

# ***Outcome of transcatheter closure of Post MI ventricular septal defects***

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# ***Disclosure Statement of Financial Interest***

## ***Saibal Kar***

Within the past 12 months, I or my spouse/partner have had a financial interest/arrangement or affiliation with the organization(s) listed below.

### **Affiliation/Financial Relationship**

- Grant/Research Support
- Consulting Fees/Honoraria
- Other Financial Benefit

### **Company**

- Abbott Vascular, Boston Scientific, Gore Medical, CardioKentix, St Jude Medical
- Abbott Vascular, Boston Scientific, Gore Medical, Coherex
- Coherex, Biosensors International

# Background

- Ventricular septal defect is a rare (0.3%) mechanical complication of myocardial infarction.. It results in poor cardiac output, multiorgan failure and death.
- Bimodal presentation with a higher incidence in the first 24 hours, and then again 3-5 days after initial MI.
- Natural history data suggest a 90% mortality rate within 2 months of diagnosis in patient who do not undergo defect closure. (3)

Egidy Assenza et al. Circ 2013  
Lee WY, et al. Arch Intern Med. 1962

# Treatment options

- Expert opinion and guideline suggest that immediate cardiothoracic surgical repair should be considered.
- Surgical mortality is very high.
- Transcatheter closure of post-MI VSD is an increasingly utilized alternative approach for this population..

Cooley DA., et al. Surgery, 1957

Deja MA., et al. Eur J Cardiothorac Surg, 2000

Papdopoulos N., et al. Ann Thorac Surg, 2009

Egidy Assenza et al. Circ 2013

# Transcatheter closure

- There is a paucity of data of the outcomes following transcatheter closure.
- Survival rates in the literature of transcatheter survival is reported to be 35-75%, and vary significantly in this heterogeneous population.

Sathananthan, J., et al., J Invasive Cardiol, 2013.

Calvert, PA., et al., Circulation, 2014.

Thiele, H., et al., Eur Heart J, 2009.

Assenza, GE., et al., Circ Cardiovasc Interv, 2013.



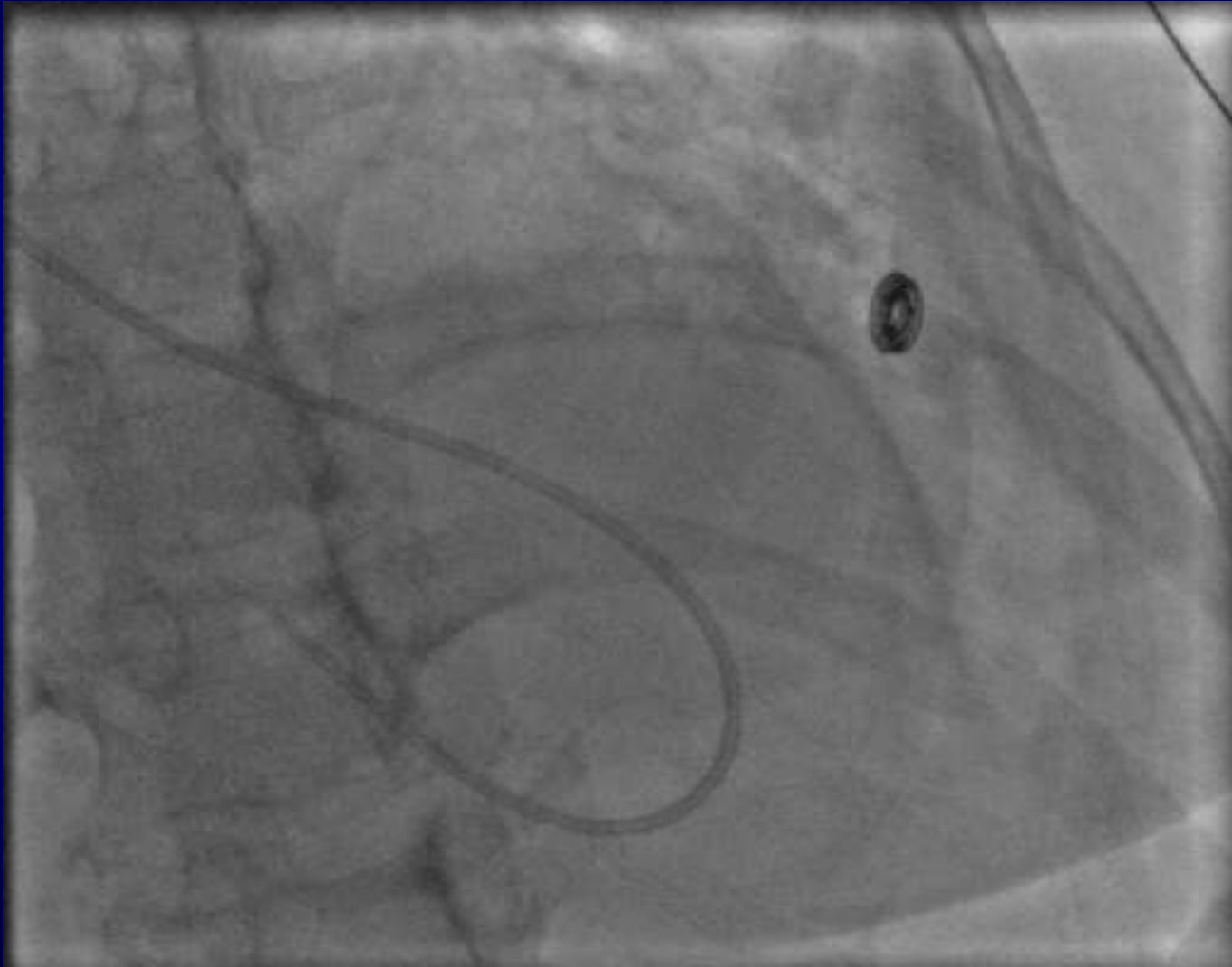
# Post MI VSD : challenges

- Patient are usually in shock
- Septum is friable, and there may be more than one defect
- Cross the defect with balloon floatation catheter
- Careful during creation of AV loop (prevent cheese cutting of septum)
- Often multiple devices are required

# Baseline LVG

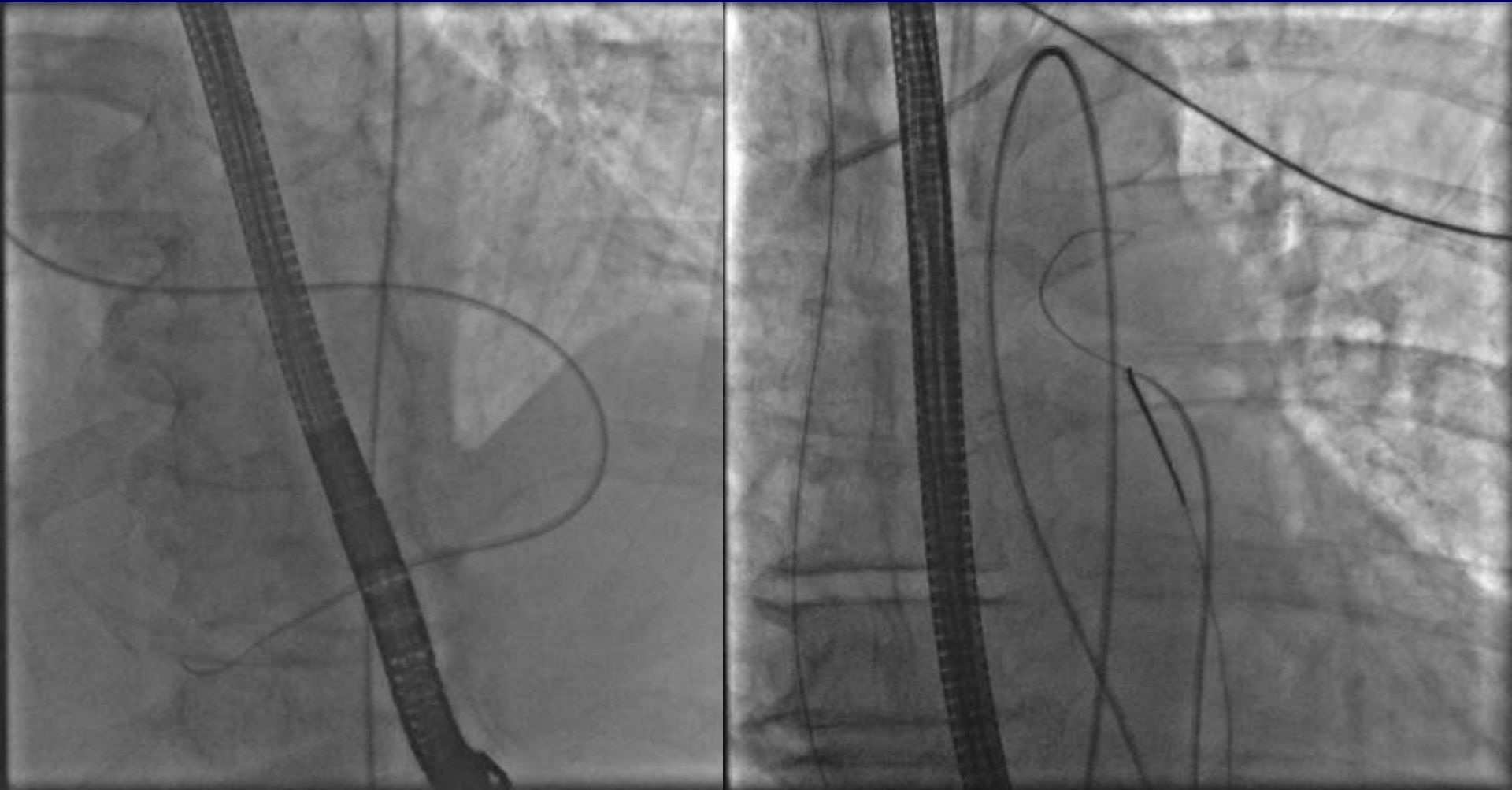


# Crossing the defect with the Balloon tipped catheter

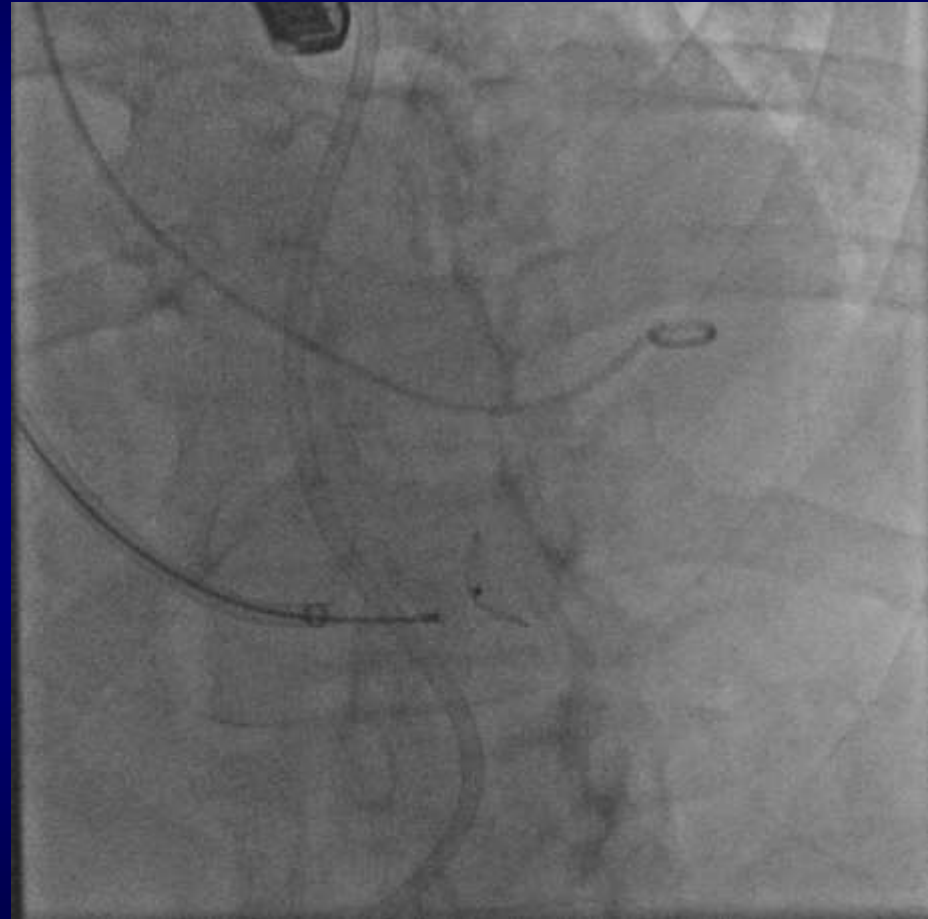




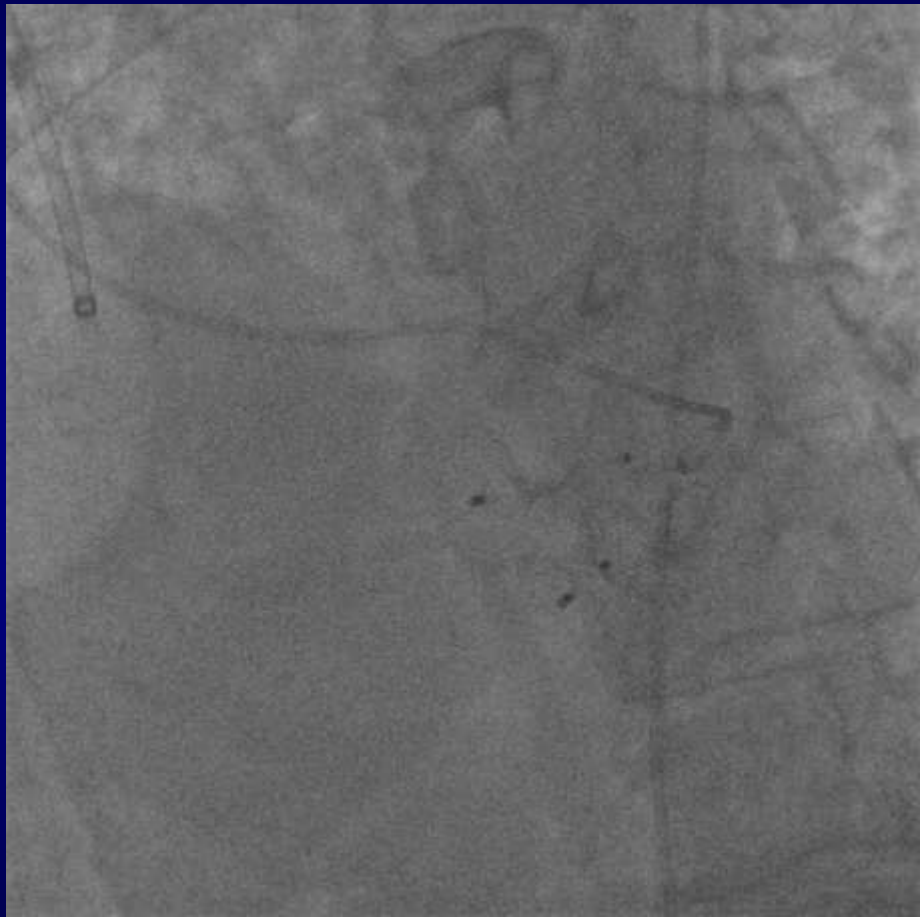
# Crossing the defect with a balloon tipped catheter and creating Arteriovenous Loop



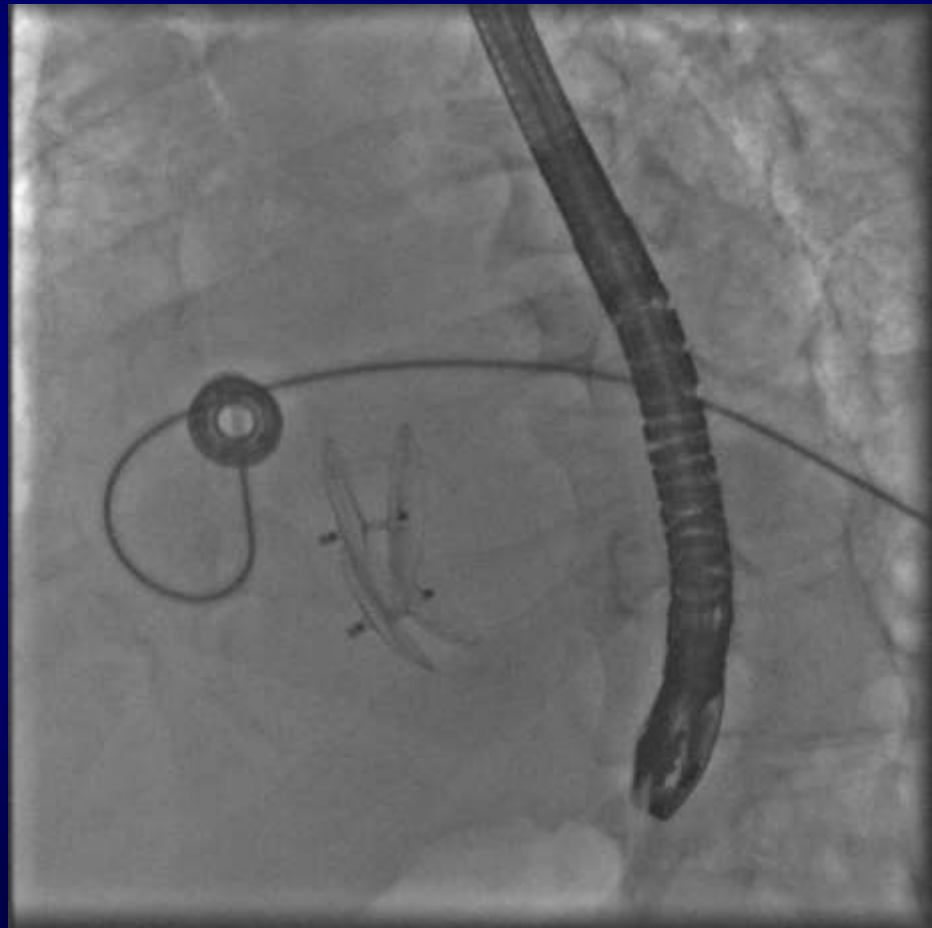
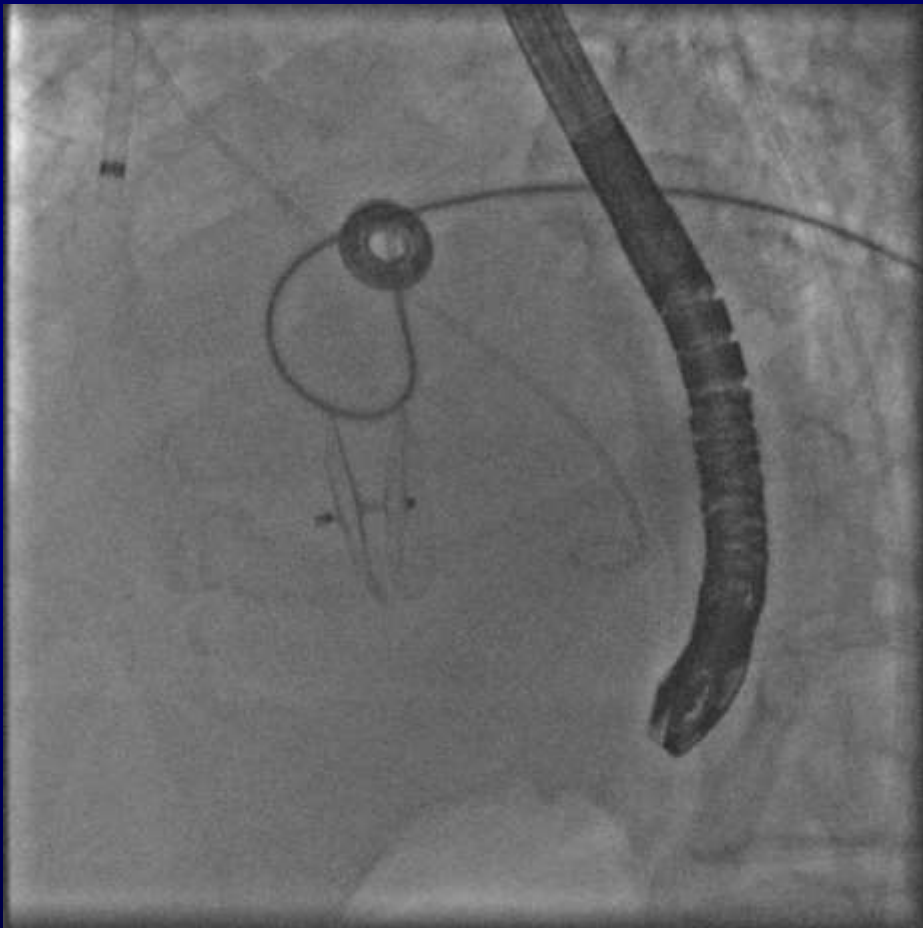
# Deployment of a VSD occluder 16mm and Final LVG



# Residual shunt closure



# Use of two Cribriform devices to close Post MI VSD



# Residual shunt closure





# Transcatheter Post-MI VSD Closure Experience at Cedars Sinai Medical Center





# Cedars Sinai Experience

21 patients between 2005 to 2015.

On Presentation	N = 21
Age (SD)	72 ± 12
Male (%)	15 (71%)
Heart Failure (%)	15 (71%)
Average NYHA FC (SD)	3.7 ± 0.5
Cardiogenic Shock (%)	14 (67%)
IABP / Impella (%)	11 (52%)
Intubation (%)	9 (43%)
Transfer from outside hospital (%)	12 (57%)

Comorbidities	N = 21
HTN (%)	15 (71%)
HL (%)	10 (48%)
DM (%)	6 (29%)
CKD (%)	4 (19%)
Active tobacco use (%)	2 (10%)
Prior MI (%)	14 (67%)

# Result

Closure Device	N = 26
Amplatzer Cribriform	10 (38%)
Amplatzer ASD	5 (19%)
Amplatzer VSD	10 (38%)
Amplatzer Ductal Occluder	1 (4%)
Average Size 1 <sup>st</sup> Device (SD)	19 ± 8 mm
Average Size 2 <sup>nd</sup> Device (SD)	22 ± 8 mm

- 26 closure devices were implanted in 18 patients.
- Failure to close 3 (14%)
- Three (14 %) patients underwent a second transcatheter closure
- Two (10%) patients had procedure-related death.
- The primary outcome of survival to discharge and 30 day survival occurred in 13 (62%) patients. The average length of hospitalization was 15 ± 18 days and follow-up was 157 ± 194 days.





# Conclusions

- Ventricular septal defect(VSD) is a rare mechanical complication of MI.
- Prognosis is very poor
- Transcatheter closure is a feasible and effective alternative to surgery.
- Early and often multiple interventions and closure devices are required to salvage these critically sick patients.