

---

# **Pathophysiology: Heart Failure**

Mat Maurer, MD  
Associate Professor of Clinical Medicine

---

## **Objectives**

At the conclusion of this seminar, learners will be able to:

1. Define heart failure as a clinical syndrome
2. Define and employ the terms preload, afterload, contractility, remodeling, diastolic dysfunction, compliance, stiffness and capacitance.
3. Describe the classic pathophysiologic steps in the development of heart failure.
4. Delineate four basic mechanisms underlying the development of heart failure
5. Interpret pressure volume loops / Starling curves and identify contributing mechanisms for heart failure state.
6. Understand the common methods employed for classifying patients with heart failure.
7. Employ the classes and stages of heart failure in describing a clinical scenario

## Heart Failure

---

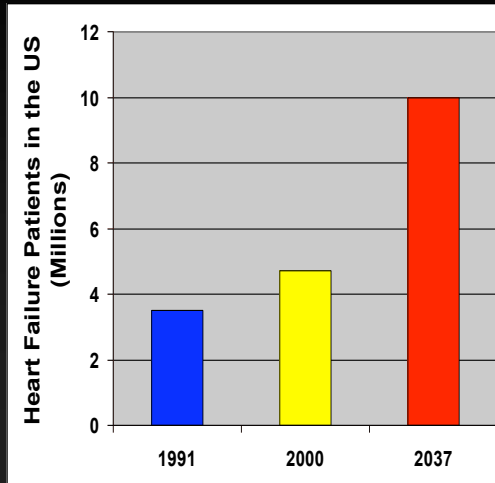
- Not a disease
- A syndrome
  - From "syn" meaning "together" and "dromos" meaning "a running".
  - A group of signs and symptoms that occur together and characterize a particular abnormality.
- Diverse etiologies
- Several mechanisms

## Heart Failure: Definitions

---

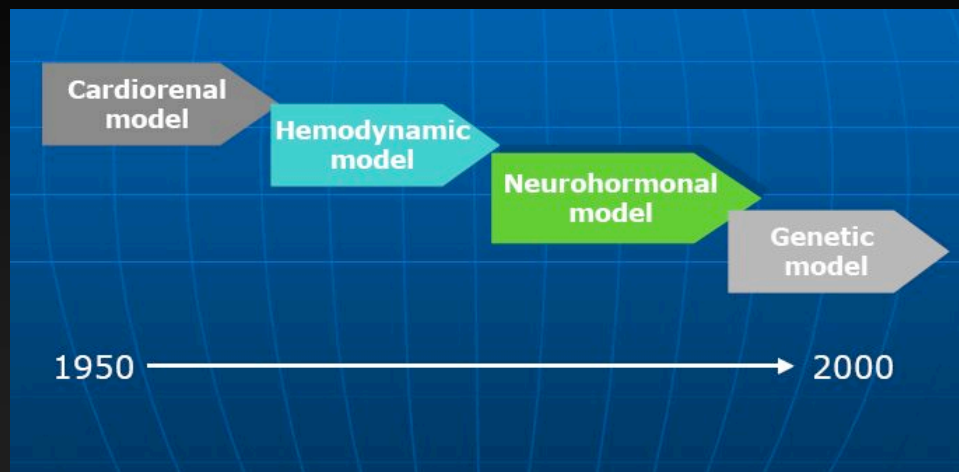
- An inability of the heart to pump blood at a sufficient rate to meet the metabolic demands of the body (e.g. oxygen and cell nutrients) at rest and during effort or to do so only if the cardiac filling pressures are abnormally high.
- A complex clinical syndrome characterized by abnormalities in cardiac function and neurohormonal regulation, which are accompanied by effort intolerance, fluid retention and a reduced longevity
- A complex clinical syndrome that can result from any structural or functional cardiac disorder that impairs the ability of the ventricle to fill with or eject blood.

# Epidemiology Heart Failure: The Problem

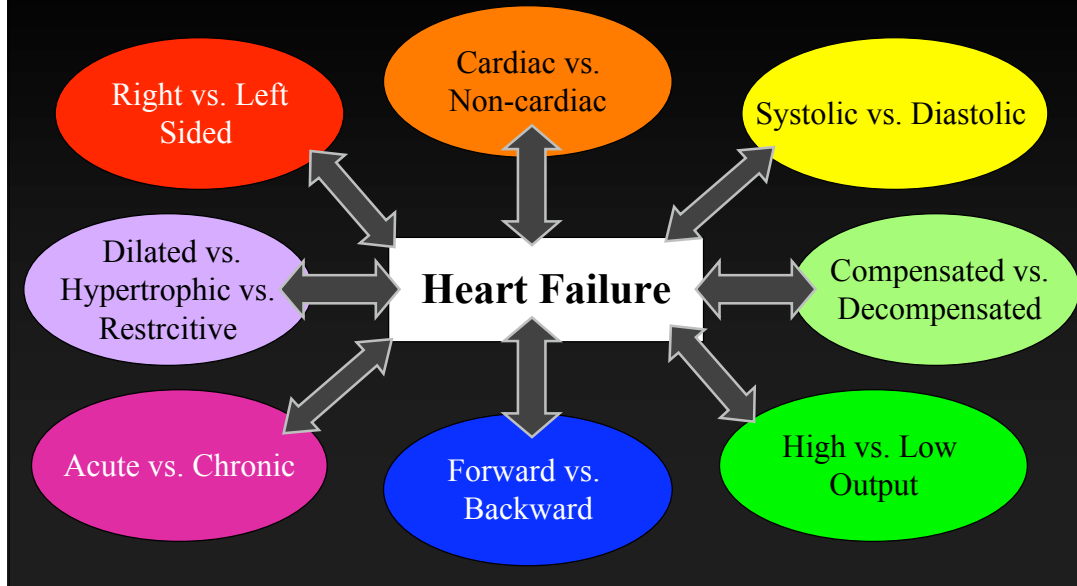


- 3.5 million in 1991, 4.7 million in 2000, estimated 10 million in 2037
- Incidence: 550,000 new cases/year
- Prevalence: 1% ages 50--59, >10% over age 80
- More deaths from HF than from all forms of cancer combined
- Most common cause for hospitalization in age >65

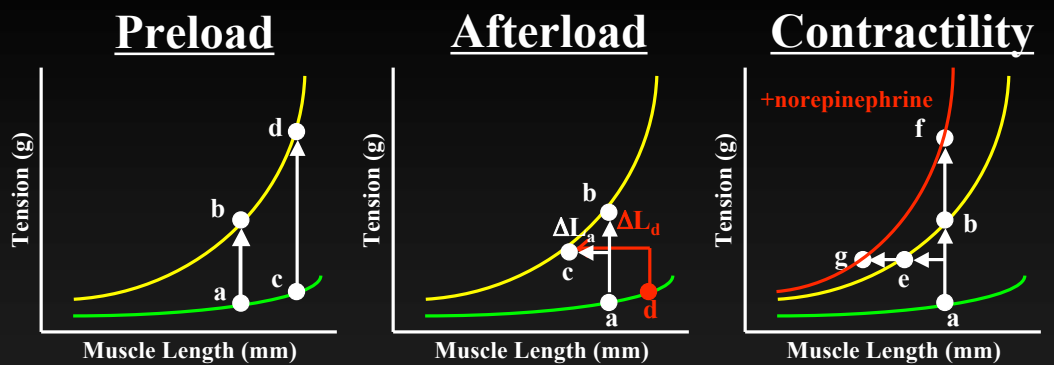
# Heart Failure Paradigms



# Heart Failure: Classifications



# Cardiac Muscle Function

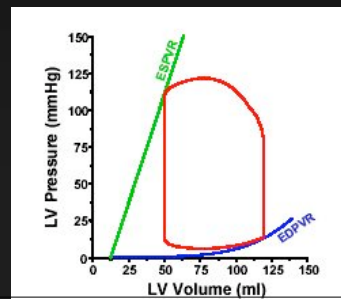
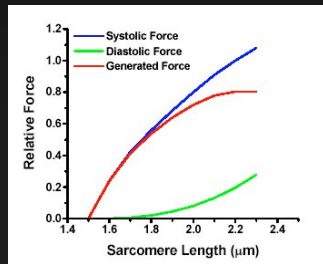
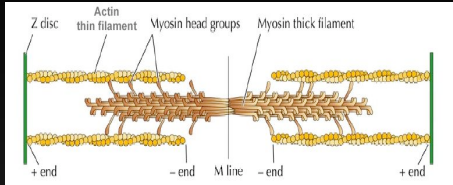


- The length of a cardiac muscle fiber prior to the onset of contraction.
- Frank Starling

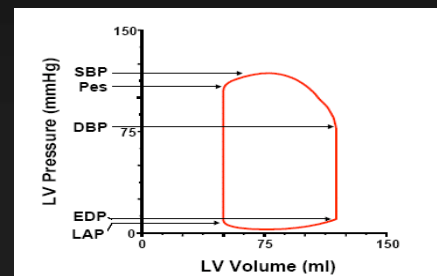
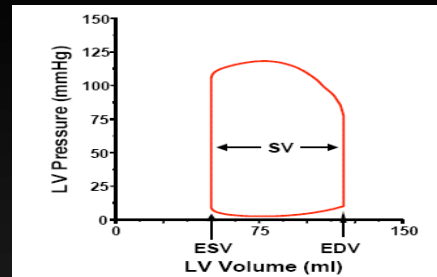
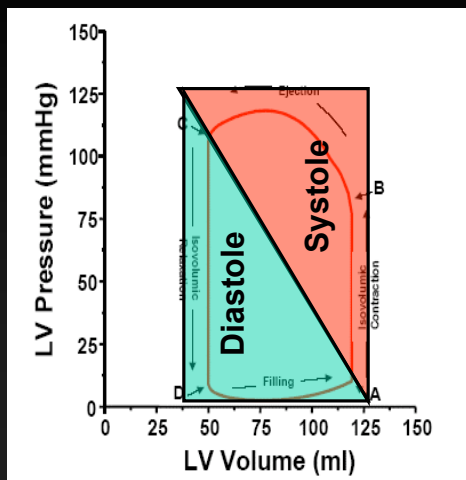
- The force against which a cardiac muscle fiber must shorten.
- Isotonic Contraction

- The force of contraction independent of preload and afterload.
- Inotropic State

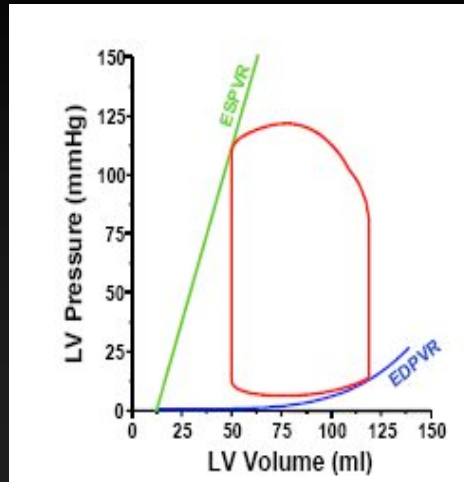
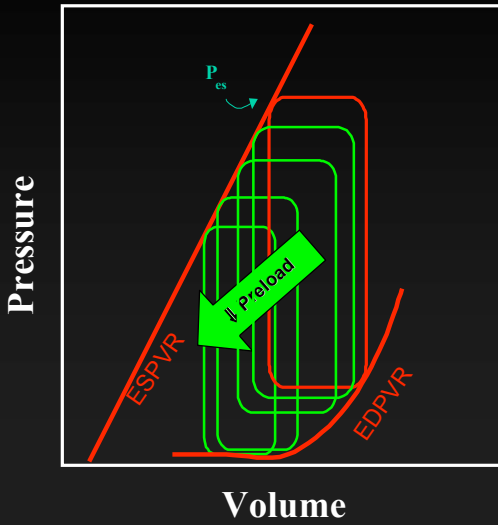
# From Muscle to Chamber



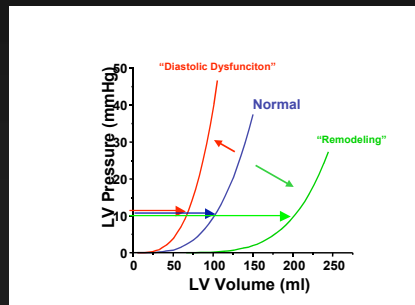
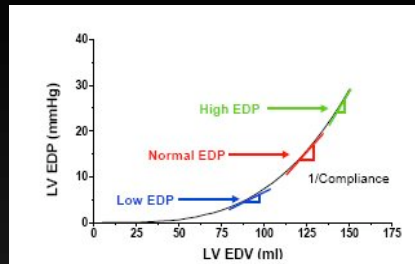
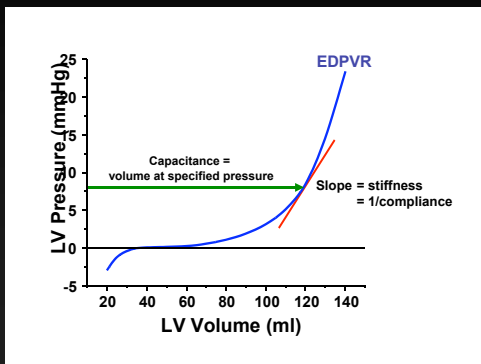
# The Pressure Volume Loop



# The Pressure Volume Loop



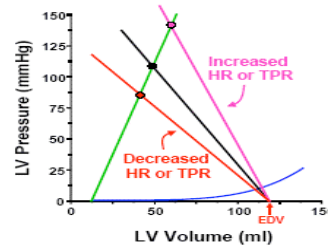
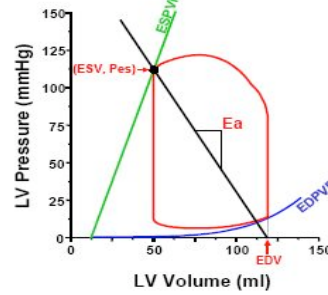
# Compliance/Stiffness vs Capacitance



# Afterload (Arterial Properties)

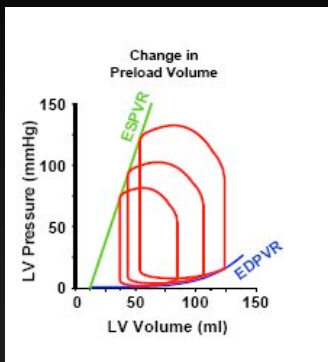
## Ea (Arterial Elastance)

- If
  - $TPR = [MAP - CVP] / CO$ , and
  - $CO = SV * HR$
- Substituting the second equation into the first we obtain:
  - $TPR = [MAP - CVP] / (SV * HR)$
- Making two simplifying assumptions.
  1. CVP is negligible compared to MAP.
  2. MAP is approximately equal to the end-systolic pressure in the ventricle ( $P_{es}$ ).
- Then,
  - $TPR = P_{es} / (SV * HR)$
- Which can be rearranged to:
  - $P_{es} / SV \cong TPR * HR$ .



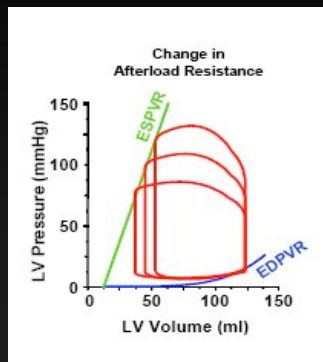
# Cardiac Chamber Function

## Preload



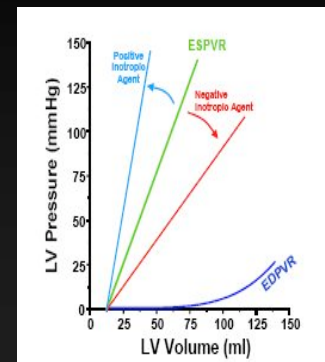
- EDV
- EDP
- Wall stress at end diastole

## Afterload



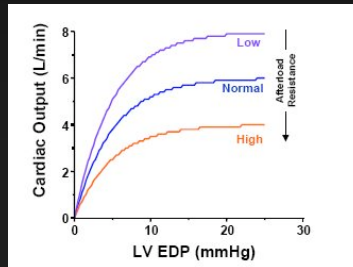
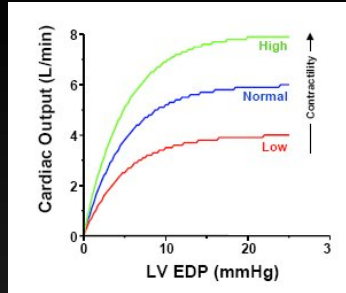
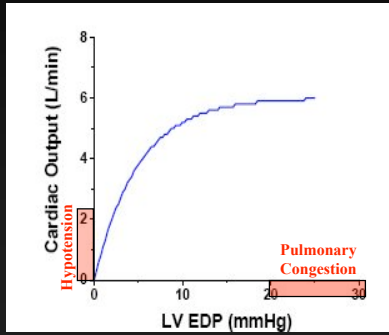
- Aortic Pressure
- Total peripheral resistance
- Arterial impedance
- Wall stress at end systole

## Contractility

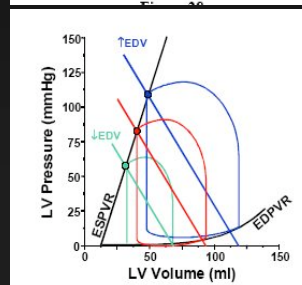
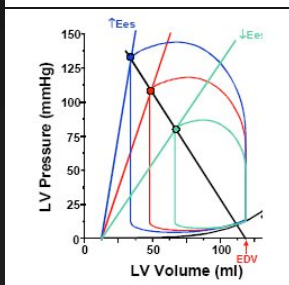
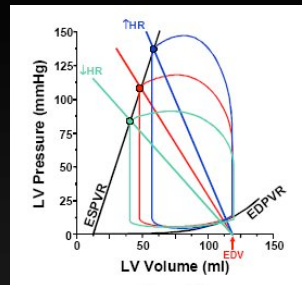
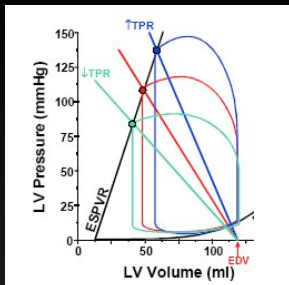


- Pressure generated at given volume.
- Inotropic State

# Frank Starling Curves

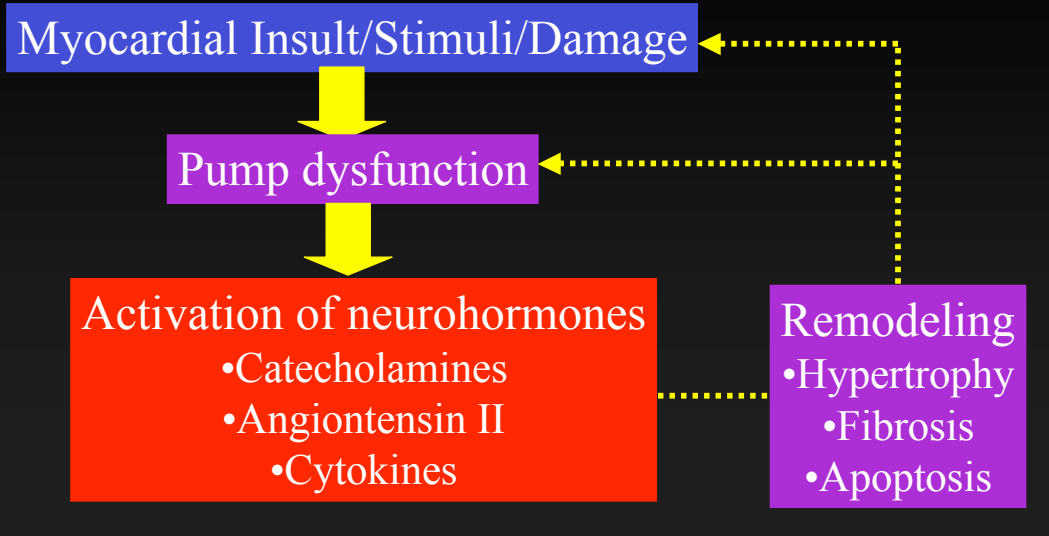


# Pathophysiology - PV Loop

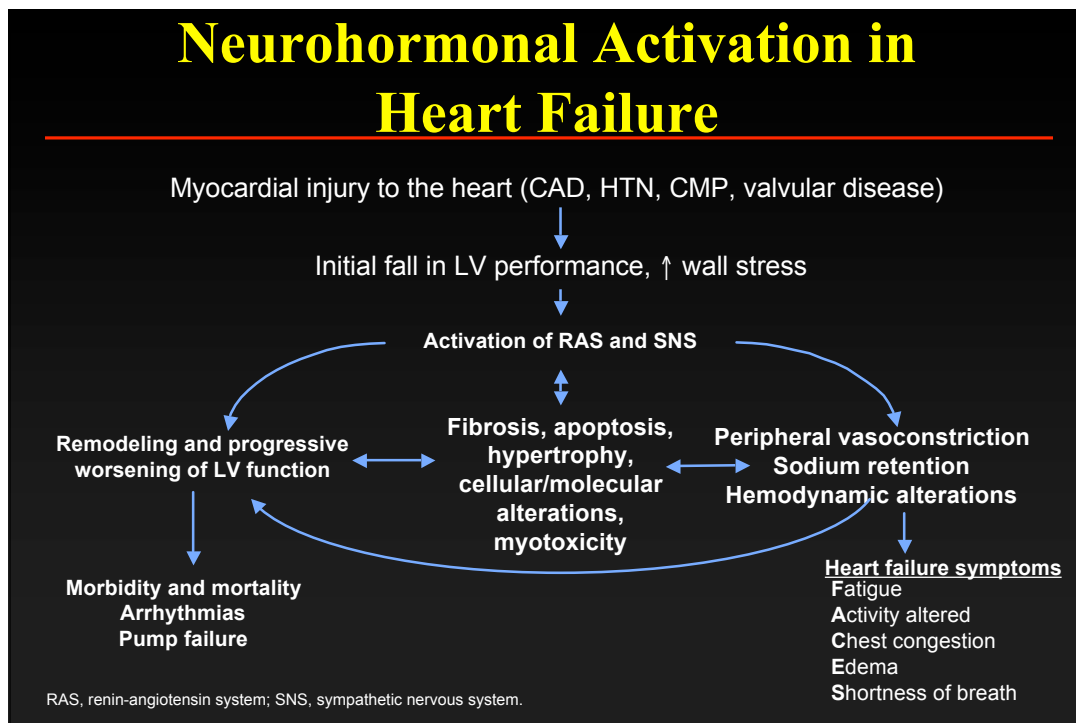




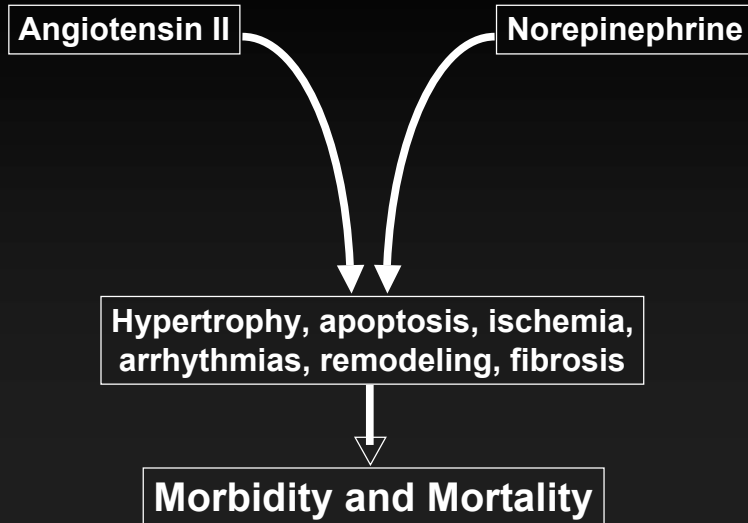
# Pathophysiology of Heart Failure



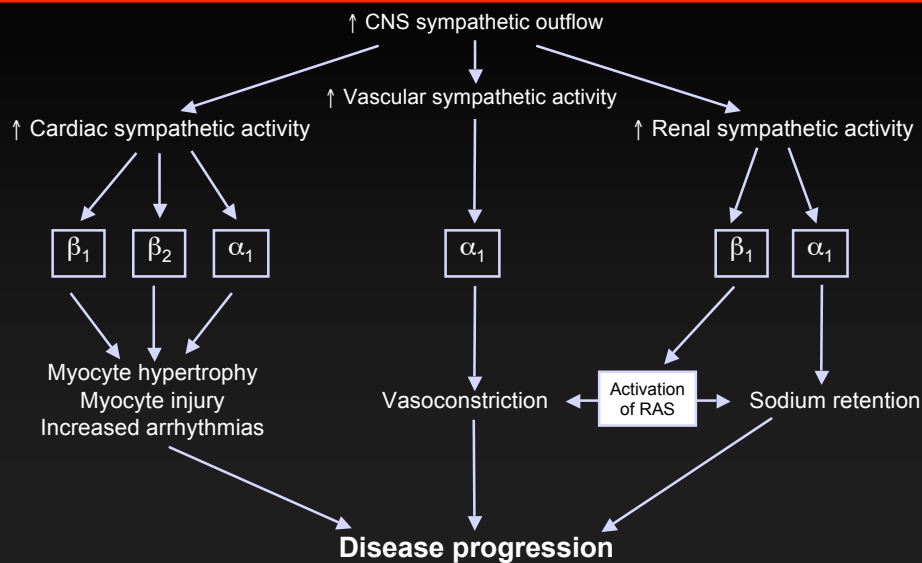
# Neurohormonal Activation in Heart Failure



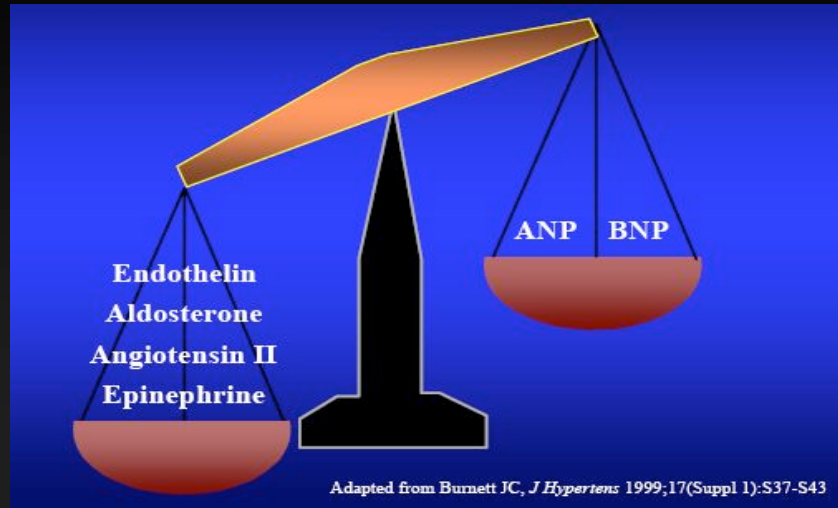
## Neurohormonal Activation in Heart Failure



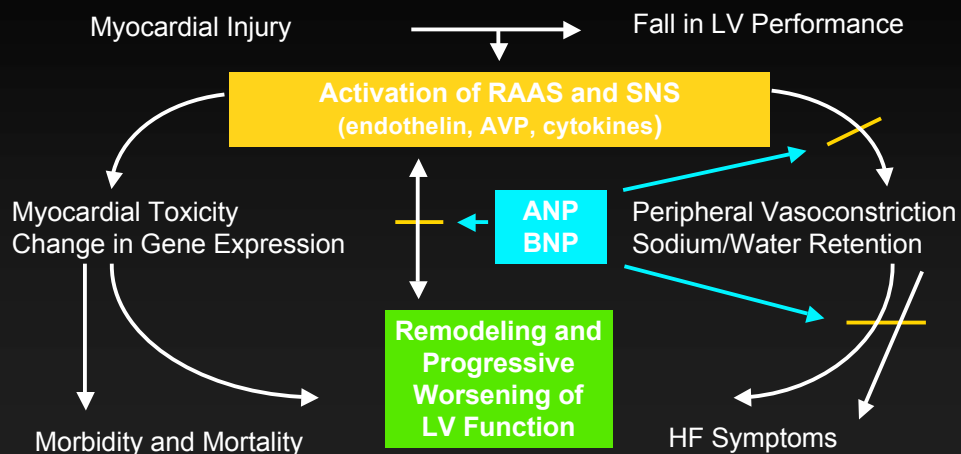
## Adrenergic Pathway in Heart Failure Progression



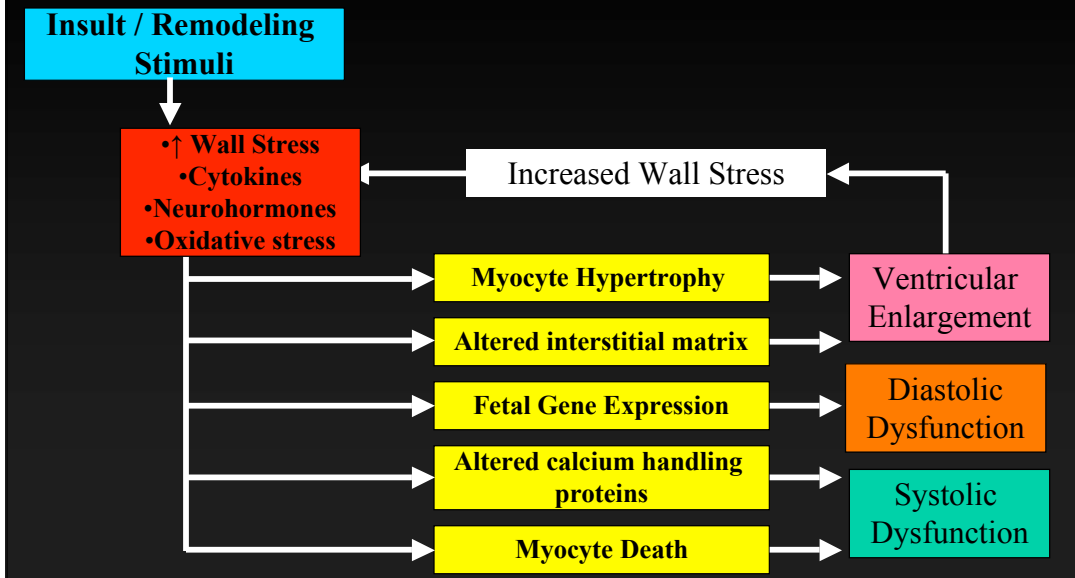
## Neurohormonal Balance in Heart Failure



## Neurohormones in Heart Failure



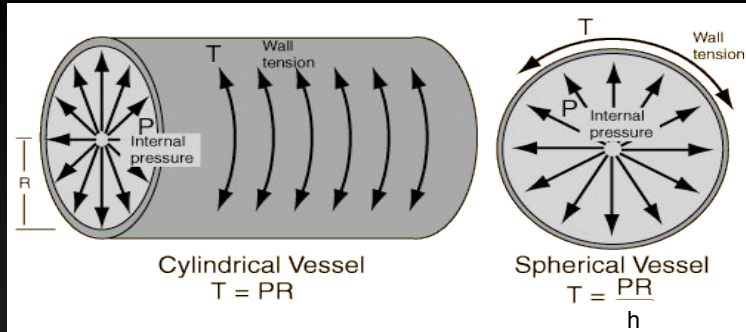
## Pathophysiology of myocardial remodeling:



## Acute and Chronic Responses – Benefits and Harm

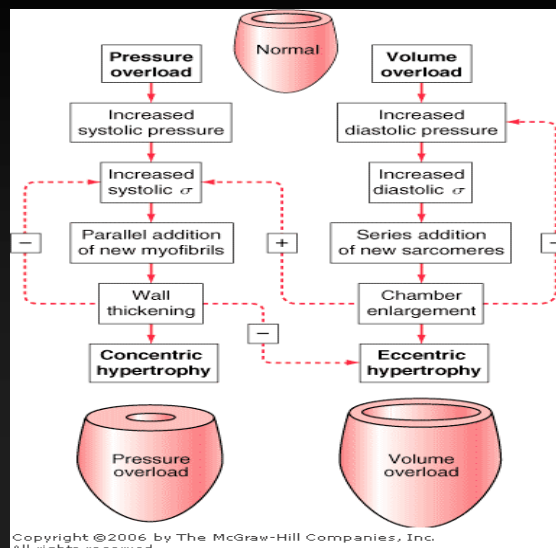
Response	Short-term Effects (mainly adaptive; hemorrhage, acute heart failure)	Long-term Effects (mainly deleterious; chronic heart failure)
Salt and water retention	Augments preload	Pulmonary congestion, anasarca
Vasoconstriction	Maintains pressure for perfusion of vital organs (brain, heart)	Exacerbates pump dysfunction, increases cardiac energy expenditure
Sympathetic stimulation	Increases heart rate and ejection	Increases energy expenditure
Cytokine activation	Vasodilatation	Skeletal muscle catabolism, deterioration of endothelial function, impaired contraction, LV remodeling.
Hypertrophy	Unloads individual muscle fibers	Deterioration and death of cardiac cells: cardiomyopathy of overload
Increased collagen	May reduce dilatation	Impairs relaxation

# Laplace's Law

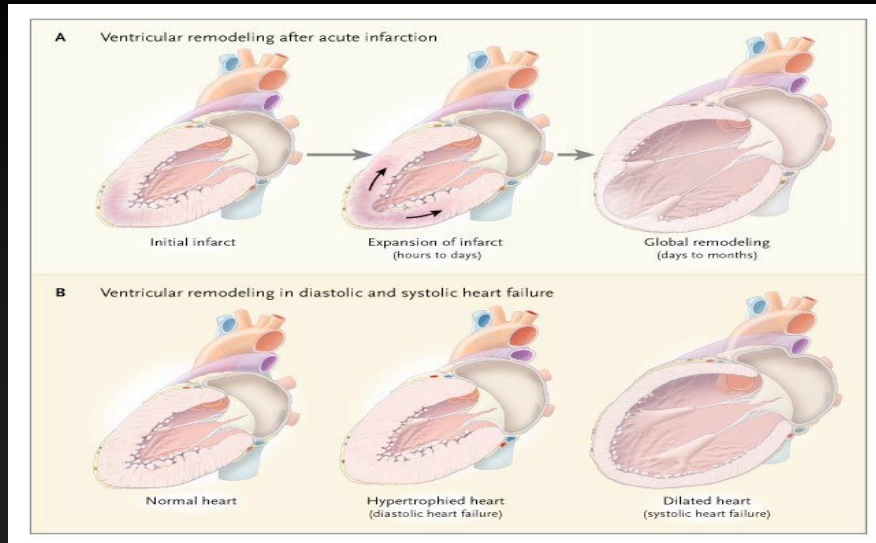


Where P = ventricular pressure, r = ventricular chamber radius and h = ventricular wall thickness

# Remodeling – Concentric vs. Eccentric



## Ventricular Remodeling

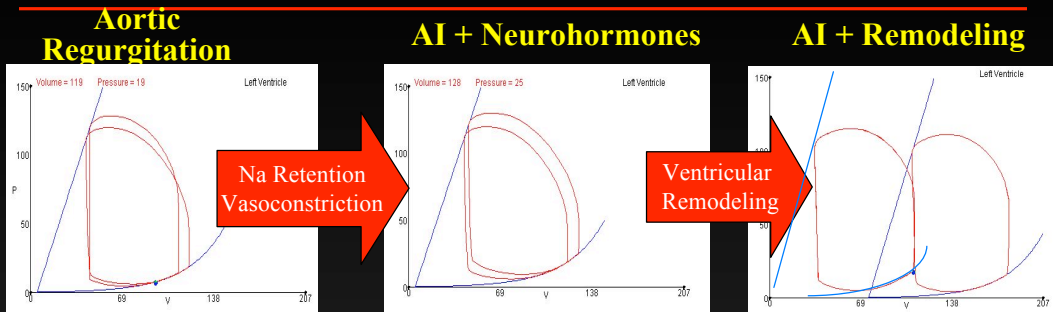


## Pathophysiology of Heart Failure

### Four Basic Mechanisms

1. Increased Blood Volume (Excessive Preload)
2. Increased Resistant to Blood Flow (Excessive Afterload)
3. Decreased contractility
4. Decreased Filling

# Increased Blood Volume

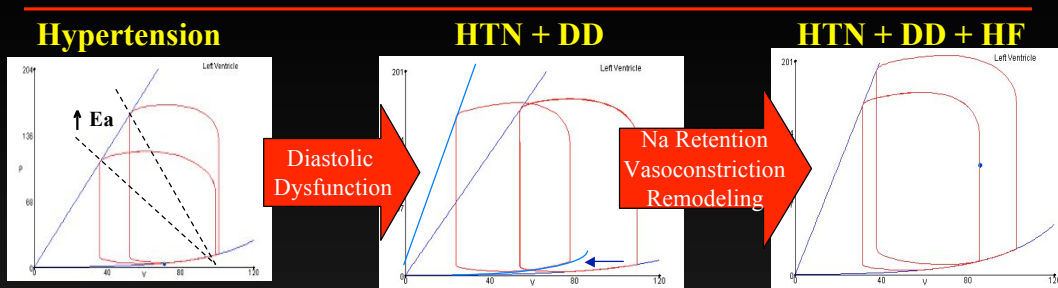


## Etiologies

- Mitral Regurgitation
- Aortic Regurgitation
- Volume Overload
- Left to Right Shunts
- Chronic Kidney Disease

Parameter	Normal	AI	AI+Neuro-Hormones	AI + Remodeling
BP (mm Hg)	140/75	128/50	130/50	104/45
SV (ml)	64	80	82	63
Cardiac Output (L/min)	3.8	3.0	4.3	2.6
PCWP (mm Hg)	10	15	25	20

# Increased Afterload

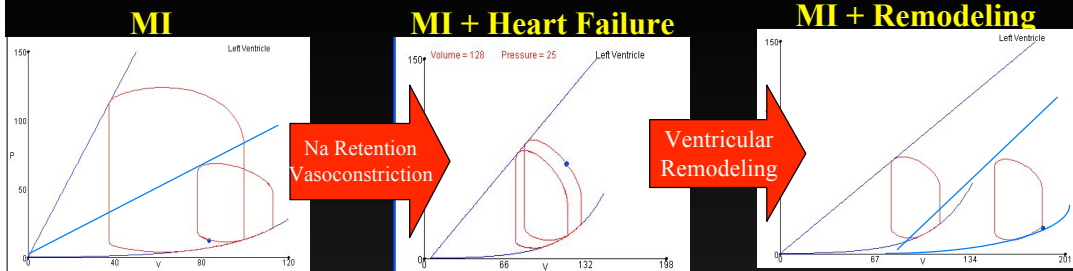


## Etiologies

- Aortic Stenosis
- Aortic Coarctation
- Hypertension

Parameter	Normal	HTN	HTN + DD	HTN + Heart failure
BP (mm Hg)	131/76	150/100	140/92	161/105
SV (ml)	57	52	49	57
Cardiac Output (L/min)	4.0	3.6	3.4	4.0
PCWP (mm Hg)	10	10	13	23

# Decreased Contractility

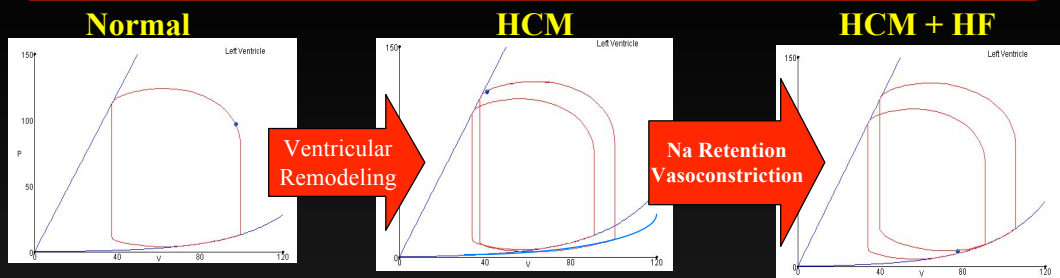


## Etiologies

- Ischemic Cardiomyopathy
  - Myocardial Infarction
  - Myocardial Ischemia
- Myocarditis
- Toxins
  - Anthracycline
  - Alcohol
  - Cocaine

Parameter	Normal	MI	MI + Neurohormones	MI + Remodeling
BP (mm Hg)	124/81	80/40	90/54	87/44
SV (ml)	60	42	46	46
Cardiac Output (L/min)	4.2	3.0	3.2	3.2
PCWP (mm Hg)	12	17	25	23

# Decreased Filling



## Etiologies

- Mitral Stenosis
- Constriction
- Restrictive Cardiomyopathy
- Cardiac Tamponade
- Hypertrophic Cardiomyopathy
- Infiltrative Cardiomyopathy

Parameter	Normal	HCM	HCM + HF
BP (mm Hg)	124/81	95/47	105/53
SV (ml)	63	50	55
Cardiac Output (L/min)	4.4	3.5	3.8
PCWP (mm Hg)	10	17	26

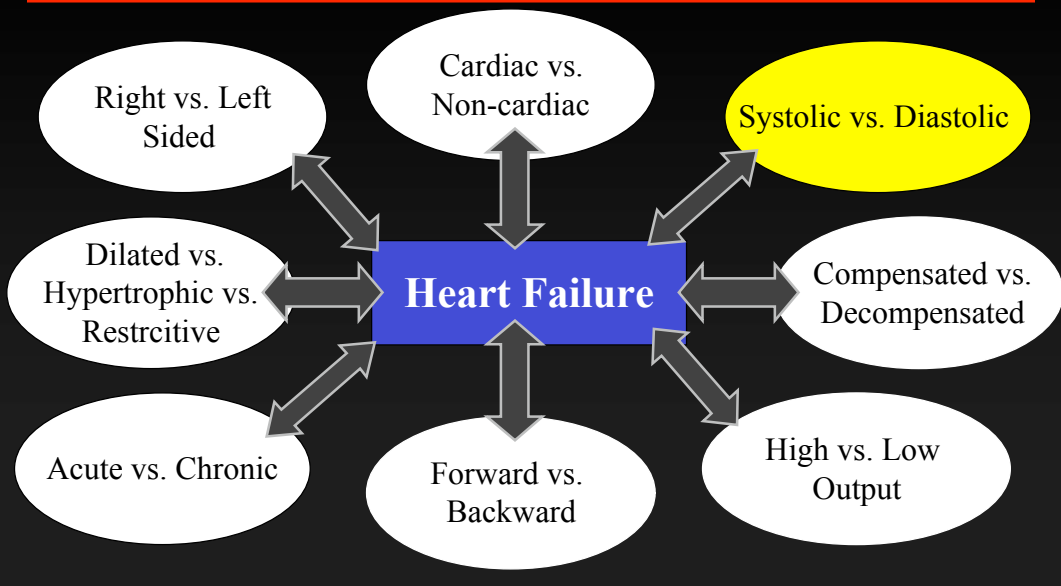


---


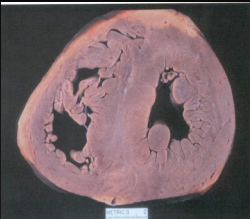
## Part II

---

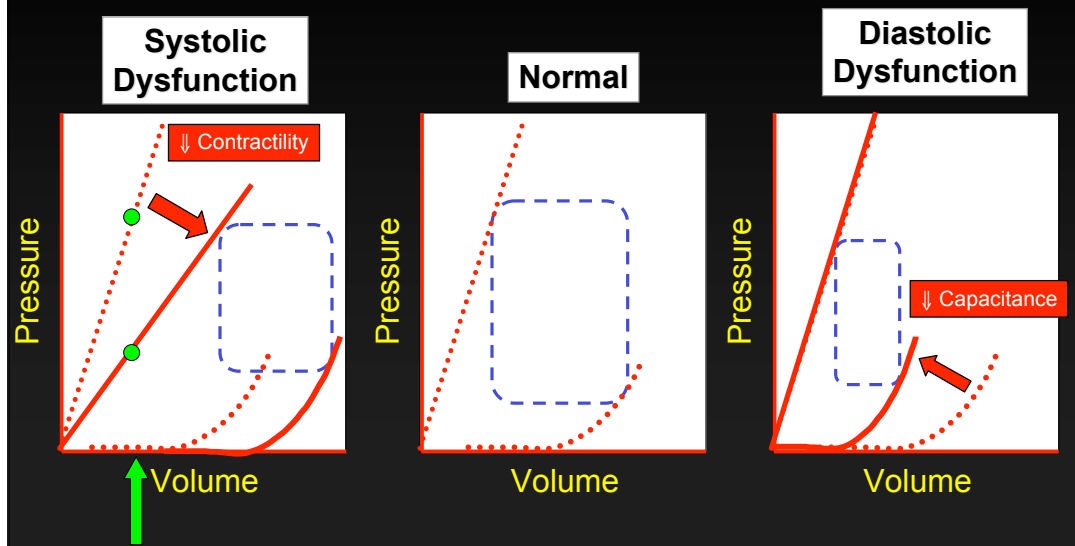
### Heart Failure: Classifications



# Types of Heart Failure

	SHF	Diastolic
		
Pathophysiology	Impaired Contraction	Impaired filling
Demographics	All ages	> 60 years
1° Cause	Coronary Artery Disease	Hypertension

# Systolic Versus Diastolic Failure



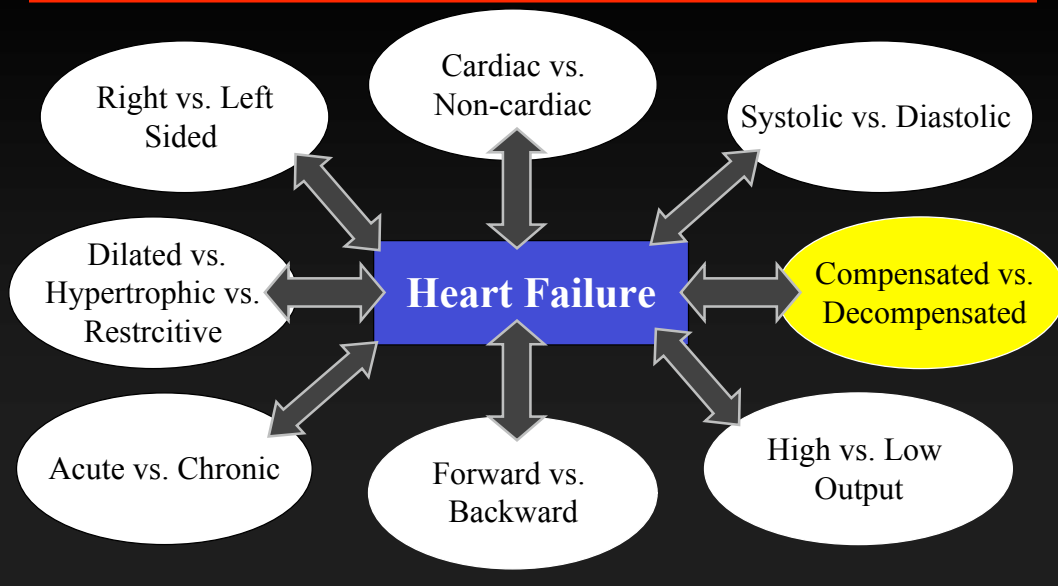
# Systolic Versus Diastolic Failure

**Table 2. Characteristics of Patients with Diastolic Heart Failure and Patients with Systolic Heart Failure.\***

Characteristic	Diastolic Heart Failure	Systolic Heart Failure
Age	Frequently elderly	All ages, typically 50–70 yr
Sex	Frequently female	More often male
Left ventricular ejection fraction	Preserved or normal, approximately 40% or higher	Depressed, approximately 40% or lower
Left ventricular cavity size	Usually normal, often with concentric left ventricular hypertrophy	Usually dilated
Left ventricular hypertrophy on electrocardiography	Usually present	Sometimes present
Chest radiography	Congestion with or without cardiomegaly	Congestion and cardiomegaly
Gallop rhythm present	Fourth heart sound	Third heart sound
Coexisting conditions		
Hypertension	+++	++
Diabetes mellitus	+++	++
Previous myocardial infarction	+	+++
Obesity	+++	+
Chronic lung disease	++	0
Sleep apnea	++	++
Long-term dialysis	++	0
Atrial fibrillation	+	+
	(usually paroxysmal)	(usually persistent)

\* A single plus sign denotes "occasionally associated with," two plus signs "often associated with," three plus signs "usually associated with," and a zero "not associated with."

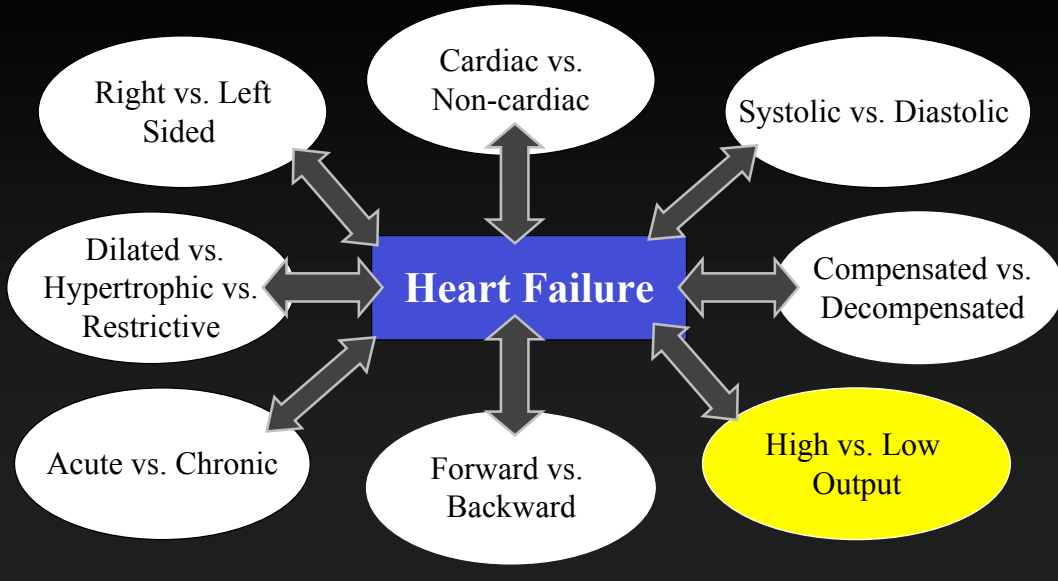
# Heart Failure: Classifications



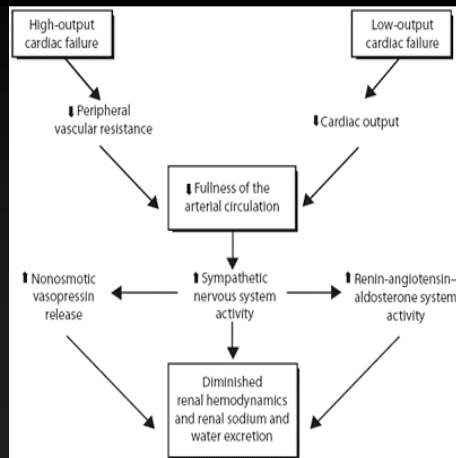
# Decompensated Heart Failure



# Heart Failure: Classifications



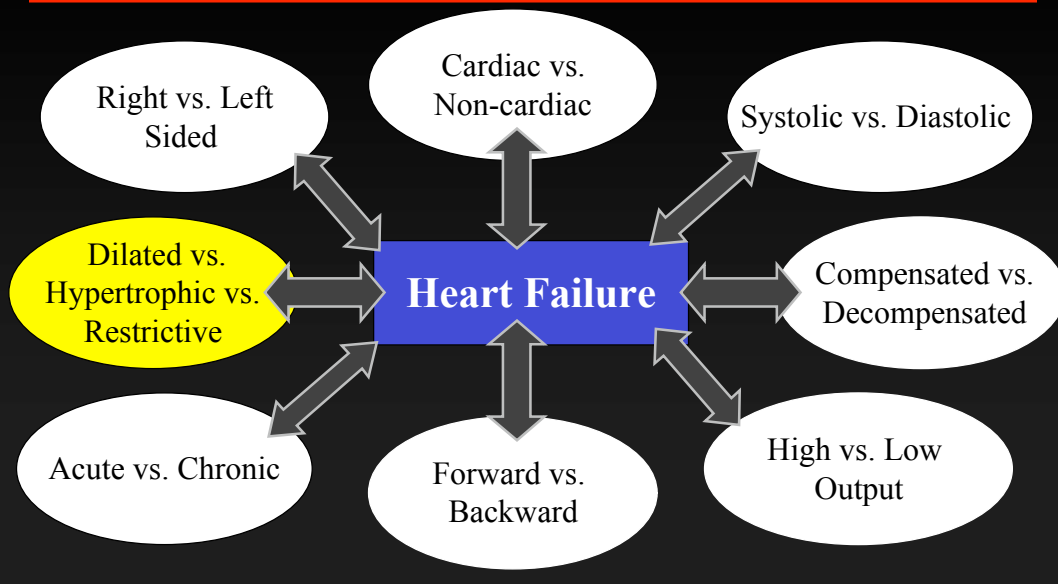
## High vs. Low Output Failure



### • Causes:

- Anemia
- Systemic arteriovenous fistulas
- Hyperthyroidism
- Beriberi heart disease
- Paget disease of bone
- Glomerulonephritis
- Polycythemia vera
- Carcinoid syndrome
- Obesity
- Anemia
- Multiple myeloma
- Pregnancy
- Cor pulmonale
- Polycythemia vera

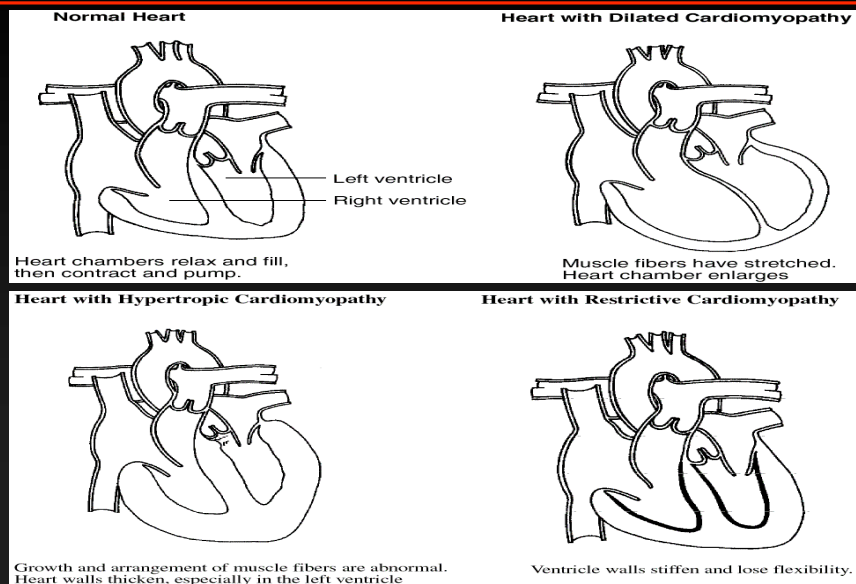
## Heart Failure: Classifications



## Dilated vs. Hypertrophic vs. Restrictive

Type	Definition	Sample Etiologies
Dilated	Dilated left/both ventricle(s) with impaired contraction	Ischemic, idiopathic, familial, viral, alcoholic, toxic, valvular
Hypertrophic	Left and/or right ventricular hypertrophy	Familial with autosomal dominant inheritance
Restrictive	Restrictive filling and reduced diastolic filling of one/both ventricles, Normal/near normal systolic function	Idiopathic, amyloidosis, endomyocardial fibrosis

## Dilated vs. Hypertrophic vs. Restrictive



## Clinical Manifestations

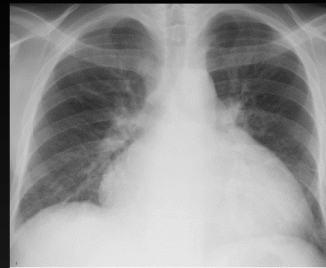
### Symptoms

- Reduced exercise tolerance
- Shortness of breath
- Congestion
- Fluid retention
- Difficulty in sleeping
- Weight loss

Variable	Sensitivity	Specificity
Hx of HF	62	94
Dyspnea	56	53
Orthopnea	47	88
Rales	56	80
S3	20	99
JVD	39	94
Edema	67	68

## Diagnosis of heart failure

- Physical examination
- Chest X ray
- EKG
- Echocardiogram
- Blood tests: Na, BUN, Creatinine, BNP
- Exercise test
- MRI
- Cardiac catheterization



## NYHA Classification

Class	Patient Symptoms
I Mild	<ul style="list-style-type: none"> <li>No limitation of physical activity</li> <li>No undue fatigue, palpitation or dyspnea</li> </ul>
II Mild	<ul style="list-style-type: none"> <li>Slight limitation of physical activity</li> <li>Comfortable at rest</li> <li>Less than ordinary activity results in fatigue, palpitation, or dyspnea</li> </ul>
III Moderate	<ul style="list-style-type: none"> <li>Marked limitation of physical activity</li> <li>Comfortable at rest</li> <li>Less than ordinary activity results in fatigue, palpitation, or dyspnea</li> </ul>
IV Severe	<ul style="list-style-type: none"> <li>Unable to carry out any physical activity without discomfort</li> <li>Symptoms of cardiac insufficiency at rest</li> <li>Physical activity causes increased discomfort</li> </ul>

## ACC/AHA Staging System

**STAGE A** High risk for developing HF

**STAGE B** Asymptomatic LV dysfunction

**STAGE C** Past or current symptoms of HF

**STAGE D** End-stage HF

Hunt, et al. *J Am Coll Cardiol.* 2001; 38:2101-2113.



## ACC/AHA Staging System

Stage	Patient Description
<b>A</b>	High risk for developing heart failure <ul style="list-style-type: none"><li>• Hypertension</li><li>• Coronary artery disease</li><li>• Diabetes mellitus</li><li>• Family history of cardiomyopathy</li></ul>
<b>B</b>	Asymptomatic heart failure <ul style="list-style-type: none"><li>• Previous myocardial infarction</li><li>• Left ventricular systolic dysfunction</li><li>• Asymptomatic valvular disease</li></ul>
<b>C</b>	Symptomatic heart failure <ul style="list-style-type: none"><li>• Known structural heart disease</li><li>• Shortness of breath and fatigue</li><li>• Reduced exercise tolerance</li></ul>
<b>D</b>	Refractory end-stage heart failure <ul style="list-style-type: none"><li>• Marked symptoms at rest despite maximal medical therapy (e.g., those who are recurrently hospitalized or cannot be safely discharged from the hospital without specialized interventions)</li></ul>

## Goals of Treatment

1. Identification and correction of underlying condition causing heart failure.
2. Elimination of acute precipitating cause of symptoms.
3. Modulation of neurohormonal response to prevent progression of disease.
4. Improve long term survival.

## **Etiologies**

---

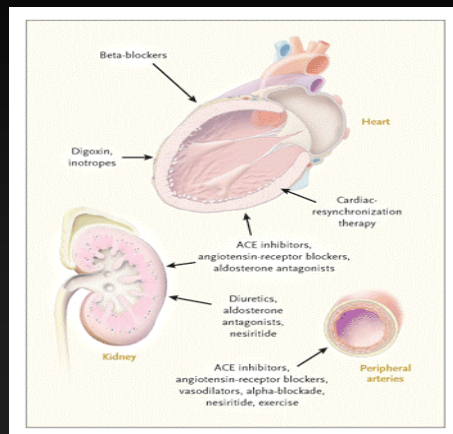
- Ischemic cardiomyopathy
- Valvular cardiomyopathy
- Hypertensive cardiomyopathy.
- Inflammatory cardiomyopathy
- Metabolic cardiomyopathy
- General system disease
- Muscular dystrophies.
- Neuromuscular disorders.
- Sensitivity and toxic reactions.
- Peripartal cardiomyopathy

## **Precipitants /Associated Factors**

---

- **Inappropriate reduction in the intensity of treatment, including**
  - Dietary sodium restriction,
  - Physical activity reduction,
  - Drug regimen reduction, or,
  - most commonly, a combination of these measures.
- **Ischemia**
- **Hypertension**
- **Anemia**
- **Volume Overload**
- **Increased Metabolic Demand**
  - Infection
  - Thyroid Disease
- **Arrhythmia**
- **Asthma/COPD**

# Targets of Treatment



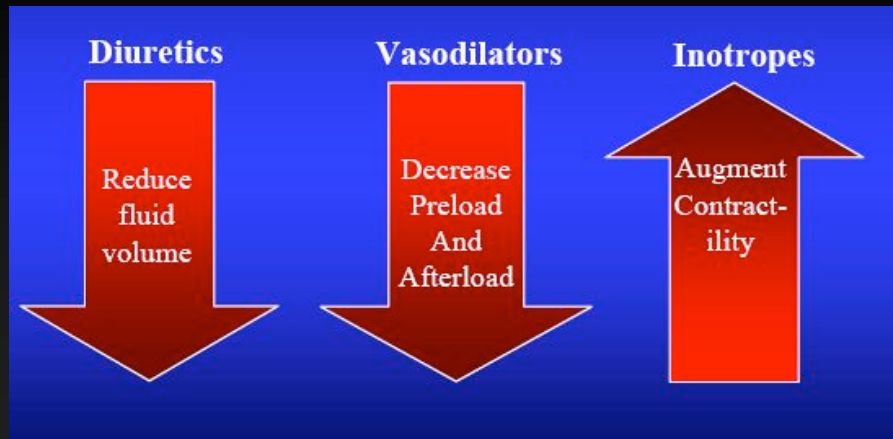
## Standard Pharmacological Therapy

- ACE inhibitors
- Angiotensin Receptor Blockers
- Beta Blockers
- Diuretics
- Aldosterone Antagonists
- Statins
- Vasodilators
- Inotropes

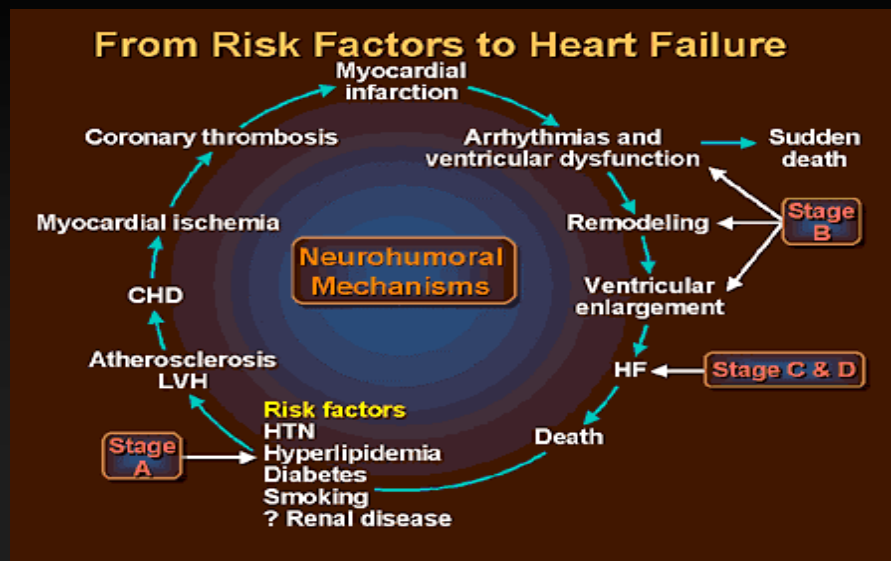
# Treatment

Stage	Patient Treatment
<b>A</b> High risk for developing heart failure	<ul style="list-style-type: none"> <li>• Optimal pharmacologic therapy (OPT)</li> <li>• Aspirin, ACE inhibitors, statins, b-blockers, a-b-blockers (carvedilol) diabetic therapy</li> </ul>
<b>B</b> Asymptomatic heart failure	<ul style="list-style-type: none"> <li>• OPT</li> <li>• ICD if left ventricular (LV) dysfunction (systolic) present</li> </ul>
<b>C</b> Symptomatic heart failure	<ul style="list-style-type: none"> <li>• OPT</li> <li>• ICD if LV dysfunction (systolic) present</li> <li>• CRT (if QRS wide, LVEF≤35%)</li> </ul>
<b>D</b> Refractory end-stage heart failure	<ul style="list-style-type: none"> <li>• OPT</li> <li>• Intermittent IV inotropes</li> <li>• ICD as a bridge to transplantation</li> <li>• CRT</li> <li>• Other devices (LVAD, pericardial restraint)</li> </ul>

# Treatment of Acute Heart Failure



# ACC/AHA Staging System



## Summary

---

- Complex Clinical Syndrome
- Multiple Etiologies and Classification Systems
- Physiologic Understanding Essential

<http://www.columbia.edu/itc/hs/medical/heartsim/>