

CHFS Heart Failure Order Set Admission Admit to: Dr. to assume MRP □ Diagnosis: Heart Failure ☐ Date of Admission: _____ (yyyy-mm-dd) Allergies or hypersensitivities? ☐ None Known ☐ Yes: Refer to organization's allergy documentation/process □ Code Status: □ Full Resuscitation ☐ DNR ☐ Primary Care Provider: ☐ Inform Primary Care Provider of patient's hospitalization for HF **Precautions** Antibiotic Resistant Organism (ARO) Screening and Management Clinical Protocol Consults Note: If patient at nutritional risk based on the Malnutrition Screening Tool assessment, ensure dietitian consulted. ☐ Cardiologist - Reason: _____ ☐ Pharmacist - Reason: _____ ☐ Dietitian - Reason: ☐ PT for early ambulation RRT - Reason: ☐ Internist - Reason: ☐ SW for discharge planning ☐ OT to screen for frailty Palliative Care Service - Reason: _____ - Reason: _____ **Diet/Nutrition** ☐ NPO, no PO medications ☐ NPO, medications with sips ☐ Diabetic _____ kJ ☐ Cardiac¹ ☐ Renal **Restrictions**: Litres fluid in 24 hours (1.5 or 2 Litres¹) ☐ 2 g Na in 24 hours¹ ☐ ____ **Activity** ☐ Activity as tolerated, encourage early mobilization² ☐ Early ambulation, aim to ambulate three times per day Vitals/Monitoring **Vitals** Weigh patient on admission: Weight: kg kg ☑ Weigh daily in morning¹ after voiding, before breakfast ☑ Vitals, SpO₂, Pain Score as per policy/procedure Submitted by: ☐ Read Back PRINTED NAME YYYY-MM-DD HH:MM Practitioner:

Last Update: 06-20 V3 Page 1 of 14

YYYY-MM-DD HH:MM

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Submitted by:

Practitioner:

Document allergies on approved form and ensure medication reconciliation has been reviewed as per organizational process

CHFS Heart Failure Order Set Vitals/Monitoring Continued... **Monitoring** ☐ Telemetry for 48 hours, then reassess □ CAM score q h and PRN Assess falls risk and implement falls prevention strategies as per policy/procedure **Capillary Blood Glucose Monitoring** ☑ For patient with diabetes, Capillary Blood Glucose monitoring as per applicable diabetes management order set ☐ Capillary Blood Glucose _____ (frequency) Respiratory **Oxygen Therapy** ***Supplemental oxygen is recommended in patients who are hypoxemic to achieve an oxygen saturation greater than 90%***1,3 \square Greater than 92%^{5,6} \square SpO₂: _____ - ____ % ■ 88 - 92%^{4,5} ☐ Target SpO₂: **Patient with Obstructive or Central Sleep Apnea** ☐ Patient to use own PAP machine at patient's prescribed settings^{4,7} after RRT/BioMed equipment check ☑ Request RRT to assess PAP machine, prescribed settings, and to enable O₂ entrainment if O₂ required Lab Investigations Lab Investigations on Admission (if not already done in ED) Hematology, Coagulation \square CBC^{1,3} □ APTT \square INR Ferritin³ ☐ Transferrin saturation³ ☐ Serum iron³ Chemistry Note: Digoxin level may be considered if not performed in the past 6 months. If digoxin level ordered, ensure level is not in toxic range. ☐ Electrolytes^{1,3} \square BNP^{1,3} ☐ A1C^{1,3} ☐ Lactate ☐ Creatinine^{1,3} ☐ Ca^{1,3} ☐ NT-proBNP^{1,3} ☐ TSH^{1,3} ☐ Glucose^{1,3} \square Mg^{1,3} ☐ Troponin³ ☐ Digoxin level Albumin ☐ Uric Acid ☐ ALT, ALP, Bilirubin³ ☐ HDL, LDL, Total Cholesterol, Triglycerides³ ☐ LDH **Lab Investigations Day 2 and Onwards** ***Consider daily electrolytes, creatinine while patient is receiving IV diuretic therapy***8 ☐ Daily Electrolytes, Creatinine

Last Update: 06-20 V3 Page 2 of 14

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CHFS Heart Failure Order Set Diagnostics ☐ 12-Lead ECG^{1,3} ☐ 15-Lead ECG ☐ CXR PA + Lateral^{1,3} Reason: ***Repeat echocardiogram only if no recent assessment of LV function performed in past 12 months and clinical status change warrants investigation*** ☐ Echocardiogram¹⁻³ Reason: ___ Reason: IV Therapy ☐ Saline lock; flush as per policy/procedure mL/h **Heart Failure Medications Diuretics** ***IV diuretics are recommended as first-line therapy for patients with pulmonary or peripheral congestion***1 ***If symptomatic hypotension arises, consider holding diuretics and reassessing for volume overload*** ***Assess daily volume status and manage diuretics accordingly*** ☐ furosemide _____ mg IV for 1 dose STAT (max 200 mg/dose) furosemide _____ mg IV q _____ h ☐ furosemide _____ mg/h IV continuous infusion (5 – 20 mg/h) furosemide _____ mg PO q _____ h bumetanide _____ mg PO q _____ h (0.5 mg; max 10 mg in 24 hours) mg PO q24h, administer 30 minutes prior to loop diuretic (2.5 mg; max 20 mg in 24 hours)² **Angiotensin-Converting Enzyme-Inhibitors (ACEI)** perindopril _____ mg PO q24h (initiation dose 2 – 4 mg; target regimen 4 – 8 mg q24h)¹ ☐ ramipril _____ mg PO q12h (initiation dose 1.25 – 2.5 mg; target regimen 5 mg q12h)¹ Angiotensin Receptor Blockers (ARB) For Patient Intolerant to ACEI andesartan _____ mg PO q24h (initiation dose 4 – 8 mg; target regimen 32 mg q24h)¹⁻³ ☐ valsartan mg PO q12h (initiation dose 40 mg; target regimen 160 mg q12h)¹-3 **Angiotensin Receptor Neprilysin Inhibitors (ARNI)** ***Patients who remain symptomatic despite triple therapy, consider changing ACEI/ARB to an ARNI***1,9 ***Concomitant use with an ACFI or ARB is contraindicated: if an ACEI was administered, wait 36 hours before administering ARNI***1,9 acubitril 24 mg/valsartan 26 mg, 1 tab PO q12h (target regimen sacubitril 97 mg/valsartan 103 mg, 1 tab q12h) accubitril 49 mg/valsartan 51 mg, 1 tab PO q12h (target regimen sacubitril 97 mg/valsartan 103 mg, 1 tab q12h) sacubitril 97 mg/valsartan 103 mg, 1 tab PO q12h Submitted by: Read Back PRINTED NAME YYYY-MM-DD HH·MM Practitioner: PRINTED NAME YYYY-MM-DD HH:MM SIGNATURE

Last Update: 06-20 V3 Page 3 of 14



CHFS Heart Failure Order Set

Heart Failure Medications Continued					
Beta-Blockers					
bisoprolol mg PO q24h (initiation dose 1.25 mg; target regin	nen 10 mg q24h)¹				
carvedilol mg PO q12h	. ,				
(initiation dose 3.125 mg; target regimen 25 mg q12h [if weight greater th	nan 50 kg, target regimen 50 mg q12h])¹				
	G. G G G I 3/				
Mineralocorticoid Receptor Antagonist (MRA)					
pelerenone mg PO q24h (initiation dose 25 mg; target reg	imen 50 mg g2/h\1-3				
spironolactone mg PO q24h (initiation dose 12.5 mg; target reg					
	at regimen so mg qz my				
Vasodilators					
The combination of isosorbide dinitrate and hydralazine are recomme with HFrEF with advanced symptoms or patients unable to	o tolerate ACEI, ARB or ARNI therapy1				
hydralazine mg PO q8h (initiation dose 25 mg; target regin	÷ · /				
And ☐ isosorbide dinitrate mg PO q8h (initiation dose 20	mg; target regimen 40 mg q8h)¹				
Sinoatrial Node Modulator					
***Ivabradine can be considered in patients with HFrEF who are in sinus r					
77 beats per minute and have had a previous HF hospitalization within the of beta-blockers***	e past year despite being at the maximally tolerated dose				
ivabradine mg PO q12h (initiation dose 2.5 – 5 mg¹ [if greater than 75 years old, initial dose 2.5 mg²]	ng]; target regimen 7.5 mg q12h¹)				
Digoxin					
Digoxin may be considered in patients in sinus rhythm who co	ntinue to be symptomatic with triple therapy1,3				
☐ digoxin mg PO q24h (0.125 – 0.25 mg)					
Sodium-glucose Cotransporter 2 (SGLT2) Inhibitor					
***SGLT2 inhibitors should be started once medically stable or upon dis routine therapy in acute hea					
	Dapagliflozin may be considered in patients with mild to moderate HFrEF (LVEF less than/equal to 40%) regardless of concomitant type 2 diabetes ¹¹ (do not use in patients with type 1 diabetes)				
Caution should be exercised when combining SGLT2 inhibitors, ARNI, and diuretics because of their concomitant effects to promote diuresis11					
Note: If serum creatinine is increasing, dapagliflozin should not be initiated	d or be reassessed if initiated.				
☐ dapagliflozin 10 mg PO q24h ¹²					
☐ Initiate on (yyyy-mm-dd) at (hh:mm)					
Electrolyte Management					
Electrolyte Management					
Non-Critical Care Potassium Oral Replacement Clinical Protocol					
Ш					
Out-with-d burn	П				
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Last Update: 06-20 V3 Page 4 of 14



CHFS Heart Failure Order Set

Glycemic Management						
	If applicable, prescriber to complete diabetes management order set					
1	-	t Order Set (NPO Patient)				
	-	t Order Set (Patient Eating Meals	3)			
	emia Managemen	· · · · · · · · · · · · · · · · · · ·	•			
0	0					
Smoking (
	•	atment combined with counselling	g is more effective than pharma	acological treatment alone***13		
	•	apy In-patient Clinical Protocol	analawia Aida la matiant Ordan (C-4		
ii applicable,	prescriber to com	plete Smoking Cessation Pharma	acologic Alds In-patient Order (Sei.		
VTE Propl	hylaxis					
VTE Pharm	acological Pro	phylaxis Not Required				
☐ No pharm	acological prophy	laxis: On therapeutic anticoagula	tion			
☐ No pharm	acological prophy	laxis: Fully mobile and expected	length of stay 24-48 hours and	l no additional risk factors		
	•	laxis: Bleeding/high risk of bleedi	•			
☐ No pharm	acological prophy	laxis - Reason:				
	acological Pro					
☐ Initiate pre	escribed anticoagu	ulant on (yy	yy-mm-dd) at (hh:mm)			
LMWH ¹⁴						
☐ dalteparin	5,000 units Subc	utaneous q24h				
☐ enoxapari	in 40 mg Subcutar	neous q24h				
Unfractionat	Unfractionated Heparin					
☐ heparin 5,	,000 units Subcuta	aneous q h (q8-12h)				
VTE Mecha	nical Prophyla	xis				
	If mechanical	prophylaxis is used alone, reasse	ess daily for conversion to antic	coagulant prophylaxis		
	-	neumatic compression devices14	Į.			
☐ Apply bila	teral, calf-length e	elastic compression stockings ¹⁴				
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Last Update: 06-20 V3 Page 5 of 14



CHFS Heart Failure Order Set

Discharge Planning					
Strategies to redu	***Strategies to reduce readmission rates include early patient discharge planning and scheduling of follow-up appointments prior to discharge1-3				
Note: Refer to heart failure discharge		is prior to discriarge			
Appointments to be Arranged P Arrange for the following appointm Cardiologist/Internist: - Dr. PCP: If patient does not have a PCP, en PCP is available, notify MD/NP for a	ent(s) for patient to be seen sure they are connected to a	within week(s) of within week(s) of within: day(s) a PCP before discharge as pe	discharge discharge s)		
Referrals to be Arranged Prior t If barriers (e.g. financial) to obtaini Smoking Cessation Program Arrange for the following referral(s	ng discharge medication(s) o				
Cardiac Rehabilitation Program Heart Function Clinic Home and community care Palliative care service	Reason: Reason: Reason:	within:	s) week(s)		
Submitted by: ID PRIN	TED NAME	YYYY-MM-DD HH:MM	☐ Read Back		
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Last Update: 06-20 V3 Page 6 of 14



CHFS Heart Failure Order Set						
☑ Discharge☑ Discharge	Discharge ☑ Discharge date: (yyyy-mm-dd)					
New Yor□ Left Ven□ Discharg	tricle Ejection F je Weight:	ation (NYHA) Fraction (LVEF kg	-):%	tion ¹⁶ :		
☑ Ensure dis☐ Ensure a f(24-72 hour	Discharge Information ☑ Ensure discharge Medication Reconciliation process has been completed as per policy/procedure¹7 ☐ Ensure a follow-up phone call to patient/caregiver has been arranged to be done within hours of discharge (24-72 hours)¹8-20					
	copy of the Pat			mation page(s), the patient's cer as per policy/procedure ^{17,20}		
For Community Health Care Providers ☑ Ensure a copy of this document, the patient's care plan, the Discharge Summary, the Medication Reconciliation form, Letter to the PCP and other relevant documents have been provided to the following as per policy/procedure¹7,20-22: ☑ PCP ☐ Home care service ☐ Specialist:						
Patient's community pharmacy ☐ Patient Education and Self-management Initiate and complete the applicable patient education checklist²³ with patient/caregiver and ensure discharge instructions have been provided as per policy/procedure. Use teach-back technique to assess and confirm patient/caregiver understanding²⁴.²⁵ Ensure the following education is provided at a level appropriate for the patient/caregiver¹⁶: Advance care directives						
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Last Update: 06-20 V3 Page 7 of 14



PATIENT AND FAMILY MD INFORMATION

CHFS Heart Failure Discharge Patient Information

This 'Heart Failure Discharge and Patient Information' section is to provide instructions to the patient, and when completed, should be printed and given directly to the patient for their review and to take with them upon discharge.

Instructions

- Review the information provided to you before you go home and again when you arrive home.
- · Bring your Medication Reconciliation form and plan of care to your Pharmacist at your next visit.
- Bring this document, the Medication Reconciliation form and plan of care to your Primary Care Provider, e.g. family doctor or nurse practitioner.

Information

- The Canadian Heart Failure Society Patient Resources: https://heartfailure.ca/education/patient-resources
- The Canadian Cardiovascular Society Heart Failure Program: http://www.ccs.ca/en/guidelines/heart-failure-program
- Heart and Stroke Foundation:
 https://www.heartandstroke.ca/-/media/pdf-files/canada/health-information-catalogue/en-living-with-heart-failure.ashx?rev=3238e9abfabc4027b4b56a042a5d804e&hash=1B4D04630249286D09B8544119E36772
- How to stop smoking: Smokers Helpline: 1-877-513-5333 http://www.smokershelpline.ca
- Finding a family doctor or nurse practitioner: Health Care Connect: 1-800-445-1822 https://www.ontario.ca/page/find-family-doctor-or-nurse-practitioner

Home Care Services

 If home care services arrangements have been s 		
coordinator within hours, please p	hone the following number	per:
• If respiratory services arrangements have been r	nade for you, and you h	ave questions or concerns, please phone
the following number:		
Diagnostic Tests		
		nould have an assessment of LV function within last 12 days from discharge from ED***8
☐ Echocardiogram Reason:		Phone Number:
Arranged by hospital: Date:	Time:	or ☐ Patient will be notified
☐ Patient to arrange test. Test to be done in	week(s)	or month(s)
Cardiac MRI Reason:		Phone Number:
Arranged by hospital: Date:	Time:	or ☐ Patient will be notified
☐ Patient to arrange test. Test to be done in	week(s)	or month(s)
☐ Patient to arrange test. Test to be done in	week(s)	or month(s) Phone Number:
☐ Patient to arrange test. Test to be done in ☐ Arranged by hospital: Date:		Phone Number:

Last Update: 06-20 V3 Page 8 of 14



PATIENT AND FAMILY MD INFORMATION

CHFS Heart Failure Discharge Patient Information

Appointments		
☐ Heart Failure Clinic:		Phone Number:
☐ Arranged by hospital: Date:	Time:	_ or ☐ Patient will be notified
☐ Patient to arrange appointment to be seen in _	day(s)	or week(s)
		Phone Number:
☐ Arranged by hospital: Date:	Time:	_ or ☐ Patient will be notified
☐ Patient to arrange appointment to be seen in _	day(s)	or week(s)
☑ Cardiologist/Internist - Dr		Phone Number:
☐ Arranged by hospital: Date:	Time:	_ or ☐ Patient will be notified
\square Patient to arrange appointment to be seen in $_$	day(s)	or week(s)
☑ Cardiac Rehabilitation Program:		Phone Number:
☐ Arranged by hospital: Date:	Time:	or ☐ Patient will be notified
☐ Patient to arrange appointment to be seen in _	day(s)	or week(s)
☐ Diabetes Clinic:		Phone Number:
☐ Arranged by hospital: Date:		
☐ Patient to arrange appointment to be seen in _	day(s)	or week(s)
☐ Smoking Cessation Program:		Phone Number:
☐ Arranged by hospital: Date:	Time:	_ or ☐ Patient will be notified
☐ Patient to arrange appointment		
		Phone Number:
Arranged by hospital: Date:	Time:	or Patient will be notified
☐ Patient to arrange appointment to be seen in _	day(s)	or week(s)

Last Update: 06-20 V3 Page 9 of 14



Order Set Development and Implementation Consideration

The CHFS acknowledges the partnership with Think Research and the important contribution of the following hospitals' heart failure order sets in the development of the present document: Alberta Health Services, the St-Boniface Hospital (Winnipeg), and the Sunnybrook Health Sciences Centre (Toronto).

Updated

This order set was last updated in May 2020.

Abbreviations

ACEI = Angiotensin-Converting Enzyme Inhibitor

ARB = Angiotensin II Receptor Blocker

BioMed = Biomedical Engineering

BNP = Brain Natriuretic Peptide

CAM = Confusion Assessment Method

DKA = Diabetic Ketoacidosis

ED = Emergency Department

EF = Ejection Fraction

GFR = Glomerular Filtration Rate

HF = Heart Failure

HFrEF = Heart Failure with Reduced Ejection Fraction

LV = Left Ventricle

LVEF = Left Ventricle Ejection Fraction

NT-proBNP = Prohormone of BNP

PAP = Positive Airway Pressure

PCP = Primary Care Provider

Patient Care Considerations

- Antiplatelet Therapy: Antiplatelet therapy (e.g. acetylsalicylic acid) is recommended in patients with HF who have had or at risk for atherosclerotic cardiovascular events.¹
- **BNP and NT-proBNP:** BNP and NT-proBNP are natriuretic peptide (NP) biomarkers that are used to establish the presence and severity of HF.⁹ NP screening can be helpful in establishing if a patient is at risk for HF and if echocardiography is necessary. The following table provides information regarding the NP levels and diagnosis of HF¹:

	Age	HF is unlikely	HF is possible but alterr diagnoses to be consid	HE IS VERV IIKEIV I
BNP	All	Less than 100 pg/mL	100 – 150 pg/mL	Greater than 500 pg/mL
NT-proBNP	Less than 50		300 – 450 pg/mL	Greater than 450 pg/mL
	50 – 75	Less than 300 pg/mL	450 – 900 pg/mL	Greater than 900 pg/mL
	Greater than 75	⁷ 5	900 – 1800 pg/mL	Greater than 1800 pg/mL

Note: BNP levels may increase early after initiation of ARNI therapy as BNP is a substrate for neprilysin. ^{9,29} Prognostic value of BNP typically resumes after the first 6 months of therapy. Though NT-proBNP is not a substrate of neprilysin, its level may lower early after initiation of ARNI therapy; however, it retains its prognostic value during this time. ^{29,30}

- Choosing an ACEI in HF: Determining which ACEI to prescribe in patients with HF depends on several factors, including: ejection fraction (EF), stroke volume (SV), systolic blood pressure (SBP), diastolic blood pressure (DBP), mean arterial pressure (MAP), renal function, adverse effects, and mortality. Many factors go into determining which ACEI to choose for patients with HF and more research needs to be done to determine if there is an ACEI that is superior to others, particularly in reducing rehospitalization and cardiac death.³¹
- **Discharge Checklist:** Key considerations for discharging a patient include the following1:
 - · Symptoms and Disease:
 - Intercurrent cardiac illness adequately diagnosed and treated
 - · Presenting symptoms resolved
 - Chronic oral HF therapy initiated, titrated, and optimized (or plan for same)

Last Update: 06-20 V3 Page 10 of 14



· Stability:

- Return to "dry" weight and stable for greater than 24 hours
- · Vital signs resolved and stable for greater than 24 hours, especially blood pressure and heart rate
- Greater than 30% decrease in natriuretic peptide level from time of admission and relatively free from congestion

Transition:

- Communication to primary care provider and/or specialist physician and/or multidisciplinary disease management program (ideally patient to be seen by cardiologist or internist within 7 days of leaving the hospital)
- Clear discharge plan for laboratory tests, follow-up, and other testing
- Education initiated, understood by patient, and continued education planned; this includes:
- · Formal education session on HF management for patient and family members
- · Education on controlling sodium intake, weighing self, and recognizing symptoms of worsening HF
- Education on algorithms to adjust diuretics in patients with recurrent fluid retention
- MRAs and Potassium and Kidney Function: MRAs can increase serum potassium, especially when a patient has a dehydrating illness where renal dysfunction can worsen. This requires patients to have kidney function (e.g. creatinine, GFR) and potassium, to be closely monitored when on these medications.¹
- Patients at Risk for Hypercapnia⁴: Chronic Obstructive Pulmonary Disease (COPD) is the most common disease to cause hypercapnia⁴; other patients at risk for hypercapnic respiratory failure include those with cystic fibrosis (CF), non-CF bronchiectasis (often in association with COPD or severe asthma), severe kyphoscoliosis or severe ankylosing spondylitis, severe lung scarring from old tuberculosis (especially with thoracoplasty), morbid obesity (BMI > 40 kg/m²), musculoskeletal disorders with respiratory muscle weakness (on home mechanical ventilation), overdose of opioids, benzodiazepines or other respiratory depressant drugs. The target SpO₂ in patients with COPD who are at risk of hypercapnia is 88-92%. The target SpO₂ in patients with other risk factors for hypercapnia is 88-92%; this is based on expert opinion which was extrapolated from observational studies.
- **SGLT2 Inhibitor:** The Canadian Cardiovascular Society (CCS) recommend SGLT2 inhibitors, such as dapagliflozin, be used in patients with mild to moderate HF due to reduced LVEF (less than/equal to 40%) and without concomitant diabetes, to improve symptoms and quality of life and to reduce the risk of hospitalization and cardiovascular mortality (Conditional Recommendation, High-Quality Evidence).¹¹ This recommendation is based of the results of the Dapagliflozin on Incidence of Worsening Heart Failure or Cardiovascular Death in Patients with CHF (DAPA-HF) trial.¹²
- Sleep Apnea: Obstructive sleep apnea (OSA) and central sleep apnea (CSA) are the main types of sleep disordered breathing (SDB). Around 40% of patients with HF have CSA and 11% have OSA. Many patients with HF with SDB go undiagnosed, likely due to limited resources and awareness. It is recommended that clinicians treating patients with HF refer to experienced sleep physicians and sleep laboratories to help differentiate between OSA and CSA.¹
- Supplemental O₂ and Target Ranges: In acutely ill adults, evidence shows that liberal O₂ therapy increases mortality without improving other patient-important outcomes. Supplemental O₂ might become unfavourable above an SpO₂ range of 94-96%.³² A systematic review and meta-analysis by Chu et al shows that patients treated liberally with O₂ had a dose-dependent increased risk of short-term and long-term mortality.³² Individual randomised controlled trials have suggested an increased risk of respiratory failure, new shock episodes, recurrent myocardial infarction, arrhythmia, and other cardiovascular adverse events as potential mechanisms of harm with liberal O₂ therapy.³² An upper level of 96% avoids the potential risks of hyperoxia and allows for patient improvement to be recognized earlier during monitoring so that O₂ can be down-titrated.⁶

Last Update: 06-20 V3 Page 11 of 14



Administration/Organizational Considerations

- Advance Care Planning Discussions: Advance Care Planning discussions may be undertaken by different health care
 professionals, e.g. MD/NP, nurse, SW. Localization of this content will involve alignment with the facility's resources,
 workflows, and policy/procedure. Advance Care Planning discussions may trigger a process leading to the creation of a
 separate legal Advance Directive document.³³ If the patient has an Advance Directive, it should be incorporated into the
 patient's Advance Care Plan in alignment with the facility's policy/procedure, and applicable law.
- Code Status: Facilities should localize code status orders in alignment with policy/procedure and applicable law.
- **Malnutrition Screen:** Facilities should have a process in place to screen all patients for malnutrition with a simple assessment tool such as the Canadian Nutrition Screening Tool.
- **Risk Scores:** HF prognostic and risk scores can be easily accessed and calculated, and when possible should be incorporated into practice. Organizations are to determine what risk scores are appropriate in their setting and implement them into practice as they see fit.

References

Key references 1-33

All medication guidance has been reviewed using Lexicomp and Compendium of Pharmaceuticals and Specialties (eCPS).

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Last Update: 06-20 V3 Page 12 of 14



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Last Update: 06-20 V3 Page 13 of 14



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Last Update: 06-20 V3 Page 14 of 14