



Moltes novetats en 5 anys. Les noves guies d'insuficiència cardíaca

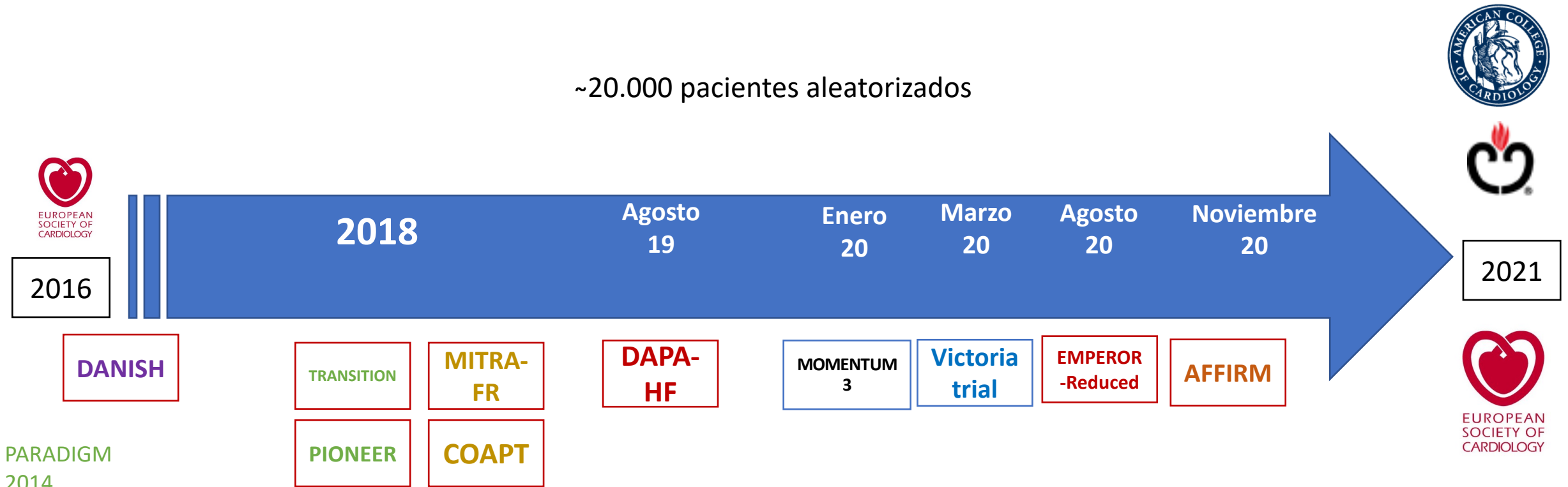
Insuficiència cardíaca amb FEr.

Ana García Álvarez
Hospital Clínic Barcelona

Sessió d'actualització en cardiologia, 4 d'octubre 2021

Publicaciones con impacto en las nuevas guías

~20.000 pacientes aleatorizados



PARADIGM
2014

Clases de recomendación y niveles de evidencia

Classes of recommendations

	Definition	Wording to use
Class I	Evidence and/or general agreement that a given treatment or procedure is beneficial, useful, effective.	Is recommended or is indicated
Class II	Conflicting evidence and/or a divergence of opinion about the usefulness/ efficacy of the given treatment or procedure.	
Class IIa	Weight of evidence/opinion is in favour of usefulness/efficacy.	Should be considered
Class IIb	Usefulness/efficacy is less well established by evidence/opinion.	May be considered
Class III	Evidence or general agreement that the given treatment or procedure is not useful/effective, and in some cases may be harmful.	Is not recommended

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Levels of evidence

Level of evidence A	Data derived from multiple randomized clinical trials or meta-analyses.
Level of evidence B	Data derived from a single randomized clinical trial or large non-randomized studies.
Level of evidence C	Consensus of opinion of the experts and/or small studies, retrospective studies, registries.

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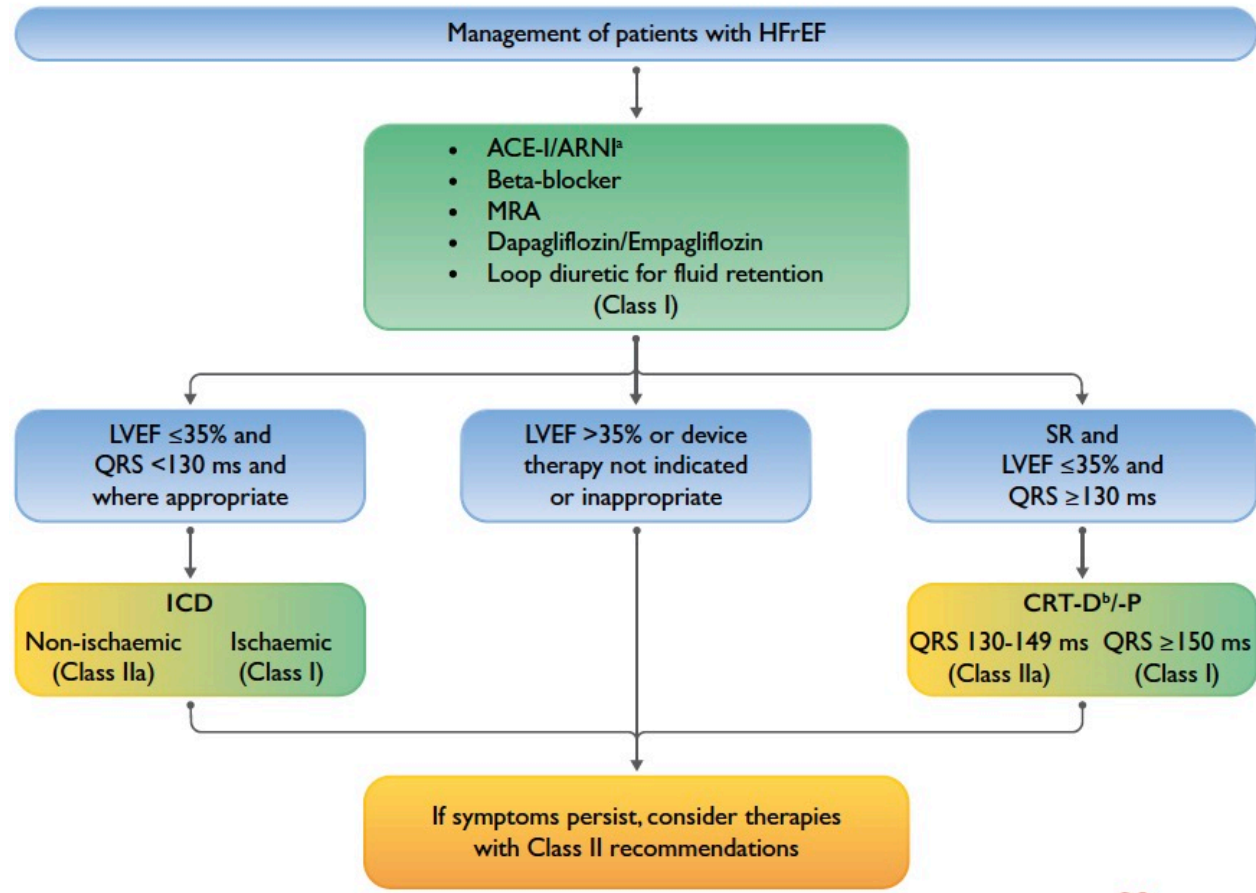
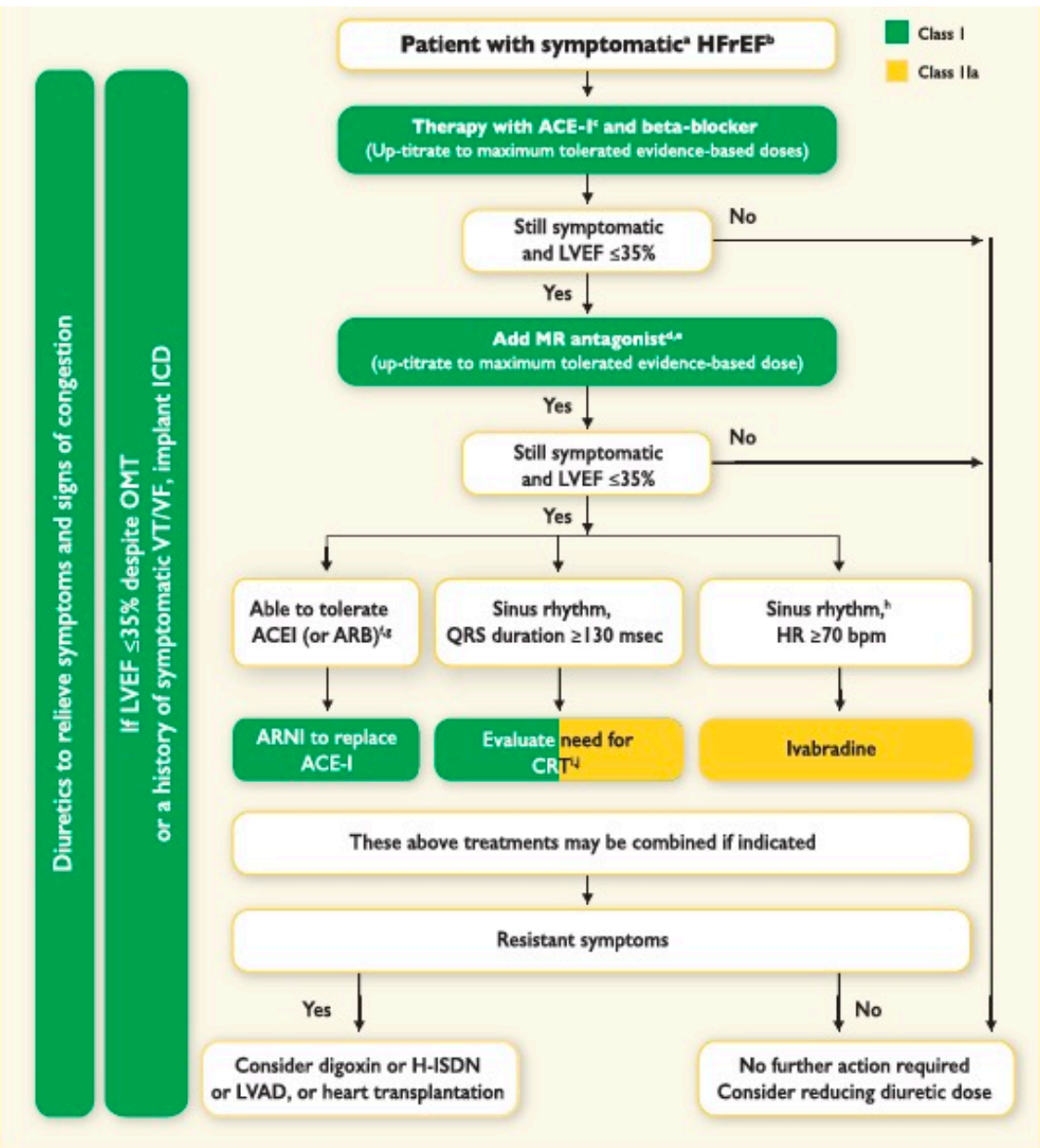
Definición

Type of HF		HFrEF	HFmrEF	HFpEF
CRITERIA	1	Symptoms ± Signs ^a	Symptoms ± Signs ^a	Symptoms ± Signs ^a
	2	LVEF ≤40%	LVEF 41–49% ^b	LVEF ≥50%
	3	—	—	Objective evidence of cardiac structural and/or functional abnormalities consistent with the presence of LV diastolic dysfunction/raised LV filling pressures, including raised natriuretic peptides ^c

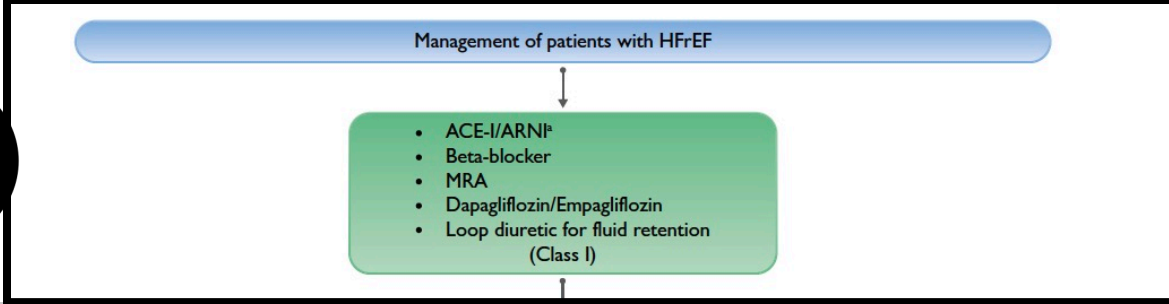
✓ Sin cambios respecto a las guías ESC 2026

2016

2021



1



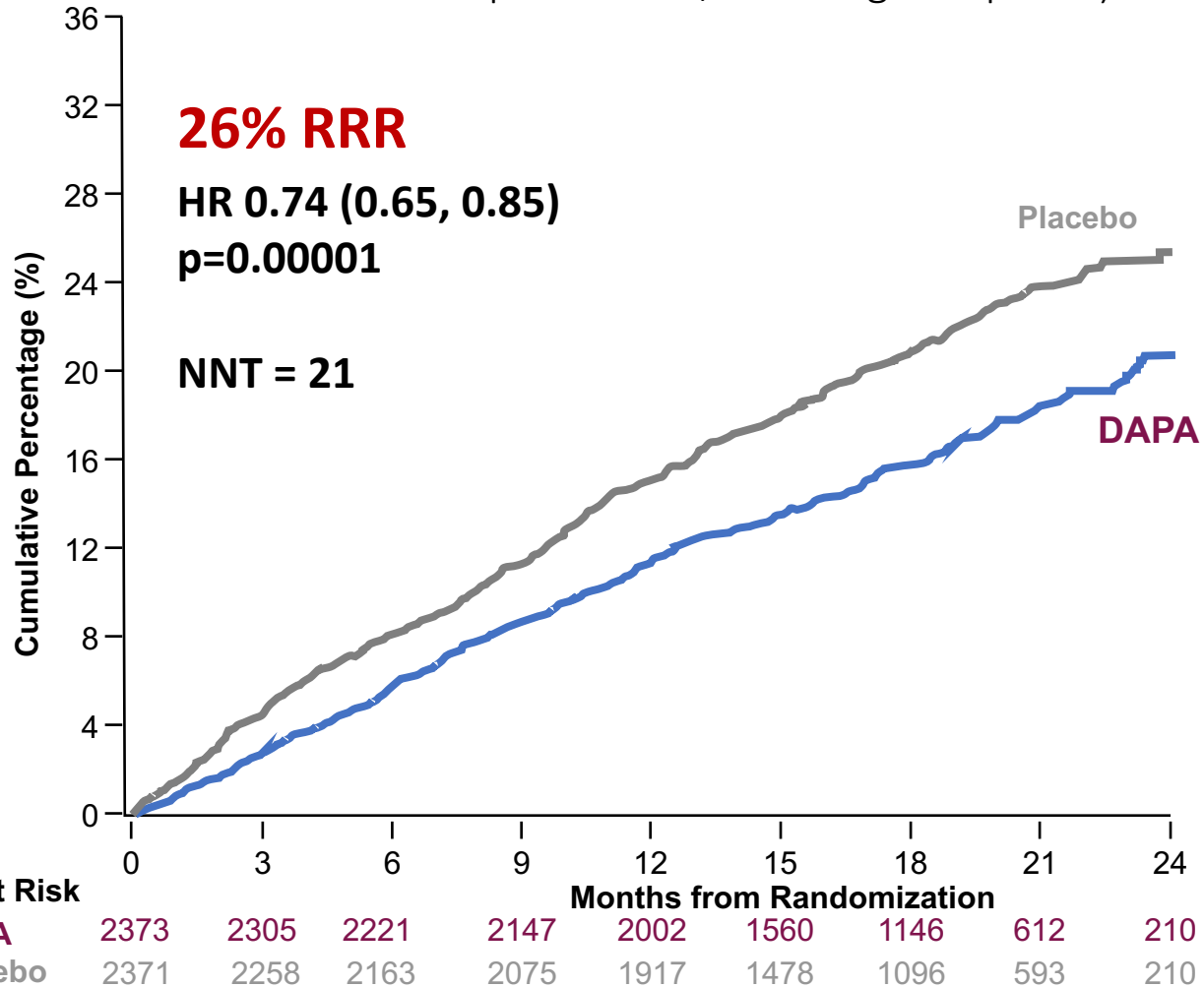
Recommendations	Class ^a	Level ^b
An ACE-I is recommended for patients with HFrEF to reduce the risk of HF hospitalization and death. ^{110–113}	I	A
A beta-blocker is recommended for patients with stable HFrEF to reduce the risk of HF hospitalization and death. ^{114–120}	I	A
An MRA is recommended for patients with HFrEF to reduce the risk of HF hospitalization and death. ^{121,122}	I	A
Dapagliflozin or empagliflozin are recommended for patients with HFrEF to reduce the risk of HF hospitalization and death. ^{108,109}	I	A
Sacubitril/valsartan is recommended as a replacement for an ACE-I in patients with HFrEF to reduce the risk of HF hospitalization and death. ¹⁰⁵	I	B



Dapagliflozin in Patients with Heart Failure and Reduced Ejection Fraction

J.J.V. McMurray, S.D. Solomon, S.E. Inzucchi, L. Køber, M.N. Kosiborod, F.A. Martinez, P. Ponikowski,

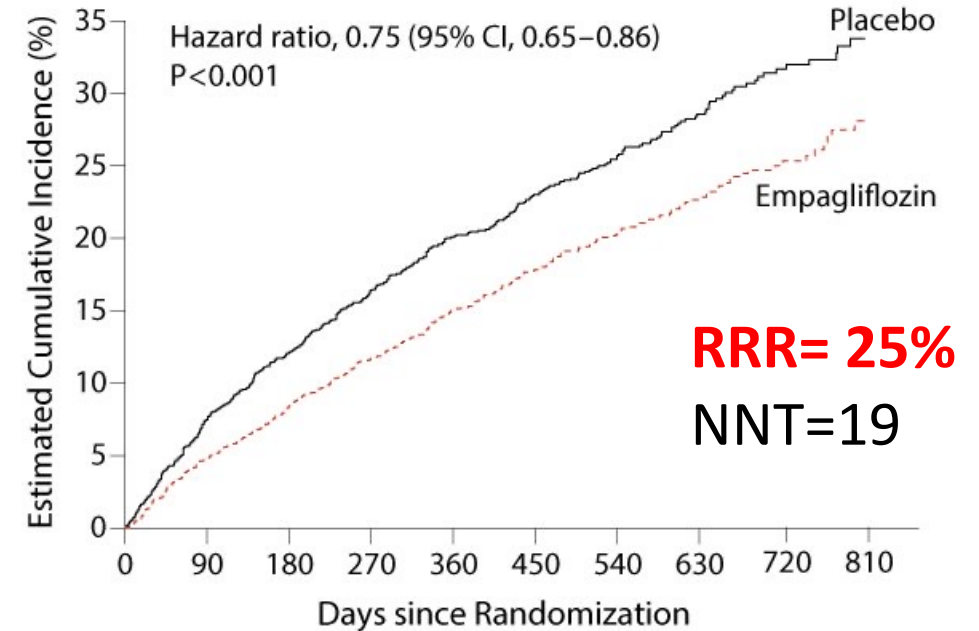
DAPA-HF: Muerte CV u Hospitalización/visita urgente por IC).



Cardiovascular and Renal Outcomes with Empagliflozin in Heart Failure

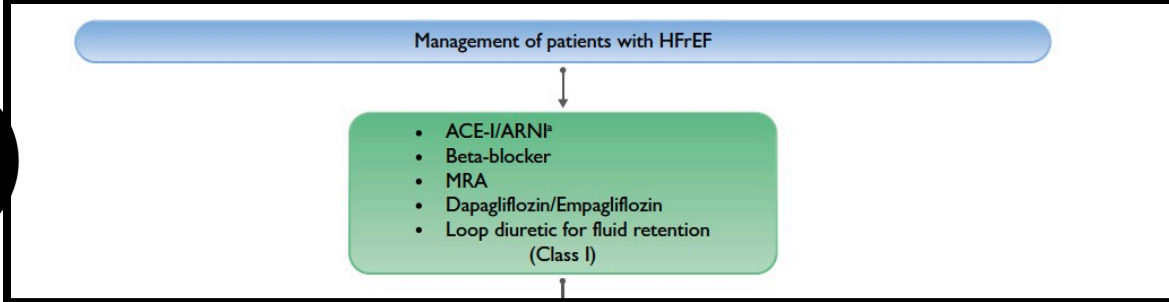
M. Packer, S.D. Anker, J. Butler, G. Filippatos, S.J. Pocock, P. Carson, J. Januzzi,

EMPEROR-Reduced: Muerte CV u Hospitalización por IC



	Empaglifozina (n=1863)		Placebo (n=1867)		HR
	Nº de eventos (%)	Eventos/100 pac/año	Nº de eventos (%)	Eventos/100 pac/año	
Hosp por IC	246 (13.2%)	10.7	342 (18.3%)	15.5	0.69 (0.59-0.81)
Muerte CV	187 (10.0%)	7.6	202 (10.8%)	8.1	0.92 (0.75-1.12)

1



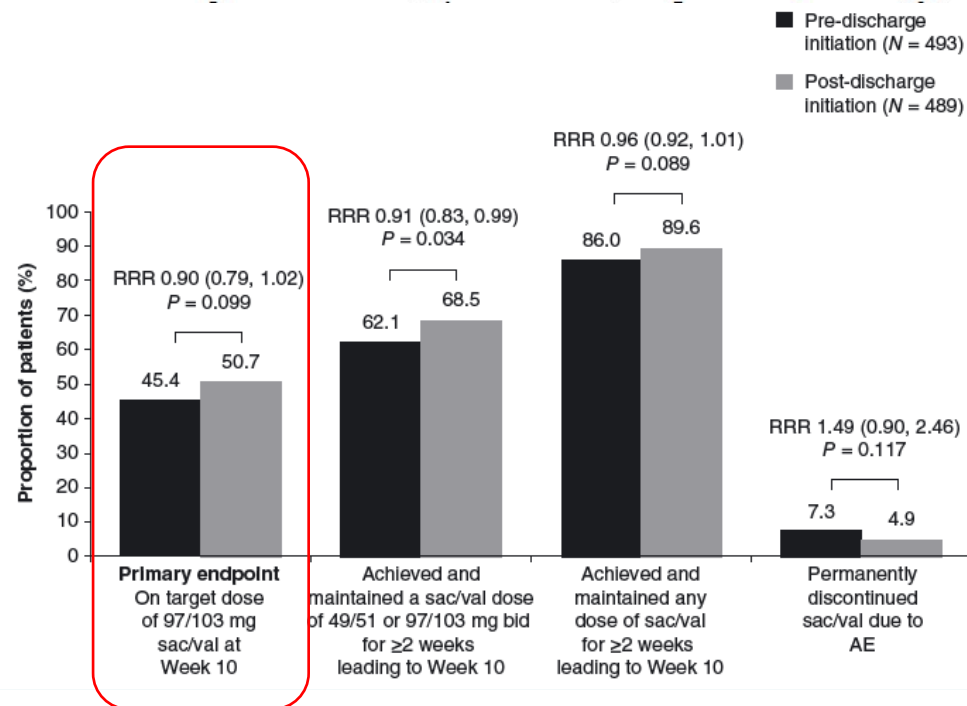
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Dapagliflozin or empagliflozin are recommended for patients with HFrEF to reduce the risk of HF hospitalization and death. ^{108,109}	I	A
Sacubitril/valsartan is recommended as a replacement for an ACE-I in patients with HFrEF to reduce the risk of HF hospitalization and death. ¹⁰⁵	I	B



talizations by 42% compared to enalapril.^{106,107,131} As such, initiation of sacubitril/valsartan in ACE-I naive (i.e. *de novo*) patients with HFrEF may be considered (class of recommendation IIb, level of evidence B). Patients being commenced on sacubitril/valsartan should have an

Initiation of sacubitril/valsartan in haemodynamically stabilised heart failure patients in hospital or early after discharge: primary results of the randomised TRANSITION study

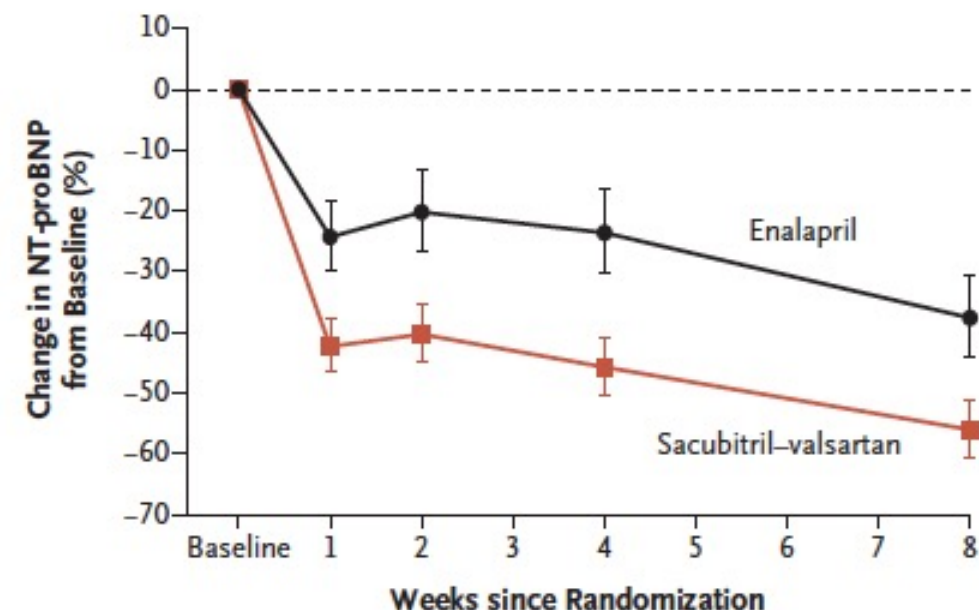
Rolf Wachter^{1*}, Michele Senni², Jan Belohlavek³, Ewa Straburzynska-Migaj⁴,



- ✓ 286 pacientes (29%) eran IC de novo.
- ✓ 242 pacientes (24%) eran naïve.

Angiotensin–Neprilysin Inhibition in Acute Decompensated Heart Failure

Eric J. Velazquez, M.D., David A. Morrow, M.D., M.P.H.,
PIONEER. New Engl J Med 2018.

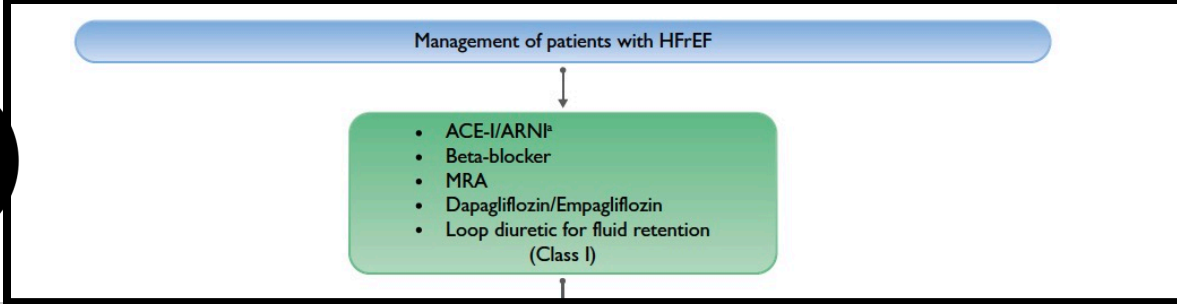


No. at Risk

	Baseline	1	2	4	8
Enalapril	394	359	351	350	348
Sacubitril-valsartan	397	355	363	365	349

- ✓ 303 pacientes (34.4%) eran IC de novo.
- ✓ 459 pacientes (52.1%) eran naïve.

1



Recommendations	Class ^a	Level ^b
An ACE-I is recommended for patients with HFrEF to reduce the risk of HF hospitalization and death. ^{110–113}	I	A
A beta-blocker is recommended for patients with stable HFrEF to reduce the risk of HF hospitalization and death. ^{114–120}	I	A
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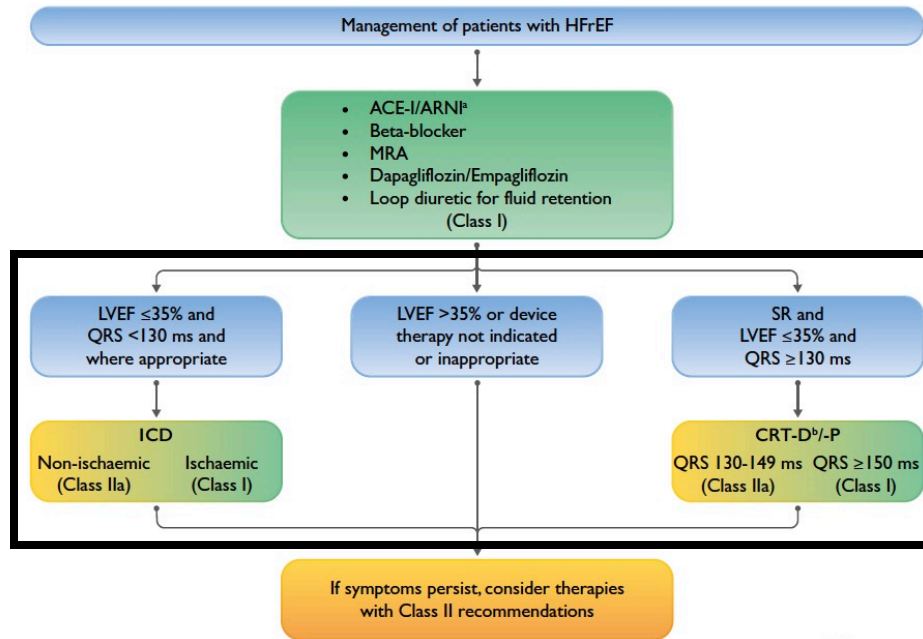


talizations by 42% compared to enalapril.^{106,107,131} As such, initiation of sacubitril/valsartan in ACE-I naive (i.e. *de novo*) patients with HFrEF may be considered (class of recommendation IIb, level of evidence B). Patients being commenced on sacubitril/valsartan should have an

Loop diuretics		
Diuretics are recommended in patients with HFrEF with signs and/or symptoms of congestion to alleviate HF symptoms, improve exercise capacity, and reduce HF hospitalizations. ¹³⁷	I	C
ARB		
An ARB ^c is recommended to reduce the risk of HF hospitalization and CV death in symptomatic patients unable to tolerate an ACE-I or ARNI (patients should also receive a beta-blocker and an MRA). ¹³⁸	I	B



2



Secondary prevention

An ICD is recommended to reduce the risk of sudden death and all-cause mortality in patients who have recovered from a ventricular arrhythmia causing haemodynamic instability, and who are expected to survive for >1 year with good functional status, in the absence of reversible causes or unless the ventricular arrhythmia has occurred <48 h after a MI.^{162–164}

I	A
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Primary prevention

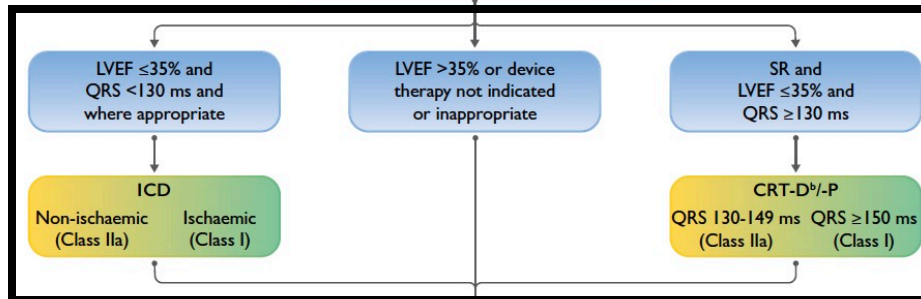
An ICD is recommended to reduce the risk of sudden death and all-cause mortality in patients with symptomatic HF (NYHA class II–III) of an ischaemic aetiology (unless they have had a MI in the prior 40 days—see below), and an LVEF ≤35% despite ≥3 months of OMT, provided they are expected to survive substantially longer than 1 year with good functional status.^{161,165}

I	A
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Management of patients with HFrEF

- ACE-I/ARNI
- Beta-blocker
- MRA
- Dapagliflozin/Empagliflozin
- Loop diuretic for fluid retention (Class I)



If symptoms persist, consider therapies with Class II recommendations

An ICD should be considered to reduce the risk of sudden death and all-cause mortality in patients with symptomatic HF (NYHA class II–III) of a non-ischaemic aetiology, and an LVEF $\leq 35\%$ despite ≥ 3 months of OMT, provided they are expected to survive substantially longer than 1 year with good functional status.^{161,166,167}



IIa

A

Patients should be carefully evaluated by an experienced cardiologist before generator replacement, because management goals, the patient's needs and clinical status may have changed.^{168–172}

IIa

B

2

The NEW ENGLAND JOURNAL of MEDICINE

ESTABLISHED IN 1812

SEPTEMBER 29, 2016

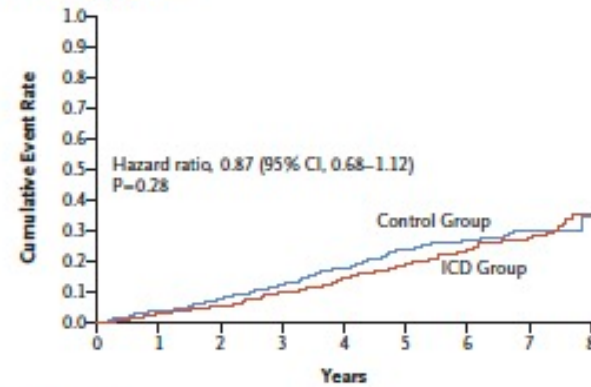
VOL. 375 NO. 13

Defibrillator Implantation in Patients with Nonischemic Systolic Heart Failure

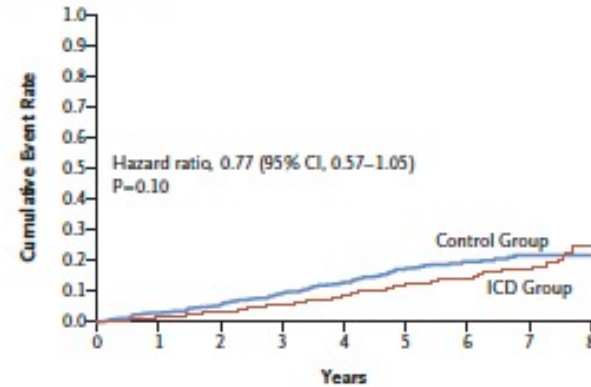
Lars Køber, M.D., D.M.Sc., Jens J. Thune, M.D., Ph.D., Jens C. Nielsen, M.D., D.M.Sc., Jens Haarlo, M.D., D.M.Sc.,

DANISH Trial

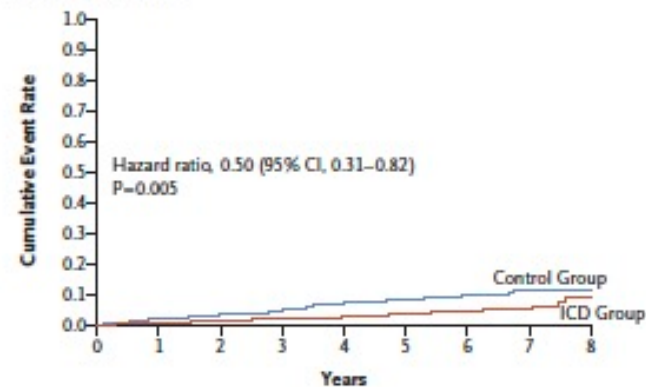
A Death from Any Cause



B Cardiovascular Death



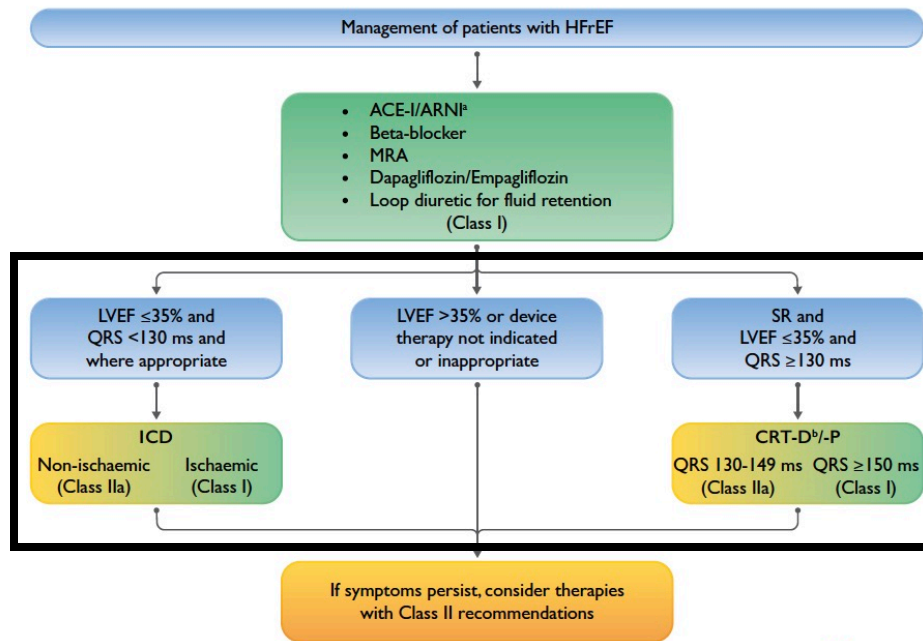
C Sudden Cardiac Death



No. at Risk

Control Group	560	540	517	438	344	248	169	88	12
ICD Group	556	540	526	451	358	272	186	107	17

2



CRT is recommended for symptomatic patients with HF in SR with a QRS duration ≥ 150 ms and LBBB QRS morphology and with LVEF $\leq 35\%$ despite OMT in order to improve symptoms and reduce morbidity and mortality.^{205–215}

I

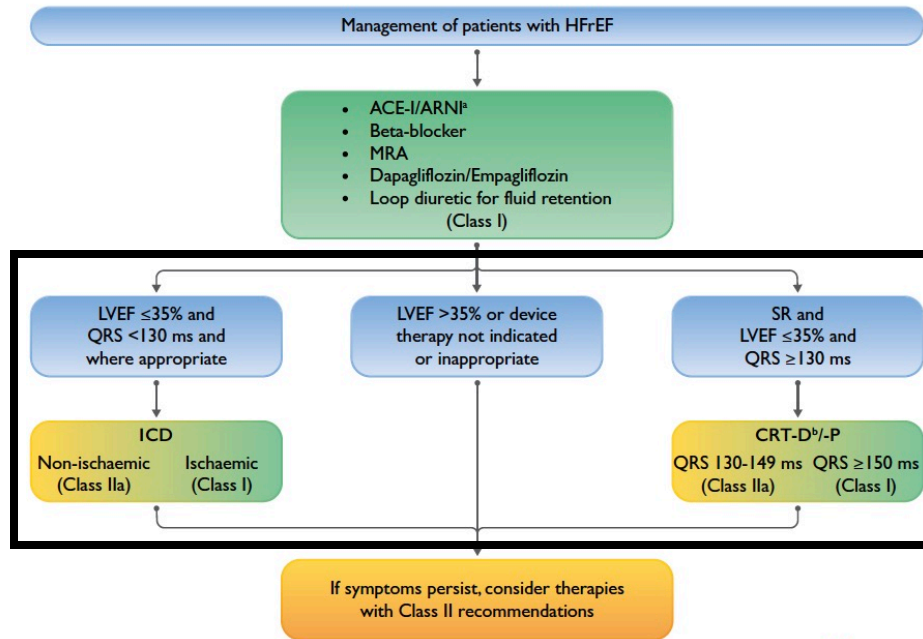
A

CRT rather than RV pacing is recommended for patients with HFrEF regardless of NYHA class or QRS width who have an indication for ventricular pacing for high degree AV block in order to reduce morbidity. This includes patients with AF.^{216–219}

I

A

2



CRT should be considered for symptomatic patients with HF in SR with a QRS duration ≥ 150 ms and non-LBBB QRS morphology and with LVEF $\leq 35\%$ despite OMT in order to improve symptoms and reduce morbidity and mortality.^{205–215}

IIa

B

CRT should be considered for symptomatic patients with HF in SR with a QRS duration of 130–149 ms and LBBB QRS morphology and with LVEF $\leq 35\%$ despite OMT in order to improve symptoms and reduce morbidity and mortality.^{211,220}



IIa

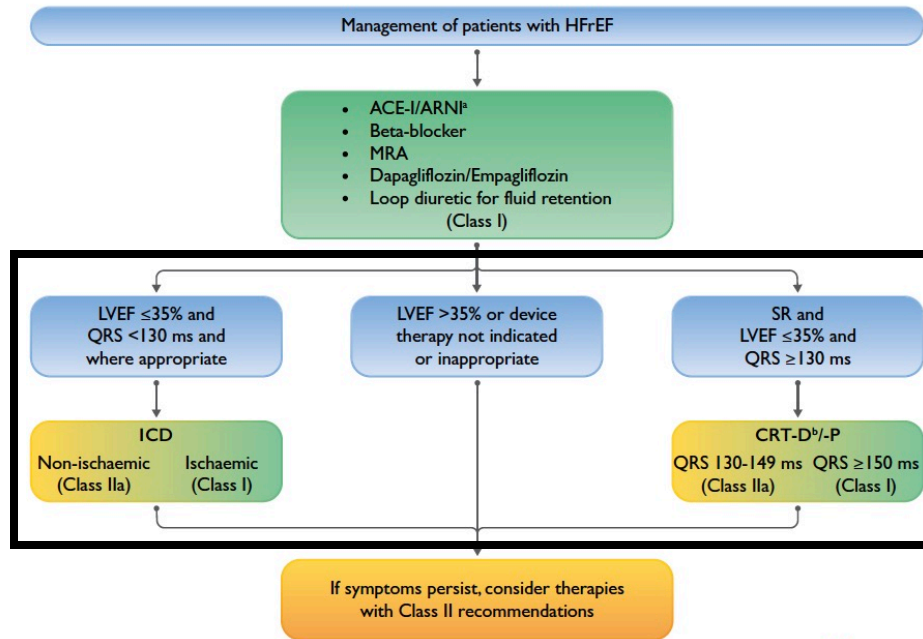
B

Patients with an LVEF $\leq 35\%$ who have received a conventional pacemaker or an ICD and subsequently develop worsening HF despite OMT and who have a significant proportion of RV pacing should be considered for 'upgrade' to CRT.²²¹

IIa

B

2



CRT may be considered for symptomatic patients with HF in SR with a QRS duration of 130–149 ms and non-LBBB QRS morphology and with LVEF $\leq 35\%$ despite OMT in order to improve symptoms and reduce morbidity and mortality.^{208,213}

IIb

B

CRT is not recommended in patients with a QRS duration <130 ms who do not have an indication for pacing due to high degree AV block.^{222–224}

III

A

Management of patients with HFrEF

- ACE-I/ARNI^a
- Beta-blocker
- MRA
- Dapagliflozin/Empagliflozin
- Loop diuretic for fluid retention (Class I)

LVEF ≤35% and QRS <130 ms and where appropriate

LVEF >35% or device therapy not indicated or inappropriate

SR and LVEF ≤35% and QRS ≥130 ms

ICD
Non-Ischaemic (Class IIa) Ischaemic (Class I)

CRT-D^b/-P
QRS 130-149 ms (Class IIa) QRS ≥150 ms (Class I)

If symptoms persist, consider therapies with Class II recommendations

3

I_f channel inhibitor

Ivabradine should be considered in symptomatic patients with LVEF ≤35%, in SR and a resting heart rate ≥70 b.p.m. despite treatment with an evidence-based dose of beta-blocker (or maximum tolerated dose below that), ACE-I/(or ARNI), and an MRA, to reduce the risk of HF hospitalization and CV death.¹³⁹

IIa

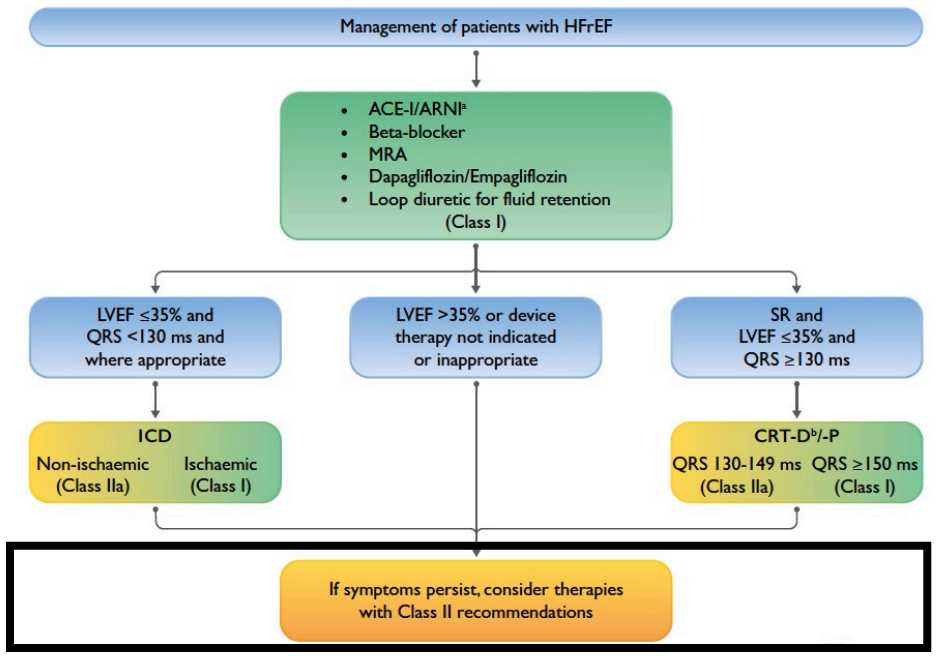
B

Ivabradine should be considered in symptomatic patients with LVEF ≤35%, in SR and a resting heart rate ≥70 b.p.m. who are unable to tolerate or have contraindications for a beta-blocker to reduce the risk of HF hospitalization and CV death. Patients should also receive an ACE-I (or ARNI) and an MRA.¹⁴⁰

IIa

C

3



Digoxin

Digoxin may be considered in patients with symptomatic HFrEF in sinus rhythm despite treatment with an ACE-I (or ARNI), a beta-blocker and an MRA, to reduce the risk of hospitalization (both all-cause and HF hospitalizations).¹⁴⁴

IIb	B
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Soluble guanylate cyclase receptor stimulator

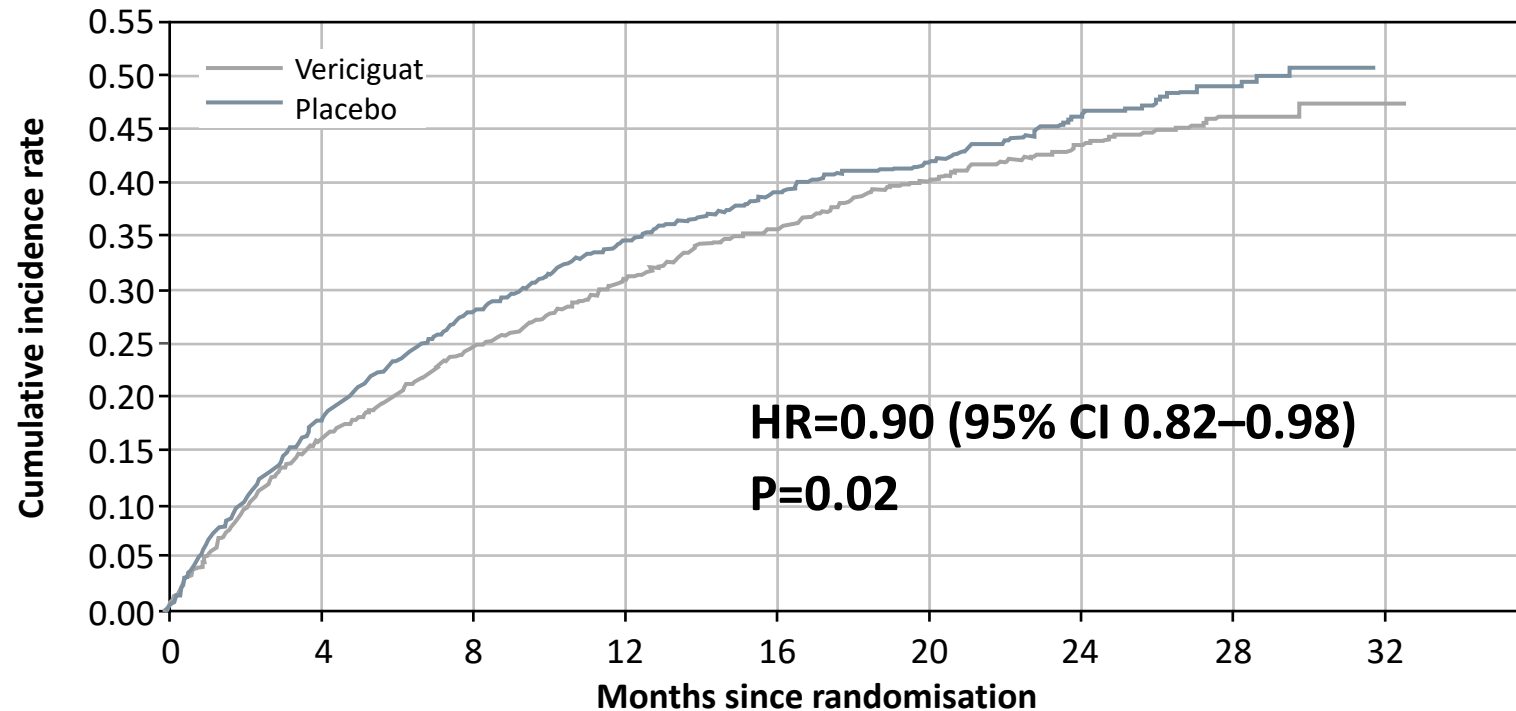
Vericiguat may be considered in patients in NYHA class II–IV who have had worsening HF despite treatment with an ACE-I (or ARNI), a beta-blocker and an MRA to reduce the risk of CV mortality or HF hospitalization.¹⁴¹

IIb	B
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Vericiguat in Patients with Heart Failure and Reduced Ejection Fraction

Paul W. Armstrong, M.D., Burkert Pieske, M.D., Kevin J. Anstrom, Ph.D., Justin Ezekowitz, M.B., B.Ch.,

VICTORIA Trial



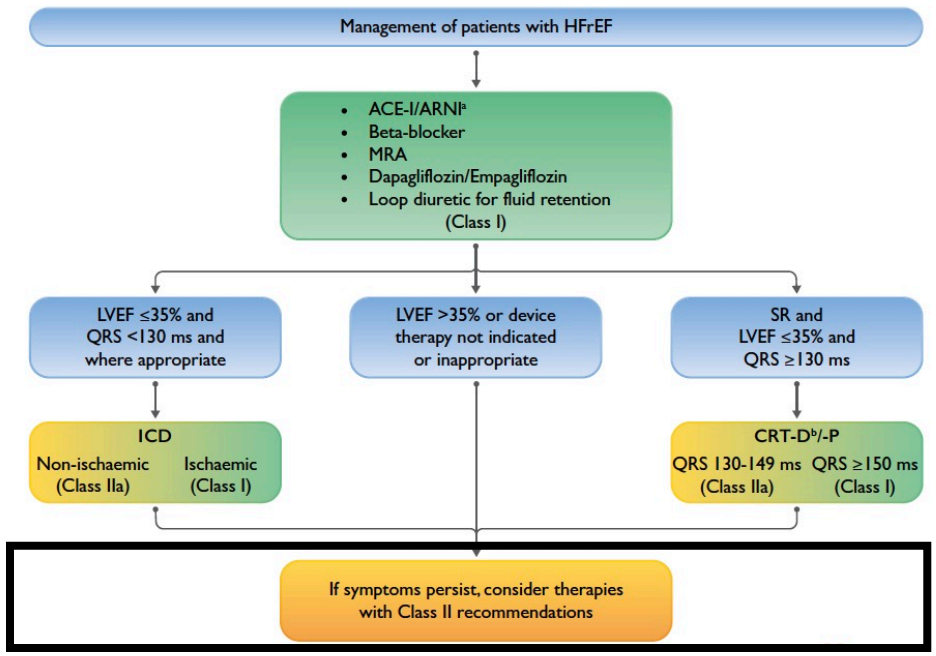
Tasa de eventos
33.6% x100 pac-año
vs. 37.8% con
placebo.

RRR=10%

RRA= 4.2% por año.

NNT= 24.

3



Hydralazine and isosorbide dinitrate

Hydralazine and isosorbide dinitrate should be considered in self-identified black patients with LVEF ≤35% or with an LVEF <45% combined with a dilated left ventricle in NYHA class III–IV despite treatment with an ACE-I (or ARNI), a beta-blocker and an MRA to reduce the risk of HF hospitalization and death.¹⁴²

IIa

B

Hydralazine and isosorbide dinitrate may be considered in patients with symptomatic HFrEF who cannot tolerate any of an ACE-I, an ARB, or ARNI (or they are contraindicated) to reduce the risk of death.¹⁴³

IIb

B

1

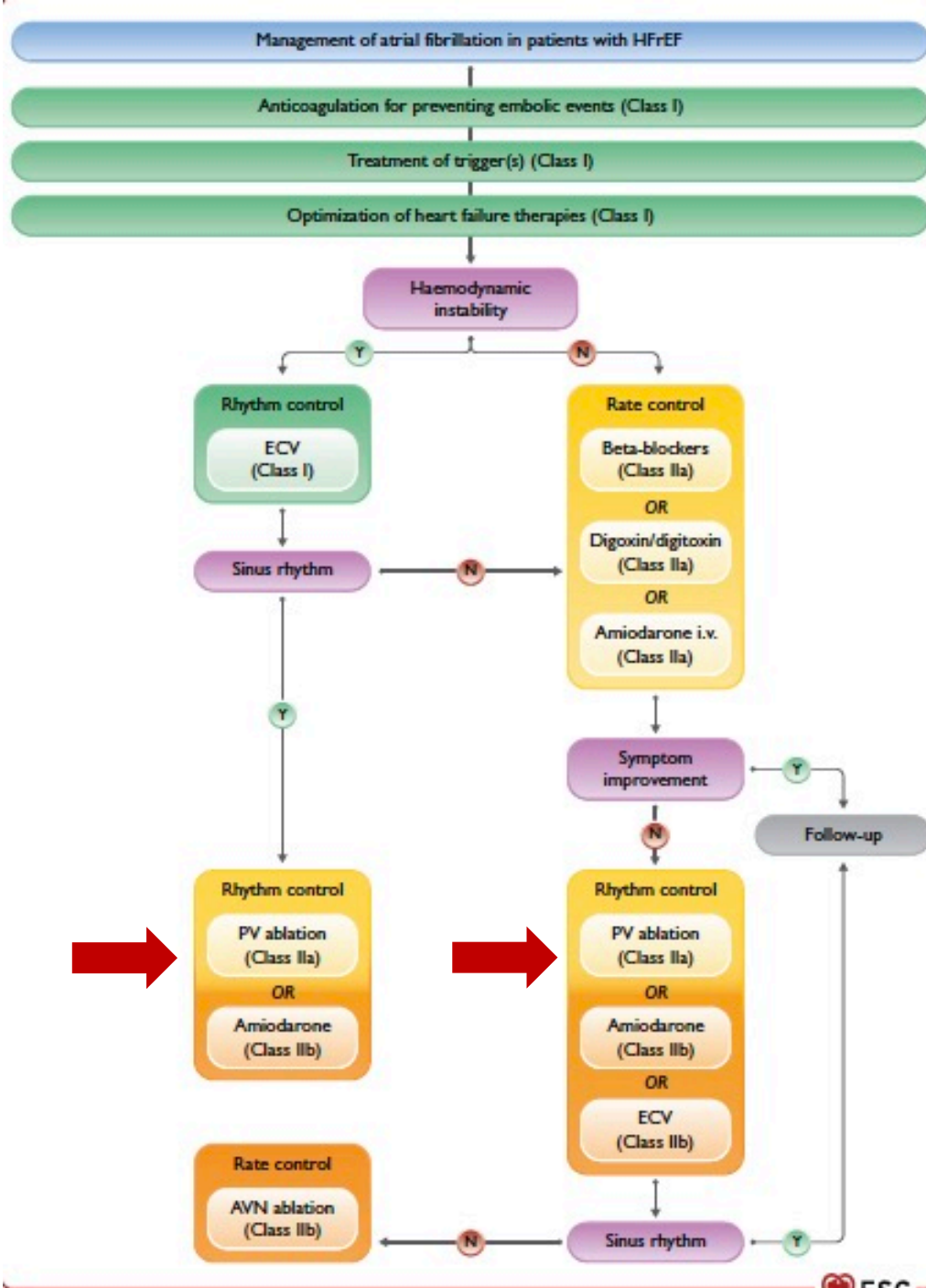
To reduce mortality - for all patients			
ACE-I/ARNI	BB	MRA	SGLT2i

2

To reduce HF hospitalization/mortality - for selected patients	
Volume overload	
Diuretics	
SR with LBBB ≥ 150 ms	SR with LBBB 130–149 ms or non LBBB ≥ 150 ms
CRT-P/D	CRT-P/D
Ischaemic aetiology	Non-ischaemic aetiology
ICD	ICD

3

Atrial fibrillation	Atrial fibrillation	Coronary artery disease	Iron deficiency
Anticoagulation	Digoxin PVI	CABG	Ferric carboxymaltose
Aortic stenosis	Mitral regurgitation	Heart rate SR > 70 bpm	Black Race
SAVR/TAVI	TEE MV Repair	Ivabradine	Hydralazine/ISDN
			ACE-I/ARNI intolerance
			ARB



DOACs are recommended in preference to VKAs in patients with HF, except in those with moderate or severe mitral stenosis or mechanical prosthetic heart valves.^{528,558}

I	A
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AF catheter ablation

In cases of a clear association between paroxysmal or persistent AF and worsening of HF symptoms, which persist despite MT, catheter ablation should be considered for the prevention or treatment of AF.^{552–554,557}

IIa	B
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1

To reduce mortality - for all patients			
ACE-I/ARNI	BB	MRA	SGLT2i

2

To reduce HF hospitalization/mortality - for selected patients	
Volume overload Diuretics	
SR with LBBB ≥ 150 ms CRT-P/D	SR with LBBB 130–149 ms or non LBBB ≥ 150 ms CRT-P/D
Ischaemic aetiology ICD	Non-ischaemic aetiology ICD

3

Atrial fibrillation Anticoagulation	Atrial fibrillation Digoxin PVI	Coronary artery disease CABG	Iron deficiency Ferric carboxymaltose
Aortic stenosis SAVR/TAVI	Mitral regurgitation TEE MV Repair	Heart rate SR > 70 bpm Ivabradine	Black Race Hydralazine/ISDN
		ACE-I/ARNI intolerance ARB	



It is recommended that all patients with HF be periodically screened for anaemia and iron deficiency with a full blood count, serum ferritin concentration, and TSAT.

I

C

Intravenous iron supplementation with ferric carboxymaltose should be considered in symptomatic patients with LVEF <45% and iron deficiency, defined as serum ferritin <100 ng/mL or serum ferritin 100–299 ng/mL with TSAT <20%, to alleviate HF symptoms, improve exercise capacity and QOL.^{720,722,724}

IIa

A

Intravenous iron supplementation with ferric carboxymaltose should be considered in symptomatic HF patients recently hospitalized for HF and with LVEF <50% and iron deficiency, defined

IIa

B

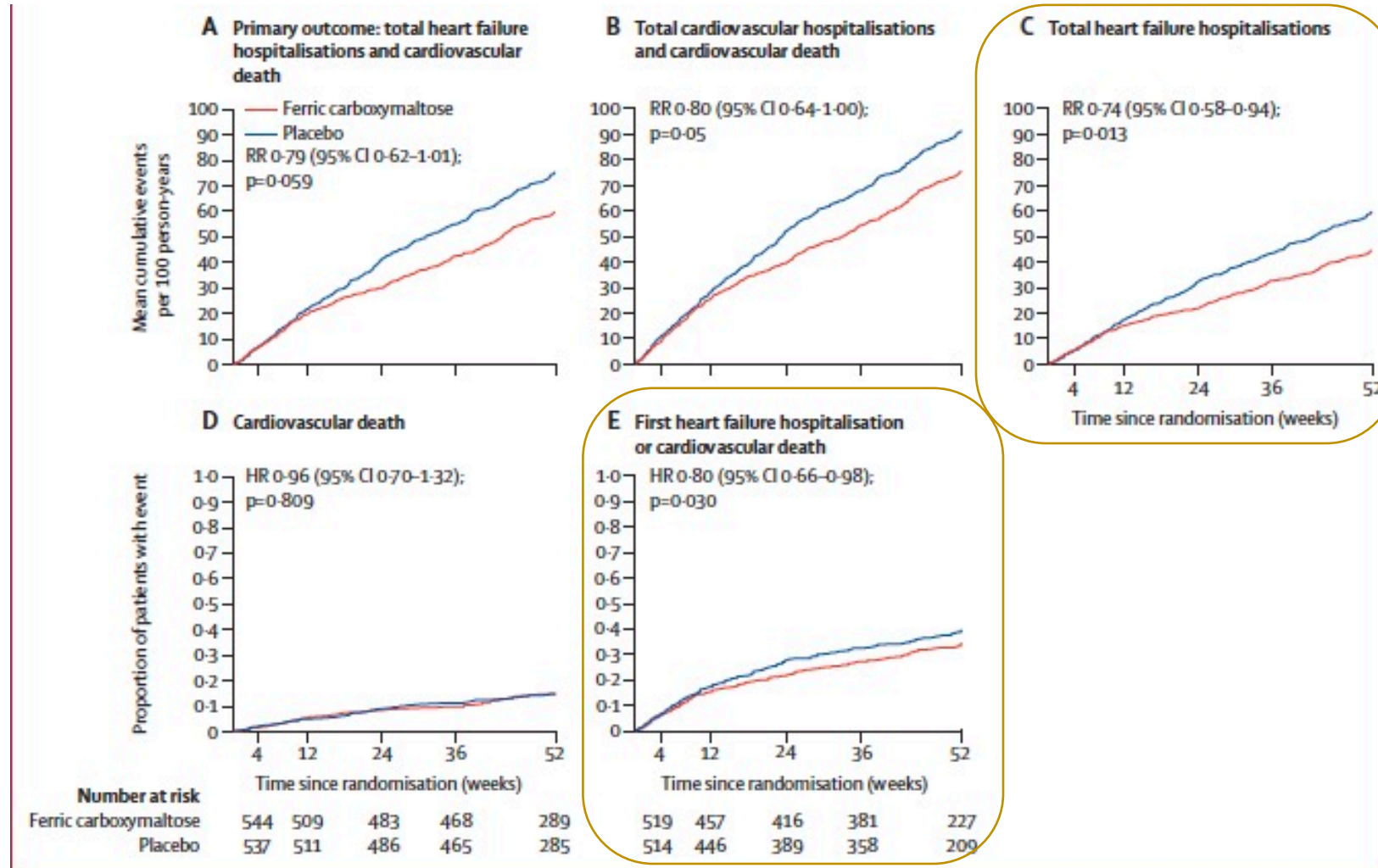
as serum ferritin <100 ng/mL or serum ferritin 100–299 ng/mL with TSAT <20%, to reduce the risk of HF hospitalization.⁵¹²



Ferric carboxymaltose for iron deficiency at discharge after acute heart failure: a multicentre, double-blind, randomised, controlled trial



Piotr Ponikowski, Bridget-Anne Kirwan, Stefan D Anker, Theresa McDonagh, Maria Dorobantu, Jarosław Drozd, Vincent Fabien,



AFFIRM Trial
 Lancet 2020.

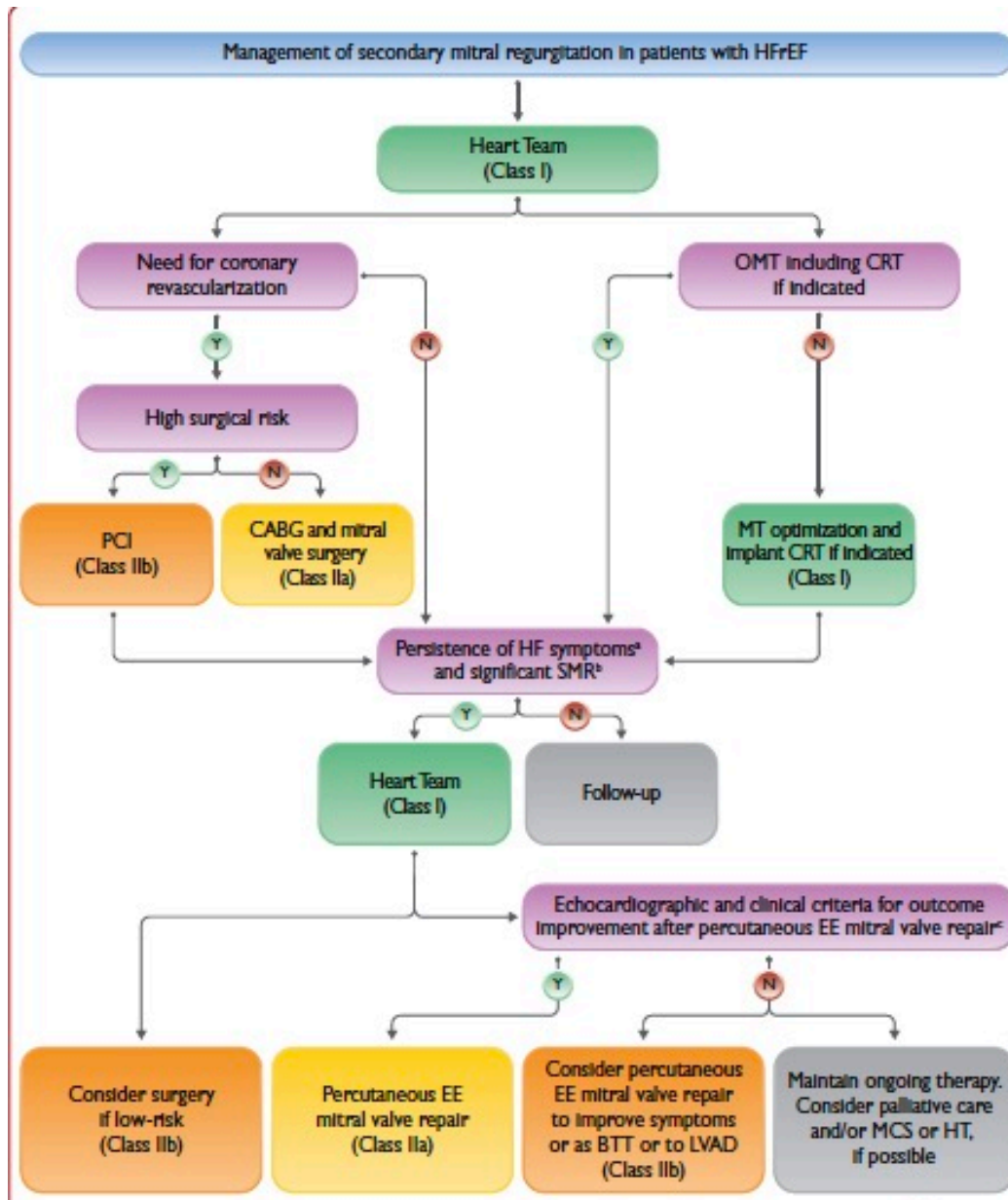
1

To reduce mortality - for all patients			
ACE-I/ARNI	BB	MRA	SGLT2i

2

To reduce HF hospitalization/mortality - for selected patients				
Volume overload				
Diuretics				
SR with LBBB ≥ 150 ms		SR with LBBB 130–149 ms or non LBBB ≥ 150 ms		
CRT-P/D		CRT-P/D		
Ischaemic aetiology		Non-ischaemic aetiology		
ICD		ICD		
Atrial fibrillation	Atrial fibrillation	Coronary artery disease	Iron deficiency	
Anticoagulation	Digoxin PVI	CABG	Ferric carboxymaltose	
Aortic stenosis	Mitral regurgitation	Heart rate SR > 70 bpm	Black Race	ACE-I/ARNI intolerance
SAVR/TAVI	TEE MV Repair	Ivabradine	Hydralazine/ISDN	ARB

3



Secondary mitral regurgitation

Percutaneous edge-to-edge mitral valve repair should be considered in carefully selected patients with secondary mitral regurgitation, not eligible for surgery and not needing coronary revascularization, who are symptomatic^c despite OMT and who fulfil criteria^d for achieving a reduction in HF hospitalizations.⁶¹²

IIa

B

Criteria COAPT*:

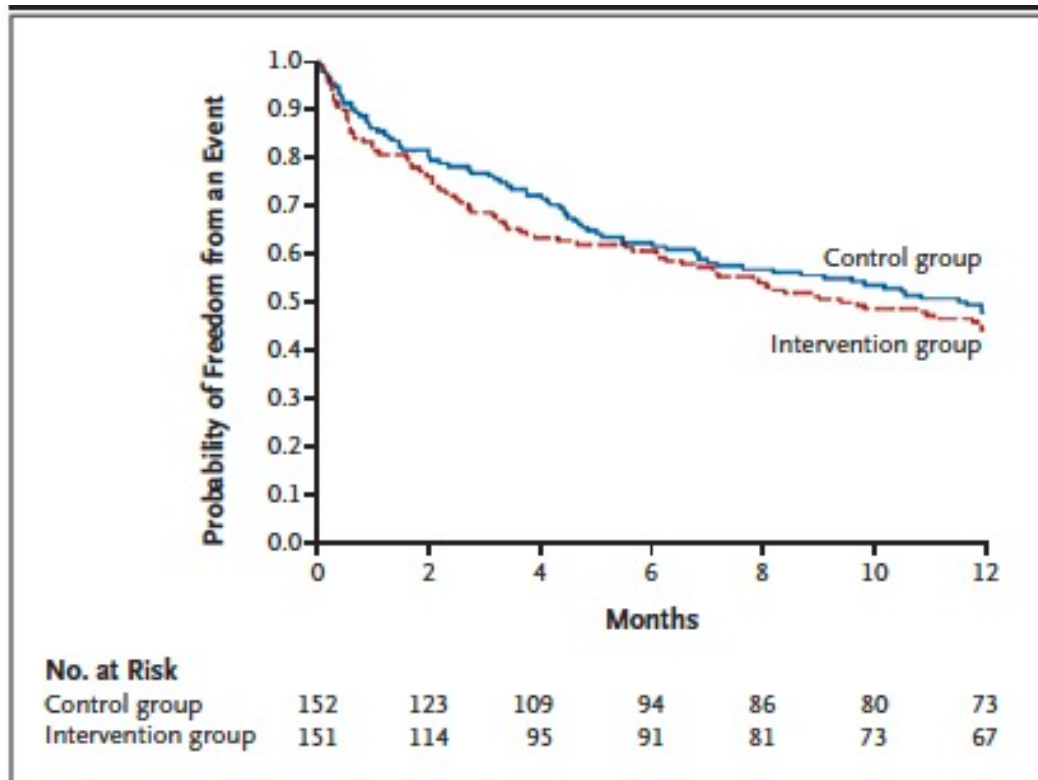
- FEVI <20%.
- DTSVI <70 mm.
- PAPS <70 mmHg.
- Ausencia de dx severa del VD e insuficiencia tricuspídea severa.
- Ausencia de inestabilidad hemodinámica.

*Stone. N Engl J Med 2018.

Percutaneous Repair or Medical Treatment for Secondary Mitral Regurgitation

J.-F. Obadia, D. Messika-Zeitoun, G. Leurent, B. Lung, G. Bonnet, N. Piriou, T. Lefèvre, C. Piot, F. Rouleau,

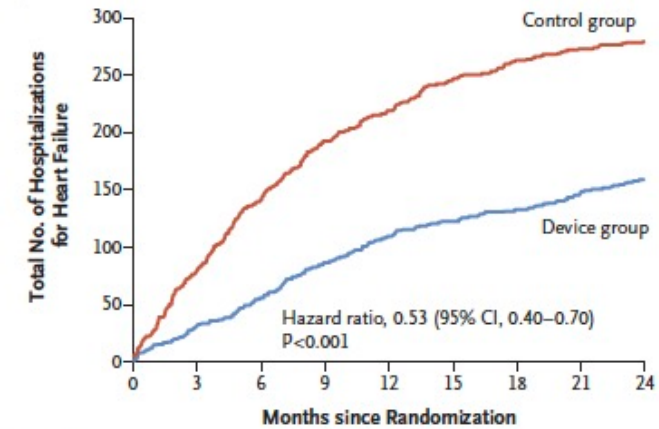
MITRA-FR Trial



Transcatheter Mitral-Valve Repair in Patients with Heart Failure

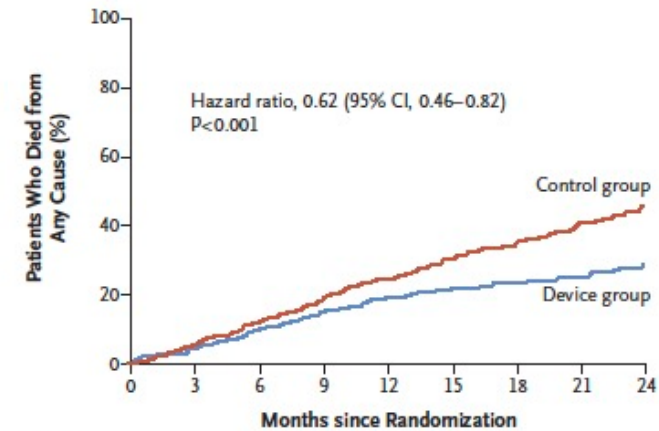
G.W. Stone, J.A. Lindenfeld, W.T. Abraham, S. Kar, D.S. Lim, J.M. Mishell,

A Hospitalization for Heart Failure



No. at Risk	0	3	6	9	12	15	18	21	24
Control group	312	294	271	245	219	176	145	121	88
Device group	302	286	269	253	236	191	178	161	124

C Death from Any Cause



No. at Risk	0	3	6	9	12	15	18	21	24
Control group	312	294	271	245	219	176	145	121	88
Device group	302	286	269	253	236	191	178	161	124

COAPT Trial.
N Engl J Med 2018

1

To reduce mortality - for all patients

ACE-I/ARNI BB MRA SGLT2i

2

To reduce HF hospitalization/mortality - for selected patients

Volume overload
Diuretics

SR with LBBB ≥ 150 ms CRT-P/D	SR with LBBB 130-149 ms or non LBBB ≥ 150 ms CRT-P/D			
Ischaemic aetiology ICD	Non-Ischaemic aetiology ICD			
Atrial fibrillation Anticoagulation	Atrial fibrillation Digoxin PVI	Coronary artery disease CABG	Iron deficiency Ferric carboxymaltose	
Aortic stenosis SAVR/TAVI	Mitral regurgitation TEE MV Repair	Heart rate SR > 70 bpm Ivabradine	Black Race Hydralazine/ISDN	ACE-I/ARNI intolerance ARB

3

4

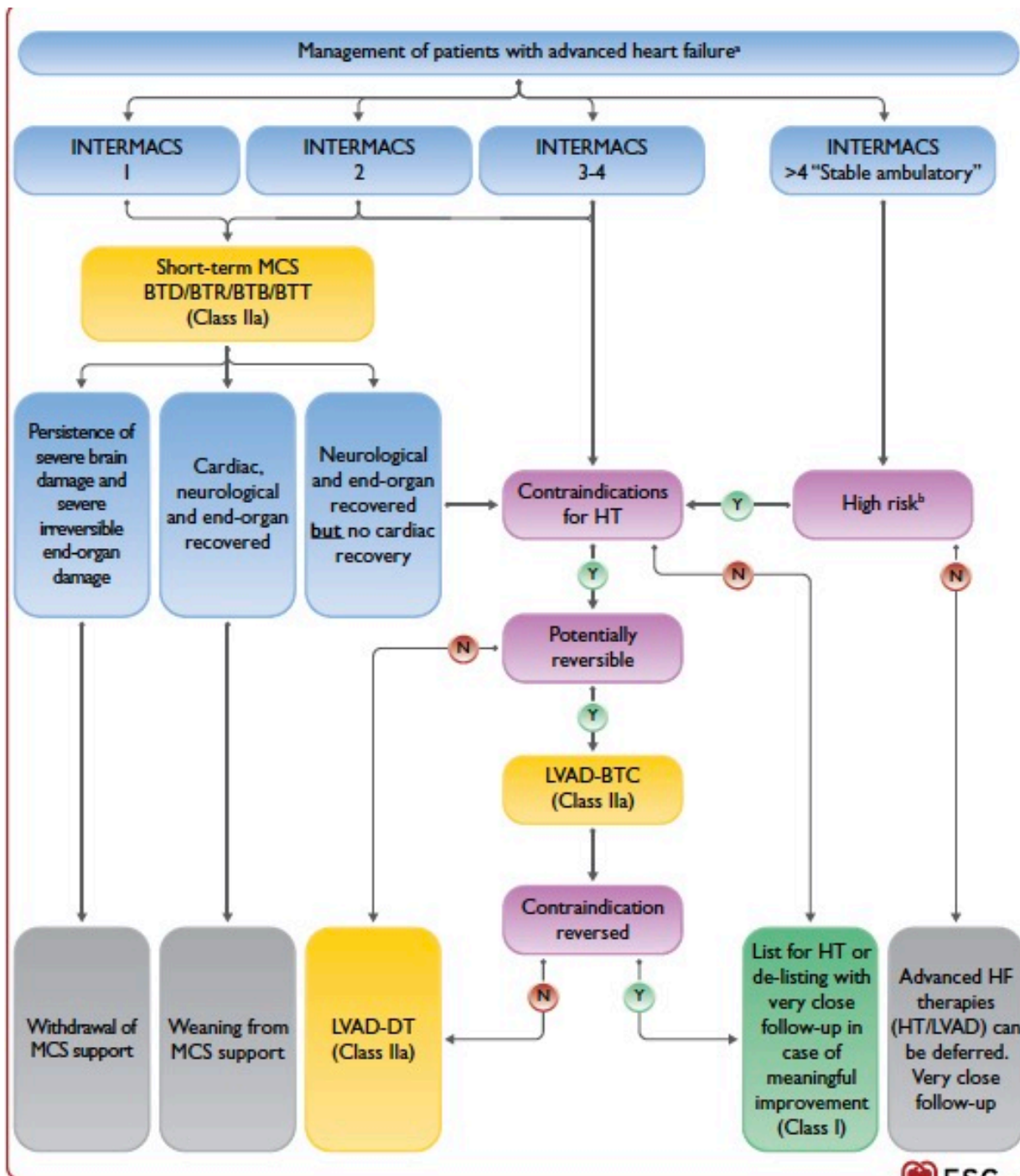
For selected advanced HF patients

Heart transplantation MCS as BTT/BTC Long-term MCS as DT

To reduce HF hospitalization and improve QOL - for all patients

Exercise rehabilitation

Multi-professional disease management



Patients being considered for long-term MCS must have good compliance, appropriate capacity for device handling and psychosocial support.^{414–416}

I	C
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Heart transplantation is recommended for patients with advanced HF, refractory to medical/device therapy and who do not have absolute contraindications.

I	C
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Long-term MCS should be considered in patients with advanced HF_rEF despite optimal medical and device therapy, not eligible for heart transplantation or other surgical options, and without severe right ventricular dysfunction, to reduce the risk of death and improve symptoms.^{378,396,397,401,402,404,417}

IIa	A
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Long-term MCS should be considered in patients with advanced HF_rEF refractory to optimal medical and device therapy as a bridge to cardiac transplantation in order to improve symptoms, reduce the risk of HF hospitalization and the risk of premature death.^{398–400,402,404}

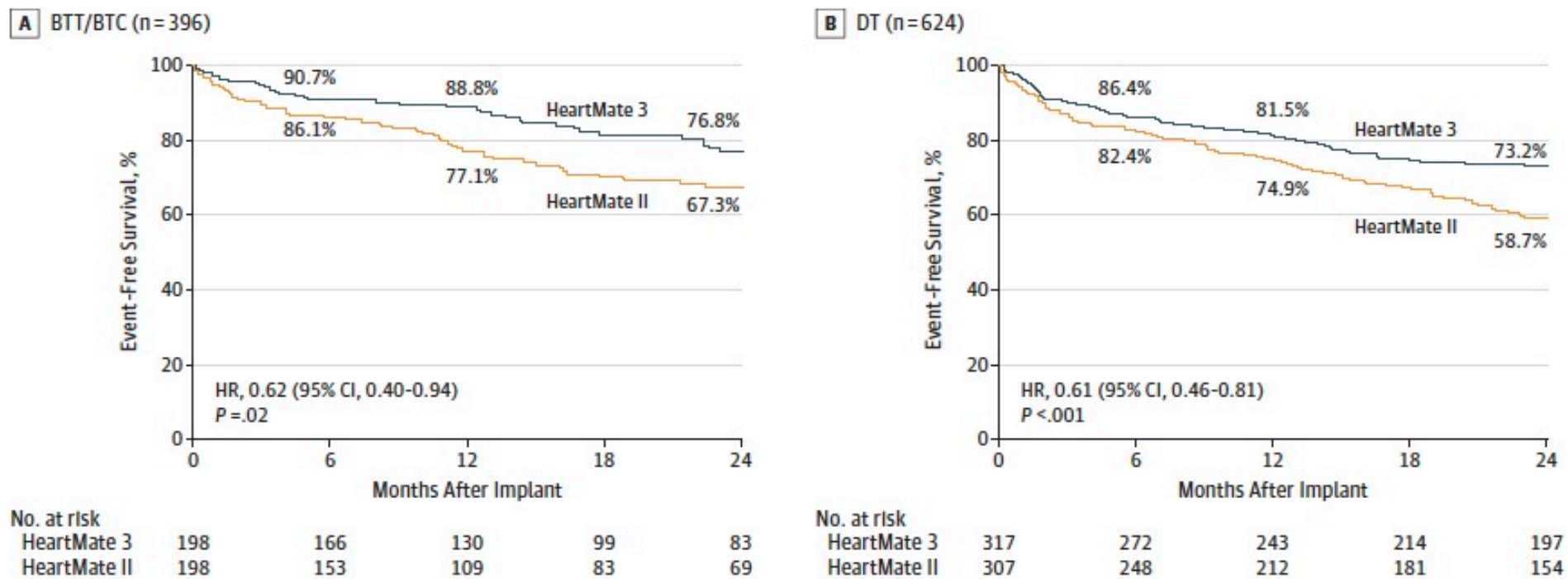
IIa	B
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Association of Clinical Outcomes With Left Ventricular Assist Device Use by Bridge to Transplant or Destination Therapy Intent

The Multicenter Study of MagLev Technology in Patients Undergoing Mechanical Circulatory Support Therapy With HeartMate 3 (MOMENTUM 3) Randomized Clinical Trial

Daniel J. Goldstein, MD; Yoshifumi Naka, MD, PhD; Douglas Horstmanshof, MD; Ashwin K. Ravichandran, MD, MPH;

Figure 1. Primary End Point Analysis



Survival at 2 years free of disabling stroke (defined as a modified Rankin score greater than 3) or reoperation to replace or remove a malfunctioning device in the bridge to transplant (BTT)/bridge to transplant candidacy (BTC) and destination therapy (DT) cohorts. HR indicates hazard ratio.

1

To reduce mortality - for all patients

ACE-I/ARNI BB MRA SGLT2i

2

To reduce HF hospitalization/mortality - for selected patients

Volume overload
Diuretics

SR with LBBB ≥ 150 ms CRT-P/D	SR with LBBB 130-149 ms or non LBBB ≥ 150 ms CRT-P/D			
Ischaemic aetiology ICD	Non-Ischaemic aetiology ICD			
Atrial fibrillation Anticoagulation	Atrial fibrillation Digoxin PVI	Coronary artery disease CABG	Iron deficiency Ferric carboxymaltose	
Aortic stenosis SAVR/TAVI	Mitral regurgitation TEE MV Repair	Heart rate SR > 70 bpm Ivabradine	Black Race Hydralazine/ISDN	ACE-I/ARNI intolerance ARB

3

4

For selected advanced HF patients

Heart transplantation MCS as BTT/BTC Long-term MCS as DT

5

To reduce HF hospitalization and improve QOL - for all patients

Exercise rehabilitation

Multi-professional disease management

Recomendaciones generales

Recommendations	Class ^a	Level ^b
It is recommended that HF patients are enrolled in a multidisciplinary HF management programme to reduce the risk of HF hospitalization and mortality. ^{309,314,315,316}	I	A
Self-management strategies are recommended to reduce the risk of HF hospitalization and mortality. ³⁰⁹	I	A
Either home-based and/or clinic-based programmes improve outcomes and are recommended to reduce the risk of HF hospitalization and mortality. ^{310,317}	I	A
Influenza and pneumococcal vaccinations should be considered in order to prevent HF hospitalizations. ^{315,316}	IIa	B

Recommendations	Class ^a	Level ^b
Exercise is recommended for all patients who are able in order to improve exercise capacity, QOL, and reduce HF hospitalization. ^{c 324–328,335–337}	I	A
A supervised, exercise-based, cardiac rehabilitation programme should be considered in patients with more severe disease, frailty, or with comorbidities. ^{95,324–327,338}	IIa	C
Non-invasive HTM may be considered for patients with HF in order to reduce the risk of recurrent CV and HF hospitalizations and CV death. ³⁷⁴	IIb	B

ACE-I/ARNI

BB

MRA

SGLT2i



Rosano GMC, et al. Eur J Heart Fail. 2021. PMID: 33932268

Muchas gracias!

anagarci@clinic.cat