Updates in Diagnosis & Management of CHF

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- CHF is reaching an epidemic level in U.S. and continues to worsen over time
- Reasons are:
 - -个HTN
 - \uparrow Dm with sedentary lifestyle and obesity
 - Aging population

Epidemiology

- More than 5 million people have Heart Failure
- 580,000 new cases each year



Statistics

Heart Failure

- Incidence 10/1000 after age 65 years
- 75% have antecedent HTN
- After age 40, lifetime risk of developing CHF is 20% for men & women
 - Mortality after CHF hospitalization
 - 30 days 10.4%
 - 1 year 30%
 - 5 years 50%

CHF – Mortality

- (2006) underlying cause of 60,000 deaths mentioned in >282,000 death certificates (1 in 8 deaths)
- 5 year survival lower in men (41%) vs. women (55%)
- Estimated direct & indirect cause of CHF in U.S. for 2010 = \$39.2 Billion

CHF - Definition



A clinical syndrome characterized by symptoms & signs of increased tissue/organ water and decreased tissue/organ perfusion

- Most patients with clinically evident HF have treatable causes
- Factors that increase HF are:
 - Age, HTN, DM, CAD, family history of cardiomyopathy and cardiotoxins
- Modifiable causes are:
 - HTN 66% of patients
 - DM 33% of patients, 2-5% risk of developing
 CM more common in diastolic dysfunction
 - CAD 60% of patients mostly systolic dysfunction

Diagnostic Testing

Dx based on
- Hx
- PE
- Lab
- Imaging



 Most patients present in E.R. with significant dyspnea or fluid overload

Then we determine etiology of CHF





12 lead EKG CXR





Labs: lipid profile, U/A, CMP, CBC, MG+ Cardiac enzymes, drug screen, ETOH TSH - \uparrow or \checkmark can be primary cause of CHF BNP level - to determine if dyspnea from CHF or other causes

Testing

- Single most useful test = 2D echo with doppler looking for:
 - LVEF normal or reduced
 - Structural LV abnormalities such as LVH or wall motion abnormalitites
 - Other structural abnormalities such as valve disorder, pericardial disorder or RV problems



Labs

Naturetic peptides (BNP)

- Synthesized & released from heart
- 个 BNP associated with low LVEF, LVH, elevated LV filling pressure, acute MI, ischemia, pulm embolus, COPD
- BNP is sensitive to age / gender / wt / renal function

Labs

Naturetic peptides (BNP)

- Elevated levels support abnormal LV function or hemodynamics causing Sx of CHF
- BNP levels lower with NI LVEF
- Levels parallel clinical CHF severity & decrease with aggressive CHF Rx

<100 – normal 100- 400 – moderate >400 - severe

Further Work-Up After Stabilization

- PA catheter
- Nuclear stress test
- R + L Heart cardiac catheterization



Treatment

- Moderate sodium restriction
- Fluid restriction
- Daily weights
- Influenza / pneumococcal vaccine
- No heavy labor
- Encourage physical activity except with acute exacerbation

Diuretics

- For patients with concurrent or prior Cx of CHF & ↓ LVEF who have evidence of fluid retention
- Interferes with sodium retention by inhibiting reabsorption of sodium at specific sites in renal tubule
- Loop diuretics (bumetanide, furosemide, torsemide) increase sodium excretion up to 20-25%; maintain efficacy unless renal function severely impaired

Diuretics

- Thiazide diuretics increase fractional sodium excretion 5-10% of filtered load, lose effectiveness if CRCL < 40 ml/min
- Metolazone (Zaroxolyn)
- Produce symptomatic benefits more rapidly than any other Rx for CHF
- Maintain diuresis until fluid overload is eliminated even if hypotension or azotemia develop if pt is asymptomatic
- Monitor electrolytes

ACE-I

- For all Pts with current or prior Sx of CHF and ↓ EF unless C/I
- Target RAAS by reducing formation of angiotensin, which causes blood vessel constriction and increase in BP
- Favorable effects on survival

ACE-I

- No difference among available ACE-I on effect on symptoms or survival
- Adverse effects
 - Hypotension
 - worsening renal function
 - Cough (5-50%)
 - Angioedema (<1%)</p>



- For pts with current or prior Sx of CHF &
 ↓ EF who are ACE-I intolerant
- Works on RAAS to block action of angiotensin's effects on blood vessels
- Angioedema much less likely

 If impaired renal function is >2.0 or TK ->5.0 then consider combination of hydrolazive & nitrates



Aldosterone Antagonists

 For selected pts with moderately to severe symptoms of CHF and ↓ LVEF who can be monitored for renal function & potassium concentration

Ideal creatinine:

- $-Men \leq 2.5 \text{ mg/dL}$
- Women ≤ 2.0 mg/dL
- K⁺ < 5.0 mEq/L

Aldosterone Antagonists

 Targets RAAS by helping reduce salt & fluid; reduce blood volume

 Risk of hyperkalemia, worsening of renal function

2 Agents Used

- Spironolactone (Aldactone)
 - ↓ mortality by 30% and hospitalizations by 30% on stage III and IV
 - Eplerenone (Inspra) used in post MI and ↓ mortality

Beta Blockers

- For all stable pts with current or prior Sx of LHF and ↓ EF.
- The following 3 beta blockers have been proven to decrease mortality and are indicated for use in CHF:
 - Bisoprolol
 - Carvedilol
 - Sr metoprolol succinate

Beta Blockers

- Slows heart rate, lowers BP, helps counteract heart's tendency to compensate for cardiomyopathy by pumping faster
- Risks of Tx
 - Fluid retention
 - Worsening CHF
 - Bradycardia
 - Heart block
 - Fatigue
 - Hypotension

Digitalis

- Can be beneficial in pts with current or prior symptoms of CHF & ↓ LVEF to decrease hospitalizations for CHF
- Causes heart to beat more strongly by increasing force of contractions by inhibiting Na⁺/K⁺ AtPase
- Risks of Tx
 - Cardiac arrhythmias
 - GI symptoms (nausea, vomiting)
 - Neurological problems (visual disturbances, confusion)

CHF & Supraventricular Arrhythmias

- 10-30% of pts with chronic CHF have atrial fibrillation – poor long term prognosis
- Afib exerts effects by:
 - Loss of atrial enhancement of ventricular filling may compromise cardiac output
 - Elevating heart rate increased demand, decreased coronary perfusion d/t shortening of ventricular filling time

CHF & Supraventricular Arrhythmias

- Afib exerts effects by:
 - Rapid ventricular response causes reduction of cardiac contraction & relaxation
 - Stasis of blood in atria can cause pulmonary and systemic emboli

Other Tx that can be used

- Continuous loop diuretics
- Dobutamine or milrinone + inotropic agents
- Nesitiride (Natrecor)
- Ultrafiltration

Transition to Outpatient Management

- A. ACE or ARB
- B. Beta blockers
- C. Counseling (smoke, exercise)
- D. Dieting education, devices
- E. Euvolemia
- F. F/U appointment

Management of Chronic CHF

NYHA Classification

(S) I Sx with strenuous activity
(O) II Sx with ordinary activity
(M) III Sx with most activity
(A) IV Sx with any activity/rest

Stages of CHF

A - No symptoms Predisposed to CHF, such as due to
- CAD/HTN or DM
- LVEF normal
- No LVH

B - No symptoms LVH present Reduced LVEF

Stages of CHF

- C Current or past symptoms with underlying structural heart disease
- D Refractory CHF needing special advanced Tx



Assessment of Functional Capacity

- Inquire about type, severity & duration of symptoms occurring during activities of daily living; inquire about specific tasks
- What tasks can patient no longer perform?
- Measurement of distance patient can walk in 6 minutes



Assessment of Volume Status

- Body weight @ each visit
- JVD → most reliable sign of volume overload
- Peripheral edema
- Rates generally reflect rapidity of CHF onset, not degree of volume overload

Factors Precipitating Hospitalization for CHF

- Noncompliance w Rx/Na⁺ or fluid restriction
- Acute MI
- Afib
- Recent starting of Θ inotrope such as verapamil / nifedipine / diltiazem or beta blocker

Factors Precipitating Hospitalization for CHF

- Pulm embolus
- NSAIDS → cause sodium retention / peripheral vasoconstriction / decrease efficacy & enhance toxicity of diuretics & ACE-I
- ETOH/Illicits
- Enodcrine abnormalities (Dm, thyroid)
- Concurrent infection (pneumonia)

CHF Prognosis Worsened With:

- Low LVEF
- Worsening NYHA status
- Degree of hyponatremia
- Low hematocrit
- Wide QRS on EKG
- Chronic hypotension
- Resting tachycardia
- Renal insufficiency
- Intolerance to conventional Tx
- Refractory volume overload



ICD

 For 2° prevention in pts with current or prior Sx of CHF and ↓ EF with Hx of cardiac arrest, VF or hemodynamically destabilizing VT



ICD

- For 1° prevention of SCD to reduce total mortality in patients with ischemic dilated cardiomyopathy or ischemic heart Dz at least 40 days post-MI, LVEF < 35% & NYHA II-III Sx while on chronic, optimal med Tx and who have reasonable expectation of survival with good functional status > 1 year
- Or after 9 months for dilated cardiomyopathy

CRT -

Cardiac Resynchronization Therapy

- Cardiac dyssynchrony = QRS duration <u>></u> 0.12 sec
- These patients should receive CRT, with or without ICD, unless contraindicated if they also have LVEF 35%, sinus rhythm, NYHA III or ambulatory NYHA IV Sx despite optimum Rx

CRT – Cardiac Resynchronization Therapy

- About 1/3 of patients affected
- Dyssynchrony causes suboptimal ventricular filling, prolonged duration of mitral regurgitation and paradoxical septal motion; associated with increased mortality.

CRT –

Cardiac Resynchronization Therapy

 Electrical activation of R&L ventricles in synchronized manner with biventricular pacing enhances ventricular contractions and reduces degree of mitral regurgitation, improves cardiac function and hemodynamics

CRT -

Cardiac Resynchronization Therapy

- CRT & optimal medical Tx shows improvement in quality of life, functional class, exercise capacity, 6-minute walk and LVEF
- 32% reduction of hospitalization for CHF, 25% reduction of all cause mortality within 3 months
- Based on studies on patients in NSR, not afib

EECP

- Enhanced external counterpulsation
- Uses 3 sets of inflating pneumatic cuffs attached to pts legs that rapidly inflate and deflate
- Applied to calves, lower thigh, upper-thigh; timed to heart beat



EECP

- 1-hour sessions for 35 days
- Improves blood pressure, blood flow, exercise capacity and duration, NYHA class & quality of life



- CHF with normal LVEF & abnormal diastolic function
- Prevalent among elderly females with HTN, Dm or both, often with CAD and afib
- Have slowed ventricular relaxation 个 LV filling pressure
- No valvular disease (aortic stenosis or mitral regurg)

- Principles of Rx
 - BP / HR / blood volume / myocardial ischemia control
 - Treat other Dz like CAD / HTN / aortic stenosis
 - Diuretics to control pulmonary congestion
 - Class IIB beta blockers, ACE-I / ARB / CCB may minimize Sx
 - Digoxin not well established

Morbidity / Mortality

- 15-20 million CHF pts (1/3 1/2 of CHF patients)
- 5-8%, annual mortality vs. 10-15% for systolic CHF; age matched controls – 1%
- 1 year readmission rates 50%

Dx of 1° Diastolic CHF

- Simultaneously requires
 - Presence of signs or symptoms of CHF
 - Presence of normal or mildly abnormal (LVEF ≥ 45%) LV systolic function
 - Evidence of abnormal LV relaxation, filling, diastolic distensibling or diastolic stiffness
 - Dx cannot be made at bedside

Treatment

- Goal: ↓ diastolic pressure
- Reduce pulmonary congestion by decreasing LV volume, maintaining synchronous atrial contraction and increasing duration of diastole by ↓ HR

Treatment

- Decrease total blood volume by fluid and salt restriction, and use of diuretics (usually at lower doses than for systolic CHF)
- Decrease central blood volume with nitrates
- Blunt neurohormonal activation with ACE-I/ARBS/Aldosterone antagonists
- Start with low doses to avoid hypotension
- Trials underway for future Tx

Heart Transplant

- End stage CHF
- LVAD LV assist device surgically implanted, bridge to transplant
- Survival rates
 - 88% 1st year post transplant
 - 72% @ 5 years
 - 50% @ 10 years
 - 16% @ 20 years
- About 2,000 heart transplants performed yearly in U.S.

LVAD (left ventricular assist device)

- Take oxygenated blood from LA or LV go through pump and return to aorta
- Short or long term use
- Bridge to transplantation (short term)
- Destination therapy long term treatment with lower mortality. However a number of patients (50%) still died at 1 year.



HEART INSTI

Heart Transplantation

- Successful since 1980's with immunosuppression treatment
- >2000 transplants a year
- Survival: 1 yr- 85%, 5 yr- 70%, 10 yrs- 50%
- Selection is important:
 - Optimal medical treatment
 - Survival < 1 year</p>
 - Good compliance and F/U

Heart Transplantation

- Contrainidications
 - Severe HTN
 - Infection
 - COPD
 - PVD or CVD
 - Psychiatric Dx
 - Liver disease
 - Age >70
 - DM end organ Dx
 - Malignancy with ψ L/E