



# Atrial Imaging Using Cardiac MRI



## Name

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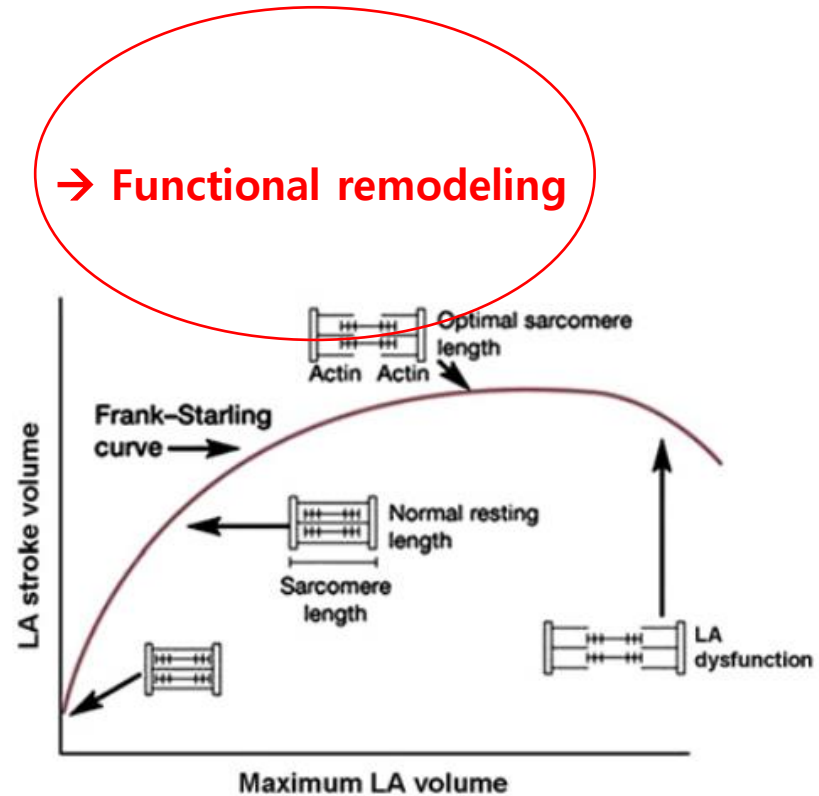
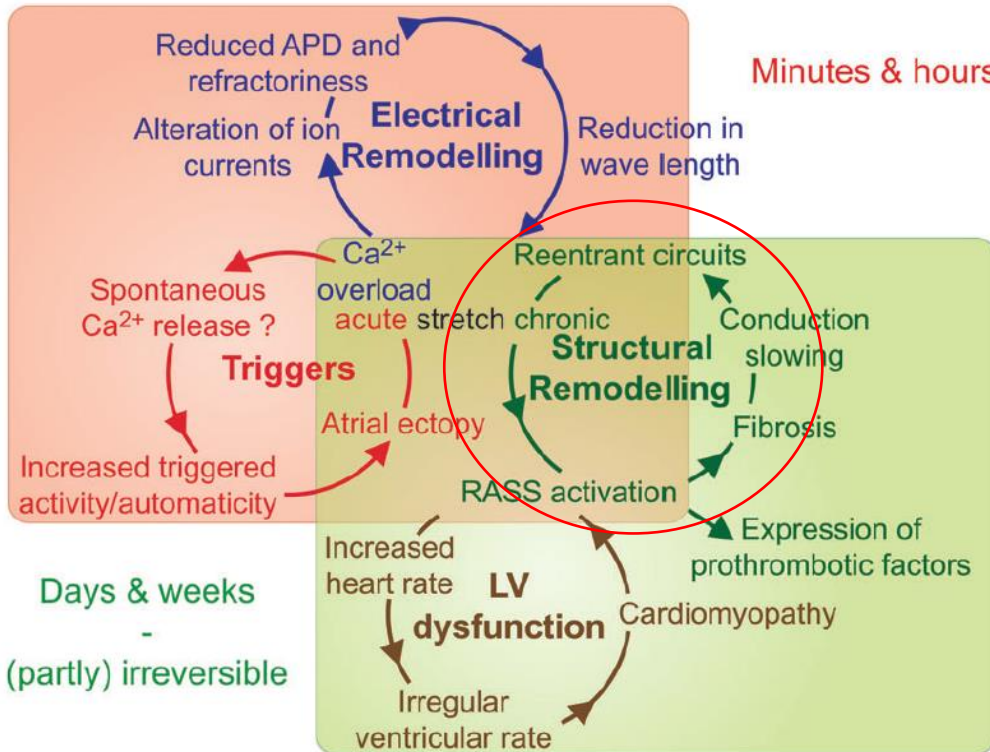
# **Korean Heart Rhythm Society COI Disclosure**

*Name of First Author: Jung Myung Lee*

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



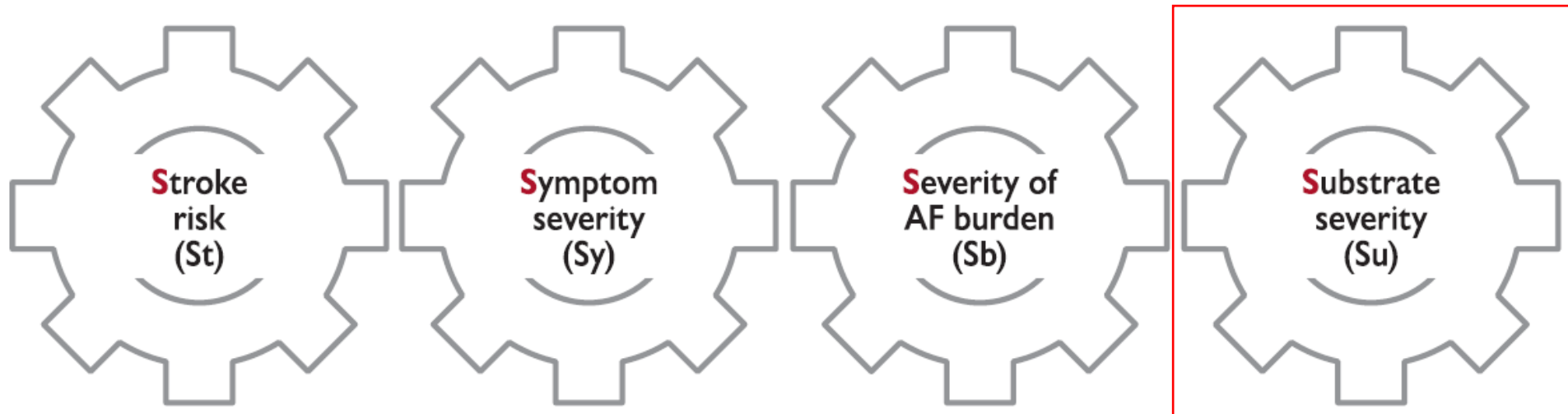
# LA remodeling accompanied with AF



**Figure 1** Frank–Starling law applied to the left atrium (LA).



# Structured characterization of AF



- Stroke risk (St)**
- Truly low risk of stroke
  - Yes
  - No

- Symptom severity (Sy)**
- Asymptomatic/mildly symptomatic
  - Moderate
  - Severe or disabling

- Severity of AF burden (Sb)**
- Spontaneously terminating
  - AF duration and density of episodes per unit of time

- Substrate severity (Su)**
- Comorbidities/ cardiovascular risk factors
  - Atrial cardiomyopathy (atrial enlargement / dysfunction / fibrosis)

<p><b>CHA<sub>2</sub>DS<sub>2</sub>-VASc score</b></p>	<p><b>EHRA symptom score</b></p> <p>QoL questionnaires</p>	<ul style="list-style-type: none"> <li>• <b>Temporal pattern of AF</b> (Paroxysmal, Persistent, Long-standing persistent, Permanent)</li> <li>• <b>Total AF burden</b> (total time in AF per monitoring period, the longest episode, number of episodes, etc.)</li> </ul>	<ul style="list-style-type: none"> <li>• <b>Clinical assessment</b> Incident AF risk scores, AF progression risk scores</li> <li>• <b>Imaging</b> (TTE, TOE, CT, cardiac MRI), biomarkers</li> </ul>
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# Substrate status in guiding AF treatment

Substrate
(+1 if >75 years old)
<b>Comorbidity/CV risk factors</b>
0 = No
1 = Single
2 = Multiple (2 or more)
<b>LA enlargement/dysfunction</b>
0 = No
1 = Mild-moderate
2 = Severe
<b>LA fibrosis</b>
0 = No
1 = Mild
2 = Moderate-severe
<b>0-2</b>
<b>3-4</b>
<b>5 or more</b>

Anatomy, size  
Structure, fibrosis  
Functions

All green => Rhythm control

1 Yellow, 2 Green => Rhythm control can be attempted

Red => Consider rate control



# LA imaging modalities in AF

Left atrial remodelling associated with AF

Anatomy

Dilatation and change in geometry

Structure

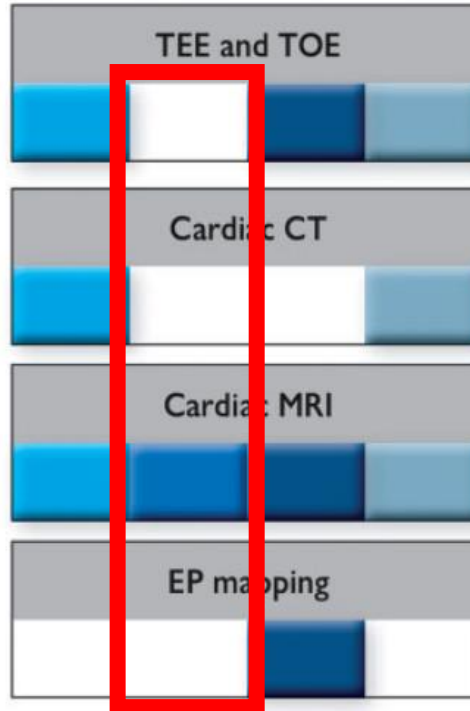
Fibrosis

Function

Altered electrophysiology, LA reservoir, conduit and booster pump function

LA/LAA thrombus detection

Value of LA imaging techniques in AF



+

Value of imaging beyond LA

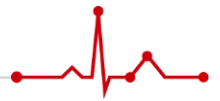
LV size, geometry and function assessment

Heart valves morphology and function

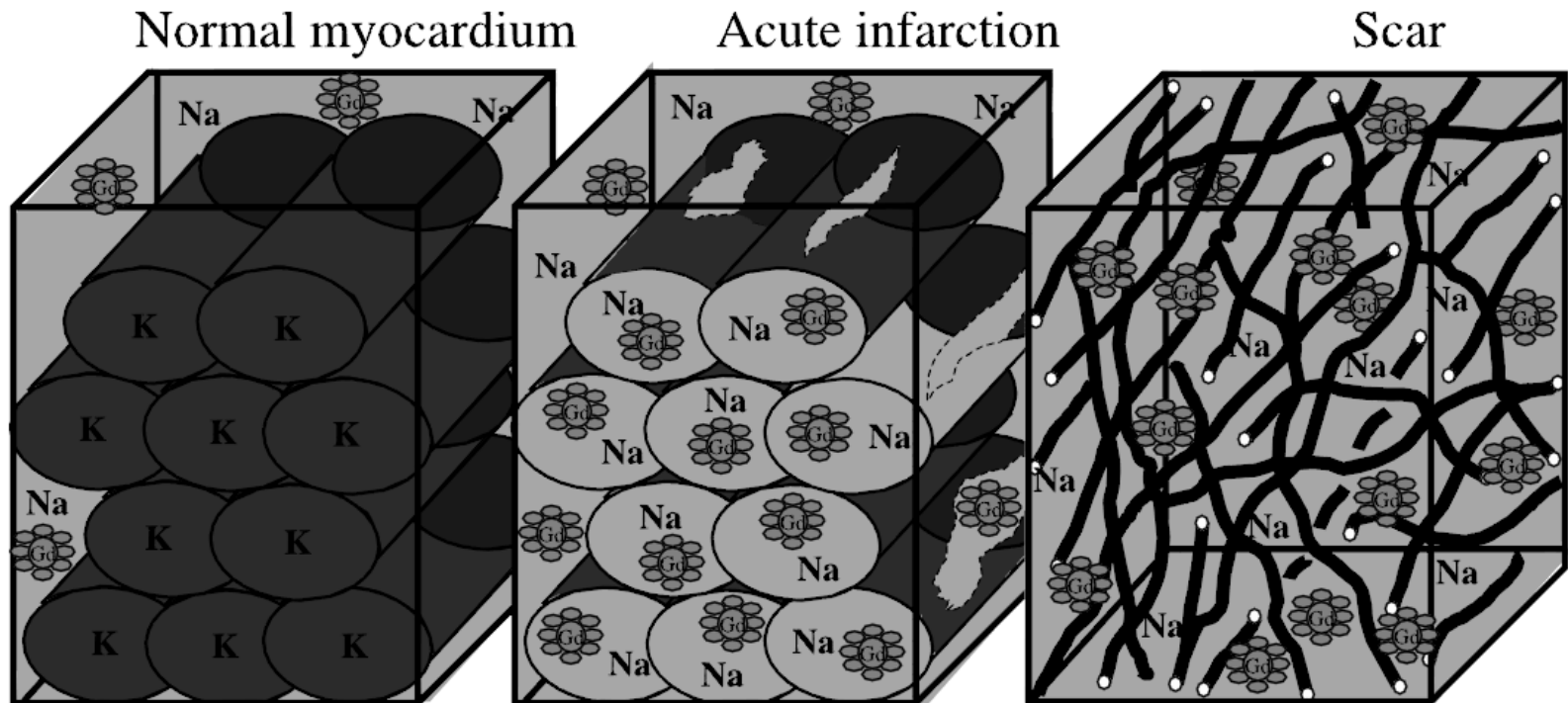
Right-heart chambers and pericardium imaging

**Advanced/Investigation imaging:**

- Echocardiographic TDI and LA strain, etc.
- MRI delayed enhancement or T1 imaging
- CT imaging of substrate, etc.



# Late gadolinium enhancement

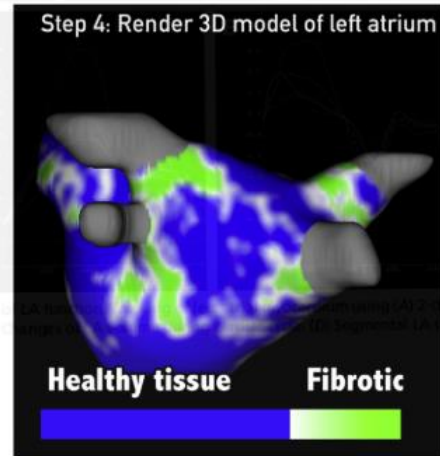
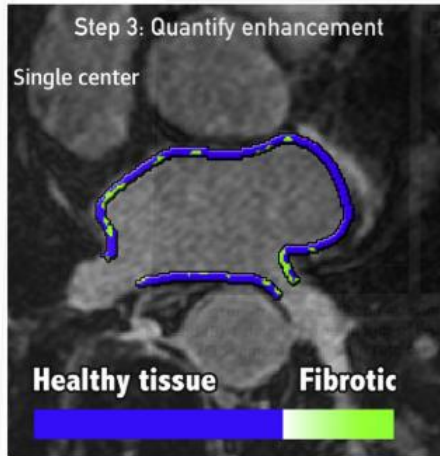


Intact cell membrane    Ruptured cell membrane    Collagen matrix

**Figure 2.** Potential mechanisms of hyper-enhancement in acute and chronic myocardial infarcts.



# Acquiring atrial LGE



Following acquisition of high-resolution LGE-MRI scans, the endocardial borders of the left atrium (LA) are defined, including an extent of pulmonary vein (PV) sleeves, by manually tracing the PV-LA blood pool in each slice of the LGE-MRI volume. Next, the endocardial border is morphologically dilated (by 4 pixel layers, 2.5 mm) with manual adjustment to create a shell of the epicardial LA surface (step 2). The endocardial segmentation is subtracted from the epicardial layer to define the wall segmentation, with manual exclusion of the mitral valve and extension of the left ventricle. The next step is the quantification of fibrosis based on the relative intensity (signal intensity) of LGE. Finally, a 3-dimensional model of the LA is rendered with the maximum enhancement intensities being projected on the model surface

Challenges:  
Thin atrial wall  
Irregular HB  
Respiration  
No uniform protocol

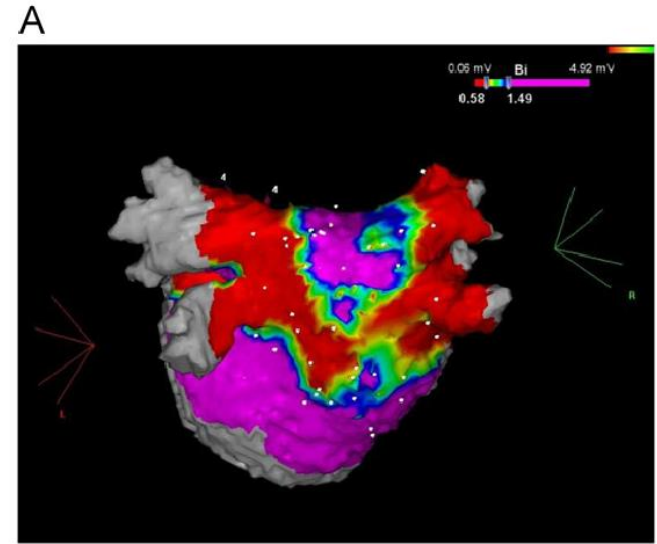
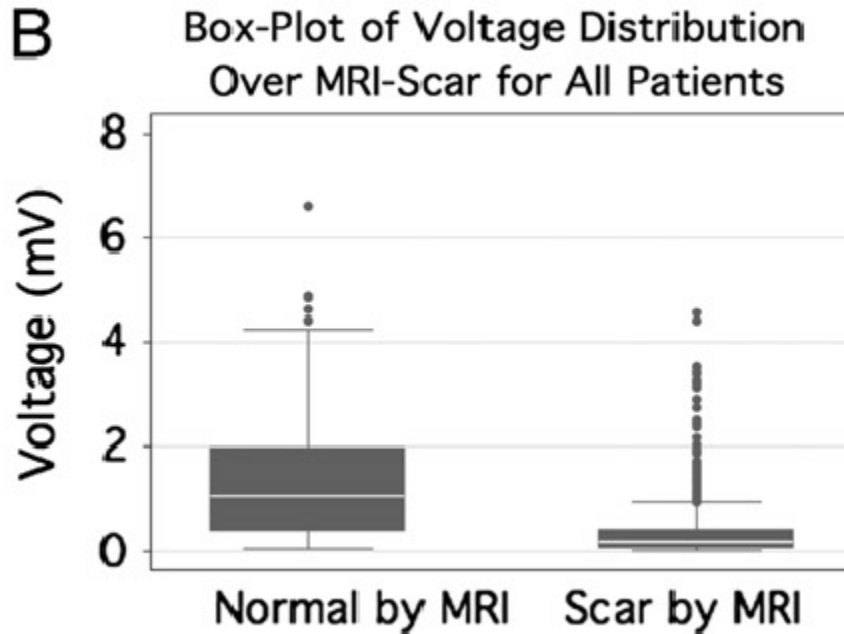




# LGE and low voltage area

a significant association between LGE and areas of low voltage

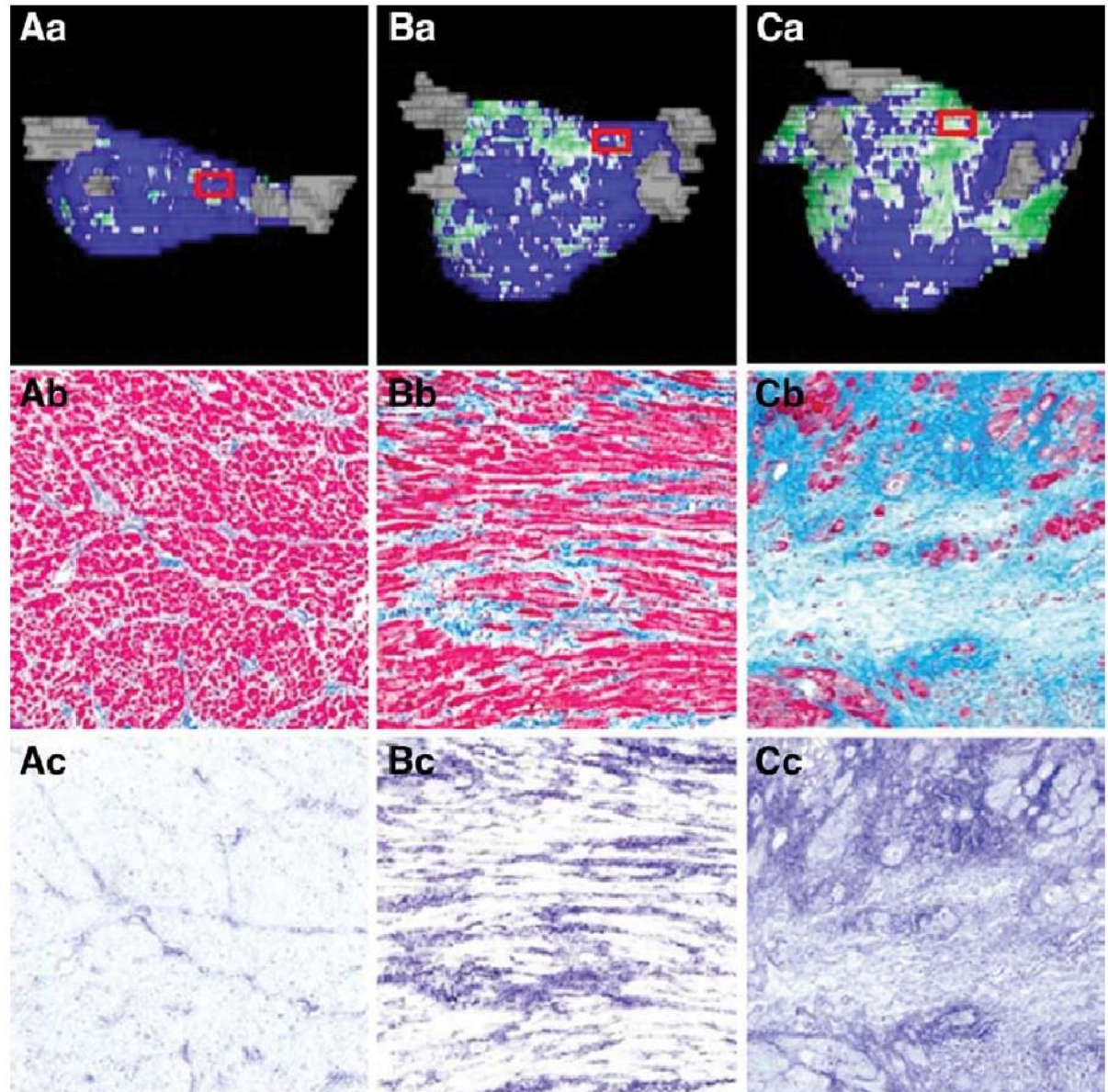
- Significant association between scar identified by DE- MRI and low-voltage regions of the LA





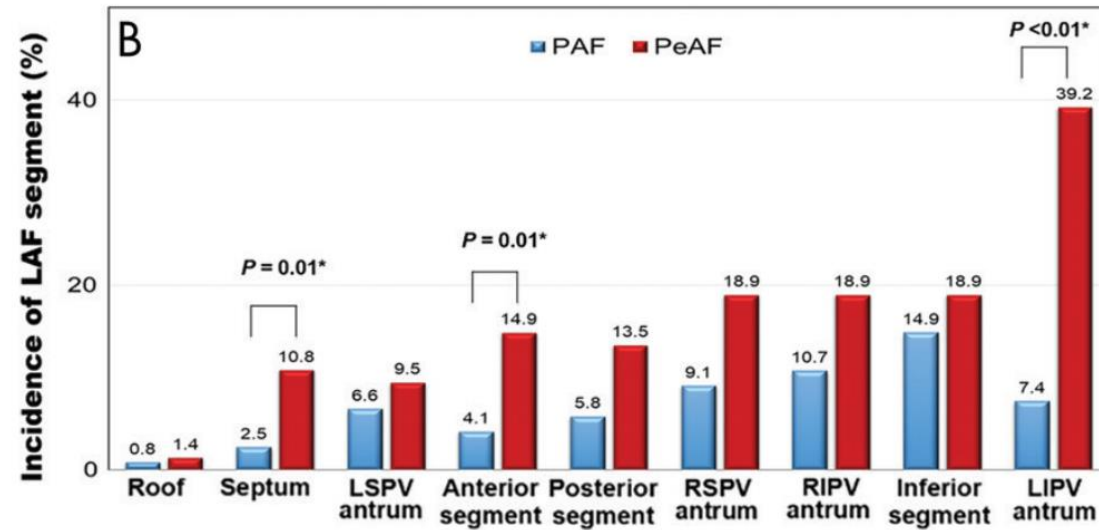
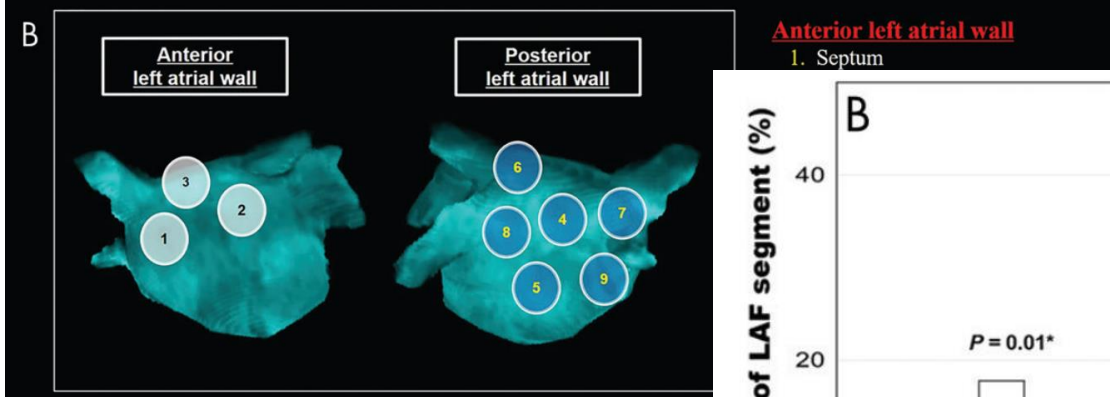
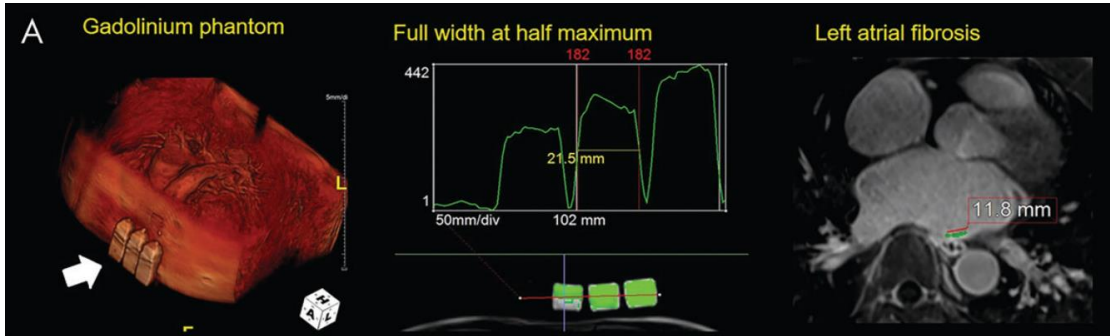
# LGE and fibrosis on biopsies

LA LGE is found to be matched with corresponding LA wall areas demonstrating fibrosis on surgical biopsies.





# LGE pattern and AF type



Korea University

195 patients

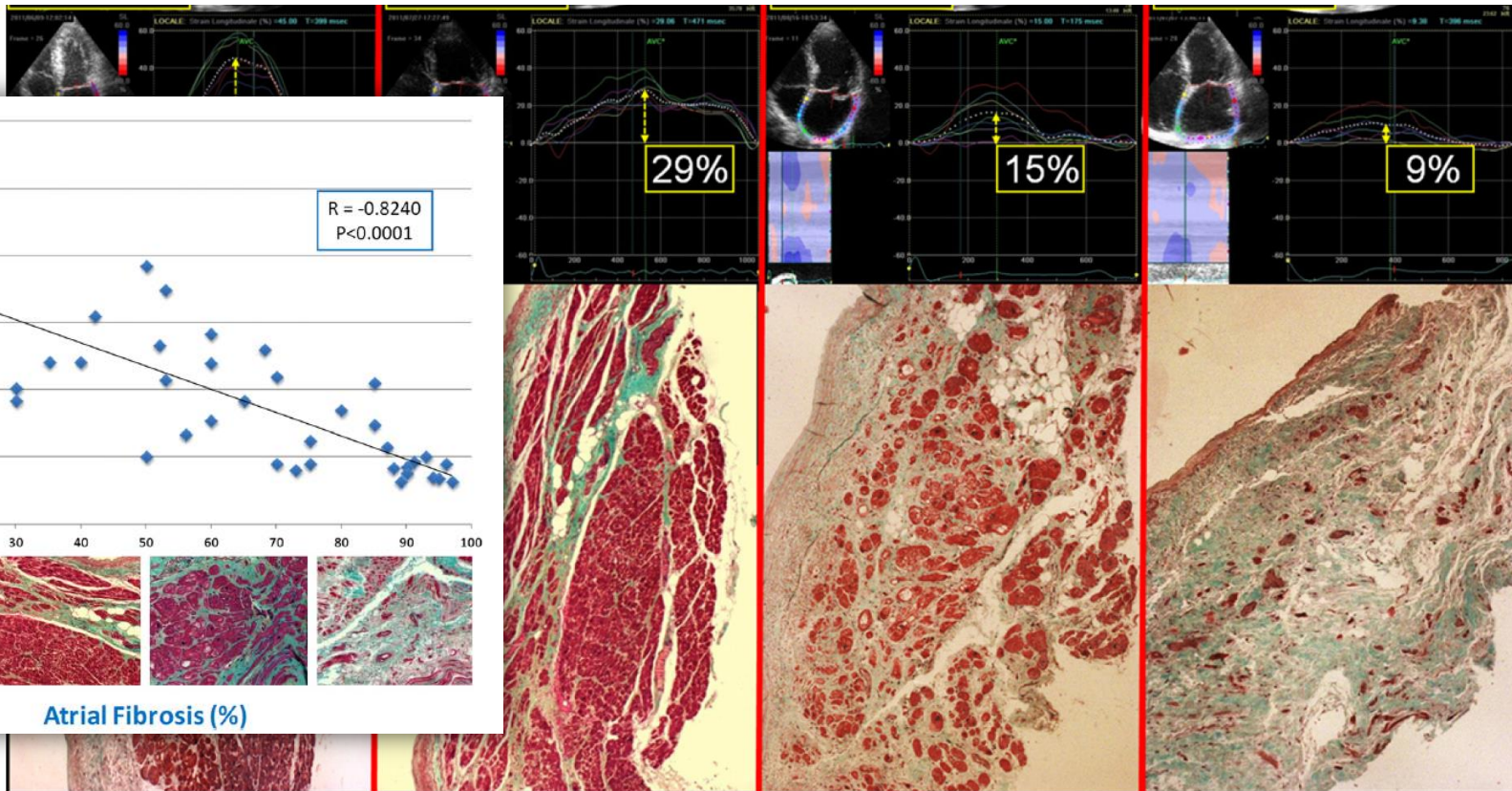
114 (58.4%) had at least one left atrial LGE segment.

The mean number of LGE segments was higher in the PeAF group than in the PAF group. The incidence of LGE at the LIPV antrum was higher in the PeAF group than in the PAF group



# Atrial fibrosis associated to function

Global PALS showed the best diagnostic accuracy to detect LA fibrosis (area under the curve 0.89) than LAVI, LAEF, E/E' (46 pts, OP for MR)

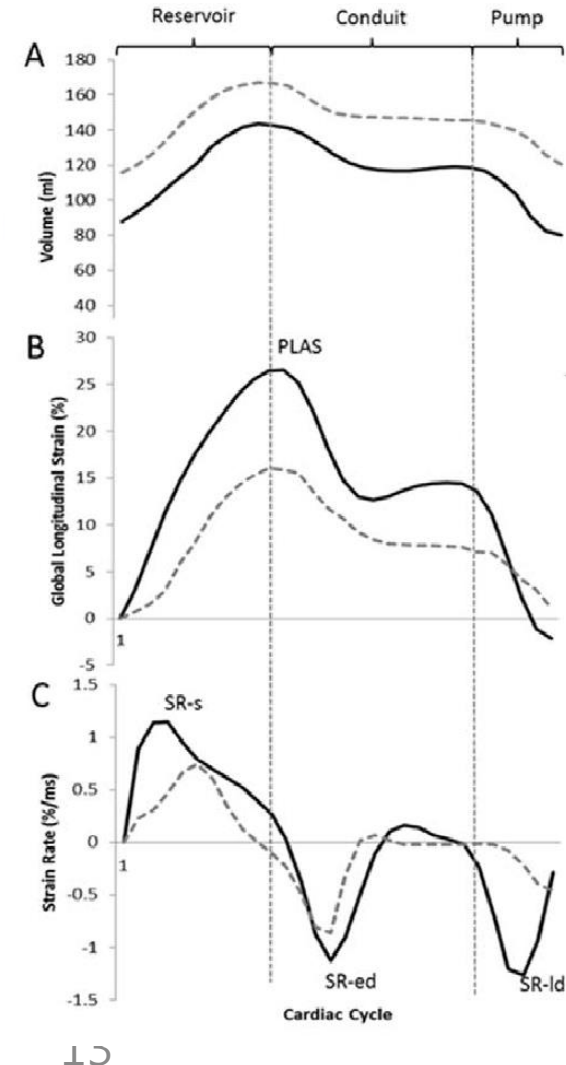
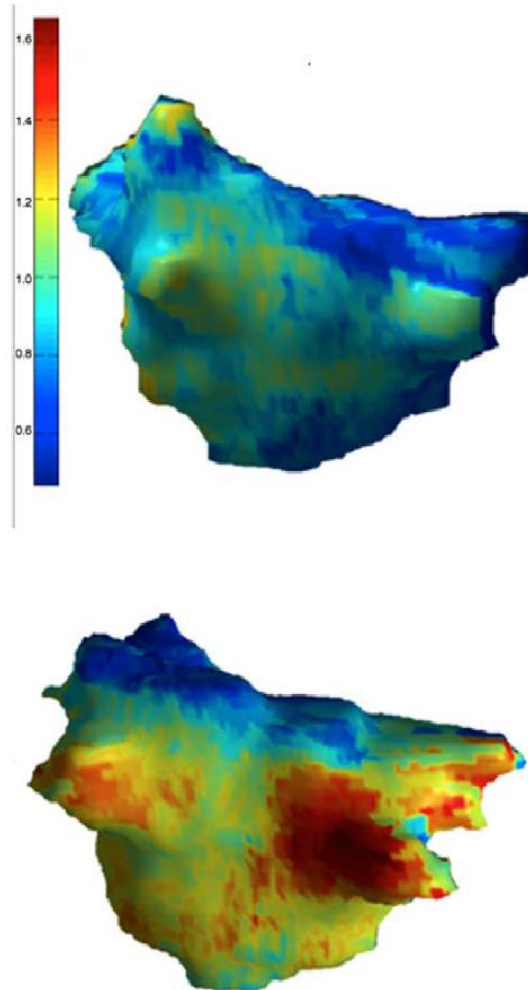




# LGE and LA function

Increased LA enhancement is associated with decreased LA reservoir, conduit, and booster pump functions.

Phasic measurement of LA function using feature-tracking cardiac magnetic resonance may add important information about the physiological importance of LA fibrosis





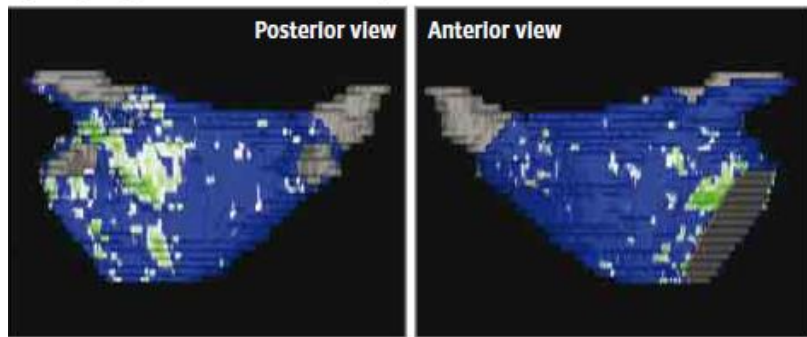
# LGE and prognosis of AF: Prediction of RFCA outcome by atrial fibrosis

272 of 329 pts

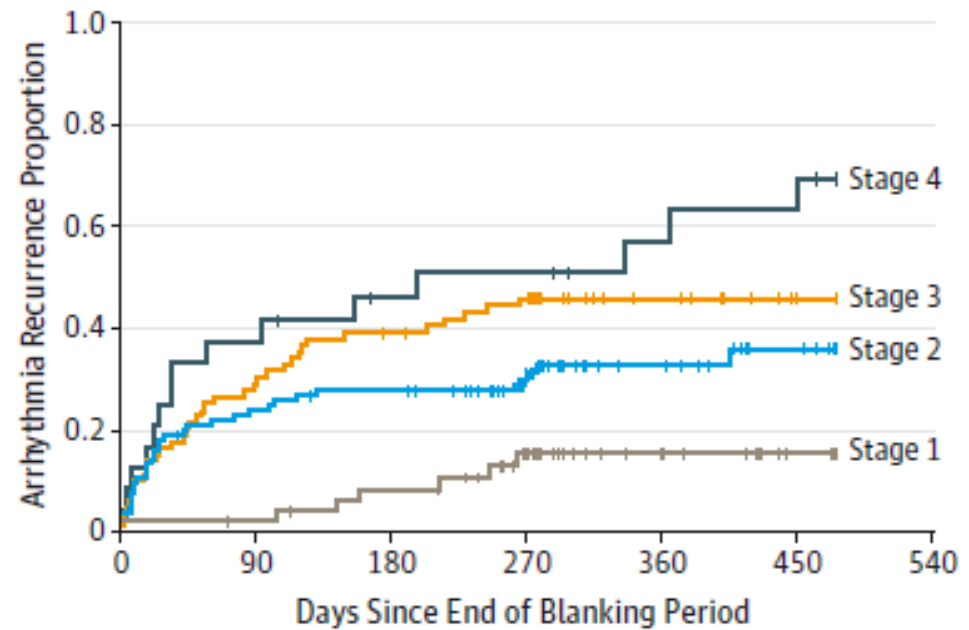
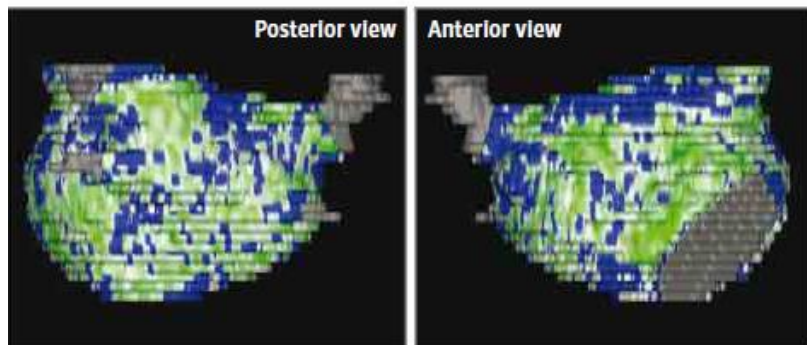
(57 pts (17%) excluded d/t poor MRI quality)

The addition of fibrosis to a recurrence prediction model → C statistic increasing from 0.65 to 0.69

**B** Stage 2 (≥10% - <20% of atrial wall)



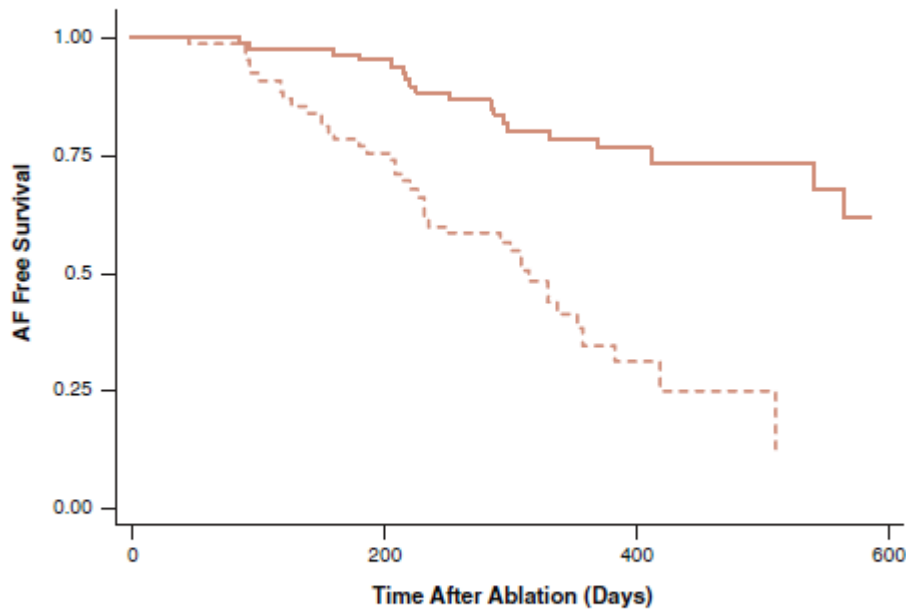
**D** Stage 4 (≥30% of atrial wall)



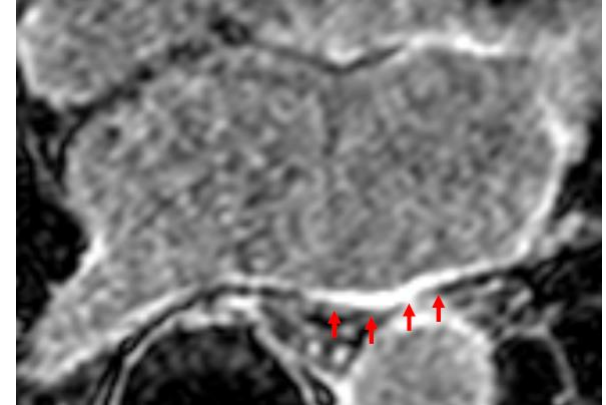


# Prediction of RFCA outcome by MRI

**FIGURE 2** Kaplan-Meier Atrial Fibrillation-Free Survival Curves After Pulmonary Vein Isolation in Participants Stratified by Median Left Atrial Late Gadolinium Enhancement Extent



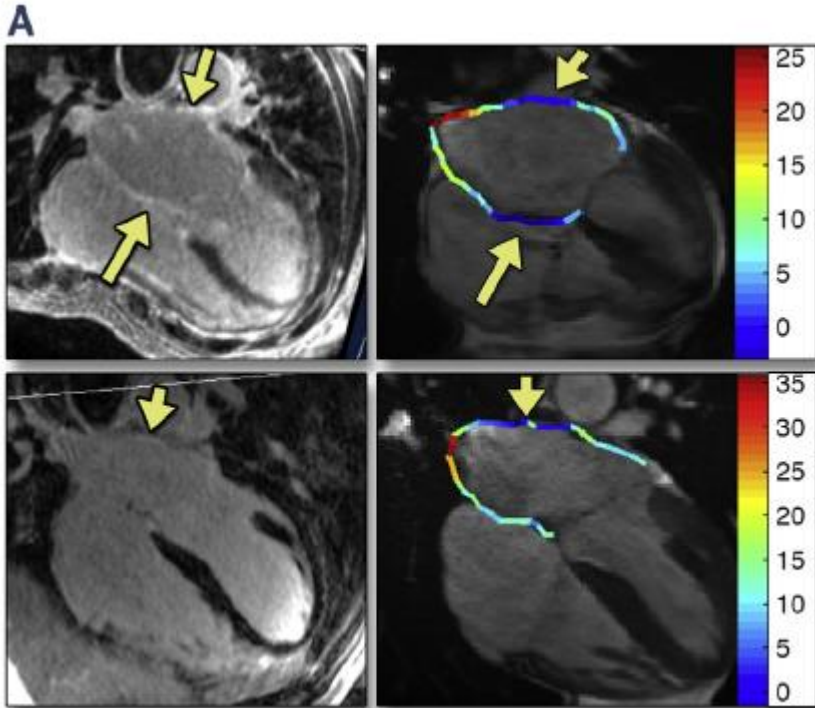
Time to first atrial fibrillation (AF) recurrence in 86 participants (52%) with late gadolinium enhancement (LGE) extent  $\leq 35\%$  (**solid line**) and 79 participants (48%) with LGE extent  $> 35\%$  of left atrial myocardium (**dashed line**) ( $p < 0.001$ ).



Regardless of AF persistence at baseline, participants with LGE  $< 35\%$  have favorable outcomes, whereas those with LGE  $> 35\%$  have a higher rate of AF recurrence in the first year after ablation



# LA function measured by MRI. Strain

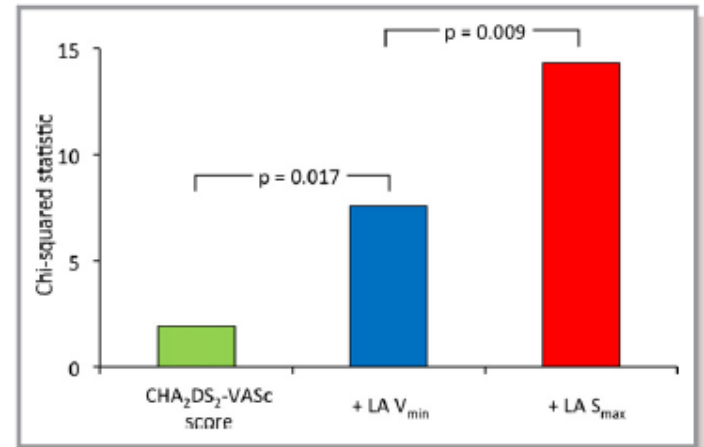


**Low strain region = LA fibrosis region**

JACC 2017' letter. <http://dx.doi.org/10.1016/j.jcmg.2016.01.015>

169 PAF pts.

LA strain measured by MRI had additive value in stroke prediction.



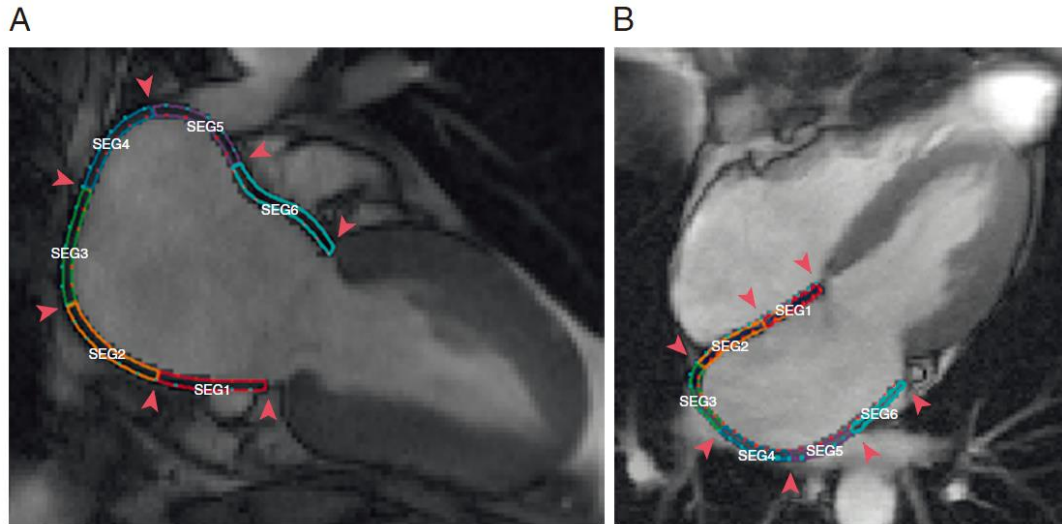
**Figure 4.** Incremental value of left atrial (LA) strain for diagnosis of stroke. The addition of the LA minimum volume ( $V_{min}$ ) to the model on the basis of the CHA<sub>2</sub>DS<sub>2</sub>-VASc score resulted in significant improvement in the diagnostic value for stroke. The value was further increased by adding the LA global longitudinal maximum strain ( $S_{max}$ ).



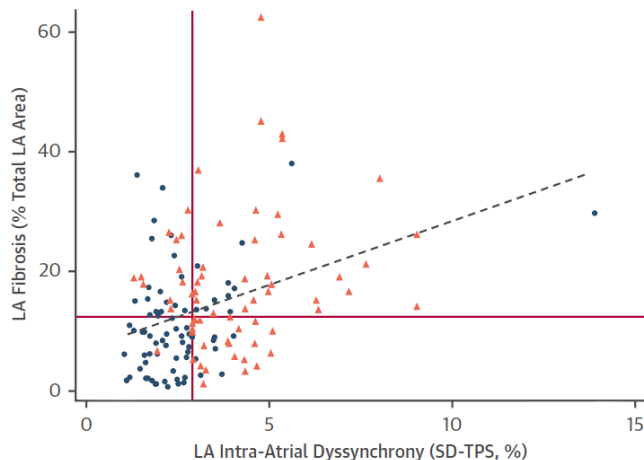


# Intra-atrial dyssynchrony during sinus rhythm is an independent predictor of recurrence after RFCA

**FIGURE 2** Quantification of LA Regional Function Using Cine CMR

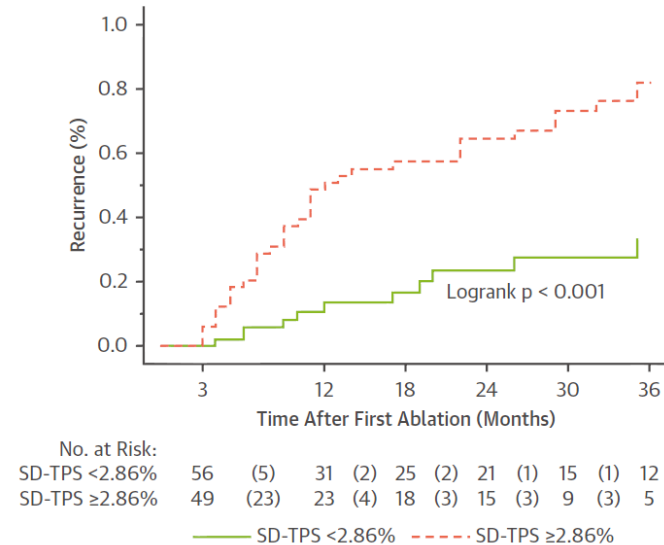


**FIGURE 4** Correlation Between Intra-Atrial Dyssynchrony and LA Fibrosis



Patients with AF recurrence after ablation (n = 101) had significantly higher SD-TPS than those without (n = 107; 3.9% vs. 2.2%; p < 0.001). Multivariable cox analysis showed that SD-TPS was associated with recurrence after adjusting for clinical risk factors, AF type, LA structure and function, and fibrosis (p < 0.001). Furthermore, ROC analysis showed SD-TPS improved prediction of recurrence better than clinical risk factors, LA structure and function, and fibrosis.

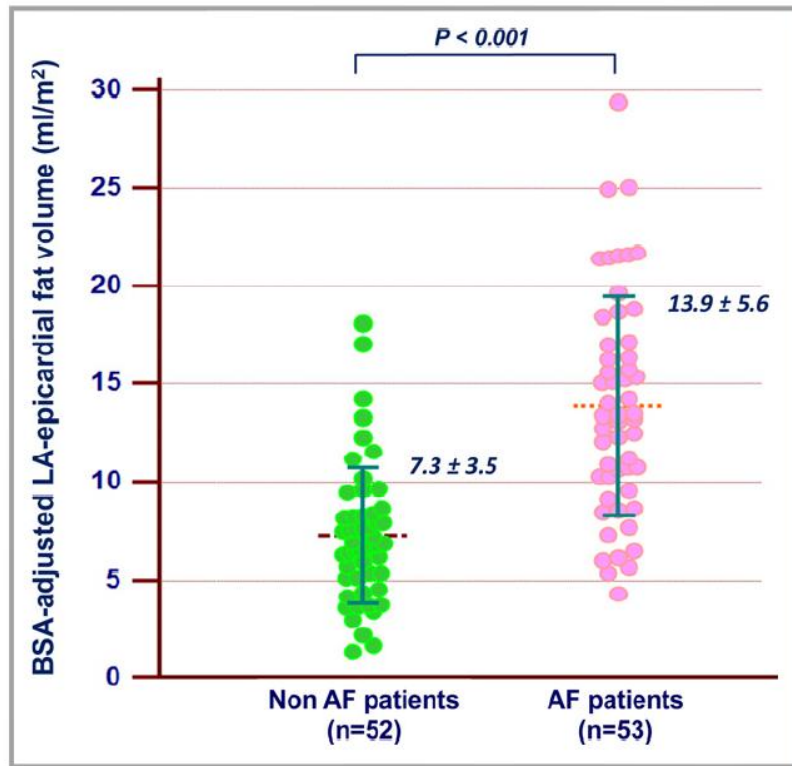
**FIGURE 5** Time to Atrial Fibrillation Recurrence in the Test Group



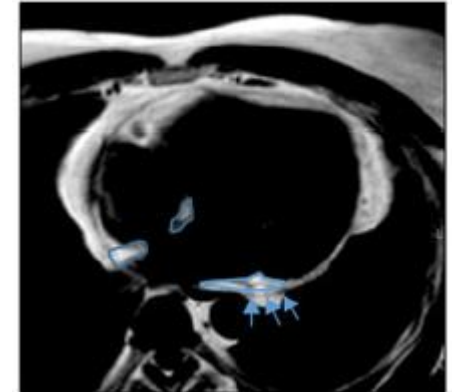


# LA epicardial fat volume by MRI

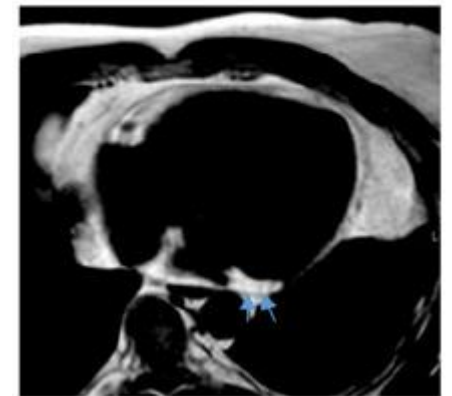
53 Hx of AF  
52 control



Fat only image

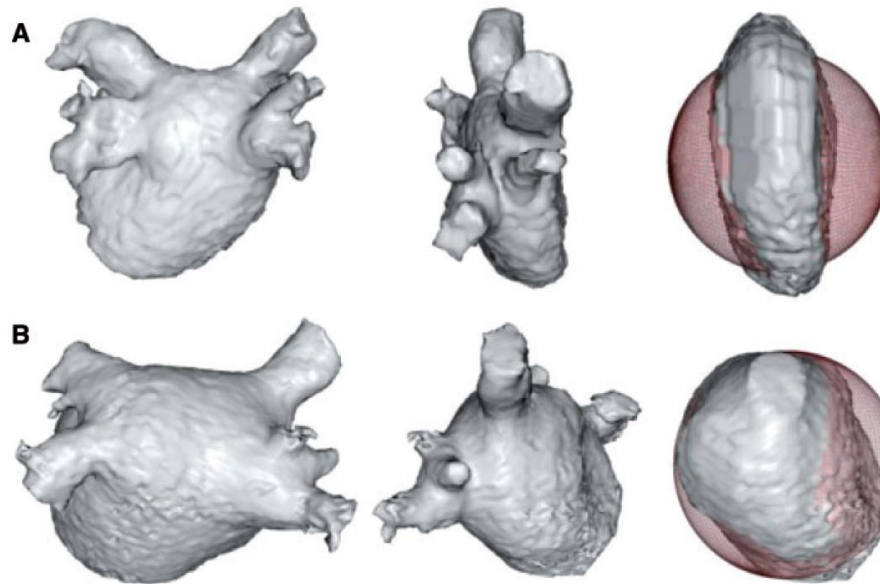


Fat only image

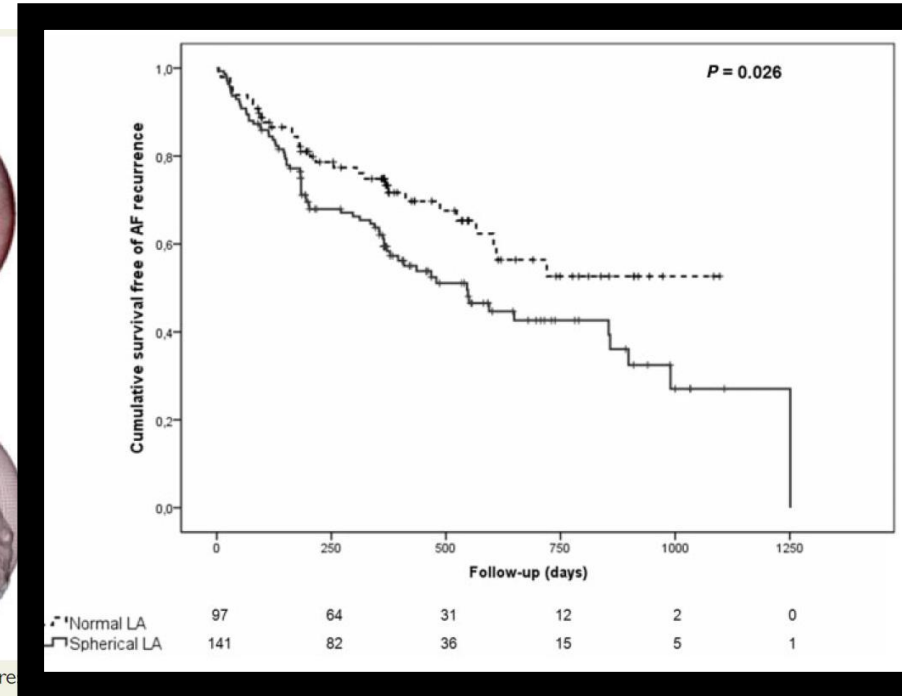




# Sphericity index



**Figure 1** Left atrial sphericity (LASP) categories: examples of discoid (A) and spherical-shaped LA (B) corresponding to the LASP categories. Posterior and right lateral views of 3D model are displayed. LAV, LA volume.



LA sphericity (HR 1.87,  $P = 0.035$ ) was an independent predictors for AF recurrence.

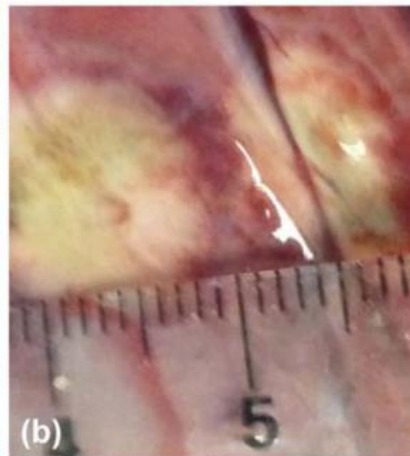
A combined clinical-imaging score [Left Atrial Geometry and Outcome (LAGO)] including five items (AF phenotype, structural heart disease,  $CHA_2DS_2\text{-VASc} \leq 1$ , LAD, and LA sphericity) classified patients at low ( $\leq 2$  points) and high risk ( $\geq 3$  points) of procedural failure (35% vs. 82% recurrence at 3-year follow-up, respectively; HR 3.10,  $P < 0.001$ ).



# Can CMR guide ablation?



(a) Photo of LV epicardial lesions



(b) 2x zoom of 2 lesions and gap

All ablations were visibly enhanced in non-contrast-enhanced T1w imaging using TI = 700 ms. T1w enhancement agreed with regions of necrosis in gross pathology and histology.



(c) T1w

(d) Early CE

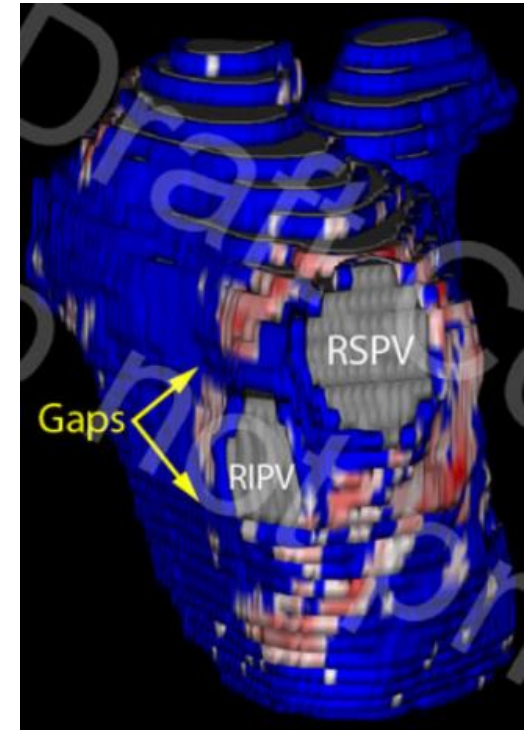
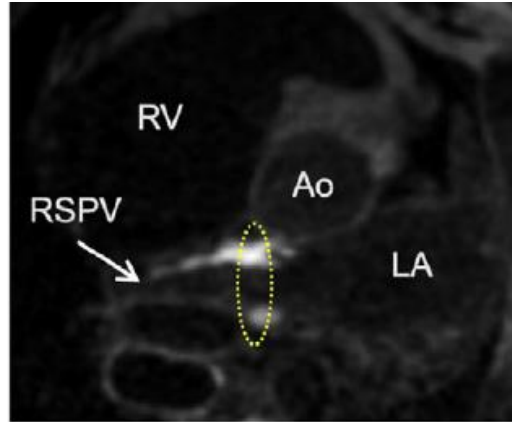
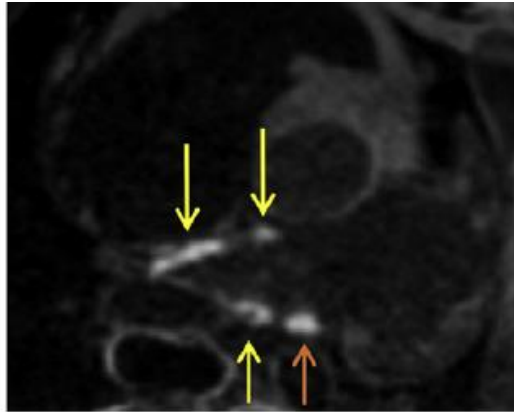
(e) Ex vivo (w/contrast)

(f) Volume rendering



# Can CMR guide AF ablation?

Non-contrast-enhanced T1-weighted (TWILITE) imaging acutely post radiofrequency ablation.



incomplete encirclement with gaps

Controversial results reported on the ability of LGE-CMR in identification of ablation gaps in patients undergoing a repeat ablation. In a study on 10 patients undergoing repeat ablation, no association was found between scar gaps in CMR and PV reconnection sites during electroanatomic mapping. However, in another study in 15 patients undergoing repeat AF ablation, the sites of electrical PV reconnection matched with a CMR gap in 79% of PVs.



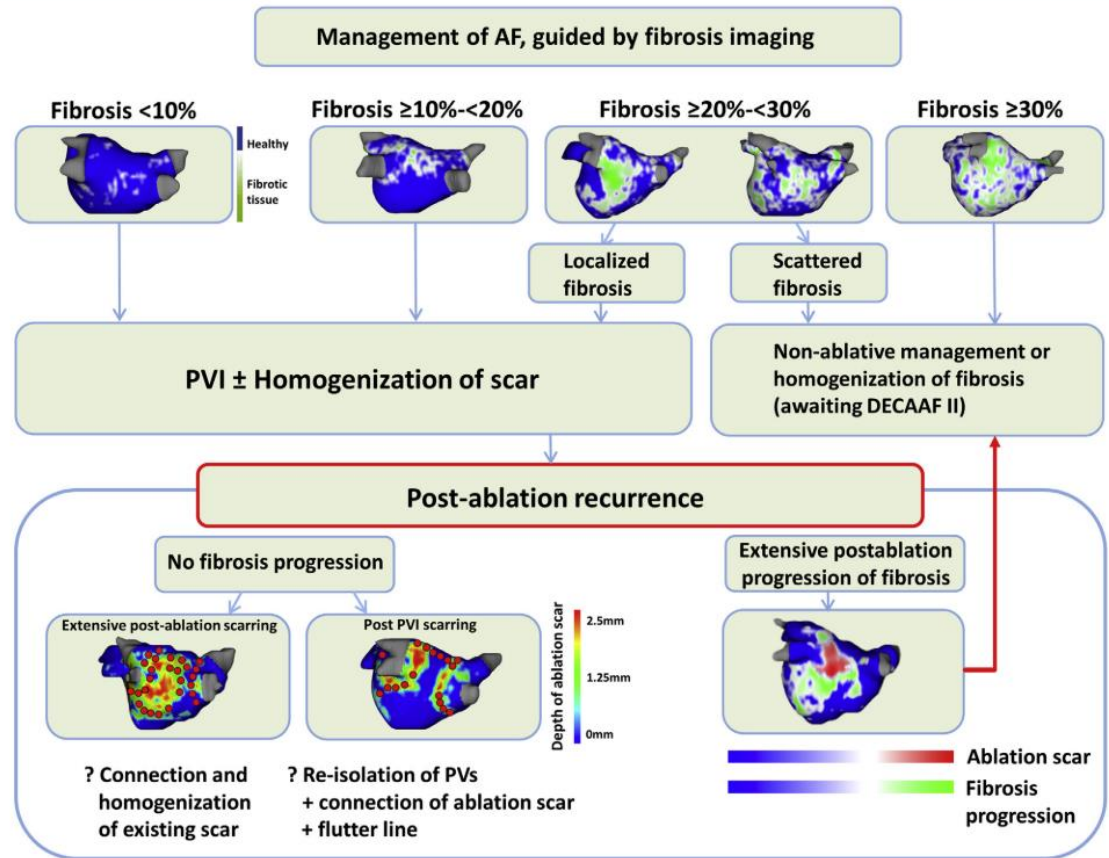
# Can CMR guide AF ablation?

Analysis of CMR images of patients from the DECAAF study reveals that in addition to preablation LGE, residual nonablated atrial fibrosis is also associated with poor outcomes.

Therefore, an approach of ablating areas of LGE detected by preablation CMR in addition to PVI may potentially eliminate the substrate and triggers.

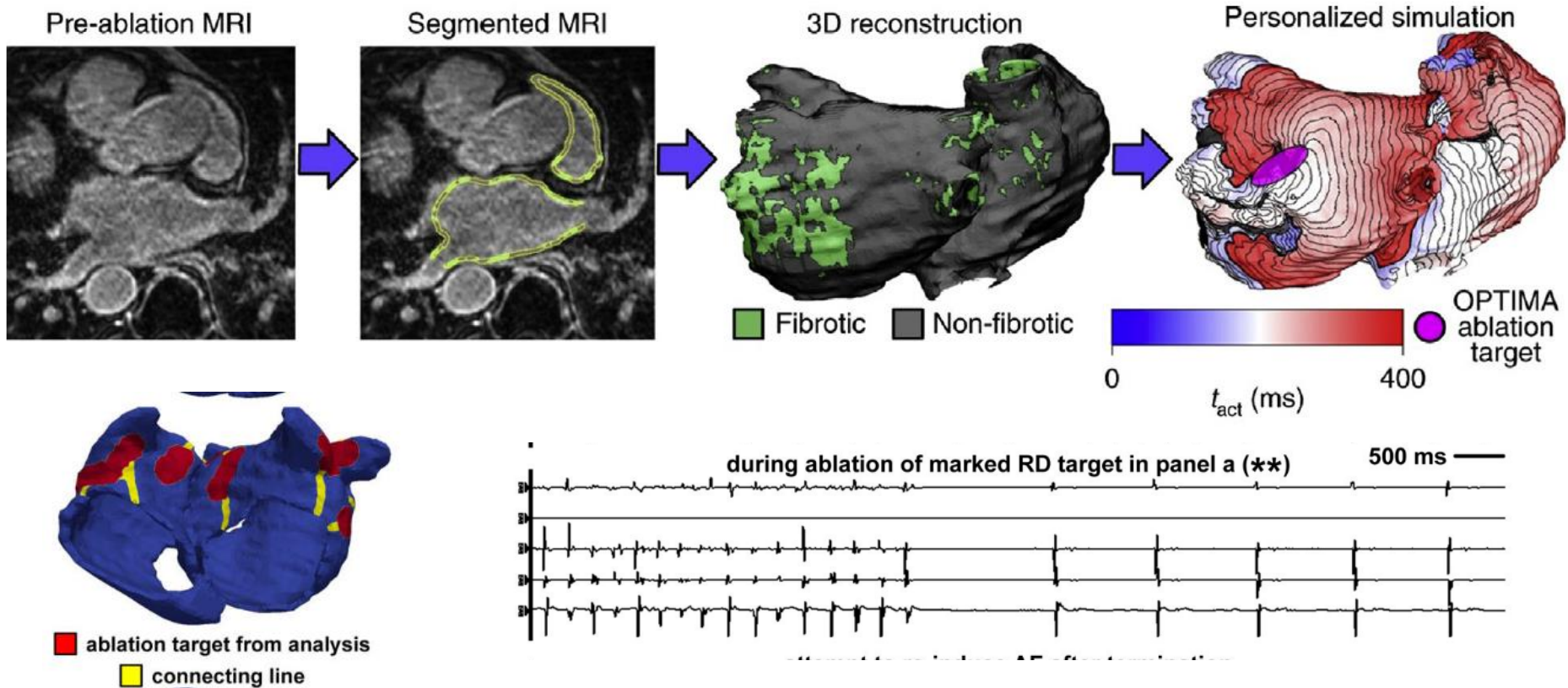
Ongoing DECAAF II trial (NCT02529319).

**FIGURE 2** Management of AF, Guided by Fibrosis Imaging





# Can CMR guide AF ablation?



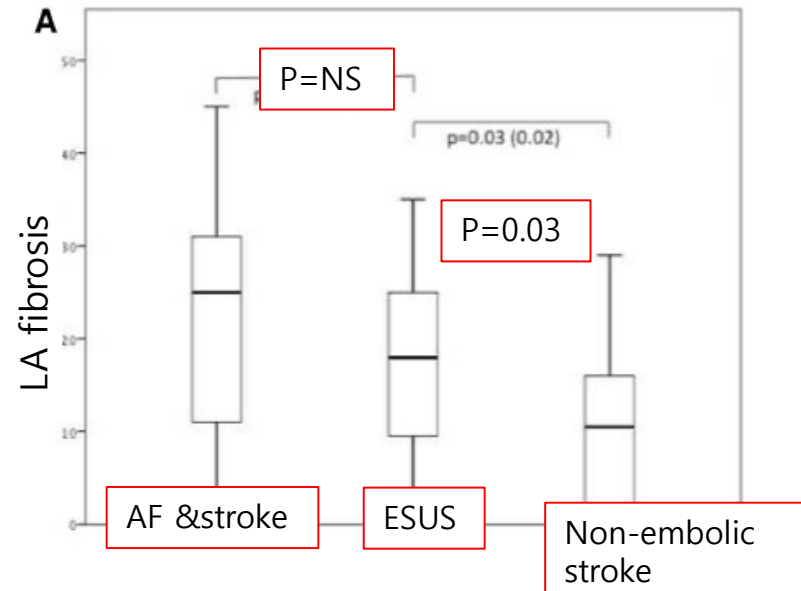
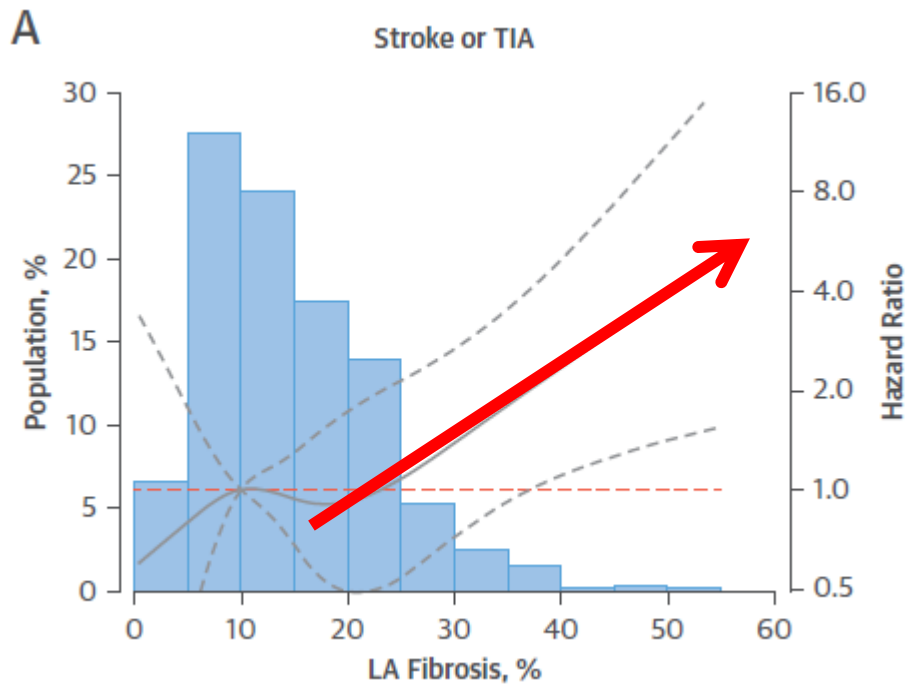
computational model of the atria of patients identifies fibrotic tissue that if ablated will not sustain AF. integrated the target-ablation sites in a clinical-mapping system, and tested its feasibility in 10 patients with persistent AF.



# Can MRI guide stroke prevention in AF?

## CARDIAC MAGNETIC RESONANCE AND ASSESSMENT FOR RISK OF STROKE

1,228 pts with AF.  
62 stroke or TIA  
Retrospective.  
LA fibrosis well correlated with stroke



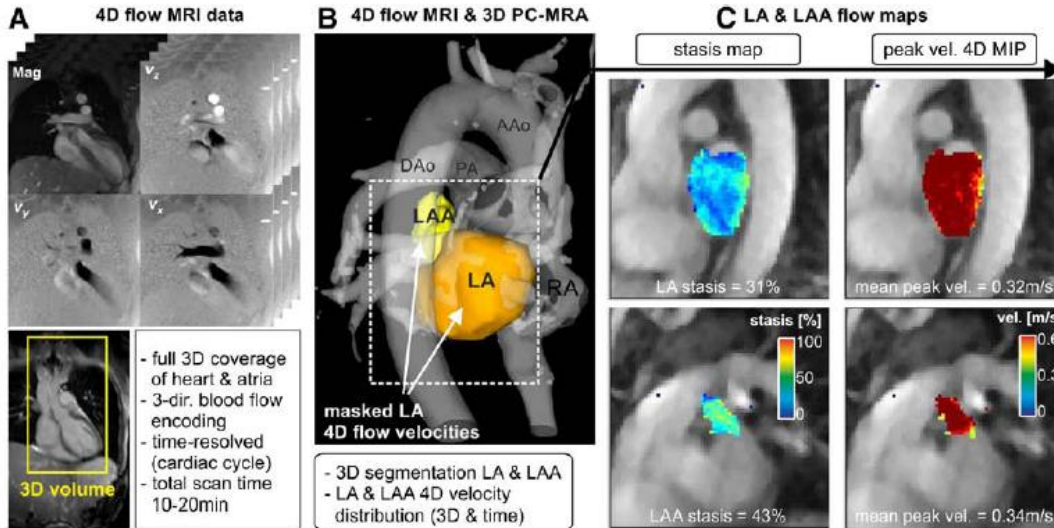
- Total 111 pts.
- ESUS a higher LA fibrosis ( $p=0.03$ )
- AF and ESUS showed a similar value of atrial fibrosis.
- an atrial disease may be associated with stroke.



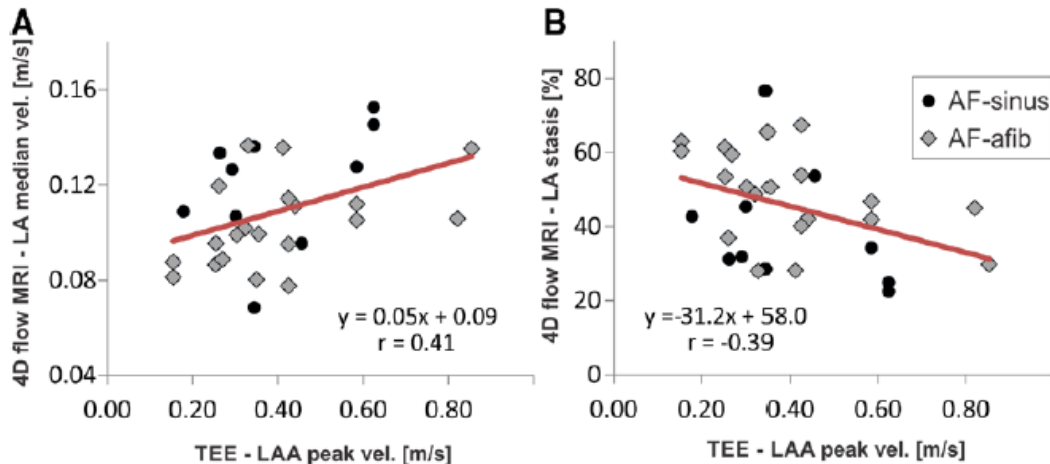
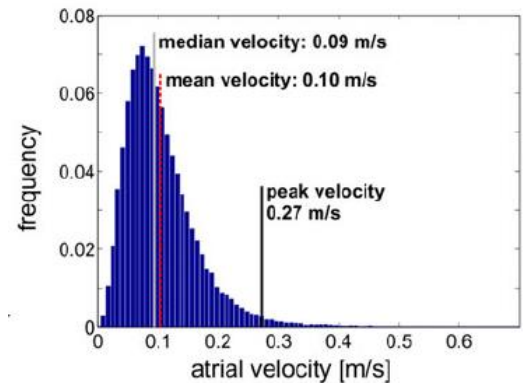
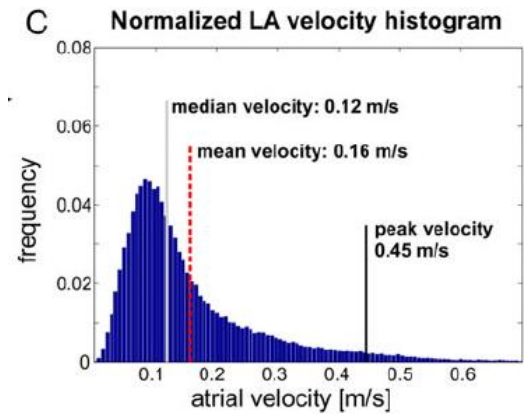


# Can MRI guide stroke prevention in AF?

## LAA flow by 4D MRI



Flow velocity was lower in AF than NSR

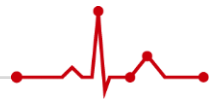


MRI-measured LAA flow velocity  
Correlated with TEE-measured velocity

M Markl et al.

Circ Cardiovasc Imaging. 2016 Sep;9(9):e004984

EJH – Cardiovascular Imaging (2016) 17, 1259–1268



## Summary

- CMR can give information on atrial structure, function, tissue characteristics, and blood flow
- Recent studies suggests AF as a part of LA myopathy. CMR has the potential of changing AF management by assessment of LA myopathy.
- It also can be used in management of patients with AF in preablation prognostication, and planning of ablation strategies.
- CMR can be used in assessment of stroke risk and, therefore, potentially can be incorporated in decision making about anticoagulation