## ANTIANGINAL DRUGS

## LEARNING OUTCOMES

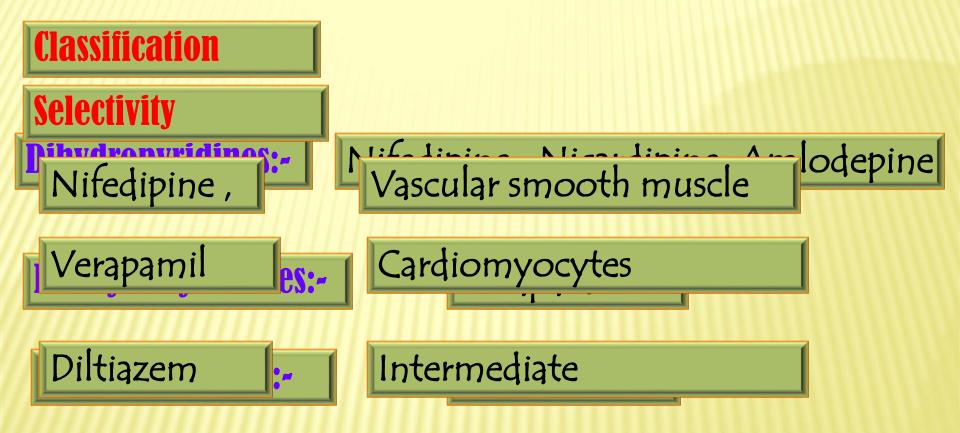
Recognize variables contributing to a balanced myocardial supply versus demand

Expand on the drugs used to alleviate acute anginal attacks versus those meant for prophylaxis & improvement of survival

Detail the pharmacology of nitrates, other vasodilators, and other drugs used as antianginal therapy



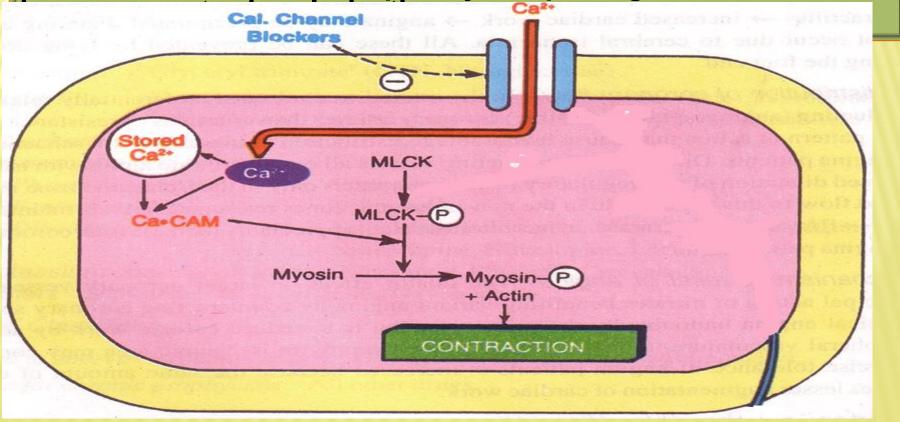
#### Calcium channel blockers



## **Mechanism of Action**

Binding of calcium channel blockers [CCBs] to the L-type Ca channels 

their frequency of opening



## **Antianginal Action**

**Cardiomyocyte Contraction** → **Cardiac** work through their –ve inotropic & chronotropic action (verapamil & diltiazem) → **Impocardial oxygen demand** 

- **→ VSMC Contraction → →** After load **→ →** cardiac work **→**
- **→** myocardial oxygen demand

Coronary dilatation - myocardial oxygen supply

## Therapeutic Uses

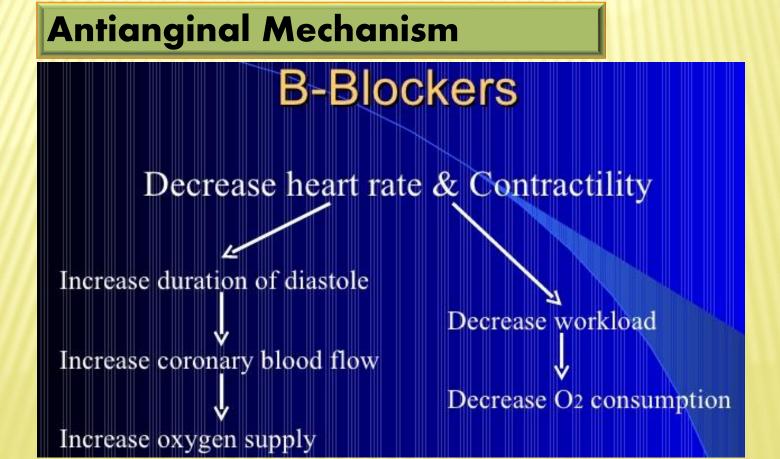
Short acting dihydropyridine should be avoided ??

Can be combined to β-AR blockers???

IN LINSTARI F ANGINA: Soldom added in refractory cases Can be combined with nitrates???

Dihydropyridenes useful antianginal if with CHF??

**Examples Atenolol, Bisoprolol, Metoprolol** ( $\beta_1$  – Selective)



**Indications in angina** 

In stable angina

Regular prophylaxis, selective are preferred?

First choice for chronic use?

Can be combined with nitrates?

Can be combined with dihydropyridine CCB?

| Verapamil? |

In variant angina

**Contraindicated?** 

**Indications in angina** 

In Unstable angina

Halts progression to MI, improve survival

In Myocardial infarction

**Reduce infarct size** 

**Reduce morbidity & mortality** 

→ reduce **02 demand** 

→reduce **arrhythmias** 

β- blockers should be withdrawn gradually?

Given to diabetics with ischemic heart disease?

#### MINICASE



Which antianginal drug is the best choice for the case of Helmi? And Why?

## **MINICASE**

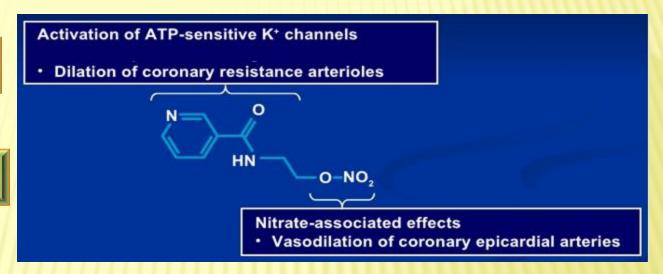


If Helmi does not respond to monotherapy, what other drug should be added to his regimen?

## Potassium channel openners

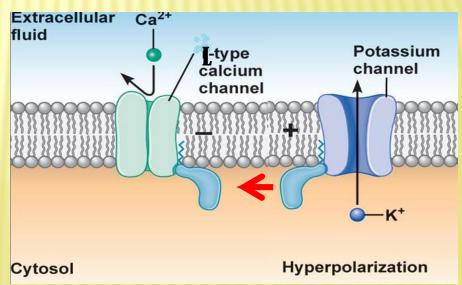
**Nicorandil** 

Mechanism



It has dual mechanism of action;

- 1. Opens K<sub>ATP</sub> channels
- (> arteriolar dilator)
- 2. NO donor as it has a nitrate moiety (> venular dilator)



## **Pharmacodynamic Effects**

## As K channel openner

- As nitric oxide donor
- s opening of K channels
- → hyperpolarization → vasodilatation
- NO ↑ cGMP/PKG → vasodilatation
- On cardiomyocytes opening of A charmers Trepolarization
- **→ ↓** cardiac work

#### **Indications**

Prophylactic 2nd line therapy in stable angina & refractory variant angina

#### **ADRs**

Flushing, headache, Hypotension, palpitation, weakness Mouth & peri-anal ulcers, nausea and vomiting.

#### THINK-PAIR-SHARE

A 55 - year - old woman complained to her physician of palpitations, flushing of the face, and vertigo. The woman, suffering from diabetes mellitus, was giving herself three daily doses of insulin. She had been recently diagnosed with exertional angina for which nitrate therapy was started with transdermal nitroglycerin and oral isosorbide mononitrate. After 3 weeks of therapy, her anginal attacks were less frequent but not completely prevented. Which would be an appropriate next therapeutic step for this patient?

## Metabolically Acting Agents

e.g. Trimetazidine 02 requirement of glucose pathway is Myocytes Glucose FFA Duri ls rise, blun Acyl-CoA Pyruvate B-oxidation Trimetazidine Acetyl-CoA

Reduces 02 demand without altering hemodynamics

Energy for contraction

# Trimetazidine

Indications

Used as an add on therapy

**ADRs** 

GIT disturbances

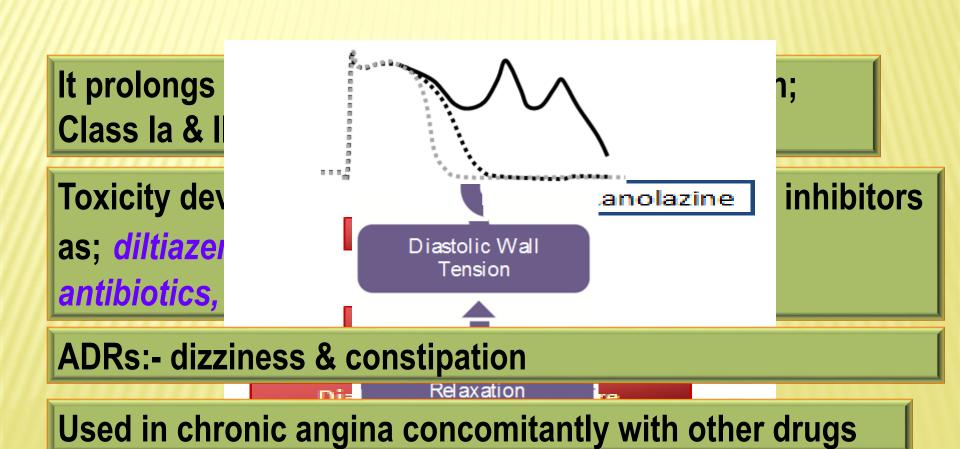
Contraindications

Hypersensitivity reaction

Pregnancy & lactation

## Ranolazine

Inhibits the late sodium current which increases during ischemia



## MINICASE



Which antihyperlipidemic drug should be prescribed to Helmi?

# Ivabradine

### Ivabradine Selectively blocks I<sub>f</sub>

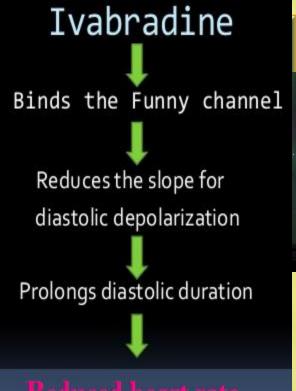
Ivabradine reduces slope of depolarization, slowing HR, reducing myocardila work & 02 demand

econasi

Used in combination with beta blocked heart failure with LVEF lower than 35 inadequately controlled by beta block whose heart rate exceeds 70/min

ADR:- luminous phenomena

I<sub>f</sub> current is a -60 ± pacemaker ( Potential (mV)



# Agents that improve prognosis

- -Aspirin / other antiplatelet agents
- -ACE inhibitors
- -Statins
- -β -blockers

Halt progression
Prevent acute insult
Improve survival

## MEMORY MATRIX

# In the following table indicate increase, decrease or no effect with signs $\uparrow$ , $\downarrow$ , – respectively

Drug/Class	HR	BP	Wall Tension	Contract- ility	O <sub>2</sub> Supply
Beta-blockers					
CCBs					
Verap/Dilt					
Dihydropyridines					
Nitrates					
Ranolazine					