Atrial Fibrillation and Anesthesia Considerations from an EP Perspective

Jacqueline Joza, MD FRCPC Cardiac Electrophysiologist McGill University Health Centre jacqueline.joza@mcgill.ca



Disclosures

Speaker has no conflict of interest

I also hope that I don't put you to sleep...

Because that's your job!



Objectives

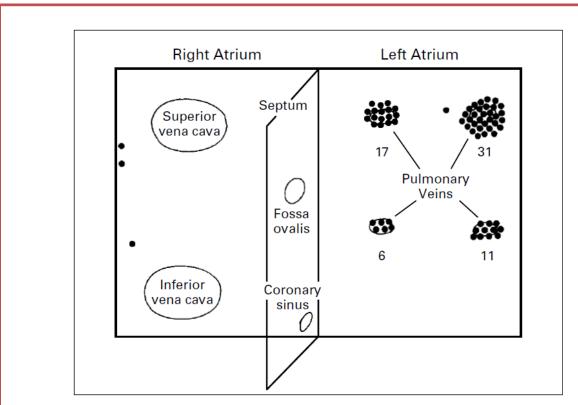
- 1. Indications and special patient populations in atrial fibrillation ablation
- 2. Mechanisms of atrial fibrillation and evolving techniques and technologies in ablation
- 3. Anesthetic considerations and techniques during ablation-Improved ablation lesion formation
 - -EP medications
 - -Esophageal considerations



NEJM, 1998

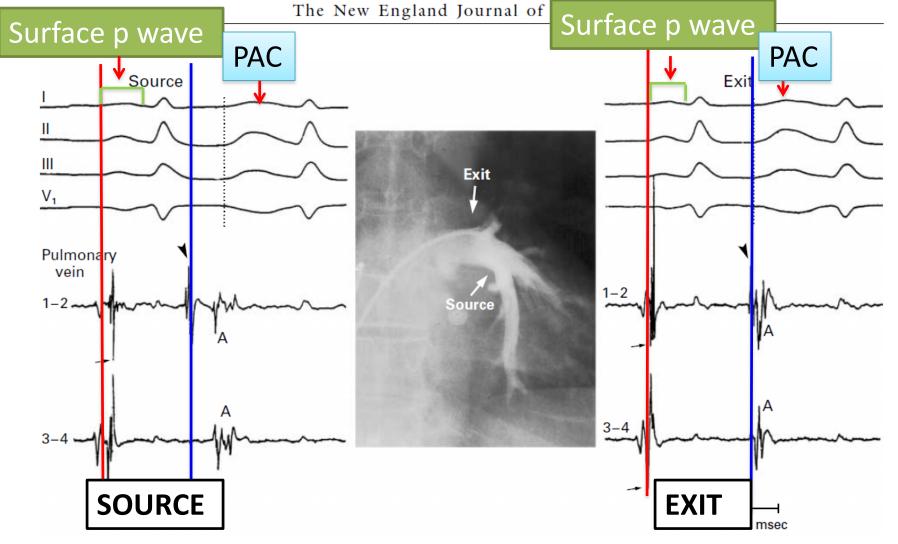
SPONTANEOUS INITIATION OF ATRIAL FIBRILLATION BY ECTOPIC BEATS ORIGINATING IN THE PULMONARY VEINS

Michel Haïssaguerre, M.D., Pierre Jaïs, M.D., Dipen C. Shah, M.D., Atsushi Takahashi, M.D., Mélèze Hocini, M.D., Gilles Quiniou, M.D., Stéphane Garrigue, M.D., Alain Le Mouroux, M.D., Philippe Le Métayer, M.D., and Jacques Clémenty, M.D.



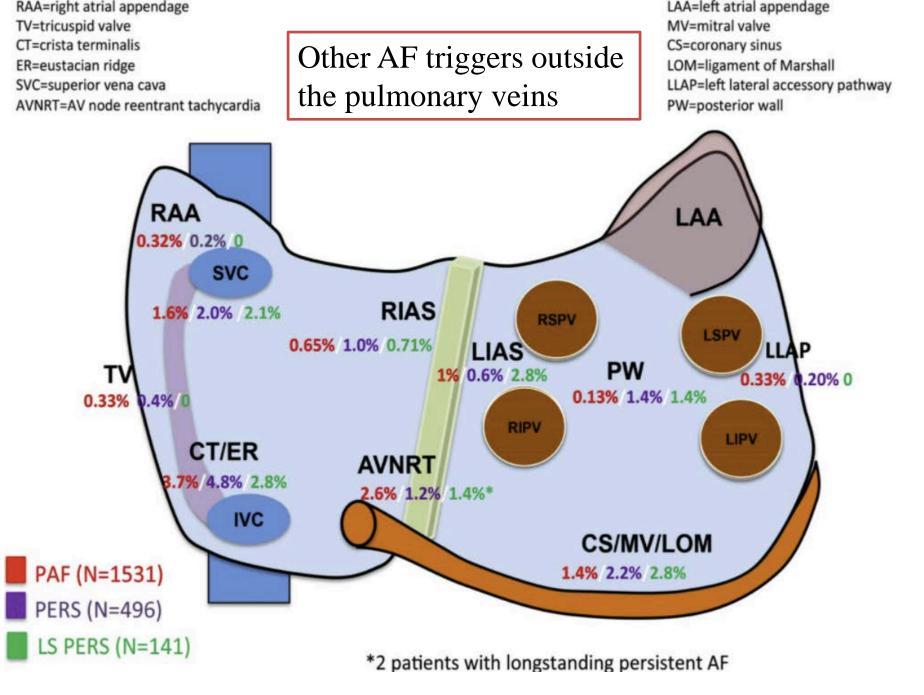
In patients with paroxysmal AF, the vast majority of atrial premature beats **originate in the pulmonary veins** (94% of all triggers).

Figure 1. Diagram of the Sites of 69 Foci Triggering Atrial Fibrillation in 45 Patients. Note the clustering in the pulmonary veins, particularly in both superior pulmonary veins. Numbers indicate the distribution of foci in the pulmonary veins.



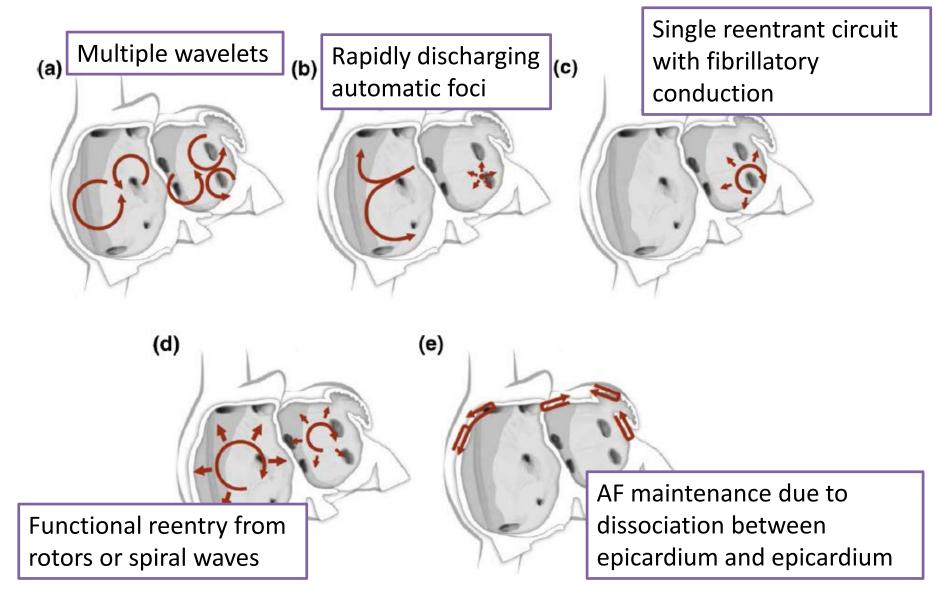


The electrogram showed characteristic changes in timing depending on the position of the recording catheter in the specific pulmonary vein. With an increasingly distal catheter position (toward the source), the spike was recorded progressively later during sinus rhythm (left-hand panel, arrows) and correspondingly earlier during ectopic activity (arrowhead). Conversely, in a proximal position at its exit into the left atrium (right-hand panel), the spike was not as delayed during sinus rhythm (arrows) nor as precocious during ectopic activity (arrowhead). The application of radio-frequency energy at the source of ectopic activity eliminated the local spike during sinus rhythm and ectopic beats and atrial fibrillation on a short-term basis. The dotted lines mark the onset of the ectopic P wave, and 1–2 and 3–4 are bipolar recordings from the distal and proximal poles of the mapping catheter. A indicates near-field atrial activity. The radiograph (center panel) shows the position of electrographic recordings inside the pulmonary vein at the source and exit.



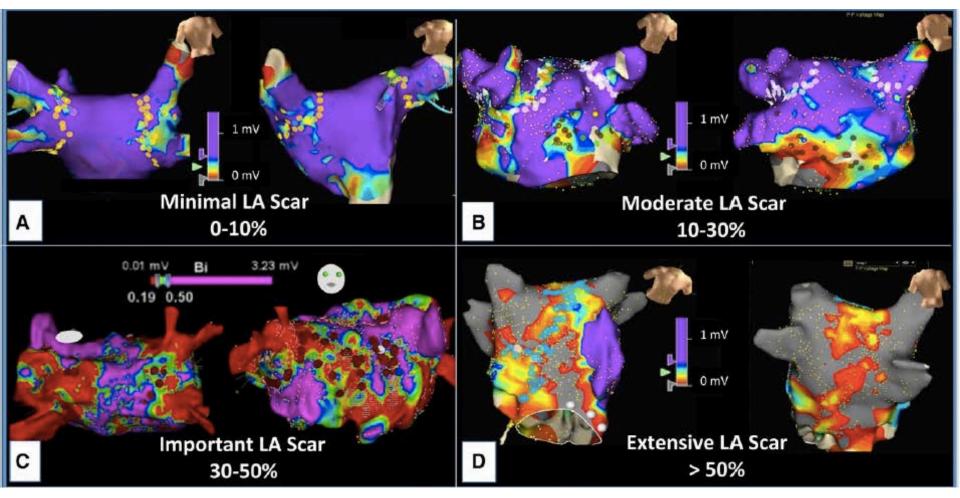
Santangeli et al. Heart rhythm July 2017. 14;7. 1087

Hypotheses regarding mechanisms of atrial fibrillation



2017 HRS expert consensus statement on catheter and surgical ablation of atrial fibrillation. J Interv Card Electrophysiol (2017) 50:1-55

Atrial Fibrosis: AF is a progressive disease



-Electroanatomic mapping allows direct contact with endocardial tissue and can reveal presence of scar (low voltage areas), not detectable by any imaging method -In contrast to paroxysmal AF, an important proportion of patients with persistent AF have regional increase in atrial fibrosis that is associated with greater frequency of AF Arentz, Circulation Arrhythmia and Electrophysiology. 2016.

Patient Populations: Who is at Risk for AF?

REVEAL-AF study: 385 patients with no history of AF (CHADS2 score of \geq 3 or 2 with 1 extra risk factor) underwent LINQ monitor implant





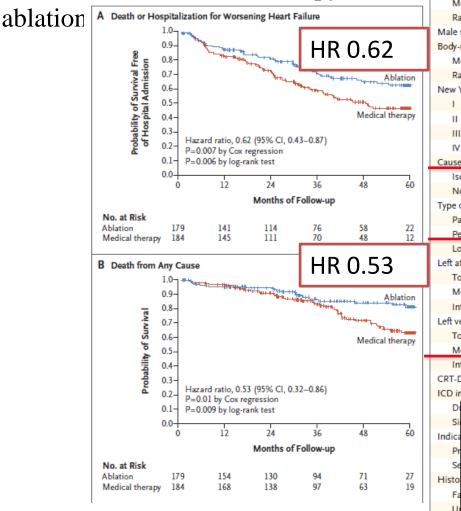
29% detection rate of AF lasting ≥ 6 min. AF would have gone undetected in most pts had monitoring been limited to 30 days. AF incidence was higher in older and more obese patients Table 2. Predictive Value of Baseline Characteristics for Atrial Fibrillation Onset

Characteristic	Hazard Ratio (95% CI) ^a	P Value
Age, y	1.08 (1.05-1.11)	<.001
Body mass index	1.04 (1.01-1.08)	.02
Male sex	1.11 (0.77-1.61)	.56
Diabetes	1.09 (0.74-1.59)	.66
Heart failure	1.08 (0.69-1.69)	.73
Hypertension	1.23 (0.58-2.60)	.58
Renal impairment	0.92 (0.64-1.32)	.65
Chronic obstructive pulmonary disease	0.73 (0.45-1.20)	.22
Stroke	0.86 (0.54-1.38)	.53
Coronary artery disease	0.78 (0.53-1.15)	.21
Sleep apnea	0.72 (0.45-1.17)	.19
Family history of atrial fibrillation	1.97 (0.76-5.14)	.16
Vascular disease	0.89 (0.56-1.43)	.63

^a Obtained from the Cox proportional hazards model.

Special Patient Populations Undergoing AF Ablation

CASTLE-AF: 398 heart failure pts with ICDs were enrolled Randomized to medical therapy vs

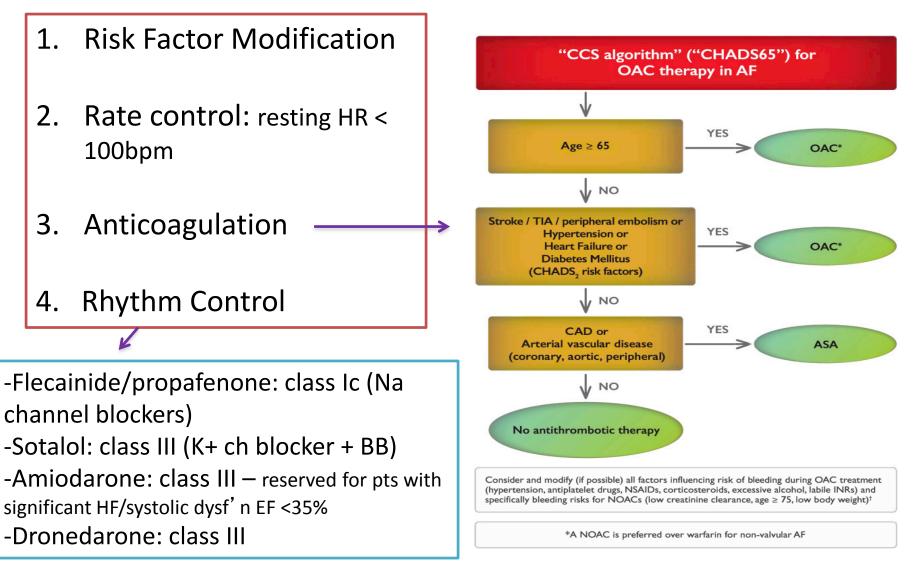


Characteristic	Treatn	nent Type
	Ablation (N=179)	Medical Therapy (N=184)
Age — yr		
Median	64	64
Range	56-71	56-73.5
Male sex — no. (%)	156 (87)	155 (84)
Body-mass index†		
Median	29.0	29.1
Range	25.9-32.2	25.9-32.3
New York Heart Association class — no./total no. (%)		
- I	20/174 (11)	19/179 (11)
11	101/174 (58)	109/179 (61)
III	50/174 (29)	49/179 (27)
IV	3/174 (2)	2/179 (1)
Cause of heart failure — no. (%) ±		
Ischemic	72 (40)	96 (52)
Nonischemic	107 (60)	88 (48)
Type of atrial fibrillation — no. (%)		
Paroxysmal	54 (30)	64 (35)
Persistent	125 (70)	120 (65)
Long-standing persistent (duration >1 year)	51 (28)	55 (30)
Left atrial diameter		
Total no. of patients evaluated	162	172
Median — mm	48.0	49.5
Interquartile range — mm	45.0-54.0	5.0-55.0
Left ventricular ejection fraction		
Total no. of patients evaluated	164	172
Median — %	32.5	31.5
Interquartile range — %	25.0-38.0	27.0-37.0
CRT-D implanted — no. (%)§	48 (27)	52 (28)
ICD implanted — no. (%) §	131 (73)	132 (72)

Significant reduction in composite outcome of death and hospitalization for heart failure and in all-cause mortality alone

(40)

4 Pillars of AF Management



2016 Focused Update of the Canadian Cardiovascular Society Guidelines for the Management of Atrial Fibrillation. CJC 32 (2016) 1170-1185

Risk of Thromboembolism

Table 2. Event rates (95% CI) and hazard ratios for hospital admission and death due to thromboembolism according to components of CHA2DS2-VASc score at 5-years follow-up

Risk Factor	Annual Risk (95% CI)	Hazard Ratio (95% CI)	Р
CHA_2DS_2 -VASc = 0	0.69 (0.59-0.81)	1.0	
CHA_2DS_2 -VASc = 1			
- Heart failure	2.35 (1.30-4.24)	3.39 (1.84-6.26)	< 0.0001
 Diabetes mellitus 	2.28 (1.42-3.66)	3.31 (2.00-5.46)	< 0.0001
- Hypertension	1.60 (1.26-2.01)	2.32 (1.75-3.07)	< 0.0001
- Age 65-74	2.13 (1.85-2.46)	3.07 (2.48-3.80)	< 0.0001
- Vascular disease	1.40 (0.91-2.15)	2.04 (1.29-3.22)	0.002
- Female sex	0.86 (0.70-1.06)	1.25 (0.96-1.63)	0.10

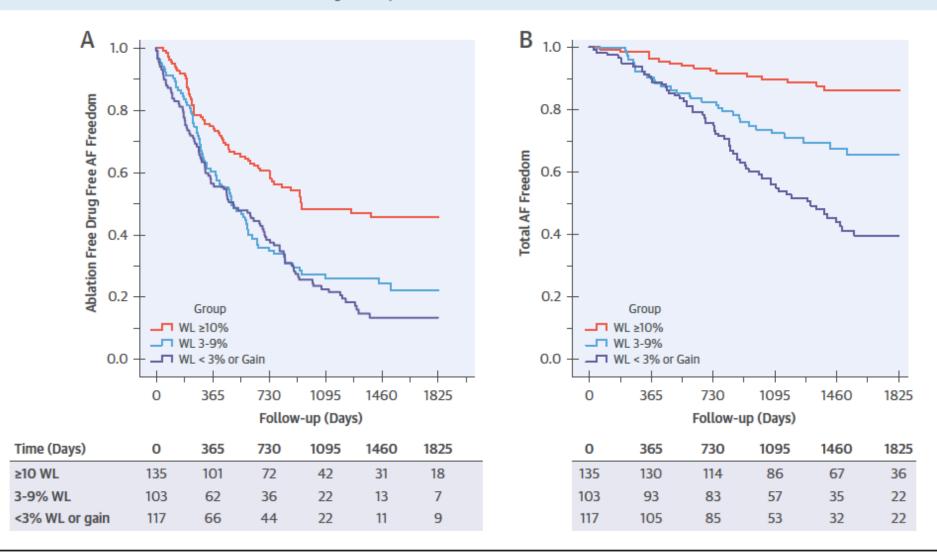
CHA₂DS₂-VASc, Congestive Heart Failure, Hypertension, Age (≥75 years), Diabetes, Stroke/Transient Ischemic Attack, Vascular Disease, Age (65-74 years), Sex (Female); CI, confidence interval. Modified from Olesen et al.²² with permission from BMJ Publishing Group Ltd.

Oral anticoagulant therapy is justified when the annual risk of stroke exceeds 1.5% OAC for patients age ≥ 65 (even without other criteria) ASA for patients with vascular disease (? questions remain)

2016 Focused Update of the Canadian Cardiovascular Society Guidelines for the Management of Atrial Fibrillation. CJC 32 (2016) 1170-1185

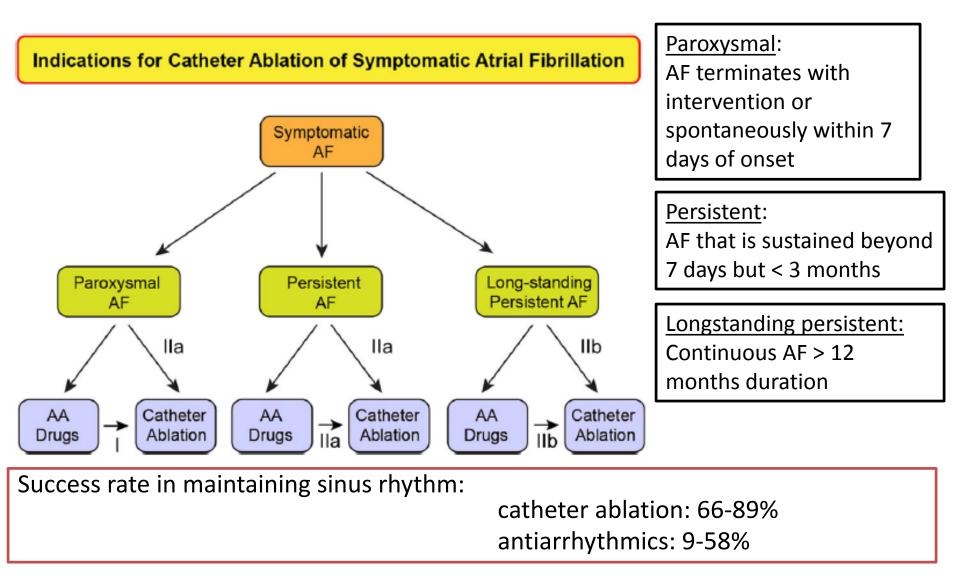
Modifiable Risk Factors: Weight loss

FIGURE 2 Atrial Fibrillation Freedom Outcome According to Group



(A) Kaplan-Meier curve for AF-free survival without the use of rhythm control strategies. (B) Kaplan-Meier curve for AF-free survival for total AF-free survival (multiple ablation procedures with and without drugs). Abbreviations as in Figure 1. Pathak et al. Weight management and AF. JACC Vol. 65, No 20. 2015.

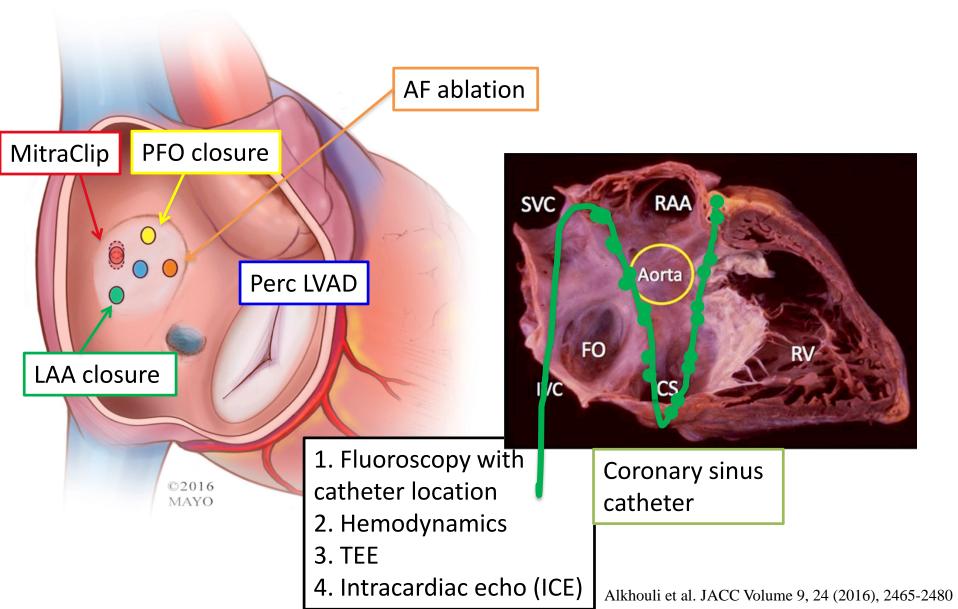
Indications for AF ablation

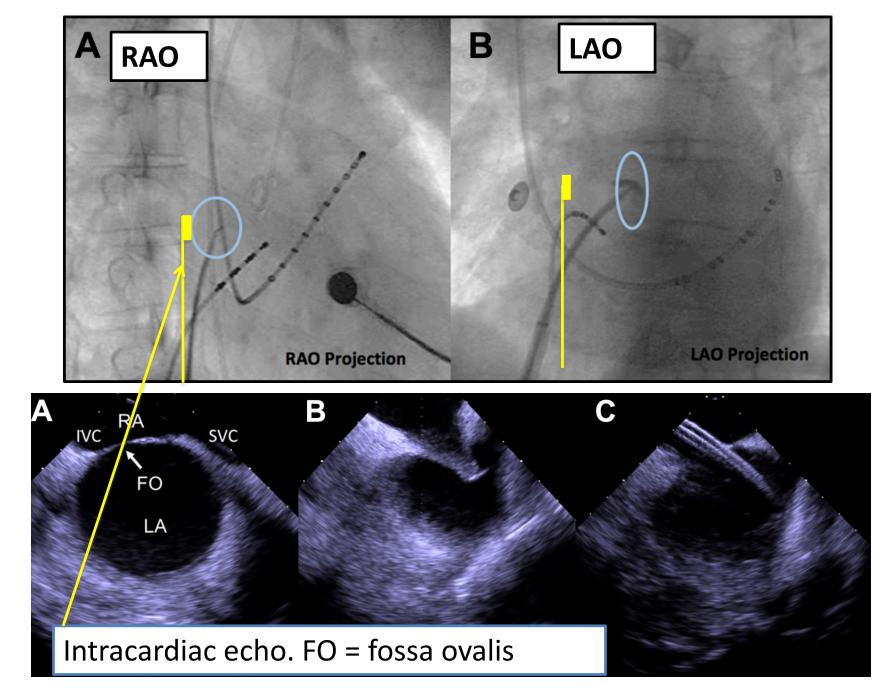


2017 HRS expert consensus statement on catheter and surgical ablation of atrial fibrillation. J Interv Card Electrophysiol (2017) 50:1-55 Contemporary AF managemenet. Andrade et al. CJC 33 (2017) 965-976

AF Ablation Procedure

Site-Specific Transseptal Puncture for Various Intracardiac Interventions





Alkhouli et al. JACC Cardiovascular Interventions. Volume 9, 24 (2016), 2465-2480

Intraprocedural Anticoagulation

Typically 5-10,000 units of heparin is bolused before transseptal

ACT during procedure = target 300-350 seconds

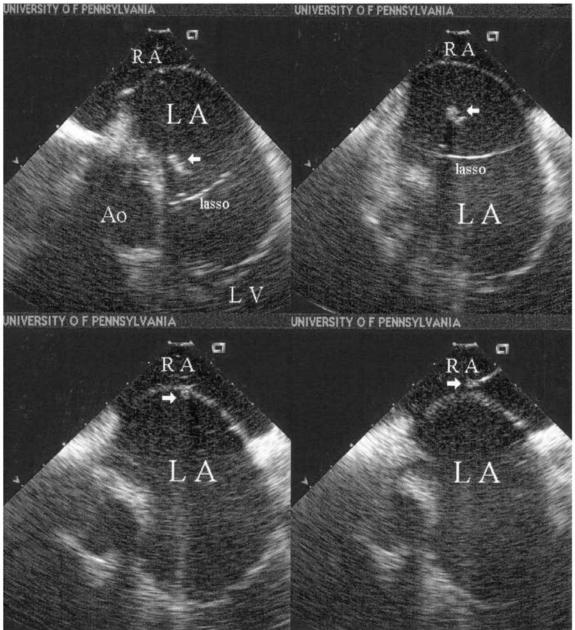
Heparin given in boluses or continuously IV

Protamine may be given at termination of procedure

<u>Periprocedural OAC</u> varies between operator.
-coumadin will typically not be stopped
-NOAC held for 1 or 2 doses, and heparin IV or a NOAC will be restarted 6 hours post sheath removal

Why is Anticoagulation Given Even Prior to Transseptal Puncture?

and restarted so soon post procedure?

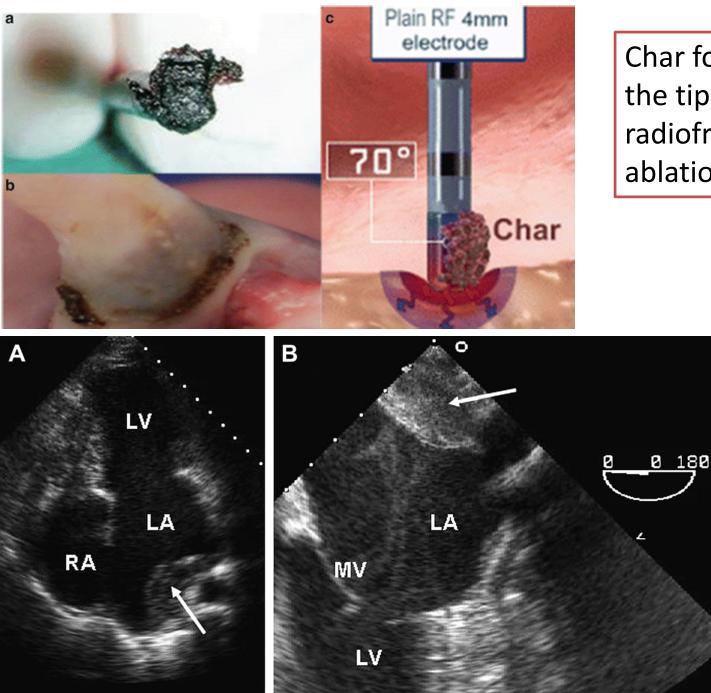


Intracardiac echo

A single linear mobile thrombus (8x4mm) attached at the sheath of the lasso catheter, pulling the sheath/lasso with thrombus back into the right atrium.

In 50%, LA thrombi occurred before RF energy application.





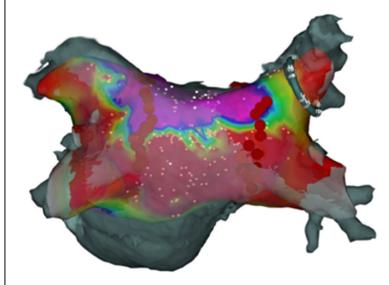
Char formation on the tip of the radiofrequency ablation catheter

> A. Transthoracic echo (apical 4-ch) demonstrating posterior LA thrombus 3 days post AF ablation B. TEE

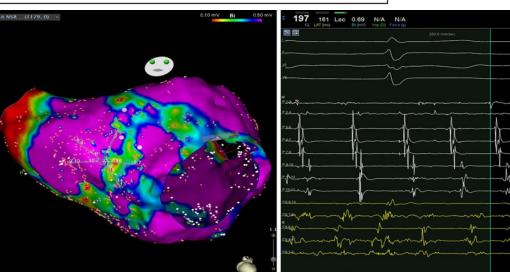
Thrombogenic milieu post abl with increased Ddimers and slow flow due to stunning

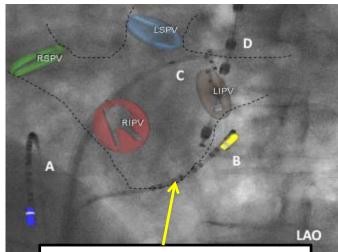
European Journal of Echocardiography, Volume 7, Issue 5, 1 October 2006, Pages 383-

AF Ablation Procedure: Mapping



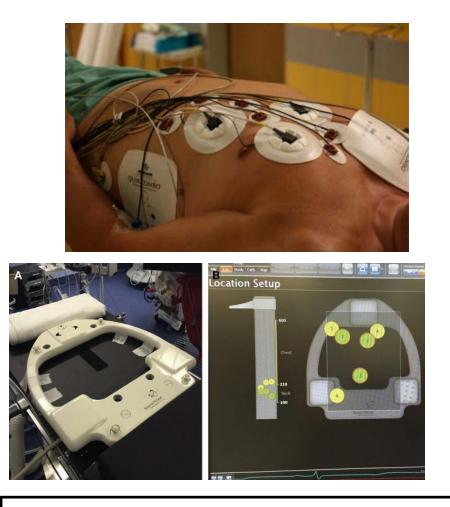
A left atrial electroanatomic map is created using a multielectrode mapping catheter (lasso or pentaray).





Coronary sinus catheter

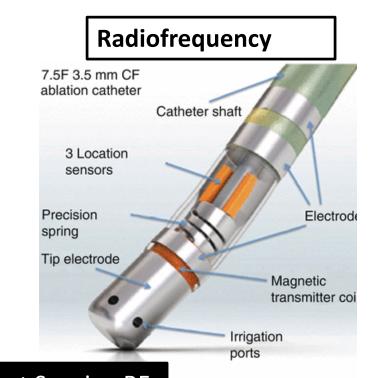
AF Ablation Procedure: importance of being still



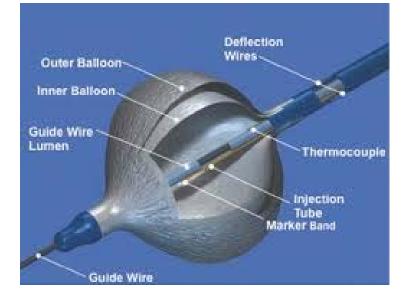
Once the mapping catheters have been introduced into the LA after transseptal, and the electroanatomic map has been started, it is essential that the patient does not move.

The mapping and ablation catheters have magnetic sensors that link to patches on the patient's front and back with a reference under the table/at torso level.

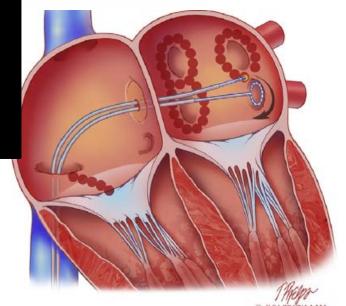
Mapping is usually started 30 minutes after achieving femoral access... typically at times when induction paralytic agent starts to wear off.

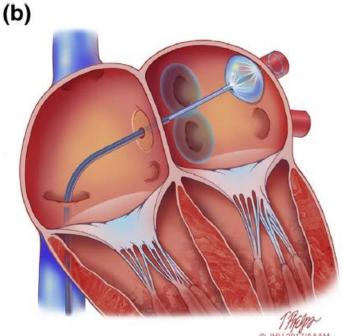


Cryoablation

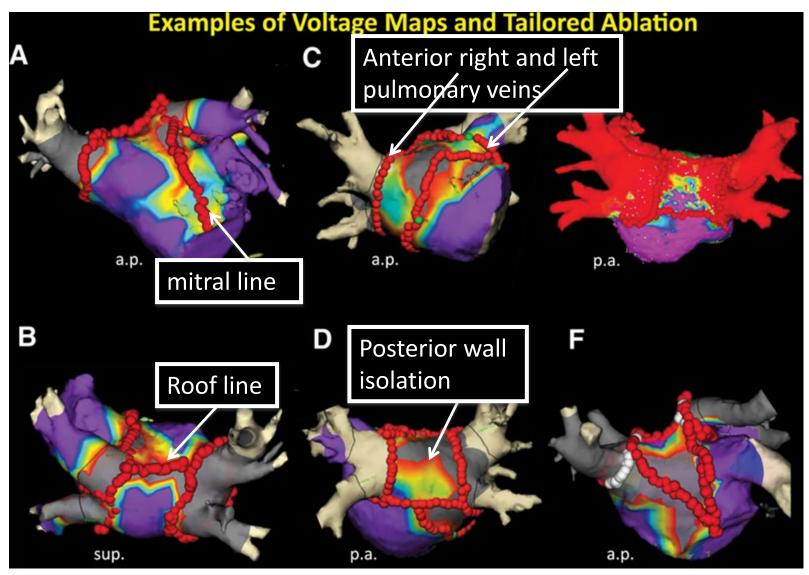


Contact Sensing RF catheters since 2014 = more efficacious lesions and safer procedure

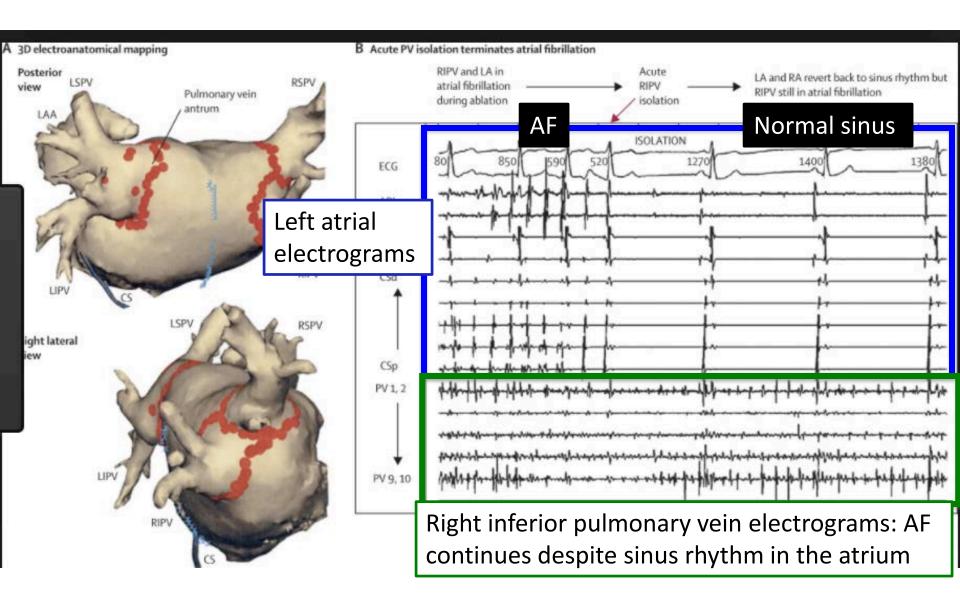




Radiofrequency Lesion Sets



Different lesion sets are created depending on location of scar, persistent vs paroxysmal AF, recurrent atrial tachycardias/flutters (red=scar, purple=healthy)



Lesion Creation and Stability

Achieving durable pulmonary vein isolation **necessitates the creation of transmural, contiguous ablation lesions** encircling the veins

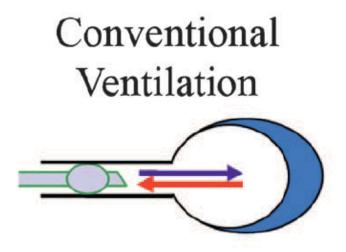
Requires the ablation catheter to maintain adequate contact with the endocardial surface to have **sufficient contact force and time-duration** to create a transmural lesion

3 things have been demonstrated to improve AF ablation outcomes:

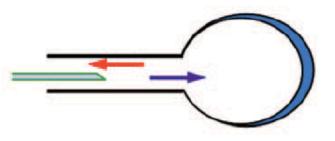
1.Incorporation of cardiac imaging (CT or MRI)2.Use of a steerable sheath3....

Anesthesia saves the day!

Jet Ventilation



High Frequency Jet Ventilation



- Gas flow is slow, bidirectional, and sequential.
- Dead space can be an issue.
- Airway sealing is mandatory.

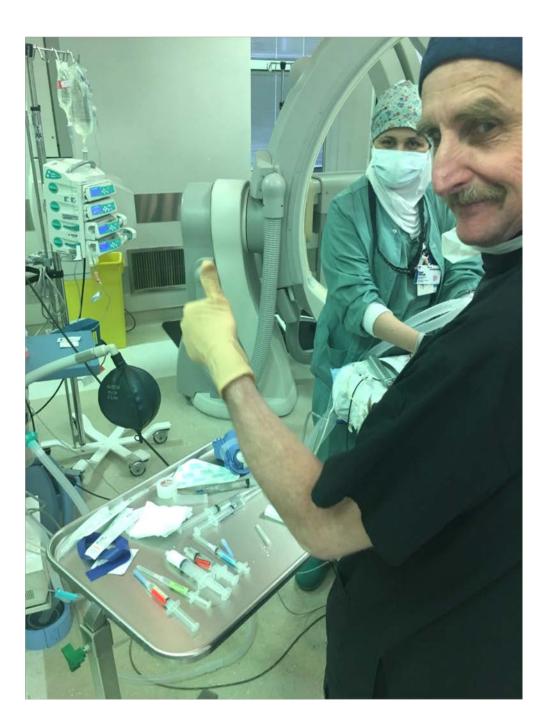
- High velocity insufflation of gas through a narrow nozzle into the open airway.
- Gas flow is fast, coaxial,

partially simultaneous, inward only.

- Dead space is less relevant.
- Airway sealing contraindicated.

Thumbs up for Jet

-Dr. David Bracco



Efforts to enhance catheter stability improve atrial fibrillation ablation outcome

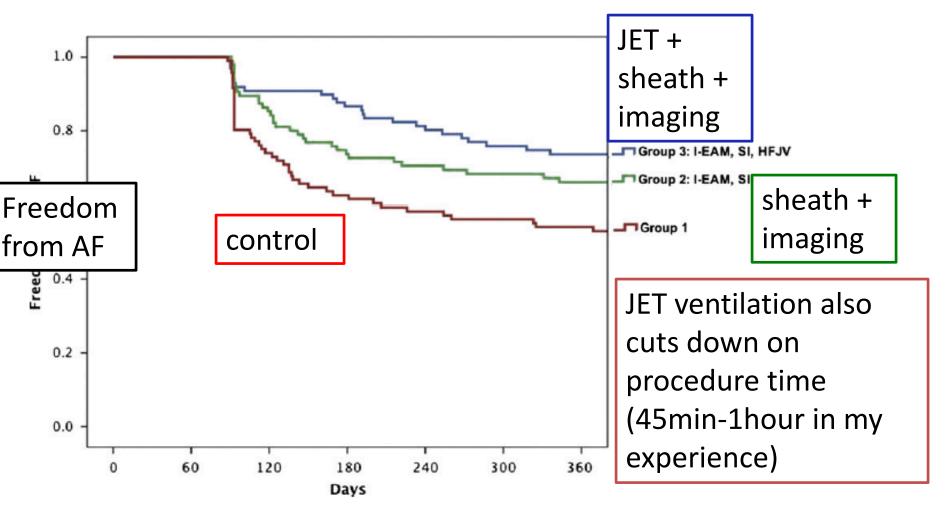
Mathew D. Hutchinson, MD, FHRS,^{*} Fermin C. Garcia, MD,^{*} Jeff E. Mandel, MD,MS,[†] Nabil Elkassabany, MD,[†] Erica S. Zado PA-C, FHRS,^{*} Michael P. Riley, MD,^{*} Joshua M. Cooper, MD,^{*} Rupa Bala, MD,^{*} David S. Frankel, MD,^{*} David Lin, MD,^{*} Gregory E. Supple, MD,^{*} Sanjay Dixit, MD, FHRS,^{*} Edward P. Gerstenfeld, MD, FHRS,^{*} David J. Callans, MD, FHRS,^{*} Francis E. Marchlinski, MD, FHRS^{*}

Table 1 Demographic data	Control	sheath + imaging	JET + sheath + imaging	
	Group 1	Group 2 (I-EAM, SI)	Group 3 (I-EAM, SI, HFJV)	Р
Sex: Man	66%	85%	75%	.008
Age (y)	57 ± 11	60 ± 10	59 ± 10	.03
BMI (kg/m ²)	28.5 ± 5.8	29.1 ± 4.8	31.2 ± 5.4	<.001
CAD	13%	16%	16%	.79
Hypertension	48%	64%	64%	.03
Diabetes mellitus	9%	10%	15%	.36
TIA/CVA	5%	3%	4%	.78
MR (≥moderate)	1%	4%	6%	.17
CM	6%	14%	15%	.09
CHF	2%	2%	6%	.19
OSA	12%	16%	20%	.30
LVEF (%)	62 ± 8	60 ± 10	$59 \pm 10 \\ 4.5 \pm 0.8$.22
LA diameter (cm)	4.2 ± 0.8	4.4 ± 0.7		.002
AF type Paroxysmal Persistent Permanent	83 6 11	70 26 4	61 28 11	.002

AF = atrial fibrillation; BMI = body mass index; CAD = coronary artery disease; CHF = congestive heart failure; HFJV = high-frequency jet ventilation; I-EAM = 3-dimensional image integration; LA = left atrium; LVEF = left ventricular ejection fraction; MR = mitral regurgitation; OSA = obstructive sleep apnea; SI = steerable introducer; TIA/CVA = transient ischemic attack/cerebrovascular accident.

300 patients undergoing AF ablation

Despite a sicker population, Jet ventilation added significant improvement to 1-year freedom from AF (52% vs 66% vs 74%)



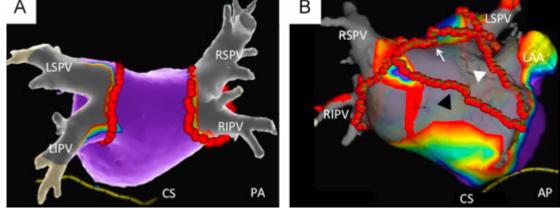
Hutchinson Heart Rhythm, Vol 10, No 3, March 2013

Apnea

EP may request apnea be performed **where difficulty with catheter stability is encountered:**

-typically left atrial roof, Marshall ridge (anterior to the left pulmonary veins), and at the anterior aspect of the right superior pulmonary vein

-apnea significantly improves catheter stability and lesion formation



During Ablation: Fluctuations in HR and BP

Adenosine is frequently given post pulmonary vein isolation to confirm isolation

Adenosine acts to hyperpolarize cells, thereby increasing conduction times and uncovering potentially dormant conduction/Afib

A brief period of bradycardia and heart block is observed with resultant brief period of hypotension.

Isoproterenol (non-selective β adrenoreceptor agonist) is given post ablation in attempts to induce atrial tachycardias or atrial fibrillation.

Hypotension may be caused with isoproterenol alone or when atrial pacing is performed at the same time

Potential Complications of AF Ablation

Signs and symptoms of complications within a month postablation

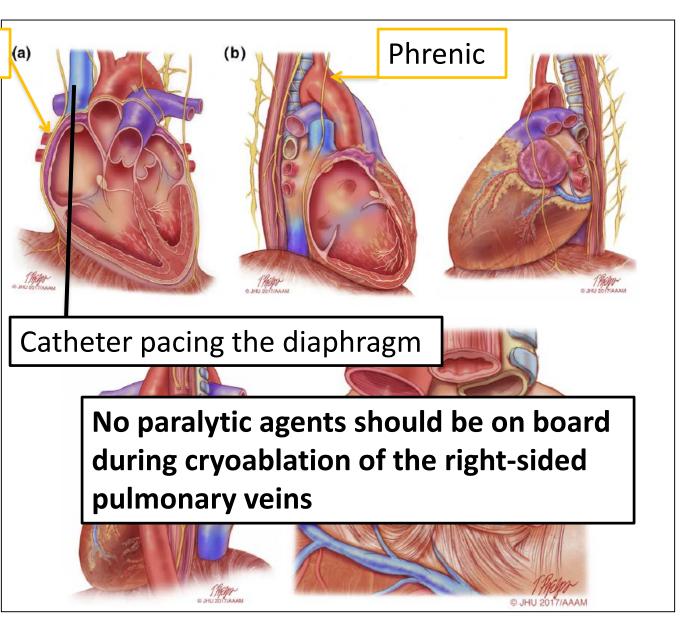
Back pain	Musculoskeletal, retroperitoneal hematoma
Chest pain	Pericarditis, pericardial effusion, coronary stenosis (ablation related), pulmonary vein stenosis, musculoskeletal (after cardioversion), worsening reflux
Cough	Infectious process, bronchial irritation (mechanical, cryoballoon), pulmonary vein stenosis
Dysphagia	Esophageal irritation (related to transesophageal echocardiography), atrioesophageal fistula
Early satiety, nausea	Gastric denervation
Fever	Infectious process, pericarditis, atrioesophageal fistula
Fever, dysphagia, neurological symptoms	Atrial esophageal fistula
Groin pain at site of access	Pseudoaneurysm, AV fistula, hematoma
Headache	Migraine (related to anesthesia or transseptal access, hemorrhagic stroke), effect of general anesthetic
Hypotension	Pericardial effusion/tamponade, bleeding, sepsis, persistent vagal reaction
Hemoptysis	PV stenosis or occlusion, pneumonia
Neurological symptoms	Cerebral embolic event, atrial esophageal fistula
Shortness of breath	Volume overload, pneumonia, pulmonary vein stenosis, phrenic nerve injury
Signs and symptoms of complicat	ions more than a month postablation
Fever, dysphagia, neurological symptoms	Atrial esophageal fistula
Persistent cough, atypical chest pain	Infectious process, pulmonary vein stenosis
Neurological symptoms	Cerebral embolic event, atrial esophageal fistula
Hemoptysis	PV stenosis or occlusion, pneumonia

Phrenic nerve paralysis:

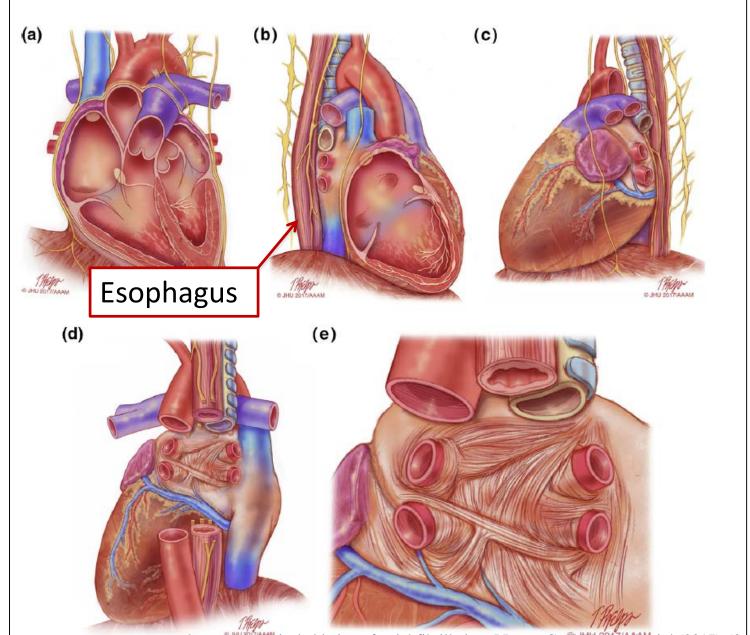
Typically resolves but may be permanent. Occurs during radiofrequency ablation at the right superior pulmonary vein and during cryoablation of the right-sided veins.

Phrenic

Diaphragmatic pacing is performed with a catheter at the SVC/RA



Esophageal Injury:



2017 HRS expert consensus statement on catheter and surgical ablation of atrial fibrillation. J Interv Card Electrophysiol (2017) 50:1-55

Not only does the esophagus have a variable position within the thoracic cavity, but can also vary temporally and based on patient position

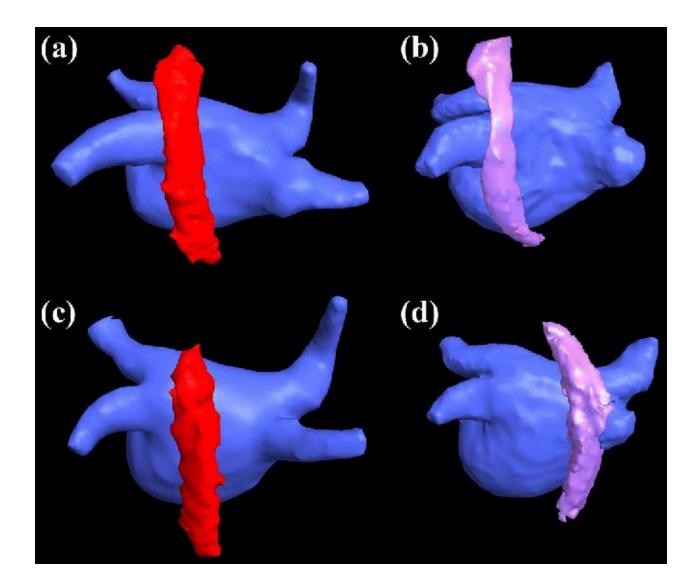
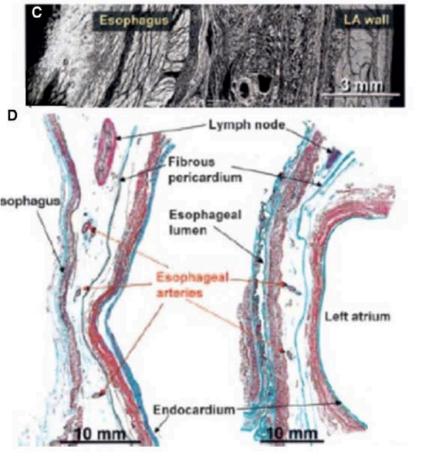




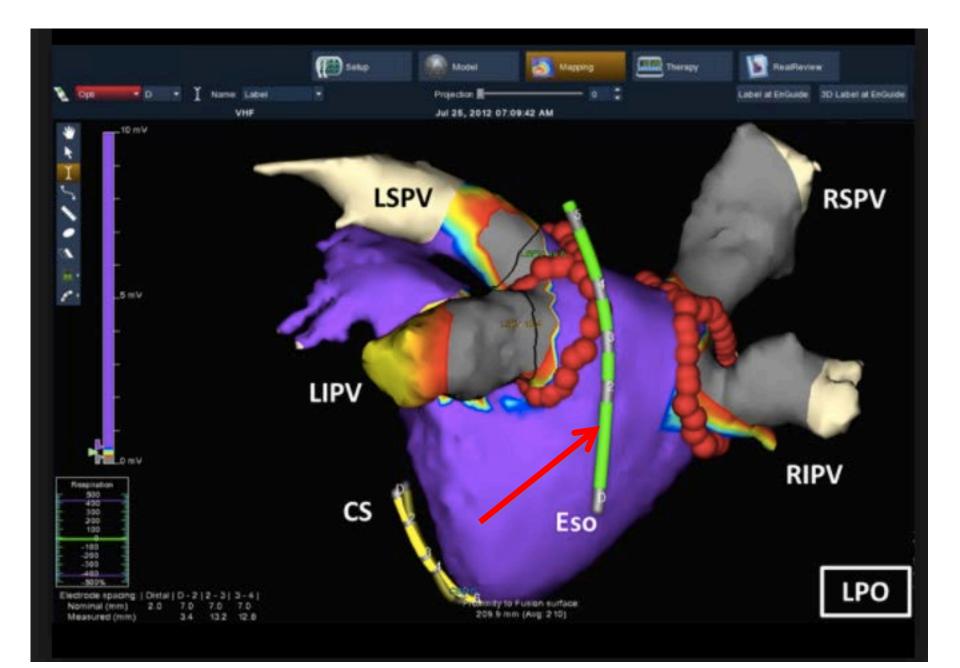
Figure 2. Different aspects of esophageal thermal lesions related to atrial fibrillation ablation as seen on upper endoscopy. A, Erythema without disruption of the esophageal mucosa. B, Esophageal ulcer showing mucosal disruption and injury. C, Necrotizing esophageal ulcer.



Esophageal heating is a mechanism for injury, but fistulas do occur in cryoablation as well

Can an esophageal temperature probe help to prevent fistulas?

Kapur et al. Circulation. 2017;136:1247–1255.



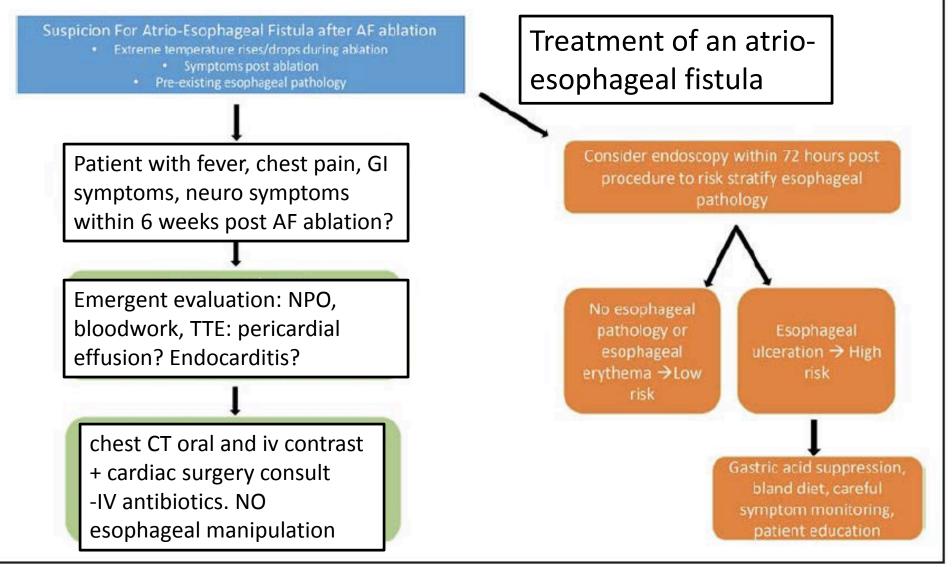
lesions occurred in pts with esophageal temp > 41° C (every 1° C increase in esophageal temperature = 1.36 increased odds of esophageal lesion)

Is there harm associated with temperature probes?

30% vs 2.5% incidence of esophageal injury in the temperature probe group (n=80 patients)

-Some harm has been associated with the type of temperature probe: temp probes with noninsulated metal thermistors may function as lightning rods, attracting electric current form the ablation catheter and potentiating heat transfer to the esophagus.

-Others suggest that use of a temp probe with insulated thermocouples was as safe compared with ablations performed without a temperature probe



If a patient presents with fever, chest pain, GI or neuro sx within 6 weeks of AF ablation: NO TEE, NO ENDOSCOPY. DO CT SCAN, DIAGNOSE ESOPHAGEAL FISTULA AND SURGERY (NOT STENTING)

Pericarditis

Most patients experience pleuritic chest pain post procedure as a result of transmural ablation lesions, lasting up to 48-72 hours

Significant improvement with Ketorolac given at time of procedure, prior to leaving the operating room

Conclusions

- Early atrial fibrillation originates at the muscular sleeves of the pulmonary veins. With time, scarring in the left atrium can occur resulting in a more complex ablation procedure
- 2. New techniques in atrial fibrillation, including contact force sensing catheters, new-generation cryo-balloons, and mapping software have translated into higher success rates with reduced complications. Sicker patients are being ablated
- 3. Importance of Anesthesia and EP communication during ablation: issues with paralysis, patient movement, improved catheter stability (JET ventilation/apnea)
- 4. Esophageal considerations to reduce complications
- 5. Specific EP medications used during ablation
- 6. And...



Big thank you to all our EP anesthesia teams across Canada, and our team here at McGill





Dr. Bondi, Dr. Bracco, Dr. Predescu, Dr. Owen, Dr. Guzzo, Dr. Donatelli, Dr. Ah-kye, and Dr. Baldini, and the rest of the EP anesthesia team

