Successful management of a pregnant woman with COVID-19 and multiple severe complications

Muhammad Ilham Aldika Akbar,1 Khanisyah Erza Gumilar,1 Brahmana Askandar Tjokroprawiro,1 Renata Alya Ulhaq2

1Department of Obstetrics and Gynecology, Medical Faculty, Universitas Airlangga, Surabaya, Indonesia
2Midwifery Study Program, Medical Faculty, Universitas Airlangga, Surabaya, Indonesia

Correspondence to
Dr Brahmana Askandar Tjokroprawiro; brahmanaaskandar@gmail.com

Accepted 29 August 2021

SUMMARY
We report a case of a 36-year-old gravida 2 para 1 woman at 38 weeks of gestation. A caesarean section was performed for severe pre-eclampsia, intrauterine growth restriction and oligohydramnios. The patient suffered postoperative bleeding, and exploratory laparotomy was performed. Uterine atonia, Couvelaire uterus and left adnexal haematoma were found, requiring a supracervical hysterectomy. As COVID-19 pneumonia and superimposed bacterial infection developed, the patient was mechanically ventilated in the intensive care unit. Remdesivir and meropenem were initially administered, but were changed to levofloxacin and ciprofloxacin following antibiotic sensitivity tests. Blood culture grew Enterococcus galinarum. Meanwhile, bleeding of the incisional wound occurred, which was controlled by the cessation of heparin therapy and regular wound care. With intensive monitoring and multidisciplinary management, the patient's condition improved, and she was discharged from the hospital on day 25 from admission.

BACKGROUND
Since it was first reported in China at the end of 2019, the COVID-19 pandemic has affected massive numbers worldwide.1 Up until March 2021, COVID-19 has affected more than 223 countries, with a total of 116521281 confirmed cases and 1589548 deaths.2 Indonesia has become the epicentre of COVID-19 in the South Asian region. Until now, there have been 1386556 confirmed cases of COVID-19 in Indonesia and 37547 deaths (a case fatality rate of 2.7%).1,4 The COVID-19 pandemic can considerably affect vulnerable populations, such as pregnant women, which can lead to increased maternal and neonatal mortality and morbidity. Pregnant women are more susceptible to severe manifestations of respiratory tract infections due to physiological changes in the immune and cardiovascular systems that occur during pregnancy.5 Pregnant women infected with SARS-CoV-2 are at an increased risk of maternal death, admission to intensive care unit (ICU), ventilator use, preterm birth and newborn admission to the neonatal ICU.6 The presence of comorbidities, such as chronic hypertension, obesity and gestational diabetes, is a risk factor of severe COVID-19 in pregnancy.6

We present a complex case of pregnancy further complicated by COVID-19, severe pre-eclampsia (PE) and postpartum haemorrhage (PPH), ultimately resulting in a hysterectomy. In addition to COVID-19, the patient also suffered from bacterial pneumonia, requiring long-term care in the ICU with multiple drug therapy. During treatment, the surgical wound in the abdomen was bleeding due to the adverse effect of heparin administered for anticoagulation. This case is unique because of its complexity, the course of the disease, side effects of therapy and successful multidisciplinary management.

CASE PRESENTATION
A 36-year-old pregnant woman, gravida 2 para 1, at 38–39 weeks of gestation was admitted to the hospital with several pregnancy complications, including severe PE, obesity class 3 (body mass index: 41.84 kg/m2), oligohydramnios (amniotic fluid index=3), suspicion of intrauterine growth restriction (IUGR) and COVID-19 with an estimated fetal weight of 2200 g. During the current pregnancy, the patient had attended 10 regular antenatal visits; her blood pressure (BP) was normal (systolic: 90–110 mm Hg and diastolic: 60–80 mm Hg). At 38 weeks, however, her BP reached 170/110 mm Hg with urinary protein of +2, establishing the diagnosis of severe PE. The patient was initially treated with the anticonvulsant MgSO4 intravenously (4 g of intravenous bolus maintained with a dose of 1 g/hour). At the time of admission, the patient was screened for COVID-19, and the IgG antibody to SARS-CoV-2 was reactive, although her chest radiograph revealed normal findings. A reverse transcription PCR (RT-PCR) test for SARS-CoV-2 was performed to confirm the diagnosis of COVID-19 infection. Fetal wellbeing tests showed an abnormal non-stress test (low variability and late deceleration), oligohydramnios and suspected IUGR. For the severe PE, IUGR and worsening fetal condition, caesarean section was performed, followed by stabilisation. We found meconium-stained amniotic fluid, and the baby was born with a birth weight of 2190 g (<5th percentile), height of 38 cm, and 1 min and 5 min appearance, pulse, grimace, activity and respirations (Apgar) scores of 7 and 8, respectively. After surgery, the maternal COVID-19 molecular test showed a negative result (on admission day 1 (D1)).

On the day after caesarean section, day 2 (D2), the maternal condition deteriorated and the patient’s BP and urine output decreased (BP: 100/60 mm Hg, urine output: 160 mL/13 hours). The patient was given fluid resuscitation, but her condition did not improve. On D3, her BP was still low (96/54 mm Hg), and laboratory examination revealed severe anaemia (haemoglobin: 3.8 g/
An ultrasound scan of the abdomen revealed significant free fluid, leading to the diagnosis of internal bleeding. Therefore, an exploratory laparotomy was performed. The uterus was found to be enlarged, atonic and resembled a Couvelaire uterus (figure 1). A Couvelaire uterus is a purplish-blue uterus caused by uterine bleeding that penetrates the myometrium. A hematoma was found in the left adnexa, which was considered the source of intrauterine bleeding. Initially, we tried to perform modified B-lynch procedure, but it was unsuccessful; therefore, we decided to perform a supracervical hysterectomy. We chose not to perform total hysterectomy in this case because the source of bleeding was in the uterine corpus. The cervix was maintained to partially preserve sexual function. The total blood loss during the surgery was 1200 mL with a fluid input of 2300 mL. The patient received a total of 3 units of whole blood (WB) transfusion before surgery and 2 units of WB during surgery.

The patient was then treated in the ICU using the ventilator mode pressure-synchronised intermittent mandatory ventilation (PSIMV), positive end expiratory pressure of 6 cm of water, and a fraction of inspired oxygen of 100%, with a global coma scale (GCS) of E4VxM6, and oxygen saturation of 97%–100%. The ventilator support was continued after surgery to maintain adequate oxygen delivery, as the patient had severe anaemia which limited oxygen delivery to the tissues. Another consideration was the risk of transfusion-related acute lung injury, which can occur as a result of massive transfusion. On laboratory examination, we found anaemia, hypoalbuminaemia, thrombocytopenia, hyponatraemia and hypokalaemia (table 1). The patient was administered non-steroidal anti-inflammatory drugs, ceftriaxone and albumin and underwent packed red cell

<table>
<thead>
<tr>
<th>Table 1: Laboratory results during treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Variables</td>
</tr>
<tr>
<td>------------------------------------------------</td>
</tr>
<tr>
<td>Hb (g/dL)</td>
</tr>
<tr>
<td>Hct (%)</td>
</tr>
<tr>
<td>PLT (10^9/L)</td>
</tr>
<tr>
<td>PT/PTT (s)</td>
</tr>
<tr>
<td>RBG (mg/dL)</td>
</tr>
<tr>
<td>AST/ALT (U/L)</td>
</tr>
<tr>
<td>BUN/SC (mg/dL; U/L)</td>
</tr>
<tr>
<td>Albumin (mg/dL)</td>
</tr>
<tr>
<td>Ca/Mg (mg/dL)</td>
</tr>
<tr>
<td>Fibrinogen/D-dimer (mg/dL; µg/mL)</td>
</tr>
<tr>
<td>Procalcitonin (ng/mL)</td>
</tr>
<tr>
<td>IgG/IgM</td>
</tr>
<tr>
<td>RT-PCR</td>
</tr>
</tbody>
</table>

ALT, alanine transaminase; ALT, aspartate transaminase; APTT, activated partial thromboplastin time; BUN, blood urea nitrogen; BB, bacte...
transfusion and serum electrolyte correction. After 3 days of treatment in the ICU (D4), ronchi were detected on auscultation of the lungs, which was suspected to be due to pulmonary oedema and intravenous meropenem, while repeated blood culture and SARS-CoV-2 molecular tests were performed. Again, the SARS-CoV-2 molecular test was negative. However, on D7, the RT-PCR for SARS-CoV-2 was positive, and the patient was diagnosed with COVID-19 with superimposed bacterial pneumonia. The pulmonologists commenced administration of remdesivir 100 mg three times per day for COVID-19 pneumonia. Over the next few days, blood culture showed no growth of microbes. During treatment in the ICU, the patient’s BP was consistently normal and the fever had started to resolve. Wound care was maintained for a week secondary to unstable respiratory function and oxygenation parameter during this period.

On D8, the patient’s condition improved, with a GCS of 14 (completely conscious). Therefore, the patient was removed from ventilator and maintained on a non-rebreathing mask (NRBM) with adequate oxygen saturation (97%–98%). On D9, because of increased D-dimer levels, the patient was administered heparin at 400 units/hour intravenously as a prophylactic anticoagulant. Over the next 2 days (D9 and D10), the patient had fever (temperature of 37.8°C) and was administered antipyretics. On D11, the incisional wound in the abdomen was bleeding (figure 3), although the vital signs were normal and the fever had started to resolve. Wound care was performed two times per day using sterile water. Moreover, on D11, the patient was started on convalescent plasma transfusion 200 mg two times per day for 2 days. We performed an ultrasound to evaluate the source of bleeding in the incisional wound and found a haematoma superficial to fascia in the surgical site. The bleeding was suspected to be caused by heparin therapy, and the administration of heparin was stopped on D13. Following cessation of heparin, no incisional site bleeding was observed. We continued to monitor the progression of COVID-19, and the fourth (D13) and fifth (D19) RT-PCR results from the nasopharyngeal- oropharyngeal swab samples were negative for SARS-CoV-2. On D18, blood culture showed the presence of *Enterococcus gallinarum*, which was sensitive to ampicillin, ciprofloxacin and levofloxacin. Therefore, the antibiotics were changed to levofloxacin 750 mg once a day and ciprofloxacin 200 mg once a day. The patient’s condition improved during the next several days. On D24, the patient had stable vital signs: normal BP (126/73 mm Hg) and good saturation with O₂ from nasal prongs (95%–96%), and the incisional wound had healed. The patient was discharged on the next day (D25). The timeline of this case is shown in figure 4.

Written informed consent was obtained from the patient for the publication of this case report and its accompanying images.

**DISCUSSION**

Although RT-PCR is the gold standard to diagnose COVID-19, in this case, we found that on initial examination the molecular test showed negative results twice. On admission, the patient was suspected of COVID-19 based on clinical signs and results of the rapid antibody test, although the chest radiographic findings were normal. Although this screening method is not ideal, this had to be performed due to limited RT-PCR test in the hospital. On the third swab (D7), the RT-PCR test showed a positive result, and the patient was diagnosed with COVID-19 pneumonia. This case suggests that RT-PCR in SARS-CoV-2 is not 100% sensitive, and there is a possibility of it being false negative. In a meta-analysis involving 12057 patients, the initial false-negative rate of RT-PCR in SARS-CoV-2 was from 2% up to 54%.[7, 8] In this case, the possibility of an initial false-negative result may be related to the disease being in the incubation period, and the number of viral copies being low is to be detected in the molecular test. Therefore, in the management of pregnant patients with suspected COVID-19 based on clinical and laboratory signs, the RT-PCR test should be repeated periodically to minimise the risk of false-negative results.[7, 8]

Severe PE was also found in this case, which may be explained by either the coexistence of true PE with COVID-19 or a severe PE-like syndrome induced by COVID-19. COVID-19 can induce clinical manifestations that resemble severe PE, including hypertension, renal and hepatic involvement, and proteinuria. The differentiation between these two conditions...
is difficult. The soluble fms-like tyrosine kinase 1, placental growth factor level and uterine artery Doppler index may differentiate between these conditions. In this case, we treated the condition as severe PE with COVID-19, and MgSO4 treatment was undertaken to prevent seizures. The diagnosis of PE instead of PE-like syndrome of COVID-19 was established based on the presence of hypertension, proteinuria, hypoalbuminaemia and IUGR in accordance with the International Society for the Study of Hypertension in Pregnancy classification. The presence of severe PE increases the risk of PPH, which in this case necessitated hysterectomy because of failed conservative treatment during surgery. However, COVID-19 itself has not been proven to increase the risk of PPH as per a retrospective study in the USA.

In this case, COVID-19 pneumonia also coexisted with a bacterial infection. The diagnosis of bacterial pneumonia rather than pulmonary oedema was established based on the clinical signs of fever and tachypnoea, pulmonary radiography, a high serum procalcitonin level and a positive blood culture. Cardiac causes were excluded based on the normal echocardiographic findings. Management was aimed at maintaining adequate oxygenation with mechanical ventilation (ventilator mode PSIMV) during treatment in the ICU with a target oxygen saturation of >95%. After the respiratory function was stable, the ventilator was removed and replaced with an NRBM, and eventually a simple mask or nasal O2. The blood culture later grew E. gallinarum, VanC-harbouring enterococci which have low-level resistance to vancomycin. These enterococci have less impact on nosocomial infection control and only occasionally cause clinical disease. Sixty percent of infection by E. gallinarum manifest as sepsis within 24 hours. In a group of mechanically ventilated female patients recovery rate with low incidence of serious adverse events with parturum women with remdesivir administration, showed a high study by Burwick time of recovery in adults with severe COVID-19 infection. A treatment of COVID-19 pneumonia and has been shown by the rationale of convalescent plasma therapy is to provide specific antibodies against the virus to help the immune system eradicate the pathogen.

The management of COVID-19 in pregnancy with respiratory compromise is to provide adequate respiratory support. In this case, the patient required mechanical ventilation due to respiratory distress and unstable oxygenation. The goal of respiratory support is to maintain an oxygen saturation of >95% and reduce tachypnoea and dyspnoea. Oxygen therapy should be administered if the SpO2 values fall below 95% in pregnant women and below 90% in non-pregnant women. The threshold of oxygen therapy in pregnant women is higher due to increased oxygen demand and placental transfer of oxygen to the fetus.

In our case, the incisional site bleeding was attributed to heparin therapy commenced for venous thromboembolism prophylaxis. Our patient had increased D-dimer levels, which is considered prognostic for thrombosis in COVID-19. Coagulopathy has been reported in >50% of patients with severe manifestations of COVID-19. The International Society on Thrombosis and Haemostasis has recommended the prophylactic administration of anticoagulants in patients with COVID-19 to prevent thrombosis and organ damage. Pregnancy has been considered as a prothrombotic state due to the physiological changes. The presence of COVID-19 increases vascular inflammation, which places pregnant women at risk of thrombosis. Therefore, anticoagulants should be administered in the management of pregnant women with COVID-19, especially in patients with elevated D-dimer (>1g/mL). The Society for Maternal-Fetal Medicine supports prophylactic heparin use in critically ill or mechanically ventilated pregnant women with COVID-19. Clinical studies have shown that the risk of a major haemorrhagic event during heparin therapy for deep venous thrombosis or pulmonary embolism is approximately 4%. Heparin therapy in COVID-19 requires intensive monitoring to balance the need for anticoagulation with the risk of bleeding. The decision to use anticoagulants should be based on an analysis of the potential risks and benefits. Risks include postoperative bleeding, as seen in our case, while the benefit of anticoagulation in patients with COVID-19 is the decreased risk of thromboembolic events. In our case, the patient also had thrombocytopenia, which could have contributed to the PPH and the Couvelaire uterus.

This complex case showed a limitation on the diagnosis and management of COVID-19 in pregnancy with multiple complications. COVID-19 can cause multiple direct or indirect complications in pregnancy. The diagnosis of COVID-19 should not be based solely on a molecular test, especially in the very early stage of the disease. The complete assessment of clinical findings, laboratory results, chest imaging and other parameters may establish the diagnosis of COVID-19 infection, despite a negative molecular test result. COVID-19 can be complicated by PE which significantly increases the risk of adverse pregnancy outcomes such as PPH. In COVID-19, the presence of complications such as PE leading to PPH requires intensive postsurgery observation to minimise the risk of PPH. A postmortem study of 621 patients found that bacterial superinfection was present in 32% of cases, mainly (95%) in the form of pneumonia. If bacterial infection is suspected, blood and sputum cultures
should be performed, with antibiotic sensitivity testing, to establish the diagnosis. Finally, anticoagulation should be used with caution in pregnant patients with COVID-19 as patients who undergo surgery may be at increased risk of bleeding.

**Patient’s perspective**

Pregnancy with multiple complications should be treated in a tertiary hospital with adequate facilities and an experienced multidisciplinary team, which is why I could be saved.

**Learning points**

- The RT-PCR test for the detection of SARS-CoV-2 should be repeated if the initial test is negative and clinical suspicion remains high. Sequential tests can enhance the diagnostic yield.
- COVID-19 may induce clinical manifestations that resemble severe pre-eclampsia (PE), including hypertension, renal and liver involvement, and proteinuria.
- COVID-19 complicated with PE can increase the risk of postpartum haemorrhage and may require close observation after surgery.
- The possibility of superimposed bacterial infection in COVID-19 pneumonia should always be considered in prolonged hospitalised patients.
- Prophylactic heparin in pregnant women with COVID-19 needs intensive monitoring of clinical signs of the side effects of heparin to balance the need for anticoagulants with the risk of bleeding, especially in patients who need surgical intervention.

**Contributors**

MIAA was involved in conceptualisation and drafting. BAT was involved in drafting, editing and formatting. KEG was involved in data collection and editing. RAU was involved in data collection and editing. KEG was involved in drafting, editing and formatting. KEG was involved in data collection.

**Funding**

The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

**Competing interests**

None declared.

**Patient consent for publication**

Obtained.

**Provenance and peer review**

Not commissioned; externally peer reviewed.

**Open access**

This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/.

**ORCID iD**

Brahmana Askandar Tijokoprawiro http://orcid.org/0000-0003-1658-3477

**REFERENCES**
