Definition of Alpha and Beta Blockers

**Alpha blockers** generally help relax the muscles, which in turn can lead to the opening of blood vessels for smooth circulation. Alpha blockers work by keeping the hormones of norepinephrine or noradrenaline at bay, leading to smoother blood flow through open veins.

**Beta blockers** work by blocking the hormone called epinephrine (better known as adrenaline). This hormone often causes increased heart rate, which can lead to increased blood pressure levels. Beta blockers prevent this from happening by reducing the heart rate, thereby reducing blood flow. Blood pressure is decreased because of the dilation of the blood vessels.

Classification of Alpha and Beta Blockers

Adrenoceptor blockers are classified based upon their selectivity toward adrenoreceptors.
Alpha blockers

**Sympathetic drugs in hypertension**

- **α agonists**
  - **α₁ agonists**
    - Phenylephrine (causes hypertension)
  - **α₂ agonists**
    - α-methyldopa
    - Clonidine

  - **Non-selective**: phenoxybenzamine, phentolamine
  - **α₁-selective**: prazosin, terazosin, doxazosin, alfuzosin, indoramin, urapidil, bunazosin, tamsulosin
  - **α₂-selective**: yohimbine

<table>
<thead>
<tr>
<th><strong>α-Methyldopa (Aldomet®)</strong></th>
<th><strong>Clonidine</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>Decrease central sympathetic outflow</td>
<td>Decrease central sympathetic outflow</td>
</tr>
<tr>
<td>Prodrug; metabolized to methylnorepinephrine</td>
<td></td>
</tr>
<tr>
<td>Decreases central sympathetic outflow, cardiac output, and vascular resistance</td>
<td>Decreases central sympathetic outflow, cardiac output, and vascular resistance</td>
</tr>
<tr>
<td>Compensatory reaction: salt retention</td>
<td>Compensatory reaction: salt retention</td>
</tr>
</tbody>
</table>
| **Idiosyncratic reaction:**  
  - Hematologic immunotoxicity (positive Coombs test) → hemolytic anemia  
  - Sedation | **Idiosyncratic reaction:**  
  - Rebound hypertension if discontinued (restart it, or use phentolamine, an alpha blocker)  
  - Sedation |
| Previously used extensively for pregnancy; second most common | Not used in pregnancy |

Beta Blockers
- **Non-selective**: nadolol, penbutolol, pindolol, propranolol, timolol, sotalol, metoprolol, carteolol, carvedilol*, labetalol*

- **β₁-selective**: acebutolol, atenolol, bisoprolol, esmolol, metoprolol, betaxolol, nebivolol

- **β₂-selective**: butoxamine

### Notable β-Blocker

<table>
<thead>
<tr>
<th></th>
<th>Propranolol</th>
<th>Metoprolol</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Prototypical β-blocker</strong></td>
<td>• Short-acting, poor BP control</td>
<td>• Prototypical cardiac β-blocker</td>
</tr>
<tr>
<td><strong>Used in anxiety and stage fright also</strong></td>
<td>• Can be used to fool a lie detector test!</td>
<td>• Used twice daily</td>
</tr>
<tr>
<td><strong>β₁ selective</strong></td>
<td>• Most commonly used in MI period</td>
<td>• More β₁ selective, less BP control</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>Labetalol</th>
<th>Bisoprolol</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Nonselective third generation</strong></td>
<td>• Wide therapeutic margin (200—2400 mg/day)</td>
<td>• Once daily β-blocker</td>
</tr>
<tr>
<td><strong>Excellent BP control</strong></td>
<td>• Most used BP med in pregnancy</td>
<td>• More β₁ selective, used post-MI</td>
</tr>
<tr>
<td><strong>Caution: Endothelial dysfunction</strong></td>
<td></td>
<td>• Good BP control</td>
</tr>
</tbody>
</table>

### β-Blockers with additional activity

<table>
<thead>
<tr>
<th></th>
<th>Nebivolol</th>
<th>Carvedilol</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>'Novel' third generation selective β-blocker</strong></td>
<td>• β₁ selective</td>
<td>• Nonselective third generation β-blocker</td>
</tr>
<tr>
<td><strong>Also, has nitric oxide activity — direct vasodilator</strong></td>
<td>• Excellent BP control</td>
<td>• Also, has alpha activity</td>
</tr>
<tr>
<td><strong>Caution: Endothelial dysfunction</strong></td>
<td></td>
<td>• Used in heart failure</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Poor BP control</td>
</tr>
</tbody>
</table>

**Note:**

- The names of α-blockers generally end with -in, while those of β-blockers generally end with –olol.

- Phenoxybenzamine is a long-acting irreversible α-blocker.

- Phentolamine is a short-acting reversible α-blocker.

- In the sympathetic nervous system, the transmitter in effector organs is norepinephrine, while, in the parasympathetic nervous system, the transmitter in effector organs is acetylcholine (Ach). Alpha and beta blockers have an antagonistic action on the sympathetic nervous system.
Adrenoceptors – Alpha and Beta Receptors

**α1 Receptors**
- Location: at the gastrointestinal tract and bladder sphincter, vascular smooth muscles of skin and splanchnic regions, and radial muscle of iris
- Function: generally produce smooth muscle constriction
- Mechanism of action: act via stimulation of IP/$^{3}$/Ca$^{3+}$

**α2 Receptors**
- Present in presynaptic nerve terminals, platelets, fat cells, and the wall of the gastrointestinal tract.
- Function: generally produce relaxation/dilation.
- Mechanism of action: act via inhibition of adenylate cyclase and decreasing the concentration of cAMP (cyclic adenosine monophosphate).

**β1 Receptors**
- Location: sinoatrial node, atrioventricular node, atrial and ventricular muscle, His-Purkinje system, and kidney.
- Mechanism of action: act via stimulation of adenylate cyclase and thereby increasing the concentration of cAMP (cyclic adenosine monophosphate).

**β2 Receptors**
- Location: smooth vessels of skeletal muscle, blood vessels, gastrointestinal tract, uterus, liver, and urinary tract
- Mechanism of action: act via stimulation of adenylate cyclase and increasing the concentration of cAMP.

Effect of Adrenoceptors on Organ Systems

Understanding the action of alpha and beta receptors on various organ systems will aid in remembering the effect of alpha and beta blockers on those organ systems because they have opposite actions to the alpha/beta-agonists.

<table>
<thead>
<tr>
<th>Receptor/organ system</th>
<th>Actions</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>α1 receptors</strong></td>
<td></td>
</tr>
<tr>
<td>Eyes</td>
<td>Contraction (mydriasis) of the iris dilator muscle.</td>
</tr>
<tr>
<td>Bladder</td>
<td>Constriction of bladder sphincter, Control of micturition and urine flow. Note: α-blockers increase the urine flow by promoting the relaxation of the bladder muscles.</td>
</tr>
<tr>
<td>Prostate</td>
<td>Cause ejaculation by prostate contraction. α-blockers are used to treat benign prostatic hyperplasia (BPH) induced urinary obstructions because it causes the relaxation of the bladder muscles (the opposite actions to the alpha agonists). α-blockers also produce impaired ejaculation due to their α-receptor antagonism.</td>
</tr>
<tr>
<td>Kidney</td>
<td>Decrease renin secretion.</td>
</tr>
<tr>
<td>Veins and arterioles (skin)</td>
<td>Contraction of smooth muscles of the peripheral blood vessels.</td>
</tr>
<tr>
<td><strong>α2 receptors</strong></td>
<td></td>
</tr>
<tr>
<td>Platelets</td>
<td>Increase the platelet aggregability.</td>
</tr>
<tr>
<td>β1 receptors</td>
<td>Heart</td>
</tr>
<tr>
<td>--------------</td>
<td>-------</td>
</tr>
<tr>
<td>β2 receptors</td>
<td>Veins and arterioles</td>
</tr>
<tr>
<td></td>
<td>Bladder</td>
</tr>
<tr>
<td></td>
<td>Bronchioles</td>
</tr>
<tr>
<td></td>
<td>Kidney</td>
</tr>
<tr>
<td></td>
<td>Liver</td>
</tr>
</tbody>
</table>

### Alpha Adrenergic Blocking Agents

These drugs block the action of alpha-adrenoceptors. They are commonly used in the treatment and management of **hypertension** and **benign prostatic hyperplasia** (BPH).

- **Phenoxybenzamine** and **phentolamine** are two nonselective alpha-adrenergic blocking agents. Thus, they act as both α1 and α2 receptors.
- **Phenoxybenzamine** binds covalently with the adrenergic receptors (irreversible and non-competitive). Due to irreversible binding, phenoxybenzamine has a longer duration of action. It decreases blood pressure by preventing constriction of the peripheral blood vessels. However, due to increased cardiac output, phenoxybenzamine does not cause a prolonged drop in blood pressure, so it is not widely used for this purpose.
- **Phentolamine** is a reversible and competitive type of alpha-adrenergic blocking agents. It has a shorter duration of action and it is used in the treatment of **pheochromocytoma**.
- **Prazosin**, **terazosin**, **doxazosin**, and **tamsulosin** have selective antagonistic action on α1 receptors.
- **Prazosin** has 1,000 times more selectivity action on α1 receptors. Due to selectivity in action, marked orthostatic hypotension and tachycardia are generally observed with nonselective alpha-adrenergic blocking agents such as phenoxybenzamine and phentolamine.
- **Prazosin** is an important drug in the treatment of hypertension and BPH. Terazosin and doxazosin have similar actions. Terazosin has a higher bioavailability (80%) than prazosin (50%).
- Postural hypotension is not observed with **tamsulosin**.
- Before the discovery of phosphodiesterase-5-inhibitors such as sildenafil, **yohimbine** was used to treat impotence (erectile dysfunction) in men, but safer and effective alternatives are now available.

**Alpha-blockers (both selective and non-selective) are not recommended as monotherapy in hypertension due to the availability of other effective antihypertensives.**

### Adverse Effects of Alpha Adrenergic Blocking Agents

Adverse effects of alpha blockers are mainly due to their antagonistic action/blocking effects on alpha receptors.
- **Orthostatic hypotension**: This results from the pooling of blood in the veins of the legs. Fainting can also result due to the reduced supply of blood to the brain. This is the most common side effect of alpha-blockers. It is more common with the use of nonselective than -selective alpha blockers.
- **Dizziness and headache.**
- **Reflex tachycardia**: Increased heart rate due to the stimulation of baroreceptors.
- **Nasal stiffness**: Occurs due to alpha-receptor blockage.
- Since alpha receptors have a role in the contraction of smooth muscle of the prostate, which induces ejaculation in males, blockage of alpha receptors inhibits the **ejaculation process** in males.

### Beta-Adrenergic Blocking Agents

All beta blockers or beta-adrenoceptor blockers antagonize the action of beta receptors.

Understanding the action of **beta blockers** on various organ systems will aid in remembering the effect of beta-agonists on those organ systems because they have opposite actions to the beta agonists.

**Note:**
- **Cardioselective β blockers** (atenolol, metoprolol, acebutolol, esmolol, bisoprolol, betaxolol) have a selective action on **β1 receptors**.
- **Cardioselective β blockers** are safer to use in patients with **asthma** (as they don’t cause bronchoconstriction), **diabetes**, and peripheral vascular disease.
- Beta blockers do not cause postural hypotension as they do not have any action on alpha receptors.
- Beta blockers act by reducing **cardiac output** (volume of blood pumped by the heart per minute), thereby reducing blood pressure.

### Effects of Beta-Adrenergic Blocking Agents

**Effect on Cardiovascular System**

- Decreases **cardiac output**
- Decreases heart rate (produce **bradycardia**)
- Decreases force of contraction
- Decreases **total peripheral resistance**
- Negative chronotropic and inotropic actions
- **Decreases renin release** from the kidneys, which is thought to be their mechanism of action in reducing blood pressure

*Always remember:*

**Note:**

- Action by beta blockers on β2 receptors is considered to be undesired because nonselective beta blockers cause bronchoconstriction and decrease insulin secretion and glycogenolysis.
- **Cardioselective beta-blockers** always act on β1 receptors.

**Effect on Pulmonary System**

Nonselective beta-blockers such as propranolol can produce **bronchoconstriction** or
exacerbate **asthma** in asthmatics. Due to this, propranolol should be avoided in **asthmatics** and patients suffering from **COPD**.

- Beta-blockers always show effects opposite to beta agonists.
- β1 agonists such as salmeterol and salbutamol have a **bronchodilatory effect** on the **lungs**.

**Effects on the Eyes**

Beta-blockers reduce **intraocular pressure**. Thus, they are used to treat glaucoma. They also reduce the production of aqueous humor in the eyes (timolol).

**Metabolic effects**

Beta-blockers increase the levels of insulin in the body, so if given to a diabetic patient who is on insulin therapy, drastic hypotension may result. Thus, beta blockers are **contraindicated in diabetes**.

Beta-blockers also block **glycogenolysis** and **gluconeogenesis**.

**Clinical uses of beta-adrenergic blocking agents**

- **Hypertension** (carvedilol, labetalol, propranolol)
- **Angina pectoris** (propranolol)
- **Myocardial infarction** (propranolol and esmolol)
- Glaucoma (timolol, betaxolol, carteolol – applied topically)
- **Migraine** (propranolol)
- Performance anxiety (propranolol)
- **Hyperthyroidism** (propranolol)

**Adverse effects and toxicity of beta blockers**

- **Bradycardia**
- **AV blockage**
- Severe asthma attacks (propranolol)
- **Hypoglycemia**
- **Arrhythmias** (upon abrupt stoppage of therapy with beta blockers)
- Sexual dysfunction (propranolol)
- Fatigue
- Vivid dreams (propranolol)

The Renin-Angiotensin-Aldosterone System

Blockers of the renin-angiotensin-aldosterone system

Beta-blockers control and regulate blood pressure. The kidney and the central nervous system are the critical components of this action. Peripheral baroreceptors and the autonomic nervous system also play important roles.

Medical therapy for MI patients: ACE inhibitors (ACEI) and angiotensin receptor blockers (ARBs)

ACE inhibitors and ARBs decrease blood pressure and decrease the work of the heart by dilating arteries. Side effects include cough, dizziness, and low blood pressure. Pregnancy is contraindicated.

The kidney is crucial to blood pressure regulation via the juxtaglomerular apparatus and renin release. Renin initiates a biochemical sequence that eventually converts angiotensinogen, which is produced in the liver, into angiotensin, a strong vasoconstrictor. Angiotensin stimulates the release of aldosterone from the adrenal gland, which causes the kidney to retain salt (NaCl) and water. Angiotensin stimulates the release of antidiuretic hormone from the pituitary gland, which causes the kidney to retain water.

CE inhibitors may lower blood pressure too much and cause allergic reactions. This system is part of the body’s defense against dehydration and/or blood loss. Blood volume should be restored to normal as quickly as possible.

<table>
<thead>
<tr>
<th>Commonly used ACEI and All blockers</th>
<th>Initial daily dose(s)</th>
<th>Target dose</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Captopril</strong></td>
<td>6.25 mg tid</td>
<td>50 mg tid</td>
</tr>
<tr>
<td><strong>Enalapril</strong></td>
<td>2.5 mg bid</td>
<td>10—20 mg bid</td>
</tr>
<tr>
<td>Fosinopril</td>
<td>5—10 mg daily</td>
<td>40 mg daily</td>
</tr>
<tr>
<td><strong>Lisinopril</strong></td>
<td>2.5—5 mg daily</td>
<td>20—40 mg daily</td>
</tr>
<tr>
<td>Perindopril</td>
<td>2 mg daily</td>
<td>8—16 mg daily</td>
</tr>
<tr>
<td>Quinapril</td>
<td>5 mg bid</td>
<td>20 mg bid</td>
</tr>
<tr>
<td></td>
<td><strong>Ramipril</strong></td>
<td><strong>Trandolapril</strong></td>
</tr>
<tr>
<td>----------------</td>
<td>--------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>Dose</td>
<td>1.25—2.5 mg daily</td>
<td>1 mg daily</td>
</tr>
<tr>
<td>Dose</td>
<td>10 mg daily</td>
<td>4 mg daily</td>
</tr>
</tbody>
</table>

**AE`s with ACEI and ARB — first-line Rx**

<table>
<thead>
<tr>
<th>ACEI's</th>
<th>ARB's</th>
</tr>
</thead>
<tbody>
<tr>
<td>Kidney damage especially in individuals with prior damage</td>
<td></td>
</tr>
<tr>
<td>Excessive drop in blood pressure</td>
<td></td>
</tr>
<tr>
<td>Tongue and facial swelling due to allergic reaction</td>
<td></td>
</tr>
</tbody>
</table>

**Summary**

1. Alpha blockers work on the blood muscles to open up the blood vessels, while the beta blockers work on the heart to ease the flow of blood.
2. Alpha blockers work on norepinephrine or noradrenaline, while beta blockers work on epinephrine or adrenaline.
3. Alpha blockers affect only blood pressure levels, while beta blockers affect both the heart and blood pressure.
4. Beta blockers can cause weight gain, while alpha blockers do not.

**References**


**Legal Note:** Unless otherwise stated, all rights reserved by Lecturio GmbH. For further legal regulations see our [legal information page.](#)