

Date (dd-Mon-yyyy)

## Heart Failure Acute Admission Adult

Select orders by placing a ( $\checkmark$ ) in the associated box

Time (hh:mm)

Last Name (Legal)		First Name (Legal)			
Preferred Name 🗆 Last 🗆 First			DOB	(dd-Mon-yyyy)	
PHN	ULI □ Same as PHN		s PHN	MRN	
Administrative Gender □ N □Non-binarv/Prefer not to di			se (X)	☐ Female ☐ Unknown	

	□Non-binary/Prefer	not to disclose (X) 🛛 Unknown						
<ul> <li>To be added to General Admission Orders</li> <li>Notify Primary Care Provider and Heart Function Clinic (HFC), if HFC patient, on next business day</li> <li>Daily morning weights (record on chart before 0900 hours) – teach patient to do and record</li> </ul>								
<ul> <li>Oxygen delivered as required to keep Sp0</li> <li>2000 mL fluid restriction ORn</li> <li>2000 mg sodium diet ORn</li> <li>2000 mg sodium diet ORn</li> <li>Ambulate - Early Mobilization (done within 4)</li> <li>Lab/Tests – Specific to Heart Failure</li> <li>Electrocardiogram</li> <li>Chest X-Ray:Posterior Anterior an</li> <li>Transthoracic Echocardiogram as soor</li> <li>Creatinine, electrolytes, daily x</li> <li>BNP or NT-proBNP on admission (<i>if not</i></li> <li>MP or NT-proBNP within 48 hours prior</li> </ul>	nL Other ( <i>specify</i> ) ng Other ( <i>specify</i> ) 8 <i>hours</i> ) d Lateral <b>or</b> □ Portable n as possible if not performed with days already completed in emergency depart	nin the past 12 months						
Heart Failure Specific Medications								
<ul> <li>Current Canadian standard of care for medical therapy for HFrEF is Angiotensin Neprilysin Inhibitor (ARNi), Beta Blocker, Mineralocorticoid Receptor Antagonist (MRA) and Sodium-Glucose Cotransporter-2 Inhibitor (SGLT2i), see Figure 1. SGLT2i class is currently not on AHS formulary for this indication and is therefore not included here.</li> <li>Medication review and optimization of evidence based therapies is a critical component of heart failure patient discharge planning.</li> <li>Avoid 'non-dihydropyridine' calcium channel blockers, nonsteroidal anti-inflammatory drugs and COX II inhibitors if possible.</li> </ul>								
Refer to Best Possible Medication	History (BPMH) before initiation	ig below medications						
Diuretics (Refer to Tables 1 & 2: Acute Heart Fa	ailure Diuretic Dosing, Recommenda	tions & Practical Tips)						
Choose ONE OR I furosemid	e mg PO daily e mg IV twice daily x _ le mg / hour IV continuc one mg PO da	days. Reassess daily. us x 1 day. Reassess daily.						
Refer to Tables 3 & 4: Modified CCS Care of Pati	ent with Reduced Ejection Fraction							
Can patient tolerate an Angiotensin Converti	ng Enzyme Inhibitor (ACEI)?							
Yes (Angiotensin Converting Enzyme Ir	hibitor (ACEI))							
Choose ONE Choose ONE Choos								
□ No (Angiotensin Receptor Blocker (ARI	3))							
Choose ONE Choose ONE Choose ONE Choose ONE CR								
Prescriber Name (print)	Prescriber Signature	Prescriber Designation						

	Table 1: Acute Heart Failure (AHF) – Diuretic Dosing							
eGFR*	Patient	Initial IV Dose^	Maintenance Dose					
Greater than or equal to 60 mL/min/1.73 m2	New-onset HF or no current diuretic therapy	Furosemide 20 to 40 mg 2 to 3 times daily	Lowest diuretic dose that allows clinical					
			stability is the ideal dose					
Less than 60 mL/min/1.73 m2	New-onset HF or no current diuretic therapy	Furosemide 20 to 80 mg 2 to 3 times daily						
	Established HF or chronic oral diuretic therapy	Furosemide dose IV equivalent of oral dose						

\*eGFR is calculated from the Cockroft-Gault, CKD-EPI, or Modification of Diet in Renal Disease formula. ^IV continuous furosemide at doses of 5 to 20 mg per hour is also an option

#### Table 2: Recommendations & Practical Tips for Diuretic Use

#### Recommendations

\*IV diuretics should be given as first-line therapy for patient with pulmonary or peripheral congestion.

\*For patients requiring IV diuretic therapy, furosemide may be dosed intermittently (eg, twice daily) or as a continuous infusion

#### Practical Tips

\*When acute congestion is cleared, the lowest dose that is compatible with stable signs and symptoms should be used.

\*Target 0.5 to 1 kg of weight loss per 24-hour period while a patient with volume overload is actively diuresing. Patients who are losing less than 0.5 kg per day despite at least 40 mg of IV furosemide will need a reassessment of fluid status and might be diuretic resistant.

# \*When transitioned from IV to oral diuretic therapy, the stability of symptoms, weight, and hemodynamics should be observed for approximately 24 hours before hospital discharge.

\*To transition a patient to oral diuretics, be aware that the oral version of furosemide has approximately 50% bioavailability compared with IV furosemide.

\*Add another type of diuretic with different site of action (thiazides, spironolactone).

2017 Comprehensive Update of the Canadian Cardiovascular Society Guidelines for the Management of Heart Failure

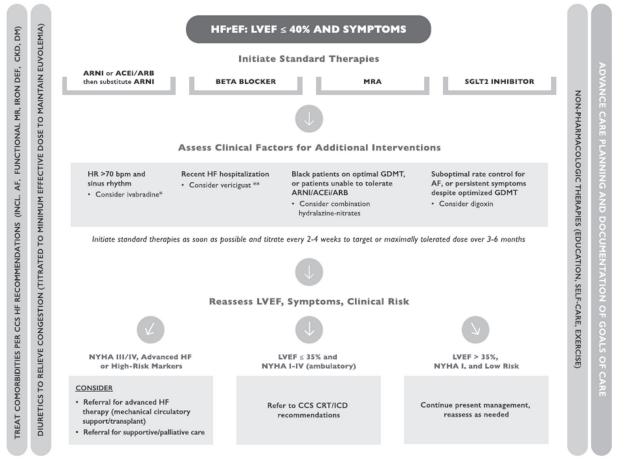


Figure 1: Simplified Treatment Algorithm for Management of HF with Reduced Ejection Fraction (HFrEF) <u>CCS/CHFS Heart Failure 2021 Guidelines Update</u>



Alberta Health Services			Last Name (L		First Nam	
	3		Preferred Na	me 🗆 Last 🗆 First	DOB	(dd-Mon-yyyy)
Heart Failure A	cute Adı	mission Adult	PHN	ULI 🗆 S	ame as PHN	MRN
Heart Failure Spe	ecific Medi	cations continued		e Gender □ N /Prefer not to di		<ul><li>□ Female</li><li>□ Unknown</li></ul>
Beta Blockers		isoPROLol 1.25 mg PO or	veo daily			
Choose ONE	JOR□b OR□c	visoPROLol mg arVEDilol 3.125 mg PO tw	PO once daily.			
Sinus Node Inhit	oitors	0				
Choose ONE	{ OR □   OR □	vabradine restrictions in Table vabradine 2.5 mg PO twice vabradine 5.0 mg PO twice vabradine 7.5 mg PO twice	e daily. e daily.	estrictions		
Vineralocorticoi	d Recepto	r Antagonists (MRA)				
		<ul> <li>□ spironolactone 12.5 mg</li> <li>□ spironolactone 25 mg</li> </ul>	PO once daily.			
Choose ONE		Eplerenone only IF patie intolerant to spironolactor				
	Refe OR	r to Eplerenone restrictions ir □ eplerenone 25 mg PO □ eplerenone 50 mg PO	once daily.	ion Restriction	IS	
Angiotensin Rec	eptor Nep	rilysin Inhibitor (ARNI)				
formulary restriction	ons. <sup>**</sup> see Ta	<b>ESTO)</b> only <b>IF</b> patient was ble 4: Medication Restrictions				
a HF specialist (In	ternal Med	an 40% done within the pa icine, Cardiologist) for opti <i>pination with ACEI or ARB the</i>	mization of this e			
	Refe	r to sacubitril-valsartan (ENT	RESTO) restrictio	ns in Table 4:	Medicatio	n Restrictions
		□ sacubitril-valsartan 24 Start date (dd-Mon-yyyy) _	mg - 26 mg (EN	TRESTO) PO		aily.
Choose ONE		□ sacubitril-valsartan 49 Start date (dd-Mon-yyyy) _		Time (hh:n	nm)	
		□ sacubitril-valsartan 97 Start date (dd-Mon-yyyy) _		Time (hh:n	nm)	
If converting patient dose to start drug.	to sacubitri	l-valsartan (ENTRESTO) fron	n ACEI: Stop ACE	El, wait at leas	t 36 hours	after last ACEI
-		l-valsartan (ENTRESTO) fron ′e been due.	n ARB: Stop ARB	, no washout j	period neo	essary, start drug
Vasodilators: Nit	rates					
Choose ONE	$\int$	□ nitroglycerin patch Patch on at (hh:mm)		ur apply daily <sup>(hh:mm)</sup>		

mg PO once daily.

### **Prior to Discharge**

Review vaccine history and eligibility criteria

□ Influenza vaccine, 0.5 mL IM x 1

• If indicated, when patient is no longer febrile or acutely ill, with verbal informed consent, during vaccination season, if NOT already vaccinated.

□ pneumococcal polysaccharide vaccine, 0.5 mL IM x 1

• If indicated, when patient is no longer febrile or acutely ill, with verbal informed consent.

**OR** isosorbide mononitrate

		vith Reduced Ejection F	
Drug Class	Specific Agent	Start Dose (orally)	Target Dose (orally)
Angiotensin receptor- neprilysin inhibitor (ARNI)	sacubitril/valsartan**	24/26 mg Daily	97/103 mg BID
ACE inhibitors (ACEI)	enalapril	1.25 to 2.5 mg BID	10 mg BID / 20 mg BID in NYHA class IV
	lisinopril	2.5 to 5 mg Daily	20 to 35 mg Daily
	ramipril	1.25 to 2.5 mg BID	5 mg BID
	perindopril	2 to 4 mg Daily	4 to 8 mg Daily
	trandolapril	1 to 2 mg Daily	4 mg Daily
Angiotensin receptor	Candesartan	4 to 8 mg Daily	32 mg Daily
Blocker (ARB)	valsartan	40 mg BID	160 mg BID
Beta-blockers	bisoPROLol	1.25 mg Daily	10 mg Daily
	carVEDilol	3.125 mg BID	25 mg BID / 50 mg BID (greater than 85 kg)
	MetoPROLol CR/XL (not available in Canada)	12.2 to 25 mg Daily	200 mg Daily
Mineralocorticoid	spironolactone	12.5 mg Daily	50 mg Daily
receptor antagonists (MRA)	eplerenone **	25 mg Daily	50 mg Daily
Sodium-glucose	dapagliflozin	10 mg Daily	10 mg Daily
Cotransporter-2 Inhibitor (SGLT2i) *	empagliflozin	10 mg Daily	10 to 25 Daily
	canagliflozin	100 mg Daily	100 to 300 mg Daily
Sinus node inhibitors	Ivabradine**	2.5 to 5 mg BID	7.5 mg BID
Soluble guanylate cyclase (sGC) stimulator	vericiguat (not available in Canada)	2.5 mg Daily	10 mg Daily
Vasodilators	hydralazine	10 to 37.5 mg TID	75 to 100 mg TID to QID
	Isosorbide dinitrate (Isosorbide mononitrate 30 to 120 mg Daily may be ordered as a long acting formulation)	10 to 20 mg TID	40 mg TID
Cardiac glycosides	digoxin	0.0625 to 0.125 mg Daily	N/A: monitor for toxicity

Table 3: Modified 2021 CCS/CHFS Evidence Based Heart Failure Drugs/Dosage for

Source: CCS/CHFS Heart Failure Guidelines Update: Defining A New Pharmacologic Standard of Care for Heart Failure with Reduced Ejection Fraction. McDonald M et al. Can Journal Cardiol 2021; 37: 531-546.

\* SGLT2i - This class is currently not on AHS formulary for this indication \*\* Refer to Table 4: Medication Restrictions

Version Date: May 18, 2021

Alberta Health		Last Name (Legal)		First Name (Legal)		
	Services	Preferred Name 🗆 L	ast 🗆 First	DOB	(dd-Mon-yyyy)	
Heart	Failure Acute Admission Adult	PHN	ULI 🗆 Sar	me as PHN	MRN	
		Administrative Geno			<ul><li>Female</li><li>Unknown</li></ul>	
Additio	onal Orders					
Prescri	per Name (print)	Prescriber Signature		Prescrit	per Designation	
		<u> </u>				

#### **Table 4: Medication Restrictions**

#### sacubitril-valsartan (ENTRESTO) restrictions:

Only use sacubitril-valsartan (ENTRESTO) for:

- 1. Heart failure patients on Entresto prior to admission; or
- 2. The treatment of heart failure in patients with the following criteria:
  - a. Reduced left ventricular ejection fraction [less than 40%]; and
  - b. New York Heart Association class II or III heart failure symptoms despite at least 4 weeks of treatment with:
    - a stable dose of an ACE inhibitor or an ARB; and
      - in combination with a beta-blocker and other recommended therapies, including an aldosterone antagonist [if tolerable]; and
  - c. Those patients who have plasma BNP greater than or equal to 150 pg/mL or NT-proBNP greater than or equal to 600 pg/mL, or if the patient has been hospitalized for heart failure within the past 12 months and has plasma BNP greater than or equal to 100 pg/mL or NT-proBNP greater than or equal to 400 pg/mL levels

\*All new starts must be ordered by a specialist in Cardiology or Internal Medicine as per Alberta Blue Cross requirements.

#### eplerenone restrictions:

Only use eplerenone for:

1. Patients on eplerenone prior to admission; or

2. Patients with New York Heart Association (NYHA) Class II chronic heart failure (HF) with left ventricular systolic dysfunction (LVSD) with ejection fraction (EF) equal to or less than 35% and who are intolerant to spironolactone (e.g., gynecomastia, loss of libido, menstrual irregularities)

#### ivabradine restrictions:

Only use ivabradine for:

1. Heart failure patients on ivabradine prior to admission; or

- 2. The treatment of heart failure in patients with the following criteria:
  - a. Reduced left ventricular ejection fraction (LVEF) of 35% or less; and
  - b. New York Heart Association (NYHA) class II or III heart failure symptoms despite at least 4 weeks of optimal treatment with:
    - a stable dose of an ACE inhibitor or an ARB in combination with a beta-blocker; and if tolerated, a MRA; and
  - c. Patients who are in sinus rhythm with a resting heart rate of 77 beats per minute (bpm) or more, using either an ECG or by continuous monitoring; **and**
- 3. Heart rate reduction in computed tomography coronary angiography (CTCA) with the following criteria:
  - a. Use of a beta blocker is deemed unsafe; or
  - b. Target heart rate cannot be achieved despite two beta-blocker doses, which may include an output trial of beta blocker as one of those doses **and**

\*All new starts must be ordered by a specialist in Cardiology or Internal Medicine. Ivabradine should be initiated and titrated under the supervision of a physician who is experienced with the treatment of patients with chronic heart failure

Source: AHS Provincial Drug Formulary

Please refer to Heart Failure guidelines at www.ccs.ca for further information.

Version Date: May 18, 2021

## Alberta Health Services

## Heart Failure Pathway: Transition to Community Care Admission to Discharge Checklist

Last Name (Legal)			First Name (Legal)		
Preferred Name  Last  First			DOB(dd-Mon-yyyy)		
PHN	ULI □ Same as PHN		s PHN	MRN	
Administrative Gender			se (X)	□ Female □ Unknown	

	Date (dd-Mon-yyyy)	Time (hh:mm)	Completed	Not indicated*	Initials			
	Echocardiogram within the past 12 months	Ejection Fraction %						
	<b>Consultations</b> (For all consultations, utilize the most appropriate/available health care provider(s) at your site to deliver services)							
	Screen for Malnutrition							
	Screen for Frailty							
	Screen for Cognitive status							
c	Refer to Transition/Discharge Services if ant	ticipated need at discharge						
dmission	Consider involving the following healthcare Social Worker Speech Language Pathologist for swallow							
at A	Heart Failure (HF) Education and Self-Ca	echnique to	reinforce le	earning				
start at A	Ambulate – Early Mobilization (done within 48)	-						
Sta	Provide and review HF education resources							
	<ul> <li>□ Management Guide</li> <li>□ Nutrition a</li> <li>□ Signs and Symptoms of HF</li> <li>□ Managing</li> <li>□ HF Medicines</li> <li>□ Online Pa</li> <li>□ Weight Chart</li> </ul>							
	Dietitian to provide/arrange for education requestion r							
	Discharge Plan							
	Determine HF Risk and recommended follow Clinic as per HF Risk Stratification (Form 2103		/					
	Complete Heart Failure Discharge Managen	nent Plan <i>(Form 21041)</i>						
	Follow-up as Required							
	Assess tobacco use of patient Provide tobacco cessation counselling ar Refer to tobacco cessation program when							
	Notify Primary Care Provider & Heart Functi (include designated supportive living and home care,	where appropriate)						
	Provide above healthcare providers with Dis HF Discharge Management Plan (Form 21041							

\*Check 'Not Indicated' only if item would NOT benefit the patient. Identify reason by placing a (✓) in the appropriate box.

□ Recently completed

□ End-of-life

□ Deceased

□ Service/assessment is unavailable

□ Other, Specify reason(s): \_

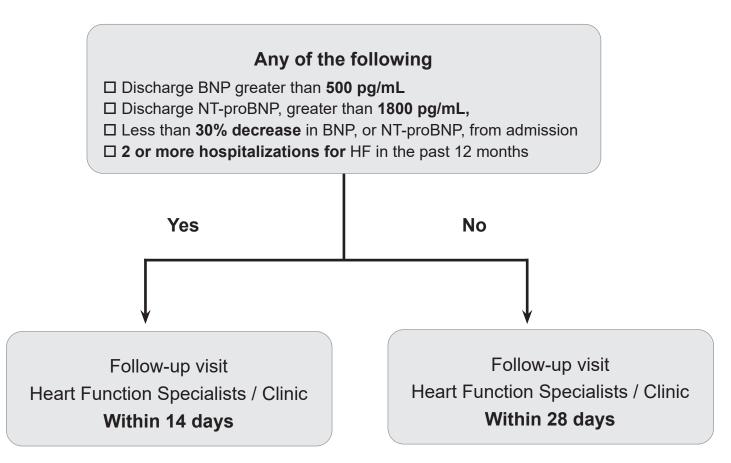


Last Name (Legal)	Name (Legal)		First Name (Legal)		
Preferred Name  La	ast 🗆 First		DOB	(dd-Mon-yyyy)	
PHN	ULI 🗆 Sa	ame as PHN MRN		MRN	
Administrative Geno			se (X)	<ul><li>□ Female</li><li>□ Unknown</li></ul>	

### Heart Failure Risk Stratification

■ Identify recommended time to follow-up visit with Heart Function Clinic/Specialist and record on Heart Failure Discharge Management Plan (*Form 21041*)

Hospital Discharge Heart Failure Patient



## **For All Patients**

Follow Up in Primary Care Clinic Within 14 Days



Heart Failure Discharge Management Plan

Last Name (Legal)		Firs	First Name (Legal)		
Preferred Name □ L	ast 🗆 First		DOB	(dd-Mon-yyyy)	
PHN	ULI □ Same as PHN		s PHN	MRN	
Administrative Gend □Non-binary/Prefer			se (X)	<ul><li>□ Female</li><li>□ Unknown</li></ul>	

		dministrative ]Non-binary/P		□ Male □ Fe to disclose (X) □ Un			
Driver this Mercenerge Plan with we							
Bring this Management Plan with yo           Nutrition         A salt restricted diet of 2000 (1 teaspoonful = 2300 mg)	•	ncouraged					
Medications Prescription given □ No □ Yes							
Discharge medication list faxed to com □ No □ Yes							
(Talk to your doctor or pharmacist before taking	g any non-prescription or her	bal medicines	)				
What you need to know			<b>D</b> .				
<ul> <li>Empty bladder, wear same amount of</li> <li>Recognize the signs of fluid buildup: (</li> </ul>	<ul> <li>Daily Weight Discharge weight:</li></ul>						
	<ul> <li>Monitor for signs and symptoms of heart failure</li> <li>Weight gain, swelling, shortness of breath, fatigue/confusion, persistent coughing or wheezing, heart palpitations, chest pain (angina)</li> </ul>						
<ul> <li>Your medications and the importance</li> <li>Signs, symptoms and actions to ten Plan;</li> </ul>	<ul> <li>Review heart failure patient education handouts. Be familiar with</li> <li>Your medications and the importance of taking medicines as instructed;</li> <li>Signs, symptoms and actions to take for the red, yellow and green zones in your Heart Failure Action Plan;</li> <li>Healthy nutrition and lifestyle choices</li> </ul>						
$\Box$ Driving $\Box$ No restrictions $\Box$ No	strenuous Grad valid license Do n not go back to work for			Do not drive for	weeks		
Follow-up	Location	Phone n	umber	Date (dd-Mon-yyyy)	Time (hh:mm)		
Primary Care Provider (within 14 days of discharge)							
Heart Function Clinic/Specialist within:							
□ 14 days   □ 28 days							
Obtain Influenza and/or oneumococcal vaccines at pharmacy, primary care provider or health clinic f needed							
□ Reviewed above content with patier	nt/family/caregiver and o	copy of form	n provid	led			
Health Care Provider (Last Name, First Name	e)		Desigi	nation	Initial		
Signature			Date (d	dd-Mon-yyyy)	<u> </u>		