

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

Classification

Selectivity

Dihydropyridines:-

Nifedipine ,

Nifedipine, Nifedipine

Vascular smooth muscle

Amlodipine

Verapamil

es:-

Cardiomyocytes

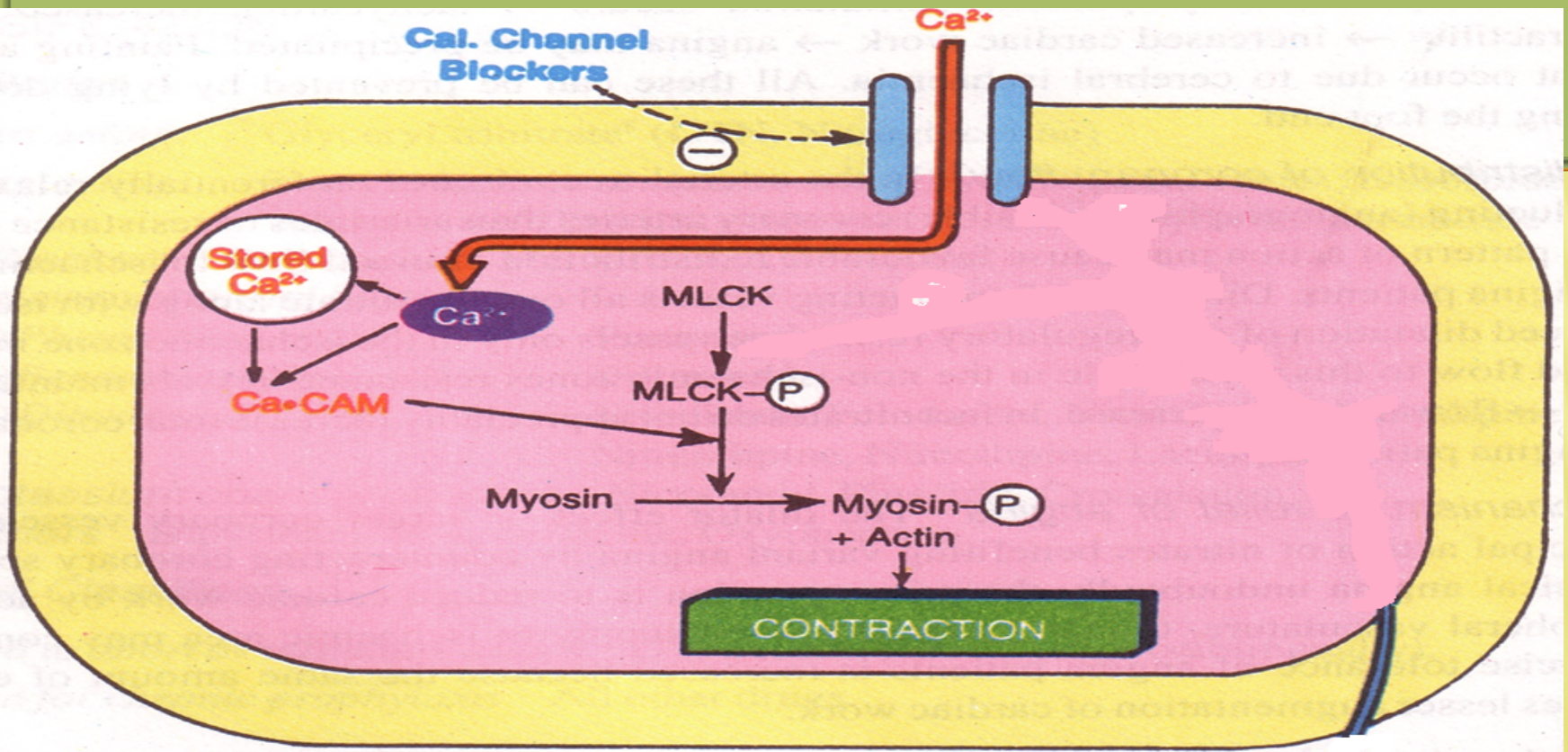
Diltiazem

:-

Intermediate

Mechanism of Action

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



Antianginal Action

↓ **Cardiomyocyte Contraction** → ↓ cardiac work through their –ve inotropic & chronotropic action (verapamil & diltiazem) → **↓ myocardial oxygen demand**

↓ **VSMC Contraction** → ↓ Afterload → ↓ cardiac work → **↓ myocardial oxygen demand**

Coronary dilatation → **↑ myocardial oxygen supply**

Therapeutic Uses

IN VARIANT ANGINA

Short acting dihydropyridine should be avoided ??

▶ Attacks prevented (> 60%) /
sometimes variably aborted

Can be combined to β -AR blockers???

IN UNSTABLE ANGINA:

Can be combined with nitrates???

Seldom added in refractory cases

Dihydropyridenes useful antianginal if with CHF??

Beta Adrenoceptor Blockers

Examples Atenolol, Bisoprolol, Metoprolol (β_1 – Selective)

Antianginal Mechanism

B-Blockers

Decrease heart rate & Contractility

Increase duration of diastole

Increase coronary blood flow

Increase oxygen supply

Decrease workload

Decrease O₂ consumption

Beta Adrenoceptor Blockers

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

Beta Adrenoceptor Blockers

Indications in angina

In Unstable angina

Halts progression to MI, improve survival

In Myocardial infarction

Reduce infarct size

Reduce morbidity & mortality

→ reduce **O₂ demand**

→ reduce **arrhythmias**

Beta Adrenoceptor Blockers

β- blockers should be withdrawn gradually?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?

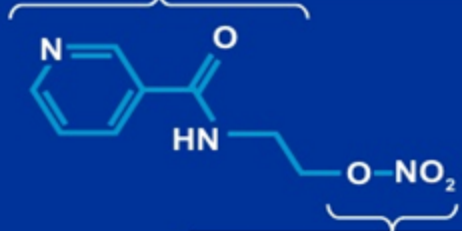
Potassium channel openers

Nicorandil

Mechanism

Activation of ATP-sensitive K⁺ channels

- Dilation of coronary resistance arterioles

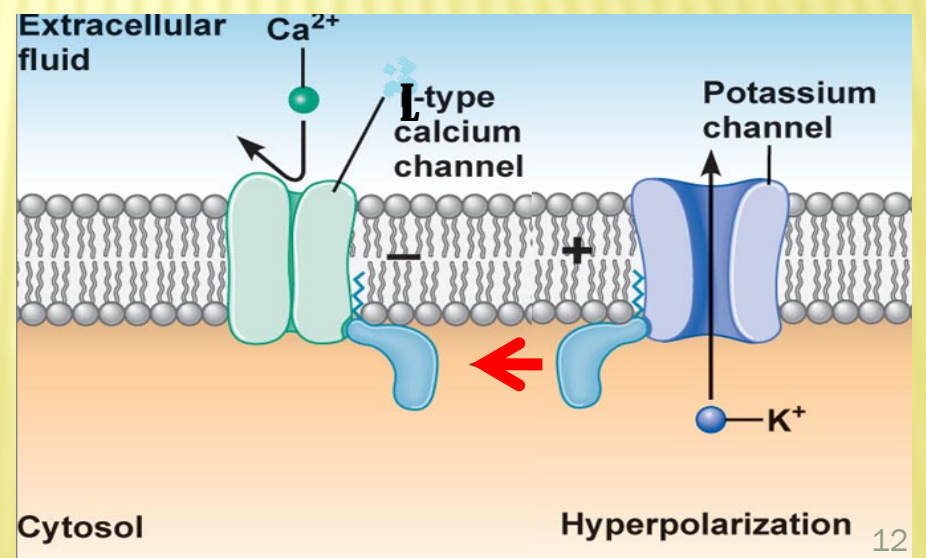


Nitrate-associated effects

- Vasodilation of coronary epicardial arteries

It has dual mechanism of action;

1. Opens K_{ATP} channels (> arteriolar dilator)
2. NO donor as it has a nitrate moiety (> venular dilator)



Pharmacodynamic Effects

As K channel opener

As nitric oxide donor → opening of K channels

→ hyperpolarization → vasodilatation

NO ↑ cGMP/PKG → vasodilatation

On cardiomyocytes opening of K channels → repolarization

→ ↓ 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

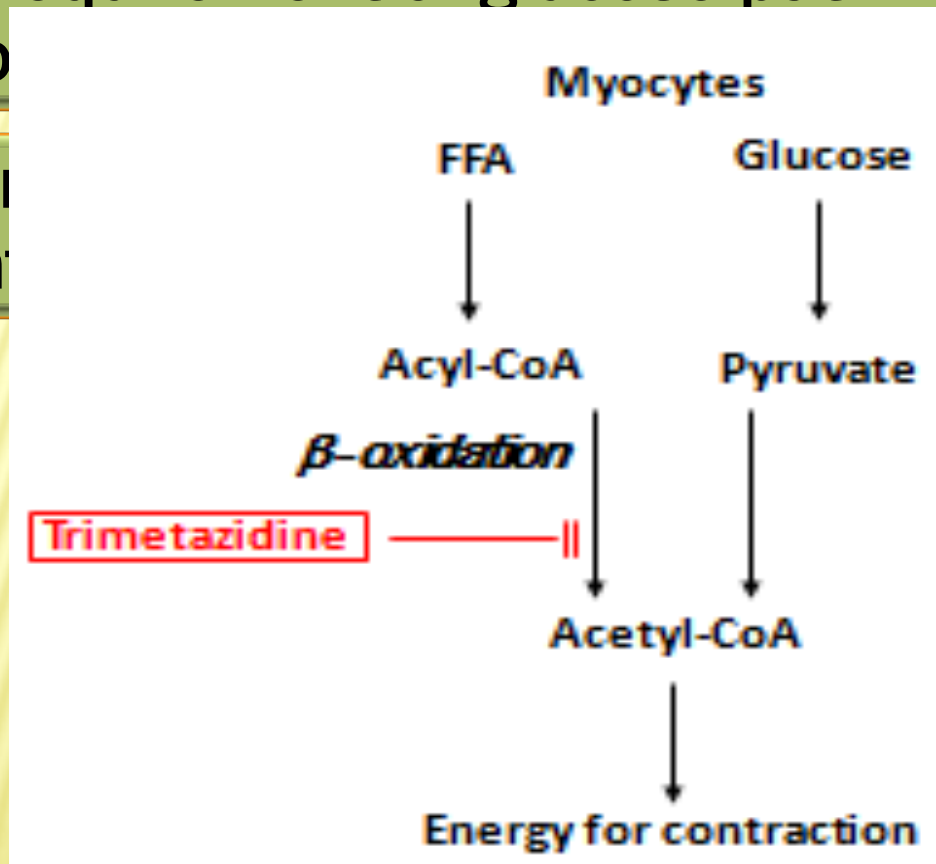
e.g. Trimetazidine

O₂ requirement of glucose pathway is

low

During
blunt

is rise,



Reduces O₂ demand without altering hemodynamics

Trimetazidine

Indications

Used as an add on therapy

ADRs

GIT disturbances

Contraindications

Hypersensitivity reaction

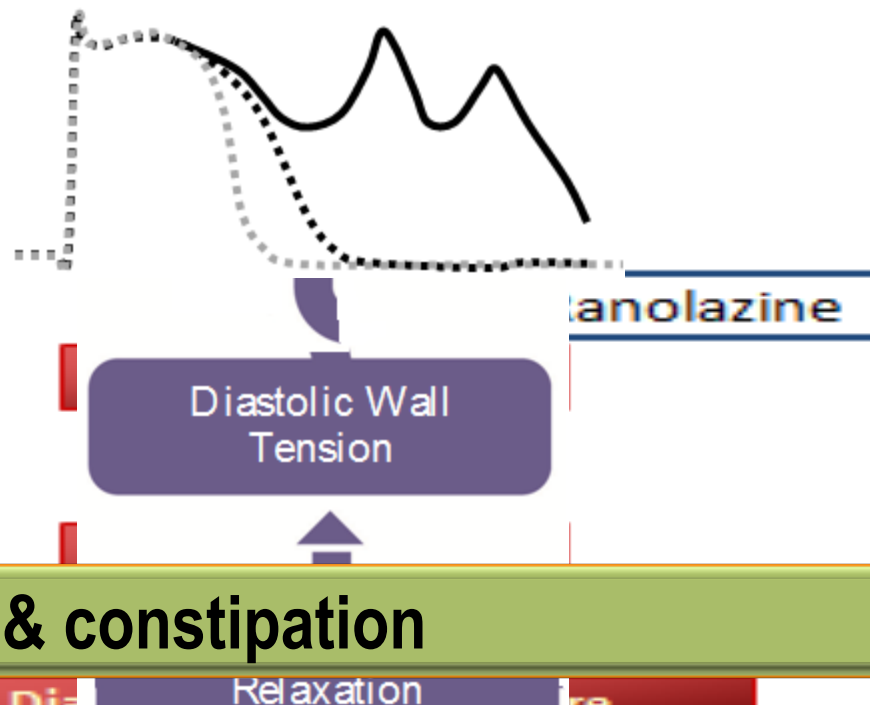
Pregnancy & lactation

Ranolazine

Inhibits the late sodium current which increases during ischemia

It prolongs
Class Ia & II

Toxicity develops with
as; *diltiazem*,
antibiotics,



ADRs:- dizziness & constipation

Used in chronic angina concomitantly with other drugs

Minicase



Which antihyperlipidemic drug should be prescribed to Helmi?

Ivabradine

Ivabradine Selectively blocks I_f

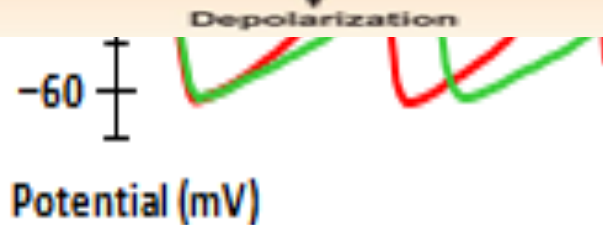
Ivabradine reduces slope of depolarization, slowing HR, reducing myocardial work & O₂ demand

Ivabradine

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

ADR:- luminous phenomena

I_f current is a pacemaker



Ivabradine

Binds the Funny channel

Reduces the slope for diastolic depolarization

Prolongs diastolic duration

Reduced heart rate

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	Contractility	O ₂ Supply
Beta-blockers					
CCBs					
Verap/Dilt					
Dihydropyridines					
Nitrates					
Ranolazine					