

# Managing Heart Failure in 2023 and Beyond - Optimizing Medical and Device Therapy

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September 13, 2023

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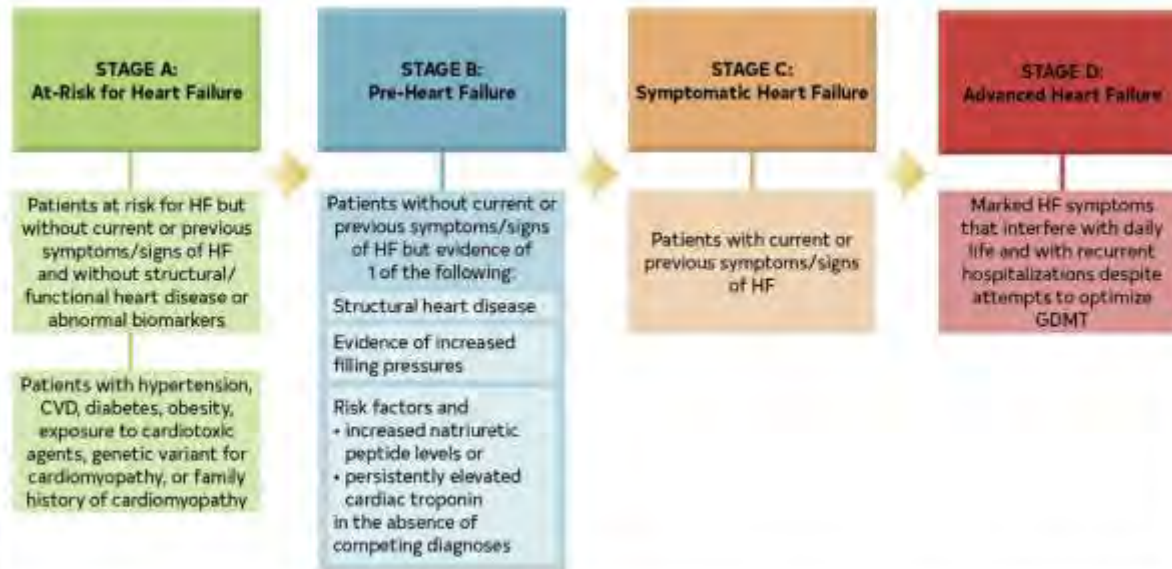
**I have no disclosures**



# Management



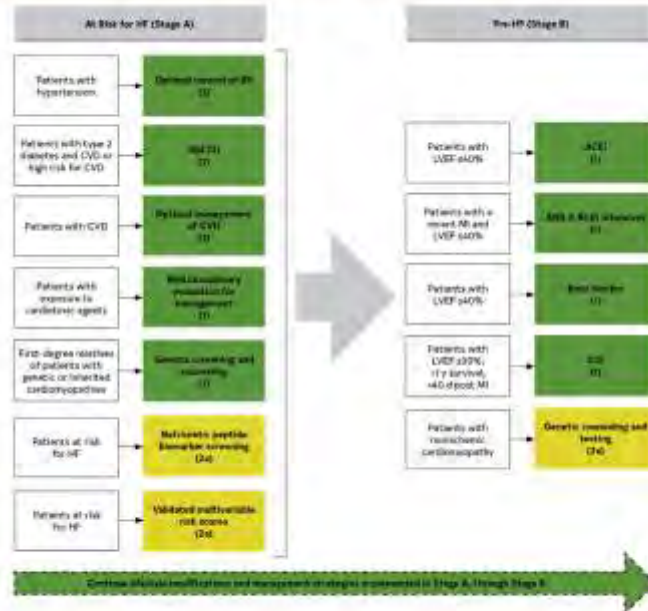
FIGURE 1 ACC/AHA Stages of HF



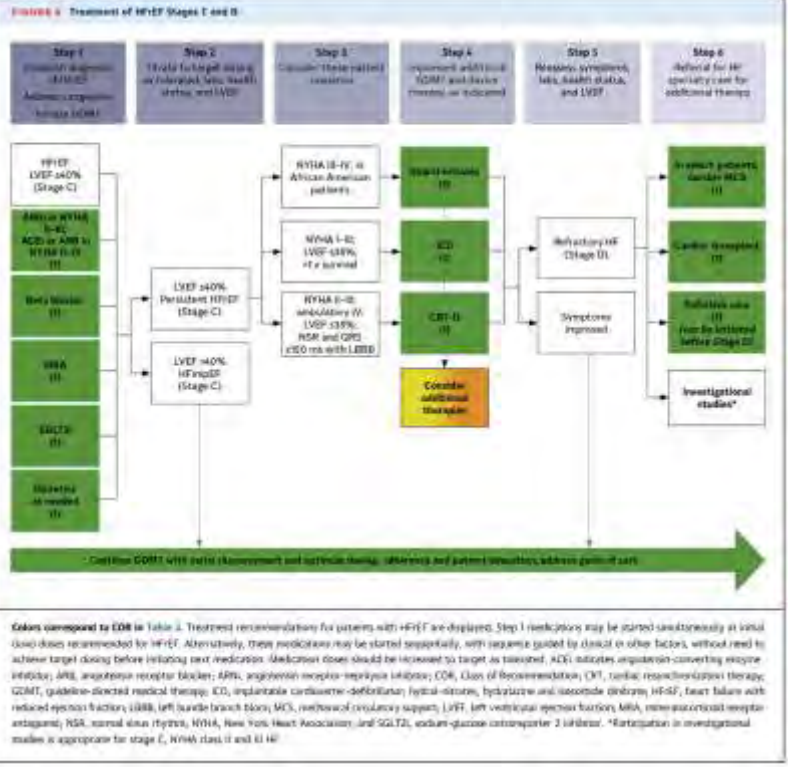
The ACC/AHA stages of HF are shown. ACC indicates American College of Cardiology; AHA, American Heart Association; CVD, cardiovascular disease; GDMT, guideline-directed medical therapy; and HF, heart failure.



**FIGURE 8. Recommendations (Class I and II) for Patients at Risk of HF (Stage A) and Those With Pre-HF (Stage B)**



Colors correspond to CDR in Table 2. CDR 1 and CDR 2a for patients at risk for HF (Stage A) and those with pre-HF (Stage B) are shown. Management strategies implemented in patients at risk for HF (Stage A) should be continued through Stage B. ACEI indicates angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blocker; BP, blood pressure; CDR, Class of Recommendation; CVD, cardiovascular disease; HF, heart failure; KD, potassium-activated adenosine triphosphatase; LVEF, left ventricular ejection fraction; MI, myocardial infarction; and SGLT2, sodium-glucose cotransporter 2 inhibitor.





# 4 Pillars of GDMT





## Beta Blockers

metoprolol succinate

carvedilol

bisoprolol

## Mineralocorticoid blockers

Spironolactone

eplerenone





## Ace Inhibitors / Angiotensin Receptor Antagonist

### ACE-I

Lisinopril

Captopril

enalapril

### ARB

Candesartan

Valsartan

# Angiotensin Receptor/Neprilysin Inhibitor (ARNI)



PARADIGM

PIONEER

PARAGON-HF



# SGLT2 Inhibitors

EMPA-REG

CANVAS

DELIVER

DAPA-HF

EMPEROR-REDUCED

CREDENCE

DAPA-CKD

EMPA-KIDNEY

\*Meta-analysis of SGLT2i studies  
showed a 23% RRR CV death/HFH

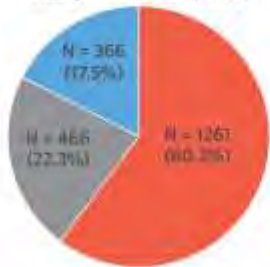
\*Meta-analysis showed 38% RRR in  
kidney disease progression



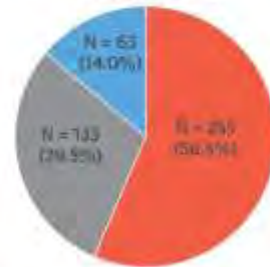
“The estimated cumulative effect of these 4 medications includes a 73% relative reduction in mortality over 2 years. Similarly, compared with conventional dual therapy with a beta-blocker and an angiotensin-converting enzyme inhibitor or angiotensin receptor blocker, treating a 55-year-old patient with quadruple therapy is projected to extend life by > 6 years.”



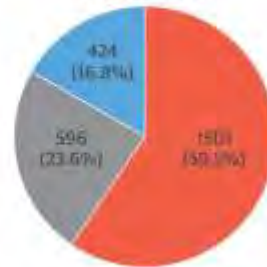
Angiotensin-Converting Enzyme Inhibitor (ACEI)/Angiotensin II Receptor Blocker (ARB)



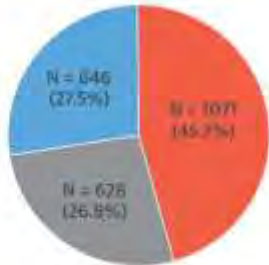
Angiotensin Receptor-Nephrilysin Inhibitor (ARNI)



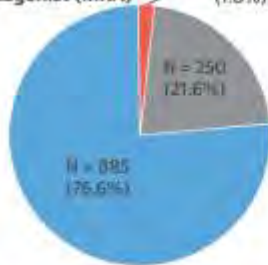
ACEI/ARB/ARNI



Beta-Blocker



Mineralocorticoid Receptor Antagonist (MRA)



■ <50%   ■ 50 to <100%   ■ ≥100%

Greene, S.J. et al. J Am Coll Cardiol. 2018;72(4):351-66.

<25% received all 3 GDMT classes

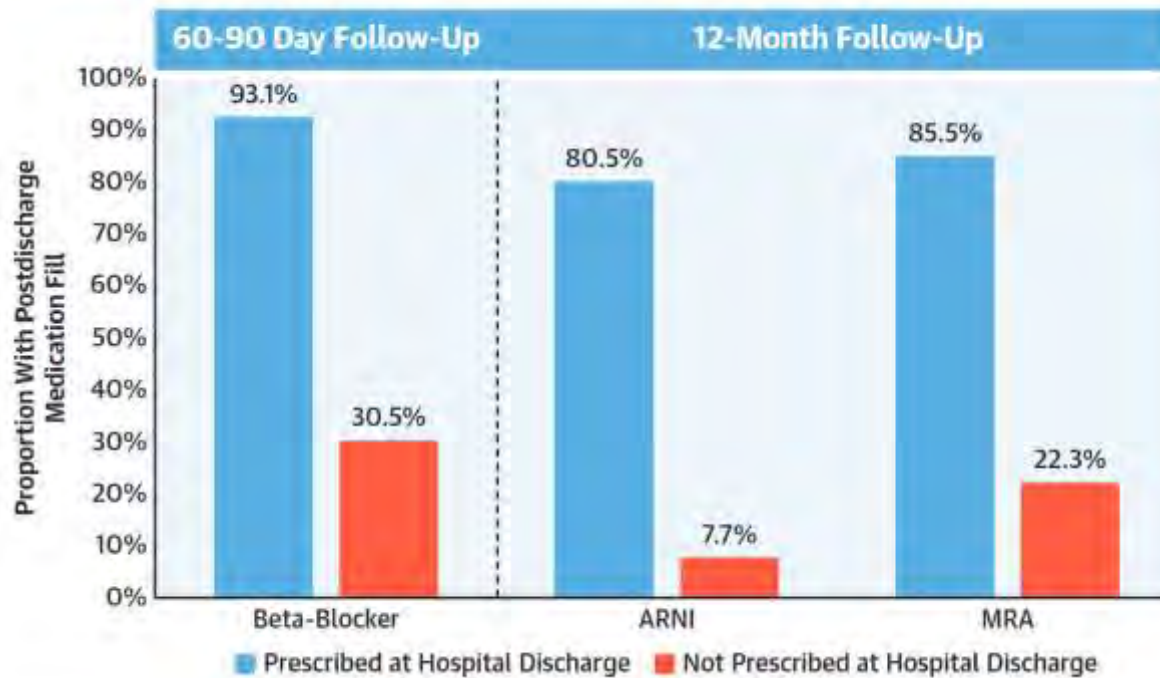
When prescribed, most doses were <50% of target

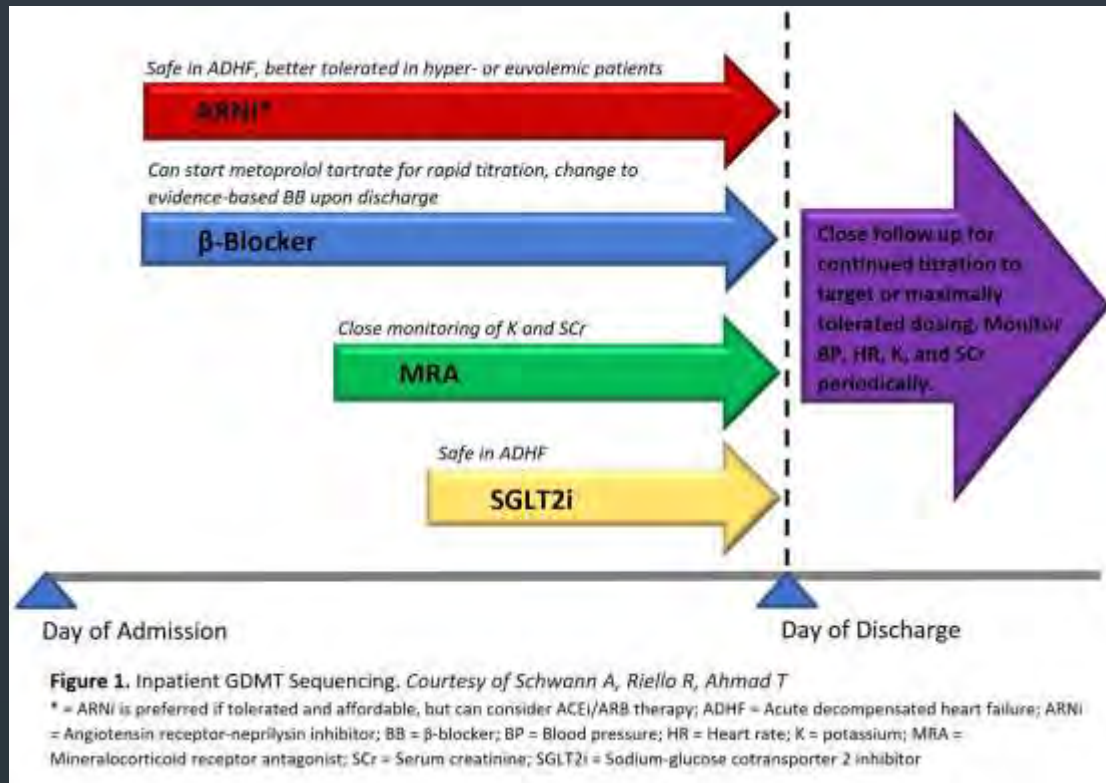
Only 1% received target dose of all three classes

CHARM-HF Study



**FIGURE 2** Postdischarge Use of GDMT by Prescription Status at Hospital Discharge







# STRONG-HF

High intensity Care Group

1, 2, 3 and 6 weeks post hospital follow up  
seen again at 90 days and 180 days

HICG

Lower re-hospitalization for HF

Lower all cause mortality

Improved QOL scores

\*8% ARR of primary outcomes

\*Results consistent across ALL  
LVEF levels

\*No difference in side effect  
profile

\*At 90 days, 36.4% in control  
group on full dose meds vs  
0.4% in standard group





COR	LOE	RECOMMENDATIONS
1	B-NR	1. In patients with HFrEF requiring hospitalization, preexisting GDMT should be continued and optimized to improve outcomes, unless contraindicated (1-5).
1	B-NR	2. In patients experiencing mild decrease of renal function or asymptomatic reduction of blood pressure during HF hospitalization, diuresis and other GDMT should not routinely be discontinued (6-11).
1	B-NR	3. In patients with HFrEF, GDMT should be initiated during hospitalization after clinical stability is achieved (2,3,5,12-18).
1	B-NR	4. In patients with HFrEF, if discontinuation of GDMT is necessary during hospitalization, it should be reinitiated and further optimized as soon as possible (19-22).



**So why are we hesitant to  
uptitrate GDMT aggressively?**



Hypotension

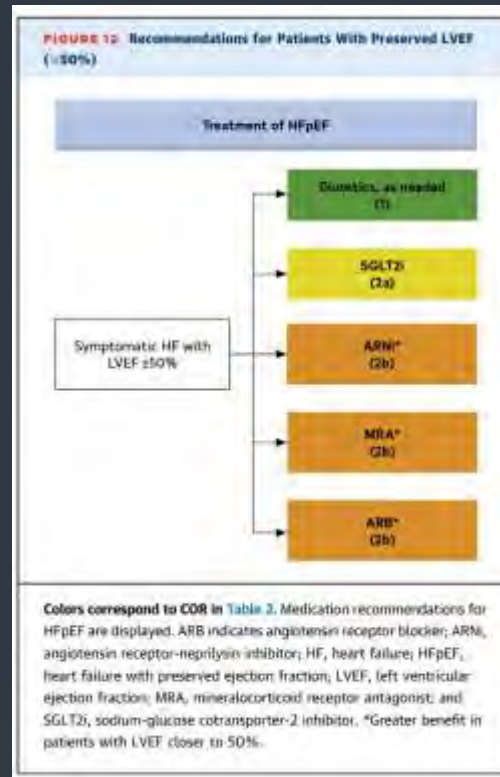
Renal dysfunction

Lack of time

Comfort level

Medication cost/coverage

# HFpEF





COR	LOE	RECOMMENDATIONS
1	B- $\bar{r}$	1. In patients with high-risk HF, particularly those with recurrent hospitalizations for HFrEF, referral to multidisciplinary HF disease management programs is recommended to reduce the risk of hospitalization (1-4).
1	B-NR	2. In patients hospitalized with worsening HF, patient-centered discharge instructions with a clear plan for transitional care should be provided before hospital discharge (5,6).
2a	B-NR	3. In patients hospitalized with worsening HF, participation in systems that allow benchmarking to performance measures is reasonable to increase use of evidence-based therapy, and to improve quality of care (7-10).
2a	B-NR	4. In patients being discharged after hospitalization for worsening HF, an early follow-up, generally within 7 days of hospital discharge, is reasonable to optimize care and reduce rehospitalization (11,12).



# GLP-1 agonists

Semaglutide

approved for chronic weight loss

BMI >30

BMI >27 and one major comorbidity

SELECT

Reduced risk of MACE by 20%



# Outpatient Monitoring Devices



# Cardiomems

Champion and GUIDE-HF

Approved for Class II/III  
with/without hospitaliza  
elevated BNP

reduce hospitalization and  
improve symptoms







# Cardiac Contractility Modulator

Approved for:

NYHA III on optimal GDMT

Not a CRT candidate

LVEF 25-45%

Improvements in 6MWT, QOL, NYHA class





# Others...

ICDs with OptiVol

Wearable defibrillators

External lung fluid monitors...



**Thank You**



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