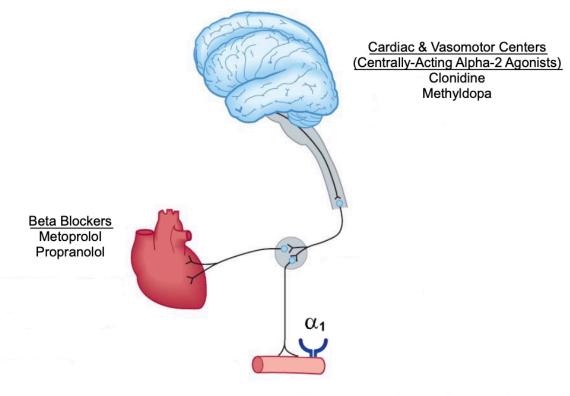
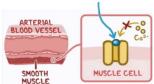
Antihypertensive Agents

Sites of Action of Major Antihypertensive Agents



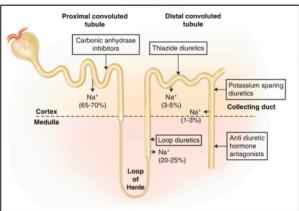
Calcium Channel Blockers
Nifedipine
Amlodipine



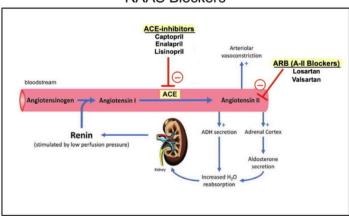
Alpha-1 Blockers Terazosin Prazosin Direct-Acting Vasodilators
Hydralazine
Minoxidil



Diuretics



RAAS Blockers



Antihypertensive Agents

I. Diuretics

• MOA: diuretics increase urine volume \rightarrow decrease blood volume \rightarrow decrease blood pressure.

A. Thiazide Diuretics

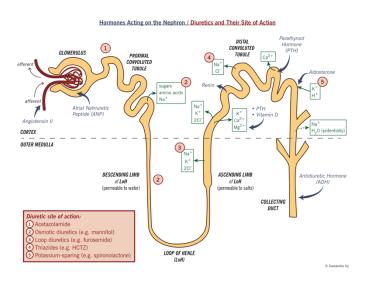
- Hydrochlorothiazide (HCTZ), chlorthalidone (Diuril), indapamide (Lozol), and metolazone (Zaroxolyn) are thiazide diuretics.
- MOA: Thiazide diuretics block Na/H₂O reabsorption in the distal tubule, which accounts for approx. 5-10% of the Na/H₂O reabsorption by the nephron.
- SEs: hypokalemia, hyponatremia, hypomagnesemia, hypercalcemia, hyperglycemia, hyperuricemia, and dehydration.
- Use with caution in patients with a history of gout unless patient is on uric acid-lowering agents, e.g., allopurinol (Zyloprim) 300 mg PO daily.

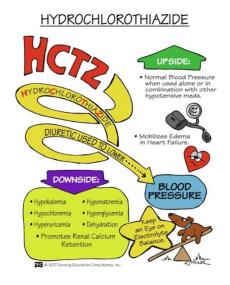
B. Loop Diuretics

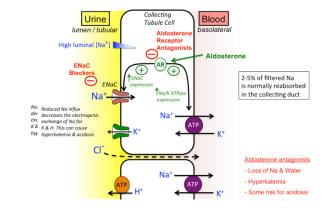
- MOA: Furosemide (Lasix), bumetanide (Bumex), and torsemide (Demadex) block Na/H₂O reabsorption in the ascending Loop of Henle, which accounts for 20-25% of Na/H₂O reabsorption by the nephron.
- Bumetanide is 40 times more potent than furosemide; so,
 40 mg of furosemide = 1 mg of bumetanide.
- Loop diuretics are preferred in patients with symptomatic heart failure (HF) and in patients with moderate-severe CKD (i.e., CrCl < 30 ml/min).

C. Potassium Sparing Diuretics

- Triamterene (Dyrenium), amiloride (Midamor), spironolactone (Aldactone), and Eplerenone (Inspra) are K-sparing diuretics.
- Spironolactone and eplerenone are also aldosterone antagonists.
- MOA: K-sparing diuretics block Na⁺ channels
 (ENaC) in the late distal tubule → inhibits Na⁺
 reabsorption from the lumen → reduces
 intracellular Na⁺ in tubular epithelial cells →
 decreases function of the Na⁺/K⁺ ATPase pump → retains K⁺ and H⁺ ion.







- D. Side Effects of Thiazide and Loop Diuretics
 - Hypokalemia may be prevented or treated with K-sparing diuretics or KCL supplementation.
 - (a) K-sparing diuretics may be added to a thiazide or a Loop diuretic to prevent K⁺ depletion.
 - Dyazide and Maxzide-25 are brand name combination products containing HCTZ 25

Brana name combination	
products containing HCTZ 25	
mg and triamterene 37.5 mg, dos	sed once daily.

- | K+, Na+, Mg2 †Uric acid and can trigger gout attack Thiazides Hydrochlorothiazide • †LDL, total cholesterol and total triglycerides Thiazide-like Chlorthalidone, Indapamide ↓K+, Na+, Mg2+, Ca2+ Furosemide, Torsemide, Ototoxicity Loop diuretics Bumetanide Hyperuricemia Dizziness Hyperkalemia Potassium sparing Spironolactone, Eplerenone, Hyponatremia Gynecomastia (Spironolactone) Triamterene, Amiloride Weakness, fatigue, dizziness
- (b) KCL supplements may be separately dosed with a thiazide or a Loop diuretic to prevent and treat hypokalemia.
 - K-Dur 20 mEq SR (sustained-release) tablets.
 - KCl 10% liquid 20 mEq for immediate release and absorption.
 - KCl IV infusions (KCL riders): KCL 40 mEq in 500 ml NS IVPB, infused over 4 hours.
- (2) Hypomagnesemia may be prevented or treated with magnesium supplementation.
 - Slo-Mag 84 mg SR (sustained-release) tablets.
 - Magnesium oxide 400 mg immediate-release tablets.
 - Magnesium Sulfate IV infusion: 1-2 grams IVPB, infused over 1-2 hours.
- (3) Hypocalcemia may be prevented or treated with calcium supplementation.
 - Note: Thiazide diuretics are calcium-sparing.
 - Calcium carbonate (TUMS) 800-1200 mg PO daily.
 - Calcium gluconate 1-2 grams IVPB, infused over 1-2 hours.

II. Beta Blockers

- MOA: Beta blockers (BB) block beta-1 receptors on the SA node, AV node, atria, and ventricles of the heart
 → reduce heart rate (SA node), reduce conduction velocity (AV node), and decrease myocardial contractility (atria, ventricles) → decrease blood pressure.
- A. Selective Beta-1 Blockers: Metoprolol (Lopressor, Toprol XL) and Atenolol (Tenormin), and intravenous Esmolol (Brevibloc).
- Mechanism of the β₁ Adrenoceptor

 Exterior

 Call Membrane

 T Hour rate

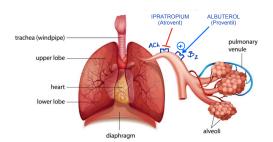
 T All rode conduction

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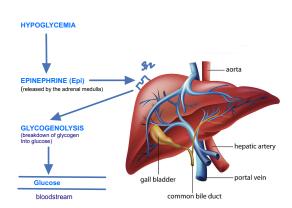
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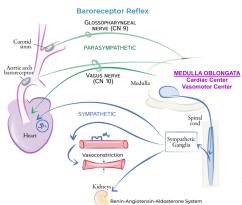
 T Contractility
- Selective Beta-1 blockers are also called "cardioselective" beta blockers.
- B. Non-Selective Beta Blocker: Propranolol (Inderal)
 - Avoid propranolol in patients with COPD/asthma, since propranolol blocks beta-2 receptors in the airways and competes with albuterol (beta-2 agonist) for beta-2 receptor sites in the bronchioles
 → induces bronchoconstriction.



- B. Non-Selective Beta Blocker: Propranolol (cont.)
 - Propranolol also blocks beta-2 receptors in the liver → blocks glycogenolysis → prevents glucose replacement in diabetics during hypoglycemic episodes.
 - During hypoglycemic episodes, epinephrine (Epi) is released into the bloodstream by the adrenal medulla to stimulate beta-2 receptors in the liver to initiate glycogenolysis. Non-selective beta blockers block glycogenolysis and prevent glucose replacement during hypoglycemic episodes in diabetics.
 - Cautionary Note: All beta blockers (i.e., selective and non-selective) will mask the signs and symptoms of hypoglycemia in diabetics.



- III. Alpha-1 Blockers: Prazosin (Minipress), Doxazosin (Cardura), and Terazosin (Hytrin).
 - MOA: Terazosin blocks alpha-1 receptors on blood vessels
 → vasodilation → decreases BP.
 - SE: orthostatic hypotension → vertigo → syncope.
 - Terazosin is dosed once daily at bedtime (QHS) to prevent orthostatic hypotension.
 - Terazosin is also indicated for BPH (benign prostatic hyperplasia).

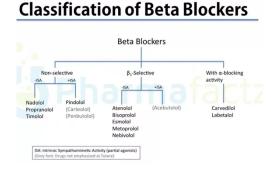


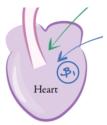
IV. Alpha-Beta Blockers: Labetalol (Trandate) and Carvedilol (Coreg).

 MOA: Labetalol (Trandate) → blocks alpha-1 receptors (blood vessels) and blocks beta-1 & beta-2 receptors.





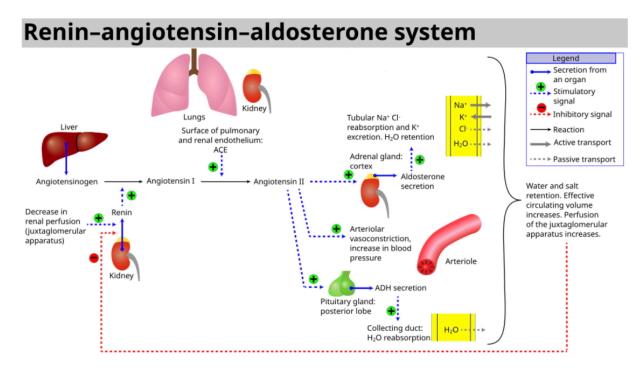




- → blocks beta-1 (SA node) → decreases HR
- → blocks beta-1 (AV node) → decreases conduction velocity
- → blocks beta-1 (atria, ventricles) → decreases contractility

→ decreases BP

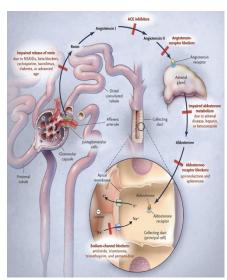
- V. ACE-Inhibitors (ACEi): Captopril (Capoten), Enalapril (Vasotec), and Lisinopril (Prinivil).
 - MOA: ACEi's inhibit ACE → reduce conversion of A-I to A-II → reduce production of A-II
 - → reduce vasoconstriction → reduce BP
 - \rightarrow reduce aldosterone secretion \rightarrow reduce Na/H₂O reabsorption (distal tubule) \rightarrow reduce BP
 - \rightarrow reduce ADH (vasopressin) secretion \rightarrow reduce H₂O reabsorption (collecting duct) \rightarrow reduce BP



- SEs: (1) hyperkalemia, (2) cough, and (3) angioedema.
- Caution: ACEi's are contraindicated in pregnancy.

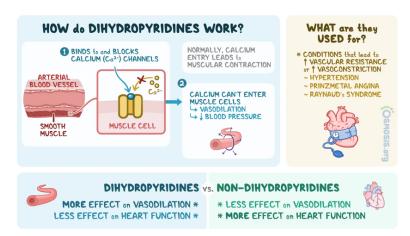
VI. Angiotensin-II Receptor Blockers (ARB): Losartan (Cozaar) and Valsartan (Diovan).

- MOA: ARBs block A-II receptor sites on blood vessels, adrenal cortex, and posterior pituitary gland.
 - → reduce vasoconstriction → reduce BP
 - → reduce aldosterone secretion → reduce Na/H₂O reabsorption (distal tubule) → reduce BP
 - → reduce ADH secretion → reduce H₂O reabsorption (collecting duct) → reduce BP
 - SEs: (1) hyperkalemia, (2) cough, and (3) angioedema.
 Note: The incidence of cough and angioedema is significantly less frequent with ARBs than with ACEi's.
 - When switching from an ACEi to an ARB due to cough or angioedema, allow a 6-week washout period before starting an ARB.
 - Caution: ARBs and ACEi's are contraindicated in pregnancy.



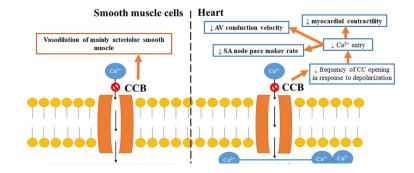
VII. Calcium Channel Blockers (CCB)

- A. Dihydropyridines: Nifedipine (Procardia), Amlodipine (Norvasc), and intravenous Nicardipine (Cardene).
 - MOA: Nifedipine blocks calcium influx into vascular smooth muscle
 → vasodilation → decreases BP.
 - Nifedipine may cause reflex tachycardia (+++) in response to baroreceptor stimulation in the aortic bodies and carotid sinuses.



B. Non-Dihydropyridines: Diltiazem (Cardizem) and Verapamil (Calan)

- MOA: Non-dihydropyridine CCBs block calcium influx into cardiac muscle → decrease myocardial contractility → decrease BP.
- MOA: Non-dihydropyridine CCBs block calcium influx into nodal tissue → decrease HR (SA node) and decrease conduction velocity (AV node) → decrease BP.



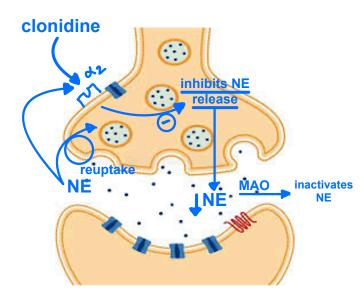
- Since non-dihydropyridine CCBs inhibit nodal tissue, they may also be used to treat atrial fibrillation and SVT (supraventricular tachycardia).
- Non-dihydropyridine CCBs, especially verapamil, should be used with caution in patients with heart failure since they may reduce contractility in an already "weakened heart."
- Nifedipine has the greatest potency (+++) for vasodilation and reflex tachycardia.
- Verapamil has the greatest potency (+++) for AV blocking effect and causing a negative inotropic effect (i.e., decreased contractility) on the heart.

EFFECTS	Nifedipine (Procardia)	Diltiazem (Cardizem)	Verapamil (Calan)
Vasodilation	(+++)	(+)	(+/-)
Reflex Tachycardia	(+++)	(+)	0
Negative Inotropic	0	(+)	(+++)

VIII. Centrally-Acting Alpha-2 Agonists: Clonidine (Catapres) and Methyldopa (Aldomet).

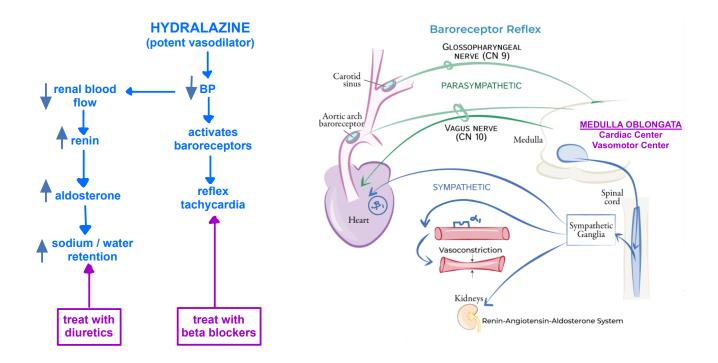
- MOA: Clonidine stimulates presynaptic alpha-2 receptors in the cardiac and vasomotor centers → decreases NE release
 - → decreases sympathetic outflow to the heart (decreases HR, decreases conduction velocity, decreases myocardial force of contraction) and decreases sympathetic to blood vessels (vasodilation)
 - → decreases blood pressure.
- SE: sedation / drowsiness

MEDULLA OBLONGATA			
ARAS Cardiac			
(consciousness)	Center		
Respiratory	Vasomotor		
Center	Center		



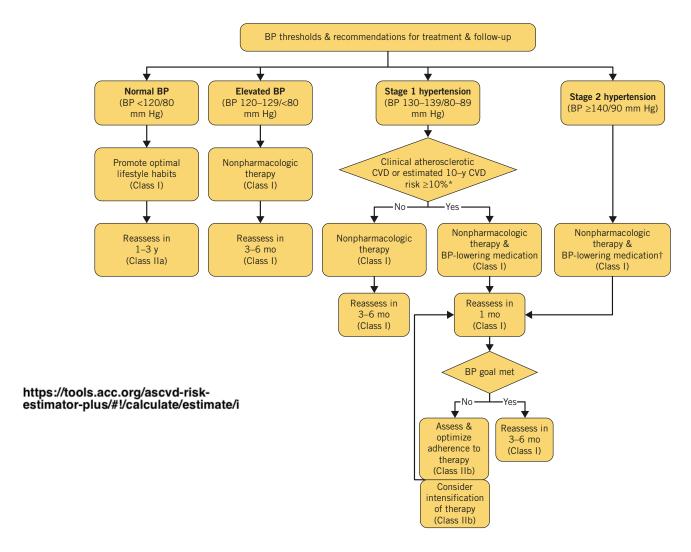
IX. Direct-Acting Vasodilators: Hydralazine (Apresoline) and Minoxidil (Loniten).

- Hydralazine is a 3rd or 4th-line agent in the stepped-care treatment of hypertension, since direct-acting vasodilators are very potent vasodilators.
- SEs: reflex tachycardia and Na/H₂O retention.



ACC/AHA: Clinical Practice Guidelines (2017)

TABLE 1. Comparing BP classifications ^{4,7}						
If the patient's systolic and diastolic BPs fall into different categories, classify the patient's hypertension according to the highest category.						
Systolic BP (mm Hg) Diastolic BP (mm Hg) 2017 guideline JNC 7						
<120	<80	Normal	Normal			
120-129	<80	Elevated	Dueleumentemeire			
130-139	80-89	Stage 1 hypertension	- Prehypertension			
140-159	90-99	Ct 2 hti	Stage 1 hypertension			
≥160	≥100	Stage 2 hypertension	Stage 2 hypertension			



INITIAL TREATMENT RECOMMENDATIONS

- In the absence of specific compelling indications: ACE-I or ARB, CCB, and thiazide diuretic.
- General non-black population, including those with diabetes, initial pharm treatment should include: ACE-I or ARB, CCB, and thiazide diuretic.
- General black population, initial treatment should include: CCB and thiazide diuretic.
- All patients with CKD and HTN, initial tx should include: ACE-I or ARB → improve kidney outcomes
- In all hypertensive patients, if goal BP is not reached within a month of initiating treatment, you may
 (1) increase the dose of the initial drug <u>OR</u> (2) add a 2nd drug from a different class <u>OR</u>
 (3) discontinue 1st drug and select a drug from a different class.

Considerations for individualizing antihypertensive therapy

Indication or contraindication	Antihypertensive drugs						
Compelling indications (major improvement in outcome independent of blood pressure)							
Heart failure with reduced ejection fraction	ACE inhibitor or ARB, beta blocker, diuretic, aldosterone antagonist*						
Postmyocardial infarction	ACE inhibitor or ARB, beta blocker, aldosterone antagonist						
Proteinuric chronic kidney disease	ACE inhibitor or ARB						
Angina pectoris	Beta blocker, calcium channel blocker						
Atrial fibrillation rate control	Beta blocker, nondihydropyridine calcium channel blocker						
Atrial flutter rate control	Beta blocker, nondihydropyridine calcium channel blocker						
Likely to have a favorable	effect on symptoms in comorbid conditions						
Benign prostatic hyperplasia	Alpha blocker						
Essential tremor	Beta blocker (noncardioselective)						
Hyperthyroidism	Beta blocker						
Migraine	Beta blocker, calcium channel blocker						
Osteoporosis	Thiazide diuretic						
Raynaud phenomenon	Dihydropyridine calcium channel blocker						
Contraindications							
Angioedema	Do not use an ACE inhibitor						
Bronchospastic disease	Do not use a non-selective beta blocker						
Liver disease	Do not use methyldopa						
Pregnancy (or at risk for)	Do not use an ACE inhibitor, ARB, or renin inhibitor (eg, aliskiren)						
Second- or third-degree heart block	Do not use a beta blocker, nondihydropyridine calcium channel blocker unless a functioning ventricular pacemaker						
Drug classes that may hav	re adverse effects on comorbid conditions						
Depression	Generally avoid beta blocker, central alpha-2 agonist						
Gout	Generally avoid loop or thiazide diuretic						
Hyperkalemia	Generally avoid aldosterone antagonist, ACE inhibitor, ARB, renin inhibitor						
Hyponatremia	Generally avoid thiazide diuretic						
Renovascular disease	Generally avoid ACE inhibitor, ARB, or renin inhibitor						

<u>Antihypertensives in Pregnancy</u> (UpToDate)

Drug	Class	Initial dose	Usual effective dose range	Maximum suggested total daily dose	Comments
Labetalol	Combined alpha and beta blocker	100 mg 2 times daily, increase by 100 mg twice daily every 2 to 3 days as needed	200 to 800 mg in 2 divided doses	2400 mg	Can cause bronchoconstriction. Avoid in patients with asthma, chronic obstructive lung disease, heart failure, bradycardia (heart rate <60 beats per minute), or greater than first-degree heart block. The dosing interval can be increased to 3 times daily if blood pressure is increased prior to the next prescribed dose.
Hydralazine NOTE: Due to reflex tachycardia, monotherapy with oral hydralazine is not recommended; hydralazine may be combined with methyldopa or labetalol if needed as add-on therapy	Peripheral vasodilator	Begin with 10 mg 4 times per day, increase by 10 to 25 mg/dose every 2 to 5 days	50 to 100 mg in 2 to 4 divided doses	200 mg*	
Nifedipine extended release (ER) [¶]	Calcium channel blocker	30 to 60 mg once daily as an extended release tablet, increase at 7 to 14 day intervals	30 to 90 mg once daily	120 mg	Do not administer sublingually. Based upon clinical experience of UpToDate contributors, some patients better tolerate nifedipine ER administered in 2 divided doses, which may serve to minimize its peak to trough effects (eg, instead of increasing the dose to 60 mg once daily, it may be desirable in some patients to increase to 30 mg 2 times daily).
Methyldopa	Centrally acting alpha agonist	250 mg 2 to 3 times daily, increase every 2 days as needed $^{\Delta}$	250 to 1000 mg in 2 to 3 divided doses	3000 mg	Sedation is a common side effect.

2017 Guideline for the Prevention, Detection, Evaluation, and Management of High Blood Pressure in Adults

Oral Antihypertensive Drugs (1 of 3)

Class	Drug	Usual Dose, Range (mg per day)*	Daily Frequency	Comments	
Primary Agents					
Thiazide or	Chlorthalidone	12.5-25	1	Chlorthalidone preferred based on prolonged	
thiazide-type	Hydrochlorothiazide	25-50	1	half-life and proven trial reduction of CVD	
diuretics	Indapamide	1.25-2.5	1	Monitor for hyponatremia and hypokalemia, uric acid and calcium levels.	
	Metolazone	2.5-10	1	Use with caution in patients with history of acute gout unless patient is on uric acid-lowering therapy.	
ACE Inhibitors	Benazepril	10-40	1 or 2	Do not use in combination with ARBs or direct	
	Captopril	12.5-150	2 or 3	renin inhibitor	
	Enalapril	5-40	1 or 2	Increased risk of hyperkalemia, especially in patients with CKD or in those on K+ supplements	
	Fosinopril	10-40	1	or K+-sparing drugs	
	Lisinopril	10-40	1	May cause acute renal failure in patients with	
	Moexipril	7.5-30	1 or 2	severe bilateral renal artery stenosis	
	Perindopril	4-16	1	Do not use if history of angioedema with ACE	
	Quinapril	10-80	1 or 2	inhibitors.	
	Ramipril	2.5-10	1 or 2	Avoid in pregnancy	
	Trandolapril	1-4	1		
ARBs	Azilsartan	40-80	1	Do not use in combination with ACE inhibitors or	
	Candesartan	8-32	1	direct renin inhibitor	
	Eprosartan	600-800	1 or 2	Increased risk of hyperkalemia in CKD or in those on K+ supplements or K+-sparing drugs	
	Irbesartan	150-300	1	May cause acute renal failure in patients with	
	Losartan	50-100	1 or 2	severe bilateral renal artery stenosis	
	Olmesartan	20-40	1	Do not use if history of angioedema with ARBs.	
	Telmisartan	20-80	1	Patients with a history of angioedema with an	
	Valsartan	80-320	1	ACEI can receive an ARB beginning 6 weeks after ACEI discontinued. • Avoid in pregnancy	
CCB-	Amlodipine	2.5-10	1	Avoid use in patients with HFrEF; amlodipine or	
dihydropyridines	Felodipine	5-10	1	felodipine may be used if required	
	Isradipine	5-10	2	Associated with dose-related pedal edema, which	
	Nicardipine SR	5-20	1	is more common in women than men	
	Nifedipine LA	60-120	1]	
	Nisoldipine	30-90	1		
CCB-	Diltiazem SR	180-360	2	Avoid routine use with beta blockers due to	
nondihydropyridines	Diltiazem ER	120-480	1	increased risk of bradycardia and heart block	
	Verapamil IR	40-80	3	Do not use in patients with HFrEF	
	Verapamil SR	120-480	1 or 2	 Drug interactions with diltiazem and verapamil (CYP3A4 major substrate and moderate inhibito 	
	Verapamil-delayed onset ER (various	100-480	1 (in the evening)	Table is continued in the next two pages	

forms)

2017 Guideline for the Prevention, Detection, Evaluation, and Management of High Blood Pressure in Adults

Oral Antihypertensive Drugs (2 of 3)

Class	Drug	Usual Dose, Range (mg per day)*	Daily Frequency	Comments
Secondary Agent	ts			
Diuretics-loop	Bumetanide	0.5-4	2	Preferred diuretics in patients with symptomatic
	Furosemide	20-80	2	HF. Preferred over thiazides in patients with moderate-to-severe CKD (e.g., GFR <30 mL/min)
•	Torsemide	5-10	1	inouerate-to-severe GND (e.g., GFK \50 III./ IIIIII)
Diuretics-	Amiloride	5-10	1 or 2	Monotherapy agents minimally effective
potassium sparing	Triamterene	50-100	1 or 2	antihypertensives Combination therapy of potassium sparing diuretic with a thiazide can be considered in patients with hypokalemia on thiazide monotherapy
				• Avoid in patients with significant CKD (e.g., GFR <45 mL/min)
Diuretics— aldosterone	Eplerenone	50-100	12	Preferred agents in primary aldosteronism and resistant hypertension
antagonists	Spironolactone	25-100	1	 Spironolactone associated with greater risk of gynecomastia and impotence compared to eplerenone Common add-on therapy in resistant hypertension Avoid use with K+ supplements, other K+-sparing diuretics or significant renal dysfunction
				Eplerenone often requires twice daily dosing for adequate BP lowering
Beta blockers-	Atenolol	25-100	12	Beta blockers are not recommended as first-line
cardioselective	Betaxolol	5-20	1	agents unless the patient has IHD or HF
	Bisorolol	2.5-10	1	Preferred in patients with bronchospastic airway
	Metoprolol tartrate	100-400	2	disease requiring a beta blocker
	Metoprolol succinate	50-200	1	Bisoprolol and metoprolol succinate preferred in patients with HFrEF Avoid abrupt cessation
Beta blockers— cardioselective and vasodilatory	Nebivolol	5-40	1	Induces nitric oxide-induced vasodilation Avoid abrupt cessation
Beta blockers—	Nadolol	40-120	1	Avoid in patients with reactive airways disease
noncardioselective	Propranolol IR	160-480	2	Avoid abrupt cessation
	Propranolol LA	80-320	1	
Beta blockers-	Acebutolol	200-800	2	Generally avoid, especially in patients with IHD or HF
intrinsic	Carteolol	2.5-10	1	Avoid abrupt cessation
sympathomimetic activity	Penbutolol	10-40	1	
astivity	Pindolol	10-60	2	Table is continued in the next page

2017 Guideline for the Prevention, Detection, Evaluation, and Management of High Blood Pressure in Adults

Oral Antihypertensive Drugs (3 of 3)

Class	Drug	Usual Dose, Range (mg per day)*	Daily Frequency	Comments			
Secondary Agent	Secondary Agents (continued from previous page)						
Beta blockers—	Carvedilol	12.5-50	2	Carvedilol preferred in patients with HFrEF			
combined alpha- and beta-receptor	Carvedilol phosphate	20-80	1	Avoid abrupt cessation			
beta-receptor	Labetalol	200-800	2				
Direct renin inhibitor	Aliskiren	150-300	1	Do not use in combination with ACE inhibitors or ARBs			
				Aliskiren is very long acting			
				Increased risk of hyperkalemia in CKD or in those on K+ supplements or K+ sparing drugs			
				May cause acute renal failure in patients with severe bilateral renal artery stenosis			
				Avoid in pregnancy			
Alpha-1 blockers	Doxazosin 1-8 1 • Associate	Associated with orthostatic hypotension,					
	Prazosin	2-20	2 or 3	especially in older adults			
	Terazosin	1–20	1 or 2	May consider as second-line agent in patients with concomitant BPH			
Central alpha1-	Clonidine oral	0.1-0.8	2	Generally reserved as last-line due to significant			
agonist and other	Clonidine patch	0.1-0.3	1 weekly	CNS adverse effects, especially in older adults			
centrally acting drugs	Methyldopa	250-1000	2	Avoid abrupt discontinuation of clonidine, which may induce hypertensive crisis; clonidine must be			
	Guanfacine	0.5-2	1	tapered to avoid rebound hypertension			
Direct vasodilators	Hydralazine	250-200	2 or 3	Associated with sodium and water retention and			
	Minoxidil	5-100	1 -3	reflex tachycardia; use with a diuretic and bet a blocker			
				Hydralazine associated with drug-induced lupus- like syndrome at higher doses			
				Minoxidil associated with hirsutism and requires a loop diuretic. Can induce pericardial effusion			

^{*}Dosages may vary from those listed in the FDA approved labeling (available at http://dailymed.nlm.nih.gov/dailymed/index.cfm). Adapted with permission from Chobanian AV, Bakris GL, Black HR, et al. The Seventh Report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure: the JNC 7 report. JAMA. 2003; 289:2560-72 Table 18



Interdependent and Interacting Factors in Blood Pressure Regulation

