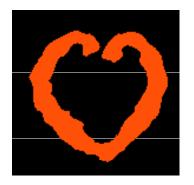


Department of Pediatric Cardiology & Adult with Congenital Heart Disease

IRCCS- Policlinico San Donato San Donato Milanese – Milano



Massimo Chessa – Cosa possiamo fare senza la cardiochirurgia. Come è possibile evitare l'intervento cardiochirurgico e riparare l'anomalia anatomofunzionale.





In the modern era of emergent pharmacological and percutaneous therapy, ventricular wall ruptures are rare with interventricular septal rupture accounting for 0.2% of all myocardial ruptures

Crenshaw BS, Circulation 2000; 101: 27-32.





Percutaneous repair of acutely ruptured myocardium is challenging

The first experience by Landzberg and Lock involved a single centre experience of percutaneous closure of post infarction VSD's using the older closure devices, (the Clam-shell double umbrella and the Cardio SEAL).



Landzberg MJ. Semin Thorac Cardiovasc Surg 1998; 10: 128-132.

Crenshaw BS, Circulation 2000; **101**: 27-32. Menon V, J Am Coll Cardiol 2000; **36**: 1110-1116.

Adverse Substrate

- Defect too big
- Cardiogenic shock/Renal failure/Sepsis
- Advanced age
- Surgical turn-downs
- Co-Morbidity



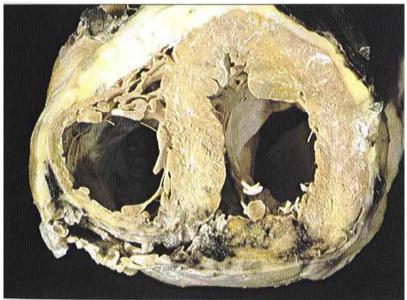
Items of concern

•VSD size and morphology

Congenital VSD

Post MI VSD







PSD

Items of concern

•VSD size and morphology



Who should we select?

- All comers
- Patients in good condition at referral
- After stabilization (IABP), observation and stenting of Infarct related artery
- Survivors for > 3 weeks (Trial of life)
- Residual/recurrent defects



Ideal Patient

Residual defect with chronic heart failure

Survivors of "Trial of Life"

sp



Who should we turn down?

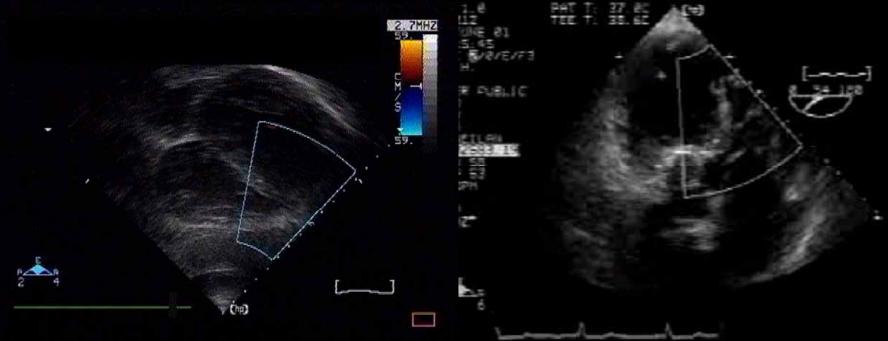
- Moribund patients despite IABP/Inotropes
- Unsuitable defect too large
- Acutely presenting large defects where early surgery may produce more survivors

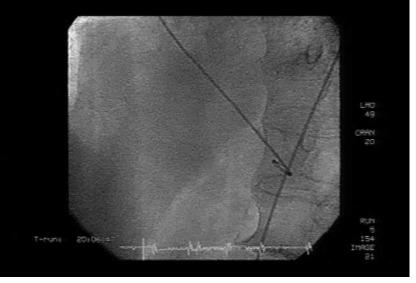
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sb

Imaging for Case Selection

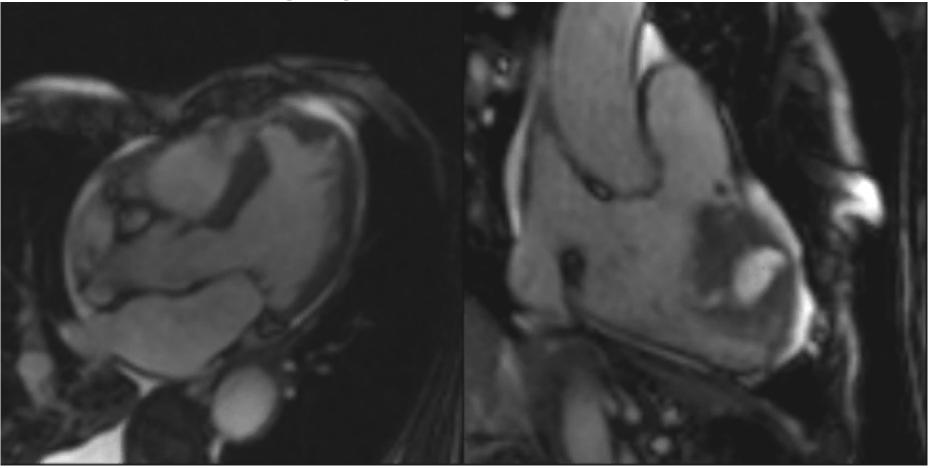








Imaging for Case Selection



Courtesy Dr Mark Turner





Trying to overcome the initial difficulties the Amplatzer Muscular Device was used.

Chessa M. J Invasive Cardiol. 2002 Jun;14(6):322-7.

A

AMPLATZER Post-Infarction Muscular VSD Occluder (AGA Medical Corp.)

AMPLATZER® Muscular VSD Occluder (Post Myocardial Infarction)

ORDER NUMBER	DEVICE SIZE (DIAMETER OF WAIST)	DEVICE SIZE (LENGTH)	MINIMUM RECOMMENDED SHEATH SIZE
9-VSDMUSCPI-016	16 mm	10 mm	9 French, 45° or 180° Curve
9-VSDMUSCPI-018	18 mm	10 mm	9 French, 45° or 180° Curve
9-VSDMUSCPI-020	20 mm	10 mm	10 French, 45° Curve
9-VSDMUSCPI-022	22 mm	10 mm	10 French, 45° Curve
9-VSDMUSCPI-024	24 mm	10 mm	10 French, 45° Curve





AMPLATZER Post-Infarction Muscular VSD Occluder (AGA Medical Corp.)

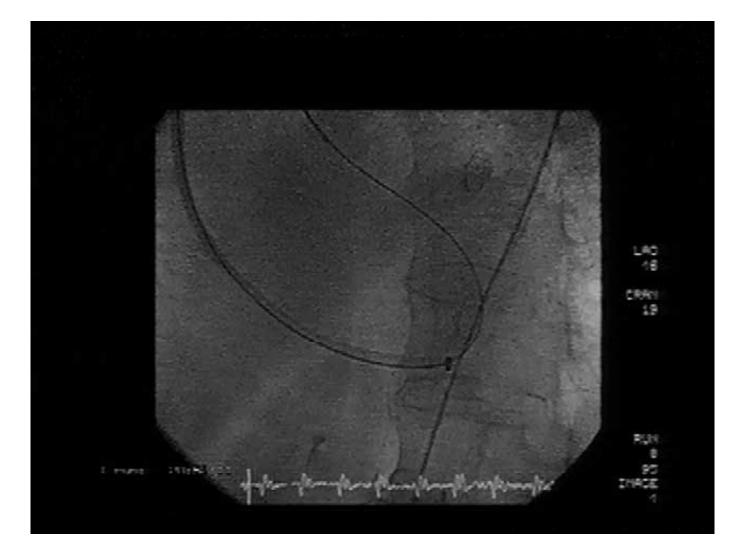
The Amplatzer postinfarction muscular VSD device (PIMVSD) is a nitinol construct with sizes ranging from 16-24 mm and a connecting waist of 10 mm in length.

The LV and RV disks are 5 mm larger than the waist. Amplatzer devices are self-centering and obtain occlusion of the VSD via in situ thrombosis of the waist. In addition, Amplatzer devices are retrievable after deployment for repositioning if the initial result is unsatisfactory.

They are not retrievable, however once the procedure is completed.

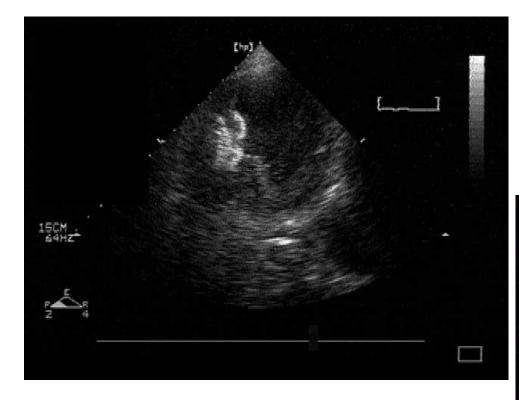


Acute post-MI VSD: device implantation





Early after the procedure



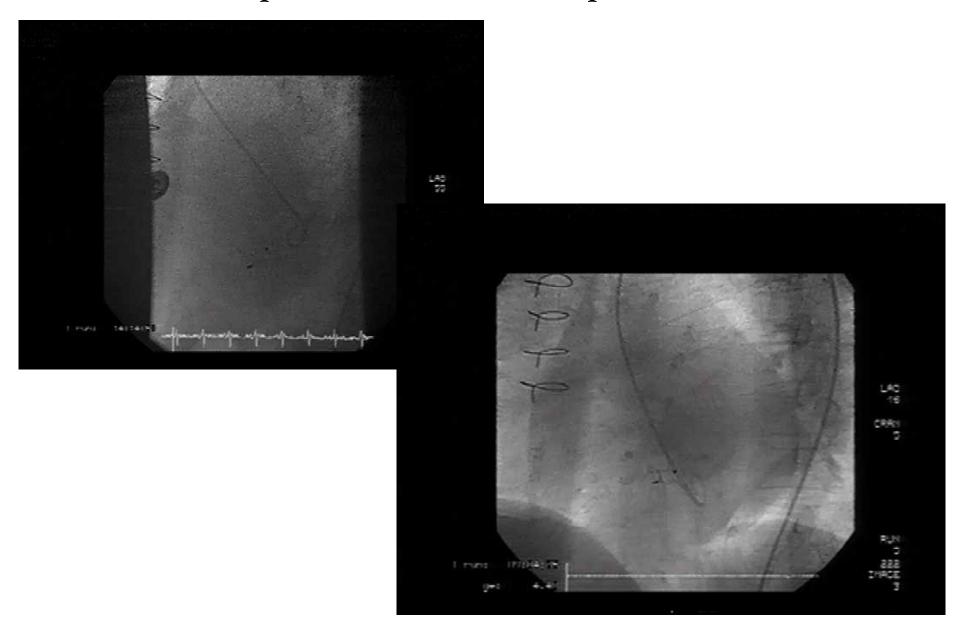
48 hours later







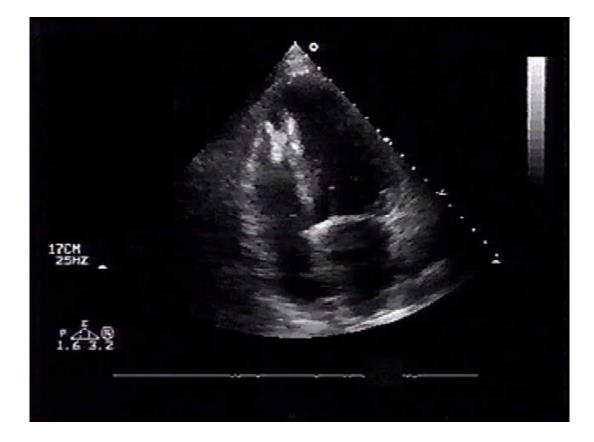
Acute post-MI VSD: device implantation





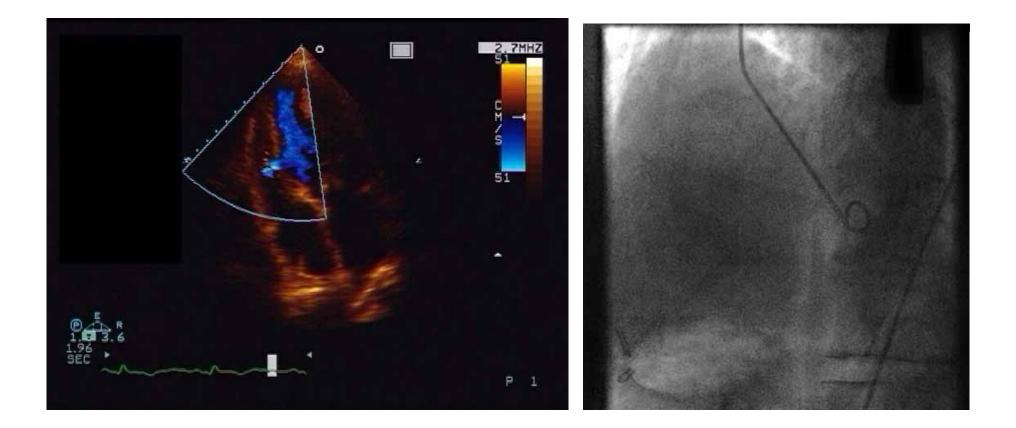
SD SD





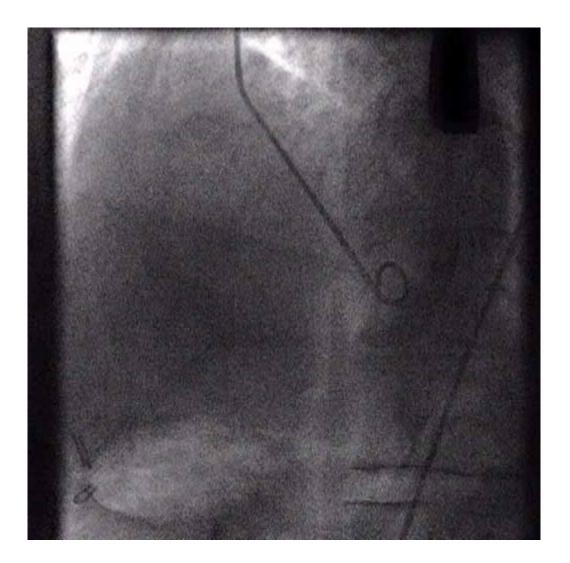


Non acute post-MI VSD





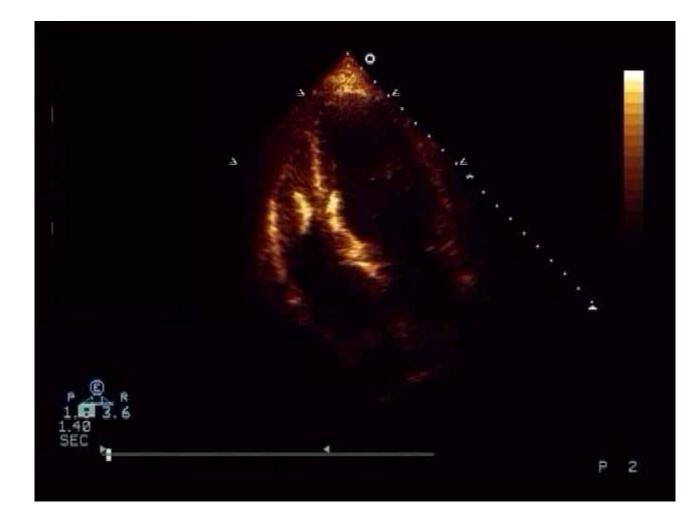








Echo post procedure





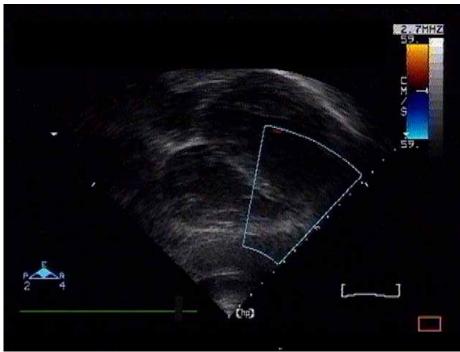
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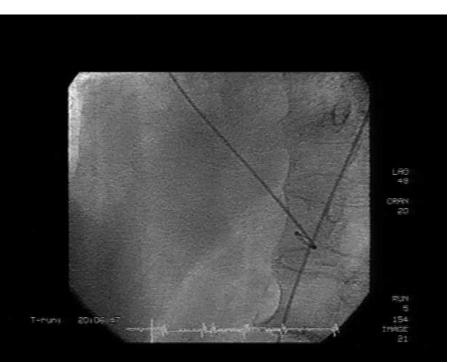




What we learned

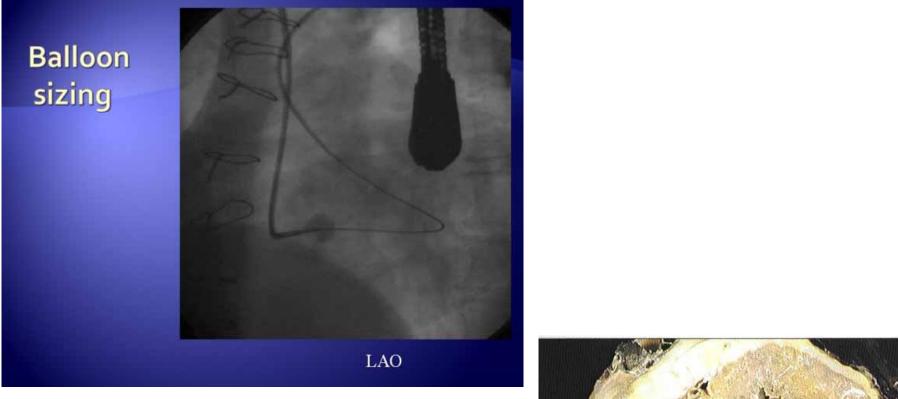
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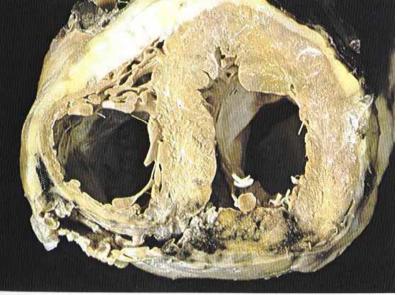






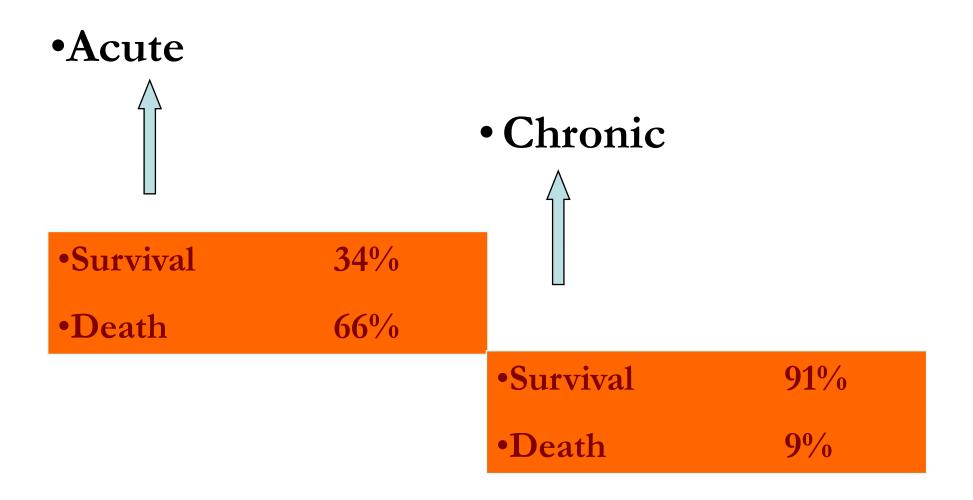
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What we learned





What we learned

Complications

1. Arrhythmias

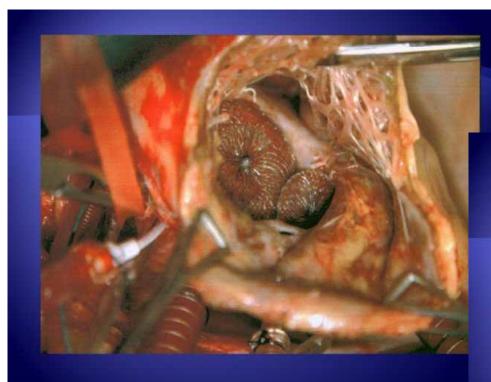
2.Avulsion of tricuspid septal leaflet while passing sheath though VSD

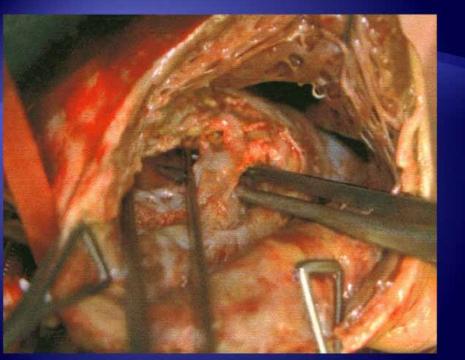
3. Haemolysis managed medically



What we learned

Cross-over to surgery













*ASO device implantation in acute phase of MI gave poor results.

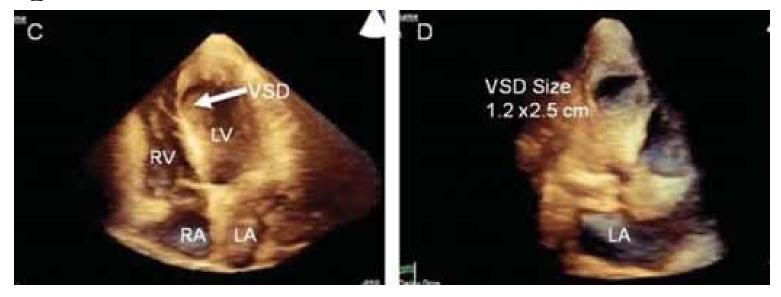
*Good results are obtained for "non acute"VSD's or residual defects after surgery

* Percutaneous closure may be a bridge to surgery





*3D Imaging both before and during will further improve the efficacy of the procedure







*Improvements in device and delivery systems are also required



Thank you for your attention



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