A simple surgical technique for closure of apical muscular ventricular septal defect

Amit Mishra, MCh, FPCS, Ritesh Shah, MD, Manan Desai, MCh, Ajay Chourasiya, MCh, Hardik Patel, MBBS, Nilesh Oswal, MD, and Dayesh Rodricks, DPT

Objective: Ventricular septal defect (VSD) is among the most common congenital heart diseases encountered in pediatric cardiac patients. Apical muscular VSD constitutes nearly 2% of defects, which may or may not be associated with other congenital heart defects. The purpose of our study is to present our innovative and simple surgical technique using custom-made low-profile polytetrafluoroethylene (PTFE) single disc device for closing multiple apical muscular and isolated apical muscular VSD.

Method: Between January 2010 and July 2013, 17 patients with isolated or multiple apical muscular VSDs with or without associated heart diseases underwent operation at our institute. The apical VSD was closed using our custom-made low-profile single disc polytetrafluoroethylene device. The operative technique and the technique used to prepare the single disc device are detailed.

Results: Seventeen patients of ages ranging from 3 months to 7 years underwent operation over 3 years. One 8-month-old patient with transposition of the great arteries with multiple VSDs died after 35 days due to severe pulmonary artery hypertension and sepsis. Another newborn infant with infracardiac total anomalous pulmonary venous connection with a 4-mm apical VSD also died after 3 days because this VSD could not be identified. All other patients are doing well on follow-up.

Conclusions: The technique described by us has the advantage of apical VSD closure through the left ventricle without left ventriculotomy. Our technique is simple and cost-effective. (J Thorac Cardiovasc Surg 2014;148:2576-9)

CASE DETAILS

Seventeen patients with multiple and isolated apical VSD with or without associated heart defects underwent operation at our institution between January 2010 and July 2013. All patients were diagnosed on the basis of clinical history, examination, electrocardiography, and 2-dimensional echocardiography. Cardiac catheterization was done to assess the operability and, if needed, angiography to confirm the anatomy of the defects. The hospital ethics committee approved the study protocol. Written consent was obtained from both parents, who were informed about the nature, the advantages, and the disadvantages of the procedure.

MATERIAL AND METHOD

The disc device (Figure 1, A and B) was prepared before surgery in the operating room.

The device has 2 parts: a disc and a shaft. The circular disc is made from 2 patches: 1 patch of expanded PTFE (ePTFE) (Gor-Tex patch, W. L. Gore, Flagstaff, Ariz) of thickness 0.6 mm and another patch of PTFE felt (CR Bard, Inc, Delran, NJ) of thickness 0.8 mm that are sutured to each other. The total thickness of the disc is nearly 1.4 mm and its diameter is nearly twice the maximum diameter of the VSD. The shaft is also made from a 2- to 2.5-cm long ePTFE tube (CR Bard, Inc, Delran, NJ) of the same diameter as the maximum size of the VSD. The graft is sutured to the center of the disc in an end-to-side fashion on the Teflon felt side (septal surface of device) using continuous monofilament polypropylene suture and is reinforced at 2 to 3 points. Two extra disc devices with graft sizes 1 mm above and below the required size of the device are also kept ready.
The device is kept in normal saline (0.9%) before use to avoid any air trapping between the 2 discs.

Surgical Technique

Intraoperative transesophageal echocardiography (TEE) and/or transthoracic echocardiography was performed in all patients to confirm the size, number, and position of VSD in relation to the moderator band (Figure 2). The VSD size was measured in 4-chamber and long axis view, the larger of the 2 values was considered for device selection.

After cardioplegic arrest and right atriotomy, the interatrial septum was widely opened. In all cases, a right angle forcep was used to gently probe the apical part of the ventricular septum at the expected site of the apical VSD (Figure 2, A). This angle forcep was passed through the defect and an assistant retracted the mitral valve through the interatrial septum (Figure 2, B). After confirming its position through the defect, the device was caught and held with the help of a right angle forcep (Figure 2, C) and its shaft was gently pulled through the ventricular septum (Figure 2, D), taking care to avoid injuring the neighboring anterior papillary muscle or the moderator band. An illuminating dental mirror was used to confirm the latter and accurate positioning of the device (Figure 2, F). The shaft of the device was sutured at its base to the septal muscles adjacent to the VSD/moderator band using 2 interrupted pledgeted polypropylene sutures closing the lumen of the graft (Figure 2, E). The remaining portion of the shaft was excised. In 1 patient 2 disc devices were used because the VSD was 16 mm in size with 2 openings in front and behind the moderator band. The procedure takes approximately 15 to 20 minutes.

Associated cardiac defects were corrected in the usual fashion and after weaning from cardiopulmonary bypass (CPB), TEE/epicardial echocardiography was routinely performed to confirm adequate repair. Routine weaning from cardiopulmonary bypass (CPB), TEE/epicardial echocardiography was performed to confirm adequate repair. Routine weaning from cardiopulmonary bypass (CPB), TEE/epicardial echocardiography was performed to confirm adequate repair.

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Step-up of 8.2% for which nothing was done because the patient was hemodynamically stable. CPB was reinstituted in 2 patients. In 1 patient who had moderate mitral regurgitation on TEE, the oversize disc was flattening and was interfering with the closure of mitral valve, so the disc was reduced in size. In that patient the diameter of the disc was nearly 2.5 times the diameter of the VSD. In another patient CPB was reinstituted because there was an additional undiagnosed posterior muscular VSD. Both patients recovered uneventfully. No patient had heart block arrhythmia, hemolysis, or any neurologic complications. Median intensive care unit stay was 4 days (range, 2-35 days). Median hospital stay was 11 days (range, 10-35 days). Median follow-up was 12 months (range, 6-30 months) and was complete. At last follow-up, no patient has required reintervention or developed a device-related complication. Follow-up echocardiograms reveal no residual left-to-right shunt.

DISCUSSION

VSD is the most common congenital heart disease encountered in pediatric cardiology clinics. Nearly 2% of VSDs are apical; they may be single or multiple. Apical VSD may be isolated but is usually associated with some other congenital heart diseases. Various standard surgical techniques and interventions are available to close different types of VSD. Technique for closure of apical VSD is still a subject of debate. There is a rapid increase in percutaneous and perventricular closure of muscular VSD. The anatomy of apical septum and echocardiographic assessment was explained in detail by Kumar and colleagues.1 Because the apical septum is the most distal part of the septum and it has trabeculations on the right ventricular side, it is difficult to visualize the margins of the defect clearly from the right ventricular aspect of the septum through the tricuspid valve. The space at the apex is also limited, hence it is difficult to close the defect from the right atrium.2,3

The technique of closing an apical VSD through left ventriculotomy was described by Aaron and colleagues.4 The identification of apical VSD through left ventriculotomy is easy because the left ventricular side of the septum is smooth with no papillary muscle attachment. An apical VSD usually has a solitary opening on the left ventricular (LV) side of the septum. Left ventriculotomy carries the risk of late ventricular aneurysm formation, myocardial dysfunction, and arrhythmias.5 Right ventriculotomy is less dangerous but it may be difficult to find the true margins of the defect between the coarse trabeculations, especially in patients who have pulmonary stenosis.

Earlier we were also closing apical septal defects through right ventriculotomy. Right ventriculotomy may increase the risk of postoperative right ventricular dysfunction,6 especially when apical VSD is associated with conditions like tetralogy of Fallot, transposition of the great arteries, VSD with pulmonary stenosis, and multiple VSD with single
pulmonary artery. Ootaki and colleagues\textsuperscript{7} described the sandwich technique for closing trabecular VSD in 11 patients. However 3 of their patients had significant residual shunt. We believe that it is difficult to stitch at the apical portion of the septum. The only advantage we see of our technique is that the shaft of the device obliterates the VSD and thus the incidence of residual shunt is least with our technique.

Initial attempts at intraoperative device closure of muscular VSD in patients in whom percutaneous closure was contraindicated (eg, small infants) had unsatisfactory results with mortality and failure rates as high as 14\% to 25\% and 20\% to 40\%, respectively.\textsuperscript{8-11} Hybrid procedures and percutaneous device closures are increasingly gaining popularity in closing muscular VSDs, with good results in isolated VSD as well as in patients with associated congenital heart defects.\textsuperscript{12} Percutaneous device closure of VSD has its own limitations, including some anatomical variations of muscular defects like high posterior muscular defects, distal apical defects, and misaligned defects where it is difficult to get access and there is no space for deployment of the right ventricle side of the disc,\textsuperscript{12,13} which is not a limitation of our device. Percutaneous VSD closure is also associated with ventricular arrhythmias, LV pseudoaneurysm formation,\textsuperscript{14} LV dysfunction,\textsuperscript{16} incomplete right ventricle disc expansion with the screw of the disc protruding into the pericardium,\textsuperscript{12,15} and hemolysis in the presence of residual shunt.\textsuperscript{13}

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**FIGURE 1.** A, Photograph of disc device. B, Diagram of disc device. ePTFE, Expanded polytetrafluoroethylene; PTFE, polytetrafluoroethylene.

**FIGURE 2.** A, A right angle forceps probed into the VSD. B, Position confirmed while an assistant retracts the mitral valve. C, Right angle forceps now holds the shaft of the device. D, The shaft of the device is pulled through the VSD avoiding injury to mitral subvalvular apparatus and moderator band. E, The device fixed to the RV side of the septum obliterating the graft. F, Position of VSD confirmed with the help of dental mirrors.
The PTFE device (GORE HELEX Septal Occluder)\textsuperscript{17} is available for easily closing atrial septal defects using the transcatheter technique. The PTFE is also used to close the VSD in operating theaters. We combined the PTFE and ePTFE—both of which are readily available in operating theaters—to prepare our indigenous device. The PTFE (Teflon felt) material is soft and spongy, which provides a cushion effect on the septal surface of the device.

The device works in 3 ways for prevention of left to right shunt across the VSD. First, the graft itself obliterates the VSD, because the size of the graft is the same as that of VSD. Second, the disc will also cover the LV side of septum as well as the surrounding area of the VSD. Third, the right ventricular side of the graft is closed with pledgeted stitches with surrounding septal muscles; hence any residual left to right shunt cannot occur around the graft. Multiple apical VSDs usually have a single opening on the left side septum, so closure of apical VSD will also close all other apical VSDs. If any additional VSD is present in the vicinity of the apical VSD, it will also be covered by the disc. The LV surface is smooth and does not have any attachment of papillary muscles so the disc gets firmly adhered to the LV side of the septum. The high pressure in the LV also keeps the disc in its position. Only if the diameter of the disc is more than twice the diameter of the VSD may we see the flattering movement of the disc, as noticed in 1 of our patients. Our disc device being low profile does not interfere with the geometry/contraction of the LV.

Our custom-made device can be tailored and prepared in-house in 10 minutes. Finally, the cost of our single disc device is $100, whereas the cost of a commercially available device is nearly $1300. The use of our device would drastically bring down the expense of the total procedure and make it significantly more affordable for patients of lower economic strata.

CONCLUSIONS

The technique we described has the advantage of apical VSD closure through the LV without left ventriculotomy. Our technique is simple and cost-effective.

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