







# Does VT ablation impact VF?

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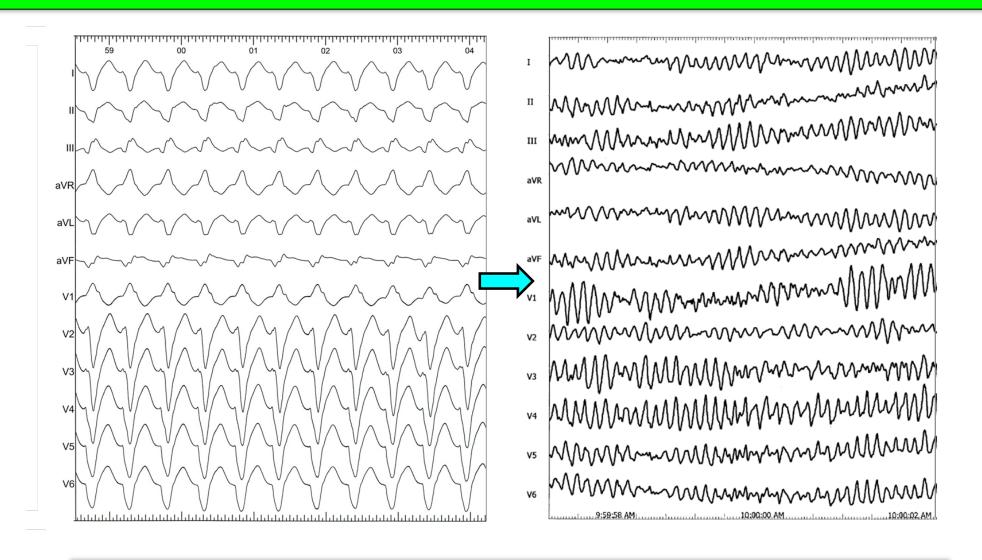
### **DISCLOSURES**

### **FINANCIAL DISCLOSURES (all modest)**:

<u>Consulting & Speaker</u>: Abbott, Biosense Webster, Boston Scientific, Meda Pharma, Medtronic, MicroPort, Stereotaxis

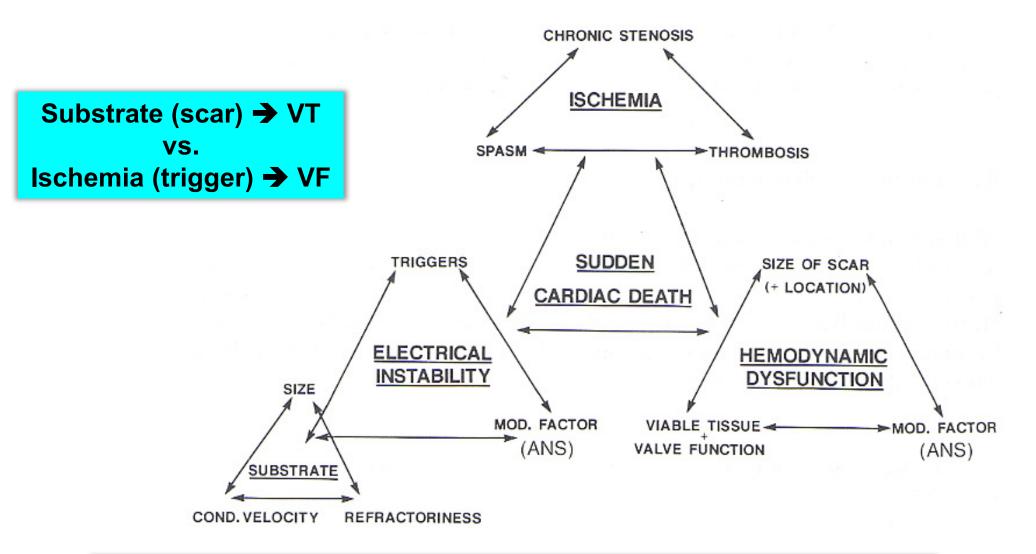
# Introduction - Background

# Ventricular Tachycardia & Ventricular Fibrillation



VF: may occur right away or may be triggered by a VT episode

# Pathophysiology of Ventricular Arrhythmias



Wellens HJJ et al. In: Sudden Cardiac Death. Kluwer Academic Publishers. 1991. pp:285-96

# IMMEDIATE CORONARY ANGIOGRAPHY IN SURVIVORS OF OUT-OF-HOSPITAL CARDIAC ARREST

TABLE 2. ANGIOGRAPHIC DATA IN THE 84 PATIENTS WHO UNDERWENT ANGIOGRAPHY.\*

VARIABLE	VALUE
Normal coronary arteries — no. (%)	17 (20)
Clinically insignificant coronary artery disease (≤50 percent stenosis) — no. (%)	7 (8)
Clinically significant coronary artery disease — no. (%)	60 (71)
Single-vessel disease	22
Two-vessel disease	13
Three-vessel disease	24
Isolated left main coronary artery disease	1
Left ventricular ejection fraction — %	$33.9 \pm 10.5$
Left ventricular end-diastolic pressure — mm Hg	25.3±9.5

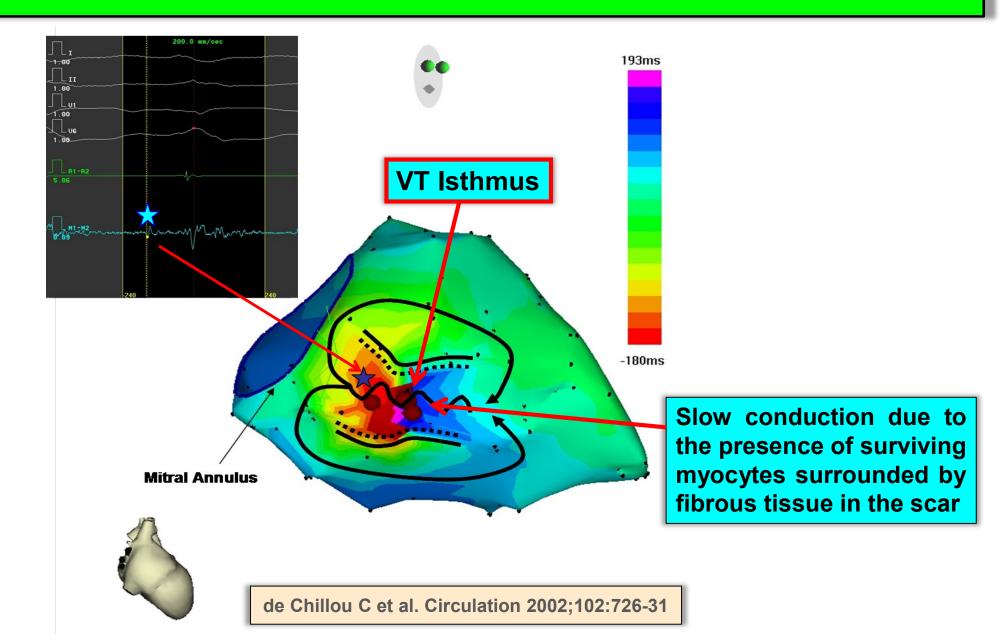
Acute coronary vessel occlusion = 48%

TABLE 3. Types of Coronary-Artery Lesions and Results of Percutaneous Transluminal Coronary Angioplasty (PTCA) in the 60 Patients with Clinically Significant Coronary Artery Disease.

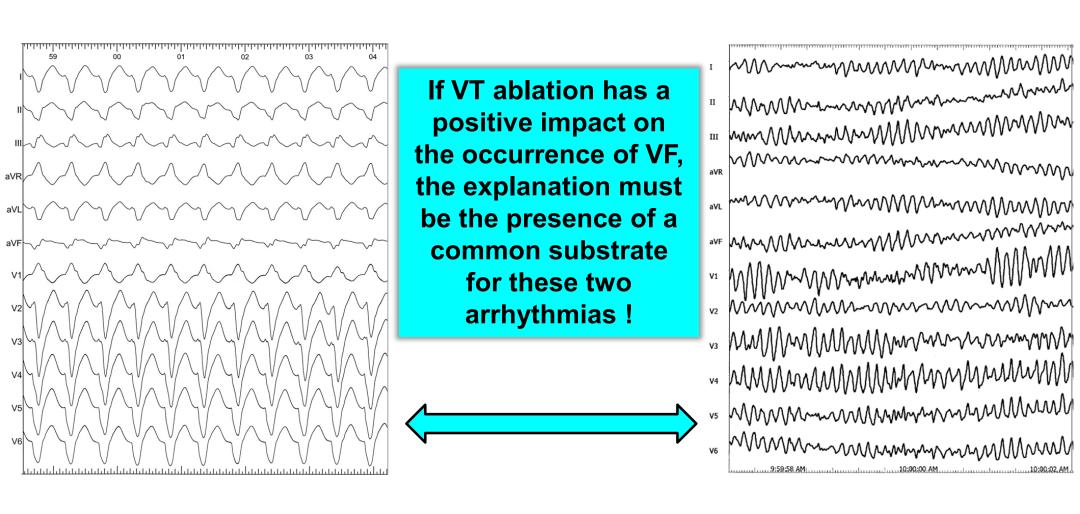
VARIABLE	VALUE
Type II lesion — no. (%) IIA	18 (30) 7
IIB	11
Type I lesion — no. (%)	2 (3)
Recent coronary-artery occlusion — no. (%) PTCA attempted — no.	40 (67) 37
Median interval between admission and PTCA — min (10th-90th percentile)	32 (18–55)
Median duration of procedure — min (10th-90th percentile)	62 (40–120)
PTCA successful — no.	28
Stent implanted — no.	5
Intraaortic balloon inserted — no.	9

Spaulding C. N Engl J Med 1997;336:1629-33

# Post-infarct mappable VT

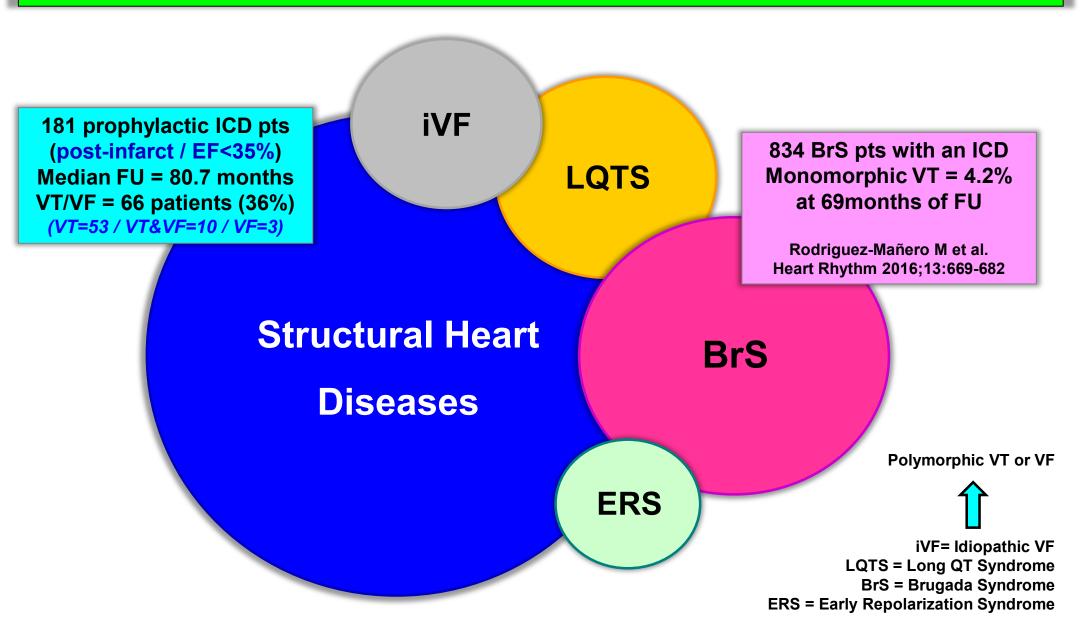


# Ventricular Tachycardia & Ventricular Fibrillation



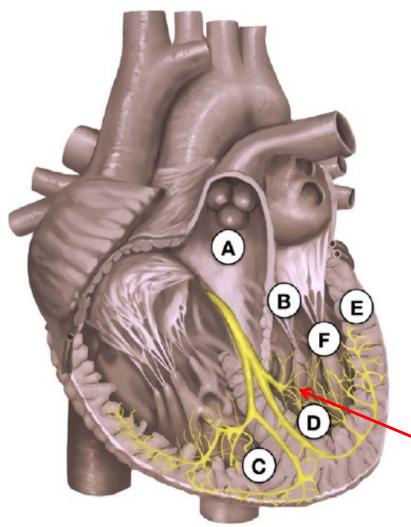
VT and VF: two different entities or the two sides of the same coin?

# Ventricular Tachycardia & Ventricular Fibrillation



# VF Mapping and Ablation: Triggers & Mechanisms

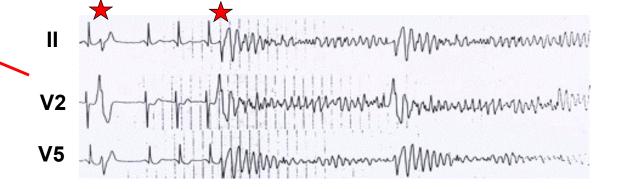
## VF Ablation Targeting PVC Triggers



#### **Location of PVC triggers**

	Anatomical Site	n (%)	Conditions
A	RVOT	13 (10%)	IVF, BrS
(B)	LVOT	9 (7%)	IVF, DCM
(C)	<b>Purkinje</b> RV-Purkinje LV-Purkinje Both-Purkinje	73 (59%) 15 53 5	IVF, LQTS, ER, IHD, BrS, DCM
(E)	Myocardium	16 (14%)	LQTS, ER, IVF, DCM
(F) F	Papillary Muscle	13 (10%)	IVF, DCM

Anderson RD et al. Heart, Lung and Circulation 2019;28:110-122



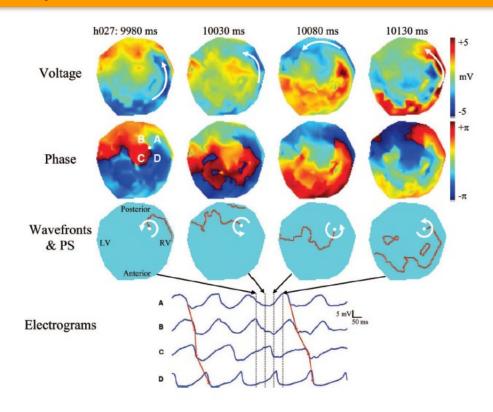
# In vivo mechanistically-based VF mapping in man

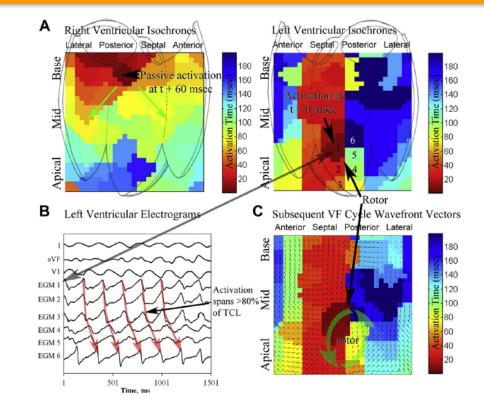
**Epicardial electrode array during open heart** surgery in CAD (10 pts)

→ Coexistence of a small number of rotors and multiple wavelets

→ Stable rotors in sustained VF episodes
 requiring defibrillation vs. unstable rotor and foci.

requiring defibrillation vs. unstable rotor and foci in self-terminated VF episodes





Nash MP et al. Circulation 2006;114:536-42

Krummen DE et al. Circulation 2014;63:2712-21

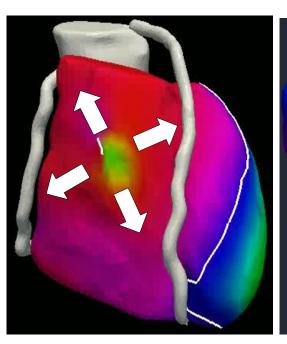
# Drivers Maintaining Human VF -> BSM

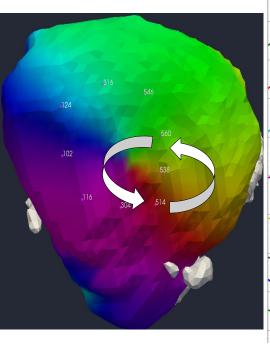


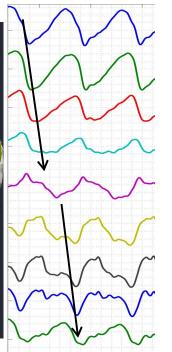
Focal Breakthrough

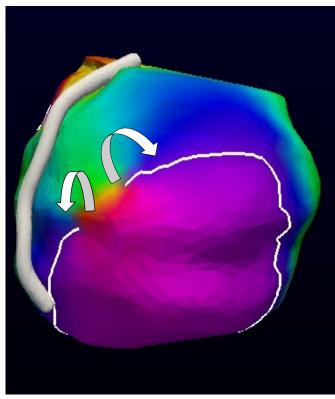
'Focal' Reentry

Figure of 8 Reentry









EGM covering >75% of VF cycle length
Gradient of VF Cycle Length (≥5ms) surrounding rotor areas → confirm their driving role

Post-infarct VF → Figure of 8 reentry in most cases

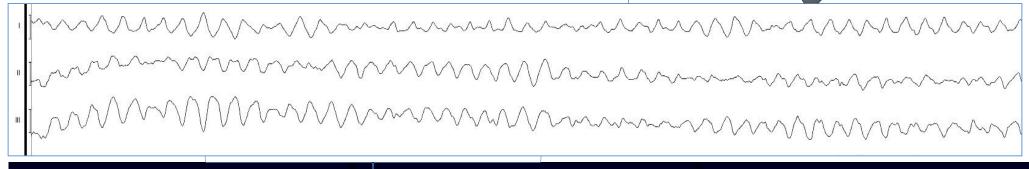
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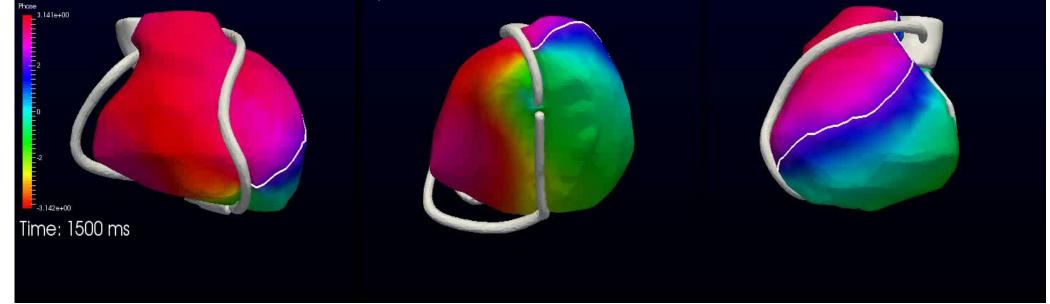


Courtesy: Pr. Michel Haïssaguerre

67 year old man / inferior MI / Induced VF 24sec



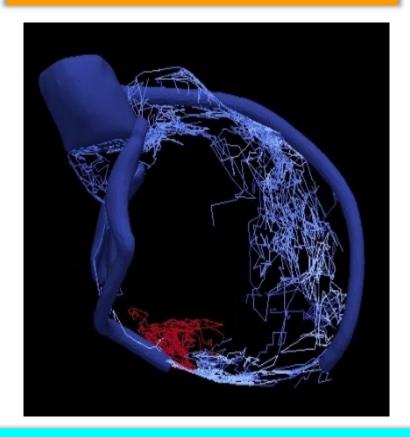




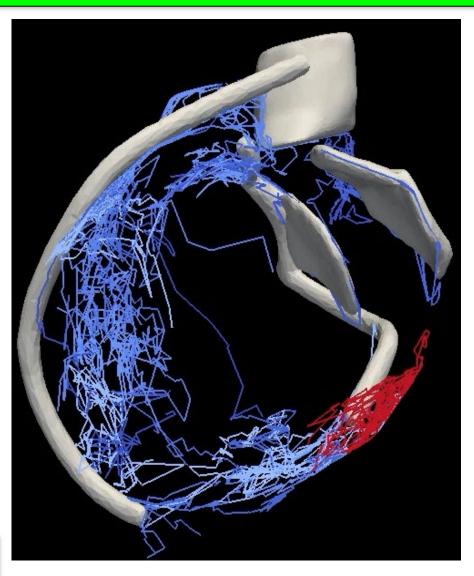
# Drivers Maintaining Human VF → BSM



# Trajectory of VF reentry: all around LV scar



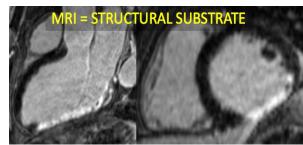
VF drivers share common pathways with coexisting mapped VT circuits

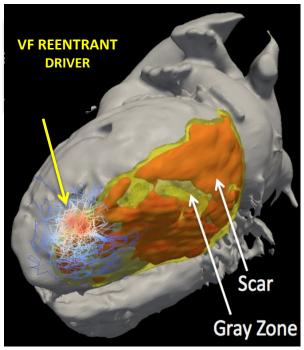


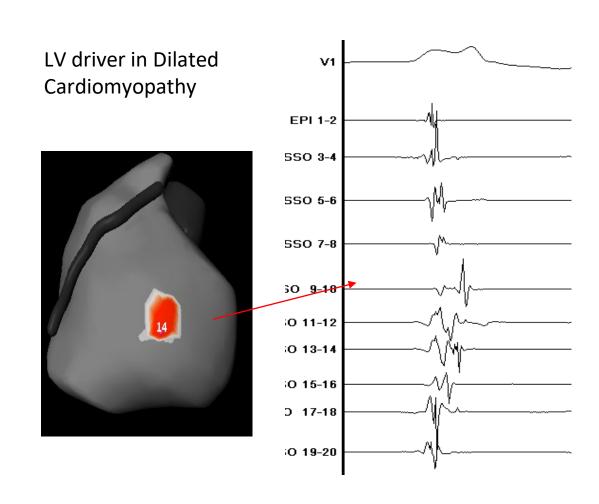
Courtesy: Pr. Michel Haïssaguerre

# Drivers Maintaining Human VF -> BSM









Main VF driver co-locate with areas of abnormal fragmented EGM during SR

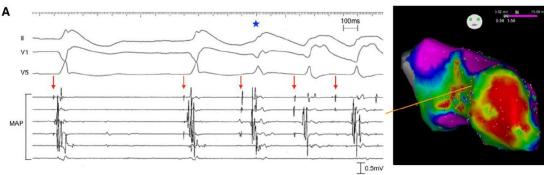
Courtesy : Dr. Mélèze Hocini

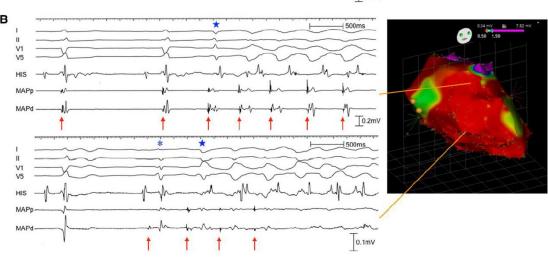
# Does VT Ablation Impact VF ? Literature Data

# Catheter Ablation of Refractory VF Storm after MI

# VT triggers originate from the scar border zone and from the dense scar

The triggering VPBs were found to originate from the surviving Purkinje tissue in the dense scar area (a bipolar voltage <0.5 mV) in 15 patients (14%) and from the scar border zone (a bipolar voltage of 0.5–1.5 mV) in 88 patients (80%; Figure 1). Although VPBs were found to originate from the normal voltage area (a bipolar voltage >1.5mV) in the remaining 7 patients (6%), these sites also correlated with the territory of infarction. The

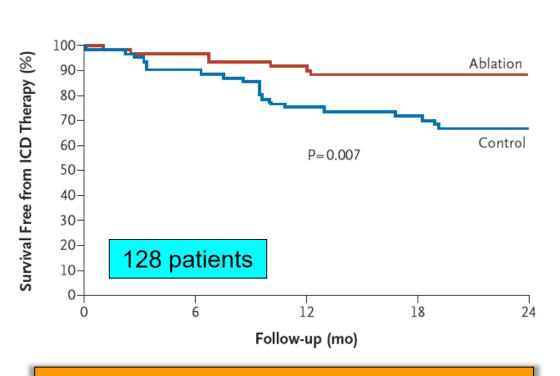


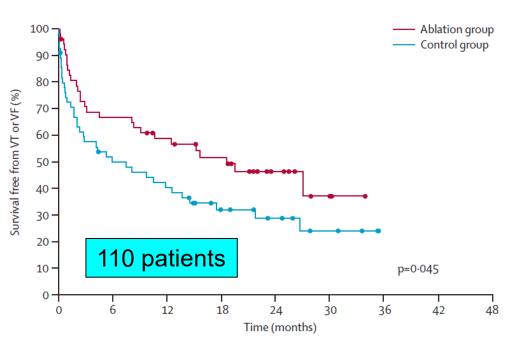


### Randomized trials: ICD alone vs. ICD + VT ablation

#### **SMASH-VT**

#### **VTACH**





Decrease by 70% of the incidence of appropriate ICD shocks, p=0.003

Decrease by 46% of the incidence of appropriate ICD shocks, p=0.045

Reddy V et al. N Engl J Med 2007;2:474-82

Kuck KH et al. Lancet 2010;375:31-40

# Retrospective study – Impact of VT ablation on VF

Courtesy : Pr. Pierre Jaïs

686 patients with a first VT ablation procedure (2010-2017)

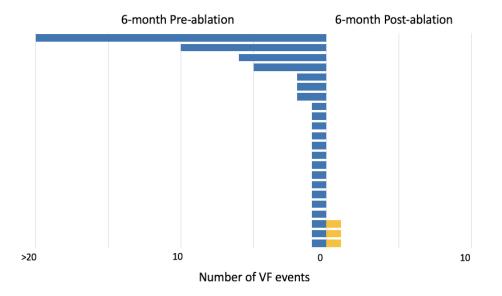
21 patients met the following inclusion criteria: ICD + documented VT + VF episodes during the 6-month period preceding LAVA-guided ablation

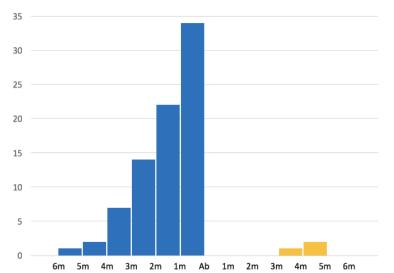
- → Post-infarct =10/21 (48%)
- → Dilated cardiomyopathy = 8/21 (38%)
- → ARVD/C = 2/21 (9%)
- **→** Myocarditis = 1/21 (5%)

80 VF events in the preceding 6 months

Kitamura T, Jaïs P et al. CircA&E 2019 (in press)

NUMBER OF VF EVENTS 6-MONTHS BEFORE AND AFTER VT ABLATION TARGETING LAVA





# **Conclusions**

- Several studies (randomized or observational) have reported a reduction of ICD shocks and/or a low incidence of SCD after VT ablation
- One retrospective dedicated study observed a significant reduction of VF burden after ablation of scar-related VT
- The most likely explanation for this favorable outcome is the following: VT and VF are sharing, at least in part, the same arrhythmogenic substrate
- □ This explanation is consistent with a number of VF mapping studies (endocardial / epicardial / BSM) in human