



Treating Hypertension: New Guidelines

What number did they just say ?!?!

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Hypertension Overview

1 billion people affected world wide and 32 million in the US

Two-thirds of those above 60yo have hypertension

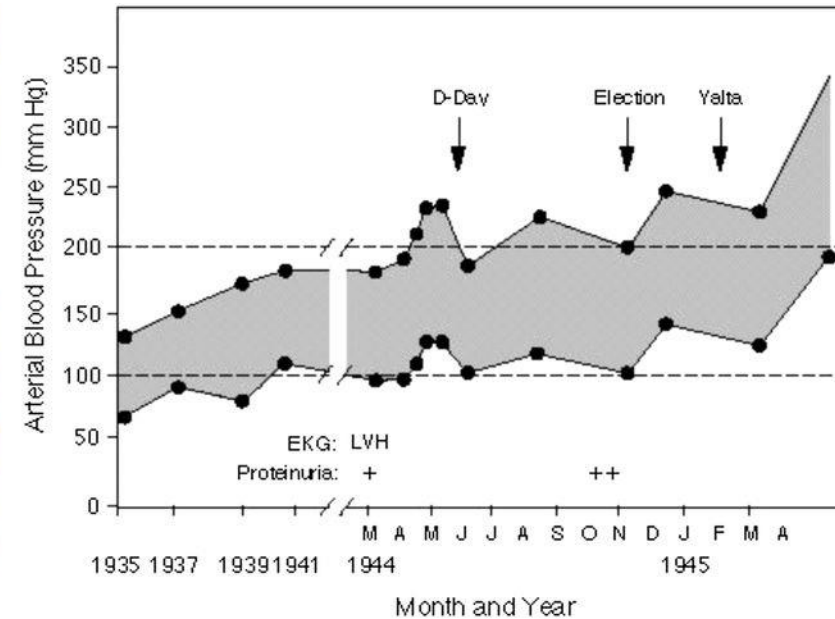
By 50yo, isolated systolic hypertension (ISH) is the most common form of hypertension and is associated with the greatest risk of organ injury

One of the most common causes of preventable death

Complications include heart disease, stroke, kidney disease, age-related dementia



Franklin D. Roosevelt, the 32nd president of the USA, shortly before his fatal cerebral haemorrhage (left) and his blood pressure during his presidency until his death (right; from Ref. 2).



Benefits of hypertension treatment: VA Cooperative Study Group

143 subjects were recruited

Baseline DBP 115 to 130mmHg

SBP typically 190 to 220 at baseline in both groups

Patients randomized to any drug or placebo

Methyl-dopa and:

Hydralazine

Reserpine

HCTZ

Subject were followed for 16 to 21 months

Treatment group achieved an average BP of 143/91

Event numbers:

Placebo group: 39%

Treatment group: 3%

Table 4.—Incidence of Mortality and Morbidity

| | Placebo-Treated Patients | Actively Treated Patients |
|------------------------------------|-----------------------------|------------------------------|
| Deaths | 4 | 0 |
| Class A events | 10 | 0 |
| Subtotal | 14 | 0 |
| Other treatment failures | 7 | 1 |
| Total terminating events | 21 | 1 |
| Class B events (nonterminating) | 6 | 1 |
| Total | 27 | 2 |

JAMA, Dec 11, 1967 • Volume 182, No 11



Benefits of hypertension treatment: Studies of “milder” hypertension

Medical Research Council (MRC) Study of hypertension

17,354 subjects recruited with baseline DBP between 90 and 109
Average BP was 159/98 at baseline

Subjects randomized to placebo or treatment
Treatment goal was DBP < 90
Thiazide diuretic or beta blocker used

Subjects followed for 5 years

Average blood pressure in the treatment group was 137/87



Benefits of hypertension treatment: Studies of “milder” hypertension

TABLE VI—Main events for both sexes together. Numbers and rates per 1000 patient years

| | Active treatment† | | Placebos | | % Difference‡ (95 % confidence limits) | Absolute difference/ 1000 patient years§ (95 % confidence limits) |
|----------------------------|-------------------|------|----------|------|--|---|
| | No | Rate | No | Rate | | |
| Strokes | | | | | | |
| Fatal | 18 | 0.4 | 27 | 0.6 | 34 | 0.2 |
| Non-fatal | 42 | 1.0 | 82 | 1.9 | 49 | 0.9 |
| Total | 60 | 1.4 | 109 | 2.6 | 45 (25, 60) | 1.2 (0.6, 1.7) |
| Coronary events | | | | | | |
| Fatal | 106 | 2.5 | 97 | 2.3 | -9 | -0.2 |
| Non-fatal | 116 | 2.7 | 137 | 3.2 | 16 | 0.5 |
| Total | 222 | 5.2 | 234 | 5.5 | 6 (-13, 21) | 0.3 (-0.7, 1.3) |
| All cardiovascular events* | 286 | 6.7 | 352 | 8.2 | 19 (5, 31) | 1.6 (0.4, 2.7) |
| All cardiovascular deaths | 134 | 3.1 | 139 | 3.3 | 4 (-22, 24) | 0.1 (-0.6, 0.9) |
| Non-cardiovascular deaths | 114 | 2.7 | 114 | 2.7 | 0 (-29, 23) | 0.0 (-0.7, 0.7) |
| All deaths | 248 | 5.8 | 253 | 5.9 | 2 (-16, 18) | 0.1 (-0.9, 1.2) |

*Not necessarily equal to the total of strokes plus coronary events because it also includes “other relevant deaths” and death due to other cardiovascular causes such as ruptured aneurysms.

†Randomised either to bendroflumazide or to propranolol.

‡Percentage difference between rates on active and on placebo therapy.

§Absolute difference between rates on active treatment and on placebo therapy.

Benefits of hypertension treatment: Studies of systolic hypertension in older patients

SHEP:

- 4700 subjects with isolated systolic hypertension (DBP < 90)
- Baseline BP 160-219/<90
- Randomized to SBP reduction of > 20mmHg
- Achieved avg BP 143/68
- 28% reduction in CHD, 36% reduction in CVA

Table 1. Antihypertensive Drug Trials: Risk Reductions for Coronary Heart Disease*

| | Active | | | Control | | | Risk Reductions, % (95% CI) | | P | |
|--------------------|--------|-------|--------|---------|-------|--------|--------------------------------|-----------|-------|-------|
| | Fatal | Total | N | Fatal | Total | N | Fatal | Total | Fatal | Total |
| 14 previous trials | 316 | 671 | 18 487 | 356 | 771 | 18 407 | 11 (-4-24) | 14 (4-22) | .12 | .007 |
| SHEP | 59 | 104 | 2365 | 74† | 142 | 2371 | 21 | 28 | .19 | .01 |
| STOP-Hypertension | 10 | 31 | 812 | 20 | 32 | 815 | 49 | 3 | .07 | .91 |
| MRC (older adults) | 85 | 128 | 2183 | 110 | 159 | 2213 | 22 | 19 | .08 | .08 |
| 17 trials | 470 | 934 | 23 847 | 560 | 1104 | 23 806 | 16 (5-26) | 16 (8-23) | .006 | .0001 |

*SHEP indicates Systolic Hypertension in the Elderly program; STOP, Swedish Trial in Old Patients; MRC, Medical Research Council; and CI, confidence interval.

†Includes one death due to myocardial infarction based on death certificate coding classification as a death of indeterminate cause in the SHEP final result.

Table 3. Antihypertensive Drug Trials: Risk Reductions for Total Vascular Mortality*

| | Active | | Control | | Risk Reductions, % (95% CI) | P |
|--------------------|-----------------|--------|-----------------|--------|--------------------------------|--------|
| | Vascular Deaths | N | Vascular Deaths | N | | |
| 14 previous trials | 489 | 18 487 | 613 | 18 407 | 21 (11-30) | .0002 |
| SHEP | 101† | 2365 | 130† | 2371 | 23 | .05 |
| STOP-Hypertension | 17 | 812 | 41 | 815 | 57 | .001 |
| MRC (older adults) | 161 | 2183 | 180 | 2213 | 10 | .35 |
| 17 trials | 768 | 23 847 | 964 | 23 806 | 21 (13-28) | <.0001 |

*See Table 1 footnotes for expansion of abbreviations.

†Includes 11 deaths in the active treatment group and 10 in the control group due to either cardiovascular diseases or unknown causes based on death certificate coding. These were classified as deaths of indeterminate cause in the SHEP final results.

STOP:

- 1600 subjects with SBP between 180 and 230
- Baseline BP 195/102
- Randomized to SBP drop of > 20mmHg
- Achieved avg BP 166/85
- 46% reduction in stroke, 43% reduction in mortality

JNC 7 Recommendations (2003)

| CLASSIFICATION OF BLOOD PRESSURE (BP)* | | | |
|--|----------|-----|----------|
| CATEGORY | SBP mmHg | | DBP mmHg |
| Normal | <120 | and | <80 |
| Prehypertension | 120–139 | or | 80–89 |
| Hypertension, Stage 1 | 140–159 | or | 90–99 |
| Hypertension, Stage 2 | ≥160 | or | ≥100 |

* See *Blood Pressure Measurement Techniques* (reverse side)

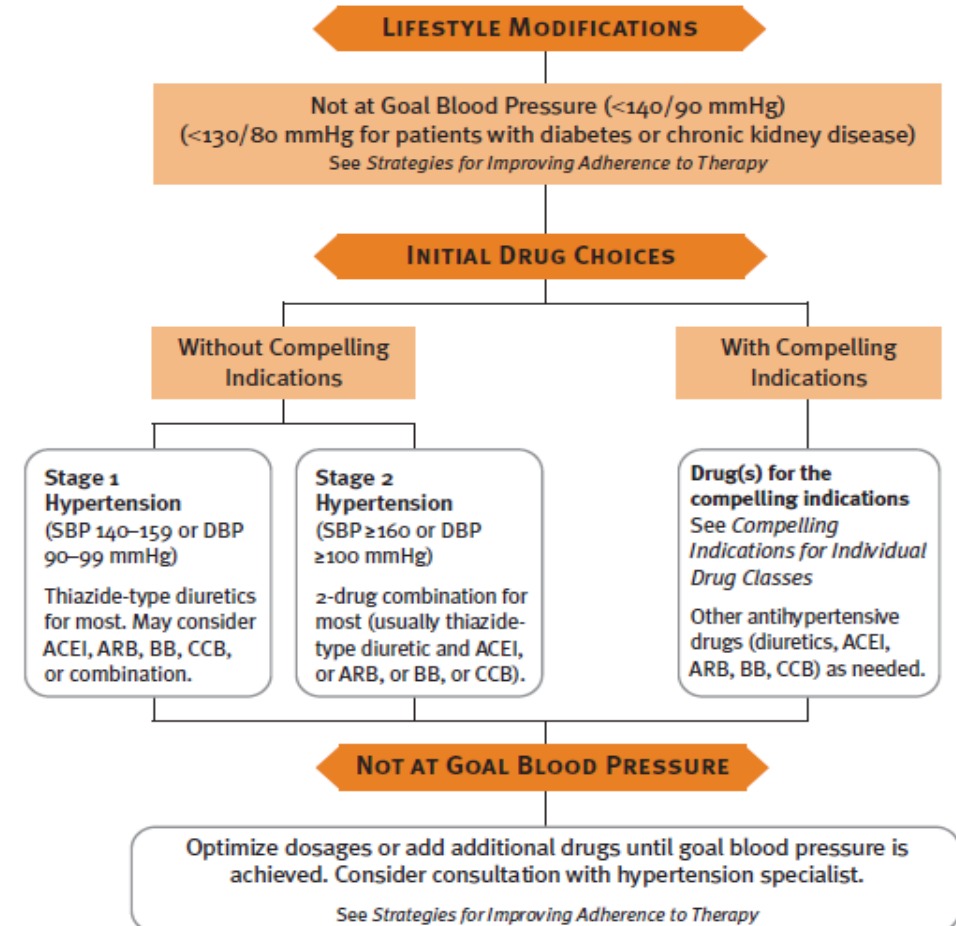
Key: SBP = systolic blood pressure DBP = diastolic blood pressure

TREATMENT

PRINCIPLES OF HYPERTENSION TREATMENT

- Treat to BP <140/90 mmHg or BP <130/80 mmHg in patients with diabetes or chronic kidney disease.
- Majority of patients will require two medications to reach goal.

ALGORITHM FOR TREATMENT OF HYPERTENSION





From: 2014 Evidence-Based Guideline for the Management of High Blood Pressure in Adults Report From the Panel Members Appointed to the Eighth Joint National Committee (JNC 8)

JAMA. 2014;311(5):507-520. doi:10.1001/jama.2013.284427

Table 1. Comparison of Current Recommendations With JNC 7 Guidelines

| Topic | JNC 7 | 2014 Hypertension Guideline |
|-------------------------------------|--|--|
| Methodology | Nonsystematic literature review by expert committee including a range of study designs Recommendations based on consensus | Critical questions and review criteria defined by expert panel with input from methodology team Initial systematic review by methodologists restricted to RCT evidence Subsequent review of RCT evidence and recommendations by the panel according to a standardized protocol |
| Definitions | Defined hypertension and prehypertension | Definitions of hypertension and prehypertension not addressed, but thresholds for pharmacologic treatment were defined |
| Treatment goals | Separate treatment goals defined for “uncomplicated” hypertension and for subsets with various comorbid conditions (diabetes and CKD) | Similar treatment goals defined for all hypertensive populations except when evidence review supports different goals for a particular subpopulation |
| Lifestyle recommendations | Recommended lifestyle modifications based on literature review and expert opinion | Lifestyle modifications recommended by endorsing the evidence-based Recommendations of the Lifestyle Work Group |
| Drug therapy | Recommended 5 classes to be considered as initial therapy but recommended thiazide-type diuretics as initial therapy for most patients without compelling indication for another class Specified particular antihypertensive medication classes for patients with compelling indications, ie, diabetes, CKD, heart failure, myocardial infarction, stroke, and high CVD risk Included a comprehensive table of oral antihypertensive drugs including names and usual dose ranges | Recommended selection among 4 specific medication classes (ACEI or ARB, CCB or diuretics) and doses based on RCT evidence Recommended specific medication classes based on evidence review for racial, CKD, and diabetic subgroups Panel created a table of drugs and doses used in the outcome trials |
| Scope of topics | Addressed multiple issues (blood pressure measurement methods, patient evaluation components, secondary hypertension, adherence to regimens, resistant hypertension, and hypertension in special populations) based on literature review and expert opinion | Evidence review of RCTs addressed a limited number of questions, those judged by the panel to be of highest priority. |
| Review process prior to publication | Reviewed by the National High Blood Pressure Education Program Coordinating Committee, a coalition of 39 major professional, public, and voluntary organizations and 7 federal agencies | Reviewed by experts including those affiliated with professional and public organizations and federal agencies; no official sponsorship by any organization should be inferred |

Abbreviations: ACEI, angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blocker; CCB, calcium channel blocker; CKD, chronic

kidney disease; CVD, cardiovascular disease; JNC, Joint National Committee; RCT, randomized controlled trial



Benefits of hypertension treatment:

Treatment goals

Based on these studies, reduction of blood pressure below 140/90 led to a 16% reduction in coronary events and a 40% reduction in stroke

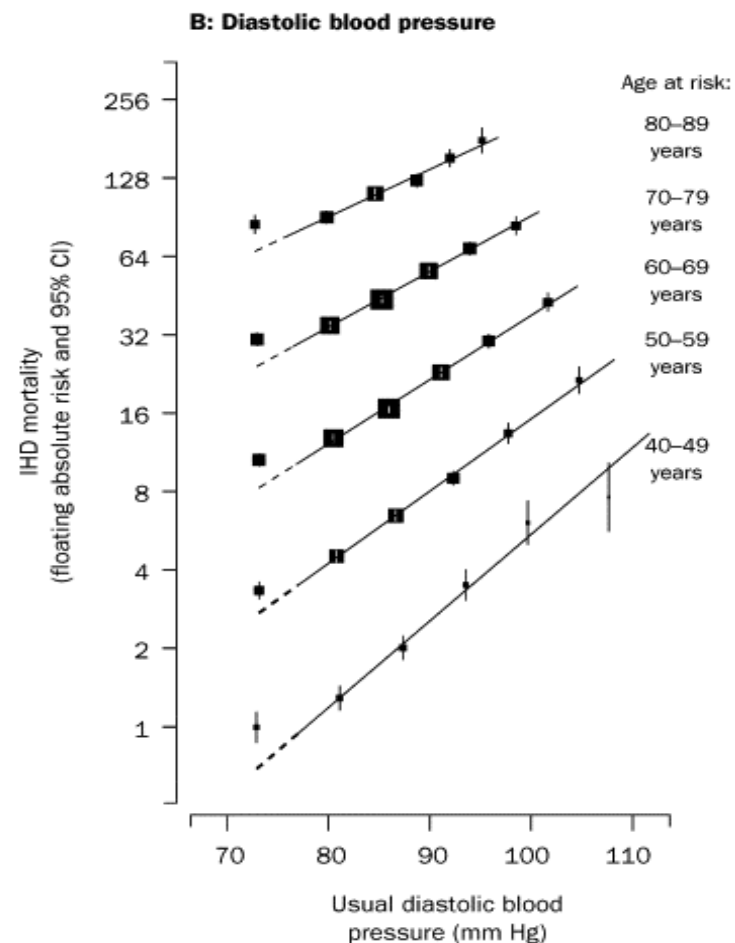
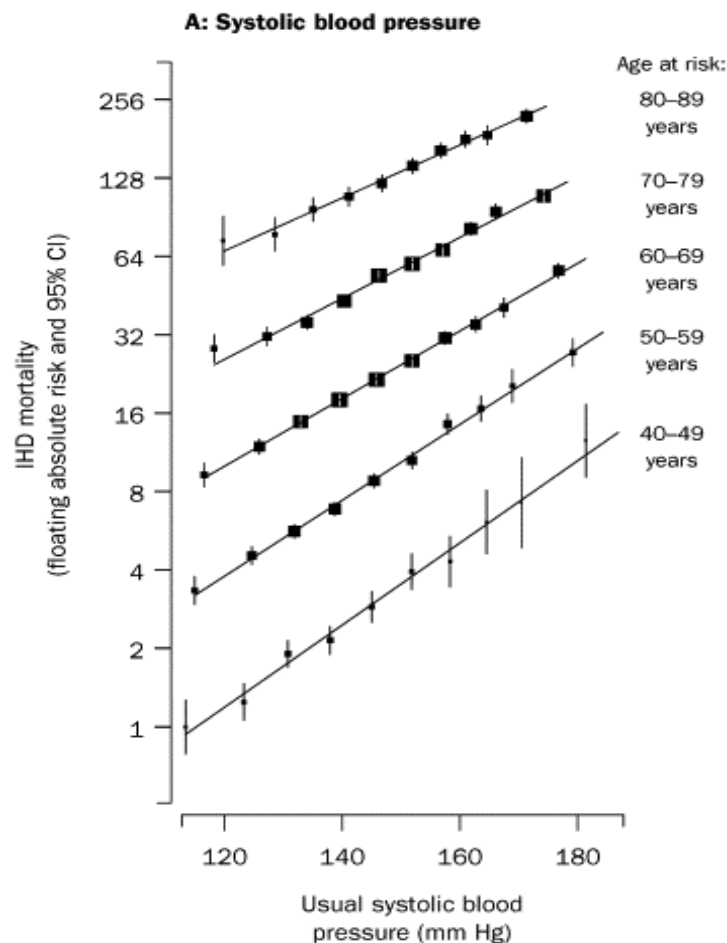
Previous treatment goals:

< 60yo - < 140 / < 90

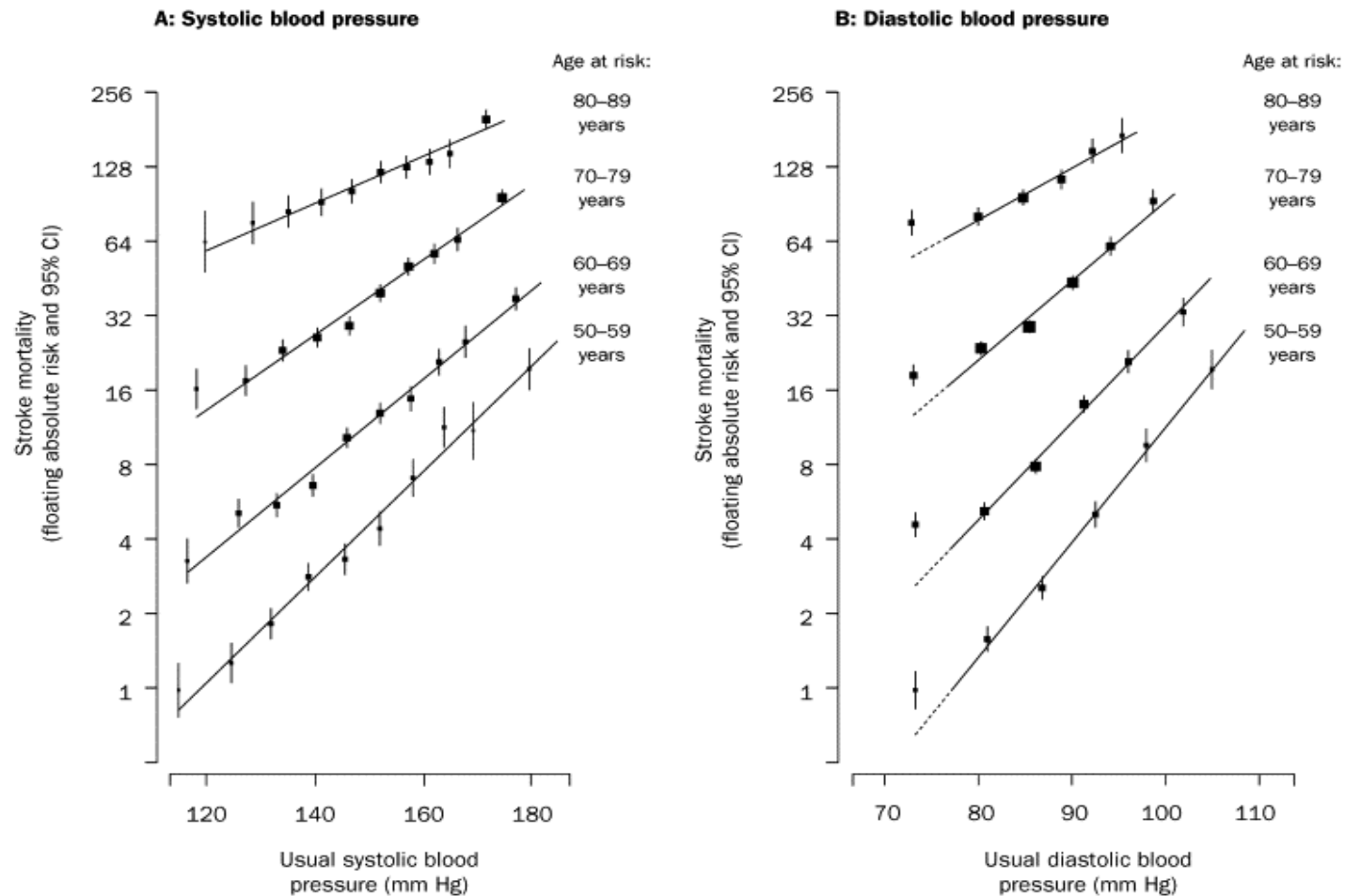
> 60yo - < 150 / < 90 except for certain populations such as chronic kidney disease with or without proteinuria or diabetics



Hypertension: What do observational studies say?



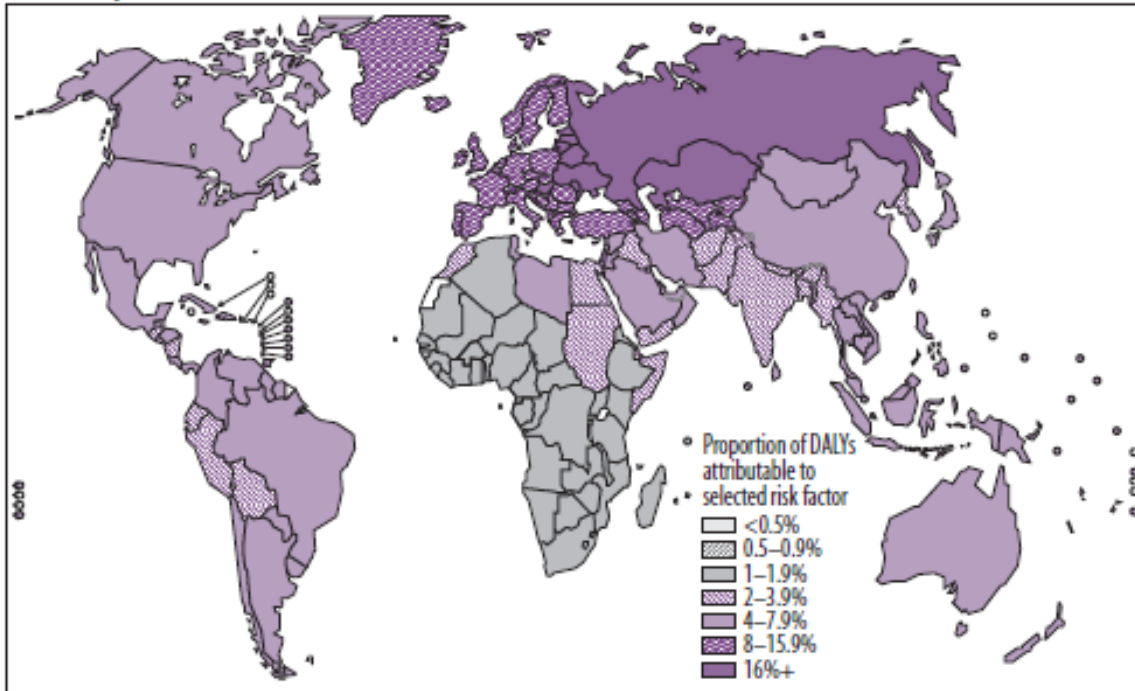
Hypertension: What do observational studies say?



Hypertension: What do observational studies say?

Figure 4.3 Burden of disease attributable to diet-related risk factors and physical inactivity
(% DALYs in each subregion)

A. Blood pressure



Impact of suboptimal
blood pressures
(SBP>115mmHg):

7.1 million (13%) deaths

49% ischemic heart
disease

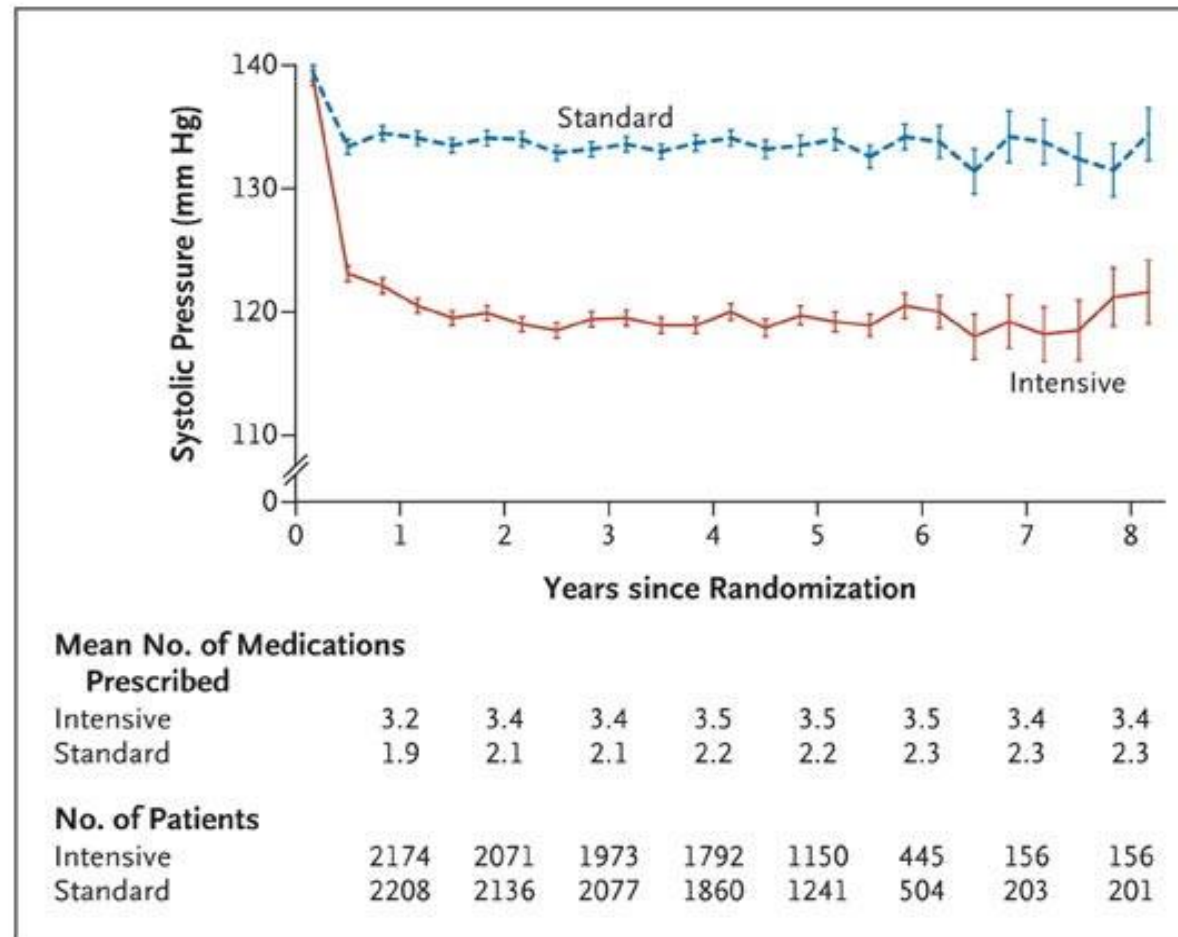
62% cerebrovascular
disease

64 million disability
adjusted life years lost

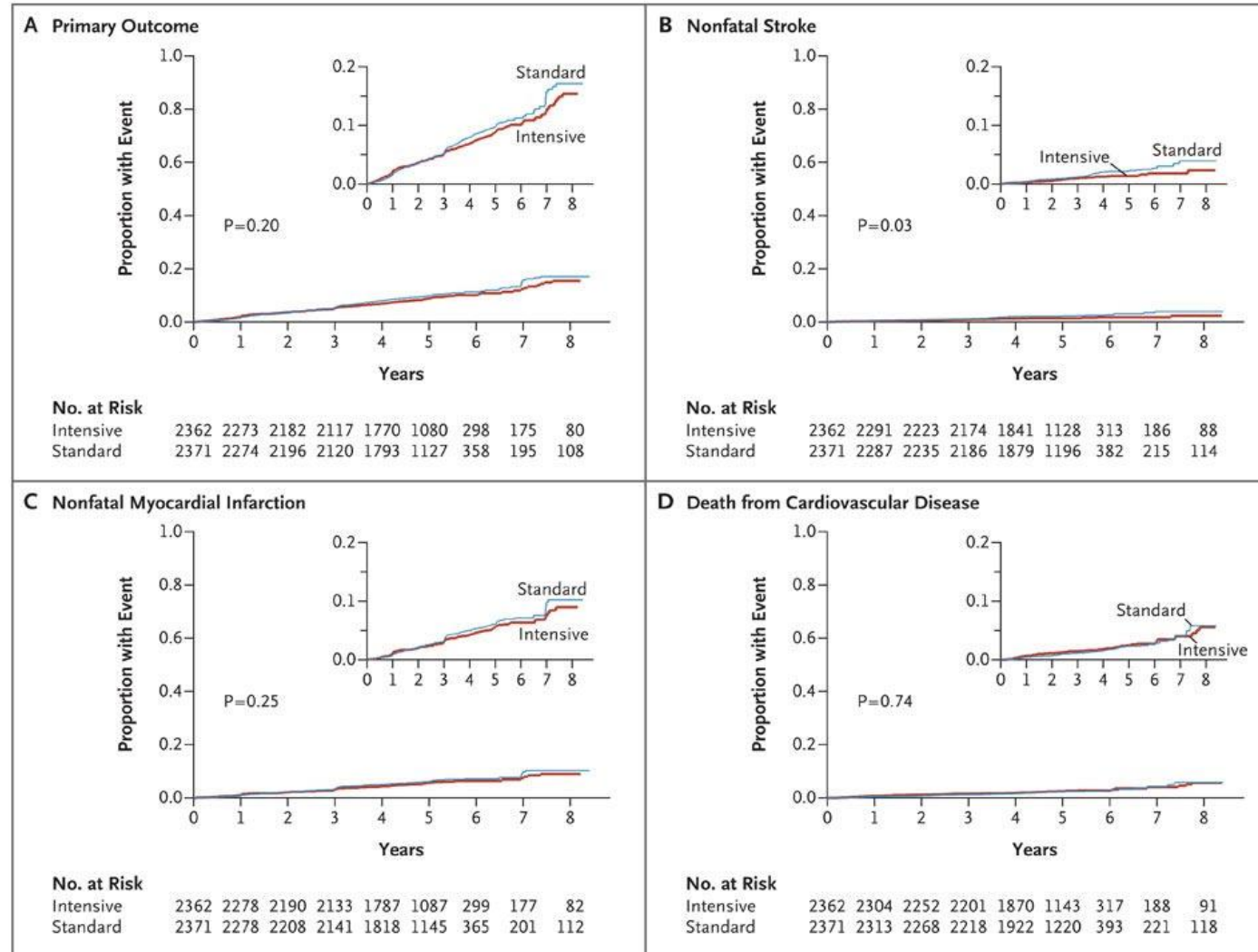
Half of these are
associated with a
SBP<145mmHg

Is an induced systolic blood pressure goal below 120mmHg beneficial?

The ACCORD Experience



Is a induced systolic blood pressure goal below 120mmHg beneficial? The ACCORD Experience





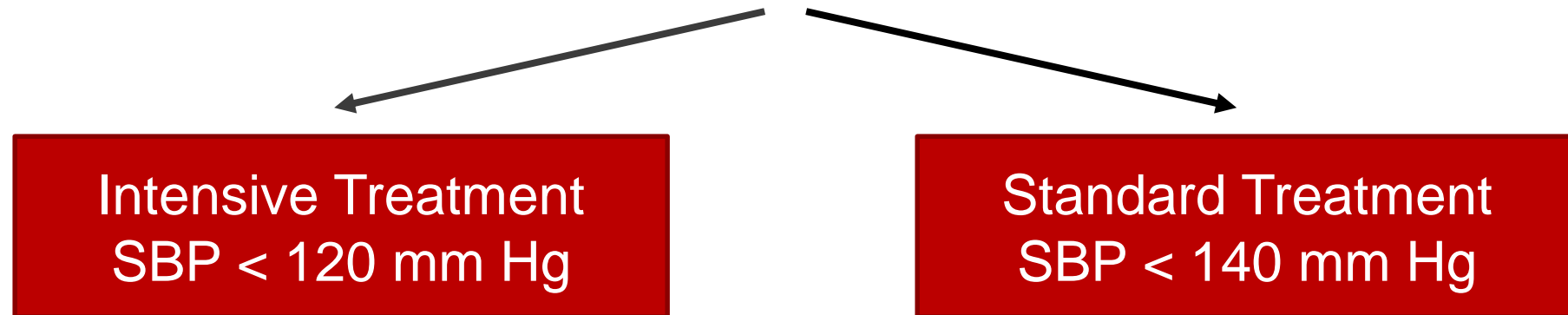
Systolic Blood Pressure Intervention Trial



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SPRINT Research Question

SPRINT was designed as a randomized controlled clinical trial to examine the effect of a more intensive high blood pressure treatment strategy than is currently recommended (standard treatment).



SPRINT Major Inclusion Criteria

≥50 years old

Systolic blood pressure : 130 – 180 mm Hg

Additional cardiovascular disease (CVD) risk (one or more of the following)

- Presence of clinical or subclinical CVD (not stroke)

- Chronic kidney disease (CKD), defined as eGFR 20 – 59 mL/min/1.73m²

- Framingham Risk Score for 10-year CVD risk ≥ 15%

 - Not needed if eligible based on preexisting CVD or CKD

- Age ≥ 75 years



SPRINT Major Exclusion Criteria

Stroke

Diabetes mellitus

Polycystic kidney disease

Congestive heart failure (symptoms or EF < 35%)

Proteinuria >1g/d

CKD with eGFR < 20 mL/min/1.73m² (MDRD)

Adherence flags



SPRINT Pre-specified Subgroups of Special Interest

Age (<75 vs. ≥75 years)

Gender (Men vs. Women)

Race/ethnicity (Black vs. non-Black)

Renal Disease (eGFR <60 vs. ≥60 mL/min/1.73m²)

CVD (CVD vs. no prior CVD)

Level of BP (Baseline SBP tertiles: ≤132, 132≤145, ≥145 mm Hg)



SPRINT Primary Outcome

The primary outcome was a composite of

Non-fatal myocardial infarction (MI)

Acute coronary syndrome not resulting in MI (non-MI ACS)

Non-fatal stroke

Non-fatal acute decompensated heart failure (HF)

Cardiovascular disease death



SPRINT Additional Outcomes

CVD secondary outcomes:

Individual categories of MI, non-MI ACS, all stroke, all heart failure, CVD mortality, all-cause mortality, primary outcome + all-cause mortality

Renal outcomes:

Main secondary outcome: >50% decline in eGFR or ESRD in CKD subgroup

Additional secondary outcomes:

Non-CKD subgroup

Incidence of decreased eGFR (>30% decrease in eGFR to <60 mL/min/1.73m²)

All trial participants

Incidence of albuminuria : doubling of urinary albumin/creatinine (<10 to >10mg/g)



SPRINT Additional Outcomes

Dementia and cognitive function outcomes:

Main secondary outcome: Incident dementia (all-cause)

Additional secondary outcomes:

Mild cognitive impairment

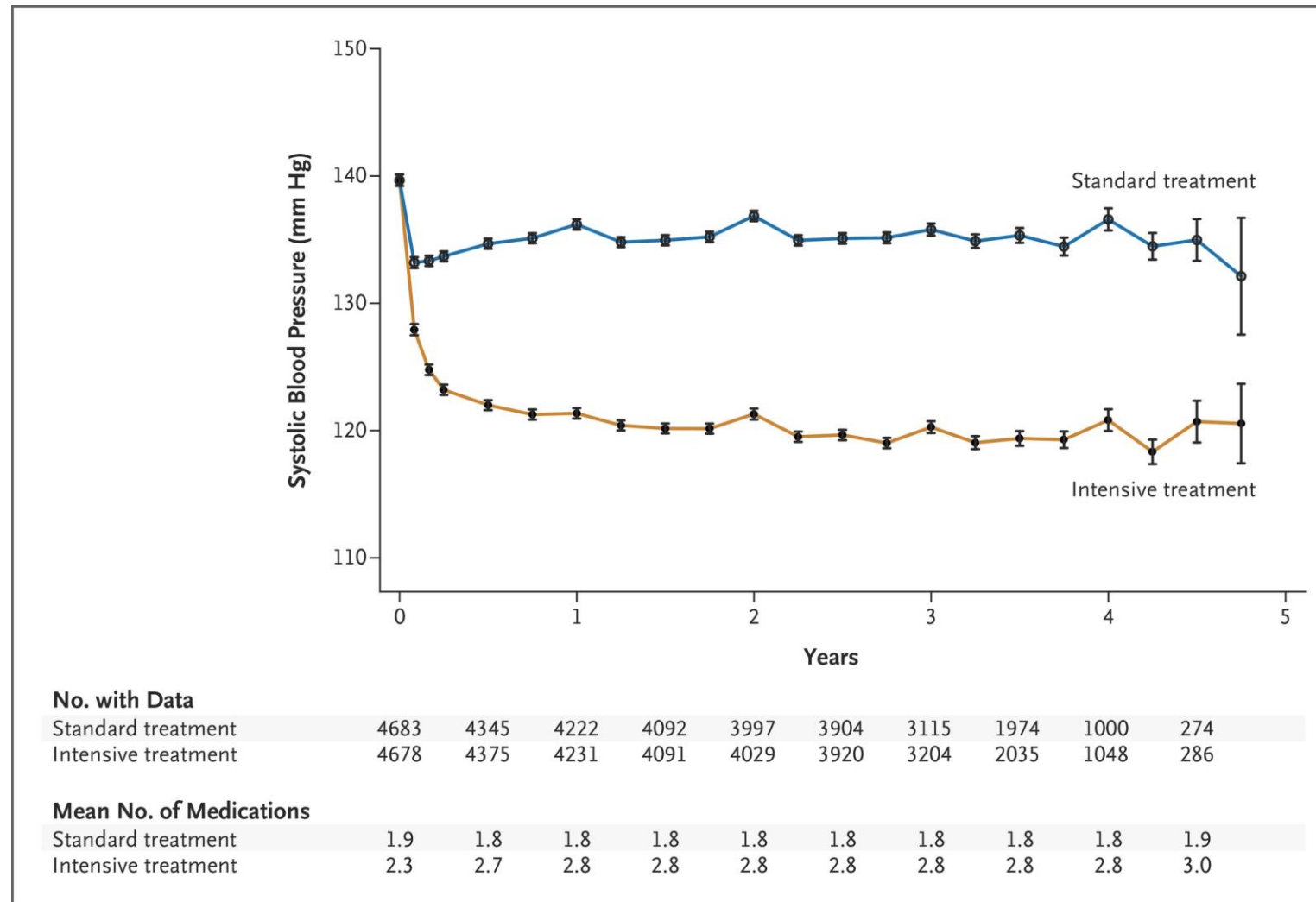
MRI

Health-related quality of life assessments

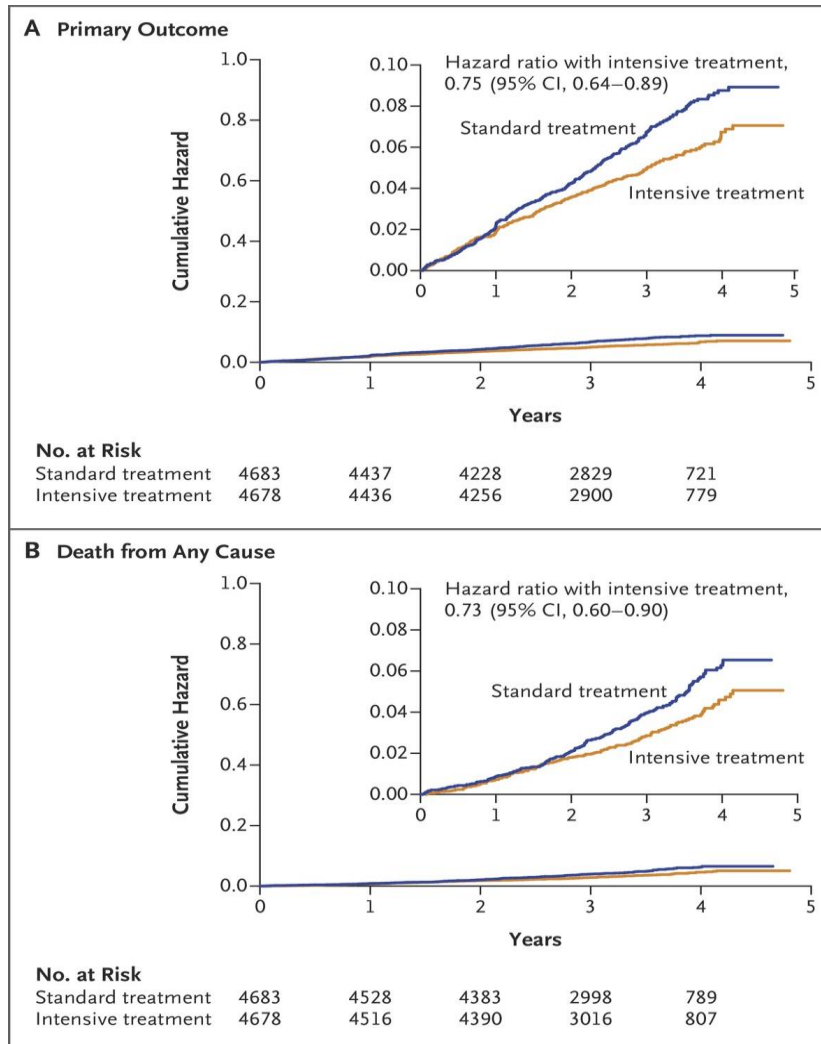
Other, including economic analyses



SPRINT: Systolic Blood Pressure in the Two Treatment Groups over the Course of the Trial



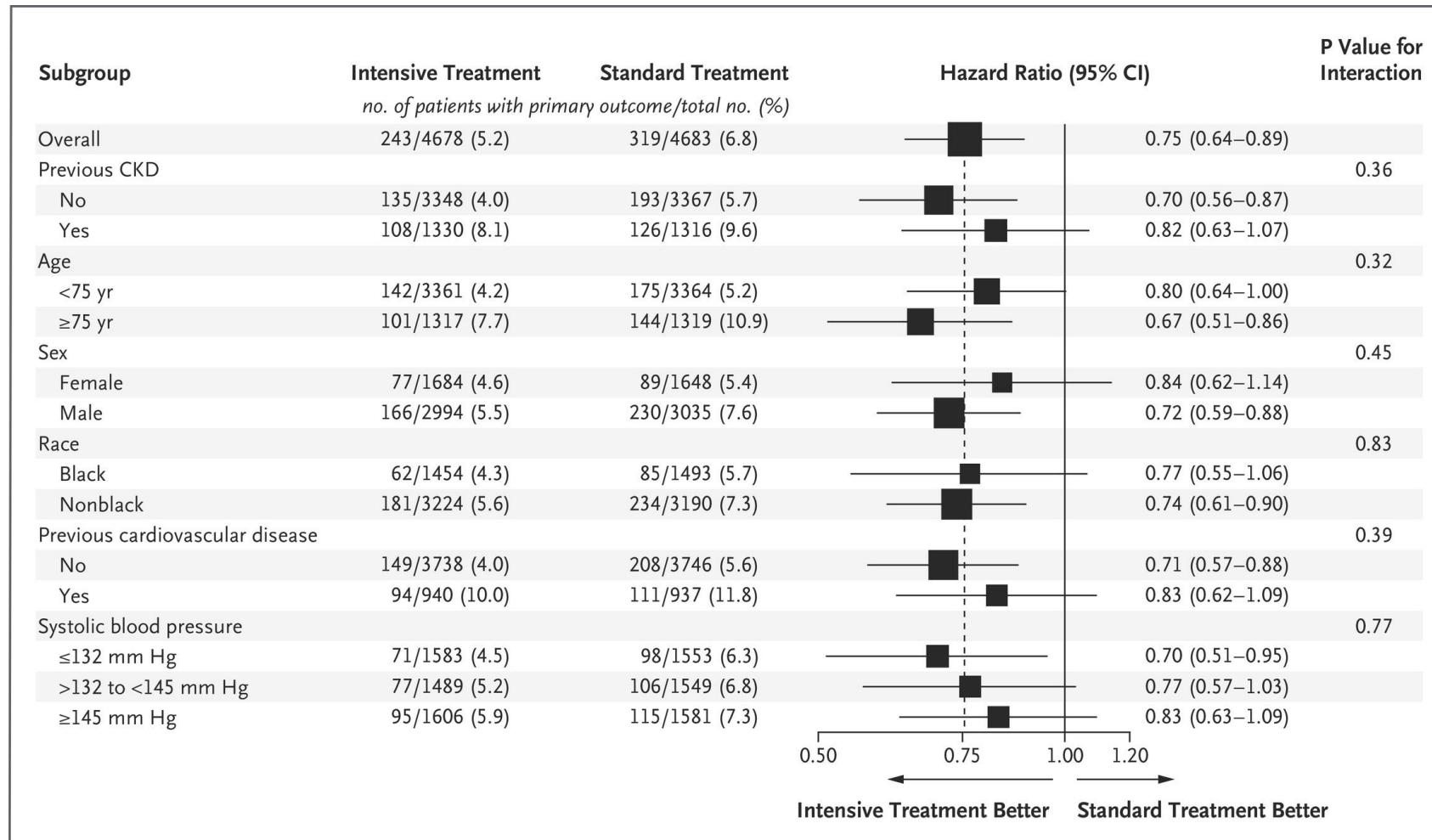
SPRINT: Primary Outcome and Death from Any Cause



SPRINT: Primary Outcome and its Components: Event Rates/Hazard Ratios

| | <i>Intensive</i> | | <i>Standard</i> | | | |
|------------------------|----------------------|---------------------|----------------------|---------------------|--------------------------|------------------|
| | <i>No. of Events</i> | <i>Rate, %/year</i> | <i>No. of Events</i> | <i>Rate, %/year</i> | <i>HR (95% CI)</i> | <i>P value</i> |
| Primary Outcome | 243 | 1.65 | 319 | 2.19 | 0.75 (0.64, 0.89) | <0.001 |
| All MI | 97 | 0.65 | 116 | 0.78 | 0.83 (0.64, 1.09) | 0.19 |
| Non-MI ACS | 40 | 0.27 | 40 | 0.27 | 1.00 (0.64, 1.55) | 0.99 |
| All Stroke | 62 | 0.41 | 70 | 0.47 | 0.89 (0.63, 1.25) | 0.50 |
| All HF | 62 | 0.41 | 100 | 0.67 | 0.62 (0.45, 0.84) | 0.002 |
| CVD Death | 37 | 0.25 | 65 | 0.43 | 0.57 (0.38, 0.85) | 0.005 |

SPRINT: Forest Plot of Primary Outcome According to Subgroups



SPRINT: Renal Disease Outcomes

| | | Intensive | | Standard | | HR (95% CI) | P |
|---|-------------------------------|------------------|-------------|-----------------|-------------|--------------------------|-------------|
| | | Events | %/yr | Events | %/yr | | |
| Participants with CKD at Baseline | | | | | | | |
| | Primary CKD outcome | 14 | 0.33 | 15 | 0.36 | 0.89 (0.42, 1.87) | 0.76 |
| | ≥50% reduction in eGFR* | 10 | 0.23 | 11 | 0.26 | 0.87 (0.36, 2.07) | 0.75 |
| | Dialysis | 6 | 0.14 | 10 | 0.24 | 0.57 (0.19, 1.54) | 0.27 |
| | Kidney transplant | 0 | - | 0 | - | - | . |
| | Secondary CKD Outcome | | | | | | |
| | Incident albuminuria** | 49 | 3.02 | 59 | 3.90 | 0.72 (0.48, 1.07) | 0.11 |
| | | | | | | | |
| Participants without CKD at Baseline | | | | | | | |
| | Secondary CKD outcomes | | | | | | |
| | ≥30% reduction in eGFR* | 127 | 1.21 | 37 | 0.35 | 3.48 (2.44, 5.10) | <.0001 |
| | | | | | | | |
| | Incident albuminuria** | 110 | 2.00 | 135 | 2.41 | 0.81 (0.63, 1.04) | 0.10 |

*Confirmed on a second occasion ≥90 days apart

**Doubling of urinary albumin/creatinine ratio from <10 to >10 mg/g

Serious Adverse Events* (SAE) During Follow-up

| | Number (%) of Participants | | |
|---|----------------------------|-------------|---------------|
| | Intensive | Standard | HR (P Value) |
| | 1793 (38.3) | 1736 (37.1) | 1.04 (0.25) |
| All SAE reports | | | |
| SAEs associated with Specific Conditions of Interest | | | |
| Hypotension | 110 (2.4) | 66 (1.4) | 1.67 (0.001) |
| Syncope | 107 (2.3) | 80 (1.7) | 1.33 (0.05) |
| Injurious fall | 105 (2.2) | 110 (2.3) | 0.95 (0.71) |
| Bradycardia | 87 (1.9) | 73 (1.6) | 1.19 (0.28) |
| Electrolyte abnormality | 144 (3.1) | 107 (2.3) | 1.35 (0.020) |
| Acute kidney injury or acute renal failure | 193 (4.1) | 117 (2.5) | 1.66 (<0.001) |

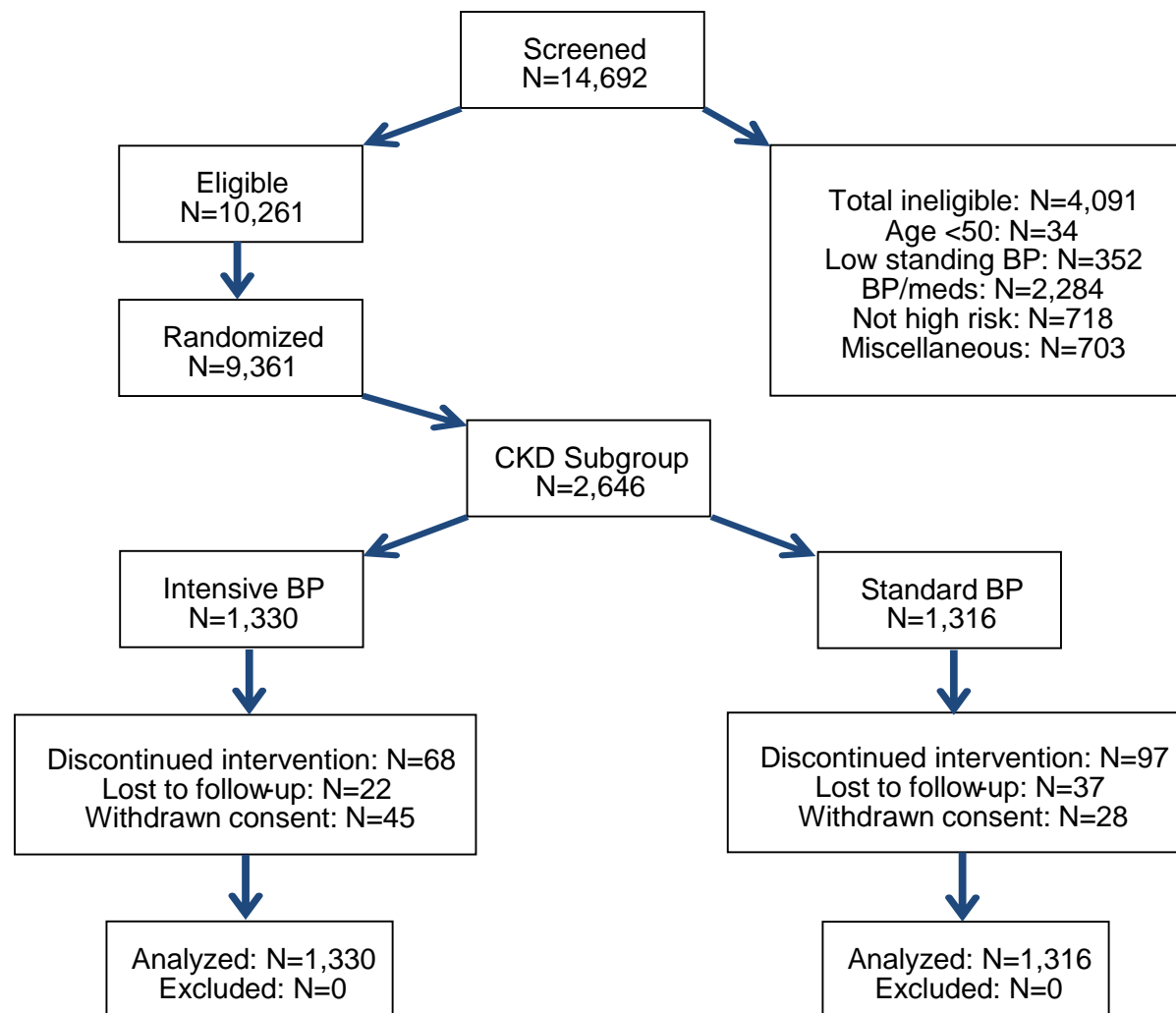
Number (%) of Participants with a Monitored Clinical Measure During Follow-up

| | Number (%) of Participants | | |
|---|----------------------------|------------|---------------|
| | Intensive | Standard | HR (P Value) |
| Laboratory Measures¹ | | | |
| Sodium <130 mmol/L | 180 (3.9) | 100 (2.2) | 1.76 (<0.001) |
| Potassium <3.0 mmol/L | 114 (2.5) | 74 (1.6) | 1.50 (0.006) |
| Potassium >5.5 mmol/L | 176 (3.8) | 171 (3.7) | 1.00 (0.97) |
| | | | |
| Signs and Symptoms | | | |
| Orthostatic hypotension² | 777 (16.6) | 857 (18.3) | 0.88 (0.013) |
| Orthostatic hypotension with dizziness | 62 (1.3) | 71 (1.5) | 0.85 (0.35) |

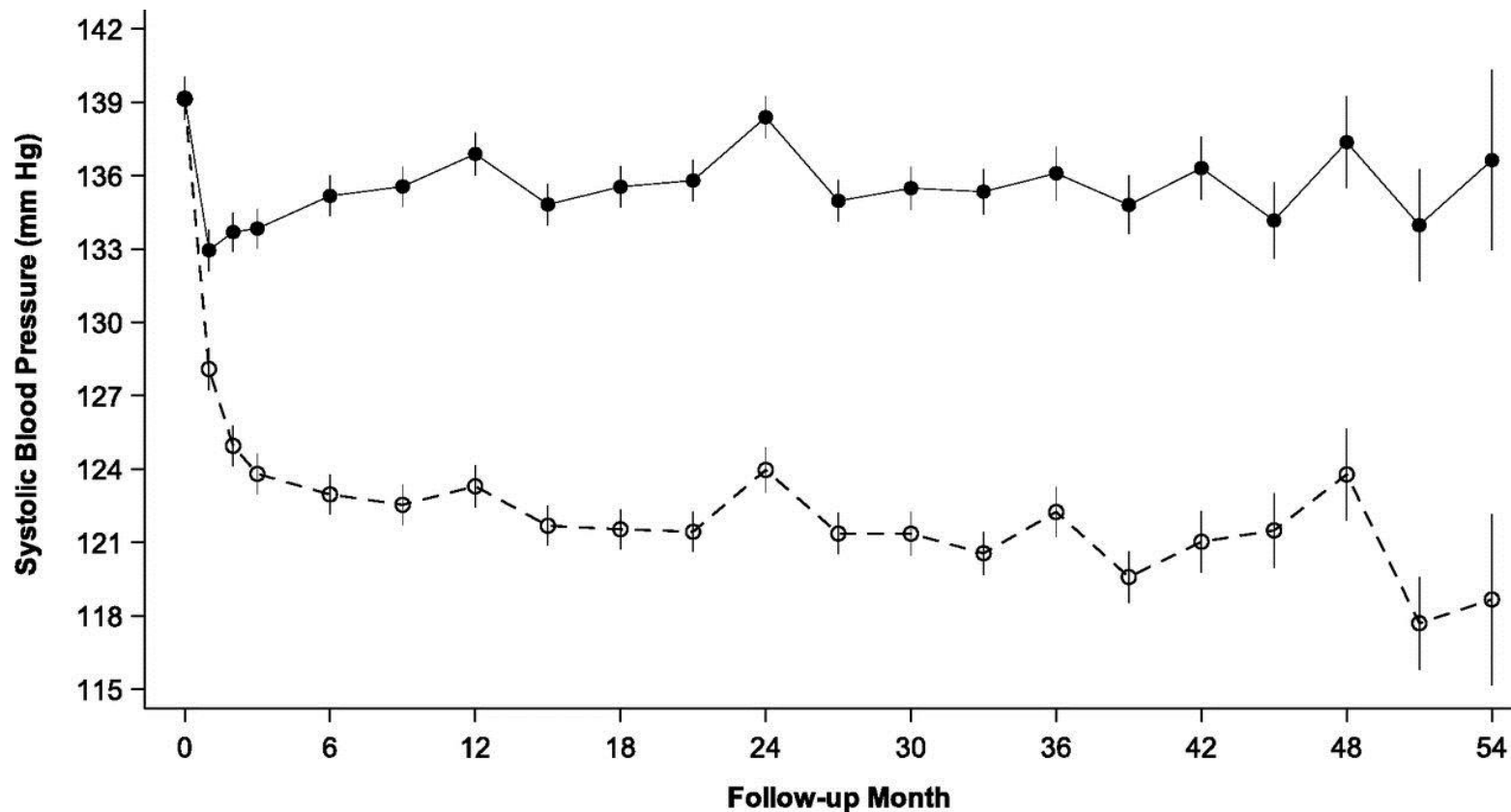
1. Detected on routine or PRN labs; routine labs drawn quarterly for first year, then q 6 months

2. Drop in SBP ≥20 mmHg or DBP ≥10 mmHg 1 minute after standing (measured at 1, 6, and 12 months and yearly thereafter)

SPRINT: SPRINT CKD Results



SPRINT: SPRINT CKD Results



| | Number With Data | | | | | | | | | |
|------------|---------------------|------|------|------|------|------|-----|-----|-----|-----|
| Standard: | 1316 | 1215 | 1156 | 1117 | 1087 | 1022 | 766 | 480 | 230 | 46 |
| Intensive: | 1330 | 1246 | 1194 | 1145 | 1136 | 1054 | 804 | 515 | 268 | 58 |
| | Mean Number of Meds | | | | | | | | | |
| Standard: | 2.1 | 2.0 | 2.0 | 2.0 | 2.1 | 2.0 | 2.1 | 2.1 | 2.1 | 2.0 |
| Intensive: | 2.1 | 2.9 | 3.0 | 3.0 | 3.0 | 3.0 | 2.9 | 2.9 | 3.0 | 3.1 |

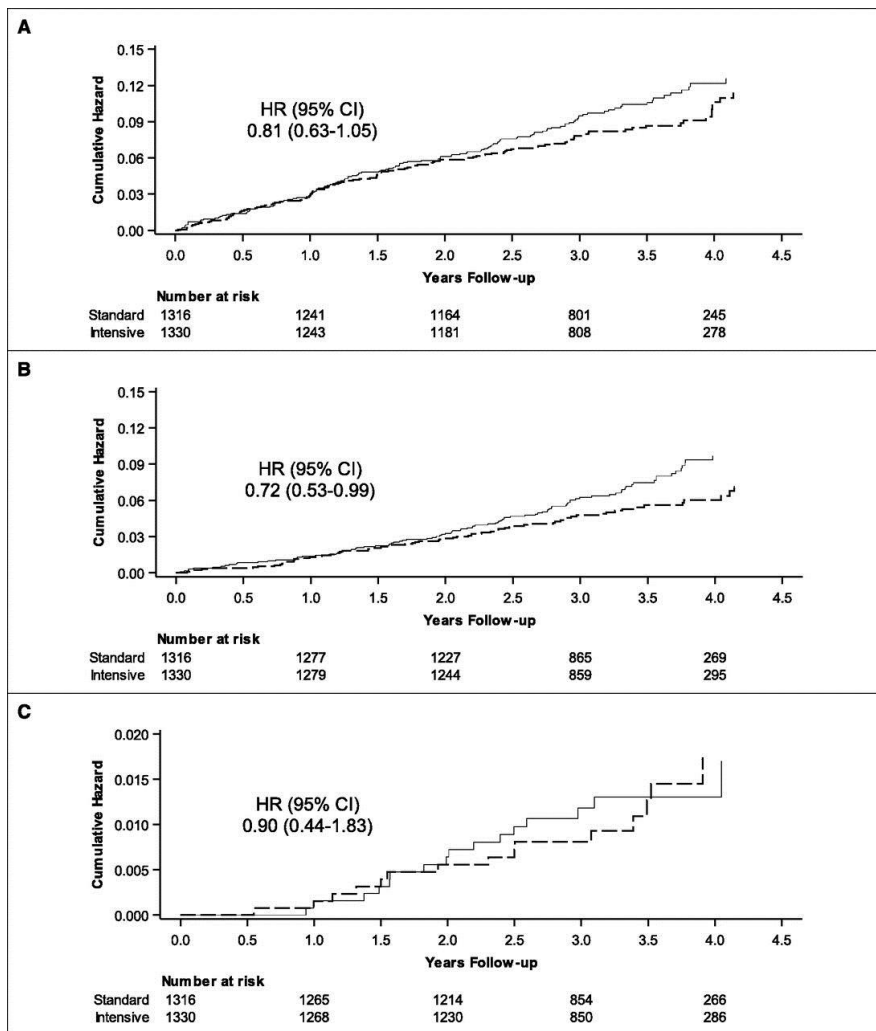


SPRINT: SPRINT CKD Results

| Outcome | Intensive Treatment, <i>n</i> =1330 | | Standard Treatment, <i>n</i> =1316 | | Intensive Treatment Versus Standard Treatment | |
|---|-------------------------------------|------|------------------------------------|------|---|----------------|
| | Events | %/yr | Events | %/yr | HR (95% CI) | <i>P</i> Value |
| Primary ^a outcome | 112 | 2.68 | 131 | 3.19 | 0.81 (0.63 to 1.05) | 0.12 |
| Myocardial infarction | 44 | 1.03 | 45 | 1.07 | 0.94 (0.62 to 1.44) | 0.79 |
| Acute coronary syndrome | 15 | 0.35 | 11 | 0.26 | 1.35 (0.60 to 3.08) | 0.47 |
| Stroke | 27 | 0.63 | 27 | 0.64 | 0.99 (0.57 to 1.70) | 0.96 |
| Heart failure | 41 | 0.96 | 52 | 1.24 | 0.72 (0.47 to 1.10) | 0.13 |
| CVD death | 18 | 0.41 | 30 | 0.70 | 0.57 (0.31 to 1.02) | 0.06 |
| All-cause death | 70 | 1.61 | 95 | 2.21 | 0.72 (0.53 to 0.99) | 0.04 |
| Primary outcome or all-cause death | 152 | 3.62 | 179 | 4.35 | 0.82 (0.66 to 1.02) | 0.08 |
| Primary outcome or cardiovascular procedure | 127 | 3.06 | 161 | 3.98 | 0.81 (0.63 to 1.05) | 0.12 |



SPRINT: SPRINT CKD Results



Kaplan-Meier curves for pre-specified outcomes in SPRINT participants with CKD.

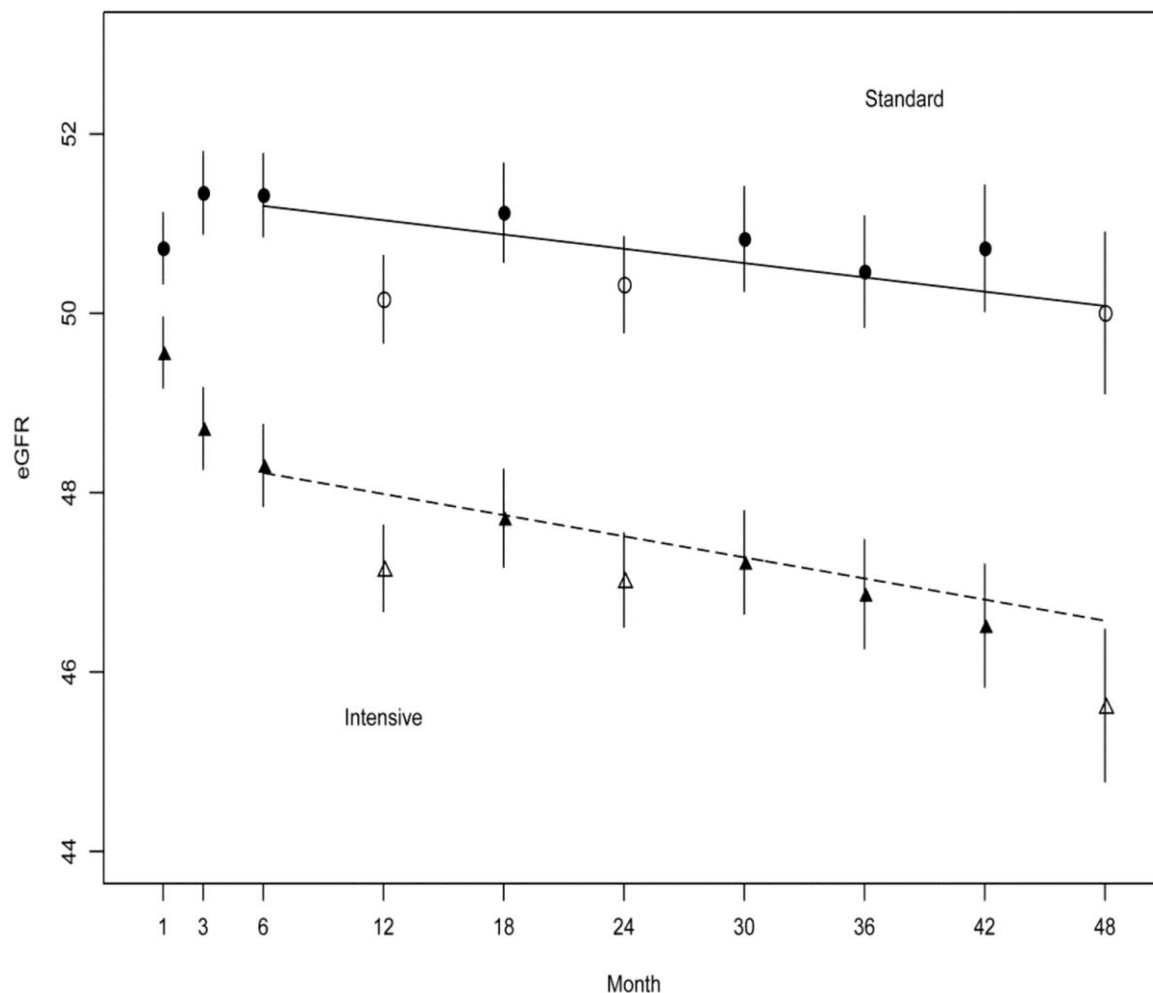
Panel A shows the primary cardiovascular outcome, defined as the composite of myocardial infarction, acute coronary syndrome, stroke, acute decompensated heart failure, and death from cardiovascular causes.

Panel B shows the all-cause death outcome.

Panel C shows the main kidney outcome, defined as the composite of a decrease in eGFR of $\geq 50\%$ from baseline (confirmed by repeat testing ≥ 90 days later) or the development of ESRD.

The broken lines depict the intensive group; the solid lines depict the standard group.

SPRINT: SPRINT CKD Results

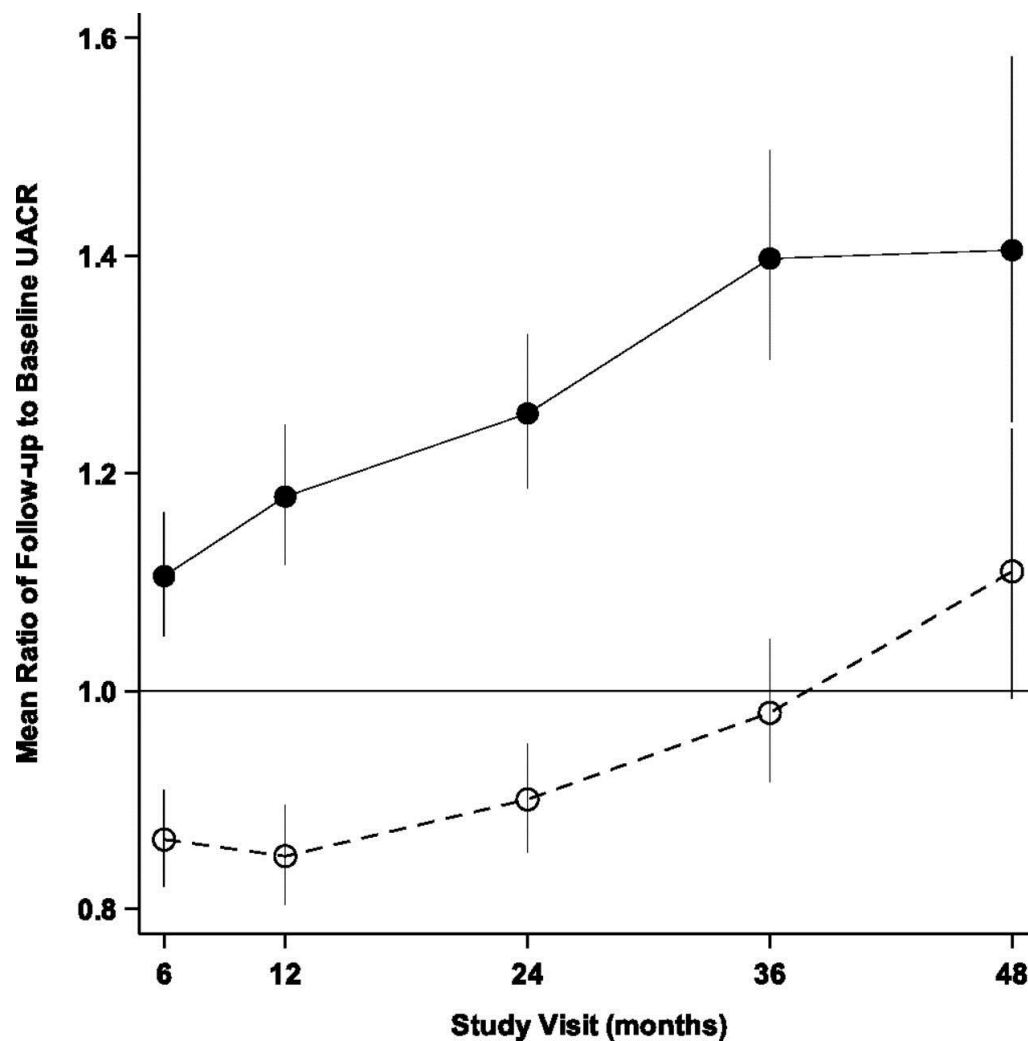


Two phases of eGFR changes during follow-up in the SPRINT participants with CKD. The rate of change in eGFR using the values at 6 months after randomization as the baseline was **-0.47 ml/min per 1.73 m² per year in the intensive group** (broken line and triangles) and **-0.32 ml/min per 1.73 m² per year in the standard group** (solid line and circles; $P=0.03$). Open symbols denote fasting visits; closed symbols denote nonfasting visits.

SPRINT: SPRINT CKD Results

| eGFR Reduction from Baseline, % ^a | No. of Events (% per 1 yr) ^b | | Intensive Treatment Versus Standard Treatment | |
|--|---|--|--|----------------|
| | Intensive Treatment, <i>n</i> =1330 | Standard Treatment, <i>n</i> =1316 | HR (95% CI) | <i>P</i> Value |
| 50 | 10 (0.25) | 12 (0.31) | 0.79 (0.34 to 1.83) | 0.58 |
| 40 | 30 (0.74) | 19 (0.49) | 1.51 (0.85 to 2.68) | 0.16 |
| 30 | 92 (2.33) | 44 (1.15) | 2.03 (1.42 to 2.91) | <0.01 |

SPRINT: SPRINT CKD Results



Urinary albumin-to-creatinine ratio (UACR) in the SPRINT participants with CKD. Geometric mean ratios of postrandomization to baseline UACR with 95% CIs. The broken line and open circles depict the intensive treatment group; the solid line and closed circles depict the standard treatment group. The horizontal line at 1.0 depicts equality of means (*i.e.*, no change in UACR).



SPRINT: Questions

Generalizability of the SPRINT population

16.8 million (7.6%) of US adults

8.2 million (16.7%) of those with treated HTN

Cognitive outcomes

Long term renal effects

AKI events



Whelton PK, et al.

2017 High Blood Pressure Clinical Practice Guideline: Executive Summary

**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**

Executive Summary

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

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THE OHIO STATE UNIVERSITY

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AHA/ACC 2017 Hypertension Guidelines: New Elements

Classification of hypertension

Previous guidelines (32% of US adults have hypertension)

| | | | |
|-----------------|----------------|----|----------------|
| Normal | SBP<120 | or | DBP<80 |
| Prehypertension | SBP 120 to 139 | or | DBP 80-89 |
| Stage I | SBP 140 to 159 | or | DBP 90-99 |
| Stage II | SBP \geq 160 | or | DBP \geq 100 |

New guidelines (46% of US adults have hypertension)

| | | | |
|----------|----------------|----|-----------|
| Normal | BP<120/80 | | |
| Elevated | 120-129/<80 | | |
| Stage I | SBP 130-139 | or | DBP 80-89 |
| Stage II | SBP \geq 140 | or | DBP >90 |



AHA/ACC 2017 Hypertension Guidelines: New Elements

Two different thresholds for starting blood pressure medications

$\geq 130/80$

$\geq 140/90$

Utilization of cardiac risk factor scoring system (Pooling Cohort 10-year CVD Risk Estimator)

Same blood pressure goals for all individuals regardless of age or comorbidities

Specific instructions for monitoring blood pressure in the office and at home



Pooled Cohort Estimator of Cardiac Risk

Risk Factors for ASCVD

| | | | |
|-------------------|--|--|---|
| Gender | <input checked="" type="button" value="Male"/> <input type="button" value="Female"/> | Systolic BP | <input type="text"/> mmHg |
| Age | <input type="text"/> years | Receiving treatment for high blood pressure (if SBP > 120 mmHg) | <input checked="" type="button" value="No"/> <input type="button" value="Yes"/> |
| Race | <input type="text" value="White or other"/> | Diabetes | <input checked="" type="button" value="No"/> <input type="button" value="Yes"/> |
| Total Cholesterol | <input type="text"/> mg/dL | Smoker | <input checked="" type="button" value="No"/> <input type="button" value="Yes"/> |
| HDL Cholesterol | <input type="text"/> mg/dL | | |

Table 23. BP Thresholds for and Goals of Pharmacological Therapy in Patients With Hypertension According to Clinical Conditions

| Clinical Condition(s) | BP Threshold, mm Hg | BP Goal, mm Hg |
|--|---------------------|----------------|
| General | | |
| Clinical CVD or 10-year ASCVD risk $\geq 10\%$ | $\geq 130/80$ | $< 130/80$ |
| No clinical CVD and 10-year ASCVD risk $< 10\%$ | $\geq 140/90$ | $< 130/80$ |
| Older persons (≥ 65 years of age; noninstitutionalized, ambulatory, community-living adults) | ≥ 130 (SBP) | < 130 (SBP) |
| Specific comorbidities | | |
| Diabetes mellitus | $\geq 130/80$ | $< 130/80$ |
| Chronic kidney disease | $\geq 130/80$ | $< 130/80$ |
| Chronic kidney disease after renal transplantation | $\geq 130/80$ | $< 130/80$ |
| Heart failure | $\geq 130/80$ | $< 130/80$ |
| Stable ischemic heart disease | $\geq 130/80$ | $< 130/80$ |
| Secondary stroke prevention | $\geq 140/90$ | $< 130/80$ |
| Secondary stroke prevention (lacunar) | $\geq 130/80$ | $< 130/80$ |
| Peripheral arterial disease | $\geq 130/80$ | $< 130/80$ |

ASCVD indicates atherosclerotic cardiovascular disease; BP, blood pressure; CVD, cardiovascular disease; and SBP, systolic blood pressure.



Figure 4. Blood Pressure (BP) Thresholds and Recommendations for Treatment and Follow-Up

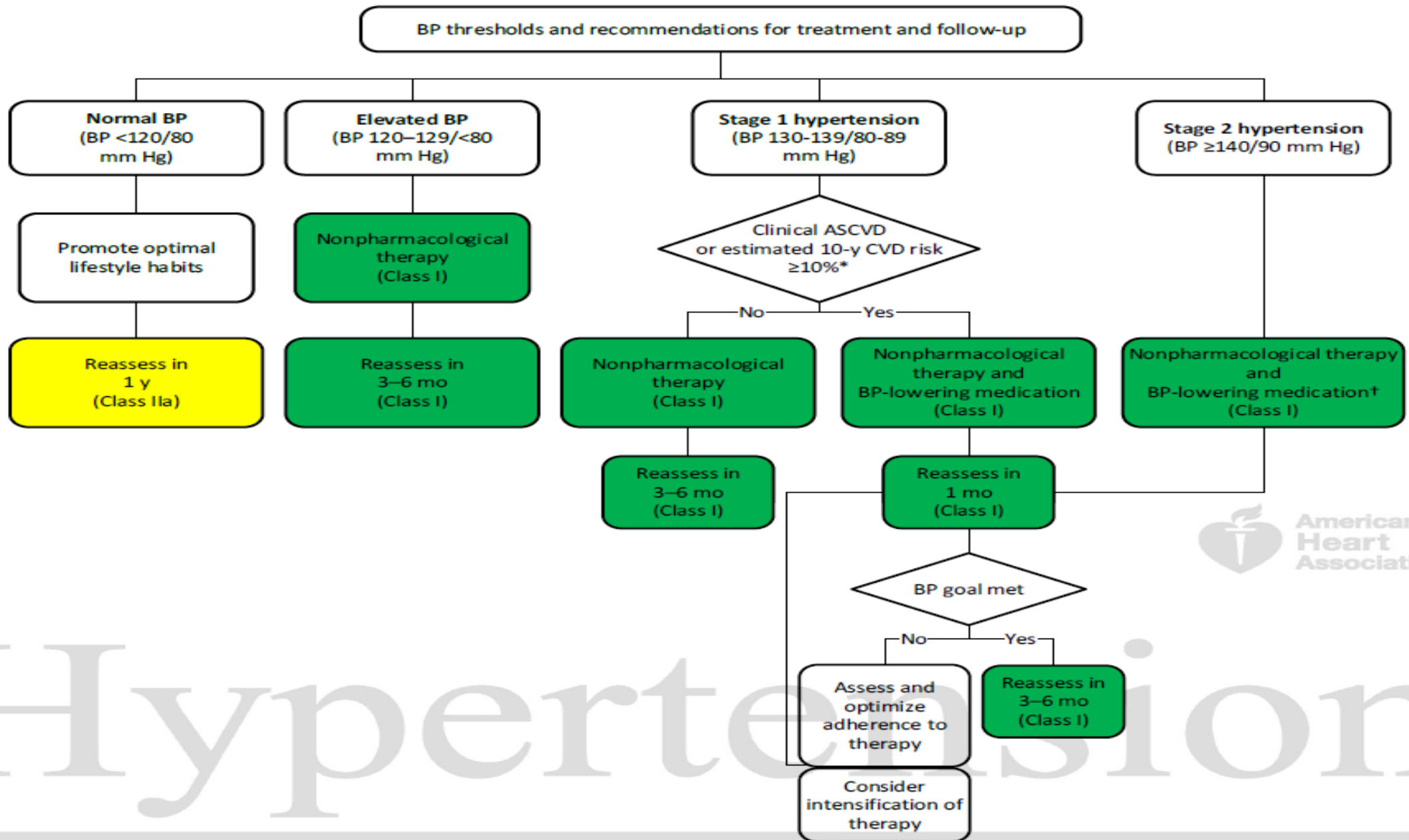


Table 15. Best Proven Nonpharmacological Interventions for Prevention and Treatment of Hypertension

| | Nonpharmacological Intervention | Dose | Approximate Impact on SBP | | |
|----------------------------------|---------------------------------|--|---------------------------|--------------|-----------|
| | | | Hypertension | Normotension | Reference |
| Weight loss | Weight/body fat | Best goal is ideal body weight, but aim for at least a 1-kg reduction in body weight for most adults who are overweight. Expect about 1 mm Hg for every 1-kg reduction in body weight. | -5 mm Hg | -2/3 mm Hg | (1) |
| Healthy diet | DASH dietary pattern | Consume a diet rich in fruits, vegetables, whole grains, and low-fat dairy products, with reduced content of saturated and total fat. | -11 mm Hg | -3 mm Hg | (6, 7) |
| Reduced intake of dietary sodium | Dietary sodium | Optimal goal is <1500 mg/d, but aim for at least a 1000-mg/d reduction in most adults. | -5/6 mm Hg | -2/3 mm Hg | (9, 10) |

| | | | | | |
|--------------------------------------|----------------------|--|------------|------------|----------|
| Enhanced intake of dietary potassium | Dietary potassium | Aim for 3500–5000 mg/d, preferably by consumption of a diet rich in potassium. | -4/5 mm Hg | -2 mm Hg | (13) |
| Physical activity | Aerobic | <ul style="list-style-type: none"> ● 90–150 min/wk ● 65%–75% heart rate reserve | -5/8 mm Hg | -2/4 mm Hg | (18, 22) |
| | Dynamic resistance | <ul style="list-style-type: none"> ● 90–150 min/wk ● 50%–80% 1 rep maximum ● 6 exercises, 3 sets/exercise, 10 repetitions/set | -4 mm Hg | -2 mm Hg | (18) |
| | Isometric resistance | <ul style="list-style-type: none"> ● 4 × 2 min (hand grip), 1 min rest between exercises, 30%–40% maximum voluntary contraction, 3 sessions/wk ● 8–10 wk | -5 mm Hg | -4 mm Hg | (19, 30) |
| Moderation in alcohol intake | Alcohol consumption | <p>In individuals who drink alcohol, reduce alcohol[†] to:</p> <ul style="list-style-type: none"> ● Men: ≤2 drinks daily | -4 mm Hg | -3 mm Hg | (22-24) |

Points of controversy

Reliance on meta-analysis vs randomized controlled trials (RCTs)

Focus on systolic blood pressure

Use of Pooling Cohort for risk assessment

 SPIRNT utilized Framingham Risk Score > 15%

 New guideline suggest > 10%

Unknown long-term effects on kidney disease and cognitive function

 SPRINT ASK is currently gathering data

American College of Physicians and American Academy of Family Physicians have not endorsed the new guidelines

Universal acceptance of the new guidelines is still being considered





Summary of ACP and AAFP guidelines

Recommendation 1:

ACP and AAFP recommend that clinicians **initiate treatment in adults aged 60 years or older with systolic blood pressure persistently at or above 150 mm Hg to achieve a target systolic blood pressure of less than 150 mm Hg** to reduce the risk for stroke, cardiac events, and possibly mortality. (Grade: strong recommendation, high-quality evidence).

ACP and AAFP recommend that clinicians select the treatment goals for adults aged 60 years or older based on a periodic discussion of the benefits and harms of specific blood pressure targets with the patient.





Summary of ACP and AAFP guidelines

Recommendation 2:

ACP and AAFP recommend that clinicians **consider initiating or intensifying pharmacologic treatment in adults aged 60 years or older with a history of stroke or transient ischemic attack to achieve a target systolic blood pressure of less than 140 mm Hg to reduce the risk for recurrent stroke.** (Grade: weak recommendation, moderate-quality evidence).

ACP and AAFP recommend that clinicians select the treatment goals for adults aged 60 years or older based on a periodic discussion of the benefits and harms of specific blood pressure targets with the patient.



Summary of ACP and AAFP guidelines

Recommendation 3:

ACP and AAFP recommend that clinicians **consider initiating or intensifying pharmacologic treatment in some adults aged 60 years or older at high cardiovascular risk, based on individualized assessment, to achieve a target systolic blood pressure of less than 140 mm Hg.** (Grade: weak recommendation, low-quality evidence).

ACP and AAFP recommend that clinicians select the treatment goals for adults aged 60 years or older based on a periodic discussion of the benefits and harms of specific blood pressure targets with the patient.





Summary

The new blood pressure guidelines have many benefits

- Increased recognition of the problem of hypertension

- Emphasis on individual risk assessment

- Specific non-pharmacologic recommendations for BP reduction

- Encouraging home BP evaluation

Long term cognitive and renal outcomes associated with more intensive blood pressure targets are still to be defined

Goal of blood pressure therapies should be customized after discussions with the patient

