BP Guideline in 2020

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Introduction



NICE guideline

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





Articles

Worldwide trends in blood pressure from 1975 to 2015: a pooled analysis of 1479 population-based measurement studies with 19.1 million participants





NCD Risk Factor Collaboration (NCD-RisC)*

Summary

Background Raised blood pressure is an important risk factor for cardiovascular diseases and chronic kidney disease. We estimated worldwide trends in mean systolic and mean diastolic blood pressure, and the prevalence of, and number of people with, raised blood pressure, defined as systolic blood pressure of 140 mm Hg or higher or diastolic blood pressure of 90 mm Hg or higher.

Methods For this analysis, we pooled national, subnational, or community population-based studies that had measured blood pressure in adults aged 18 years and older. We used a Bayesian hierarchical model to estimate trends from 1975 to 2015 in mean systolic and mean diastolic blood pressure, and the prevalence of raised blood pressure for 200 countries. We calculated the contributions of changes in prevalence versus population growth and ageing to the increase in the number of adults with raised blood pressure.

Findings We pooled 1479 studies that had measured the blood pressures of 19 · 1 million adults. Global age-standardised mean systolic blood pressure in 2015 was 127.0 mm Hg (95% credible interval 125.7–128.3) in men and 122.3 mm Hg (121.0-123.6) in women; age-standardised mean diastolic blood pressure was 78.7 mm Hg (77.9-79.5) for men and 76.7 mm Hg (75.9-77.6) for women. Global age-standardised prevalence of raised blood pressure was 24.1% (21·4-27·1) in men and 20·1% (17·8-22·5) in women in 2015. Mean systolic and mean diastolic blood pressure



Lancet 2017; 389: 37-55

Published Online November 15, 2016 http://dx.doi.org/10.1016/ 50140-6736(16)31919-5

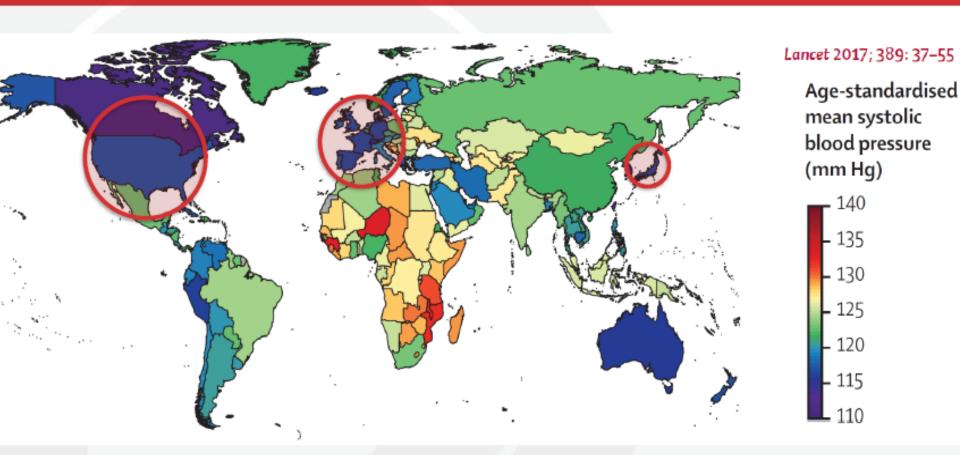
This online publication has been corrected. The corrected version first appeared at thelancet.com on September 24, 2020

See Comment page 3

*NCD Risk Factor Collaboration members are listed at the end of the paper

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Introduction





- 1.39 billion estimated with hypertension in 2010
- 349 million from HIC
- **1.04 billion** from LMIC *Circulation.* 2016;134:441–450

Introduction

The ISH 2020 Global Hypertension Practice Guidelines were thus developed based on evidence criteria,

- a) to be used globally
- to be fit for application in low-resource and highresource settings by advising on **ESSENTIAL** and
 OPTIMAL standards of care; and
- to be concise, simplified and easy to use by clinicians, nurses and community health workers, as appropriate.



Case

خانم ۴۶ ساله متاهل جهت چکاب مراجعه کرده سابقه بیماری خاصی در گذشته نم ی دهد در طَی دو سال اخیر BP در منزل حدود۱۳۵/۸۵ ست سابقه HTN در پدر دارد کارمند بانک است ۲ دختر دانشجو دارد

PE: RR: 16/min, BP:140/85 mmHg, PR: 74/min, T: 36 C, BMI: 28

در معاینه نکته خاصی ندارد. HBPMیک ماه قبل:

Day time average: 130/90

Night time average: 125/80

24 hs average: 130/85

سوال ١

در مورد تشخیص بیمار کدام مورد صحیح است؟

- A. مبتلا به HTN است
- B. جهت تشخیص قطعی نیاز به کنترل مجدد BP در مطب است
 - C. جهت تشخیص قطعی نیاز به HBPM است
 - D. جهت تشخیص قطعی نیاز به تکرار ABPM است

Definition of Hypertension

ESSENTIAL

Classification of hypertension based on Office blood pressure (BP) measurement

Category	Systolic (mmHg)		Diastolic (mmHg)
Normal BP	< 130	and	< 85
High-normal BP	130-139	and/or	85-89
Grade 1 Hypertension	140-159	and/or	90-99
Grade 2 Hypertension	≥160	and/or	≥100



ISH vs ACC/AHA Guidelines

International Society of Hypertension	Systolic (mmHg)		Diastolic (mmHg)
Normal BP	<130	and	<85
High-normal BP	130-139	and/or	85-89
Grade 1 Hypertension	140-159	and/or	90-99
Grade 2 Hypertension	≥160	and/or	≥100

Heart Association	E of		DBP
Normal	<120 mm Hg	and	<80 mm Hg
Elevated	120-129 mm Hg	and	<80 mm Hg
Hypertension			
Stage 1	130-139 mm Hg	or	80-89 mm Hg
Stage 2	≥140 mm Hg	or	≥90 mm Hg

Definition of Hypertension

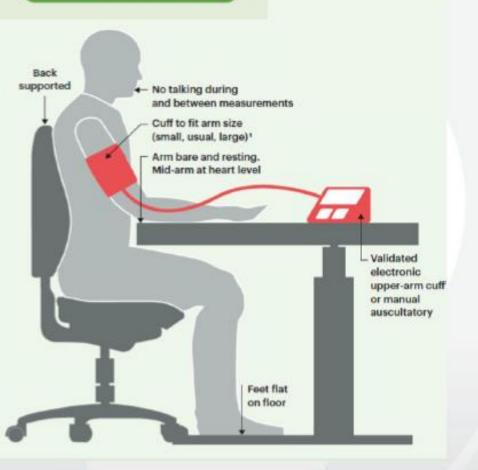
ESSENTIAL

Hypertension based on Office-, Ambulatory (ABPM)and Home Blood Pressure (HBPM) measurement

		SBP / DBP (mmHg)
Office BP		≥ 140 and/or ≥ 90
ABPM	Day Time (or awake) average Night Time (or asleep) average	≥ 130 and/or ≥ 80 ≥ 135 and/or ≥ 85 ≥ 120 and/or ≥ 70
НВРМ		≥ 135 and/or ≥ 85



ESSENTIAL





Office Blood Pressure Measurement

- 2-3 office visits at 1-4-week intervals.
- Whenever possible, the diagnosis should not be made on a single visit (unless BP ≥180/110 mmHg and CVD).
- If possible and available the diagnosis of hypertension should be confirmed by outof-office measurement.

ESSENTIAL

OFFICE BP MEASUREMENT

Conditions

Position

- Setting
- · Body position
- Talking



Device

Cuff

- Validated electronic upper-arm cuff (<u>www.stridebp.org</u>)
- Alternatively manual auscultatory device
- Cuff size

Protocol

Interpretation

- Average 2nd-3rd measurement
- 2-3 office visits required

www.ish-world.com















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About us BP Monitors Training Literature





STRIDE BP is an international scientific non-profit organization founded by hypertension experts with the mission of improving the diagnosis and management of hypertension.

Read more →















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EN ES ZH

Q Search



STRIDE BP has approved **310** and recommends as preferred only **170** of the over **4,000** electronic blood pressure monitors currently available on the market











→ Download















VALIDATED DEVICES FOR OFFICE / CLINIC BP MEASUREMENT

Preferred devices (35)

A&D UM-101

A&D UM-211

Accoson Greenlight 300

Andon iHealth BP3

Andon iHealth Clear BPM1 *

Andon iHealth Ease BP3L *

Andon iHealth Neo BP5S *

Andon iHealth Track KN-550BT *

Andon KD-558BR *

Andon KD-5920 *

Andon KD-5923 *

Andon KD-5965

Artsana Pic Indolor Professional

Dinamap ProCare 400

Erkameter 125 PRO

InBody BPBIO250

InBody HBP570 *

Microlife 3AS1-2

Microlife VSA (BP3GP1-1L) *

Microlife WatchBP Office

Microlife WatchBP Office (BP3SK1-3B) *

Microlife WatchBP Office ABI

Microlife WatchBP Office AFIB *

Microlife WatchBP Office Central *

Nissei DM3000

Omron HBP T105

Omron HBP-1300

Omron HBP-1320

Omron HBP-9030 *

Omron M3500

Raycome RBP-1200

RisingSun RS-651

Rossmax AC1000f

Suntech CT40

Welch Allyn ProBP 2000

Validated devices (10)

- Andon iHealth View BP7S
- 2. Omron HEM-907
- 3. Andon KD-723
- Rossmax Mandaus II
- 5. BpTRU BPM-100
- 6. Spengler Pro M
- 7. Dinamap ProCare
- 8. Suntech 247
- 9. Mindray Datascope Accutorr Plus
- 10. Welch Allyn Vital Signs

ESSENTIAL

BP Measurement Plan according to Office BP levels

Office blood pressure levels (mmHg)

<130/85	130-159/85-99	>160/100
 Remeasure within 3 years (1 year if other risk factors). 	If possible confirm with out-of-office measurement.	 Confirm within a few days/weeks.
other had ractors).	 Alternatively confirm with repeated office visits. 	



OPTIMAL

Office Blood Pressure

Initial evaluation

 Measure BP in both arms. Difference >10 mmHg: use arm with higher BP; >20 mmHg: consider further investigation.

Standing BP

- In treated patients when symptoms of postural hypotension.
- At first visit in elderly and diabetics.

Unattended BP

- More standardized. Lower BP levels with uncertain threshold.
- Out-of-office BP again needed in most cases



سوال ۲

در مورد تعریف Unattended BP کدام مورد صحیح است؟

- A. کنترل فشارخون در مطب بدون حضور همراه
- B. کنترل فشارخون بطور اتوماتیک در مطب بدون حضور هر فردی در اتاق
 - .C کنترل فشارخون بطور اتوماتیک در مطب توسط پرستار ماهر در حضور همراه
- D. کنترل فشارخون در منزل و اطلاع آن بوسیله همراه بدون حضور بیمار

Automated OBP (AOBP) technique or Unattended BP

Methodology adopted in SPRINT (SBP Intervention
 Trial), used a fully automated oscillometric device with the patient sitting alone in the examination room for
 minutes, after which 3 readings were taken automatically at 1-minute intervals with all 3 values averaged.

Automated OBP (AOBP) technique or Unattended BP

 Human involvement was reduced to the minimum, eliminating the "white coat effect," & AOBP readings correlated more closely with those of ABPM than conventional office recordings.

OPTIMAL

Clinical Use of Home and Ambulatory BP Monitoring

Conditions

Device

Protocol

Position

Cuff

Interpretation



OPTIMAL



	Home BP Monitoring	Ambulatory BP Monitoring	
Conditions	As for office blood pressure (see above).	Routine working day.	
Position	As for office BP (see above).	Avoid strenuous activity. Arm still and relaxed during each measurement.	
Device	Validated electronic (oscillometric) upper-arm cuff device (www.stridebp.org, and Section 11: Resources)		
Cuff	Size according to the individual's arm circumference		
Measurement protocol		 24-hour monitoring at 15 – 30 min intervals during daytime and nighttime. At least 20 valid daytime and 7 nighttime BP readings are required. If less, the test should be repeated. 	
Interpretation	 Average home blood pressure after excluding readings of the first day ≥ 135 or 85 mmHg indicates hypertension. 	 24-hour ambulatory blood pressure ≥ 130/80 mmHg indicates hypertension (primary criterion). Daytime (awake) ambulatory blood pressure ≥ 135/85 mmHg and nighttime (asleep) ≥ 120/70 mmHg indicates hypertension 	

OPTIMAL

White-coat Hypertension

- Intermediate CV risk.
- If low total CV risk and no organ damage, drug treatment may not be prescribed.
- Follow with lifestyle changes.

Masked Hypertension

- Similar CV risk as sustained hypertensives.
- Drug treatment may be required aiming to normalise out-of-office BP.



Diagnostic and Clinical Tests

ESSENTIAL

- Medical History (BP, risk factors, co-morbidities, signs/symptoms of secondary hypertension...)
- Physical Examination (circulation, heart, other systems)
- Lab Investigations (Na+, K+, creatinine, eGFR, dipstick lipids, Fasting Glucose where available)
- 12 lead ECG (AF, LV hypertrophy, IHD...)

OPTIMAL

 Additional tests to consider (extended biochemistry, cardiac/kidney/brain/vascular imaging, fundoscopy...)



Cardiovascular Risk Factors

- More than 50% of hypertensive patients have additional CV risk factors
- Most commonly: Met Syn, T2DM, lipid disorders, ↑ uric acid
- CV risk assessment is important and should be assessed in all hypertensive patients
- Consider increased risk with: chronic inflammatory disease,
 COPD, psychiatric disorders, psycho-social stressors



Cardiovascular Risk Factors

Other risk factors, HMOD, or disease	High-normal SBP 130-139 DBP 85-89	Grade 1 SBP 140-159 DBP 90-99	Grade 2 SBP ≥ 160 DBP ≥ 100
No other risk factors	Low	Low	Moderate High
1 or 2 risk factors	Low	Moderate	High
≥3 risk factors	Low Moderate	High	High
HMOD, CKD grade 3, diabetes mellitus, CVD	High	High	High



Hypertension-mediated Organ Damage

HMOD Assessment

ESSENTIAL

- Serum creatinine
- eGFR
- Dipstick urine test
- 12-lead ECG

OPTIMAL

- Brain
- Eyes
- Heart
- Kidneys
- Arteries

Serial assessment of HMOD

may help to determine efficacy of treatment



سوال ۳

دراین بیمار چه اقدام درمانی توصیه می کنید؟

- LSM .A
- B. والزارتان ۸۰ میلیگرم روزانه
 - C. لوزاتن H روزی ۱ عدد
- D. والزوميكس ۵/۸۰ روزي ۱ عدد
- E. هیدروکلروتیازید ۱۲/۵ میلیگرم روزانه

Exacerbators & Inducers of Hypertension

Most common medications that can increase BP

- Non-selective or traditional NSAIDs
- Combined oral contraceptive pill
- Select anti depressant medications including tricyclic antidepressants and SNRIs
- Acetaminophen when used almost daily and for prolonged periods



Exacerbators & Inducers of Hypertension

- The effect of Anti-retroviral therapy is unclear as studies demonstrate either no effect on BP or some increase.
- Alcohol raises BP regardless of the type of alcoholic drink.
- Limited evidence on herbal and other substances.
- Ma Huang, Ginseng at high doses and St. John's Wort reported to increased BP.



Non-pharmacological Treatment

- Healthy lifestyle choices can prevent or delay the onset of high BP and can reduce CV risk
- Lifestyle modification is often the first line of antihypertensive treatment.
- Modifications in lifestyle can also enhance the effects of antihypertensive treatment.





Non-pharmacological Treatment - Diet

- Reducing salt added when preparing foods and at the table. Avoid or limit consumption of high salt foods.
- Eating a diet rich in whole grains, fruits, vegetables, polyunsaturated fats and dairy products, such as DASH diet.
- Reducing food high in sugar, saturated fat and trans fats.
- Increasing intake of vegetables high in nitrates (leafy vegetables and beetroot). Other beneficial foods and nutrients include those high in magnesium, calcium and potassium (avocados, nuts, seeds, legumes and tofu).

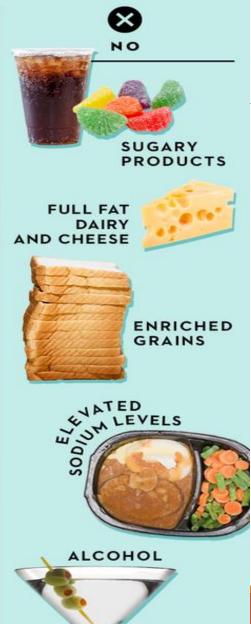






THE **DASH** DIET WHAT CAN I EAT?





Non-pharmacological Treatment - Diet

- Moderate consumption of healthy drinks (coffee, green and black tea, Karkadé (Hibiscus) tea, pomegranate juice, beetroot juice and cocoa.
- Moderation of alcohol consumption and avoidance of binge drinking.
- Reduce weight and avoid obesity.
- Be careful with complementary, alternative or traditional medicines – little/no evidence.



Non-pharmacological Treatment - Lifestyle

Smoking cessation.



 Engage in regular moderate intensity aerobic and resistance exercise, 30 minutes on 5 – 7 days per week or HIIT (High Intensity Interval Training).

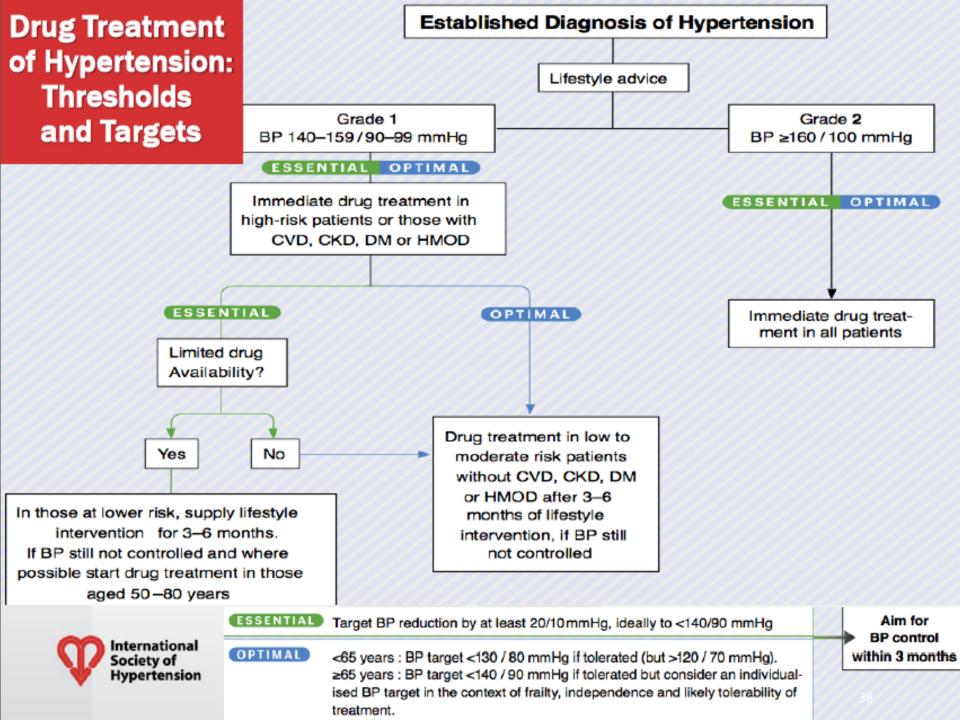


Reduce stress and introduce mindfulness.



Reduce exposure to air pollution and cold temperature.





ESSENTIAL

- · Use whatever drugs are available with as many of the ideal characteristics (see Table 9) as possible.
- · Use free combinations if SPCs are not available or unaffordable
- Use thiazide diuretics if thiazide-like diuretics are not available
- Use alternative to DHP-CCBs if these are not available or not tolerated (i.e. Non-DHP-CCBs: diltiazem or verapamil).

ESSENTIAL and OPTIMAL

Consider beta-blockers at any treatment step when there is a specific indication for their use, e.g. heart failure, angina, post-MI, atrial fibrillation, or younger women with, or planning pregnancy.



Drug choice & Sequencing

Ideally Single Pill Combination Therapy (SPC)

OPTIMAL

Step 1

Dual low-dose# combination

A + C a, b, c

Step 2

Dual full-dose combination

A + C a, b

Step 3

Triple combination

A + C + D

Step 4

(Resistant Hypertension) Triple Combination + Spironolactone or other drug*

A + C + DAdd Spironolactone (12.5 - 50 mg o.d.)d

- a) Consider monotherapy in low risk grade 1 hypertension or in very old (≥80 yrs) or frailer patients.
- **b)** Consider A + D in post-stroke, very elderly, incipient heart failure or CCB intolerance. its.
- c) Conside A screenshot of a cell phone
- d) Caution Description automatically generated tassium sparing diuretics when estimated GFR <45 mi/min/1./3m* or K* >4.5 mmol/L.
- A = ACE-Inhibitor or ARB (Angiotensin Receptor Blocker)
- C = DHP-CCB (Dihydropyridine -Calcium Channel Blocker)
- D = Thiazide-like diuretic

Drug Treatment of Hypertension

Summary 1

In established hypertension, uncontrolled by lifestyle measures:

Drug Treatment Threshold

≥140/90 mmHg (raising to ≥160/100 mmHg for those at lowest risk)

Drug Treatment Target

OPTIMAL

<65 years: <130/80 mmHg

≥65 years: <140/90 mmHg

[ESSENTIAL]

reduce BP by ≥20/10 mmHg



Drug Treatment of Hypertension

Summary 2

OPTIMAL

- (i) Uptitration to target, of the following:
 Low dose A+C → Full dose A+C → A+C+D
 - → A+C+D + spironolactone
- (ii) Consider other initial combinations for specific patient subgroups
- (iii) Use SPC's where possible
- (iv) Use thiazide-like diuretics preferentially

ESSENTIAL

 Where less ideal agents are available, focus on effective BP lowering (≥20/10 mmHg)



سوال ۴

دراین بیمار قواصل ویزیت چند ماه یک بار است؟

- 1 .A
- ۲ .B
- ٣ .C
- 9 .D

سوال ۵

دراین بیمار بعد از کنترل BP فواصل ویزیت چند ماه یک بار می شود؟

- **M-1.A**
- ۶-۳ .B
- 17-8 .C
- D. بسته به سن بیمار و نوع دارو فواصل ویزیت تغییر می کند

Comorbidities of Hypertension

- Well established common comorbidities include CAD, stroke, CKD, Heart failure, COPD and HIV/AIDS.
- Emerging uncommon comorbidities include rheumatic/inflammatory diseases and psychiatric diseases.
- Uncommon comorbidities are largely underestimated by guidelines and often treated with self-prescribed drugs frequently interfering with BP control.



Comorbidities of Hypertension

In patients with **common comorbidities** the therapeutic strategy depends on CV risk profile and includes:

- Lifestyle changes (diet, exercise, body weight, smoking).
- BP control to target.
- Effective treatment of CV risk factors (LDL-C, Fasting Glucose, SUA).
- Antiplatelet therapy in patients with CVD.



Comorbidities of Hypertension

Additional co-morbidit	Recommended Drugs	Warning
Rheumatic	 RAS-inhibitors and CCBs ± Diuretics 	High doses of
disorders	 Biologic drugs not affecting blood pressure 	NSAID's
	should be preferred	
	(where available)	
Psychiatric	RAS-inhibitors and diuretics	Avoid CCBs if
disorders	Beta-blockers (not metoprolol) if drug-induced	orthostatic
	tachycardia (antidepressant, antipsychotic	hypotension (SRI's)
	drugs).	
	 Lipid-lowering drugs/Antidiabetic drugs 	
	according to risk profile	



Resistant Hypertension

- Suspect resistant hypertension if office BP >140/90 mmHg on treatment with at least 3 antihypertensives (in maximal or maximally tolerated doses) including a diuretic.
- Exclude pseudo-resistant hypertension (white-coat effect, non-adherence to treatment, incorrect BP measurements, errors in antihypertensive therapy) and substance-induced hypertension as contributors.
- Optimise health behaviours and lifestyle.



Resistant Hypertension

- Consider changes in the diuretic-based treatment prior to adding the fourth antihypertensive medication.
- Add a low dose of spironolactone (if serum potassium is <4.5 mmol/L and eGFR is >45 ml/min/1.73 m²).
- Consider amiloride, doxazosin, eplerenone, clonidine and beta-blockers as alternatives to spironolactone. If unavailable, consider any antihypertensive class not already in use.
- Optimally, consider referring to a specialist centre with sufficient expertise/resources.



Secondary Hypertension

ESSENTIAL

Basic screening for secondary hypertension thorough history + physical examination (clinical clues) + basic blood biochemistry (including serum sodium, potassium, eGFR, TSH) + dipstick urine analysis.

OPTIMAL

Arrange other investigations for secondary hypertension (additional biochemistry/imaging/others) based on information from history, physical examination and basic clinical investigations and/or if feasible refer to a specialist centre



Hypertensive Emergencies

Management

- Requires immediate BP lowering to prevent or limit further HMOD
- Sparse evidence to guiding management recommendations largely consensus based.
- Time to lower BP and magnitude of BP reduction depends on clinical context.
- IV Labetalol and nicardipine generally safe to use in all hypertensive emergencies



	Clinical presentation	Timeline and target BP	1st line treatment
	Malignant hypertension with or	Several hours,	Labetalol
	without TMA or acute renal failure	MAP - 20 % to - 25 %	Nicardipine
	Hypertensive encephalopathy	Immediate,	Labetalol
		MAP - 20 % to - 25 %	Nicardipine
	Acute ischemic stroke and BP > 220	1 h,	Labetalol
	mmHg systolic or >120 mmHg	MAP - 15 %	Nicardipine
	diastolic		
Hyportoneive	Acute ischemic stroke with indication for thrombolytic therapy	1 h, MAP – 15 %	Labetalol
Hypertensive	and BP > 185 mmHg systolic or >	MAF - 15 /0	Nicardipine
Hypertensive Emergencies	110 mmHg diastolic		
	Acute <u>hemorrhagic</u> stroke and	Immediate,	Labetalol
	systolic BP >180 mmHg	systolic 130 < BP	Nicardipine
		< 180 mmHg	
	Acute coronary event	Immediate,	Nitroglycerine
		systolic BP < 140 mmHg	Labetalol
	Acute cardiogenic pulmonary	Immediate,	Nitroprusside or
	edema	systolic BP <140 mmHg	Nitroglycerine
			(with loop diuretic)
	Acute aortic disease	Immediate,	Esmolol and Nitroprusside or
		systolic BP <120 mmHg	Nitroglycerine or Nicardipine
International		and heart rate <60 b.p.m.	
Society of Hypertension	Eclampsia and severe pre-	Immediate, systolic	Labetalol or Nicardipine and
	eclampsia/HELLP	BP < 160 mmHg and diastolic BP < 105 mmHg	Magnesium sulphate

rdipine etalol Nitroprusside rdipine etalol rdipine

Nitroprusside Urapidil

Urapidil

Urapidi (with loop diuretic)

Labetalol or

Metoprolol

Alternative

Nitroprusside

Urapidil

Nitroprusside

Ethnicity, Race and Hypertension

Populations from ASIA

 Morning & nighttime hypertension vs Europeans

EAST ASIAN populations

- Likelihood of salt-sensitivity + mild obesity in hypertensive patients
- Stroke prevalence (esp. hemorrhagic) & nonischemic HF vs Western populations

SOUTH ASIAN populations (Indian subcontinent)

Risk for CV & metabolic diseases (CAD & T2DM)

Management of hypertension

SOUTH EAST ASIA: Standard treatment until more evidence becomes available



Hypertension Management at a Glance

Diagnosis

Normal BP level (<130/85 mmHg)

Remeasure after 3 years (1 year in those with other risk factors)

* Use a validated upper arm-ouff device with appropriate ouff size for the individual patient.

Single office BP measurement*

High-normal BP level (130-139/85-89 mmHg)

Take 2 more readings - use the average of 2nd-3nd

≥130/85 mmHq

Remeasure in 2-3 office visits. If possible confirm with home or ambulatory BP monitoring

Repeated office BP ≥140/90 mmHg indicates hypertension, particularly if home BP ≥ 135/85 mmHg or 24h ambulatory BP ≥ 130/80 mmHg

ESSENTIAL

Evaluation

History & Physical Exam

- Exclude drug-induced hypertension
- Evaluate for organ damage
- Assess total cardiovascular risk
- Search for symptoms/signs of secondary hypertension

Lab Tests

- Serum sodium, potassium & creatinine
- Lipid profile & alucose
- Urine dipatick
- 12 lead ECG

Additional Tests

 If necessary for suspected organ damage or secondary hypertension

Hypertension BP level

(≥140/90mmHg)

Treatment

Grade 1 Hypertension:

140-159/90-99 mmHg 1. Start lifestyle interventions

- 2. Start drug treatment in:
- High-risk patients (CVK.CKD. diabetes, organ damage.
 - or aged 50-80 years) All others with persistent BP

elevation after 3-6 months of lifestyle intervention

≥160/100 mmHg

Grade 2 Hypertension:

2. Start lifestyle intervention

1. Start drug treatment immediately

Lifestyle Interventions

- Stop smoking
- Regular exercise
- · Lose weight
- · Salt reduction
- Healthy diet and drinks
- Lower alcohol intake

Drug Therapy Steps

Use any drugs available and include as many of those below as possible. Consider monotherapy in low-risk grade 1 hypertension and in patients aged >80 years or frail. Simplify regimen with once daily dosing and single pill combinations.

Non-Black Patients

- 1. Low dose ACEI/ARB* + dCCB
- 2. Increase to full dose
- Add thiazide/thiazide-like diuretic
- Add spironolactone or, if not tolerated or contraindicated, amiloride, doxazosin, eplerenone, clonidine or beta-blocker

Black Patients

Low dose ARB* + dCCB or dCCB

- + thiazide/thiazide-like diuretic
- 2. Increase to full dose
- 3. Add diuretic or ARB /ACEI
- 4. Add spironolactone or, if not toler-
- ated or contraindicated, amiloride, doxazosin, eplerenone, clonidine or beta-blocker

No ACEI/ARB in women with or planning pregnancy



Monitoring

- Reduce BP by at least 20/10 mmHg, ideally to < 140/90 mmHg
- Individualize for elderly based on frailty

Monitor

- BP control
- (achieve target within 3 months)
- Adverse effects
- Long-term adherence

Referral

 If BP still uncontrolled, or other issue, refer to care provider with hypertension expertise

Hypertension Management at a Glance

Diagnosis

Office BP measurement

(3 readings - use the average of 2nd-3rd)*

Confirm with home or ambulatory BP monitoring

≥130/85mmHg

Home BP <135/85mmHg or 24h ambulatory BP < 130/80 mmHg

Remeasure after 1 year

Home BP ≥135/85 mmHq or 24 h ambulatory $BP \ge 130/80 \, \text{mmHg}$

Hypertension diagnosis

OPTIMAL

Evaluation

History & Physical Exam

Exclude drug-induced

the arm with the higher BP.

- hypertension Evaluate for organ damage
- Consider additional CV risk factors
- Assess total cardiovascular risk

<130/85 mmHa

Remeasure after 3 years

(1 year in those with other risk factors)

* Use a validated automated upper arm-cuff device with appropriate cuff size for the indi-

vidual patient. At first visit measure BP in both arms simultaneously. If consistent difference use

- · Search for symptoms/signs of secondary hypertension
- Check adherence

Lab Tests

 Serum sodium, potassium & creatinine, uric acid

- Lipid profile & glucose
- Urine dipstick
- 12 lead ECG

Additional Tests

 If necessary for suspected organ damage or secondary hypertension

Treatment

Grade 1 Hypertension:

140-159/90-99 mmHg Start lifestyle interventions

- Start drug treatment:
- Immediately: In high-risk patients (CVD, CKD, diabetes or organ damage)
- After 3—6 months of lifestyle intervention: In low-moderate risk patients with persistent BP elevation

Grade 2 Hypertension:

- ≥160/100 mmHg
- 1. Start drug treatment immediately
- Start lifestyle intervention

Lifestyle Interventions

- Stop smoking
- Regular exercise
- Lose weight
- Salt reduction
- Healthy diet and drinks
- Lower alcohol intake
- Lower stress
- Reduce exposure to air pollution

Drug Therapy Steps Simplify regimen with once daily dosing and single pill combinations. Consider monotherapy in low-risk grade 1 hypertension and in patients aged >80 years or frail

Non-Black Patients

- 1. Low dose ACEI/ARB* + dCCB
- 2. Increase to full dose
- Add thiazide-like diuretic

or beta-blocker

Add spironolactone or, if not tolerated or contraindicated, amiloride, doxazosin, eplerenone, clonidine

No ACEI/ARB in women with or planning pregnancy.

- Black Patients Low dose ARB* + dCCB or dCCB
- + thiazide-like diuretic
- 2. Increase to full dose
- Add diuretic or ACE/ARB
- 4. Add spironolactone or, if not tolerated or contraindicated, amiloride, doxazosin, eplerenone, clonidine or beta-blocker



Monitoring

Target

- BP <130/80 mmHg
- Individualise for elderly based on frailty

Monitor

- BP control (achieve target within 3 months)
- Adverse effects
- Long-term adherence

Referral

 If BP still uncontrolled, or other issue, refer to care provider with hypertension expertise

ISH vs ACC/AHA Guidelines

- Blood pressure definitions of normal blood pressure stages of hypertension are different.
- Inclusion of high-normal blood pressure category.
- Blood pressure value thresholds for treatment are therefore different (i.e., treatment initiated at lower blood pressure in ACC/AHA guidelines).
- Adoption of ESSENTIAL VS. OPTIMAL throughout ISH guidelines.

