Case Report

Management of Pregnancy in 31-Year-Old Woman with Persistent Atrial Flutter After Cone Reconstruction for Ebstein’s Anomaly

E. Susilowati1,*, V. Tedjamulia2, R. Myrtha3
1Internship Doctor Programme, Indonesia
2Cardiovascular Department, Mardi Waluyo Hospital, Blitar, Indonesia
3Cardiovascular Department, Faculty of Medicine, Universitas Sebelas Maret, Surakarta, Indonesia

ARTICLE INFO

Article history:
Submitted 31st January 2020
Reviewed 18th February 2020
Accepted 16th March 2020
Available online March 2020

*Corresponding author: elean.eliana@gmail.com

Keywords:
Atrial Flutter
Ebstein Anomaly
Cone reconstruction

ABSTRACT

There is a growing number of mothers suffering innate heart disease which could potentially risk their gestation period. Ebstein’s anomaly refers to a rare innate disease in the heart which accounts for less than 1% of most cases of innate heart disease, and only 5% of the patients surviving beyond 50 years of age [7,8]. It is characterized by the attachment of septal and posterior leaflet to the myocardium underlying it with the dislocation of the functional tricuspid annulus downward, resulting in the dilation of atrialized portion of right ventricle (RV) and true tricuspid annulus [9]. Several methods have been tried for the surgical repairment of the patients suffering the anomaly Ebstein.

Introduction

The statistical data of mothers suffering from innate heart disease during their gestation period is rising, because more than 90% grew with it when they reached adulthood. Most common problems during gestation and delivery are bleeding, arrhythmias, failure of the heart, and infrequently maternal death [8]. Ebstein’s anomaly refers to a rare innate disease in the heart which accounts for less than 1% of most cases of innate heart disease, and only 5% of the patients surviving beyond 50 years of age [7,8]. It is characterized by the attachment of septal and posterior leaflet to the myocardium underlying it with the dislocation of the functional tricuspid annulus downward, resulting in the dilation of atrialized portion of right ventricle (RV) and true tricuspid annulus [9]. Several methods have been tried for the surgical repairment of the patients suffering the anomaly Ebstein.

© 2020 Cardiovascular Cardiometabolic Journal
E-mail: ccj@journal.unair.ac.id
The managing preparations mainly focused on the repairment of the monocusp of the tricuspid valve with discerning plication of the atrialized right ventricle. Cone reconstruction was an option to repair tricuspid valve in Ebstein anomaly patients. A leaflet tissue of 360 degrees and made to surround the real annulus is the result of this strategy. Actually, after conducting CR, the risk of arrhythmia for young patients with EA is very small. However, despite the low risk arrhythmia, a persistent arrhythmia, such as AFL could still occur after CR. Nevertheless, both AFL and pregnancy could influence to thromboembolic problems. Pregnancy, especially after first stability, raised the risk of thromboembolism up to 1.5 times. It is presumed that AFL patients have a similar risk of thromboembolism to those with atrial fibrillation. This case report focuses on elaborating suitable pregnancy management of persistent atrial flutter patients after Cone reconstruction for Ebstein’s Anomaly.

Case Presentation

31 years old woman consulted for pregnancy planning due to the history of Ebstein’s anomaly post her CR. The patient was asymptomatic and was clinically stable with blood pressure of 100/70 mmHg, heart rate of 60 bpm, respiratory rate of 18 tpm, and BMI of 25.11. Physical examination findings were TR murmur, without signs and symptoms RV failure. Electrocardiography showed counterclockwise typical atrial flutter. Electrocardiography showed counter-clockwise typical AFL (Figure 1). Chest X-Ray demonstrated normal aortic and pulmonal segment, cardiac waist (+), downward cardiac apex, cardiomegaly with CTR 69% (Figure 2). Echocardiography finding showed post Cone reconstruction, mild tricuspid regurgitation, and EF of 76%. Her laboratory tests showed no remarkable finding. Her routine medications were bisoprolol 1x 2.5 and Aspirin 1x80mg.

Figure 1. AFL type 1 patterns on ECG
Figure 2. Chest X-Ray post Cone Reconstruction

Figure 3. Echocardiography post CR

Figure 4. Sinus Rhythm Pattern on ECG before Cone Reconstruction
Past medical history showed that patient was diagnosed with ebstein's anomaly without other concomitant congenital heart disease, no recorded history of atrial flutter. Her ECG before CR was sinus rhythm with incomplete LBBB (Figure 4). Her previous chest X-ray showed cardiomegaly with
rounded cardiac apex, with CTR of 56%, and the absence of cardiac waist (Figure 5). Patient undergone successful cone procedure with mild residual TR. Echo finding demonstrated the typical features of Ebstein anomaly. Severe TR was observed with displaced insertion of posterior leaflet far toward RV apex, without coaptation. It showed RA and RV dilatation, D shaped LV, and normal LV with EF of 64% (Figure 6).

Discussion

Women with congenital heart disease require careful monitoring before, during, and after pregnancy. In pregnancy, hemodynamic and hormonal alteration could disturb the arrhythmias and the prothrombic state that contribute to thromboembolic problems [1]. Haemodynamic changes in the course of pregnancy include plasma volume raise, heart rate, and decrease of aortic pressure and systemic vascular resistance. Plasma volume increase in the peak of 32 weeks of gestation, it counts higher by 40 to 50 percent than the pre-gestation period. Rate of heart also reaches its peak at 32 weeks of pregnancy; almost higher by 20% than in the pre-gestation period [12]. Variability of heart rate was oppressed during pregnancy, this impaired autonomic nerve activity and the overload volume of the heart and operational wound could count for the presence of tachyarrhythmia during pregnancy [14]. Pulmonary artery pressure stays the same during pregnancy, because the increased volume of pulmonary blood will decrease pulmonary vascular resistance. In the first trimester, there would be rapid rise of arterial pressure; louder pulse of the jugular venous with heartbeat that splits secondly. In the late pregnancy, hypotension could be encountered because of the inferior vena cava compression by the enlarged uterus.

Hypercoagulable state also happened in pregnancy. Factors of anticoagulant such as coagulation factor I, V, VII, VIII, X and XII; von Willebrand factor; and plasma fibrinogen are activated. Fibrinolytic inhibitors are also activated, such as plasminogen activator inhibitor (PAI-1 and PAI-2) [6]. Because of this hypercoagulability, patients with arrhythmia require thorough care. AFL increased the risk of thromboembolism and may make the heart function worse [1,13]. Presumably, AFL patients have a similar risk of thromboembolism to those with atrial fibrillation [13], while risk of ischemic stroke only significantly increased at CHA2DS2-VASc score of >4, [5,15] or in CHADS2 score of more than 2 [15].

Thromboprophylaxis is needed for high risk pregnant woman, but careful consideration on the medication choice should be made. Oral anticoagulant use during pregnancy could increase the risk of bleeding in the time of delivery, also fetal deaths which could be resulted from sudden miscarriage. Heparin was the preferred option since it does not cross the placenta and relatively safe [1]. Vitamin K antagonists are still tolerable for most cases (without history of congenital heart disease) from second trimester untill 1month before expected delivery [13].

Vitamin K antagonists, that are associated with warfarin embryopathy, cross the placenta in the first trimester state, the increasing risk of CNS abnormalities, fetal and neonatal bleeding near the delivery time, and losing gestation. LMWH is recommended rather than fondaparinux in the time of gestation. There was no LMWH or UFH that was detected in the breast milk in discernible quantity or crosses the placenta [2]. Hypodermic LMWH in a maintained dose could be used in the course of the gestation’s first trimester and the last month. Dabigatran which was said to be one of the new antagonists of oral thrombin, should not be used because a high dose of fetotoxicity was detected. High risk patients with atrial fibrillation seemed to be
least effective to both single or dual antiplatelet
therapy (clopidogrel and acetylsalicylic acid) than warfarin \[15\].

Beside the increasing risk of thromboembolic in
pregnant woman and especially with overt AFL, AFL itself during pregnancy results in some serious
undesirable events. Studies showed that there is a
raise in maternal mortality and low birth weight
during pregnancy in AFL patients \[1,4\]. The increase
of maternal mortality found in 12% of pregnancies
complicated with AFL. Thus, it is proven that the
association between serious results in pregnant
women suffering structural heart disease and atrial
arrhythmias. \[11\] Lapses of suffered tachycardia, in
particular the atrial flutter, are intolerable and could
inflict fetal hypoperfusion in structural heart disease
\[15\].

Ebstein anomaly tends to be related to tricuspid
ejection, a RV that functions abnormally and a
dilated right atrium. Patients may also have multiple
accessory pathways. When there are RV size and
systolic function that are reasonable, pregnancy is
tolerated, as it is related to the absence of
significant cyanosis or arrhythmias \[14\]. As the
majority antiarrhythmic medication could be
regarded as harmful to the fetus, starting AFL
medical treatment shows complicated choice in its
practice \[2\]. Beta-blockers cross the placenta and
are associated with intra-uterine growth retardation,
neonatal respiratory depression, bradycardia, and
hypoglycemia, particularly for the exposure in 12-24
weeks of gestation \[13\].

Calcium channel blocker may also be administered
to treat atrial arrhythmias, however, since oral
Verapamil is an FDA class C, intreavenous
Verapamil is more likely to be avoided due to the
hypotensive effect. Diltiazem was associated with
teratogenicity. Digoxin is an FDA class C
medication, while Amiodarone is opposed because
it influenced the function of fetal thyroid in the
period of gestation \[1\]. The hemodynamic changes
in gestation time influence the pharmacokinetics
of the antiarrhythmic drugs, for example, a higher
dose of digoxin is required for the adequate serum
level during pregnancy, compared for those used
out of pregnancy \[4\]. Despite the weakness of each
antiarrhythmic drugs, their use should be
considered if weigh the benefit. The guideline for
the management of cardiovascular diseases in
gestation time from ESC in 2011 suggested Beta-
blocker for the first choice of rate control in
arrhythmia during pregnancy. Strenuous exercise is
also recommended because it is more effective
than Digoxin. Prophylactic medications of antiarrhythmic (propafenone, flecainide, and sotalol)
are considerable in the case of severe symptoms
despite the rate-controlling of the drugs. Those
drugs should be reconsidered in the AFL with the
presence of abnormal heart structure. \[15\]

In patients with persistent tachyarrhythmia,
including AFL, the high possibility of recurrent
arrhythmias in gestation period leads to the need of
performing a pre-pregnancy ablation \[6,15,16\]. It is
known that AFL mechanism involves a critical
isthmus of slow conduction restricted by anatomic
structures. The CTI is involved in the re-entry circuit
in typical AFL, and the implemented RF energy to
this isthmus has a chance to restore. Radiofrequency ablation proved to have high
success rate to fight AFL, which is known to have
been one of the atrial tachyarrhythmias. This results
is noted with some smaller studies presenting 91%
higher success with 1 procedure \[1,7\]. The Canadian
Heart Rhythm Society and European Heart Rhythm
Association elaborated that in both symptomatic
and recurrent AFL, it is considerable to perform
catheter ablation given the high success and
relatively low complication rates for CTI dependent
AFL. Catheter ablation for the management of CTI-dependent AFL had a significant success of 97% and a recurrence probability of 10% over the 14 months, while catheter ablation for non-CTI dependent AFL technically has higher difficulty with a wide percentage from 73 to 100 percent in the success reports [3].

It is proved that radiofrequency catheter ablation of right sided atrial tachyarrhythmia is associated with high acute rate and significant long term recurrences, and repeated radiofrequency catheter ablation of recurrent atrial tachyarrhythmia was effective in most of the patients. However, the arrhythmogenic substrate in patients with history of previous surgery proved to be complicated, and aiming for these arrhythmias needs 3D mapping that is detailed to raise the success rate of the ablation. [17]

**Conclusion**

Considering the efficacy and weighing the benefits, radiofrequency ablation with detailed 3D mapping prior to pregnancy is the most beneficial and safest choice for this 31 year-old woman with persistent typical AFL with a history Cone reconstruction due to Ebstein Anomaly.

**Acknowledgement**

The working of this paper has no conflict of interest.

**References**


