# Cardiovascular Diseases in Family Medicine

## János Nemcsik MD, PhD

specialist of internal medicine, family medicine, occupational medicine, hypertension care

Department of Family Medicine Semmelweis University Budapest

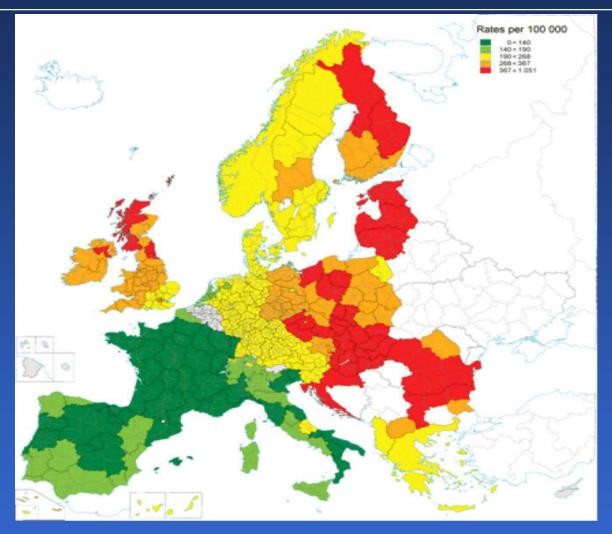
NGNB Med. Medical Services Ltd. 1148 Budapest, Örs vezér tér 23.

Email: janos.nemcsik@gmail.com

# Why the prevention of CV diseases is necessary?

- Atherosclerotic CVD, especially CHD, remains the leading cause of premature death worldwide.
- CVD affects both men and women; of all deaths that occur before the age of 75 years in Europe, 42% are due to CVD in women and 38% in men.
- CVD mortality is changing, with declining age-standardized rates in most European countries, which remain high in Eastern Europe.
- Prevention works: 50% of the reductions seen in CHD mortality is related to changes in risk factors, and 40% to improved treatments.

## Regional variation in cardiovascular mortality within Europe



Age-standardized mortality from ischaemic heart disease in European regions (men; age group 45–74 years; year 2000)

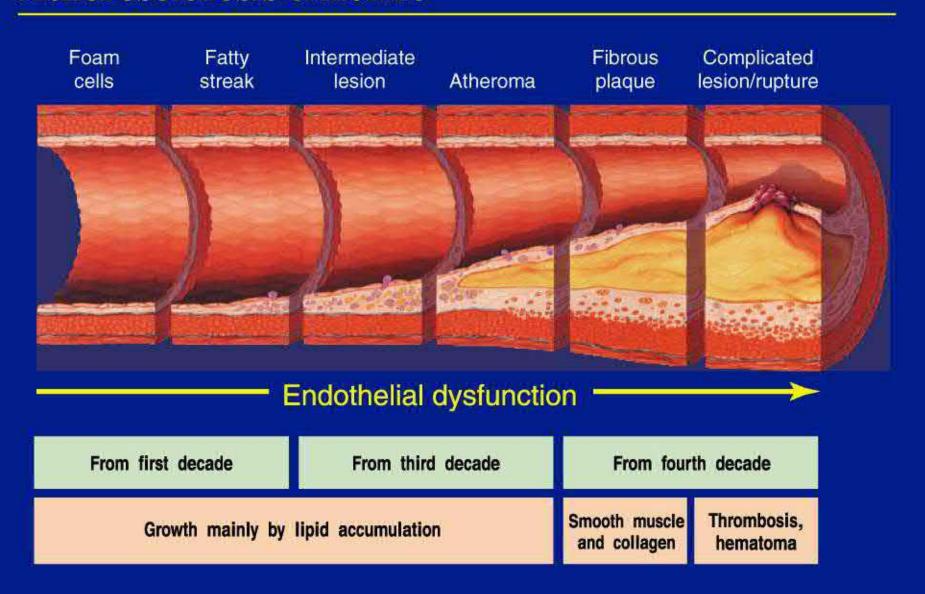
Eur Heart J. 2008;29(10):1316-1326. doi:10.1093/eurheartj/ehm604

# Mortality: Cardiovascular diseases and diabetes, deaths per 100,000

http://apps.who.int/gho/data/?vid=2510

	male	female
Hungary	416	241
USA	190	122
Spain	140	86
France	128	69
Russia	772	414
Mali	419	393

## Atherosclerosis timeline



## **Risk Factors for Cardiovascular Disease**

## Modifiable

- Smoking
- Dyslipidaemia
  - raised LDL-C
  - low HDL-C
  - raised triglycerides
- Raised blood pressure
- Diabetes mellitus
- Obesity
- Dietary factors
- Thrombogenic factors
- Lack of exercise
- Excess alcohol consumption

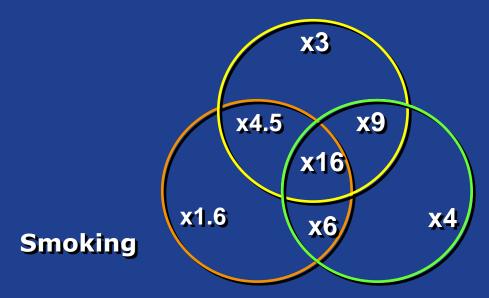
## Non-modifiable

- Personal history of CHD
- Family history of CHD
- Age
- Gender

**15**:1300-1331.

## Levels of Risk Associated with Smoking, Hypertension and Hypercholesterolaemia

Hypertension (SBP >195 mmHg)



Serum cholesterol level (>8.5 mmol/L, 330 mg/dL)

## Preventive medicine

# 2016 European Guidelines on cardiovascular disease prevention in clinical practice

Eur Heart J. **2016**;37:2315-2381.

e policies of workplaces, educational ctive.<sup>532</sup>

own to have a limited effect.

ary care to prevent alcohol abuse has

ve alcohol intake can be limited by re-

d opening hours of outlets and by inr sales and servings.<sup>495</sup>

needed with regard to potential coneffects of alcohol consumption.

## ronment

r CV preve... ×

ticles in the EU are motorized road trafland residential heating using oil, coal or eans living in urban areas are exposed to y standards. In particular, young and old high risk of CVD are more prone to the ution on the circulation and the heart. used a policy package to be implemenneasures to reduce harmful emissions d agriculture. Further efforts to reduce

mand and taken up by national accord

most enective mutualscipilitary care.

## Recommendation for cardiovascular disease prevention in primary care

Recommendation	Classa	Levelb
It is recommended that GPs, nurses and allied health professionals within primary care deliver CVD prevention for high-risk patients.	ı	U

<sup>&</sup>lt;sup>a</sup>Class of recommendation.

The physician in general practice is the key person to initiate, coordinate and provide long-term follow-up for CVD prevention. In most countries, GPs deliver >90% of consultations and provide most public health medicine, including preventive care and chronic disease monitoring. In the case of CVD prevention, they have a unique role in identifying individuals at risk of CVD and assessing their eligibility for intervention based on their risk profile. How to maximize attendance rates and adherence, particularly in those who are at highest risk, remains an issue.

As mentioned in section 2.2, a systematic approach is recom-

<sup>&</sup>lt;sup>b</sup>Level of evidence.

## How to stay healthy?

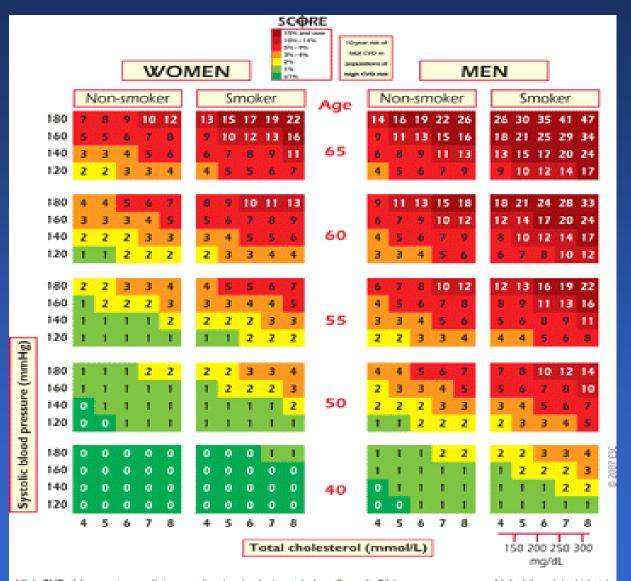
- No use of tobacco.
- Adequate physical activity: at least 30 min five times a week.
- Healthy eating habits.
- No overweight.
- Blood pressure below 140/90 mmHg.
- Blood cholesterol below 5 mmol/L (190 mg/dL).
- Normal glucose metabolism.
- Avoidance of excessive stress.

## Psychosocial risk factors

Contribute both to the risk of developing CVD and the worsening of clinical course and prognosis of CVD:

- -low socio-economic status
- -lack of social support
- -stress at work and in family life
- -depression, anxiety, hostility
- -type D personality

## SCORE: 10-year risk of fatal CV disease in high-risk population



High CVD risk countries are all those not listed under the low risk chart (Figure 4). Of these, some are at very high risk, and the high-risk chart may underestimate risk in these. These countries are Armenia, Azerbaijan, Belarus, Bulgaria, Georgia, Kazakhstan, Kyngyzstan, Latvia, Lichuania, Macedonia PYR, Moldova, Russia, Ukraine, and Utbekistan.

Calculate score at: www.hearscore.org



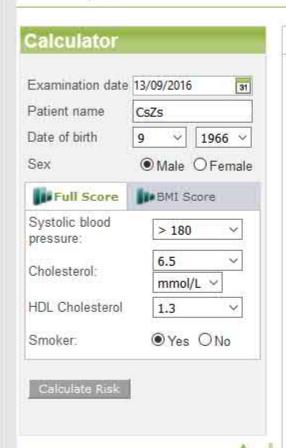


## HeartScore\*



HOME





HeartScore is based on the SCORE Risk Charts, which evaluate CVD risk for patients between 40 and 65 with SBP between 100 - 180 mmHg and Cholesterol between 3 - 8 mmol/L (105 - 305 mg/dl). Please note that patients with examination data over these value range are automatically at higher risk.

Patient Advice

CVD Prevention Guidelines

## Patient Advice



Patient p

The Patient Advice tab consolidates the advices given to the patient at the date of the examination.

This page allows you to have graphical displays of your patient's risk evaluation on the date of the examination.

The CVD Prevention Guidelines tab includes recommendations from the European Guidelines on CVI
Prevention

### What is CVD risk?

CVD risk means you risk of dying of a heart attack, stroke or other circulatory problem

Actual Total CVD Risk Level | What makes up your risk | Personalized health advice

### Your results

Examination date 13 September 2016

Patient name NJ

Age 50 (9/1966)

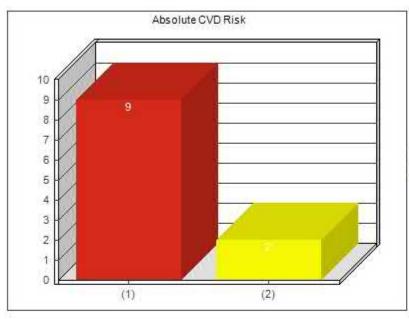
Sex Male

Risk factors	Your results	What you should aim for:	
Systolic blood pressure	sure 181 140 or less		
Cholesterol	6.5 mmol/L	5 mmol/L or less	
HDL Cholesterol	1.3 mmol/L	Greater than 1 mmol/L	
Smoker	Yes	No	

### Actual Total CVD Risk Level

The total cardiovascular disease risk level (left bar below) shows you the percentage risk of having a fatal cardiovascular event, such as a stroke or heart attack. Based on examination results, your total CVD risk is 9%.

However, by becoming aware of your risk factors and taking a few preventive actions, you can reach the treatment goals and reduce your risk to 2% as shown by the treatment goal level (right bar below).



- (1) Your current risk is 9%
- (2) Your risk if you reach your treatment goals will approach 2%

At present, your risk of dying from a heart attack or a stroke within the next ten years is increased. You can reduce this risk further by becoming aware of your risk factors and by changing your lifestyle.

Your Risk Age: because of your risk factors your risk is similar to a 80 year old person with no risk factors, this is called your risk age. You can reduce your risk age by reducing your risk factors

## What makes up your risk

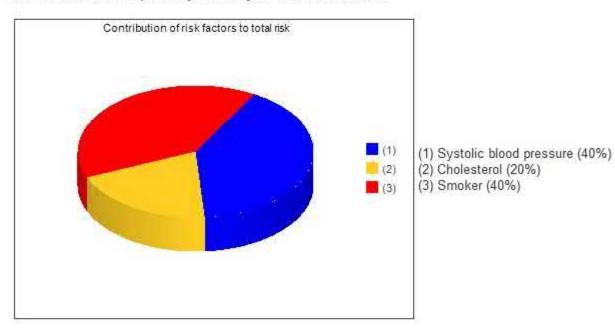
Cardiovascular disease is generally due to a combination of several risk factors. The more risk factors you have, the greater the chance of having a heart attack or stroke. The pie chart below shows you the distribution of your modifiable risk factors and the impact they have on your total CVD risk level.

and nak taraler by becoming aware or your nak factors and by changing your mestyle.

Your Risk Age: because of your risk factors your risk is similar to a 80 year old person with no risk factors, this is called your risk age. You can reduce your risk age by reducing your risk factors

## What makes up your risk

Cardiovascular disease is generally due to a combination of several risk factors. The more risk factors you have, the greater the chance of having a heart attack or stroke. The pie chart below shows you the distribution of your modifiable risk factors and the impact they have on your total CVD risk level.



#### Personalized health advice

For most people walking 30 minutes per day and eating plenty of of vegetables, fruit, cereals and fish helps lower risk.

#### Smoker

You are noted to be a smoker. If you can stop smoking this would greatly reduce your risk. Many smokers who want to quit find nicotine chewing gum and patches helpful

Smoking increases your risk of many diseases: it is an extra good reason to quit. If you can manage to stop, you will have halved your risk of a heart attack or stroke: no drug is this good at reducing risk. I will do all that I can to help you.

which evaluate CVD risk for patients between 40

and 65, with SBP between 100 - 180 mmHg and Cholesterol between 3 - 8 mmol/L (105 - 305

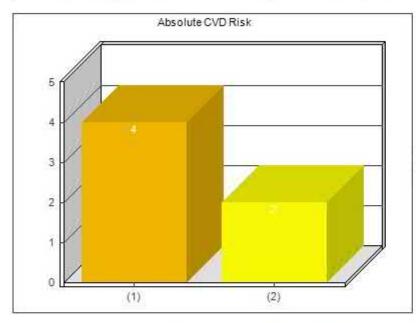
Risk factors Your results What you should aim for: 140 05 1000

Patient p

#### Actual Total CVD Risk Level

The total cardiovascular disease risk level (left bar below) shows you the percentage risk of having a fatal cardiovascular event, such as a stroke or heart attack. Based on examination results, your total CVD risk is 4%.

However, by becoming aware of your risk factors and taking a few preventive actions, you can reach the treatment goals and reduce your risk to 2% as shown by the treatment goal level (right bar below).



- (1) Your current risk is 4%
- (2) Your risk if you reach your treatment goals will approach 2%

At present, your risk of dying from a heart attack or a stroke within the next ten years might seem low, but this risk is 3 times higher than it could be. You can reduce this risk further by becoming aware of your risk factors and by changing your lifestyle.

Your Risk Age: because of your risk factors your risk is similar to a 64 year old person with no risk factors, this is called your risk age. You can reduce your risk age by reducing your risk factors

### What makes up your risk

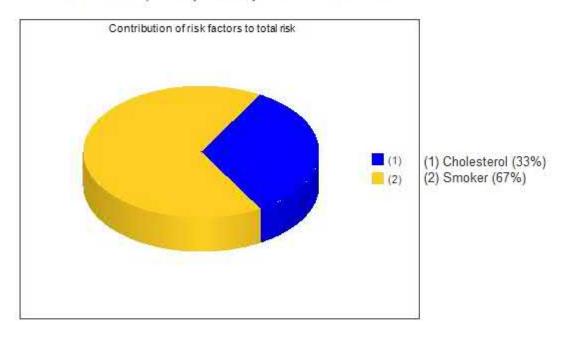
Cardiovascular disease is generally due to a combination of several risk factors. The more risk factors you have, the greater the chance of having a heart attack or stroke. The pie chart below shows you the distribution of your modifiable risk factors and the impact they have on your total CVD risk level.

3 times higher than it could be. You can reduce this risk further by becoming aware of your risk factors and by changing your lifestyle.

Your Risk Age: because of your risk factors your risk is similar to a 64 year old person with no risk factors, this is called your risk age. You can reduce your risk age by reducing your risk factors

### What makes up your risk

Cardiovascular disease is generally due to a combination of several risk factors. The more risk factors you have, the greater the chance of having a heart attack or stroke. The pie chart below shows you the distribution of your modifiable risk factors and the impact they have on your total CVD risk level.



### Personalized health advice

For most people walking 30 minutes per day and eating plenty of of vegetables, fruit, cereals and fish helps lower risk.

### Smoker

You are noted to be a smoker. If you can stop smoking this would greatly reduce your risk. Many smokers who want to quit find nicotine chewing gum and patches helpful

### Systolic blood pressure

Your blood pressure is 120 mmHg, and that is within the normal range.

#### Chalastaral

# The proper treatment of hypertension works...

- Reduction of stroke- 40%
- Reduction of acute coronary events- 20%
- Reduction of acute heart failure- 50%

Mancia G et al, Journal of Hypertension. 2007 Jun;25(6):1105-87.

## Hypertension Management

2013 Practice guidelines for the management of arterial hypertension of the European Society of Hypertension (ESH) and the European Society of Cardiology (ESC)

Journal of Hypertension 2013, 31:1281–1357

## **Blood pressure categories**

Category	SBP		DBP
Optimal	<120	and	<80
Normal	120-129	and/or	80-84
High normal	130-139	and/or	85-89
Grade 1 hypertension	140-159	and/or	90-99
Grade 2 hypertension	160-179	and/or	100-109
Grade 3 hypertension	≥180	and/or	≥110
Isolated systolic hypertension	≥140	and	<90

The blood pressure (BP) category is defined by the highest BP level, whether systolic blood pressure (SBP) or diastolic blood pressure (DBP). Isolated systolic hypertension should be graded 1, 2 or 3 according to SBP values in the ranges indicated.























Whelton PK, et al.

2017 High Blood Pressure Clinical Practice Guideline

## 2017 ACC/AHA/AAPA/ABC/ACPM/AGS/APhA/ASH/ASPC/NMA/PCNA Guideline for the Prevention, Detection, Evaluation, and Management of High Blood Pressure in Adults

A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines

### WRITING COMMITTEE MEMBERS

Paul K. Whelton, MB, MD, MSc, FAHA, Chair Robert M. Carey, MD, FAHA, Vice Chair

Wilbert S. Aronow, MD, FACC, FAHA\* Donald E. Casey, Jr, MD, MPH, MBA, FAHA† Karen J. Collins, MBA‡ Cheryl Dennison Himmelfarb, RN, ANP, PhD, FAHA§ Sondra M. DePalma, MHS, PA-C, CLS, AACC Samuel Gidding, MD, FACC, FAHA¶ Kenneth A. Jamerson, MD# Daniel W. Jones, MD, FAHA† Eric J. MacLaughlin, PharmD\*\*

Bruce Ovbiagele, MD, MSc, MAS, MBA, FAHA†

Sidney C. Smith, Jr, MD, MACC, FAHA++

Crystal C. Spencer, JD‡

Randall S. Stafford, MD, PhD‡‡

Sandra J. Taler, MD, FAHA§§

Randal J. Thomas, MD, MS, FACC, FAHA

Kim A. Williams, Sr, MD, MACC, FAHA†

Jeff D. Williamson, MD, MHS¶¶

Jackson T. Wright, Jr, MD, PhD, FAHA##



## 2017 High Blood Pressure Clinical Practice Guideline

## Table 6. Categories of BP in Adults\*

AHA 2017 HT guid... ×

		DBP
<120 mm Hg	and	<80 mm Hg
120–129 mm Hg	and	<80 mm Hg
	•	
130–139 mm Hg	or	80–89 mm Hg
≥140 mm Hg	or	≥90 mm Hg
	120–129 mm Hg  130–139 mm Hg  ≥140 mm Hg	120–129 mm Hg and 130–139 mm Hg or

BP indicates blood pressure (based on an average of ≥2 careful readings obtained on ≥2 occasions, as detailed in Section 4); DBP, diastolic blood pressure; and SBP systolic blood pressure.

## References

- Lewington S, Clarke R, Qizilbash N, et al. Age-specific relevance of usual blood pressure to vascular mortality: a
- meta-analysis of individual data for one million adults in 61 prospective studies. Lancet. 2002;360:1903-13. Rapsomaniki E, Timmis A, George J, et al. Blood pressure and incidence of twelve cardiovascular diseases: lifetim
- risks, healthy life-years lost, and age-specific associations in 1.25 million people. Lancet. 2014;383:1899-911. Ettehad D, Emdin CA, Kiran A, et al. Blood pressure lowering for prevention of cardiovascular disease and death: systematic review and meta-analysis. Lancet. 2016;387:957-67.
- Guo X, Zhang X, Guo L, et al. Association between pre-hypertension and cardiovascular outcomes: a systematic

# The NEW ENGLAND JOURNAL of MEDICINE

ESTABLISHED IN 1812

**NOVEMBER 26, 2015** 

VOL. 373 NO. 22

## A Randomized Trial of Intensive versus Standard Blood-Pressure Control

The SPRINT Research Group\*

#### ABSTRACT

#### BACKGROUND

The most appropriate targets for systolic blood pressure to reduce cardiovascular morbidity and mortality among persons without diabetes remain uncertain.

#### METHODS

We randomly assigned 9361 persons with a systolic blood pressure of 130 mm Hg or higher and an increased cardiovascular risk, but without diabetes, to a systolic blood-pressure target of less than 120 mm Hg (intensive treatment) or a target of less than 140 mm Hg (standard treatment). The primary composite outcome was myocardial infarction, other acute coronary syndromes, stroke, heart failure, or death from cardiovascular causes.

#### RESULTS

At 1 year, the mean systolic blood pressure was 121.4 mm Hg in the intensive-treatment group and 136.2 mm Hg in the standard-treatment group. The intervention was stopped early after a median follow-up of 3.26 years owing to a significantly lower rate of the primary composite outcome in the intensive-treatment group than in the standard-treatment group (1.65% per year vs. 2.19% per year; hazard ratio with intensive treatment, 0.75: 95% confidence interval [CI], 0.64 to

The members of the writing committee (Jackson T. Wright, Jr., M.D., Ph.D., Jeff D. Williamson, M.D., M.H.S., Paul K. Whelton, M.D., Joni K. Snyder, R.N., B.S.N., M.A., Kaycee M. Sink, M.D., M.A.S., Michael V. Rocco, M.D., M.S.C.E., David M. Reboussin, Ph.D., Mahboob Rahman, M.D., Suzanne Oparil, M.D., Cora E. Lewis, M.D., M.S.P.H., Paul L. Kimmel, M.D., Karen C. Johnson, M.D., M.P.H., David C. Goff, Jr., M.D., Ph.D., Lawrence J. Fine, M.D., Dr.P.H., Jeffrey A. Cutler, M.D., M.P.H., William C. Cushman, M.D., Alfred K. Cheung, M.D., and Walter T. Ambrosius, Ph.D.) assume responsibility for the overall content and integrity of the article. The affiliations of the members of the writing group are listed in the Appendix. Address reprint requests to Dr. Wright at the Division of Nephrology and Hypertension, University Hospitals Case Medical Center, Case

of 15% or greater on the basis of the Framingham risk score; or an age of 75 years or older. Patients with diabetes mellitus or prior stroke were excluded. Detailed inclusion and exclusion criteria are listed in the Supplementary Appendix. All participants provided written informed consent. RANDOMIZATION AND INTERVENTIONS Eligible participants were assigned to a systolic blood-pressure target of either less than 140 mm Hg (the standard-treatment group) or less than 120 mm Hg (the intensive-treatment group). Randomization was stratified according to clinical site. Participants and study personnel were aware of the study-group assignments, but outcome adjudicators were not.

iar risk was defined by one of more of the for-

lowing: clinical or subclinical cardiovascular dis-

ease other than stroke; chronic kidney disease,

excluding polycystic kidney disease, with an estimated glomerular filtration rate (eGFR) of 20 to

less than 60 ml per minute per 1.73 m<sup>2</sup> of body-

surface area, calculated with the use of the four-

variable Modification of Diet in Renal Disease

equation; a 10-year risk of cardiovascular disease

After the participants underwent randomization, their baseline antihypertensive regimens

Omron Healthcare). Lifestyle modification was encouraged as part of the management strategy. Retention in the study and adherence to treatment were monitored prospectively and routinely throughout the trial.26

## STUDY MEASUREMENTS

Demographic data were collected at baseline. Clinical and laboratory data were obtained at

ment group were adjusted on a monthly basis to target a systolic blood pressure of less than 120 mm Hg. For participants in the standardtreatment group, medications were adjusted

role in the study.

to target a systolic blood pressure of 135 to 139 mm Hg, and the dose was reduced if systolic blood pressure was less than 130 mm Hg on a single visit or less than 135 mm Hg on two consecutive visits. Dose adjustment was based

on a mean of three blood-pressure measurements at an office visit while the patient was

seated and after 5 minutes of quiet rest; the measurements were made with the use of an

automated measurement system (Model 907,

Pharmaceuticals; helther company had any other

3 months and every 3 months thereafter. Medi-

cations for participants in the intensive-treat-

Participants were seen monthly for the first

# 2018 ESC/ESH Guidelines for the management of arterial hypertension

The Task Force for the management of arterial hypertension of the European Society of Cardiology (ESC) and the European Society of Hypertension (ESH)

Authors/Task Force Members: Bryan Williams\* (ESC Chairperson) (UK), Giuseppe Mancia\* (ESH Chairperson) (Italy), Wilko Spiering (The Netherlands), Enrico Agabiti Rosei (Italy), Michel Azizi (France), Michel Burnier (Switzerland), Denis L. Clement (Belgium), Antonio Coca (Spain), Giovanni de Simone (Italy), Anna Dominiczak (UK), Thomas Kahan (Sweden), Felix Mahfoud (Germany), Josep Redon (Spain), Luis Ruilope (Spain), Alberto Zanchetti<sup>†</sup> (Italy), Mary Kerins (Ireland), Sverre E. Kjeldsen (Norway), Reinhold Kreutz (Germany), Stephane Laurent (France), Gregory Y. H. Lip (UK), Richard McManus (UK), Krzysztof Narkiewicz (Poland), Frank Ruschitzka (Switzerland), Roland E. Schmieder (Germany), Evgeny Shlyakhto (Russia), Costas Tsioufis (Greece), Victor Aboyans (France), Ileana Desormais (France)

Table 3 Classification of office blood pressure and definitions of hypertension grade b

Category	Systolic (mmHg)		Diastolic (mmHg)
Optimal	<120	and	<80
Normal	120–129	and/or	80–84
High normal	130–139	and/or	85–89
Grade 1 hypertension	140–159	and/or	90–99
Grade 2 hypertension	160–179	and/or	100–109
Grade 3 hypertension	≥180	and/or	≥110
Isolated systolic hypertension <sup>b</sup>	≥140	and	<90

BP = blood pressure; SBP = systolic blood pressure.

The same classification is used for all ages from 16 years.

<sup>&</sup>lt;sup>a</sup>BP category is defined according to seated clinic BP and by the highest level of BP, whether systolic or diastolic.

blsolated systolic hypertension is graded 1, 2, or 3 according to SBP values in the ranges indicated.

# How to measure the blood pressure?

- Allow the patient to sit for 3-5 minutes before beginning BP measurement.
- Take at least two BP measurements spaced 1-2 min apart, and additional measurements if the first two are quite different.
- Repeated measurements in atrial fibrillation.
- Use standard bladder (12-13 cm), but have larger and smaller as well for large and thin arms.
- Measure at first visit BP 1 and 3 min after assumption of the standing position in elderly participants, diabetic patients and in other conditions in which orthostatic hypertension may be frequent or suspected.

## BLOOD PRESSURE MEASUREMENT 😪



### PREPARATION BEFORE TAKING BP

The portions should be sesting comfortably in a quiet environment for 5 minutes in u chair. The patient should have an empty bladder and not have eaten, ingested coffeine, smoked, or engaged in physical activity at least 30 minutes prior to the mansurement. There should be no talking during the procedure by the potient or observer.

Inflatable bledder width should be about 40% of arm circumberance and bladder length should be about 80-100% of the individual's arm discumfunction.

For esscultation, the lower edge of the cult should be 2-3 cm shove the abov crease and the bladder should be centered ever the brocked arresy.

### **BID YOU KNOW?**

Using too large a cuff loods to folsely low readings and using too small a cuff. folially high readings. Markings on the cuff deply indicate the ideal arm circurferences. couponigte for the cuff size.

### Ideally, use validated upper-arm electronic devices.

For electronic devices, apply the cuff as: recommended by the manufacturer and record the EF exactly as displayed on the automated device.



Empty bladder

Leas unoxessed

#### Auscultation

If only this method is available, the proparation is as above.

For ausculatotory measurements, the cuff should be at heart level. Increess the pressure repliefly to 30 run Hg chave the level at which the bracked or radial pulse is extinguished, plant the stathoscope head over the brochial orters. deflate the cuff by approximately 2 mmHg per heartheat, and deterrine symble (appearance of Korotkoff sounds) and diestobs (disappearance of Karatkoff sounds). If the Lorotkoff sounds persist towards zero, use the point of maffing of the sounds to indicate dissole by

> Record the BF to the closest 2 mmHq. Axid ternind digit preferance (rounding) up or down to a zero for the last digit).



### GOOD PRACTICE

On the initial visit, readings should be taken in each arm and the higher arm should be used for subsequent measurements.

lwo or more restlings should be taken of each visit and the mean calculated.

Clinical indications for HBPM or ABPM Suspicion of white-coat hypertension

Grade 1 hypertension in the office

High office BP in individuals without asymptomatic organ damage and at low total CV risk

Suspicion of masked hypertension

High normal BP in the office

Normal office BP in individuals with asymptomatic organ damage or at high total CV risk

Identification of white-coat effect in hypertensive patients

Considerable variability of office BP over the same or different visits

Autonomic, postural, postprandial, siesta-induced and druginduced hypotension

Elevated office BP or suspected preeclampsia in pregnant women Identification of true and false resistant hypertension

Specific indications for ABPM

Marked discordance between office BP and home BP

Assessment of dipping status

Suspicion of nocturnal hypertension or absence of dipping, such as in patients with sleep apnoea, CKD or diabetes Assessment of BP variability

ABPM, ambulatory blood pressure monitoring; BP, blood pressure; CKD, chronic kidney disease; CV, cardiovascular; HBPM, home blood pressure monitoring.

## Normal ranges of blood pressure

Category	SBP (mmHg)		DBP (mmHg)
Office BP	≥140	and/or	≥90
Ambulatory BP	2031000	1-11-11-11-11-11-11-11-11-11-11-11-11-1	1000
Daytime (or awake)	≥135	and/or	≥85
Night-time (or asleep)	≥120	and/or	≥70
24-h	≥130	and/or	>80
Home BP	≥135	and/or	>85

BP, blood pressure; DBP, diastolic blood pressure; SBP, systolic blood pressure.

# How often should BP be measured?

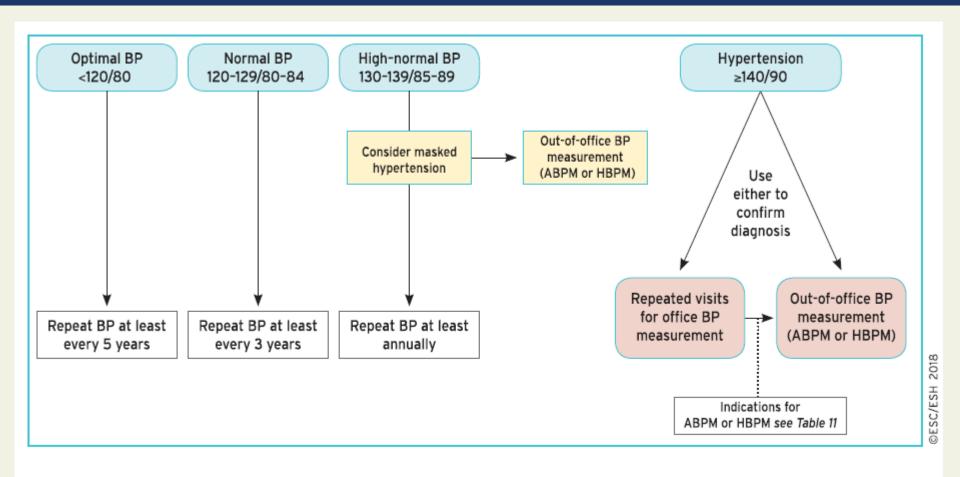


Figure 2 Screening and diagnosis of hypertension. ABPM = ambulatory blood pressure monitoring; BP = blood pressure; HBPM = home blood pressure monitoring.

## Personal and family medical history

- Duration and previous level of high BP, including measurements at home
- 2. Secondary hypertension
  - (a) Family history of CKD (polycystic kidney)
  - (b) History of renal disease, urinary tract infection, haematuria, analgesic abuse (parenchymal renal disease)
  - (c) Drug/substance intake, for example oral contraceptives, liquorice, carbenoxolone, vasoconstrictive nasal drops, cocaine, amphetamines, glucocorticosteroids and mineralocorticosteroids, NSAIDs, erythropoietin, cyclosporine
  - (d) Repetitive episodes of sweating, headache, anxiety, palpitations (pheochromocytoma)
  - (e) Episodes of muscle weakness and tetany (hyperaldosteronism)
  - (f) Symptoms suggestive of thyroid disease
- 3. Risk factors
  - (a) Family and personal history of hypertension and CVD
  - (b) Family and personal history of dyslipidaemia
  - (c) Family and personal history of diabetes mellitus (medications, blood glucose levels, polyuria).
  - (d) Smoking habits
  - (e) Dietary habits
  - (f) Recent weight changes; obesity
  - (g) Amount of physical exercise
  - (h) Snoring; sleep apnoea (information also from partner)
  - (i) Low birth weight.
- 4. History and symptoms of organ damage and CVD
  - (a) Brain and eyes: headache, vertigo, impaired vision, TIA, sensory or motor deficit, stroke, carotid revascularization
  - (b) Heart: chest pain, shortness of breath, swollen ankles, myocardial infarction, revascularization, syncope, history of palpitations, arrhythmias, especially atrial fibrillation
  - (c) Kidney: thirst, polyuria, nocturia, haematuria
  - (d) Peripheral arteries: cold extremities, intermittent claudication, pain-free walking distance, peripheral revascularization
  - (e) History of snoring/chronic lung disease/sleep apnoea
  - (f) Cognitive dysfunction
- 5. Hypertension management
  - (a) Current antihypertensive medication
  - (b) Past antihypertensive medication
  - (c) Evidence of adherence or lack of adherence to therapy
  - (d) Efficacy and adverse effects of drugs

## Signs suggesting secondary hypertension

features of Cushing syndrome
skin stigmata of neurofibromatosis (pheochromocytoma)
palpation of enlarged kidneys (polycystic kidney)
auscultation of abdominal murmurs (renovascular hypertension)
auscultation of precordial chest murmurs (aortic coarctation,
aortic disease: upper extremity artery disease)
left-right arm blood pressure difference (aortic coarctation,
subclavian artery stenosis)

## Laboratory, other investigations

Routine tests

Haemoglobin and/or haematocrit

Fasting plasma glucose

Serum total cholesterol, low-density lipoprotein cholesterol, high-density lipoprotein cholesterol

Fasting serum triglycerides

Serum potassium and sodium

Serum uric acid

Serum creatinine (with estimation of GFR),

Urine analysis: microscopic examination; urinary protein by dipstick test; test for microalbuminurea

12-lead EGG.

Additional tests, based on history, physical examination and findings from routine laboratory tests

Haemoglobin A<sub>1c</sub> [if fasting plasma glucose is >5.6 mmol/l (102 mg/dl) or previous diagnosis of diabetes]

Quantitative proteinuria (if dipstick test is positive); urinary potassium and sodium concentration and their ratio

Home and 24-h ABPM

Echocardiogram

Holter monitoring in case of arrhythmias

Carotid ultrasound

Peripheral artery/abdominal ultrasound

Pulse wave velocity

Ankle-brachial index

Fundoscopy

Extended evaluation (mostly domain of the specialist)

Further search for cerebral, cardiac, renal and vascular damage, mandatory in resistant and complicated hypertension

Search for secondary hypertension when suggested by history, physical examination or routine and additional tests

ABPM, ambulatory blood pressure monitoring; ECG, electrocardiogram; GFR, glomerular filtration rate.

## CV risk and blood pressure

Hypertension disease staging	Other risk factors, HMOD, or disease	BP (mmHg) grading					
		High normal SBP 130-139 DBP 85-89	Grade 1 SBP 140-159 DBP 90-99	Grade 2 SBP 160-179 DBP 100-109	Grade 3 SBP ≥180 or DBP ≥110		
Stage 1 (uncomplicated)	No other risk factors	Low risk	Low risk	Moderate risk	High risk		
	1 or 2 risk factors	Low risk	Moderate risk	Moderate to high risk	High risk		
	≥3 risk factors	Low to Moderate risk	Moderate to high risk	High Risk	High risk		
Stage 2 (asymptomatic disease)	HMOD, CKD grade 3, or diabetes mellitus without organ damage	Moderate to high risk	High risk	High risk	High to very high risk		
Stage 3 (established disease)	Established CVD, CKD grade ≥4, or diabetes mellitus with organ damage	Very high risk	Very high risk	Very high risk	Very high risk		

©ESC/ESH 2018

# Initiation of lifestyle changes and antihypertensive drug treatment

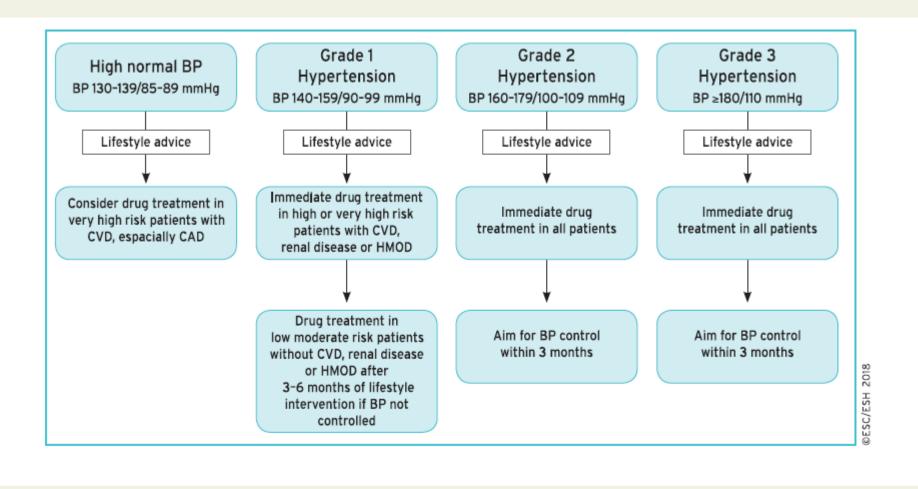


Figure 3 Initiation of blood pressure-lowering treatment (lifestyle changes and medication) at different initial office blood pressure levels. BP = blood pressure; CAD = coronary artery disease; CVD = cardiovascular disease; HMOD = hypertension-mediated organ damage.

### How to treat hypertension?

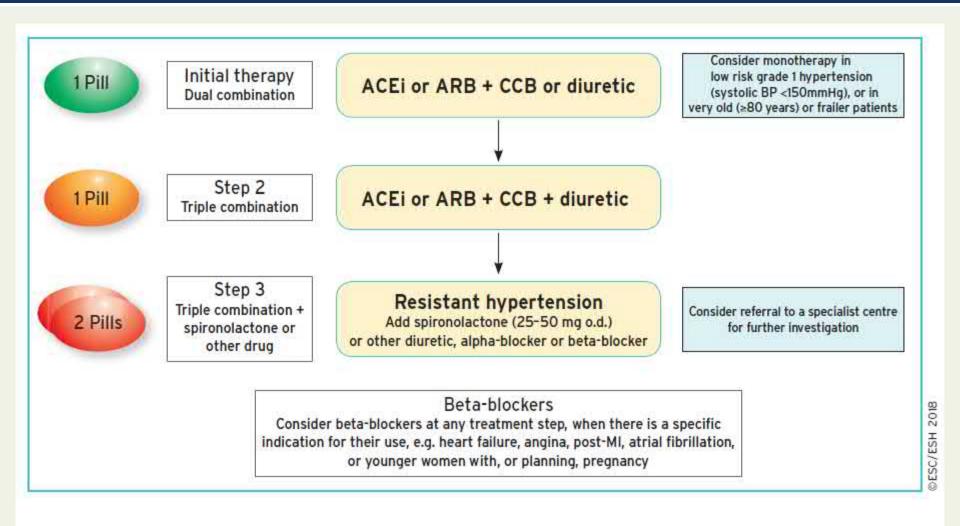


Figure 4 Core drug treatment strategy for uncomplicated hypertension. The core algorithm is also appropriate for most patients with HMOD, cerebrovascular disease, diabetes, or PAD. ACEi = angiotensin-converting enzyme inhibitor; ARB = angiotensin receptor blocker; CCB = calcium channel blocker; HMOD = hypertension-mediated organ damage; MI = myocardial infarction; o.d. = omni die (every day); PAD = peripheral artery disease.

### Treatment target ranges

Table 23 Office blood pressure treatment target range

Age group	Office SBP treatment target ranges (mmHg)							
	Hypertension	+ Diabetes	+ CKD	+ CAD	+ Stroke <sup>a</sup> /TIA			
18 - 65 years	Target to 130 or lower if tolerated Not <120	Target to 130 or lower if tolerated Not <120	Target to <140 to 130 if tolerated	Target to 130 or lower if tolerated Not <120	Target to 130 or lower if tolerated Not <120	70–79		
65 - 79 years <sup>b</sup>	Target to 130-139 if tolerated	Target to 130-139 if tolerated	Target to 130-139 if tolerated	Target to 130-139 if tolerated	Target to 130-139 if tolerated	70–79		
≥80 years <sup>b</sup>	Target to 130-139 if tolerated	Target to 130-139 if tolerated	Target to 130-139 if tolerated	Target to 130-139 if tolerated	Target to 130-139 if tolerated	70–79		
Office DBP treatment target range (mmHg)	70–79	70–79	70–79	70–79	70–79			

CAD = coronary artery disease; CKD = chronic kidney disease (includes diabetic and non-diabetic CKD); DBP = diastolic blood pressure; SBP = systolic blood pressure; TIA = transient ischaemic attack.

aRefers to patients with previous stroke and does not refer to blood pressure targets immediately after acute stroke.

<sup>&</sup>lt;sup>b</sup>Treatment decisions and blood pressure targets may need to be modified in older patients who are frail and independent.

### Lifestyle changes

- 1. salt restriction to 5–6 g/day;
- 2. moderation of alcohol consumption to no more than 20–30 g of ethanol per day in men and 10–20 g/day in women;
- 3. high consumption of vegetables and fruits and low fat dairy products;
- 4. reduction of weight to a BMI of 25 kg/m<sup>2</sup> and waist circumference to less than 102 cm in men and less than 88 cm in women;
- 5. at least 30 min of moderate dynamic exercise on 5 to 7 days per week.

# Compelling and possible contraindications to the use of antihypertensive drugs

Drug Compelling Possible

Diuretics Gout Metabolic syndrome;

glucose intolerance;

pregnancy; hypercalcaemia;

hypokalaemia

b-Blockers Asthma; A-V block Metabolic syndrome; glucose

intolerance; athletes and physically active patients; chronic obstructive

pulmonary disease (except for

vasodilator b-blockers)

**Calcium antagonists (dihydropyridines)** 

Calcium antagonists (verapamil, diltiazem)

Tachyarrhythmia; heart failure

A-V block (Grade 2 or 3, trifascicular block); severe left ventricular dysfunction; heart failure

# Compelling and possible contraindications to the use of antihypertensive drugs

**Drug** Compelling

Pregnancy; angioneurotic

oedema; hyperkalaemia;

bilateral renal artery stenosis

ARBs Pregnancy; hyperkalaemia;

bilateral renal artery

stenosis

**Possible** 

Women with childbearing

potential

Women with childbearing

potential

**Mineralocorticoid** 

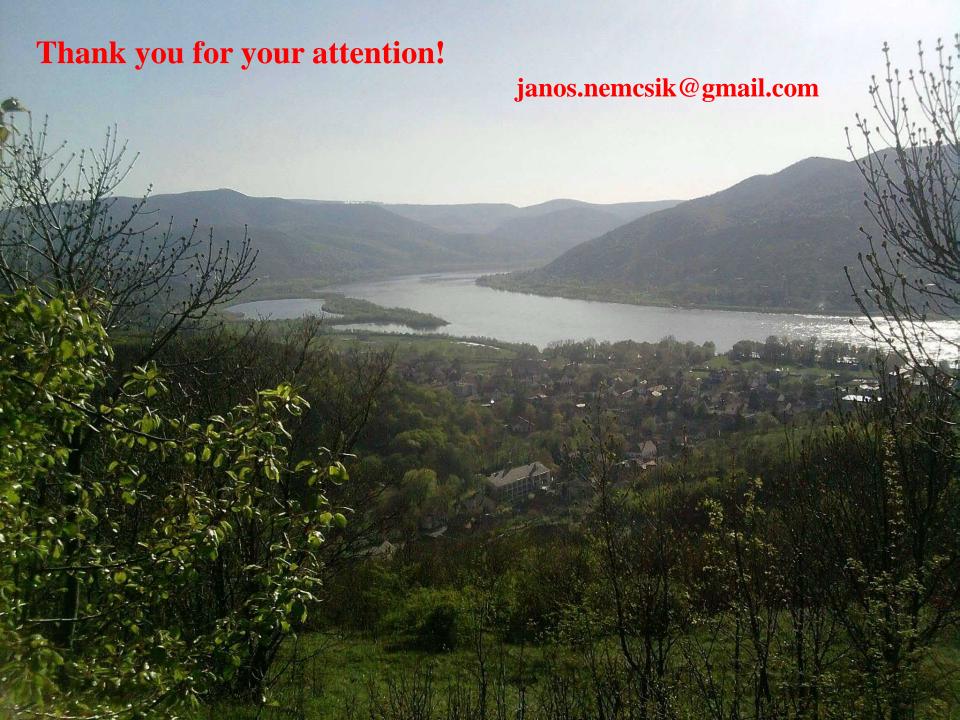
**ACE** inhibitors

receptor antagonists Acute or severe renal failure

(eGFR <30 ml/min); hyperkalaemia

## Main take-home messages

- keep the patients healthy;
- screen and treat hypertension, diabetes, dyslipidaemia and additional risk factors (smoking, anxiety, hyperuricaemia, obesity, alcohol...) to avoid cardiovascular events;
- be up-to-date



```
Risk factors
Male sex
Age (men ≥55 years; women ≥65 years)
Smoking
Dyslipidaemia
Total cholesterol >4.9 mmol/L (190 mg/dL), and/or
Low-density lipoprotein cholesterol >3.0 mmol/L (115 mg/dL),
and/or
High-density lipoprotein cholesterol: men <1.0 mmol/L
(40 \text{ mg/dL}), women <1.2 mmol/L (46 \text{ mg/dL}), and/or
Triglycerides >1.7 mmol/L (150 mg/dL)
Fasting plasma glucose 5.6–6.9 mmol/L (102–125 mg/dL)
Abnormal glucose tolerance test
Obesity [BMI ≥30 kg/m<sup>2</sup>]
Abdominal obesity (waist circumference: men ≥102 cm;
women ≥88 cm) (in Caucasians)
```

Family history of premature CVD (men aged <55 years; women aged <65 years)

#### Asymptomatic organ damage

Pulse pressure (in the elderly) ≥60 mmHg

Electrocardiographic LVH (Sokolow–Lyon index >3.5 mV;

RaVL >1.1 mV; Cornell voltage duration product >244

mV\*ms), or Echocardiographic LVH [LVM index: men

>115 g/m<sup>2</sup>; women >95 g/m<sup>2</sup> (BSA)]

Carotid wall thickening (IMT > 0.9 mm) or plaque

Carotid-femoral PWV >10 m/s

Ankle-brachial index < 0.9

Microalbuminuria (30–300 mg/24 h), or albumin–creatinine ratio (30–300 mg/g; 3.4–34 mg/mmol) (preferentially on morning spot urine)

#### **Diabetes mellitus**

Fasting plasma glucose  $\geq$ 7.0 mmol/L (126 mg/dL) on two repeated measurements, and/or

HbA1c > 7% (53 mmol/mol), and/or

Post-load plasma glucose >11.0 mmol/L (198 mg/dL)

#### **Established CV or renal disease**

Cerebrovascular disease: ischaemic stroke; cerebral haemorrhage; transient ischaemic attack CHD: myocardial infarction; angina; myocardial revascularization with PCI or CABG Heart failure, including heart failure with preserved EF Symptomatic lower extremities peripheral artery disease

CKD with eGFR <30 mL/min/1.73m<sup>2</sup>; proteinuria (>300 mg/24 h).

Advanced retinopathy: haemorrhages or exudates, papilloedema