## HEART FAILURE HOSPITALIZATION PATHWAY TOOLKIT: KEY TABLES AND FIGURES FOR POINT-OF-CARE

This toolkit serves as a companion to the **2019 ACC Expert Consensus Decision Pathway on Risk Assessment, Management, and Clinical Trajectory of Patients Hospitalized with Heart Failure** and its **2024 Focused Update.** 

The goal of the pathway is to help clinicians consider the short-term and long-term outlook for their patients hospitalized with heart failure (HF)—to institute therapies to reduce symptoms and optimize outcomes, to ensure that those plans are conveyed clearly to caregivers after discharge, and to engage patients to share in decisions and become active participants in their care.

The toolkit provides the tables and figures from the documents, adapted to help clinicians implement key principles from the pathway at the point of care by posting them for reference, filling in provided forms and checklists to help standardize processes, or using these figures as templates for your institution's EHR programming teams.



## Heart Failure Hospitalization Pathway Toolkit: Key Tables and Figures for Point-of-Care

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## Major Nodes and Tools

## Roadmap for Risk Assessment, Management, and Clinical Trajectory of Patients Hospitalized with Heart Failure

Node

## ools

TRIAGE IN ED

ADMISSION

DAILY TRAJECTORY CHECK TRANSITION TO ORAL THERAPIES

DISCHARGE DAY POST DISCHARGE FOLLOW-UP

- Risk Stratification of Acute HF in the ED (Figure 2)
- Predictors of Risk in Emergency (Table 1)
- Factors Contributing to HF Decompenstation (Table 2)
- Clinical Evidence of Congestion (<u>Table 3</u>)
- Hemodynamic Profiles (<u>Figure 3</u>)
- Comorbidities to Consider (<u>Table 4</u>)
- Risk Assessment During Hospitalization (<u>Table 5</u>)
- Interventions for Patients at High Risk (Table 6)

- Trajectory
   Assessment and
   Next Steps
   (Figure 4)
- Evaluation of Initial and Residual Congestion (Figure 5)
- Diuretic Therapy by Trajectory (Figure 6)
- Clinical Trajectories (<u>Figure 7a</u>)
- Diuretic Dosing (Table 7)

- Risk Assessment During Hospitalization (<u>Table 5</u>)
- Titration of GDMT for HFrEF (Figure 7b)
- Education for Patients and Caregivers (Figure 10)
- Focused Discharge Handoff (Figure 11)
- Checklist for Communication to Continuing Care Providers (Figure 12)
- Checklist for Follow-up Phone Call within 48-72 Hours (Figure 13)
- First post-Discharge Visit Checklist (Figure 14)

## PALLIATIVE CARE

- Aspects of Palliative Care (Figure 15)
- Goals of Care/Advanced Care Planning (Table 9)







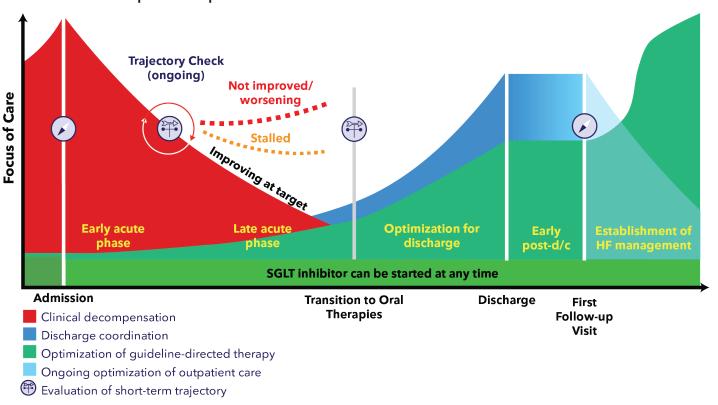
## Pathway Summary Graphic

## Figure 1

This shows the degree of focus on clinical decompensation (*red*), discharge coordination (*blue*), ongoing coordination of outpatient care (*light blue*), and optimization of guideline-directed medical therapy (*green*), with ongoing assessment of the clinical course (*circle with arrows*), and key time points for review and revision of the long-term disease trajectory for the HF journey (*compass signs*).

For optimization of guideline directed medical therapy, refer to the 2024 ECDP for Optimization of Heart Failure Treatment and the 2022 ACCF/AHA/HFSA Heart Failure Guidelines.

## Graphic Depiction of Course of Heart Failure Admission







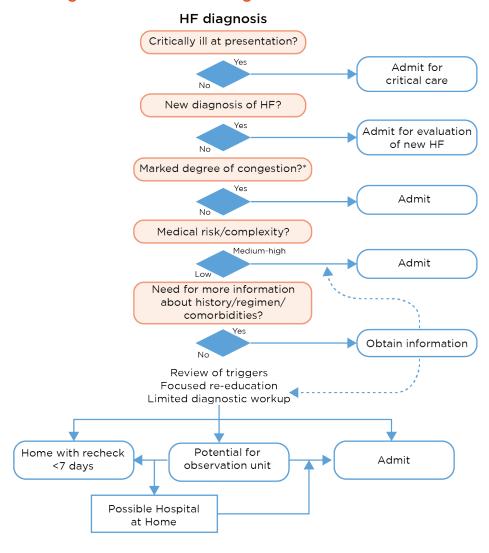


## Risk Stratification of Acute Heart Failure in the Emergency Department

## Figure 2

Use this triage algorithm about admission and initial therapy to guide thought processes during admission evaluations rather than as a formal description of admission criteria and administrative processes surrounding admission.

## **High Likelihood of HF Diagnosis**



<sup>\*</sup> Marked leg edema, ascites, or scrotal or perineal edema may be clinical signs of marked congestion. The degree of radiographic and biochemical abnormalities may also indicate the degree of congestion.

Abbreviations: ED = emergency department; HF = heart failure







## Predictors of Risk in Emergency Care Studies Evaluating Patients With Acute Heart Failure

## **Table 1**Use this table to evaluate a patient's risk and help determine additional next steps.

Immediate Risk (measures of acute severity)	<ul> <li>☐ Hypoxia</li> <li>☐ Shock/hypoperfusion</li> <li>☐ Respiratory distress</li> <li>☐ Anuria</li> <li>☐ Acute and worsening condition (sepsis, stroke, acute coronary syndrome, hemodynamically significant arrhythmia)</li> </ul>
Intermediate Risk (predictors of events through 30 days)	<ul> <li>New onset heart failure</li> <li>Low BP without shock or hypoperfusion</li> <li>□ Tachycardia</li> <li>□ Kidney dysfunction</li> <li>□ Hyponatremia</li> <li>□ Elevated cardiac troponin without ACS</li> <li>□ Degree of BNP elevation</li> <li>□ Liver dysfunction</li> </ul>
Lower Risk	<ul> <li>□ Normal BP and HR</li> <li>□ Brisk response to initial intravenous diuretic with diuresis and symptom relief</li> <li>□ Rapid resolution of symptoms in the ED</li> <li>□ Normal kidney and liver function without recent decline</li> <li>□ Normal BNP and cardiac troponin</li> </ul>

ACS = acute coronary syndrome; BP = blood pressure; BNP = B-type natriuretic peptide; HR = heart rate; ED = emergency department







## Common Factors That Can Contribute to Worsening Heart Failure

## Table 2

Use this table to support the evaluation of patients for factors, both cardiac and non-cardiac, that may contribute to worsening heart failure.

Common Factors That Can Contribute to Worsening Heart Failure
☐ Acute myocardial ischemia
☐ Uncontrolled hypertension
☐ Atrial fibrillation and other arrhythmias
☐ Medications with negative inotropic effect
☐ Medications that increase sodium retention (NSAIDs, thiazolidinediones, steroids)
☐ Non-adherence with medication regimen, sodium or fluid restriction
□ Anemia
☐ Acute infections (upper respiratory infection, pneumonia, urinary tract infections)
☐ Additional acute cardiovascular diagnoses (aortic valve disease, endocarditis, myopericarditis)







## Clinical Evidence of Congestion

## Table 3

Use this table to identify signs and symptoms of congestion, which may be tracked as targets during decongestion and may serve as sentinel symptoms for recurrent congestion after discharge.

Clinical Evidence of Congestion		
Symptoms	Signs <sup>†</sup>	
Orthopnea	Elevated jugular venous pressure	
Dyspnea on minimal exertion	• Rales <sup>‡</sup>	
Paroxysmal nocturnal dyspnea	Pleural effusion <sup>‡</sup>	
Nocturnal cough*	Increased intensity of pulmonary component of second sound	
Bendopnea	Third heart sound	
Abdominal swelling	Murmurs of mitral and/or tricuspid regurgitation	
Early satiety	Pulsatile hepatomegaly	
Anorexia, nausea	Ascites§	
Right upper quadrant pain	Pre-sacral, scrotal, or perineal edema	
Peripheral swelling	Peripheral edema	
Rapid weight gain		

<sup>\*</sup> Often when supine; † JVP is the most sensitive sign. Rales may not always be present; ‡ Not common in chronic HF;§ May be difficult to distinguish from central adiposity





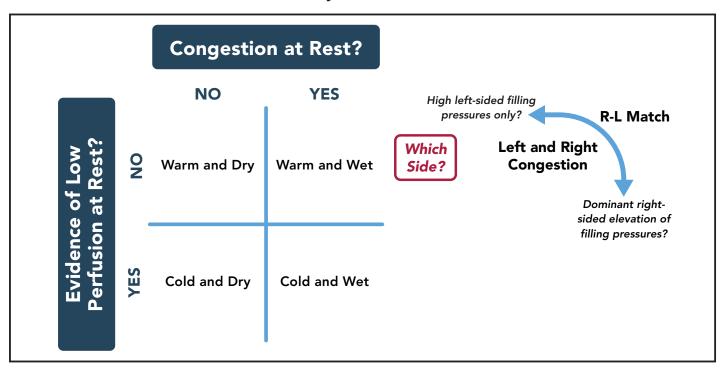


## Classification of Patients Presenting With Acutely Decompensated Heart Failure

## Figure 3

Use this figure to identify signs and symptoms of congestion, which may be tracked as targets during decongestion and may serve as sentinel symptoms for recurrent congestion after discharge.

## Hemodynamic Profiles





## Key Comorbid Conditions to Consider

## **Table 4**

Evaluation of patient comorbidities is a key component of patient assessment. Consider this list of comorbidities and their therapies in relation to their role in HF decompensation and as independent targets for intervention.

	Key Comorbid Conditions to Consider
ır	☐ Coronary artery disease/acute coronary syndrome
cula	☐ Atrial fibrillation/flutter
ovas	☐ Cerebrovascular disease, TIA/stroke
Cardiovascular	☐ Peripheral vascular disease
	☐ Structural valvular heart disease
	☐ Hypertension
	☐ Diabetes mellitus
	☐ Chronic kidney disease
e	☐ Chronic lung disease
Systemic Disease	☐ Liver disease
Dis	□ Infection
emic	□ Sleep apnea
Syste	☐ Anemia/iron deficiency
Ś	☐ Rheumatologic diseases
	□ Amyloidosis
	□ Cancer
	☐ Thyroid disease
al ons	□ Obesity
General Conditions	☐ Malnutrition
Cor	☐ Frailty, deconditioning
	☐ Dementia/cognitive decline
_	□ Depression
Psychosocial	☐ Substance abuse
sou	☐ Tobacco abuse
Psyc	☐ Alcohol abuse
а.	☐ Inadequate social support
	□ Nonadherence





## Assessing Risk During Hospitalization

## Table 5

Use this table to identify risk factors that are most likely to be modifiable. These should help to set patient goals, guide inhospital management, and re-assess integrated risk at discharge. (*Table will continue on the next page*)

**Chronic History Prior to Admission** 

Modification of bolded/italicized items decreases risk. Note that the references for risk factors are provided as examples and are not meant to list all sources of validation.

☐ Older Age (robust in all models	
□ Number of Previous HF hospita	lizations
☐ Comorbidities, especially diabet	tes, COPD, liver disease, cancer, dementia
□ Frailty	
☐ Known low LVEF in HFrEF	
☐ RV dysfunction	
Assessment at Admission	Re-Assessment at Discharge
Class IV symptoms	Effective decongestion improves prognosis.
Non-adherence to medications or salt/fluid restriction	Focused education during hospitalization with increased home and community support may improve adherence.
Progressively higher risk with higher admission natriuretic	Larger % reduction (>30-60%) in NP levels associated with better outcomes.
peptide (NP) levels	Progressively higher risk with higher discharge NP levels.
Renal dysfunction markers:	
Elevated serum creatinine or low clearance	Risk increased, but small increases in creatinine accompanying successful decongestion are associated with better prognosis.
Additional risk of high BUN	High BUN at discharge increases risk.
<ul> <li>Low spot urine sodium after first IV diuretic dose</li> </ul>	Low total urinary sodium excretion may be a more important marker than total urine output during hospitalization.
Diuretic resistance with high outpatient doses	Diuretic resistance in-hospital associated with longer LOS and worse outcomes. High risk if discharged on high loop diuretic doses.
Degree of congestion at	Residual congestion after treatment confers high risk.
admission not predictive of outcome except longer length of stay with greater excess volume	<ul> <li>High measured filling pressures</li> <li>Orthopnea</li> <li>Edema</li> <li>Composite congestion scores</li> <li>Lack of hemoconcentration</li> </ul>







## Assessing Risk During Hospitalization

## **Table 5** (Continued)

Assessment at Admission	Re-Assessment at Discharge
Hemodynamic profile of "Cold and Wet" at admission	Discharge with either cold or wet profile associated with higher risk.
Low systolic blood pressure	Low systolic blood pressure at discharge also identifies high risk.
Troponin elevation	Risk if elevated at any time during hospitalization.
Hyponatremia	Lower sodium at discharge predicts higher risk.
Increased risk at admission if: • No RAS therapy • No beta blocker therapy	Discontinuation of ACEI/ARB in hospital for hypotension or kidney dysfunction is associated with poor outcomes.  Unknown impact of re-initiation after discontinuation for circulatory and/or renal reasons.  Discharge without RAS inhibition or discharge without beta-blocker associated with high risk.

## **Unexpected In-hospital Events Conferring Additional Risks**

- · Resuscitation or Intubation
- Intravenous inotropic therapy even if brief

**Integrated Risk at Transition to Discharge =** 

- Admission Risk
- In-hospital Trajectory
- Unexpected Events

ACEI = angiotensin-converting enzyme inhibitor; ARB = angiotensin receptor blocker; BUN = blood urea nitrogen; COPD = chronic obstructive pulmonary disease; eGFR = estimated glomerular filtration rate; HF = heart failure; IV = intravenous; LOS = length of stay; LV = left ventricular; NP = natriuretic peptide; RAS = renin-angiotensin system; RV = right ventricular.







## Interventions for Patients at High Risk of Unfavorable Outcomes

## Table 6

Once you have assessed a patient's comorbidities and other risk factors, and determined them to be at high risk for unfavorable outcomes, use this table to guide additional steps.

## **Interventions for Patients at High Risk of Unfavorable Outcomes**

- Discussion of prognosis
- Evaluation for advanced therapies\* if appropriate
- Review/revision of goals of care and advanced directives
- Consideration before interventions<sup>†</sup> that may be difficult to discontinue
- Education regarding palliative care and hospice options

† Intravenous inotropic therapy, temporary circulatory support, mechanical ventilation, dialysis



<sup>\*</sup> Transplantation, mechanical circulatory support



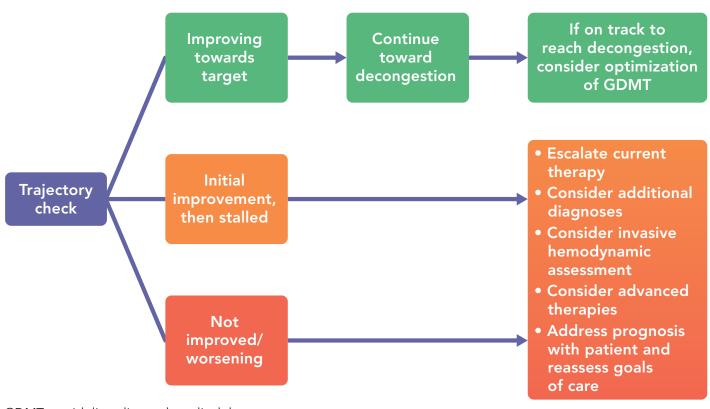


## Clinical Trajectories and Their Implications for Therapy

## Figure 4

Use figures 4 (this page) and 7a (next page) to assess a patient's daily clinical trajectory. These trajectories translate into different management strategies throughout the hospitalization and post-discharge.

## Trajectory Assessment and Next Steps



GDMT= guideline-directed medical therapy



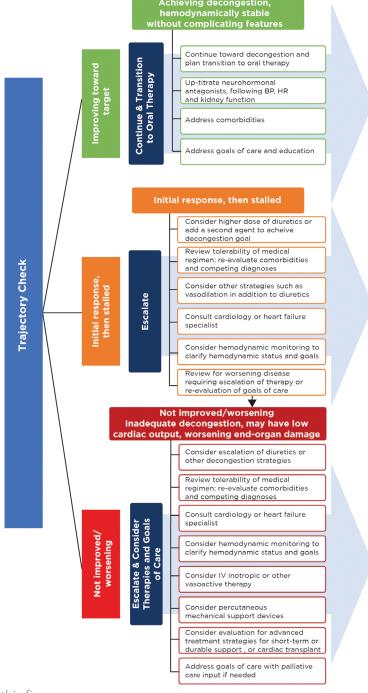


## **Clinical Trajectories**

## Figure 7a

Identification of patient's clinical trajectory translates into different management strategies throughout hospitalization and post-discharge.

→ To optimize GDMT, refer to the <u>2024 ECDP for Optimization of Heart Failure Treatment.</u>





Lick to download this figure.







## Evaluation of the Degree of Clinical Congestion, With Common Reasons for Residual Congestion Listed in the Text Box

## Figure 5

Inpatient trajectories are primarily defined by the pace and extent of decongestion. Use the figure below for evaluation of the success of de-congestion for a patient.

# Decongestion

## Freedom from clinical congestion

No peripheral edema

No rales

No dyspnea on minimal exertion

No hepatomegaly or congestive GI symptoms

No orthopnea or bendopnea

Jugular venous pressure ≤ 6-8 mm Hg

No hepatojugular reflex

## **Common Reasons for Residual Congestion**

Low cardiac output state

Dominant right heart failure

Advanced kidney disease

Symptomatic hypotension

Limitations to patient engagement in self-care

Lack of improvement in signs/symptoms of HF Lack of decrease in natriuretic peptide levels Lack of decrease in weight

GI = Gastrointestinal; HF = heart failure

Congestion





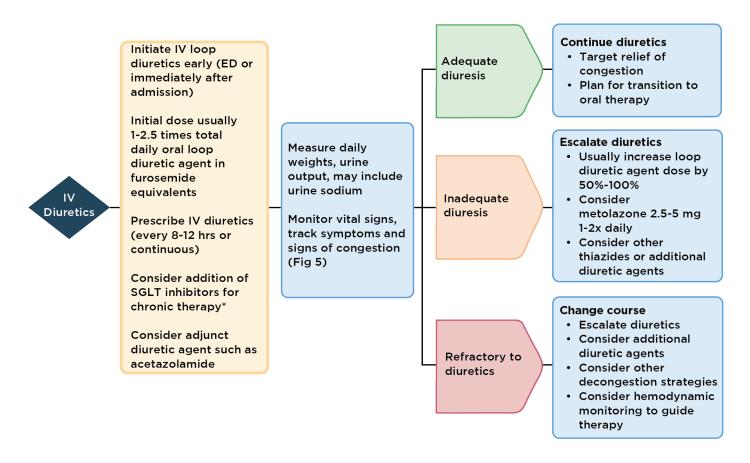


## Diuretic Therapy in Different Clinical Trajectories

Figure 6

Use this figure to guide therapy at different clinical trajectories.

## Guidance on Diuretic Therapy



BP = blood pressure; ER = emergency room; IV = intravenous







## **Diuretic Dosing**

Table 7

Use the table to help establish an effective diuretic regimen.

## **Guidance on Diuretic Dosing**

Class	Drug	Usual Outpatient Dosing (Maximum†)	Usual Inpatient Dosing* (Maximum <sup>†</sup> )
Loop diuretics	Bumetanide	0.5–2 mg orally once to twice daily (10 mg/day)	0.5-4 mg IV once to three times daily (5 mg/dose)  Or  0.5-2 mg/hour IV infusion (4 mg/hour)
	Furosemide	20–80 mg orally once to twice daily (600 mg/day)	40-160 mg IV once to three times daily (200 mg/dose)  Or 5-20 mg/hour IV infusion (40 mg/hour)
	Torsemide	10-40 mg orally once daily (200 mg/day)	N/A <sup>‡</sup>
Thiazide-type diuretics	Chlorothiazide	N/A	0.5-1 g IV once to twice daily (2 g/day)
	Hydrochlorothiazide	25-50 mg orally once daily (100 mg/day)	25–50 mg orally once to twice daily (100 mg/day)
	Chlorthalidone	25–50 mg orally once daily (100 mg/day)	12.5–25 mg orally once to twice daily (100 mg/day)
	Metolazone	2.5-5 mg orally once daily (20 mg/day)	2.5–5 mg orally once to twice daily (20 mg/day)

<sup>\*</sup> For patients receiving loop diuretics prior to admission, the oral dose should be changed to an intravenous dose of 1-2.5 times the home dose. For patients naïve to therapy, the lower end of the dosing interval should be used.

IV = intravenous; N/A = not applicable



<sup>† &</sup>quot;Usual" dose ranges reflect approved product labeling and safety and efficacy results from large, randomized controlled trials. Higher ranges may be considered on the basis of observational data and clinical experience.

<sup>‡</sup> Torsemide is not available as an intravenous formulation in the United States; oral therapy may be initiated prior to discharge to assess patient response.



## Assessing Risk During Hospitalization

## Table 5

Use this table after the patient has undergone clinical stabilization and is being transitioned to oral therapy for discharge. This assessment should help clarify goals and guide plans for longitudinal follow-up after discharge. The integrated riskfor discharge includes non-modifiable risk factors from admission, residual congestion or unexpected events during hospitalization, and lack of guideline-directed therapies for chronic heart failure. The transition node offers further opportunity for additional adjustment of these therapies to be continued after discharge. (Table will continue on the next page.)

Modification of bolded/italicized items decreases risk. Note that the references for risk factors are provided as examples and are not meant to list all sources of validation.

Cł	nronic History Prior to Admission
<ul> <li>□ Older Age (robust in all models)</li> <li>□ Number of Previous HF hospita</li> <li>□ Comorbidities, especially diabet</li> <li>□ Frailty</li> <li>□ Known low LVEF in HFrEF</li> <li>□ RV dysfunction</li> </ul>	
Assessment at Admission	Re-Assessment at Discharge
Class IV symptoms	Effective decongestion improves prognosis.
Non-adherence to medications or salt/fluid restriction	Focused education during hospitalization with increased home and community support may improve adherence.
Progressively higher risk with higher admission natriuretic peptide (NP) levels	Larger % reduction (>30-60%) in NP levels associated with better outcomes.  Progressively higher risk with higher discharge NP levels.
Renal dysfunction markers:	
Elevated serum creatinine or low clearance	Risk increased, but small increases in creatinine accompanying successful decongestion are associated with better prognosis.
Additional risk of high BUN	High BUN at discharge increases risk.
Low spot urine sodium after first IV diuretic dose	Low total urinary sodium excretion may be a more important marker than total urine output during hospitalization.
Diuretic resistance with high outpatient doses	Diuretic resistance in-hospital associated with longer LOS and worse outcomes. High risk if discharged on high loop diuretic doses.
Degree of congestion at admission not predictive of outcome except longer length of stay with greater excess volume	Residual congestion after treatment confers high risk.  • High measured filling pressures  • Orthopnea  • Edema  • Composite congestion scores  • Lack of hemoconcentration





## **THERAPIES**

## Assessing Risk During Hospitalization

## **Table 5** (Continued)

Modification of bolded/italicized items decreases risk. Note that the references for risk factors are provided as examples and are not meant to list all sources of validation.

Assessment at Admission	Re-Assessment at Discharge	
Hemodynamic profile of "Cold and Wet" at admission	Discharge with either cold or wet profile associated with higher risk.	
Low systolic blood pressure	Low systolic blood pressure at discharge also identifies high risk.	
Troponin elevation	Risk if elevated at any time during hospitalization.	
Hyponatremia	Lower sodium at discharge predicts higher risk.	
Increased risk at admission if: No RAS therapy	Discontinuation of ACEI/ARB in hospital for hypotension or kidney dysfunction is associated with poor outcomes.	
· No beta blocker therapy	Unknown impact of re-initiation after discontinuation for circulatory and/or renal reasons.	
	Discharge without RAS inhibition or discharge without beta-blocker associated with high risk.	

## **Unexpected In-hospital Events Conferring Additional Risks**

- Resuscitation or Intubation
- · Intravenous inotropic therapy even if brief

**Integrated Risk at Transition to Discharge =** 

- Admission RiskIn-hospital TrajectoryUnexpected Events

ACEI = angiotensin-converting enzyme inhibitor; ARB = angiotensin receptor blocker; BUN = blood urea nitrogen; COPD = chronic obstructive pulmonary disease; eGFR = estimated glomerular filtration rate; HF = heart failure; IV = intravenous; LOS = length of stay; LV = left ventricular; NP = natriuretic peptide; RAS = renin-angiotensin system; RV = right ventricular.





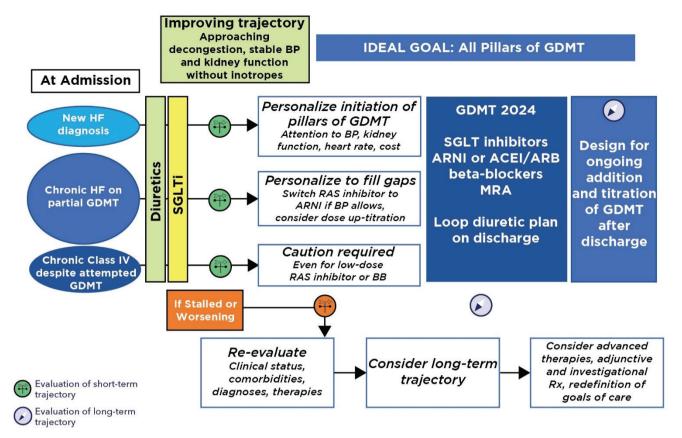


## Titration of GDMT in HFrEF

## Figure 7b

Use this figure to guide titration of GDMT in HFrEF based on patient's initial presentation and clinical trajectory.

→ To optimize GDMT, refer to the 2024 ECDP for Optimization of Heart Failure Treatment.



Titration of GDMT in HFrEF by initial presentation and trajectory. Patients with decompensated HFrEF should be diuresed and started on SGLT inhibitor unless contraindicated or cost prohibitive. Those with an improving trajectory (denoted by the green weathervane icon) should have optimization of GDMT. Patients may have a new HF diagnosis, in which case initiation of all 4 pillars of GDMT should be attempted. Patients with chronic HF on partial GDMT should have personalized therapy to fill in gaps, considering a switch from an ACE inhibitor/ARB to ARNI if appropriate. Caution is required for patients with chronic Class IV HF with decompensated HF; these patients may tolerate even low doses of beta-blockers and RAS poorly, although an attempt at titration may be made. Patients whose short-term trajectory is stalled, or worsening (denoted by the orange weathervane icon) should have re-evaluation of comorbidities and consideration of other diagnoses. The long-term trajectory (denoted by the compass icon) should be reevaluated, with consideration of goals of care, candidacy for advanced therapies, and experimental treatments. The ideal goal is initiation of all 4 pillars of GDMT for HFrEF in the hospital on a baseline of diuretic therapy. A plan for ongoing addition and titration of GDMT after discharge should be fashioned as well.

ACE = angiotensin-converting enzyme; ARB = angiotensin receptor blocker; ARNI = angiotensin receptor/neprilysin inhibitor; BB = beta-blocker; BP = blood pressure; d/c = discharge; GDMT = guideline-directed medical therapy; MRA = mineralocorticoid antagonist; RAS = renin-angiotensin system; SGLT = sodium-glucose cotransporter.





## Education for Patients, Families, and Caregivers

## Figure 10

Use this checklist to guide education for the patient, family, and caregivers. Education provided should be culturally appropriate, and delivered verbally and in written form.

Current medications
• Indication
<ul> <li>Dose/frequency</li> </ul>
Potential side effects
Potential adherence barriers
Activity level
Dietary sodium restrictionmg/day
Fluid restriction $\square$ YesL/day or $\square$ No
Daily weight monitoring
Has scale  ☐ Yes  ☐ No
• Records daily weights $\square$ Yes $\square$ No
Assessment for peripheral edema
Smoking cessation counseling for current or recent smokers
Substance use counseling, if applicable
List of warning signs of decompensation
What to bring to each outpatient appointment
• List of meds
Recordings of daily weights
Who to call for increased weight / worsening symptoms / ICD discharge
Diuretic management plan
Plans for continuation of care
Cardiology specialty clinic follow-up appointment/





## Model Focused Discharge Handoff

## Figure 11

Use this form to convey crucial patient information to continuing care providers to support direct communication. This form is specifically designed to travel with the patient. Areas that are shaded can be formatted as selection menus if this form is being used as a template in a clinical decision tool.

## **FOCUSED DISCHARGE HANDOFF**

Name Age MRN Admission D	ate/Dis	charge Date//
HF TYPE: ☐ HFrEF ☐ HFpEF (≥50) ☐ HFmrEF (41-49) Last LVEF	Triggers for H	ospitalization
<b>HF ETIOLOGY:</b> □ Ischemic □ Non-isch □ Other		
Arrhythmia history		
	-	
CONDITION AT DISCHARGE:   Discharged to Rehab or other facility		
D/C BP/ HR Rhythm Sinus AF paced Weight At D/Clbs Est. target	weightlbs	
Residual congestion at D/C Y N Edema 0-4+ JVP Rales Y N A	scites 🗆 Y 🗆 N	
If still wet, limited by ☐ Dominant R heart failure ☐ Kidney function ☐ Hypotension ☐ Oth	ner	
Kidney Function Discharge BUN/Cr eGFR Worst CR in hospital	Baseline_	
Biomarkers Admission BNP/proBNP "Dry" BNP/proBNP (if known)		
0	• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •
Comorbidites:		
Psychosocial Factors  Other hospital events □ code □ sepsis □ dialysis □ intubation IV inotropes used?		
Code Status   Full code   Full code but discussed   DNR/DNI   DNI only   POLST	iveeas aiscussion	
DISCHARGE HF MEDICATIONS:		
Diuretic Loopdose/freq Metolazone dose/freq Other: dose/freq		
In-hospital effective loop dosemg IV   Daily   BID   TID   Drip atmg/hr Metolaz		N
Triggers for rescue dose: Iflbs up, or(sentinel symptom)		
Rescue doseorally and/or metolazonefordays before rechec	k	
K+ replacementmeQ/day Plan for more K+ with rescue dose?		
GUIDELINE DIRECTED MEDICAL THERAPY:		PLAN
RAS meds ACEImg/day ARBmg/day ARNImg/day Dose in hospit.	al? UY UN	PLAN  Start Titrate
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RAS meds ACEImg/day ARBmg/day ARNImg/day Dose lin hospit.  If not given or dose l, why? ☐ Hypotension ☐ Dizziness ☐ Kidney fx ☐ Cough  If no ARNI, why? ☐ Hypotension ☐ Cost ☐ Hyperkalemia ☐ Angioedema  Beta-blockermg/day Dose l in hospital? ☐ Y ☐ N	al? OY ON	☐ Start ☐ Titrate
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RAS meds ACEImg/day ARBmg/day ARNImg/day Dose lin hospit.  If not given or dose l, why?   Hypotension   Dizziness   Kidney fx   Cough    If no ARNI, why?   Hypotension   Cost   Hyperkalemia   Angioedema  Beta-blockermg/day Dose l in hospital?   Y   N  If not given or dose l, why?   Hypotension   bradycardia  MRA   Y   N   If not, why not?   Hypotension   Kidney function   Hyperkalemia	el? Y N	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Titrate
RAS meds       ACEImg/day       ARBmg/day       ARNImg/day       Dose lin hospit.         If not given or dose l, why?       Hypotension □ Dizziness □ Kidney fx □ Cough         If no ARNI, why?       Hypotension □ Cost □ Hyperkalemia □ Angioedema         Beta-blockermg/day       Dose l in hospital? □ Y □ N         If not given or dose l, why?       □ Hypotension □ bradycardia         MRA □ Y □ N       If not, why not? □ Hypotension □ Kidney function □ Hyperkalemia         SGLTi □ Y □ N       If not, why not? □ Kidney function □ DKA □ UTI/GMI □ Cost		□ Start □ Titrate □ △ ARNI □ Start □ Titrate
RAS meds       ACEImg/day       ARBmg/day       ARNImg/day       Dose lin hospit.         If not given or dose l, why?       Hypotension □ Dizziness □ Kidney fx □ Cough         If no ARNI, why?       Hypotension □ Cost □ Hyperkalemia □ Angioedema         Beta-blockermg/day       Dose l in hospital? □ Y □ N         If not given or dose l, why?       □ Hypotension □ bradycardia         MRA □ Y □ N       If not, why not? □ Hypotension □ Kidney function □ Hyperkalemia         SGLTi □ Y □ N       If not, why not? □ Kidney function □ DKA □ UTI/GMI □ Cost		Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Titrate
RAS meds       ACEImg/day       ARBmg/day       ARNImg/day       Dose lin hospit.         If not given or dose l, why?       Hypotension □ Dizziness □ Kidney fx □ Cough         If no ARNI, why?       Hypotension □ Cost □ Hyperkalemia □ Angioedema         Beta-blockermg/day       Dose l in hospital? □ Y □ N         If not given or dose l, why?       □ Hypotension □ bradycardia         MRA □ Y □ N	<b>□Y</b> □N	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Titrate
RAS meds       ACEImg/day       ARBmg/day       ARNImg/day       Dose lin hospit.         If not given or dose l, why?       Hypotension □ Dizziness □ Kidney fx □ Cough         If no ARNI, why?       Hypotension □ Cost □ Hyperkalemia □ Angioedema         Beta-blockermg/day       Dose l in hospital? □ Y □ N         If not given or dose l, why?       □ Hypotension □ bradycardia         MRA □ Y □ N       If not, why not? □ Hypotension □ Kidney function □ Hyperkalemia         SGLTi □ Y □ N       If not, why not? □ Kidney function □ DKA □ UTI/GMI □ Cost	□Y □N arfarin	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Titrate
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RAS meds ACEmg/day ARBmg/day ARNImg/day Dose lin hospit.  If not given or dose l, why?   Hypotension   Dizziness   Kidney fx   Cough    If no ARNI, why?   Hypotension   Cost   Hyperkalemia   Angioedema    Beta-blockermg/day   Dose l in hospital?   Y   N    If not given or dose l, why?   Hypotension   bradycardia    MRA   Y   N   If not, why not?   Hypotension   Kidney function   Hyperkalemia    SGLTi   Y   N   If not, why not?   Kidney function   DKA   UTI/GMI   Cost    Other HF meds   Hydral/nitrates   Y   N   Ivabradine   Y   N   Digoxin   Y   N   Inotropes    Anticoagulation for   AF   DVT/PE   Mech valve   LV thrombus with   DOAC   Was    Antiplatelet for   ACS   PCI   Stroke/TIA with   ASA   Clopidogrel   Prasugrel	□Y □N arfarin	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Titrate
RAS meds ACEmg/day ARBmg/day ARNImg/day Dose lin hospit.  If not given or dose l, why?   Hypotension   Dizziness   Kidney fx   Cough     If no ARNI, why?   Hypotension   Cost   Hyperkalemia   Angioedema     Beta-blockermg/day   Dose l in hospital?   Y   N     If not given or dose l, why?   Hypotension   bradycardia     MRA   Y   N   If not, why not?   Hypotension   Kidney function   Hyperkalemia     SGLTI   Y   N   If not, why not?   Kidney function   DKA   UTI/GMI   Cost     Other HF meds   Hydral/nitrates   Y   N   Ivabradine   Y   N   Digoxin   Y   N   Inotropes     Anticoagulation for AF   DVT/PE   Mech valve   LV thrombus with   DOAC   Water     Antiplatelet for ACS   PCI   Stroke/TIA with   ASA   Clopidogrel   Prasugrel     Antiarrhythmic   Amiodarone   Dofetilide   Sotalol   Other     See patient discharge document and full discharge summary for complete medication list	□Y □N arfarin □Ticagrelor	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Titrate □ Start □ Continue
RAS meds ACEImg/day ARBmg/day ARNImg/day Dose lin hospit.  If not given or dose l, why?   Hypotension   Dizziness   Kidney fx   Cough	□Y □N arfarin	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Titrate □ Start □ Continue
RAS meds ACEImg/day ARBmg/day ARNImg/day Dose lin hospit.  If not given or dose l, why?   Hypotension   Dizziness   Kidney fx   Cough	□Y □N arfarin □ Ticagrelor  Appt date and time	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Continue
RAS meds   Mode   Mod	□Y □N arfarin □ Ticagrelor  Appt date and time	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Continue
RAS meds ACEmg/day ARBmg/day ARNImg/day Dose lin hospit.  If not given or dose l, why?   Hypotension   Dizziness   Kidney fx   Cough	□Y □ N  arfarin □ Ticagrelor  Appt date and time  Results sent to	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Continue
RAS meds   ACE  mg/day   ARB mg/day   ARN  mg/day   Dose 1 in hospit.	□Y□N  arfarin □ Ticagrelor  Appt date and time  Results sent to  Phone Number	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Continue
RAS meds ACEImg/day ARBmg/day ARNImg/day Dose lin hospit.  If not given or dose l, why?   Hypotension   Dizziness   Kidney fx   Cough	□Y□N  arfarin □ Ticagrelor  Appt date and time  Results sent to  Phone Number  Phone Number	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Continue
RAS meds ACEmg/day ARBmg/day ARNImg/day Dose lin hospit.  If not given or dose l, why?   Hypotension   Dizziness   Kidney fx   Cough	□Y□N  arfarin □ Ticagrelor  Appt date and time  Results sent to  Phone Number  Phone Number	Start □ Titrate □ △ ARNI □ Start □ Titrate □ Start □ Continue









## Checklist for Communication to Continuing Care Providers

## Figure 12

Use this checklist as a guide for communication with continuing care providers about pertinent patient information and potential issues.

HC	Reason for admission Sentinel symptoms Congestion status  • Admission, discharge, and target weight  • Admission and discharge kidney function  • Diuretic dosing  • Rescue dosing Unexpected events
<b>PL</b>	ANNED THERAPIES AND MONITORING  Plan for initiation, titration, and optimization of GDMT  ARNI/ACEI/ARB  Beta blockers  Aldosterone mineralocorticoid antagonists  SGLT inhibitor  Ivabradine  Hydralazine/isosorbide
	Adjustment of diuretics Plan to monitor electrolytes and kidney function Follow-up for pending or planned diagnostic tests Plan for EP consult if sudden death risk or potential candidate for device therapy OR AF Recommendations for when to assess response to therapy and in what venue (in person vs telehealth) COVID, Pneumovax and Influenza vaccination
FO	Kidney function Diabetes Sleep-disordered breathing Depression Anemia IV iron given? Other





## Checklist for Follow-Up Phone Call

## Figure 13

Use this checklist to help organize and streamline the follow-up phone call to ensure that it is comprehensive yet focused.

## Early Post-Discharge: Checklist for 48-72 Hour Follow-Up Phone Call

TOPIC	VITAL QUESTION	CAUSE FOR IMMEDIATE CONCERN	TEACHING POINTS TO BE COVERED I CALL/CLINIC, USING TEACH BACK
Symptoms Sentinel symptom from hospitalization Shortness of breath Orthopnea Edema	How is?  □ Same □ Better □ worse than at discharge	Alert If WORSE	Do you know what symptoms you should be paying attention to?
Dizziness	Are you having trouble with dizziness? Yes No Is it just when you first stand up or does it last longer?	FREQUENT DIZZINESS	Review dizziness as potential symptom of concern
Daily Weights	Are you weighing yourself daily?	ALERT If no weights or if weight increase > trigger	Importance of weights as short- term indication of fluid balance. Review diuretic plan from discharg Do you have a plan for what to do your weight increases?
Medications (Refer to discharge list)	Do you have these medications prescribed at discharge?	ALERT If Not obtained, Or not taking correctly	Types and purposes of HF medications
Salt restriction	Are you watching your salt intake?		Review contribution of salt to fluid retention Common high-salt items How to read labels
Fluid restriction (for patients who have one)	Are you keeping track of your fluid intake?  \( \text{Yes} \) No What is your daily limit?  What are you doing to stay within your limit?		Review contribution of fluid to symptoms, Importance of fluid restriction for fluid balance and how to account for fluids in food as well beverages.  Reassure this is often not a sign of dehydration in heart failure  Present alternatives such as frozen fruit, etc
Follow-up	When is your follow-up appointment? and in what venue (in person, telehealth)?	NO follow-up appt or no	
	vende (in person, colonedia).	way to get there	





## FIrst Post-Discharge Visit Checklist

## Figure 14

Consider the key components listed in this checklist to guide the first post-discharge visit to reassess clinical status, review medications, provide additional education, and address issues that may lead to worsening HF.

History	Medications
Discharge summary reviewed.	Comprehensive medication reconciliation
Etiology of cardiomyopathy identified.	Beta-blocker
Precipitant of exacerbation identified.	- Dose optimized?
Heart failure compensated?	ARNI/ACEI/ARB
- NYHA class.	- Dose optimized?
- Weight log reviewed?	- Eligible to switch to ARNI?
- Symptoms reviewed?	Mineralocorticoid antagonist
<ul> <li>Important concomitant disease states</li> </ul>	- Dose optimized?
- CKD	SGLT inhibitor
- Diabetes	Diuretic agents
- Hypertension	- Dose adjustment?
- AF	• Ivabradine? (Consider initiation if heart rate remains
- COPD	elevated despite beta-blocker optimization)
- OSA	Interventional therapies (if applicable)
- Others	Revascularization
Goals of Care changed	• CRT
Physical Exam	• ICD
Vital signs	AV node or AF ablation
BMI	Valvular intervention
Orthostatic blood pressure	Remote monitoring
Jugular venous distention	Patient education
• Rales +/-	Importance of adherence
<ul><li>"cold/warm", "wet/dry" profile</li></ul>	Medication education
S3 present/absent	Dietary education
•	Activity education
Diagnostic Testing	Smoking cessation
<ul><li>Basic metabolic panel</li><li>Complete blood count</li></ul>	Alcohol consumption
BNP or NT-proBNP	Follow-up appointment scheduled
Liver function panel (per discretion of clinician)	☐ In-person
Iron studies (per discretion of clinician)	☐ Telehealth
High-sensitivity Troponin, sST2,	Consultations
Gal-3 (per discretion of clinician)	Home health services
• 12-lead ECG	Cardiac rehab referral
Chest x-ray (per discretion of clinician)	<ul> <li>Advanced heart failure clinic referral</li> </ul>
Review LVEF (%). If not available, attain TTE	Palliative/hospice referral
• Follow-up EF:	
- 40 days post-MI	
- 3 months post-NICM	
Ischamia Evaluation Mondad?	





## PALLIATIVE CARE

## **Aspects of Palliative Care**

## Figure 15

Consider the aspects of palliative care listed in the figure to help set goals around advance care planning.

## Advanced Care Planning

- Identification of surrogate decision-maker
- Exploration of values and general preferences
- · Execution of living will

## Goals of Care Discussions

- · Resuscitation status
- · Return to ICU
- · Inactivation of defibrillation
- · Services, limitations, and family role during hospice care
- · End of life at home or in skilled facility

## Determination of Need for Specialist Palliative Care

- · Complicated advanced care planing
- Symptom control
- · Disagreement between clinicans and patient/family
- · Marked caregiver or family distress
- · Hospice referral

## Caregiver/Family Support

· Referral to social, psychological, and respite services

This figure reflects recent data and publications, including:

- 1. A study demonstrating that a palliative care referral trigger tool decreased hospital readmission rates up to 90 days postdischarge while also increasing the completion of advanced directives.
- 2. The 2022 AHA/ACC/HFSA HF Guideline, which advocates a need-based approach to palliative care, with emphasis on shared decision-making, caregiver and bereavement support, and hospice care.

HF = heart failure; ICU = intensive care unit





PALLIATIVE CARE

## Goals of Care/Advance Care Planning

## Table 9

The table is intended to guide clinician-patient discussions on goals of care and advance care planning.

## Advance Care Planning Discussion

## **Assess Readiness to Discuss Goals of Care**

## **Assess Understanding of Prognosis**

## **Confirm/Discuss Goals of Care**

- Confirm/Elicit patient values and preferences pertaining to quality of life and life prolongation (cultural, religious).
- Discuss aspects of what the patient would consider an unacceptable quality of life.
- Discuss benefits/burdens of reasonable therapeutic options.

## **Confirm/Establish Surrogate Decision Maker**

Person best able and willing to represent patient's values and preferences and patient's best interests.

## **Establish/Reassess Code Status**

- · Based on Goals of Care Discussion.
- Do Not Attempt Resuscitation (DNAR).
- Full Code.
- · Attempt shock without other measures.

## **Discuss Management of Defibrillator when appropriate**

Pacing function is often left intact even if defibrillation is deactivated.

## Determine need for specialist palliative care consultation





## HEART FAILURE HOSPITALIZATION PATHWAY TOOLKIT

## **Links to Additional Resources**

## **■ ACC Resources:**

TreatHF App:

Tools.ACC.org/TreatHF/

HF Advance Care Planning Toolkit:

<u>CvQuality.ACC.org/clinical-toolkits/advance-care-planning-toolkit</u>

**HF Condition Center:** 

CardioSmart.org/Heart-Conditions/Heart-Failure

HF Action Plan:

CardioSmart.org/MyHFActionPlan

**HF Infographics:** 

CardioSmart.org/~/media/Images/Infographics/2016/Heart%20Failure%20resize.ashx

CardioSmart.org/~/media/Images/Infographics/Heart-Failure-Journey.ashx

2022 AHA/ACC/HFSA Guideline for the Management of Heart Failure:

https://www.jacc.org/doi/10.1016/j.jacc.2021.12.012?

ga=2.74773586.1174326798.1737484184-1586257890.1733853887

2024 ECDP for Optimization of HF Treatment:

https://www.jacc.org/doi/10.1016/j.jacc.2023.12.024?

ga=2.3585904.1174326798.1737484184-1586257890.1733853887

2024 ECDP Focused Update of the 2019 ACC ECDP on Heart Failure Hospitalization:

https://www.jacc.org/doi/10.1016/j.jacc.2024.06.002?

ga=2.6160243.1174326798.1737484184-1586257890.1733853887

2019 ACC ECDP on Heart Failure Hospitalization:

https://www.jacc.org/doi/pdf/10.1016/j.jacc.2019.08.001?

ga=2.74896466.1174326798.1737484184-1586257890.1733853887

## **■ External Resources:**

**HFSA Advanced Care Training Module:** 

HFSA.org/wp-content/uploads/2018/03/HFSA-Module-9-03.14.2018-LR.pdf

