

The atrioventricular complete heart block diagnosed on the preoperative routine test for caesarian section

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Abstract

Objective: The management of women presenting with complete heart block during pregnancy remains very challenging. Until now, there is not an established consensus for the most appropriate anaesthetic technique for caesarean section in women with complete atrioventricular block.

Case(s): On our case, the atrioventricular complete heart block was diagnosed on the preoperative routine test for Caesarian Section due to cephalo-pelvic disproportion. The patient had no regular antenatal check ups at a local hospital. Her parents reported rare episodes of syncope during childhood and adolescence and one more episode two years before. During pregnancy she did not report any syncope episode except from being tired. For obstetric reasons caesarean section was performed successfully under spinal anaesthesia with continuous monitoring during intraoperative time without a pacemaker. Even though the patient reacted well during administration of atropine a temporary pacemaker was found to be in case we would need it. A healthy baby boy of 3350 gram was delivered. During postpartum period the patient did not have any complaints or syncope episodes. It was strongly recommended to her a regular follow up to cardiology department.

Conclusion: As suggested by our case, asymptomatic atrioventricular complete heart block in pregnancy can be managed successfully without pacemaker. However, careful monitoring, is necessary by the pregnancy heart team with a cardiologist, anaesthetist and obstetrician, with experience in the management of high risk pregnancies. Management of the risk for cardiovascular and obstetrical complications is difficult in pregnant women with complete heart block. Asymptomatic complete heart block in late pregnancy should be managed without pacemaker by the pregnancy heart team with a cardiologist, anaesthetist and obstetrician, with experience in the management of high risk pregnancies.

Keywords: Pregnancy, complete atrio ventricular heart block, temporary pacing, obstetrical complication

Introduction

Management of the risk for cardiovascular and obstetrical complications is difficult in pregnant women with complete heart block. Complete congenital or acquired atrioventricular block can be detected for the first time during pregnancy and delivery but only few cases have been reported in the literature. The first case was reported in 1914 by Nanta.^[1] The acquired variety is very rare during pregnancy as this type is mostly seen after 50 years of age.^[2] During pregnancy acquired heart block may be due to myocarditis, collagen vascular diseases, following infective endocarditis of aortic valve with root abscess or as a complication of cardiac surgery. In acquired heart block, heart rate is usually 40 or less per minute with wide QRS in ECG. They are usually symptomatic with

temporary loss of consciousness in the form of presyncope or syncope.^[3] The prognosis is generally worse, but ultimately depends on the underlying cause.^[4] Complete atrioventricular block detected for the first time during pregnancy is usually congenital. With an incidence of 1 in 20,000 live born infants^[5], congenital complete atrioventricular block is a rare disease. The aetiology is not completely understood. However, complete atrioventricular block may be isolated or combined with congenital heart diseases in up to 53% of affected individuals.^[6] It may be caused by cardiac malformations or damaged by maternal antibodies. Family history of complete atrioventricular block is one of the risk factors for congenital atrioventricular block.^[7-8]

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Case(s)

A 25-years -old women, in the 40 week of pregnancy was admitted to the hospital with labor pains since 3-4 h. The patient had no rregular antenatal check ups at a local hospital. She had no known history of heart disease or any other illnesses. She deneid any complaint before pregnancy and she had no evidence of prior ECG. She did not have syncope, palpitations and dyspnea on efforts, besides fatigue in late pregnancy. She had no known history of heart disease or any other illnesses Her parents reported rare episodes of syncope during childhood and adolescence and one more episode two year before. On examination, she was conscious oriented with pulse rate of 46-53 beats/min and blood pressure of 140/70 mmHg. Her respiratory rate was 20 per minute, and chest was bilaterally clear. Listen with the stethoscope diaphragm were heard loud first heart sound ,normal second heart sound and no murmurs. The electrocardiogram showed sinus bradycardia in which not every narrow QRS complex followed a P wave, suggestive of Complete Heart Block . Echocardiography showed normal heart. All routine laboratory tests were normal. Serum electrolyte Na+, K+, Ca2+, Mg2+ were normal. The fetal heart rate was regular, and she was having regular uterine contractions of moderate intensity every 4-5 min. On vaginal examination, os was closed with a firm cervix of around 4 cm. She had a abnormally shaped pelvis ,android pelvis. The cardiotocography had recorded regular fetal heart rate about 142 beats/minute. An doppler ultrasound showed a fetus about 3 kg with good biophysical profile .Her fetus had normal development, with no signs of growth retardation . The decision for cesarean section was taken in view of cephalopelvic disproportion. The diagnosis of complete heart block was confirmed by 12-lead ECG. After consulting together cardiologist, anaesthetist and obstetrician the woman was taken up for caesarean section under spinal anesthesia. under continuous ECG monitoring, pulse oximeter and noninvasive blood pressure. Patient and her husband were properly explained the condition. A temporary pacemaker was kept on standby to withstand any haemodynamic variations. Patient ,parents and her husband were properly explained the condition. The spinal anaesthesia was planned to keep the haemodynamics stable and drugs causing bradycardia were avoided. A healthy boy of 3350 kg with normal heart rate and normal Apgar score was delivered. The patient was transferred to the Intensive Care Unit for better continuous monitoring. During postpartum period the patient did not have any complains or syncope episodes. Postoperative period was uneventful and patient was discharged in good health on 6th postoperative day. It was strongly recommended

to her a regular follow up to cardiology department. Follow-up for next 3 months was uneventful, and she is still under regular follow-up.

Discussion

Guidelines for implantation of cardiac pacemakers have been established by a task force formed by the American College of Cardiology, the American Heart Association the Heart Rhythm Society^[9] and The European Society of Cardiology.^[10] A pacemaker is indicated in the presence of symptoms like syncope, chest pain, dyspnea, palpitations, heart rate less than 40/minute, Q-T interval prolongation, wide QRS complex, ventricular dysfunction, or heart failure.^[11] Temporary transvenous pacing is traditionally an emergency procedure to stabilize patients suffering from hemodynamically unstable bradyarrhythmia.^[12] In patients with bradycardia and indications for pacemaker implantation, the importance of shared decision making and patient-centered care is endorsed and emphasized in this guideline in which treatment decisions are based not only on the best available evidence, but also on the patient's goals of care and preferences [9]. Women with complete atrioventricular heart block without a permanent pacemaker normally receive temporary pacing for labour and birth.^[12] In women without a permanent pacemaker, temporary pacemakers have been inserted for labour and caesarean delivery probably to withstand any haemodynamic variations. Temporary pacing before delivery appeared to be beneficial for women in same cases.^[13] But review of cases for temporary pacemaker showed that the insertion of temporary pacemaker is not without risk. In patients with temporary pacemaker for bradyarrhythmias during the waiting period for permanent pacemaker implantations, bedrest might not prevent adverse events, such as cardiovascular events and complications associated with temporary pacemaker.^[14]

Complications such as irradiation, bleeding, infection or embolism, malfunction leading to sudden hemodynamic instability are common.^[15] Now guidelines for management of women with cardiac disease are established to prevent unnecessary morbidity and expense of the procedure. According to 2018 European Society of Cardiology Guidelines states that isolated congenital complete heart block has a favourable outcome during pregnancy especially with narrow QRS ^[16] temporary pacemaker is unnecessary in stable patients but recommended in selected women with symptoms due to bradycardia and syncope.

Our patient was detected with congenital complete heart block and she was almost asymptomatic. She had no suffered of heart disease or any other illnesses. Her electrocardiogram showed complete heart block with heart

rate about 42 beats/minute and QRS duration 94 ms. Her parents reported rare episodes of syncope during childhood and adolescence and one more episode two year before. During pregnancy she did not report any syncope episode except from being tired. Isolated heart block is relatively benign with narrow QRS complexes on ECG, and heart rate may increase with atropine or sympathomimetics. Patients with complete atrioventricular block who are asymptomatic, with narrow complex in ECG, ventricular rate between 40 and 60 and there is rise in heart rate with exercise or atropine, usually tolerate the pregnancy and delivery without any unfavourable events. Isolated congenital complete heart block in the mother has a favourable outcome during pregnancy, especially when the escape rhythm has a narrow QRS complex.^[17]

Our case had narrow QRS complex in ECG, responded to atropine. However, antenatal care needs to be by the pregnancy heart team with a cardiologist, anaesthetist and obstetrician, with experience in the management of high risk pregnancies. In our case the fetus had normal development, with no signs of growth retardation. Feto-maternal outcome is favourable in asymptomatic cases and in uncomplicated bradyarrhythmias without significant underlying heart disease.^[18] Our case was complicated with obstetrical complications such as cephalo-pelvic disproportion and she was taken up for caesarean section under spinal anesthesia.

On the other hand there are no specific recommendations concerning the most appropriate anaesthetic technique for immediate Caesarian Section in women with congenital complete heart block. Ideally, developing an anaesthetic plan for a high-risk cardiac patient should begin preconception or early in pregnancy so the patient can better comprehend the risks involved with pregnancy and the anaesthetic plan that may be best for the expectant mother.^[19] On our case, the atrioventricular complete heart block was diagnosed on the preoperative routine test for Caesarian Section due to cephalo-pelvic disproportion. There are quite a few anaesthetic problems in patients with complete heart block undergoing incidental surgeries. These include bradycardia, hypotension, arrhythmias, cardiac arrest or even sudden death.^[4] Studies suggested that regional anaesthesia is safe in pregnant women with cardiac disease undergoing caesarean section. Fortunately in our case was not necessary the implementation of standardized procedures for temporary pacing.

Conclusion

As suggested by our case, asymptomatic atrioventricular complete heart block in pregnancy can be managed successfully without pacemaker. However, careful monito-

ring, is necessary by the pregnancy heart team with a cardiologist, anaesthetist and obstetrician, with experience in the management of high risk pregnancies. Management of the risk for cardiovascular and obstetrical complications is difficult in pregnant women with complete heart block. Asymptomatic complete heart block in late pregnancy should be managed without pacemaker by the pregnancy heart team with a cardiologist, anaesthetist and obstetrician, with experience in the management of high risk pregnancies.

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