



Endocarditis during Pregnancy in a Woman with Interventricular Septum Defect

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Background

Infective endocarditis is a rare but life-threatening infection during pregnancy and in the postpartum period. For this reason, a high index of suspicion leading to early diagnosis and institution of appropriate management is required [1].

In the current era, risk factors are changing, with a marked decrease in rheumatic heart disease, as an underlying cardiac risk factor, and a concomitant increase in congenital heart disease and intravenous drug use.

The presence of Ventricular Septum Defect (VSD) is one of the predisposing factors; in fact, VSD, according to several registries, is the most frequent congenital heart disease in pregnant women [2] (Figure 1). Over time, maternal and fetal outcomes have improved and this is likely related to advances in maternal-fetal monitoring, better surgical techniques, and availability of nontoxic and effective antimicrobial agents.

Staphylococcus, *Streptococcus*, and *Enterococcus* species account for between 80% and 90% of all cases of IE worldwide [3]. In particular, *S. gordonii* is a rare cause of IE that has been sparsely reported in the literature. *S. gordonii* are Gram-positive, alpha-hemolytic chains of cocci that play an important role in the alkalization of the oral cavity and protective biofilm production [4]. Once in the blood stream, *S. gordonii* appear to have the virulence factors that are pathogenic in the development of IE. The cell wall of *S. gordonii* contains a serine-rich glycoprotein, GspB, which mediates binding to human platelets [4]. After adherence to the platelet, the combination has the potential to attach to the fibronectin-rich extracellular matrix of the cardiac valves and subsequently form valvular vegetations.

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Case Presentation

A 40-year-old patient was admitted with fever at 18 weeks' gestation in her fourth pregnancy.

Medical history included a small Ventricular Septal Defect (VSD) which was diagnosed in childhood and never required surgical intervention. The patient underwent regular follow up: Her last cardiology assessment in 2019 was normal.

One month prior to admission (March 2020) she suffered from pain in the upper right dental arch and fever; she was treated with amoxicillin for 5 days (April 2nd to 7th) with benefit. Four days after stopping the therapy, fever reappeared and inflammatory markers increased; a one-week antibiotic cycle with amoxicillin and ceftriaxone was administered; but again, a recrudescence of fever occurred after the suspension of therapy.

The following examinations were performed:

1. Abdomen ultrasound (April 8th): No significant alterations.
2. IgM and IgG dosage for toxoplasma (14.4): Negative
3. Coproculture (20.4): Negative.

On April 20th, a dental granuloma was removed. Few days after, the patient had recurrence of fever.

Laboratory test were performed (Table 1) and blood cultures were obtained, testing positive for *Streptococcus gordonii*. The patient was finally admitted to our department. Body mass index was 19 at admission.

Table 1: Laboratory investigations.

Investigation	Value	Reference range
Hemoglobin	9.6g/dL	9.5-15g/dL
White cell count	10 × 10 ⁹ /L	5.9-16.9 × 10 ⁹ /L
C-reactive protein	123mg/L	0.4-8.1 mg/L
Arterial blood gas	pH 7.5	7.39-7.45

A blood culture was repeated on day 1, confirming positivity for *S. gordonii* (Figure 2). Transthoracic echocardiography (Figure 3) confirmed the presence of VSD but no evidence of endocardial vegetation was found. According to ESC guidelines, we performed a transesophageal echocardiography. No vegetation was detected but we noted a hypoechogenic thickening near the VSD (Figure 4, 5), probably due to turbulent flow. We think it's very likely that this zone can favor bacterial growth, which was prevented by the presence of rapid flow at this level; however, microlesions created by the turbulent flow itself may have triggered transient bacterial permanence.

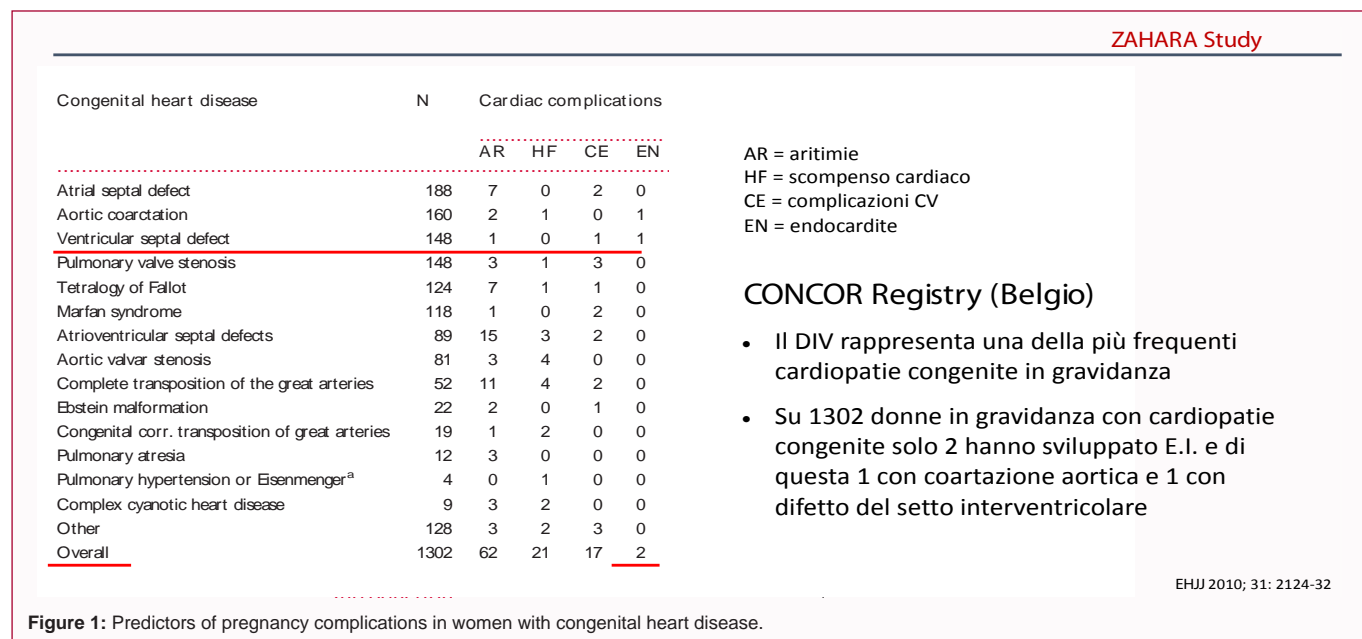
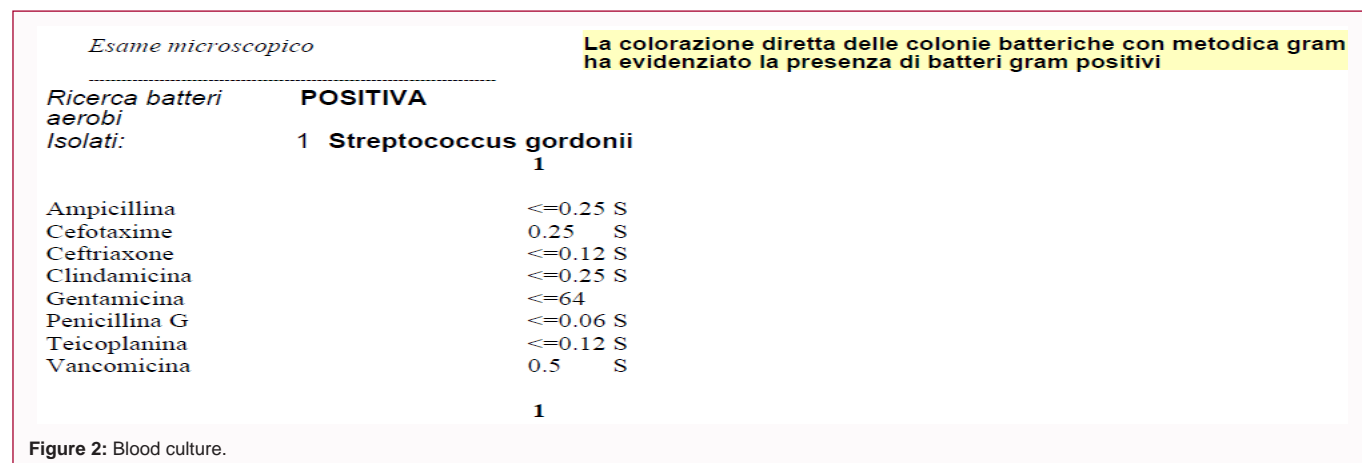
The patient was treated with intravenous ceftriaxone 2 g once daily and blood culture, 4 weeks later, were negative.

Obstetric ultrasounds showed no fetal suffering. The patient was dismissed in good conditions.

Discussion

We report the case of a 40-year-old patient who presented with fever at 18 weeks and 3 days of gestation in her fourth pregnancy. A diagnosis of infective endocarditis was made. The patient described developed *Streptococcus gordonii* infective endocarditis, which is a rare but aggressive causative organism in infective endocarditis. Infective endocarditis in pregnancy is a rare but serious condition with significant fetal and maternal morbidity and mortality [5]. Early diagnosis with a multidisciplinary team approach is essential to improve outcomes.

Several studies showed that, even today, maternal and fetal mortality is higher than 10% [6]. For this reason, in our case, given the positive blood culture even in the absence of a clear endocarditic vegetation (see echocardiogram findings), we prescribed targeted antibiotic therapy. The patient was dismissed 4 weeks after completion of antibiotic treatment, with negative blood culture. During the pregnancy, the baby showed a delay in surfactant development requiring corticosteroid therapy; shortly after the delivery, she developed a pneumonia with respiratory failure requiring prolonged stay in neonatal ICU. Happily, she totally recovered after 14 days.

**Figure 1:** Predictors of pregnancy complications in women with congenital heart disease.**Figure 2:** Blood culture.

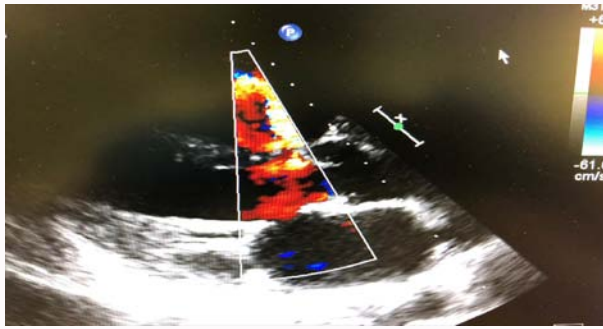


Figure 3: TTE showing the ventricular septum defect.



Figure 5: Hypoechoic thickening near the VSD.



Figure 4: Hypoechoic thickening near the VSD.

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