# Drug Therapy Of Heart Failure 

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## Learning objectives

By the end of this lecture, students should be able to:

- Describe the different classes of drugs used for treatment of acute \& chronic heart failure \& their mechanism of action
- Understand their pharmacological effects, clinical uses, adverse effects \& their interactions with other drugs.


## HEART FAILURE



Inability of the heart to maintain an adequate cardiac output to meet the metabolic demands of the body.

## GAUSES OF MENRT FMDURE

disorder of
coronary arteries
high
blond pressurie

HEART FAILURE
cardionnyopathy

Acute or Chronic

## Symptoms of Heart failure

- Tachycardia
- Decreased exercise tolerance (rapid fatigue)
- Dyspnea (pulmonary congestion)
- Peripheral edema
- Cardiomegaly.



## Pathophysiology of CHF

## $\downarrow$ Force of contraction

Low C.O.
$\downarrow$ Renal blood flow

Activate renin-angiotensin-
Aldosterone system
$\downarrow$ Carotid sinus firing

Activate sympathetic system
$\uparrow$ Sympathetic discharge


## $\uparrow$ Preload

$\uparrow$ Afterload

## Factors affecting cardiac output and heart failure

## 1- Preload

2- Afterload
3- Cardiac contractility

## Drugs used in treatment of heart failure

I- Drugs that decrease preload:
1 - Diuretics
2 - Aldosterone antagonists
3 - Venodilators
II- Drugs that decrease afterload:
1 - Arteriodilators

## Drugs used in treatment of heart failure

III- Drugs that decrease both preload \& afterload: (Combined arteriolo- \& venodilators)

1- Angiotensin converting enzyme (ACE) inhibitors
2- Angiotensin receptor antagonists
3- $\alpha_{1}$-adrenoceptor antagonists
4- Direct vasodilators

Drugs used in treatment of heart failure

## IV- Drugs that increase heart contractility:

1- Cardiac glycosides (digitalis)
2- $\beta$-adrenoceptor agonists
3- Phosphodiesterase inhibitors

## I- Drugs that decrease preload

## 1-Diuretics:

Mechanism of action in heart failure :
reduce salt and water retention
$\downarrow$
decrease ventricular preload and venous pressure
$\downarrow$
reduction of cardiac size
$\downarrow$
Improvement of cardiac performance

I- Drugs that decrease preload

## 1-Diuretics:

## Chlorothiazide

- first-line agent in heart failure therapy
- used in volume overload (pulmonary and/ or peripheral edema)
- used in mild congestive heart failure.


## I- Drugs that decrease preload

## 1-Diuretics:

## Furosemide

- a potent diuretic
- used for immediate reduction of pulmonary congestion \& severe edema associated with :
- acute heart failure
- moderate \& severe chronic failure


## I- Drugs that decrease preload

## 2-Aldosterone antagonists:

## Spironolactone

- non-selective antagonist of aldosterone receptor
- a potassium sparing diuretic
- improves survival in advanced heart failure


# I- Drugs that decrease preload 2-Aldosterone antagonists: 

## Eplerenone

- a new selective aldosterone receptor antagonist
(does not inhibit other hormones such as estrogens \& androgens)
- indicated to improve survival of stable patients with congestive heart failure.


## I- Drugs that decrease preload

3-Venodilators:

> A- Nitroglycerine B- Isosorbide dinitrate

- used I.V. for severe heart failure when the main symptom is dyspnea due to pulmonary congestion \& edema
- dilates venous blood vessels \& reduce preload.


## II- Drugs that decrease afterload

## 1- Arteriodilators:

## Hydralazine

- used when the main symptom is rapid fatigue due to low cardiac output
- reduce peripheral vascular resistance


## III- Drugs that decrease both preload \& afterload

1-Angiotensin converting enzyme (ACE) inhibitors:

- considered as first-line drugs for
chronic heart failure along with diuretics
- first-line drugs for hypertension therapy.


## Angiotensin converting enzyme inhibitors MECHANISM OF ACTION

VASOCONSTRICTION
VASODILATATION


SYMPATHETIC
1 Angiotensinogen


## BRADYKININ

Breakdown

## ACE Inhibitors

## Pharmacological actions:

1 - Decrease peripheral resistance (Afterload)
2 - Decrease Venous return (Preload)
3 - Decrease sympathetic activity
4- Inhibit cardiac and vascular remodeling associated with chronic heart failure Decrease in mortality rate

## ACE Inhibitors

## Captopril, Enalapril and Ramipril

 Pharmacokinetics:- rapidly absorbed from GIT after oral administration
- food reduce their bioavailability


## Enalapril , Ramipril

- prodrugs, converted to their active metabolites in liver
- have long half-life \& given once daily.


## ACE Inhibitors

Adverse effects:
1- acute renal failure, especially in patients with renal artery stenosis

2- hyperkalemia, especially in patients with renal insufficiency or diabetes

3- severe hypotension in hypovolemic patients
(due to diuretics, salt restriction or gastrointestinal fluid loss).

## ACE Inhibitors

## Adverse effects:

4- dry cough sometimes with wheezing

5- angioneurotic edema (swelling in the nose, throat, tongue, larynx)

6- dysgeusia (reversible loss or altered taste).

## ACE Inhibitors

## Contraindications:

- during the second \& third trimesters of pregnancy
(due to the risk of : fetal hypotension renal failure \& malformations)
- renal artery stenosis.


## III- Drugs that decrease both preload \& afterload

## 2- Angiotensin receptor blockers (ARBs) :

## Losartan, Valsartan, Irbesartan

Mechanism of action:

- block AT 1 receptors
- decrease action of angiotensin II


# III- Drugs that decrease both preload \& afterload 

## 3- $\alpha-A D R E N O C E P T O R$ BLOCKERS :

## Prazosin

- blocks $\alpha$ - receptors in arterioles and venules
- decrease both afterload \& preload


# III- Drugs that decrease both preload \& afterload 

## 4- Direct acting vasodilators:

## Sodium nitroprusside

- given I.V. for acute or severe heart failure
- acts immediately and effects lasts for 1-5 min.

IV- Drugs that increase contractility 1-Cardiac glycosides (digitalis):

## Digoxin

-increases the force of myocardial contraction (+ve inotropic effect)
Mechanism of action :

- Inhibit $\mathrm{Na}^{+} / \mathrm{K}^{+}$ATPase enzyme
(the sodium pump)


## MECHANISM OF ACTION OF DIGOXIN



## IV- Drugs that increase contractility

1- Cardiac glycosides (digitalis) :

## Digoxin

Therapeutic uses:

- Congestive heart failure
- has narrow therapeutic index

1- Cardiac glycosides (digitalis) : Digoxin

Adverse effects (Cardiac):

- digitalis-induced arrhythmias
- extrasystoles
- coupled beats (Bigeminal rhythm)
- ventricular tachycardia or fibrillation
- cardiac arrest


## Bigeminal PVC's: every other beat is a PVC.



## IV- Drugs that increase contractility

 1- Cardiac glycosides (digitalis):
## Digoxin

Adverse effects (non-cardiac):
GIT:
anorexia, nausea, vomiting, diarrhea
CNS:
headache, visual disturbances, drowsiness.

IV- Drugs that increase contractility 1- Cardiac glycosides (digitalis) : Digoxin

Factors that increase its toxicity:

- Renal diseases
- Hypokalemia
- Hypomagnesemia
- Hypercalemia


## IV- Drugs that increase contractility

## 2- $\beta$-Adrenoceptor agonists:

## Dobutamine

- Selective $\beta_{1}$ agonist
- Uses : Treatment of acute heart failure in cardiogenic shock


## IV- Drugs that increase contractility

 3- phosphodiesterase -III inhibitors: Milrinone
## Mechanism of action:

Inhibits phosphodiesterase -III (cardiac \& B. Vessels)

Increase cardiac
Contractility
dilatation of arteries \& veins
(reduction of preload \& afterload)

# IV- Drugs that increase contractility 3- phosphodiesterase -III inhibitors : <br> <br> Milrinone 

 <br> <br> Milrinone}

## Therapeutic uses:

- used only IV for management of


## acute heart failure

- not safe or effective in the longer ( > 48 hours) treatment of patients with heart failure


# IV- Drugs that increase contractility 3- phosphodiesterase -III inhibitors : Milrinone 

## Adverse effects:

- Hypotension and chest pain (angina)


## Chemical interaction:

- furosemide should not be administered in I.V. lines containing milrinone due to formation of a precipitate
- Enoximone \& Vesnarinone new drugs in clinical trials.


## The use of $\beta$-adrenoceptor blockers in heart failure

- The elevated adrenergic activity in chronic heart failure patients cause structural remodeling of the heart (cardiac dilatation \& hypertrophy)
$\beta$-blockers:
- reduce the progression of chronic heart failure
- not used in acute heart failure.


## The use of $\beta$-adrenoceptor blockers in heart failure

Mechanism of action of $\beta$-blockers in HF:
1- attenuate cardiac remodeling
2- slow heart rate, which allows the left ventricle to fill more completely
3- decrease renin release
reduce mortality \& morbidity of patients with HF

## The use of $\beta$-adrenoceptor blockers in heart failure

- Second generation:
cardioselective ( $\beta_{1}$-receptors)
e.g. Bisoprolol, Metoprolol
- Third generation:
have vasodilator actions ( $\alpha$ - blocking effect)
e.g. Carvedilol, Nebivolol.


## New drugs for heart failure

1- Natriuretic Peptides:

## Nesiritide

- BNP is secreted by the ventricles in response to stretch
- elevated BNP is associated with advanced heart failure (compensatory mechanism in HF)

Cardiac distension
Sympathetic stimulation
Angiotensin Il
Endothelin


## Natriuretic Peptides

## Nesiritide

- a purified preparation of human BNP, manufactured by recombinant DNA technology
- $\uparrow$ cyclic-GMP in vascular smooth muscle, leading to smooth muscle relaxation \& reduction of preload and afterload
- indicated (IV) for the treatment of patients with acute decompensated heart failure (ADHF) who have dyspnea at rest or with minimal activity.


## New drugs for heart failure

2- Calcium sensitisers:

## Levosimendan

- used in the management of ADHF


## New drugs for heart failure <br> Levosimendan

## mechanism of action:

- Calcium sensitization
(improves cardiac contractility without increasing oxygen consumption)
- potassium-ATP channel opening
(cause vasodilation, improving blood flow to vital organs)
These effects reduce the risk of worsening ADHF or death compared with dobutamine.


## Management of chronic heart failure

> Reduce work load of the heart

- Limit patient activity
- Reduce weight
- Control hypertension
> Restrict sodium
> Stop smoking.


## Heart Failure Functional Classification

## NYHA <br> Class

## Symptoms

Cardiac disease, but no symptoms \& no limitation in ordinary physical activity, e.g. no shortness of breath when walking, climbing stairs etc

Mild symptoms (mild shortness of breath \&/or angina), slight limitation during ordinary activity

Marked limitation in activity due to symptoms, even during less-than-ordinary activity, e.g. walking short distances (20-100 m). Comfortable only at rest

Severe limitations. Experiences symptoms even while at rest. Mostly bedbound patients.

## Management of chronic heart failure

|  | For Survival/Morbidity | For Symptoms |
| :--- | :--- | :--- |
| NYHA I | Continue ACE inhibitor/ARB if ACE <br> inhibitor intolerant, continue <br> aldosterone antagonist if post-MI <br> add beta-blocker if post-MI | reduce / stop diuretic |
| NYHA II | ACE inhibitor as first-line treatment/ARB <br> if ACE inhibitor intolerant <br> add beta-blocker <br> and aldosterone antagonist if post MI | +/- diuretic depending on <br> fluid retention |
| NYHA III | ACE inhibitor plus ARB or ARB <br> alone if ACE intolerant <br> beta- blocker <br> add aldosterone <br> antagonist | + diuretics + digitalis <br> If still symptomatic |
| NYHA <br> IV | Continue ACE inhibitor/ARB <br> beta-blocker <br> Aldosterone antagonist | +diuretics + digitalis <br> + consider temporary <br> inotropic support |

Congestive Heart Failure in Black patients

## Hydralazine/isosorbide dinitrate fixed dose combination

- FDA approved to add to standard therapy for black Americans with congestive heart failure
(due to poor response to ACE inhibitors)
- should be considered for patients intolerant to ACE inhibitors \& ARBs due to renal dysfunction


## Acute decompensated heart failure (ADHF)

A sudden worsening of the signs \& symptoms of heart failure, which typically includes:

- difficulty breathing (dyspnea)
- leg or feet swelling
- fatigue
- ADHF is a common \& potentially serious cause of acute respiratory distress.


## Management of acute heart failure

Ultrafiltration: Aqua/natriuresis

Nitrates, nitroprusside, dobutamine: Arterial vasodilation


Bilevel or continuous
positive airway pressure:
Preload reduction

Dobutamine, dopamine, milrinone: Increased inotropy

Nitrates, morphine:
Venodilation


Furosemide:
Natriuresis

## Thank you



