بسم الله الرحمن الرحيم

Adrenergic receptors blockers
(antagonists)
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1- α adrenergic blockers

Pharmacological actions:

- Blocking α₁ receptors will relax smooth muscles of blood vessels leading to vasodilatation and decrease peripheral resistance and hypotension.
- \square Hypotension \rightarrow reflex tachycardia.
- □Blocking of the pre-synaptic α₂ receptors will increase Norepinephrine release leading to tachycardia (beta 1 effect).
- Prostates: Blockade of α₁ receptors will relax the prostatic smooth muscles and can improve the urinary obstructive symptoms of benign prostatic hyperplasia (BPH).
- \triangleright N.B. Prostate exhibits a high proportion of α_{1A} receptors.
- N.B. Blood vessels contain more α₁B receptors

A) Non- selective a -blockers.

- 1- Competitive blockers: Phentolamine
- 2- Non-competitive blockers: irreversible antagonist e.g., phenoxybenzamine.

<u>Uses:</u> both drugs are used for <u>short-term</u> in hypertension with pheochromocytoma (adrenaline secreting tumor of the adrenal medulla).

Adverse effects:

- Postural hypotension: Dizziness, vertigo and syncope after the initial dose.
- **2.** Marked tachycardia (overcome by concurrent use of β blocker).
- 3. Nasal congestion and pulsating headache.
- Decreased libido and sexual dysfunction.

B) Selective α₁ blockers

Prazosin, terazosin, doxazosin, and tamsulosin.

Uses:

- 1- Mild to moderate hypertension (Prazosin, terazosin can be used; however, Doxazosin is the longest acting of these drugs (excreted in feces).
- 2- Congestive heart failure as they decrease preload and afterload.
- 3- Decrease obstructive symptoms in benign prostatic hyperplasia (BPH); all can be used but alfuzosin and tamsulosin are better as more selective for α_{1A} –receptors (present on prostatic smooth m.), causing less vasodilator side effects as hypotension and headache (blood vessels contain alpha 1b receptors).

Common advantages of all selective α₁- blockers

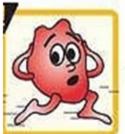
- a) Less tachycardia (compared to non selective alpha blockers).
- b) Beneficial favorable effect on plasma lipids.

Adverse effects of Selective α_1 blockers

- First dose phenomena: Dizziness, vertigo and syncope due to marked postural hypotension; 30-90 min after the initial dose avoided by using low doses and at bedtime.
- Reflex Tachycardia avoided by β blocker.
- Nasal congestion avoided by topical nasal decongestants.
- Fluid retention (prazosin) avoided by diuretic
- Decreased libido and sexual dysfunctions.



Orthostatic hypotension



Tachycardia



Vertigo



Sexual dysfunction

Selective Alpha-2 Receptor Blocker

MOA: Antagonist at α_2 -prejunctional receptors in the CNS, increasing sympathetic outflow.

Clinical uses:

- **1-Yohimbine:** treatment of postural hypotension and sexual dysfunctions induced by clonidine.
- 2- Mirtazapine: used as antidepressant drug.

N.B. ergot alkaloids (like ergotamine) has weak alpha blocking actions and a direct potent smooth muscle stimulatory actions. They produce vasoconstriction (treat acute migraine) and uterine contraction (treat postpartum hemorrhage)

β- Adrenergic Receptors Blockers

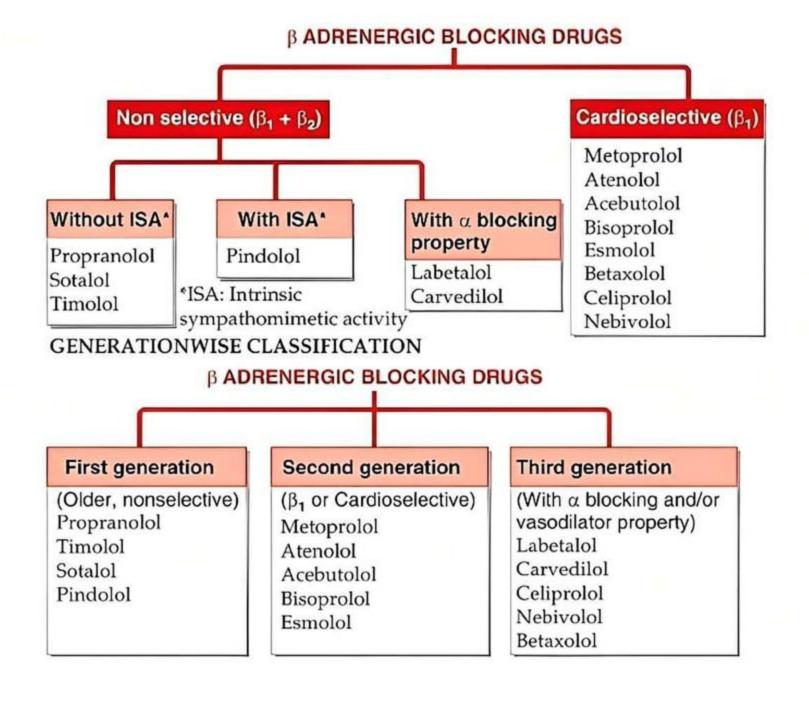
Beta 1 blocking

- 1- Decrease heart Rate
- 2- Decrease cardiac contractility
- 3- Decrease AV conduction
- 4- Decrease renin secretion from kidney

Beta 2 blocking

- 1- Bronchoconstriction
- 2- vasoconstriction
- 3- Hypoglycemia
- 4- prevent skeletal muscle tremors

Beta 3 blocking Inhibit lipolysis



1- <u>Heart</u>: Blocking β₁ results in:

- -ve Inotropic (decrease cardiac contraction)
- -ve Chronotropic (decreased heart rate)
- -ve Dromotropic (Decreased atrio-ventricular conduction)
 - $-\downarrow$ cardiac output (COP) $\rightarrow \downarrow$ BP (antihypertensive)
 - $-\downarrow$ Cardiac work & \downarrow O₂ consumption (antianginal)
 - ↓ Automaticity, & ↓ conductivity in SA and AV nodes
 - $\rightarrow \downarrow$ HR (Antiarrhythmic).

2- Blood vessels (β_2 blocking):

- Blocks V.D. effect (due to β₂ blocking)→ Unopposed α → V.C.→ ↓ blood flow to various tissues.
 - ■So, not used in vasospastic angina and peripheral vascular disease.
 - \checkmark **↓ I.O.P.**: **↓** secretion of aqueous humor in the eye.
 - ✓ Constrict blood vessels in hemangiomas
- **3-Bronchi** (β_2) \rightarrow Bronchospasm
- Not used in patients with chronic obstructive COPD or BA

- 4- Metabolic effects of BBs
- \triangleright Non-selective β -blockers:
 - **1- hypoglycemia**: blocking hepatic $\beta_2 \rightarrow \downarrow$ Glycogenolysis and augments the hypoglycemic action of insulin.
 - 2- Masking hypoglycemic reactions "symptoms" (blocking β_1); Beta blockers mask the signs of hypoglycemia (tachycardia, anxiety and tremors).
- Selective β₁ blockers: Better tolerated in diabetic patients. However, still masking hypoglycemic reactions particularly tachycardia and palpitation.
- > Beta blockers increase blood lipids.

Adverse Effects & Contraindications

1 Bronchoconstriction

Nonselective β blockers are potentially dangerous in asthma and COPD (as severe bronchospasm may cause asphyxia).

So, Non- selective BBs are contraindicated in asthmatics.

2. Rebound effects:

Never stop suddenly (can worsen angina, hypertension or precipitate arrhythmia) due to **up regulation of receptors**; withdraw gradually (1week).

- 3. Bradycardia and heart block.
- 4. Sexual impairment: impotence (psychogenic)

5. Disturbances in metabolism:

Mask manifestations of hypoglycemic reactions in diabetes and augment the hypoglycemia. They used cautiously in diabetics (especially those treated by insulin).

- 6. Worsen acute heart failure: by -ve inotropic effect.
 - 7-C.N.S: Sedation, sleep disturbance, depression, fatigue, night mares, weakness (with lipid soluble as propranolol).
- 8- Unlike alpha blockers, most beta blockers ↑ Low density lipoproteins (LDL).

1- Propranolol, Nadolol, Timolol

- Non-selective beta blocker (first generation)
- Chronic use is associated with hyperlipidemia (↑ LDL-C and/or triglycerides (↑TGs).
- They are hazardous in bronchial asthma, diabetes, peripheral vascular disease & hyperlipidemia.

1. Propranolol:

- It is the prototype of this class;
- It is highly lipid soluble so more crossing BBB and produces sedation.
- 2. Nadolol: has the longest half-life (24h) and no CNS entry.
- Timolol: is used mainly in glaucoma (topically in eyes).

2- Atenolol, Metoprolol, Bisoprolol:

- Cardioselective β-blockers (second generation)
- Less effect on blood vessels, bronchioles, and metabolism than first generation.
- Atenolol has no CNS entry

<u>Advantages:</u>

Safer in asthma, diabetes, and peripheral vascular disease.

3- Pindolol and acebutolol

- BBs with <u>intrinsic sympathomimetic activity</u> (ISA)
- Less bradycardia compared to propranolol.
- Most suitable BBs for patients with bradycardia or heart block
- Used in <u>hypertensive patients with moderate</u> <u>bradycardia</u>

4- Carvedilol and labetalol

 Have combined α₁ & β-blocking activity (third generation-non selective Vasodilator BBs).

<u>Labetalol :</u>

- It is used I.V. in <u>hypertensive emergencies</u> and orally in <u>chronic hypertension</u>.
- It is safe to be used in severe hypertension of pregnancy

Carvedilol:

- It has an additional antioxidant property
- Useful in chronic stabilized heart failure.

5- Esmolol

- It is <u>ultra-short acting</u> BB
- Rapidly metabolized by esterases of RBCs with t_{1/2} 10 minutes
- Useful in arrhythmias associated with cardiac surgery.
- **6- Nebivolol** Highly selective β_1 and α_1 receptor blockers of 3rd generation BB.
- Causes vasodilation through increasing the production of <u>endothelial NO</u>.
- It is used whenever the <u>erectile sexual</u> <u>dysfunction</u> is associated with cardiac disease because it improves both conditions (e.g., hypertension and erectile dysfunction).

Therapeutic uses of BBs

1- Cardiovascular uses

- A. Mild-moderate hypertension: all BB except esmolol (ultrashort).
- B. Chronic stable angina: All except BBs with ISA.
- C. Supraventricular arrhythmias: Correct and treat only tachycardia; protect the ventricle.
- D. Prophylactic after myocardial infarction: to decrease the incidence of sudden death. (Propranolol, metoprolol or timolol are used).
- E. Chronic heart failure: Low dose of carvedilol, metoprolol or bisoprolol can decrease the mortality rate in patients with heart failure.
 - Carvidilol is preferred because it has <u>antiarrhythmic</u>, <u>antiremodeling</u> and <u>antioxidant</u> effects.

2- Endocrinal uses:

A. <u>Hyperthyroidism</u>: propranolol is used to control tachycardia & inhibits the conversion of T₄ to T₃ which is the most active form.

B. <u>Pheochromocytoma</u>: To control the associated arrhythmia, but it must be given after α-blockers to avoid marked increase in BP.

3- Hepatic uses:

Prevention of **oesophageal variceal** bleeding in patients with portal hypertension caused by liver cirrhosis (Carvedilol and propranolol can be used).

- 4- Neurological uses: Propranolol can be used in the following conditions.
- 1. Tremors
- 2. Alcohol withdrawal manifestations (to decrease anxiety).
- 3. Acute panic symptoms with tachycardia.
- 4. Anxiety associated with psychosomatic disorders.
- 5. Prophylactic uses in migraine.

5- Topical uses:

- Eye drops timolol, betaxolol and carteolol reduces intraocular pressure and are used in open angle glaucoma.
- <u>Note</u>: sufficient amount of BBs may be absorbed from the eye to cause serious systemic adverse effects.
- Topical propranolol cream can be used for treatment of infantile haemangioma (as it constricts blood vessels).

Characteristics of Some Beta Blockers

Drugs	β 1-Selective	ISA	Sedation	Blood Lipids
Acebutolol	+	++	+	_
Atenolol	+	_	-	↑ ↑
Metoprolol	+	_	+	↑ ↑
Pindolol	_	++	+	_
Propranolol	_	-	+++	↑ ↑
Timolol	_	_	++	↑ ↑

