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## Maternal and perinatal outcomes in the COVID-19 Omicron wave in comparison with the Delta wave: a multicentre observational study

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### ABSTRACT

**Objective.** The aim of the study was to compare maternal and neonatal outcomes of the COVID-19 in Omicron and Delta wave.

**Materials and Methods.** In this prospective observational multicentre study, we included unvaccinated patients who were hospitalized to deliver and who gave birth while infected. Patients who gave birth from May 15, 2021, to November 15, 2021 were allocated to the Delta group, and those who gave birth from November 15, 2021, to June 01, 2022 were included in the Omicron group.

The maternal and foetal outcomes were compared between the two groups. The significance level was set at  $p < 0.05$ .

**Results.** We included 84 patients in the Delta group and 45 patients in the Omicron group. We noted more asymptomatic COVID-19 ( $p = 0.001$ ), less dyspnoea ( $p = 0.021$ ), and less need for oxygen ( $p = 0.008$ ) in the Omicron group. Caesarean section delivery was seen in 62 patients from the Delta group, and vaginal delivery was seen in 20 patients from the Omicron group with  $p = 0.029$ . Even if the incidence of complications and clinical deterioration after delivery were comparable in both groups, the length of hospital stay and the number of deaths were significantly higher in the Delta group with  $p = 0.038$  and  $p = 0.024$ , respectively. The foetal outcomes were comparable.

**Conclusions.** SARS-CoV-2 infection during the Omicron wave had a lower requirement for oxygen support and improved maternal outcomes in comparison with the Delta wave.

## INTRODUCTION

Pregnant women are vulnerable to viral infections such as COVID-19 infection [1]. Heavy morbidity and high rates of mortality [2] may be due to immunological and physiological changes during pregnancy [3], particularly when infection occurs in the peripartum period with unvaccinated patients [4]. Since the beginning of the pandemic, several waves with different SARS-CoV-2 variants have created a tsunami of COVID-19 worldwide [5]. The omicron variant of SARS-CoV-2 (B.1.1.529) spread rapidly across the world, out-competing the previous variants such as the Delta variant (B.1.617.2) [6]. Omicron, first detected in November 2021, appears to cause less severe acute illness than previous variants, at least in vaccinated populations [6]. The role of vaccination in reducing the COVID-19 severity should be considered. To date, scientists think that this omicron wave indicates the end of the pandemic even if COVID-19 persists [7]. However, we must exercise caution regarding the impact of this new variant in specific and high-risk patients, such as pregnant women. The impact of this new variant on maternal and foetal outcomes is not yet well known.

The aim of the study was to compare maternal and perinatal outcomes of the Omicron COVID-19 wave with the Delta wave among unvaccinated pregnant women, infected at delivery.

## MATERIALS AND METHODS

After obtaining patients' oral consent and local ethics committee approval, a multicentre observational cohort study was conducted to compare the maternal and neonatal outcomes of the Omicron COVID-19 wave with the Delta wave among unvaccinated pregnant women. This study was conducted in four (level 2 or level 3) maternity hospitals from 3 regions in the south of Tunisia (Sfax, Medenine, and Tataouine) from May 15, 2021, to June 01, 2022.

We included only unvaccinated patients who were hospitalized for delivery and gave birth while infected with COVID-19. The patients included had a positive reverse transcriptase-polymerase chain reaction (RT-PCR nasopharyngeal swab) test result for severe acute respiratory syndrome Coronavirus 2 within the 5 days preceding delivery. We excluded cases of foetal loss defined as spontaneous

antepartum foetal death <14 weeks of gestation (WG) and cases of late miscarriage 14-24 WG. Vaccinated patients and those whose management did not adhere to the standard protocol were also excluded. We also excluded patients with recurrent SARS-CoV-2 infections.

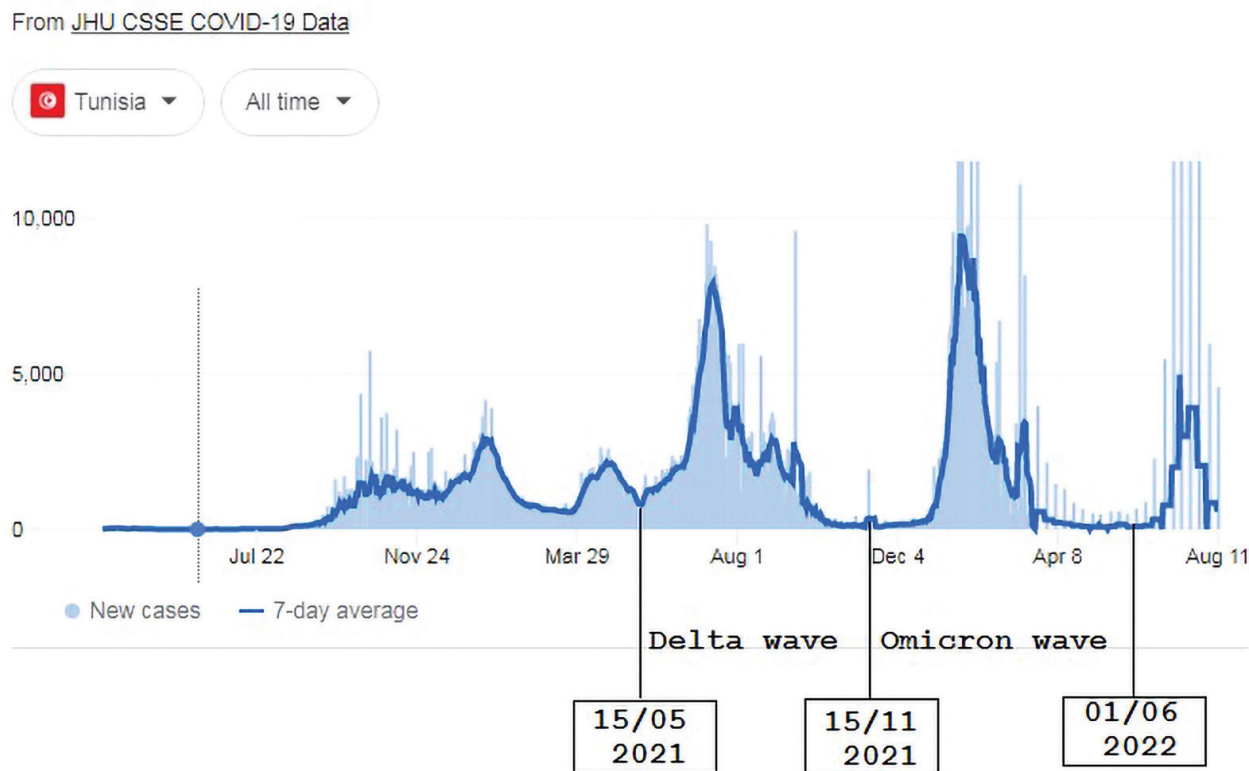
The variables were:

- Demographic parameters: age, body mass index (BMI), term of pregnancy, parity, and previous co-morbidities.
- Clinical and biological features of the COVID-19 infection before delivery and the need for oxygen supplementation define the severity of the infection.
- Maternal adverse outcomes after delivery include increased need for oxygen supplementation after delivery, referral to the intensive care unit (ICU), maternal complications, and maternal deaths.
- Clinical deterioration was defined by an increased need for O<sub>2</sub> supplementation after delivery, or referral to ICU, and maternal complications (acute respiratory distress syndrome: ARDS, postpartum haemorrhage, thromboembolic events, septic shock, and pregnancy-related complications including retro-placental haematoma; haemolysis, elevated liver enzymes, and low platelets syndrome: HELLP syndrome, and acute fatty liver of pregnancy). For severe COVID-19 requiring advanced oxygen support or intensive care before delivery, clinical deterioration is defined by the incidence of a severe complication or death. We looked at the length of hospital stay and the incidence of maternal deaths.

Neonatal outcomes considered were neonatal ICU (NICU) admission, prematurity, vertical transmission, and stillbirth or neonatal deaths.

All patients enrolled in this study were not vaccinated and had the same management protocol. All maternity hospitals participating in this study adhere to the INAES (Instance Nationale d'Evaluation et d'Accréditation en Santé) guidelines [8]. To compare maternal and perinatal outcomes between Omicron and Delta waves among pregnant women who tested positive at the moment of birth, patients were divided into 2 groups with reference to the epidemiologic situation and COVID-19 waves that occurred in Tunisia (**Figure 1**):

- Delta Group: patients who gave birth during the delta wave, lasting from May 15, 2021, to November 15, 2021.



**Figure 1.** COVID-19 waves and groups definition.

- Omicron Group: patients who gave birth during the Omicron waves, lasting from November 15, 2021, to the June 01, 2022.

The COVID-19 waves were called Delta and Omicron because, in these periods, the predominant variants circulating in our country were respectively the Delta variant (B.1.617.2) and the Omicron variant (B.1.1.529) [9].

Then, statistical analyses were conducted using the SPSS 23.0 (SPSS, Chicago, IL, USA) statistical package. We distinguished two groups according to the COVID-19 wave among positive pregnant women. The comparison between groups was achieved by Student's t-test and Chi<sup>2</sup> test for continuous variables and categorical variables, respectively. The Fisher exact test was used when the Chi<sup>2</sup> test was not applicable. The Mann-Whitney U test was used for non-parametric continuous variables. The significance level was set at  $p < 0.05$ .

## RESULTS

Of the 189 patients participating in the study, 129 deliveries were included: 84 deliveries in the Delta wave and 45 deliveries in the Omicron wave. Sixty patients were excluded: three for foetal loss (< 14

WG), four because of late miscarriage (14-24 WG), and 53 because of completed or uncompleted vaccination. No patient was excluded because of recurrent SARS-CoV-2 infection. All foetal losses and late miscarriages occurred in the Delta wave, and all vaccinated patients were admitted in the Omicron wave. The 129 patients included were recruited mainly from 3 regions in the south of Tunisia: Sfax ( $n = 102$ ), Medenine ( $n = 19$ ), and Tataouine ( $n = 8$ ).

Demographic parameters (age, BMI, comorbidities, parity, and term of pregnancy) were comparable in both groups (Table 1). Asymptomatic COVID-19 was seen in 22% and 3.5% of patients in the Omicron wave and Delta wave, with  $p = 0.001$ . In the Omicron group, we noted lower rates of cough ( $p = 0.032$ ), fever ( $p = 0.001$ ), and dyspnoea ( $p = 0.021$ ). The other clinical features (asthenia, digestive troubles, otorhinolaryngological symptoms, preeclampsia, anaemia, and cytolysis) were comparable in both groups (Table 2).

The Delta group had a higher rate of patients requiring oxygen supplementation before delivery (47 patients: 55.9%) than the Omicron group (15 patients: 33.3%), with  $p = 0.008$ . However, the oxygen flow administered (in patients who needed oxygen supplementation) and the need for ICU

**Table 1.** Demographic parameters.

	Delta Group n = 84	Omicron Group n = 45	P-value
Age	30.5 ± 4.7	31.4 ± 5.7	0.309
Age > 35 years	21 (25%)	15 (33.3%)	0.211
BMI	28.6 ± 3.6	28.6 ± 3.1	0.966
BMI > 30 kg/m <sup>2</sup>	31 (36.9%)	18 (40%)	0.437
With comorbidities	10 (11.9%)	2 (4.4%)	0.086
Term of pregnancy (WG)	35.6 ± 3.8	36.6 ± 3.0	0.130
Multiparity (> 2)	59 (70.2%)	39 (86.6%)	0.051

**Table 2.** COVID-19 severity before delivery.

	Delta Group n = 84	Omicron Group n = 45	P-value
Asymptomatic	3 (3.5%)	10 (22.2%)	<b>0.001</b>
Cough	64 (76.1%)	27 (60%)	<b>0.032</b>
Fever	58 (69%)	17 (37.7%)	<b>0.001</b>
Headache and asthenia	51 (60.7%)	25 (55.5%)	0.314
Dyspnoea	31 (36.9%)	7 (15.5%)	<b>0.021</b>
Digestive signs	18 (21.4%)	6 (13.3%)	0.208
Sore throat, rhinorrhoea, anosmia and ageusia)	9 (10.7%)	2 (4.4%)	0.194
Preeclampsia	18 (21.4%)	12 (26.6%)	0.375
Anaemia	11 (13%)	11 (24.4%)	0.133
Cytolysis	11 (13%)	7 (15.5%)	0.419
Thrombopenia (< 50,000)	0 (0%)	1 (2.2%)	0.176
CT-Scan > 50% (yes/no)	7/5	3/0	0.266
Need for O <sub>2</sub> before delivery	47 (55.9%)	15 (33.3%)	<b>0.008</b>
O <sub>2</sub> < 6 L/min	37 (82.2%)	13 (86.6%)	
O <sub>2</sub> = 6-15 L/min	7 (14.9%)	0	0.233
O <sub>2</sub> > 15 L/min*	3 (6.4%)	2 (13.4%)	
ICU before delivery	2 (2.3%)	3 (6.6%)	0.230

\*Optiflow or CPAP.

admission were comparable in both groups (Table 2). The incidence of vaginal delivery was higher in the Omicron group (55.5%) in comparison with the Delta group (26.1%) with  $p = 0.029$ . Caesarean section was performed mainly for foetal distress in the Delta group (51.6%) and for obstetrical reasons in the Omicron group (59%),  $p = 0.001$  (Table 3). After delivery, the incidence of clinical deterioration, an increased need for oxygen, maternal complications, and referral to the ICU were comparable in both groups. Furthermore, the Delta group had a significantly longer hospital stay and a higher number of deaths (Table 3). Foetal and neonatal outcomes regarding prematurity, vertical transmission, neonatal critical care unit admission, and stillbirth or neonatal deaths were comparable in both groups (Table 4).

## DISCUSSION

This study showed that the COVID-19 severity in unvaccinated pregnant women was reduced during the Omicron wave. We noted more asymptomatic and minor forms, which influenced the mode of delivery and the indications of caesarean deliveries and reduced postpartum morbidity and mortality.

The main strength of this study is that it provides data related to maternal and perinatal outcomes associated with COVID-19 during the Delta and the Omicron waves among unvaccinated women from the south of Tunisia. To date, there is little data on the severity of infection by the emergent Omicron variant in unvaccinated pregnant women compared with previous variants. Our study showed that even if the Omicron variant is less

**Table 3.** Maternal outcomes after delivery.

	Delta Group n = 84	Omicron Group n = 45	P-value
Mode of delivery (caesarean/vaginal)	62/22	25/20	<b>0.029</b>
Indications of caesarean delivery			
Foetal distress	32 (51.6%)	2 (8%)	
Obstetrical indications	18 (29%)	13 (59%)	
Severe preeclampsia	2 (3.2%)	1 (4%)	<b>0.001</b>
Maternal life-saving	9 (14.5%)	5 (11.1%)	
Other circumstances	1 (1.6%)	5 (11.1%)	
Clinical deterioration (%)	37 (44%)	14 (31.1%)	0.106
Increased need for O <sub>2</sub> after delivery	32 (38%)	13 (28.8%)	0.198
Postpartum referral to ICU	24 (28.5%)	9 (20%)	0.198
Complications (yes/no)	34 (40.4%)	15 (33.3%)	0.168
ARDS	16	8	
Postpartum haemorrhage	4	5	
Thromboembolic events	1	2	0.065
Septic shock	4	0	
Pregnancy related complication	9	0	
Length of hospital stay (days)	8.6 ± 6.5	6.3 ± 4.4	<b>0.038</b>
Maternal deaths	12 (14.2%)	1 (2.2%)	<b>0.024</b>

**Table 4.** Neonatal outcomes.

	Delta Group n = 84	Omicron Group n = 45	P-value
Severe prematurity: delivery < 28 WG	4 (4.7%)	0	0.175
Premature delivery: 28-34 WG	26 (30.9%)	10 (22.2%)	0.199
Prematurity < 34 WG	30 (35.7%)	10 (22.2%)	0.084
Vertical transmission	3 (3.5%)	1 (2.2%)	0.565
Admission in neonatal ICU	24 (28.5%)	10 (22.2%)	0.287
Neonatal deaths and stillbirth	7 (8.3%)	2 (4.4%)	0.333

dangerous than the Delta, unvaccinated parturients may need oxygen support and sometimes ICU admissions. This emphasizes the importance of vaccination [4].

The main limitation of the study was that variant sequencing data was not available for all patients. So, the time of infection was used to indicate the predominant variant. The second limit is that the patients were not recruited simultaneously (the Omicron wave came after), which can affect the management of the patients because health workers (anaesthesiologists, obstetricians, midwives, and nurses) have gained more experience from previous waves [10].

The Omicron variant, first identified in Botswana in November 2021, is rapidly becoming the dominant circulating variant. It has a significant growth in contagion over the Delta, leading to rapid spread with higher incidence levels [11]. In our study, the number of patients admitted

during the omicron wave was lower than that of patients admitted during the delta wave. This may be explained by the exclusion of vaccinated patients. We excluded vaccinated pregnant women because previous studies showed the effectiveness of the BNT162b2 vaccine against severe COVID-19 requiring hospital admission [12]. COVID-19 has been mild in comparison to the Delta since the emergence and spread of the omicron variant [11]. Several studies in the general population [13, 14], and in pregnant women [15, 16] evoked lower severity with a significant decrease in deaths [17]. These findings were comparable with our results, as we noted more asymptomatic patients and less need for oxygen supplementation. However, we should mention that the omicron variant appeared after the globalization of the vaccine, and even if the majority of studies comparing delta with omicron selected unvaccinated patients, the indirect effect

of vaccination in the general population should be discussed. It was reported that mRNA-based COVID-19 vaccines are associated with a reduction in SARS-CoV-2 infections requiring hospital admission, not only among vaccinated individuals but also among unvaccinated adults [18]. In our population, pregnant women were hesitant about vaccination. However, the general population has been widely and completely vaccinated since September 2021 [19]. The COVID-19 severity can impact the mode of delivery and the maternal and foetal outcomes as a consequence. Maternal hypoxemia due to COVID-19 pneumonia, ARDS, and villous infarction (thrombotic lesions in the placenta) [20], often seen in the Delta wave, can lead to emergent caesarean section delivery for foetal distress or maternal life-saving [21]. However, vaginal delivery can be accepted only in patients who have no need or low flow of oxygen [22].

This may be the explanation for our obstetrical results (mode of delivery and indications for caesarean delivery) [23].

Even if enhanced recovery strategies after caesarean section reduce the risks of postpartum complications and improve maternal outcomes [24], the physiological stress induced by surgery is known to increase the rate of complications in infected pregnant women. This might explain the higher length of hospital stay after delivery in the delta group, in which caesarean delivery was the main mode of delivery.

In our study, foetal outcomes were comparable in both groups. However, prematurity (< 34 WG) was higher in the delta wave. These results are compatible with those of previous studies [16] which reported that preterm birth was significantly increased during the Delta wave (15.4%) compared with the pre-Delta period (4.9%) and omicron wave (2.8%). The incidence of prematurity in the delta wave in our region seems to be comparable with previous waves [25]. Nevertheless, maternal oxygen support, which seems to be increased with the delta variant, was reported as a risk factor for adverse foetal outcomes [26]. This may argue in favor of a higher incidence of adverse perinatal outcomes in foetuses with maternal COVID-19 infection in the Delta wave [27]. It was also reported that early gestational age at infection is one of the main determinants of adverse foetal outcomes [26]. However, in our study, we included only infected parturients at delivery. In

the literature, some habits known to be involved in adverse foetal outcomes were not investigated in our study, like coffee drinking [28], alcohol intake [29], and sleep deprivation [30]. It was also reported that maternal vascular malperfusion, including decidual arteriopathy, was significantly more frequent after SARS-CoV-2 infection, with the highest rates in the Delta era [27]. This may explain the poor foetal outcomes that could be associated with the Delta variant.

Recent and novel evidence suggests that the cytokine storm underlies the activation of the coagulation system which may lead to villous infarction and decidual arteriopathy with thrombosis [31]. MicroRNAs (miRNAs), small non-coding RNAs, serve as gene expression regulators and are involved in balancing the pro-/anticoagulant and pro-/anti-inflammatory factors maintaining homeostasis [32]. The miRNAs can also reduce the viral load by degradation of viral RNA and reducing the expression of ACE2 receptors, besides mitigating the deleterious consequences of the exaggerated secretion of cytokines [33]. The miRNAs are expressed in the placenta and are involved in regulating trophoblast differentiation, migration, invasion, proliferation, apoptosis, vasculogenesis, angiogenesis, and cellular metabolism [34]. As a result, changes in the expression of selective miRNAs in COVID-19 may play an important role in both placental-induced diseases, such as intrauterine growth restriction [35]. However, the impact of SARS CoV-2 variants (Delta variant and Omicron variant) on placental miRNA dysregulation is not yet well known and further studies are needed.

## CONCLUSIONS

SARS-CoV-2 infection during the Delta wave was associated with clinical signs of severity, increased need for oxygen support, and higher maternal mortality compared with infection during the Omicron wave. Obstetrical outcomes may be influenced by COVID-19 severity. This may explain the improved maternal and perinatal outcomes during the Omicron wave. However, underestimating the risks of the Omicron variant is inexcusable and may adversely affect the vaccination rate among pregnant women, who are at an increased risk of adverse maternal and perinatal events related to COVID-19 severity.

## COMPLIANCE WITH ETHICAL STANDARDS

### *Authors contribution*

A.J.: Conceptualization, investigation, methodology, writing – review & editing. M.K.: Writing – original draft. O.B.: Data curation, formal analysis, investigation. Y.E.: Article revision. O.D.: Investigation. K.C., K.K.: Supervision, validation, visualization.

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### *Study registration*

None.

### *Disclosure of interests*

The authors declare that they have no conflict of interests.

### *Ethical approval*

It was obtained from the HCUH (local Ethics Committee).

### *Informed consent*

It was obtained from all patients included in the study.

### *Data sharing*

Data are available under reasonable request to the corresponding author.

## REFERENCES

1. Anand VV, Yogaraj GA, Priya S, Raj PP, Priyadharshini CB, Sridevi PN. A cross-sectional study on COVID19 mortality among people below 30 years of age in Tamilnadu-2020. *Clin Epidemiol Glob Health*. 2021;12:100827. doi: 10.1016/j.cegh.2021.100827
2. Keita H, James A, Bouvet L, Herrmann E, Le Gouez A, Mazoit JX, et al. Clinical, obstetrical and anaesthesia outcomes in pregnant women during the first COVID-19 surge in France: a prospective multicentre observational cohort study. *Anaesth Crit Care Pain Med*. 2021;40(5):100937. doi: 10.1016/j.accpm.2021.100937.
3. Benhamou D, Keita H, Ducloy-Bouthors AS, Bonnet MP, Bonnin M, Bouthors AS, et al. Coagulation changes and thromboembolic risk in COVID-19 obstetric patients. *Anaesth Crit Care Pain Med*. 2020;39(3):351-3. doi: 10.1016/j.accpm.2020.05.003.
4. Jarraya A, Kammoun M, Amouri S, Elleuch S, Khanfir F, Chaabene K, et al. Impact of COVID-19 vaccination among pregnant women requiring hospital admission: prospective observational research. *Ital J Gynaecol Obstet*. 2023;35(2):211-8. doi: 10.36129/jog.2022.53.
5. Mohapatra RK, Tiwari R, Sarangi AK, Sharma SK, Khandia R, Saikumar G, et al. Twin combination of omicron and delta variant triggering a Tsunami wave of ever high surges in COVID-19 cases: a challenging global threat with a special focus on Indian sub-continent. *J Med Virol*. 2022;94(5):1761-5. doi:10.1002/jmv.27585.
6. Antonelli M, Pujol JC, Spector TD, Ourselin S, Steves CJ. Risk of long COVID associated with delta versus omicron variants of SARS-CoV-2. *Lancet*. 2022;399(10343):2263-4. doi:10.1016/S0140-6736(22)00941-2.
7. Daria S, Islam MR. The SARS-CoV-2 omicron wave is indicating the end of the pandemic phase but the COVID-19 will continue. *J Med Virol*. 2022;94(6):2343-5. doi: 10.1002/jmv.27635.
8. Instance Nationale de l'Evaluation et de l'Accréditation en Santé. *Les Guides de l'INEAS*. 2021. Tunis, Tunisia.
9. Ayadi W, Taktak A, Gargouri S, Smaoui F, Chtourou A, Skouri-Gargouri H, et al. Development of a simple genotyping method based on indel mutations to rapidly screen SARS-CoV-2 circulating variants: Delta, Omicron BA.1 and BA.2. *J Virol Methods*. 2022;307:114570. doi: 10.1016/j.jviromet.2022.114570.
10. Del Piccolo L, Raffaelli R, Garzon S, Bosco M, Casarin J, Ciccarone F, et al. IPSICO survey on the psychological impact of COVID-19 on healthcare providers in obstetrics: a study design. *Ital J Gynaecol Obstet*. 2020;32(4):276-86. doi: 10.36129/jog.32.04.07.
11. Vitiello A, Ferrara F, Auti AM, Di Domenico M, Boccellino M. Advances in the Omicron vari-

- ant development. *J Intern Med.* 2022;292(1):81-90. doi: 10.1111/joim.13478.
12. Collie S, Champion J, Moultrie H, Bekker LG, Gray G. Effectiveness of BNT162b2 vaccine against omicron variant in South Africa. *NEJM.* 2022;386(5):494-6. doi: 10.1056/NEJMc2119270.
  13. Sigal A, Milo R, Jassat W. Estimating disease severity of Omicron and Delta SARS-CoV-2 infections. *Nat Rev Immunol.* 2022;22(5):267-9. doi:10.1038/s41577-022-00720-5.
  14. Abdullah F, Myers J, Basu D, Tintinger G, Ueckermann V, Mathebula M, et al. Decreased severity of disease during the first global omicron variant covid-19 outbreak in a large hospital in tshwane, south africa. *Int J Infect Dis.* 2022;116:38-42. doi: 10.1016/j.ijid.2021.12.357.
  15. Birol İter P, Prasad S, Mutlu MA, Tekin AB, O'Brien P, von Dadelszen P, et al. Maternal and perinatal outcomes of SARS-CoV-2 infection in unvaccinated pregnancies during Delta and Omicron waves. *Ultrasound Obstet Gynecol.* 2022;60(1):96-102. doi: 10.1002/uog.24916.
  16. İter PB, Prasad S, Mutlu MA, Tekin AB, O'Brien P, von Dadelszen P, et al. Maternal and perinatal outcomes of SARS-CoV-2 infection in unvaccinated pregnancies during Delta and Omicron waves. *Ultrasound Obstet Gynecol.* 2022;60(1):96-102. doi: 10.1002/uog.24916.
  17. Tekin AB, Yassa M, İter PB, Yavuz E, Önden B, Usta C, et al. COVID-19 related maternal mortality cases in associated with Delta and Omicron waves and the role of lung ultrasound. *Turk J Obstet Gynecol.* 2022;19(2):88-97. doi: 10.4274/tjod.galenos.2022.36937.
  18. Salo J, Hägg M, Kortelainen M, Leino T, Saxell T, Siikanen M, et al. The indirect effect of mRNA-based COVID-19 vaccination on healthcare workers' unvaccinated household members. *Nat Commun.* 2022;13(1):1162. doi:10.1038/s41467-022-28825-4.
  19. Mejri N, Berrazega Y, Ouertani E, Rachdi H, Bohli M, Kochbati L, Boussem H. Understanding COVID-19 vaccine hesitancy and resistance: another challenge in cancer patients. *Support Care Cancer.* 2022;30(1):289-93. doi: 10.1007/s00520-021-06419-y.
  20. Baral G, Shrestha O, Baral RS. Thrombotic pathology in placenta of COVID positive pregnancy. *J Nepal Health Res Coun.* 2021;19(1):206-8. doi: 10.33314/jnhrc.v19i1.3403.
  21. Debrabandere ML, Farabaugh DC, Giordano C. A review on mode of delivery during COVID-19 between December 2019 and April 2020. *Am. J. Perinatol.* 2021;38(04):332-41. doi: 10.1055/s-0040-1721658.
  22. Di Mascio D, Buca D, Berghella V, Khalil A, Rizzo G, Odibo A, et al. Counseling in maternal-fetal medicine: SARS-CoV-2 infection in pregnancy. *Ultrasound Obstet Gynecol.* 2021;57(5):687-97. doi: 10.1002/uog.23628.
  23. Franchi M, Bosco M, Garzon S, Laganà AS, Cromi A, Barbieri B, et al. Management of obstetrics and gynaecological patients with COVID-19. *Ital J Gynaecol Obstet.* 2020;32(1):6-19. doi: 10.36129/jog.32.01.01.
  24. Jarraya A, Zghal J, Abidi S, Smaoui M, Kolsi K. Subarachnoid morphine versus TAP blocks for enhanced recovery after caesarean section delivery: a randomized controlled trial. *Anaesth Crit Care Pain Med.* 2016;35(6):391-3. doi: 10.1016/j.accpm.2015.10.012.
  25. WAPM (World Association of Perinatal Medicine) Working Group on COVID-19. Maternal and perinatal outcomes of pregnant women with SARS-CoV-2 infection. *Ultrasound Obstet Gynecol.* 2021;57(2):232-41. doi: 10.1002/uog.23107.
  26. Di Mascio D, Sen C, Saccone G, Galindo A, Grünebaum A, Yoshimatsu J, et al. Risk factors associated with adverse fetal outcomes in pregnancies affected by Coronavirus disease 2019 (COVID-19): a secondary analysis of the WAPM study on COVID-19. *J Perinat Med.* 2020;48(9):950-8. doi: 10.1515/jpm-2020-0355.
  27. Shanes ED, Miller ES, Otero S, Ebbott R, Aggarwal R, Willnow AS, et al. Placental Pathology After SARS-CoV-2 Infection in the Pre-Variant of Concern, Alpha/Gamma, Delta, or Omicron Eras. *Int J Surg Pathol.* 2023;31(4):387-97. doi: 10.1177/10668969221102534.
  28. Parazzini F, Chiaffarino F, Chatenoud L, Tozzi L, Cipriani S, Chiantera V, et al. Maternal coffee drinking in pregnancy and risk of small for gestational age birth. *Eur J Clin Nutr.* 2005;59(2):299-301. doi: 10.1038/sj.ejcn.1602052.
  29. Chiaffarino F, Parazzini F, Chatenoud L, Ricci E, Sandretti F, Cipriani S, et al. Alcohol drinking and risk of small for gestational age birth. *Eur J Clin Nutr.* 2006;60(9):1062-6. doi: 10.1038/sj.ejcn.1602419.
  30. Abeysena C, Jayawardana P, DE A Seneviratne R. Maternal sleep deprivation is a risk factor for small for gestational age: a cohort study. *Aust N Z J Obstet Gynaecol.* 2009;49(4):382-7. doi: 10.1111/j.1479-828X.2009.01010.x.



31. Baral G, Shrestha O, Baral RS. Thrombotic pathology in placenta of COVID positive pregnancy. *J Nepal Health Res Counc.* 2021;19(1):206-8. doi: 10.33314/jnhrc.v19i1.3403.
32. Mortazavi-Jahromi SS, Aslani M. Dysregulated miRNAs network in the critical COVID-19: An important clue for uncontrolled immunothrombosis/thromboinflammation. *Int Immunopharmacol.* 2022;110:109040. doi: 10.1016/j.intimp.2022.109040.
33. Ahmed JQ, Maulud SQ, Dhawan M, Choudhary OP, Jalal PJ, Ali RK, et al. MicroRNAs in the development of potential therapeutic targets against COVID-19: A narrative review. *J Infect Public Health.* 2022;15(7):788-99. doi: 10.1016/j.jiph.2022.06.012.
34. Hayder H, O'Brien J, Nadeem U, Peng C. MicroRNAs: crucial regulators of placental development. *Reproduction.* 2018;155(6):R259-271. doi: 10.1530/REP-17-0603.
35. Chiofalo B, Laganà AS, Vaiarelli A, La Rosa VL, Rossetti D, Palmara V, et al. Do miRNAs play a role in fetal growth restriction? A fresh look to a busy corner. *Biomed Res Int.* 2017;2017:6073167. doi: 10.1155/2017/6073167.