

**ECHO IDAHO**

Managing Heart Failure  
in Primary Care

# Heart Failure Diagnosis and Workup

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Chris Longenecker, MD reported a financial relationship with Gilead Sciences as being on their Advisory Board on HIV. This relationship was deemed irrelevant in his role as a panelist in this series. None of the other planners or presenters for this educational activity have relevant financial relationship(s) to disclose with ineligible companies whose primary business is producing, marketing, selling, re-selling, or distributing healthcare products used by or on patients.

Spencer Carter, MD, speaker for this educational activity, reported financial relationships as being on the advisory board for Pfizer, BridgBio, and Alnylam. All of the relevant financial relationships listed for these individuals have been mitigated.



**University of Idaho**  
School of Health and Medical  
Professions

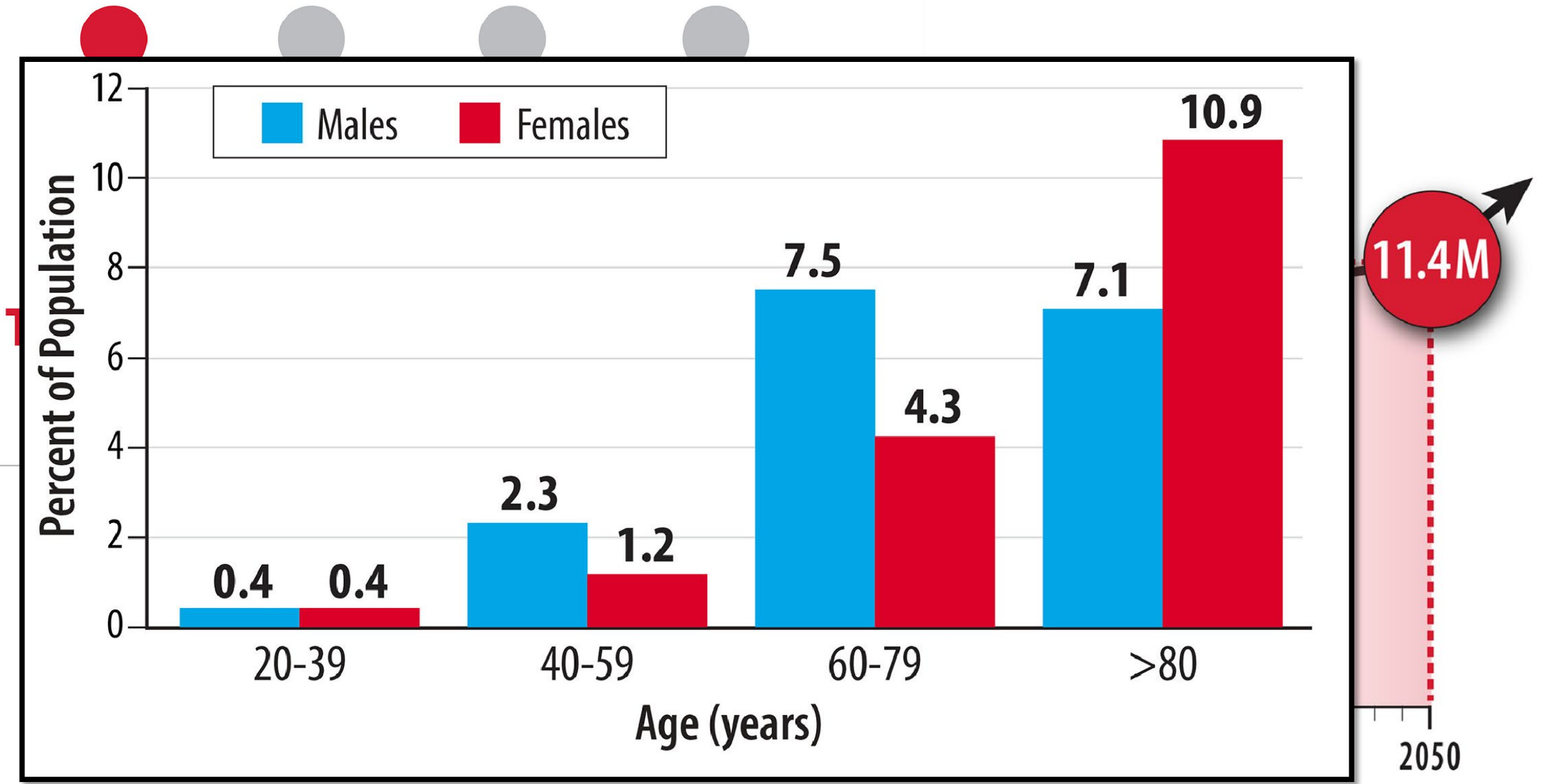


# Disclosures

- Medical advisory board- Pfizer, BridgeBio, Alnylam

# Learning Objectives

- Diagnosis of Heart Failure
- Heart Failure Schema
- Searching for Etiology



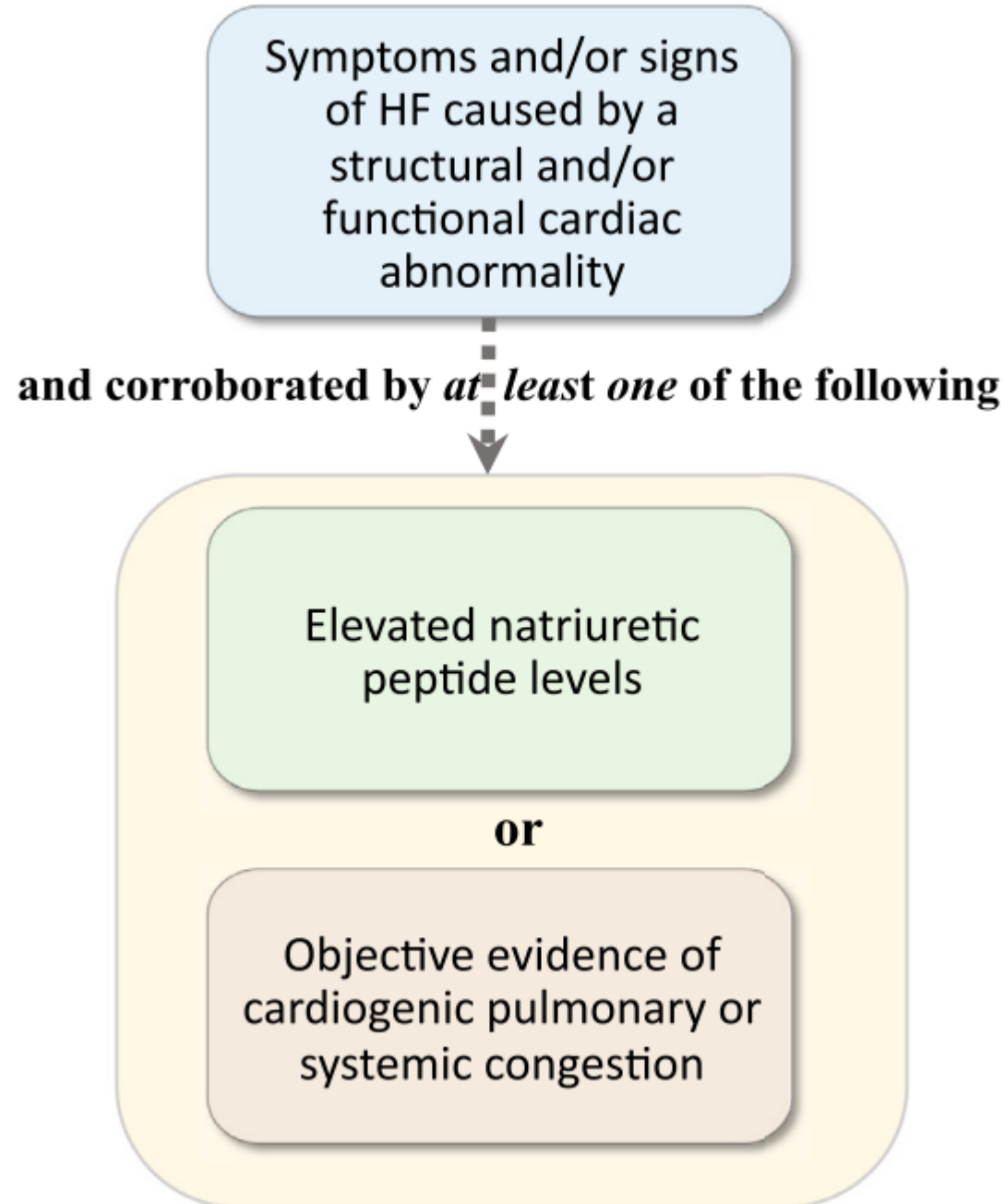
# Heart Failure Diagnosis

- Heart Failure is a clinical syndrome
- 2011 ACC definition

*“HF is a complex clinical syndrome that results from any structural or functional impairment of ventricular filling or ejection of blood. The cardinal manifestations of HF are dyspnea and fatigue, which may limit exercise tolerance, and fluid retention, which may lead to pulmonary and/or splanchnic congestion and/or peripheral edema. Some patients have exercise intolerance but little evidence of fluid retention, whereas others complain primarily of edema, dyspnea, or fatigue.”*

# Heart Failure

# Universal

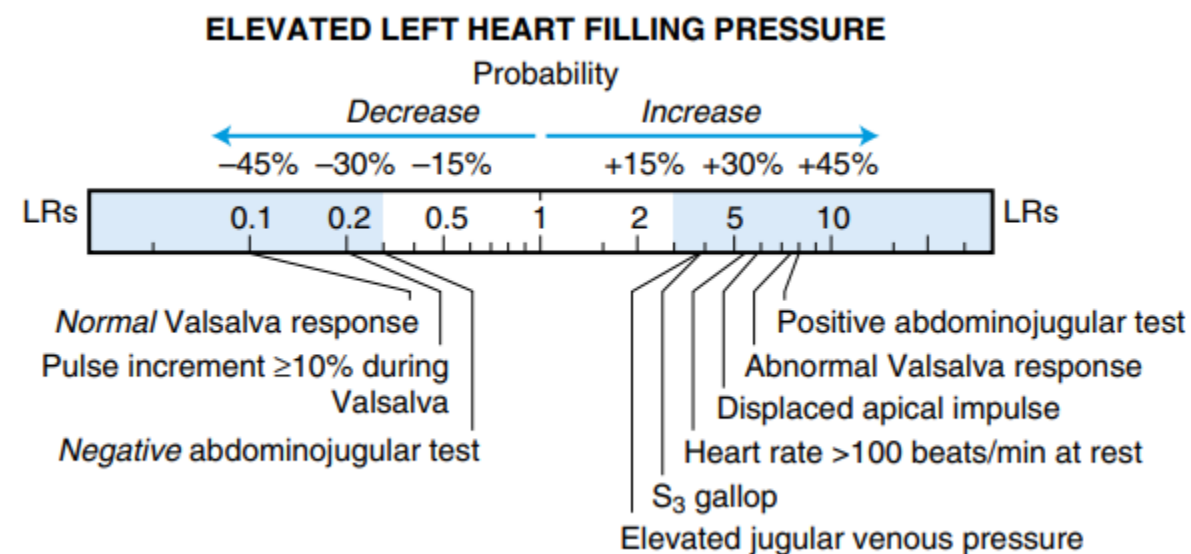


# Heart Failure Symptoms

- Symptoms- dyspnea on exertion, orthopnea, bendopnea, swelling in extremities
- Signs- third heart sound, elevated JVP, displaced PMI
- Less common signs/symptoms- significant edema, rales, pleural effusions, tachycardia, oliguria, narrow PP, cold extremities

**EBM BOX 48.1****Congestive Heart Failure—Elevated Left Heart Filling Pressures\***

Finding (Reference) <sup>†</sup>	Sensitivity (%)	Specificity (%)	Likelihood Ratio <sup>‡</sup> if Finding Is	
			Present	Absent
<b>Vital Signs</b>				
Heart rate >100 beats/min at rest <sup>15</sup>	6	99	5.5	NS
Abnormal Valsalva response <sup>16</sup>	95	88	7.6	0.1
Pulse increase of ≥10% during Valsalva strain <sup>17</sup>	11	54	0.2	1.7
<b>Lung Examination</b>				
Crackles <sup>12,15,18,19</sup>	12-23	88-96	NS	NS
<b>Heart Examination</b>				
Elevated jugular venous pressure <sup>12,15,19</sup>	10-58	96-97	3.9	NS
Positive abdominojugular test <sup>19-21</sup>	55-84	83-98	8.0	0.3
Supine apical impulse lateral to MCL <sup>18</sup>	42	93	5.8	NS
S <sub>3</sub> gallop <sup>12,15,18,22</sup>	12-37	85-96	3.9	0.8
S <sub>4</sub> gallop <sup>12,23</sup>	35-71	50-70	NS	NS
<b>Other Findings</b>				
Edema <sup>12,15</sup>	10	93-96	NS	NS





# How Helpful is the BNP?

**Table 8.** Natriuretic Peptide Levels Supporting Definition of HF

	Ambulatory	Hospitalized/ Decompensated
BNP, pg/mL	$\geq 35$	$\geq 100$
NT-proBNP, pg/mL	$\geq 125$	$\geq 300$

In the ED

- BNP >100 has a sensitivity of ~95%
- NT-proBNP <300 has a negative likelihood ratio of 0.09
- Specificity is lacking, even at levels in the 1000s
  - PE, PH, ARDS, renal dysfunction

# Description

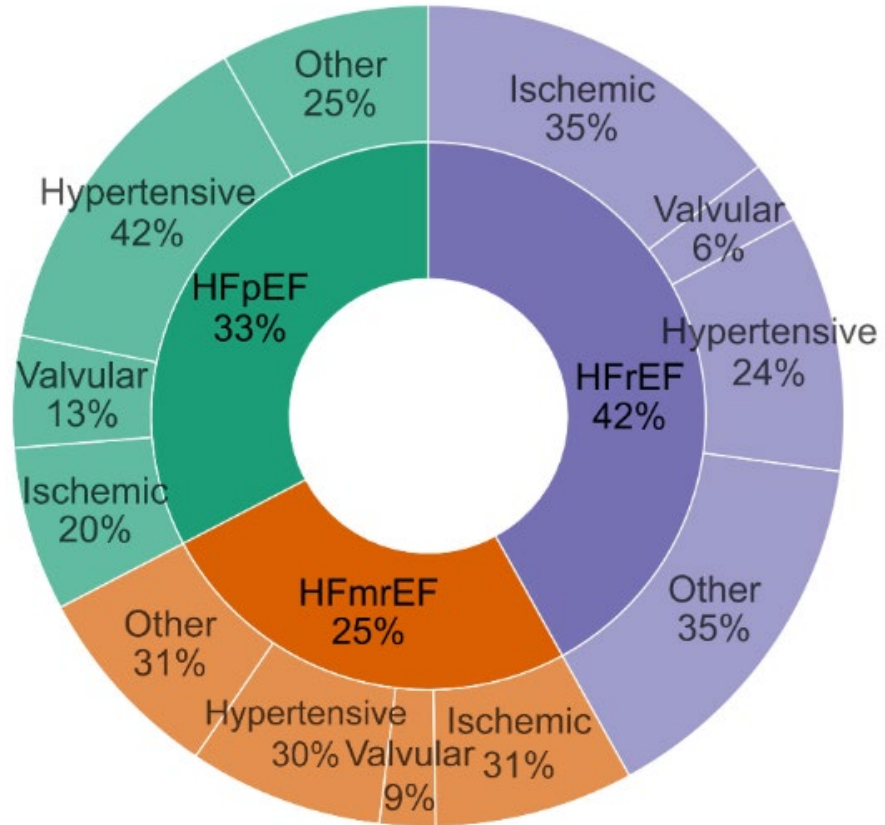
- Stage
  - 1- at risk, 2- pre-HF, 3-classic CHF, 4- advanced HF with sx at rest
- NYHA
  - 1- no sx, 2-mild sx, 3-moderate sx, 4- severe symptoms

# Key Points

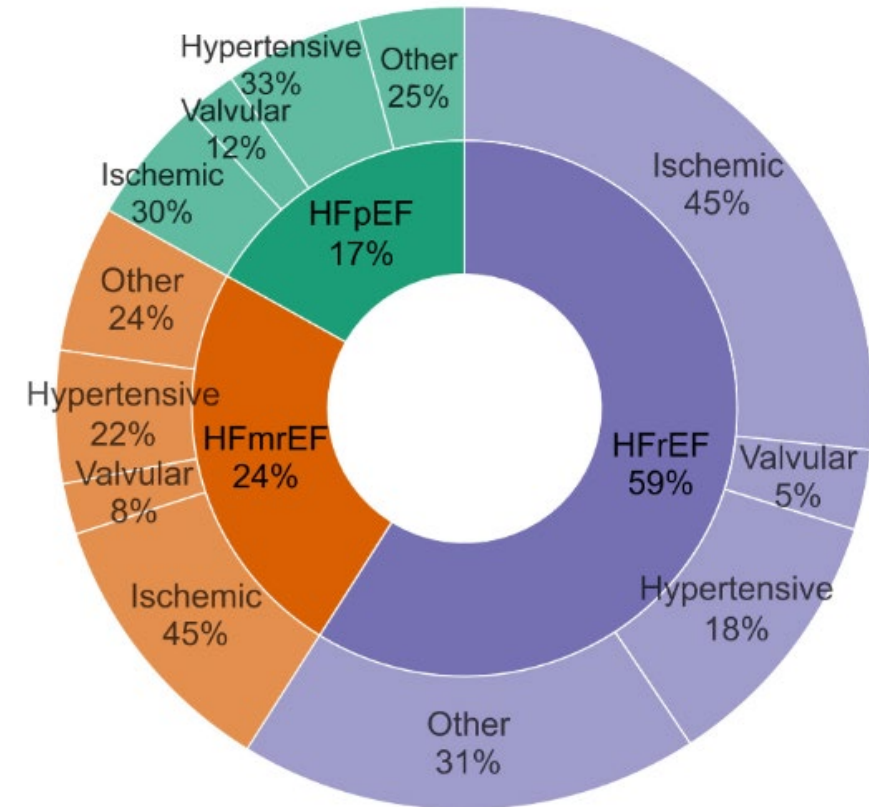
- Heart failure is a clinical diagnosis supported by physical exam and laboratory work

# Workup- Establish Etiology and Guide Therapy

**C** Female



**C** Male



# Workup- Goal is to Establish Etiology (and Treatment Plan)

- History- PMH, chest pain, family history, “red flags”, infections, palpitations, pregnancies
- Echocardiogram
  - Ejection fraction- establishes treatment pathway
  - Valvular, ischemic, infiltrative, intra-cardiac filling pressures, pericardial dz
- ECG
  - Potential insight into etiology and treatment (eg LBBB→chagas diagnosis, CRT treatment)
- Labs: CMP, CBC, iron studies, TSH, HIV, Hepatitis, Lipids
- Extras: Genetic testing? Extended rhythm monitoring? Autoimmune?

### **HF with reduced EF (HFrEF):**

- HF with LVEF  $\leq 40\%$

### **HF with mildly reduced EF (HFmrEF):**

- HF with LVEF 41–49%

### **HF with preserved EF (HFpEF):**

- HF with LVEF  $\geq 50\%$

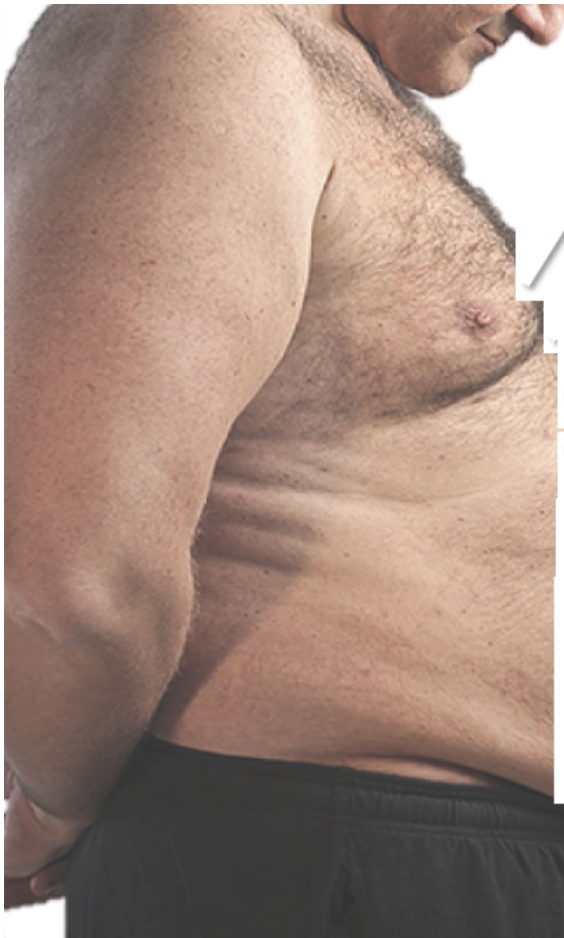
### **HF with improved EF (HFimpEF):**

- HF with a baseline LVEF  $\leq 40\%$ , a  $\geq 10$  point increase from baseline LVEF, and a second measurement of LVEF  $> 40\%$

# Ischemic Evaluation- What are the Basics?

- HFrEF/HFmrEF- virtually all patients
- Ischemic evaluation- choice is dependent on pre-test probability and what is available
  - High risk features → left heart cath
    - Typical anginal symptoms, heavy smoking/DM2, older age
  - Low to moderate risk (young, no risk factors) → non-invasive
    - Stress cMRI can help establish non-ischemic etiology. CTA coronary, PET generally good sensitivity/specificity
    - SPECT, dobutamine stress echo, stress ecg not adequate
- HFpEF- testing if moderate to high test probability (common given co-morbidities)

# HFpEF Pathophysiology



## Lung

Chest wall restriction, reduced vital capacity,  
impaired ventilation and diffusion  
Obstructive sleep apnea  
Pulmonary hypertension



## Heart

Direct and indirect myocardial lipotoxicity  
Worsened cardiac mechanics  
Diastolic dysfunction; increased filling pressures/  
volume overload, increased afterload



## Liver

Non-alcoholic fatty liver disease  
Promotes generalized  
inflammatory state



## Visceral adiposity

Inflammatory cytokines  
Adverse neurohormones  
Increased BNP clearance



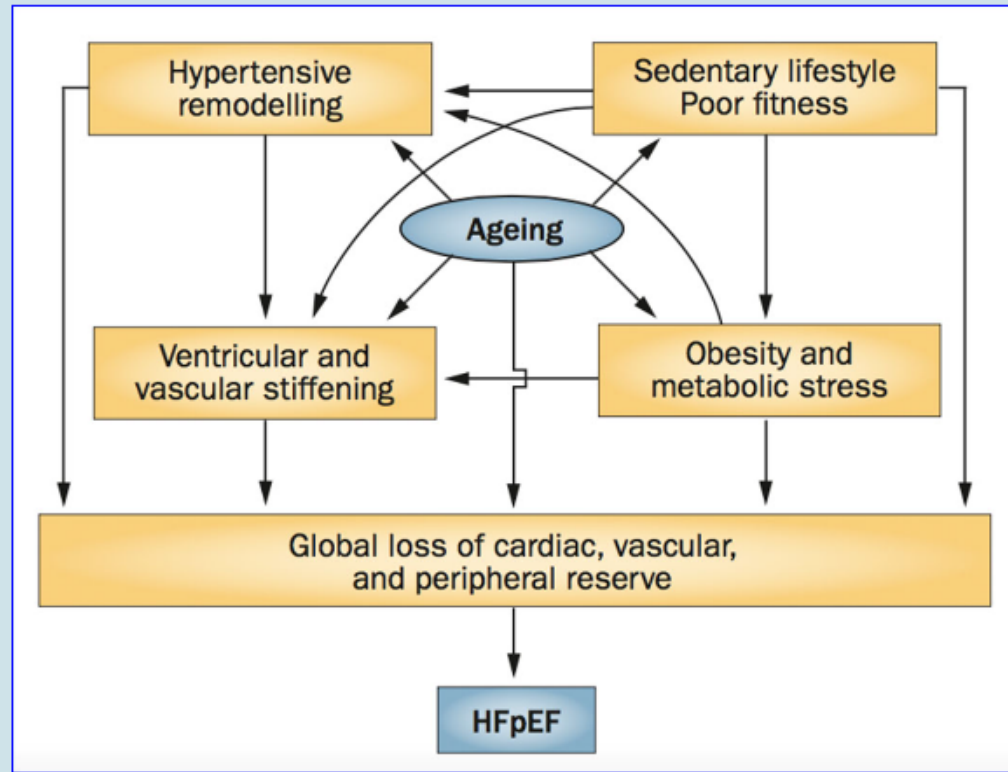
## Kidney

Direct toxic effects of perinephric fat  
Glomerulomegaly with  
glomerular dysfunction



## Skeletal muscle

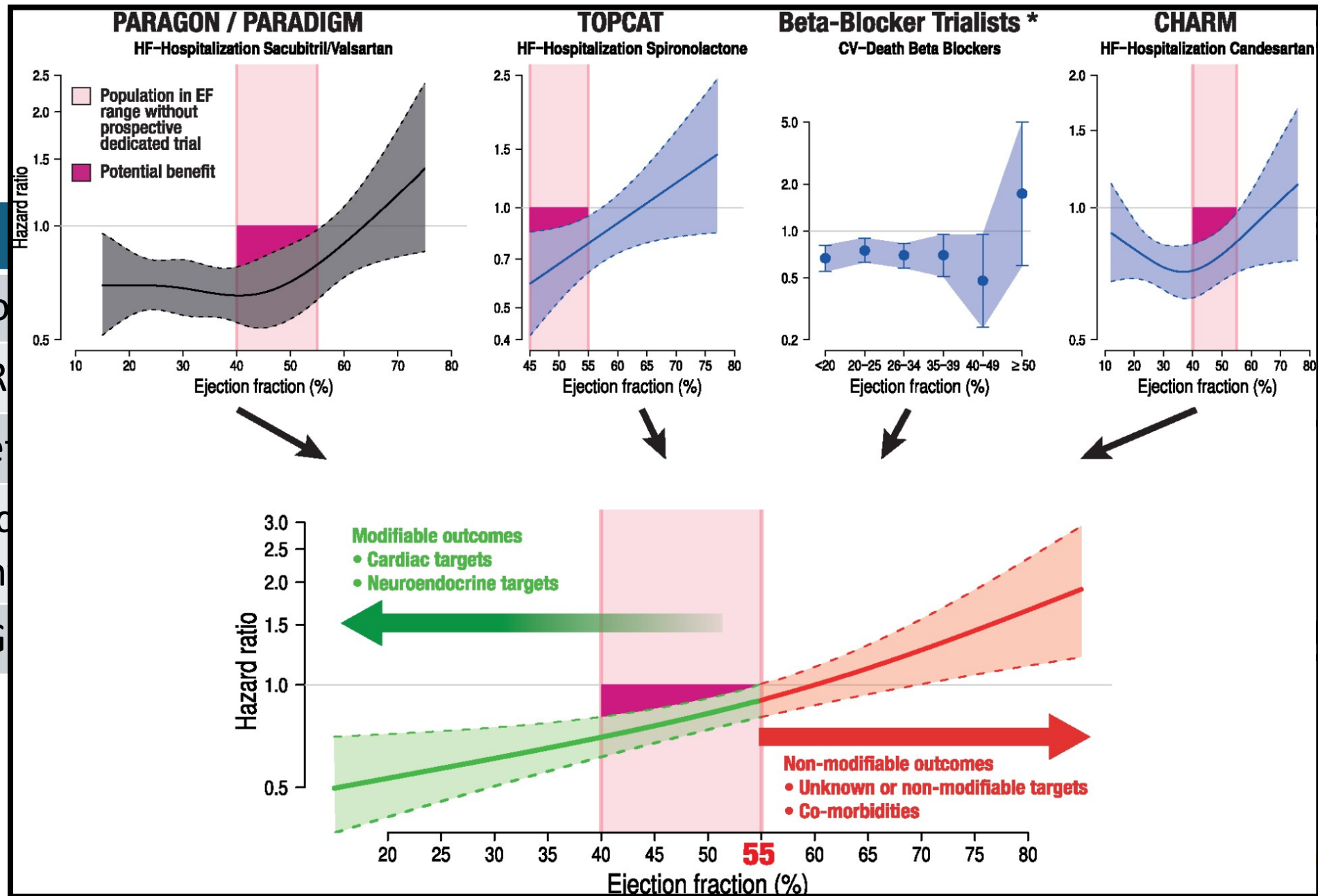
Increased adipose infiltration  
Impaired perfusion  
Decreased diffusive O<sub>2</sub> transport  
Mitochondrial dysfunction



Kitzman D, Shah SJ. *JACC* 2016; Borlaug B. *Nat Rev Cardiol* 2014



No  
AR  
Be  
Alc  
An  
SG



Relative risk  
on 72.9%

Relative risk  
on 25.5%

9

# Description

- Stage
  - A- at risk, B- pre-HF, C-classic CHF, D- advanced HF with sx at rest
- NYHA
  - 1- no sx, 2-mild sx, 3-moderate sx, 4- severe symptoms
- Ejection Fraction
  - HFrEF, HFmrEF, HFpEF, HFimpEF
- Etiology

“this is a 44yo male from Idaho with HFrEF (EF 25% 2/2025 cMRI), NICM (noCAD LHC 2022) 2/2 myocarditis and underlying TTN mutation. Stage C, NYHA 2, compensated on exam”

# When to Refer for Advanced Care Quickly

<b>I</b>	Need for <b>inotropes</b>
<b>N</b>	<b>New</b> York Heart Association Class IV
<b>E</b>	Worsening <b>end-organ</b> dysfunction
<b>E</b>	<b>Ejection fraction</b> <20%
<b>D</b>	<b>Defibrillator</b> shocks for ventricular arrhythmias
<b>H</b>	Recurrent <b>HF</b> hospitalizations
<b>E</b>	<b>Escalating</b> diuretic dose
<b>L</b>	<b>Low blood pressure</b>
<b>P</b>	<b>Progressive</b> intolerance of GDMT

# Questions?