

Delayed Sternal Closure in Congenital Heart Surgery: A Risk-Benefit Analysis

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ABSTRACT

Background: Delayed sternal closure commonly is used after pediatric cardiac surgery. Its benefits include relieving cardiac compression and stabilizing postoperative critically ill patients.

Methods: We retrospectively reviewed the records of 72 patients, who had undergone delayed sternal closure, among 1,254 patients operated for congenital heart diseases. Indications of delayed sternal closure, perioperative hemodynamic and metabolic status, postoperative infection, and mortality were reported.

Results: Transposition of great arteries was the most common preoperative cardiac pathology (26.3%). Bleeding and hemodynamic instability were the most frequent indications for delayed sternal closure, representing 38.8% and 34.7%, respectively. The mean duration of open chest was 3.45 days \pm 1.46 days. The mean duration of ICU stay was 20.95 days \pm 20.06 days. Two patients had deep sternal wound infection. Sepsis was found in 39 patients (54.1%), and the most common causative organism was coagulase negative (30.5%). ICU stay was a significant risk factor for sepsis ($P = .003$); duration of open sternum, period of mechanical ventilation (MV), and total hospital stay were not statistically significant risk factors. Sternal closure time (SCT) was affected by period of hemodynamic instability ($P = .036$). Bypass time, clamping time, and nonsurgical bleeding did not significantly affect SCT. The mortality rate was 15.2% ($N = 11$).

Conclusion: Delayed sternal closure is a simple and effective technique that could prevent postoperative cardiac compression in hemodynamic instability states after pediatric cardiac operations.

INTRODUCTION

Delayed sternal closure is a therapeutic modality to manage hemodynamic instability resulting from cardiac compression by sternal closure.

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MATERIALS AND METHODS

We retrospectively reviewed records of 72 patients, who had undergone delayed sternal closure, among 1,254 patients operated for congenital heart diseases at King Abdul Aziz University Hospital, from 2013 to 2018. Preoperative cardiac pathology is shown in Table 1.

Recorded were indication of DSC, time to sternal closure, and pre- and post-closure cardiopulmonary and metabolic status, including hemodynamic state and lactate level. Other variables, including ventilator support duration, mean intensive care unit (ICU) and hospital stay times, mortality, wound infections, and bloodstream infections were studied.

For myocardial protection, we used antegrade cold blood cardioplegia without epicardial ice slush. The initial dose was 30ml/kg and subsequent doses were 20 ml/kg with intervals of 20 minutes. Systemic temperature was drafted to 30°C to 32°C. In deep hypothermic circulatory arrest, the patient was cooled to 18°C with an ice pack on the head. All cases routinely were ultrafiltrated, during CPB with the exception of one case of coarctation repair.

The decision to leave the sternum open was based on hemodynamic state, lactate level, presence of myocardial edema, non-surgical bleeding, and conduction of extracorporeal membrane oxygenation (ECMO).

Sterna were kept open without the use of a retractor. Skin was not sutured. We closed wounds with a latex membrane (Esmark bandage). If there was a marked disproportion between cardiac size and thoracic cavity, we fixed a piece of bypass circuit tube between the two sternal bars to keep steadily open the chest. Dressing was changed daily in ICU.

Regarding antibiotic prophylaxis, we used cefazolin and vancomycin then according to culture. Postoperatively, we used Fentanyl at 1-4 μ /kg/hr for sedation and Pancuronium at 0.25-0.75 μ /kg/m as a muscle relaxant. When hemodynamic and respiratory parameters improved, lactate level lowered and negative fluid balance settled, a trial of sternal closure was attempted. We closed the sternum by interrupted simple stainless steel wires (size 1 to 4, according to weight). Muscle, subcutaneous tissue and skin were closed as one layer by interrupted mattress 3/0-4/0 polypropylene sutures.

Table 1. Cardiac Pathology

VSD	2
IAA	5
Tetralogy of Fallot	7
Truncus arteriosus	7
Tricuspid atresia	1
LVOT tunnel type	1
TGA	19
Mitral insufficiency	1
Coarctation	5
Univentricular heart	2
Mitral atresia	2
TAPVD	3
DORV	3
Hypoplastic left heart syndrome	5
Pulmonary atresia	6
DOLV	1
AVSD	2

RESULTS

Seventy-two patients with DSC were studied. All were left open at the end of the operation, with no patients opened in ICU. We had 30 (41.6%) females and 42 (58.3%) males. The mean age was 317.14 ± 753.28 (4-4256 days). There were no patients with previous sternotomy. TGA was the most common preoperative cardiac pathology (26.3%). Mean bypass time was 90.4 minutes \pm 50.5 minutes (15-319 minutes) and mean clamping time was 55.77 minutes \pm 38.01 minutes (6-215 minutes). Total circulatory arrest was conducted in 25 patients, and the mean duration was 22.72 minutes \pm 12.98 minutes (3-51 minutes). Indications of leaving the sternum open in OR are summarized in Table 2. We used ECMO in 5 patients. Peritoneal dialysis was performed in 38 patients (52.7%). Echocardiography routinely was done in ICU on the first postoperative day and immediately before closure. It showed poor cardiac function in 73.36% of patients on the first postoperative day. The mean duration of open chest was 3.45 days \pm 1.46 days (1-7 days). The mean period of mechanical ventilation was 9.25 days \pm 8.37 days (3-56 days). The mean duration of ICU stay was 20.95 days \pm 20.06 days (4-120 days) and total hospital stay was 26.48 days \pm 22.67 days (1-124 days). Sepsis was found in 39 patients (54.1%), and the most common causative organisms were coagulase negative (30.5%), klebsiella pneumonia (16.6%), and pseudomonas (13.8%).

Table 4 shows postoperative infection data. Two patients had deep sternal wound infection; one of them died from septic shock. We had 11 cases (15.2%) of mortality. Five patients were supported by ECMO. Causes of death are shown in Table 3.

Statistical regression showed ICU stay as a significant risk factor for sepsis ($P = .003$), while duration of sternal closure,

Table 2. Indications of Open Chest

Indication	Number
Non-surgical bleeding	28
Cardiac edema	12
Hemodynamic instability	25
High inotropes	16
Respiratory (low saturation)	7
Long bypass time	8
ECMO	5

period of MV, and total hospital stay were not statistically significant risk factors. Sternal closure time was affected by period of hemodynamic instability ($P = .036$). Bypass time, clamping time, and nonsurgical bleeding did not significantly affect SCT.

DISCUSSION

Cardiomedastinal disproportion was first described by Riahi and colleagues in 1975 [Riahi 1975]. Factors accounting for this problem include myocardial edema from reperfusion injury and cardiac manipulation, bleeding, arrhythmias, and ventricular dysfunction. This condition often is associated with poor myocardial protection and prolonged bypass time [Das 2011]. In the surgical era of shifting from palliative to corrective operations, DSC was more common. Furnary and colleagues reported the beneficial effect of DSC on hemodynamics. Cardiac index increased up to 59% and systolic blood pressure rose up to 18% with the opening of the sternum [Yasa 2010].

It has become an acceptable practice in pediatric centers to leave the sternum open electively in OR especially after long bypass time or after Norwood procedures [Johnson 2010]. All patients included in our study electively were left open. Our decision was based on hemodynamic parameters, TEE, and lactate level. TGA and HLHS were the most common cardiac pathology reported in most of series of DSC [Iyer 1997; Riphagen 2005; Tabbutt 1997].

In our study, TGA was the most common diagnosis accounting for 26.3% of our cases. Some investigators [Das 2011; Johnson 2010] limit their studies about DSC to patients with HLHS, in an effort to decrease variability. We included 5 patients with HLHS in our series. One of patients died because of post-ECMO multiorgan failure. Bleeding and hemodynamic instability represent the most frequent indications for DSC [Hurtado 2018]. They accounted for 38.8% and 34.7% of our indications, respectively. When to close the sternum still is debatable. Some centers delay closure for at least 3 days [Tabbutt 1997].

Riphagen et al emphasized the feasibility of early DSC within 24 hours without hemodynamic, respiratory, or metabolic compromise. The advantages of their policy are lower

Table 3. Causes of Death

Cause of death	ECMO Used	ECMO Not Used
	No.	No.
Septic shock	1	2
Low cardiac output	2	0
Multisystem organ failure	1	0
DIC	1	1
Pulmonary hemorrhage	0	1
Respiratory failure	0	2
Total	5	6

nosocomial infections and ventilator associated complications [Riphagen 2005].

However, this aggressive approach should be taken with caution since premature closure of the sternum with repeat DSC has more adverse effects than prolonging the period of DSC [Harder 2013].

We decide for closure when (1) the patient becomes hemodynamically stable; (2) no malignant arrhythmias are present; (3) arterial blood gases are acceptable; (4) negative fluid balance state; (5) lactate level decreases; and (6) echocardiography shows improved cardiac function and no significant residual shunt across septae or significant gradient across LVOT, RVOT, or valves. Mean closure time in our series was 3.45 days \pm 1.46 days. The only factor that proved to affect sternal closure time in our study was the period of hemodynamic instability. Various methods have been described for sternal closure. These include primary skin closure, use of sternal retractor, silicone sheet, mediastinal packing, and stenting the sternum by a piece of bypass circuit tube [Iyer 1997]. Staged closure using a binder clip was described by Fuchigami et al [Fuchigami 2016]. We usually use latex membrane (Esmark) for closure. This way, the mediastinum is isolated from the atmosphere.

With any cases of marked cardiac edema, we stent the sternum with a piece of bypass circuit tube. Complications of DSC include sternal wound infection, mediastinitis, and sepsis. We had 2 patients (2.7%) who developed deep sternal wound infection. The reported incidence of surgical site infection in the pediatric population after DSC ranges from 0% to 28%. Woodward et al reported 1.53% incidences of mediastinitis based on a multicenter study that included 8,774 patients [Woodward 2011]. They stated that DSC was not associated with increased risk of mediastinitis. We think that elective opening and closing of the sternum in OR together with a short period of open chest (3.45 days \pm 1.46 days in our study) explained our low incidence of SSI. We found that duration of ICU stay was a significant risk factor for sepsis. This finding is similar to that previously reported by Johnson et al who evaluated 1,283 patients operated in 45 centers for Stage 1 palliation for HLHS. DSC was performed in 74%. They stated that longer length of stay was associated with a higher rate of infection [Johnson 2010]. On the other hand, sternal closure time did not significantly influence our rate

of deep sternal wound infection. This is an interesting point in our results. We confirm the previous studies in this regard [Erek 2012; Tabbutt 1997; Woodward 2011]. Many authors reported higher incidence of infection with a longer period of open chest [Anderson 2002; Furnary 1992; Nelsonm 2016]. We agree with Harder et al [Harder 2013] that repeat DSC rather than prolonged period of open chest influences the rate of infection. This raises the importance of avoiding premature closure of the sternum.

Blood stream infection increases four-fold after DSC [Das 2011]. For more information, look to Table 5. Özker et al reported prolonged antibiotic use due to postoperative infection in 52.6% of patients [Özker 2012]. Sepsis was found in 39 of our patients (54.1%). The most common organism isolated in our patients was Gram negative, which is in agreement with literature [Pollock 1990; Tabbutt 1997; Woodward 2011].

Our mortality rate was 15.2% (11 patients), which is comparable to the range of 11% to 36% reported in literature [Alexi 1995; Hurtado 2018; Özker 2012]. For more information, look to Table 5. Özker et al and Hurtado et al reported mortality of 22% and 34.2%, respectively [Hurtado 2018; Özker 2012].

Higher rates of infection and mortality were reported if the sternum was opened in CICU, when compared with those kept open electively in OR [Pollock 1990; Samir 2002]. This highlights the importance of predicting patients in need of DSC at the end of an operation. Samir et al analyzed risk factors that may predict the need for DSC among 312 neonates, who underwent open heart surgery. They found that an age of less than 7 days, diagnosis of IAA or TAPVD, aortic clamping time of more than 98 minutes and bypass time exceeding 185 minutes, and central venous saturation after bypass lower than 51% are significant risk factors. Other factors, including prematurity, preoperative inotropic or ventilator support, circulatory arrest, type of cardioplegia, and use of ultrafiltration were not statistically significant risk factors [Samir 2002].

Edwards and Baker found the rate of infection is increased with long bypass time, long duration of low COP state, and excessive bleeding [Edwards 1983]. We found that ICU stay was the only significant factor that influenced the rate of infection. Nelson-McMillan analyzed STS congenital heart surgery database of 6,127 operations with sternum left open in 100 centers. They reported infection rate 18.7% in patients with the sternum left open compared with 6.6% in patients without this in the same age and procedural groups. DSC in ICU location was not associated with increased risk of infection. Within the first few days, day-to-day increments in risk of infection were small [Nelsonm 2016]. Harder et al found that use of ECMO and multiple periods of DSC were independent risk factors of SSI [Harder 2013]. We used ECMO in 10 patients. Three of them developed sepsis and 5 died. Mean duration of open chest in the remaining 5 survivors was 5.25 days.

CONCLUSION

Delayed sternal closure is a simple and effective method that could relieve cardiac compression in a state of hemodynamic

Table 4. Postoperative infection data

Patient	Diagnosis	Site of infection	Organism	Result
1	VSD-coarctation	Blood, pleural fluid	coagulase -ve staph	
2	Tricuspid atresia	Respiratory	polymrph	
3	Mitral insufficiency	Respiratory	Gram -ve bacilli	
4	TGA	Respiratory	Enterobacter	
5	Univentricular heart type isomeric	Blood, sputum	coagulase -ve staph	
6	Coarctation	Sputum	klebsella pneumonia	
7	Truncus arteriosus	Tracheal aspirate	MRSA	
8	Coarctation	Bronchoalveolar lavage	Pseudomonas argunosa, sphengomonas paucimobilis	
9	DORV	Peritoneal fluid	coagulase -ve staph	Died
10	VSD	Tracheal aspirate pseudomonas aregenosa	STENOTRPHOMONAS MALTOPHILIA	
11	Hypoplastic left heart syndrome	Wound swab	MRSA	
12	Pulmonary atresia VSD (PA/VSD)	Tracheal aspirate	Pseudomonas argunosa	
13	TGA	Tracheal aspirate	Klebsiella pneumonia	
14	TGA	Nasopharyngeal aspirate	Klebsiella pneumonia	
15	TAPVD-intracardiac	Tracheal aspirate	Klebsiella pneumonia	
16	Univentricular heart LV type	Blood	Coagulase negative staph	
17	Tetralogy of Fallot	Urine Peritoneal fluid coagulase negative staph	Enterococcus fecalis, E.coli	Died
18	Tetralogy of Fallot	Sputum blood staph epidermdis	Yeast, acinetobacter baumannii	
19	HLHS	Sputum Blood - sputum	YEAST CELLS coagulase negative	
20	Coarctation	Sputum	enterbacter cloacea	
21	Coarctation	Sputum	strept viridian, psudomonas	
22	Truncus arteriosus	Blood	Coagulase negative staph	
23	TGA	Blood	burkholderia gladioli	
24	PA/VSD	Blood	Pleural fluid Klebsiella , candida albican coagulase negative staph	Died
25	DORV	Urine	klebseilla pneumonia	
26	Truncus arteriosus	Tracheal aspirate	H. influenza	
27	TGA	Tracheal aspirate	stenotrophomonas maltophilia PSEDOMONAS ARGUNOSA	
28	Pulmonary atresia IVS (PA/IVS)	Tracheal aspirate	MRSA	
29	TOF	Tracheal aspirate	Pseudomonas	
30	DOLV	Blood	coagulase -ve staph	
31	TOF	Tracheal aspirate	Enterobacter aeriginosa	
32	AVSD-Complete	Nasopharyngeal swab blood	MORAXELLA CATARRALIS chyreseobactrium menengiosepticum	
33	TGA	Blood	Coagulase -ve Staph	
34	Interrupted aortic arch (IAA)	Wound	Pseudomonas	
35	Interrupted aortic arch (IAA)	Tracheal aspirate	K. pneumonia	Died
36	HLHS	Tracheal aspirate	ENTEROBACTER CLOACAE)	

Table 5. Incidence, sites of infection and mortality reported in pediatric series are shown

Study	Total number of bypass cases	Number of DSC	Mean age	Mean SCT	Infection	Superficial SSI	Deep SSI	Mortality
Alexi-Meskishvili et al, 1995(6)	1252	113(9%)	38% newborn 32% infants 30% children	In 86% within the first 6 days of operation	0	0	1 mediastinitis	41(36.2%)
Tabbutt et al, 1997(13)	2559	10 days	3.4 days	4	5	7 (3.9%) mediastinitis	34(19%)	
Iyer et al, 1997(5)	3718	150(4.03%)	229 ± 51 days	3.86 ± 0.29	-	15	0	17 (11.3%)
Samir K et al, 2002 (9)	-	312	Median 11.9 d	- Min.3d	-	-	-	30 (21.4%)
Riphagen et al, 2005 (10)	585	66 (11.3%)	Median 5d	Median 21 hrs in 60 patients	-	5	0	13(19.7%)
Johnson J et al, 2010 (4)	1283 HLHS (45 centers)	74%	Median 6 d	-	155	51	17 mediastinitis	-
Hurtado-Sierra, 2010 (14)	2325	259 (11%)	-	2.3±1.4 d	-	-	6 (2.3%) mediastinitis	22.4%
Özker, 2012 (20)	1100	38 (3.45%)	38.5±85d	2.9±2.3d	30	6	4 mediastinitis	13(34.2%)
Erek E, 2012 (17)	188	91(48.4%)	Median 9.5±11d	-	35	3	3	18
Das S, 2011(16)	110 HLHS	67 (61%)	Median 4d	4-5d	20	-	1 mediastinitis	14
Harder EE, 2013(12)	-	375	-	Median 9d	-	12	27 mediastinitis	33
Nilson-McMillan, 2016 (11)	89120 (100 centers)	6127	8d	-	505 (8.2%)	384 (6.3%)	107 (1.8%) mediastinitis	-
Göçen, 2017 (21)	43 arterial switch	22.67±75.75	43	2.7±2.3 d	6	1	0	1(2.32%)

instability after congenital heart surgery. A low incidence of surgical site infection is a favorable outcome to adopt it as a therapeutic modality.

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