

# Disclosures

## Consulting/Advisory Board:

Abbott, Akcea, Astra Zeneca, Amgen, Alnylam, Boehringer Ingelheim, Cardiol Therapeutics, Merck, Novartis, Pfizer, Servier

## Speaker:

Astra Zeneca, Boehringer Ingelheim, Eli- Lilly, Novartis, Servier

## Clinical Trials:

Amgen, Astra Zeneca, Bayer, Boehringer Ingelheim, Merck, Novartis

## Research Grants:

Novartis

## Educational Grants:

Servier



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Canadian Heart Failure Society  
Société canadienne d'insuffisance cardiaque

# CCS CHFS Clinical Practice Update HF Phenotype Pathway *Preview*

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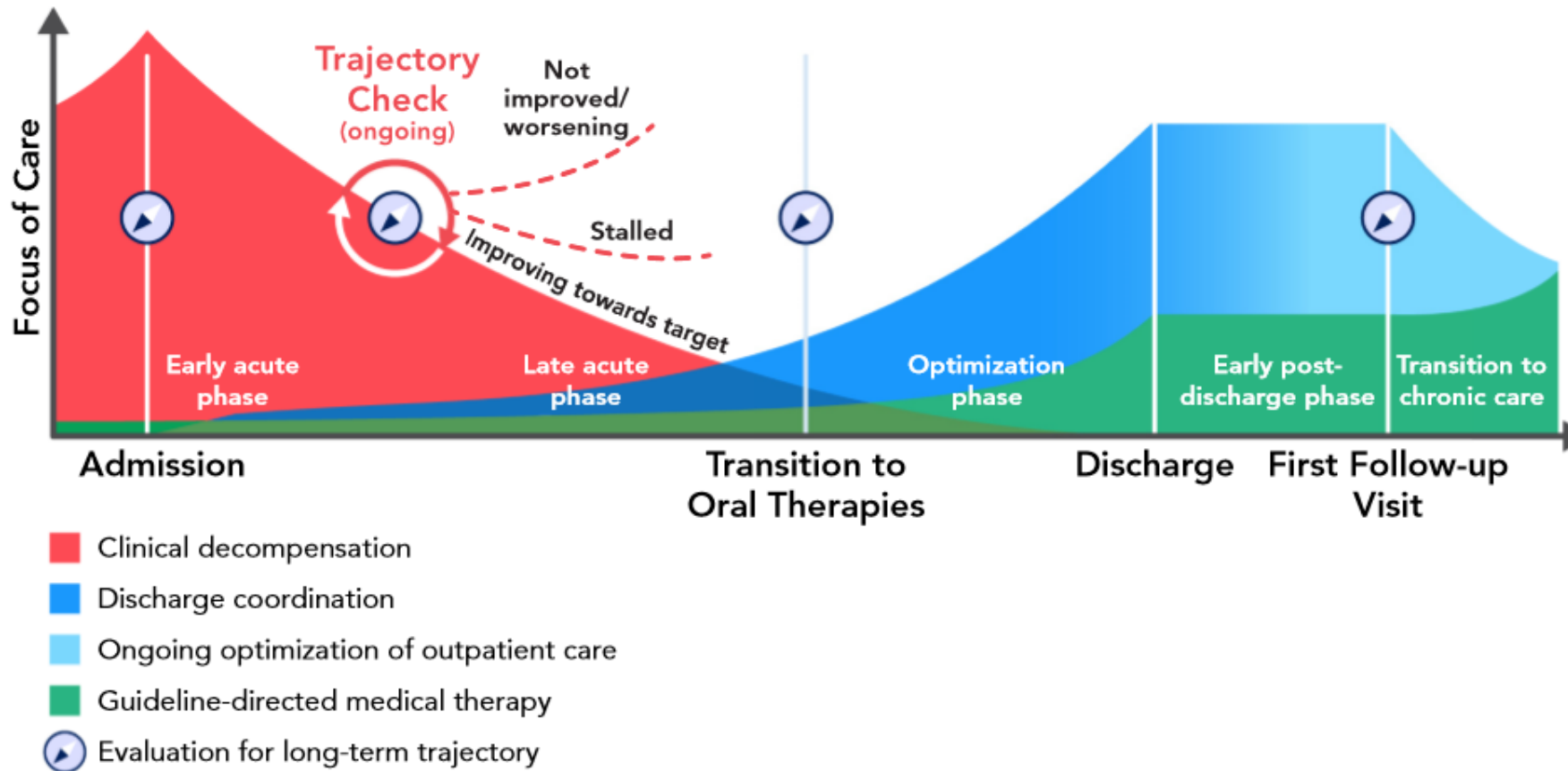
# Objectives

- ▶ Preview/*crowdsource* the CHFS phenotype pathway/algorithm for the practical application of HF therapies
- ▶ Reveal the CHFS inpatient HF order set, discharge tool and patient diary

## NOTE:

Final CCS CHFS CPU HF Phenotype Pathway will be e-published in the **Canadian Journal of Cardiology** simultaneous with the CHFS Spotlight podium presentation at the **Canadian Cardiovascular Congress**, October 22-25, 2020

# Clinical Course of Heart Failure

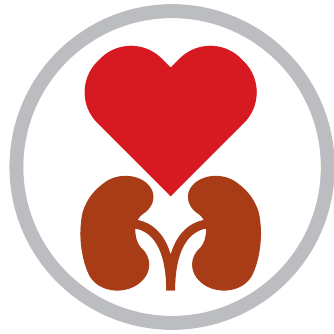


2019 ACC Expert Consensus Decision Pathway on Risk Assessment, Management, and Clinical Trajectory of Patients Hospitalized With Heart Failure  
 Steven M. Hollenberg, Lynne Warner Stevenson, Tariq Ahmad, Vaibhav J. Amin, Biykem Bozkurt, Javed Butler, Leslie L. Davis, Mark H. Drazner, James N. Kirkpatrick, Pamela N. Peterson, Brent N. Reed, Christopher L. Roy, Alan B. Storrow, J Am Coll Cardiol. 2019 Oct, 74 (15) 1966-2011

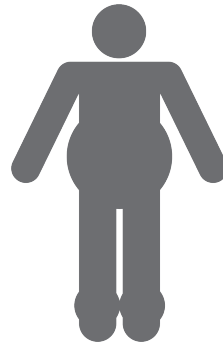
# Phenotypes



T2DM



CRS/Stalled



Wet  
Hypertensive



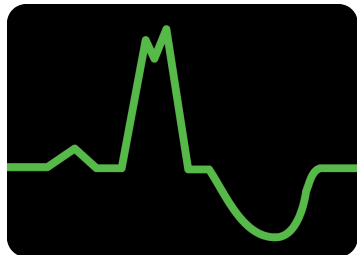
Elderly



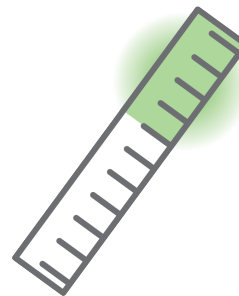
Afib



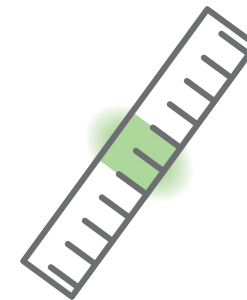
High  
Heart Rate



LBBB



HFpEF



HFmrEF



De novo



# Type 2 Diabetes

- ▶ In a patient with Type 2 DM and LVEF 35% and NYHA 2-3 symptoms how would you prioritize their GDMT (assuming Health Canada/ regulatory approval)

- A) RASi    BB    MRA    SGLT2i
- B) RASi    SGLT2i    BB    MRA
- C) SGLT2i    RASi    BB    MRA

RASi = (Ace or ARB or ARNi)

BB = (B Blocker)

MRA = (Mineralocorticoid receptor antagonist)

SGLT2i = (Sodium glucose co-transporter 2 inhibitors)



# Type 2 Diabetes

- ▶ In a patient with Type 2 DM and LVEF 35% and NYHA 2-3 symptoms how would you prioritize their GDMT (assuming Health Canada/ regulatory approval)

What did the majority around the CPU table say?

A) RASi BB MRA SGLT2i

B) RASi SGLT2i BB MRA

C) SGLT2i RASi BB MRA



# How About Without Type 2 Diabetes ?

- ▶ In a patient with LVEF 35% and NYHA 2-3 symptoms how would you prioritize their GDMT (assuming Health Canada/ regulatory approval)

- A) RASi    BB    MRA    SGLT2i
- B) RASi    SGLT2i    BB    MRA
- C) SGLT2i    RASi    BB    MRA

RASi = (Ace or ARB or ARNi)

BB = (B Blocker)

MRA = (Mineralocorticoid receptor antagonist)

SGLT2i = (Sodium glucose co-transporter 2 inhibitors)



# How About Without Type 2 Diabetes ?

- ▶ In a patient with LVEF 35% and NYHA 2-3 symptoms how would you prioritize their GDMT (assuming Health Canada/ regulatory approval)
  - ▶ What did the majority around the CPU table say?
- 
- A) RASi BB MRA SGLT2i
  - B) RASi SGLT2i BB MRA
  - C) SGLT2i RASi BB MRA



# How About Without Type 2 Diabetes ?

- ▶ In a patient with LVEF 35% and NYHA 2-3 symptoms how would you prioritize their GDMT (assuming Health Canada/ regulatory approval)
  - ▶ In the last case which 2 drugs would you start simultaneously?
- 
- A) RASi BB MRA SGLT2i
  - B) RASi SGLT2i BB MRA
  - C) SGLT2i RASi BB MRA
  - D) SGLT2i RASi BB MRA



# High Heart Rate

- ▶ In a euvolemic patient with a BP of 90/60 on metoprolol 100 mg po BID and a Heart Rate of 98 bpm (sinus) LVEF 35% and NYHA 2-3 symptoms how would you address the residual risk of elevated heart rate?
- A) Uptitrate Bblocker
  - B) Add ivabradine



# High Heart Rate

- ▶ In a euvolemic patient with a BP of 90/60 on metoprolol 100 mg po BID and a Heart Rate of 98 bpm (sinus) LVEF 35% and NYHA 2-3 symptoms how would you address the residual risk of elevated heart rate?
  - ▶ What did the majority around the CPU table say?
- A) Uptitrate Bblocker
- B) Add ivabradine



# De novo HFrEF



▶ In a patient admitted with newly diagnosed ADHF with LVEF 35% how would you prioritize their in hospital GDMT (assuming **NO** regulatory restriction) ?

- A) ACEi    BB    MRA then switch to ARNI as outpatient
- B) ACEi    BB    MRA no switch to ARNI as outpatient until reassess LVEF
- C) ARNI    BB    MRA

ACEi = Ace Inhibitor

BB = Beta blocker

MRA = Mineralcorticoid receptor antagonis

ARNI = Angiotensin receptor neprilysin inhibitor



# De novo HFrEF



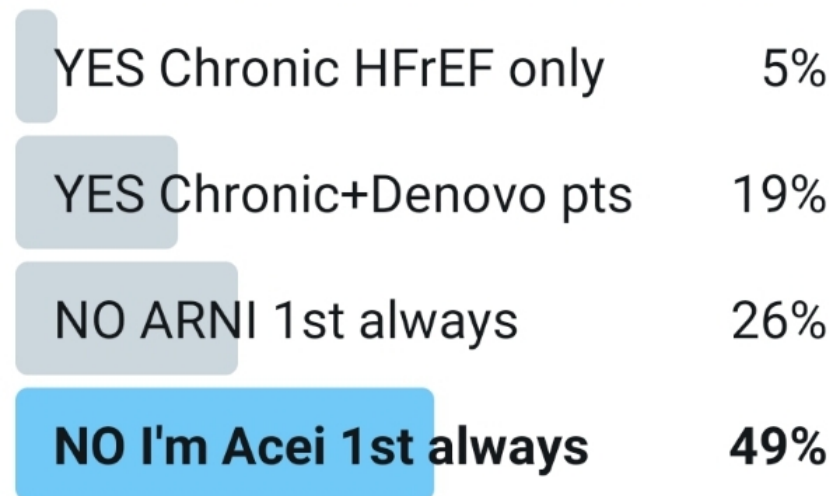
- ▶ In a patient admitted with newly diagnosed ADHF with LVEF 35% how would you prioritize their in hospital GDMT (assuming **NO** regulatory restriction) ?
  - ▶ What did the majority around the CPU table say?
- 
- A) ACEi    BB    MRA then switch to ARNI as outpatient
  - B) ACEi    BB    MRA no switch to ARNI as outpatient until reassess LVEF
  - C) **ARNI    BB    MRA**



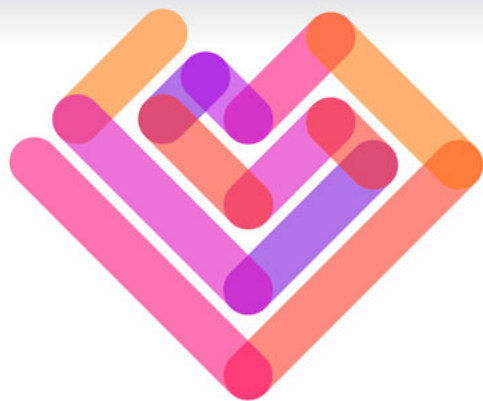
**Shelley Zieroth** @ShelleyZie... · 1d ✓

🌟 Polltime pre [HFUpdate.ca](https://HFUpdate.ca) next wknd: Did [#covid19](#) 📈 your use of ARNI first line over Acei to reduce hosp+ HCU in HFrEF pts?

[@IAmDrIbrahim](#) [@JJheart\\_doc](#)  
[@GiuseppeGalati\\_](#) [@ValleAlfonso](#)  
[@hvanspall](#) [@gcfmd](#) [@BiykemB](#)  
[@AndrewJSauer](#) [@DrRajivsankar](#)  
[@mmamas1973](#) [@DrMarthaGulati](#)



50% already use  
ARNI 1<sup>st</sup> line



# CollabCare



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

CHFS Heart Failure Order Set	ACTION
<b>Diagnostics</b> <b>Investigations on Admission (if not already done in ED)</b> <input type="checkbox"/> 12-Lead ECG <sup>1,2</sup> <input type="checkbox"/> 15-Lead ECG <input type="checkbox"/> CXR PA + Lateral <sup>1,2</sup> Reason: _____ <b>***Repeat echocardiogram only if no recent cardiac imaging performed in past 12 months and clinical status change warrants investigation***</b> <input type="checkbox"/> 2D Echocardiogram <sup>1-3</sup> Reason: _____ <input type="checkbox"/> _____ Reason: _____	<p style="writing-mode: vertical-rl; transform: rotate(180deg);">Reference Document Only © 2015 Think Research Corporation. All rights reserved. Unauthorized use, reproduction or disclosure is prohibited.</p>
<b>IV Therapy</b> <input type="checkbox"/> Saline lock; flush as per policy/procedure <input type="checkbox"/> _____ at _____ mL/h	
<b>Heart Failure Medications</b> <b>Diuretics</b> <b>***IV diuretics are recommended as first-line therapy for patients with pulmonary or peripheral congestion***</b> <b>***If symptomatic hypotension arises, consider holding diuretics and reassessing for volume overload***</b> <input type="checkbox"/> furosemide _____ mg IV for 1 dose STAT (max 200 mg/dose) <input type="checkbox"/> furosemide _____ mg IV q _____ h <input type="checkbox"/> furosemide _____ mg/h IV continuous infusion (5 – 20 mg/h) <input type="checkbox"/> furosemide _____ mg PO q _____ h <input type="checkbox"/> bumetanide _____ mg PO q _____ h (0.5 mg; max 10 mg in 24 hours) <input type="checkbox"/> metolazone _____ mg PO q24h, administer 30 minutes prior to loop diuretic (2.5 mg; max 20 mg in 24 hours) <sup>2</sup> <input type="checkbox"/> _____	
<b>Angiotensin-Converting Enzyme-Inhibitors (ACEI)</b> <input type="checkbox"/> perindopril _____ mg PO q24h (initiation dose 2 – 4 mg; target regimen 4 – 8 mg q24h) <sup>1</sup> <input type="checkbox"/> ramipril _____ mg PO q12h (initiation dose 1.25 – 2.5 mg; target regimen 5 mg q12h) <sup>1</sup> <input type="checkbox"/> _____	
<b>Angiotensin Receptor Blockers (ARB) For Patient Intolerant to ACEI</b> <input type="checkbox"/> candesartan _____ mg PO q24h (initiation dose 4 – 8 mg; target regimen 32 mg q24h) <sup>1-3</sup> <input type="checkbox"/> valsartan _____ mg PO q12h (initiation dose 40 mg; target regimen 160 mg q12h) <sup>1-3</sup> <input type="checkbox"/> _____	
<b>Angiotensin Receptor Neprilysin Inhibitors (ARNI)</b> <b>***Patients who remain symptomatic despite triple therapy, consider change ACEI/ARB to an ARNI***<sup>1,2</sup></b> <b>***Concomitant use with an ACEI or ARB is contraindicated;</b> <b>if an ACEI was administered, wait 36 hours before administering ARNI***<sup>1,2</sup></b> <input type="checkbox"/> sacubitril 24 mg/valsartan 26 mg, 1 tab PO q12h (target regimen sacubitril 97 mg/valsartan 103 mg, 1 tab PO q12h) <input type="checkbox"/> sacubitril 49 mg/valsartan 51 mg, 1 tab PO q12h (target regimen sacubitril 97 mg/valsartan 103 mg, 1 tab PO q12h) <input type="checkbox"/> sacubitril 97 mg/valsartan 103 mg, 1 tab PO q12h	

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

CHFS Heart Failure Order Set	ACTION
<b>Heart Failure Medications Continued...</b> <b>Beta-Blockers</b> <input type="checkbox"/> bisoprolol _____ mg PO q24h (initiation dose 1.25 mg; target regimen 10 mg q24h) <sup>1</sup> <input type="checkbox"/> carvedilol _____ mg PO q12h (initiation dose 3.125 mg; target regimen 25 mg q12h [if weight greater than 50 kg, target regimen 50 mg q12h]) <sup>1</sup> <input type="checkbox"/> _____	<p style="writing-mode: vertical-rl; transform: rotate(180deg);">Reference Document Only © 2015 Think Research Corporation. All rights reserved. Unauthorized use, reproduction or disclosure is prohibited.</p>
<b>Mineralocorticoid Receptor Agonist (MRA)</b> <input type="checkbox"/> eplerenone _____ mg PO q24h (initiation dose 25 mg; target regimen 50 mg q24h) <sup>1-3</sup> <input type="checkbox"/> spironolactone _____ mg PO q24h (initiation dose 12.5 mg; target regimen 50 mg q24h) <sup>1</sup>	
<b>Vasodilators</b> <b>***The combination of isosorbide dinitrate and hydralazine are recommended in addition to standard treatment for black patients with HFrEF with advanced symptoms or patients unable to tolerate ACEI, ARB or ARNI therapy***<sup>1</sup></b> <input type="checkbox"/> hydralazine _____ mg PO q8h (initiation dose 25 mg; target regimen 75 mg q8h) And <input type="checkbox"/> isosorbide dinitrate _____ mg PO q8h (initiation dose 20 mg; target regimen 40 mg q8h) <sup>1</sup>	
<b>Sinoatrial Node Modulator</b> <b>***Ivabradine can be considered in patients with HFrEF who are in sinus rhythm with a resting heart rate of greater than or equal to 77 beats per minute and have had a previous HF hospitalization within the past year***<sup>1,2</sup></b> <input type="checkbox"/> ivabradine _____ mg PO q12h (initiation dose 2.5 – 5 mg <sup>1</sup> [if greater than 75 years old, initial dose 2.5 mg]; target regimen 7.5 mg q12h) <sup>1</sup>	
<b>Digoxin</b> <b>***Digoxin may be considered in patients in sinus rhythm who continue to be symptomatic with triple therapy***<sup>1,3</sup></b> <input type="checkbox"/> digoxin _____ mg PO q24h (0.125 – 0.25 mg)	
<b>Sodium-glucose Cotransporter 2 (SGLT2) Inhibitor</b> <b>***SGLT2 inhibitors should be started once medically stable or upon discharge given risks of euglycemic DKA***</b> <b>***Dapagliflozin may be considered in patients with mild to moderate HFrEF (LVEF less than/equal to 40%) and with or without concomitant type 2 diabetes<sup>11</sup> (do not use in patients with type 1 diabetes)***</b> <b>***Caution should be exercised when combining SGLT2 inhibitors, ARNI, and diuretics because of their concomitant effects to promote diuresis***<sup>11</sup></b> <b>Note: If serum creatinine is increasing, dapagliflozin should not be initiated or be reassessed if initiated.</b> <input type="checkbox"/> dapagliflozin 10 mg PO q24h <sup>12</sup>	
<b>Electrolyte Management</b> <input type="checkbox"/> Non-Critical Care Potassium Oral Replacement Clinical Protocol <input type="checkbox"/> _____	

Submitted by: \_\_\_\_\_ ☐ Read Back

Practitioner: \_\_\_\_\_  
 ID PRINTED NAME YYYY-MM-DD HH:MM SIGNATURE





*presents*

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# HEART FAILURE

AWARENESS WEEK MAY 4-10, 2020

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