

Original Research

Maternal and Neonatal Outcomes during the COVID-19 Infection in China: Data from a Retrospective Cohort Study

Yingying Lin^{1,†}, Zhiwei Chen^{2,3,†}, Zhiwei Wu³, Yongjin Xie³, Li Xie^{3,*}, Xiaoyan Xiu^{4,*}

Academic Editor: Paolo Ivo Cavoretto

Submitted: 27 November 2023 Revised: 9 January 2024 Accepted: 23 January 2024 Published: 15 April 2024

Abstract

Background: This study aimed to investigate the relationship between coronavirus disease 2019 (COVID-19) disease and maternal and neonatal outcomes. **Methods**: This is a retrospective cohort study. This study analyzed 3615 participants from Fujian Provincial Maternity and Child Health Hospital Fujian Obstetrics and Gynecology Hospital from November 1, 2022, to January 31, 2023. All pregnant women must provide a 24-hour nucleic acid test report when they are admitted to the hospital to deliver babies. Chi-square test and linear regression analyses were used to evaluate the risk of COVID-19 infection with the maternal outcome and neonatal outcomes. **Results**: Finally, 3615 patients were included in the cohort study. 549 (15.2%) were diagnosed with COVID-19 infection. The most common symptom is fever, cough, sore throat. 51 (9.2%) newborns had positive test results. In addition, the mother with COVID-19 infection were significantly associated with a higher rate of premature rupture of membranes (PPROM) and postpartum hemorrhage. Furthermore, a mother with COVID-19 infection was significantly associated with a higher rate of low birth weight infant (LBW) and macrosomia in newborns, higher rate of respiratory distress syndrome (RDS), higher rate of intro-ventricular hemorrhage (IVH), higher rate of neonatal pneumonia, and a higher rate of aspiration of amniotic fluid and meconium syndrome (AAFMS) (all p < 0.05). **Conclusions**: This study proves that COVID-19 infection is a potential risk factor for adverse pregnancy outcomes. The COVID-19 epidemic continues, requiring targeted public health measures to reduce the infection rate in pregnant women and the poor prognosis of mothers and children.

Keywords: China; COVID-19; outcomes; pregnancy; SARS-CoV-2

1. Introduction

The emergence of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) caused coronavirus disease 2019 (COVID-19), posing a massive threat to public health [1]. It has led to more than 6.3 million deaths by midway through 2022 [2]. SARS-CoV-2 took the respiratory tract as the leading invasion site, with cough, shortness of breath, and fever as the main symptoms. In addition, it can also cause signs in the digestive system, cardiovascular system, nervous system, reproductive system, immune system, and urinary system [3–6]. The severity of COVID-19 varies by population. Studies have indicated that the elderly, men, and individuals with other cardiovascular commodities are at increased risk of developing critical illness [7,8]. Pregnancy is considered an independent risk factor for severe COVID-19 due to immunological and physical changes during pregnancy [9]. Therefore, studies about COVID-19 in pregnancy and the pregnant outcomes provide essential information for obstetric care.

With the spread of the epidemic, for the particular population of pregnant women, the public is paying more attention to whether the novel coronavirus can lead to ver-

tical transmission from mother to child and the impact of the virus on pregnancy complications and adverse pregnancy outcomes in pregnant women. Some studies demonstrated that the risk of severe disease after COVID-19 infection was not significantly higher than in pregnant womento-non-pregnant women in the early days of the COVID-19 pandemic [10-12]. However, recent studies revealed COVID-19 infection had a higher risk of developing severe maternal disease and the fetus [13–16]. Papageorghiou et al. [17] found that COVID-19 in pregnancy is strongly associated with preeclampsia, especially in non-parturient women. Pre-pregnancy vaccination also has an impact on the outcome of a pregnancy infected with COVID-19. Villar et al. [18] found that infection with COVID-19 of pregnancy is associated with an increased risk of severe maternal morbidity and death, especially in symptomatic and unvaccinated women, and vaccination reduces the risk of serious complications and death in pregnant women. Pregnant women may still need to be vaccinated. In addition, diabetes mellitus and overweight or obesity were risk factors for COVID-19 diagnosis in pregnancy [19]. Currently, there is a controversy about the symptoms of COVID-19 infection and its impact on pregnancy outcomes. Opti-

¹Department of Healthcare, Fujian Maternity and Child Health Hospital, 350001 Fuzhou, Fujian, China

²College of Clinical Medicine for Obstetrics & Gynecology and Pediatrics, Fujian Medical University, 350001 Fuzhou, Fujian, China

³Department of Gynecology and Obstetrics, Fujian Maternity and Child Health Hospital, 350001 Fuzhou, Fujian, China

⁴Department of Health Education, Fujian Maternity and Child Health Hospital, 350001 Fuzhou, Fujian, China

^{*}Correspondence: 309381162@qq.com (Li Xie); xiuxiaoyan1982@fjmu.edu.cn (Xiaoyan Xiu)

[†]These authors contributed equally.

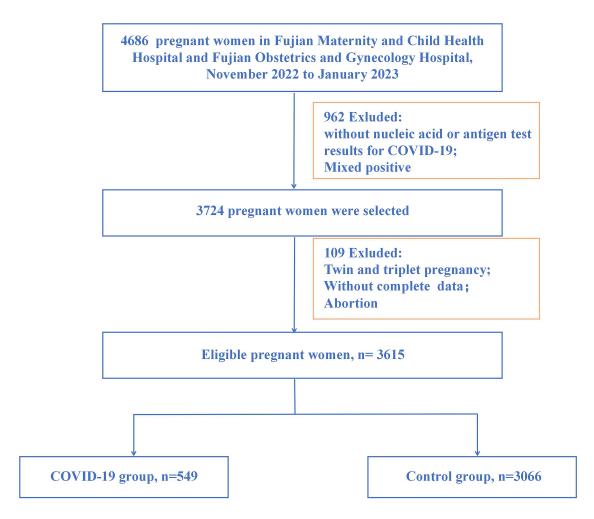


Fig. 1. Flowchart of participants in the study. COVID-19, coronavirus disease 2019.

mal management of the mother and newborn remains unknown. Therefore, it is still necessary to carry out research on COVID-19 infection and pregnancy outcome.

There is growing evidence that COVID-19 also poses a higher risk of severe illness in pregnant women, and threats to this population can, unfortunately, be overlooked. In addition, a few studies are comparing pregnant women with COVID-19 and without COVID-19 in China mainland due to the government's strict prevention and control policies [20,21]. Under prevention and control protection, the COVID-19 infection rate of pregnant women is shallow, so research on pregnant women cannot be provided. Unfortunately, at the end of 2022, with the national development and adjustment of the epidemic prevention and control policy, the COVID-19 infection rate of the population increased, and the number of pregnant women also increased significantly. Therefore, it is urgent to study the symptoms of pregnant women after COVID-19 infection and the impact on pregnancy outcomes in China.

In this study, we conducted a large cohort study to analyze the relationship between pregnant women with or without COVID-19 infection and the clinical outcomes with maternal-neonatal outcomes in China. We aimed to evaluate whether COVID-19 infection is a risk factor for pregnancy complications and adverse neonatal effects to provide best practices regarding infection control in China.

2. Materials and Methods

2.1 Participants

This is a retrospective cohort study. This study analyzed 3615 participants from Fujian Provincial Maternity and Child Health Hospital and Fujian Obstetrics and Gynecology Hospital from November 1, 2022, to January 31, 2023 (Fig. 1). Pregnant women must meet the following criteria: (1) single pregnancy; (2) with nucleic acid or antigen test; (3) complete clinical data. Those women were excluded due to the following criteria: (1) the functions of vital organs such as heart, liver and kidney are severely deficient; (2) fetal anomalies; (3) people who do not have a fever at admission but are at risk of intrapartum fever. Maternal and newborn data were obtained from outpatient and inpatient records. The study was approved by the Hospital Ethics Committee of Fujian Provincial Maternity and Children's Health Hospital, an affiliated hospital of Fujian Medical University (2023KY005).



Table 1. Maternal demographic and characteristics of the participants.

	0 1			p-value	
Characteristic	Total (N = 3615)	COVID-19 (N = 549)	No COVID-19 (N = 3066)		
Maternal age (years)	30.2 ± 4.3	30.4 ± 4.4	30.2 ± 4.3	0.319	
Gravida	2(1,3)	2 (1, 3)	2(1, 3)	0.617	
Parity	1 (1, 2)	1 (1, 2)	1 (1, 2)	0.637	
Pre-BMI (kg/m ²)				0.740	
Underweight	510 (14.1)	82 (14.9)	428 (14.0)		
Normal weight	2440 (67.5)	363 (66.1)	2077 (67.7)		
Overweight	665 (18.4)	104 (19.0)	561 (18.3)		
Education				0.208	
Junior school or under	393 (10.9)	71 (12.9)	322 (10.5)		
High school	444 (12.3)	62 (11.3)	382 (12.5)		
Undergraduate or higher	2778 (76.8)	416 (75.8)	2362 (77.0)		
Occupation				0.160	
White-collar workers and civil servant	563 (15.6)	80 (14.6)	483 (15.8)		
Professionals	893 (24.7)	155 (28.2)	738 (24.1)		
Domestic works	472 (13.1)	69 (12.6)	403 (13.1)		
Laborers	321 (8.9)	36 (6.6)	285 (9.3)		
Student	15 (0.4)	2 (0.4)	13 (0.4)		
Servant	162 (4.5)	30 (5.5)	132 (4.3)		
Other	1189 (32.9)	177 (32.2)	1012 (33.0)		
Hypertension	221 (6.1)	39 (7.1)	182 (5.9)	0.293	
Diabetes mellitus	41 (1.1)	9 (1.6)	32 (1.0)	0.225	

Continuous variables are presented as mean \pm SD (standard deviation range) and categorical variables as n (%). Notes: BMI, body mass index; COVID-19, coronavirus disease 2019.

2.2 Procedures

According to the local prevention and control policies, all pregnant women must provide a 24-hour nucleic acid test report to deliver babies when admitted to the hospital. When the mother occurred symptoms of fever, sore throat, cough, shortness of breath, anosmia, ageusia, diarrhea, rhinorrhoea, myalgias, and vomiting, nucleic acid testing is required again.

Testing for SARS-CoV-2 was done using real-time polymerase chain reaction (RT-PCR); RealStar SARS-CoV-2 RT-PCR Kit, cobas SARS-CoV-2 Test (DAAN GENE, Guangzhou, Guangdong, China). Turnaround time from specimen collection to result from reporting was 24 h. Neonates with mother under COVID-19 infection were tested for SARS-CoV-2 by RT-PCR on a nasopharyngeal swab sample at 12–24 h, 2–7 days, and 8–28 days of life and as indicated at subsequent visits. These time points were chosen to allow repeat testing and routine neonatal care.

2.3 Definition

Symptomatic COVID-19 pregnancies were defined as the pregnancies had COVID-19 positive tests and the common symptom, such as fever, cough, sore throat, pain, and fatigue. Non-symptomatic COVID-19 pregnancies were defined as the pregnancies had positive tests but not the symptom [22]. Newborn COVID-19 refers to the period be-

tween delivery of the mother and 28 days after birth when the fetus tests positive for COVID-19. Pregnancy complication was defined as the adverse events that occurred during pregnancy or delivery times, including gestational diabetes mellitus (GDM), gestational hypertension, preterm premature rupture of membranes (PPROM), abnormal amniotic fluid (such as oligohydramnios and polyhydramnios), placental abruption, postpartum hemorrhage, and perineal laceration. An adverse neonatal outcome was defined as the presence of any of the following: neonatal death, intro-ventricular hemorrhage (IVH), patent ducts arterioles (PDA), respiratory distress syndrome (RDS), neonatal pneumonia, neonatal jaundice and aspiration of amniotic fluid and meconium syndrome (AAFMS).

2.4 Statistical Analysis

The counting data used t-tests, and the measurement data were analyzed with Pearson chi-squared tests by SPSS version 26.0 (IBM, Armonk, NY, USA). Pearson chi-squared tests were used to assess the differences in demographic and pregnancy characteristics by COVID-19 infection. Spearman rank correlation coefficients were analyzed to evaluate the relationships between the symptoms of COVID-19 disease and maternal and neonatal outcomes. In all statistical tests, the differences were considered statistically significant at p-values < 0.05.



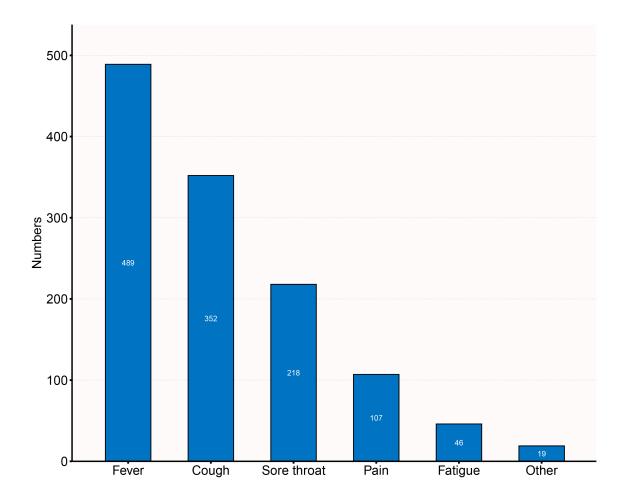


Fig. 2. Percentage of women with symptoms under COVID-19.

Table 2. Serial RT-PCR testing results for neonates.

24 h of life $(N = 549)$	2-7 days of life (N = 545)	8-28 days of life (N = 532)
549 (100%)	545 (100%)	532 (100%)
0	0	0
4 (0.7%)	13 (2.4%)	34 (6.4%)
545 (99.3%)	532 (97.6%)	498 (93.6%)
	549 (100%) 0 4 (0.7%)	549 (100%) 545 (100%) 0 0 4 (0.7%) 13 (2.4%)

Data are n (%) or n/N (%). RT-PCR, real-time polymerase chain reaction.

3. Results

3.1 The Clinical Characteristic of Pregnancy

Finally, a total of 3615 patients were included in the cohort study. The maternal outcomes and neonatal outcomes are shown in Table 1. The mean maternal age was 30.2 ± 4.3 years. The median and quartile gravid are 2(1,3) times, parity 1(1,2) times. According to the pre-BMI (body mass index), 510 (14.1%) pregnancies were underweight, 2440 (67.5%) were normal weight, and 665 (18.4%) were overweight. Regarding the education level, 393 (10.9%) women were in junior school or under, 444 (12.3%) were in high school, and 2778 (76.8%) were undergraduate or

higher. In the item of occupation, 563 (15.6%) women were white-collar workers and civil servants, 893 (24.7%) were professionals, 472 (13.1%) were domestic workers, 321 (8.9%) laborers, 15 (0.4%) student, 162 (4.5%) servant and 1189 (32.9%) others. In addition, we analyzed the basic disease before pregnancy. 222 (6.1%) had hypertension, and 41 (1.1%) had diabetes mellitus. There 549 (15.2%) were diagnosed with COVID-19 infection, and 3066 (84.8%) were without COVID-19 infection. Furthermore, there were no significant differences between the COVID-19 infection cohort regarding the basic information.



Table 3. The association between pregnancy outcomes and COVID-19 infection.

Variable	COVID-19 (N = 549) n (%)	No COVID-19 (N = 3066) n (%)	<i>p</i> -value	
Maternal outcomes				
PPROM	173 (31.5%)	766 (25.0%)	0.001	
Placental abruption	14 (2.6%)	85 (2.8%)	0.769	
Abnormal amniotic fluid	32 (5.8%)	191 (6.2%)	0.719	
GA at delivery (weeks)	38.2 ± 2.6	38.3 ± 2.5	0.114	
Cesarean delivery	215 (39.2%)	1119 (36.5%)	0.233	
Postpartum hemorrhage	24 (4.4%)	62 (2.0%)	0.001	
Perineal laceration	244 (44.4%)	1349 (44.0%)	0.846	
Neonatal Mortality	7 (1.3%)	54 (1.8%)	0.415	
Birth weight			0.022	
LBW	52 (9.5%)	231 (7.5%)		
Normal birth weight	469 (85.4%)	2736 (89.2%)		
Macrosomia	28 (5.1%)	99 (3.2%)		
Apgar score (1 min)	9.8 ± 1.3	9.7 ± 1.4	0.865	
Apgar score (5 min)	9.8 ± 1.2	9.8 ± 1.3	0.699	
Apgar score (10 min)	9.7 ± 1.2	9.8 ± 1.3	0.633	
NICU admission	39 (7.2%)	174 (5.8%)	0.200	
RDS	17 (3.1%)	43 (1.4%)	0.004	
IVH	25 (4.6%)	87 (2.8%)	0.033	
PDA	13 (2.4%)	108 (3.5%)	0.166	
Neonatal pneumonia	60 (10.9%)	236 (7.7%)	0.011	
Neonatal jaundice	47 (8.6%)	336 (11.0%)	0.093	
AAFMS	47 (8.6%)	127 (4.1%)	< 0.001	

Notes: COVID-19, coronavirus disease 2019; GA, gestational age; PPROM, preterm premature rupture of membranes; LBW, low birth weight infant; NICU, neonatal intensive care unit; RDS, respiratory distress syndrome; IVH, intraventricular haemorrhage; PDA, patent ductus arteriosus; AAFMS, aspiration of amniotic fluid and meconium syndrome.

3.2 The Symptomatic Characteristics of Women with COVID-19 Infection

Of the 549 with COVID-19 infection, 541 (98.5%) had symptoms. The specific symptoms and distribution are shown in Fig. 2. Fever is the most common of these 541 women with symptoms, affecting 489 (89.1%) women. 352 (64.1%) of 541 women had a cough, followed by 218 (39.7%) with a sore throat, 107 (19.5%) with pain, 46 (8.4%) with fatigue, and 19 (3.5%) with other symptoms, such as vomiting, diarrhea, weight change, and insomnia.

3.3 The Infection Rate of Neonates with Mothers under COVID-19 Infection

The RT-PCR results from a nasopharyngeal swab obtained at birth were available for all 549 neonates initially identified (Table 2). 545 (99.3%) neonates had a negative RT-PCR result, and 4 (0.7%) neonates had positive results. 545 neonates had repeat RT-PCR testing at 2–7 days of life, and 13 (2.4%) neonates had positive results. 532 neonates had recurrence RT-PCR testing at 8–28 days of life, of whom 34 (6.4%) had a positive test result.

3.4 The Relationships between the COVID-19 Infection and Maternal/Neonatal Outcomes

The relationships between COVID-19 infection and pregnancy outcomes are shown in Table 3. Some maternal outcomes were significantly associated with COVID-19 disease. Mothers with COVID-19 infection were significantly associated with a higher rate of PPROM and a higher rate of postpartum hemorrhage. In addition, some neonatal outcomes were significantly associated with COVID-19 infection. Mothers with COVID-19 infection were significantly associated with a higher rate of low birth weight infant (LBW) and macrosomia in newborns, a higher rate of RDS, a higher rate of IVH, a higher rate of neonatal pneumonia, and a higher rate of AAFMS (all p < 0.05).

3.5 The Relationships between the Maternal/Neonatal Outcomes and Symptoms of COVID-19 Infection

According to Table 4, we found that the symptoms of COVID-19 infection, such as fever, cough, sore throat, pain, and fatigue, were significantly associated with maternal outcomes. It suggested that positive symptoms indicated a higher rate of gestational hypertension (r = -0.141, p = 0.001). The mothers with pain, sore throat, and fatigue had a higher rate of PPROM (r = 0.091, p = 0.033; r = 0.114,



Table 4. Relationships between the COVID-19 symptoms and maternal-neonatal outcomes.

GDM r 0.015 −0.001 0.045 0.012 0.066 −0.039 Gestational hypertension r −0.141** −0.043 −0.033 −0.031 −0.035 0.027 p 0.001 0.310 0.442 0.470 0.414 0.534 PPROM r −0.063 0.021 0.091* 0.114** −0.042 0.024** PROM r −0.063 0.021 0.091* 0.114** −0.042 −0.042 −0.043 −0.042 −0.013 −0.048 0.034 PROM r 0.018 0.019 −0.051 −0.013 −0.048 0.034 Abnormal amniotic fluid r 0.028 −0.042 −0.011 −0.048 0.032 Abnormal amniotic fluid r 0.028 −0.042 −0.011 −0.048 0.032 Cesarean delivery r r −0.018 0.008 −0.082 −0.039 −0.021 −0.059 Cesarean delivery r r	Variable Variable		Symptoms	Fever	Pain	Sore throat	Cough	Fatigue
Gestational hypertension r −0.141** −0.043 −0.033 −0.031 −0.035 0.021 PPROM r −0.063 0.021 0.091* 0.114** −0.042 0.204** PPROM r −0.063 0.021 0.091* 0.114** −0.042 0.001* Placental abruption r 0.018 0.019 −0.051 −0.013 −0.048 0.034 Abnormal amniotic fluid r 0.028 −0.042 −0.101* −0.054 0.018 −0.045 Cesarean delivery r 0.018 0.008 −0.082 −0.039 −0.057 0.284 Cesarean delivery r 0.018 0.008 −0.082 −0.039 −0.057 0.284 Cesarean delivery r 0.018 0.008 −0.082 −0.039 −0.051 −0.059 p 0.676 0.885 0.055 0.359 0.631 0.172 Postpartum hemorrhage r 0.066 0.885 0.023 0	GDM	r	0.015	-0.001	0.045	0.012	0.066	-0.039
Gestational hypertension r -0.141*** -0.043 -0.033 -0.031 -0.035 0.021 PPROM r -0.063 0.021 0.091** 0.114** -0.042 0.204*** PPROM r -0.063 0.021 0.091** 0.114*** -0.042 0.001** Placental abruption r 0.018 0.019 -0.051 -0.013 -0.048 0.034 Abnormal amniotic fluid r 0.028 -0.042 -0.101* -0.054 0.018 -0.046 Cesarean delivery r 0.018 0.008 -0.022 0.019 -0.021 -0.059 Postpartum hemorrhage r 0.018 0.008 -0.082 -0.039 -0.051 -0.059 Postpartum hemorrhage r 0.0676 0.855 0.055 0.359 0.631 0.172 Postpartum hemorrhage r 0.0696 0.082 0.023 0.006 0.015 <0.001		p	0.731	0.973	0.293	0.774	0.120	0.360
PROM p 0.001 0.310 0.442 0.470 0.414 0.534 PPROM r -0.063 0.021 0.091* 0.114** -0.042 0.204*** Placental abruption r 0.018 0.019 -0.051 -0.013 -0.048 0.034 Abnormal ammiotic fluid r 0.028 -0.042 -0.101* -0.054 0.018 -0.046 Abnormal ammiotic fluid r 0.028 -0.042 -0.101* -0.054 0.018 -0.046 Cesarean delivery r -0.018 0.008 -0.082 -0.039 -0.021 -0.059 Postpartum hemorrhage r 0.0676 0.855 0.055 0.359 0.631 0.172 Postpartum hemorrhage r 0.069 0.039 -0.032 0.006 0.015 <0.001	Gestational hypertension		-0.141**	-0.043	-0.033	-0.031	-0.035	0.027
PPROM r -0.063 0.021 0.091* 0.114** -0.042 0.204** Placental abruption r 0.143 0.630 0.033 0.008 0.327 <0.001 Placental abruption r 0.018 0.019 -0.051 -0.013 -0.048 0.034 Abnormal amniotic fluid r 0.028 -0.042 -0.101* -0.054 0.018 -0.046 Postpartion r 0.018 0.008 -0.082 -0.039 -0.021 -0.059 Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450** Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450** Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450** Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450** Postpartum hemorrhage r 0.024	• •	p		0.310	0.442	0.470	0.414	0.534
Placental abruption	PPROM	r	-0.063	0.021	0.091*	0.114**	-0.042	0.204**
Placental abruption		p	0.143	0.630	0.033	0.008	0.327	< 0.001
Abnormal ammiotic fluid r 0.028 -0.042 -0.101* -0.054 0.018 -0.046 Cesarean delivery r -0.018 0.008 -0.082 -0.039 -0.021 -0.059 Postpartum hemorrhage r -0.018 0.008 -0.082 -0.039 -0.021 -0.059 Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450*** Perineal laceration r 0.069 0.039 -0.034 0.030 0.025 -0.058 Perineal laceration r 0.069 0.039 -0.034 0.030 0.025 -0.058 Mortality r 0.016 0.365 0.431 0.491 0.558 0.174 Mortality r 0.013 0.040 0.026 0.040 0.017 -0.058 MICU admission r -0.032 -0.020 0.011 -0.073 -0.036 0.153** Apgar score (1 min) r 0.037 0.010	Placental abruption	-	0.018	0.019	-0.051	-0.013	-0.048	
Abnormal ammiotic fluid r 0.028 -0.042 -0.101* -0.054 0.018 -0.046 Cesarean delivery r -0.018 0.008 -0.082 -0.039 -0.021 -0.059 Postpartum hemorrhage r -0.018 0.008 -0.082 -0.039 -0.021 -0.059 Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450** Postpartum hemorrhage r 0.069 0.039 -0.031 0.006 0.015 <0.001 Perineal laceration r 0.069 0.039 -0.034 0.030 0.025 -0.058 Pointality r 0.013 0.040 0.026 0.040 0.017 -0.058 Miculatity r 0.013 0.040 0.026 0.040 0.017 -0.035 Miculatity r 0.013 0.040 0.026 0.040 0.017 -0.035 Miculatity r 0.033 0.040 0.0	-	p	0.667	0.658	0.237	0.753	0.261	0.425
Cesarean delivery r -0.018 0.008 -0.082 -0.039 -0.021 -0.057 Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450*** Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450*** Postpartum hemorrhage r 0.069 0.039 -0.034 0.030 0.025 -0.058 Postpartum hemorrhage r 0.069 0.039 -0.034 0.030 0.025 -0.058 Postpartum hemorrhage r 0.069 0.039 -0.034 0.030 0.025 -0.058 Postpartum hemorrhage r 0.069 0.039 -0.034 0.030 0.025 -0.058 Postpartum hemorrhage r 0.069 0.039 -0.024 0.030 0.030 0.025 -0.058 Dold 0.010 0.040 0.017 -0.035 0.042 0.001 0.049 0.035 0.011 -0.073 0	Abnormal amniotic fluid		0.028	-0.042	-0.101*	-0.054	0.018	-0.046
Postpartum hemorrhage p 0.676 0.855 0.055 0.359 0.631 0.172 Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450** Perineal laceration r 0.069 0.039 -0.034 0.030 0.025 -0.058 Mortality r 0.013 0.040 0.026 0.040 0.017 -0.035 Mortality r 0.013 0.040 0.026 0.040 0.017 -0.035 NICU admission r -0.032 -0.020 0.011 -0.073 -0.036 0.153*** Apgar score (1 min) r 0.037 0.010 -0.010 0.001 0.049 -0.031 Apgar score (5 min) r 0.082 0.017 -0.008 0.012 -0.001 0.008 Apgar score (5 min) r 0.082 0.017 -0.008 0.012 -0.001 0.008 Apgar score (5 min) r 0.087* 0.022 -		p	0.515	0.322	0.018	0.209	0.675	0.284
Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450** Perineal laceration r 0.069 0.039 -0.034 0.030 0.025 -0.058 Perineal laceration r 0.069 0.039 -0.034 0.030 0.025 -0.058 Mortality r 0.013 0.040 0.026 0.040 0.017 -0.035 Mortality r 0.013 0.040 0.026 0.040 0.017 -0.035 MICU admission r -0.032 -0.020 0.011 -0.073 -0.036 0.153*** Apgar score (1 min) r 0.037 0.010 -0.010 0.001 0.049 -0.031 Apgar score (5 min) r 0.082 0.017 -0.008 0.012 -0.001 0.008 Apgar score (10 min) r 0.082 0.017 -0.008 0.012 -0.001 0.008 Apgar score (10 min) r 0.082 0.017	Cesarean delivery	r	-0.018	0.008	-0.082	-0.039	-0.021	-0.059
Postpartum hemorrhage r 0.024 0.074 0.097* 0.118** 0.104* 0.450** Perineal laceration r 0.069 0.039 -0.034 0.030 0.025 -0.058 Perineal laceration r 0.069 0.039 -0.034 0.030 0.025 -0.058 Mortality r 0.013 0.040 0.026 0.040 0.017 -0.035 Mortality r 0.013 0.040 0.026 0.040 0.017 -0.035 MICU admission r -0.032 -0.020 0.011 -0.073 -0.036 0.153*** Apgar score (1 min) r 0.037 0.010 -0.010 0.001 0.049 -0.031 Apgar score (5 min) r 0.082 0.017 -0.008 0.012 -0.001 0.008 Apgar score (10 min) r 0.082 0.017 -0.008 0.012 -0.001 0.008 Apgar score (10 min) r 0.082 0.017	•	p	0.676	0.855	0.055	0.359	0.631	0.172
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Postpartum hemorrhage	r	0.024		0.097*	0.118**	0.104*	0.450**
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		p	0.570	0.082	0.023	0.006	0.015	< 0.001
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Perineal laceration	r	0.069	0.039	-0.034	0.030	0.025	-0.058
NICU admission r -0.032 -0.020 0.011 -0.073 -0.036 0.153** p 0.453 0.651 0.800 0.089 0.402 <0.001 Apgar score (1 min) r 0.037 0.010 -0.010 0.001 0.049 -0.031 p 0.391 0.817 0.816 0.977 0.249 0.467 Apgar score (5 min) r 0.082 0.017 -0.008 0.012 -0.001 0.008 p 0.054 0.691 0.854 0.773 0.984 0.857 Apgar score (10 min) r 0.087* 0.022 -0.014 0.028 0.008 0.047 p 0.042 0.607 0.743 0.519 0.851 0.269 RDS r -0.263** -0.174** -0.030 -0.049 -0.018 -0.018 p 0.561 0.839 0.033 <0.001 0.003 0.938 PDA r 0.018 0.015 -0.047 -0.029 0.016 -0.004 p 0.679 0.718 0.277 0.503 0.704 0.923 Neonatal pneumonia r -0.013 0.026 0.022 0.042 0.050 0.001 p 0.763 0.549 0.608 0.321 0.239 0.988 Neonatal jaundice r -0.081 0.022 0.030 0.004 -0.016 0.024 p 0.057 0.603 0.484 0.925 0.706 0.569 AAFMS r 0.034 0.020 0.050 0.090* 0.061 0.005		p	0.106	0.365	0.431	0.491	0.558	0.174
NICU admission $\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Mortality	r	0.013	0.040	0.026	0.040	0.017	-0.035
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		p	0.762	0.356	0.544	0.346	0.690	0.420
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	NICU admission	r	-0.032	-0.020	0.011	-0.073	-0.036	0.153**
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		p	0.453	0.651	0.800	0.089	0.402	< 0.001
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Apgar score (1 min)	r	0.037	0.010	-0.010	0.001	0.049	-0.031
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		p	0.391	0.817	0.816	0.977	0.249	0.467
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Apgar score (5 min)	r	0.082	0.017	-0.008	0.012	-0.001	0.008
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		p	0.054	0.691	0.854	0.773	0.984	0.857
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Apgar score (10 min)	r	0.087*	0.022	-0.014	0.028	0.008	0.047
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$		p	0.042	0.607	0.743	0.519	0.851	0.269
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	RDS	r	-0.263**	-0.174**	-0.030	-0.049	-0.018	-0.018
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		p	< 0.001	< 0.001	0.486	0.250	0.675	0.668
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	IVH	r	0.025	-0.009	0.091*	0.198**	0.127**	-0.003
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		p	0.561	0.839	0.033	< 0.001	0.003	0.938
Neonatal pneumonia r -0.013 0.026 0.022 0.042 0.050 0.001 p 0.763 0.549 0.608 0.321 0.239 0.988 Neonatal jaundice r -0.081 0.022 0.030 0.004 -0.016 0.024 p 0.057 0.603 0.484 0.925 0.706 0.569 AAFMS r 0.034 0.020 0.050 $0.090*$ 0.061 0.005	PDA	r	0.018	0.015	-0.047	-0.029	0.016	-0.004
p 0.763 0.549 0.608 0.321 0.239 0.988 Neonatal jaundice r -0.081 0.022 0.030 0.004 -0.016 0.024 p 0.057 0.603 0.484 0.925 0.706 0.569 AAFMS r 0.034 0.020 0.050 0.090* 0.061 0.005		p	0.679	0.718	0.277	0.503	0.704	0.923
Neonatal jaundice r -0.081 0.022 0.030 0.004 -0.016 0.024 p 0.057 0.603 0.484 0.925 0.706 0.569 AAFMS r 0.034 0.020 0.050 0.090* 0.061 0.005	Neonatal pneumonia	r	-0.013	0.026	0.022	0.042	0.050	0.001
p 0.057 0.603 0.484 0.925 0.706 0.569 AAFMS r 0.034 0.020 0.050 0.090* 0.061 0.005		p	0.763	0.549	0.608	0.321	0.239	0.988
AAFMS r 0.034 0.020 0.050 0.090* 0.061 0.005	Neonatal jaundice	r	-0.081	0.022	0.030	0.004	-0.016	0.024
		p	0.057	0.603	0.484	0.925	0.706	0.569
p 0.421 0.637 0.242 0.035 0.153 0.907	AAFMS	r	0.034	0.020	0.050	0.090*	0.061	0.005
		p	0.421	0.637	0.242	0.035	0.153	0.907

**p < 0.01, *p < 0.05; COVID-19, coronavirus disease 2019; PPROM, preterm premature rupture of membranes; GDM, gestational diabetes mellitus; NICU, neonatal intensive care unit; RDS, respiratory distress syndrome; IVH, intraventricular haemorrhage; PDA, patent ductus arteriosus; AAFMS, aspiration of amniotic fluid and meconium syndrome.

p=0.008; r=0.204, p<0.001). Significant relationships exist between the pain symptom and the higher rate of abnormal amniotic fluid (r=-0.101, p=0.018). In addition, the symptom of pain, sore throat, cough, and fatigue were significantly related to a higher rate of postpartum hemorrhage (r=0.097, p=0.023; r=0.118, p=0.006; r=0.104, p=0.015 and r=0.450, p<0.001).

According to Table 4, the mothers with the sign of fatigue had a higher rate of neonatal intensive care unit

(NICU) admission (r = 0.153, p < 0.001). It suggested that positive symptoms and pain indicated a higher rate of RDS (r=-0.263, p < 0.001; r=-0.174, p < 0.001). The mothers with the sign of pain, sore throat, and cough had a higher rate of IVH (r = 0.091, p = 0.033; r = 0.198, p < 0.001; r = 0.127, p = 0.003). Significant relationships exist between the sore throat symptom and the higher rate of AAFMS (r = 0.090, p = 0.035).



4. Discussion

Pregnancy is considered an independent risk factor for severe COVID-19 due to immunological and physical changes during pregnancy [9]. The progression of COVID-19 mainly depends on the viral entry into the host cells after binding to the angiotensin-converting enzyme 2 (ACE2). This factor may make pregnant women more susceptible to COVID-19 because ACE 2 is highly expressed in the placenta throughout the pregnancy [23]. During the COVID-19 pandemic, concerns arose about whether infection by the SARS-CoV-2 virus adversely affects pregnancy outcomes. Under the prevention and control protection in China Mainland, the COVID-19 infection rate among pregnant women is shallow. Therefore, few studies about COVID-19 infection in pregnancy and pregnancy outcomes exist. Unfortunately, at the end of 2022, with the national development and adjustment of the epidemic prevention and control policy, the COVID-19 infection rate of the population increased, and the number of pregnant women also increased significantly. Therefore, it is urgent to study the symptoms of pregnant women after COVID-19 infection and the impact on pregnancy outcomes in China.

In the present study, 549 (15.2%) were diagnosed with COVID-19 infection. Among them, 489 (89.1%) fever is the most common symptom, followed by 352 (64.1%) cough, 218 (39.7%) sore throat, 107 (19.5%) pain, 46 (8.4%) fatigue and 19 (3.5%) other symptoms. Similarly, studies have indicated that the most common clinical signs of COVID-19 are fever, fatigue, cough, expectoration, anorexia, sputum production, and shortness of breath [24,25]. Thus, the symptoms of pregnant women with COVID-19 infection are similar to those of non-pregnant women.

We found that mothers with COVID-19 infection had slightly higher cesarean section rates than non-infected mothers, but the difference was not statistically significant. The current findings are different. The research of Jafari et al. [26] shows that the rate of cesarean delivery in pregnant women infected with COVID-19 infection varied from 42.9%–85%, significantly higher than that of without COVID-19 disease pregnant women. And, a survey of obstetricians by the Japanese Obstetrics Association showed that obstetricians are more inclined to use cesarean section for pregnant women with COVID-19 [27]. However, Eleje et al. [28] found that the cesarean section rate during the COVID-19 period was significantly less than the period prior to the pandemic. Moreover, we revealed a higher rate of postpartum hemorrhage with COVID-19 infection. The study by Auger et al. [29] also shows an increased risk of COVID-19 infection and postpartum hemorrhage. In addition, we found pregnancies with COVID-19 infection were significantly associated with higher PPROM. It is similar to previous studies. Zhu et al. [30] reported a study with 9 cases of pregnant women with COVID-19 infection and 10 newborns delivered, among which 3 cases had PPROM.

Also, Du *et al.* [31] reported an 11% increased risk of premature rupture of membranes and fetal distress during the covid-19 pandemic (95% confidence interval (95% CI), 1.04 to 1.18; p < 0.01) and 14% (95% CI, 1.01 to 1.29; p < 0.05).

We demonstrated that some neonatal outcomes were significantly associated with COVID-19 infection. Mothers with COVID-19 infection were significantly associated with higher rates of LBW and macrosomia in newborns, higher rate of RDS, higher rate of IVH, higher rate of neonatal pneumonia, and a higher rate of AAFMS. Similarly, there is a rise in preterm and low birth weight cases due to COVID-19 infection [32]. Symptomatic illness in pregnancy is related to adverse maternal and neonatal outcomes such as higher cesarean delivery rates, preterm delivery, and low birth weight. In addition, 4 (0.7%) newborns are positive at the 12–24 h. However, the four newborns could not confirm whether it was caused by vertical transmission. There is still debate regarding the rate of vertical transmission of COVID-19 infection to newborns and its possible mechanisms-transplacental or during passage through the birth canal. In a study conducted by Edlow et al. [33] in 2020, no evidence of placental infection or standard vertical transmission of COVID-19 was found in the 64 patients who tested positive for COVID-19. Also, a retrospective cohort [34] analysis reported data for 101 newborns from mothers that tested positive for SARS-CoV-2. There was no evidence of vertical transmission during the first 25 days of life, despite the newborns rooming in with mothers and direct breastfeeding practices.

This is a large sample of data in South China, which is representative. However, there are some limitations to this study. Firstly, this study did not include data on vaccination, so it is impossible to assess the impact of vaccination on pregnant women. Secondly, though the case number of the present study was over the previous studies, it is a retrospective single-center study, rather than a prospective study. Multi-center or randomized controlled trial is necessary for the future.

5. Conclusions

In conclusion, our data prove that specific symptoms of pregnant women were similar to non-pregnant, including fever, cough, sore throat, pain, and fatigue. In addition, some maternal complications and neonatal outcomes were significantly associated with COVID-19 infection. It suggests that pregnancy with COVID-19 disease in China has some adverse consequences for pregnant women and newborns. Therefore, infected pregnant women need to be alert to these pregnancy complications, such as PPROM and postpartum hemorrhage. Clinicians should strengthen clinical monitoring to provide better care for mothers and children, and provide infected pregnant women with appropriate counseling and advice on PPROM and postpartum hemorrhage. The COVID-19 epidemic continues, requir-



ing targeted public health measures to reduce the infection rate in pregnant women and the poor prognosis of mothers and children.

Availability of Data and Materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Author Contributions

YL, ZC, LX and XX designed the research study. ZW and YX collected the data and performed the research. YL analyzed the data. All authors contributed to editorial changes in the manuscript. All authors read and approved the final manuscript. All authors have participated sufficiently in the work and agreed to be accountable for all aspects of the work.

Ethics Approval and Consent to Participate

The study was conducted in accordance with the Declaration of Helsinki, and the protocol was approved by the Ethics Committee Fujian Provincial Maternity and Children's Hospital, an affiliated hospital of Fujian Medical University (approval number: 2023KY005). This study is a retrospective study, the data are anonymous, and the requirement for informed consent was therefore waived.

Acknowledgment

We would like to thank all the women who kindly agreed to participate in this study.

Funding

Joint Funds for the innovation of science and Technology, Fujian province (Grant number 2020Y9147). Joint Funds for the innovation of science and Technology, Fujian province (2020Y9165).

Conflict of Interest

The authors declare no conflict of interest.

References

- [1] Zhu N, Zhang D, Wang W, Li X, Yang B, Song J, *et al.* A Novel Coronavirus from Patients with Pneumonia in China, 2019. The New England Journal of Medicine. 2020; 382: 727–733.
- [2] WHO Coronavirus (COVID-19) Dashboard | WHO Coronavirus (COVID-19) Dash board With Vaccination Data. 2022. Available at: https://covid19.who.int/ (Accessed: 22 May 2022).
- [3] Li MY, Li L, Zhang Y, Wang XS. Expression of the SARS-CoV-2 cell receptor gene ACE2 in a wide variety of human tissues. Infectious Diseases of Poverty. 2020; 9: 45.
- [4] Wang N, Qin L, Ma L, Yan H. Effect of severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2) on reproductive system. Stem Cell Research. 2021; 52: 102189.
- [5] Chowdhury MA, Hossain N, Kashem MA, Shahid MA, Alam A. Immune response in COVID-19: A review. Journal of Infection and Public Health. 2020; 13: 1619–1629.

- [6] Hassanein M, Radhakrishnan Y, Sedor J, Vachharajani T, Vachharajani VT, Augustine J, et al. COVID-19 and the kidney. Cleveland Clinic Journal of Medicine. 2020; 87: 619–631.
- [7] Bonafè M, Prattichizzo F, Giuliani A, Storci G, Sabbatinelli J, Olivieri F. Inflamm-aging: Why older men are the most susceptible to SARS-CoV-2 complicated outcomes. Cytokine & Growth Factor Reviews. 2020; 53: 33–37.
- [8] Madjid M, Safavi-Naeini P, Solomon SD, Vardeny O. Potential Effects of Coronaviruses on the Cardiovascular System: A Review. JAMA Cardiology. 2020; 5: 831–840.
- [9] Majumder J, Minko T. Recent Developments on Therapeutic and Diagnostic Approaches for COVID-19. The AAPS Journal. 2021; 23: 14.
- [10] Elshafeey F, Magdi R, Hindi N, Elshebiny M, Farrag N, Mahdy S, et al. A systematic scoping review of COVID-19 during pregnancy and childbirth. International Journal of Gynaecology and Obstetrics: the Official Organ of the International Federation of Gynaecology and Obstetrics. 2020; 150: 47–52.
- [11] Fan C, Lei D, Fang C, Li C, Wang M, Liu Y, et al. Perinatal Transmission of 2019 Coronavirus Disease-Associated Severe Acute Respiratory Syndrome Coronavirus 2: Should We Worry? Clinical Infectious Diseases: an Official Publication of the Infectious Diseases Society of America. 2021; 72: 862–864.
- [12] Yan J, Guo J, Fan C, Juan J, Yu X, Li J, *et al.* Coronavirus disease 2019 in pregnant women: a report based on 116 cases. American Journal of Obstetrics and Gynecology. 2020; 223: 111.e1–111.e14.
- [13] Verma S, Carter EB, Mysorekar IU. SARS-CoV2 and pregnancy: An invisible enemy? American Journal of Reproductive Immunology (New York, N.Y.: 1989). 2020; 84: e13308.
- [14] Cosma S, Borella F, Carosso A, Sciarrone A, Cusato J, Corcione S, et al. The "scar" of a pandemic: Cumulative incidence of COVID-19 during the first trimester of pregnancy. Journal of Medical Virology. 2021; 93: 537–540.
- [15] Patberg ET, Adams T, Rekawek P, Vahanian SA, Akerman M, Hernandez A, et al. Coronavirus disease 2019 infection and placental histopathology in women delivering at term. American Journal of Obstetrics and Gynecology. 2021; 224: 382.e1– 382.e18.
- [16] Villar J, Ariff S, Gunier RB, Thiruvengadam R, Rauch S, Kholin A, et al. Maternal and Neonatal Morbidity and Mortality Among Pregnant Women With and Without COVID-19 Infection: The INTERCOVID Multinational Cohort Study. JAMA Pediatrics. 2021; 175: 817–826.
- [17] Papageorghiou AT, Deruelle P, Gunier RB, Rauch S, García-May PK, Mhatre M, et al. Preeclampsia and COVID-19: results from the INTERCOVID prospective longitudinal study. American Journal of Obstetrics and Gynecology. 2021; 225: 289.e1–289.e17
- [18] Villar J, Soto Conti CP, Gunier RB, Ariff S, Craik R, Cavoretto PI, et al. Pregnancy outcomes and vaccine effectiveness during the period of omicron as the variant of concern, INTERCOVID-2022: a multinational, observational study. Lancet (London, England). 2023; 401: 447–457.
- [19] Eskenazi B, Rauch S, Iurlaro E, Gunier RB, Rego A, Gravett MG, *et al.* Diabetes mellitus, maternal adiposity, and insulin-dependent gestational diabetes are associated with COVID-19 in pregnancy: the INTERCOVID study. American Journal of Obstetrics and Gynecology. 2022; 227: 74.e1–74.e16.
- [20] Chen A, Acharya G, Hu M, Gao X, Cheng G, Jiang L, et al. Association of maternal SARS-CoV-2 infection at the time of admission for delivery with labor process and outcomes of vaginal birth: A cohort study. Acta Obstetricia et Gynecologica Scandinavica. 2024; 103: 103–110.
- [21] Lv A, BianBaZhuoMa, DeQiong, DaWaZhuoMa, PuBuZhuoMa, Yao D, et al. Effect of COVID-19 infection on



- pregnant women in plateau regions. Public Health. 2024; 229: 57-62.
- [22] Jenabi E, Bashirian S, Khazaei S, Masoumi SZ, Ghelichkhani S, Goodarzi F, et al. Pregnancy outcomes among symptomatic and asymptomatic women infected with COVID-19 in the west of Iran: a case-control study. The Journal of Maternal-fetal & Neonatal Medicine: the Official Journal of the European Association of Perinatal Medicine, the Federation of Asia and Oceania Perinatal Societies, the International Society of Perinatal Obstetricians. 2022; 35: 4695–4697.
- [23] Azinheira Nobrega Cruz N, Stoll D, Casarini DE, Bertagnolli M. Role of ACE2 in pregnancy and potential implications for COVID-19 susceptibility. Clinical Science (London, England: 1979). 2021; 135: 1805–1824.
- [24] Xu XW, Wu XX, Jiang XG, Xu KJ, Ying LJ, Ma CL, et al. Clinical findings in a group of patients infected with the 2019 novel coronavirus (SARS-Cov-2) outside of Wuhan, China: retrospective case series. BMJ (Clinical Research Ed.). 2020; 368: m606.
- [25] Chen N, Zhou M, Dong X, Qu J, Gong F, Han Y, *et al.* Epidemiological and clinical characteristics of 99 cases of 2019 novel coronavirus pneumonia in Wuhan, China: a descriptive study. Lancet (London, England). 2020; 395: 507–513.
- [26] Jafari M, Pormohammad A, Sheikh Neshin SA, Ghorbani S, Bose D, Alimohammadi S, et al. Clinical characteristics and outcomes of pregnant women with COVID-19 and comparison with control patients: A systematic review and meta-analysis. Reviews in Medical Virology. 2021; 31: 1–16.
- [27] Komatsu H, Banno K, Yanaihara N, Kimura T, Board Members of Japan Society of Obstetrics and Gynecology. Prevention and practice during the COVID-19 emergency declaration period in

- Japanese obstetrical/gynecological facilities. The Journal of Obstetrics and Gynaecology Research. 2020; 46: 2237–2241.
- [28] Eleje GU, Ugwu EO, Enebe JT, Okoro CC, Okpala BC, Ezeora NC, et al. Cesarean section rate and outcomes during and before the first wave of COVID-19 pandemic. SAGE Open Medicine. 2022; 10: 20503121221085453.
- [29] Auger N, Ukah UV, Wei SQ, Healy-Profitós J, Lo E, Dayan N. Impact of Covid-19 on risk of severe maternal morbidity. Critical Care (London, England). 2023; 27: 344.
- [30] Zhu H, Wang L, Fang C, Peng S, Zhang L, Chang G, et al. Clinical analysis of 10 neonates born to mothers with 2019-nCoV pneumonia. Translational Pediatrics. 2020; 9: 51–60.
- [31] Du M, Yang J, Han N, Liu M, Liu J. Association between the COVID-19 pandemic and the risk for adverse pregnancy outcomes: a cohort study. BMJ Open. 2021; 11: e047900.
- [32] Maeda Y, Nakamura M, Ninomiya H, Ogawa K, Sago H, Miyawaki A. Trends in intensive neonatal care during the COVID-19 outbreak in Japan. Archives of Disease in Childhood. Fetal and Neonatal Edition. 2021; 106: 327–329.
- [33] Edlow AG, Li JZ, Collier ARY, Atyeo C, James KE, Boatin AA, et al. Assessment of Maternal and Neonatal SARS-CoV-2 Viral Load, Transplacental Antibody Transfer, and Placental Pathology in Pregnancies During the COVID-19 Pandemic. JAMA Network Open. 2020; 3: e2030455.
- [34] Dumitriu D, Emeruwa UN, Hanft E, Liao GV, Ludwig E, Walzer L, et al. Outcomes of Neonates Born to Mothers with Severe Acute Respiratory Syndrome Coronavirus 2 Infection at a Large Medical Center in New York City. JAMA Pediatrics. 2021; 175: 157–167.

