
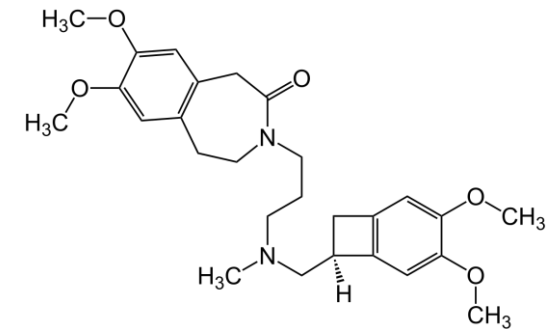


3er Curs
D'ACTUALITZACIÓ EN
INSUFICIÈNCIA
CARDÍACA

Hotel Hilton Barcelona
Avda Diagonal 589-591, 08014 BCN
12 de novembre de 2021



Amb la col·laboració de:
NOVARTIS

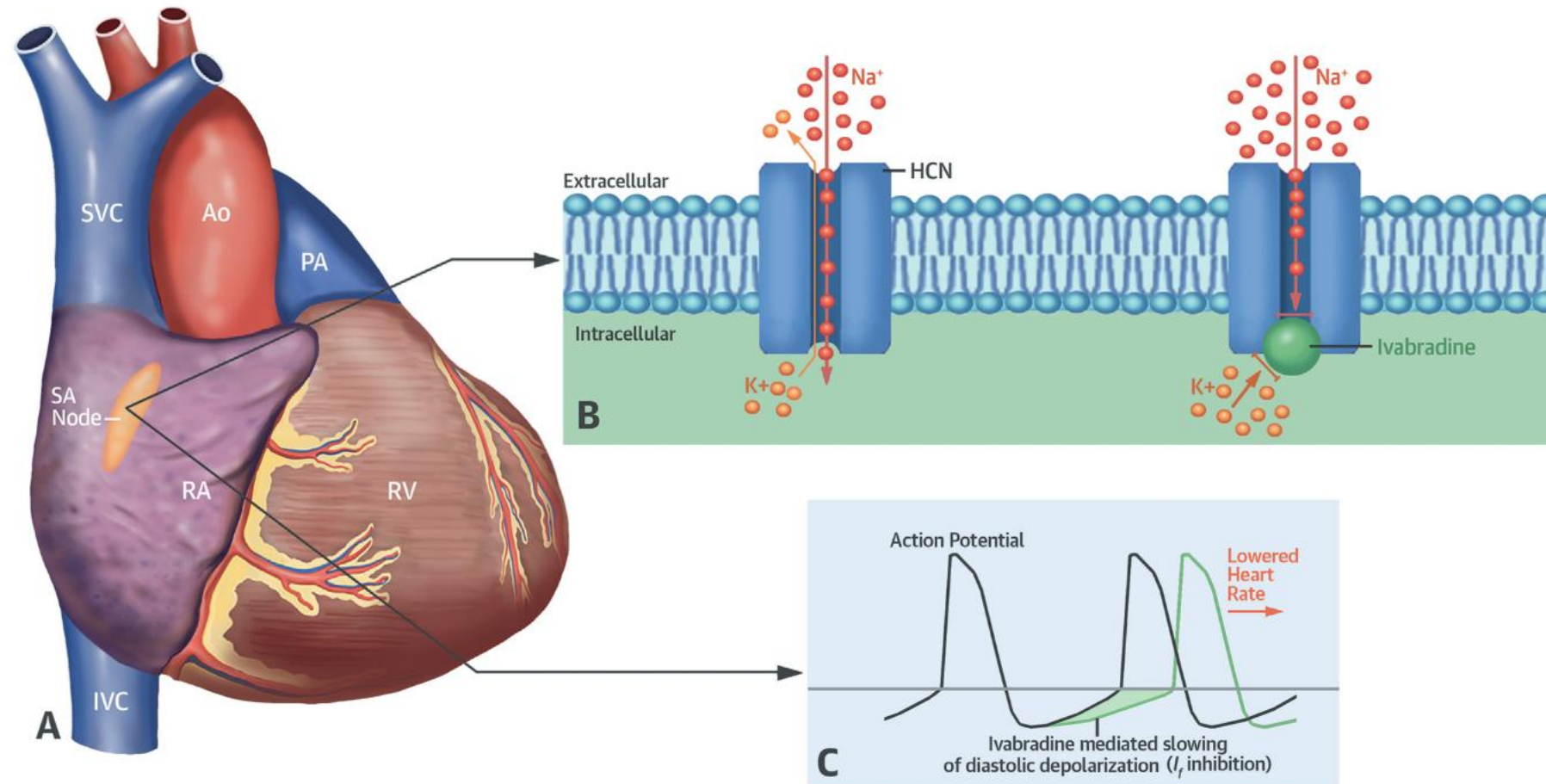


Papel de la Ivabradina

Santiago Cudini
Hospital de Figueres

12 de Novembre de 2021

Mecanismo de acción de la Ivabradina



Evidencia del fármaco en IC

Ivabradine and outcomes in chronic heart failure (SHIFT): a randomised placebo-controlled study

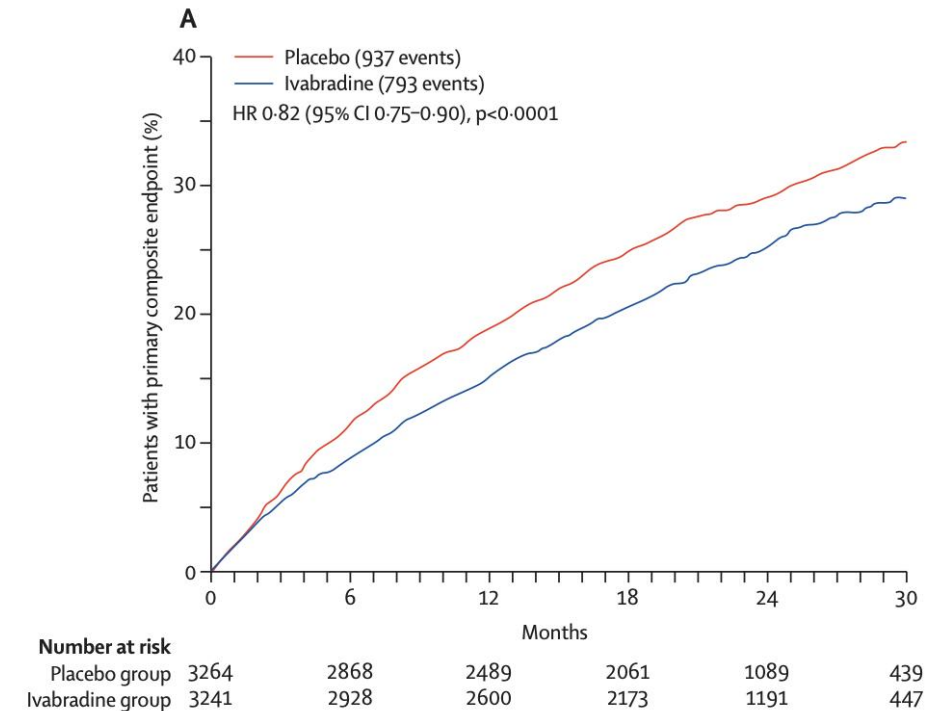
6558 Pacientes con IC sintomática
FEVI < 35%

Ingreso en los últimos 12 meses

RS con FC >70lpm
TMO con BB a dosis máxima
tolerada

Endpoint
Combiinado

Mort CV y
Reingreso



¿Qué característica tenía la población sustituta en SHIFT?

	Ivabradine group (n=3241)	Placebo group (n=3264)
NYHA class		
Class II	1585 (49%)	1584 (49%)
Class III	1605 (50%)	1618 (50%)
Class IV	50 (2%)	61 (2%)

Treatment at randomisation

β blocker	2897 (89%)	2923 (90%)
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**IECA/ARAI
BB
ARM**

ICD	92 (3%)	115 (4%)
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Reasons for failure to reach target dose*†

Hypotension	933 (44%)	952 (45%)
Fatigue	676 (32%)	670 (32%)
Dyspnoea	284 (14%)	302 (14%)
Dizziness	267 (13%)	245 (12%)
Bradycardia	134 (6%)	125 (6%)
Other	199 (9%)	219 (10%)
Patients not receiving β blocker	344 (11%)	341 (10%)
Reasons for non-prescription of β blocker		
Chronic obstructive pulmonary disease	126 (37%)	109 (32%)
Hypotension	59 (17%)	68 (20%)
Asthma	35 (10%)	39 (11%)
Cardiac decompensation	23 (7%)	31 (9%)
Dizziness or bradycardia	24 (7%)	17 (5%)
Fatigue	17 (5%)	20 (6%)
Raynaud or peripheral arterial disease	16 (5%)	20 (6%)
Other	44 (13%)	37 (11%)

2021 ESC Guidelines for the diagnosis and treatment of acute and chronic heart failure

I_f-channel inhibitor

Ivabradine should be considered in symptomatic patients with LVEF $\leq 35\%$, 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 $\leq 35\%$, 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

IC sintomática
FEVI < 35%
RS con FC >70lpm

Con BB a dosis máxima
tolerada o sin BB con menor
evidencia
MAS IECA (¿ARNI?) y ARM

The NEW ENGLAND JOURNAL of MEDICINE

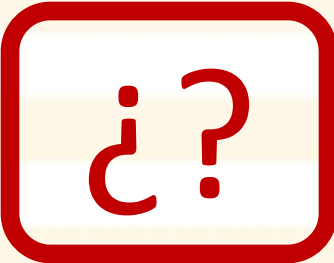
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Angiotensin–Neprilysin Inhibition versus Enalapril in Heart Failure

Characteristic	LCZ696 (N=4187)	Enalapril (N=4212)
Treatments at randomization — no. (%)		
Diuretic	3363 (80.3)	3375 (80.1)
Digitalis	1223 (29.2)	1316 (31.2)
Beta-blocker	3899 (93.1)	3912 (92.9)
Mineralocorticoid antagonist	2271 (54.2)	2400 (57.0)
Implantable cardioverter–defibrillator	623 (14.9)	620 (14.7)
Cardiac resynchronization therapy	292 (7.0)	282 (6.7)



Management of HFrEF

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

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

Conclusión

- Debemos iniciar e intentar titular hasta dosis máximas los BB antes de comenzar Ivabradina.
- Esta recomendada para un grupo seleccionado de pacientes.
- Existe un GAP en la evidencia debido a como ha ido evolucionando el TM en IC.

Muchas gracias.