

Ablation de FA

Pour qui et quand ?

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Disclosures

- ◆ Consulting and Speaker's fees from Biotronik, Medtronic, Boston Scientific, Abbott, Microport



European Society
of Cardiology

European Heart Journal (2020) 00, 1–125
doi:10.1093/eurheartj/ehaa612

ESC GUIDELINES

2020 ESC Guidelines for the diagnosis and management of atrial fibrillation developed in collaboration with the European Association of Cardio-Thoracic Surgery (EACTS)

The Task Force for the diagnosis and management of atrial fibrillation of the European Society of Cardiology (ESC)

Developed with the special contribution of the European Heart Rhythm Association (EHRA) of the ESC

Authors/Task Force Members: Gerhard Hindricks* (Chairperson) (Germany), Tatjana Potpara* (Chairperson) (Serbia), Nikolaos Dagres (Germany), Elena Arbelo (Spain), Jeroen J. Bax (Netherlands), Carina Blomström-Lundqvist (Sweden), Giuseppe Boriani (Italy), Manuel Castella¹ (Spain), Gheorghe-Andrei Dan (Romania), Polychronis E. Dilaveris (Greece), Laurent Fauchier (France), Gerasimos Filippatos (Greece), Jonathan M. Kalman (Australia), Mark La Meir¹

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Table 18 Principles of antiarrhythmic drug therapy

Principles

AAD therapy aims to reduce AF-related symptoms

Efficacy of AADs to maintain sinus rhythm is modest

Clinically successful AAD therapy may reduce rather than eliminate AF recurrences

If one AAD 'fails', a clinically acceptable response may be achieved by another drug

Drug-induced proarrhythmia or extracardiac side-effects are frequent

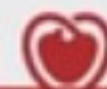
Safety rather than efficacy considerations should primarily guide the choice of AAD

Recommendations for long-term antiarrhythmic drugs (1)

Reduces CV hospitalizations & death
Most solid safety data.

Recommendations	Class	Level
Flecainide or propafenone is recommended for long-term rhythm control in AF patients with normal LV function and without structural heart disease , including significant LVH and myocardial ischaemia.	I	A
Dronedarone is recommended for long-term rhythm control in AF patients with: • Normal or mildly impaired (but stable) LV function, or • HFpEF, ischaemic, or valvular heart disease.	I	A
Amiodarone is recommended for long-term rhythm control in all AF patients, including those with HFrEF . However, owing to its extracardiac toxicity, other AADs should be considered first whenever possible.	I	A
Sotalol may be considered for long-term rhythm control in patients with normal LV function or with ischaemic heart disease if close monitoring of QT interval, serum potassium levels, creatinine clearance, and other proarrhythmia risk factors is provided.	IIb	A

Recommendations for rhythm control/**catheter ablation** of AF (1)



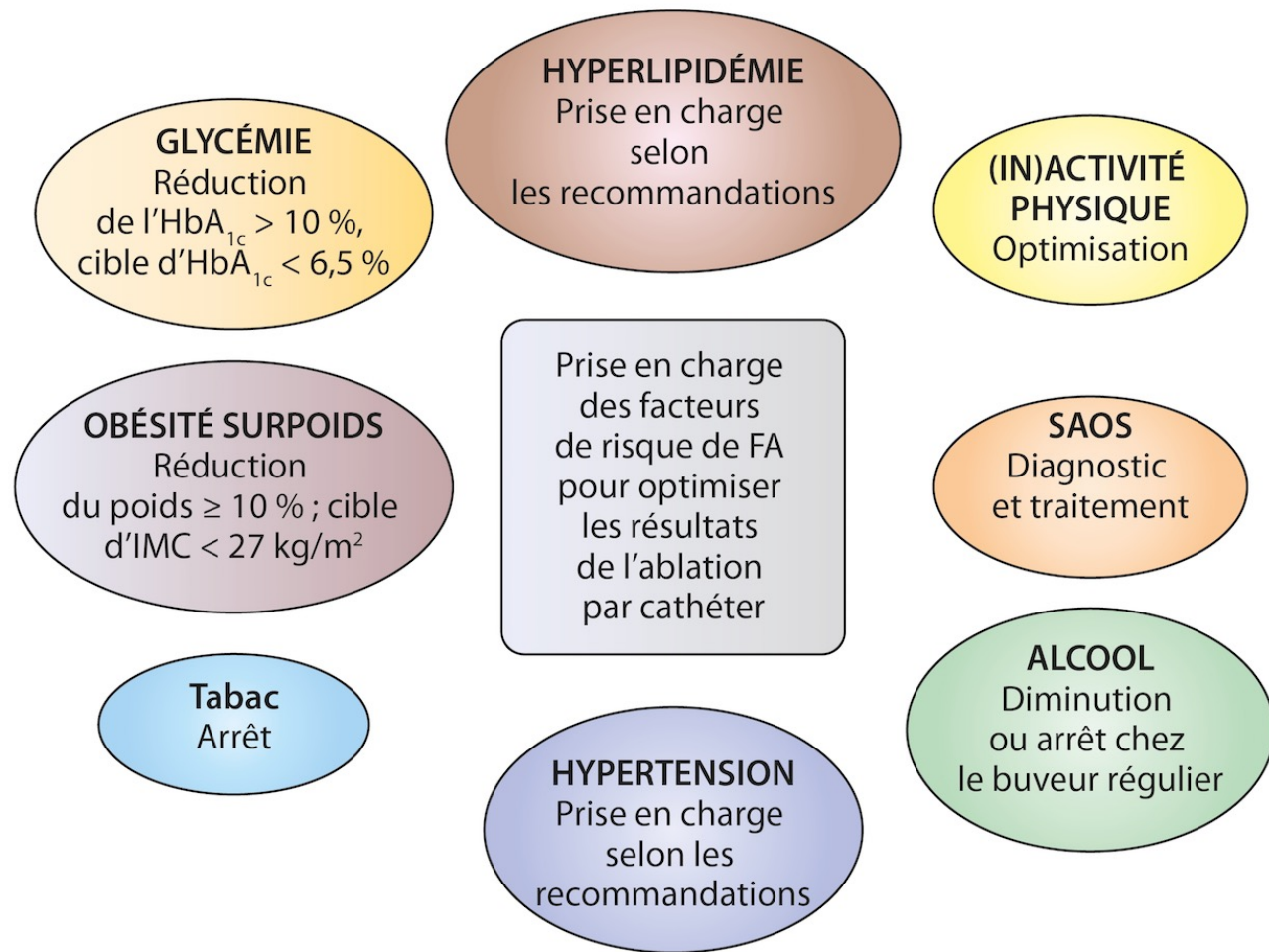
ESC

European Society
of Cardiology

New !

Recommendations	Class	Level
<i>General recommendations</i>		
For the decision on AF catheter ablation , it is recommended to take into consideration the procedural risks and the major risk factors for AF recurrence following the procedure and discuss them with the patient.	I	B
Repeated PVI procedures should be considered in patients with AF recurrence provided the patient's symptoms were improved after the initial PVI.	IIa	B

New !



**Recommandations de style de vie et prise en charge
des facteurs de risque et des comorbidités chez les patients avec FA**

La prise en charge optimale du SAOS peut être envisagée pour réduire
l'incidence de la FA, sa progression, ses récives et les symptômes

IIb

Le traitement du SAOS devrait être optimisé pour réduire les
et améliorer les résultats du traitement de la FA

IIa

Figure 3. Recommandations 2020 concernant la prise en charge de la FA⁽⁸⁾.

Syndrome d'apnées obstructives du sommeil

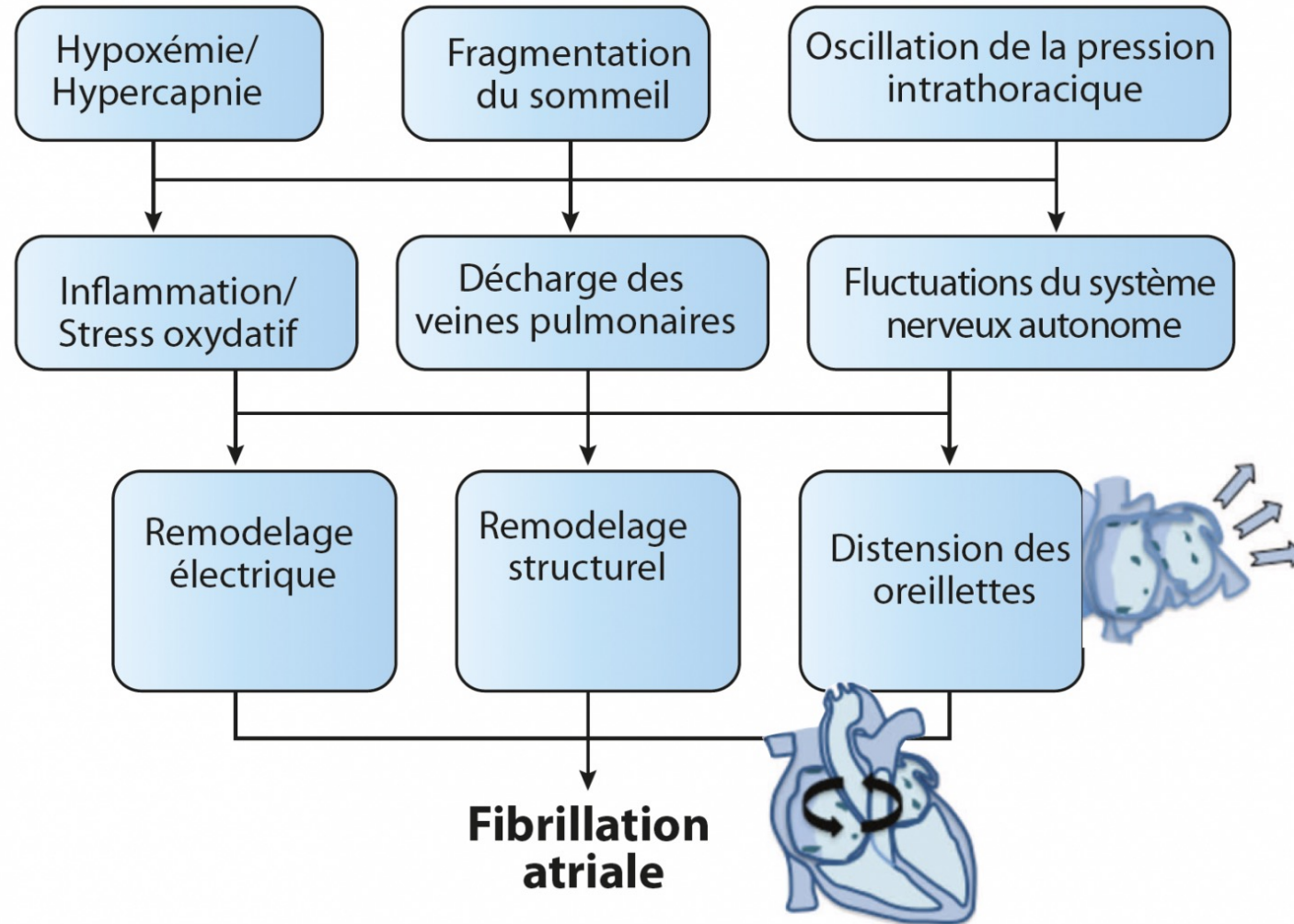


Figure 1. Mécanismes physiopathologiques du syndrome d'apnées obstructives du sommeil contribuant à la fibrillation atriale.

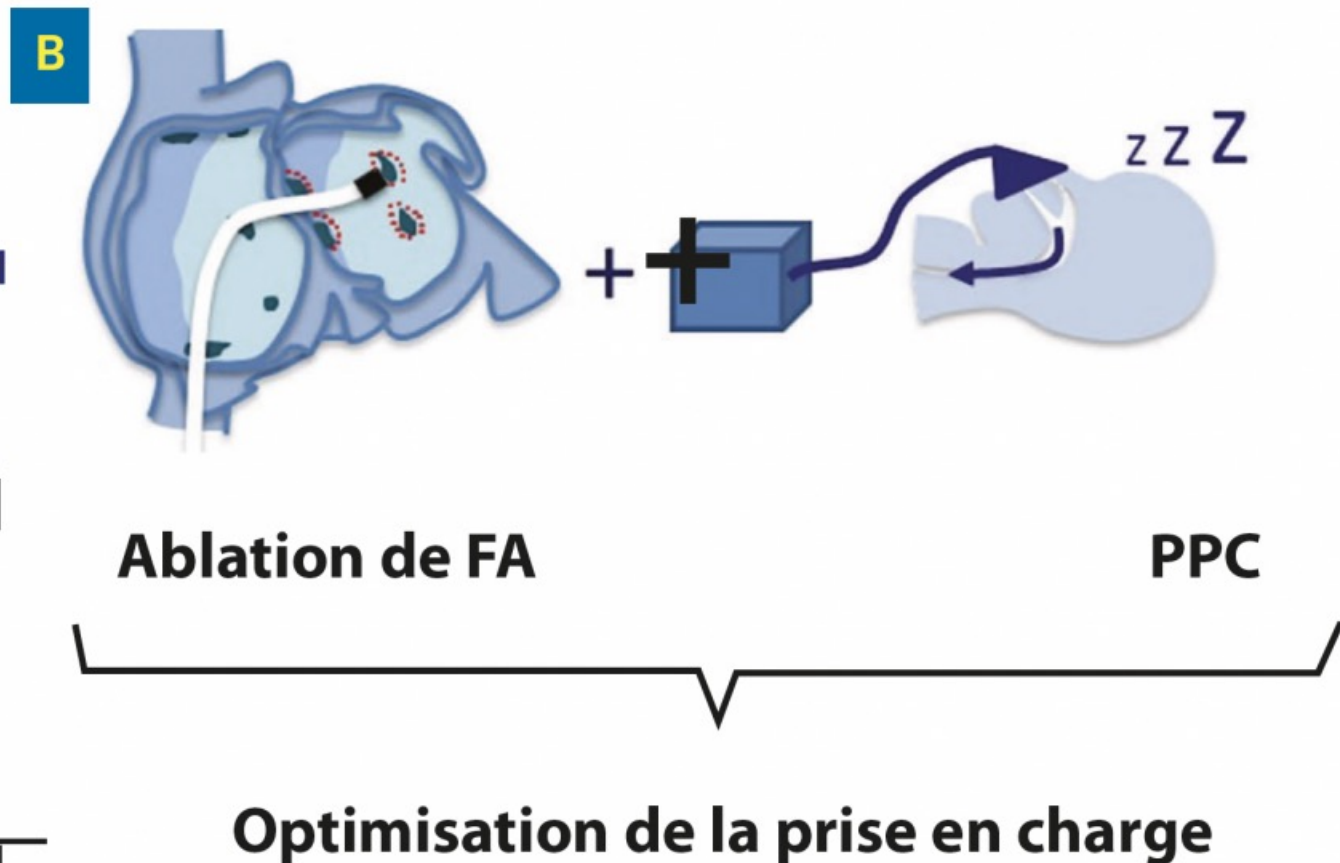
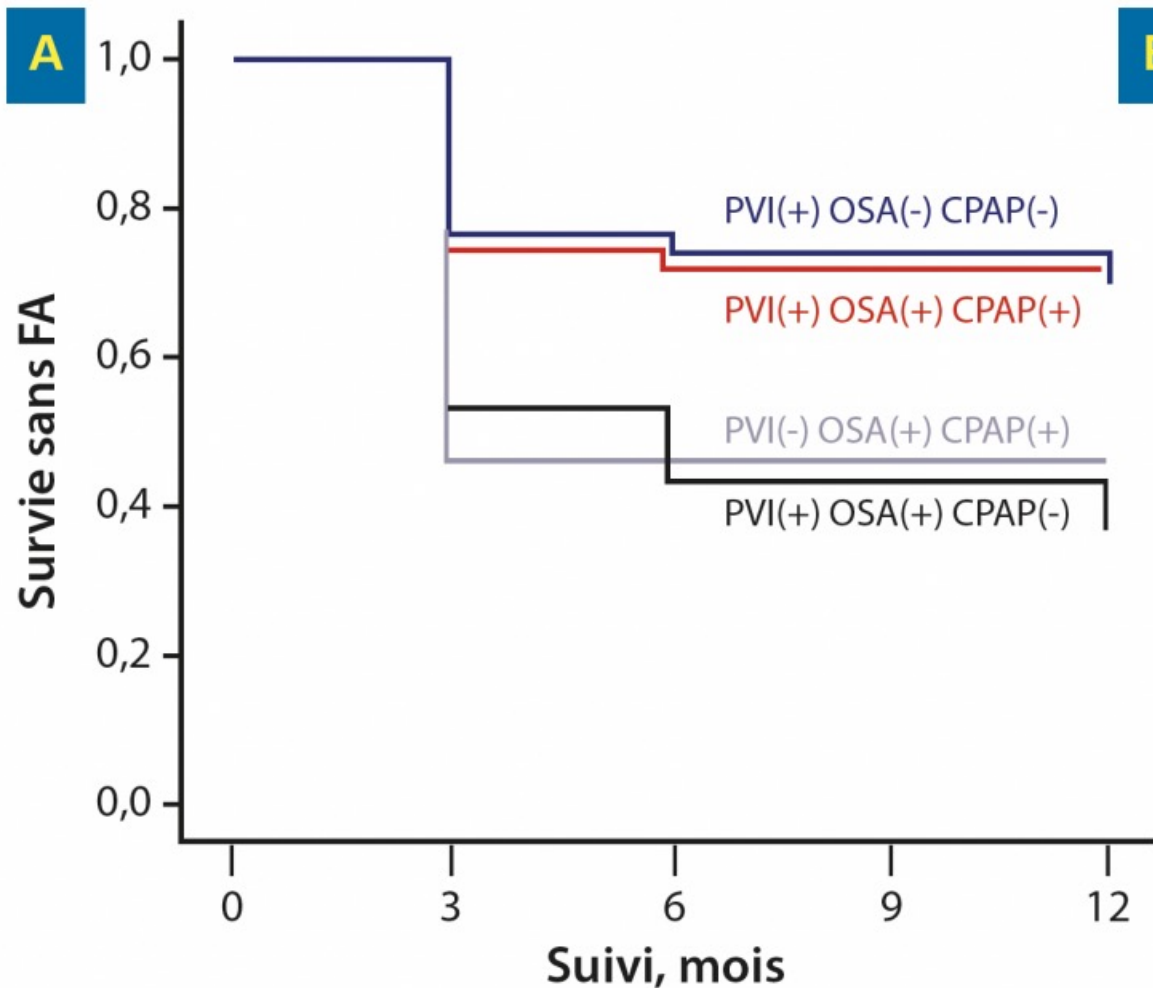
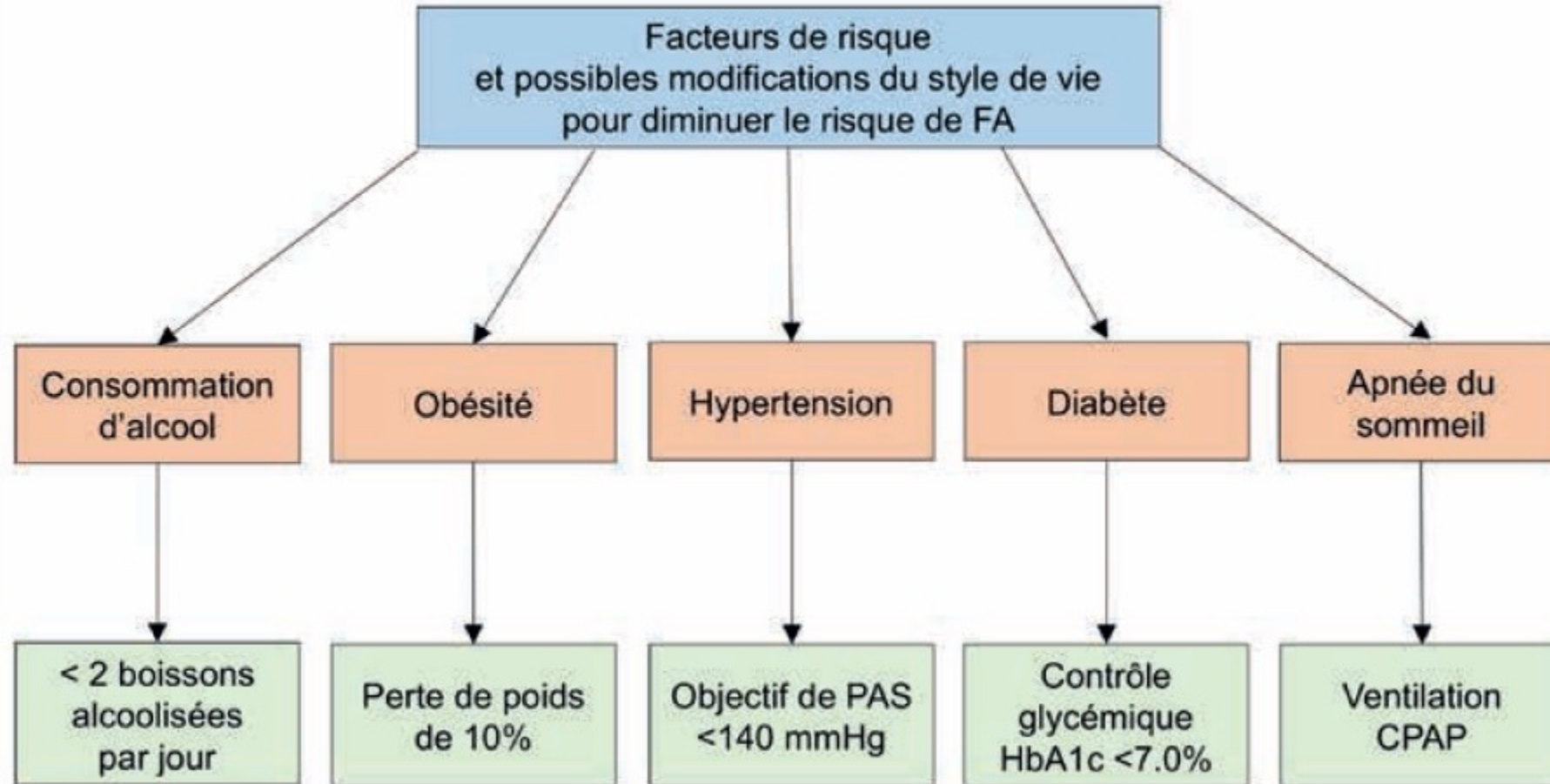


Figure 2. A. Courbe de survie en fonction du type de traitement. D'après Fein *et al.* 2013⁽⁷⁾. **B.** L'ablation de fibrillation atriale associée au traitement du SAOS permet une optimisation de la prise en charge du patient.

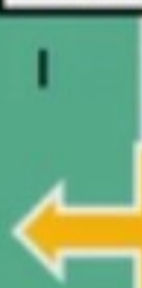
Figure 1. Prise en charge des facteurs modifiables du risque de fibrillation atriale (FA).



Recommendations for rhythm control/catheter ablation of AF (2)

Recommendations	Class	Level
<i>AF catheter ablation after failure of drug therapy</i>		
AF catheter ablation for PVI is recommended for rhythm control after one failed or intolerant class I or III AAD, to improve symptoms of AF recurrences in patients with	I	A
• Paroxysmal AF, or		
• Persistent AF without major risk factors for AF recurrence, or		
• Persistent AF with major risk factors for AF recurrence		Change IIa
AF catheter ablation for PVI should be considered for rhythm control after one failed or intolerant to beta-blocker treatment to improve symptoms of AF recurrences in patients with paroxysmal and persistent AF .	IIa	B

Results from
CAPTAF &
CABANA trials

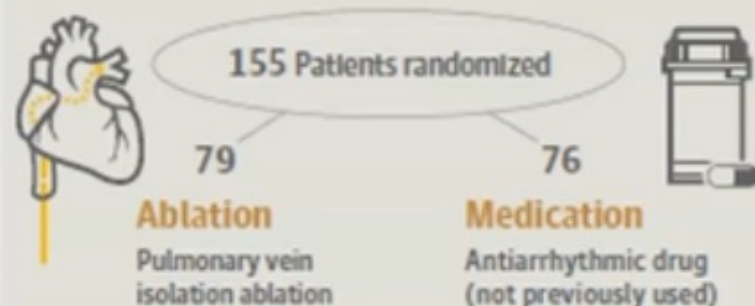


Effect of Catheter Ablation vs Antiarrhythmic Medication on Quality of Life in Patients With Atrial Fibrillation

The CAPTAF Randomized Clinical Trial

Carina Blomstrom-Lundqvist, MD, PhD; Sigfus Gaurason, MD, PhD; Jonas Schweder, MD, PhD; Steen M. Jensen, MD, PhD; Lennart Bergfeldt, MD, PhD; Göran Knebel, MD, PhD; Aggers Rubeck, MD, PhD; Helena Malmberg, MD, PhD; Pekka Rautanen, MD, PhD; Stefan Linnerholm, MD, PhD; Niklas Höglund, MD, PhD; David Mossell, MD

INTERVENTION



PRIMARY OUTCOME

Quality of life based on SF-36 General Health subscale score (range, 0 [worst] to 100 [best])



Blomstrom-Lundqvist, et al JAMA 2019

FINDINGS

Change in SF-36 General Health subscale score

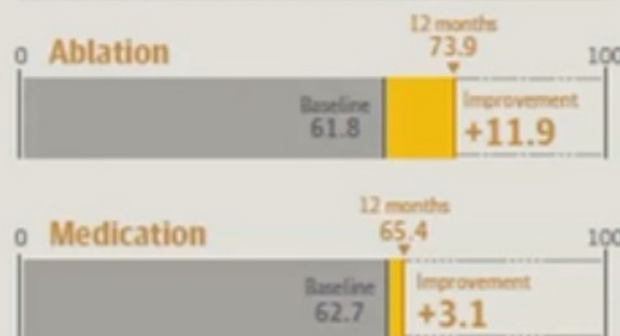
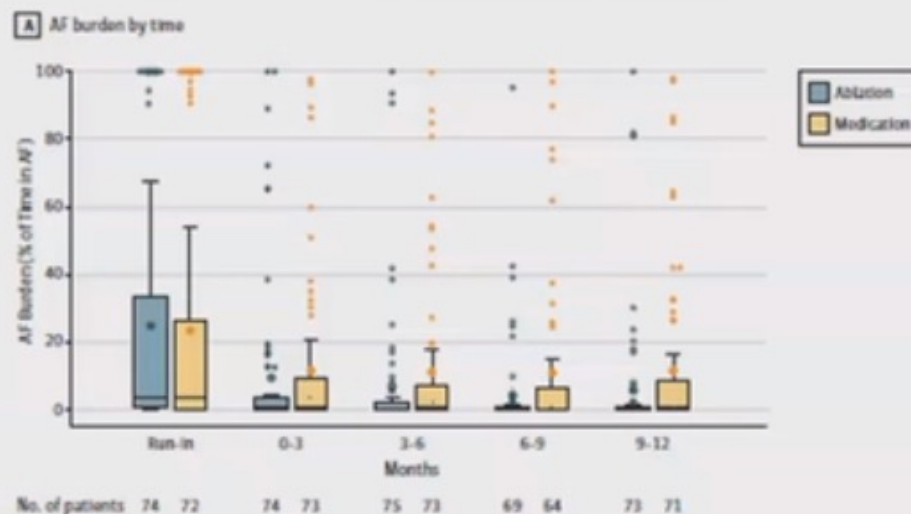


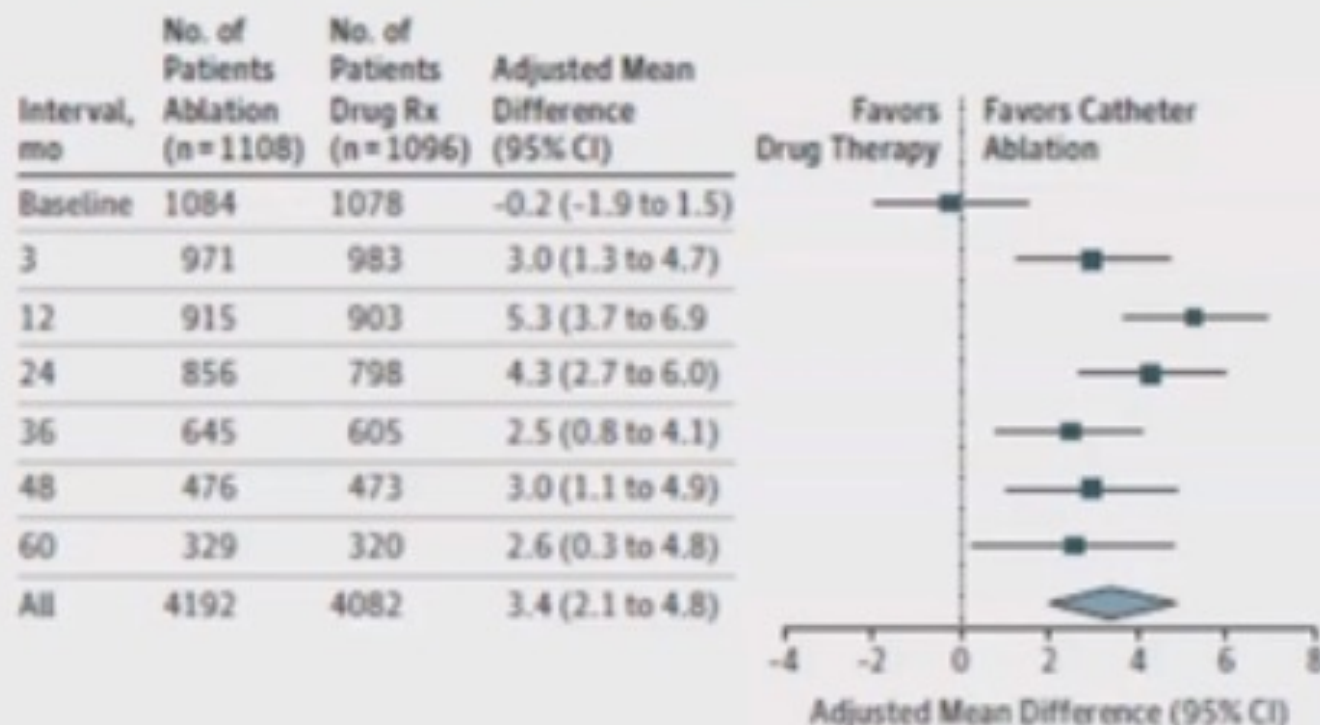
Figure 3. Secondary End Point Atrial Fibrillation Burden From the Implantable Cardiac Monitor



Improvement in QoL and reduction in AF burden - significantly greater in AF ablation vs AAD group

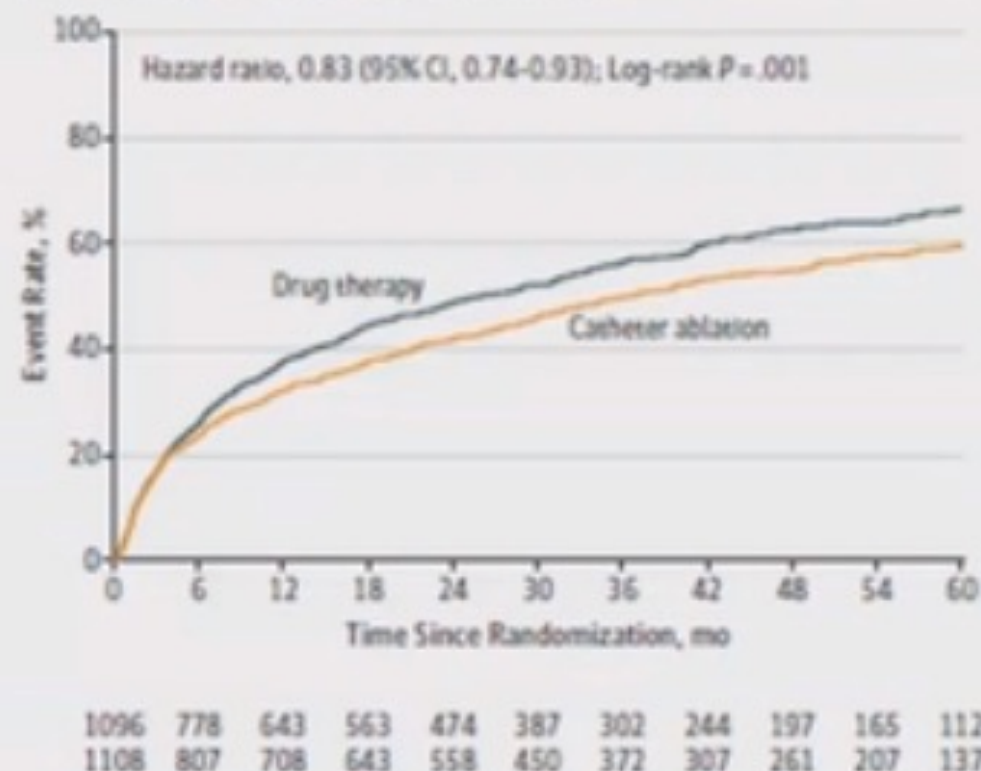
Effect of Catheter Ablation vs Medical Therapy on Quality of Life Among Patients With Atrial Fibrillation The CABANA Randomized Clinical Trial

B Between-group AFEQT summary score difference



Effect of Catheter Ablation vs Antiarrhythmic Drug Therapy on Mortality, Stroke, Bleeding, and Cardiac Arrest Among Patients With Atrial Fibrillation The CABANA Randomized Clinical Trial

B Mortality or cardiovascular hospitalization



Recommendations for rhythm control/catheter ablation of AF (3)

Recommendations	Class	Level
<i>First-line therapy</i>		
AF catheter ablation for PVI should/may be considered as first-line rhythm control therapy to improve symptoms in selected patients with symptomatic:		
• Paroxysmal AF episodes, or	IIa	B
• Persistent AF without major risk factors for AF recurrence.	IIb	C
as an alternative to AAD class I or III, considering patient choice, benefit, and risk.		

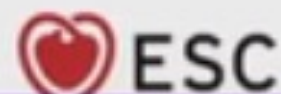
New !

Recommendations for rhythm control/catheter ablation of AF (4)

Recommendations	Class	Level
<i>First-line therapy (continued)</i>		
AF catheter ablation:		
• Is recommended to reverse LV dysfunction in AF patients when tachycardia-induced cardiomyopathy is highly probable, independent of their symptom status.	I	
• Should be considered in selected AF patients with HF with reduced LVEF to improve survival and reduce HF hospitalization .	IIa	B
AF catheter ablation for PVI should be considered as a strategy to avoid pacemaker implantation in patients with AF-related bradycardia or symptomatic pre-automaticity pause after AF conversion considering the clinical situation.	IIa	C

Change
IIa

Recommendations for rhythm control/catheter ablation of AF (4)



Recommendations

First-line therapy

AF catheter ablation

3 RCTs on AF ablation vs AAD in HF

- DiBase, AATAC, Circulation 2016
- Marrouche, CASTLE-AF, NEJM 2018
- Kuck, AMICA, Circ Arrh Electrophys 2019

1 substudy RCT on AF abl vs AAD

- Packer, CABANA subanalysis, HR 2020

Reduction in all-cause mortality and hospitalizations with ablation

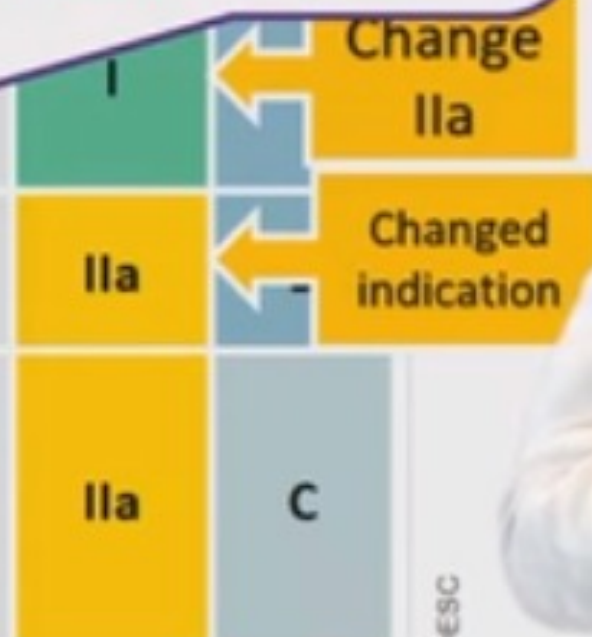
No difference in LVEF

Reduction in "death, stroke, bleeding, CA", and mortality with ablation in HF pats.

- Is recommended to **reverse or delay** in patients with **tachycardia-induced cardiomyopathy** is highly probable, independent of their symptom status.

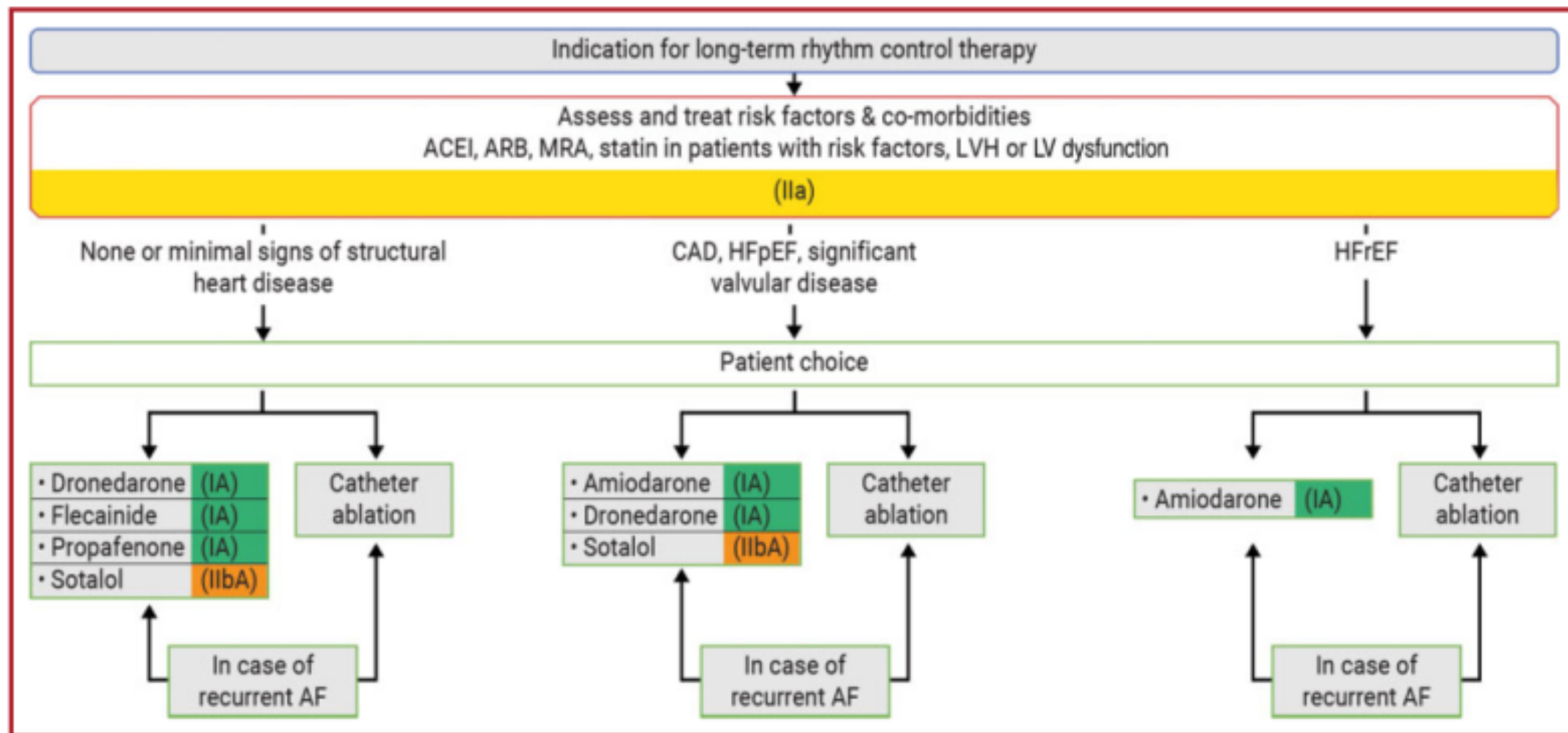
- Should be considered in **selected AF patients** with **HF with reduced LVEF** to **improve survival and reduce HF hospitalization**.

AF catheter ablation for **PVI** should be considered as a **strategy to avoid pacemaker** implantation in patients with **AF-related bradycardia** or **symptomatic pre-automaticity pause after AF conversion** considering the clinical situation.



Recommendations	Class	Level
Women with symptomatic paroxysmal or persistent AF should be offered timely access to rhythm control therapies , including AF catheter ablation , when appropriate for medical reasons.	Ila	B

New !



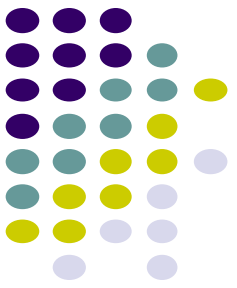
Vis ma vie de *Rythmologue*

Groupement Hospitalier de Territoire

Grand Paris Nord-Est

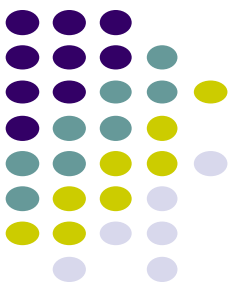
Aulnay-sous-Bois - Le Raincy-Montfermeil * Montreuil





Le jeune patient des urgences

- Homme
- 40 ans
- 75 Kg pour 1m79
- Pas de facteurs de risque
- Palpitations 2 heures (après un repas et 2 bières)
- FA 180 bpm aux urgences
- Réduction spontanée



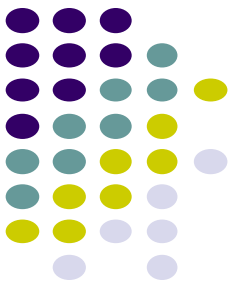
Que disent les recommandations?

11.2 Long-term antiarrhythmic drug therapy

The aim of antiarrhythmic drug therapy is improvement in AF-related symptoms.^{41,580} Hence, the decision to initiate long-term antiarrhythmic drug therapy needs to balance symptom burden, possible adverse drug reactions, and patient preferences. The principles of antiarrhythmic drug therapy outlined in the 2010 ESC AF guidelines³⁶⁹ are still relevant and should be observed:

- (1) Treatment is aimed at reducing AF-related symptoms;
- (2) Efficacy of antiarrhythmic drugs to maintain sinus rhythm is modest;
- (3) Clinically successful antiarrhythmic drug therapy may reduce rather than eliminate the recurrence of AF;
- (4) If one antiarrhythmic drug 'fails', a clinically acceptable response may be achieved with another agent;
- (5) Drug-induced pro-arrhythmia or extracardiac side-effects are frequent;
- (6) Safety rather than efficacy considerations should primarily guide the choice of antiarrhythmic drug.

Une première FA



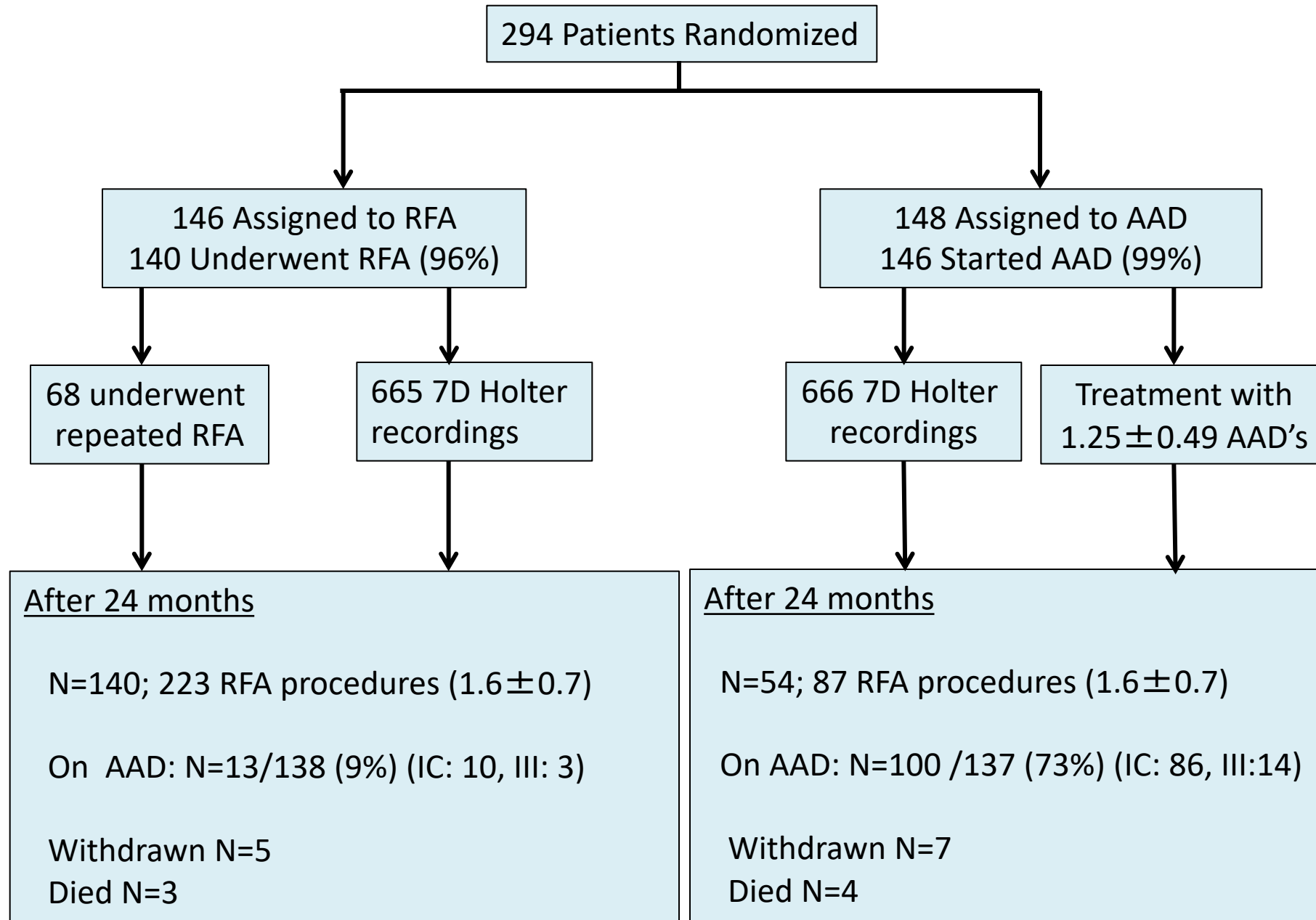
- Hôtesse de l'air
- 40 ans
- 58 kg/1m74
- Pas de FDR
- Adressée en USIC le 2.1.2014 par SAMU pour FA rapide 200 bpm responsable d'un OAP (VNI 1h)
- ETT FE et OG nles
- Régularisation après 2 jours par amiodarone
- Revue en cs sous amiodarone / AOD

Ablation de FA en première intention : résultats de MANTRA PAF

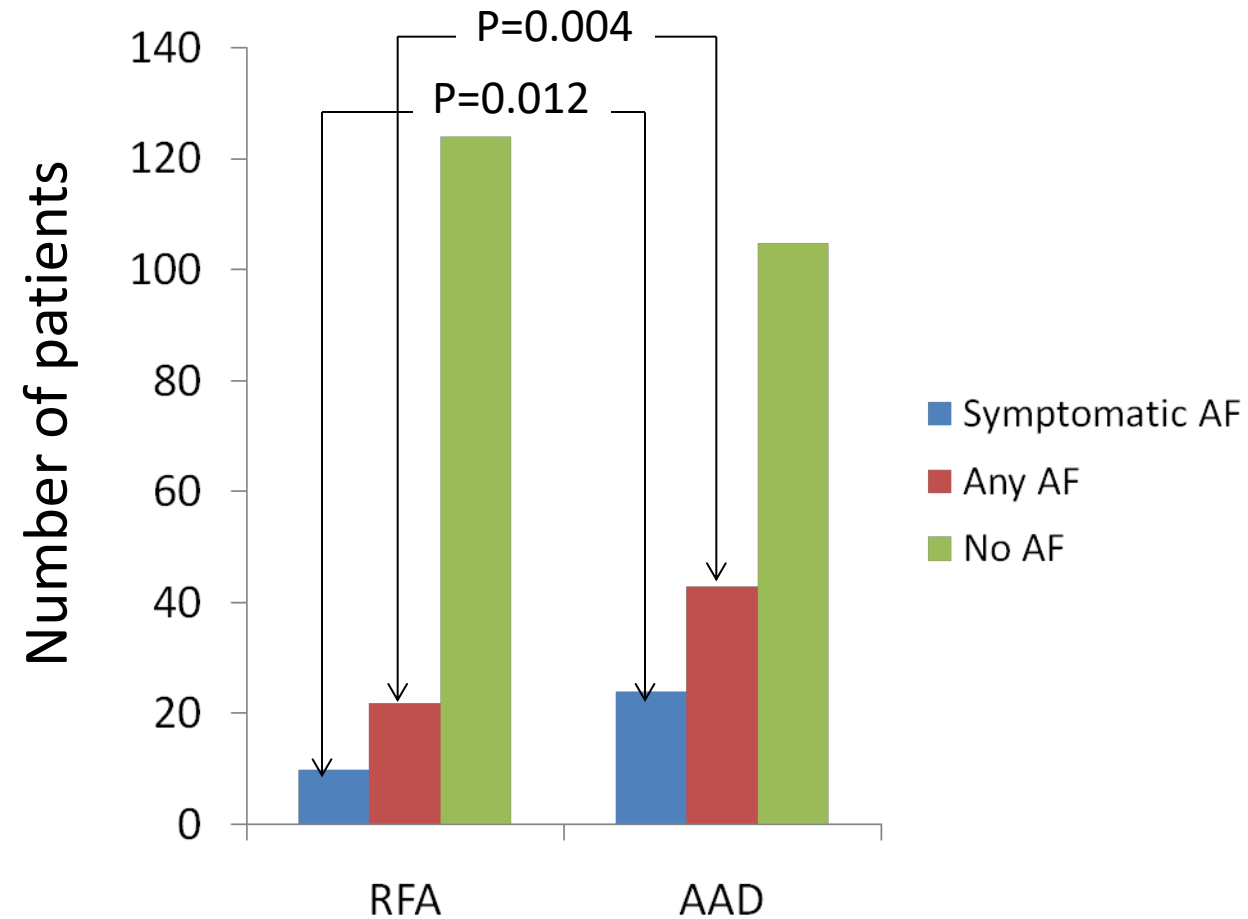
- Randomized controlled multicenter trial
- Antiarrhythmic drug therapy (Class IC or III) (AAD) *versus* pulmonary vein isolation (RFA)
- Primary endpoints:
 - Cumulative burden of AF
 - AF-burden at each 7-day Holter recording
- Secondary endpoints:
 - Freedom from any AF after 24 months
 - Freedom from symptomatic AF after 24 months
 - Burden of symptomatic AF after 3, 6, 12, 18, and 24 months
 - Atrial flutter
 - Quality-of life after 12 and 24 months
 - Serious adverse events

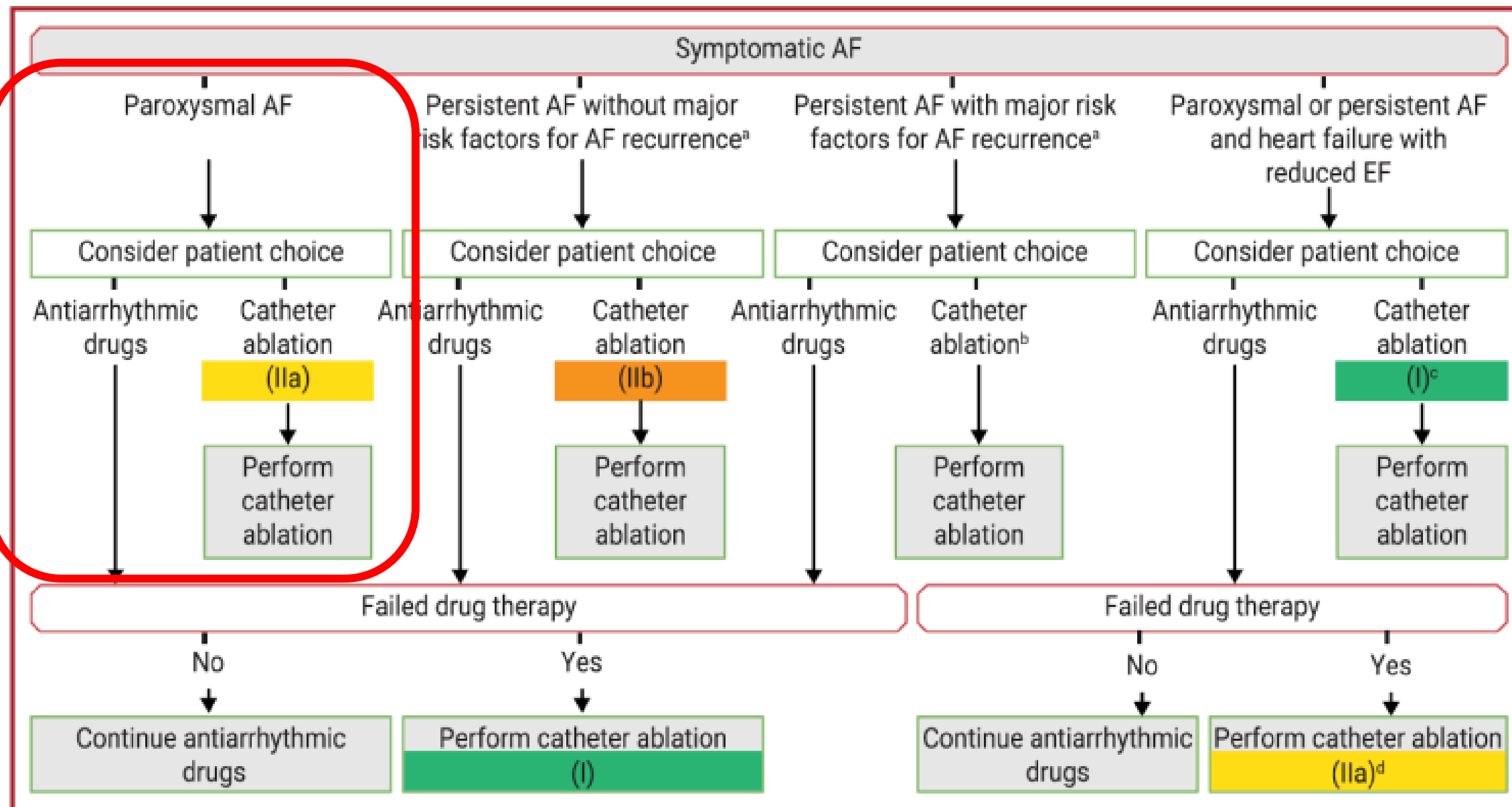
Nielsen JC, Johannessen A, Raatikainen P et al. Radiofrequency ablation as initial therapy in paroxysmal atrial fibrillation. New Engl J Med 2012; 367: 1587-95.





AF after 24 Months

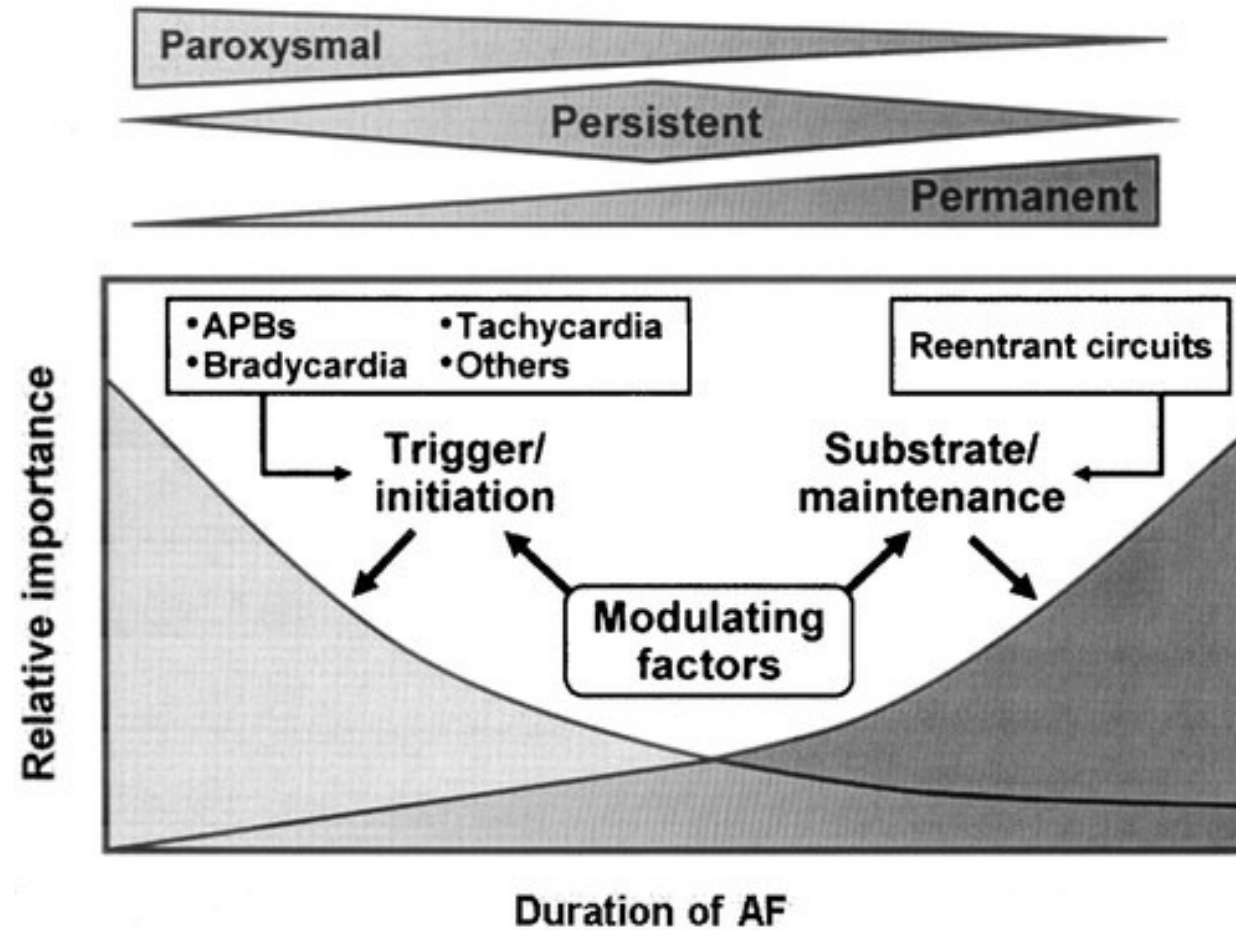




Catheter ablation of atrial fibrillation and atrial fibrillation surgery (1)

Recommendations	Class	Level
Catheter ablation of symptomatic paroxysmal AF is recommended to improve AF symptoms in patients who have symptomatic recurrences of AF on antiarrhythmic drug therapy (amiodarone, dronedarone, flecainide, propafenone, sotalol) and who prefer further rhythm control therapy, when performed by an electrophysiologist who has received appropriate training and is performing the procedure in an experienced centre.	I	A
Ablation of common atrial flutter should be considered to prevent recurrent flutter as part of an AF ablation procedure if flutter has been documented or occurs during the AF ablation.	IIa	B
Catheter ablation of AF should be considered as first-line therapy to prevent recurrent AF and to improve symptoms in selected patients with symptomatic paroxysmal AF as an alternative to antiarrhythmic drug therapy, considering patient choice, benefit, and risk.	IIa	B
All patients should receive oral anticoagulation for at least 8 weeks after catheter (IIaB) or surgical (IIaC) ablation.	IIa	B C
Anticoagulation for stroke prevention should be continued indefinitely after apparently successful catheter or surgical ablation of AF in patients at high-risk of stroke.	IIa	C
When catheter ablation of AF is planned, continuation of oral anticoagulation with a VKA (IIaB) or NOAC (IIaC) should be considered during the procedure, maintaining effective anticoagulation.	IIb	B C
Catheter ablation should target isolation of the pulmonary veins using radiofrequency ablation or cryotherapy balloon catheters.	IIa	B

Maintaining SR and reduction of AF Disease Progression



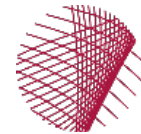
ORIGINAL ARTICLE

Early Rhythm-Control Therapy in Patients
with Atrial Fibrillation

P. Kirchhof, A.J. Camm, A. Goette, A. Brandes, L. Eckardt, A. Elvan, T. Fetsch,
I.C. van Gelder, D. Haase, L.M. Haegeli, F. Hamann, H. Heidbüchel,
G. Hindricks, J. Kautzner, K.-H. Kuck, L. Mont, G.A. Ng, J. Rekosz, N. Schoen,
U. Schotten, A. Suling, J. Taggeselle, S. Themistoclakis, E. Vettorazzi, P. Vardas,
K. Wegscheider, S. Willems, H.J.G.M. Crijns, and G. Breithardt, for the
EAST-AFNET 4 Trial Investigators*

p.kirchhof@bham.ac.uk / p.kirchhof@uke.de





Patients at risk for cardiovascular events ($\approx$ CHA₂DS₂VASc score ≥ 2)
and with recent onset atrial fibrillation ('**early AF**', ≤ 1 year duration or first documented by ECG)

Randomization

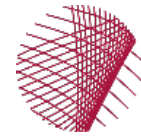
Early Rhythm Control

anticoagulation, rate control and
either antiarrhythmic drug therapy or AF ablation
In case of recurrent AF:
Re-ablation or adaptation of antiarrhythmic drugs

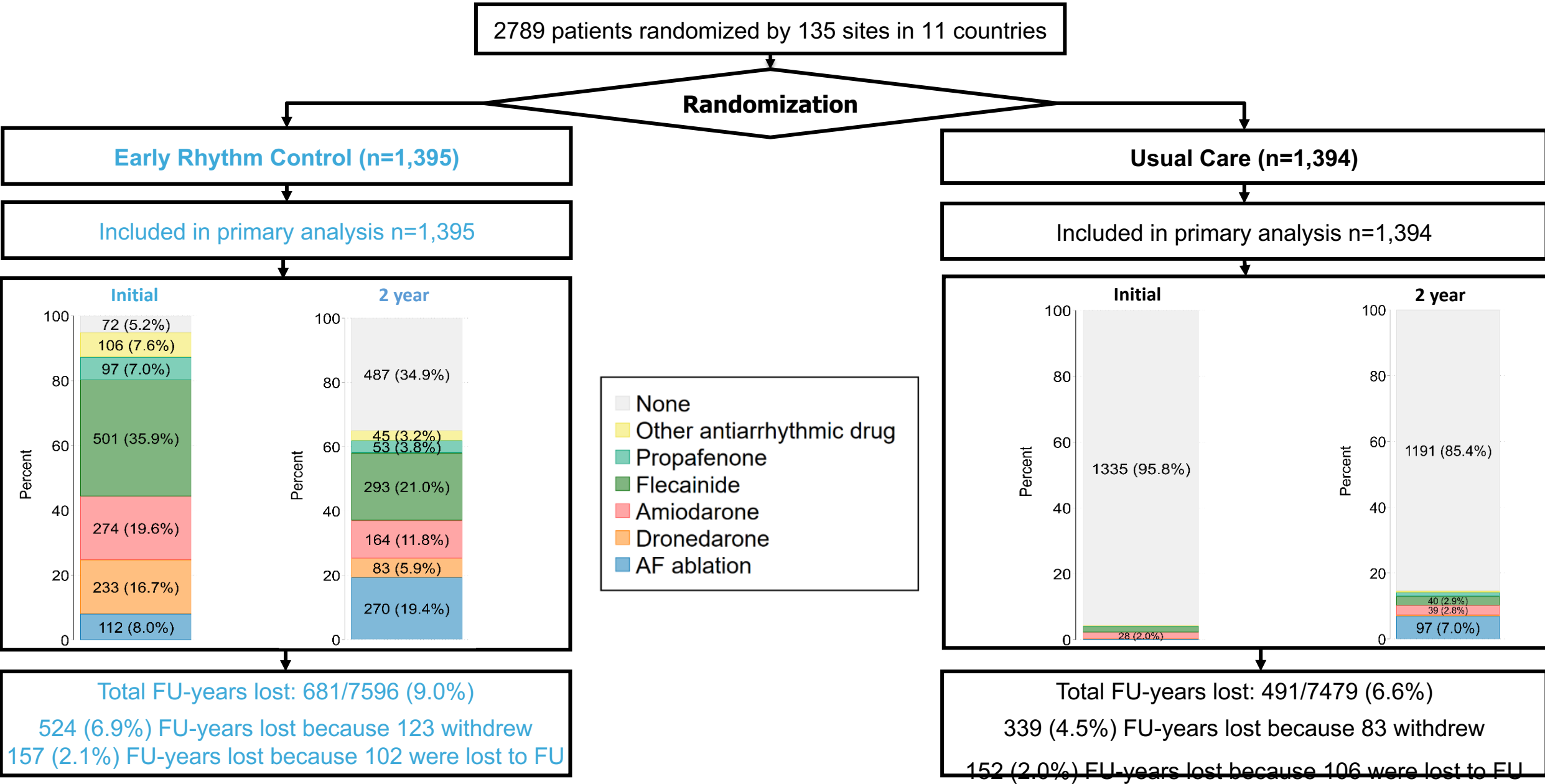
Usual Care

anticoagulation, rate control,
supplemented by rhythm control
only in symptomatic patients
on optimal rate control therapy

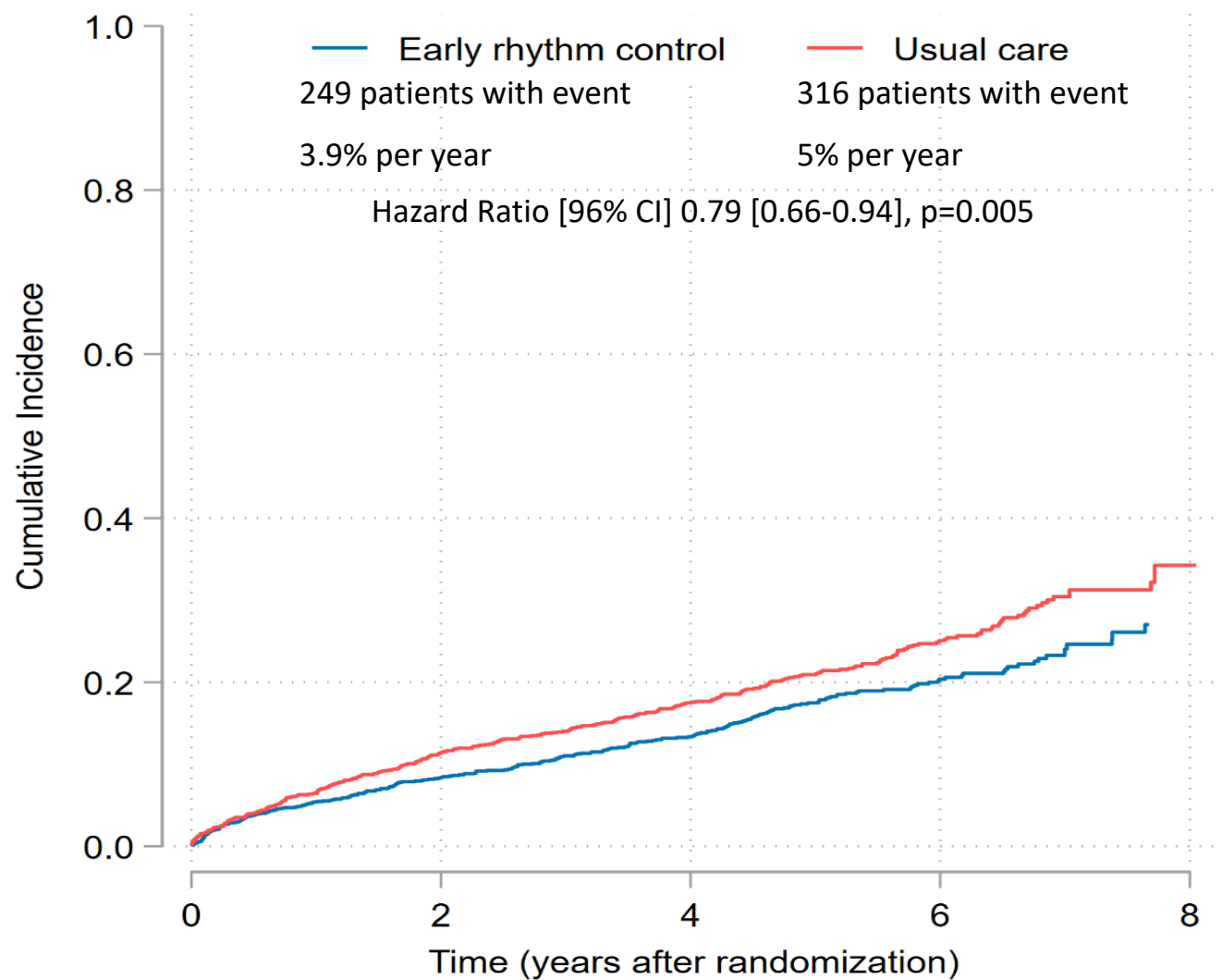
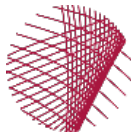
therapy of concomitant cardiovascular diseases (both randomized groups)
in-person follow-up at 1 and 2 years
all patients were followed up until the end of the study



EAST – AFNET 4 CONSORT diagram



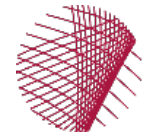
EAST – AFNET 4 Analysis of first primary outcome



	Patients with event in Early Rhythm Control (n=1395)	Patients with event in Usual Care (n=1394)	Uncorrected Hazard Ratio [95% CI]
Cardiovascular death	67	94	0.72 [0.52-0.98]
Stroke	40	62	0.65 [0.44-0.97]
Hospitalization with worsening of heart failure	139 (2.1)	169	0.81 [0.65-1.02]
Hospitalization with acute coronary syndrome	53 (0.8)	65	0.83 [0.58-1.19]

Patients at risk

Early rhythm control	1395	1193	913	404	26
Usual care	1394	1169	888	405	34



EAST – AFNET 4 Safety outcomes

	Early Rhythm Control (n=1395)	Usual Care (n=1394)
Occurrence of a primary safety outcome	231 (16.6%)	223 (16.0%)
Occurrence of stroke	40 (2.9%)	62 (4.4%)
Occurrence of death	138 (9.9%)	164 (11.8%)
Occurrence of a serious adverse event of special interest related to rhythm control therapy (detailed listing of events given in lines below)	68 (4.9%)	19 (1.4%)
Serious adverse events related to antiarrhythmic drug therapy		
Non-fatal cardiac arrest	1 (0.1%)	1 (0.1%)
Drug toxicity of AF-related drug therapy	10 (0.7%)	3 (0.2%)
Drug-induced bradycardia	14 (1.0%)	5 (0.4%)
AV block	2 (0.1%)	0 (0.0%)
Torsade de Pointes tachycardia	1 (0.1%)	0 (0.0%)
Serious adverse events related to AF ablation		
Pericardial tamponade	3 (0.2%)	0 (0.0%)
Bleeding related to AF ablation, major	6 (0.4%)	0 (0.0%)
Bleeding related to AF ablation, non-major	1 (0.1%)	2 (0.1%)
Other serious adverse events of special interest related to rhythm control therapy		
Blood pressure related (hypotension, hypertension; except syncope)	1 (0.1%)	0 (0.0%)
Hospitalization for AF	11 (0.8%)	3 (0.2%)
Other cardiovascular event	5 (0.4%)	1 (0.1%)
Other event	1 (0.1%)	3 (0.2%)
Syncope	4 (0.3%)	1 (0.1%)
Hospitalization for worsening of heart failure with decompensated heart failure	3 (0.2%)	0 (0.0%)
Implantation of a pacemaker, defibrillator, cardiac resynchronization device, or other device	8 (0.6%)	4 (0.3%)

Embargoed for 9:34am CT 11-16-20

Early Intervention for Atrial Fibrillation: The EARLY-AF Study

Jason G. Andrade, Jean Champagne, Marc W. Deyell, Vidal Essebag,
Sandra Lauck, Carlos A. Morillo, John L. Sapp, Allan Skanes,
Patricia Theoret-Patrick, George Wells, Atul Verma



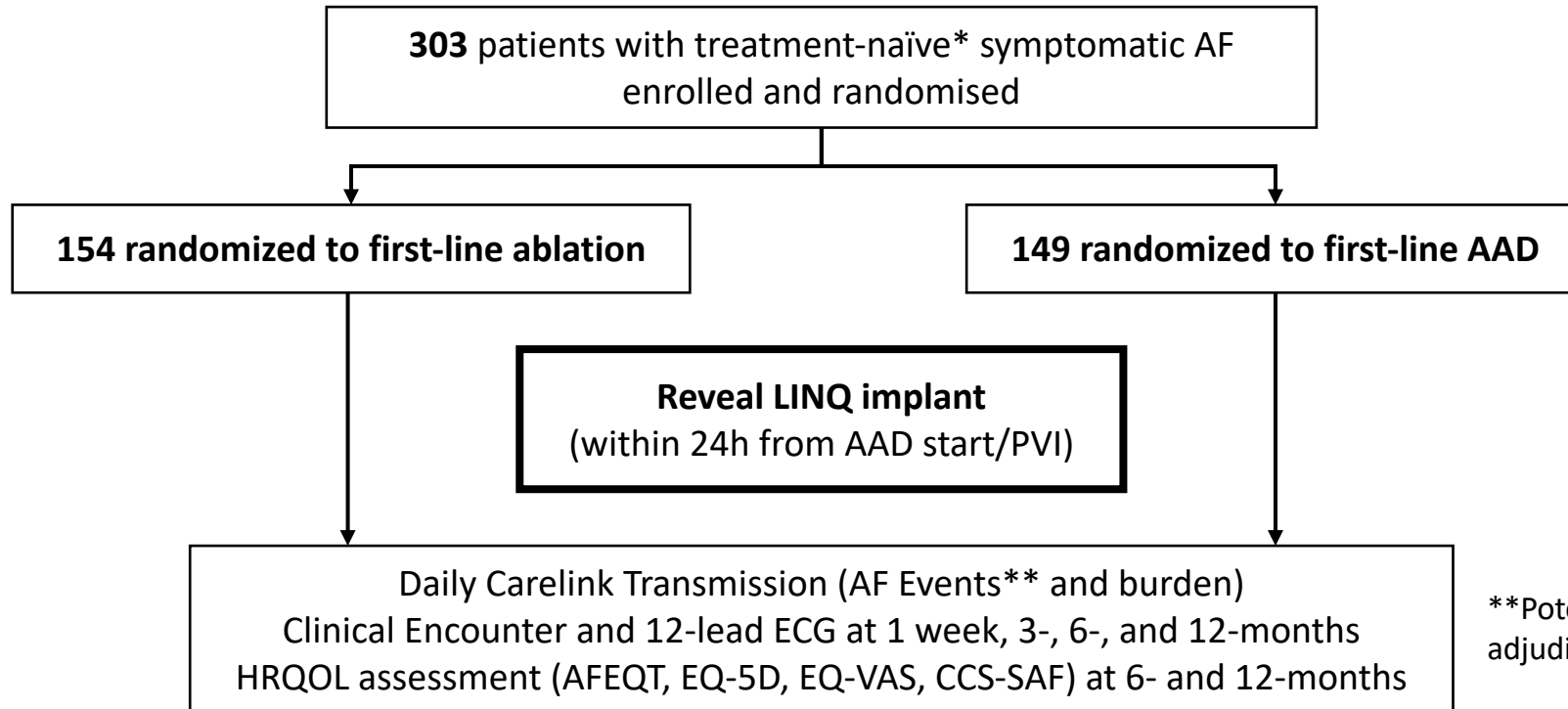
#AHA20



American Heart Association®
Scientific Sessions



Patient flow



*Enrollment Permitted if:

1. AAD Treatment Naïve

- Never treated with an AAD

2. Current AAD use

- Treatment < 6m but **below** therapeutic threshold

3. Previous AAD use

- Treatment initiated, discontinued, and washed out >6m

4. Temporary AAD use

- Treatment at therapeutic dose for a period <4 weeks

No adverse drug effects or inefficacy

**Potential arrhythmia events detected by the device were stored for adjudication by an independent, blinded clinical end-point committee.



Treatment Groups

Antiarrhythmic Drug Group

Used	First N (%)	Second N (%)	Third N (%)	Any time N (%)	Median dose in mg/day
Flecainide	114 (76.5%)	10 (6.7%)	0	124 (83.2%)	200 (125, 250)
Propafenone	7 (4.7%)	9 (6.0%)	2 (1.3%)	18 (12.1%)	600 (450, 600)
Sotalol	23 (15.4%)	17 (11.4%)	2 (1.3%)	42 (28.2%)	160 (160, 240)
Dronedarone	5 (3.4%)	7 (4.7%)	0	12 (8.1%)	800 (800, 800)
Amiodarone	0	3 (2.0%)	4 (2.7%)	7 (4.7%)	200 (200, 200)
Total	149 (100%)	46 (30.9%)	8 (5.4%)		

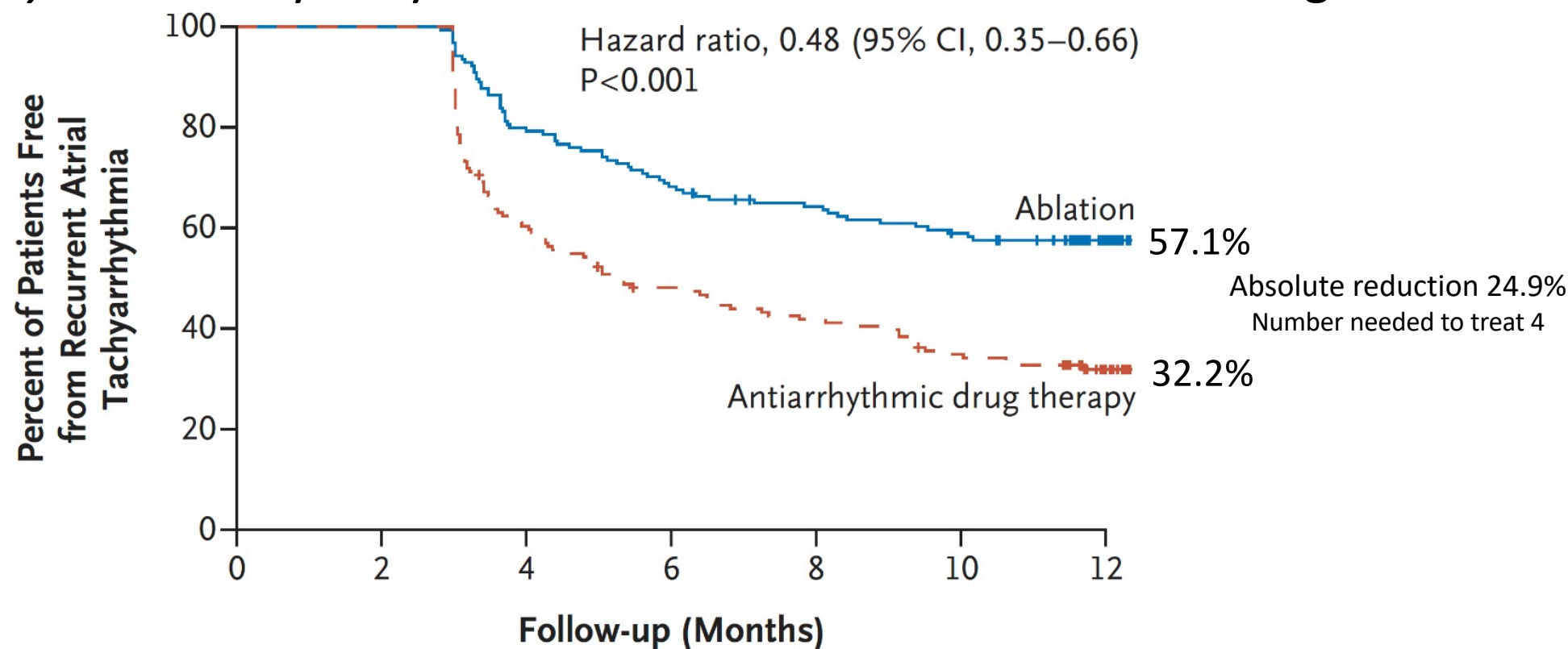
Cryoballoon Ablation Procedure

Procedural Characteristic	No. of Patients (%) n=152
Procedure Duration, in minutes	106 (89,131)
Fluoroscopy exposure, in minutes	18.9 (12.6,27.0)
Left atrial time, in minutes	73.7 (56.3,94.4)
Ablation duration, in minutes	26.1 (23.1, 30.1)
23-mm Cryoballoon, N (%)	5 (3.2%)
Acute pulmonary reconnection, N (%)	6 (3.9%)



Primary Outcome

Freedom from *any* atrial tachyarrhythmia on continuous cardiac monitoring



No. at Risk

Ablation	154	154	123	105	96	86	55
Antiarrhythmic drug therapy	149	149	89	69	60	49	27

Adverse Events

Serious Adverse Events - HR 0.81 (0.25–2.59)

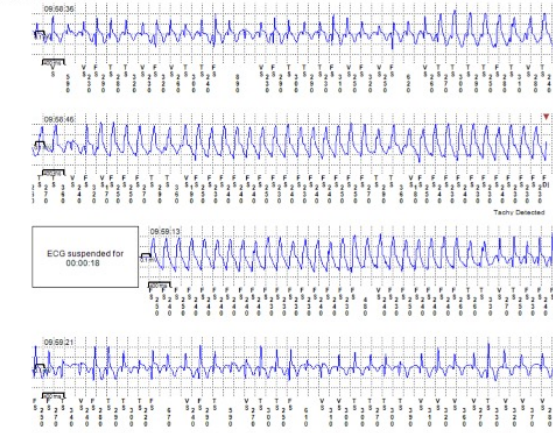
- **Ablation – 5 (3.2%)**
 - 3 phrenic nerve injuries
 - 2 pacemakers for bradycardia
- **Antiarrhythmic drugs – 6 (4.0%)**
 - 2 wide-complex tachycardia
 - 2 pacemakers for bradycardia
 - 1 heart failure
 - 1 syncope

Any Safety Endpoint - HR 0.59 (0.29–1.21)

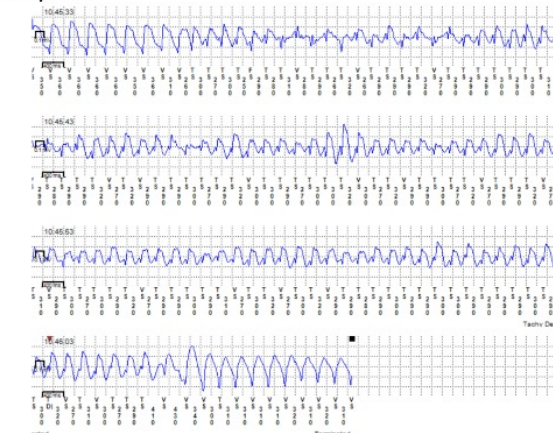
- **Ablation – 14 (9.1%)**
- **Antiarrhythmics – 24 (16.1%)**

Event	Ablation Group (N = 154)	Antiarrhythmic Drug Group (N = 149)
Any serious adverse event related to the trial regimen — no. of patients (%) ^a	5 (3.2)	6 (4.0)
Any safety end point — no.		
Patients	14	24
Events	15	27
Death — no.	0	0
Cardiac event — no.		
Pericardial effusion for which drainage was warranted or tamponade	0	1†
Pericardial effusion for which drainage was not warranted	0	0
Pericarditis	0	0
Exacerbation of heart failure	0	1
Syncope	1	2
Wide-complex tachycardia or proarrhythmic event	0	2
Bradycardia or atrioventricular block for which pacemaker insertion was warranted	2	2
Acute coronary syndrome	0	2
Neurologic event — no.		
Stroke	0	0
Transient ischemic attack	0	1
Vascular event — no.		
Arteriovenous fistula	0	0
Hematoma for which intervention was warranted	0	0
Hematoma for which intervention was not warranted	1	0
Pseudoaneurysm for which intervention was warranted	0	0
Deep-vein thrombosis	1	0
Pulmonary event — no.		
Persistent phrenic nerve palsy	3‡	0
Pneumonia	1	0
Self-limited hemoptysis	1	1
Gastrointestinal event — no.		
Esophageal injury or perforation	0	0
Gastrointestinal upset such as indigestion or diarrhea	2	1
Adverse drug reaction leading to dose modification or discontinuation — no.		
Prolongation of QT interval	0	1
Presyncope	0	5
Tremor	0	1
Visual disturbance	0	1
Mild cognitive impairment	0	1
Insomnia	0	1
Other event — no.		
Erectile dysfunction	0	1
Rash	0	1
Epistaxis	2	0
Joint pain	0	2
Migraine	1	0

Wide-Complex Tachycardia 1



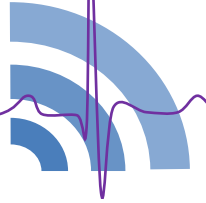
Wide-Complex Tachycardia 2





Conclusions

- First-line ablation was associated with significant reductions in arrhythmia outcomes:
 - Time to first recurrence of **any** AF/AFL/AT
 - Time to first recurrence of **any** AF
 - Time to first recurrence of **symptomatic** AF/AFL/AT
 - Time to first recurrence of **symptomatic** AF
 - Total AF burden
 - Days with AF
- } Continuous cardiac monitoring with implantable loop recorders
- First-line ablation was associated with meaningful improvements in quality of life and symptoms
 - Adverse events were similar between contemporary cryoballoon ablation and AAD therapy



Thank you!



The NEW ENGLAND
JOURNAL of MEDICINE

ORIGINAL ARTICLE

Cryoablation or Drug Therapy for Initial Treatment of Atrial Fibrillation

Jason G. Andrade, M.D., George A. Wells, Ph.D., Marc W. Deyell, M.D.,
Matthew Bennett, M.D., Vidal Essebag, M.D., Ph.D., Jean Champagne, M.D.,
Jean-Francois Roux, M.D., Derek Yung, M.D., Allan Skanes, M.D.,
Yaariv Khaykin, M.D., Carlos Morillo, M.D., Umjeet Jolly, M.D., Paul Novak, M.D.,
Evan Lockwood, M.D., Guy Amit, M.D., Paul Angaran, M.D., John Sapp, M.D.,
Stephan Wardell, M.D., Sandra Lauck, Ph.D., Laurent Macle, M.D., and
Atul Verma, M.D., for the EARLY-AF Study Investigators*

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- Julia Cadrin-Tourigny, Chris Cheung, Jeff Healey, Corinne Pearce, Rose Pfeffer

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- Paul Dorian (Chair), Ben Glover, Lehana Thabane



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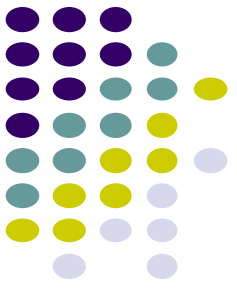


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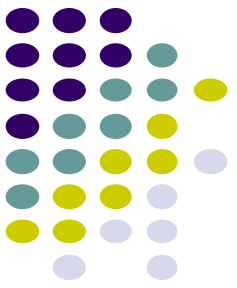
CANet.

Revue 6 mois après ablation FA



- Asymptomatique
- Pas de traitement
- ECG : RS
- Au sol et pas autorisée à voler...

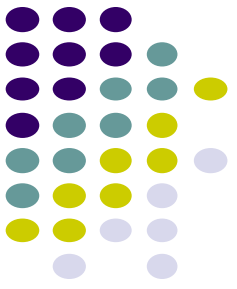
Cas



- Patient de 40 ans
- Diabète de type 2
- Obésité BMI 42 kg/m²
- HTA traitée par ARA2 – amlodipine
- Hospitalisé pour OAP sous VNI
- FA 180 bpm
- FE 25%

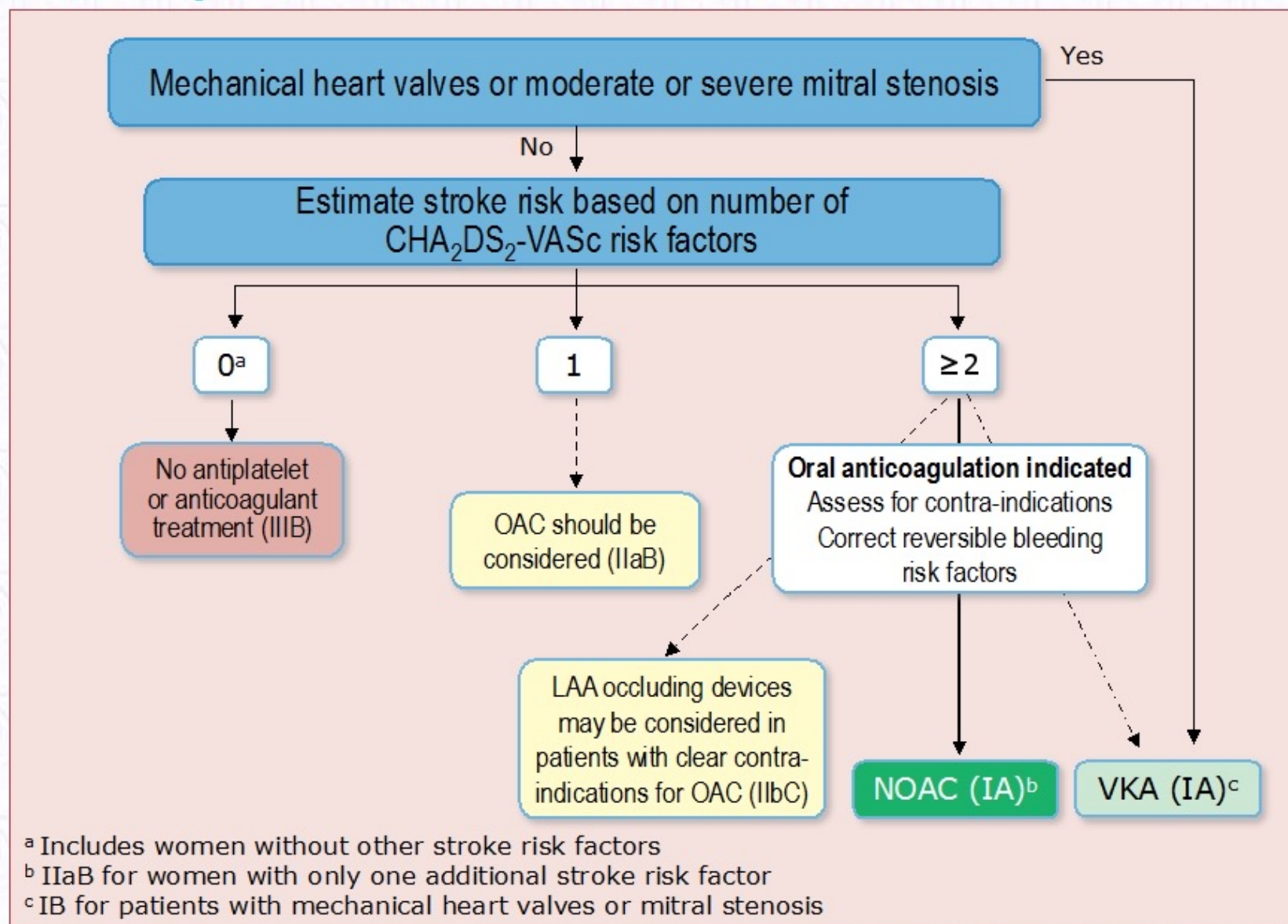
Suite

- Coro Nle
- Ralentissement FA
- FE 45% sous traitement



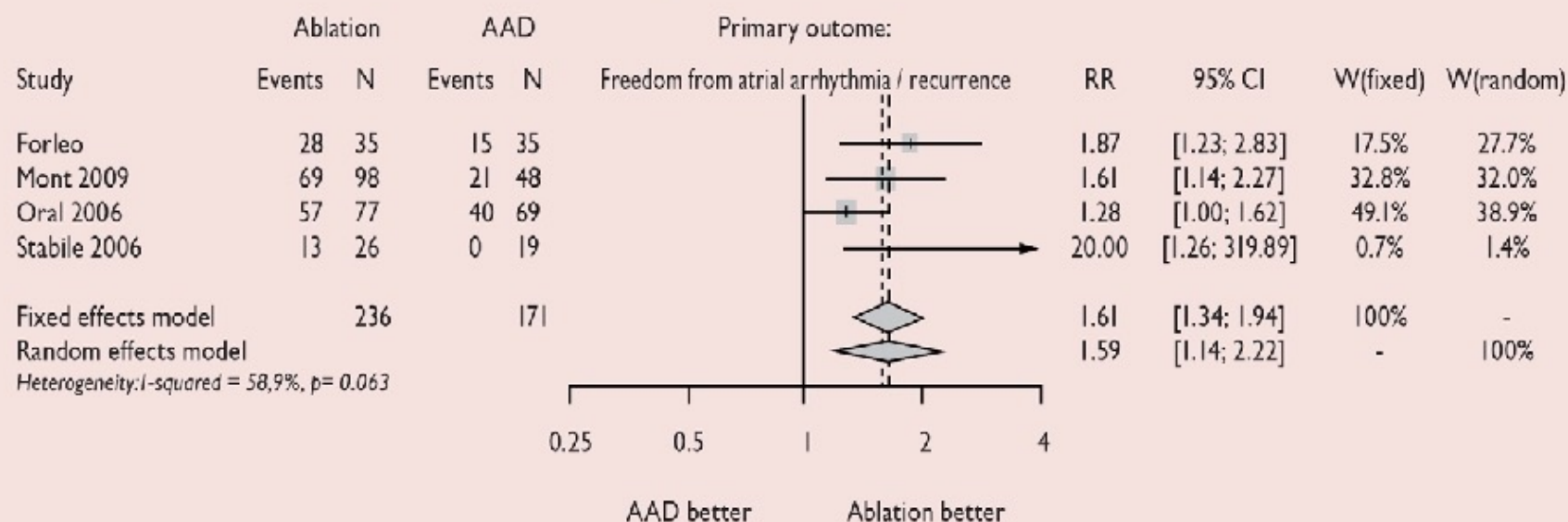


Stroke prevention in atrial fibrillation

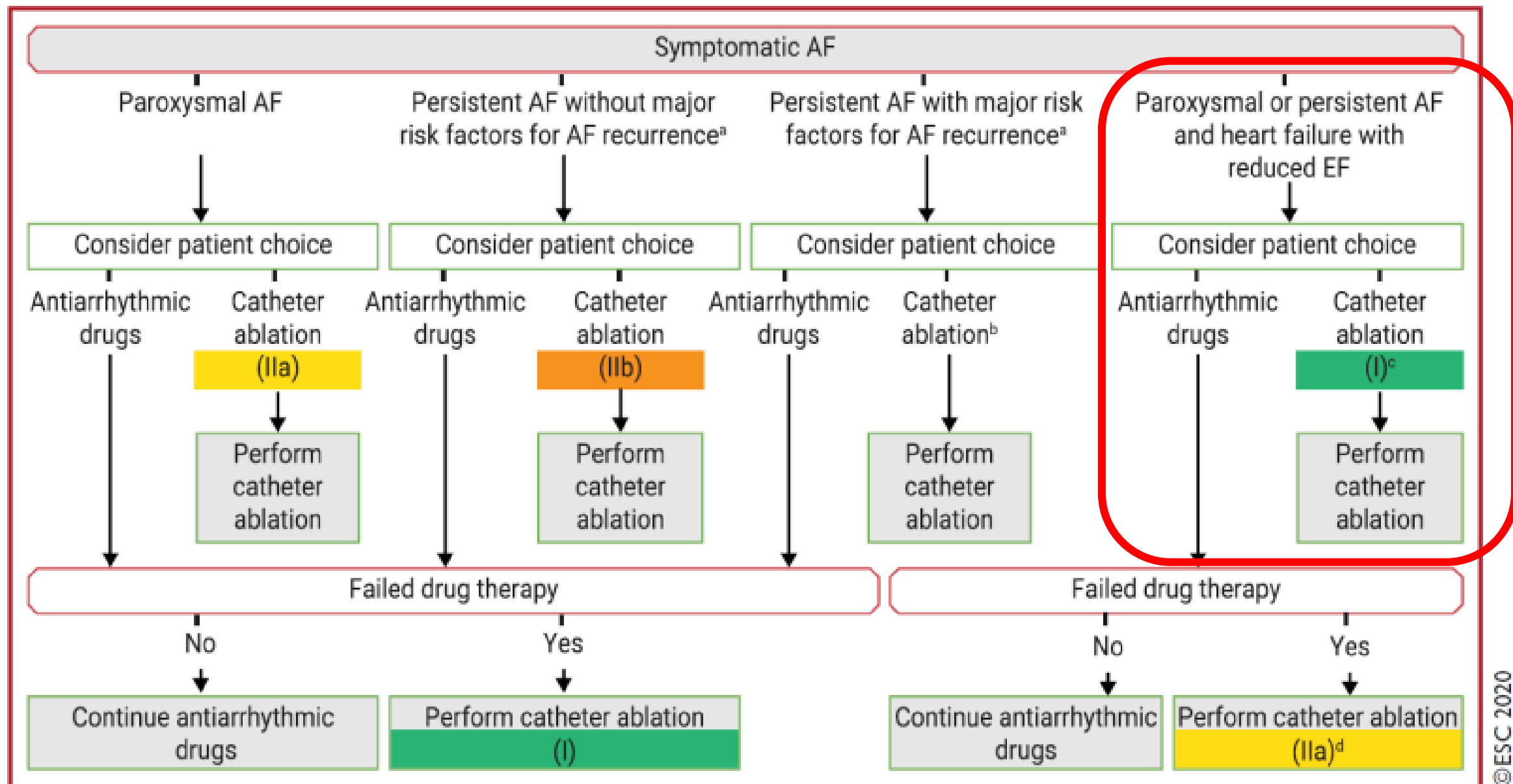


Rhythm outcome after catheter ablation compared to cardioversion and AAD in patients with persistent or long-standing persistent atrial fibrillation

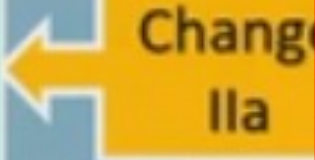
Freedom from recurrence of atrial fibrillation or atrial arrhythmias, comparing catheter ablation with antiarrhythmic drug therapy in patients with persistent or long-standing persistent atrial fibrillation



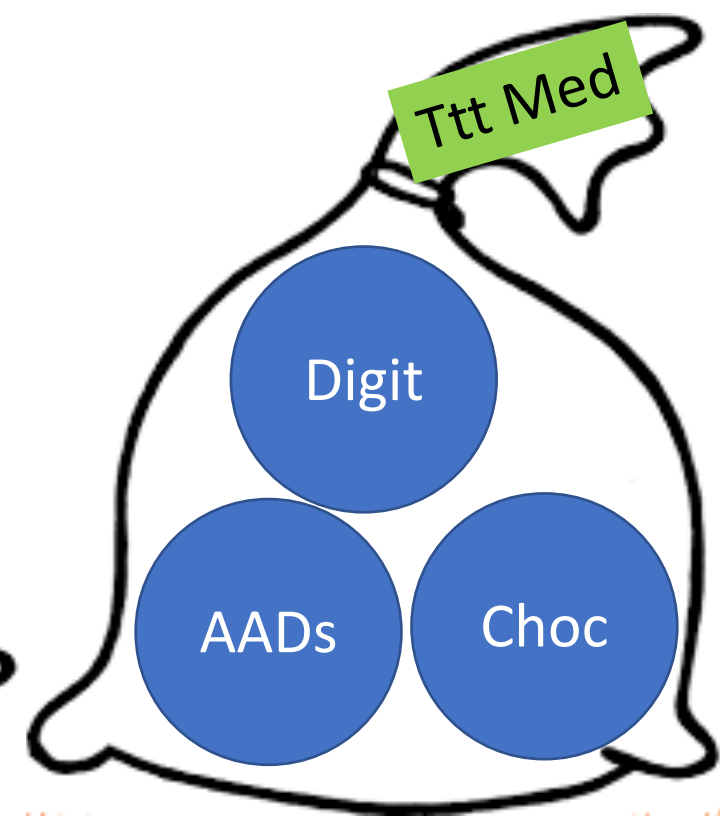
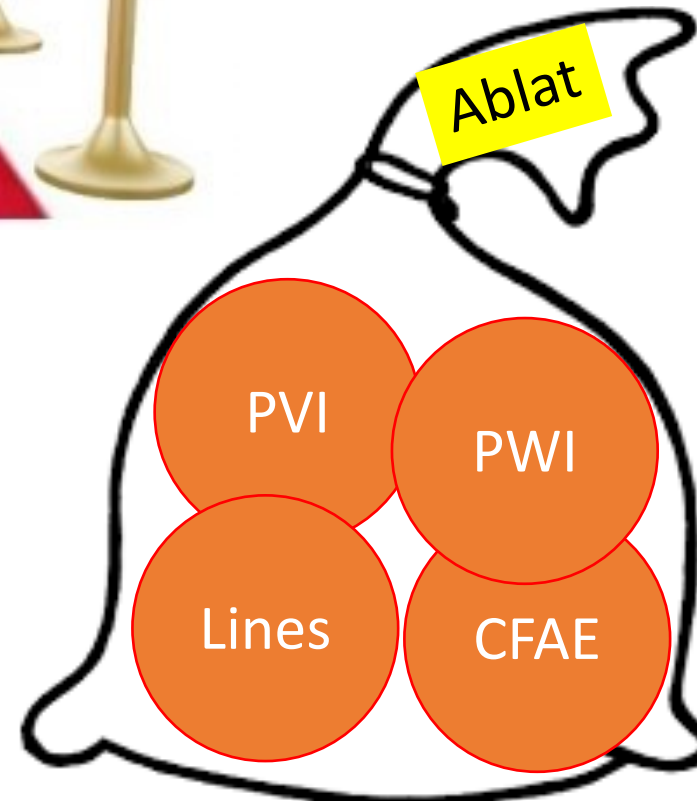
AAD = antiarrhythmic drug therapy; CI = confidence interval; N = number of patients; RR = risk ratio; W = study weighting.



Recommendations for rhythm control/catheter ablation of AF (4)

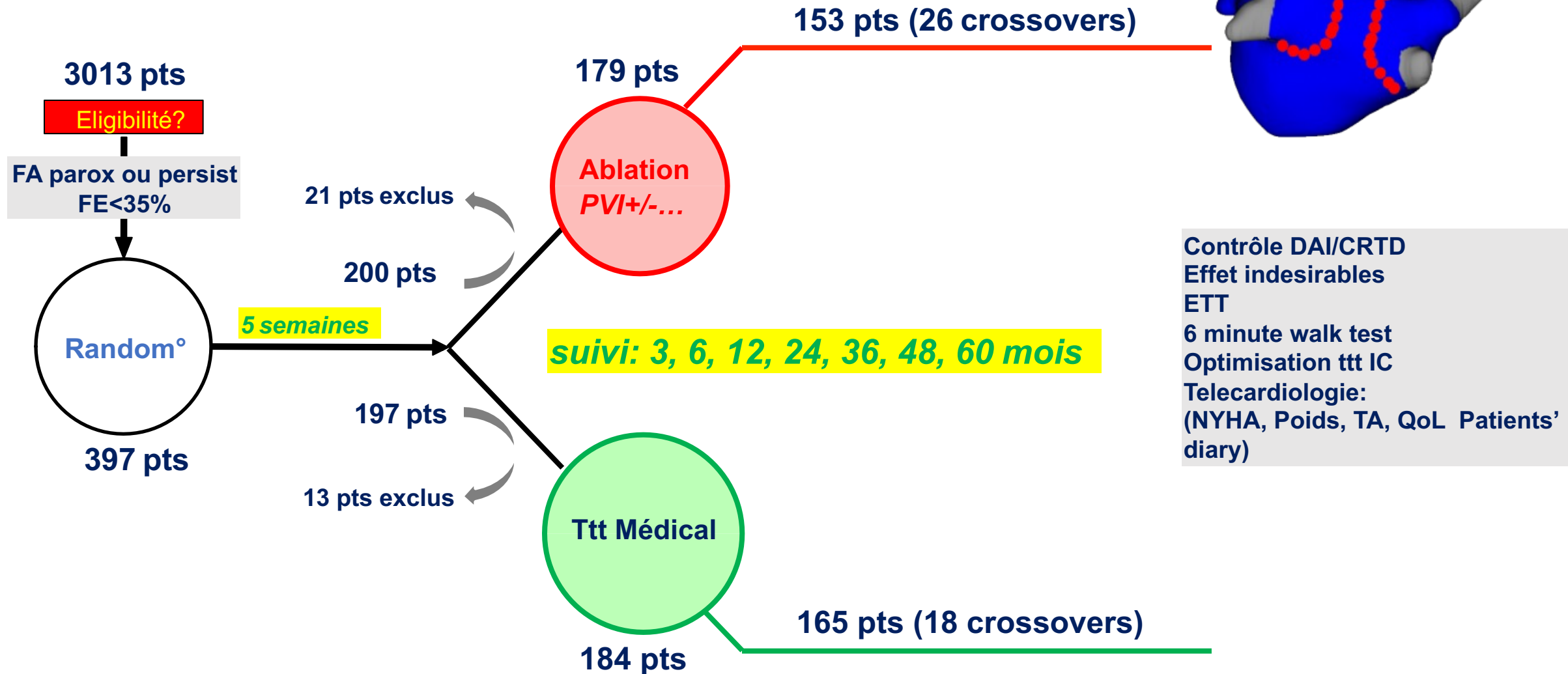
Recommendations	Class	Level
<i>First-line therapy (continued)</i>		
AF catheter ablation:		
• Is recommended to reverse LV dysfunction in AF patients when tachycardia-induced cardiomyopathy is highly probable, independent of their symptom status.	I	 Change IIa
• Should be considered in selected AF patients with HF with reduced LVEF to improve survival and reduce HF hospitalization .	IIa	B
AF catheter ablation for PVI should be considered as a strategy to avoid pacemaker implantation in patients with AF-related bradycardia or symptomatic pre-automaticity pause after AF conversion considering the clinical situation.	IIa	C

Et **ci** CASTLE-**AF** arriva....



Study Design— CASTLEAF

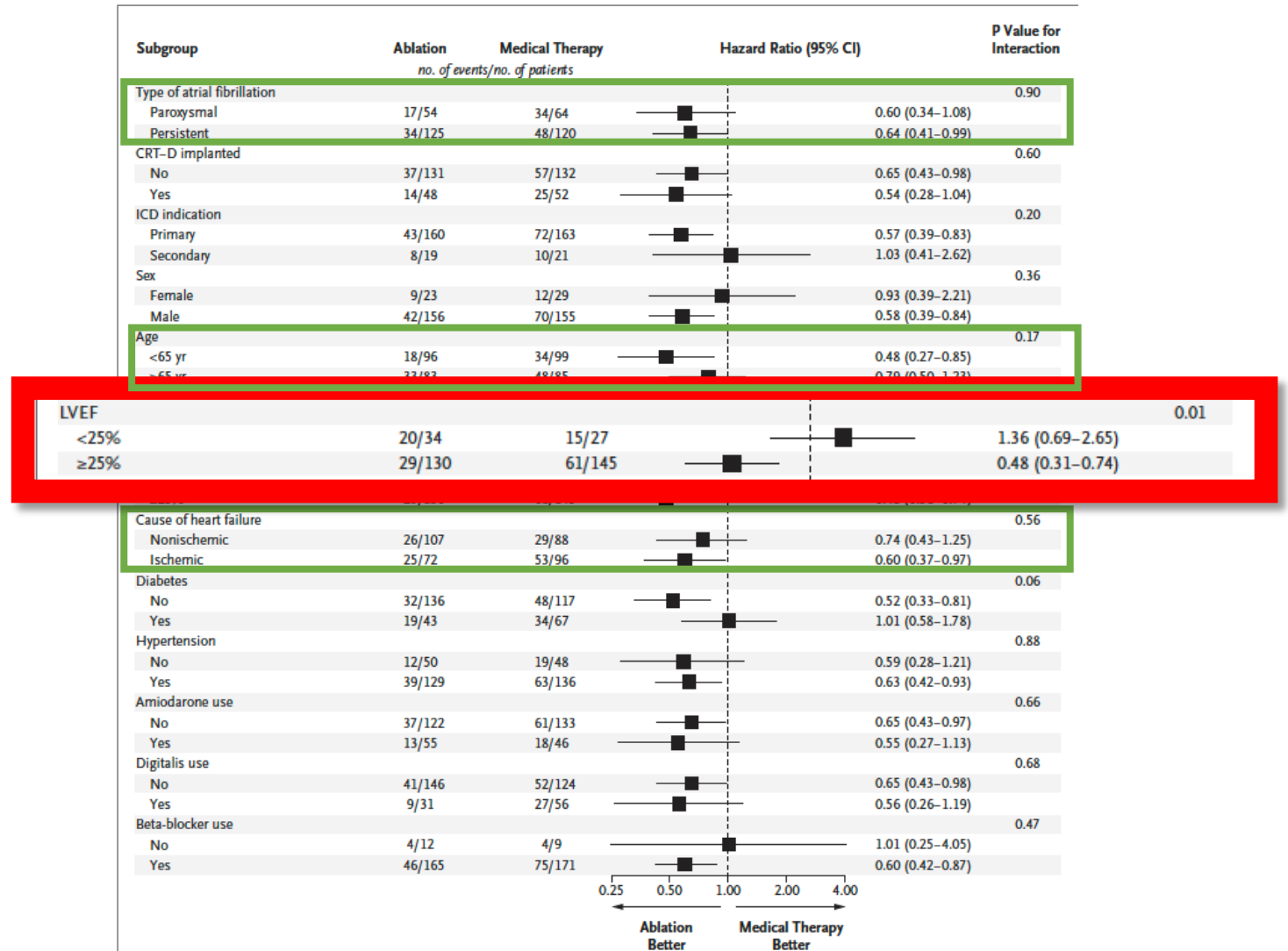
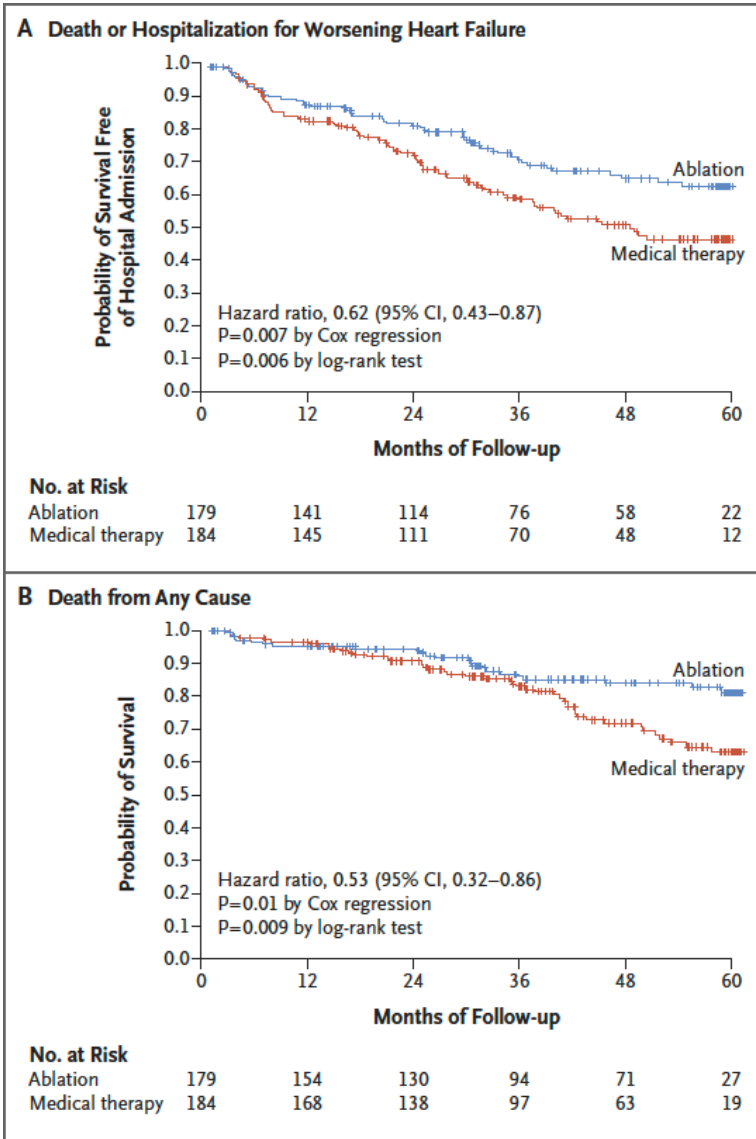
- Essai thérapeutique contrôlé randomisé (31 sites, 9 pays),



Patients CASTLE AF

	Ablation (179 patients)	Medical (184 patients)
Age – years	64 (5671)	64 (5673.5)
New York Heart Association class		
I (%)	11	11
II (%)	58	61
III (%)	29	27
IV (%)	2	1
Left ventricular ejection fraction – %	32.5 (25.038.0)	31.5 (27.037.0)
Left atrial diameter– mm	48	49.5
Current type of atrial fibrillation		
Paroxysmal (%)	30	35
Persistent (%)	70	65
Long standing Persistent (%)	28	29
CRTD implanted (%)	27	28

Castle AF Résultats



Safety

Table 5. Safety Outcomes for Patients With AF Undergoing Catheter Ablation

circae;2/4/349/TBL5824789T5

Outcomes	t	n/N	%
Mortality			
Death overall	65	42/5781	0.7
Procedure-related	64	0/5192	0.0
Vascular access complications			
Arteriovenous fistula	32	1/2885	0.0
Bleeding	33	1/2960	0.0
Hematoma	38	17/3719	0.5
Pneumothorax	34	0/2974	0.0
Femoral artery pseudoaneurysm	34	15/3032	0.5
Periprocedure events			
Stroke, ischemic	62	17/5665	0.3
TIA	60	13/5467	0.2
Cardiac tamponade	63	45/5723	0.8
PE	60	3/5496	0.1
DVT	56	1/4758	0.0
Other embolism	57	10/5347	0.2
LA-esophageal fistula	60	0/5496	0.0
Other fistula	58	3/5407	0.1
Pericardial effusion	64	36/5719	0.6
PV stenosis*	65	91/5831	1.6
AV block	60	1/5496	0.0
CHF exacerbation	60	0/5496	0.0
Need for a pacemaker	46	4/3902	0.1
Total No. of patients with events	28	97/1964	4.9

Safety

Table 6. Safety Outcomes for Patients With AF Receiving AAD Therapy

circae;2/4/349/TBL6824789T6

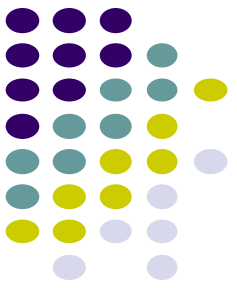
Safety Outcomes	Overall		
	t	n/N	%
Mortality			
Death overall	33	120/4291	2.8
Sudden death	21	18/2900	0.6
Treatment-related death	22	15/3179	0.5
Not treatment-related death	20	40/3023	1.3
Adverse events			
CV events	10	58/1572	3.7
Bradycardia	19	44/2349	1.9
GI	16	97/1499	6.5
Neuropathy	4	48/969	5.0
Thyroid dysfunction	5	19/576	3.3
Torsades	12	16/2238	0.7
Q-T* prolongation	12	5/2034	0.2
Total No. of patients with events	24	989/3318	29.8
Discontinuations			
Total	32	1035/4347	23.8
Due to AE	32	384/3682	10.4
Due to inefficacy	12	229/1694	13.5
Due to noncompliance	4	19/457	4.2

Table 5. Safety Outcomes for Patients With AF Undergoing Catheter Ablation

circae;2/4/349/TBL5824789T5

Outcomes	t	n/N	%
Mortality			
Death overall	65	42/5781	0.7
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Arteriovenous fistula	32	1/2885	0.0
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Hematoma	38	17/3719	0.5
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Cas

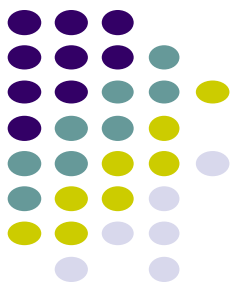


- Patient de 40 ans
- Diabète de type 2
- Obésité BMI 42 kg/m²
- HTA traitée par ARA2 – amlodipine
- Hospitalisé pour OAP sous VNI
- FA 180 bpm
- FE 25%

Après 2 ablations et une chirurgie de l'obésité:

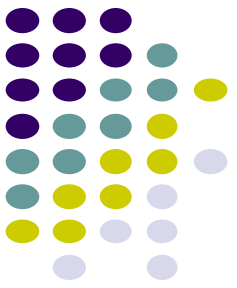
RS

Ce n'est pas qu'une histoire de rythme



In addition to antiarrhythmic drug therapy and catheter ablation (see Chapter 11.3), management of concomitant cardiovascular conditions can reduce symptom burden in AF and facilitate the maintenance of sinus rhythm.^{203,204,296,312} This includes weight reduction, blood pressure control, heart failure treatment, increasing cardiorespiratory fitness, and other measures (see Chapter 7).

Cas



- Patient de 40 ans
- Obésité BMI 40 kg/m²
- HTA traitée par ARA2 – amlodipine
- Hospitalisé pour OAP sous VNI
- FA 180 bpm
- FE 30%

Résultat après ablation ??

Weight reduction in patients with atrial fibrillation

Recommendations	Class	Level
In obese patients with AF, weight loss together with management of other risk factors should be considered to reduce AF burden and symptoms.	IIa	B

Management of respiratory diseases in patients with atrial fibrillation

Recommendations	Class	Level
Correction of hypoxaemia and acidosis should be considered as initial management for patients who develop AF during an acute pulmonary illness or exacerbation of chronic pulmonary disease.	IIa	C
Interrogation for clinical signs of obstructive sleep apnoea should be considered in all AF patients.	IIa	B
Obstructive sleep apnoea treatment should be optimized to reduce AF recurrences and improve AF treatment results.	IIa	B



A qui proposer l'ablation de FA ?



Quels patients ne pas proposer en ablation ?

- Asymptomatiques.
- Trop âgés (ou mauvais état général).
- Trop de comorbidités.
- Cardiopathie sous-jacente (Valvulopathie +++)
- OG trop dilatée
- Fibrose OG
- FA persistante « très » prolongée



Ablation de FA

Comment je fais en pratique ?

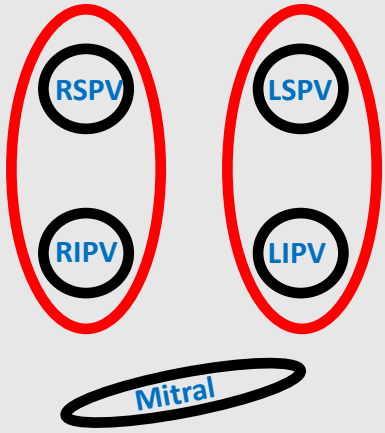
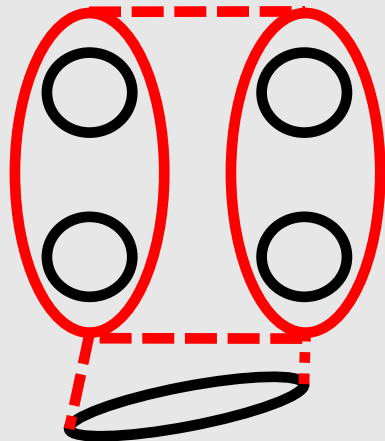
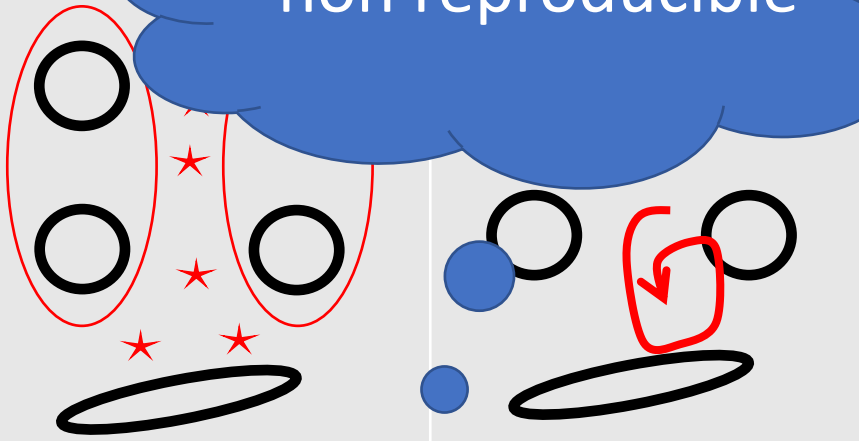
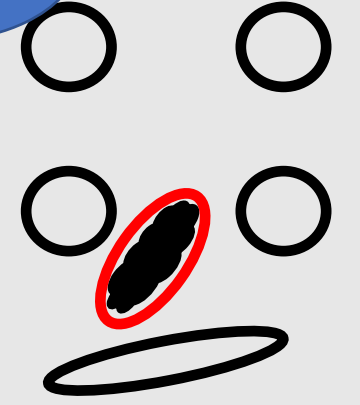
Groupement Hospitalier de Territoire

Grand Paris Nord-Est

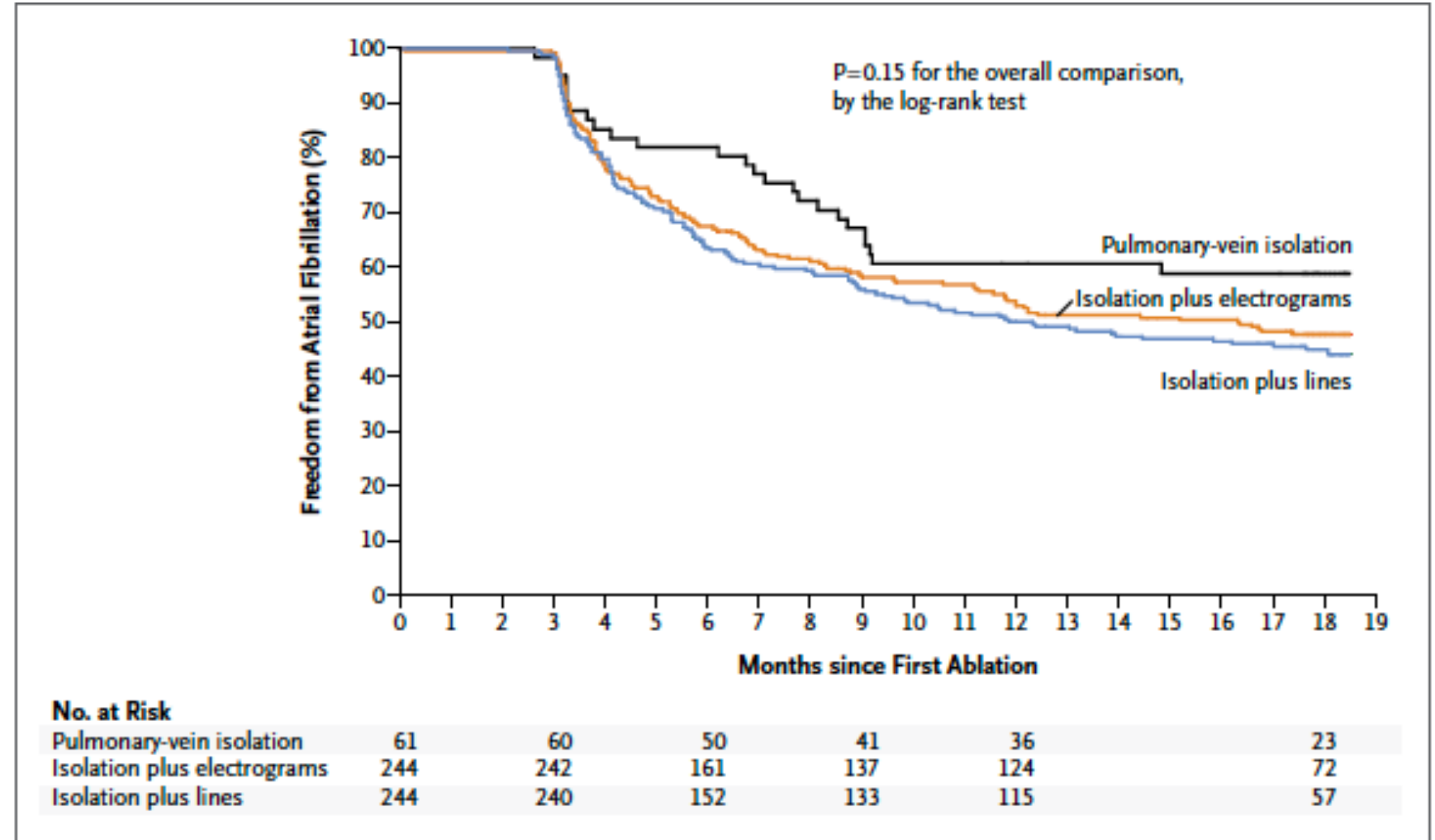
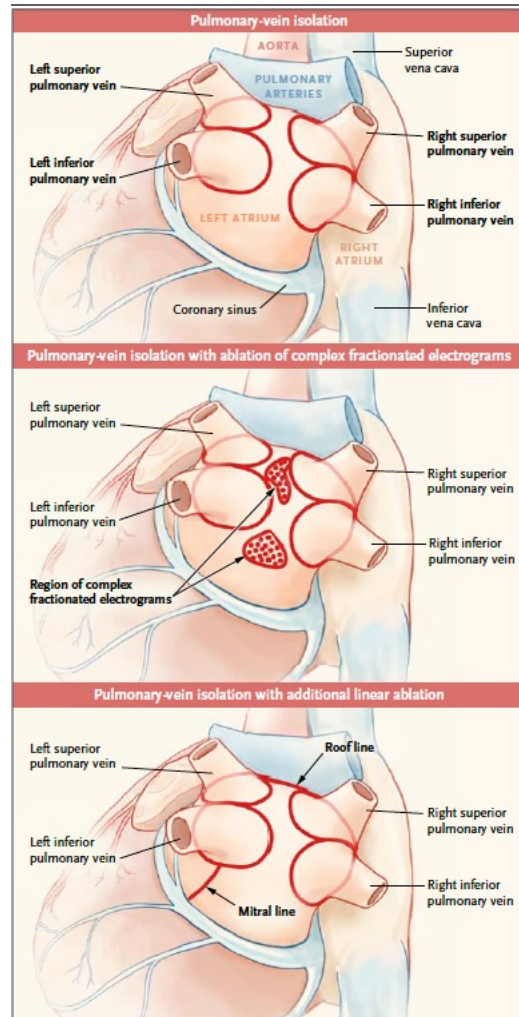
Aulnay-sous-Bois - Le Raincy-Montfermeil * Montreuil



Comment ablater la FA persistante?

PAF	(LS)PsAF		
PVI	Lines	CFA	BIFA
			
AntalPV Block	AF Conversion? SR?		

Et si isoler les veines pulmonaires suffisait



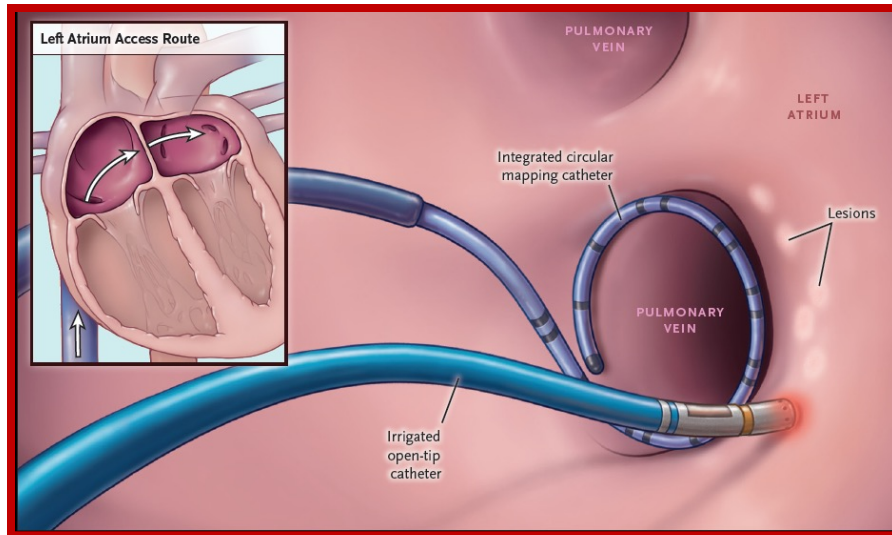
Techniques

Technique	Class-LOE
Isolation VP	I-A
Lignes	IIb-C
Isolation du mur posterieur	IIb-C
Domain Frequency Ablation	IIb-C
Ablation BIFA	IIb-B
Ablation CFAEs	IIb-B
Ablation de rotors	IIb-B
Ablation ganglions	IIb-B

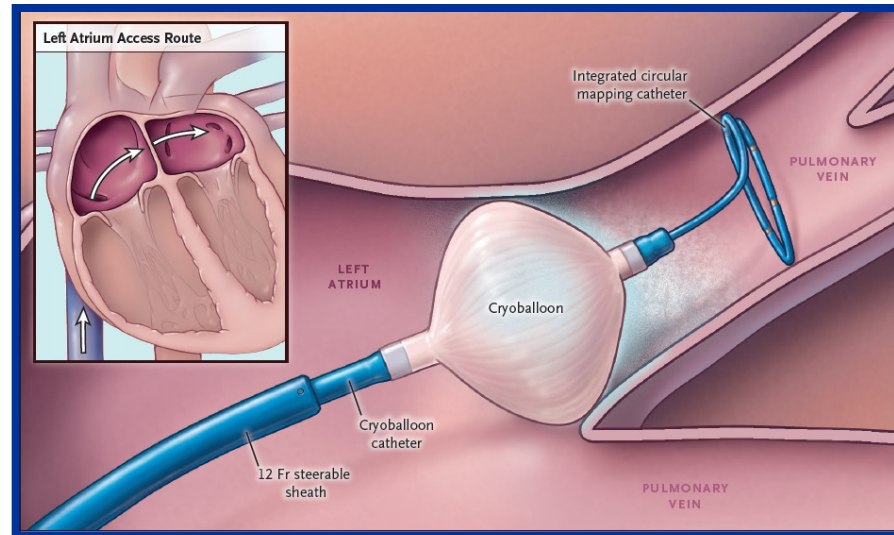
FIRE AND ICE

AF Clinical Trial

- **RFC Ablation (“FIRE”)**
 - Power was not to exceed
 - 40 W at A/I aspect
 - 30 W at P/S aspect
 - 3D electroanatomical mapping

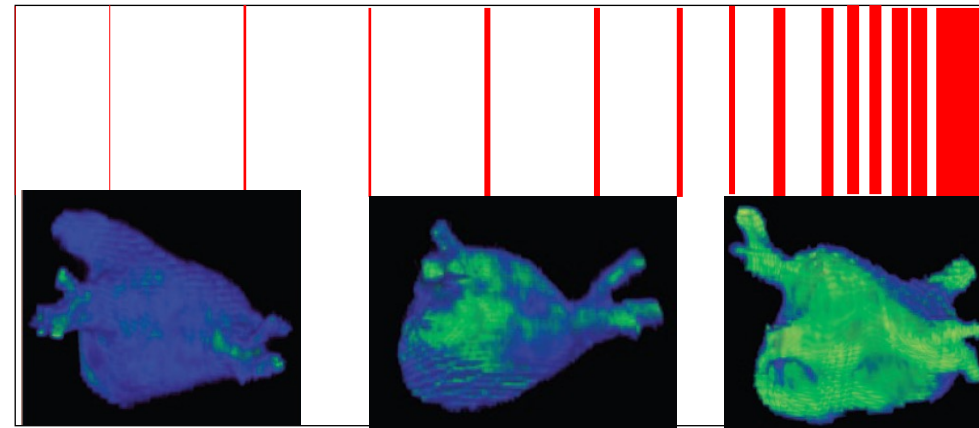


- **Cryoballoon Ablation (“ICE”)**
 - Max. freeze duration of 240s recommended
 - Bonus freeze after isolation recommended
 - Phrenic nerve pacing required



Deuxième regard vers les recommandations....

Factors that have been identified as predictors of a poorer outcome, at least in some studies, include (1) non-PAF and particularly long-term persistent AF; (2) sleep apnea and obesity; (3) increased LA size; (4) increased age; (5) hypertension; and (6) LA fibrosis as detected by cardiac MRI.³⁶⁵



SUIVI POST-ABLATION

Table 17 Key issues in follow-up after AF catheter ablation

Key issues
Recognition and management of complications Patients must be fully informed about the clinical signs and symptoms of rare but potentially dangerous ablation-related complications that may occur after hospital discharge (e.g. atrio-oesophageal fistula, pulmonary vein stenosis).
Follow-up monitoring: Useful to assess procedural success and correlate symptom status with rhythm.^{795,796} Recurrences beyond the first month post-ablation are generally predictive of late recurrences,^{797,798} but recurrent symptoms may be due to ectopic beats or other non-sustained arrhythmia^{642,799,800}; conversely the presence of asymptomatic AF after ablation is well described.^{801–803} Monitoring may be performed with intermittent ECG, Holter, Patch recordings, external or implanted loop recorder, or smart phone monitor (although the latter has not been validated for such use). Patients should be first reviewed at a minimum of 3 months and annually thereafter.¹
Management of antiarrhythmic medication and treatment of AF recurrences a. Continuing AAD treatment for 6 weeks to 3 months may reduce early AF recurrences, rehospitalizations and cardioversions during this period.^{797,804} Clinical practice regarding routine AAD treatment after ablation varies and there is no convincing evidence that such treatment is routinely needed. b. Subsequently, AADs may be weaned, ceased, or continued according to symptoms and rhythm status. Recent findings suggest that in AAD-treated patients remaining free of AF at the end of the blanking period, AAD continuation beyond the blanking period reduces arrhythmia recurrences.⁸⁰⁵
Management of anticoagulation therapy a. In general, OAC therapy is continued for 2 months following ablation in all patients.^{1,806} Beyond this time, a decision to continue OAC is determined primarily by the presence of CHA₂DS₂-VASc stroke risk factors rather than the rhythm status (section 10.2.2.6).

Effect of Catheter Ablation With Vein of Marshall Ethanol Infusion vs Catheter Ablation Alone on Persistent Atrial Fibrillation

The VENUS Randomized Clinical Trial

Miguel Valderrábano, MD; Leif E. Peterson, PhD; Vijay Swarup, MD; Paul A. Schurmann, MD; Akash Makkar, MD; Rahul N. Doshi, MD; David DeLurgio, MD; Charles A. Athill, MD; Kenneth A. Ellenbogen, MD; Andrea Natale, MD; Jayanthi Koneru, MD; Amish S. Dave, MD, PhD; Irakli Giorgberidze, MD; Hamid Afshar, MD; Michelle L. Guthrie, RN; Raquel Bunge, RN; Carlos A. Morillo, MD; Neal S. Kleiman, MD

Figure 1. Clinical Trial Conduct

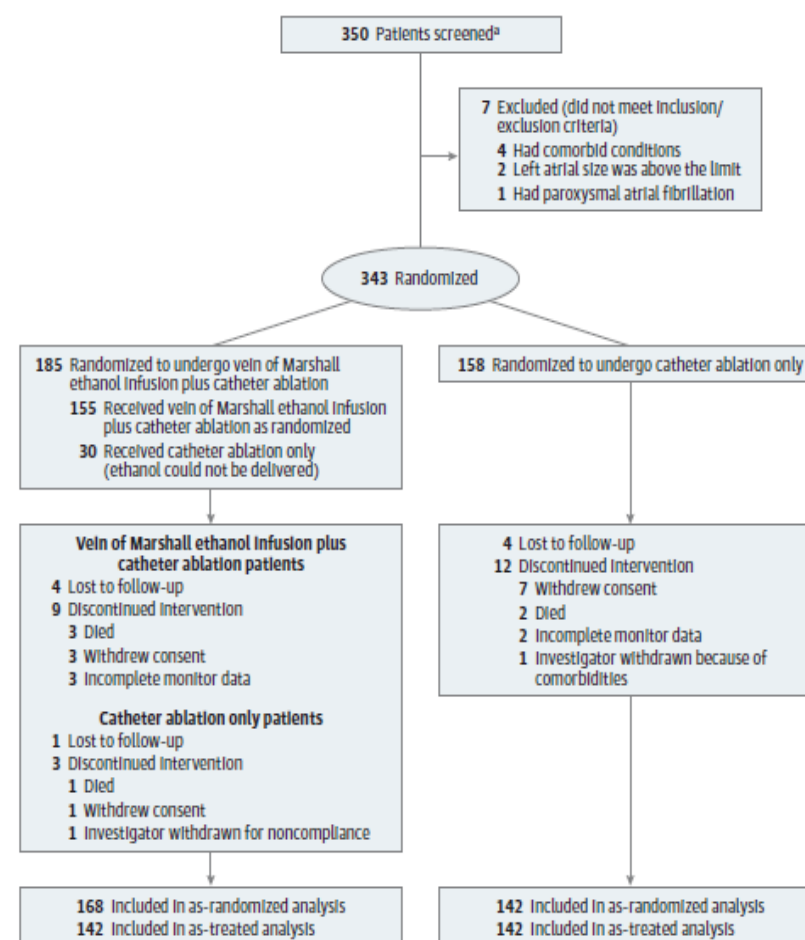
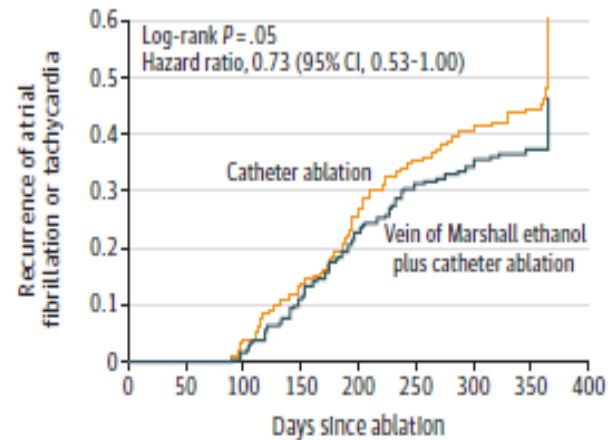


Table 1. Patient Demographic Characteristics

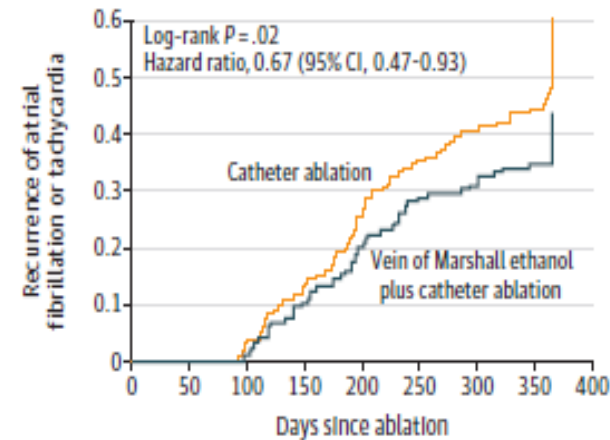
	No. (%)	
	Vein of Marshall-catheter ablation (n = 185)	Catheter ablation (n = 158)
Demographics		
Age, mean (SD), y	66.6 (9.6)	66.4 (9.9)
Sex, No. (%)		
Male	137 (74)	124 (78)
Female	48 (26)	34 (22)
Race and ethnicity		
White	169 (91)	150 (95)
Black	5 (3)	2 (1)
Hispanic	3 (2)	1 (1)
Asian	1 (1)	2 (1)
Not stated	7 (4)	3 (2)
Medical history and risk factors		
Hypertension	144 (77)	104 (66)
Diabetes	52 (28)	31 (20)
Coronary disease	52 (28)	41 (26)
Stroke-TIA	19 (10)	19 (12)
Heart failure	48 (26)	42 (27)
Body mass index ^a	31.2 (6.6)	31.9 (6.5)
CHA ₂ DS ₂ -VASC score ^b	2.9 (1.6)	2.6 (1.6)
Cardiac parameters		
Ejection fraction, %	52.1 (10.1)	53.4 (9.4)
Left atrial diameter, mm	44.8.1 (7.9)	47.0 (7.5)
Left atrial volume, mL	110.9 (46.8)	113.9 (46.3)
Time from first AF diagnosis		
<6 mo	15 (8)	10 (6)
6 mo to 2 y	76 (41)	65 (41)
>2 y	94 (51)	83 (52)
Longstanding persistent AF, No. (%) ^c	99 (54)	82 (52)

A Atrial fibrillation or tachycardia occurrence after single procedure in as-randomized analysis



No. at risk									
Vein of Marshall ethanol plus catheter ablation	185	180	174	153	129	116	108	89	68
Catheter ablation	158	157	148	132	110	95	86	69	54

B Atrial fibrillation or tachycardia occurrence after single procedure in as-treated analysis



No. at risk									
Vein of Marshall ethanol plus catheter ablation	155	151	145	129	111	100	95	77	58
Catheter ablation	158	157	148	132	110	95	86	69	54

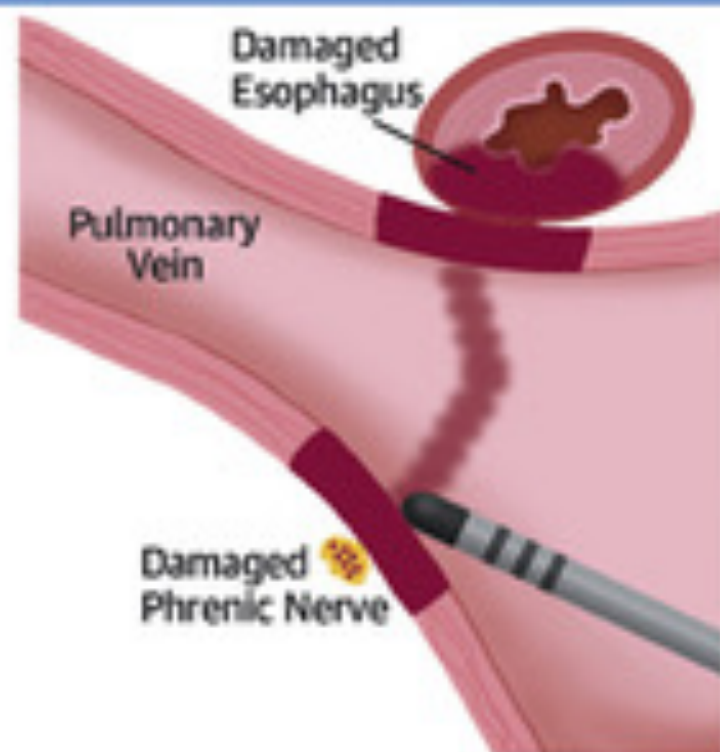
-ETUDE DANS 11 CENTRES US

-DANS 16% DES CAS L'ALCOOLISATION N'A PAS PU ETRE EFFECTUEE

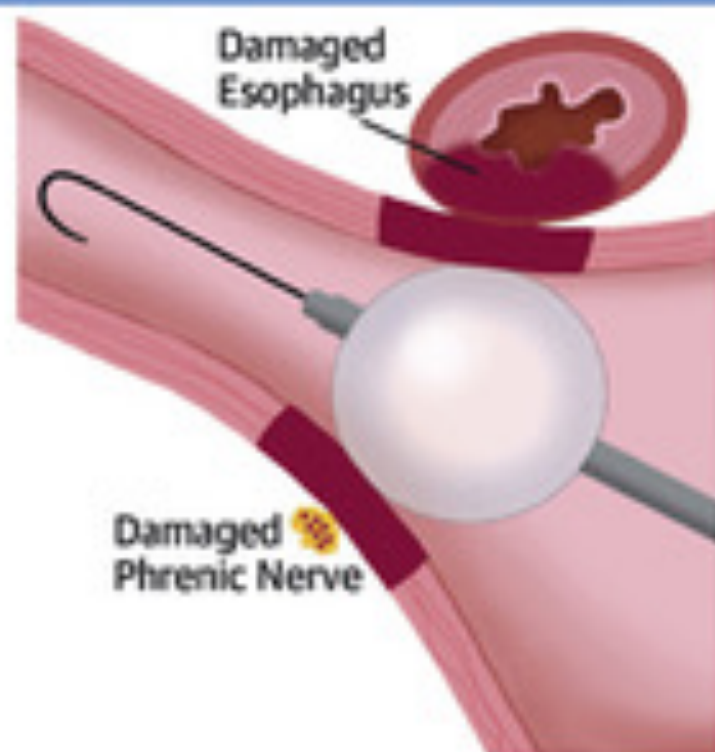
-Critère primaire de jugement : : récidence de FA/ flutter d'au moins 30 secondes après 3 mois après l'ablation = critère classique avec 1 mois d'ECG continu à 6 et 12 mois.

-TAUX DE SUCCES A 1 AN 38% PVI vs 51% dans PVI+LOM

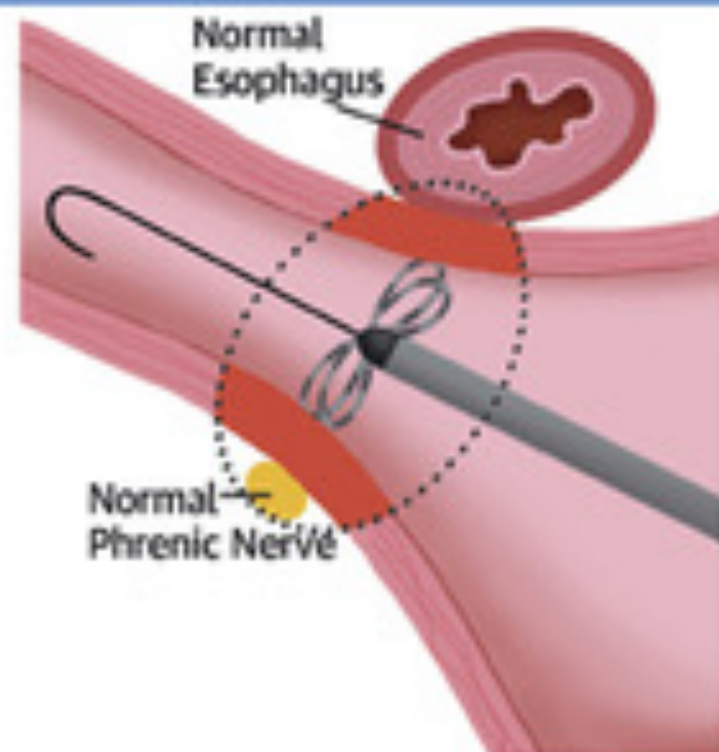
Radiofrequency Ablation



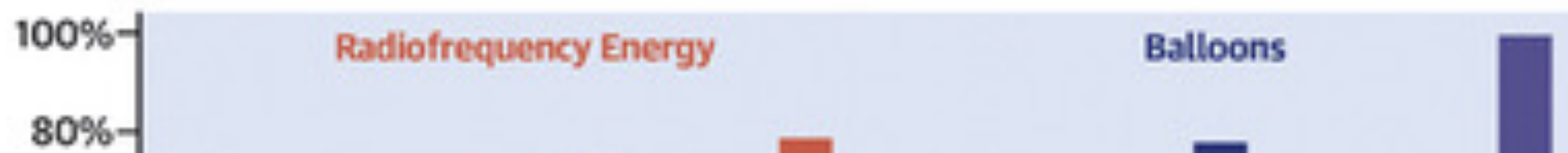
Cryoballoon Ablation



Pulsed Field Ablation



Frequency of Patients With ALL PVs Durably Isolated



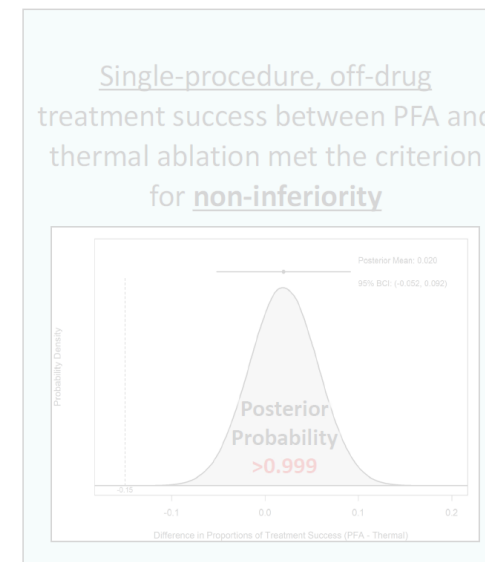
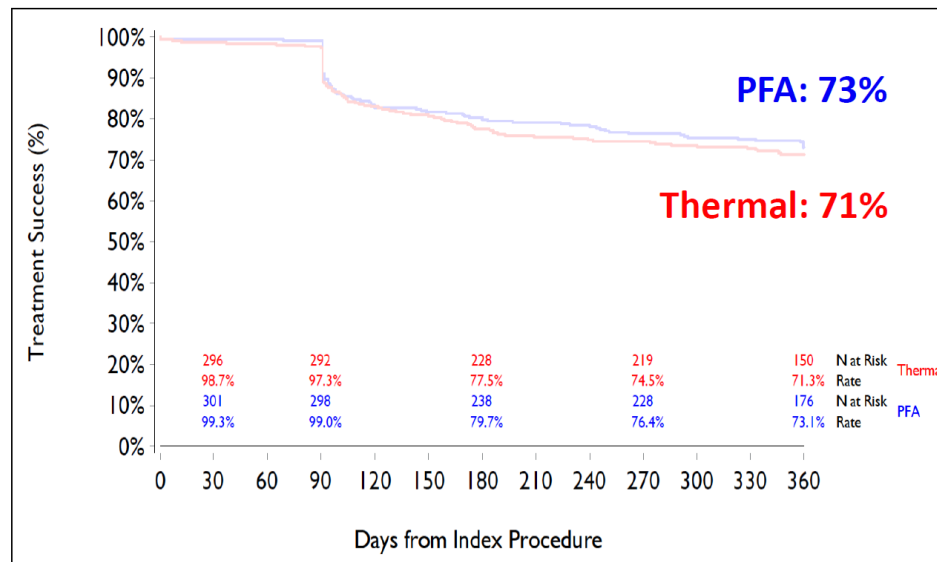
Randomized Controlled Trial for Pulsed Field Ablation versus Standard of Care Thermal Ablation for Paroxysmal Atrial Fibrillation

Primary Results of the *ADVENT* Trial

Vivek Y. Reddy MD,¹ Edward P. Gerstenfeld MD,² Andrea Natale MD,³ William Whang MD,¹ Frank A. Cuoco MD,⁴ Chinmay Patel MD,⁵ Stavros E. Mountantonakis

Primary Effectiveness Endpoint

(Acute Success + 1-Yr Freedom from Atrial Arrhythmias / Re-Ablation / Cardioversion / AAD Use)



Primary Safety Endpoint

	Serious Composite Safety Events	
	Pulsed Field Group, N = 305 n (%)	Thermal Group, N = 302 n (%)
Any Composite Safety Event	6 (2.0) †	4 (1.3)
Death	1 (0.3)	0
Myocardial infarction	0	0
Persistent phrenic nerve palsy	0	0
Stroke	0	1 (0.3)
Transient ischemic attack	1 (0.3)	0
Systemic thromboembolism	0	0
Cardiac tamponade or perforation	2 (0.7)	0
Pericarditis	1 (0.3)	0
Pulmonary edema	1 (0.3)	1 (0.3)
Vascular access complication	1 (0.3)	2 (0.7)
Heart block	0	0
Gastric motility/ pyloric spasm	0	0
Pulmonary vein stenosis	0	0
Atrio-esophageal fistula	0	0

† One patient who sustained a cardiac tamponade subsequently died; accordingly, the individual components add to more than the composite total.

Is this death spurious or specific to PFA?

- The pentaspline PFA catheter received CE-Mark Approval in March 2021
- Registry of all sites performing PFA
 - 24 EU centers / 77 operators
 - 1,568 patients
- Mortality:
 - **1 in 1,568 → 0.06%**

M.Turagam, P.Neuzil, B.Schmidt...VY.Reddy, *Circulation* 148:35–46 (2023)





Conclusion

- CASTLE AF, CABANA = Premières données quant à un effet sur la morbidimortalité de l'ablation de la FA
- Effet notamment dans l'insuffisance cardiaque
- Importance de la sélection des patients, de l'information et de l'expérience du centre et de l'opérateur.
- Simplification des procédures pourrait contribuer à de meilleurs résultats procéduraux