





Optimizing Guideline-directed Medical Therapies for Heart Failure with Reduced Ejection Fraction During Hospitalization

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Abstract

Heart failure remains a huge societal concern despite medical advancement, with an annual direct cost of over \$30 billion. While guideline-directed medical therapy (GDMT) is proven to reduce morbidity and mortality, many eligible patients with heart failure with reduced ejection fraction (HFrEF) are not receiving one or more of the recommended medications, often due to suboptimal initiation and titration in the outpatient setting. Hospitalization serves as a key point to initiate and titrate GDMT. Four evidence-based therapies have clinical benefit within 30 days of initiation and form a crucial foundation for HFrEF therapy: renin-angiotensin-aldosterone system inhibitors with or without a neprilysin inhibitor, β -blockers, mineralocorticoid-receptor-antagonists, and sodium-glucose cotransporter-2 inhibitors. The authors present a practical guide for the implementation of these four pillars of GDMT during a hospitalization for acute heart failure.

Keywords

Clinical barriers, guideline-directed medical therapy, heart failure hospitalization, acute decompensated heart failure, heart failure with reduced ejection fraction.

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In the US, the annual direct cost of heart failure (HF) is estimated at over \$30 billion, largely driven by the >1 million hospitalizations in which HF is the primary discharge diagnosis.¹⁻³ The subset of patients with HF with reduced ejection fraction (HFrEF) comprises about half of all patients with HF.⁴ Guideline-directed medical therapy (GDMT) is proven to reduce mortality and morbidity for patients with HFrEF. GDMT includes the following drug therapies: renin-angiotensin-aldosterone system inhibitors (RAAS-I), with or without a neprilysin inhibitor, β -blockers, and mineralocorticoid-receptor-antagonists (MRA).⁵

Recently, sodium-glucose cotransporter-2 inhibitors (SGLT2i) demonstrated efficacy as an important fourth pillar of GDMT.⁶ Together, this combination can add over six additional years of lifespan for HFrEF patients compared to the traditional approach of RAAS-I and β -blockers alone.⁷ However, studies highlight that many eligible HFrEF patients are not receiving one or more of the recommended medications, in the absence of contraindications or intolerance.⁸ Even among patients who are treated, less than half receive optimal doses of GDMT. Additionally, time to initiation and optimization of dosing may be exceedingly slow in the outpatient setting.

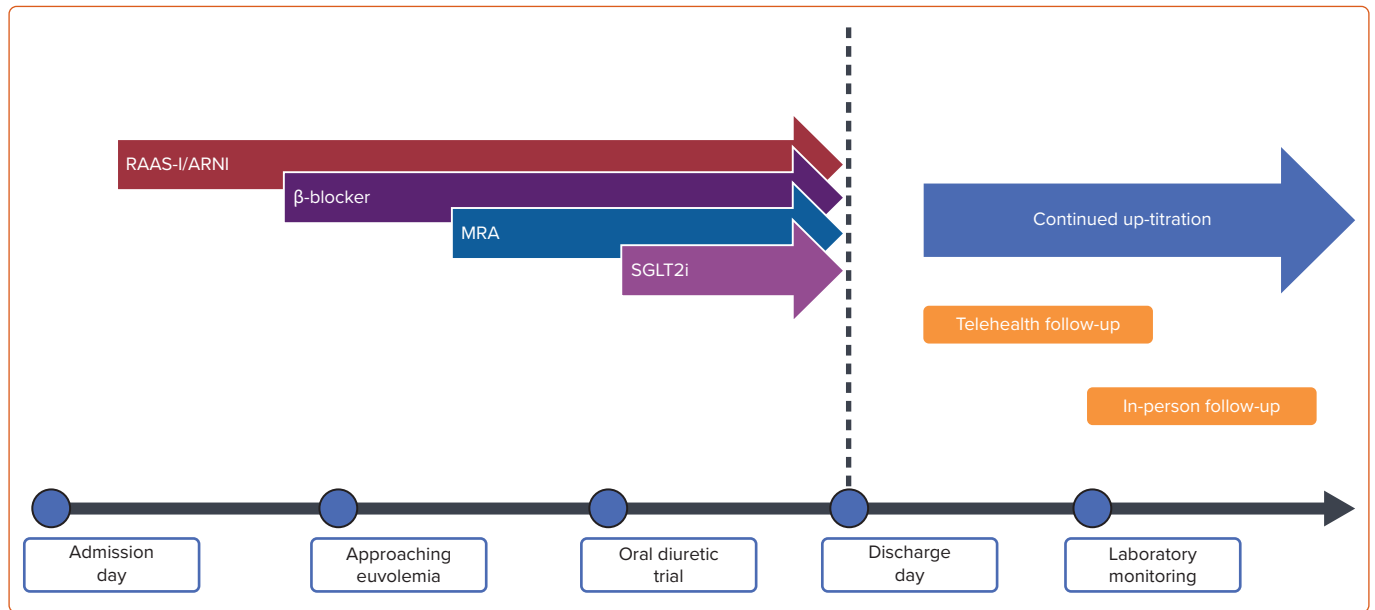
Analysis of the CHAMP registry showed that less than 1% of patients receive target doses of RAAS-I/angiotensin receptor–neprilysin inhibitor (ARNI), β -blockers, and MRA simultaneously over a 12-month period.^{8,9}

Guideline recommendations for device therapy, such as ICD and cardiac resynchronization therapy (CRT), recommend consideration only after 3 months of GDMT optimization. Yet, target doses of GDMT are rarely achieved prior to device therapy implantation.^{5,10} Barriers to optimization include lack of access and coordination of care, gaps in provider knowledge, and patient-related factors such as renal function, perceived intolerance to medications, inadequate insurance coverage, out-of-pocket expenses, and adherence.^{11,12} Hospitalization provides an opportunity to initiate and titrate medical therapy with close monitoring in patients with new-onset or acute on chronic HFrEF.¹³ Medications provided at discharge are more likely to be adhered to, continued, and further titrated in the outpatient setting.¹⁴ An acute heart failure (AHF) hospitalization is an important opportunity to initiate and titrate all four pillars of GDMT (*Figure 1*), laying a complete foundation for further outpatient optimization. We present a practical guide for the initiation and titration of evidence-based chronic HFrEF therapies during hospitalization.

Renin-Angiotensin-Aldosterone-Inhibitor/Angiotensin Receptor–Neprilysin Inhibitor

According to expert consensus, optimization of GDMT in an AHF hospitalization should occur once a positive clinical trajectory has been obtained.¹⁵ Although initially studied in the outpatient setting, angiotensin converting enzyme inhibitors (ACEi) and angiotensin receptor blockers

Figure 1: Shifting the Paradigm of Guideline-directed Medical Therapy Initiation



A suggested timeline of initiating guideline-directed medical therapy (GDMT) for patients admitted with heart failure with reduced ejection fraction during their hospitalization. ACEi = angiotensin converting enzyme inhibitor; ARB = angiotensin receptor blocker; ARNI = angiotensin receptor–neprilysin inhibitor; MRA = mineralocorticoid receptor antagonist; RAAS-I = renin-angiotensin-aldosterone system inhibitor; SGLT2i = sodium–glucose cotransporter-2 inhibitor.

(ARB) have been shown in multiple observational studies to be safely and effectively initiated during hospitalization for AHF, with associated reduction in rehospitalization and mortality.^{16,17}

ARNI is preferred to RAAS-I alone given substantial mortality benefit, similar tolerability, and long-term cost-effectiveness compared to ACEi or ARB.^{18–20} The PARADIGM-HF trial showed a 16% all-cause death reduction and 21% hazard reduction for HF first hospitalization in the ARNI group compared to the ACEi group.²¹ Of the patients entering the ARNI ‘run-in’ phase of the trial, less than 5% dropped out as a result of hypotension, renal dysfunction, and hyperkalemia – the three most common reasons for pre-trial drop-out.²² Patients who were then randomized to ARNI actually had lower rates of discontinuation compared to those randomized to ACEi.²¹ Importantly, symptomatic hypotension is the most common adverse effect of ARNI initiation and occurs about 15% of the time; it is less commonly seen with ACEi or ARB.^{21,23} A detailed review by Sauer et al. provides further practical guidance on ARNI use.²⁴

The PIONEER-HF trial suggested benefits of ARNI initiation *during* hospitalization greater than that achieved with ACEi.²³ Analysis of the trial showed a 31% lower hazard for cardiovascular (CV) death or rehospitalization events for patients that were initiated on an ARNI during hospitalization versus those initiated on the ACEi, enalapril, in the first 8 weeks.²⁵ PIONEER-HF showed that an ARNI can safely be initiated in patients with a systolic blood pressure (SBP) >100 mmHg for 6 hours, no use of vasodilators or increase in dose of intravenous diuretics in the preceding 6 hours, and no use of inotropes in the preceding 24 hours.²³

Contrary to popular belief, RAAS-I/ARNI initiation is well tolerated in cases of mild renal dysfunction, for example, the interquartile range (IQR) of baseline estimated glomerular filtration rate (eGFR) in PIONEER-HF was 47.1–71.5 ml/min/1.73m².^{23,26,27} Acute kidney injury (AKI) after starting RAAS-I occurs in ~13% of HF patients but the long-term overall benefit of RAAS inhibition tends to outweigh any reduction in renal function, which is often transient.²⁸

Experts agree that RAAS-I/ARNI should be continued if serum creatinine (SCr) increases <30% after initiation.²⁹ For changes in SCr >30%, RAAS-I/ARNI doses can be reduced or temporarily held to assess for improvement in renal function. Once renal function begins to improve, RAAS-I/ARNI should be re-trialed to assess maximally tolerated dose.

RAAS-I show modest dose responsiveness. High-dose RAAS-I (equivalent to lisinopril 20–40 mg daily) results in a 6% decrease in all-cause mortality compared to low-dose RAAS-I (equivalent to lisinopril 5–10 mg daily).³⁰ This effect is similar for ARNI with maintained advantage over ACEi or ARB even at lower doses.³¹ Clinicians should aim to initiate a RAAS-I, preferably an ARNI, as soon as clinical stability has been obtained. *Table 1* shows our proposed strategy of RAAS-I/ARNI initiation.

β-blockers

In three landmark trials, β-blockers demonstrated a reduction in all-cause mortality by 34–35%, even among patients with severe HF or recent decompensation.^{32–34} In-hospital initiation of β-blocker therapy for HFrEF has been shown to be safe, well tolerated, and associated with reduced mortality within the first 30 days of hospital discharge.³⁵ This was observed even in patients with signs and symptoms of congestion. β-blockers show the clearest dose response relationship among GDMT.³⁶ Initiating or up-titrating a β-blocker during hospitalization is an effective strategy to increase the likelihood of long-term target dose attainment.³⁷ Martinez-Sellez et al. showed the safety of early initiation of carvedilol in AHF patients at a median 3 days after admission, achieving a mean discharge dose of 23 mg/day among those who tolerated in-hospital up-titration.³⁸ No study has definitively shown the superiority of one evidence-based β-blocker versus another.³⁹ Carvedilol may have a more potent blood pressure lowering effect versus metoprolol succinate in those with more elevated blood pressure.⁴⁰ Although, for patients with lower blood pressure at baseline, β-blockers are hemodynamically well tolerated.⁴¹ In COPERNICUS, patients with a systolic blood pressure (SBP) of 85–95 mmHg at baseline had an increase in SBP with carvedilol greater than placebo.

Table 1: Optimizing Guideline-directed Medical Therapies for Heart Failure with Reduced Ejection Fraction During Hospitalization

Drug	RAAS-I/ARNI	β-blocker	MRA	SGLT2i*
Initiation criteria	<p>ARNI initiation:</p> <ul style="list-style-type: none"> • SBP >100 mmHg for 6 h • No use of IV VD or increase in dose of IV diuretics in the preceding 6 h • No use of inotropes in the preceding 24 h • eGFR >45 ml/min/1.73 m² • K <5.0 mEq/l <p>RAAS-I: See below</p>	<ul style="list-style-type: none"> • No hypoxia, symptomatic hypotension, or evidence of shock 	<ul style="list-style-type: none"> • On at least minimum dose RAAS-I and β-blocker • SCr <2.5 mg/dl in men, <2.0 mg/dl in women • K <5.0 mEq/l • No symptomatic hypotension 	<ul style="list-style-type: none"> • On at least minimum dose RAAS-I and β-blocker • eGFR >30 ml/min/1.73 m² • No symptomatic hypotension
Up-titration strategy	<ul style="list-style-type: none"> • Direct initiation of ARNI preferred strategy • If SBP 100–120 mmHg initiate sacubitril/valsartan 24/26 mg twice daily • If SBP >120 mmHg initiate sacubitril/valsartan 49/51 mg twice daily • Double dose every 1–2 days as tolerated until target dose reached or initiation of next pillar of GDMT 	<ul style="list-style-type: none"> • If SBP 90-120 mmHg or <85 kg start equivalent of carvedilol 3.125 mg twice daily • If SBP >120 mmHg or >85 kg start equivalent of carvedilol 6.25 mg twice daily • Increase every 1–2 days as tolerated until target dose reached 	<ul style="list-style-type: none"> • Initiate at equivalent of spironolactone 12.5 mg daily after the initiation of β-blockers and increase weekly 	<ul style="list-style-type: none"> • Initiate before or after MRA, prior to discharge • No dosage increase required
Potential contraindications	K >5.5 mEq/l	HR <50 BPM	K >5.5 mEq/l, SCr >2.5 mg/dl in men, SCr >2.0 mg/dl in women	T1D, eGFR <30 ml/min/1.73 m ²
Clinical considerations	<ul style="list-style-type: none"> • If SBP is <100 mmHg throughout hospitalization, prior to discharge, trial on equivalent of valsartan 20 mg twice daily or lisinopril 5 mg daily with intent to switch to ARNI when tolerated • No use of ACEi in preceding 36 h of ARNI initiation 	<ul style="list-style-type: none"> • If shock, severe pulmonary edema or SBP <90 mmHg, hold β-blocker and reinitiate at lowest dose, per above • Always ensure patient is adherent to outpatient β-blocker before continuing dose • Younger and heavier patients may tolerate more aggressive dosing • Caution in patients with pulmonary disease 	<ul style="list-style-type: none"> • Recheck of serum potassium within 7 days and within 1–2 months³¹ • Consider addition of potassium-binder if K >5.0 mEq/l 	<ul style="list-style-type: none"> • Consider early initiation and endocrinology consultation for patients with T2D on insulin or sulfonylurea. These patients may require reduced insulin doses and close glycemic monitoring • Consider submitting prior authorization, if required, early in hospitalization • Immediate reduction in eGFR by 4–6 ml/min/1.73m² is expected and not harmful
Relative risk reduction of all-cause mortality in meta-analysis ³²	RAAS-I 20% ARNI 28% ⁺³³	31%	25%	13% ⁶
Relative risk reduction of HF hospitalization in meta-analysis	RAAS-I 33% ³⁴ ARNI 49% ⁺³³	37% ³⁵	23% ³⁶	25% ⁶
Clinical benefits within 30 days of initiation demonstrated?	Yes	Yes	Yes	Yes
Landmark trials	PIONEER-HF, PARADIGM-HF, CHARM, MERIT-HF, COPERNICUS, and CIBIS-II RALES, EMPHASIS-HF, EPHEBUS Val-Heft, ELITE II, ATLAS, SOLVD-2, SOLVD			DAPA-HF, EMPEROR-REDUCED, EMPA-REG

*No consensus guidelines for in-hospital initiation but in-hospital initiation preferred by authors. †Computed versus putative placebo in analysis of PARADIGM-HF Trial. ACEi = angiotensin converting enzyme inhibitor; ARB = angiotensin receptor blocker; ARNI = angiotensin receptor–neprilysin inhibitor; BNP = brain natriuretic peptide; CV = cardiovascular; eGFR = estimated glomerular filtration rate; HF = heart failure; HR = heart rate; MRA = mineralocorticoid receptor antagonist; RAAS-I = renin-angiotensin-aldosterone system inhibitor; SBP = systolic blood pressure; SCr = serum creatinine; SGLT2i = sodium glucose cotransporter-2 inhibitor; T1D = type 1 diabetes; T2D = type 2 diabetes; VD = vasodilator.

β-blockers should not routinely be discontinued on admission for AHF. Multiple studies have shown that discontinuation of β-blockers in AHF is associated with worsening short- and long-term mortality, although hypotension/shock or severe pulmonary edema may represent indications for temporary discontinuation.⁴² Otherwise, as long as the patient confirms

adherence prior to admission, β-blockers can be safely continued during hospitalization.

β-blockers should be initiated or up-titrated so long as the patient is hemodynamically stable without marked volume overload, typically after

Figure 2: Pre-discharge Checklist

Medications	Follow-up	Patient Education
<input type="checkbox"/> GDMT initiation: ACEi/ARB/ARNI, β-blocker, MRA, SGLT2i	<input type="checkbox"/> Telehealth/in-person visit within 1 week	<input type="checkbox"/> Medication education
<input type="checkbox"/> Assessment of oral diuretic efficacy	<input type="checkbox"/> Heart failure clinic referral	<input type="checkbox"/> Nutrition counseling
<input type="checkbox"/> Iron deficiency repletion	<input type="checkbox"/> Labs: creatinine, electrolyte panel, glucose, BNP	<input type="checkbox"/> Physical exercise education
<input type="checkbox"/> Assess for potential drug–drug interactions	<input type="checkbox"/> Cardiac rehabilitation referral	<input type="checkbox"/> Daily weight and blood pressure monitoring
		<input type="checkbox"/> Substance use/tobacco cessation counseling

A suggested pre-discharge checklist to help clinicians ensure that patients who are hospitalized with heart failure with reduced ejection fraction receive optimal guideline-directed medical therapy (GDMT). ACEi = angiotensin converting enzyme inhibitor; ARB = angiotensin receptor blocker; ARNI = angiotensin receptor–neprilysin inhibitor; BNP = brain natriuretic peptide; MRA = mineralocorticoid receptor antagonist; SGLT2i = sodium glucose cotransporter-2 inhibitor.

the initiation of a RAAS-I/ARNI.⁴³ Younger patients with higher BMI and without chronic obstructive pulmonary disease tend to tolerate higher doses and may be appropriate for a higher starting dose or more aggressive up-titration.⁴⁴ Up-titration in patients may occur as tolerated toward target dose prior to discharge (*Table 1*).

Mineralocorticoid Receptor Antagonist

Only one-third of patients indicated for MRA receive them.⁸ Use of these agents has reduced CV death and HF hospitalization, with benefits observed within 30 days of initiation.⁴⁵ Addition of MRA to RAAS-I and β-blockers has been shown to be safe during hospitalization.⁴⁶ Further, long-term adherence is substantially improved with in-hospital initiation compared to delaying therapy to outpatient at clinician discretion.⁴⁷ Hospitalization represents an opportunity to safely initiate an MRA under close monitoring, with provider fear of hyperkalemia and renal function often cited as barriers to initiation.⁴⁸ Potassium binding drugs such as patiomer and sodium zirconium cyclosilicate have shown to be effective long-term options to maintain normokalemia in HF patients.⁴⁹ While these agents do not confer mortality benefit on their own, they are beneficial in patients who would have otherwise been unable to tolerate MRA therapy. Additionally, combined usage of MRA with SGLT2i or ARNI (versus ACEi) lowers the risk of hyperkalemia.^{50,51} At the dosages commonly used in trials, MRAs do not have any blood pressure lowering effect and can be safely initiated on borderline hypotensive patients.⁵² There is no clear dose response relationship with MRA, although the target dose in major trials was the equivalent of 25–50 mg/day of spironolactone.⁵³ We suggest initiating an MRA after RAAS-I/ARNI and β-blocker initiation, but prior to discharge (*Table 1*).

Sodium-Glucose Cotransporter-2 Inhibitor

The DAPA-HF and EMPEROR-Reduced trials showed the significant benefits of SGLT2i in the treatment of patients with HFrEF, including reductions in all-cause mortality risk by 13% and recurrent CV hospitalizations by 25% in HFrEF patients with or without type 2 diabetes (T2DM).⁶ These clinical benefits can be seen within 12 days of initiation.⁵⁴ Additionally, the DAPA-CKD trial showed a 44% decrease in the composite of a sustained decline in eGFR of at least 50%, end-stage kidney disease, or death from renal causes.⁵⁵ Similar results were found in the CREDENCE trial.⁵⁶ With an estimated 54% of HFrEF patients having concomitant CKD and declining renal function as one of the strongest predictors of mortality in HFrEF, SGLT2i hold substantial clinical promise as a vital fourth pillar of GDMT.⁵⁷

The novelty of SGLT2i, along with their potential impact on glycemic control for patients with T2DM contribute to relatively low rates of prescription for HFrEF compared to more established pillars of GDMT.^{58,59} An eGFR <30 ml/min/1.73 m² is one of a few exclusions to starting the medication at this time, but more than 80% of HFrEF patients are eligible

for treatment with SGLT2i.⁷ SGLT2i can be safely initiated during hospitalization and do not need to be up-titrated from its initial dosage. Sotagliflozin, a dual SGLT-1 and -2 inhibitor, displayed benefits within 1 month of initiation when started during or shortly after hospitalization.^{60–62}

SGLT2i do exhibit a modest BP lowering effect in HF patients (~1–4mmHg).^{63,64} Additionally, attention should be paid to glycemic management when initiating SGLT-2i in patients on sulfonylureas or insulin due to risk of hypoglycemia and euglycemic diabetic ketoacidosis.⁶⁴ Volume status of patients on diuretics should be reevaluated due to the risk of over-diuresis. Honigberg et al. previously described in detail the practical considerations for the initiation of SGLT2i. Given the rapid onset of clinical benefit, we favor in-hospital initiation of SGLT2i.⁶⁵ A summary of the important points for in-hospital initiation are presented in *Table 1*.

Hydralazine/Nitrate

Although the hydralazine/nitrate combination was the first treatment to show mortality benefit in HFrEF, its usage varies widely.^{66,67} Current guidelines recommend a hydralazine/nitrate combination for New York Heart Association (NYHA) class III/IV self-described African-American patients who are optimized on RAAS-I and β-blockers.⁵ Because of modest mortality benefits and relatively larger blood pressure effects, the hydralazine/nitrate combination does not need to be prioritized for in-hospital initiation and is not one of the critical four pillars of therapy.^{67,68} In patients who cannot take RAAS-I/ARNI due to renal dysfunction, hyperkalemia, or drug intolerance, the hydralazine/nitrate combination can be considered as alternate therapy, regardless of race.⁵ However, because of the demonstrated superiority of RAAS-I/ARNI, a repeat trial with a RAAS-I/ARNI should be attempted when deemed safe.⁶⁸

Loop Diuretic

Diuretic optimization is an important aspect of chronic HF treatment for symptom management and reducing the likelihood of hospitalization for worsening HF.⁶⁹ Three loop diuretics are commonly used for the treatment of HF: furosemide, bumetanide and torsemide.⁷⁰ No trials have definitively shown superiority of one versus the other; however, furosemide is most commonly used in clinical practice and generally has the lowest acquisition cost.^{71,72} In a meta-analysis, torsemide has been shown to reduce hospitalizations, lower cardiac mortality, and improve functional status compared to furosemide.⁷³ Torsemide also has the longest half-life and offers the advantage of consistent diuretic effect with once daily dosing.⁷¹ Both torsemide and bumetanide have superior bioavailability to furosemide. Guidelines recommend trialing an oral diuretic prior to discharge to verify dose effectiveness.⁵ In a study comparing patients observed on at least 24 hours of an oral diuretic regimen prior to discharge versus those not, 30-, 60-, and 90-day HF readmissions were reduced significantly.⁷⁴ Fluid restriction should be liberalized for an oral diuretic trial to better match the diuretic dose with the patient's unrestricted fluid intake.⁷⁵ The lowest effective dose of loop diuretic should be used to maintain euvoolemia, allowing for maximal up-titration of GDMT.⁷¹

Transition of Care

Although the average length of stay for an AHF admission is 5 days, an early focus on GDMT during the admission can increase the likelihood of initiation of all four agents, especially with increasing system pressures to reduce time in hospital.^{76,77} Following discharge, maintaining momentum and ensuring safety of in-hospital initiation and titration of GDMT is dependent on successful transition of care to the outpatient setting. A systematic review identified eight common components of successful programs: “telephone follow-up, education, self-management, weight

Table 2: Overcoming Common Clinical Barriers to Guideline-directed Medical Therapies Optimization

Barrier to Optimization	First-line Strategy	Second-line Strategy
Acute kidney injury	<ul style="list-style-type: none"> Reduce dose or hold RAAS-I/ARNI. Retrial once renal function improves Reduce diuretic dose to the lowest dose required to maintain euvolesmia 	<ul style="list-style-type: none"> Switch RAAS-I/ARNI to combination of hydralazine/nitrate only after multiple failed trials
Hyperkalemia	<ul style="list-style-type: none"> Remove potassium supplementation Consider addition of potassium-binders and low potassium diet Add SGLT2i Retrial with ARNI (instead of ACEi or ARB) 	<ul style="list-style-type: none"> Reduce or hold doses of RAAS-I/ARNI or MRA. Retrial one at a time Switch RAAS-I/ARNI to combination of hydralazine/nitrate only after multiple failed trials
Symptomatic hypotension	<ul style="list-style-type: none"> Reduce or remove medications that lower BP and are not guideline recommended Stagger doses of GDMT that lower BP (e.g. morning and evening doses) Reduce GDMT dose based on symptoms of hypotension, not blood pressure parameters alone Reduce diuretic dose to the lowest dose to required maintain euvolesmia 	<ul style="list-style-type: none"> Prioritize β-blocker dosage Switch carvedilol to metoprolol succinate Reduce dosage of RAAS-I/ARNI Switch ARNI to ACEi/ARB and retrial with ARNI in future Reduce SGLT2i dose and retrial at regular dose in future
Adherence	<ul style="list-style-type: none"> Medication reminders (e.g. pillboxes, smartphone apps, medication logs). Use once daily medications 	<ul style="list-style-type: none"> Post-discharge telehealth. Refer to HF-specific medication titration clinics
Cost/insurance	<ul style="list-style-type: none"> Submit prior authorization requests early in hospitalization Assess patient willingness and ability to pay and prescribe more affordable medications, if necessary 	<ul style="list-style-type: none"> Periodically reassess availability of new/higher cost medications Perform institution specific cost-effectiveness analysis

ACEi = angiotensin converting enzyme inhibitor; ARB = angiotensin receptor blocker; ARNI = angiotensin receptor–neprilysin inhibitor; BP = blood pressure; GDMT = guideline-directed medical therapy; HF = heart failure; MRA = mineralocorticoid receptor antagonist; RAAS-I = renin-angiotensin-aldosterone system inhibitor; SBP = systolic blood pressure; SGLT2i = sodium glucose cotransporter-2 inhibitor.

monitoring, sodium restriction or dietary advice, exercise recommendations, medication review, and social and psychological support.⁷⁸ Supervised exercise training programs (cardiac rehabilitation) for at least 8 weeks following discharge reduce mortality and rehospitalization.⁷⁹ Iron deficiency should be treated prior to discharge, with one dose of intravenous iron prior to discharge and additional doses as an outpatient shown to improve symptoms and reduce risk of rehospitalization.^{43,80} Additionally, laboratory values, particularly renal function markers, potassium, and glucose, should be monitored within 1–2 weeks of discharge to evaluate for adverse reactions to GDMT initiation.⁵ Brain natriuretic peptide (BNP) or N-terminal pro-hormone BNP (NT-proBNP) levels obtained at 1–2 months after GDMT initiation can be useful predictors of rehospitalization and response to therapy.^{18,81} An assessment of left ventricular systolic function by echocardiography should be performed ~3 months after optimal GDMT has been reached to assess response to therapy and indication for ICD implantation.⁴³ Further, the coronavirus disease 2019 pandemic has highlighted the opportunities

to integrate telehealth to continue the aggressive up-titration of GDMT with the use of specialized nurses and pharmacists.⁸² A suggested pre-discharge checklist is presented in *Figure 2* and strategies to overcome common clinical barriers are presented in *Table 2*.

Conclusion

Given the clinical benefits seen within 30 days of initiation of RAAS-I/ARNI, β -blockers, MRA, and SGLT2i, combined initiation of these four foundational therapies is preferable to the outdated paradigm of pursuing maximally tolerated β -blockers and RAAS-I prior to the addition of secondary agents. Additionally, due to the severe lack of optimization of GDMT in the outpatient setting, in-hospital initiation and titration of the four agents should be attempted in all patients admitted for AHF. The ability to closely monitor in the hospital setting allows for more aggressive up-titration and early recognition of adverse effects. Our strategy prioritizes the implementation of all four pillars of GDMT to meet the evolving standard of optimal treatment of HFREF. □

- Heidenreich PA, Albert NM, Allen LA, et al. Forecasting the impact of heart failure in the United States: a policy statement from the American Heart Association. *Circ Heart Fail* 2013;6:606–19. <https://doi.org/10.1161/HHF.0b013e318291329a>; PMID: 23616602.
- Urbich M, Globe G, Pantiri K, et al. A systematic review of medical costs associated with heart failure in the USA (2014–2020). *Pharmacoeconomics* 2020;38:1219–36. <https://doi.org/10.1007/s40273-020-00952-0>; PMID: 32812149.
- Blecker S, Paul M, Taksler G, et al. Heart failure-associated hospitalizations in the United States. *J Am Coll Cardiol* 2013;61:1259–67. <https://doi.org/10.1016/j.jacc.2012.12.038>; PMID: 23500328.
- Owan TE, Hodge DO, Herges RM, et al. Trends in prevalence and outcome of heart failure with preserved ejection fraction. *N Engl J Med* 2006;355:251–9. <https://doi.org/10.1056/NEJMoa052256>; PMID: 16855265.
- Yancy CW, Jessup M, Bozkurt B, et al. 2013 ACCF/AHA guideline for the management of heart failure: a report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines. *J Am Coll Cardiol* 2013;62:e147–239. <https://doi.org/10.1016/j.jacc.2013.05.019>; PMID: 23747642.
- Zannad F, Ferreira JP, Pocock SJ, et al. SGLT2 inhibitors in patients with heart failure with reduced ejection fraction: a meta-analysis of the EMPEROR-Reduced and DAPA-HF trials. *Lancet* 2020;396:819–29. [https://doi.org/10.1016/S0140-6736\(20\)31824-9](https://doi.org/10.1016/S0140-6736(20)31824-9); PMID: 32877652.
- Vaduganathan M, Claggett BL, Jhund PS, et al. Estimating lifetime benefits of comprehensive disease-modifying pharmacological therapies in patients with heart failure with reduced ejection fraction: a comparative analysis of three randomised controlled trials. *Lancet* 2020;396:121–8. [https://doi.org/10.1016/S0140-6736\(20\)30748-0](https://doi.org/10.1016/S0140-6736(20)30748-0); PMID: 32446323.
- Greene SJ, Butler J, Albert NM, et al. Medical therapy for heart failure with reduced ejection fraction: the CHAMP-HF Registry. *J Am Coll Cardiol* 2018;72:351–66. <https://doi.org/10.1016/j.jacc.2018.04.070>; PMID: 30025570.
- Gracia E, Hamid A, Butler J. Timely management of new-onset heart failure. *Circulation* 2019;140:621–3. <https://doi.org/10.1161/CIRCULATIONAHA.118.035452>; PMID: 31424994.
- Roth GA, Poole JE, Zaha R, et al. Use of guideline-directed medications for heart failure before cardioverter-defibrillator implantation. *J Am Coll Cardiol* 2016;67:1062–9. <https://doi.org/10.1016/j.jacc.2015.12.046>; PMID: 26940927.
- Hancock HC, Close H, Fuat A, et al. Barriers to accurate diagnosis and effective management of heart failure have not changed in the past 10 years: a qualitative study and national survey. *BMJ Open* 2014;4:e003866. <https://doi.org/10.1136/bmjopen-2013-003866>; PMID: 24691215.
- Piamjariyakul U, Yadrich DM, Russell C, et al. Patients' annual income adequacy, insurance premiums and out-of-pocket expenses related to heart failure care. *Heart Lung* 2014;43:469–75. <https://doi.org/10.1016/j.hrtlng.2014.05.013>; PMID: 25012635.
- Yamaguchi T, Kitai T, Miyamoto T, et al. Effect of optimizing guideline-directed medical therapy before discharge on mortality and heart failure readmission in patients hospitalized with heart failure with reduced ejection fraction. *Am J Cardiol* 2018;121:969–74. <https://doi.org/10.1016/j.amjcard.2018.01.006>; PMID: 29477488.
- Fonarow GC, Abraham WT, Albert NM, et al. Prospective evaluation of beta-blocker use at the time of hospital

- discharge as a heart failure performance measure: results from OPTIMIZE-HF. *J Card Fail* 2007;13:722–31. <https://doi.org/10.1016/j.cardfail.2007.06.727>; PMID: 17996820.
15. Hollenberg SM, Stevenson LW, Ahmad T, et al. 2019 ACC expert consensus decision pathway on risk assessment, management, and clinical trajectory of patients hospitalized with heart failure: a report of the American College of Cardiology Solution Set Oversight Committee. *J Am Coll Cardiol* 2019;74:1966–2011. <https://doi.org/10.1016/j.jacc.2019.08.001>; PMID: 31526538.
 16. Gilstrap LG, Fonarow GC, Desai AS, et al. Initiation, continuation, or withdrawal of angiotensin-converting enzyme inhibitors/angiotensin receptor blockers and outcomes in patients hospitalized with heart failure with reduced ejection fraction. *J Am Heart Assoc* 2017;6:e004675. <https://doi.org/10.1161/JAHA.116.004675>; PMID: 28189999.
 17. Sanam K, Bhatia V, Parvataneni S, et al. Discharge initiation of ACE inhibitors or ARBS is associated with significantly lower 30-day all-cause readmission in hospitalized older patients with heart failure and reduced ejection fraction. *J Am Coll Cardiol* 2014;63(12 Suppl):A562. [https://doi.org/10.1016/S0735-1097\(14\)60562-3](https://doi.org/10.1016/S0735-1097(14)60562-3).
 18. DeVore AD, Braunwald E, Morrow DA, et al. Initiation of angiotensin-neprilysin inhibition after acute decompensated heart failure: secondary analysis of the open-label extension of the PIONEER-HF Trial. *JAMA Cardiol* 2020;5:202–7. <https://doi.org/10.1001/jamacardio.2019.4665>; PMID: 31825471.
 19. Gaziano TA, Fonarow GC, Velazquez EJ, et al. Cost-effectiveness of sacubitril-valsartan in hospitalized patients who have heart failure with reduced ejection fraction. *JAMA Cardiol* 2020;5:1236–44. <https://doi.org/10.1001/jamacardio.2020.2822> PMID: 32785628.
 20. Gaziano TA, Fonarow GC, Claggett B, et al. Cost-effectiveness analysis of sacubitril/valsartan vs enalapril in patients with heart failure and reduced ejection fraction. *JAMA Cardiol* 2016;1:666–72. <https://doi.org/10.1001/jamacardio.2016.1747>; PMID: 27438344.
 21. McMurray JJ, Packer M, Desai AS, et al. Angiotensin-neprilysin inhibition versus enalapril in heart failure. *N Engl J Med* 2014;371:993–1004. <https://doi.org/10.1056/NEJMoa1409077>; PMID: 25176015.
 22. Desai AS, Solomon S, Claggett B, et al. Factors associated with noncompletion during the run-in period before randomization and influence on the estimated benefit of LCZ696 in the PARADIGM-HF trial. *Circ Heart Fail* 2016;9:e002735. <https://doi.org/10.1161/CIRCHEARTFAILURE.115.002735>; PMID: 27296397.
 23. Velazquez EJ, Morrow DA, DeVore AD, et al. Angiotensin-neprilysin inhibition in acute decompensated heart failure. *N Engl J Med* 2019;380:539–48. <https://doi.org/10.1056/NEJMoa1812851>; PMID: 30415601.
 24. Sauer AJ, Cole R, Jensen BC, et al. Practical guidance on the use of sacubitril/valsartan for heart failure. *Heart Fail Rev* 2019;24:167–76. <https://doi.org/10.1007/s10741-018-9757-1>; PMID: 30565021.
 25. DeVore AD, Braunwald E, Morrow DA, et al. Initiation of angiotensin-neprilysin inhibition after acute decompensated heart failure: secondary analysis of the open-label extension of the PIONEER-HF trial. *JAMA Cardiol* 2020;5:202–7. <https://doi.org/10.1001/jamacardio.2019.4665>; PMID: 31825471.
 26. Akerman CC, Beavers JC. Risk factors for intolerance of inpatient sacubitril/valsartan initiation. *J Pharm Pract* 2019. <https://doi.org/10.1177/0897190019878948>; PMID: 31581926; epub ahead of press.
 27. Kozuharov N, Goudev A, Flores D, et al. Effect of a strategy of comprehensive vasodilation vs usual care on mortality and heart failure rehospitalization among patients with acute heart failure: the GALACTIC randomized clinical trial. *JAMA* 2019;322:2292–302. <https://doi.org/10.1001/jama.2019.18598>; PMID: 31846016.
 28. Beldhuis IE, Streng KW, Ter Maaten JM, et al. Renin-angiotensin system inhibition, worsening renal function, and outcome in heart failure patients with reduced and preserved ejection fraction: a meta-analysis of published study data. *Circ Heart Fail* 2017;10. <https://doi.org/10.1161/CIRCHEARTFAILURE.116.003588>; PMID: 28209765.
 29. Clark AL, Kalra PR, Petrie MC, et al. Change in renal function associated with drug treatment in heart failure: national guidance. *Heart* 2019;105:904–10. <https://doi.org/10.1136/heartjnl-2018-314158>; PMID: 3118203.
 30. Khan MS, Fonarow GC, Ahmed A, et al. Dose of angiotensin-converting enzyme inhibitors and angiotensin receptor blockers and outcomes in heart failure: a meta-analysis. *Circ Heart Fail* 2017;10. <https://doi.org/10.1161/CIRCHEARTFAILURE.117.003956>; PMID: 28790052.
 31. Vardeny O, Claggett B, Packer M, et al. Efficacy of sacubitril/valsartan vs. enalapril at lower than target doses in heart failure with reduced ejection fraction: the PARADIGM-HF trial. *Eur J Heart Fail* 2016;18:1228–34. <https://doi.org/10.1002/ehfj.580>; PMID: 27283779.
 32. Packer M, Fowler MB, Roecker EB, et al. Effect of carvedilol on the morbidity of patients with severe chronic heart failure: results of the carvedilol prospective randomized cumulative survival (COPERNICUS) study. *Circulation* 2002;106:2194–9. <https://doi.org/10.1161/01.CIR.0000035653.72855.BF>; PMID: 12390947.
 33. The Cardiac Insufficiency Bisoprolol Study II (CIBIS-II): a randomised trial. *Lancet* 1999;353:9–13. [https://doi.org/10.1016/S0140-6736\(98\)1181-9](https://doi.org/10.1016/S0140-6736(98)1181-9); PMID: 10023943.
 34. Effect of metoprolol CR/XL in chronic heart failure: Metoprolol CR/XL Randomised Intervention Trial in Congestive Heart Failure (MERIT-HF). *Lancet* 1999;353:2001–7. [https://doi.org/10.1016/S0140-6736\(99\)04440-2](https://doi.org/10.1016/S0140-6736(99)04440-2); PMID: 10376614.
 35. Bhatia V, Bajaj NS, Sanam K, et al. Beta-blocker use and 30-day all-cause readmission in medicare beneficiaries with systolic heart failure. *Am J Med* 2015;128:715–21. <https://doi.org/10.1016/j.amjmed.2014.11.036>; PMID: 25554369.
 36. Fuzat M, Wojdyla D, Kitzman D, et al. Relationship of beta-blocker dose with outcomes in ambulatory heart failure patients with systolic dysfunction: results from the HF-ACTION (Heart Failure: A Controlled Trial Investigating Outcomes of Exercise Training) trial. *J Am Coll Cardiol* 2012;60:208–15. <https://doi.org/10.1016/j.jacc.2012.03.023>; PMID: 22560018.
 37. Gattis WA, O'Connor CM, Gallup DS, et al. Predischarge initiation of carvedilol in patients hospitalized for decompensated heart failure: results of the Initiation Management Predischarge: Process for Assessment of Carvedilol Therapy in Heart Failure (IMPACT-HF) trial. *J Am Coll Cardiol* 2004;43:1534–41. <https://doi.org/10.1016/j.jacc.2003.12.040>; PMID: 15120808.
 38. Martinez-Selles M, Datino T, Alhama M, et al. Rapid carvedilol up-titration in hospitalized patients with systolic heart failure. *J Heart Lung Transplant* 2008;27:914–16. <https://doi.org/10.1016/j.healun.2008.05.008>; PMID: 18656807.
 39. Fröhlich H, Zhao J, Täger T, et al. Carvedilol compared with metoprolol succinate in the treatment and prognosis of patients with stable chronic heart failure: Carvedilol or Metoprolol Evaluation Study. *Circ Heart Fail* 2015;8:887–96. <https://doi.org/10.1161/CIRCHEARTFAILURE.114.001701>; PMID: 26175538.
 40. Franz IW, Agrawal B, Biewel D, Ketelhut R. Comparison of the antihypertensive effects of carvedilol and metoprolol on resting and exercise blood pressure. *Clin Investig* 1992;70(Suppl 1):S53–7. <https://doi.org/10.1007/BF00207612>; PMID: 1350485.
 41. Rouleau JL, Roecker EB, Tendera M, et al. Influence of pretreatment systolic blood pressure on the effect of carvedilol in patients with severe chronic heart failure: the Carvedilol Prospective Randomized Cumulative Survival (COPERNICUS) study. *J Am Coll Cardiol* 2004;43:1423–9. <https://doi.org/10.1016/j.jacc.2003.11.037>; PMID: 15093878.
 42. Prins KW, Neill JM, Tyler JO, et al. Effects of beta-blocker withdrawal in acute decompensated heart failure: a systematic review and meta-analysis. *JACC Heart Failure* 2015;3:647–53. <https://doi.org/10.1016/j.jchf.2015.03.008>; PMID: 26251094.
 43. Yancy CW, Januzzi Jr JL, Allen LA, et al. 2017 ACC Expert Consensus Decision Pathway for Optimization of Heart Failure Treatment: answers to 10 pivotal issues about heart failure with reduced ejection fraction: a report of the American College of Cardiology Task Force on Expert Consensus Decision Pathways. *J Am Coll Cardiol* 2018;71:201–30. <https://doi.org/10.1016/j.jacc.2017.11.025>; PMID: 29277252.
 44. Cohen-Solal A, Jacobson AF, Piña IL. Beta blocker dose and markers of sympathetic activation in heart failure patients: interrelationships and prognostic significance. *ESC Heart Fail* 2017;4:499–506. <https://doi.org/10.1002/ehf2.12153>; PMID: 29154422.
 45. Rossi R, Crupi N, Coppi F, et al. Importance of the time of initiation of mineralocorticoid receptor antagonists on risk of mortality in patients with heart failure. *J Renin Angiotensin Aldosterone Syst* 2015;16:119–25. <https://doi.org/10.1177/1470320313482603>; PMID: 23539659.
 46. Dürstelndorf MS, Katz SD, Park H, Blecker S. Mineralocorticoid receptor antagonist use after hospitalization of patients with heart failure and post-discharge outcomes: a single-center retrospective cohort study. *BMC Cardiovasc Disord* 2019;19:194. <https://doi.org/10.1186/s12872-019-1175-3>; PMID: 31399059.
 47. Wirtz HS, Sheer R, Honarpour N, et al. Real-world analysis of guideline-based therapy after hospitalization for heart failure. *J Am Heart Assoc* 2020;9:e015042. <https://doi.org/10.1161/JAHA.119.015042>; PMID: 32805817.
 48. Ferreira JP, Rossignol P, Machu JL, et al. Mineralocorticoid receptor antagonist pattern of use in heart failure with reduced ejection fraction: findings from BIOSAT-CHF. *Eur J Heart Fail* 2017;19:1284–93. <https://doi.org/10.1002/ehfj.900>; PMID: 28580625.
 49. Sidhu K, Sanjanwala R, Zieroth S. Hyperkalemia in heart failure. *Curr Opin Cardiol* 2020;35:150–5. <https://doi.org/10.1097/HCO.0000000000000709>; PMID: 31833959.
 50. Desai AS, Vardeny O, Claggett B, et al. Reduced risk of hyperkalemia during treatment of heart failure with mineralocorticoid receptor antagonists by use of sacubitril/valsartan compared with enalapril: a secondary analysis of the PARADIGM-HF Trial. *JAMA Cardiol* 2017;2:79–85. <https://doi.org/10.1001/jamacardio.2016.4733>; PMID: 27842179.
 51. Kristensen SL, Docherty KF, Jhund PS, et al. Dapagliflozin reduces the risk of hyperkalaemia in patients with heart failure and reduced ejection fraction: a secondary analysis DAPA-HF. *Eur Heart J* 2020;41(Suppl 2):ehaa946.0939. <https://doi.org/10.1093/ehjci/ehaa946.0939>.
 52. Bazoukis G, Thomopoulos C, Tse G, Tsioufis C. Is there a blood pressure lowering effect of MRAs in heart failure? An overview and meta-analysis. *Heart Fail Rev* 2018;23:547–53. <https://doi.org/10.1007/s10741-018-9689-9>; PMID: 29527640.
 53. Berbenetz NM, Mrkobrada M. Mineralocorticoid receptor antagonists for heart failure: systematic review and meta-analysis. *BMC Cardiovasc Disord* 2016;16:246. <https://doi.org/10.1186/s12872-016-0425-x>; PMID: 27905877.
 54. Packer M, Anker SD, Butler J, et al. Effect of Empagliflozin on the Clinical Stability of Patients with Heart Failure and a Reduced Ejection Fraction: the EMPEROR-Reduced Trial. *Circulation* 2021;143:326–336. <https://doi.org/10.1161/CIRCULATIONAHA.120.051783>; PMID: 33081531.
 55. Wheeler DC, Stefanesson BV, Batishian M, et al. The Dapagliflozin and Prevention of Adverse Outcomes in Chronic Kidney Disease (DAPA-CKD) trial: baseline characteristics. *Nephrol Dial Transplant* 2020;35:1700–11. <https://doi.org/10.1093/ndt/gfaa234>; PMID: 32862232.
 56. Perkovic V, Jardine MJ, Neal B, et al. Canagliflozin and renal outcomes in type 2 diabetes and nephropathy. *N Engl J Med* 2019;380:2295–306. <https://doi.org/10.1056/NEJMoa1811744>; PMID: 30990260.
 57. McAlister FA, Ezekowitz J, Tarantini L, et al. Renal dysfunction in patients with heart failure with preserved versus reduced ejection fraction: impact of the new Chronic Kidney Disease-Epidemiology Collaboration Group formula. *Circ Heart Fail* 2012;5:309–14. <https://doi.org/10.1161/CIRCHEARTFAILURE.111.966242>; PMID: 22441773.
 58. Vaduganathan M, Sathiyakumar V, Singh A, et al. Prescriber patterns of SGLT2i after expansions of U.S. Food and Drug Administration labeling. *J Am Coll Cardiol* 2018;72:3370–2. <https://doi.org/10.1016/j.jacc.2018.08.2202>; PMID: 30409566.
 59. Gao Y, Peterson E, Pagidipati N. Barriers to prescribing glucose-lowering therapies with cardiometabolic benefits. *Am Heart J* 2020;224:47–53. <https://doi.org/10.1016/j.ahj.2020.03.017>; PMID: 32304879.
 60. Bhatt DL, Szarek M, Steg PG, et al. Sotagliflozin in patients with diabetes and recent worsening heart failure. *N Engl J Med*. 2021;384:117–28. <https://doi.org/10.1056/NEJMoa2030183>; PMID: 33200892.
 61. Bhatt DL, Szarek M, Pitt B, et al. Sotagliflozin in patients with diabetes and chronic kidney disease. *N Engl J Med* 2021;384:129–39. <https://doi.org/10.1056/NEJMoa2030186>; PMID: 33200891.
 62. Damman K, Beusekamp JC, Boersma EM, et al. Randomized, double-blind, placebo-controlled, multicentre pilot study on the effects of empagliflozin on clinical outcomes in patients with acute decompensated heart failure (EMPA-RESPONSE-AHF). *Eur J Heart Fail* 2020;22:713–22. <https://doi.org/10.1002/ehfj.1713>; PMID: 31912605.
 63. McMurray JJV, Solomon SD, Inzucchi SE, et al. Dapagliflozin in patients with heart failure and reduced ejection fraction. *N Engl J Med* 2019;381:1995–2008. <https://doi.org/10.1056/NEJMoa1913003>; PMID: 31535829.
 64. Honigberg MC, Vardeny O, Vaduganathan M. Practical considerations for the use of sodium-glucose co-transporter 2 inhibitors in heart failure. *Circ Heart Fail* 2020;13:e006623. <https://doi.org/10.1161/CIRCHEARTFAILURE.119.006623>; PMID: 32059632.
 65. Berg DD, Jhund PS, Docherty KF, et al. Time to clinical benefit of dapagliflozin and significance of prior heart failure hospitalization in patients with heart failure with reduced ejection fraction. *JAMA Cardiol* 2021. <https://doi.org/10.1001/jamacardio.2020.7585>; PMID: 33595593; epub ahead of press.
 66. Cohn JN, Archibald DG, Ziesche S, et al. Effect of vasodilator therapy on mortality in chronic congestive heart failure. Results of a Veterans Administration Cooperative Study. *N Engl J Med* 1986;314:1547–52. <https://doi.org/10.1056/NEJM198606123142404>; PMID: 3520315.
 67. Al-Mohammad A. Hydralazine and nitrates in the treatment of heart failure with reduced ejection fraction. *ESC Heart Fail* 2019;6:878–83. <https://doi.org/10.1002/ehf2.12459>; PMID: 3119890.
 68. Farag M, Mabote T, Shoib A, et al. Hydralazine and nitrates

- alone or combined for the management of chronic heart failure: a systematic review. *Int J Cardiol* 2015;196:61–9. <https://doi.org/10.1016/j.ijcard.2015.05.160>; PMID: 26073215.
69. Faris R, Flather MD, Purcell H, et al. Diuretics for heart failure. *Cochrane Database Syst Rev* 2006;(1):CD003838. <https://doi.org/10.1002/14651858.CD003838.pub2>; PMID: 16437464.
 70. Buggy J, Mentz RJ, Pitt B, et al. A reappraisal of loop diuretic choice in heart failure patients. *Am Heart J* 2015;169:323–33. <https://doi.org/10.1016/j.ahj.2014.12.009>; PMID: 25728721.
 71. Felker GM, Ellison DH, Mullens W, et al. *J Am Coll Cardiol* 2020;75:1178–95. <https://doi.org/10.1016/j.jacc.2019.12.059>; PMID: 32164892.
 72. Mentz RJ, Hasselblad V, DeVore AD, et al. Torsemide versus furosemide in patients with acute heart failure (from the ASCEND-HF trial). *Am J Cardiol* 2016;117:404–11. <https://doi.org/10.1016/j.amjcard.2015.10.059>; PMID: 26704029.
 73. Abraham B, Megaly M, Sous M, et al. Meta-analysis comparing torsemide versus furosemide in patients with heart failure. *Am J Cardiol* 2020;125:92–9. <https://doi.org/10.1016/j.amjcard.2019.09.039>; PMID: 31699358.
 74. Laliberte B, Reed BN, Devabhakthuni S, et al. Observation of patients transitioned to an oral loop diuretic before discharge and risk of readmission for acute decompensated heart failure. *J Card Fail* Oct 2017;23:746–52. <https://doi.org/10.1016/j.cardfail.2017.06.008>; PMID: 28688888.
 75. Li Y, Fu B, Qian X. Liberal versus restricted fluid administration in heart failure patients. A systematic review and meta-analysis of randomized trials. *Int Heart J* 2015;56:192–5. <https://doi.org/10.1536/ihj.14-288>; PMID: 25740394.
 76. Sud M, Yu B, Wijeyesundera HC, et al. Associations between short or long length of stay and 30-day readmission and mortality in hospitalized patients with heart failure. *JACC Heart Fail* 2017;5:578–88. <https://doi.org/10.1016/j.jchf.2017.03.012>; PMID: 28501521.
 77. Samsky MD, Ambrosy AP, Youngson E, et al. Trends in readmissions and length of stay for patients hospitalized with heart failure in Canada and the United States. *JAMA Cardiol* 2019;4:444–53. <https://doi.org/10.1001/jamacardio.2019.0766>; PMID: 30969316.
 78. Takeda A, Taylor SJ, Taylor RS, et al. Clinical service organisation for heart failure. *Cochrane Database Syst Rev* 2012;(9):CD002752. <https://doi.org/10.1002/14651858.CD002752.pub3>; PMID: 22972058.
 79. Piepoli MF, Davos C, Francis DP, Coats AJ. Exercise Training Meta-analysis of Trials in Patients with Chronic Heart Failure (ExTraMATCH). *BMJ* 2004;328:189. <https://doi.org/10.1136/bmj.37938.645220.EE>; PMID: 14729656.
 80. Ponikowski P, Kirwan B-A, Anker SD, et al. Ferric carboxymaltose for iron deficiency at discharge after acute heart failure: a multicentre, double-blind, randomised, controlled trial. *Lancet* 2020;396:1895–904. [https://doi.org/10.1016/S0140-6736\(20\)32339-4](https://doi.org/10.1016/S0140-6736(20)32339-4); PMID: 33197395.
 81. Omar HR, Guglin M. Post-discharge rise in BNP and rehospitalization for heart failure. *Herz* 2019;44:450–4. <https://doi.org/10.1007/s00059-018-4687-1>; PMID: 29516117.
 82. Thibodeau JT, Gorodeski EZ. Telehealth for uptitration of guideline-directed medical therapy in heart failure. *Circulation* 2020;142:1507–9. <https://doi.org/10.1161/CIRCULATIONAHA.120.050582>; PMID: 33074759.