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Case Report

Emergent caesarean section under mechanical circulatory support for acute severe peripartum cardiomyopathy



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ABSTRACT

Peripartum cardiomyopathy (PPCM) is an idiopathic left ventricular dysfunction in women who are in late pregnancy or the postpartum period. PPCM is a rare but sometimes fatal disease, and mechanical circulatory support is required if heart failure is refractory to conventional therapy. A 28-year-old woman in late pregnancy was admitted to our hospital due to congestive heart failure with cardiogenic shock. Her heart rate was 200 beats per minute (sinus tachycardia), and left ventricular ejection fraction (LVEF) was 10%. Additionally, fetal heart rate decreased to 80 beats per minute. It was extremely difficult to continue her pregnancy because of decompensated heart failure and fetal asphyxia; therefore, we delivered her baby via caesarean section after initiating mechanical circulatory support. With optimal medical therapy, including bromocriptine, we were able to remove mechanical circulatory support. Additionally, LVEF improved to 42%, and she was discharged with her baby who had no growth failure. This case highlights the safety and risk of caesarean section under mechanical circulatory support, and the effectiveness of bromocriptine.

<Learning objective: Mechanical circulatory support is occasionally required if heart failure is refractory to conventional medical therapy in peripartum cardiomyopathy (PPCM). Mechanical circulatory support, particularly extracorporeal membrane oxygenation needing sufficient anticoagulation, should be initiated following delivery via caesarean section because of bleeding from the uterus. Moreover, bromocriptine may be effective in improving PPCM and should be the first-line treatment in clinical practice.>

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Introduction

Peripartum cardiomyopathy (PPCM) is an idiopathic left ventricular dysfunction, similar to dilated cardiomyopathy, in women who are in late pregnancy or the postpartum period. It is reported that PPCM is a rare but sometimes fatal disease [1], particularly during subsequent pregnancy in women with a history of PPCM who had preserved left ventricular dysfunction [2]. There is no effective and specific therapy for PPCM, and mechanical circulatory support and heart transplantation are required if heart failure is refractory to conventional therapy. In fact, because PPCM is a life-threatening condition, rapid therapeutic interventions are

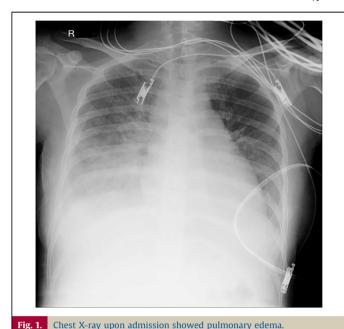
Case report

We present a 28-year-old woman, gravida 3 and para 2, at 35 weeks and 6–7 days of gestation. Her first delivery was at 24 years and her second was at 26 years. During these pregnancies, she was asymptomatic, but sinus tachycardia was indicated. At 28 years, she became pregnant a third time. Her heart rate was approximately

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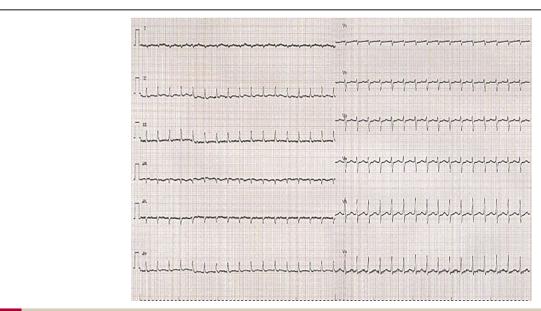
necessary, and are typically performed and managed after delivery or abortion. To the best of our knowledge, there are only a few case reports demonstrating caesarean section under mechanical circulatory support before delivery. In addition, a small randomized pilot study and several observational reports have suggested the usefulness of bromocriptine for PPCM [3–6]. We report our experience with a patient who developed PPCM that required emergent caesarean section under mechanical circulatory support, and who was prescribed bromocriptine.

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140 beats per minute at 10 weeks of gestation, and then she experienced feebleness. At 34 weeks of gestation, her heart rate increased to 170 beats per minute. The patient presented with sinus tachycardia throughout her pregnancy but without hypertension and preeclampsia. At 35 weeks and 6-7 days of gestation, she was admitted into a local clinic due to premature labor. Obstetricians administered ritodrine hydrochloride, and her heart rate continued to rise, up to 200 beats per minute, and she experienced progressive dyspnea. Her heart rate did not decrease once the medicine was discontinued; therefore, she was transferred to our hospital. Upon examination, her systolic blood pressure was 70 mmHg and oxygen saturation was 94% on 2 L/min via oxygen cannula. Respiratory sound was coarse crackle, and her chest X-ray showed pulmonary edema (Fig. 1). Her brain natriuretic peptide (BNP) level was 1000 pg/mL, but troponin I was not elevated. Hemoglobin level was 9.4 g/dL and thyroid function was as follows: thyroid stimulation hormone, $1.120~\mu IU/mL$; free T4, 1.03~ng/dL. Her electrocardiogram demonstrated a heart rate at 203 beats per minute (sinus tachycardia) (Fig. 2), and her left ventricular ejection fraction (LVEF) was 10% and left ventricular diastolic dimension was 45 mm based on echocardiogram.

Her respiratory condition rapidly deteriorated, as her arterial blood gas showed severe respiratory acidosis: pH, 6.765; PaCO₂, 94 mmHg; PaO₂, 26 mmHg. Fetal heart rate also decreased to 80 beats per minute. Because of these factors, it was extremely difficult to continue her pregnancy. We needed to deliver her baby immediately, but vaginal delivery and caesarean section were also considered dangerous due to decompensated heart failure. Therefore, we initiated mechanical circulatory support, which included intra-aortic balloon pump (IABP) counterpulsation and peripheral veno-arterial extracorporeal membrane oxygenation (ECMO), prior to delivery. Blood flow rate was maintained at approximately 3.0 L/min through ECMO. After these treatments, caesarean section was performed under general anesthesia by obstetricians. Coronary angiography, which was performed after the caesarean section, showed no stenosis, and histological examination of cardiac tissue obtained via endomyocardial biopsy during cardiac catheterization found no characteristic findings (Fig. 3). As a result, we diagnosed her with PPCM due to the absence of another identifiable cause for heart failure, and we prescribed bromocriptine (2.5 mg twice daily) with noradrenaline (up to 0.015 µg/kg/min) to increase blood pressure and landiolol (up to 80 µg/kg/min) to decrease heart rate. Following these medicinal treatments, her blood pressure gradually increased and her heart rate gradually decreased. We then tapered her off landiolol and changed her prescription to bisoprolol at a dose of 0.625 mg daily on the third day after admission. During this period, we intravenously injected furosemide (5 mg/h) to treat her pulmonary congestion. She also experienced sustained bleeding from her uterus because of administered unfractionated heparin, which kept activated clotting time from 180 to 200 s in order to maintain mechanical circulatory support, and she was in dire need of a blood transfusion. Since the insertion of a Bakri balloon (Cook Women's Health, Spencer, IN, USA) and a transcatheter arterial embolization were not effective, we performed a hysterectomy on the fifth day after admission. Her left ventricular dysfunction gradually improved, indicating that we could remove the mechanical



2. Electrocardiogram upon admission demonstrated a heart rate of 203 beats per minute (sinus tachycardia).

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