

Management of hypertension in CKD

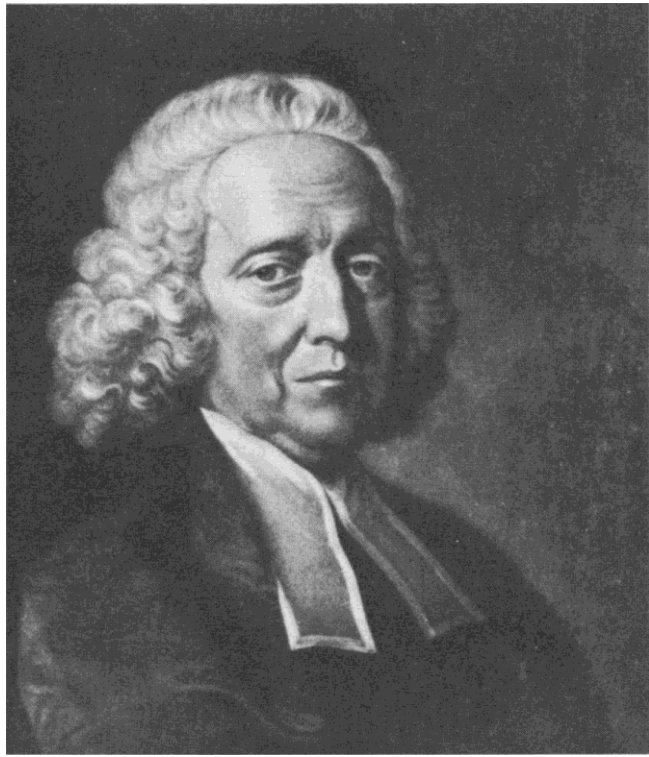
Csaba P Kovesdy, MD

University of Tennessee Health Science Center, Memphis TN

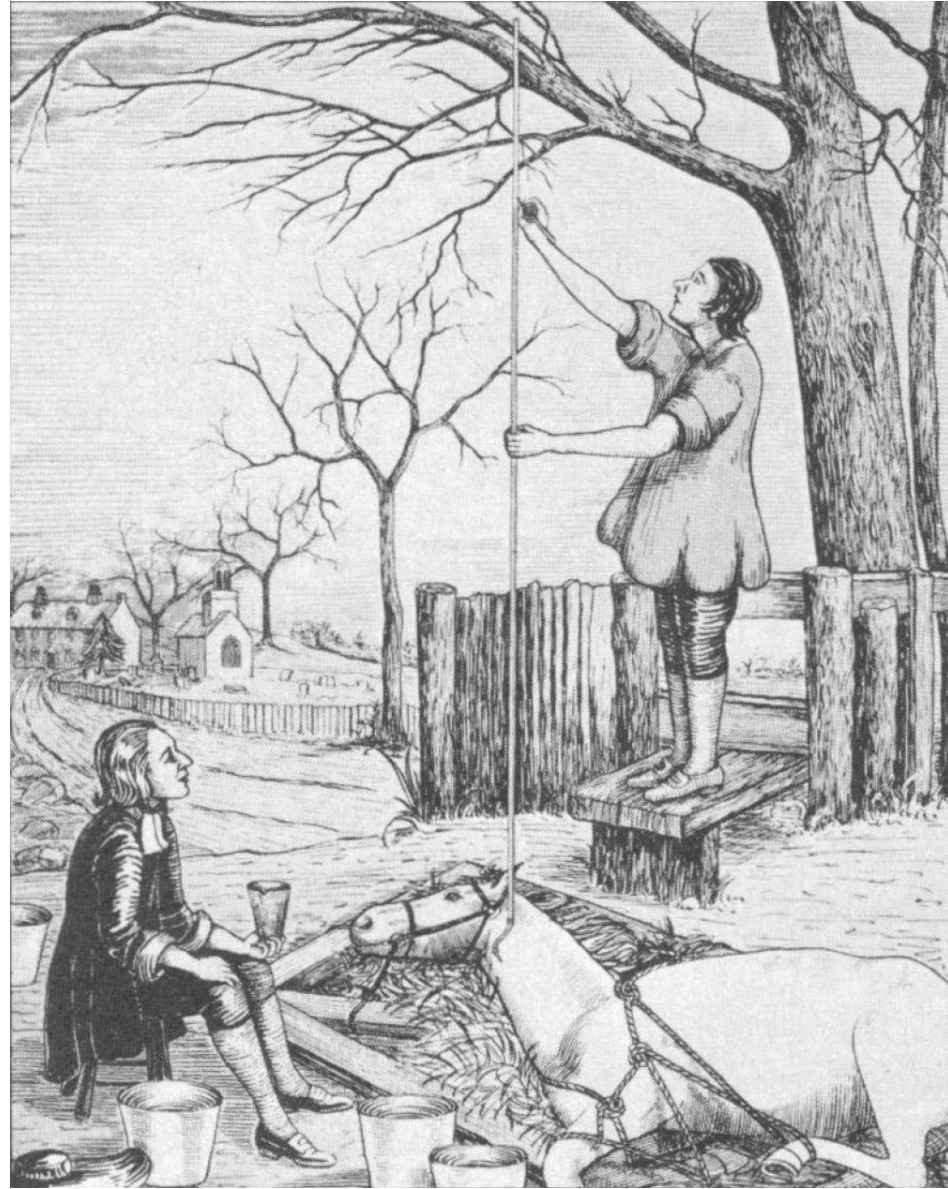
Memphis VA Medical Center, Memphis TN

Objectives

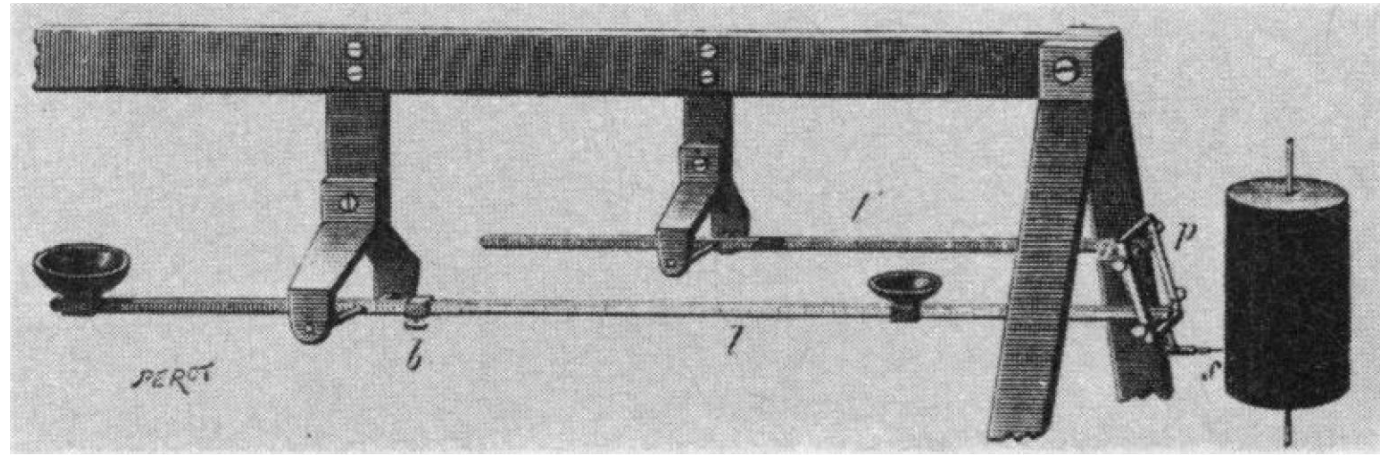
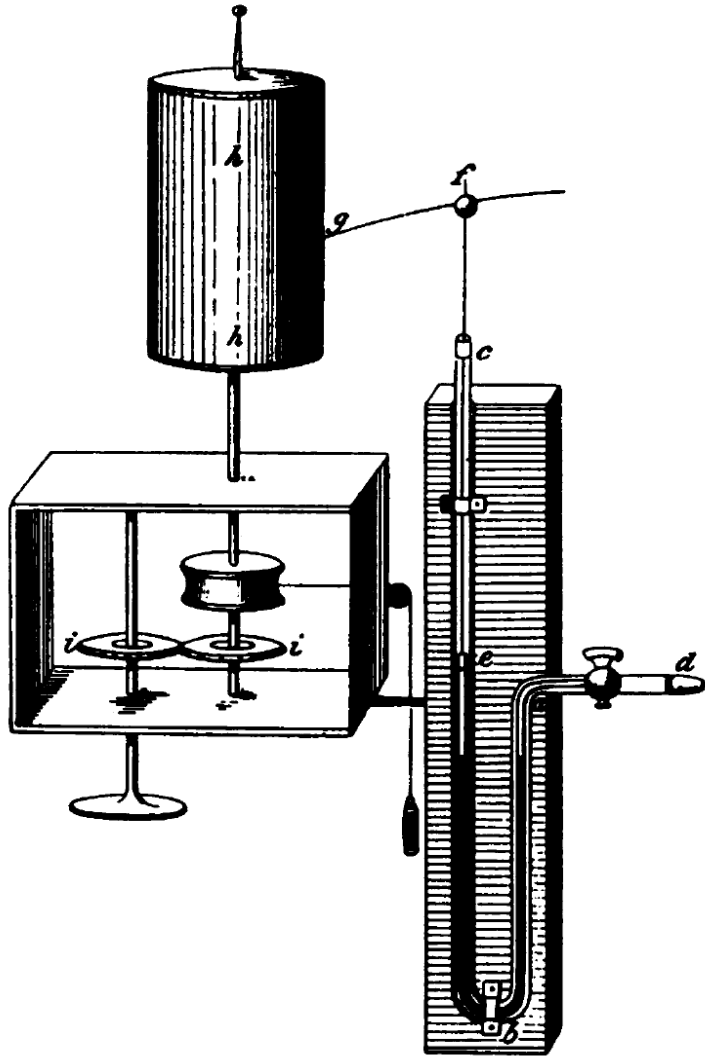
- Describe the prevalence and consequences of hypertension in the general population and in patients with chronic kidney disease
- Review the physiology and pathophysiology underlying the differences in hypertension characteristics in patients with and without chronic kidney disease
- Examine practical questions regarding hypertension treatment targets in patients with chronic kidney disease



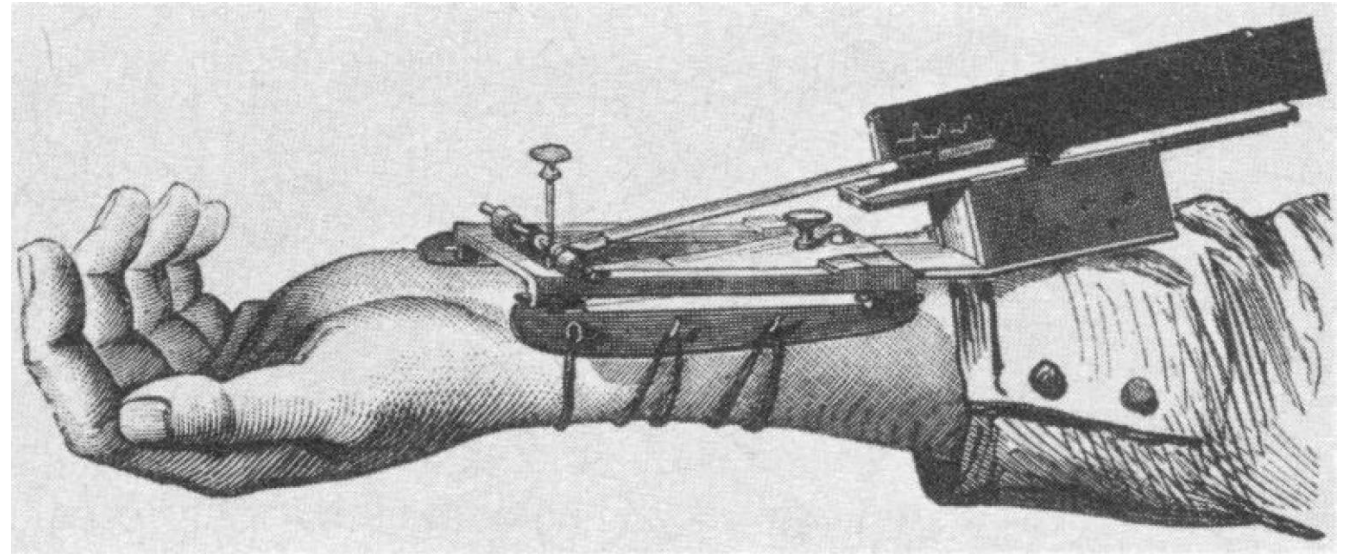
Stephen Hales (1677-1761)



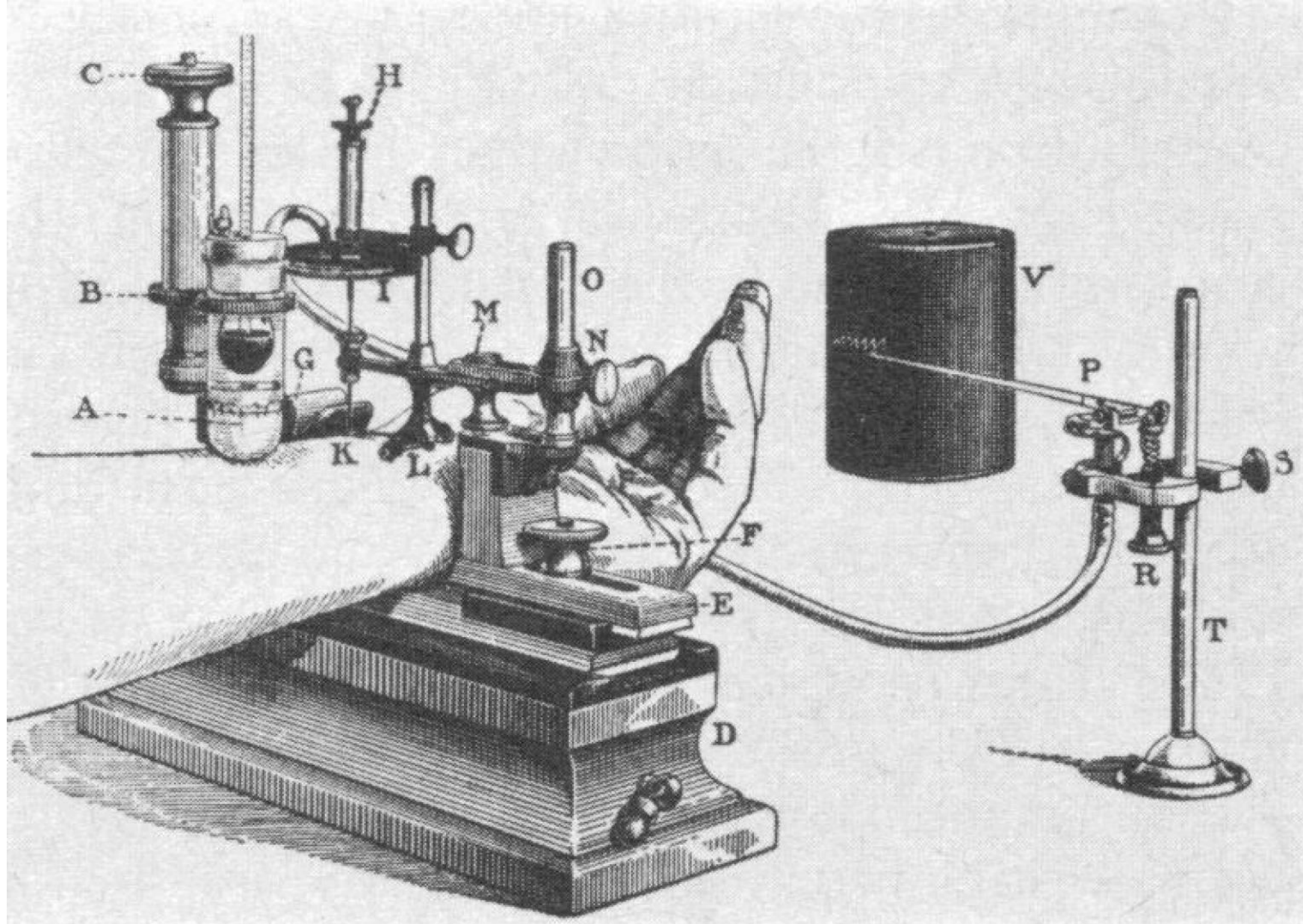
Ludwig's kymograph (1847)



Vierordt's sphygmograph (1852)



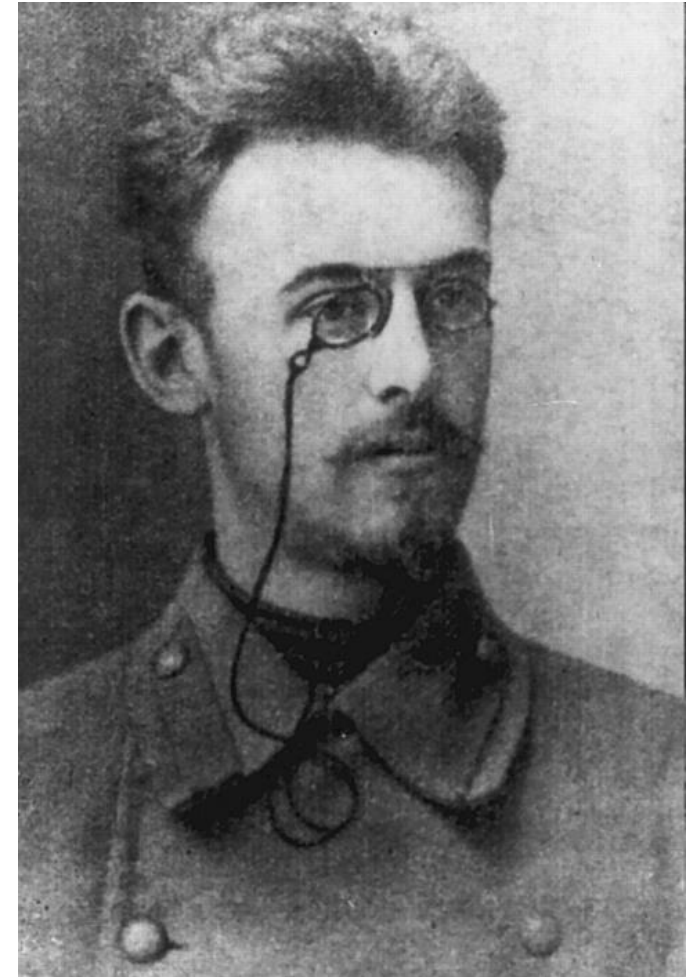
Direct sphygmograph (Marey, cca 1881)



Von Basch's sphygmomanometer and stand (cca 1881)

History of BP measurement

- Scipione Riva-Rocci: Introduction of modern circumferential BP cuff (1896)
- Friedrich von Recklinghausen: Started using larger size cuff (1901)
- Nikolai Korotkov: Introduction of the auscultatory technique (1905)



Nikolai Sergeevich Korotkov
(1874-1920)

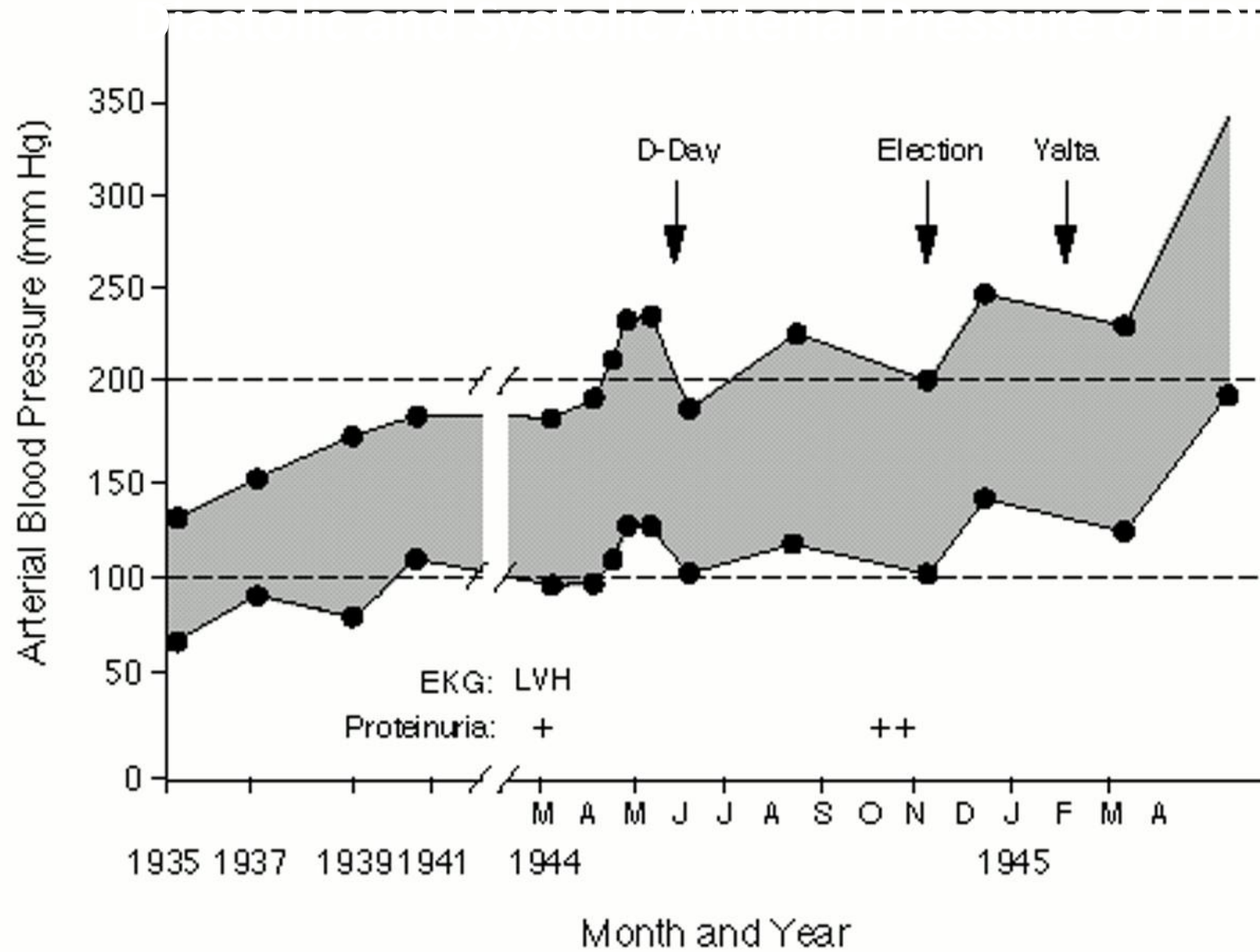
'CAME OUT OF CLEAR SKY,' SAYS PRESIDENT'S PHYSICIAN

Adm. Ross T. McIntire
Asserts There Was No
Indication of Immi-
nent Danger.

By CHARLES G. ROSS

DEATH DUE TO CEREBRAL
HEMORRHAGE --- BLOOD
VESSEL IN BRAIN BROKE

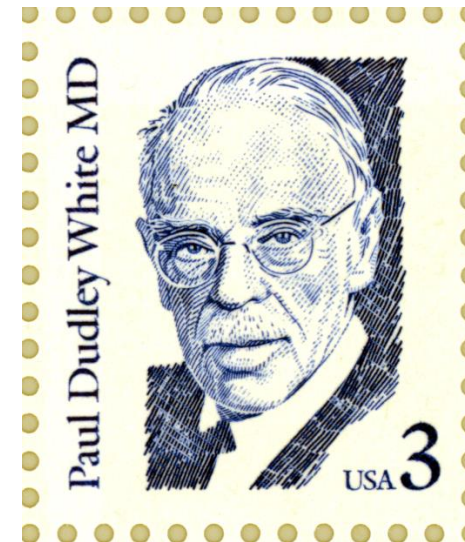
WASHINGTON, April 13 (AP).
PRESIDENT ROOSEVELT
died from what doctors call
a cerebral hemorrhage,
which means a sudden exten-



Hypertension may be an important compensatory mechanism which should not be tampered with, even were it certain that we could control it.

—*Paul Dudley White, 1937*

Paul Dudley White (June 6, 1886 – October 31, 1973), [American physician](#) and [cardiologist](#), was born in [Roxbury, Massachusetts](#), the son of Herbert Warren White and Elizabeth Abigail Dudley. He was one of the leading cardiologists of his day, and a prominent advocate of [preventive medicine](#).



Age	Blood pressure (mm Hg)	Reduction in life expectancy (years)	
		Man	Woman
35 years	130/90	4	-
	140/95	9	-
	150/100	16.5	-
45 years	130/90	3	1.5
	140/95	6	5
	150/100	11.5	8.5
55 years	130/90	1	0.5
	140/95	4	3
	150/100	6	4

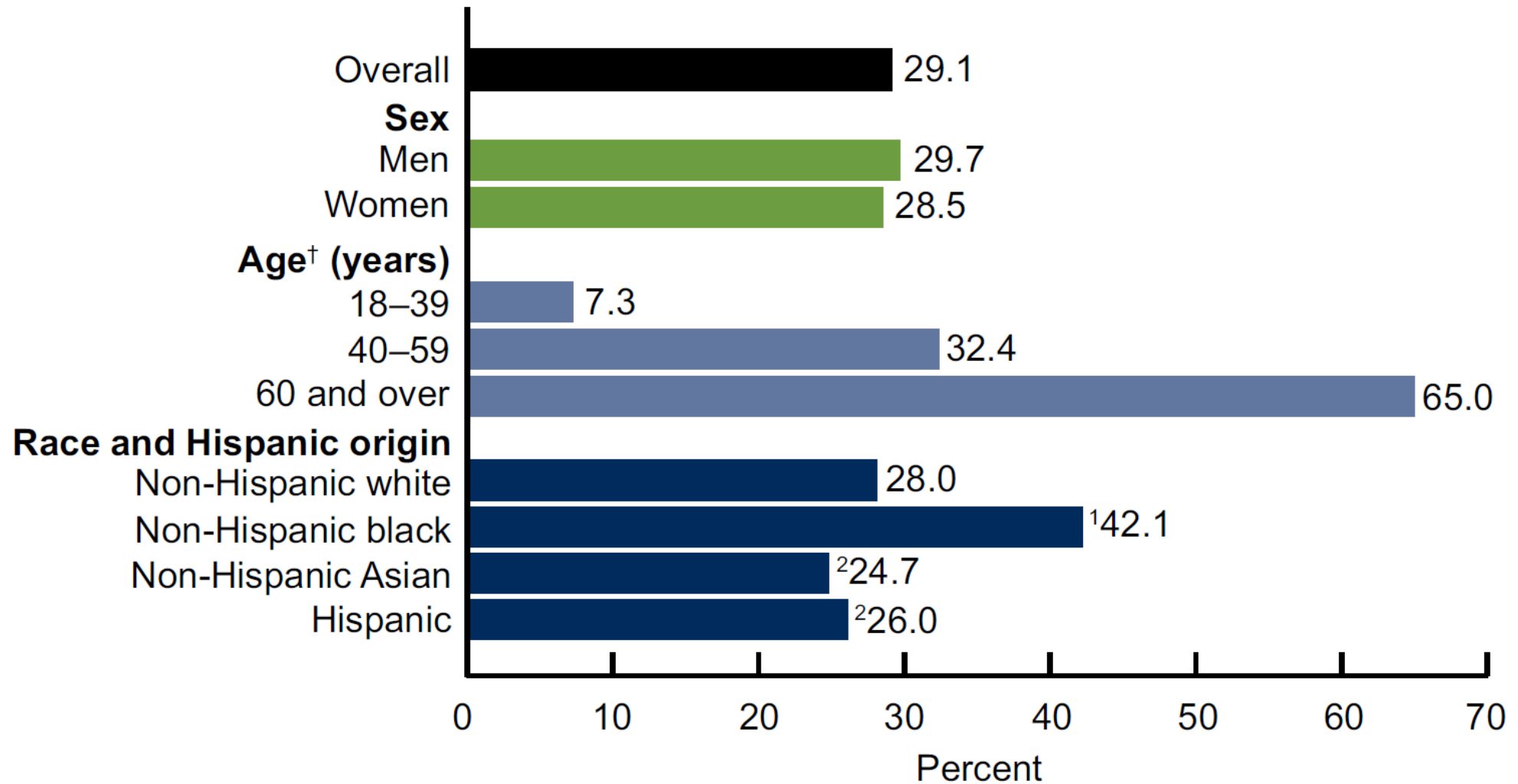
METROPOLITAN LIFE INSURANCE COMPANY. "Blood pressure: insurance experience and its implications". New York: Metropolitan Life Insurance Company; 1961.

Hypertension in Medicine: Key Moments

- Framingham Study: Kannel et al, *Ann Intern Med*. 1961 Jul;55:33-50
- First clinical trials of BP lowering
 - Hamilton M, Thompson EN: The role of blood pressure control in preventing complications of hypertension. *Lancet* 1964; 1:235-239
 - VA Cooperative Study Group: Effects of treatment on morbidity of hypertension. *JAMA* 1967; 202:1028-1033

- Relationship of BP with CV risk is “strong, continuous, independent, predictive and etiologically significant.”

Hypertension Among Adults in the United States: National Health and Nutrition Examination Survey, 2011–2012



- The 7th Report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure (JAMA, May 2003):

Blood Pressure Classification	SBP (mmHg)	DBP (mmHg)
Normal	<120	and <80
Prehypertension	120 ~ 139	or 80 ~ 89
Stage 1 Hyper	140 ~ 159	or 90 ~ 99
Stage 2 Hyper	>=160	or >=100

- K/DOQI CVD guideline: Predialysis and Postdialysis blood pressure goals should be <140/90 mmHg and <130/80 mm Hg, respectively. (AJKD, April 2005)

Target Blood Pressure

For Individuals With:	BP Goal:
Hypertension (no diabetes or renal disease)	<140/90 mmHg (JNC 7)
Diabetes Mellitus	<130/80 mmHg (ADA, JNC 7)
Renal Disease (CKD) with proteinuria >1 gram/24 hours or diabetic kidney disease	<135/85 mmHg <125/75 mmHg (NKF)

Chobanian AV et al. *JAMA*. 2003;289:2560–2571.

American Diabetes Association. *Diabetes Care*. 2002;25:134–147.

National Kidney Foundation. *Am J Kidn Dis*. 2002;39(suppl 1):S1–S266.

JNC 8, JAMA 2014, 311(5)

Special Communication

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)

Paul A. James, MD; Suzanne Oparil, MD; Barry L. Carter, PharmD; William C. Cushman, MD; Cheryl Dennison-Himmelfarb, RN, ANP, PhD; Joel Handler, MD; Daniel T. Lackland, DrPH; Michael L. LeFevre, MD, MSPH; Thomas D. MacKenzie, MD, MSPH; Olugbenga Ogedegbe, MD, MPH, MS; Sidney C. Smith Jr, MD; Laura P. Svetkey, MD, MHS; Sandra J. Taler, MD; Raymond R. Townsend, MD; Jackson T. Wright Jr, MD, PhD; Andrew S. Narva, MD; Eduardo Ortiz, MD, MPH

Hypertension is the most common condition seen in primary care and leads to **myocardial infarction, stroke, renal failure, and death** if not detected early and treated appropriately.

Patients want to be assured that BP treatment will reduce their disease burden, while clinicians want guidance on hypertension management using the best scientific evidence.

There is strong evidence to support treating HTN in persons **>60 years** to a BP goal **<150/90** mmHg and persons **30-59 years** to a diastolic goal **<90** mmHg

There is **insufficient** evidence in **<60** years for a systolic goal, or in **<30** years for a diastolic goal, so the panel recommends BP **<140/90** based on **expert opinion**.

The same thresholds and goals are recommended for adults with **diabetes** or nondiabetic **CKD** as for the general hypertensive population **<60** years.

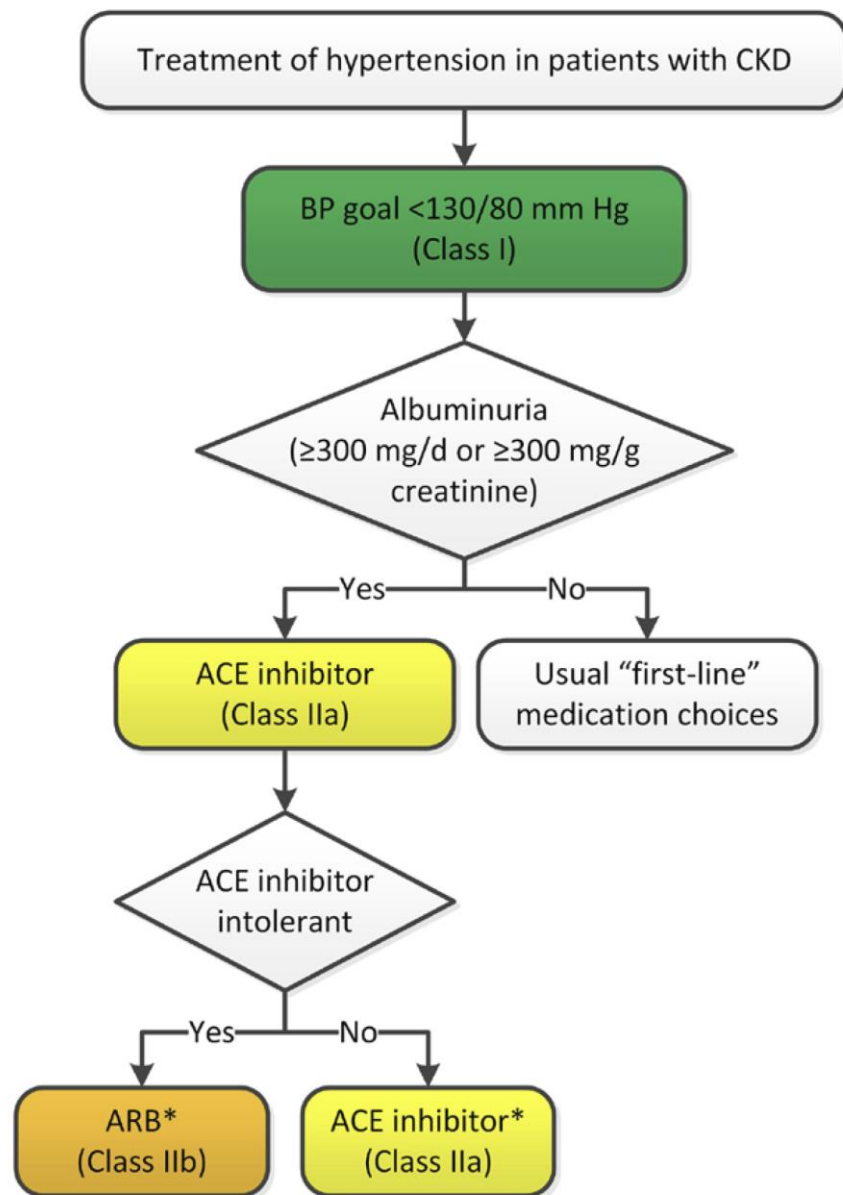
There is **moderate evidence** to support initiating drug treatment with an **ACE inhibitor, ARB, CCB, or thiazide-type diuretic** in the **non-black** hypertensive population, including those with **diabetes**.

In the **black** hypertensive population, including those with **diabetes**, a **CCB blocker or thiazide-type diuretic** is recommended as initial therapy.

There is moderate evidence to support initial or **add-on** antihypertensive therapy with an **ACE inhibitor** or **ARB** in persons with **CKD** to improve kidney outcomes.

New 2017 AHA/ACC Hypertension Guidelines

BP Category	Systolic BP		Diastolic BP	Treatment or Follow-up
Normal	<120 mm Hg	and	<80 mm Hg	Evaluate yearly; encourage healthy lifestyle changes to maintain normal BP
Elevated	120-129 mm Hg	and	<80 mm Hg	Recommend healthy lifestyle changes and reassess in 3-6 months
Hypertension: stage 1	130-139 mm Hg	or	80-89 mm Hg	<p>Assess the 10-year risk for heart disease and stroke using the atherosclerotic cardiovascular disease (ASCVD) risk calculator</p> <ul style="list-style-type: none"> • If risk is less than 10%, start with healthy lifestyle recommendations and reassess in 3-6 months • If risk is greater than 10% or the patient has known clinical cardiovascular disease (CVD), diabetes mellitus, or chronic kidney disease, recommend lifestyle changes and BP-lowering medication (1 medication); reassess in 1 month for effectiveness of medication therapy <ul style="list-style-type: none"> – If goal is met after 1 month, reassess in 3-6 months – If goal is not met after 1 month, consider different medication or titration – Continue monthly follow-up until control is achieved
Hypertension: stage 2	≥140 mm Hg	or	≥90 mm Hg	<p>Recommend healthy lifestyle changes and BP-lowering medication (2 medications of different classes); reassess in 1 month for effectiveness</p> <ul style="list-style-type: none"> • If goal is met after 1 month, reassess in 3-6 months • If goal is not met after 1 month, consider different medications or titration • Continue monthly follow-up until control is achieved



Recommendations for Treatment of Hypertension in Patients With CKD

References that support recommendations are summarized in **Online Data Supplements 37 and 38** and **Systematic Review Report**.

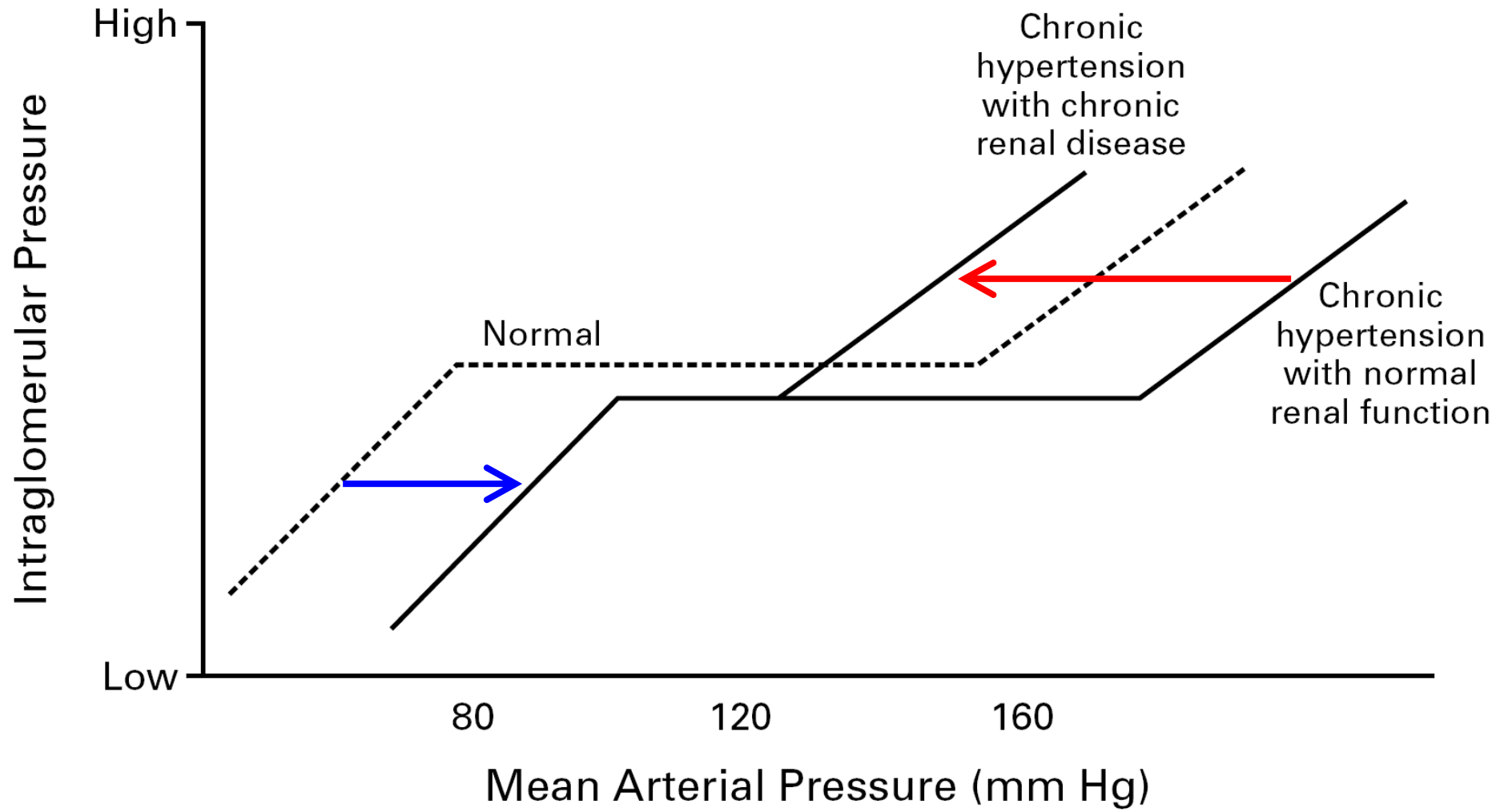
COR	LOE	Recommendations
I	SBP: B-R ^{SR}	1. Adults with hypertension and CKD should be treated to a BP goal of less than 130/80 mm Hg. ^{S9.3-1–S9.3-6}
	DBP: C-EO	
IIa	B-R	2. In adults with hypertension and CKD (stage 3 or higher or stage 1 or 2 with albuminuria [≥ 300 mg/d, or ≥ 300 mg/g albumin-to-creatinine ratio or the equivalent in the first morning void]), treatment with an ACE inhibitor is reasonable to slow kidney disease progression. ^{S9.3-3,S9.3-7–S9.3-12}
IIb	C-EO	3. In adults with hypertension and CKD (stage 3 or higher or stage 1 or 2 with albuminuria [≥ 300 mg/d, or ≥ 300 mg/g albumin-to-creatinine ratio in the first morning void]) (S9.3-7,S9.3-8), treatment with an ARB may be reasonable if an ACE inhibitor is not tolerated.

BP goal in CKD

- Recommendation 3.1.1: We suggest that adults with high BP and CKD be treated with a **target systolic blood pressure (SBP) of <120 mm Hg**, when tolerated, using standardized office BP measurement (**2B**).
 - Practice Point 3.1.1: It is potentially hazardous to apply the recommended SBP target of <120 mm Hg to BP measurements obtained in a non-standardized manner.
 - Practice Point 3.1.2: Clinicians can reasonably offer less intensive BP-lowering therapy in patients with very limited life expectancy or symptomatic postural hypotension.

BP treatment goals in CKD: What is the level of evidence?

- Paucity of conclusive clinical trials, trials with contradictory results
 - Extrapolation from other populations
 - Secondary analyses of trials examining different end points
 - Newer clinical trials that are relevant: SPRINT, ACCORD



Changes in renal vasculature

- Afferent arteriolar endothelial dysfunction leading to impaired vasodilatation
 - Hyaline arteriosclerosis
 - Myointimal hyperplasia
-
- Result is impaired autoregulatory capacity
 - Intraglomerular pressure begins to vary directly with changes in systemic pressure

Ditscherlein G, *Hypertension* 1985;7:II-29–II-32.

Bidani et al, *Hypertension* 1994;24:309-16.

Pelayo and Westcott, *J Clin Invest* 1991;88:101-5.

Hayashi et al, *J Hypertens* 1996;14:1387-401.

Palmer BF, *Am J Med Sci* 2001;321:388-400.

Table 1. Factors Increasing Susceptibility to Renal Hypoperfusion.

Failure to decrease arteriolar resistance

Structural changes in renal arterioles and small arteries

Old age

Atherosclerosis

Chronic hypertension

Chronic kidney disease

Malignant or accelerated hypertension

Reduction in vasodilatory prostaglandins

Nonsteroidal antiinflammatory drugs

Cyclooxygenase-2 inhibitors

Afferent glomerular arteriolar vasoconstriction

Sepsis

Hypercalcemia

Hepatorenal syndrome

Cyclosporine or tacrolimus

Radiocontrast agents

Failure to increase efferent arteriolar resistance

Angiotensin-converting–enzyme inhibitors

Angiotensin-receptor blockers

Renal-artery stenosis

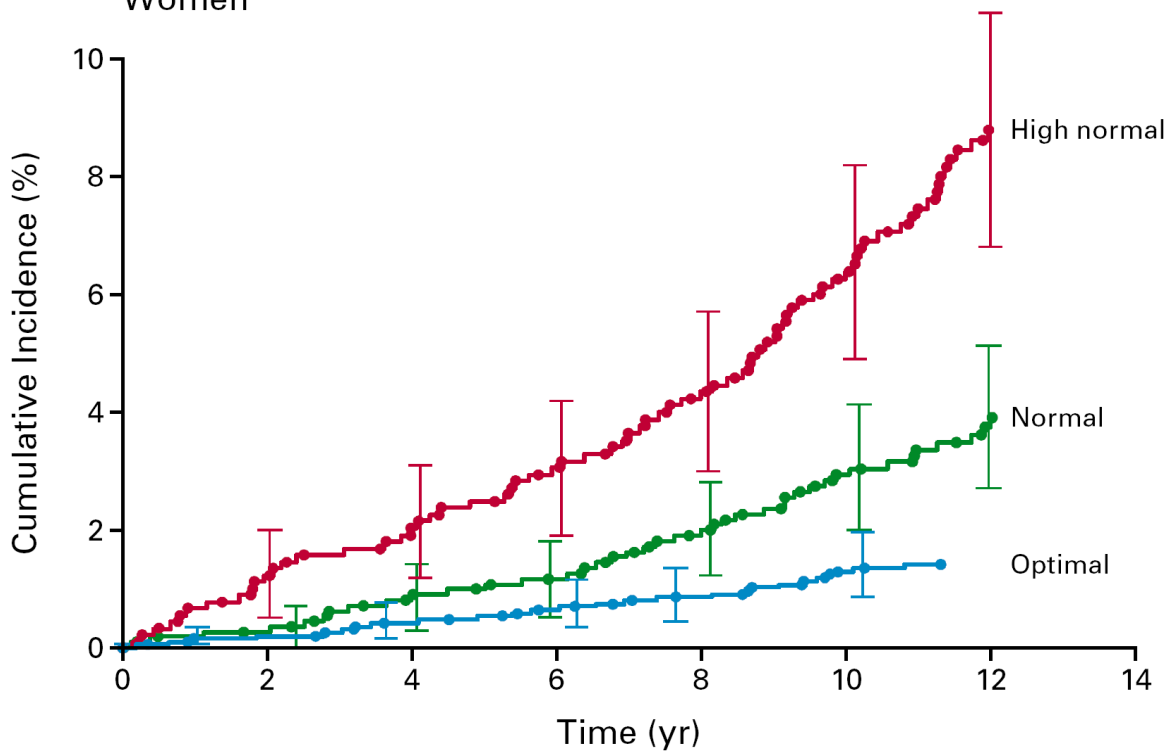
Abuelo JG, N

Engl J Med 2007;357:797-805.

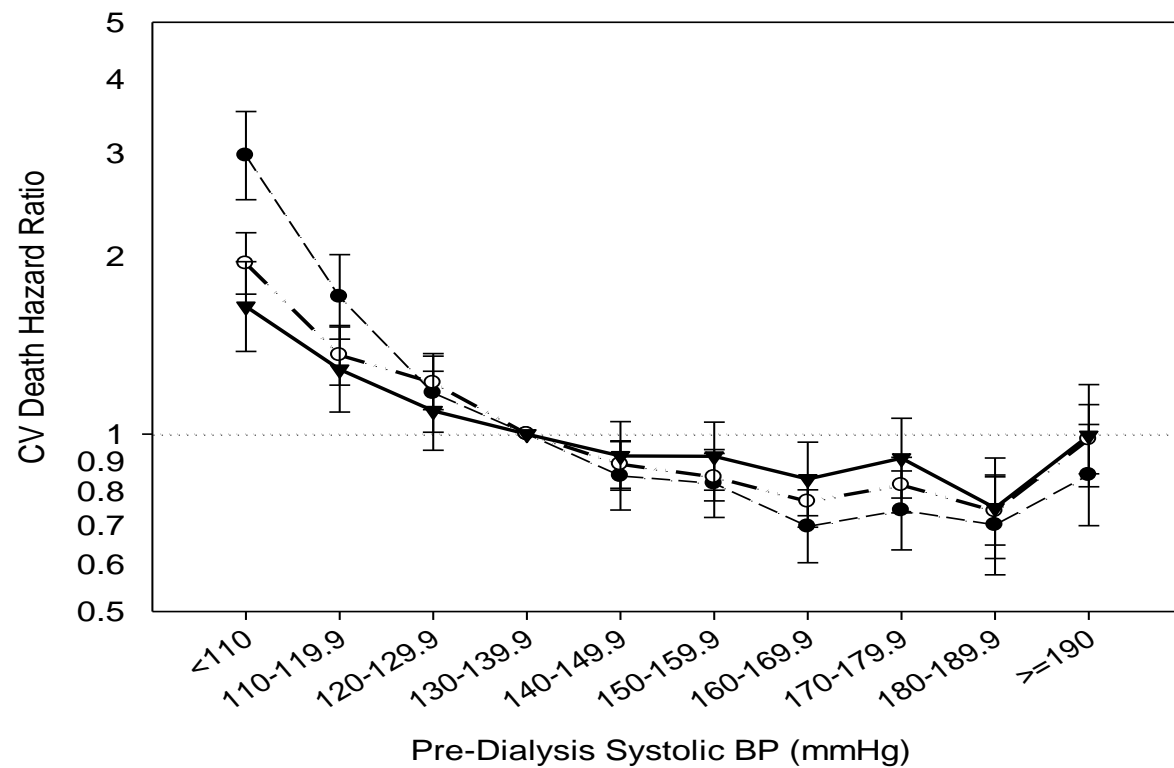


APPLES VS ORANGES

Women



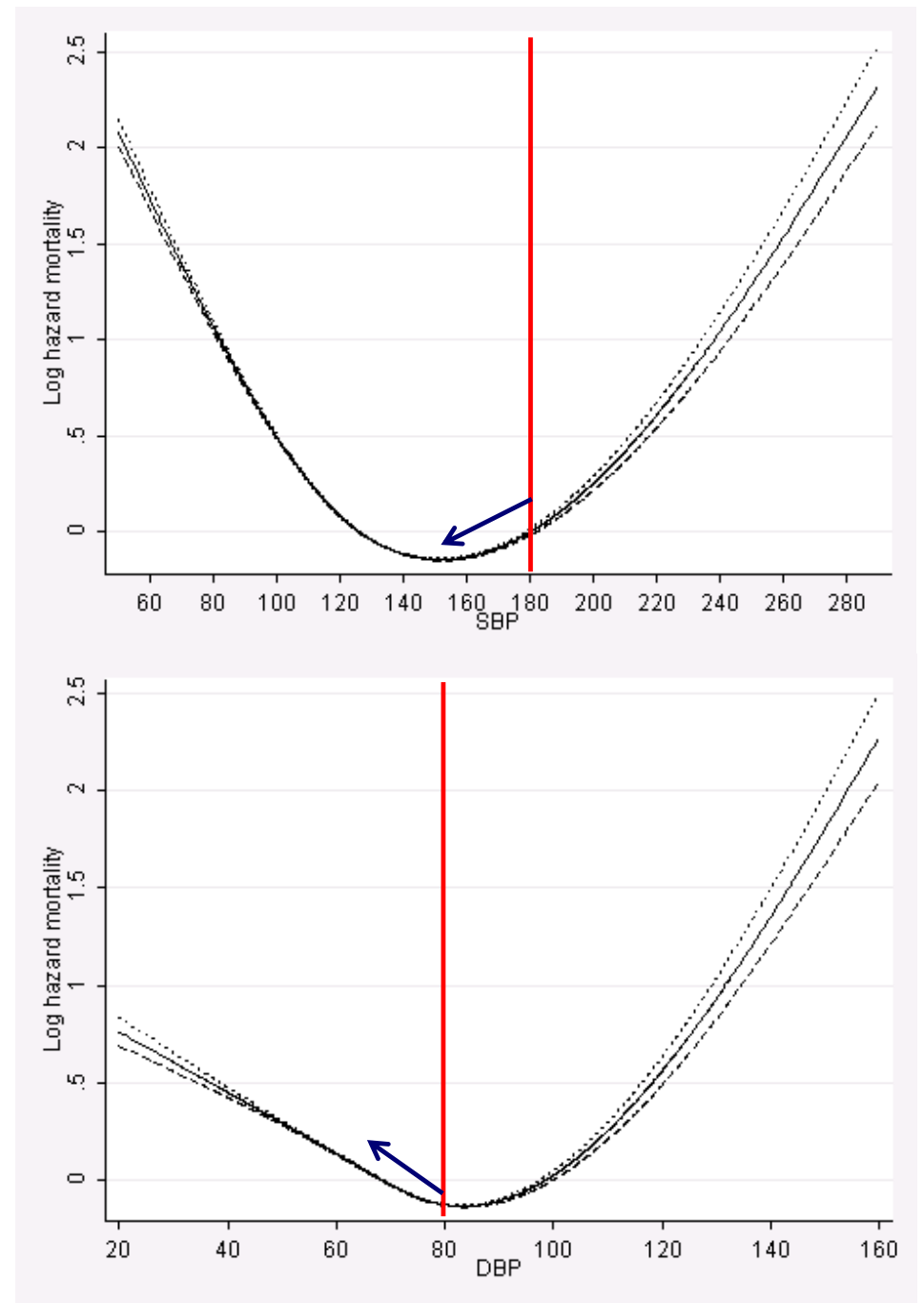
Vasan et al, *N Engl J Med* 2001;345:1291-7



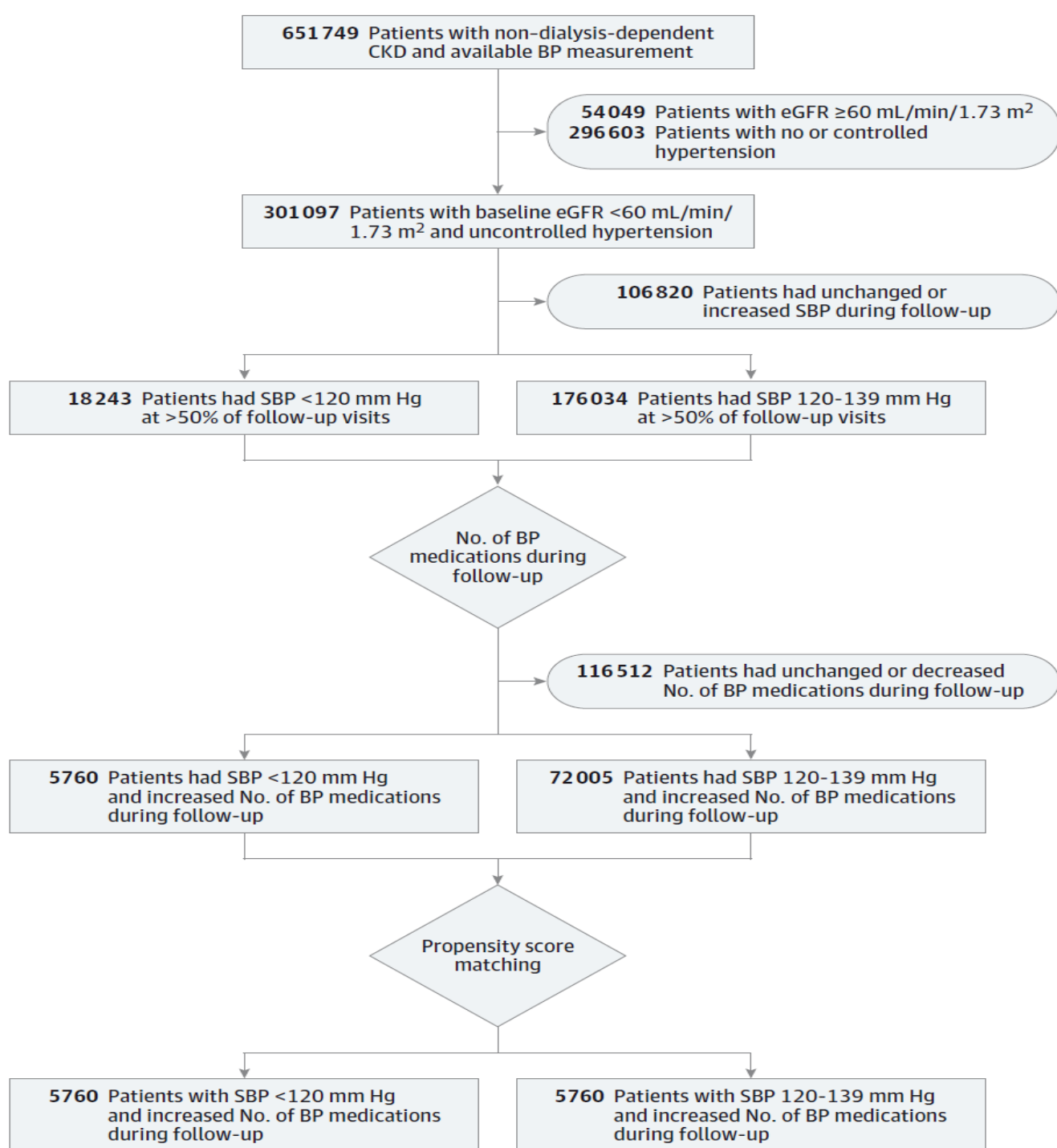
Kalantar-Zadeh et al, *Hypertension* 2005;45:811-7

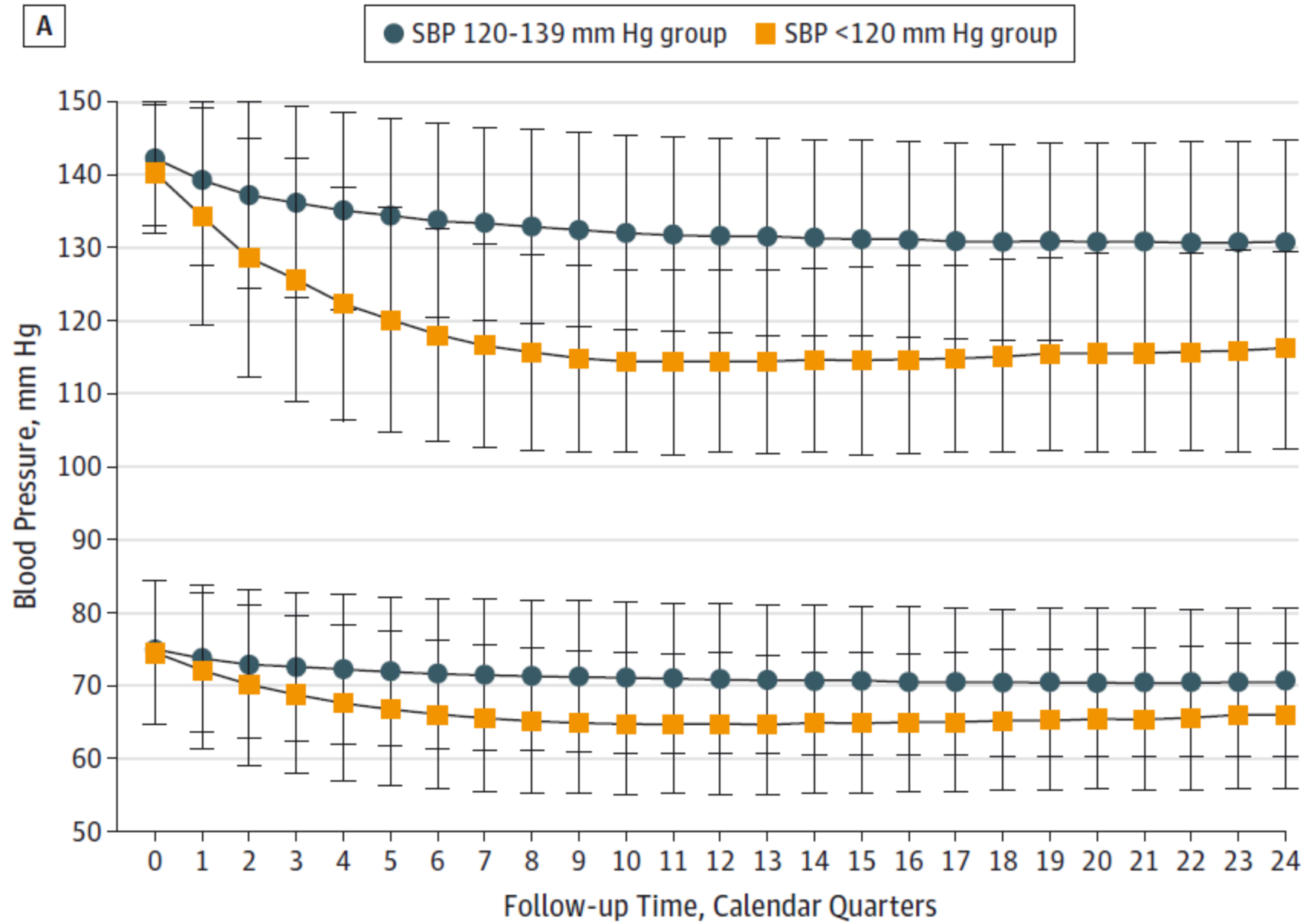
- The dilemma of BP lowering on the J-curve:
 - Lowering SBP
 - Also lowers DBP

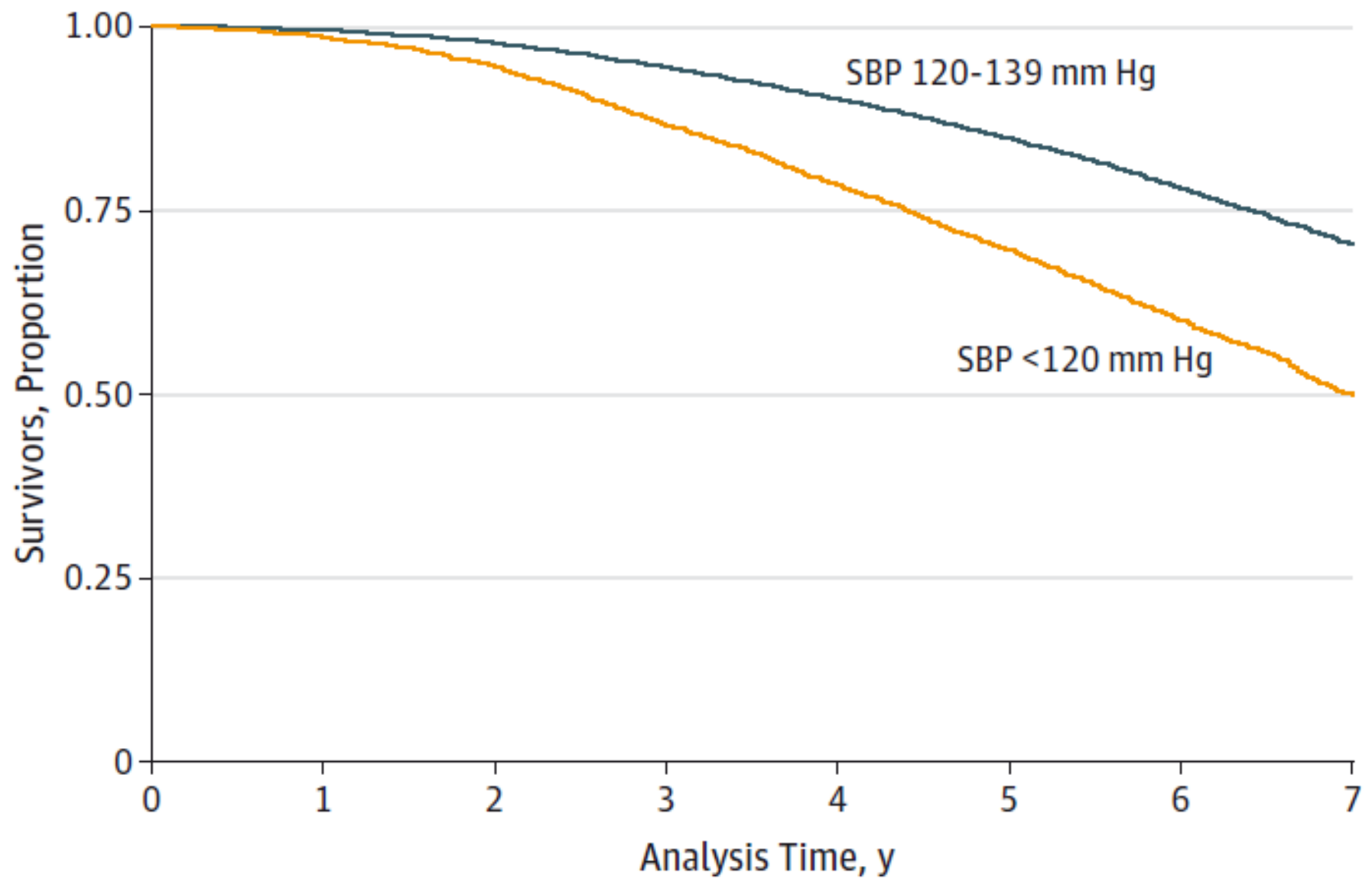
Kovesdy et al, *Ann Intern Med* 2013;159(4):233-42

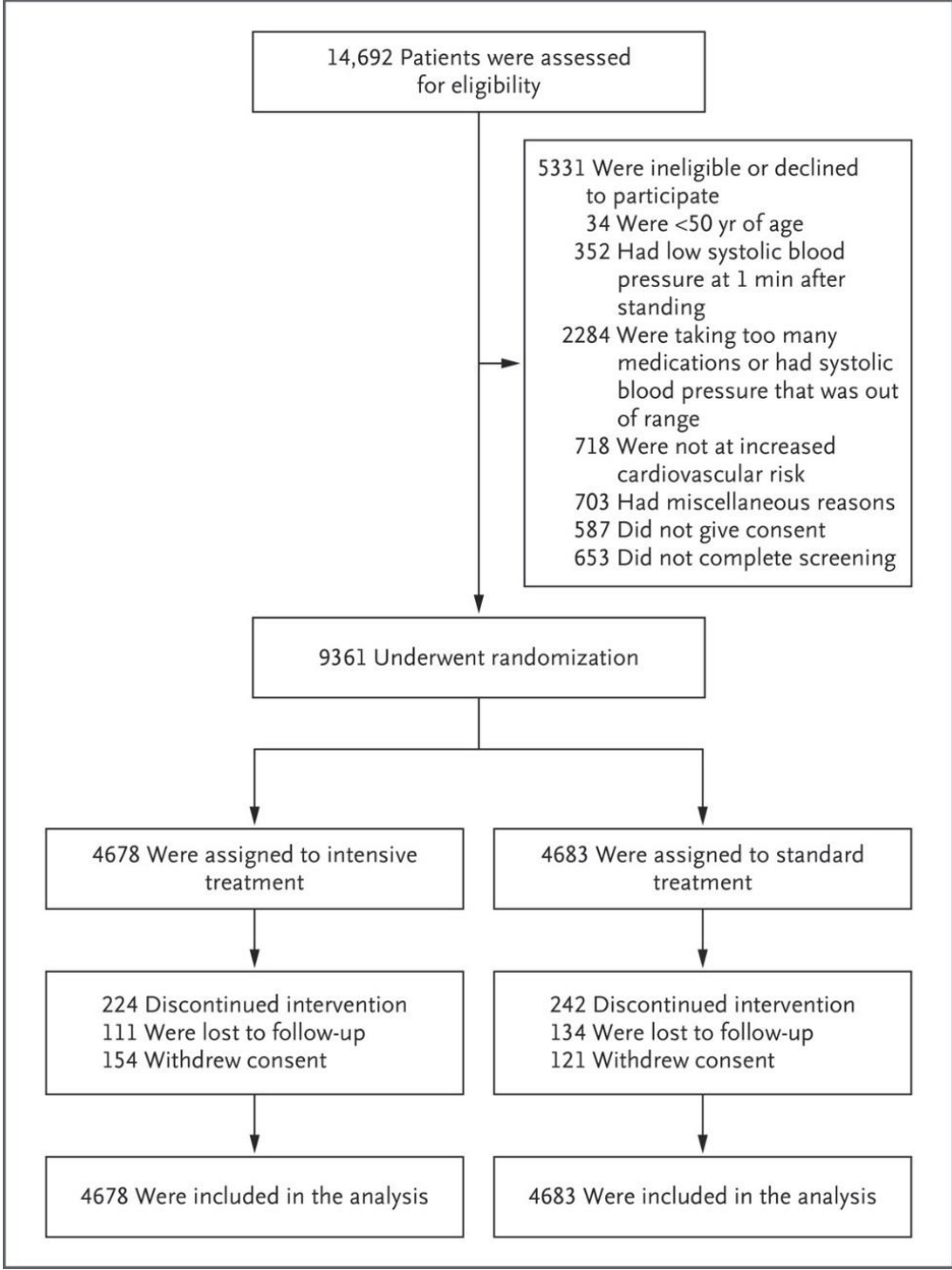


		SBP (mmHg)														
		<80	80-89	90-99	100-109	110-119	120-129	130-139	140-149	150-159	160-169	170-179	180-189	190-199	200-210	≥210
DBP (mmHg)	<40	2.56	2.42	2.55	2.15	1.73	1.69	1.91								
	40-49	2.99	2.69	2.31	1.77	1.58	1.39	1.37	1.30	1.50	1.83					
	50-59	3.25	2.88	2.24	1.77	1.51	1.27	1.14	1.17	1.27	1.32	1.63	1.20			
	60-69		3.11	2.32	1.82	1.48	1.23	1.09	1.09	1.12	1.13	1.28	1.36	1.00		
	70-79			2.05	1.70	1.34	1.14	1.01	1.01	1.04	1.07	1.12	1.19	1.11	1.17	1.26
	80-89				1.82	1.27	1.08	0.98	Ref.	1.01	1.07	1.13	1.22	1.43	1.25	1.35
	90-99					1.57	1.26	1.08	1.10	1.15	1.18	1.25	1.23	1.16	1.38	1.04
	100-109							1.53	1.16	1.31	1.33	1.37	1.30	1.62	1.40	1.42
	110-119									1.11	1.28	1.81	1.35	1.89	1.85	1.71
	≥120											1.62			2.44	2.06



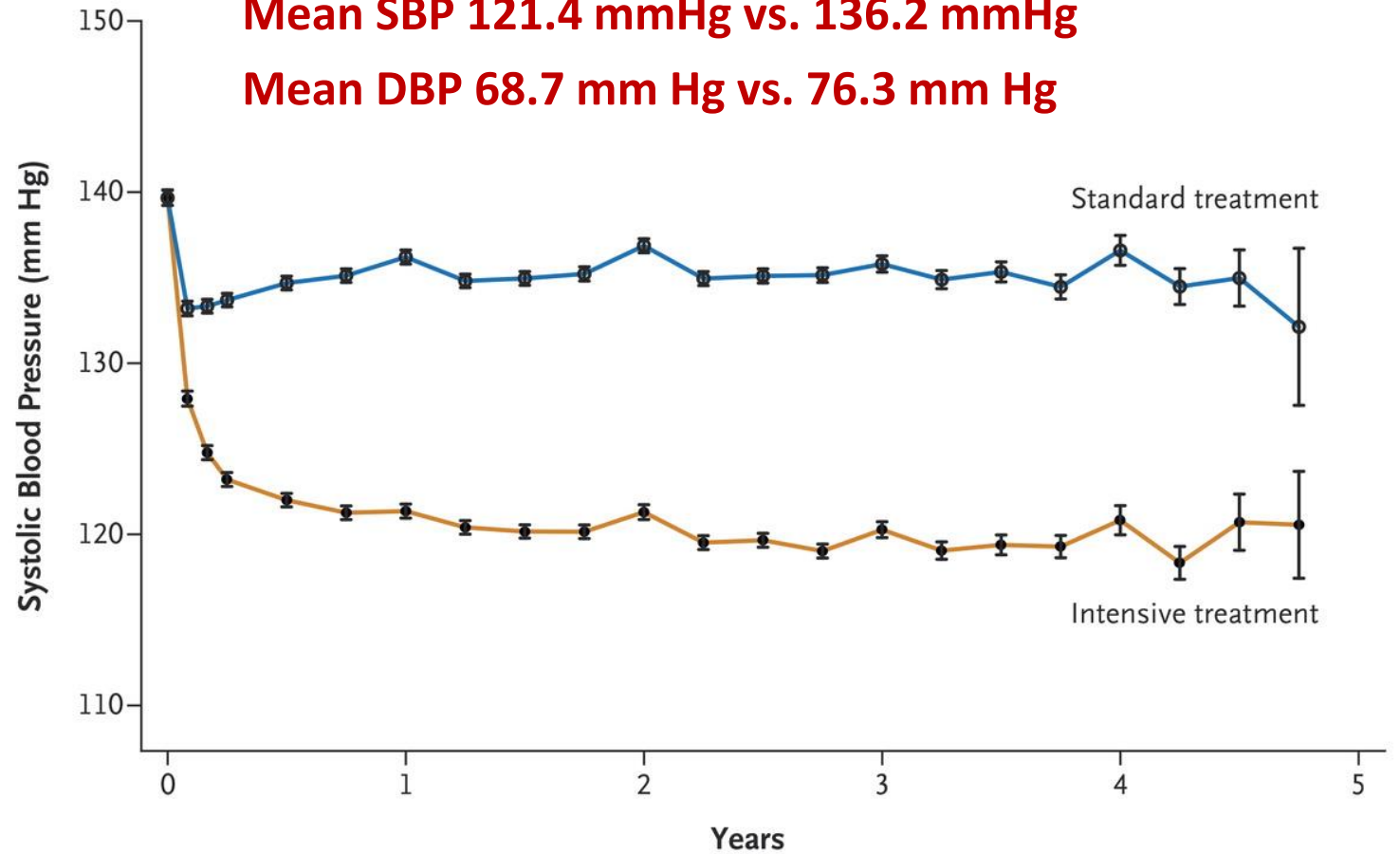








Mean SBP 121.4 mmHg vs. 136.2 mmHg
Mean DBP 68.7 mm Hg vs. 76.3 mm Hg



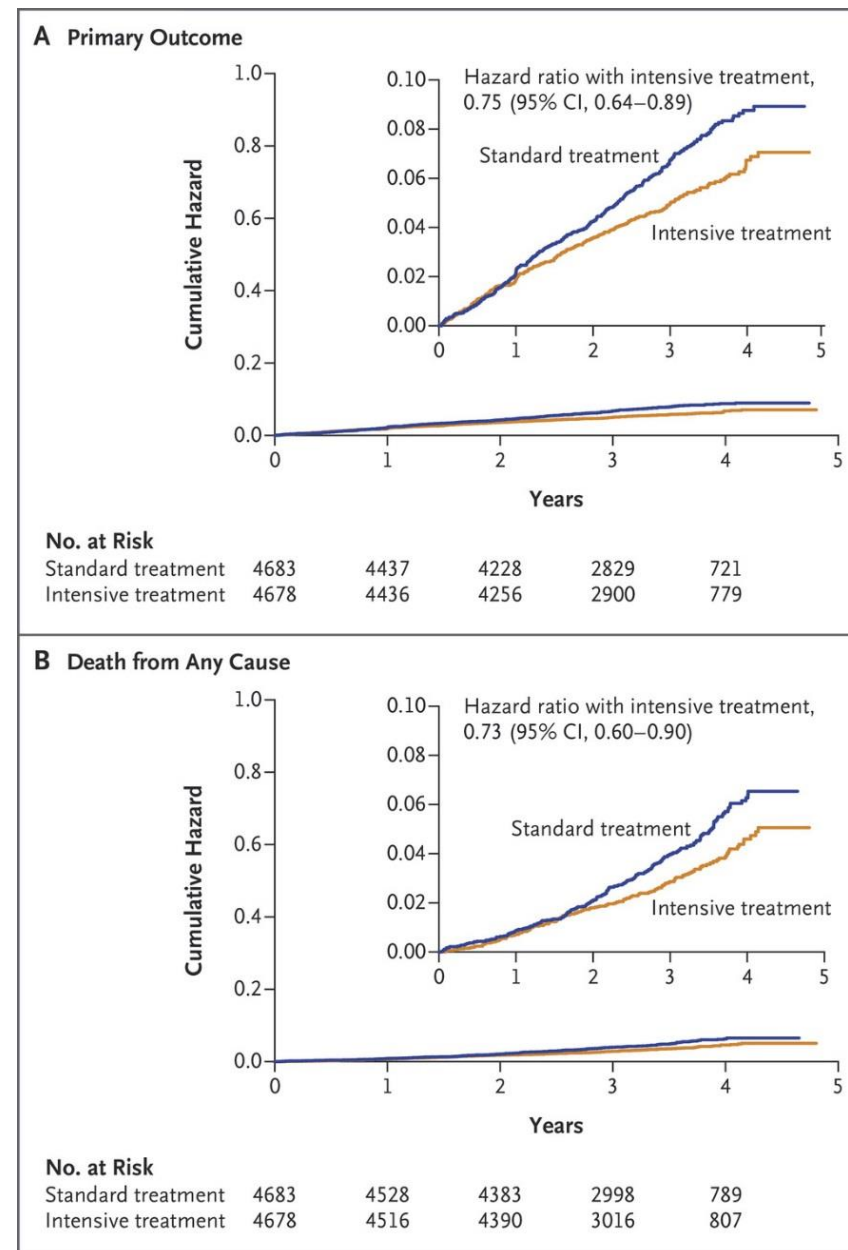
No. with Data

Standard treatment	4683	4345	4222	4092	3997	3904	3115	1974	1000	274
Intensive treatment	4678	4375	4231	4091	4029	3920	3204	2035	1048	286

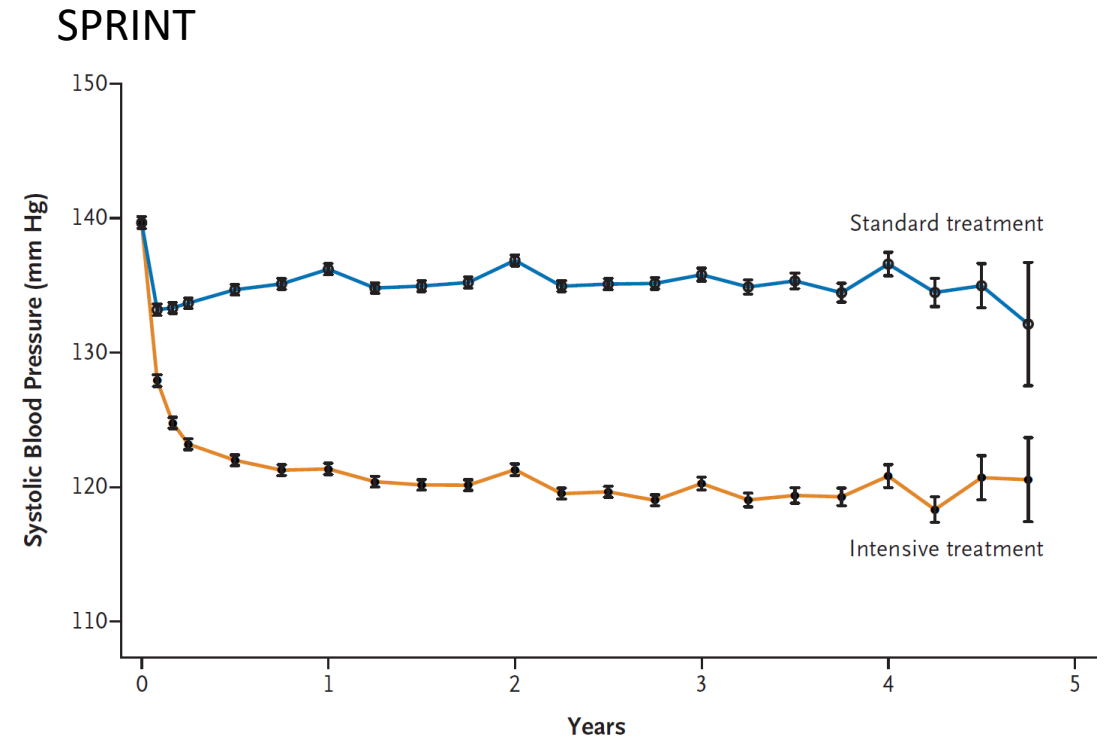
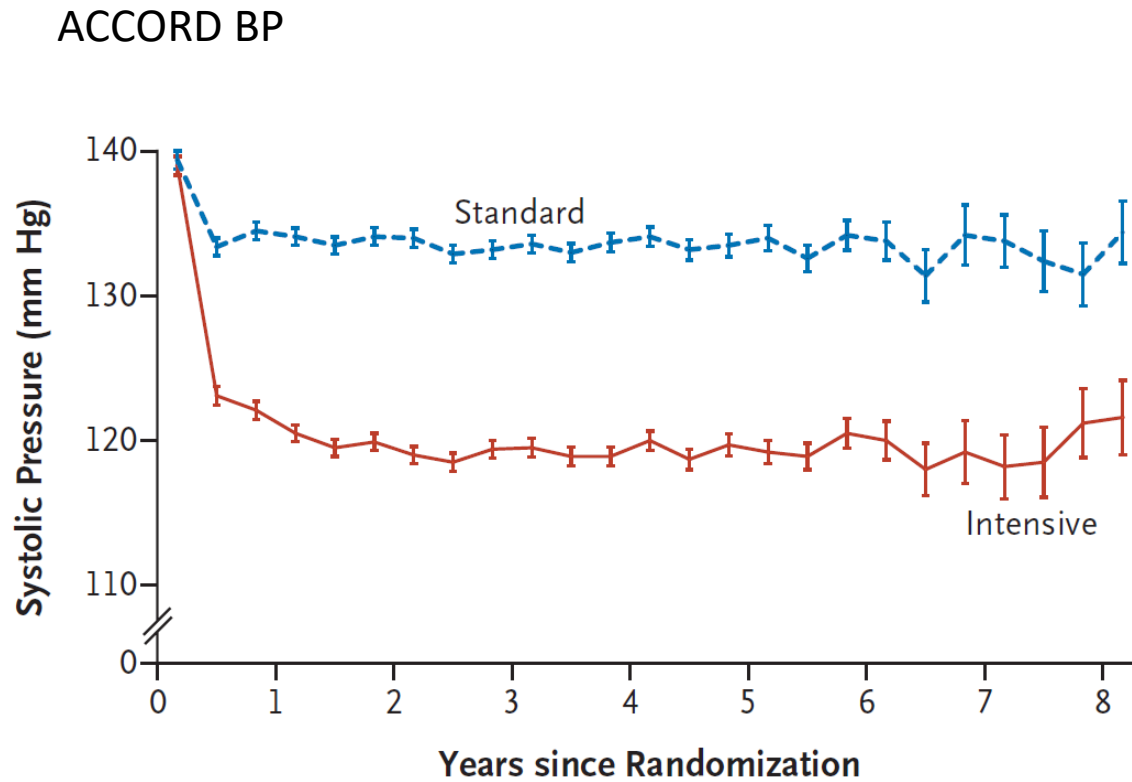
Mean No. of Medications

Standard treatment	1.9	1.8	1.8	1.8	1.8	1.8	1.8	1.8	1.8	1.9
Intensive treatment	2.3	2.7	2.8	2.8	2.8	2.8	2.8	2.8	2.8	3.0

- Median follow-up of 3.26 years
- Significantly lower rate of the primary composite outcome in the intensive-treatment group
 - 1.65% per year vs. 2.19% per year
 - Hazard ratio 0.75; 95% CI 0.64 to 0.89; $P < 0.001$
- All-cause mortality hazard ratio 0.73; 95% CI, 0.60 to 0.90; $P = 0.003$



Are differences in results a result of biases inherent of observational studies?



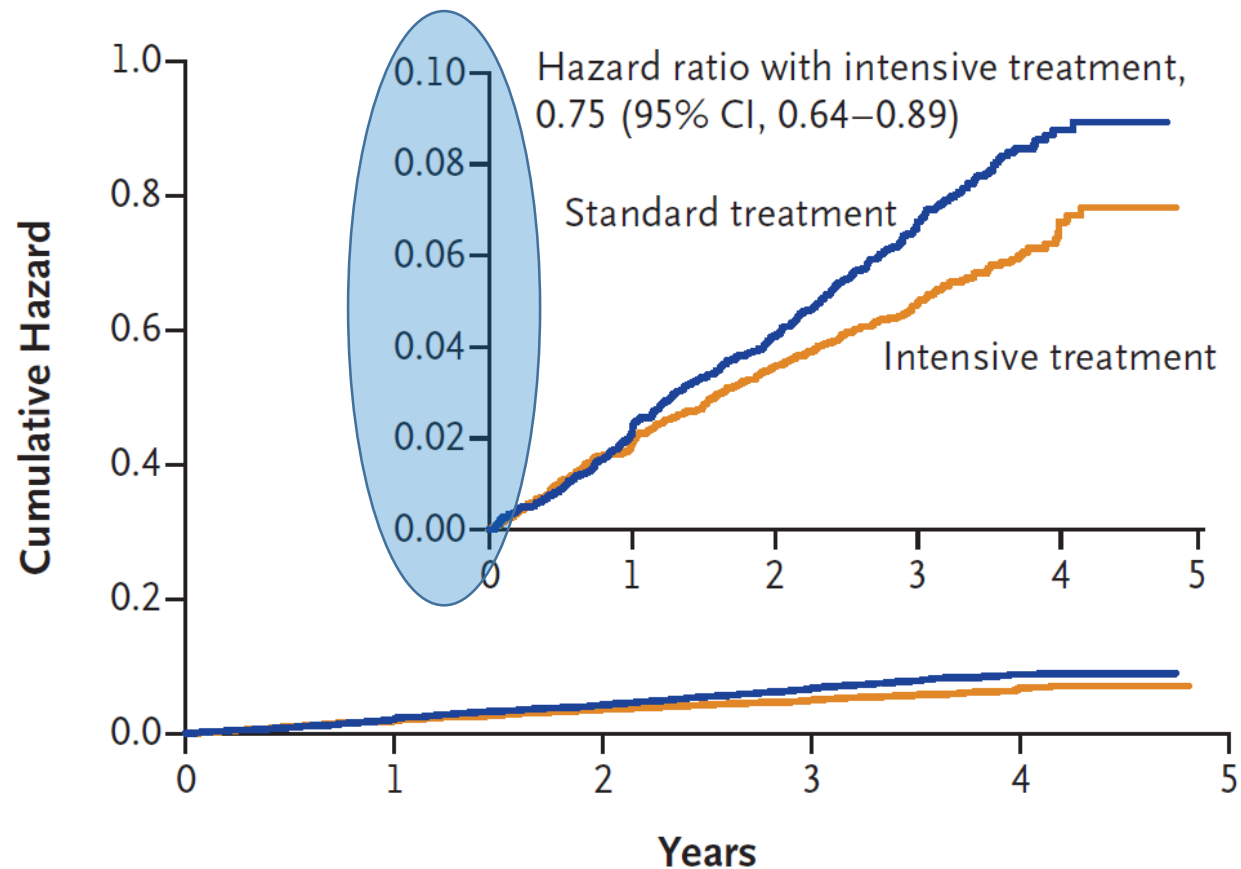
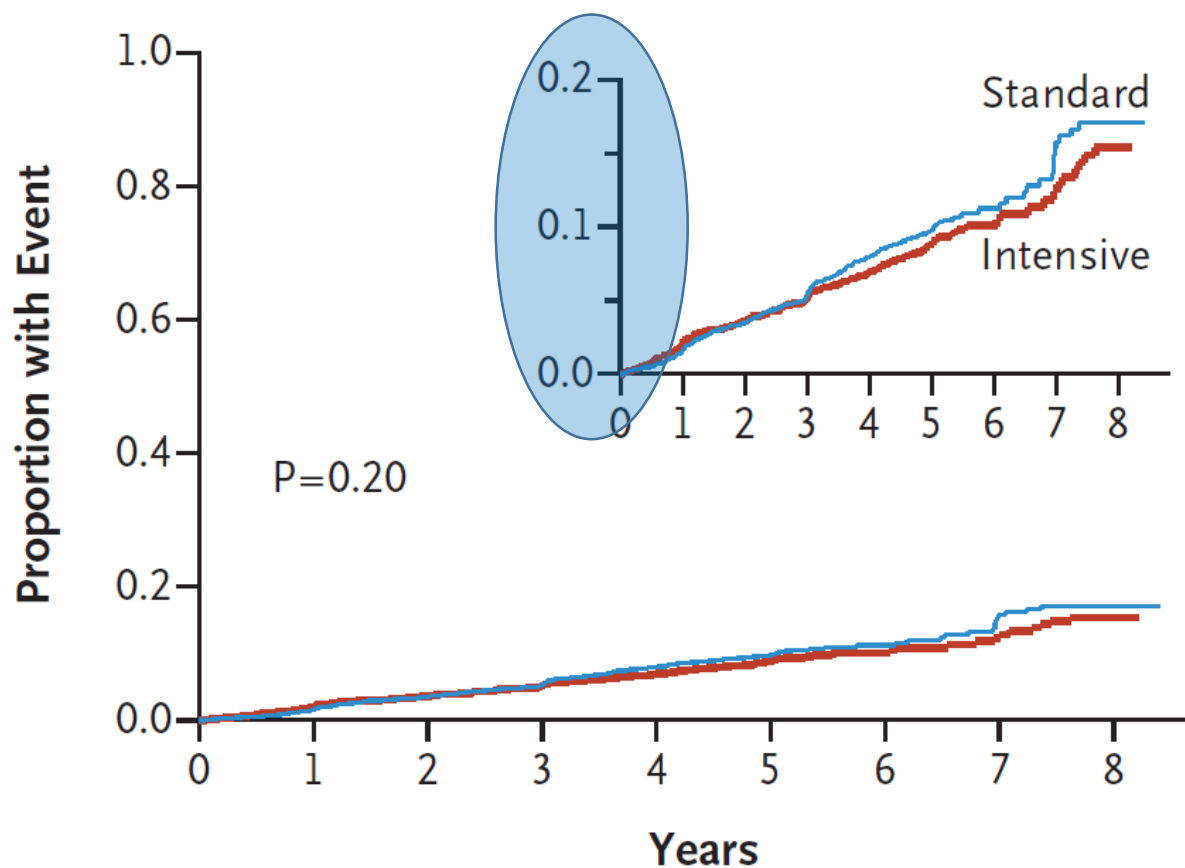
The ACCORD Study Group. N Engl J Med 2010;362:1575-85

The SPRINT Research Group. N Engl J Med 2015. DOI: 10.1056/NEJMoa1511939



ACCORD VS. SPRINT

APPLES VS ORANGES

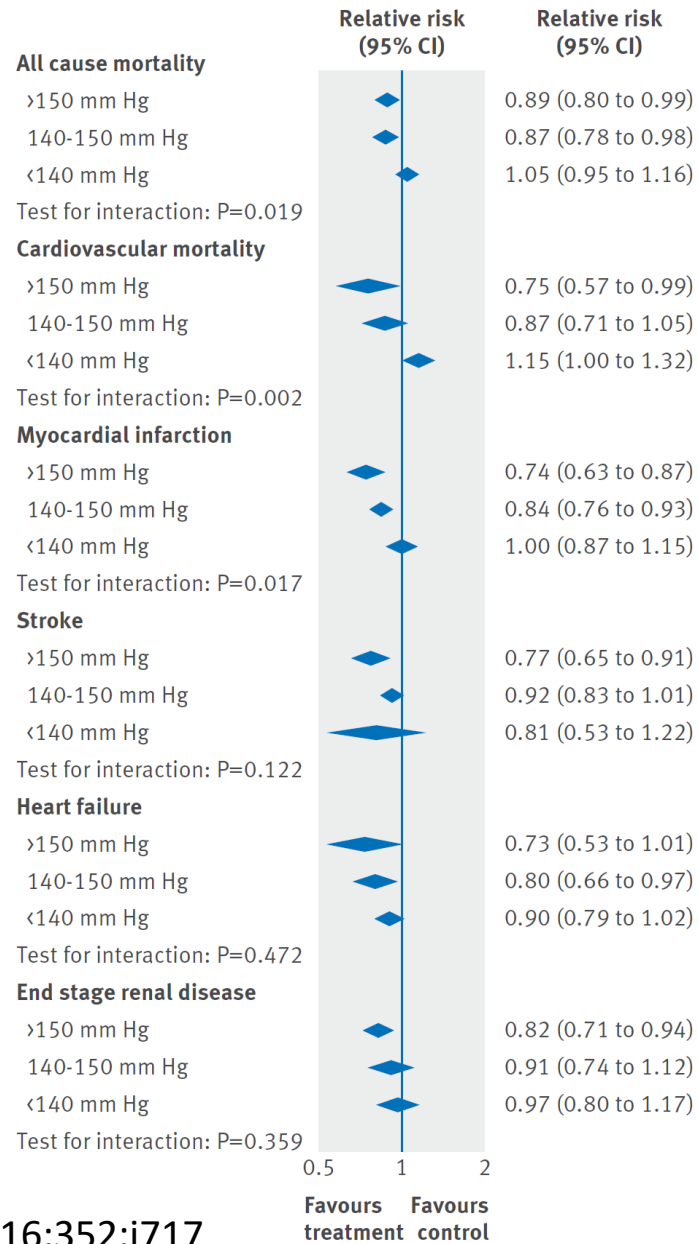


ACCORD Study Group. N Engl J Med 2010;362:1575-85

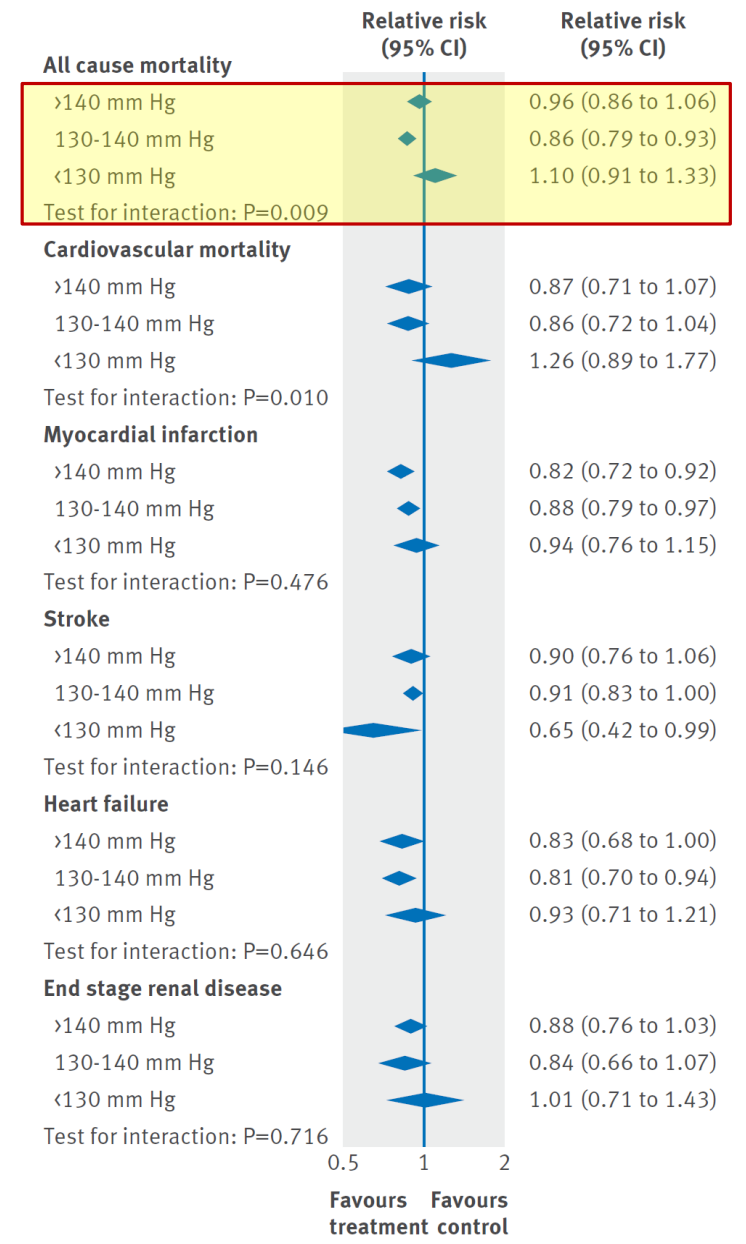
SPRINT Research Group. N Engl J Med 2015;26;373(22):2103-16

Strict BP targets in DM

Baseline SBP



Treated SBP



SBP target in SPRINT: Possible explanations for discrepant results?

- “Strict” vs. “conventional” SBP target
 - Achieved SBP of 121 vs 136 mmHg
 - Achieved DBP of 69 vs 76 mmHg
- BP measurement method
 - Examiner left the room and an automated device measured the patients’ BP after several minutes
- This method can result in measured BP levels that are ~5-20 mmHg lower compared to methods applied in clinical practice
 - Achieved SBP may have been equivalent to levels associated with ideal outcomes in observational studies and in DM trials (SBP ~130-140 mmHg)



SPRINT vs. all-comers

- SPRINT

- N=2,646 (CKD subgroup)
- Age 68 years (overall)
- Black race 31% (overall)
- Baseline SBP 140 mmHg
- eGFR 48 ml/min/1.73m² (CKD)
- Baseline number of AHD 1.8±1
- Deaths 6.2% (2% per year) (CKD)

- VA-CKD observational modeling

- N=77,765
- Age 74 years
- Black race 7%
- Baseline SBP 142 mmHg
- eGFR 49 ml/min/1.73m²
- Baseline number of AHD 2 (1-2)
- Deaths 25% (4.4% per year)

SPRINT Research Group. N Engl J Med 2015;26;373(22):2103-16

Kovesdy CP et al, JAMA Intern Med 2014;174(9):1442-1449

Kovesdy CP et al., Adv Chronic Kidney Dis. 2012 Jan;19(1):11-8.

Relative effect (i.e., hazard ratio) of intensive BP control may vary across CKD stages.

(Original report)



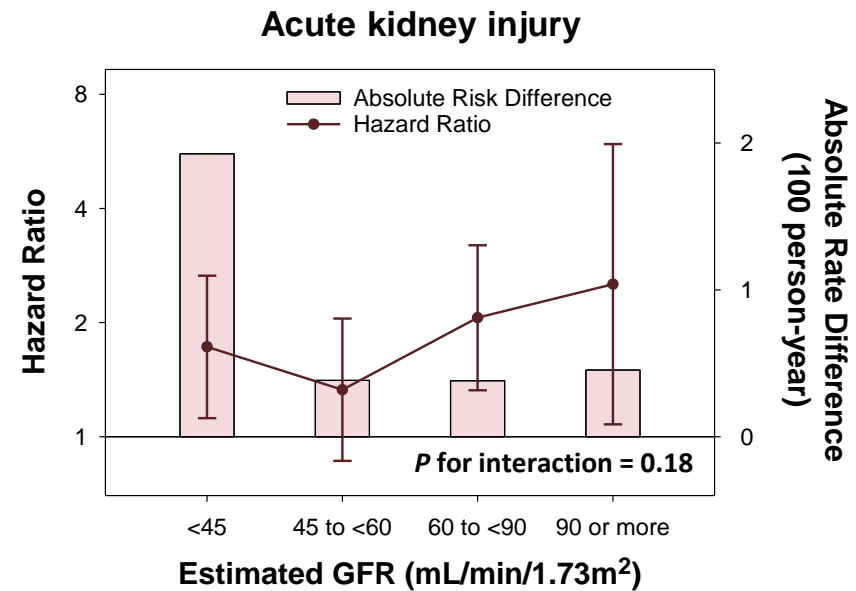
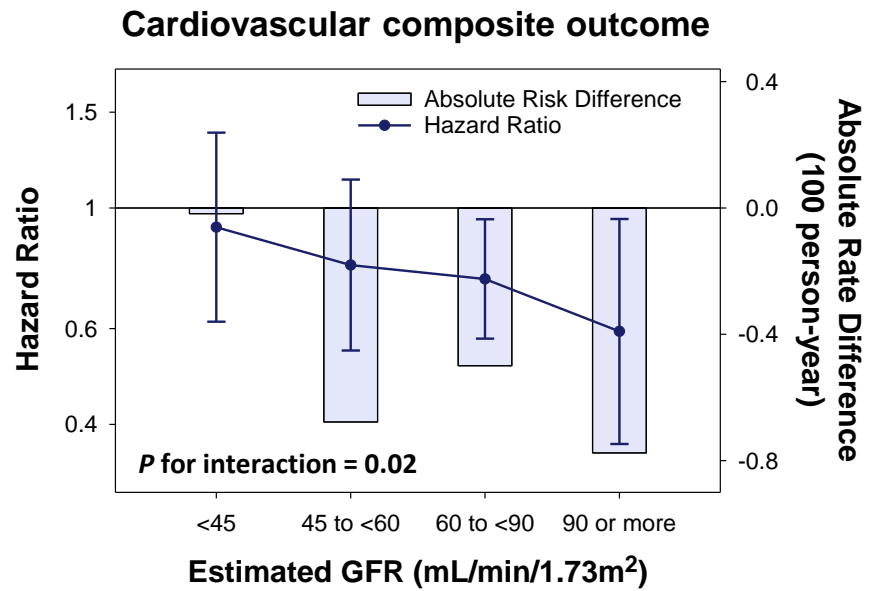
But what if ...

CKD stages

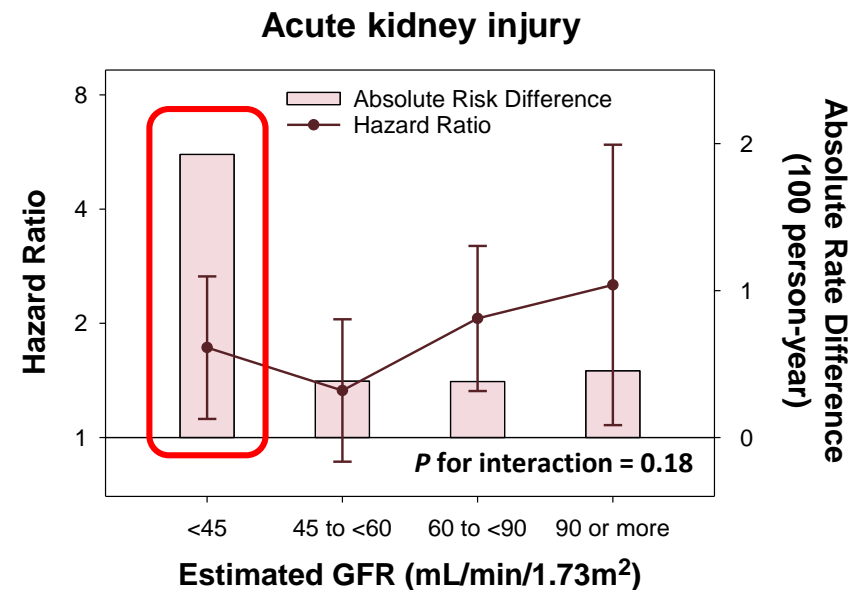
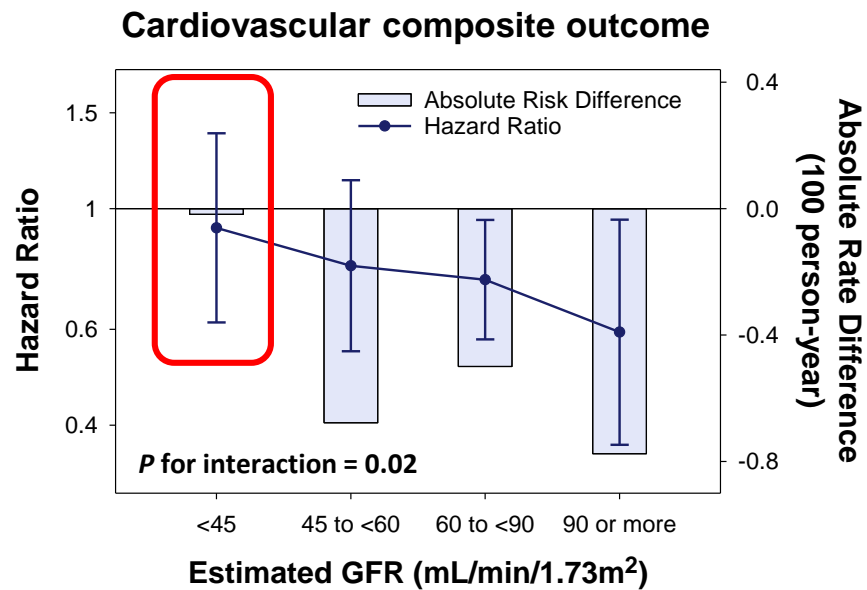
- 1
- 2
- 3a
- ≥3b

Additionally, there were no information on AKI risk in CKD vs. non-CKD.

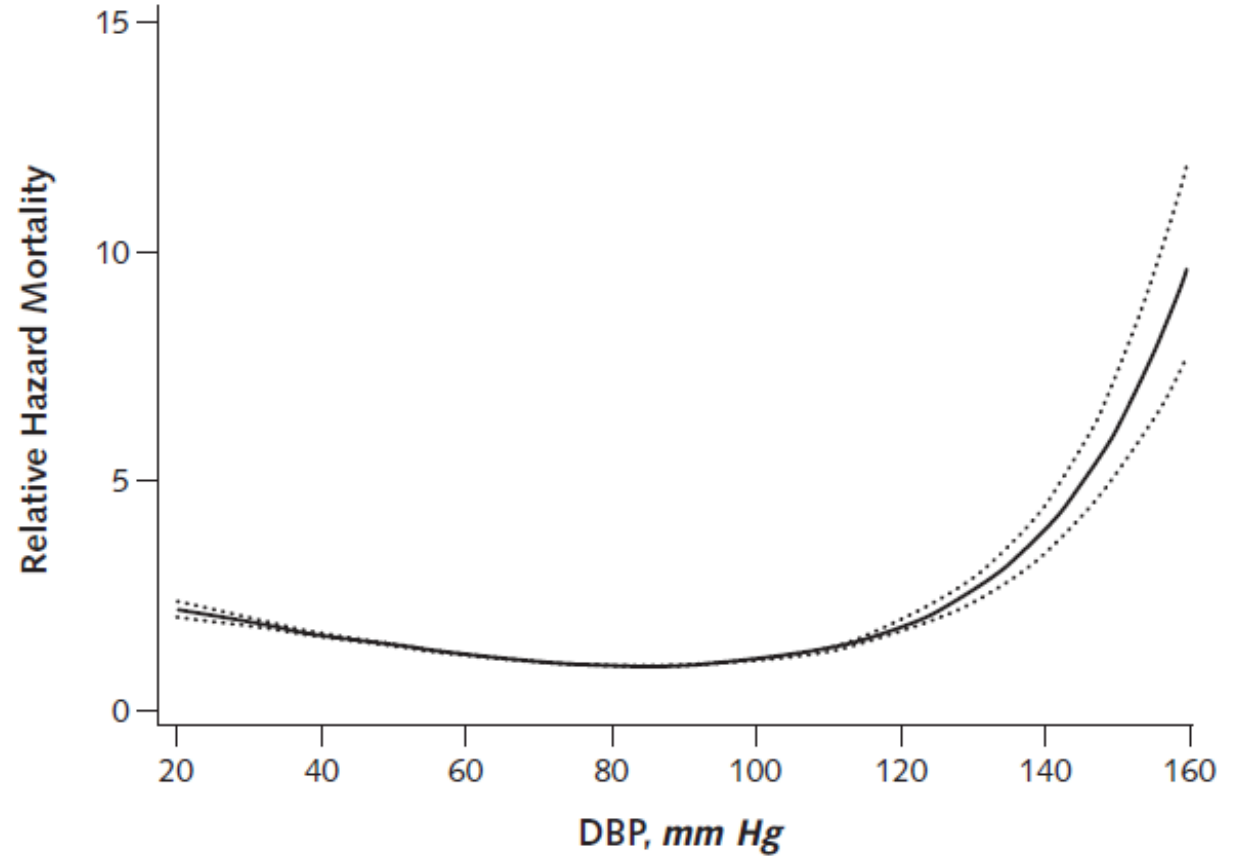
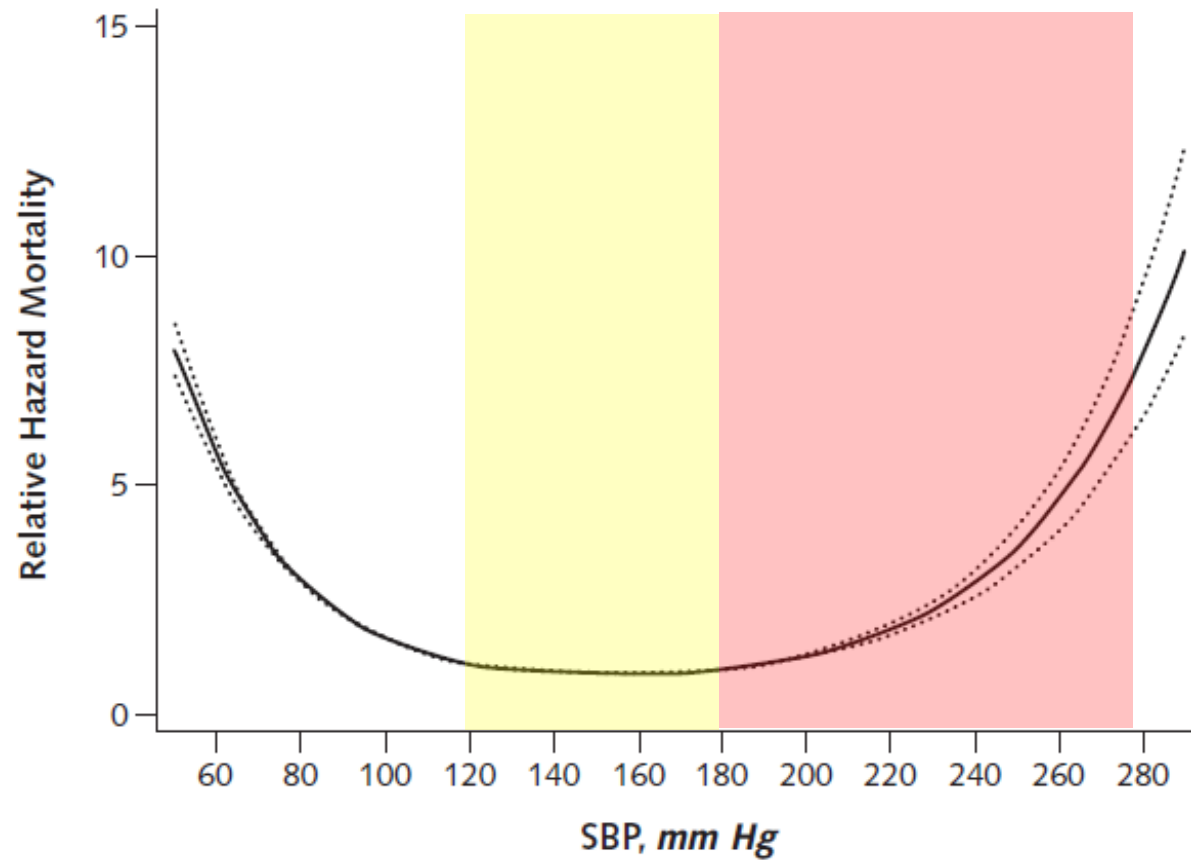
Intensive BP control may provide little benefit but potential harm among patients with advanced CKD
- Secondary analysis of the SPRINT study -



***Intensive BP control may provide little benefit but potential harm among patients with advanced CKD
- Secondary analysis of the SPRINT study -***



Multivariable-adjusted relative hazards (hazard ratios [95% CIs]) of all-cause mortality associated with SBP and DBP



Treating BP in CKD: Magnitude of benefit

- Baseline BP: 186/121 mmHg
 - Reduction in SBP: 43 mmHg
 - Reduction in DBP: 30 mmHg
- NNT to prevent 1 AE: 2.8
- NNT to prevent 1 death: 17.5

• Base

A storm in a tea cup

- M
- ir
- M
- co
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g
intensive
57
0

Kovesdy CP, JAMA Intern Med. 2017 Oct 1;177(10):1506-1507

VA Cooperative Study Group, JAMA. 1967;202(11):1028-1034

Malhotra R et al., JAMA Intern Med. 2017 Oct 1;177(10):1498-1505

Lv et al., CMAJ 2017 Oct 1;177(10):1498-1505

Potential benefits of lowering blood pressure

Reduction of all-cause mortality

Reduction of cardiovascular disease and mortality

Reduction of chronic kidney disease progression and the development of end-stage renal disease

Potential risks of lowering blood pressure

Reduction in organ perfusion due to altered auto-regulation:
- renal perfusion: increased risk of AKI
- cerebral perfusion: increased dizziness, falls

Significant reduction in diastolic blood pressure leading to myocardial hypoperfusion and a potential of increased all-cause mortality

Increased patient burden: polypharmacy, cost

Balance

Managing Hypertension in Patients with CKD: A Marathon, Not a SPRINT

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^{*}Division of Nephrology, Department of Medicine, Stanford University, Stanford, California; [†]Division of Nephrology, Internal Medicine School of Medicine, University of Utah, Salt Lake City, Utah; [‡]Division of Nephrology and Hypertension, Department of Medicine, Vanderbilt University, Nashville, Tennessee; and [§]Department of Clinical Science, University of Texas Southwestern, Dallas, Texas

Questions?