

Clinical and Echocardiographic Outcome of Percutaneous Closure of Ventricular Septal Defects



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Abstract

Background: Ventricular septal defects (VSD) are the most common congenital cardiac anomalies. Hemodynamically significant defects require closure. Surgery is the gold standard for most defects. Recently, transcatheter closure has provided an acceptable option in selected cases. We aim to evaluate the outcomes of percutaneous closure of VSD and assess the effect of device closure on echocardiographic left-sided chamber parameters.

Methods: A retrospective cross-sectional study included patients who underwent percutaneous VSD closure between 2015 and 2021. Clinical data, angiographic, and echocardiographic images before and after closure were reviewed.

Results: Among 38 patients who had VSD closure during the study period, 34 had isolated defects. The most common defect type was perimembranous (27 patients, 71%). Median age was 6.5 years (0.5-49), and median weight was 19.5 kgs (5-70). Median VSD size was 4 mm (3-10). Most patients were symptomatic (26, 68%). Following closure, all symptomatic patients reported improved symptoms. There were no serious complications during a mean follow up period of 18months (± 19). All echocardiographic parameters of left sided volume overload decreased when compared to baseline within the first month of the procedure (left ventricular Z-score dropped from 1.53 ± 0.87 to 0.31 ± 0.68 , $p=0.0001$, left atrial Z-score dropped from 2.78 ± 0.71 to 1.54 ± 1.13 , $p=0.0001$).

Conclusion: Percutaneous closure of VSD in the current era is safe and effective. In addition to improved symptoms of heart failure and exercise intolerance, there is a significant reduction of left atrial and left ventricular size within a short period after closure.

Keywords: Ventricular Septal Defects; Percutaneous Closure; Echocardiography

Introduction

Isolated ventricular septal defects (VSD) account for approximately 40% of all congenital heart lesions [1]. The natural history of VSDs is variable. It has been estimated that spontaneous closure may occur in 12-84% of the defects, with higher rates associated with smaller defects [2-5]. Persistent small defects usually remain asymptomatic and require no treatment unless complicated, while larger defects present early in life with congestive heart failure symptoms and warrant closure [2,5].

Surgery is the traditional modality to close VSDs. However, for the last two decades, advances in percutaneous closure have been described, with several device designs and modifications, resulting in improved success and decreased complication rates [6-9].

At our institution, percutaneous closure of VSD has been done for selected patients since 2015. Indications for closure included

symptoms of heart failure or exercise intolerance, left ventricular or left atrial dilatation on echocardiography, and in a few patient, significant residual defect following surgical closure of VSDs.

Our aim in this study is to assess the outcome of percutaneous closure of VSD as a therapeutic modality and to describe the echocardiographic changes of the left ventricular and the left atrial size following percutaneous VSD closure.

Methodology

A Retrospective cross-sectional study included (38) patients who had elective VSD closure at Jordan University Hospital between 2015- 2021. Revision of patient files, echocardiographic and angiographic images were done. Data reviewed included demographic and clinical data, including age at procedure, indication of procedure, type and size of VSD, presence of other congenital anomalies, type of device used to close the

defect, approach for closure, complications, and follow up data. Echocardiographic data included parameters of volume overload: Left ventricular end diastolic dimension (LV), left atrial (LA) dimension as well as left atrial to aortic root dimension ratio (LA/AO), in addition to ejection fraction. The measurements are done in the standard M-Mode echocardiography of long axis parasternal views. Measurements of LV and LA dimensions are converted to Z-scores based on weight and height, while LA/AO ratio is expressed as a numeric value. Echocardiographic data was obtained in three different time points for comparison: At baseline (prior to procedure), first follow up following the VSD closure (within the first six weeks of procedure), and at last follow up. In addition to echocardiographic assessment of left sided volume overload, follow up echocardiography was done to assess device position, the relation of the device to the surrounding structures, any complications such as residual shunts, and the presence of aortic or tricuspid regurgitation.

Categorical data is presented as number (percentage), continuous data is presented as mean (\pm standard deviation), or median (range). P value of less than 0.05 was considered significant. Statistical analysis was done using online graphpad quickcalcs software (www.graphpad.com/quickcalcs, San Diego, USA).

The study was approved by institutional review board at Jordan University Hospital and by research committee of the University of Jordan.

Procedure

Procedures were performed under either deep sedation or general anesthesia. And via Femoral arterial and venous access. All

patients were given 100 IU/kg heparin and prophylactic antibiotics at the start of the procedure. All procedures were guided by trans-thoracic (TTE), or trans-esophageal echocardiography (TEE) as well as fluoroscopy. Left ventricular angiography was done, and the projection that best profiled the defect was used for sizing and as a reference for fluoroscopic guidance. Defects were crossed either retrogradely from the left ventricle, or antegradely from the right ventricle. Technique of closure of the defect has been described extensively in the literature, and it depends on type and location of the defect, type and design of the occluder device, in addition to the operator experience and preference [6-9]. Device size and type selection were based on echocardiographic and angiographic assessments. Generally, the selected waist size of the device was 2-3 mm larger than the measured VSD diameter.

Results

Thirty-eight patients underwent VSD closure at our hospital between 2015-2021. Females were 24 (63%), median age at intervention was 6.5 years (range 0.5-49), and median weight was 19.5 Kg (5-70). Defects were either perimembranous (27, 71%), or muscular (7, 18%), while two patients had multiple muscular defects, and two patients had both a muscular and perimembranous defect. In addition to the VSD closure, two patients had PDA closure. Four patients were syndromic; three had down syndrome, and one had Williams syndrome. Most patients had symptoms related to the VSD (26, 68%): Eleven patients (29%) had symptoms of heart failure and were on medical therapy before procedure, 10 (26%) children had poor weight gain, 5 patients (13%) had decreased exercise tolerance, and 12 patients had no symptoms. Indications for closure in the asymptomatic patients was left sided dilatation.

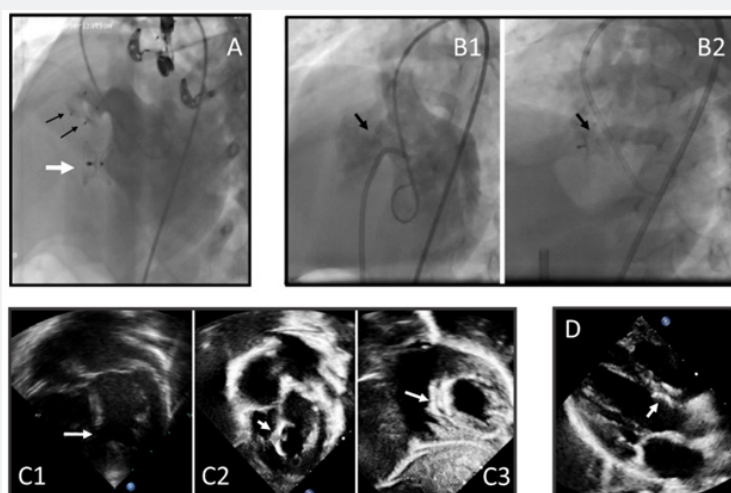


Figure 1: Images of 4 patients with different devices used for percutaneous ventricular septal defect closure: A: Angiogram of a patient with a large muscular defect closed using Amplatzer muscular VSD occluder (whit arrow), and perimembranous VSD closed using two Amplatzer Duct Occluder II within the VSD aneurysm (black arrows). B: Angiogram of a patient with a perimembranous VSD (B1), closed using Occlutech PDA occluder (B2). C: Echocardiographic images of a patient with a large muscular VSD (C1), closed using Konar MF occluder shown in 4-chamber view (C2), and subcostal view (C3) with good device alignment to the ventricular septum. D: Echocardiographic image of perimembranous defect closed using Occlutech perimembranous VSD occluder.

Total of 42 defects were closed in 38 patients (two patients had two defects closed, and one patient had three defects closed). The most commonly used device was Amplatzer duct occluder II (ADOII) (Abbott, CA, USA), in 27 defects (64%). Other devices used included LifeTech muscular device in 4 defects (LifeTech scientific, Shenzhen, China), Occlutech PDA device in 4 defects (Occlutech Holding, Schaffhausen, Switzerland), Amplatzer duct occluder I (ADOI) in 2, LifeTech Konar-MF occluder in 2, Occlutech VSD device in 2, and Amplatzer VSD device in 1 patient. Figure 1 illustrates a few images of different devices. Selection of specific device depended on type and size of defect, in addition to device availability at the time of procedure. Median defect size was 4mm (3-11mm).

All procedures were done with both fluoroscopic and echocardiographic guidance. In 19 patients (50%) TEE was used. In the other 50%, TTE was used. There were two incidences of device embolization, both retrieved and the defect closed using a larger device. Approach for crossing the defect was from right ventricle to left ventricle in 16 patients, and from left ventricle to right ventricle in 22 patients. In those with left ventricle to right ventricle approach, 14 patients had closure via arterial catheter (13 ADOII device, and 1 Konar MFO device), while 8 required arteriovenous loop.

All symptomatic patients reported improved symptoms following the procedure, and all patients who were on anti-failure medications prior to the procedure were off medications within one month of the closure. Regarding complications of the procedure, one patient had transient epistaxis that was treated conservatively, two patients developed transient intermittent junctional acceleration rhythm which resolved spontaneously within a few days. No patient developed atrioventricular block.

Echocardiographic data

For evaluation of the effect of percutaneous closure of VSD on echocardiographic left ventricular and left atrial size, we included 34 patients with isolated VSD closure. The 4 patients who were not included in the analysis had either a VSD and a PDA (2 patients), or multiple VSDs that were not closed in a single procedure (one had muscular percutaneous VSD closure followed by surgical inlet VSD closure, the other had muscular percutaneous VSD closure followed by perimembranus percutaneous VSD closure 6 months later).

At initial follow up with a mean duration of 4 weeks (± 2) following the procedure, 8 (21%) patients had residual shunting (1 moderate, 4 small, and 3 trivial), At the last follow up with mean duration of 18 months (± 19), only 6 patients (16%) had residual shunting, (4 trivial and 2 small). There was no aortic valve regurgitation, and only three patient had trivial tricuspid valve regurgitation.

All echocardiographic parameters of left sided volume overload decreased when compared to baseline (Figure 2), the major drop was noted by the first follow up within the first month of the procedure (LV Z-score dropped from 1.53 ± 0.87 to 0.31 ± 0.68 ($p=0.0001$), LA Z-score dropped from 2.78 ± 0.71 to 1.54 ± 1.13 ($p= 0.0001$), and LA/AO ratio dropped from 1.61 ± 0.27 to 1.33 ± 0.18 ($p=0.0001$)). In addition, there was further drop of LA/AO ratio at the last follow up to $1.23 (\pm 0.12)$ ($p=0.022$). Other parameters also showed further drop at last follow up with LV Z-score dropping to $0.12 (\pm 0.95)$, LA Z-score dropping to $1.24 (\pm 0.8)$, however without statistical significance ($p=0.117$ and 0.095 , respectively). There was no change in the left ventricular ejection fraction before and after the procedure (69.7 ± 5.5 at baseline, 68.8 ± 6.5 at first follow up, and 70.2 ± 4.5 at last follow up).

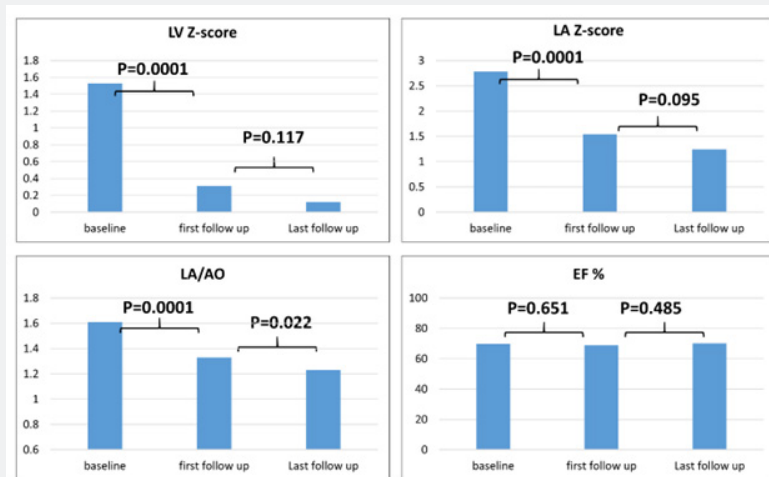


Figure 2: Echocardiographic results of left ventricular diastolic dimention (LV Z-score), left atrial dimention (LA Z-score), left atrial to aortic root ratio (LA/AO), and left ventricular ejection fraction (EF) in 34 patients who underwent percutaneous closure of isolated ventricular septal defect at baseline, at first follow up with mean duration of 1 month, and at last follow up at mean duration of 18 months after the closure showing significant decrease of left ventricular, left atrial dimention, and LA/AO ratio at first follow up, with further decrease of LA/AO ratio at last follow up, with no change in ventricular ejection fraction.

Discussion

Left to right shunt in VSDs increases pulmonary blood flow and pulmonary venous return, eventually causing left atrium and left ventricle dilatation. Larger defects typically allow a significant volume overload, following a reduction in pulmonary vascular resistance after birth, and result in early-onset heart failure. Conversely, smaller defects resulting in small shunts generally have a favorable outcome even if persisted. However, systolic dysfunction and a reduction in the compliance of both ventricles due to chronic pressure and volume overload have been demonstrated even in patients with small VSDs who may show left ventricular dilatation on echocardiography and exhibit diminished left ventricular performance during exercise with a subnormal functional capacity with age [10]. Moreover, many long-term complications such as aortic regurgitation, arrhythmia, double chamber right ventricle, and endocarditis have been observed even in small VSDs [4], making their clinical course not entirely benign.

Cardiac structural abnormalities resulting from chronic volume overload associated with VSDs are shown to be partially reversible in patients operated or had percutaneous closure during adulthood [11,12], while early VSD closure is likely to result in complete restoration of the normal cardiac structure. Thus, the reversibility of cardiac structural changes depends to some extent on the degree and the duration of volume overload [13-15].

Thresholds for defect closure nowadays include patients with echocardiographic dilatation of the left atrium, and the left ventricle. Dilated LA and LV diameters adjusted for body surface area (Z-scores) are found to be associated with significant shunting, with Qp: Qs >1.5 [16,17].

A significant reduction in LV Z- score has been demonstrated following percutaneous and surgical closure of VSDs with no significant difference, favoring device closure over surgery in some cases, as device closure has fewer side effects [18]. Our study demonstrated a significant reduction in LV and LA Z-scores, and LA/AO ratio within one month following the procedure. In addition, these parameters seem to continue to decrease gradually after that, indicating restoration of the normal heart size over time [13,18].

Limitations

Our study is limited by its retrospective nature. In addition, it is a single institution experience with a small number of patients who were followed for a relatively short period. To prove the normalization of cardiac chamber parameters and the lack of device-related complications, a larger sample size of patients and a longer follow-up period are required.

Conclusion

Percutaneous ventricular septal defect closure is a highly successful procedure given the wide variety of available devices.

The elimination of the shunt leads to normalization of cardiac size, namely left ventricle and left atrial sizes, in a short period of time, and can be achieved with a very low risk of complications.

Declarations

- Authors received no financial, technical, or other assistance for this work.
- Authors declare no conflict of interest associated with this research.
- All Authors ascertain that they have approved the content of this work.

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