.5	9.0/88.2	10.5/101.6		0.5/31.1		
24	Mother	+/+	-/+			8.0/83.9		NA	NA
	Infant	30.1/106.0	29.5/84.6	31.5/94.9	9.7/73.1				

*Bold text indicates titer >10 AU/mL (i.e., above cutoff). GA, gestational age; NA, not available; -, negative; +, positive.
 †Some mothers delivered in other hospitals, and antibody testing was conducted using other qualitative antibody-detection kits.

diagnosed (41 days since symptom onset) and by elective cesarean section at 38 weeks' gestation because of the mother's previous cesarean section. The infant had high IgM and IgG titers in the umbilical cord blood and peripheral blood on the first day of life, which gradually decreased at repeated tests thereafter (Table). The infant also had negative nucleic acid tests results in a series of specimens, including cord blood, placenta, amniotic fluid, stool, urine, peripheral blood, and gastric juice at different timepoints. The placenta sample described in case 19 was collected during surgery and sent for pathologic

examination, which revealed slight inflammation with slight fibrin deposition and lymphocyte infiltrates (Appendix Figure, <https://wwwnc.cdc.gov/EID/article/26/10/20-2328-App1.pdf>).

Although positive results in SARS-CoV-2 nucleic acid or virus-specific IgM in infants have been reported previously, evidence of vertical transmission of SARS-CoV-2 is not complete (6–8). We report the dynamic changes of SARS-CoV-2-specific antibodies in infants born to mothers with COVID-19. Five of 11 infants were seropositive for IgM at birth; however, these findings were not sufficient to confirm

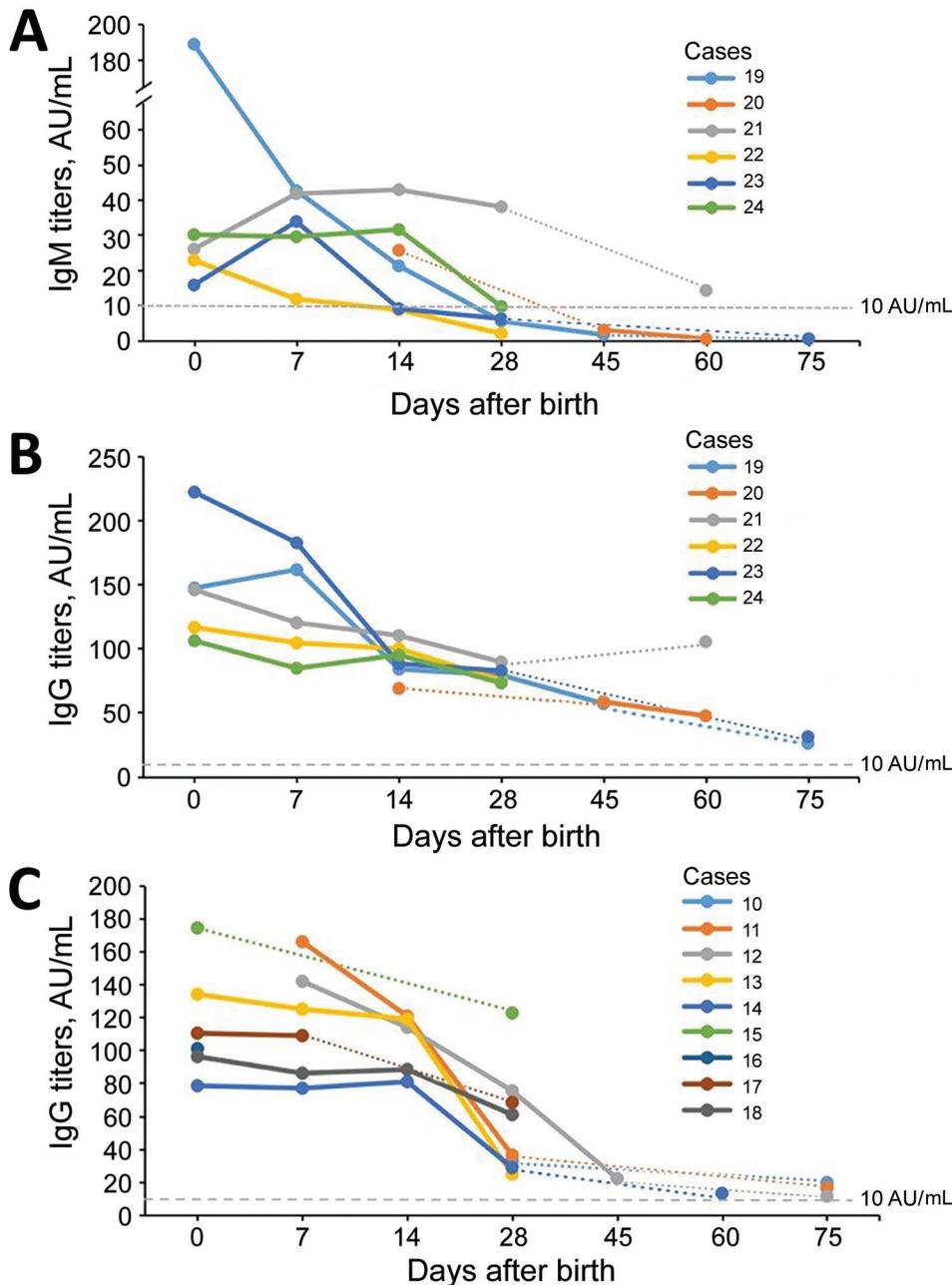


Figure. Temporal changes in severe acute respiratory syndrome coronavirus 2-specific antibodies in infants born to women with coronavirus disease, Wuhan, China. A, B) Dynamic changes of IgM (A) and IgG (B) titers in infants with positive IgM. C) Dynamic changes of IgG titers in infants with negative IgM. The IgM and IgG titers gradually decreased with time. IgG titers with positive IgM declined more slowly than those without, and the duration was as long as 75 days.

SARS-CoV-2 vertical transmission without positive nucleic acid testing. The study was also limited by a small sample size. However, these findings show a rapid rate of decline in antibody titers, suggesting lack of protective passive immunity in infants, and IgM detection in infants, supporting a growing body of evidence of possible vertical transmission. We still do not have a correlate of immunity (e.g., we do not know exactly what level of antibody titers are considered protective against infection), and whether infants testing positive by PCR at birth have higher levels of IgM or IgG remains to be seen. More work is needed to understand SARS-CoV-2 immunity in infants; such findings might have implications for potential vaccination efforts.

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About the Author

Dr. Gao is a physician in the Department of Pediatrics, Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, China. Her primary research interests include neonatology and infectious diseases.

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Address for correspondence: Ling Chen, Department of Pediatrics, Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, 1095 Jiefang Ave, Wuhan 430030, China; email: chenling@tjh.tjmu.edu.cn; Suhua Chen, Department of Obstetrics and Gynecology, Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, 1095 Jiefang Ave, Wuhan 430030, China; email: tj_csh@163.com

Culture-Competent SARS-CoV-2 in Nasopharynx of Symptomatic Neonates, Children, and Adolescents

Arnaud G. L'Huillier,¹ Giulia Torriani,¹ Fiona Pigny, Laurent Kaiser, Isabella Eckerle

Author affiliation: Geneva University Hospitals and Faculty of Medicine, University of Geneva, Geneva, Switzerland

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Children do not seem to drive transmission of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). We isolated culture-competent virus in vitro from 12 (52%) of 23 SARS-CoV-2-infected children; the youngest was 7 days old. Our findings show that symptomatic neonates, children, and teenagers shed infectious SARS-CoV-2, suggesting that transmission from them is plausible.

Children are underrepresented in coronavirus disease (COVID-19) case numbers (1,2). Severity in most children is limited, and children do not seem to be major drivers of transmission (3,4). However, severe acute respiratory syndrome

¹These authors contributed equally to this article.