

**NYHA CLASSIFICATION OF HF:**

Class I	Sx with more than ordinary activity
Class II	Sx with ordinary activity
Class III	Sx with minimal activity <ul style="list-style-type: none"> <li>• Class IIa: no dyspnea at rest</li> <li>• Class IIb: recent dyspnea at rest</li> </ul>
Class IV	Sx at rest

**GOALS OF THERAPY:**

1. PROLONG SURVIVAL \*\*\*\*\*
2. Reduce morbidity: exercise tolerance, hospitalization, exacerbations, QOL

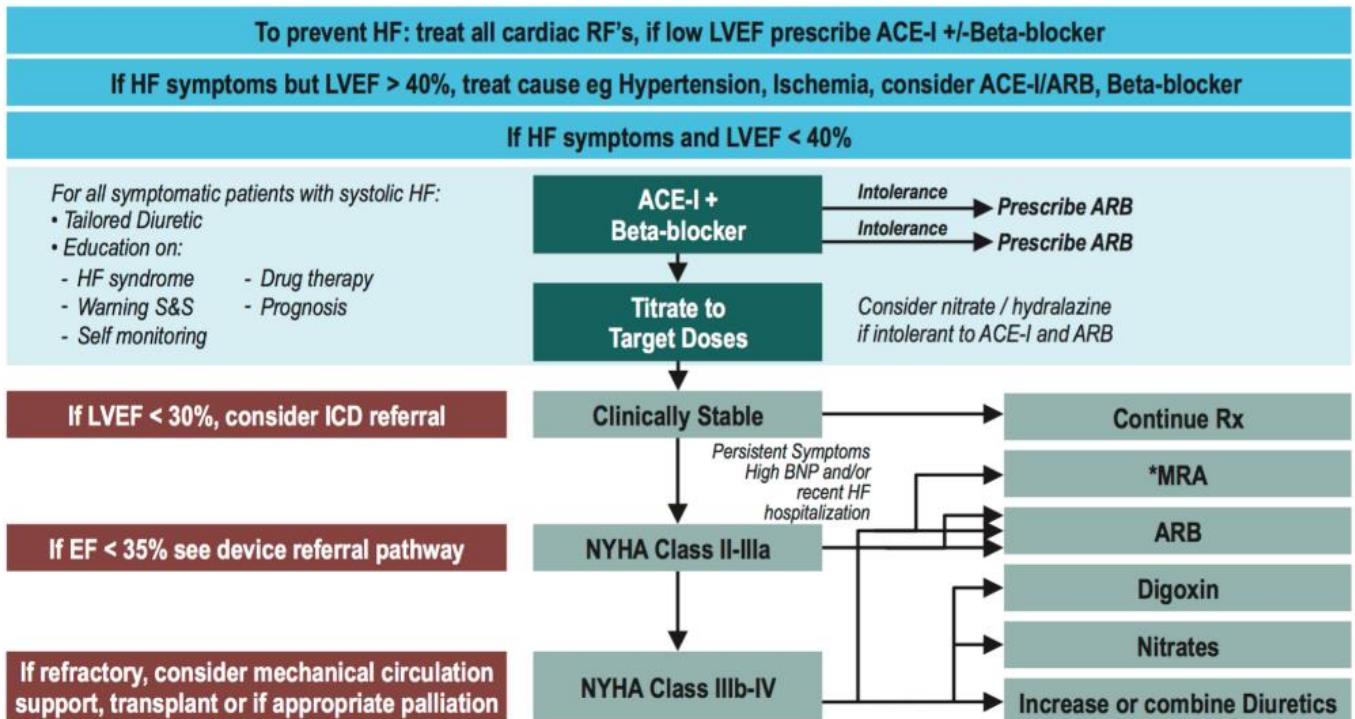
**HF PRECIPITANTS:**

- Anemia
- Ischemia
- Arrhythmia (V or A)
- Infection
- Medication non-adherence
- Drugs
  - NSAIDs
  - Corticosteroids
  - Verapamil/ diltiazem
  - VW class I antiarrhythmics
  - B-blockers
  - Glitazones / Gliptins (diabetic drugs)
    - Generally avoid if known LV dysfunction, if not then counsel re: edema
  - Anthracyclines

**ECHOCARDIOGRAM;**

	Systolic HF HFrEF	Diastolic HF/HFpEF	Valve dysfunction
EF	< 35 – 40%	≥ 40%	Sx like HF but is problem with valves <ul style="list-style-type: none"> <li>• Valves won't open = aortic, mitral or tricuspid stenosis</li> <li>• Valves won't close = prolapse</li> </ul>
Pathophys	Inability to eject blood from LV (contraction problem)	Inability to fill LV (relaxation problem)	
CXR	Enlarged heart (cardiothoracic ratio > 0.55) because LV is weak (dilated, thin walls)	LV is stiff (thick, hypertrophied) but normal heart size	
Population	Mostly male, CAD	Mostly female, HTN	
Drug efficacy (mortality)	ACEI, B-blocker, MRA, hydralazine + ISDN, ARB	N/A	
Drug efficacy (mortality)	Digoxin, diuretics, others above	Digoxin, candesartan, verapamil (?), diltiazem (?), beta blockers (?), diuretics  NOT ACEI or MRA	

**HF TREATMENT:**



ACEI													
Drugs	Drug	Start dose	Target dose										
	Captopril	6.25 – 12.5 mg tid	25 mg to 50 mg tid										
	Enalapril	1.25 – 2.5 mg bid	10 mg bid										
	Ramipril (MOST COMMON)	1.25 – 2.5 mg bid	5 mg bid or 10 mg od										
	Lisonopril (RCT showed improved exercise tolerance, but not large enough to show mortality benefit)	2.5 mg – 5 mg od	20 mg – 35 mg od										
	Trandolapril (preferred in pts with MI)	1 mg od	4 mg od										
Benefits	<ul style="list-style-type: none"> <li>• Mortality, Class I – IV = 20% RRR, NNT 3y = 18</li> <li>• Morbidity (hospitalization) = 20% RRR, NNT 3y = 42</li> </ul>												
Dosing Strategy	Start low, titrate to target doses over several weeks												
Risks/monitoring	Hypotension, hyperkalemia, renal dysfunction, cough, angioedema												
Checklist	<ul style="list-style-type: none"> <li>• Allergy/intolerance (ACEI – cough?)</li> <li>• Hypovolemia = contraindication</li> <li>• Hypotension</li> <li>• Renal dysfunction <ul style="list-style-type: none"> <li>◦ Bilateral renal artery stenosis or RAS in pt with solitary kidney = contraindication</li> <li>◦ Unilateral renal artery stenosis is ok (other kidney can compensate)</li> </ul> </li> <li>• Aortic stenosis (pts very sensitive to vasodilating drug, monitor BP closely)</li> </ul>												
ACEI/ARB and renal function	<ul style="list-style-type: none"> <li>• Generally contraindicated when SCr &gt; 200 mcmol/L</li> </ul> <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th>ACEI/ARB can worsen GFR ....</th> <th>What to do ...</th> </tr> </thead> <tbody> <tr> <td>Concurrent NSAID</td> <td>Remove NSAID</td> </tr> <tr> <td>Volume depleted / heavily diuresed</td> <td>Decrease diuretic dose</td> </tr> <tr> <td>Hyponatremia (marker for volume depletion, neurohormonal activation, severe HF)</td> <td>Increase Na intake slightly</td> </tr> <tr> <td> <ul style="list-style-type: none"> <li>• Low GFR to start with</li> <li>• Renal artery stenosis, especially bilateral</li> </ul> </td> <td>Decrease ACEI dose</td> </tr> </tbody> </table> <ul style="list-style-type: none"> <li>• ACEI can improve GFR by reducing SVR</li> <li>• In 1/3 of people, ACEI initiation doesn't affect GFR at all</li> </ul>			ACEI/ARB can worsen GFR ....	What to do ...	Concurrent NSAID	Remove NSAID	Volume depleted / heavily diuresed	Decrease diuretic dose	Hyponatremia (marker for volume depletion, neurohormonal activation, severe HF)	Increase Na intake slightly	<ul style="list-style-type: none"> <li>• Low GFR to start with</li> <li>• Renal artery stenosis, especially bilateral</li> </ul>	Decrease ACEI dose
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B-blockers			
Drugs	Drug	Start Dose	Target Dose
	Carvedilol (alpha and beta blocker + antioxidant properties)	3.125 mg bid	25 mg bid
	Bisoprolol	1.25 mg od	10 mg od
	Metoprolol CR/XL	12.5 mg bid (or 12.5 – 25 mg od)	100 mg bid (or 200 mg od)
Benefits	<ul style="list-style-type: none"> <li>• Mortality, Class I – IV = RRR 30%, NNT 1 y = 26</li> <li>• Morbidity (hospitalization) = 30% RRR, NNT 1 y = 25</li> </ul>		
Dosing strategy	Start low, work toward target doses from trials over several weeks (q 1-2 weekly increase)		
Risks/monitoring	Abrupt withdrawal, worsening HF symptoms during first 1-12 weeks (also see checklist)		
Checklist	<ul style="list-style-type: none"> <li>• Allergy/ intolerance</li> <li>• Bradycardia</li> <li>• Hypotension (NORMOTENSIVE IS FINE, even mild hypotensive is okay)</li> <li>• Heart block &gt; 1° (i.e. 2<sup>nd</sup> or 3<sup>rd</sup> degree heart block) = contraindication</li> <li>• Asthma/severe COPD</li> <li>• Severe PVD</li> <li>• Hypoglycemic risk (be cautious, counsel)</li> </ul>		

<b>Diuretics</b>	
Benefits	<ul style="list-style-type: none"> <li>• Morbidity, if fluid overloaded – Class II – IV</li> </ul>
Dosing strategy	<ul style="list-style-type: none"> <li>• Furosemide 10 – 160 mg daily               <ul style="list-style-type: none"> <li>○ Self-management strategy                   <ul style="list-style-type: none"> <li>▪ If weight gain x lbs over x days, then take furosemide for a few days until weight is normal again</li> <li>▪ If weight loss x lbs over x days, hold furosemide</li> </ul> </li> </ul> </li> <li>• HCTZ may be added for surgery</li> <li>• Add metolazone if really resistant to furosemide (better at diuresis than HCTZ)</li> </ul>
Risks/monitoring	Hypovolemia, hypokalemia, hypomagnesemia, hyperglycemia, hyperuricemia (HCTZ), hypocalcemia (furosemide)

<b>MRA</b>	
Benefits	<ul style="list-style-type: none"> <li>• Mortality, Class I – IV</li> <li>• Morbidity</li> </ul>
Dosing strategy	Add 25 mg once daily to stable Class II/IV patient already on ACEI and B-blocker
Risks/monitoring	<b>HYPERKALEMIA</b> , breast tenderness/gynecomastia, hypotension
Pre-spirolactone checklist	<ul style="list-style-type: none"> <li>• Allergy/intolerance</li> <li>• Hypotension</li> <li>• NYHA Class III – IV heart failure (this is the population to use in)</li> <li>• Hyperkalemia (note pt, may be on K supplements when on furosemide)</li> </ul>
Eplerenone checklist	<ul style="list-style-type: none"> <li>• CCS recommends in NYHA II</li> <li>• Can probably use spironolactone instead (covered by PharmaCare, cheaper)</li> </ul>

<b>ARB</b>			
Drugs	Drug	Start does	Target dose
	Candesartan (gold standard)	4 mg od	32 mg od
	Valsartan	40 mg bid	160 mg bid
Benefits	<ul style="list-style-type: none"> <li>• Morbidity (vs. placebo, and when added to standard therapy) – Class I – IV</li> <li>• Mortality (candesartan, whether or not on ACEI)</li> </ul>		
Dosing strategy	<ul style="list-style-type: none"> <li>• Start low go slow when adding to ACEI therapy</li> <li>• When switching from ACEI to ARB, may switch directly to a comparable dose</li> </ul>		
Risks/monitoring	Renal dysfunction, hypotension, hyperkalemia		

<b>Hydralazine/nitrate (used concurrently)</b>			
Drugs (use hydralazine + isosorbide OR nitropatch)	Drug	Start does	Target dose
	Hydralazine	37.5 mg TID	75 mg TID
	Isosorbide dinitrate	20 mg TID	40 mg TID
	Nitropatch	0.2 – 0.4 mg/h x 12 h/day	0.6 – 0.8 mg/h x 12 h/day
Benefits	Only statistically significant benefit in African American patients (males)		

<b>Diuretics digoxin</b>	
Benefits	Morbidity, Class II – III
Dosing strategy	0.0625 – 0.375 mg QD depending on renal function, age, tolerability
Risks/monitoring	CNS ADRs (confusion, hallucinations), diarrhea, increased toxicity during hypokalemia, monitor RENAL FUNCTION

<b>Sacubitril / Valsartan (LCZ696, Entresto)</b>	
Benefits	Add on if still uncontrolled after standard therapy (Class II – IV)