Takotsubo cardiomyopathy in early term pregnancy: a rare cardiac complication of SARS-CoV-2 infection

Pranab J Bhattacharyya, Pawan K Attri, Waseem Farooqui

DESCRIPTION
A 32-year-old primigravida at a 38-week gestation was initially admitted in cardiology isolation ward on referral by her local obstetrician for inferolateral ST-segment elevation on ECG (figure 1A) which was obtained for complaints of New York Heart Association functional class II symptoms with palpitations of a 3-day duration. Except for a blood pressure of 150/100 mm Hg on presentation, the rest of her physical examination, vital signs and medical history were insignificant. She was on amlodipine for gestational hypertension. A transthoracic echocardiogram (TTE) demonstrated hypokinetic mid and akinetic apical left ventricular (LV) segments and hypercontractile basal segments with prominent apical ballooning typical of takotsubo cardiomyopathy (TTC) (figure 1B, arrows). Two-dimensional speckle tracking echocardiography revealed LV global longitudinal strain (GLS) of −13.9 and ejection fraction (EF%) of 38% (figure 1C). Blood tests showed elevated troponin I (37.00 pg/mL, normal <26.20) and N-terminal pro b-type natriuretic peptide (NT-proBNP) (1312.00 pg/mL, normal <115.00). Nasopharyngeal swab reverse transcription PCR (RT-PCR) test was positive for SARS-CoV-2 infection. Coronary angiography (CAG) was deferred due to her active COVID-19 status. She was further managed in the ward of our dedicated COVID-19 hospital where she was started on medical therapy with bisoprolol and enoxaparin (40mg subcutaneous twice per day for 5 days) along with oral vitamins and antibiotic as per local treatment protocol for patients with COVID-19. Regular maternal and fetal monitoring continued. As oxygen saturation in room air and respiratory rate were maintained within normal limits without any clinical evidence of pneumonia or respiratory failure, she did not require mechanical ventilation at any stage during her 7-day stay in the COVID-19 ward. Additional laboratory tests showed leucocytosis (20.48×10^9/μL, normal 4.00–11.00), neutrophilia (85.8%, normal range 37–72), lymphopenia (9.2%, normal range 20–40), neutrophil–lymphocyte ratio (9, normal 1–3), raised levels of D-dimer (0.69 μg/mL, normal <0.05), lactate dehydrogenase (456 U/L, normal range 120–246), alkaline phosphatase (188 U/L, normal range 38–126) and globulin (3.5 g/dL, normal range 2.80–3.20), low levels of albumin (3.0 g/dL, normal range 3.50–5.00) and albumin/globulin ratio (0.90, normal 1.25–1.51) and normal levels of ferritin (116 ng/mL, normal range 11–306.8), total bilirubin (0.50 mg/dL, normal range 0.20–1.3), aspartate aminotransferase (37 U/L, normal range 17–59) and alanine aminotransferase (28 U/L, normal range 4–50). On day 8, her RT-PCR test was negative and due to the onset of early labour, she was shifted to labour and delivery ward for necessary obstetrical management. Subsequently on the same day, she underwent an uneventful caesarean section delivery under spinal anaesthesia for fetal distress and associated cephalopelvic disproportion. Despite a diagnosis of TTC, an expedited delivery in our patient was not considered by the obstetrical unit as there was no evidence of clinical or haemodynamic worsening of maternal or fetal status. Repeat TTE on day 13 on transfer to the cardiology ward showed the normalisation of the LV regional wall motion abnormalities (RWMA) and significant improvement of GLS (−16.5) and EF% (51%) (figure 1D) further ratifying the diagnosis of TTC. Subsequent CAG on day 14 revealed non-obstructive coronary artery disease (CAD) involving the left anterior descending artery (figure 1E, arrows). She was finally discharged from the cardiology ward after full recovery on day 16 with aspirin, atorvastatin and bisoprolol.

As typified by this index case, TTC can mimic acute ST-segment elevation myocardial infarction and is considered to be a reversible form of cardiomyopathy characterised by a complete recovery of RWMA and LV function within weeks of presentation. Incidental CAD can be present in up to 10%
cases. Although TTC classically affects postmenopausal women, it has been infrequently reported previously in pregnant women unrelated to SARS-CoV-2 infection. This is the first reported case of TTC in pregnancy as a manifestation of SARS-CoV-2 infection during this ongoing pandemic.

TTC can be preceded by emotional or physical stressful triggers. Coronary artery vasospasm, coronary microvascular dysfunction, LV outflow tract obstruction and catecholamine surge have all been elucidated as potential mechanisms. As TTC has also been reported with viral infections, the more intense inflammation associated with COVID-19 may contribute to its development. Deranged inflammatory markers were also a notable finding in our patient described previously. Whether the inflammatory response of SARS-CoV-2 infection and any specific markers that may portend a greater likelihood of development of TTC especially in pregnancy may be a subject matter for further study. The overall prognosis of TTC is favourable, with full recovery of LV function seen in most patients by 2 months. Pregnancy women may be at greater risk for SARS-CoV-2 infection as the virus enters the cell via the ACE 2 receptor, which is upregulated in normal pregnancy.

Therefore, treating physicians dealing with the COVID-19-positive pregnant population need to remain vigilant towards this rare cardiac complication of SARS-CoV-2 infection.

Contributors PJB, PKA, and WF were involved in patient management. PJB prepared the manuscript. PJB and PKA acquired the echocardiographic images and PJB and WF performed the coronary angiogram. All authors have revised the manuscript and approved of the final draft.

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.

This article is made freely available for use in accordance with BMJ’s website terms and conditions for the duration of the covid-19 pandemic or until otherwise determined by BMJ. You may use, download and print the article for any lawful, non-commercial purpose (including text and data mining) provided that all copyright notices and trade marks are retained.

REFERENCES