Use of a Composite Survival Curve to Optimise the Surgical Strategy for Double Inlet Left Ventricle

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ABSTRACT

Objectives: Palliative surgery has improved the long-term survival of patients with Double inlet left ventricle (DILV). Neonates and infants with DILV presenting with reduced pulmonary blood flow (RPBF) are often offered a systemic arterial-to-pulmonary artery shunt (SAPAS). There is evidence that SAPAS in these patients, may not improve survival compared to the natural history. An objective assessment of the efficacy of a treatment requires a comparison of the treatment outcome with the natural history. Our aim was to review the literature to determine whether SAPASs in patients with DILV and RPBF improves survival compared to the natural history.

Methods: We reviewed the literature and used the most externally valid data to create a composite survival curve that facilitates the comparison of surgical outcome and natural history.

Results: The data suggests that SAPASs in patient with DILV and RPBF, may not improve long term survival compared to the natural history. There is also evidence that palliative surgery in general in patients older than 1 year of age with increased or balanced pulmonary blood flow may not improve survival compared to the natural history.

Conclusions: SAPASs probably should not be offered to patients with DILV and RPBF. In addition, cardiac surgery probably should not be offered to patients with DILV presenting for the first time after 1 year of age.

Keywords
Double inlet ventricle surgery.

Introduction
With respect to double inlet left ventricle (DILV), palliative cardiac surgery refers to the following procedures: pulmonary artery banding, systemic arterial-to-pulmonary arterial shunts (SAPAS), and systemic venous- to-pulmonary artery shunts [1].

Systemic venous-to-pulmonary artery shunts include the ‘Fontan circulation’ [2]. The introduction of palliative cardiac surgery has improved the long-term survival of patients with DILV [1,3]. Neonates and infants presenting with reduced pulmonary blood flow (RPBF) are often offered SAPASs; there is evidence that surgical outcomes in this subset of patients are worse than in patients with balanced or increased pulmonary blood flows [1]. An objective assessment of the efficacy of a treatment requires a comparison of the treatment outcome with the natural history.

We reviewed the literature and used the most externally valid data to create a composite survival curve that facilitates the comparison of surgical outcome and natural history. A pilot study suggested that there would be insufficient data to conduct a credible meta-analysis or receiver operator curve analysis; for this reason we used the ‘Proximal Similarity Model’ (PSM) to determine which natural history and surgical outcomes studies had the greatest external validity [4]. As the observed survival from a natural history curve is an estimate of the probability of survival, it can be directly compared with the postoperative Kaplan-Meier survival curves [5].
We used the most externally valid studies to generate a composite survival curve. This curve graphically demonstrates the surgical survival benefit as a function of pulmonary blood flow and patient age.

**Patients and Methods**

**The Proximal Similarity Model**

According to this model, there are essentially 3 major threats to external validity; these relate to the study subjects, the study location and the study time. The proximal similarity model is used to establish which study best represents the population of interest (i.e. which study has the greatest external validity). Table 1. Shows which parameters were utilised to decide which study had the greatest external validity with respects to natural history and surgical outcome studies.

| Area of threat to External validity | NH studies: parameters used to assess the validity | SO studies: parameters used to assess the validity |
|------------------------------------|--------------------------------------------------|--------------------------------------------------|
| 1. Subjects                         | DILV; LSS                                         | DILV; LSS; AAS                                   |
| 2. Geography                       | Anywhere                                          | Multicenter                                     |
| 3. Time                             | Any time                                          | Paper published after the year 1990             |

Abbreviations: AAS: Age At Surgery; DILV: Double Inlet Left Ventricle; LSS: Large Sample Size; NH: Natural History; SO: Surgical Outcome.

The following factors were considered when choosing which papers were to be used in the construction the surgical outcome curves:

- Study sample size.
- Median age at SAPAS
- Assessment of pulmonary blood flow
- Paper was published 10 years after the introduction of the Fontan circulation (This was to ensure that enough time had elapsed for broad adoption of the technique).

**Search strategy and selection criteria**

PUBMED and Google Scholar were systematically searched; the time frame for the search was January 1st 1966 to November 30th 2021. The search terms and search strategy are shown in Table 2. PUBMED searches utilised the “title/abstract” option and Google scholar searches utilised the “all in title” option; all search terms were combined with “AND”. Titles and abstracts were reviewed and full-text articles were examined when the abstract suggested that there was a possibility that data pertaining to natural history or long-term surgical outcomes could be present. Non-English language studies and non-human studies were also excluded. A recent large study by Hadjicosta [6] reported a 10 year mortality rate of 12% and the survival curve had almost flattened by 2 years of follow-up. To detect this level of mortality with a 85 % confidence level, a study would have to have followed-up at least 88 patients to 10 years. In view the fact that the survival curve had almost flattened out at 2 years post follow-up, we only considered surgical outcome studies that followed at least 88 patients to 2 years post-operatively. This sample size ‘cut-off’ was determined using a free online sample size calculator [7]. The search flow is shown in figure 1.

**Table 2: Search strategy.**

| Search number and terms                                                                 |
|------------------------------------------------------------------------------------------|
| 1. Double, inlet, left, ventricle, natural, history (GS,P)                              |
| 2. Double, inlet, left, ventricle, survival (GS,P)                                      |
| 3. Double, inlet, left, ventricle, long, term, survival (GS,P)                          |
| 4. Double, inlet, left, ventricle, unoperated (GS, P)                                   |
| 5. Natural, history, cardiac, malformations (GS,P)                                      |
| 6. Congenital, heart, disease, natural, survival (GS,P)                                 |
| 7. Double, inlet, left, ventricle, long, term, results (GS,P)                           |
| 8. Double, inlet, ventricle (GS)                                                       |
| 9. Fontan, long, term, outcome (GS)                                                     |
| 10. Bidirectional, cavopulmonary, Double, inlet, left, ventricle (P)                    |
| 11. Cavopulmonary, Double, inlet, left, ventricle (GS)                                  |
| 12. Glenn, Double, inlet, left, ventricle (GS)                                         |

Key: GS = Goggle scholar, P = Pubmed

**Creation of the composite graph**

The most externally valid natural history data (essentially the paper with the largest sample size of patients followed from birth till death) was used to create a natural history curve. We then examined the postoperative survival data of relevant retrieved full-texts. The most externally valid studies were used to generate the composite graphs. A paper on surgical outcome was considered to have the greatest external validity if it had all 3 of the following factors:

- The largest sample sizes
- The lowest median age at the time the SAPAS was performed.
- The paper was published after 1981 (10 years after the introduction of the Fontan circulation; this ensured adequate time for the procedure to be broadly adopted)

The chosen surgical outcome curves were then superimposed on the natural history curve.

**Results**

The search yielded 696 results; 39 relevant full-texts were obtained after examining titles and abstracts. Sixteen full-texts related to the natural history of DILV and 23 related to the surgical outcome. The references of these full texts were examined for additional
relevant publications. The important details of the studies included for consideration when developing the postoperative survival branch of the composite survival curves are shown in table 3. A list of excluded full texts that may have followed-up enough patients for 10 years, but survival data could not be determined from the texts, is shown in table 4. A list of the retrieved natural history full texts is included in table 5. The natural history curve for DILV is shown in figure 2; composite graphs are shown in figures 3 and 4.

Table 3: Surgical outcome full-texts included.

| Authors               | YOES   | TN  | AOVPBF | 10 YS(%) |
|-----------------------|--------|-----|--------|----------|
| Franken et al. [8]    | -      | 152 | No     | 93       |
| Tham et al. [9]       | 1990   | 145 | No     | 76       |
| Franklin et al. [10]  | 1973   | 191 | Yes    | 42       |

Key: AOVPBF: Assessed Outcome Versus Pulmonary Blood Flow; TN: Total Number of Patients; YOES: Year of Earliest Surgery; YS: Year Survival.

Table 4: Surgical outcome full texts excluded but may have followed up enough patients postoperatively to 10 years.

| Study                | Study Year | Number of patients |
|----------------------|------------|--------------------|
| 1. Hadjicosta et al. [6] | 2021       | 327                |

Figure 2: Natural history curve for Double inlet left Ventricle (DILV). NHx= Natural History.

Figure 3: Comparative NHx composite graph DILV. NH= Natural History, PBF= pulmonary blood flow; Sx= Surgery.

Discussion

Two methods are used to determine the natural history of a disease:
- Follow-up a group of live patients to determine when patients die.
- Perform an autopsy study on a group of patients thought to have died from the disease to determine the age pattern at the time of death.

Despite an extensive search, only one study was found that could potentially be used to generate our natural history curve. This study followed 946 patients with congenital heart disease (CHD) over 27 years [3]. This study was conducted in Central Bohemia under circumstances that may not occur again: at the time of the study, all patients with congenital heart disease were managed at one institution. Minimal surgery was performed locally and patients did not travel out of the country for surgery. Patient follow-up was rigorous and post-mortems in all children were mandatory. Together, these factors provided an ideal opportunity to study the natural history of CHD. We used the data from this study to develop our natural history curve for the first 15 years of life. There were 32 patients with DILV in this study [3,24]. The 15-year natural history mortality was 69% (+/- 11.77%; 85% confidence interval).

Only 3 surgical outcome studies included more than 88 patients at 2 years of follow-up; Franklin et al were the only group to consider how pulmonary blood flow affects surgical outcome [10]. They examined the third largest cohort of patients in the papers we retrieved and did so during an era when the treatment of DILV by shunts or pulmonary artery (PA) banding was already established. They also gave the actual number of patients at various durations of follow-up. For these reasons we considered this study to be the most externally valid and we used it to create our composite graphs. Franklin published additional data in his MD thesis [1]; specifically, he published a natural history survival curve for a subset of DILV patients with a particularly poor prognosis. This was the only data of this kind that we retrieved from our search. We used it to create our composite curve in figure 3.
A meta-analysis of randomized clinical trials (RCT’s) or observational studies is the preferred method of reviewing scientific literature. Our search did not yield any RCT’s and the observational studies found had insufficient data for a meta-analysis or receiver-operator curve analysis. We chose the PSM technique to assess bias and confounding. The most externally valid studies where then used to create the composite curves. Although Franklin [10] categorized the degree of pulmonary blood flow using cardiac catheterization and chest radiographs, the degree of pulmonary blood flow (PBF) may be pragmatically defined as follows [25]:

- Reduced PBF: arterial oxygen saturation of less than 75%
- Balanced PBF: arterial oxygen saturation between 75% to 85%
- Increased PBF: arterial oxygen saturation and greater than 85% PBF.

According to figure 3, the surgical survival for patients with DILV and RPBF is essentially the same as the worst possible natural history survival. This ‘worst possible natural history’ curve was derived from survival data from a subset of patients with DILV and multiple risk factors for poor surgical outcomes: there is evidence that RPBF, systemic outflow obstruction and aortic obstruction are risk factors for poor outcome. Franklin examined the natural history of a subset of patients with a combination of these factors [1]. As expected, they had particularly poor outcomes; these results were used to produce the worst possible natural history curve in figure 3. Figure 3 demonstrates that surgery in patients with DILV and RPBF may not significantly improve survival compared to the natural history. This is a significant and counterintuitive point as this is the subset of patients one would assume could benefit most from a SAPAS.

The actual natural history curve for DILV is flat from 1 year of age and remains flat up to 15 years of age [3]. This implies that once an infant has survived to 1 year of age, they should survive to 15 years of age. Although we do not have adequate natural history survival data beyond 15 years of age, we retrieved several reports of unoperated patients living beyond the 4th the decade of life (table 5). Considered together, the flat natural history survival curve beyond 1 year of age and the reports of unoperated patients surviving beyond the 4th decade of life, imply that there is a possibility that once a patient survives to 1 year of age they should survive beyond the 4th decade of life. If this is the case, surgery beyond 1 year of age may not improve survival compared to the natural history. To develop this argument, consider that the composite graph in figure 4 demonstrates that the surgical survival curve is essentially the same as the natural history curve when surgery is performed on patients after 1 year of age. This evidence suggests that offering cardiac surgery to any patient with DILV who presents for the first time after one year of age may not improve 15 year survival compare to the natural history. This has important implications with respects to surgical decision making in developing countries where the diagnosis of congenital heart disease is often confirmed after one year of age [26].

Prostaglandin (PGE1) has been shown to increase pulmonary blood flow in cyanotic patients with ductus dependent pulmonary blood flow [27]. PGE1 is unlikely to be effective if given after after 4 days of age [27]. The use of PGE1 was well established during the era that the data used to construct the composite graph was obtained; the median age at presentation of these patients was 1 day [1]. It is reasonable to assume that patients who had an increase in SPO2 from < 75% to 75% or greater, were not considered to be in the category of “reduced pulmonary blood flow”. In this regard, palliative surgery would be indicated in these patients. There are reports of patients who present with features of reduced pulmonary blood flow and are diagnosed as having a ‘closing’ persistent ducts; an emergency SAPAS is usually offered to these patients. A large natural history study found that 96% of persistent ducts have closed by 7 days of age; the percentage-patency curve has essentially flattened by this age, implying that the remaining ducts are unlikely to close beyond this age [28]. When this fact is considered in conjunction with the fact the prostaglandin is only effective during the first 4 days of life, the actual interval where an ‘emergency’ SAPAS would be indicated would be days 5, 6 and 7 of life. Clearly, where a neonate has been started on PGE1 before the 5th day of life, and currently has a SPO2 of >75%, an urgent

| Author                      | Study Year | Study type          | Max age of patient(s) | SpO2 (%) | Status  |
|-----------------------------|------------|---------------------|-----------------------|----------|---------|
| 1. Salame-Waxman et al. [11] | 2019       | Case report         | 28                    | 80       | Alive   |
| 2. Alpat et al. [12]        | 2018       | Case report         | 45                    | 85       | Alive   |
| 3. Herbert et al. [13]      | 2017       | Case report         | 54                    | 77       | Alive   |
| 4. Agrawal et al. [14]      | 2017       | Case report         | 41                    | -        | Alive   |
| 5. Brida et al. [15]        | 2016       | Case report         | 21                    | -        | Alive   |
| 6. Potterucha et al. [16]   | 2016       | Descriptive cohort  | 77                    | 82(median) | Alive   |
| 7. Park et al. [17]         | 2007       | Case report         | 41                    | 86.9     | Alive   |
| 8. Kaya et al. [18]         | 2007       | Case report         | 34                    | -        | Alive   |
| 9. Book et al. [19]         | 2007       | Case report         | 71                    | 75       | Alive   |
| 10. Restaino et al. [20]    | 2004       | Case report         | 57                    | -        | Alive   |
| 11. Hager et al. [21]       | 2002       | Case report         | 62                    | 88       | Alive   |
| 12. Amnash et al. [22]      | 1996       | Descriptive cohort  | 66                    | -        | Alive   |
| 13. Vitarelli et al. [23]   | 1996       | Case report         | 59                    | 78       | Alive   |
| 14. Samanez et al. [3]      | 1992       | Descriptive cohort  | 15                    | -        | Alive   |

Key: * years.
SAPAS should be performed. This should be offered whether the infant showed improvement after starting PGE1 or had a SPO2 of > 75% at the time PGE1 was started. The rational for this is that there is no way of knowing, in the latter case, whether this was an infant whose duct would remain open if PGE1 was stopped.

In conclusion our results suggest the following:

- SAPASs probably should not be offered to patients who have DILV with RPBF.
- Cardiac surgery probably should not be offered to patients with DILV who present for the first time after 1 year of age.
- An ‘Emergency SAPAS’ for a ‘closing PDA’ is probably only indicated during the first week of life.

We hope that our results will be used to optimise surgical decision making in patients with DILV.

References

1. Franklin, Rodney CG. The fate, survival and suitability for definitive surgery of infants with double inlet ventricle and tricuspid atresia. Doctoral thesis (Ph.D.), University College London (United Kingdom). 1997.
2. Fontan F, Baudet E. Surgical repair of tricuspid atresia. Thorax. 1971; 26: 240-248.
3. Samánek M. Children with congenital heart disease: probability of natural survival. Pediatr Cardiol. 1992;3: 152-158.
4. Polit DF, Beck CT. Generalization in quantitative and qualitative research: Myths and strategies. International Journal of Nursing Studies. 2010; 47: 1451-1458.
5. http://ocw.jhsph.edu/courses/fundepi/pdfs/Lecture9.pdf
6. Hadjicosta E, Franklin R, Seale A, et al. Cohort study of intervened functionally univentricular heart in England and Wales (2000-2018). Heart. 2021.
7. https://www.calculator.net/sample-sizecalculator.html.
8. Franken LC, Admiraal M, Verrall CE, et al. Improved long-term outcomes in double-inlet left ventricle and tricuspid atresia with transposed great arteries: systemic outflow tract obstruction present at birth defines long-term outcome. Eur J Cardiothorac Surg. 2017; 51: 1051-1057.
9. Tham EB, Wald R, McElhinney DB, et al. Outcome of fetuses and infants with double inlet single left ventricle. Am J Cardiol. 2008; 101: 1652-1656.
10. Franklin RC, Spiegelhalter DJ, Anderson RH, et al. Double-inlet ventricle presenting in infancy. I. Survival without definitive repair. J Thorac Cardiovasc Surg. 1991; 101: 767-776.
11. Salame-Waxman D, Meyer SL, Ebels T, et al. Natural History of Double Inlet Left Ventricle and Pulmonary Hypertension in an Adult Patient. JACC Case Rep. 2019; 1: 532-534.
12. Alpat S, Demircin M. Living with functionally univentricular heart: beating the odds? Asian Cardiovasc Thorac Ann. 2018; 26: 694-696.
13. Herbert S, Gin-Sing W, Howard L, et al. Early Experience of Macitentan for Pulmonary Arterial Hypertension in Adult Congenital Heart Disease. Heart Lung Circ. 2017; 26: 1113-1116.
14. Agrawal H, Velagapundi P, Katz N, et al. 41-year old patient with double inlet left ventricle without pulmonary stenosis presenting with Eisenmenger’s syndrome. J Am Coll Cardiol. 2017.
15. Brida M, Diller GP, Baumgartner H, et al. Double inlet left ventricle with unrestricted pulmonary blood flow and survival into adulthood, European Heart Journal. 2016; 37: 967.
16. Poterucha JT, Anavekar NS, Egbe AC, et al. Survival and outcomes of patients with unoperated single ventricle. Heart. 2016; 102: 216-222.
17. Park SJ, Kwak CH, Hwang JY. Long-term survival in double outlet ventricle combined with pulmonary stenosis and parachute mitral valve. Int Heart J. 2007; 48: 261-276.
18. Kaya MG, Tulmac M, Sen N, et al. Double-inlet left ventricle with transposition of great arteries in an asymptomatic adult. Arch Turk Soc Cardiol. 2007; 35: 136.
19. Book WM, Vaccari C, McConnell ME. Exceptional survival: Double inlet left ventricle presenting with aortic dissection. International Journal of Cardiology. 2007; 116: e65-e67.
20. Restaino G, Dirksen MS, De Roos A. Long-term survival in a case of unoperated single ventricle. Int J Cardiovasc Imaging. 2004; 20: 221-225.
21. Hager A, Kaemmerer H, Eicken A, et al. Long-term survival of patients with univentricular heart not treated surgically. J Thorac Cardiovasc Surg. 2002; 123: 1214-1217.
22. Ammash NM, Wames CA. Survival into adulthood of patients with unoperated single ventricle. Am J Cardiol. 1996; 77: 542-544.
23. Vitarelli A, Gabbarini F, Holmes heart in the adult: transesophageal echocardiographic findings and long-term natural survival. International Journal of Cardiology. 1996; 56: 301-305.
24. Samánek M, Benesová D, Goetzová J, et al. Distribution of age at death in children with congenital heart disease who died before the age of 15. Br Heart J. 1988; 59: 581-585.
25. Magoon R, Makhija N, Jangid SK. Balancing a single-ventricle circulation: 'physiology to therapy'. Indian J Thorac Cardiovasc Surg. 2020; 36: 159-162.
26. Awori MN, Ogendo SW, Gitome SW, et al. Management pathway for congenital heart disease at Kenyatta National Hospital, Nairobi. East Afr Med J. 2007; 84: 312-317.
27. Freed MD, Heymann MA, Lewis AB, et al. Prostaglandin E1 infants with ductus arteriosus-dependent congenital heart disease. Circulation. 1981; 64: 899-905.
28. Nagasawa H, Hamada C, Wakabayashi M, et al. Time to spontaneous ductus arteriosus closure in full-term neonates. Open Heart. 2016; 3: e000413.

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