Intracardiac Lateral Tunnel Fontan by using Right Atrial Wall

Md Zahidul Islam¹, AM Asif Rahim², Kazi Abul Hasan¹, Syed Imtiaz Ahsan¹

¹Department of Paediatric Cardiac Surgery, National Institute of Cardiovascular Diseases, Dhaka, ²Department of Cardiac Surgery, Chittagong Medical College Hospital, Chittagong

Abstract

The Fontan procedure is a palliative surgical procedure used in children with complex congenital heart defects. It involves diverting the venous blood from the right atrium to the pulmonary arteries without passing through the morphologic right ventricle. A 23 years old adult male from Noakhali admitted with the diagnosis of transposition of great arteries (D-TGA) with large perimembranous ventricular septal defect (PM VSD) with severe pulmonary stenosis (PS) and single ventricle morphology in National Institute of Cardiovascular Disease (NICVD), Dhaka. He underwent Intracardiac lateral tunnel Fontan by using right atrial wall. Patient recovered uneventfully in his early postoperative period. The lateral tunnel Fontan procedure results in excellent long-term outcome even when used in patients with diverse anatomic diagnoses. The incidence of atrial tachyarrhythmia is low and mainly depends on the underlying cardiac morphology and preoperative arrhythmia. The good long-term outcome after an intracardiac lateral tunnel Fontan procedure should serve as a basis for comparison with other surgical alternatives. But using right atrial wall as a baffle is rare and done first time in Bangladesh.


Key Words: Congenital heart disease, transposition of great arteries, Lateral tunnel Fontan, Right atrial wall.

Introduction

The Fontan procedure is a palliative surgical procedure used in children with complex congenital heart defects. It involves diverting the venous blood from the right atrium to the pulmonary arteries without passing through the morphologic right ventricle; i.e., the systemic and pulmonary circulations are placed in series with the functional single ventricle. It was initially described in 1971 by Dr. Francois Marie Fontan from Bordeaux, France, and later in 1973 by Dr. Guillermo Kreutzer from Buenos Aires, Argentina separately as a surgical treatment for tricuspid atresia. The Fontan procedure is used in pediatric patients who possess only a single functional ventricle, either due to lack of a heart valve (e.g. tricuspid or mitral atresia), an abnormality of the pumping ability of the heart (e.g. hypoplastic left heart syndrome or hypoplastic right heart syndrome), or a complex congenital heart disease where a bi-ventricular repair is impossible or inadvisable. The single ventricle is doing nearly twice the expected amount of work because it has to pump blood for the body and lungs. Patients typically present as neonates with cyanosis or congestive heart failure. Fontan completion is usually carried out when the patient is 2-5 years of age, but is also performed before 2 years of age.¹⁻³

There are three variations of the Fontan procedure 1) Atrio-pulmonary connection (the original technique), 2) Intracardiac total cavopulmonary connection (lateral tunnel) and 3) Extracardiac total cavopulmonary connection. The Fontan is usually done as a two staged repair. The first stage, also called a Bidirectional Glenn procedure or Hemi-Fontan (Kawashima procedure), involves redirecting oxygen-poor blood from the top of the body to the lungs. That is, the pulmonary arteries are disconnected from their existing blood supply (e.g. a shunt created during a Norwood procedure, a patent ductus arteriosus, etc.). The superior vena cava (SVC), which carries blood returning from the upper body, is disconnected from the heart and instead redirected into the pulmonary arteries. The inferior vena cava (IVC), which carries blood returning from the lower body, continues to connect to the heart. The second stage, also called Fontan completion, involves redirecting the blood from the IVC to the lungs. At this point, the oxygen-poor blood from upper and lower body flows through the lungs without being pumped (driven only by the pressure that builds up in the veins).
This corrects the hypoxia and leaves the single ventricle responsible only for supplying blood to the body.\textsuperscript{2,4}

He also complained of progressive breathlessness that increases in severity for the last few years and becomes more severe for the last 6 months with mild exertion. On examination, his body build was below average, small for age, decubitus on choice, cyanosis and clubbing were present, neck vein not engorged and there were no leg oedema or ascites. Pulse was 120 b/min, regular, blood pressure-90/60 mm of Hg, respiratory rate 22/min. On palpation, apex beat is slightly shifted laterally about 1cm from midclavicular line. On auscultation, first and second heart sounds were normal, there were a pansystolic murmur along left 3\textsuperscript{rd} or 4\textsuperscript{th} space with no radiation. Chest X-ray P/A view revealed an enlarged cardiac shadow with congested lungs more.

\textbf{Case report:}

A 23 years old adult young male admitted with bluish discoloration of whole body more prominent in nail, lips, nose and ear lobule since birth along with repeated bouts of respiratory tract infection. Intracardiac Lateral Tunnel Fontan by using Right Atrial Wall

\textbf{Fig.-1: Preoperative chest X-ray and Echocardiography.}
prominent in hilar region. Echocardiography revealed D-TGA. PM VSD, severe PS and single ventricle. CT pulmonary angiogram revealed severely stenosed pulmonary infundibulum and valve although main pulmonary artery (MPA), right pulmonary artery (RPA), left pulmonary artery (LPA), Aorta, SVC, IVC were of normal size with no evidence of other congenital anomalies. In Heart there were three chambered heart with single ventricle. Both the inlets were normal. But double outlets single ventricle was detected with large atrial septal defect (ASD) secondum. After detail counseling patient underwent open heart surgery. A median sternotomy was performed in all patients. The thymus gland was removed completely and pericardium was opened. The right and left superior vena cavae were dissected and mobilized upto the inominate vein. Also the right and left pulmonary arteries were dissected and mobilized. Azygos vein was doubly ligated and divided. Though intra-cardiac or lateral tunnel method is the ‘older’ method (introduced in 1987) and the extra-cardiac method was more recently developed (1990); we like to perform the intra-cardiac method as our preferred option. The lateral tunnel/intra-cardiac method involves sewing a piece of plastic inside the right atrium to route all blood from the lower part of the body but here in this case we use patient’s own right atrial wall instead of PTFE tube (via the inferior vena cava).
to the lungs and thereby performed intracardiac lateral tunnel Fontan by using patient’s own right Atrial wall. Patient recovered uneventfully and discharged on 10th Post operative day.

Discussion:
After Fontan completion, blood must flow through the lungs without being pumped by the heart. Therefore, children with high pulmonary vascular resistance may not tolerate a Fontan procedure. Often, cardiac catheterization is performed to check the resistance before proceeding with the surgery. This is also the reason a Fontan procedure cannot be done immediately after birth; the pulmonary vascular resistance is high *in utero* and takes months to drop.2

Regarding post operative complication In the short term, children can have trouble with pleural effusions (fluid building up around the lungs). This can require a longer stay in the hospital for drainage with chest tubes. To address this risk, some surgeons make a fenestration from the venous circulation into the atrium. When the pressure in the veins is high, some of the oxygen-poor blood can escape through the fenestration to relieve the pressure. However, this results in hypoxia, so the fenestration may eventually need to be closed by an interventional cardiologist. In the long term postoperative complication, children can have trouble with atrial flutter and atrial fibrillation because of scarring in the atrium, especially if the connection of the IVC to pulmonary arteries involved an Intracardiac baffle (instead of an extracardiac conduit). This sometimes requires treatment such as radiofrequency ablation. Other long-term risks include protein-losing enteropathy and chronic renal insufficiency, although understanding of these risks is still incomplete. Some patients require long-term anticoagulant. Four case reports of hepatocellular carcinoma following the Fontan procedure were reported by physicians at the Mayo Clinic. This group of patients may warrant screening for this malignancy, especially if the alpha-fetoprotein level is elevated. The Fontan procedure is palliative — not curative — but in many cases it can result in normal or near-normal growth, development, exercise tolerance and good quality of life.7 In 20-30% cases, patients will eventually require heart transplantation.4
of medications given to patients that underwent intra-cardiac vs. extra-cardiac Fontan. Though this probably attributable to the hospital itself and how they treat their patients, most of the patients were on aspirin (94%) and there were no other differences between the type of frequency of medications taken between patients that underwent intra-cardiac or extra-cardiac method. No difference in heart rhythm problems between patients that had undergone intra-cardiac vs. extra-cardiac Fontan at 3 years after the surgery. This is somewhat surprising that 15% of patients that had undergone the intra-cardiac Fontan had heart rhythm issues (sinus node dysfunction) whereas more (28%) patients that had underwent the extra-cardiac Fontan had heart rhythm issues. Although this is not statistically different, this is opposite than what would be expected. Two patients underwent permanent pacemaker implantation (1 lateral tunnel and 1 extra-cardiac method) and in one case for slow junctional rhythm. No difference in the post-operative blood pressure in various parts of the atrium and in the Fontan pathway (“Fontan pressure” and transpulmonary gradient) between patients that underwent the intra-cardiac vs. extra-cardiac Fontan. The authors provide brief discussion how these pressures can be predictive of early Fontan failure but they found no difference between these two methods in the first 24 hours after the Fontan.2,4

Kaplan-Meier estimated survival was 93% at 5 years and 91% at 10 years, and freedom from failure was 90% at 5 years and 87% at 10 years. Freedom from new supraventricular tachyarrhythmia was 96% at 5 years and 91% at 10 years; freedom from new bradyarrhythmia was 88% at 5 years and 79% at 10 years and ultimately high rates of early survival 3-5 years after the Fontan for both the intra-cardiac (97%) and extra-cardiac (91%) methods.2,4

Conclusion:
The lateral tunnel Fontan procedure results in excellent long-term outcome even when used in patients with diverse anatomic diagnoses. The incidence of atrial tachyarrhythmia is low and mainly depends on the underlying cardiac morphology and preoperative arrhythmia. The good long-term outcome after an intracardiac lateral tunnel Fontan procedure should serve as a basis for comparison with other surgical alternatives. Here we use patient’s own right atrial wall instead of Gore-tex tube which is completely new concept in our situation and no such evidence based study on use of right atrial wall in intracardiac lateral tunnel Fontan has been performed so far. Our patient recovered uneventfully so this may be an acceptable procedure in near future in this operation instead of using Gore-tex tube.

Conflict of Interest - None.

References: