



HFpEF Masterclasses
in centers of expertise



FRANCE

7th November 2024 - DAY 1

8th November 2024 - DAY 2

Management of atrial fibrillation and coronary artery disease in patients with HFpEF

Pr. Nathan Mewton, Lyon

HCL
HOSPICES CIVILS
DE LYON

L'INSTITUT DE
CARDIOLOGIE

ALI MACGRAW

RYAN O'NEAL

LOVESTORY

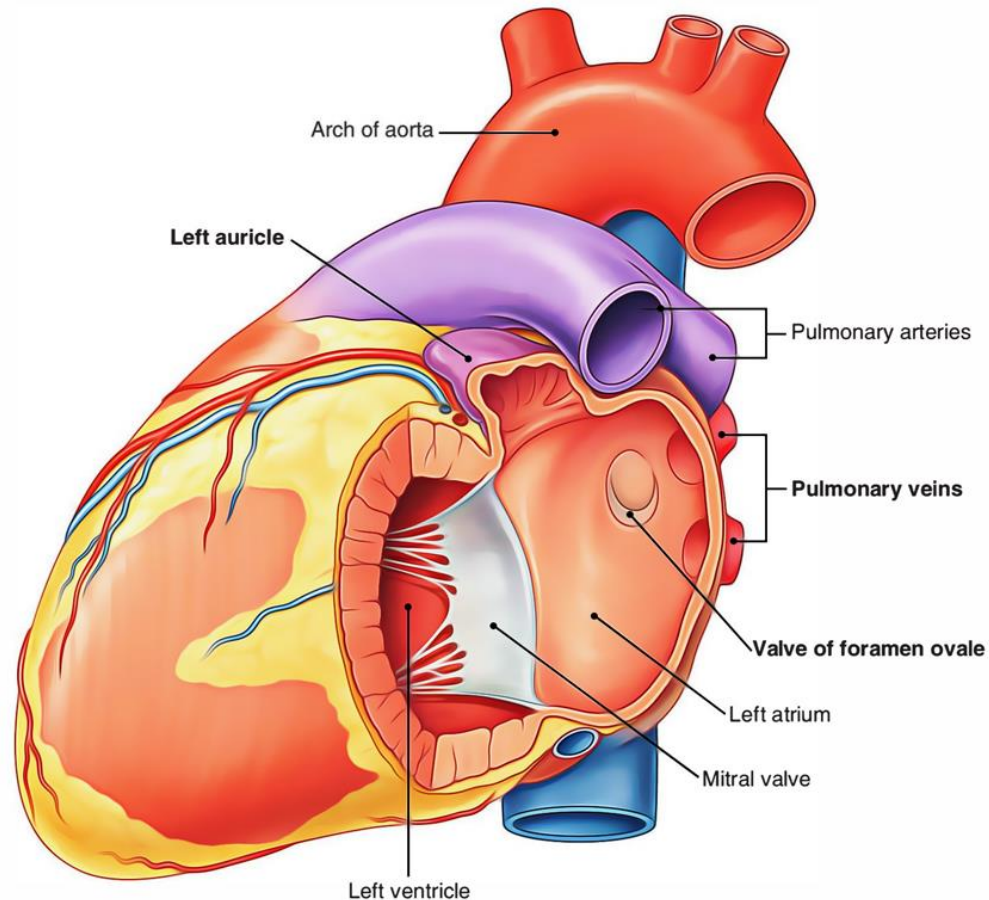
L'AMOUR C'EST N'AVOIR JAMAIS À DIRE QU'ON EST DÉSOLÉ

HFPEF

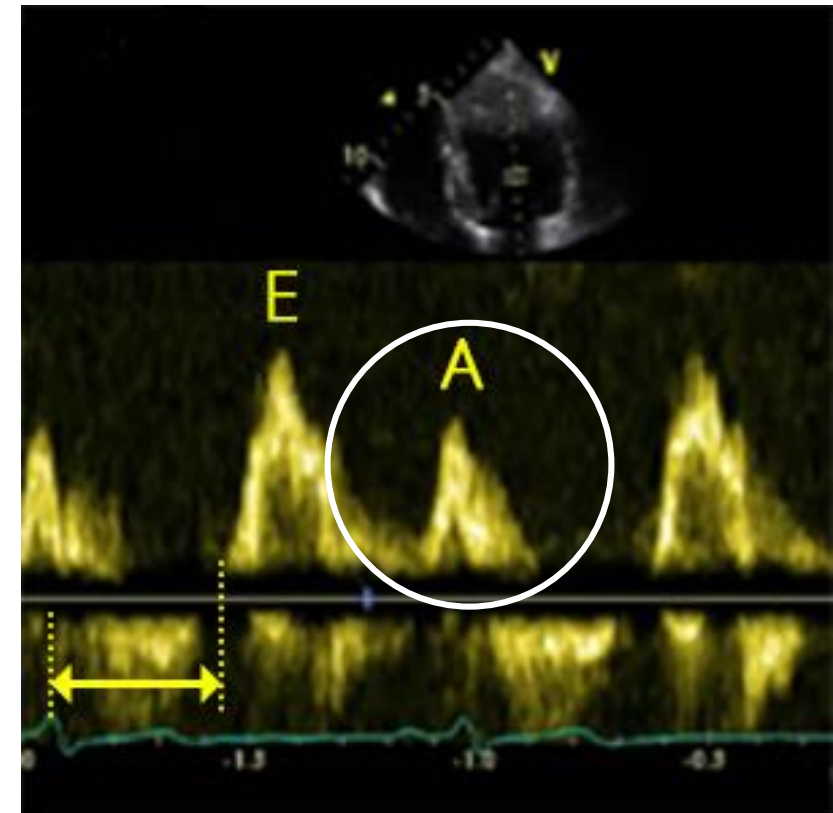
AF



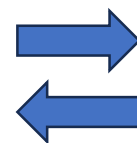
Healthy Left Atrium



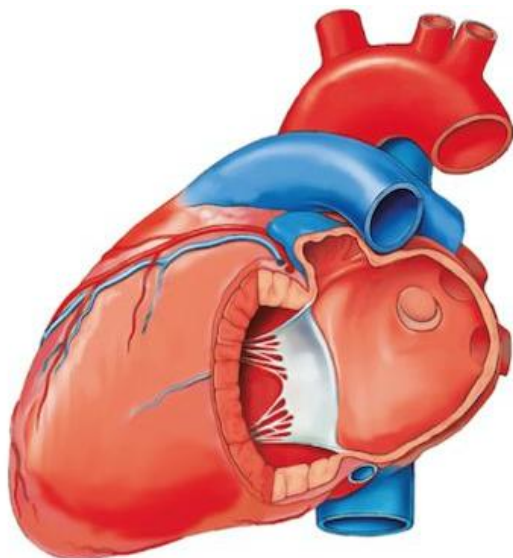
- Index volume $< 34 \text{ ml/m}^2$
- Atrial Diastolic pressure = 0-5 mmHg
- Atrial Systolic pressure = 10-15 mmHg
- Atrial systole produces the A wave
- Improves LV filling by 25-30%



Atrial Myopathy



Atrial Fibrillation



Atrial dilatation
Atrial cardiomyopathy?

Genetic

Aging

Hypertension

Coronary artery disease

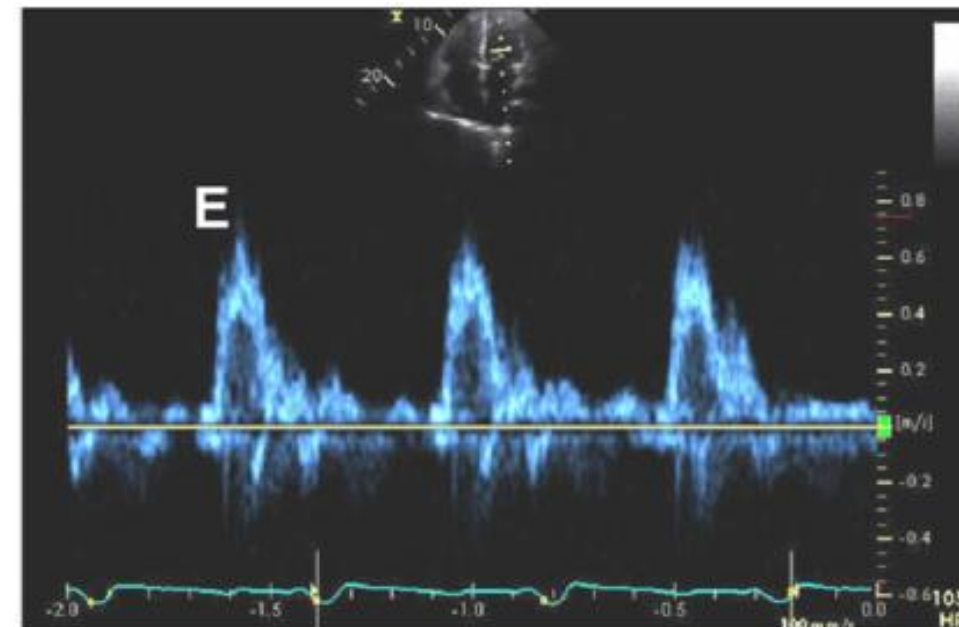
Valvular heart disease

Type 2 diabetes

Obesity

Chronic respiratory disorders

Risk factors and predisposing comorbidities



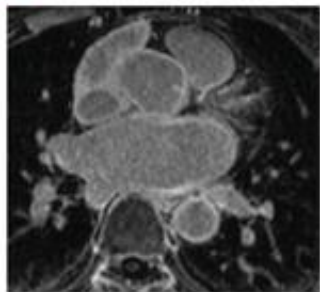
Loss of LA contraction
Reduction in LV filling by 20-30%
LVEF reduction up to 10%

Assessment of Left Atrial Myopathy in Heart Failure

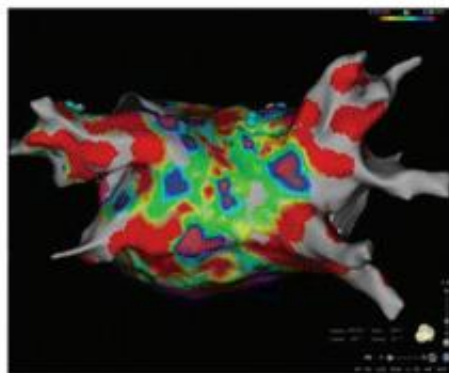
LA Structure



LA Dilation

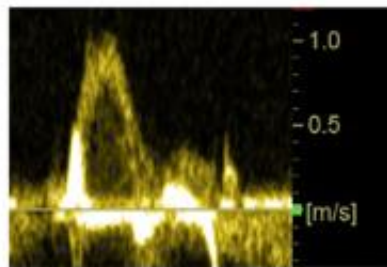


Fibrosis on CMR

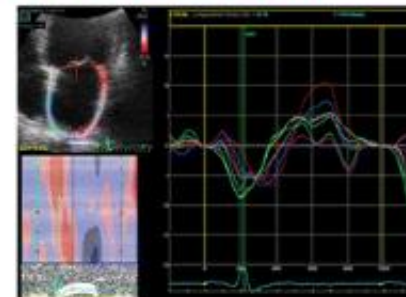


Areas of Low Voltage

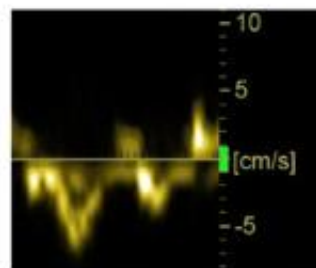
LA Function



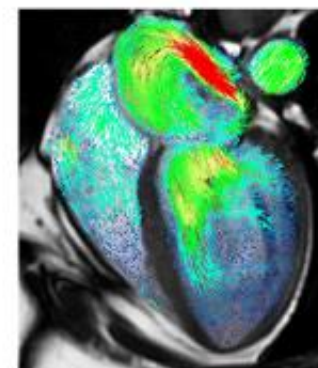
Low A wave velocity



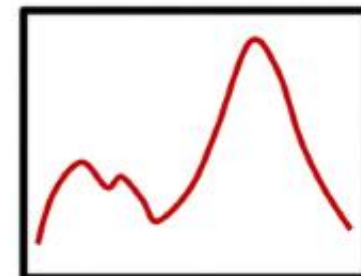
Reduced Strain



Low a' tissue Doppler velocity

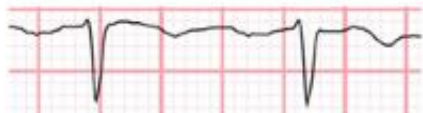


LA Stasis on 4D Flow CMR

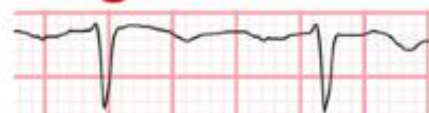


PCWP: Large v waves in absence of significant MR

Supportive Diagnostics

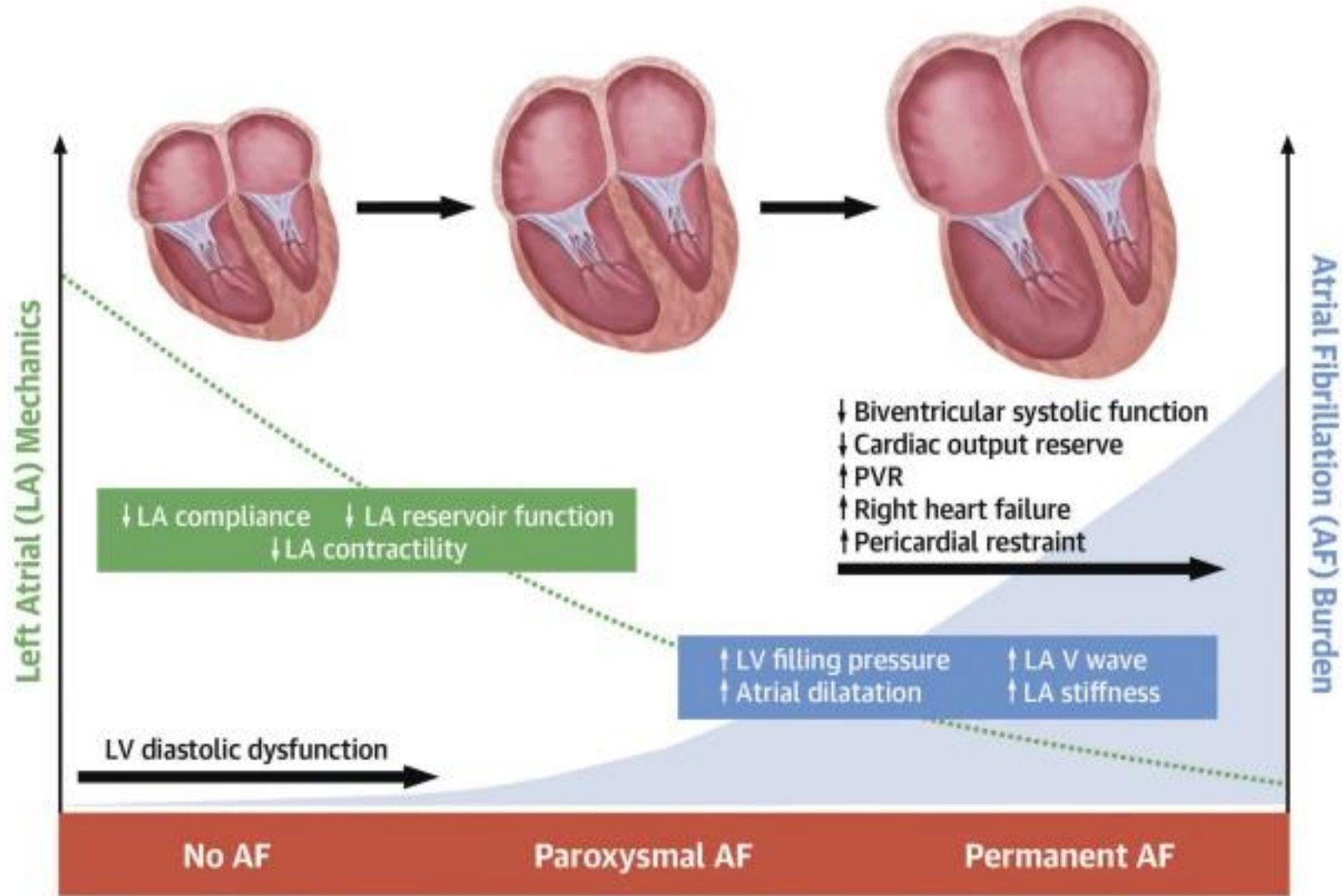


High AF burden



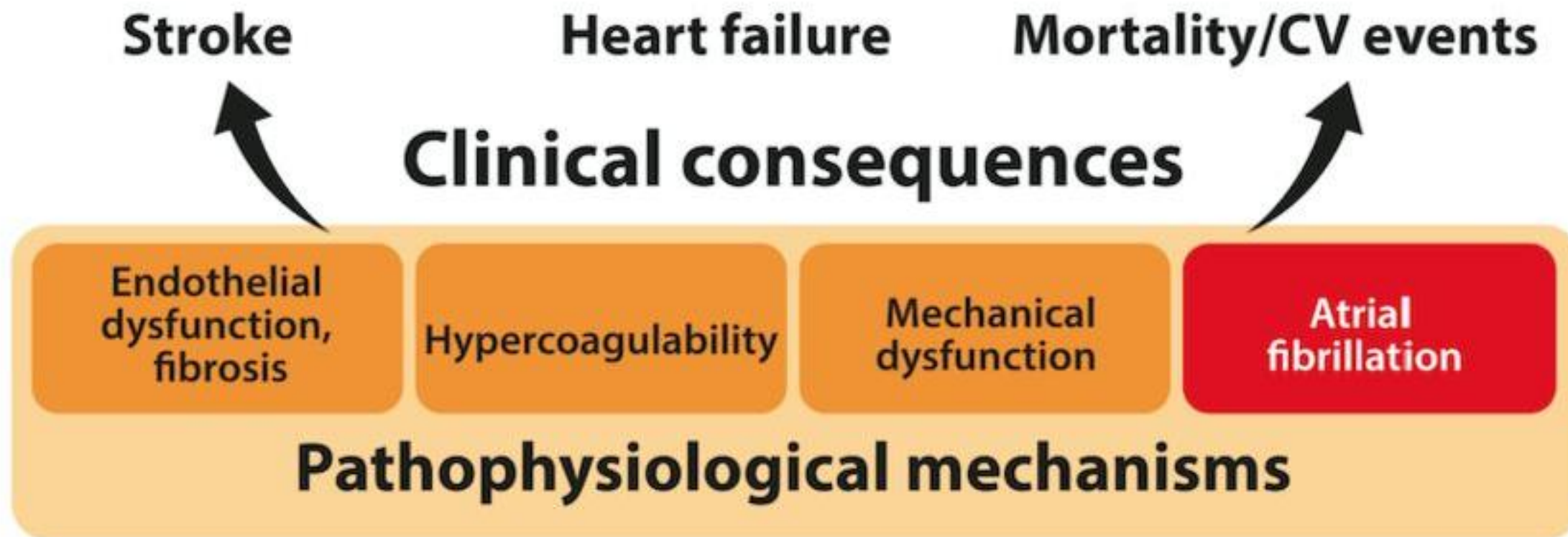
Low amplitude P-waves in sinus rhythm

CENTRAL ILLUSTRATION: Progressive Left Atrial Myopathy and Atrial Fibrillation Burden in Heart Failure With Preserved Ejection Fraction



Reddy, Y.N.V. et al. J Am Coll Cardiol. 2020;76(9):1051-64.

Clinical Consequences of AF



Characterizing atrial fibrillation in patients with and without heart failure across the ejection fraction spectrum: Incidence, prevalence, and treatment strategies

Valeria Valente¹ , **Giulia Ferrannini^{1,2}**, **Lina Benson¹**, **Paolo Gatti^{1,2}**,
Federica Guidetti¹, **Michael Melin^{1,3}**, **Frieder Braunschweig^{1,4}**, **Cecilia Linde^{1,4}**,
Ulf Dahlström⁵, **Lars H. Lund^{1,4}**, **Marat Fudim^{6,7}**, and **Gianluigi Savarese^{1*}** 

Methods

Study population: SwedeHF registry matched 1:1 by sex, year of birth and, for individuals <90 years old, county of residence with a non-HF cohort chosen at random from the Swedish population.

195 106 patients (97 553 with and 97 553 without HF), of which 63% were men, median age 75 years (Q1–Q3: 66–82).

Cohort 1 : prevalence of AF overall by 3-year periods of time and treatment management

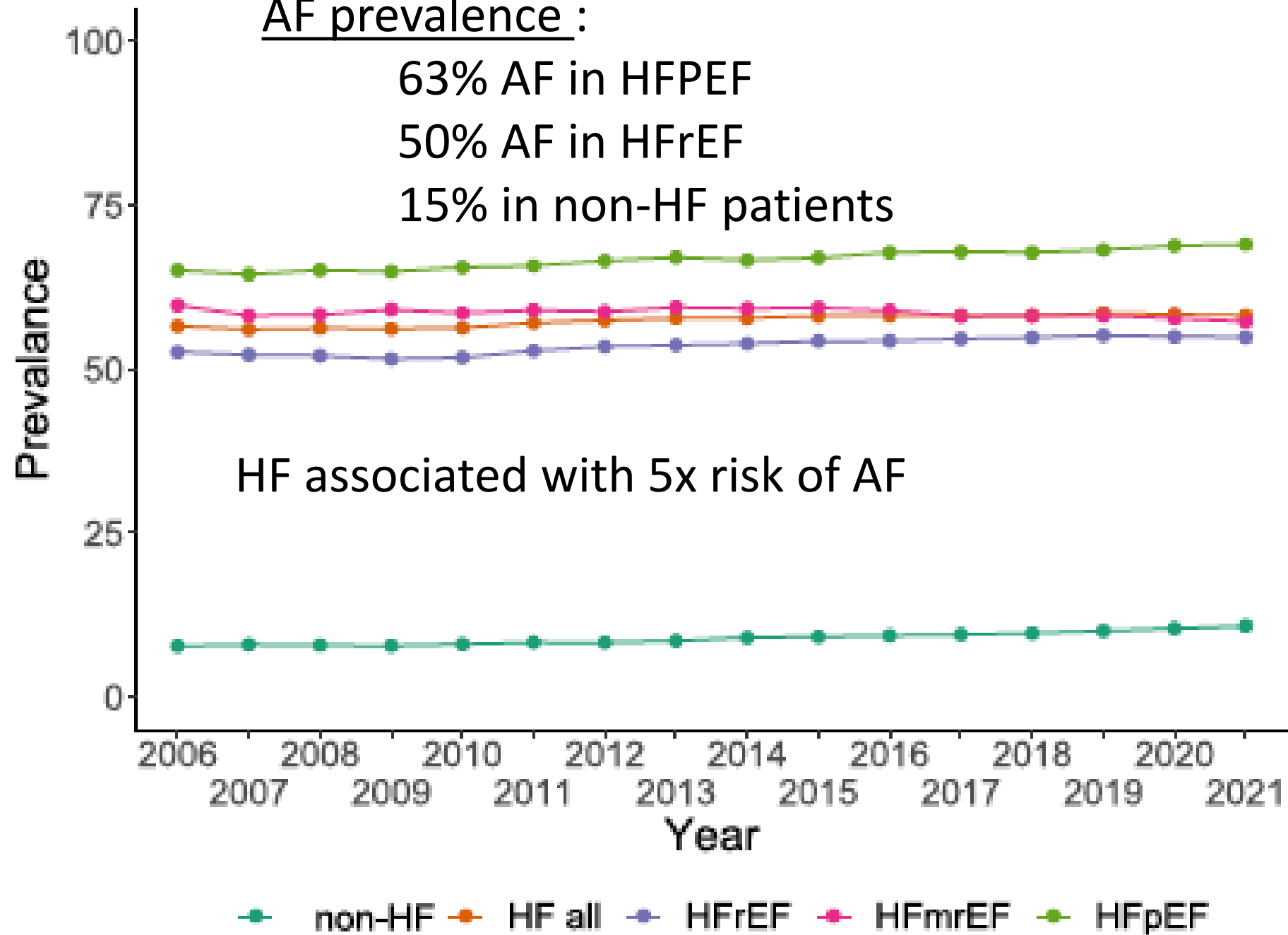
Cohort 2 : incidence of AF over time and predictors

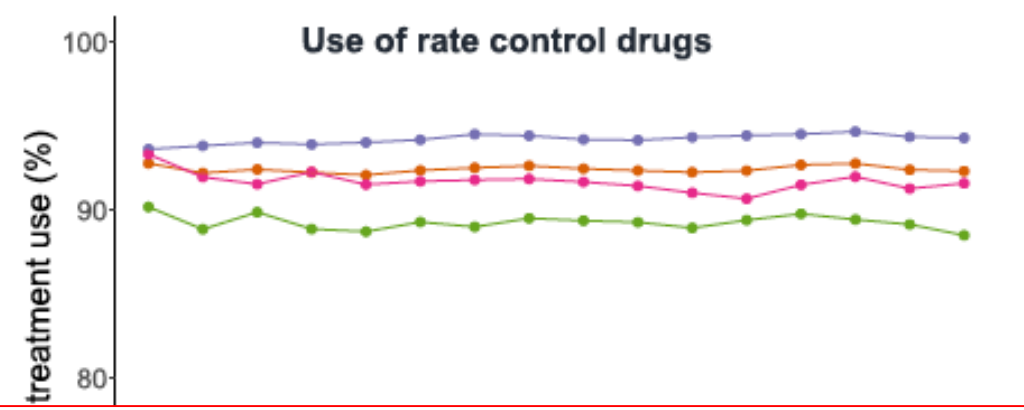
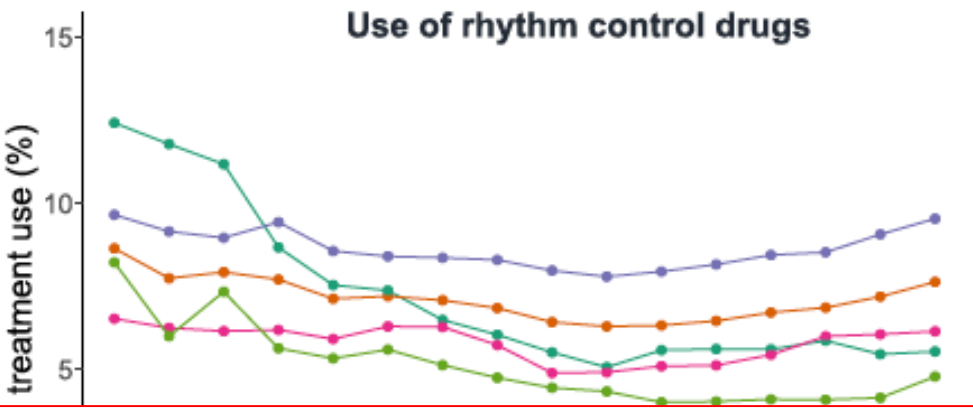
AF prevalence :

63% AF in HFpEF

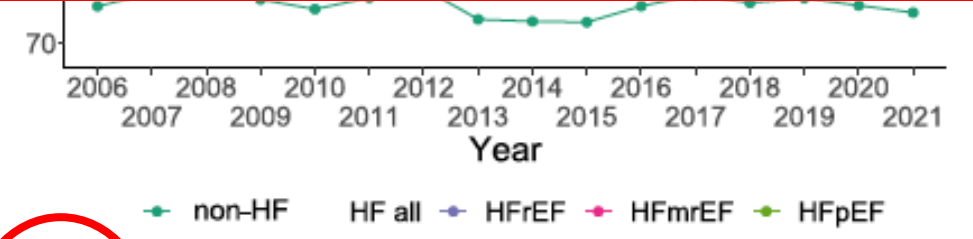
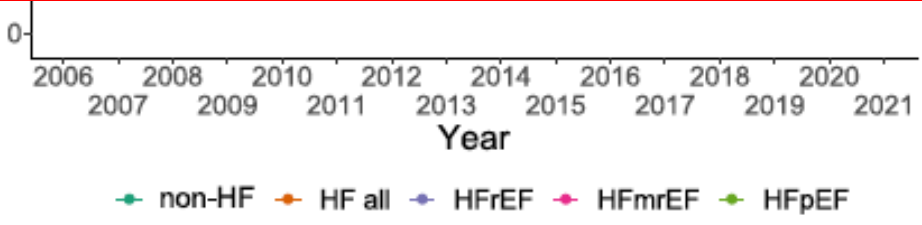
50% AF in HFrEF

15% in non-HF patients

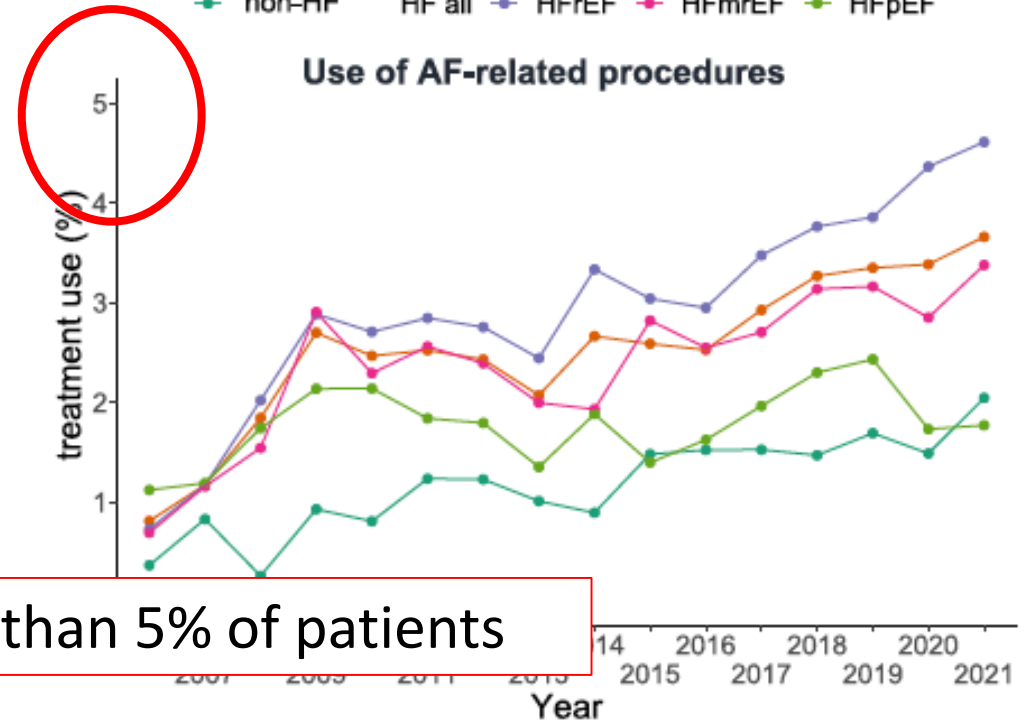
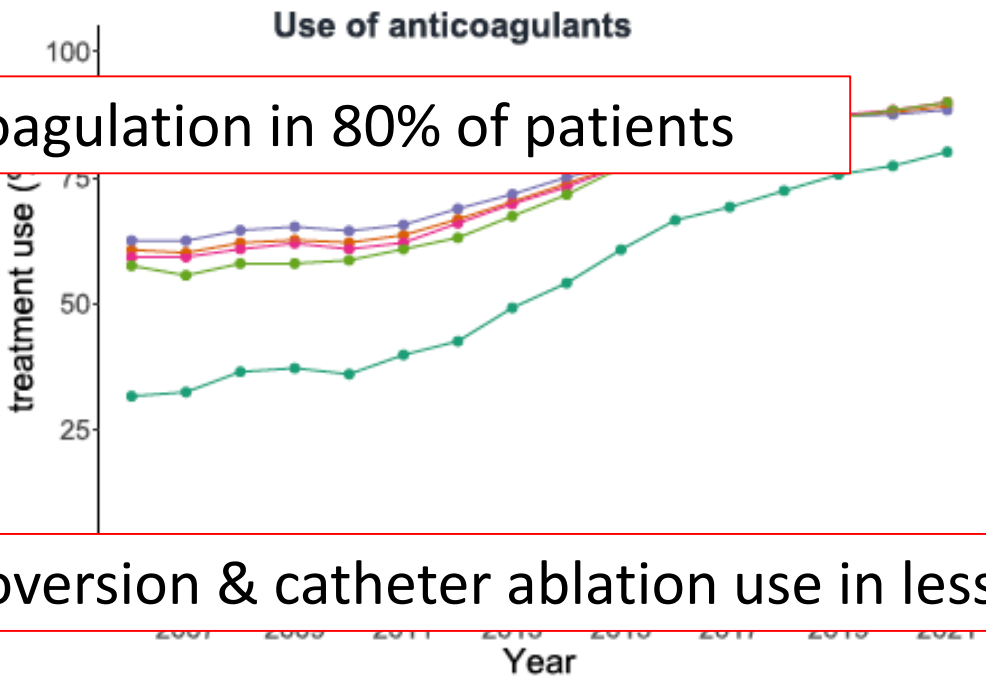




Rate control drugs used in > 90% of HF patients and <15% antiarrhythmic drugs

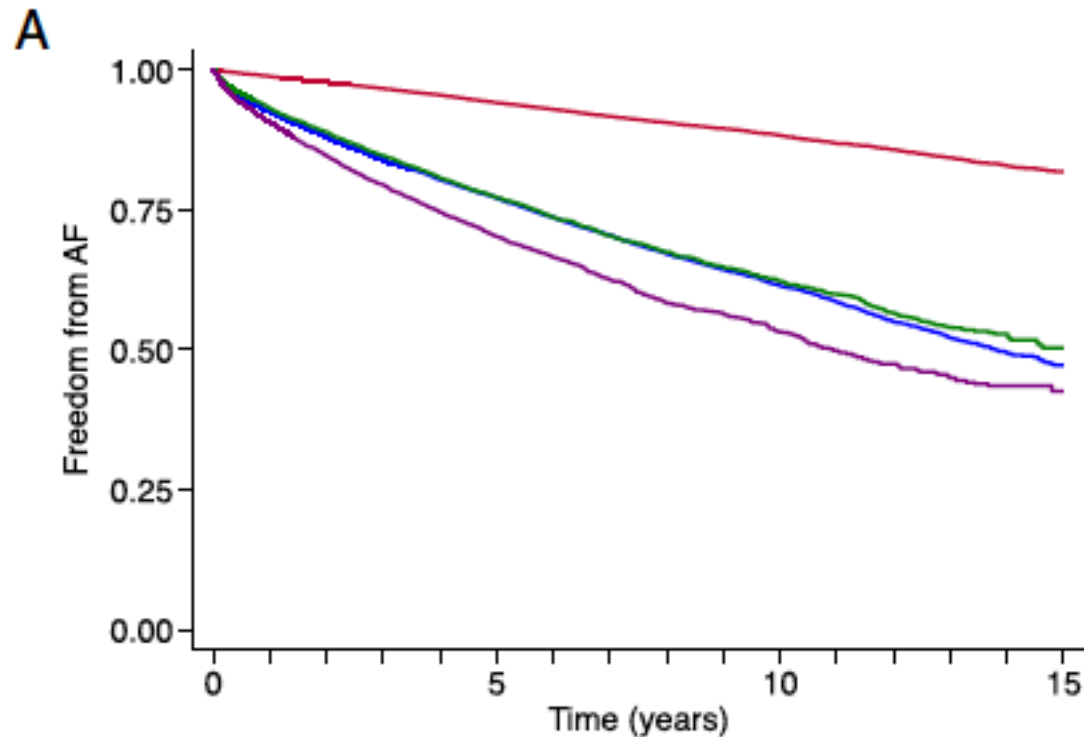


Anticoagulation in 80% of patients



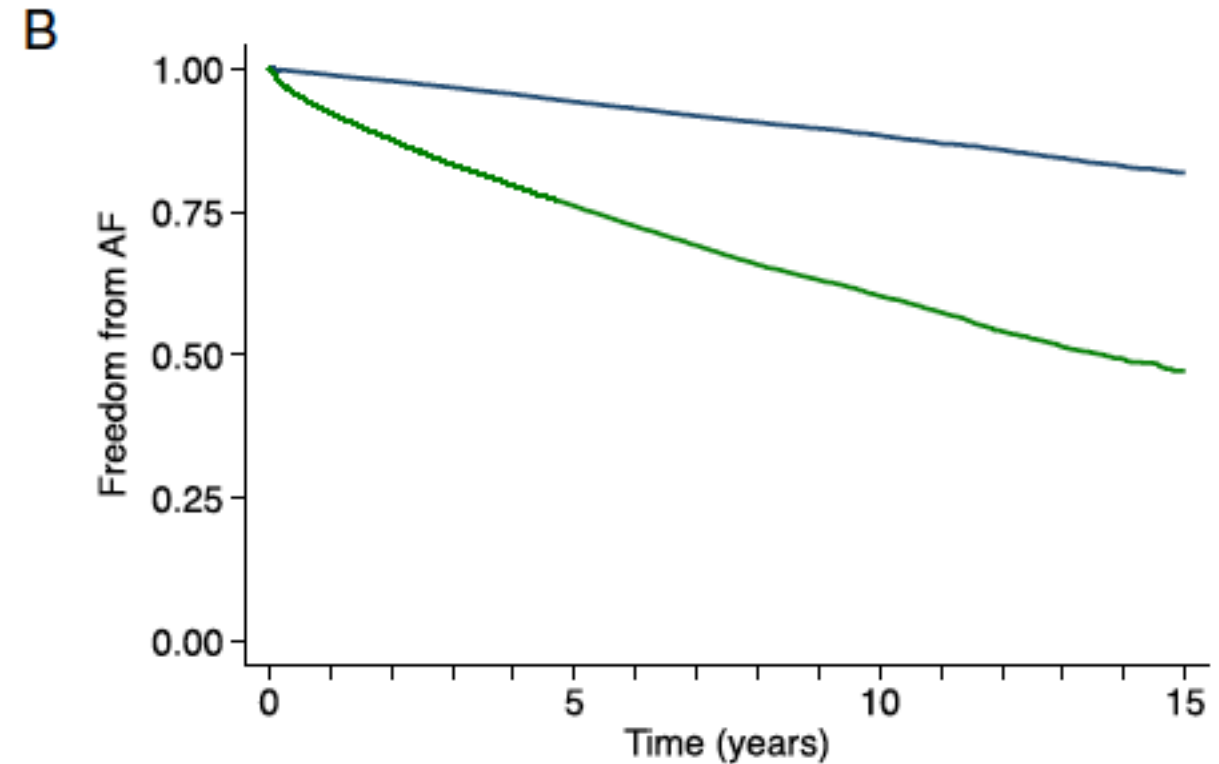
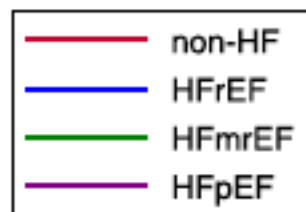
Cardioversion & catheter ablation use in less than 5% of patients

Incidence of AF in HF patients = 0.03% pt / year



Number at risk

non-HF 43081	19937	8002	650
HFfrEF 25471	8257	2515	174
HFmrEF 9771	3016	884	61
HFpEF 7846	2034	550	36



Number at risk

non-HF 43081	19937	8002	650
HF 43088	13307	3949	271

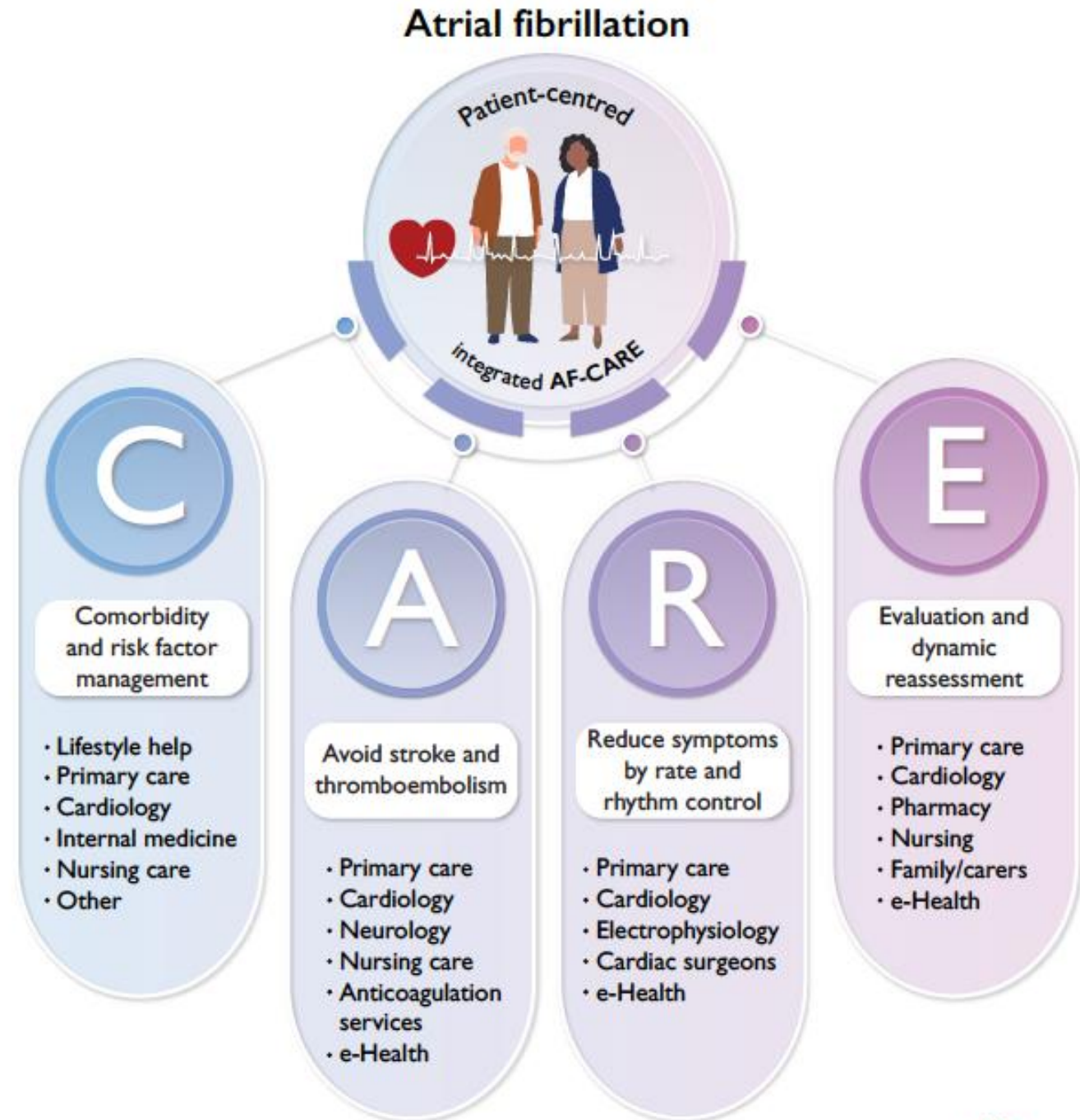
HFPEF independent predictor of AF (HR=3.12)
Higher in HFPEF vs HFfrEF

AF Management ESC Guidelines 2024

Integrated AF-CARE

The first treatment of AF in HFPEF is the HF treatment !

Van Gelder et al. EHJ 2024



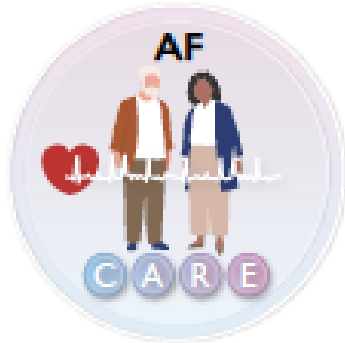
ANTICOAGULATION NEW SCORE = CHADS₂-VA

Table 10 Updated definitions for the CHA₂DS₂-VA score

CHA ₂ DS ₂ -VA component		Definition and comments	Points awarded ^a
C	Chronic heart failure	Symptoms and signs of heart failure (irrespective of LVEF, thus including HFpEF, HFmrEF, and HFrEF), or the presence of asymptomatic LVEF ≤40%. ^{261–263}	1
H	Hypertension	Resting blood pressure >140/90 mmHg on at least two occasions, or current antihypertensive treatment. The optimal BP target associated with lowest risk of major stroke or events is 120–129/70–79 mmHg (or keep as low as reasonably achievable).	1
A	Age 75 years or above	Age is an independent determinant of ischaemic stroke risk and risk is a continuum, but for reasons of practicality, two points are given for age ≥75 years.	2
D	Diabetes mellitus	Diabetes mellitus (type 1 or type 2), as defined by accepted criteria, ²⁶⁶ or treatment with glucose lowering therapy.	1
S	Prior stroke, TIA, or arterial thromboembolism	Previous thromboembolism is associated with an elevated risk of recurrence and therefore weighted 2 points.	2
V	Vascular disease	Coronary artery disease, including myocardial infarction, angina, history of coronary revascularization (percutaneous or surgical), and significant CAD on angiography or cardiac imaging. ²⁶⁷ OR Peripheral vascular disease, including: intermittent claudication, previous revascularization for PVD, percutaneous or surgical intervention on the abdominal aorta, and complex aortic plaque on imaging (defined as features of mobility, ulceration, pedunculation, or thickness ≥4 mm). ^{268,269}	1
A	Age 65–74 years	1 point is given for age between 65 and 74 years.	1

No more gender difference!

Recommendations	Class ^a	Level ^b
Oral anticoagulation is recommended in patients with clinical AF at elevated thromboembolic risk to prevent ischaemic stroke and thromboembolism. ^{239,240}	I	A
A CHA ₂ DS ₂ -VA score of 2 or more is recommended as an indicator of elevated thromboembolic risk for decisions on initiating oral anticoagulation.	I	C
Oral anticoagulation is recommended in all patients with AF and hypertrophic cardiomyopathy or cardiac amyloidosis, regardless of CHA ₂ DS ₂ -VA score, to prevent ischaemic stroke and thromboembolism. ^{270–276}	I	B
Individualized reassessment of thromboembolic risk is recommended at periodic intervals in patients with AF to ensure anticoagulation is started in appropriate patients. ^{277–280}	I	B



Equality in healthcare provision (gender, ethnicity, socioeconomic) (Class I)

Education for patients, families and healthcare professionals (Class I)

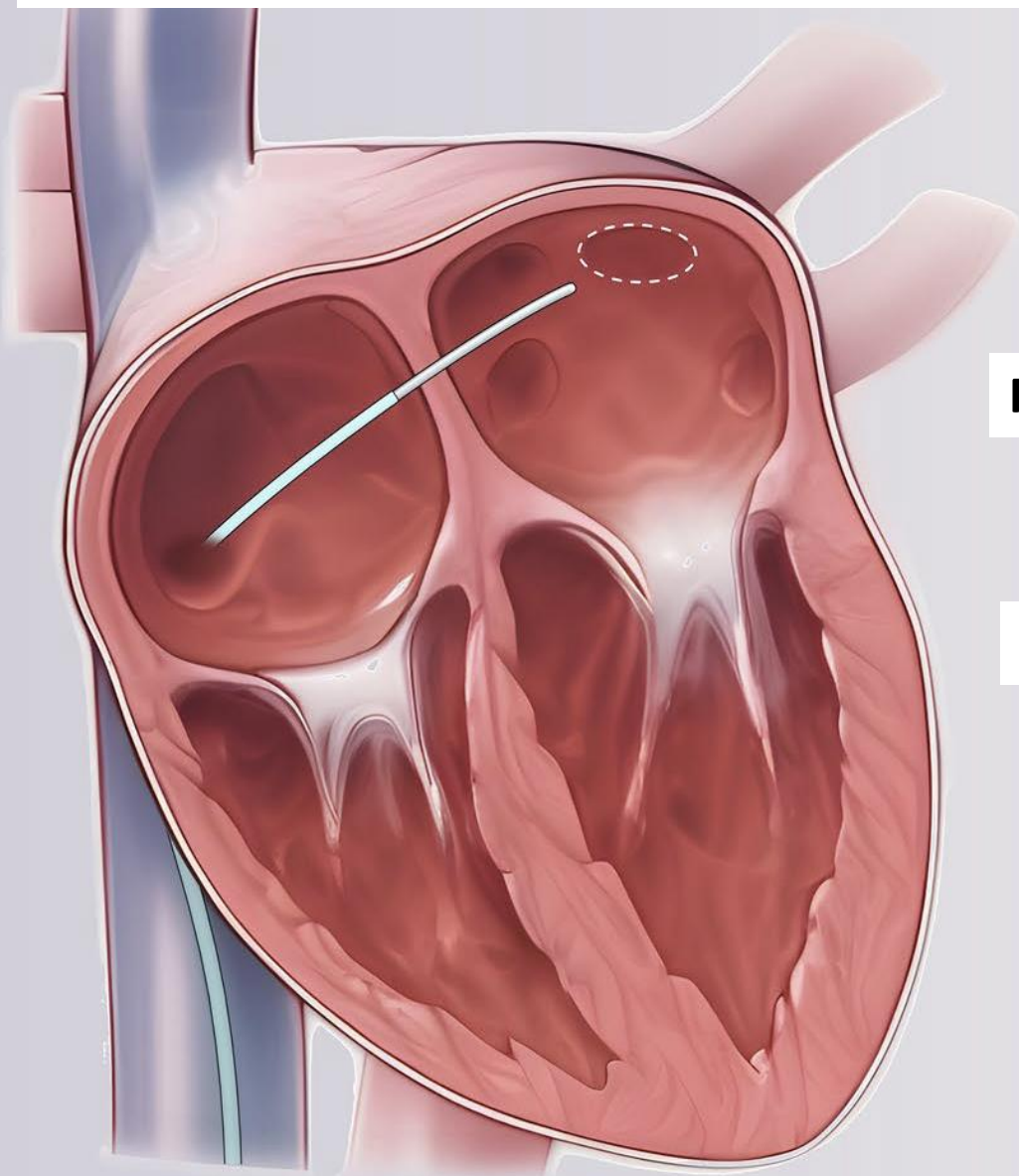
Patient-centred AF management with a multidisciplinary approach (Class IIa)



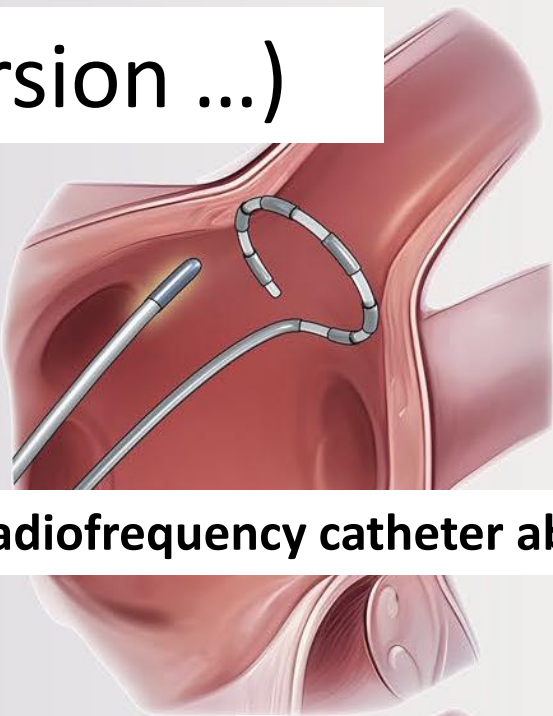
Comorbidity and risk factor management

Hypertension	Heart failure	Overweight or obese	Obstructive sleep apnoea	Alcohol
Blood pressure lowering treatment (Class I)	Diuretics for congestion (Class I)	Weight loss (target 10%) ^a (Class I)	Management of OSA ^a (Class IIb)	Reduce to ≤3 drinks per week (Class I)
Diabetes mellitus	Appropriate HFrEF medical therapy (Class I)	Bariatric surgery if rhythm control ^a (Class IIb)	Exercise capacity	Other risk factors/comorbidities
Effective glycaemic control ^a (Class I)	SGLT2 inhibitors (Class I)		Tailored exercise programme (Class I)	Identify and manage aggressively ^a (Class I)

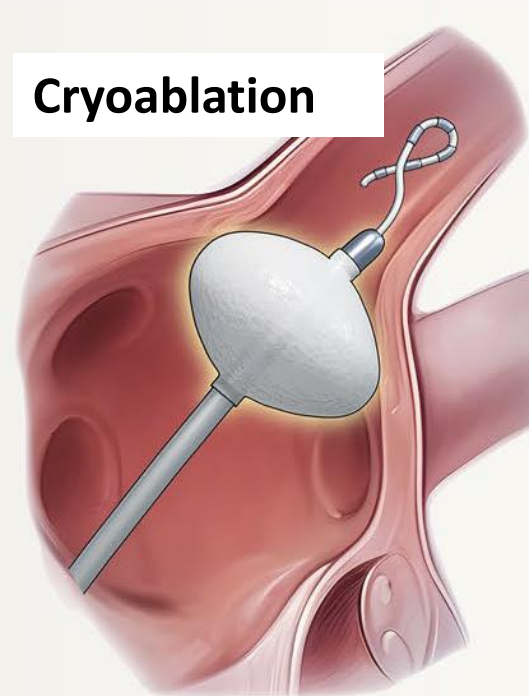
AF Ablation (and cardioversion ...)



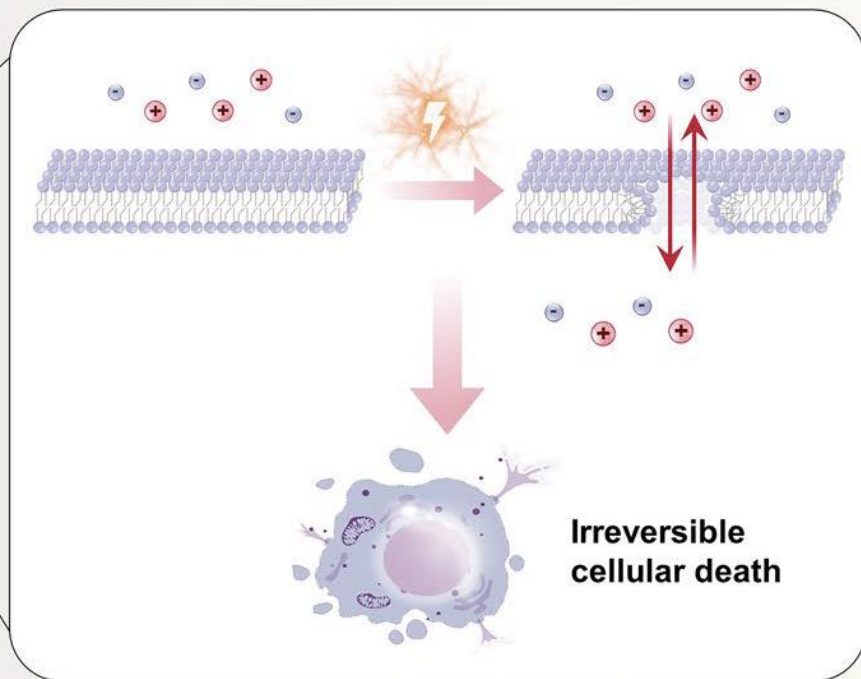
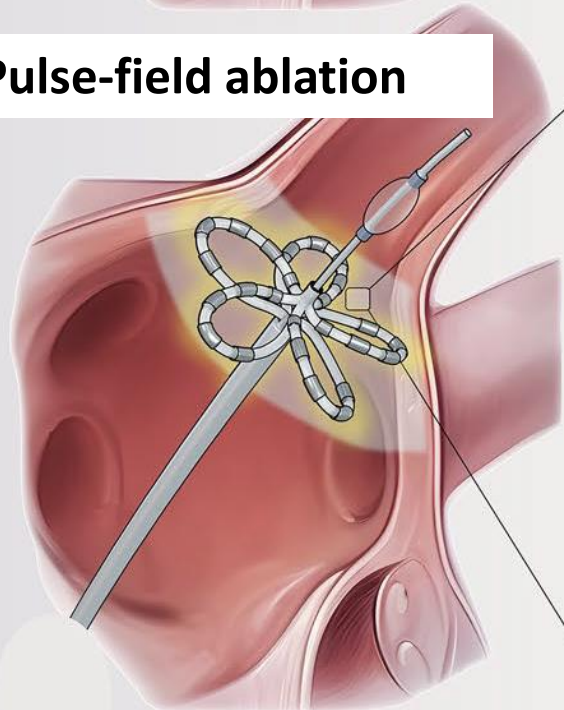
Radiofrequency catheter ablation

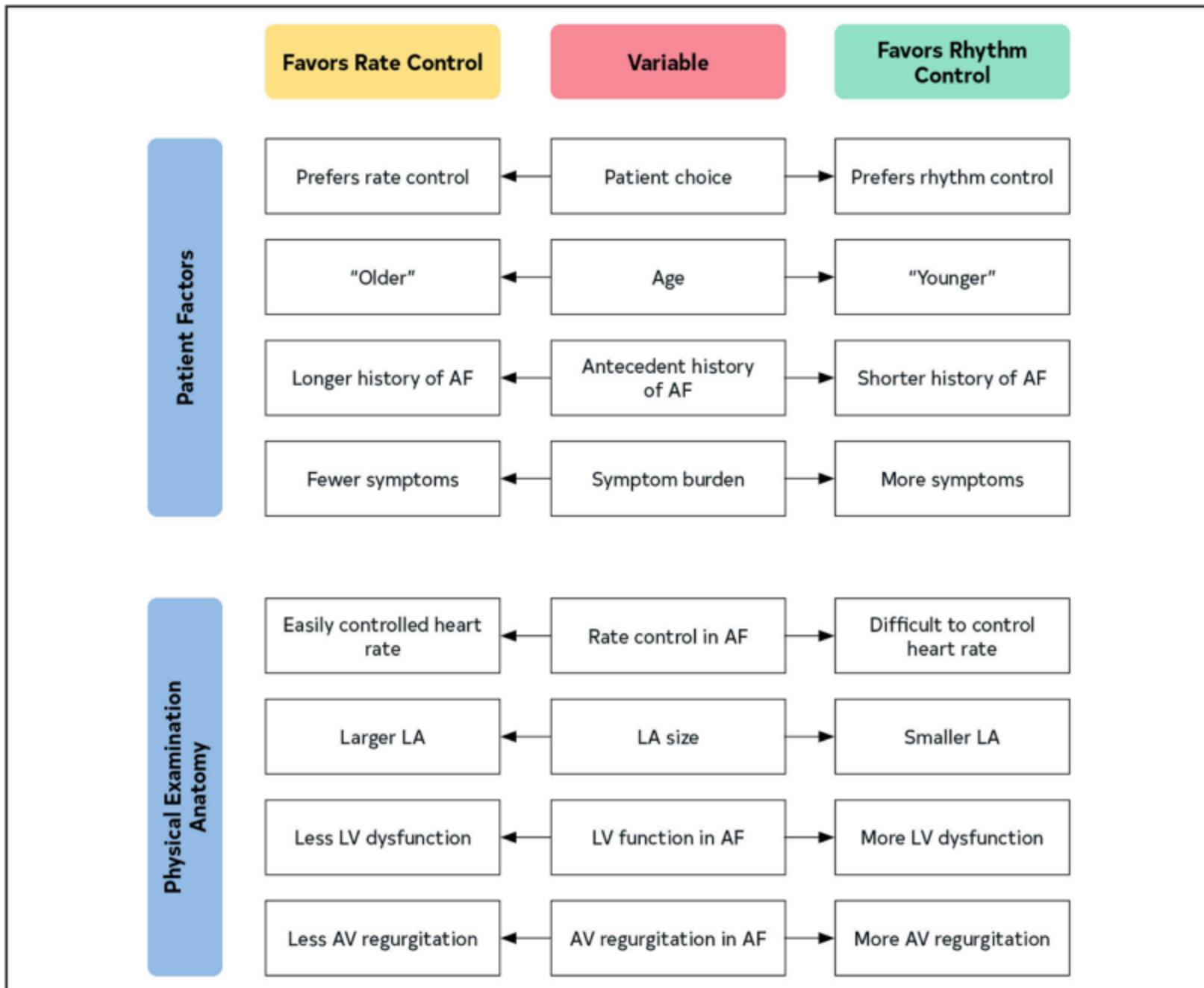


Cryoablation



Pulse-field ablation





No randomised studies for catheter ablation in HFPEF

- Ongoing CABA-HFPEF study
- East AFNET4 trial : early rhythm control with catheter ablation in pts with AF <1 year improves cardiovascular outcomes and symptoms compared to usual care.
- Chinese registry : Catheter ablation associated with improved outcomes in ChadsVasc pts ≤ 4 , but no difference in pts with ChadsVasc > 4



Reduce symptoms by rate and rhythm control

See patient pathways for:

First-diagnosed AF

Paroxysmal AF

Persistent AF

Permanent AF

Consider:

Rate control drugs

Cardioversion

Antiarrhythmic drugs

Catheter ablation

Endoscopic/hybrid ablation

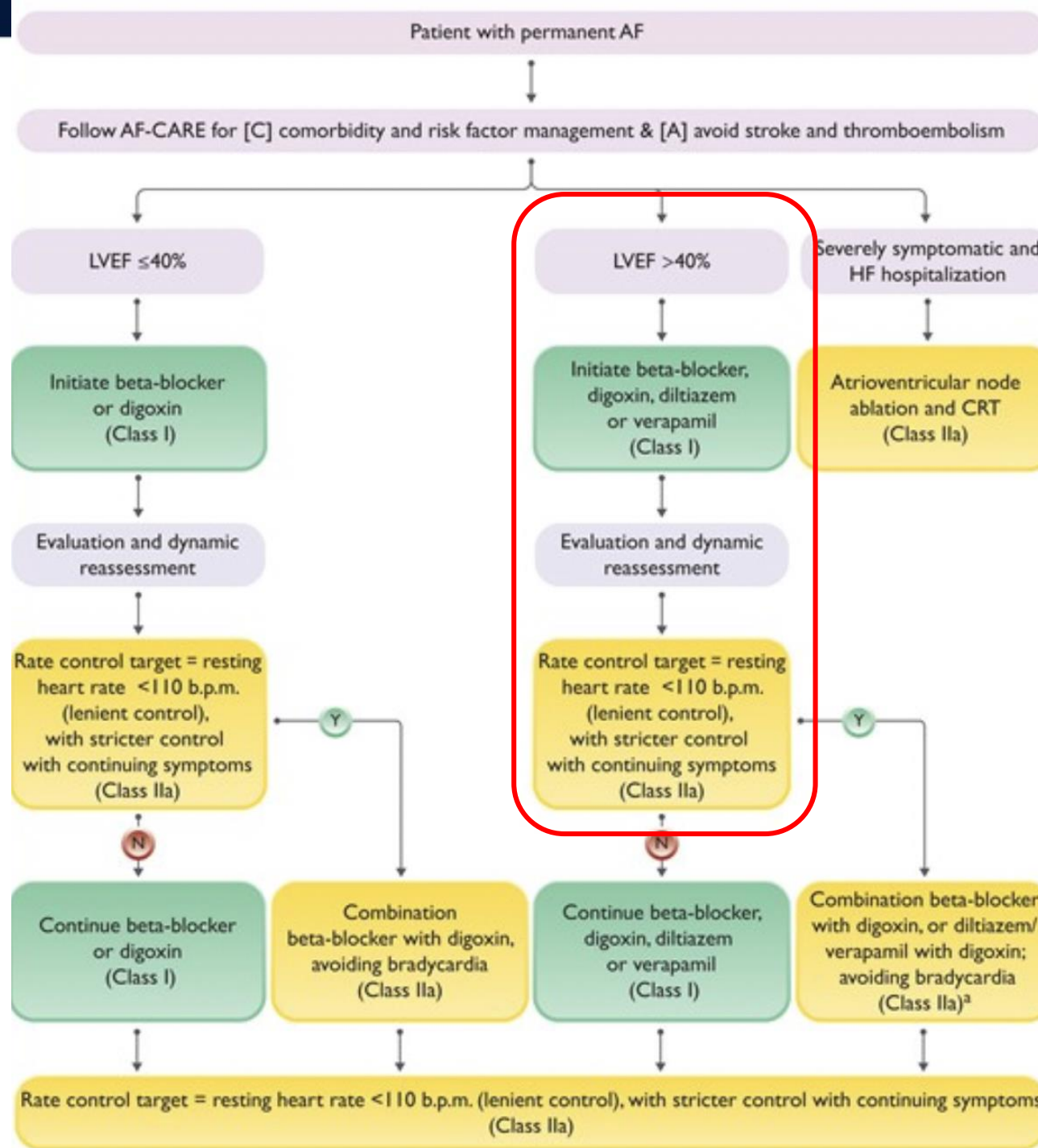
Surgical ablation

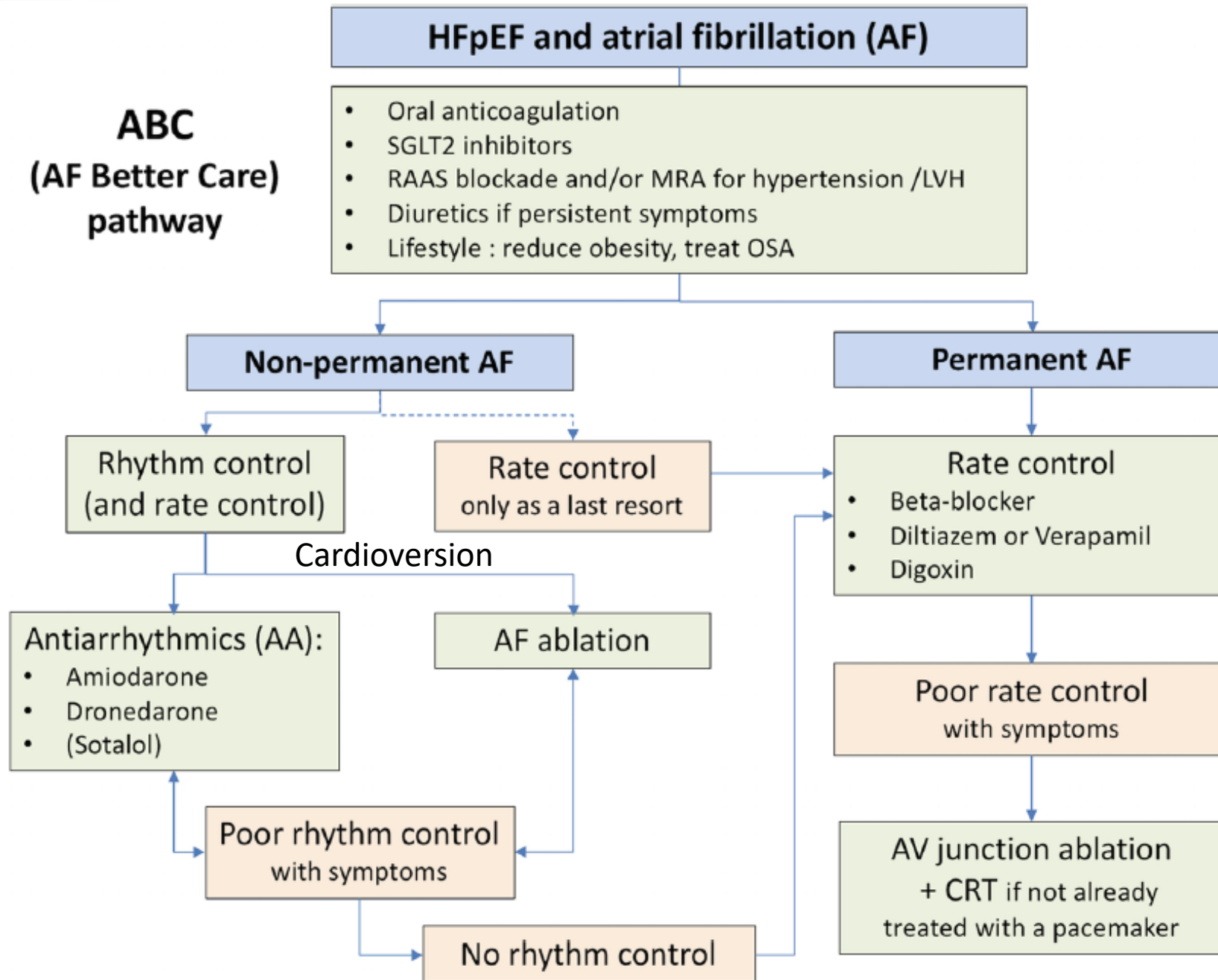
Ablate and pace

Rhythm control favored in patients with paroxysmal AF, low atrial damage

Rate control favored in patients with permanent AF, enlarged LA, many comorbidities

AF → HF : better outcome / HF → AF : worse outcome





Prevalence of Coronary Artery Disease and Coronary Microvascular Dysfunction in Patients With Heart Failure With Preserved Ejection Fraction

Christopher J. Rush, MB, ChB, PhD^{1,2}; Colin Berry, MB, ChB, PhD^{1,2}; Keith G. Oldroyd, MB, ChB, MD^{1,2}; [et al](#)

» [Author Affiliations](#) | [Article Information](#)

JAMA Cardiol. 2021;6(10):1130-1143. doi:10.1001/jamacardio.2021.1825

51% of HFPEF patients with significant coronary artery disease

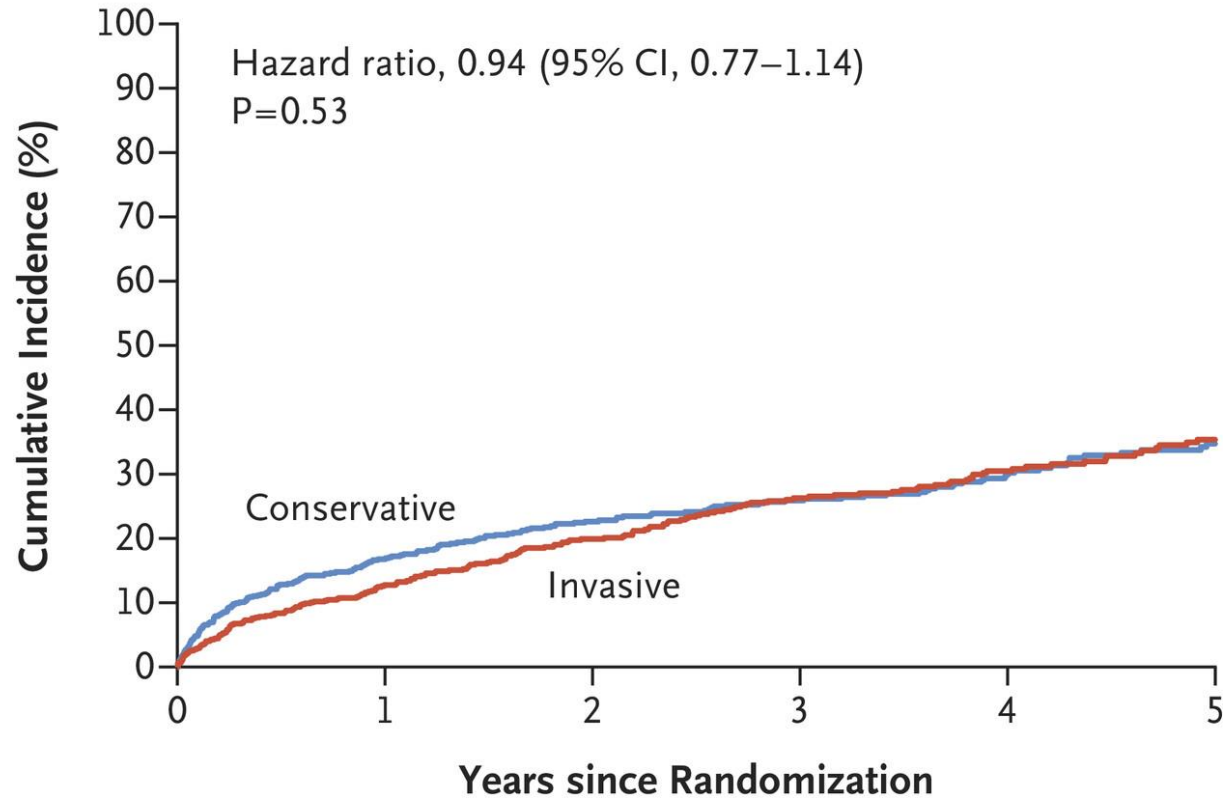
±75% of HFPEF patients have cardiac microvascular disease (CMD)

} 91%
with
CAD
or
CMD

Systematic rule-out of CAD is performed by 37% of doctors, while 35 % look for CAD only if symptoms or high CAD risk .

The SENIOR-RITA trial : prospective, multicenter, open-label, randomized controlled trial.

A Primary Outcome



No. at Risk

Conservative	765	553	417	315	236	89
Invasive	753	570	418	305	232	100

1518 NSTEMI patients >75 yrs randomized to an invasive strategy compared to a conservative treatment strategy.

No difference in primary endpoint between invasive strategy & conservative strategy.

No difference in HF Hospitalization between groups (11%)

No RCTs on invasive strategy in HFPEF patients

Recommendations for management of chronic coronary syndrome patients with chronic heart failure (1)

Recommendations	Class	Level
Managing CCS in heart failure patients		
In HF patients with LVEF >35% and suspected CCS with low or moderate (>5%–50%) pre-test likelihood of obstructive CAD, CCTA or functional imaging is recommended.	I	C
In HF patients with LVEF >35% and suspected CCS with very high (>85%) pre-test likelihood of obstructive CAD, ICA (with FFR, iFR, or QFR when needed) is recommended.	I	C
In patients with HFpEF with persistent angina or equivalent symptoms and normal or non-obstructive epicardial coronary arteries, PET or CMR perfusion or invasive coronary functional testing should be considered to detect or rule out coronary microvascular dysfunction.	IIa	B

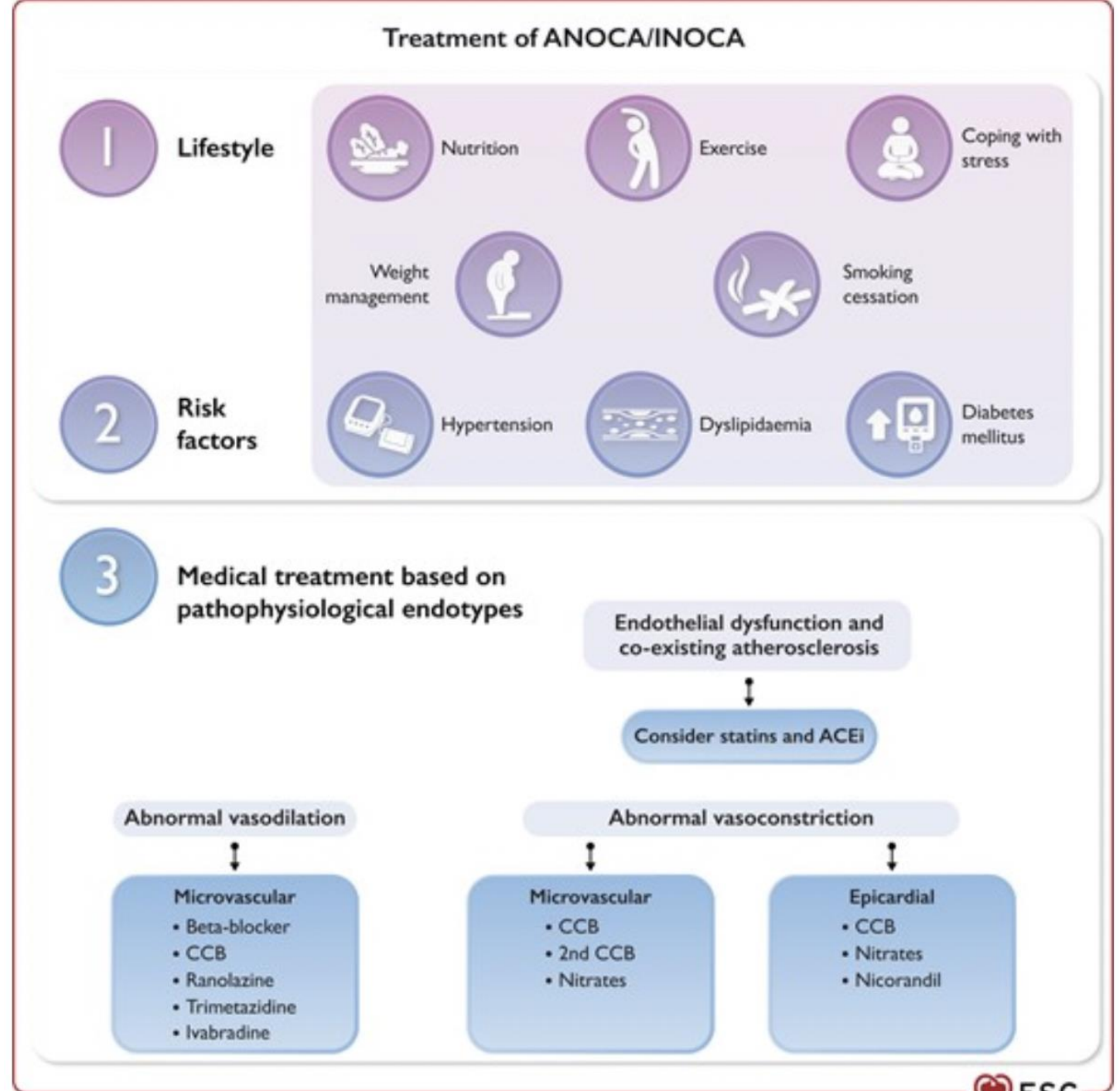
Recommendations for management of chronic coronary syndrome patients with chronic heart failure (2)

Recommendations	Class	Level
Managing heart failure in CCS patients		
It is recommended that CCS patients with HF be enrolled in a multidisciplinary programme to reduce the risk of HF hospitalization and to improve patient outcomes.	I	A
An ACE-I, an MRA, an SGLT2 inhibitor (dapagliflozin or empagliflozin) and a beta-blocker are recommended for CCS patients with HF to reduce the risk of HF hospitalization and death.	I	A
An SGLT2 inhibitor (dapagliflozin or empagliflozin) is recommended in patients with Heart Failure with mildly reduced Ejection Fraction (HFmrEF) to reduce the risk of HF hospitalization or cardiovascular death.	I	A
An ARB is recommended for CCS patients with CCS and HFrEF unable to tolerate an ACE-I or ARNI to reduce the risk of HF hospitalization and cardiovascular death.	I	B
Sacubitril/valsartan is recommended as a replacement for an ACE-I or ARB in CCS patients with HFrEF to reduce the risk of HF hospitalization and of cardiovascular and all-cause death.	I	B
Diuretics are recommended in CCS patients with HF and signs and/or symptoms of congestion to alleviate symptoms, improve exercise capacity, and reduce HF hospitalizations.	I	B

Optimal HF drugs GDMT recommended for all HF patients with CCS IA

Microvascular Disease Management in HFPEF

ESC CCS 2024 Guidelines



Take Home Messages

- AF and CAD (& CMD) are present in more than 2/3 of HFPEF patients
- In both conditions, **global management** of the patient with **HF drugs and health style change** are **first line** treatments → **first treat HF !!!**
- If AF appears as a driver of the HF in case of moderate atrial cardiomyopathy features : rhythm control should be considered (cardioversion, ablation, amiodarone)
- If AF is permanent with significant atrial damage : rate control to be preferred
- In both AF & CAD conditions, personalized management targeted with the patient using tools/therapies at your disposal is the best management