



## Outcomes of Ablation of Multiple PVC Morphologies



Ji Hyun Lee, MD, PhD, CEPS

Seoul National University Bundang Hospital

# Korean Heart Rhythm Society

## COI Disclosure

*Ji Hyun Lee*

The authors have no financial conflicts of interest  
to disclose concerning the presentation



# Contents

- 1. Clinical significance of PVCs with multiple morphologies**
- 2. Ablation treatment of multiple PVCs**



# What do PVCs with multiple morphologies indicate ?

1. Higher likelihood of PVC induced cardiomyopathy

Predictors of PVC-induced CMP in univariable analysis (n=107)

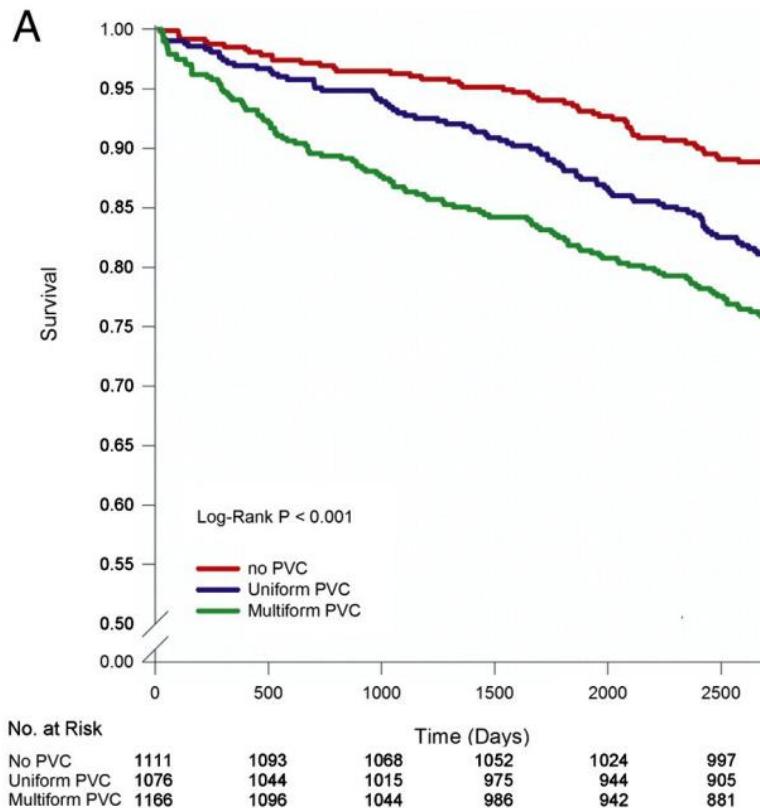
Variable	P	Unadjusted odds ratio
Sex	.01	4.25
Symptom duration	.008	1.01
PVC burden	.0001	1.08
<u>Pleomorphic PVCs</u>	<u>.006</u>	<u>3.11</u>
Interpolation	.022	3.3
Asymptomatic status	.0001	5.4
QRS duration of PVCs	.0001	1.07
Interquartile CoV < 31%	.0001	8.3



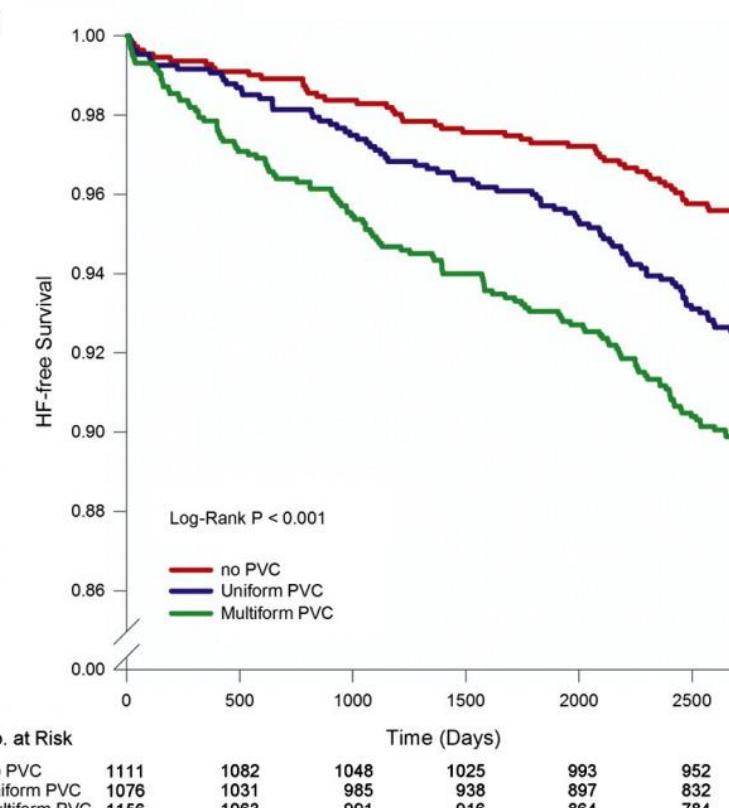
# What do PVCs with multiple morphologies indicate ?

2. Might be associated with underlying subclinical heart disease

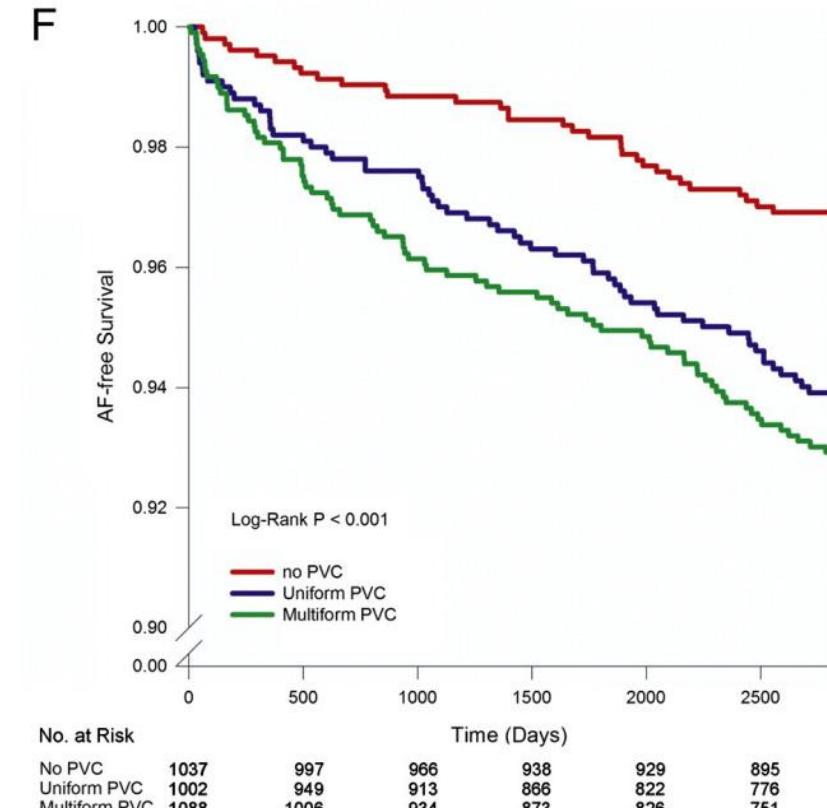
Survival



HF-free survival

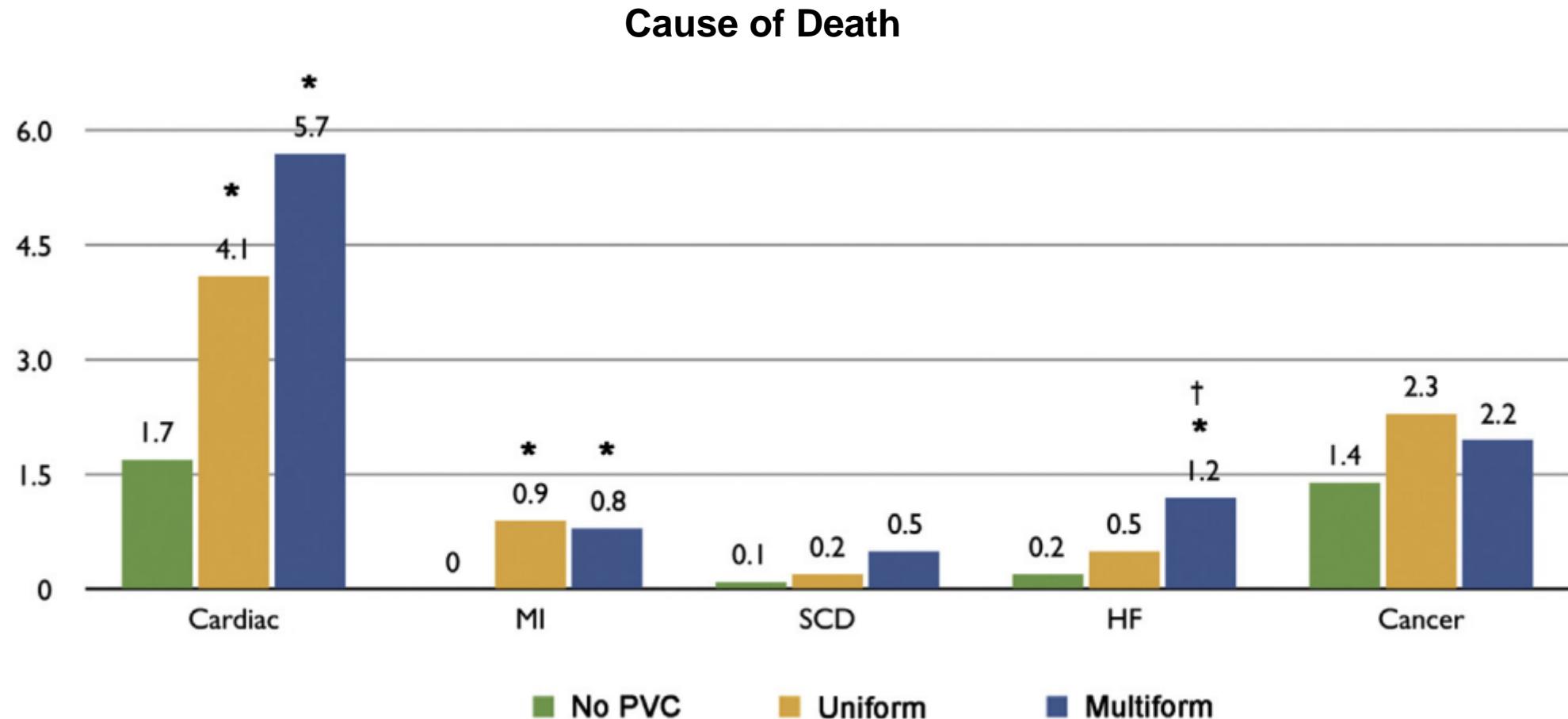


AF-free survival



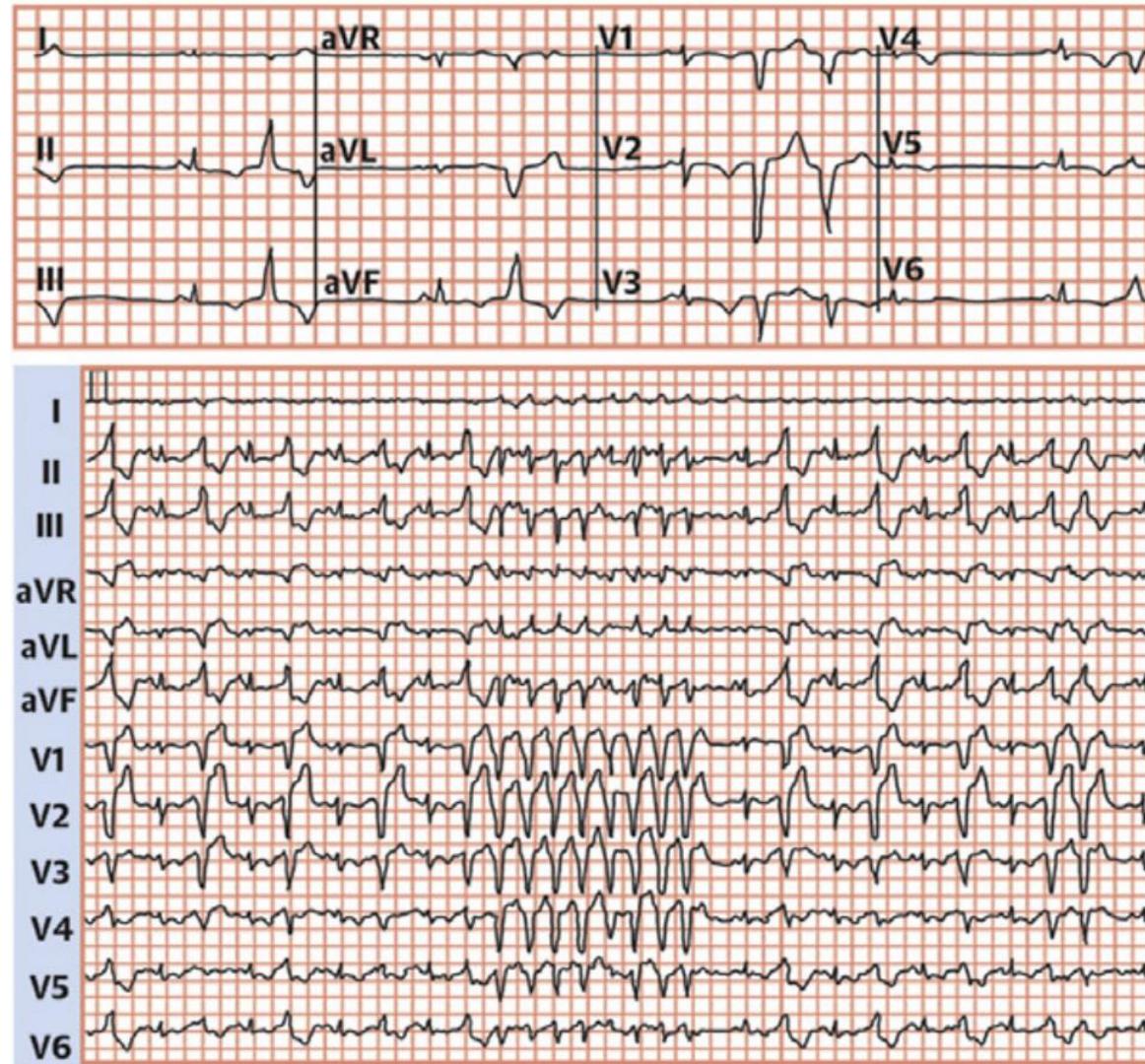
# What do PVCs with multiple morphologies indicate ?

2. Might be associated with underlying subclinical heart disease



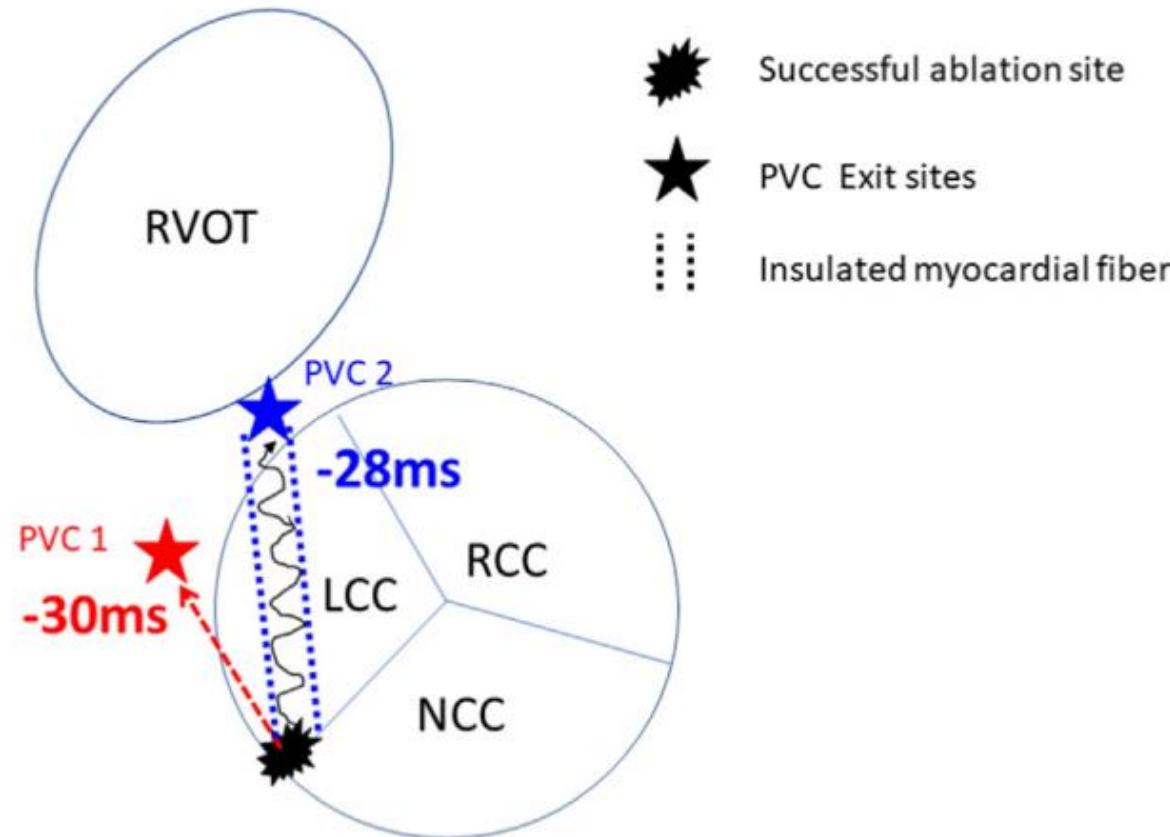
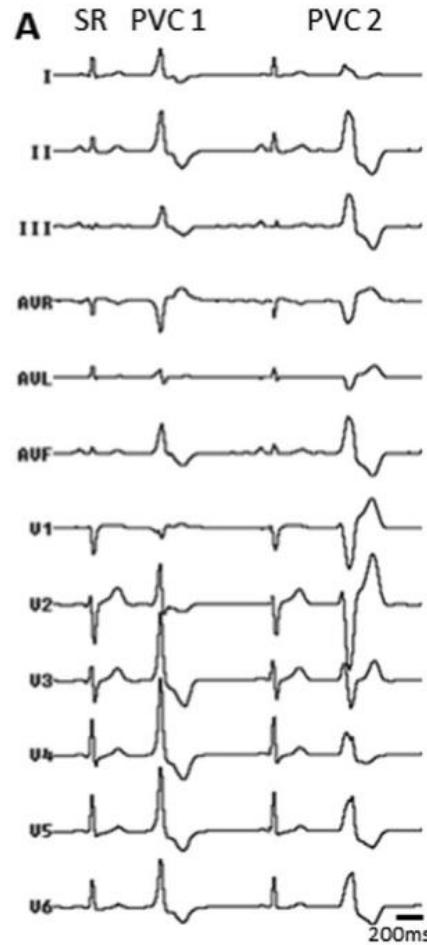
# What do PVCs with multiple morphologies indicate ?

3. Increase the pretest probability for larger arrhythmic substrate as ARVD/C,



# What do PVCs with multiple morphologies indicate ?

4. Occasionally, single focus generating different morphology (OT, papillary m, moderate band etc)



# PVCs with multiple morphologies

Have higher likelihood of incident AF, HF and cardiac death.

PVCs with multiple morphologies are bad in every way.

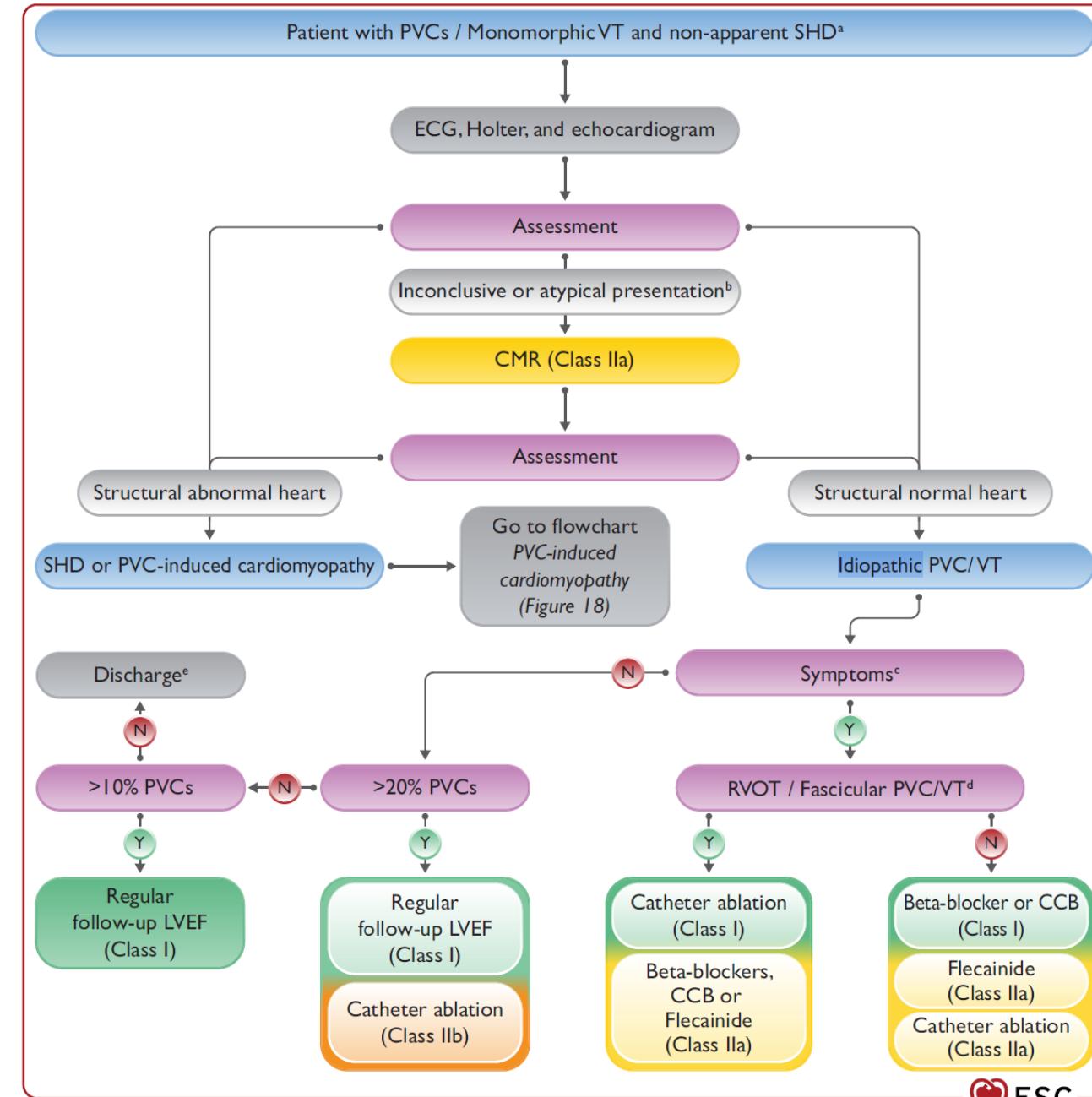


# Contents

1. Clinical significance of multiple PVCs
2. Ablation treatment of multiple PVCs



# ESC guideline



ESC

European Heart Journal (2022) 43, 3997–4126

KHRS 2023

# ESC guideline recommendation for idiopathic PVC/VT

	<b>Ablation</b>	<b>Beta-blocker</b>	<b>CCB</b>	<b>Flecainide</b>	<b>Amiodarone</b>
RVOT/fascicular PVC/VT: Symptomatic, normal LV function	Class I	Class IIa	Class IIa	Class IIa	Class III
PVC/VT other than RVOT/fascicular: Symptomatic, normal LV function	Class IIa	Class I	Class I	Class IIa	Class III
RVOT/fascicular PVC/VT: LV dysfunction	Class I	Class IIa	Class III <sup>a</sup>	Class IIa <sup>b</sup>	Class IIa
PVC/VT other than RVOT/fascicular: LV dysfunction	Class I	Class IIa	Class III <sup>a</sup>	Class IIa <sup>b</sup>	Class IIa
PVC: Burden >20%, asymptomatic, normal LV function	Class IIb				Class III

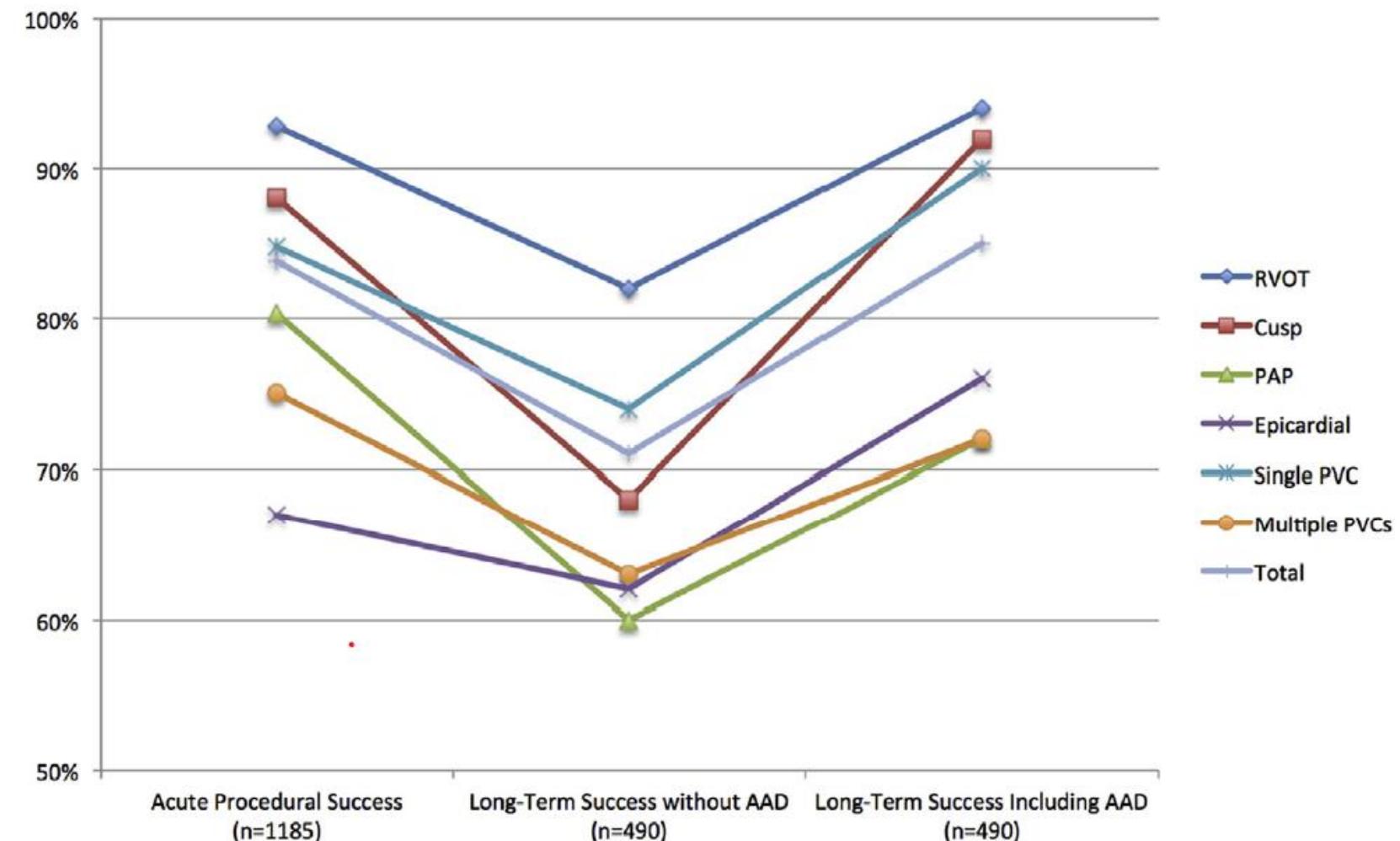


# Outcomes of Ablation of PVC

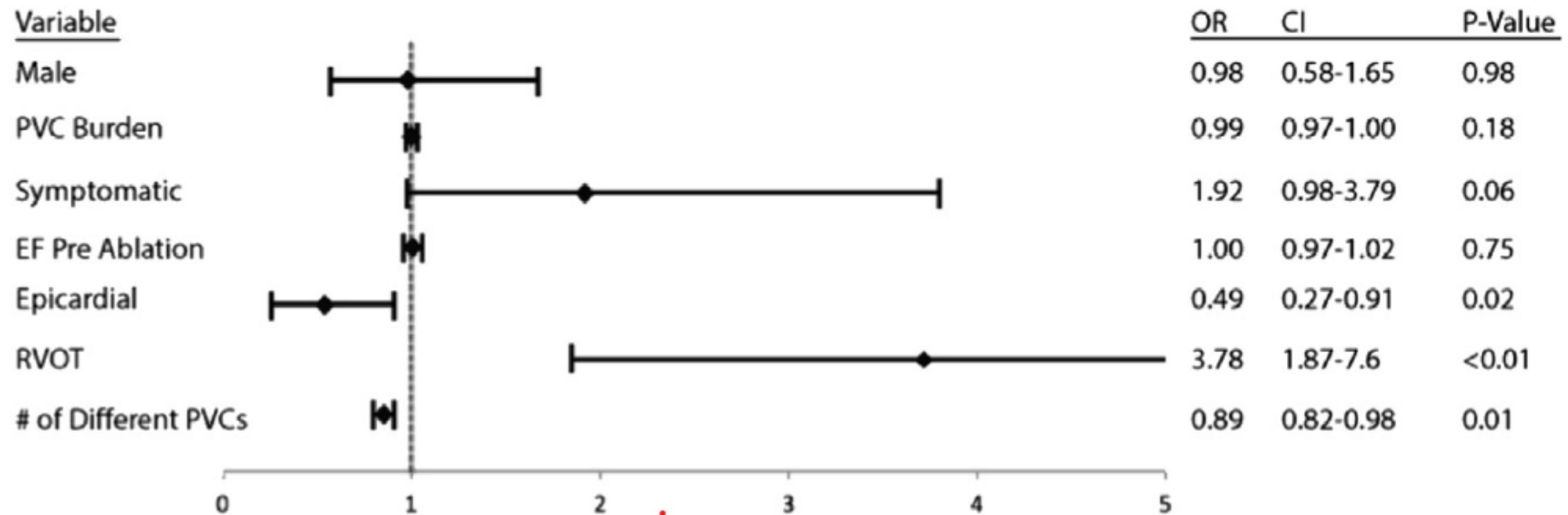
N= 1,185 from 8 centers in the US and Europe

Acute procedure success:  
Elimination of the target PVCs  $\geq$  30 min after ablation

Long-term success:  
80% decrease of PVC burden.



# Predictor of acute procedural success



# **Idiopathic PVC with high recurrence**

1. Papillary muscle PVCs
2. Epicardial PVCs
3. Aortic cusp PVCs
4. Para-Hisian PVC
5. Moderate band PVC



# Characteristics of patients with monomorphic vs pleomorphic PVCs

100 consecutive patients referred for ablation of idiopathic PVCs

	Monomorphic (n = 69)	Pleomorphic (n = 31)	P value
Age, y, mean ± SD	51 ± 14	54 ± 17	.29
Male sex, n (%)	41 (59)	12 (39)	.055
Comorbidities			
Coronary artery disease, n (%)	2 (3)	3 (10)	.15
Hypertension, n (%)	29 (42)	12 (39)	.75
Diabetes mellitus, n (%)	3 (4)	5 (16)	.045
Stroke, n (%)	5 (7)	0 (0)	.12
Hyperlipidemia, n (%)	22 (32)	8 (26)	.54
Valve disease, n (%)	2 (3)	1 (3)	.95
Pacemaker, n (%)	2 (3)	0 (0)	.34
ICD, n (%)	2 (3)	1 (3)	.93
PVC burden			
Overall burden, %, mean ± SD	17.6 ± 12.5	19.9 ± 11.3	.80
Site of predominant PVC site of origin			
Endocardial, n (%)	51 (74)	16 (52)	.028
Intramural, n (%)	12 (17)	6 (19)	.81
Epicardial, n (%)	6 (9)	9 (29)	.008
Predominant PVC region			
Outflow tract, n (%)	53 (77)	20 (65)	.20
Papillary muscle, n (%)	3 (4)	4 (13)	.12
Para-Hisian, n (%)	5 (7)	2 (6)	.89
Mitral annulus, n (%)	4 (6)	1 (3)	.59
Fascicular, n (%)	1 (1)	2 (6)	.18
Other, n (%)	3 (4)	2 (6)	.66
Procedural details			
Repeat procedure, n (%)	14 (20)	6 (19)	.91
Successful ablation, n (%)	62 (90)	22 (71)	.017
Echocardiographic data			
Baseline LV end-diastolic diameter, mm, mean ± SD	51 ± 7	52 ± 6	.80
Baseline LVEF, %, mean ± SD	57 ± 10	55 ± 29	.32

ICD = implantable cardioverter-defibrillator; LV = left ventricle; LVEF = left ventricle ejection fraction; PVC = premature ventricular complex; SD = standard deviation.



# Characteristics of patients with monomorphic vs pleomorphic PVCs

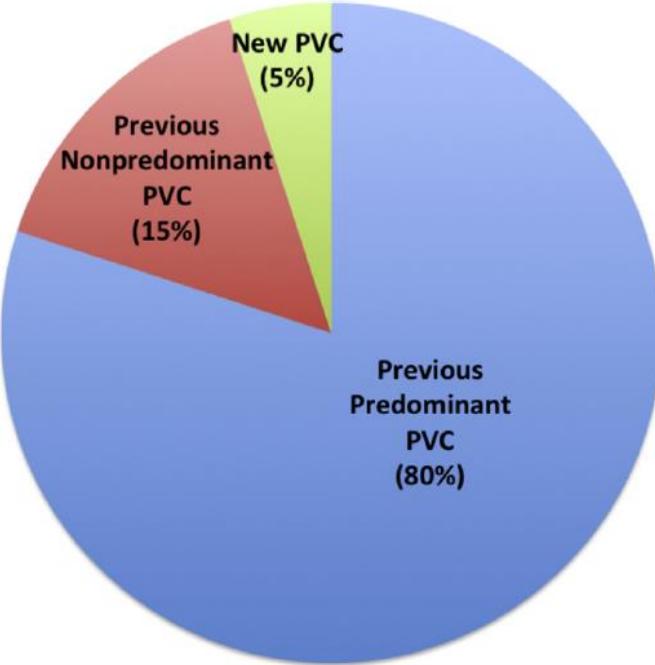
100 consecutive patients referred for ablation of idiopathic PVCs

	Preablation, %, mean $\pm$ SD	Follow-up, %, mean $\pm$ SD	P value
Overall PVC burden	19.8 $\pm$ 9.8	3.1 $\pm$ 4.6	<.0001
Predominant PVC	18.0 $\pm$ 9.6	0.6 $\pm$ 1.3	<.0001
Nonpredominant PVCs	1.8 $\pm$ 2.6	2.1 $\pm$ 4.6	.77
Nonpredominant 1	1.0 $\pm$ 1.1	2.0 $\pm$ 4.6	.31
Nonpredominant 2	0.7 $\pm$ 2.2	0.04 $\pm$ 0.1	.14
New PVCs	N/A	0.5 $\pm$ 1.6	N/A

N/A = not applicable; PVC = premature ventricular contraction; SD = standard deviation.



# Repeat ablation (n=20)

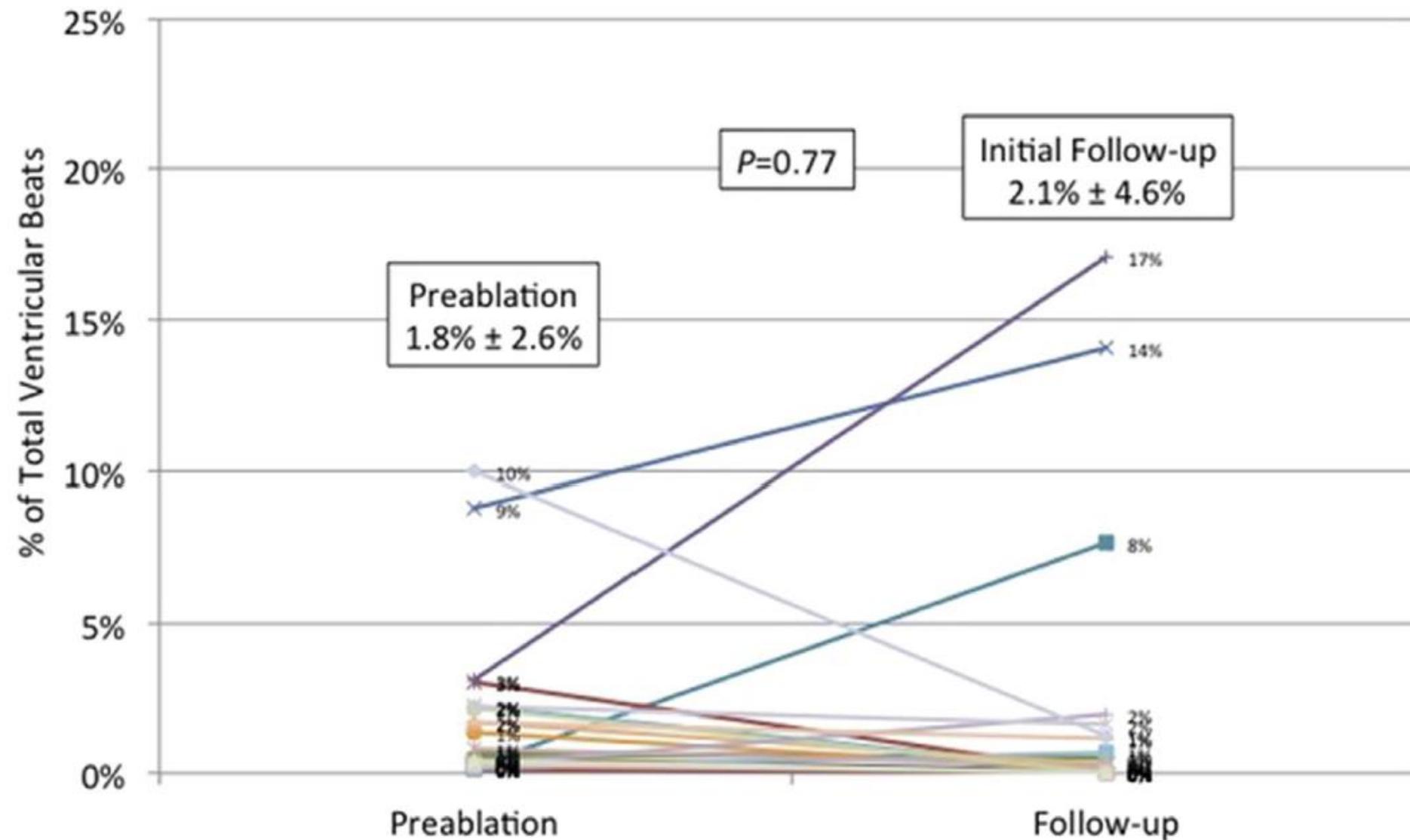


Patient #	Initial procedure predominant origin	Pleomorphic PVCs, n	Procedure success	Interval between procedures, mo	2nd procedure reason	2nd procedure origin (if not same)
1	Tricuspid annulus	0	Yes	7.7	Return of previously predominant PVC	N/A
2	Intramural OFT	0	Yes	45.9, 63.8, 68.2	Return of previously predominant PVC	N/A
3	Epicardial OFT	882	No	12.0	Previously predominant PVC	N/A
4	Endocardial OFT	7	Yes	16.3	Return of previously predominant PVC	N/A
5	PMP	207	No	38.6	Nonpredominant	GCV (Epicardial OFT)
6	Endocardial OFT	9	Yes	6.7	Return of previously predominant PVC	N/A
7	Para-Hisian	0	Yes	10.3	Return of previously predominant PVC	N/A
8	Intramural OFT	57	Yes	23.2, 23.7, and 37.1	Return of previously predominant PVC	N/A
9	Epicardial OFT	58	Yes	58.3	Return of previously predominant PVC	N/A
10	ALP and PMP	292	Yes	13.3	Return of previously predominant PVC	N/A
11	ALP	180	No	35.4	Nonpredominant	PMP
12	Intramural OFT	194	No	36.2	Previously predominant PVC	N/A
13	Epicardial OFT	0	No	69.3	Previously predominant PVC	N/A
14	Epicardial OFT	0	Yes	31.9	Return of previously predominant PVC	N/A
15	Endocardial MA	0	Yes	44.4	New	Para-Hisian
16	Endocardial OFT	0	Yes	95.1	Return of previously predominant PVC	N/A
17	Endocardial OFT	138	Yes	42.9	Return of previously predominant PVC	N/A
18	Endocardial OFT	13	No	42.0	Nonpredominant	ALP
19	Para-Hisian	134	Yes	6.6	Return of previously predominant PVC	N/A
20	Epicardial OFT	1805	Yes	15.0	Return of previously predominant PVC	N/A

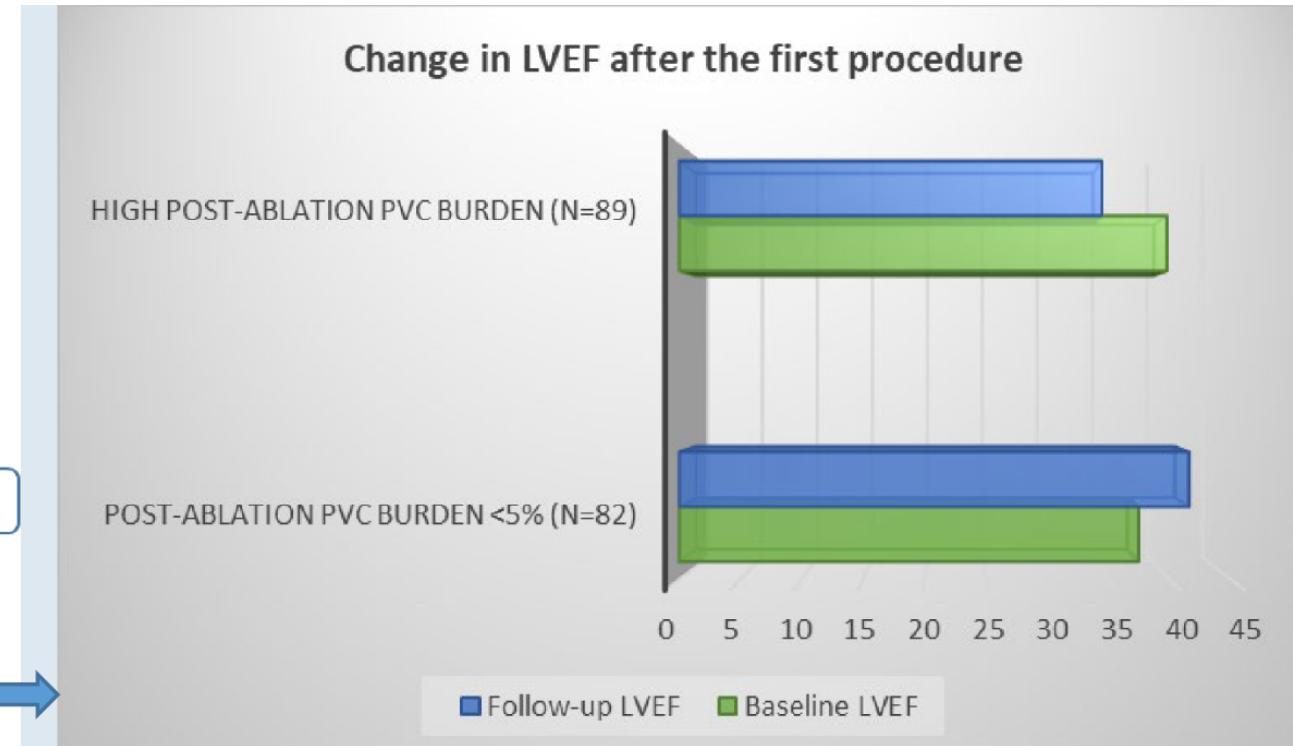
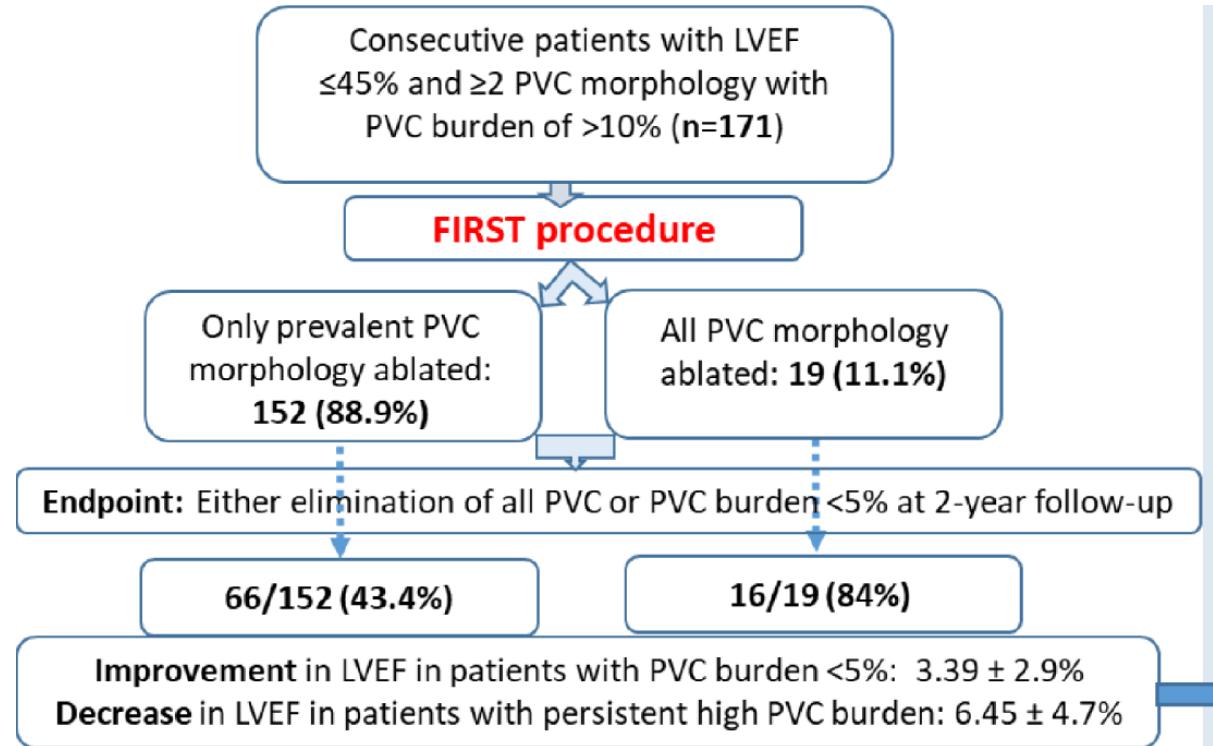
ALP = anterolateral papillary muscle; GCV = great cardiac vein; MA = mitral annulus; N/A = not applicable; OFT = outflow tract; PMP = posteromedial papillary muscle; PVC = premature ventricular complex.



# Change in non-predominant PVC burden



# Predominant PVC ablation alone vs all PVC ablation



# Outcomes of Ablation of Multiple PVC Morphologies

Depends on.

1. Individual PVC's location
2. Burden of non-predominant PVCs
3. Operator's experience

The burden of non-predominant PVCs does not change much after successful ablation of the predominant PVC.

Repeat ablation usually has been performed targeting previous predominant PVC.

Reduction of PVC burden is associated with increase of LV dysfunction in PVC induced CMP.



# Acknowledgement

## Electrophysiologists

- Il-young Oh, MD, PhD
- Youngjin Cho, MD, PhD
- Ji Hyun Lee, MD, PhD

## EP fellow (rotation)

- Ji-Suck Park, MD
- Hyung-bum Ahn, MD
- Do-Hyun Kim, MD
- Woong-su Yoon, MD
- Soo-Young Lee, MD
- Jina Choi, MD

## EP assistant

- Sung-wook Kim, RT, CEPS
- Su-ji Kim, RT, CEPS
- Jin-hyung Kim, RN
- Chan-yang Kim, MT
- Hu-lim Kim, MT
- Ga-hyuk Park, RT
- Eun-sung Yoon, MT
- Su-min Lee, RT
- Hyo-mi Chang, MT
- Yu-ri Choi, MT
- Ji-yun Whang, MT, CEPS/CCDS

## CIED lab

- Ji-Hye Yoo, RN
- Jung-Hwa Lee, RN
- Jin Ju Yang, MT

## EP PA

- Myung-sun Moon, RN
- Ok Choi, RN, CEPS

## EP research

- Eun-jung An, HIM
- Yun-ju Kim, RN
- Eun-ji Yoon, HIM
- Minji Yeo, RN