

Current Therapy for Hypoplastic Left Heart Syndrome and Related Single Ventricle Lesions

ABSTRACT: Universally fatal only 4 decades ago, the progress in the 3-stage palliation of hypoplastic left heart syndrome and related single right ventricular lesions has drastically improved the outlook for these patients. Although the stage II operation (hemi-Fontan or bidirectional Glenn) and stage III Fontan procedure have evolved into relatively low-risk operations, the stage I Norwood procedure remains one of the highest-risk and costliest common operations performed in congenital heart surgery. Yet, despite this fact, experienced centers now report hospital survivals of >90% for the Norwood procedure. This traditional 3-stage surgical palliation has seen several innovations in the past decade aimed at improving outcomes, particularly for the Norwood procedure. One significant change is a renewed interest in the right ventricle-to-pulmonary artery shunt as the source of pulmonary blood flow, rather than the modified Blalock-Taussig shunt for the Norwood. The multi-institutional Single Ventricle Reconstruction trial randomly assigned 555 patients to one or the other shunt, and these subjects continue to be followed closely as they now approach 10 years postrandomization. In addition to modifications to the Norwood procedure, the hybrid procedure, a combined catheter-based and surgical approach, avoids the Norwood procedure in the newborn period entirely. The initial hybrid procedure is then followed by a comprehensive stage II, which combines components of both the Norwood and the traditional stage II, and later completion of the Fontan. Proponents of this approach hope to improve not only short-term survival, but also potentially longer-term outcomes, such as neurodevelopment, as well. Regardless of the approach, traditional surgical staged palliation or the hybrid procedure, survivals have vastly improved, and large numbers of these patients are surviving not only through their Fontan in early childhood, but also into adolescence and young adulthood. As this population grows, it becomes increasingly important to understand the longer-term outcomes of these Fontan patients, not only in terms of survival, but also in terms of the burden of disease, neurodevelopmental outcomes, psychosocial development, and quality of life.

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Hypoplastic left heart syndrome (HLHS) is characterized by hypoplasia of the left ventricle and systemic outflow tract obstruction.¹ Other common related functional single right ventricle (FSRV) lesions include double-outlet right ventricle with mitral atresia and unbalanced atrioventricular septal defect. Without intervention, HLHS and related FSRV lesions are essentially uniformly fatal. Since the first description of the Norwood procedure in 1981, great strides have been made in the treatment of the HLHS and related lesions.² Over the past 35 years, hospital survival for the Norwood procedure has improved from 0% to >90% in experienced centers.³

CURRENT THERAPY FOR HLHS AND OTHER FUNCTIONAL SINGLE RIGHT VENTRICLE LESIONS

Traditional surgical staged palliation consists of the Norwood procedure at birth, a stage II superior cavopulmonary connection, generally performed at 4 to 6 months of age, and a completion Fontan at 18 to 48 months of age. More recently, the hybrid procedure has been proposed as an alternative to the initial Norwood procedure. Although the physiology remains the same after a Norwood or a hybrid procedure, the hybrid is less invasive and does not require as extensive an intervention in the neonatal period. The trade-off is a more involved second-stage operation, and more intensive follow-up and more frequent interventions in the period between stage I and stage II. The following state-of-the-art review will discuss the current therapy for HLHS and related FSRV lesions, and the intermediate-term outcomes for the Fontan procedure, as well.

TRADITIONAL STAGED SURGICAL PALLIATION

The requirements for the Norwood procedure have not changed from Dr Norwood's initial description in 1980. These requirements are (1) unobstructed systemic outflow from the single right ventricle to a reconstructed aorta, (2) unobstructed pulmonary venous return into the right atrium, and (3) controlled pulmonary blood flow (PBF).⁴ Although the first 2 requirements have remained largely unchanged, there are now 2 acceptable options for the source of the measured PBF.

In the classic Norwood procedure, a modified Blalock-Taussig shunt (MBTS) provides PBF from the innominate or subclavian artery to the pulmonary arteries via a polytetrafluoroethylene tube. Because of the placement of the MBTS downstream of the neo-aortic valve, there is continuous forward flow from the systemic to the pulmonary circulation in both systole and diastole following the Norwood procedure. Coronary steal may result as diastolic retrograde flow occurs in both the coronary arteries and descending aorta.⁵ Because 70% to 80%

of coronary flow occurs during diastole, coronary insufficiency attributable to this coronary steal may play an important role in the recognized incidence of mortality between the stage I and II operations. Indeed, decreased coronary arterial flow and oxygen delivery both at rest and after administration of adenosine (simulating exercise) have been shown to be significantly decreased in patients after the Norwood procedure with MBTS in comparison with patients after anatomic repair of other congenital cardiovascular malformations.⁶

Recently, there has been renewed enthusiasm for the right ventricle-to-pulmonary artery shunt (RVPAS) as a source of PBF for the Norwood procedure. Although the initial description was by Norwood and colleagues in 1981² and first current era reports were by Kishimoto and colleagues in 1999,⁷ the RVPAS was popularized by Sano in a number of articles in the early 2000s,⁸⁻¹⁰ and is often referred to as a Sano shunt or a Sano modification of the Norwood procedure. The RVPAS eliminates the diastolic runoff and coronary arterial steal associated with the MBTS.⁵ The disadvantage of the RVPAS is the need to perform a ventriculotomy, with the potential risk to ventricular function and arrhythmia generation.

Small, nonrandomized case series comparing the 2 sources of PBF demonstrated conflicting results. Some reports showed improved hemodynamics, branch pulmonary artery growth, and decreased mortality.¹¹⁻¹⁵ Others centers, in general, those with favorable results with the MBTS, failed to show a benefit with the RVPAS.¹⁶⁻¹⁸

SINGLE VENTRICLE RECONSTRUCTION TRIAL

Because of this uncertainty about the impact of the source of PBF on the Norwood procedure, the National Heart, Lung, and Blood Institute-funded Pediatric Heart Network (www.pediatricheartnetwork.com) SVR trial (Single Ventricle Reconstruction) was undertaken starting in 2005 to compare the MBTS and the RVPAS.¹⁹ The SVR trial enrolled neonates with HLHS or related FSRV lesions from 15 North American centers. The primary outcome was a combined end point of death or cardiac transplantation 12 months after randomization. Secondary outcomes included Norwood and stage II postoperative course, right ventricular function at the stage II operation, branch pulmonary artery dimensions at stage II, and neurodevelopmental outcomes at 14 months of age.

The SVR trial enrolled 555 eligible neonates of which 549 subjects (275 MBTS and 274 RVPAS) underwent a Norwood procedure and were included in the analysis. At the 12-month primary end point, subjects in the RVPAS group had a 74% transplantation-free survival versus 64% for the MBTS group ($P=0.01$).²⁰ However, when all available follow-up data (mean 32 ± 11 months) were analyzed at the conclusion of the trial, a statistically sig-

nificant transplant-free survival advantage of the RVPAS cohort could no longer be shown ($P=0.06$). Anatomic subtype (aortic atresia, aortic stenosis, mitral atresia, or mitral stenosis) did not impact outcome. The right ventricular function by echocardiography was similar in both cohorts at 14 months. The RVPAS group required more unplanned surgical and interventional catheter-based cardiovascular interventions to address stenosis in the shunt, branch pulmonary arteries, or neo-aorta in comparison with the MBTS group ($P=0.003$).

A more in-depth competing risks analysis revealed that mortality was separated into 2 phases: early/acute and late/constant.²¹ The early/late phase inflection point occurred at 6 to 7 months, suggesting that risk factors for mortality were a function of time. The early phase, which is more closely associated with the Norwood procedure, showed obstructed pulmonary venous return, single ventricle diagnosis other than HLHS in comparison with HLHS subtypes aortic stenosis/mitral stenosis and aortic atresia/mitral atresia, lower socioeconomic status, and smaller ascending aorta to be risk factors for death. The highest hazard ratio for mortality in the early/acute phase was obstructed pulmonary venous return. The risk factors for mortality during the late/constant phase were lower gestational age and the presence of a genetic syndrome. In reviewing the influence of various subgroups, an interesting interaction between HLHS anatomic subtype, prematurity, and shunt type emerged. The benefit of the RVPAS at a mean follow-up of 2.7 ± 0.9 years was largely confined to full-term neonates with aortic atresia, in whom mortality was approximately one-third that of similar subjects undergoing an MBTS. Interestingly, preterm neonates with a patent aortic valve had a better transplant-free survival with an MBTS.

The same publication analyzed risk factors for transplantation.²¹ Cardiac transplantations occurred in 19 subjects in the SVR trial. Multivariable analysis revealed decreased pre-Norwood procedure right function as measured by lower fractional area change, HLHS versus other single ventricle diagnosis, and number of surgeries before the Norwood procedure as risk factors for transplantation.

FOLLOW-UP RESULTS FROM THE SVR TRIAL

Since the initial SVR publications in 2010, the 3-year and 6-year results have been analyzed. At 3 years, the combined death and cardiac transplantation rates for the RVPAS versus MBTS groups were 33% versus 39% ($P=0.14$).²² When all available data were examined by Kaplan-Meier analysis (mean follow-up 4.4 ± 1.0 years), there was also no difference between groups (log rank $P=0.11$). Overall, there were 100 deaths and 10 transplantations in the MBTS cohort and 86 deaths and 11 transplantations in the RVPAS group. For subjects surviv-

ing to 1 year, there were 10 events (9 deaths, 1 transplant) in the MBTS group and 25 events (18 deaths, 7 transplantations) in the RVPAS group. This loss of benefit to transplant-free survival for the RVPAS is also illustrated by the differing nonproportional hazards associated with the 2 sources of PBF over time. The RVPAS group had a lower hazard before 5 months of age (the mean age for stage II) (hazard ratio, 0.63; 95% confidence interval, 0.45–0.88), no difference from 5 months to 1 year, and a higher hazard after 1 year (hazard ratio, 2.22; 95% confidence interval, 1.07–4.62). As reflected by this increasing hazard over time, the transplantation-free survival conditional on surviving to 1 year was worse in the RVPAS group (log-rank $P=0.03$). This nonproportional hazard was seen despite no differences between stage II surgery morbidity and growth, neurodevelopmental status, right ventricular ejection fraction, and tricuspid regurgitation grade at 14 months of age or differences in the rates of Fontan completion, medical events, and morbidities. Subjects receiving the RVPAS had more interventional catheterization laboratory visits and catheter-based interventions at 3 years (both $P<0.001$), a hazard risk which increased over the study time period.

The 6-year results have been presented as an abstract, with the article pending.²³ Although the point averages continued to reflect a difference favoring the RVPAS (combined death/transplantation rate, 36%) in comparison with the MBTS (41%), the number of subjects are not sufficient to demonstrate a statistically significant difference between the 2 groups (log-rank $P=0.13$, Figure 1). Similar to the 3-year results, RVPAS subjects had a higher incidence of any catheter intervention (0.38 versus 0.23 interventions/patient-year, $P<0.001$), including balloon angioplasty ($P=0.014$), stent ($P=0.009$), and coiling ($P<0.001$). Morbidities were similar between groups. Common morbidities included pacemaker 3%, thrombosis 16%, stroke 7%, seizures 13%, protein-losing enteropathy 3%, and plastic bronchitis 0.5%. Functional performance revealed New York Heart Association class I in 71% of subjects, class II in 21%, class III in 3%, and class IV in 5%. Risk factors for death or transplantation were low birth weight (<2500 g), higher degree of pre-Norwood tricuspid regurgitation (>2.5 mm jet width), lower surgeon Norwood procedure volume, preterm birth (<37 weeks), and combined aortic atresia and preterm birth (all $P<0.01$). By echocardiography, right ventricular ejection fraction was similar in the RVPAS and MBTS groups (46% in both groups, $P=0.9$).

NEURODEVELOPMENTAL OUTCOMES

Because survivals have improved for even the most complex congenital heart malformations, such as HLHS, attention has turned to longer-term outcomes, including neurodevelopment. Neurodevelopmental outcomes were analyzed at 14 months and again at 3 years for the SVR

trial.²⁴ Three hundred twenty-one subjects were available for neurodevelopmental assessment at 14 months following randomization in the SVR trial. The Psychomotor Development Index and Mental Development Index of the Bayley Scales of Infant Development-II were administered. Overall, the average scores for the Psychomotor Development Index and the Mental Development Index were below normal means. Independent risk factors for a lower Psychomotor Development Index score included lower birth weight, the center where the Norwood was performed, longer length of stay following the Norwood procedure, and more complications between the Norwood procedure and 12 months of age. Independent risk factors for a lower Mental Development Index score included lower birth weight, center, genetic syndrome, lower maternal education, longer mechanical ventilation after the Norwood procedure, and more complications after the Norwood discharge to 12 months of age. It is notable (and perhaps frustrating) that many of these risk factors are patient related rather than easily modifiable.

Subjects from the SVR trial were also analyzed for neurodevelopmental outcome at 3 years. The Ages and Stages Questionnaire (ASQ) was used to measure neurodevelopment. The ASQ contains 5 domains: Communication, Gross Motor, Fine Motor, Problem Solving, and Personal/Social Interaction. Delay in any of these domains is defined as <2 standard deviations below the mean. All ASQ domains for the study subjects demonstrated means lower than the reference population. The percentage of subjects classified as delayed was 20% for

the Communication Scale, 30% Gross Motor, 35% Fine Motor, 24% Problem Solving, and 17% Personal/Social. Overall, at least 51% of the cohort demonstrated delay in at least 1 domain. On multivariable analysis, risk factors that were associated with >1 abnormal ASQ domain were census block with a greater percentage below the federal poverty level, vision or hearing problems, male sex, feeding therapy at 2 years of age, and more complications between the Norwood procedure and 3 years.

Behavior was measured with the Parent-Report Behavior Assessment System for Children, Second Edition, which qualifies adaptive and problem behaviors at home and at school. The Parent-Report Behavior Assessment System for Children, Second Edition includes 14 subscales that are used to generate 4 composite scores: Externalizing Problems, Internalizing Problems, Behavioral Symptoms Index, and Adaptive Skills. At-risk or frankly abnormal behavioral composite scores were seen in 7.8% to 21.9%.

Quality of life was assessed by using both the Generic and Cardiac Modules of the Pediatric Quality of Life Inventory (PedsQL). The PedsQL measures physical, emotional, social, and school performance. Subjects in the SVR trial had lower mean scores in each of the domains of the Generic PedsQL. The scores for the cardiac-specific module were not reported.

The Functional Status, 2nd Edition Revised is a parent-reported questionnaire assessing functional status for children with chronic health problems. The Functional Status, 2nd Edition Revised demonstrated lower total scores ($P<0.001$) and lower activity scores ($P<0.001$) than the reference population.

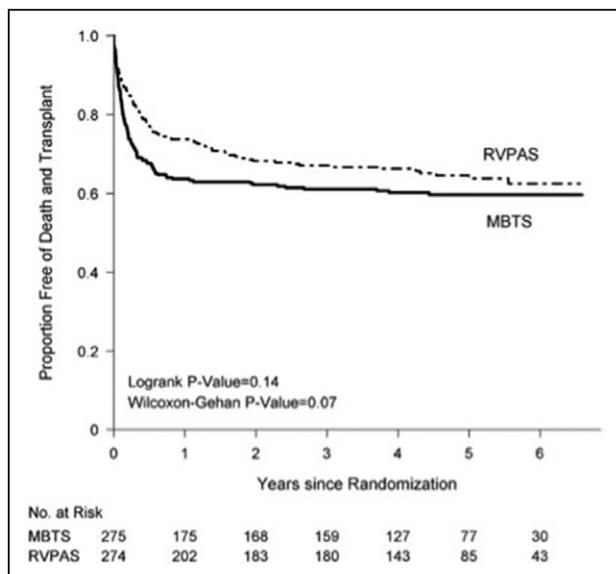


Figure 1. Comparison of the shunt types by intention-to-treat analysis in their freedom from the composite end point of death or cardiac transplantation (ie, transplantation-free survival).

MBTS indicates modified Blalock-Taussig shunt; and RVPAS, right ventricle-to-pulmonary artery shunt.²²

PRACTICE PATTERN VARIATION

Perhaps one of the more informative articles from the SVR trial, with respect to the ability to alter outcomes, addressed the practice pattern variations seen between centers in the SVR trial.²⁴ Although gestational age, birth weight, and proportion of subjects with a diagnosis of HLHS were similar across centers, there was a wide variation in preoperative, intraoperative, and postoperative management. The range of these differences were striking and included such basic elements as use of preoperative intubation, cardiopulmonary bypass time, route of feeding at discharge, and use of a home-monitoring program. Although variation in care is not unto itself problematic, the mortality following the Norwood procedure varied from 7% to 39% across centers. It seems likely that adoption of best practices would positively influence this wide range in the mortality rate.

THE HYBRID APPROACH

Hypothesized as a less invasive approach to palliate neonates with HLHS, in 1993 Gibbs et al²⁵ reported the

first stenting of the arterial duct combined with bilateral pulmonary arterial banding (bPAB) and atrial septostomy as an alternative procedure to the Norwood operation. However, based on the first experiences in 8 patients, the same group from Leeds, England did not further recommend ductal stenting as a palliation of newborns with duct-dependent systemic blood flow.²⁶ Later, in Giessen, Germany, a more successful collaborative surgical-interventional approach was achieved by surgically performed bPAB via a brief open-chest procedure, followed by a second elective percutaneous transcatheter arterial duct stenting, which was combined with atrial septum (interatrial septum) manipulation, if necessary.²⁷ Subsequently, in 1998, Hakan Akintuerk performed the first comprehensive stage II approach. Postnatal bPAB placement saved the life of the newborn, who had been admitted in cardiogenic shock 5 months earlier. Early promising results of a hybrid stage I, followed by successful comprehensive stage II and Fontan completion were published in 2002.²⁸ Independent from the experiences in Giessen, early in the new millennium, a second group of collaboratively minded interventionists and surgeons in Columbus, Ohio focused on a 1-step hybrid procedure, which entailed placing a stent within the duct by transpulmonary access immediately after bPAB during the same open-chest approach. Percutaneous manipulation of the interatrial septum was delayed until just before the palliated patient was discharged home.²⁹ In both centers, the hybrid procedure replaced the Norwood as the preferred first stage of palliation.^{29,30} In Sao Paulo, Brazil, the Columbus hybrid variant also supplanted the use of the Norwood procedure.³¹ Subsequently, the group in Toronto described an additional variant of the hybrid approach for HLHS patients with aortic atresia.³² This approach added a reverse MBTS after bPAB placement, but before ductal stent placement. Based on an institutional risk score, a recommendation is made as to whether the patient is best served by a hybrid or a Norwood procedure. Over time, multiple centers worldwide have established hybrid programs for high-risk cases.³³ In Japan³⁴ and London,³⁵ the hybrid procedure is used in some critical newborns as a palliative step for bridging newborns to a delayed Norwood procedure beyond the neonatal period with or without ductal stenting.

Irrespective of the exact technical approach, a successful hybrid requires the same 3 objectives as originally proposed by Norwood for surgical palliation: adequate systemic blood flow, sufficient protection of the pulmonary arterial circulation, and unrestrictive flow of the pulmonary venous system. After the learning curves and the acquisition of the requisite technical skills, an elective hybrid procedure treating HLHS can be performed with extremely low mortality.^{29,30}

In addition to a complete stage I procedure, the individual components of the hybrid approach can be used in specific situations. Percutaneous duct stenting is the

treatment of choice for prostaglandin-refractory ductal obstruction.³⁶ Balloon atrial septostomy with or without stent placement is also the preferred method to treat severe hypoxemia attributable to significantly obstructed or intact atrial septum. Obstructed pulmonary venous blood flow associated with a total anomalous pulmonary venous return can be palliated by transcatheter techniques.³⁷ Neonates who present in extremis because of low systemic perfusion can be resuscitated with bPABs.³⁸

Considering the wide variation of different techniques described for the hybrid approach, and the variable results, as well, it can be challenging to understand the risks and benefits. The primary benefit of a hybrid approach is as a minimally invasive alternative to a Norwood procedure with a less acute postprocedural hospital course.^{29,30} In addition to the immediate advantages, by postponing the open surgical procedure until after the neonatal period, it has been postulated that there may be potential benefits to long-term outcomes, such as neurodevelopment.^{39,40}

The current state of the art of the hybrid approach in HLHS might be summarized as follows:

- Prenatal diagnosis of HLHS is optimal for elective postnatal treatment by a trained and prepared surgical-interventional team.
- Ductal patency is maintained open with low-dose prostaglandin E1 (2–5 ng/kg-min), which decreases the risk of apnea and the rapid decrease of pulmonary vascular resistance (Rp).
- If there is no need for immediate atrial septum manipulation, pulmonary blood flow should be immediately controlled by surgically fashioned bPABs. As described by Galantowicz et al,²⁹ the pulmonary arterial banding can easily be constructed by cutting a 3.0- or 3.5-mm polytetrafluoroethylene tube graft to a small 1- to 2-mm strip, which are used in patients with body weight above or below 3 kg, respectively. After median sternotomy and partially opened pericardium, polytetrafluoroethylene bands are sewn around the left and right pulmonary artery (Figure 2). We recommend against manipulations such as transpulmonary ductal stenting or placement of a reverse MBTS, which may result in blood loss, circulatory instability, or the need for inotropic infusions. Concurrent use of additional interventions at this time have been associated with reported mortality rates between 10% and 25%.^{32,35} Conversely, the surgical approach in Giessen is simply focused on bPAB, and the surgical procedure mortality has been <1%.²⁹ In addition, time under anesthesia with slightly permissive hypercapnic ventilation can be kept short, followed by immediate spontaneous breathing and early extubation. The heart rate and systemic vascular resistance (Rs) are perioperatively controlled by continuous

infusion of the α -2 agonist clonidine, followed by early oral application of a β 1-receptor blocker (bisoprolol) together with tissue angiotensin-converting enzyme inhibitor (lisinopril) without jeopardizing the coronary perfusion pressure.⁴¹

- In all elective patients, ductal stenting is performed as an independent percutaneous approach by transfemoral vascular access. The enormous development of stent technology in Europe resulted in a CE mark for stents indicated for ductal stenting in newborns. The stent design, based on a self-expanding nitinol alloy (Sinus-SuperFlexDS, OptiMed Inc, Karlsruhe, Germany) allows duct stenting by advancing loaded stents with widths from 7 to 9 mm and lengths between 12 and 24 mm deliverable through a 4F vascular sheath. Therefore, stent placement within the duct can be performed by femoral venous or preferentially by femoral arterial access. Before stent placement, the extremely variable anatomy of the junction of the duct to descending aorta needs to be precisely delineated by brief episodes of angiography using minimal volume hand injections of contrast medium. The entire length of the duct needs to be fully covered (Figure 3). A further advantage of percutaneous stent placement is the optimal visualization of the isthmus region to determine whether an additional intervention to maintain retrograde flow to the proximal aorta is necessary. Ballooning of an obvious reverse coarctation should be performed before duct stenting, and, if stent placement becomes necessary, it should be performed after stent placement within the duct.
- To guarantee unrestricted flow of pulmonary venous return into the right atrium, interventions on the atrial septum in advance may be of benefit.

The unrestricted atrial-level left-to-right flow and ductal right-to-left flow, combined with a protected pulmonary circulation, need to be maintained until the comprehensive stage II, which is performed in most patients at an age of 4 to 5 months. The fragile interstage I period, between the hybrid and the comprehensive stage II, requires not only a successful initial hybrid palliation with a well-balanced circulation, but also appropriate and assiduous follow-up monitoring, which includes detailed instructions for the parents. Medications are titrated to balance the systemic-to-pulmonary vascular resistance ratio, and the ratio of oxygen consumption-to-demand, as well. In regard to the important role of the parents, the cardiovascular medications need to be safe, well understood, and easily administered. The respiratory rate while sleeping, oral intake, and body weight are monitored closely by the parents.^{30,41} In addition, interstage management requires close outpatient evaluation by an experienced pediatric cardiologist to detect any hemodynamic imbalance before the development of any crisis.³⁰ Based on this close outpatient monitoring, interventional procedures should

be aggressively pursued for not only detected hemodynamic abnormalities, but also any suspected issue during the interstage period. Interventions to address obstructions in the stented duct, atrial septum, or even of the descending aortic arch can be effectively performed by using various catheter techniques.³⁰ In summary, this combined parents-physician monitoring program has been able to significantly reduce the interstage morbidity and, in particular, mortality to <5%, for HLHS, which is comparable to that described for interstage surveillance programs following the Norwood procedure.⁴²

Precomprehensive stage II catheterizations have generally been replaced by magnetic resonance imaging in sedated, spontaneously breathing infants, similar to our practice for all elective heart catheterizations. Summarizing the 2-center experience for the comprehensive stage II in Giessen⁴³ and Columbus,⁴⁴ accounting for >200 patients, the comprehensive stage II can be performed with a mortality of <5%, despite the early learning curve being included in these reports. The surgical approach consists of bilateral pulmonary debanding, ductal stent removal, atrial septectomy, reconstruction of the aortic arch and bidirectional cavopulmonary connection, which combines components of the Norwood and stage II procedures into 1 operation, thereby avoiding an extensive open-heart operation in the neonatal period.³⁹ In addition, anesthetic and intensive care strategies focusing on early extubation and strategies to reduce oxygen consumption facilitate care in the immediate postoperative period. The Achilles heel of the comprehensive stage II is the variable fate of the left pulmonary artery with respect to stenosis. Refinements, such as intraoperative stenting, are warranted to improve the current surgical techniques.^{30,43} Following the comprehensive stage II, the completion Fontan circulation can be performed with the typically expected low mortality. At dedicated centers, an estimated 10-year survival of 78% can currently be achieved by the hybrid approach in an unselected population of neonates.⁴⁴

LATE FONTAN OUTCOMES

The outcomes of patients who undergo the Fontan procedure are by far better than initially expected.^{45,46} The current variation in outcomes in various reports is related to variation in the indications for Fontan completion: centers with restrictive indications will present higher survival than those with more permissive indications.⁴⁷⁻⁵¹ Technical variations have also been shown to impact outcomes, such that procedures performed in more recent decades resulted in better early and late outcomes.⁴⁸⁻⁵² Therefore, the interpretation of any outcomes after Fontan surgery should take into consideration the precise technique used and the era when it was performed. Specific long-term outcomes after Fontan for patients born with HLHS have rarely been reported. Only a limited

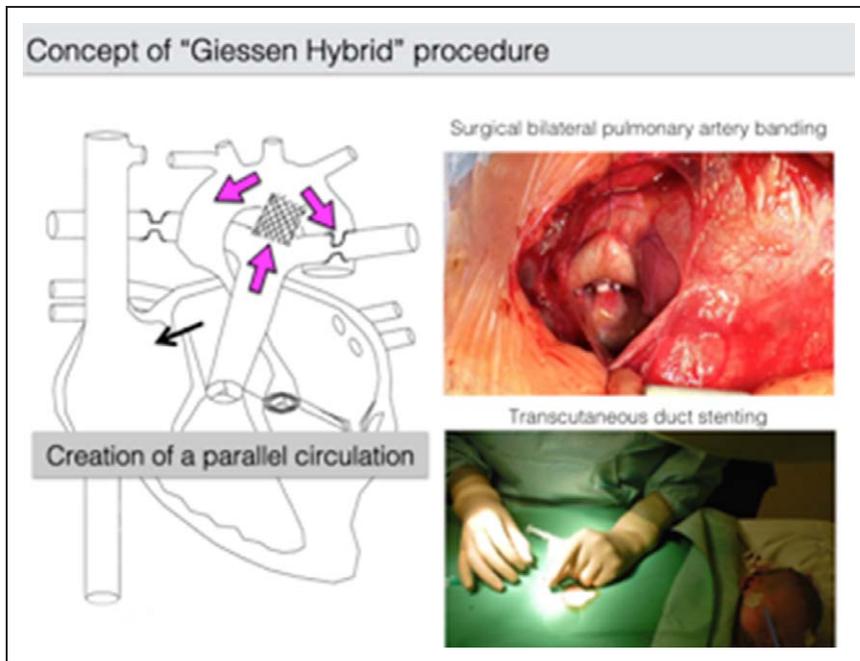


Figure 2. Line drawing (left) of bilateral pulmonary artery banding and ductal stent, with flow characteristics shown by purple arrows, as well as the left-to-right shunting at the atrial communication (black arrow) (Akintuerk et al²⁸).

The open-chest approach with right pulmonary banding (Right upper) and percutaneous heart catheterization by femoral access for ductal stenting after surgical bPAB (Right lower). bPAB indicates with bilateral pulmonary arterial banding.

number of studies reported outcomes after Norwood surgery beyond 15 years.^{53–56} In the recent years, large series of follow-up of outcomes after Fontan into the third decade have been published, and these will be the best guide to give long-term predictions to the families requiring counseling.^{47,49,57,58} The outcomes of patients with HLHS who have reached Fontan completion seem to be somewhat equivalent to the remaining population in the first 2 decades. We should therefore feel entitled to use current available data from the whole population of patients with a Fontan circulation to draw current predictions of late outcomes for patients born with HLHS.

SURVIVAL

Current expectations point toward a 30-year survival of 85% for those operated on today with, at this early stage, only minor differences in survival between those with and without previous Norwood surgery.^{59–62} Hospital mortality of Fontan completion has also decreased consistently and is identified from large multicentric data to be between 1% and 2%.^{63,64} From longitudinal data from birth, it therefore seems that two-thirds of the patients born with HLHS and operated on in best centers may hope to reach adulthood.^{53–56}

The overarching discriminant in late survival is the type of Fontan. Patients with total cavopulmonary connections, the lateral tunnel, and the extracardiac conduit seem to have similar outcomes, which are better than those with an atriopulmonary connection.^{47,59,61,62,65–67} Because the introduction of the Norwood procedure was contemporary to the introduction of the total cavopulmonary connection, the majority of those surviving initial staged operations had this latter form of Fontan and predictions of their late survival should be based on

these latter procedures. The Australia and New Zealand Fontan Registry, collating data from 1423 subjects, reported survival at 26 years of 89% for those operated on with the lateral tunnel technique and 92% at 18 years for those with an extracardiac conduit, in concordance with recent major series.^{57,59,67} Interestingly, the survival

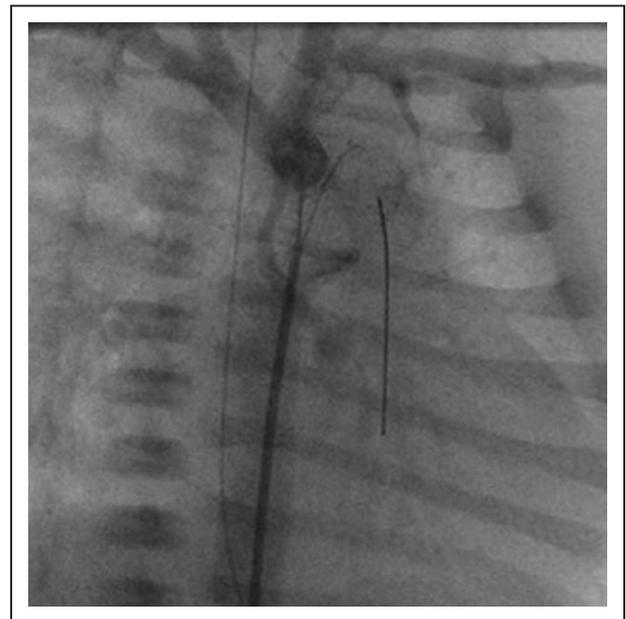


Figure 3. Right anterior oblique angiography performed through a 4F multipurpose catheter positioned at the junction of the arterial duct (stented by an open cell self-expanding SinusSuperflex-DS stent).

The extreme hypoplastic ascending aorta is connected to a rather well-developed aortic arch. A coronary soft-tip wire is seen passing retrograde through the ductal stent into the right ventricular outflow tract.

of this population has not been subjected to a sudden decline in early adulthood as was previously expected. The attrition rate of this population has been remarkably stable, and, for this reason, we feel emboldened to make prediction of survival to 30 years.⁵⁹

HEART TRANSPLANTATION

Since the conception of the Fontan operation, predictions were made that heart transplantation would be the final end-of-life option and counseling of the families invariably included this option. In reality, heart transplantation has taken only a very small part in the treatment armamentarium of this population with only 4% to 7% reaching this status within 20 years of the Fontan.^{49,68} The reasons for the infrequent use of this ultimate resource are likely the multiplicity of previous operations increasing the risk of reentry and immunoreactivity, the poor general status of the patients, and the complexity of the reconstruction necessary.^{69,70} It has been recognized that those with a Fontan circulation did not have equal access to transplantation because this access is dependent on structuring of heart transplantation programs with expertise in congenital heart diseases.⁶⁸ A large multicentric study has suggested that posttransplantation survival in patients with a previous Fontan surgery was inferior to the survival of patients with idiopathic dilated cardiomyopathy and biventricular congenital heart diseases.⁷⁰ Numerous single-center studies have since demonstrated equal late outcomes after transplantation of patients with a Fontan circulation including those with HLHS. It demonstrates that heart transplantation should be offered to patients with a Fontan circulation and that restriction of this practice to expert centers may be advisable.^{68,71–73}

BURDEN OF DISEASE

Even though survival after Fontan is now proven to be superior to previous expectations, the burden of disease is still considerable. In a study of 529 patients, who had undergone an extracardiac conduit in Australia and New Zealand, 47% of the hospital survivors experienced an adverse event within 15 years, when including events such as transplantation, reoperation, stroke, tachy- and bradyarrhythmia, thromboembolic events, bleeding, pacemaker implantation, protein-losing enteropathy, plastic bronchitis, and poor functional status.⁶⁰

It has been already demonstrated that the burden of late complications is higher in patients with HLHS.^{60,74} It does not seem at this stage that the higher severity of this condition is related to the intrinsic inability of a single right ventricle to support a Fontan circulation for several decades, but rather to the fact that patients with HLHS tend to have a higher incidence of decreased ventricular function and atrioventricular valve regurgitation, and have

a higher propensity of developing arrhythmias.^{60,74–76} Because patients with HLHS clearly have a higher burden of disease, it is likely that some difference will appear in the future in the survival rate of these patients in comparison with the remaining Fontan population.

Tachyarrhythmias and Bradyarrhythmias

Dysrhythmia has been one of the most prevalent complications after Fontan surgery with the quoted incidence varying between 13% and 54% at 20 years.^{47,77,78} Unfortunately, the relative incidence of brady- and tachyarrhythmias and their relative type have rarely been specified. The incidence of supraventricular tachycardia is higher in patients with atriopulmonary connection type of Fontan because of the progressive dilatation of their atrial cavity. It seems that patients with an extracardiac conduit might have a lower incidence of dysrhythmias than those with a lateral tunnel.^{47,79,80}

Thromboembolic Events

Both the extracardiac conduit and the lateral tunnel technique result in the exposure of polytetrafluoroethylene in the venous flow, predisposing to the formation of clots and requiring the administration of antithrombotic agents.⁸¹ There is still no formal consensus on the ideal antithrombotic agent, and patients are either placed under warfarin or aspirin. A prospective randomized trial, a propensity score–matched analysis, and a meta-analysis all point toward the equivalence of both strategies or to the superiority of aspirin in the prevention of thromboembolic events.^{81–84} The incidence of thromboembolic events seems to be limited to 18% to 21% at 10 years.^{85,86}

Protein-Losing Enteropathy and Plastic Bronchitis

These 2 feared complications consist in the loss of proteins in the lumen of either the intestine or the bronchi, and seem to occur in between 5% and 10% of cases with an incidence increasing with time.^{47,49} A few years ago, the development of these complications was associated with a risk of death of 49% at 5 years, but it seems that the introduction of an enteric-coated form of steroid has now improved this prognosis at least temporarily.^{86–88} Although various interventions have been attempted for these complications, heart transplantation has been shown to remain the most effective treatment, likely because it normalizes the systemic venous pressures of these patients.^{89,90}

LIVER AND RENAL FAILURE

The key driver of the Fontan circulation is the increased systemic venous pressure, which over time has been noted to impact liver and renal function. Within years of

Fontan completion, almost all patients are noted to develop hepatic fibrosis with some evolving to cirrhosis.⁹¹ Some rare cases of hepatocarcinomas have also been described.^{49,92} Similarly, decreased glomerular filtration has been noted in patients with a Fontan circulation.⁹³ At this stage, it is difficult to appreciate to what extent liver fibrosis and cirrhosis will affect the survival and quality of life of those with a Fontan circulation. Although the existence of liver fibrosis is widespread, it has not yet been associated with profound hepatic functional impairment. The impact of the Fontan on the liver remains nonetheless one of primary concern for the long-term survivors with a Fontan circulation.

EXERCISE CAPACITY

It has been consistently reported that patients with Fontan circulation have decreased maximal exercise capacity with peak oxygen consumption averaging $\approx 65\%$ of predicted values.^{94,95} There is a wide variation in the capacity of these individuals to participate in individual and even group sporting activities, with some being able to function at levels that are surprising for patients bearing only 1 functional ventricle. The American Heart Association has recently lifted a ban on exercise for these patients.⁹⁶ It has now been demonstrated that resistance exercise training is actually beneficial for these patients.⁹⁷ An increased leg muscle mass is responsible for increased cardiac output on exercise, acting as an additional pump at maximal exercise. There is growing belief that a lifestyle including regular exercise training will be beneficial for this population.

PREGNANCY

The majority of practitioners recommend the avoidance of pregnancy.⁹⁸ It is remarkable that some women with a Fontan circulation have nonetheless achieved successful pregnancies, even though the number of those with HLHS remain anecdotal.^{98,99} Women with a Fontan procedure, who have attempted to become pregnant or carry to delivery, have faced issues of infertility, miscarriages, and pregnancies resulting in small and premature babies. The maternal mortality has remained low, but the impact of pregnancy on long-term outcomes remains to be elucidated.

NEURODEVELOPMENTAL OUTCOMES AND QUALITY OF LIFE

The late functional outcomes of patients living with a Fontan circulation remains the topic of heated debate. The majority of studies report them to have decreased scores of quality of life, but several optimistic studies have also been published depicting a normal quality of

life and a normal level of emotional functioning.^{95,100–102} The performance of patients with HLHS do not seem to differ from those with other single ventricle conditions or from other patients with congenital heart diseases.¹⁰³ There is no doubt that we should focus on improving the quality of life and provide psychological support to this population at risk. One should not, however, see this population as being profoundly debilitated. The majority of this population is able to function normally without perceived restrictions. It was remarkable to note that, in a yet unpublished work, 28 years after an atriopulmonary Fontan performed in Australia and New Zealand, not only two-thirds of the patients were still alive, but two-thirds of those patients were working.

MEDICAL THERAPY

Large variations in medical therapy have been described.^{104,105} As an example, it has long been debated whether angiotensin-converting enzyme inhibitors, a cornerstone medical therapy of heart failure, would be of interest in single ventricle circulation. In a randomized study of enalapril performed in 230 patients of the Paediatric Heart Network, no benefits could be demonstrated.¹⁰⁶ The recent TEMPO study (Treatment With Endothelin Receptor Antagonist in Fontan Patients, a Randomized, Placebo-Controlled, Double-Blind Study Measuring Peak Oxygen Consumption) suggested the benefits of the administration of bosentan.¹⁰⁷ The ideal medical therapy for these patients remains elusive.

COMMENT AND FUTURE CONSIDERATIONS

Tremendous progress has been made over the past 3 decades in the care of children with functional single ventricle malformations. Progress has been dramatic at times, such as the development of the Norwood procedure before which HLHS was universally fatal, and incremental, but no less important in improving outcomes. Our understanding of the long-term implications of a Fontan circulation continues to evolve as the number of adolescent and young adult survivors grows.

The state of the art for the care of the single ventricle patient is a moving target, constantly changing and improving. Currently, there are a variety of valid approaches to the management of the patient with HLHS and related FSRV malformations, particularly in the first stage. For the first stage of traditional surgical staged palliation (the Norwood procedure), there are 2 alternatives for the source of PBF, the MBTS and the RVPAS. Although the 12-month end point comparing transplantation-free survival showed a benefit associated with the RVPAS and the point averages continue to favor the RVPAS at 6 years, the difference is no longer statistically significant. Concerns over the ventriculotomy required

for the RVPAS and the development of subsequent late right ventricular function do not seem to be warranted, because right ventricular function by echocardiogram is equivalent at 6 years. Thus, either approach seems reasonable at this time. Continued follow-up of this large, well-characterized cohort of patients will be important, not only to detect any late effects of shunt choice, but also to understand long-term outcomes and how they may be useful to inform early management decisions beyond shunt type.

The hybrid approach is also an equally viable alternative to traditional staged palliation. As stated in the section on the hybrid approach, one of the keys to success with this management strategy is the requirement for a dedicated, experienced, and collaborative team of cardiologists and surgeons. The results for the hybrid procedure outlined in this article are the outcomes from arguably the 2 most experienced centers in the world. The results from the SVR trial represents a 15-center sampling of large- and moderate-sized institutions with varying traditions of success with surgical single ventricle palliation. The highest performing centers in the SVR trial have results superior to those reported for the hybrid approach, whereas standard performing centers have lower survivals. The potential long-term benefits of avoiding an extensive neonatal open-heart surgery, particularly in neurodevelopment, remain to be determined. Thus, the selection of one or the other approach depends largely on local experience and resources. However, an important caveat is that optimal results for the hybrid approach require equal dedication and expertise as is required for optimal surgical results, and simply switching from one to the other or vice versa does not guarantee success.

A common theme is the need for institutional commitment, expertise, and experience for optimal results for the care of these fragile and complex patients, whether the approach is the traditional staged surgical palliation and an initial Norwood procedure or the hybrid approach. There are multiple publications demonstrating a positive correlation between volumes and outcomes for high-complexity lesions.^{108,109} However, these results are on aggregate and do not indicate that any individual center, large, moderate, or small, by virtue of volume alone ensures superior results, nor that a moderate or small program cannot achieve excellent outcomes. In addition, the concept that regionalization to the small number of centers with superior results can take all cases of FSRV malformations each year, which number >1000 cases/y in the United States alone, is unrealistic in many countries.

If neither the type of shunt nor the development of the hybrid approach has dramatically decreased mortality, and regionalization seems unlikely in some countries, what potential avenues for improvement exist? As noted above, there was wide variation in practices across cen-

ters in the SVR trial. Although this may seem unimportant, the similarly wide variation in Norwood procedure mortality across centers, from 7% to 39%, is sobering.²⁰ It is possible that determining and sharing of best practices would be effective in improving the results at all centers. The Northern New England Cardiovascular Disease Study Group and the Michigan Society of Thoracic and Cardiovascular Surgeons Quality Collaborative have demonstrated that this approach has been effective in decreasing mortality and morbidity in adult cardiac surgery.^{110,111} The Pediatric Heart Network funded a study on collaborative learning aimed at decreasing the length of intubation following congenital cardiovascular surgery, which showed the method to be effective in promoting early extubation.¹¹²

Future directions include the use of stem cell therapy to improve cardiac performance in the single ventricle patient. Several groups have either demonstrated efficacy in small numbers of patients or are starting trials on the use of stem cells in this population.^{113,114} Techniques may include intracoronary injection, intramyocardial injection, or onlay patches containing stem cells aimed at improving both systolic and diastolic function of the myocardium.

Universally fatal only 35 years ago, improvements in the treatment of HLHS and related FSRV malformations has been transformative. The survival of this population of babies born with HLHS has now surpassed the initial expectations. The current state of the art reflects these remarkable achievements. Yet, despite this progress, the enthusiasm for improvement in survivals and long-term quality of life remains. As the first generation of these patients enter adulthood, the focus must now be on improving not only their longevity, but also their quality of life. In that respect, it becomes increasingly important to address the still considerable burden of disease associated with the Fontan circulation. It is likely that these improvements will depend on not only ongoing progress in the care for those living with a single ventricle malformation, but also on the improvement of the initial stages of their palliation.

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DISCLOSURES

None.

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FOOTNOTES

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