

CHFS Heart Failure Order Set Admission Admit to: to assume MRP □ Diagnosis: Heart Failure ☐ Date of Admission: (yyyy-mm-dd) Allergies or hypersensitivities? \(\square\) None Known Yes: Refer to organization's allergy documentation/process ☐ Code Status: ☐ Full Resuscitation \square 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 Internist - Reason: RRT - Reason: ☐ OT to screen for frailty ☐ SW for discharge planning 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 \Box _ 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

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YYYY-MM-DD HH:MM

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Vitale/Bitale	n Oontineerd							
Vitals/Monitoring Continued								
Monitoring								
	Telemetry for 48 hours, then reassess							
	CAM score q h and PRN							
☐ Intake and Output	•		, ,					
	nd implement falls preventio	n strategies as per polic	y/procedure					
Capillary Blood Glu								
	abetes, Capillary Blood Gluc		pplicable diabetes manag	ement order set				
Сарінату віоосі G	lucose	(irequericy)						
Respiratory								
Oxygen Therapy								
		atients who are hypoxen	nic to achieve an oxygen s	saturation greater than 90%*** ^{1,3}				
☐ Target SpO₂:	☐ 88 - 92% ^{4,5}	☐ Greater than 92%	^{5,6} SpO ₂ :	%				
	Clinical Protocol							
Patient with Obst	ructive or Central Sleep	Apnea						
☐ Patient to use own	n PAP machine at patient's p	rescribed settings ^{4,7} afte	er RRT/BioMed equipmen	t check				
□ Request RRT	to assess PAP machine, pre	scribed settings, and to	enable O ₂ entrainment if 0	O ₂ required				
Lab Investigation	ns s on Admission (if not al	ready done in ED)						
Hematology, Coagu		ready done in ED)						
☐ CBC ^{1,3}	APTT	□INR	П					
☐ Ferritin ³	☐ Transferrin saturation ³	☐ Serum iron ³	<u> </u>					
Chemistry								
_	nay be considered if not perfo	ormed in the past 6 mon	ths. If digoxin level ordere	ed, ensure level is not in toxic				
range.	,							
☐ Electrolytes ^{1,3}	☐ Lactate	☐ BNP ^{1,3}	☐ A1C ^{1,3}					
☐ Creatinine ^{1,3}	Ca ^{1,3}	☐ NT-proBNP ^{1,3}	☐ TSH ^{1,3}					
☐ Glucose ^{1,3}	☐ Mg ^{1,3}	☐ Troponin ³	☐ Digoxin level					
Albumin	☐ ALT, ALP, Bilirubin ³	Uric Acid						
☐ HDL, LDL, Total Cholesterol, Triglycerides³ ☐ LDH ☐								
Lab Investigations	s Day 2 and Onwards							
	Consider daily electroly	tes, creatinine while pati	ent is receiving IV diuretion	c therapy ⁸				
☐ Daily Electrolytes,	Creatinine							
Submitted by:)000(1M15511111	☐ Read Back				
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Diagnostics					
Diagnostics					
12-Lead ECG ^{1,3}					
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					
☐ Echocardiogram ^{1–3} Reason:					
Reason:					
IV Therapy					
☐ Saline lock; flush as per policy/procedure ☐ at mL/h					
Heart Failure Medications					
Diuretics					
IV diuretics are recommended as first-line therapy for patients with pulmonary or peripheral congestion1					
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)					
metolazone 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					
☐ candesartan 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 ARNI1,9					
***Concomitant use with an ACEI or ARB is contraindicated;					
if an ACEI was administered, wait 36 hours before administering ARNI***1,9					
sacubitril 24 mg/valsartan 26 mg, 1 tab PO q12h (target regimen sacubitril 97 mg/valsartan 103 mg, 1 tab q12h)					
sacubitril 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:					
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Hoart Failure Mediceti	one Continued						
Heart Failure Medicati	ons continued						
Beta-Blockers hisoprolol me	n PO d24h (initiation dose 1 25 ma	ı: target regimen 10 mg g24h\1					
 □ bisoprolol mg PO q24h (initiation dose 1.25 mg; target regimen 10 mg q24h)¹ □ carvedilol mg PO q12h 							
(initiation dose 3.125 mg; target regimen 25 mg q12h [if weight greater than 85 kg, target regimen 50 mg q12h]) ¹							
Mineralocorticoid Recep	tor Antagonist (MRA)						
•	g PO q24h (initiation dose 25 mg;	target regimen 50 mg q24h) ^{1–3}					
	☐ spironolactone mg PO q24h (initiation dose 12.5 mg; target regimen 50 mg q24h)¹						
Vasodilators							
***The combination of isoson		recommended in addition to stand s unable to tolerate ACEI, ARB or A	ard treatment for black patients with				
	p PO q8h (initiation dose 25 mg; ta		Artivi iliciapy				
· ·		ation dose 20 mg; target regimen 4	0 mg q8h) ¹				
Sinoatrial Node Modulate	or						
	had a previous HF hospitalization		art rate of greater than or equal to 77 at the maximally tolerated dose of				
ivabradine mg		-piockers					
	-	al dose 2.5 mg]; target regimen 7.5	5 mg q12h¹)				
Digoxin	, ,	31 , 3	3 .				
	considered in patients in sinus rhy	thm who continue to be symptoma	atic with triple therapy***1,3				
digoxin mg P	•	, , , , , , , , , , , , , , , , , , , ,					
Sodium-glucose Cotrans	sporter 2 (SGLT2) Inhibitor						
	d be started once medically stable	or upon discharge given risks of e	euglycemic DKA; not indicated for				
Danagliflazin may be sone	• •	in acute heart failure	al to 40%) regardless of concomitant				
Dapagiiiloziii may be cons		e in patients with type 1 diabetes)*					
***Ca		ombining SGLT2 inhibitors, ARNI,					
Note: If serum creatinine is in		ant effects to promote diuresis***11 be initiated or be reassessed if ini					
dapagliflozin 10 mg PO q2			ilatou.				
	(yyyy-mm-dd) at	(hh:mm)					
Electrolyte Manageme		LI					
☐ Non-Critical Care Potassii	um Oral Replacement Clinical Pro	(OCOI					
Submitted by:			□ Pood Pook				
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CHFS Heart Failure Order Set

Glycemic Management								
If applicable, prescriber to comple	te diabetes management order s	et						
Diabetes Insulin Management Order Set (NPO Patient)								
Diabetes Insulin Management Order Set (Patient Eating Meals)								
	☑ Hypoglycemia Management Clinical Protocol							
Smoking Cessation								
	ment combined with counselling i	is more effective than phermose	plogical treatment alone***13					
☐ Nicotine Replacement Therap		is more enective than pharmact	biogical treatment alone					
If applicable, prescriber to comple	•	Nogic Aids In nationt Order Set						
applicable, presenber to comple		nogic Aids in-patient Order Oct.						
VTE Prophylaxis								
VTE Pharmacological Proph	ylaxis Not Required							
☐ No pharmacological prophylax	is: On therapeutic anticoagulation	n						
☐ No pharmacological prophylax	is: Fully mobile and expected len	gth of stay 24-48 hours and no	additional risk factors					
☐ No pharmacological prophylax	is: Bleeding/high risk of bleeding							
☐ No pharmacological prophylax	is - Reason:							
VTE Pharmacological Proph	ylaxis							
☐ Initiate prescribed anticoagula		m-dd) at (hh:mm)						
LMWH ¹⁴		,, , ,						
dalteparin 5,000 units Subcuta	ineous a24h							
enoxaparin 40 mg Subcutaneo	•							
Unfractionated Heparin	·							
heparin 5,000 units Subcutane	eous a h (a8-12h)							
VTE Mechanical Prophylaxis								
Apply bilateral intermittent pne	rophylaxis is used alone, reasses	s daily for conversion to anticoa	aguiant propnylaxis***					
☐ Apply bilateral, calf-length elas	•							
Apply bilateral, call-length elas	atic compression stockings							
Submitted by:			□ Pood Pook					
Submitted by:	PRINTED NAME	YYYY-MM-DD HH:MM	☐ Read Back					
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CHFS Heart Failure Order Set

Discharge Planning						
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 checklist.						
Appointments to be Arranged Pr	rior to Discharge					
☐ Arrange for the following appointme	ent(s) for patient to be seen ¡	post-discharge within the time fr	ame specified below:			
☐ Cardiologist/Internist: - Dr		within week(s) of di	ischarge			
☐ PCP:		within week(s) of di	scharge			
		- , ,	week(s)			
☐ If patient does not have a PCP, ens			olicy/procedure. If no			
PCP is available, notify MD/NP for	alternate provision of care a	rrangements				
Referrals to be Arranged Prior to						
If barriers (e.g. financial) to obtaining	g discharge medication(s) o	r equipment, 15 arrange referral	to:			
☐ Smoking Cessation Program						
Arrange for the following referral(s)		-	·			
☐ Cardiac Rehabilitation Program☐ Heart Function Clinic	·		week(s)			
☐ Home and community care	Reason: Reason:	within: day(s) within: day(s)	week(s) week(s)			
Palliative care service	Reason:	within: day(s)				
	_	· · ·	week(s)			
-			_			
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Practitioner:

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

CHFS Heart Failure Order Set Discharge _____ (yyyy-mm-dd) □ Length of stay (LOS): day(s) □ Discharge date: □ ☑ Discharge patient to: Home ☐ Complex Continuing Care ☐ Long Term Care Discharge diagnosis: Comorbidities: **Clinical Assessment at Discharge** New York Heart Association (NYHA) Functional Classification ¹⁶: ☐ Class I ☐ Class II ☐ Class III ☐ Class IV Left Ventricle Ejection Fraction (LVEF): ______ % ☐ Discharge Weight: _____ kg ☐ K: _____ ☐ Na: ____ Lab Values: Creatinine: **Discharge Information** ☐ Ensure discharge Medication Reconciliation process has been completed as per policy/procedure 17 Ensure a follow-up phone call to patient/caregiver has been arranged to be done within hours of discharge (24-72 hours)18-20 **For Patient** Ensure a copy of the Patient Discharge and Follow-up Information page(s), the patient's care plan, and the Medication Reconciliation form have been provided to the patient/caregiver as per policy/procedure 17,20-22 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/procedure 17,20-22: □ 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 understanding^{24,25} Ensure the following education is provided at a level appropriate for the patient/caregiver¹⁶: Advance care directives Definition of heart failure and cause Physical activity/exercises²⁶ Daily weight monitoring Diuretic monitoring and management Self-management • Diet, e.g. nutrition, fluid, salt restriction • Lifestyle, e.g. alcohol, smoking **Smoking Cessation** Heart failure risk modification Medication management · Symptoms of worsening heart failure · When to seek medical attention, e.g. specific symptoms or weight changes Advise patient to talk to their PCP about recommended vaccinations ☑ Provide applicable written education materials²⁷ in patient's preferred language and review with patient/caregiver as per policy/procedure^{21,28} Submitted by: Read Back PRINTED NAME YYYY-MM-DD HH:MM

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PRINTED NAME



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: <a href="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

	J		,	,			ed by your home care
 If respiratory 		nents have been ma	ade for you	, and you h	ave questi	ons or conc	erns, please phone
	HF seen in the ED of a	admission date or p	lanned with	nin 30 days	from disch	arge from E	
	ıram Reason:						t will be matified
_	I by hospital: Date:					-	t will be notified
	o arrange test. Test -					<u> </u>	month(s)
☐ Cardiac MR							
	I by hospital: Date:					_	t will be notified
∐ Patient to	o arrange test. Test	to be done in	W	eek(s)	or		month(s)
					Phone N	Number:	
☐ Arranged	l by hospital: Date:		Time:		or	☐ Patien	t will be notified
☐ Patient to	o arrange test. Test	to be done in	w	eek(s)	or		month(s)
Submitted by:							☐ Read Back
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CHFS Heart Failure Discharge Patient Information

Appointments						
⊠ Heart Failure Clinic:			Phone	Number:		
☐ Arranged by hospital: Date:		_Time:	or	☐ Patien	t will be notified	
☐ Patient to arrange appointm	ent to be seen in	day(s)	or		week(s)	
			Phone	Number:		
☐ Arranged by hospital: Date:	·		or	☐ Patien	t will be notified	
☐ Patient to arrange appointm	ent to be seen in	day(s)	or		week(s)	
⊠ Cardiologist/Internist - Dr			Phone	Number:		
☐ Arranged by hospital: Date:		Time:	or	☐ Patien	t will be notified	
☐ Patient to arrange appointm	ent to be seen in	day(s)	or		week(s)	
□ Cardiac Rehabilitation Program	n:		Phone	Number:		
☐ Arranged by hospital: Date:				☐ Patien	t will be notified	
☐ Patient to arrange appointm	ent to be seen in	day(s)	or		week(s)	
☐ Diabetes Clinic:			Phone			
Arranged by hospital: Date:					t will be notified	
☐ Patient to arrange appointm			or		week(s)	
☐ Smoking Cessation Program: _		- , ,		Number:		
Arranged by hospital: Date:				·	t will be notified	
☐ Patient to arrange appointm					. Will be fround	
			Phono	Numbor		
Arranged by hospital: Date:					t will be notified	
☐ Patient to arrange appointm		-	or		week(s)	
ration to arrange appointm		uay(o)	0.		#6614(6)	
Submitted by: ID Practitioner:	PRINTED NAME		YYYY-MM-	DD HH:MM	☐ Read Back	
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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 GFR = Glomerular Filtration Rate

ARB = Angiotensin II Receptor Blocker HF = Heart Failure

BioMed = Biomedical Engineering HFrEF = Heart Failure with Reduced Ejection Fraction

BNP = Brain Natriuretic Peptide LV = Left Ventricle

CAM = Confusion Assessment Method

DKA = Diabetic Ketoacidosis

NT-proBNP = Prohormone of BNP

ED = Emergency Department

EF = Ejection Fraction

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 alternative diagnoses to be considered	HF is very likely
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		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)

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· 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). 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

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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 references¹⁻³³

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

- 1. Ezekowitz JA, O'Meara E, McDonald MA, et al. 2017 Comprehensive update of the Canadian Cardiovascular Society guidelines for the management of heart failure. Can J Cardiol. 2017;33(11):1342-1433. doi:10.1016/j.cjca.2017.08.022.
- 2. 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(16):e147-239. doi:10.1016/j.jacc.2013.05.019.
- 3. Ponikowski P, Voors AA, Anker SD, et al. 2016 ESC Guidelines for the diagnosis and treatment of acute and chronic heart failure: the Task Force for the diagnosis and treatment of acute and chronic heart failure of the European Society of Cardiology (ESC). Developed with the special contribution of the Heart Failure Association (HFA) of the ESC. Eur Heart J. 2016;37(27):2129-2200. doi:10.1093/eurheartj/ehw128.
- 4. O'Driscoll BR, Howard LS, Earis J, et al. BTS guideline for oxygen use in adults in healthcare and emergency settings. Thorax. 2017;72(Suppl 1):i1-i90.
- 5. Siemieniuk RAC, Chu DK, Kim LH-Y, et al. Oxygen therapy for acutely ill medical patients: a clinical practice guideline. BMJ. October 2018:k4169. doi:10.1136/bmj.k4169.
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