

Approach to Congestive Heart Failure

A Case Based Approach

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FAMILY MEDICINE DAY – January 19th, 2025

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Disclosures

- None

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Objectives

- Case based
- Congestive heart failure types
- Lifestyle modifications
- Medical therapy
- Implantable cardiac devices

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Case 1

- Mrs. J is a 94F presents to hospital with 4 day history of shortness of breath and leg edema.
- PMHx: Hypertension. DM2.
- No prior cardiac history

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Case 1

- On examination:
 - BP 168/78 mmHg, HR 72 bpm, O2 sat 92% on R/A
 - JVP elevated
 - Normal heart sounds. Bilateral lung crackles.
 - Moderate leg edema.
- 12 Lead EKG
 - Normal sinus rhythm.
- Bedside Echocardiogram
 - Normal LV/RV size and function. No significant valvular abnormalities.

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Case 1

- Labs:
 - Hb 105, WBC 11, PLT 150
 - Creatinine 110, Na/K= normal.
 - NT pro-BNP elevated at 2000.
 - Troponin – normal.
 - CXR: Increased vascular redistribution
- **Diagnosis:**
 - Congestive heart failure with PRESERVED LV function.

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Congestive Heart Failure Subtypes

- Congestive Heart Failure with Preserved LV function (HFPeF)
 - LVEF>50%
- Congestive Heart Failure with mid-range EF (HFmEF)
 - LVEF 41-49%
- Congestive Heart Failure with REDUCED LV systolic function (HFrEF)
 - LVEF <40%

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Case 1

- Patient was treated with iv Furosemide until euvolemic
- Patient discharged home on maintenance diuretics
 - Furosemide 40mg po daily
 - Spironolactone 12.5 mg daily.
 - Empagliflozin 10mg daily

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HFPeF

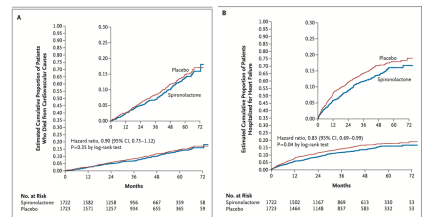
- Medical Treatment:
 - Loop diuretics (ie. Furosemide)
 - MRA inhibitors (ie. Spironolactone)
 - SGLT2 inhibitors

Medications reduce hospitalizations, but do not reduce mortality.

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Medical therapy for HFPeF

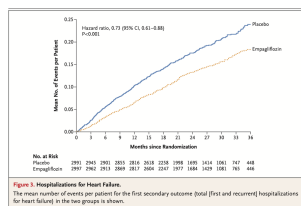
Spironolactone
TOPCAT Study
NEJM 2014



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Medical therapy for HFPeF

- Empagliflozin
EMPEROR-Preserved NEJM 2021



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Lifestyle modifications for CHF

- Diet
 - Well balanced diet, consider careful weight loss
- Salt
 - Avoid salt rich foods
 - < 2,000mg daily. (even less if hypertensive)
- Fluids
 - <1.5-2L daily
- Alcohol
 - Limit as much as possible. (<1 beverage daily)
- Avoid
 - NSAIDs

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Lifestyle modifications for CHF

- Exercise
 - Daily activities
 - Walking
 - Avoid over-exertion
 - Increase amount of activity slowly
 - Symptom limited
 - Consider referral to cardiac rehab.

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Lifestyle modifications for CHF

- Self Monitoring
 - Report the following to primary care or Cardiology
 - Change in breathing pattern. (SOB, Orthopnea, PND)
 - Worsening fatigue
 - Weight gain (1 Kg in 2-3 days)
 - Edema- increase
 - Side effects from medications
 - Irregular pulse, abnormal vital signs

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Case 2

- 62M with history of HTN, DM2 presents to ER with 4 days of worsening shortness of breath, leg edema, lightheadedness
- In ER, BP 105/74mmHg, HR 162 bpm, O2 Sat 92% on 2L O2
- JVP elevated to angle of jaw, Mild early peaking systolic murmur at the base
- Bilateral leg edema

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Case 2

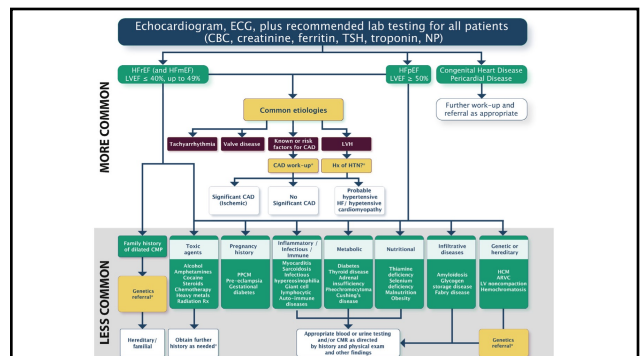
ECG strip showing sinus tachycardia with a heart rate of 162 bpm. The rhythm is regular with narrow QRS complexes. The PR interval is normal. The QRS complexes are narrow and the ST segment and T waves are unremarkable.

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Case 2

- 12 lead EKG: Atrial fibrillation with rapid ventricular response
- 2D Echocardiogram:
 - Global hypokinesis. SEVERE LV systolic function. LVEF 25-30%.
 - Mild aortic stenosis.
- Coronary angiography: Normal coronaries

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Case 2

- Patient was successfully cardioverted back to sinus rhythm
- Discharged home on Bisoprolol 5mg daily, Apixaban 5mg bid

- 3 months later....
 - Patient remains in sinus rhythm.
 - Repeat Echocardiogram: Normal LV systolic function!

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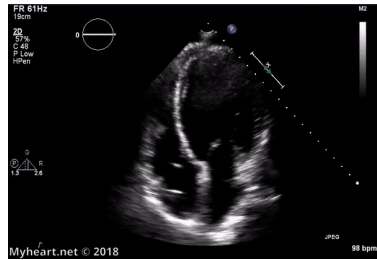
Case 3

- 72M with history of MI 2021- PCI to LAD/RCA. HTN. DM2
- Meds: ASA 81mg daily, Ramipril 5mg OD, Atorvastatin 20mg
- Presents with symptoms suggestive of CHF.
- On examination
 - BP 145/70 mmHg, HR 94 bpm, O2 96% RA
 - JVP elevated. Normal heart sounds.
 - Mild leg edema
- EKG: Sinus rhythm. Septal infarct pattern

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Case 3

- Echocardiogram:
- Severe LV systolic dysfunction. LVEF 25%

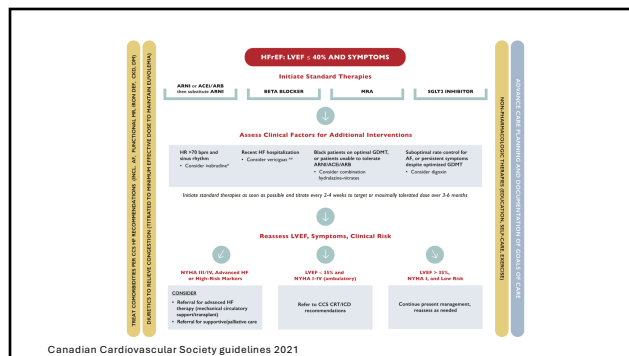


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Case 3

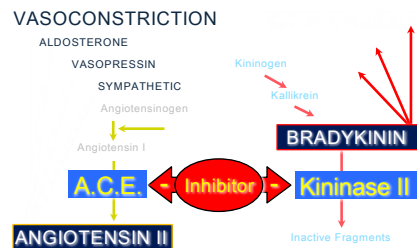
- Patient started on Furosemide for diuresis, and then initiated GDMT= Guideline Directed Medical Therapy

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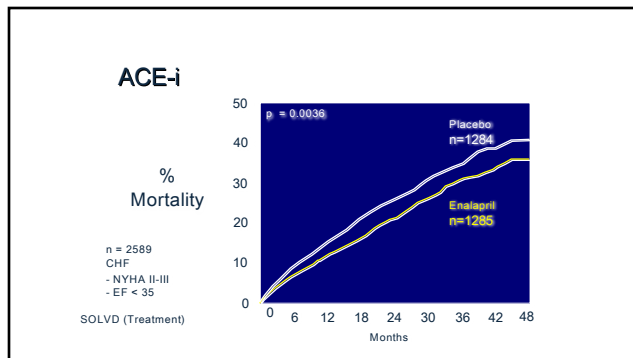


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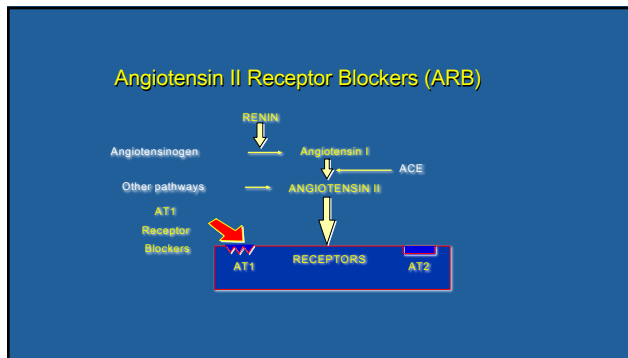
Ace Inhibitors Mechanism of Action



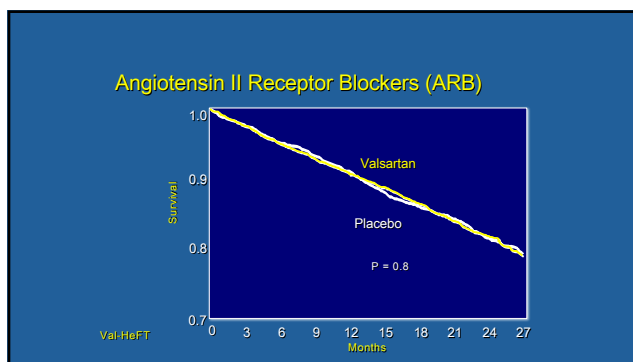
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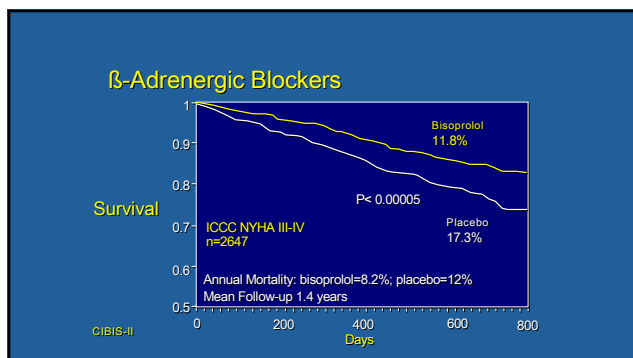
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- ### β-Adrenergic Blockers Mechanism of action
- Decreased Neurohormonal activation
 - Decreased HR
 - Antiischemic
 - Antihypertensive
 - Antiarrhythmic
 - Antioxidant, Antiproliferative

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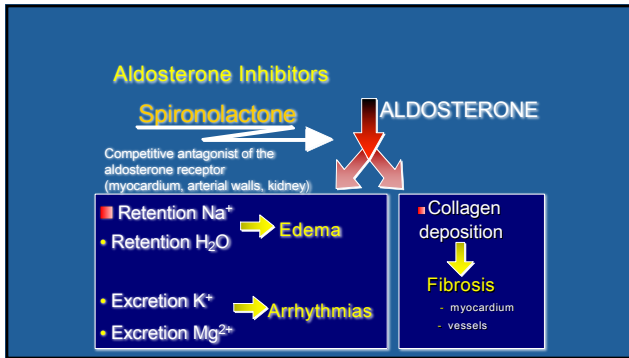
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Supplemental Table S3. Evidence-based heart failure medications and doses in the management of patients with heart failure with reduced ejection fraction

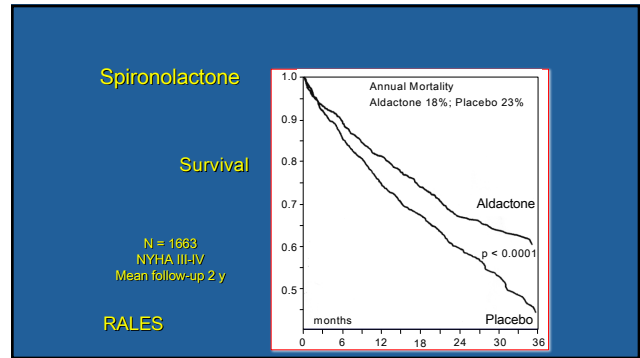
Drug	Start Dose	Target Dose
ACE Inhibitors		
Captopril	6.25-12.5 mg TID	25-50 mg TID
Enalapril	1.25-2.5 mg BID	10 mg BID
Lisinopril	2.5-5 mg OD	20-35 mg OD
Perindopril	2-4 mg OD	4-8 mg OD
Ramipril	1.25-2.5 mg BID	5 mg BID
Trandolapril	1-2 mg OD	4 mg OD
Beta-blockers		
Bisoprolol	1.25 mg OD	10 mg OD
Carvedilol	3.125 mg BID	25 mg BID*
Metoprolol CR/XL**	12.5-25 mg OD	200 mg OD
ARBs		
Candesartan	4 mg OD	32 mg OD
Valsartan	40 mg BID	160 mg BID
Aldosterone Antagonists		
Spiro lactone	12.5 mg OD	50 mg OD
Eplerenone	25 mg OD	50 mg OD
Vasodilators		
Hydralazine	37.5 mg TID	75 mg TID
Isorbide dinitrate	20 mg TID	40 mg TID

* 50 mg BID if weight is >85 kg

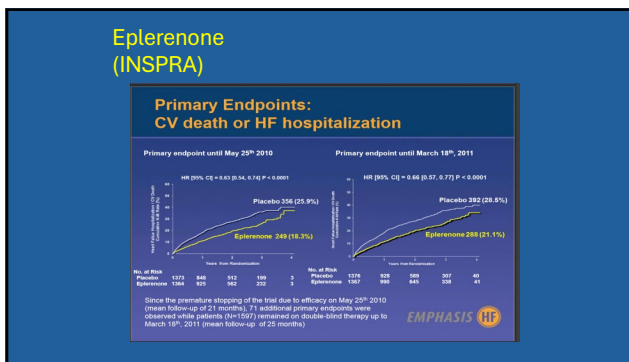
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Supplemental Table S3. Evidence-based heart failure medications and doses in the management of patients with heart failure with reduced ejection fraction

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Beta-blockers		
Bisoprolol	1.25 mg OD	10 mg OD
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Valsartan	40 mg BID	160 mg BID
Aldosterone Antagonists		
Spirolactone	12.5 mg OD	50 mg OD
Eplerenone	25 mg OD	50 mg OD
Vasodilators		
Hydralazine	37.5 mg TID	75 mg TID
Isorbide dinitrate	20 mg TID	40 mg TID

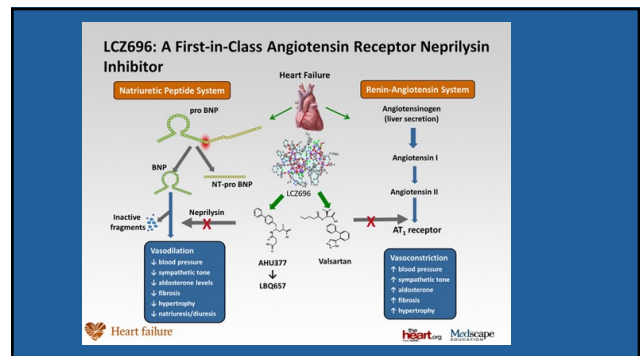
*50 mg BID if weight is >85 kg

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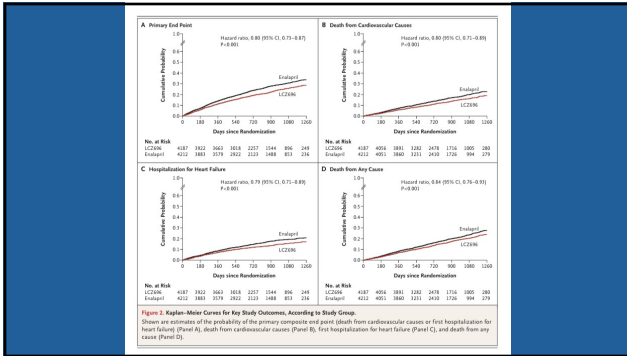
ENTRESTO

- PARADIGM-HF Trial
- LCZ696 =
 - Sacubitril (Nephrilysin inhibitor) + Valsartan (ARB)
- 8442 patients NYHA 2-4, EF<40% randomized to LCZ696 vs. Enalapril 10mg bid.

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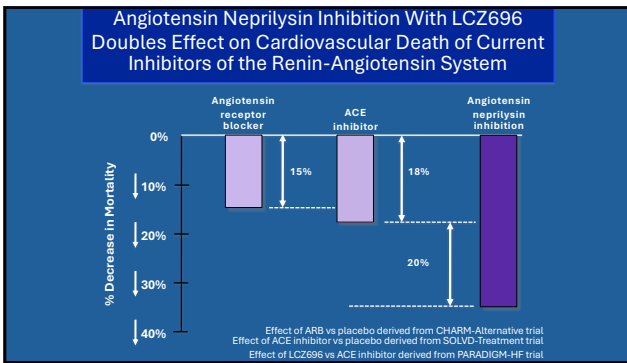


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Table 3. Adverse Events during Randomized Treatment.*

Event	LCZ696 (N=4187)	Enalapril (N=4212)	P Value
Hypotension			
Symptomatic	588 (14.0)	388 (9.2)	<0.001
Symptomatic with systolic blood pressure <90 mm Hg	112 (2.7)	59 (1.4)	<0.001
Elevated serum creatinine			
≥2.5 mg/dl	139 (3.3)	188 (4.5)	0.007
≥3.0 mg/dl	63 (1.5)	83 (2.0)	0.10
Elevated serum potassium			
>5.5 mmol/liter	674 (16.1)	727 (17.3)	0.15
>6.0 mmol/liter	181 (4.3)	236 (5.6)	0.007
Cough	474 (11.3)	601 (14.3)	<0.001
Angioedema†			
No treatment or use of antihistamines only	10 (0.2)	5 (0.1)	0.19
Use of catecholamines or glucocorticoids without hospitalization	6 (0.1)	4 (0.1)	0.52
Hospitalization without airway compromise	3 (0.1)	1 (<0.1)	0.31
Airway compromise	0	0	—

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DOSAGE FORM/ STRENGTH

ENTRESTO (sacubitril/valsartan) film-coated tablets:

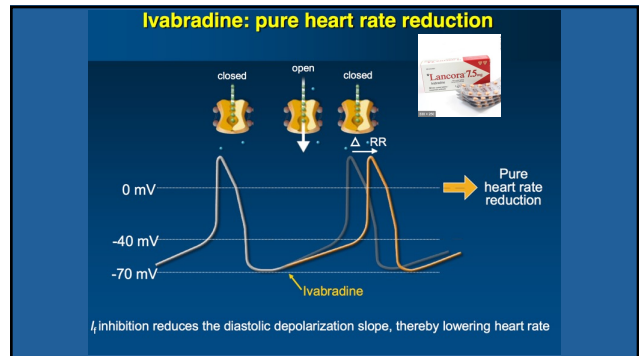
- 24.3mg sacubitril/ 25.7mg valsartan
 - Equivalent to 40mg valsartan monotherapy
- 48.6mg sacubitril/ 51.4mg valsartan
 - Equivalent to 80mg valsartan monotherapy
 - Equivalent to 100mg of LCZ696 in the Paradigm HF study
- 97.2mg sacubitril/ 102.8mg valsartan
 - Equivalent to 160mg valsartan monotherapy
 - Equivalent to 200mg of LCZ696 in the Paradigm HF study

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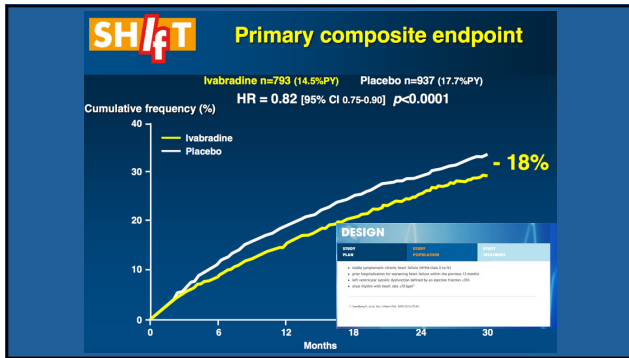
WARNINGS AND PRECAUTIONS

- Pregnancy (Serious Warning and Precaution)**
 - When used in pregnancy angiotensin receptor (AT1) blockers (ARB) can cause injury to or even death of the developing fetus. When pregnancy is detected, sacubitril/valsartan should be discontinued as soon as possible.
- Co-administration with ACEi**
 - sacubitril/valsartan must not be administered with an ACEi due to the risk of angioedema. Sacubitril/valsartan must not be initiated until at least 36 hours have elapsed following discontinuation of ACEi therapy. If treatment with sacubitril/valsartan is stopped, ACEi therapy must not be initiated until 36 hours after the last dose of sacubitril/valsartan.
- Co-administration with ARB**
 - sacubitril/valsartan should not be co-administered with any other drug formulation containing an ARB, due to the angiotensin II receptor blocking activity of sacubitril/valsartan by its valsartan moiety.

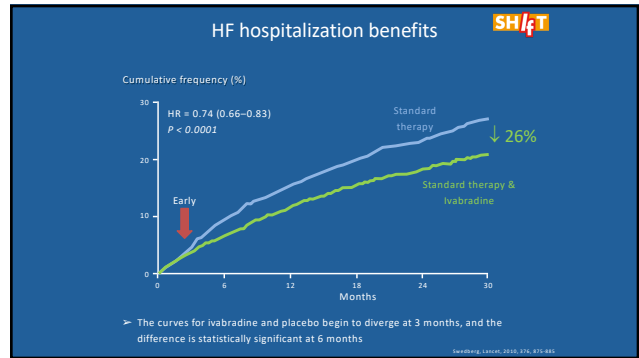
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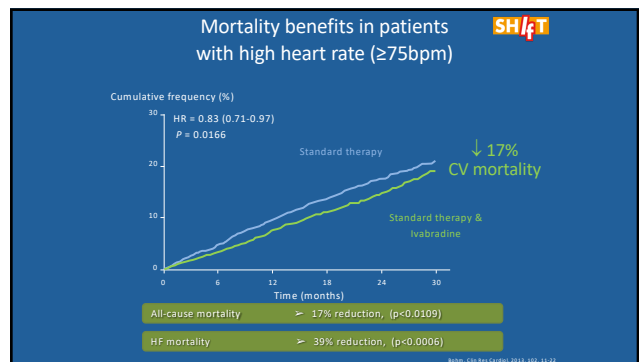
SH/FT Subgroup analysis

➤ Patients with baseline heart rate ≥ 75 bpm
 63% of the Shift patients (N = 4150)

Post-hoc analysis

- Requested by European Medicines Agency
- Ivabradine was approved in Europe and indicated for HF patient with a heart rate ≥ 75 bpm based on this analysis

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SGLT-2 Inhibitors

Dapagliflozin in Patients with Heart Failure and Reduced Ejection Fraction

John J. McMurray, M.D., Scott D. Solomon, M.D., Silvio E. Inzucchi, M.D., Lars Køber, M.D., D.M.Sc., Mikhail N. Kosiborod, M.D., Felipe A. Martinez, M.D., Piotr Ponikvarski, M.D., Ph.D., Marc S. Sabatine, M.D., M.P.H., Indar S. Anand, M.D., Jan Böhler, M.D., Ph.D., Michael Böhm, M.D., Ph.D., Chen-En Chiang, M.D., Ph.D., et al., for the DAPA-HF Trial Committees and Investigators*

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DAPA-HF Trial

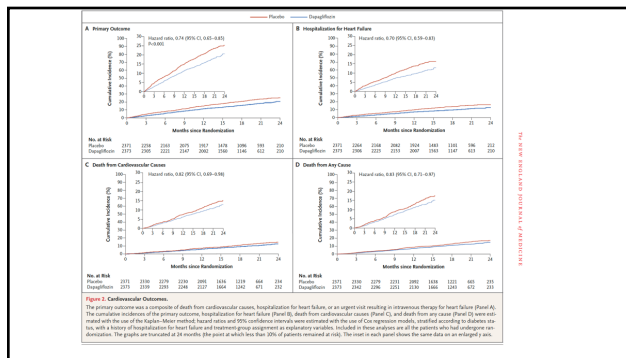
- 4744 patients with NYHA 2-4
- HFREF (LVEF <40%)
- Diabetes not necessary
- Randomized:
 - Dapagliflozin 10mg daily vs Placebo
- Primary outcome:
 - Composite: worsening HF/hospitalization or cardiovascular death

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Table 1. (Continued.)

Characteristic	Dapagliflozin (N=2373)	Placebo (N=2371)
Heart failure medication — no. (%)		
Diuretic	2216 (93.4)	2217 (93.5)
ACE inhibitor	1332 (56.1)	1329 (56.1)
ARB	675 (28.4)	632 (26.7)
Sacubitril-valsartan	250 (10.5)	258 (10.9)
Beta-blocker	2278 (96.0)	2280 (96.2)
Mineralocorticoid receptor antagonist	1696 (71.5)	1674 (70.6)
Digitalis	445 (18.8)	442 (18.6)
Glucose-lowering medication — no./total no. (%)**		
Biguanide	504/993 (50.8)	512/990 (51.7)
Sulfonylurea	228/993 (23.0)	210/990 (21.2)
DPP-4 inhibitor	161/993 (16.2)	149/990 (15.1)
GLP-1 receptor agonist	11/993 (1.1)	10/990 (1.0)
Insulin	274/993 (27.6)	266/990 (26.9)

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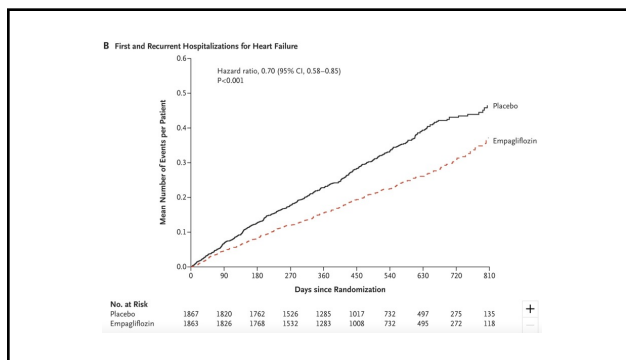
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ORIGINAL ARTICLE

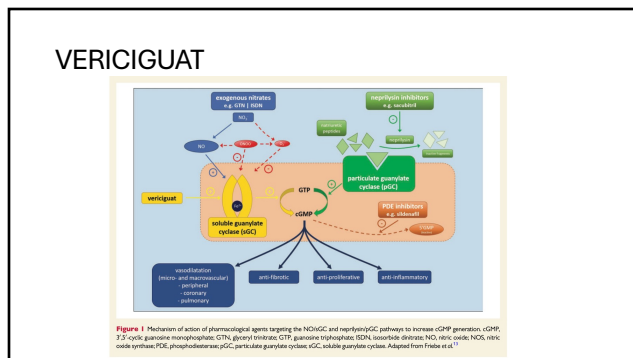
Cardiovascular and Renal Outcomes with Empagliflozin in Heart Failure

Milton Packer, M.D., Stefan D. Anker, M.D., Ph.D., Javed Butler, M.D., Gerasimos Filippatos, M.D., Stuart J. Pocock, Ph.D., Peter Carson, M.D., James Januzzi, M.D., Subodh Verma, M.D., Ph.D., Hirooyuki Tsutsui, M.D., Martina Brueckmann, M.D., Waheed Jamal, M.D., Karen Kimura, Ph.D., et al., for the EMPEROR-Reduced Trial Investigators*

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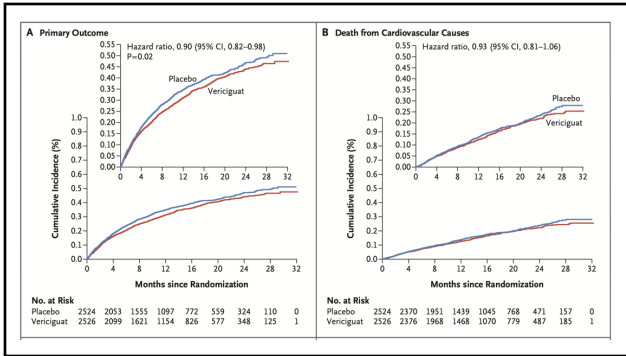


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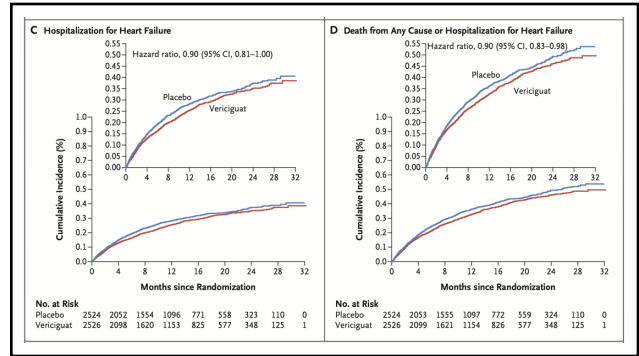
Vericiguat in Patients with Heart Failure and Reduced Ejection Fraction

Paul W. Armstrong, M.D., Burkert Pieske, M.D., Kevin J. Anstrom, Ph.D., Justin Ezekowitz, M.B., B.Ch., Adrian F. Hernandez, M.D., M.H.S., Javed Butler, M.D., M.P.H., M.B.A., Carolyn S.P. Lam, M.B., B.S., Ph.D., Piotr Ponikvar, M.D., Adriaan A. Voors, M.D., Ph.D., Gang Jia, Ph.D., Steven E. McNulty, M.S., Mahesh J. Patel, M.D., Lothar Roessig, M.D., Joerg Koglin, M.D., Ph.D., and Christopher M. O'Connor, M.D., for the VICTORIA Study Group*

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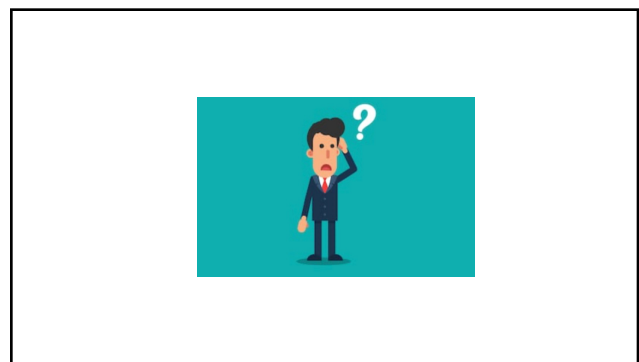


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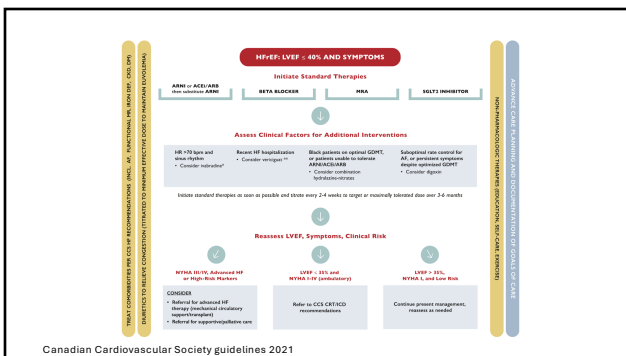
DRUGS to AVOID in HFReF

- Calcium Channel blockers
- Class Ic anti-arrhythmic medications
- NSAIDS
- Cold/cough meds with ephedrine
- Thiazolidinediones

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Case 3 - 3 months later

- Patient feeling a lot better.
- Meds:
 - Entresto 48 mg po BID. (Switched from Ramipril with 48h washout)
 - Bisoprolol 5mg po daily. (Feels fatigued on Bisoprolol)
 - Empagliflozin 10mg daily
 - Spironolactone 25mg po daily
 - ASA 81mg daily, Atorvastatin 20mg OD
- Vitals:
 - BP 95/60 mmHg, HR 82 bpm. (SINUS RHYTHM)
 - JVP not elevated. Normal heart sounds. No edema
- ?Further medication additions?

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Case 3 – 3 months later

- Patient started on Ivabradine 5mg po bid
- BP: 90/60 mmHg. HR 65 bpm.
- JVP not elevated. Normal heart sounds. No edema.
- Repeat ECHOCARDIOGRAM
 - LV systolic mildly improved. LVEF ~30%.

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Long term sequelae of Severe LV systolic dysfunction

Abnormal electrical signals in the ventricles

Ventricular Tachycardia ECG

Ventricular Fibrillation

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Implantable Cardiac Defibrillators

ChestDevices.com

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Implantable Cardiac Defibrillators

- Indications:
 - PRIMARY prophylaxis
 - SECONDARY prophylaxis

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Primary Prophylaxis Indications

RECOMMENDATION

51. We recommend consideration of primary ICD therapy in patients with:

- Ischemic cardiomyopathy, NYHA class II-III, EF \leq 35%, measured at least 1 month post MI, and at least 3 months post coronary revascularization procedure (Strong Recommendation; High-Quality Evidence); or
- Ischemic cardiomyopathy, NYHA class I, and an EF \leq 30% at least 1 month post MI, and at least 3 months post coronary revascularization procedure (Strong Recommendation; High-Quality Evidence); or
- Nonischemic cardiomyopathy, NYHA class II-III, EF \leq 35%, measured at least 3 months after titration and optimization of GDMT (Strong Recommendation; High-Quality Evidence).

52. We recommend against ICD implantation in patients with NYHA class IV symptoms who are not expected to improve with any further therapy and who are not candidates for cardiac transplantation or mechanical circulatory support (MCS) (Strong Recommendation; Moderate-Quality Evidence).

CCS Guidelines 2021

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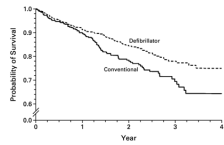
How an implantable cardiac defibrillator works

Type	Seq	ATP	Shock	Therapy	Rhythm	N° ID	Date	On	Duration	Media rate	AVV	EGM complete	Stamp
VF			35J	SI	SI	#7	25-Jul-2018	02:27	:15	---	1563	CCS	

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MADIT II- NEJM 2002

- 1232 patients ischemic cardiomyopathy. ICD vs Conventional therapy



No. at Risk
 Defibrillator 762 608 (81%) 274 (36%) 110 (15%) 0
 Conventional 490 339 (69%) 276 (56%) 81 (16%) 3
 Figure 2. Kaplan-Meier Estimates of the Probability of Survival in the Group Assigned to Receive an Implantable Defibrillator and the Group Assigned to Receive Conventional Medical Therapy. The difference in survival between the two groups was significant (nominal P=0.007, by the log-rank test).

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Case 4

- 48F admitted to hospital with congestive heart failure.
- Echocardiogram: Severe global hypokinesia, LVEF 25%.
- Coronary angiogram: Normal coronaries.
- Patient initiated on GDMT.
 - Entresto
 - Bisoprolol
 - Empagliflozin
 - Spironolactone

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Case 4

- 3 months later, patient remains stable, euvolemic
- NYHA 2.
- Repeat ECHOCARDIOGRAM:
 - LVEF 30%. No significant change
- Cardiac MRI:
 - LVEF 28%. No LGE.
- Diagnosis: Nonischemic cardiomyopathy

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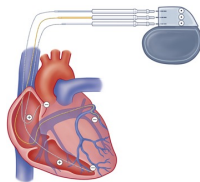
Case 4

- 12 Lead EKG: Sinus rhythm. LBBB.



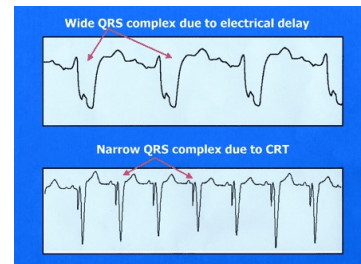
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Cardiac Synchronization Therapy



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CRT leads to improved interventricular synchrony



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Cardiac Resynchronization Therapy

- Benefits:
 - Improved symptoms
 - Reduction of mortality
 - Improved cardiac output
 - Improvement of LVEF
 - ?Reduction of arrhythmia burden



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Cardiac Resynchronization Therapy

- Benefit**
 - NYHA >1
 - LBBB (wider the better)
 - NYHA >1, EF < 50% with AV block (BLOCK-HF trial)
- No Benefit**
 - Narrow QRS (<120msec)
 - RBBB
 - Atrial fibrillation (unless AV nodal ablation)

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RECOMMENDATION

58. We recommend CRT for patients in sinus rhythm with NYHA class II, III, or ambulatory class IV HF despite optimal medical therapy, a LVEF ≤ 35%, and QRS duration ≥ 130 ms with left bundle branch block (LBBB) (Strong Recommendation; High-Quality Evidence).


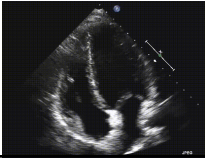
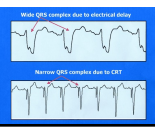
59. We suggest that CRT may be considered for patients in sinus rhythm with NYHA class II, III, or ambulatory class IV HF despite optimal medical therapy, a LVEF ≤ 35%, and QRS duration ≥ 150 ms with non-LBBB (Weak Recommendation; Low-Quality Evidence).

CCS HF guidelines 2021

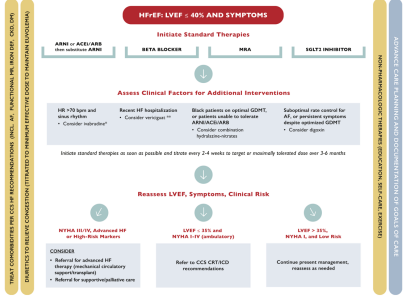
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Case 4

- Patient had successful CRT-d implantation
- 3 months later, patient feeling well.
- Repeat ECHOCARDIOGRAM. LVEF 60%!

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The End

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