	VOLUME-7, ISSUE-10, OCTOBER-2018 • PRINT ISSN No 2277 - 8160	
Sunt FOR Reserves	Original Research Paper	Surgery
Piternational	VENTRICULAR SEPTAL DEFECTS WITH NEAR SYSTEMIC PULMONARY PRESSURES -AN INSTITUTIONAL EXPERIENCE	
Dr. Jaikaran G.k	M.Ch Professor and HOD, Department of Pediatric Cardiothoracic Surgery, Institute of Child Health (ICH), Egmore, Chennai.	
Dr. Sivasankaran . K*	Final year M.Ch Post Graduate, Madras Medical College, Chennai. *Corresponding Author	
ABSTRACT Ventricular Septal Defects (VSD) with severe pulmonary hypertension (PHT) patients are the difficult ones to intervene as they carry increased risk of postoperative morbidity and mortality especially when pulmonary pressures are near to systemic pressures. We share our experience of 31 such cases managed by double flap technique and use of		

pulmonary vasodilators.

KEYWORDS : Ventricular Septal Defects , Pulmonary Hypertension , Double Flap technique.

INTRODUCTION:

Patients with VSD in which PHT is severe have traditionally thought to be inoperable because of the high risk of operation. Zhou et al reported the use of Unidirectional valve patch closure of cardiac septal defects with severe PHT. Excellent improvement in Functional state was noted. These results were corroborated by Ad et al. We share our experience of 31 such cases managed by double flap technique and use of pulmonary vasodilators.

MATERIALS:

- No of cases:31
- Duration of study: August 2012 to December 2017

INCLUSION CRITERIA:

- Primary diagnosis VSD without any coexisting pathology
- Echo showing bidirectional shunt with severe PHT
- · Cardiac catheterization showing reversibility of PHT

EXCLUSION CRITERIA:

- Cases with sub systemic pulmonary pressures
- VSD with coexisting anomalies
- Eisenmengerised VSD

METHODOLOGY :

31 patients having VSD with severe PHT were included in the study.

OUR PROTOCOL:

The children were treated with preoperative oral sildenafil; Intraoperatively half the dose of phenoxybenzamine was given directly into the main pulmonary artery just before cross clamp Intraoperatively double flap technique was followed (goretex-PTFE patch); Weaned off from bypass with infusion of phenoxybenzamine, milrinone, inotropic support; Postoperatively electively ventilated for 24-72 hours; Elective tracheostomy done if unable to wean from ventilator after 4 days; Followed up with oral Sildenafil / Bosentan and ECHO.



Figure 1 pre bypass aortic and PA pressures

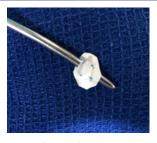


Figure 2 Flap made using goretex patch and prolene sutures



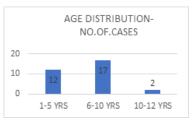
Figure 3 Intraoperative image showing closure of VSD using the patch



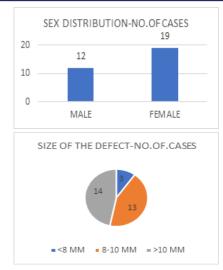
Figure 4 Post bypass Aortic and PA pressures

RESULTS:

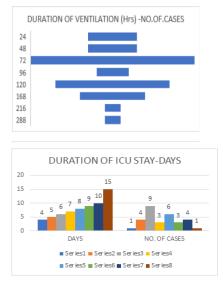
Of the 31 children, 12 were in the age group of 1-5 years, 17 in the age group of 6-10 years, 2 in the age group of 11-12 years.



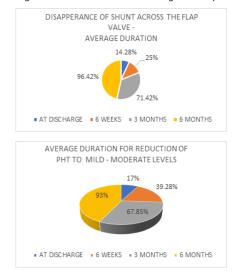
VOLUME-7, ISSUE-10, OCTOBER-2018 • PRINT ISSN No 2277 - 8160



Average duration of ventilation was 114 hours.13 children needed tracheostomy and the average duration of ICU stay was 7-8 days.



Complications encountered were PHT crisis, Right ventricular failure, Heart block, Reintubation. Mortality was encountered in 3 cases the cause being right ventricular failure, septicaemia. All the remaining 28 children were stable during follow up.



Shunt across the flap valve disappeared in 27 children by 6 months and PHT regressed to mild / moderate levels in 25 children in the same duration.

CONCLUSION:

Ventricular Septal defects with severe pulmonary hypertension can be effectively managed by double flap patch closure surgical technique along with judicious use of pulmonary vasodilators. The protocol followed in our institution using double flap technique, pulmonary vasodilators, ventilatory support for the management of patients with VSD/PHT may be used as an alternative to the use of nitric oxide in the effective postoperative management of this subgroup of patients

DISCUSSION:

Two main determinants of the Pressure gradient across VSD and Shunt volume across VSD are 1.Size of defect 2. Pulmonary vascular resistance.

Pulmonary Vascular Resistance Index

- <4 units/BSA --- Normal
- 4-5 --- Mildly elevated
- 5-8 --- Moderately elevated
- >8 --- Severely elevated

When the index is > 8 pulmonary vasodilator challenge is performed (hyperventilation / 100% Fio2 / Nitric oxide / Pulmonary vasodilators) and if the index falls to </= 7, the response is considered favourable and surgery is advised.

HEATH-EDWARD CLASSIFICATION

Pulmonary Vascular resistance in patients with VSD is corelated with histologic severity of the Hypertensive pulmonary vascular disease classified by Heath et al. Histologic reversibility of Pulmonary vascular disease after closure of VSD has not been documented. Presumably pulmonary vascular disease of grade 3 or greater severity is not reversible.

- Grade I hypertrophy of the media of small muscular arteries and arterioles.
- Grade II intimal cellular proliferation in addition to medial hypertrophy.
- Grade III progressive intimal proliferation and concentric fibrosis.
- Grade IV "plexiform lesions"
- Grade V angiomatous and cavernous lesions and hyalinization of intimal fibrosis.
- Grade VI necrotizing arteritis.

Surgical correction has to be done before irreversible damage to pulmonary vasculature occurs.the primary investigations include 1.Echocardiogram - To assess the site & size of VSD, the direction of the shunt and to assess the PHT severity.

2. Cardiac catheterisation

a)To assess RA, RV, PA & Aorta pressures b)To ascertain the shunt fraction & Shunt diameter c)To determine PVRI & to assess the reversibility of PHT with the use of 100% oxygen & vasodilators.

VSD in Patients with pulmonary hypertension may benefit from repair using Flap valve technique – flap opens right to left if right ventricular pressure exceeds left ventricular pressure in severe PHT, the right ventricle will decompress to the left ventricle, supporting the systemic cardiac output. It is inferred that as the PHT decreases late postoperatively, the flap valve

will close by cicatrix.

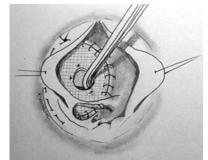


Figure 1Novak's double flap technique

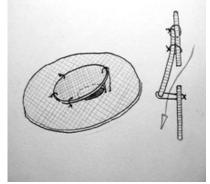


Figure 2 flap valve opening should be into LV directed towards the apex

REFERENCES:

- Kirklin JW, Harshbarger HG, Donald DE, et al: Surgical correction of ventricular septal 1. defect: anatomic and technical considerations. JThorac Surg 33:45-53, 1957.
- 2. Stirling GR, Stanley PH, Lillehei CW: The effects of cardiac bypass and ventriculotomy upon right ventricular function with report of successful closure of ventricular septal defect by use of atriotomy. Surg Forum 8:433-438, 1957.
- Heath D, Edwards JE: The pathology of hypertensive pulmonary vascular disease: a 3. description of 6 grades of structural changes in the pulmonary arteries with special reference to congenital cardiac septal defects. Circulation 18:533–543, 1958.
- Collins G, Calder L, Rose V, et al: Ventricular septal defect: clinical and hemodynamic 4. changes in the first five years of life. Am Heart J 84:695-701, 1972.
- Hoffman JIE, Rudolph AM: The natural history of ventricular septal defects in infancy. 5. Am J Cardiol 16:634-638, 1965.
- б. Bacha EA, Cao QL, Starr J, et al: Periventricular device closure of muscular ventricular septal defects on the beating heart: technique and results. J Thorac Cardiovasc Surg 126:1718-1723, 2003.
- Berner M, Behetti M, Ricou B, et al: Relief of severe pulmonary hypertension after 7. closure of a large ventricular septal defect using low dose inhaled nitric oxide. Intensive Care Med 19:75–79, 1993.
- Yeager SB, Freed MD, Keane JF, et al: Primary surgical closure of ventricular septal 8. defect in the first year of life: results in 128 infants. J Am Coll Cardiol 3:1269-1273, 1984.
- Weintraub RG, Menahem S: Early surgical closure of a large ventricular septal defect: influence on long-term growth. J Am Coll Cardiol 18:552–557, 1991.
 Zhou Q, Lai Y, Wei H, Song R, Wu Y, Zhang H. Unidirectional valve patch for repair of
- cardiac septal defects with pulmonary hypertension. Ann Thorac Surg 1995;60:1245
- 11. Double patch closure of ventricular septal defect with increased pulmonary vascular resistance Novick, William M et al. The Annals of Thoracic Surgery, Volume 66, Issue 5, 1533-1537