

CASE REPORT

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Extensive bilateral diffuse infiltrates and deterioration of lung following infection with severe acute respiratory syndrome coronavirus 2 in a pregnant woman: a case report

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Abstract

Introduction: Severe acute respiratory syndrome coronavirus 2 is the third member of the coronavirus family to cause global concern in the twenty-first century. Pregnant women are particularly at higher risk of developing severe viral pneumonia, possibly because of a partial immune suppression during their pregnancy. Under such critical and rapidly evolving circumstances, these poor findings might be helpful for the treatment of infected pregnant women with the 2019 novel coronavirus.

Case presentation: In this study, we report the case of a 33-year-old Asian pregnant woman at 25 gestational weeks with coronavirus disease 2019 who developed severe complications, including hypoxemia, acute respiratory distress syndrome, pulmonary infiltration, and bilateral pleural effusion. She died 1 month after admission to the hospital.

Conclusion: Pregnant populations are especially at higher risk of viral pneumonia development caused by severe acute respiratory syndrome coronavirus 2. Further research on the prevention and treatment of the new coronavirus is necessary.

Keywords: Maternal death, SARS-CoV-2, Pulmonary fibrosis, Case report, COVID-19

Introduction

Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) is the novel coronavirus that was first discovered in the Chinese city Wuhan in December 2019 [1]. The outbreak of the SARS-CoV-2) was declared a global pandemic by the World Health Organization (WHO) on 11 March 2020. To date, there have been approximately 5 million confirmed cases, resulting in thousands of

deaths worldwide [2]. The severity of symptoms ranges from mild symptoms to severe illness. The most frequent symptoms include cough, pharyngitis, fever, myalgia, chills, and dyspnea [3].

Severe dyspnea is caused by the progression of lung lesions in multiple lung lobes, and in some cases in advanced stages, white lung syndrome is observed. Lung lesions are manifested as airspace opacities on plain chest radiography, or ground-glass opacities or consolidation on chest radiography images, usually with a rounded morphology and a peripheral lung distribution, which is a diagnostic method for coronavirus disease 2019

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(COVID-19) [4–6]. Pregnant women are particularly at higher risk of developing severe viral pneumonia, possibly because of a partial immune suppression during their pregnancy [7]. Pneumonia was reported as the third most common cause of mortality among pregnant women [8]. Previous data on Middle East respiratory syndrome (MERS) and severe acute respiratory syndrome (SARS) suggest that adverse pregnancy outcomes may occur due to the infection [9]. According to some studies, COVID-19 in the pregnant population can cause premature delivery and intrauterine growth restriction [6].

The effect of COVID-19 on pregnancy outcomes is not clear yet. Moreover, there has been no consensus on the maternal–fetal transmission of SARS-CoV-2 to date. Also, it has not been well understood whether pre-term delivery ameliorates the symptoms of a critically ill mother. In this case report, we present the case of a 33-year-old Asian pregnant woman who was infected with SARS-CoV-2 in the late second trimester of pregnancy. Under such critical and rapidly evolving circumstances, these poor findings might be helpful for the treatment of pregnant women who are infected with SARS-CoV-2 in Iran and elsewhere.

Case presentation

A 33-year-old Asian pregnant woman para 2, gravida 3, gestational age of 25 weeks, was hospitalized with fever, chills, shortness of breath, and myalgia. She had her first baby delivered by vaginal delivery 5 years ago and her second by cesarean section 2 years ago. The patient had no underlying diseases and did not have direct contact with COVID-19 cases. Vital signs of the patient were recorded as follows: respiratory rate (RR) 26 breaths

per minute, blood pressure (BP) 110/70 mmHg, temperature (T) 39.5 °C, heart rate (HR) 110 beats per minute, and oxygen saturation (SpO₂) 94%. The laboratory results showed that lymphocyte count was lower than normal (lymphocyte count $1 \times 10^9/L$). While platelet count, hepatic enzymes, and creatinine levels were within the normal range, C-reactive protein level (CRP) was significantly increased. Coagulation function and blood biochemistry were normal (Table 1). A GeneXpert SARS-CoV-2 RNA polymerase chain reaction (PCR) test was performed, and the result was positive. The patient had no complaints about uterine contractions, hemorrhage, and symptoms of amniorrhexis. Bilateral involvement was detected on chest computed tomography (CT), and on lung auscultation, diminished breath sound was detected. A day after the hospitalization, the patient was transferred to the intensive care unit (ICU) because of the shortness of breath (RR 32 breaths per minute) and low oxygen saturation (SpO₂ 88%). CT chest on the second day of hospitalization revealed an exacerbation of pulmonary involvement. The patient was intubated 1 week after hospitalization owing to a reduction in the oxygen saturation to 80% and exacerbation of respiratory distress. Chest CT on the eighth day of hospitalization showed extensive bilateral diffuse infiltrates and deterioration of lung involvement (Fig. 1). From the beginning of hospitalization, despite the use of broad-spectrum antibiotic coverage, the patient's fever was intermittent.

One week after transferring to the ICU and due to the patient's deteriorating condition, plasmapheresis was performed twice with 10 units of fresh frozen plasma (FFP) for the patient, which did not improve her condition. Also, due to a decrease in hemoglobin (HB 8.1 g/dl),

Table 1 Laboratory characteristics by the day of hospitalization

| | Day 1 | Day 4 | Day 8 | Day 12 | Day 16 | Day 20 | Day 24 | Day 28 |
|---|-------|-------|-------|--------|--------|--------|--------|--------|
| White blood cell count, $\times 10^9/L$ | 10.1 | 9.2 | 8.1 | 7.3 | 9.0 | 8.3 | 15.2 | 20.5 |
| Lymphocyte count, $\times 10^9/L$ | 1.0 | 1.0 | 1.0 | 0.8 | 0.8 | 0.5 | 0.7 | 0.5 |
| Hemoglobin, g/L | 12.8 | 11 | 8.8 | 9.6 | 9.4 | 8.3 | 11 | 10.5 |
| Platelet count, $\times 10^9/L$ | 310 | 345 | 250 | 210 | 158 | 140 | 144 | 126 |
| C-reactive protein, mg/L | 79 | 50 | | | | | | |
| Alanine aminotransferase, U/L | 15 | | 24 | | 28 | | | 35 |
| Aspartate aminotransferase, U/L | 25 | | 30 | | 30 | | | 32 |
| Creatinine, $\mu\text{mol/L}$ | 0.6 | 0.8 | 0.9 | 1 | 1 | 1.1 | 1.5 | 1.5 |
| VBG | | | | | | | | |
| pH | 7.39 | 7.46 | 7.41 | 7.49 | 7.39 | 7.42 | 7.19 | |
| PO ₂ , mmHg | 59.5 | 42 | 40 | 35.7 | 74.9 | 38 | 43.7 | |
| PCO ₂ , mmHg | 74.6 | 41.8 | 42.3 | 46.7 | 48 | 43.4 | 68.8 | |
| HCO ₃ , mEq/L | 45.1 | 30.1 | 30.5 | 31.5 | 29.5 | 28.2 | 26.2 | |
| BE, $\mu\text{mol/L}$ | 17.5 | 6.7 | 8.1 | 12.5 | 4.6 | 3.1 | 43.4 | |
| O ₂ Sat, % | 90.4 | 80.5 | 82 | 80 | 94.9 | 72.8% | 65.4 | |

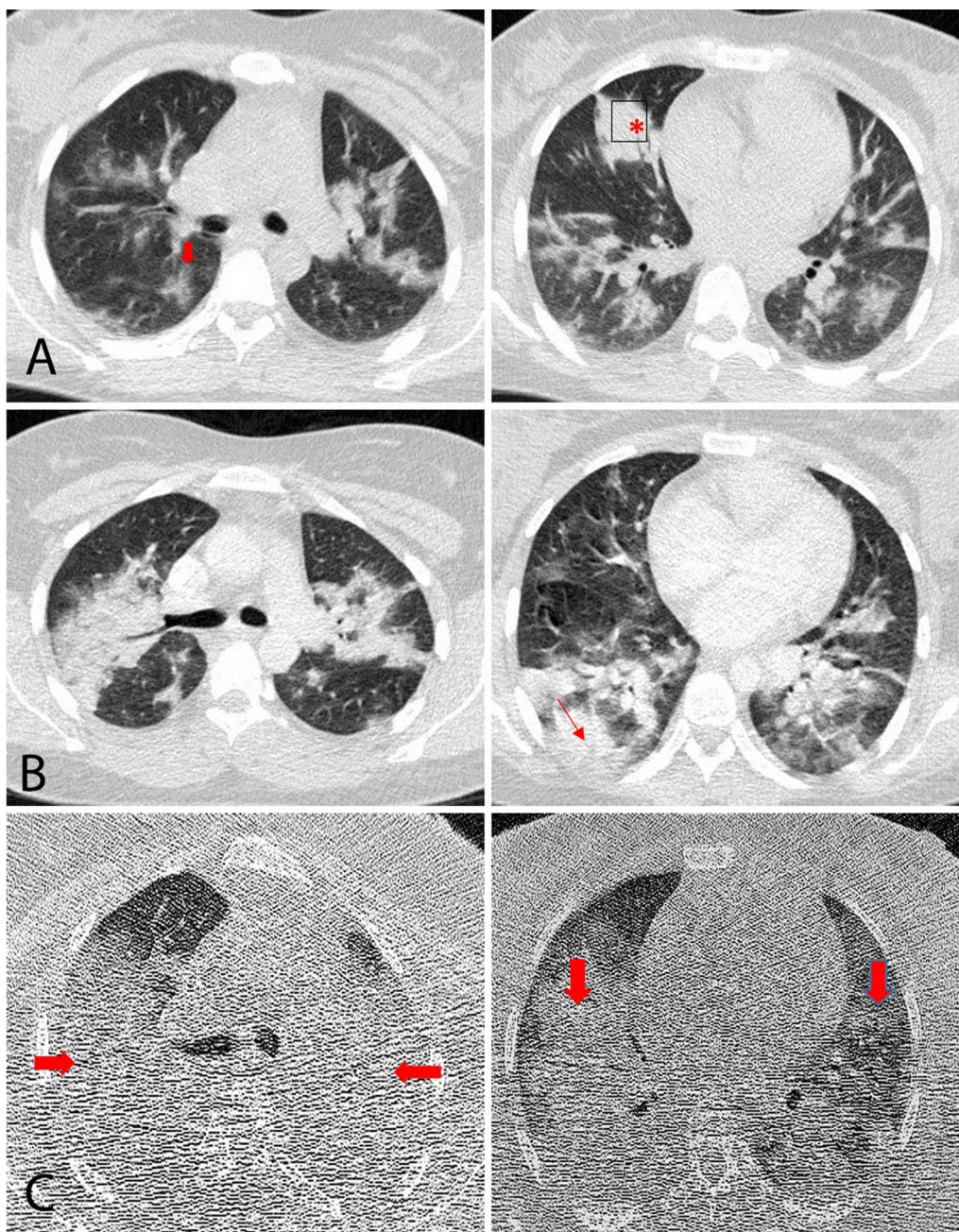


Fig. 1 **A** Axial CT image from day 1 of hospitalization showing multifocal ground-glass opacities (↓) and nodular consolidation (*) in both lungs. **B** Newly developed opacities and decreasing density of the nodular opacities (↓) detected on follow-up CT on day 2 of hospitalization. **C** Last follow-up CT image on day 8, showing extensive bilateral diffuse infiltrates (→) and deterioration of lung involvement

two units of packed cells were transfused twice. Three weeks after hospitalization, ultrasounds were performed several times to assess fetal growth, and a sharp decrease in the amniotic fluid (amniotic fluid index 2.5 cm) and fetal growth retardation were observed. To rule out the possibility of a preterm ruptured membrane, an

AmniSure test was performed twice, which was negative both times. Due to the condition of the fetus and the exacerbation of the patient’s respiratory condition, it was decided in the committee constituting of specialists in the special care and departments of lung and perinatology that the pregnancy should be terminated to

improve the patient's respiratory condition. Therefore, to induce cervical ripening, an intracervical Foley catheter was placed and extra-amniotic saline infusion was performed. After receiving three doses of vaginal misoprostol at doses of 50, 100, and 200 µg every 6 hours, a 600 g baby was born with a 1-minute Apgar score of 2 and a 5-minute Apgar score of 4, and the fetal heart rate was 145 beats per minute. The newborn was intubated and transferred to the neonatal intensive care unit (NICU) and died after 12 hours.

One week after the termination of pregnancy, due to the observation of bilateral pleural effusion on the chest CT, a chest tube was installed on both sides. Since no improvement in lung condition was observed and complete destruction of lung tissue was detected, the patient died 1 month after hospitalization.

Discussion and conclusion

Lessons learned from the previous viral outbreaks have proven the negative impacts of viral infection on maternal, neonatal, and fetal outcomes. For instance, the total mortality rate of 1918 influenza was 2.6%, while it was 37% among pregnant women [10]. It was shown that MERS or SARS infection during pregnancy correlated with a higher incidence of severe maternal and fetal complications, such as admission to the ICU, preterm delivery, maternal and fetal mortality, and intrauterine growth restriction [11]. According to a study conducted by Nan Yu and associates, the pregnancy outcomes of pregnant women with COVID-19 are considerably better than those of pregnant women with SARS [12].

As previously discussed, pregnant women are especially susceptible to the adverse complications of viral pneumonia. However, for COVID-19, data regarding the differences in the clinical features of pregnant and nonpregnant women are yet limited. Chen *et al.* have reported that the clinical signs and symptoms of pregnant women with COVID-19 are similar to those of nonpregnant women [13]. Also, in a large case series conducted in the Hubei Province of China, the incidence of severe pneumonia in the pregnant population was the same as in the general population [14]. More clinical evidence is needed to support this relationship.

Fever is the most prevalent symptom in pregnant women infected with SARS-CoV-2, observed in 78% of the cases [15]. Consistent with previous studies, our patient presented with fever, chills, dyspnea, and myalgia. It is confirmed that COVID-19 is associated with a decrease in lymphocytes, as observed in our study [13]. One study performed chest CT scans on 15 pregnant women with COVID-19 and indicated that the most common early finding was ground-glass opacities, as seen in our case [15]. Many studies have reported

that adverse outcomes and mortality are more likely to occur in patients with underlying medical conditions, whereas the present case did not have any underlying medical condition. In our study, the administration of convalescent plasma did not improve the clinical status of the patient, similar to the study of Niveditha that demonstrated that convalescent plasma has few or no clinical benefits for the treatment of COVID-19 [16]. It is not clear yet whether convalescent plasma with specific antibodies against SARS-CoV-2 that is collected from patients recovered from COVID-19 can improve the survival of critically ill patients. However, the plasma therapy has not been approved yet for use by the United States Food and Drug Administration, and numerous clinical trials are working on it [17].

Maternal complications of the present case were also in line with previous reports, which have found that pregnant women with severe or critical COVID-19 have a higher chance of developing respiratory failure, using mechanical ventilation, and maternal death, as well as fetal complications including preterm birth, intrauterine growth restriction, and intrauterine fetal death [18–21]. On the contrary, Zaigham *et al.* conducted a systematic review and, surprisingly, have suggested a lower rate of admission to the ICU, no maternal deaths, and only one neonatal death and one intrauterine fetal death [22].

In this patient, due to lack of improvement in respiratory status and progressive course of pulmonary destruction, as well as lack of fetal growth within 3 weeks and decreased immunity, it was decided to terminate the pregnancy to improve the respiratory condition. This is consistent with the rising concern that Castro *et al.* reported regarding preterm delivery in pregnant women with COVID-19. In their study, they found that 41% of deliveries in pregnant women with COVID-19 occurred before 37 weeks of pregnancy [23].

The limitation of our study was that no real-time polymerase chain reaction test was conducted on the premature infant, so we do not have any information about vertical transmission.

Since it has been proven in numerous studies that the pregnant population is especially at higher risk of viral pneumonia caused by coronaviruses, further research on the prevention and treatment of the new coronavirus is necessary.

Abbreviations

COVID-19: Coronavirus disease 2019; WHO: World Health Organization; ARDS: Acute respiratory distress syndrome; SARS-CoV-2: Severe acute respiratory syndrome coronavirus 2; RT-PCR: Real-time polymerase chain reaction; CT: Computed tomography; BP: Blood pressure; HR: Heart rate; GGO: Ground-glass opacity; T: Temperature; RR: Respiratory rate; CRP: C-reactive protein; ICU: Intensive care unit; SpO₂: Oxygen saturation; FFP: Fresh frozen plasma; NICU:

Neonatal intensive care unit; ABG: Arterial blood gas; BE: Base excess; HB: Hemoglobin.

Acknowledgements

We would like to thank the patient's family for their cooperation. Also, we wish to express our gratitude to the nurses who took care of the patient presented in this case report.

Authors' contributions

SD and PL conceived the idea. SM collected patient data and described it in the case report with a literature review. ZSh, FF, MB, SD, and MR had the primary responsibility for the writing of the manuscript. All authors read and approved the final manuscript.

Funding

This study was supported by the Student Research Committee of Mashhad University of Medical Sciences in Iran.

Availability of data and materials

The patient's information and medical records used for the case report are available from the corresponding author upon request.

Declarations

Ethics approval and consent to participate

This study did not include experiments on animals or humans. The patient's next of kin consented to the use of her personal data for the purpose of this case report.

Consent for publication

Written informed consent was obtained from the patient's next of kin for publication of this case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

Competing interests

The authors declare that they have no competing interests.

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Received: 3 June 2020 Accepted: 18 October 2021

Published online: 13 December 2021

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