

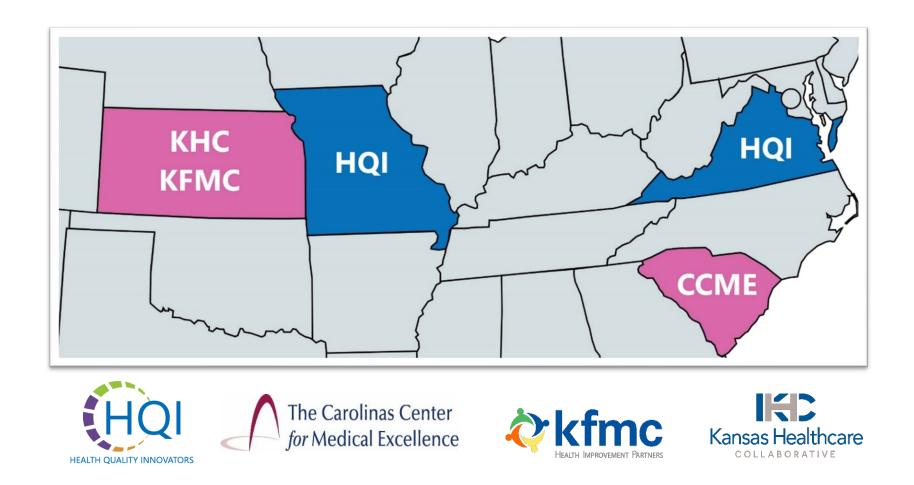


Importance of Assessing CVD Risk and Monitoring

September 7th, 2022



Health Quality Innovation Network







This HQIN **HEARTS in America** series is delivered by **HEARTS** subject matter experts. They are introducing the pillars of the <u>HEARTS Technical Package</u> while beginning the conversation about HEARTS in America.

If you would like to speak to a HEARTS Advisor, learn more about the initiative, and discuss possibilities for your organization, please connect with your HQIN Quality Improvement Advisor to begin the next steps.



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Non-physician Attendees

All non-physicians will receive a certificate of participation.





Disclosures

Disclosure Information

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Invited Faculty:

Andres Rosende, MD, MSc No financial relationships were declared

Logistics – Zoom Meeting



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Raise your hand if you want to verbally ask a question.

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You have been automatically muted with video turned off.

Purpose & Learning Objectives

- 1. Detail the CVD prevalence and impact on healthcare systems and delivery.
- 2. Introduction to the PAHO CVD Risk Assessment Tool.
- 3. Understanding the importance of completing CVD screenings and Risk Assessment Tool to engage and inform treatment and patients.

Andres Rosende, MD



Andres Rosende is a physician specializing in cardiology. He obtained a Master's degree in clinical research and epidemiology from the University of Buenos Aires, Argentina. In 2018, and after several years of clinical practice, Andres began to work in the Ministry of Health of Argentina as the Coordinator of the National Program for Cardiovascular Disease Prevention, leading the implementation of the HEARTS initiative in the country.

Since 2021, he has been working as an International PAHO Consultant for HEARTS in the Americas Initiative, specifically, overseeing the Medication and Standardized Treatment Protocols Pillar.





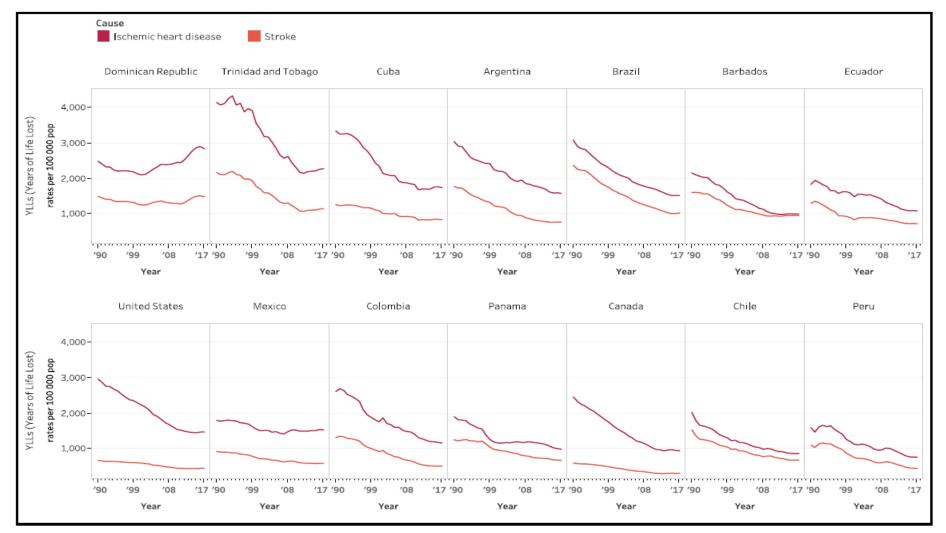
Importance of Assessing CVD Risk and Monitoring

Andres Rosende, MD, MSc

International PAHO/WHO Consultant HEARTS in the Americas



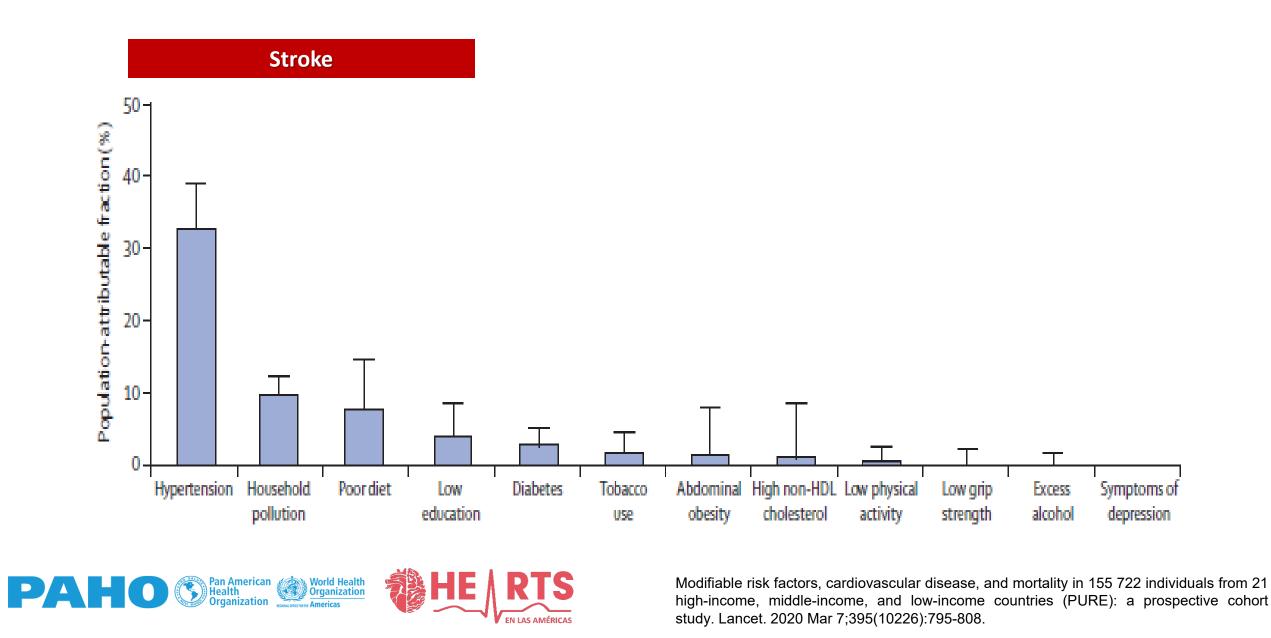
Slowdown in Premature CVD Mortality Reduction in the Americas



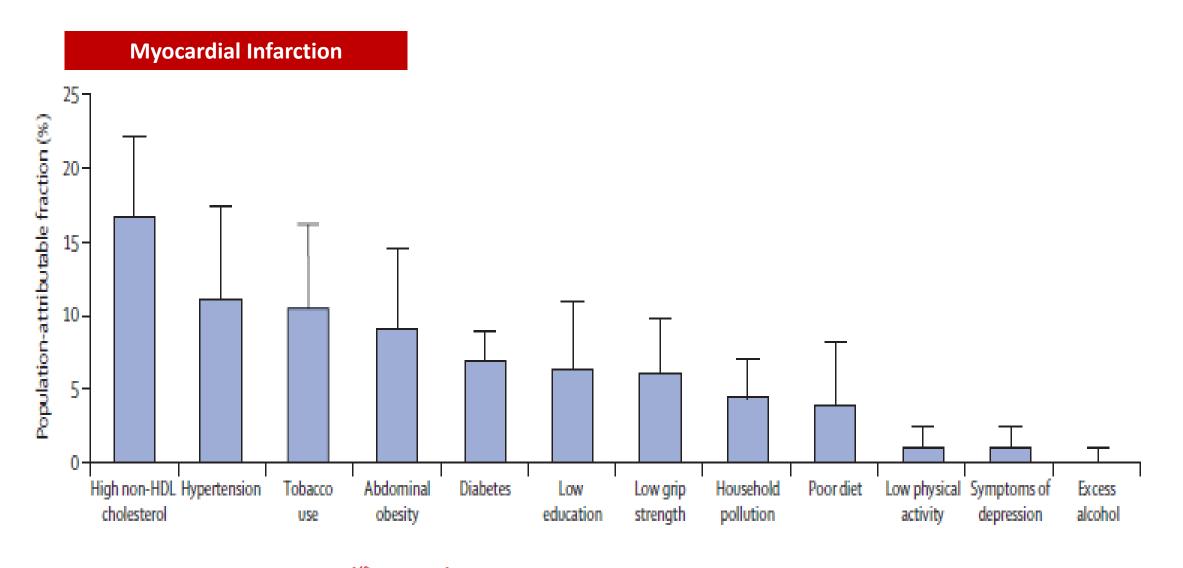


Martinez R, Soliz P, Mujica OJ, Reveiz L, Campbell NRC, Ordunez P. The slowdown in the reduction rate of premature mortality from cardiovascular diseases puts the Americas at risk of achieving SDG 3.4: A population trend analysis of 37 countries from 1990 to 2017. J Clin Hypertens (Greenwich). 2020 Aug;22(8):1296-1309.

HTN Attributable Risk Factors



HTN Attributable Risk Factors

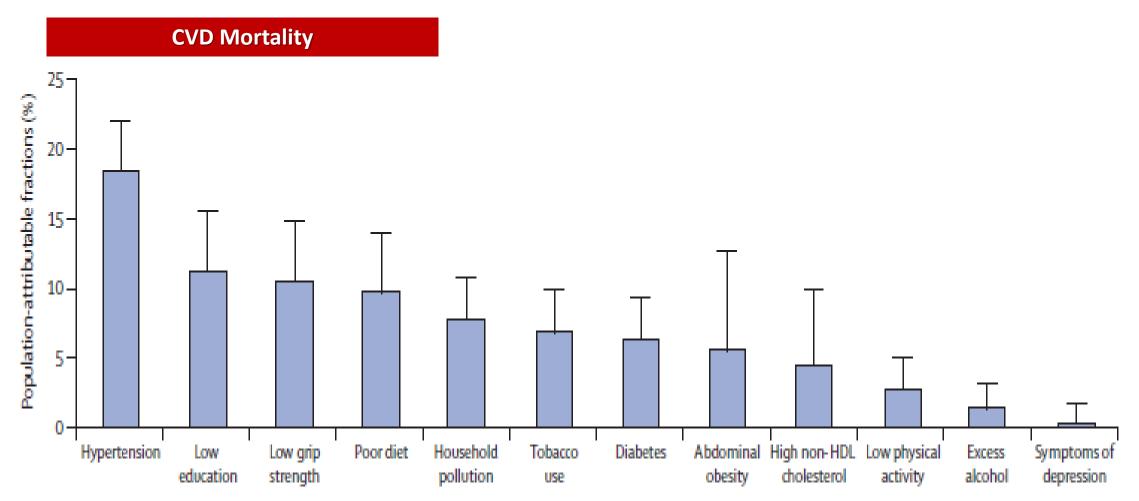


Pan American Health Organization

PA

Modifiable risk factors, cardiovascular disease, and mortality in 155 722 individuals from 21 high-income, middle-income, and low-income countries (PURE): a prospective cohort study. Lancet. 2020 Mar 7;395(10226):795-808.

HTN Attributable Risk Factors



Pan American Health Organization World Health Organization World Health Organization Organiza

PA



Healthy Lifestyle

Lifestyle changes	Average Reduction of SBP
Low sodium intake	5 mmHg
Weight loss	12 mmHg per 10 kg lost
DASH Diet	11 mmHg
Regular physical activity (>120 min/week)	6 mmHg

Midgley et al. Effect of reduced dietary sodium on blood pressure: meta-analisis of RCT. JAMA 1996; 275:1590-1597.

Neter et al. Influence of weight reduction on blood pressure: metaanalysis of RCT. Hypertension 2003; 42:878-884.



Apple et al. A clinical trial of effects of dietary patterns on blood pressure. DASH CRG. N Eng J Med 1997;336:1117-1124.

Halbert et al. The effectiveness of excercise training in lowering blood pressure: a meta-analysis of randomised, controlled clinical trials of 4 weeks or longer. J Hum Hypertens 1997;11:641-649.



Pharmacologic Monotherapy Treatment

Average SBP reduction (mmHg) compared to placebo by drug class.

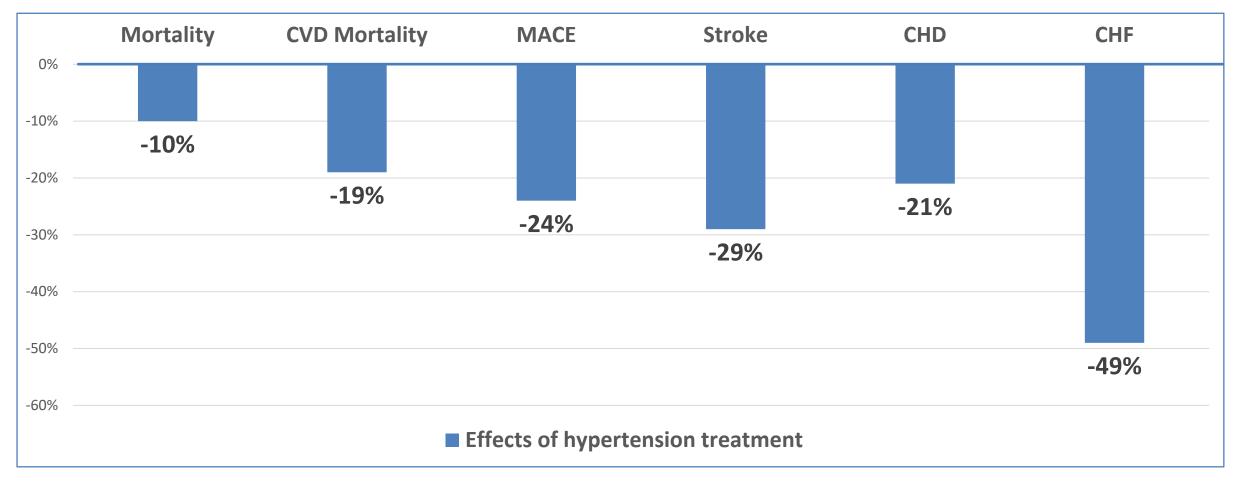
Drug	Half maximum dose			
Thiazides	8.8			
Beta Blockers	9.2			
Angiotensin Converting-Enzyme Inhibitors	8.5			
Calcium Channel Blockers	8.8			
Angiotensin Receptor Blockers	10.3			
Any drug	9.1			



Law MR, Wald NJ, et al. Value of low dose combination treatment with blood pressure lowering drugs: analysis of 354 RCT. BMJ 2003;326:1427-1434.



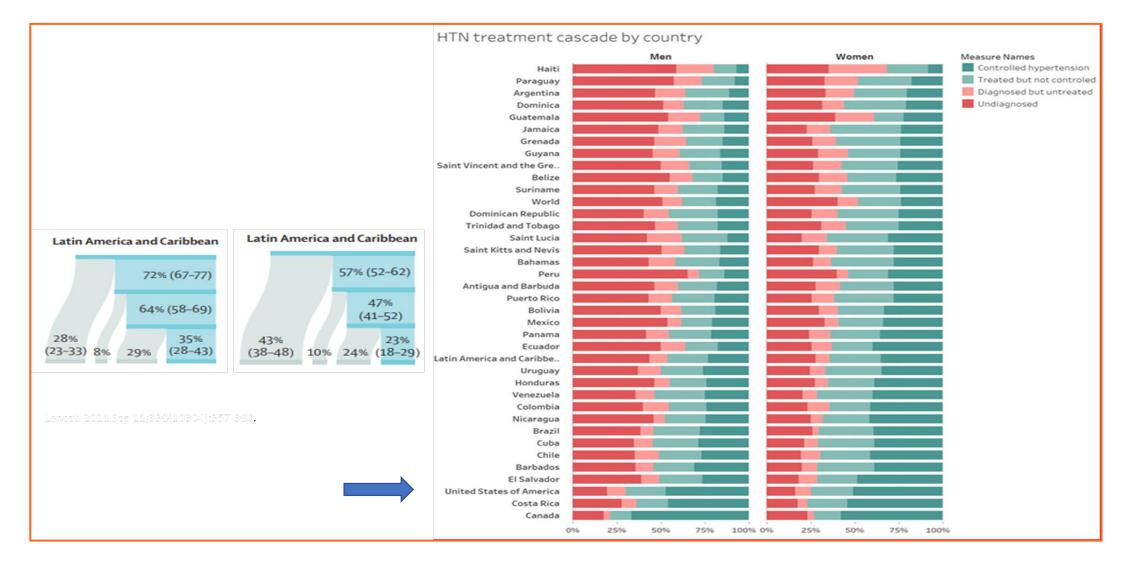
Benefits of Pharmacologic Treatment





Psaty BM, et al. Health outcomes associated with various antihypertensive therapies used as first-line agents. A network meta-analysis. JAMA 2003;289:2534-2544.

Hypertension in America

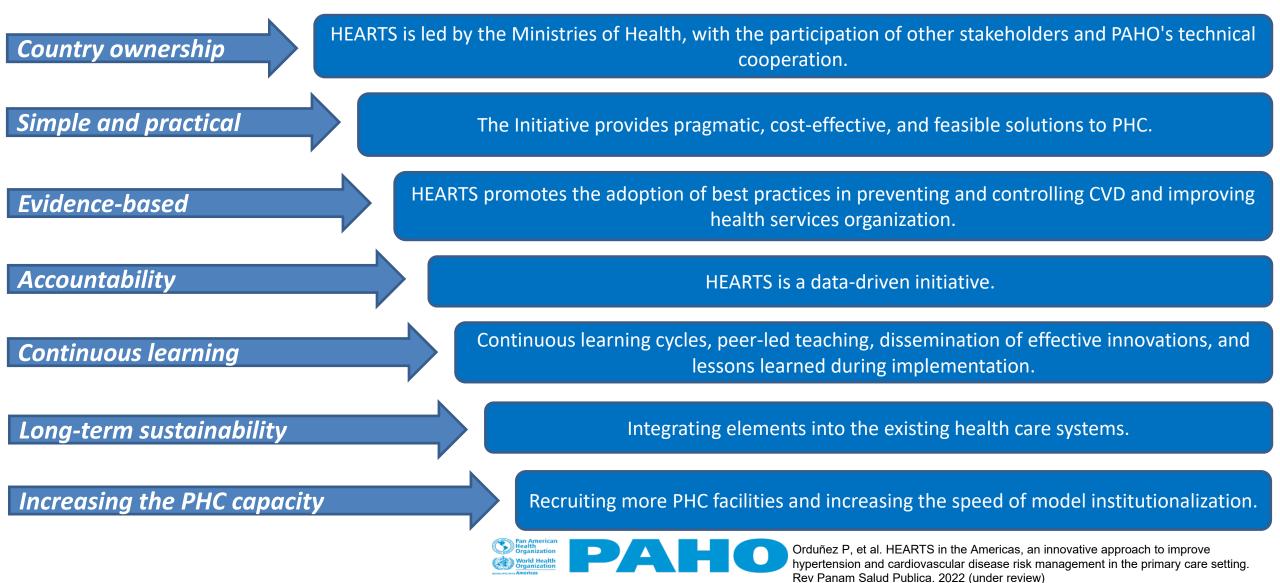




NCD Risk Factor Collaboration (NCD-RisC). Worldwide trends in hypertension prevalence and progress in treatment and control from 1990 to 2019: a pooled analysis of 1201 population-representative studies with 104 million participants. Lancet. 2021 Sep 11;398(10304):957-980.



Guiding Principles



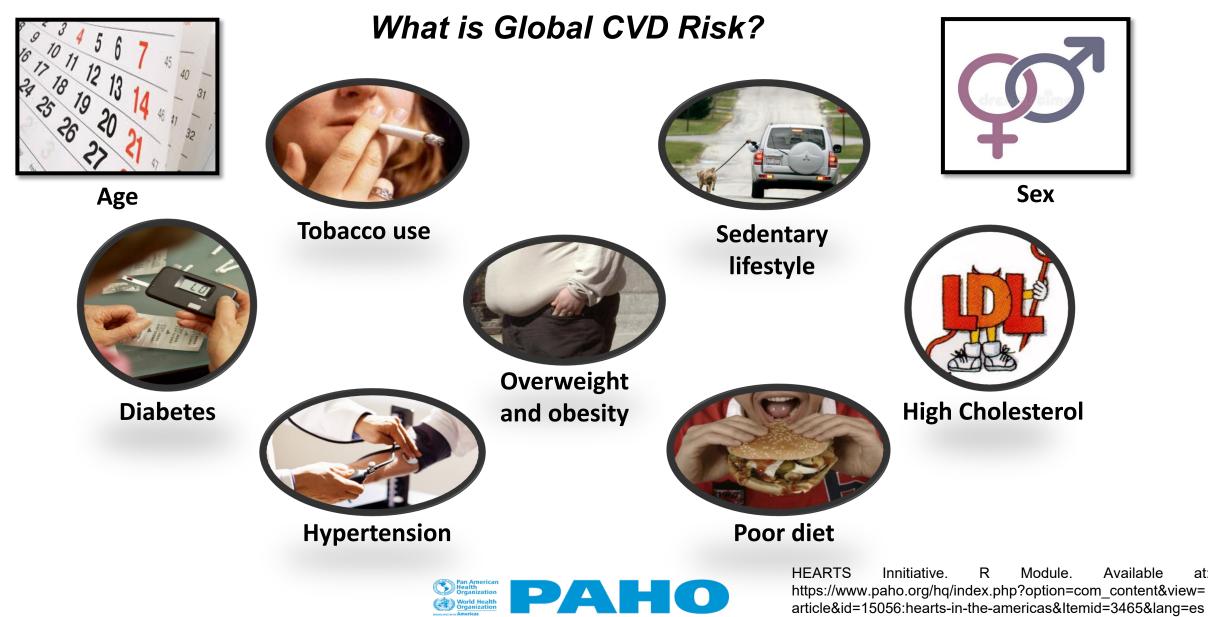


Treatment and Control of Hypertension is the TOP priority But...Is That Enough? Importance of Knowing and Treating CVD Risk





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at:



WHO Guidelines for the Pharmacological Treatment of Hypertension in Adults: What about CVD risk?

Guideline for the pharmacological treatment of hypertension in adults



WHO recommends initiation of pharmacological antihypertensive treatment of individuals with a confirmed diagnosis of hypertension and SBP of >/= 140 mmHg or DBP of >/= 90 mmHg. (Strong recommendation, moderate-high certainty evidence)

WHO recommends pharmacological antihypertensive treatment of individuals with existing cardiovascular disease and systolic blood pressure of 130-139 mmHg (Strong recommendation, moderate-high certainly evidence)

WHO suggests pharmacological treatment of individuals without cardiovascular disease but with high cardiovascular risk, diabetes mellitus, chronic kidney disease, and a SBP of 130-139 mmHg (Conditional recommendation, moderate-high certainly evidence)

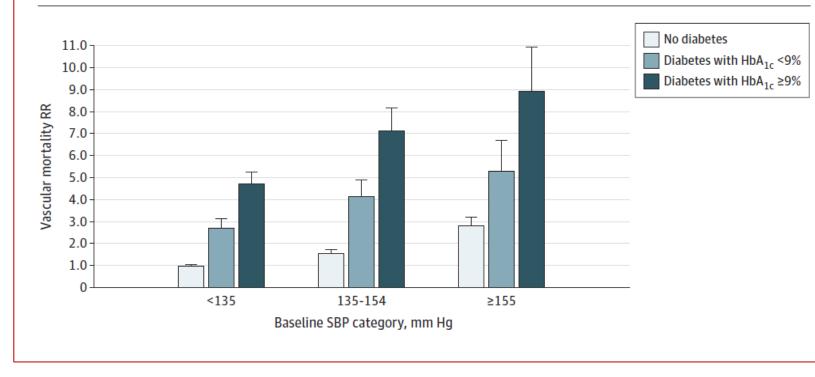
Implementation remarks: Treatment should start no later than 4 weeks after diagnosis. If BP >/= 160/90 mmHg or end organ damage is present-start without delay



World Health Organization. Guideline for the pharmacological treatment of hypertension in adults. Geneva: WHO; 2021. Available at: https://apps.who.int/iris/bitstream/handle/10665/344424/9789240033986-eng.pdf

CVD Mortality Associated with Systolic Blood Pressure and History of Diabetes.

Figure 3. Absolute Excess Vascular Mortality Between Ages 35 and 74 Years Associated With Systolic Blood Pressure (SBP) by History and Control or Previously Diagnosed Diabetes



Analyses excluded participants with prior chronic diseases (ie, ischemic heart disease, stroke, chronic kidney disease, cirrhosis, cancer, or emphysema) apart from diabetes. For the 9 groups shown, the mortality rate ratio (RR) estimates are stratified by age-at-risk (in 5-year ranges) and adjusted for sex, district of residence, highest education level attained, smoking status, alcohol intake, leisure-time physical activity, and measures of anthropometry. The error bar extending above each column extends to the upper 95% confidence limit of the RR. The mean usual SBP in the 3 SBP categories shown was 121, 139, and 158 mm Hg, respectively.

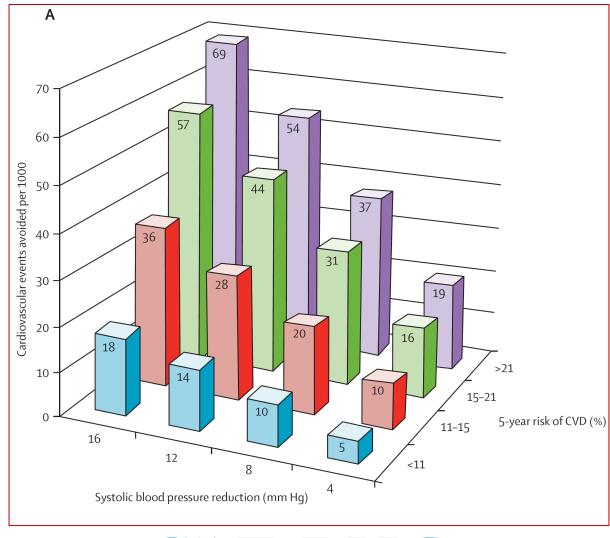


Tapia-Conyer R et al. Association of Blood Pressure With Cause-Specific Mortality in Mexican Adults. JAMA Netw Open. 2020 Sep 1;3(9):e2018141.



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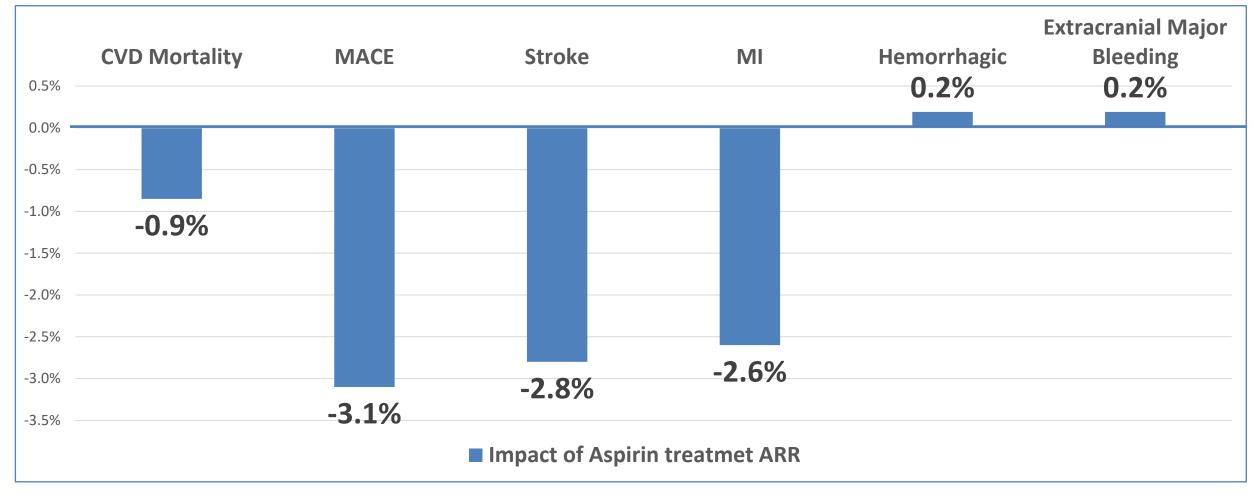
Impact of BP Reductions by CVD Risk Level







Aspirin in Secondary Prevention

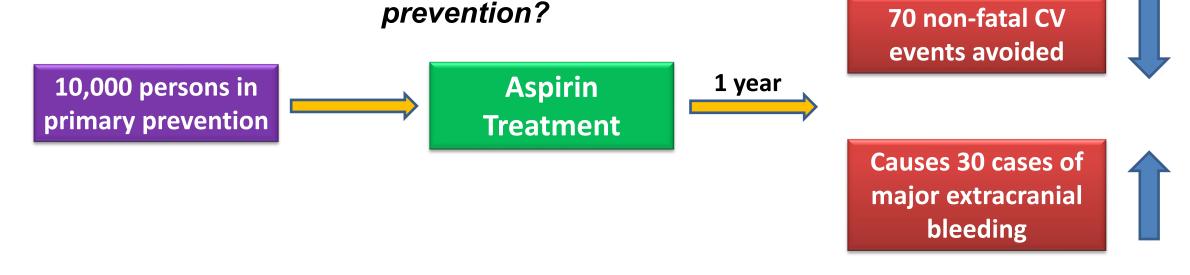




ATT Collaboration. Aspirin in the primary and secondary prevention of vascular disease: collaborative meta-analysis of individual participant data from randomized trials. Lancet. 2009;373:1849–60.



What is the impact of aspirin use in primary



Aspirin in Primary Prevention does not reduce total mortality nor cardiovascular mortality

Aspirin in Primary Prevention does not reduce Stroke

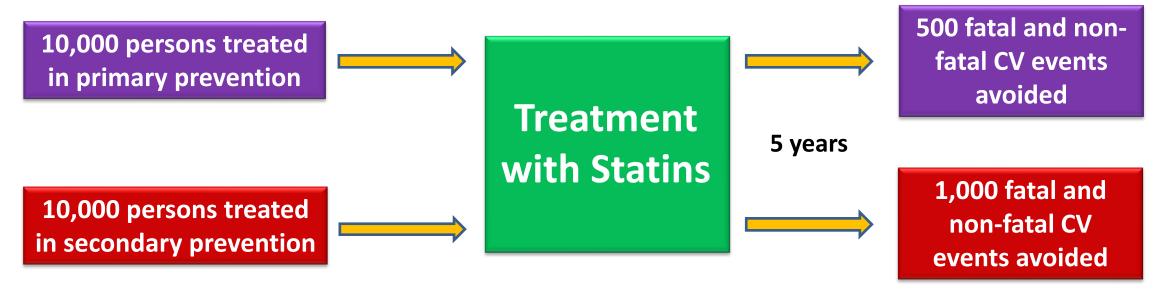
The net clinical benefit is scarce



ATT Collaboration. Aspirin in the primary and secondary prevention of vascular disease: collaborative meta-analysis of individual participant data from randomized trials. Lancet. 2009;373:1849–60.



Impact of statin therapy



Excellent safety profile

High impact over CV events and mortality

Collins R, Reith C, Emberson J, Armitage J, Baigent C, Blackwell L, et al. Interpretation of the evidence for the efficacy and safety of statin therapy. Lancet. 2016 Nov 19;388(10059):2532–61.

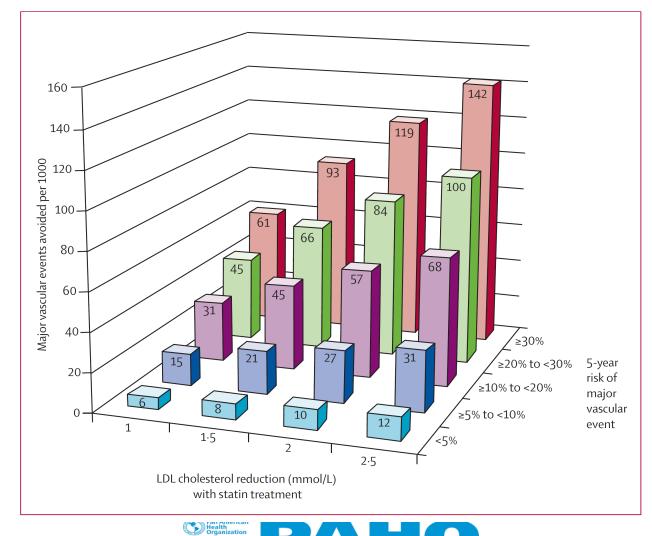


Cholesterol Treatment Trialists' (CTT) Collaboration, Baigent C,Blackwell L,Emberson J,Holland LE,Reith C,et al. Efficacy and safety of more intensive lowering of LDL cholesterol: a meta-analysis of data from 170,000 participants in 26 randomised trials. Lancet. 2010 Nov 13, 376(9753):1670–81.



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Impact of Statin Therapy by CVD Risk Level



Sundstrom J, Arima H, Woodward M, et al. Lancet. 2014;384:591-8



Why should we estimate CVD risk?

1- To establish different treatment regimens (Aspirin, Statins)

2- To establish the need of more intensive BP thresholds and targets.

3- To establish diffrent follow-up intervals for control.

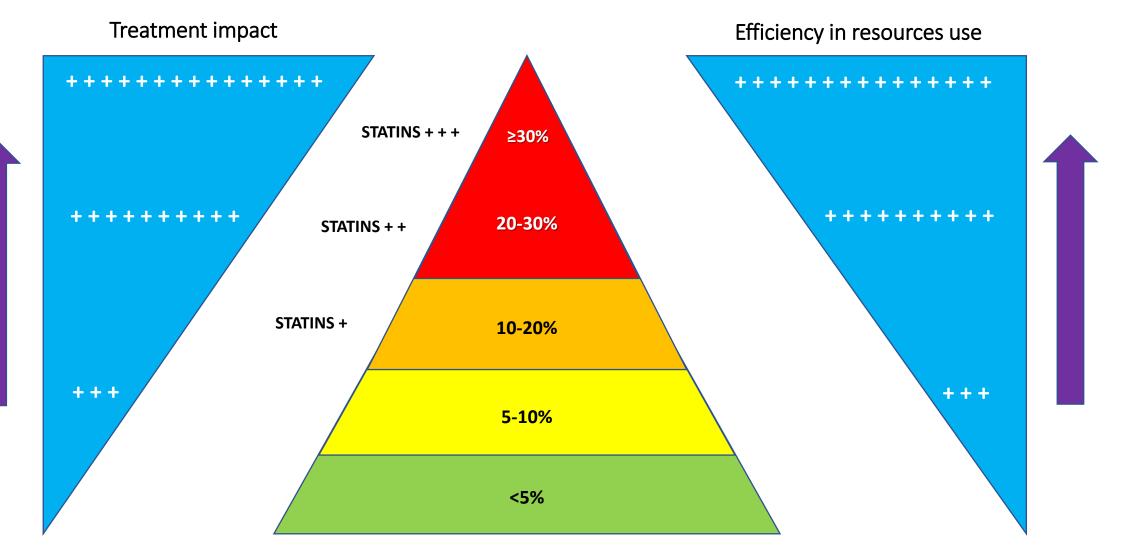
4- To assess the need for consultation with a specialist.

To improve healthcare with efficient use of resources and impact the reduction of CVD and premature death.



HEARTS Innitiative. R Module. Available at: https://www.paho.org/hq/index.php?option=com_content&view= article&id=15056:hearts-in-the-americas&Itemid=3465&Iang=es

Primary prevention



Proportion of individuals by CVD risk strata

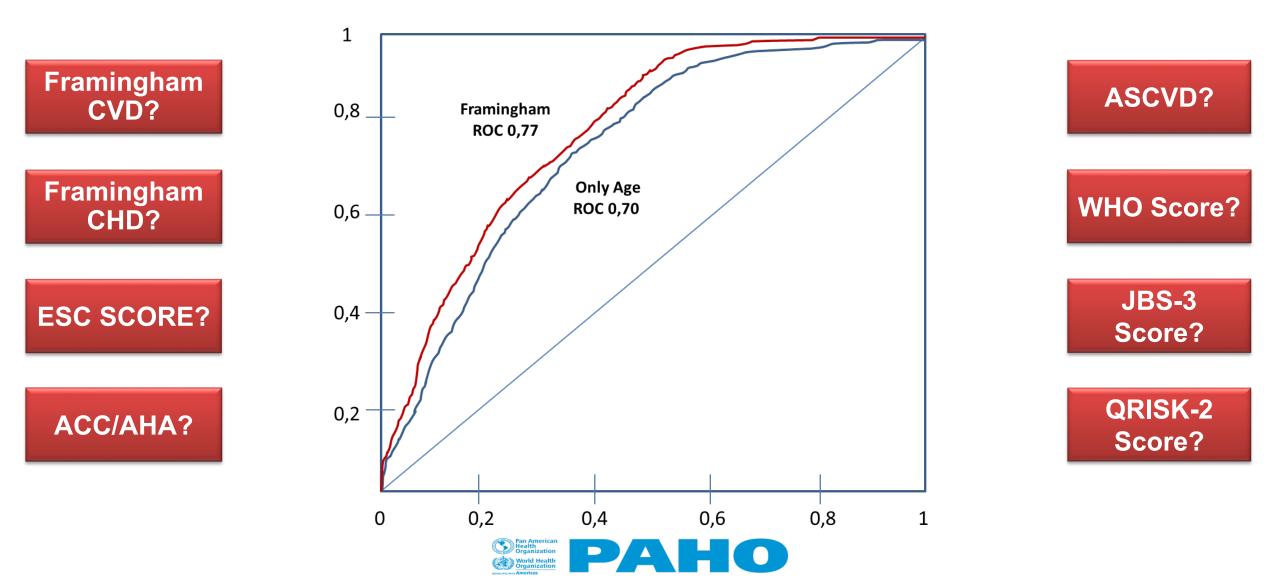




HEARTS Innitiative. R Module. Available at: https://www.paho.org/hq/index.php?option=com_content&view= article&id=15056:hearts-in-the-americas&Itemid=3465&Iang=es

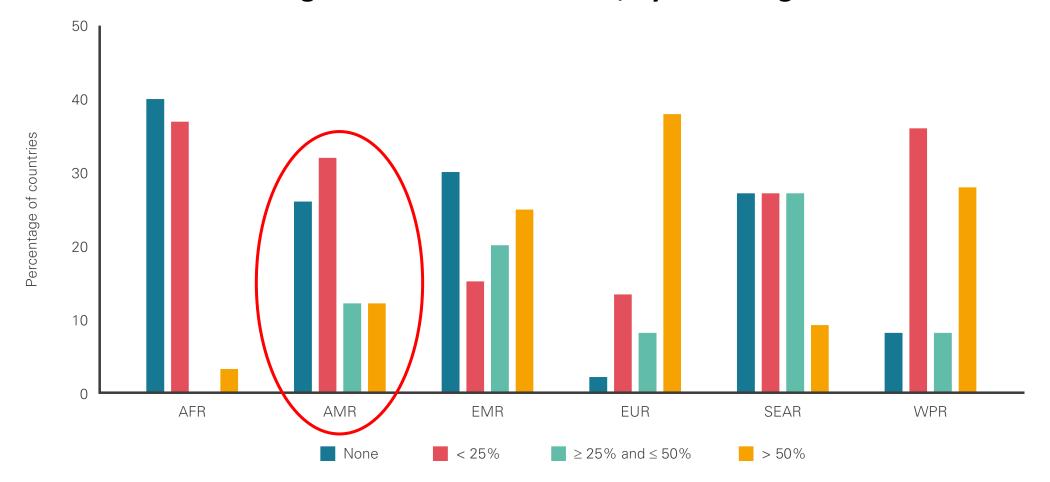


What is the best CVD risk score?





Primary health care centers offering CVR stratification for the management of patients at high risk of AMI and stroke, by WHO Region

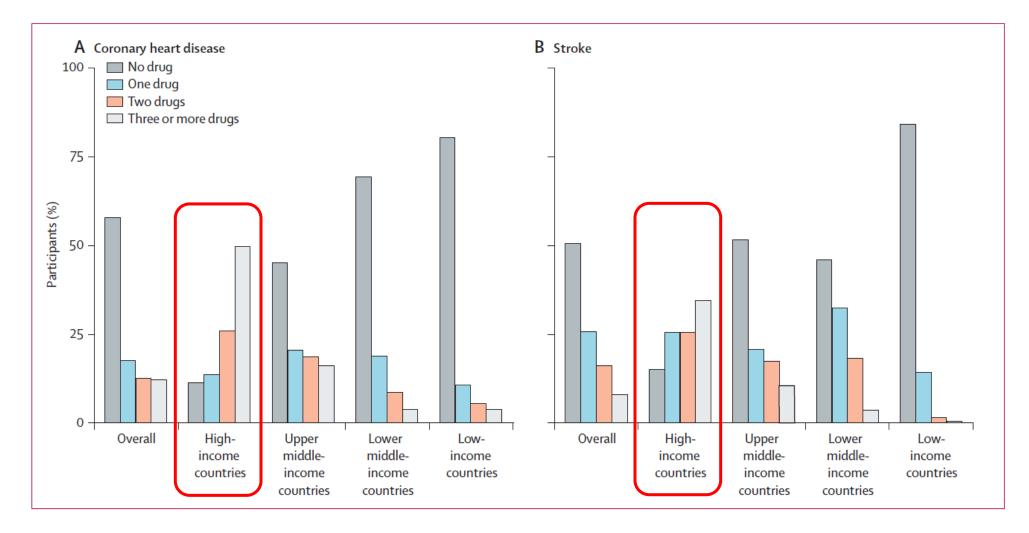




WHO. Hearts: technical package for cardiovascular disease management in primary health care. 2016



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Yusuf S, et al. Prospective Urban Rural Epidemiology (PURE) Study Investigators. Use of secondary prevention drugs for cardiovascular disease in the community in high-income, middle-income, and low-income countries (the PURE Study): a prospective epidemiological survey. Lancet. 2011 Oct 1;378(9798):1231-43.



The HEARTS APP



CVD RISK CALCULATOR

FIND OUT YOUR RISK OF HEART DISEASE AND HOW TO LOWER IT

This **FREE** app is a cardiovascular risk calculator that estimates

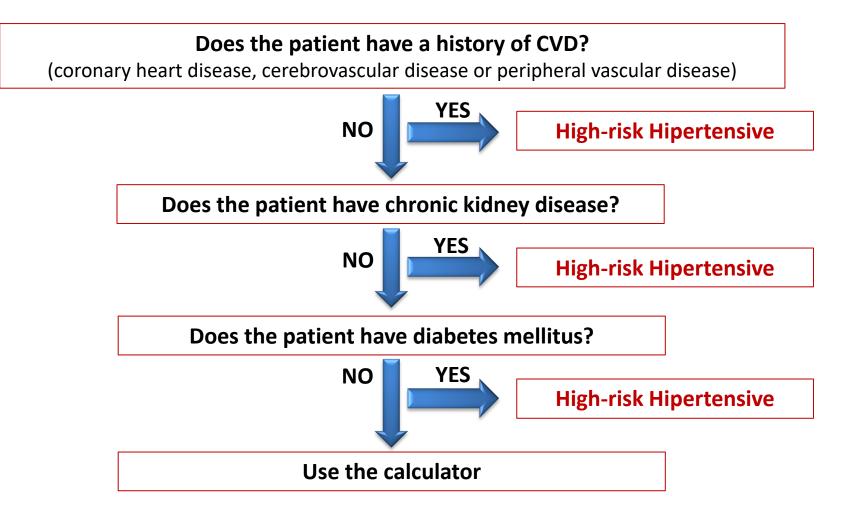
the possible 10-year risk of myocardial infarction stroke or



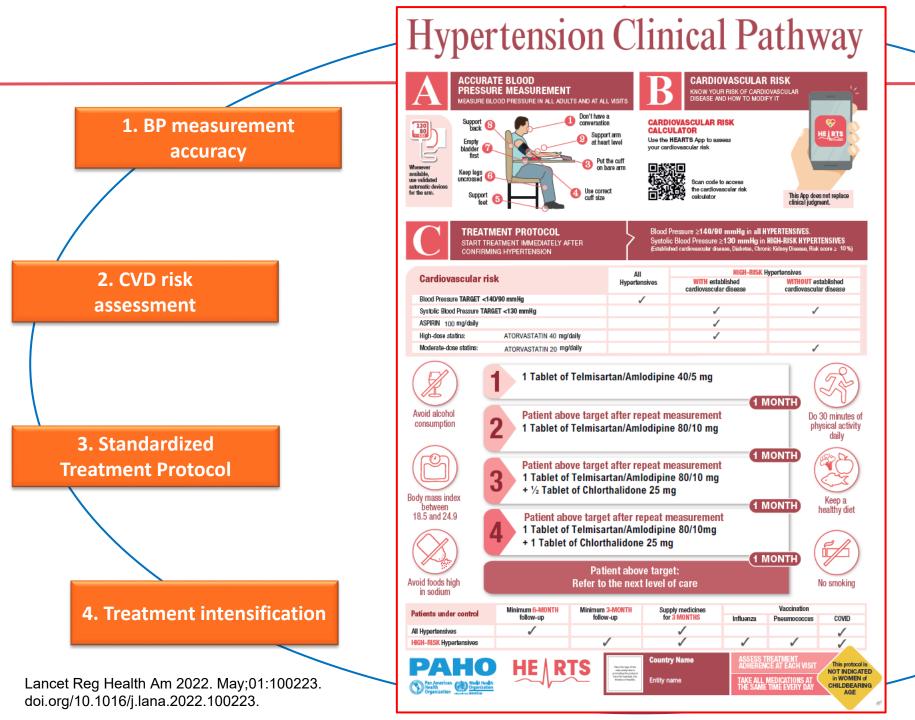
Available at: https://www.paho.org/en/hearts-americas/cardiovascular-risk-calculator-app

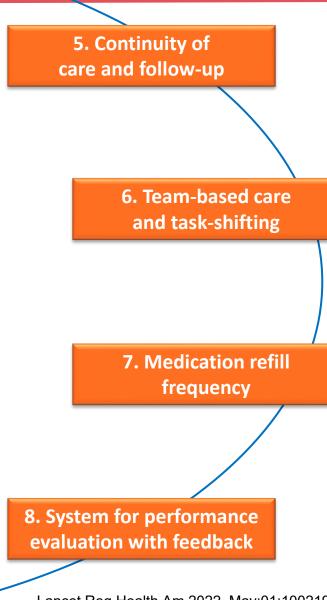
Ordunez P et al. The HEARTS app: a clinical tool for cardiovascular risk and hypertension management in primary health care. Rev Panam Salud Publica. 2022 Mar 28;46:e12.











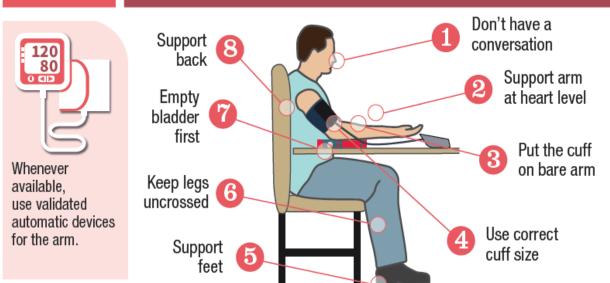
Lancet Reg Health Am 2022. May;01:100219. doi.org/10.1016/j.lana.2022.100219



Hypertension Clinical Pathway

ACCURATE BLOOD PRESSURE MEASUREMENT

MEASURE BLOOD PRESSURE IN ALL ADULTS AND AT ALL VISITS



KNOW YOUR RISK OF CARDIOVASCULAR DISEASE AND HOW TO MODIFY IT

CARDIOVASCULAR RISK

CARDIOVASCULAR RISK CALCULATOR

Use the **HEARTS** App to assess your cardiovascular risk



Scan code to access the cardiovascular risk calculator



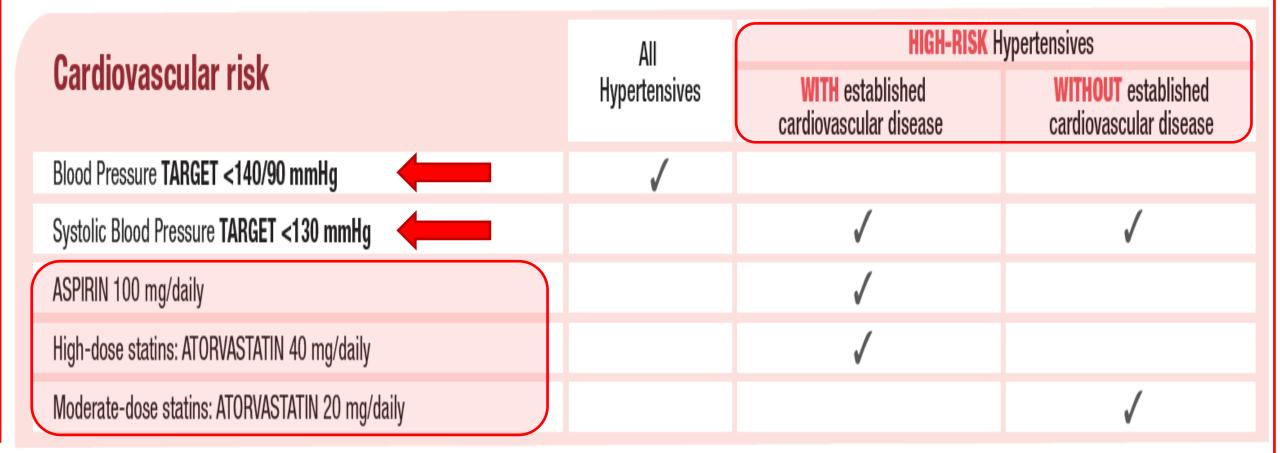


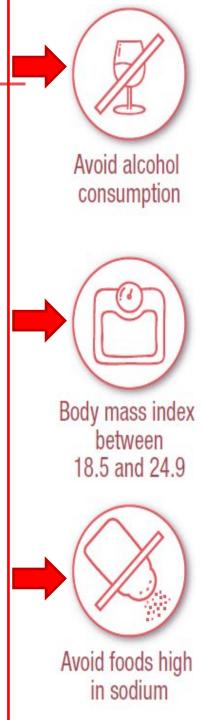
TREATMENT PROTOCOL

START TREATMENT IMMEDIATELY AFTER START TREATMENT IMMEDIATELY AFTER CONFIRMING HYPERTENSION

Blood Pressure ≥140/90 mmHg in all HYPERTENSIVES.

Blood Pressure ≥140/90 mmHg in all HYPERTENSIVES. Systolic Blood Pressure ≥130 mmHg in HIGH-RISK HYPERTENSIVES (Established cardiovascular disease, Diabetes, Chronic Kidney Disease, Risk score ≥ 10%)





3

1 Tablet of Telmisartan/Amlodipine 40/5 mg

Patient above target after repeat measurement 1 Tablet of Telmisartan/Amlodipine 80/10 mg

Patient above target after repeat measurement 1 Tablet of Telmisartan/Amlodipine 80/10 mg + 1/2 Tablet of Chlorthalidone 25 mg

Patient above target after repeat measurement 1 Tablet of Telmisartan/Amlodipine 80/10mg + 1 Tablet of Chlorthalidone 25 mg

> Patient above target: Refer to the next level of care



Patients under control	Minimum <mark>6-MONTH</mark> follow-up	Minimum <mark>3-MONTH</mark> follow-up	Supply medicines for 3 MONTHS	Vaccination Influenza Pneumococcus COVID		
All Hypertensives	\checkmark					
HIGH-RISK Hypertensives				v	V	
Country name Entity name		HEARTS	TAKE ALL MEDICATIONS AT THE SAME TIME EVERY DAY			This protocol is NOT INDICATED in WOMEN of CHILDBEARING
				T		AGE

The Hypertension Clinical Pathway is the main tool for HEARTS implementation, promoting a comprehensive approach of CVD risk management in PHC.

Rosende A, DiPette D, Brettler J, Rodríguez G, Zuniga E, Connell K, et al. HEARTS in the Americas appraisal checklist and clinical pathway for comprehensive hypertension management in primary care. Rev Panam Salud Publica. 2022;46:e125.



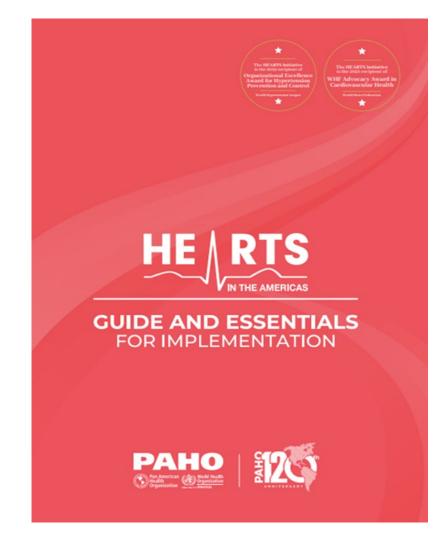


Conclusion

Hypertension treatment is the top priority, but this treatment is best tailored according to the level of CVD risk, as recommended by the HEARTS Clinical Pathway.

The HEARTS Clinical Pathway allows a pragmatic approach to CVD risk, strengthening the secondary prevention through the appropriate indication of statins and aspirin.





Questions?

Join us for our next HEARTS in America event:

Hypertension, Diabetes, and Chronic Disparities Within Healthcare

September 21st @ 12:15pm

CME Process

CME credit and certificate distribution are managed through SMA's **online process**. Within one week after the conclusion of the webinar, **please be on the lookout for an email from the Southern Medical Association (customerservice@sma.org) that will include your unique link to an online form** to complete the evaluation, attendance attestation, and claim credit. Please review the following process to receive your certificate awarding credit (for physicians), or a certificate of participation (for non-physician attendees).

- Southern Medical Association (SMA) will create an online account for you including your unique login, using the email address you provided during registration (your username/ID is your email address).
- Upon receipt of your post-meeting email, click the link provided, and please **make sure that your name and email address appear at the top of the form before completion**.
- <u>After</u> you complete and submit your evaluation and attendance documentation, your certificate will be emailed to you as a .pdf attachment from <u>customerservice@sma.org</u> within 24 hours.





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