

The Progression of Hypertension and CVD Risk Factors in Africa

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Feedback

BLOOD PRESSURE IN THE AFRICAN NATIVE.: ITS BEARING UPON THE ÆTIOLOGY OF HYPERPIESIA AND ARTERIO-SCLEROSIS

C.P. Donnison, M.B., B.S. LOND. (LATE MEDICAL OFFICER, EAST AFRICAN MEDICAL SERVICE.)

BLOOD PRESSURE IN THE AFRICAN NATIVE.

ITS BEARING UPON THE ÆTIOLOGY OF HYPERPIESIA
AND ARTERIO-SCLEROSIS.

BY C. P. DONNISON, M.B., B.S. LOND.,
LATE MEDICAL OFFICER, EAST AFRICAN MEDICAL SERVICE.

The investigations recorded here were all carried out on natives living in one of the largest of the native reserves in Kenya Colony—namely, South Kavirondo, situated on the shores of Lake Victoria Nyanza. These natives live under primitive conditions, that is, under conditions which have probably undergone no appreciable change for many centuries. They have

**BLOOD PRESSURE IN THE AFRICAN
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A series of 1000 examinations was made in apparently healthy male natives of all ages, ranging from 15 years to about 70 or 80 years.

Age group.	Cases examined.	Aver. pulse-rate.	Blood pressure.																							
			Aver. syst.	Aver. diast.	Aver. diff.																					
15-19	99	68.71	123.07	81.89	41.18																					
20-24	100	63.22	122.76	Collected Year 2005 <table border="1"> <thead> <tr> <th>N</th> <th>Heart rate</th> <th>SBP (mmHg)</th> </tr> </thead> <tbody> <tr> <td>120</td> <td>69.8</td> <td>126.8</td> </tr> <tr> <td>129</td> <td>69.3</td> <td>129.5</td> </tr> <tr> <td>138</td> <td>72.5</td> <td>132.3</td> </tr> <tr> <td>108</td> <td>70.0</td> <td>136.4</td> </tr> <tr> <td>100</td> <td>70.7</td> <td>141.8</td> </tr> <tr> <td>143</td> <td>68.2</td> <td>146.5</td> </tr> </tbody> </table>		N	Heart rate	SBP (mmHg)	120	69.8	126.8	129	69.3	129.5	138	72.5	132.3	108	70.0	136.4	100	70.7	141.8	143	68.2	146.5
N	Heart rate	SBP (mmHg)																								
120	69.8	126.8																								
129	69.3	129.5																								
138	72.5	132.3																								
108	70.0	136.4																								
100	70.7	141.8																								
143	68.2	146.5																								
25-29	100	63.57	126.37																							
30-34	115	64.55	126.05																							
35-39	100	69.46	125.55																							
40-44	93	68.52	118.32																							
45-49	96	69.17	113.19																							
50-54	100	72.20	109.79																							
55-59	100	70.73	106.59																							
60 and over	97	75.23	105.76																							



National, regional, and global trends in systolic blood pressure since 1980: systematic analysis of health examination surveys and epidemiological studies with 786 country-years and 5·4 million participants

Goodarz Danaei*, Mariel M Finucane*, John K Lin*, Gitanjali M Singh*, Christopher J Paciorek, Melanie J Cowan, Farshad Farzadfar, Gretchen A Stevens, Stephen S Lim, Leanne M Riley, Majid Ezzati, on behalf of the Global Burden of Metabolic Risk Factors of Chronic Diseases Collaborating Group (Blood Pressure)†

Summary

Lancet 2011; 377: 568–77

Published Online

February 4, 2011

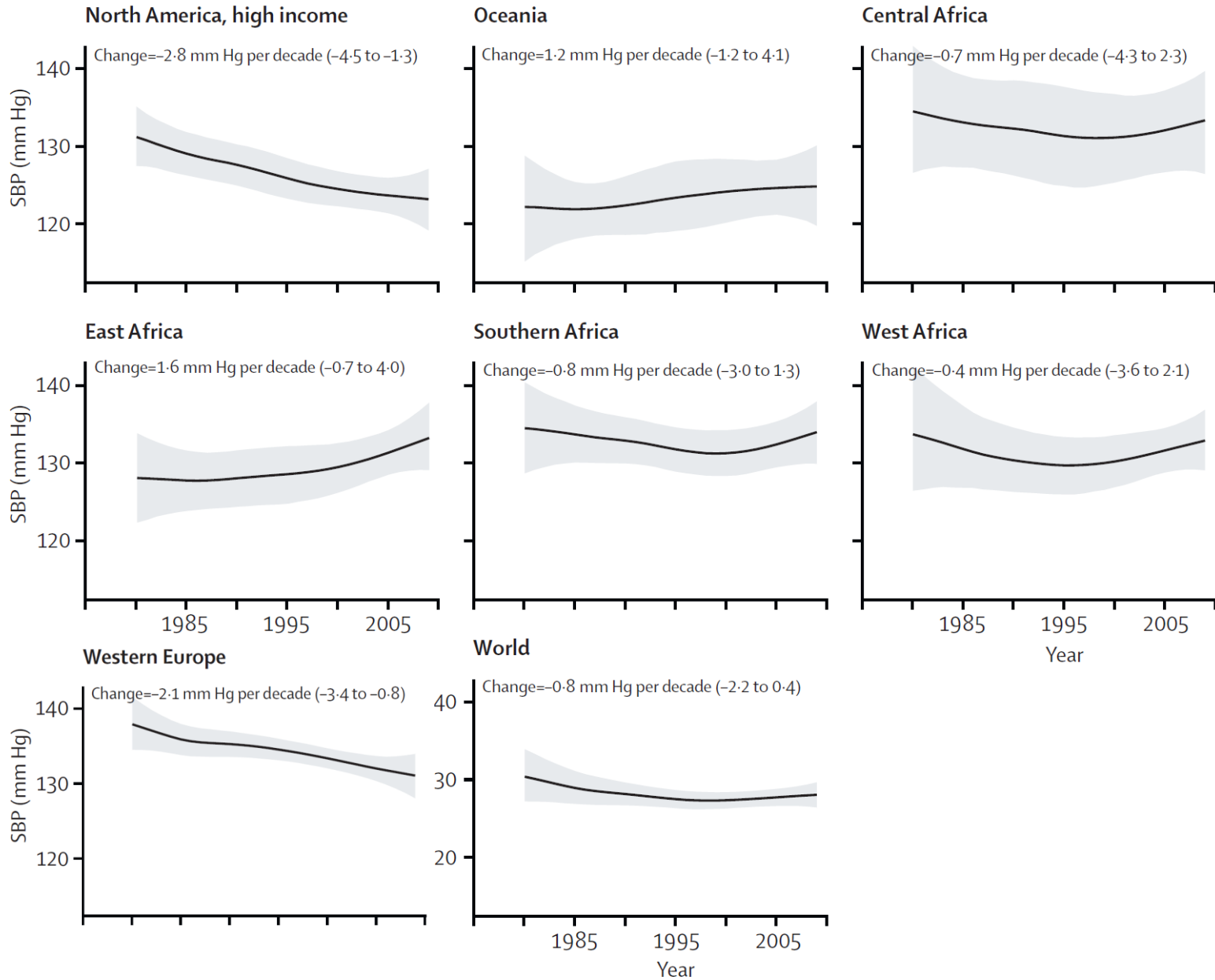
DOI:10.1016/S0140-

6736(10)62036-3

Background Data for trends in blood pressure are needed to understand the effects of its dietary, lifestyle, and pharmacological determinants; set intervention priorities; and evaluate national programmes. However, few worldwide analyses of trends in blood pressure have been done. We estimated worldwide trends in population mean systolic blood pressure (SBP).

Interpretation On average, global population SBP decreased slightly since 1980, but trends varied significantly across regions and countries. SBP is currently highest in low-income and middle-income countries. Effective population-based and personal interventions should be targeted towards low-income and middle-income countries.

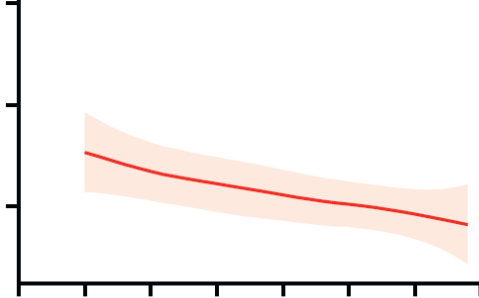
A Men



B Women

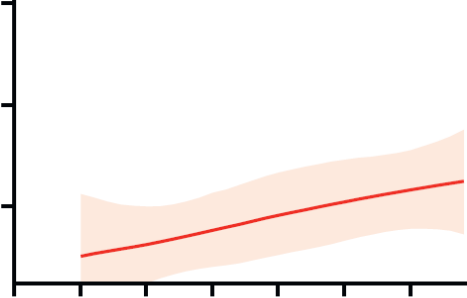
North America, high income

Change=-2.3 mm Hg per decade (-4.0 to -0.5)



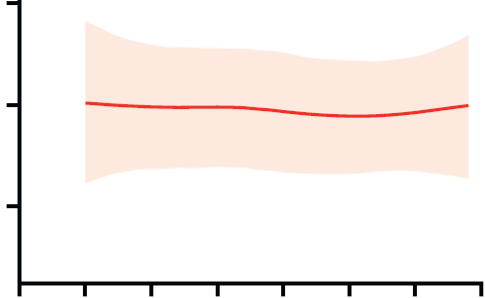
Oceania

Change=2.7 mm Hg per decade (0.0 to 5.5)



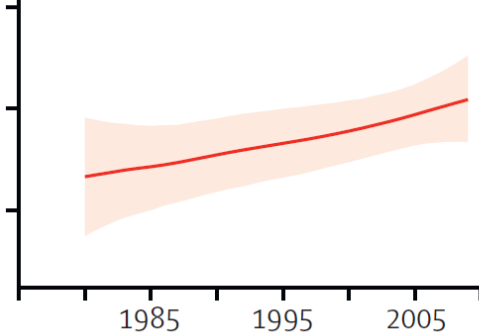
Central Africa

Change=-0.3 mm Hg per decade (-4.0 to 3.0)



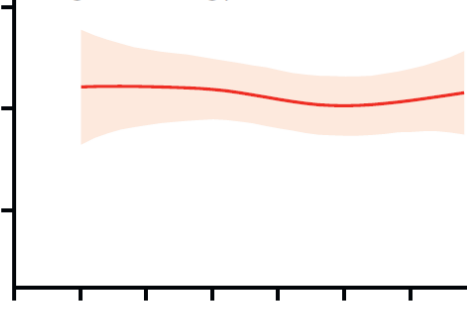
East Africa

Change=2.5 mm Hg per decade (0.2 to 5.0)



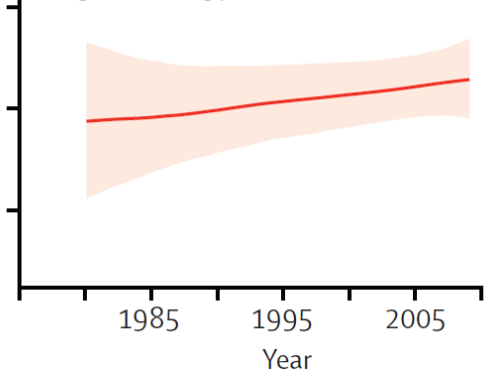
Southern Africa

Change=-0.7 mm Hg per decade (-3.0 to 1.7)



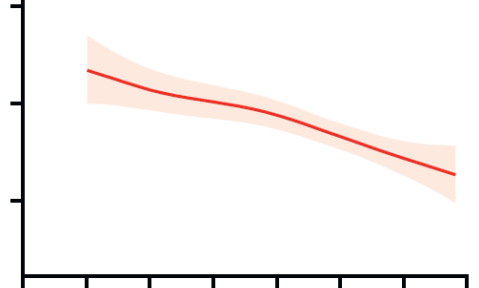
West Africa

Change=1.5 mm Hg per decade (-1.5 to 4.4)



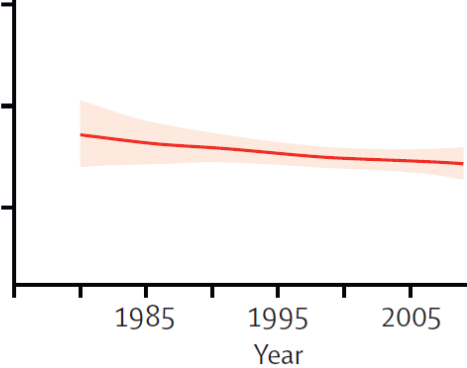
Western Europe

Change=-3.5 mm Hg per decade (-5.0 to -2.2)



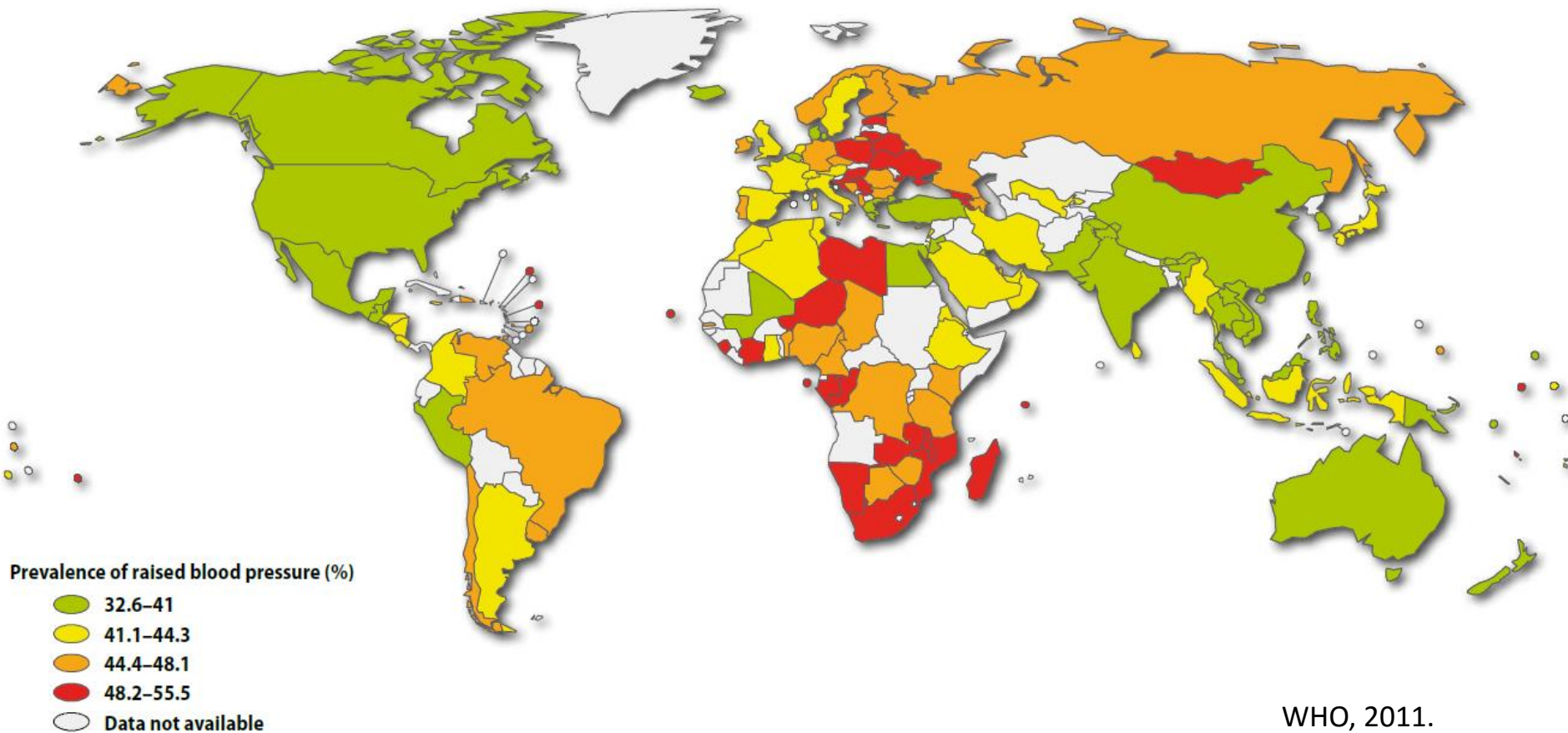
World

Change=-1.0 mm Hg per decade (-2.3 to 0.3)



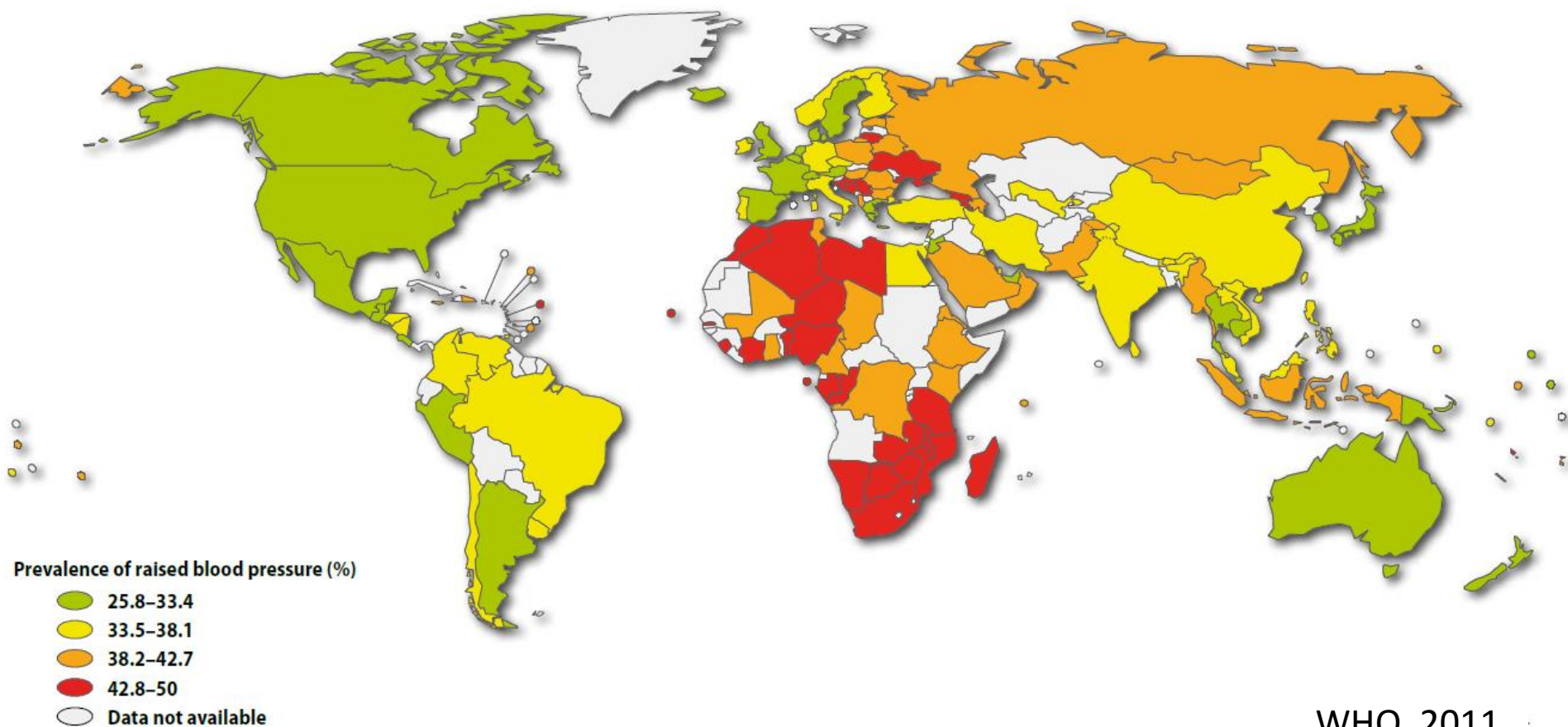
Background

Prevalence of hypertension in MEN (aged >25 yrs)



Background

Prevalence of hypertension in WOMEN (aged >25 yrs)



WHO, 2011.

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CARDIOVASCULAR DISEASE

Hypertension among older adults in low- and middle-income countries: prevalence, awareness and control

Peter Lloyd-Sherlock,^{1*} John Beard,² Nadia Minicuci,³ Shah Ebrahim⁴ and Somnath Chatterji⁵

¹School of International Development, University of East Anglia, Norwich, UK, ²Department of Ageing and Life Course, World Health Organization, Geneva, Switzerland ³National Research Council, Institute of Neuroscience, Padova, Italy, ⁴London School of Hygiene and Tropical Medicine, London, UK and ⁵Department of Health Statistics and Informatics, World Health Organization, Geneva, Switzerland

Background This study uses data from the World Health Organization's Study on Global Ageing and Adult Health (SAGE) to examine patterns of hypertension prevalence, awareness, treatment and control for people aged 50 years and over in China, Ghana, India, Mexico, the Russian Federation and South Africa.

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¹School of International Development, University of East Anglia, Norwich, UK, ²Department of Ageing and Life Course, World Health Organization, Geneva, Switzerland ³National Research Council, Institute of Neuroscience, Padova, Italy, ⁴London School of Hygiene and Tropical Medicine, London, UK and ⁵Department of Health Statistics and Informatics, World Health Organization, Geneva, Switzerland

We found high prevalences of hypertension in all countries, with the highest in Africa. Around half of older people in high-income countries. ^{21–23} Indeed, South Africa's prevalence is the highest ever reported by a nationally representative survey of people aged 50 and over for any country. It is substantially higher than recently published estimates for South Africa and 11 other sub-Saharan African countries.²⁴

Non-Communicable Diseases in Sub-Saharan Africa: The Case for Cohort Studies

Michelle D. Holmes^{1,2*}, Shona Dalal², Jimmy Volmink³, Clement A. Adebamowo⁴, Marina Njelekela⁵, Wafaie W. Fawzi⁶, Walter C. Willett⁶, Hans-Olov Adami^{2,7}

1 Channing Laboratory, Department of Medicine, Brigham Young University, Harvard Medical School, Boston, Massachusetts, United States of America, **2** Department of Epidemiology, University of Stellenbosch, Stellenbosch, South Africa, **3** Department of Epidemiology, University of Cape Town, South Africa, **4** Department of Epidemiology, University of Michigan, Ann Arbor, Michigan, United States of America, **5** Department of Health, Behavior, and Society, Johns Hopkins University, Baltimore, Maryland, United States of America, **6** Department of Nutrition, Harvard School of Public Health, Boston, Massachusetts, United States of America, **7** Department of Epidemiology, Harvard School of Public Health, Boston, Massachusetts, United States of America

Conclusions The prevalence of NCDs and their risk factors is high in some SSA settings. With the lack of vital statistics systems, epidemiologic studies with a variety of designs (cross-sectional, longitudinal and interventional) capable of in-depth analyses of risk factors could provide a better understanding of NCDs in SSA, and inform health-care policy to mitigate the oncoming NCD epidemic.

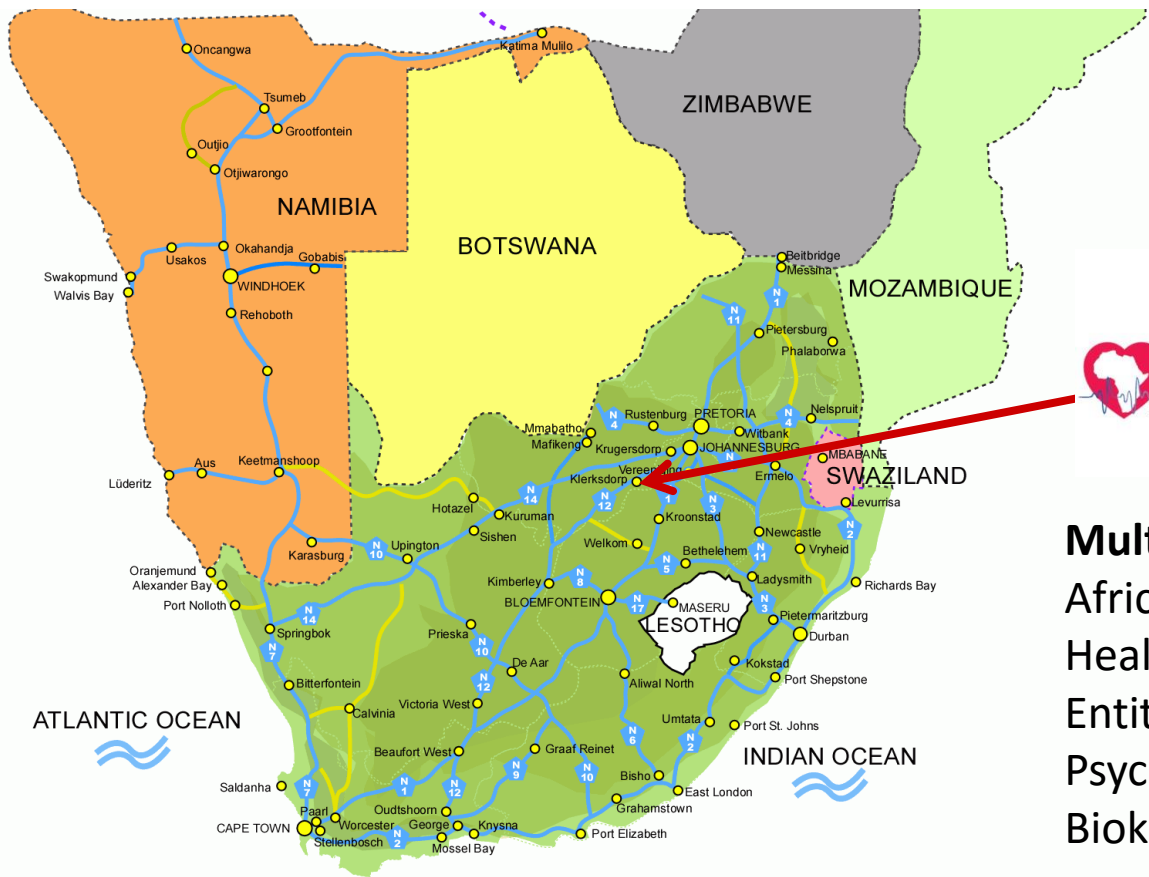
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2011;40:885–901
doi:10.1371/journal.pmed.10093/ije/dyr050

NON-COMMUNICABLE DISEASES

Non-communicable diseases in sub-Saharan Africa: what we know now

Shona Dalal,^{1*} Juan Jose Beunza,^{1,2} Jimmy Volmink,³ Clement Adebamowo,^{4,5} Francis Bajunirwe,⁶ Marina Njelekela,⁷ Dariush Mozaffarian,^{1,8} Wafaie Fawzi,^{1,9} Walter Willett,⁹ Hans-Olov Adami^{10,11} and Michelle D Holmes^{1,12}



Multidisciplinary team:
 Africa Unit for Transdisciplinary
 Health Research
 Entities: Physiology, Nutrition,
 Psychology, Pharmacy, Nursing,
 Biokinetics, Biochemistry



The Prospective Urban Rural Epidemiology (PURE) study: Examining the impact of societal influences on chronic noncommunicable diseases in low-, middle-, and high-income countries

Trial Design

Koon Teo, PhD,^a Clara K. Chow, PhD,^a Mario Vaz, MD,^b Sumathy Rangarajan, MSc,^a and Salim Yusuf, DPhil^a, The PURE Investigators-Writing group *Hamilton, Ontario, Canada; and Bangalore, India*

American Heart Journal
July 2009



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Rural



Urban



The PURE study (NWP South Africa): longitudinal data, 2005, 2010, 2015

Multidisciplinary team taking a wide range of measurements in 1000 rural and 1000 urban black South Africans (aged >30 yrs)

This data provided an excellent opportunity to

- (a) Explore and compare risk factors for HT development
- (b) Determine which conventional CV risk factors are associated with a 5-yr change in BP



Are behavioural risk factors to be blamed for the conversion from optimal blood pressure to hypertensive status in Black South Africans? A 5-year prospective study

Aletta E Schutte,^{1*} Rudolph Schutte,¹ Hugo W Huisman,¹ Johannes M van Rooyen,¹ Carla MT Fourie,¹ Nico T Malan,¹ Leoné Malan,¹ Catharina MC Mels,¹ Wayne Smith,¹ Sarah J Moss,² G Wayne Towers,³ H Salomé Kruger,³ Edelweiss Wentzel-Viljoen,³ Hester H Vorster³ and Annamarie Kruger⁴

International Journal of Epidemiology 2012;**41**:1114–1123

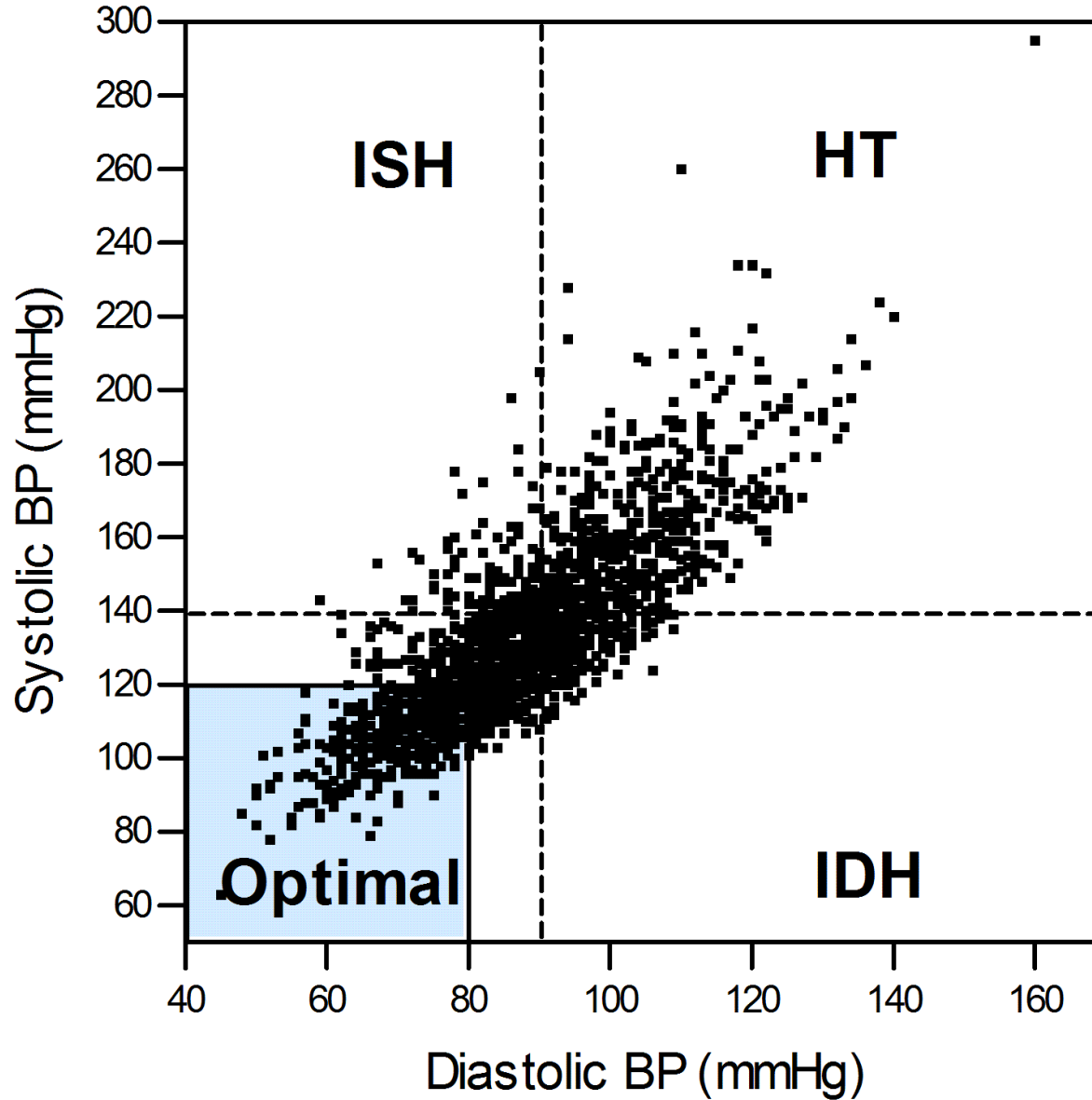
Study population with BP data:

PURE baseline
N=1994

2005

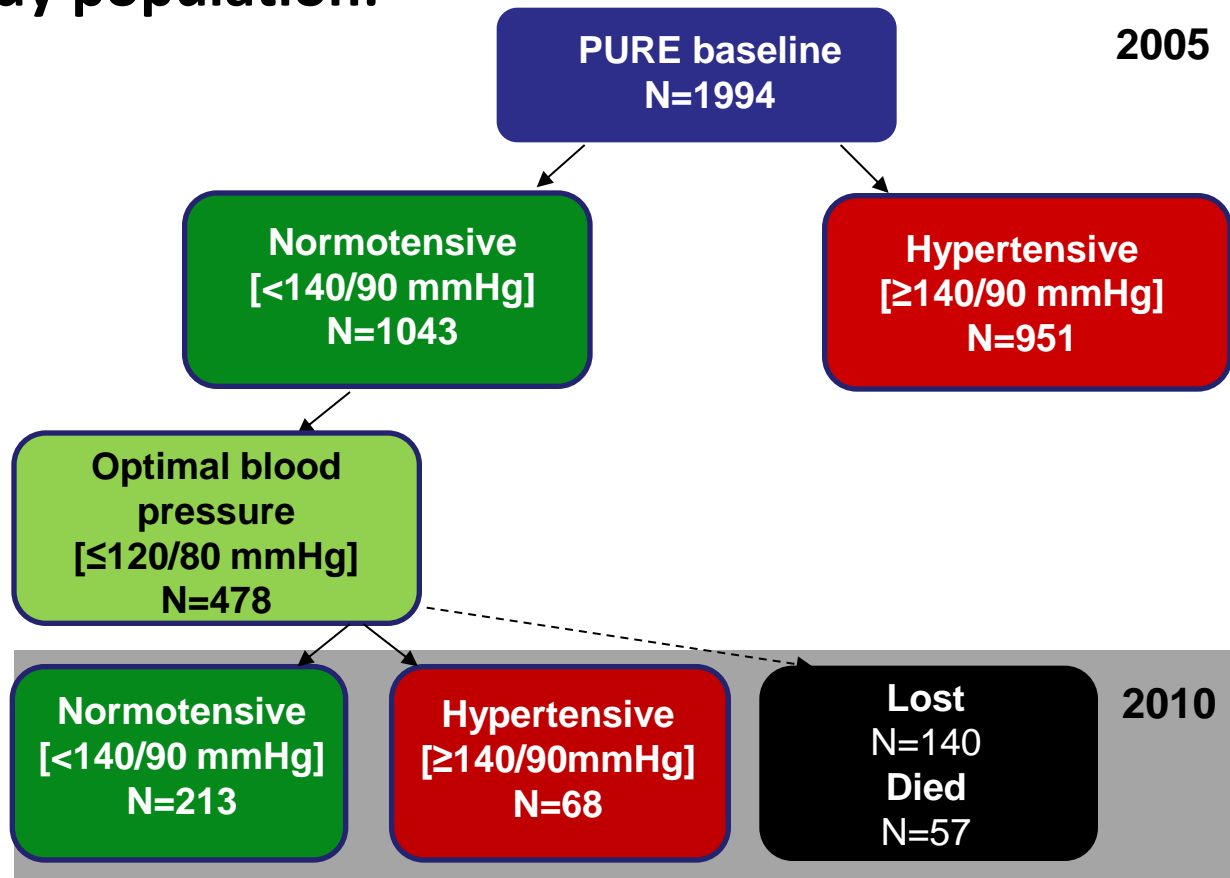
Blood pressure distribution at baseline:

2005 (N=1994)

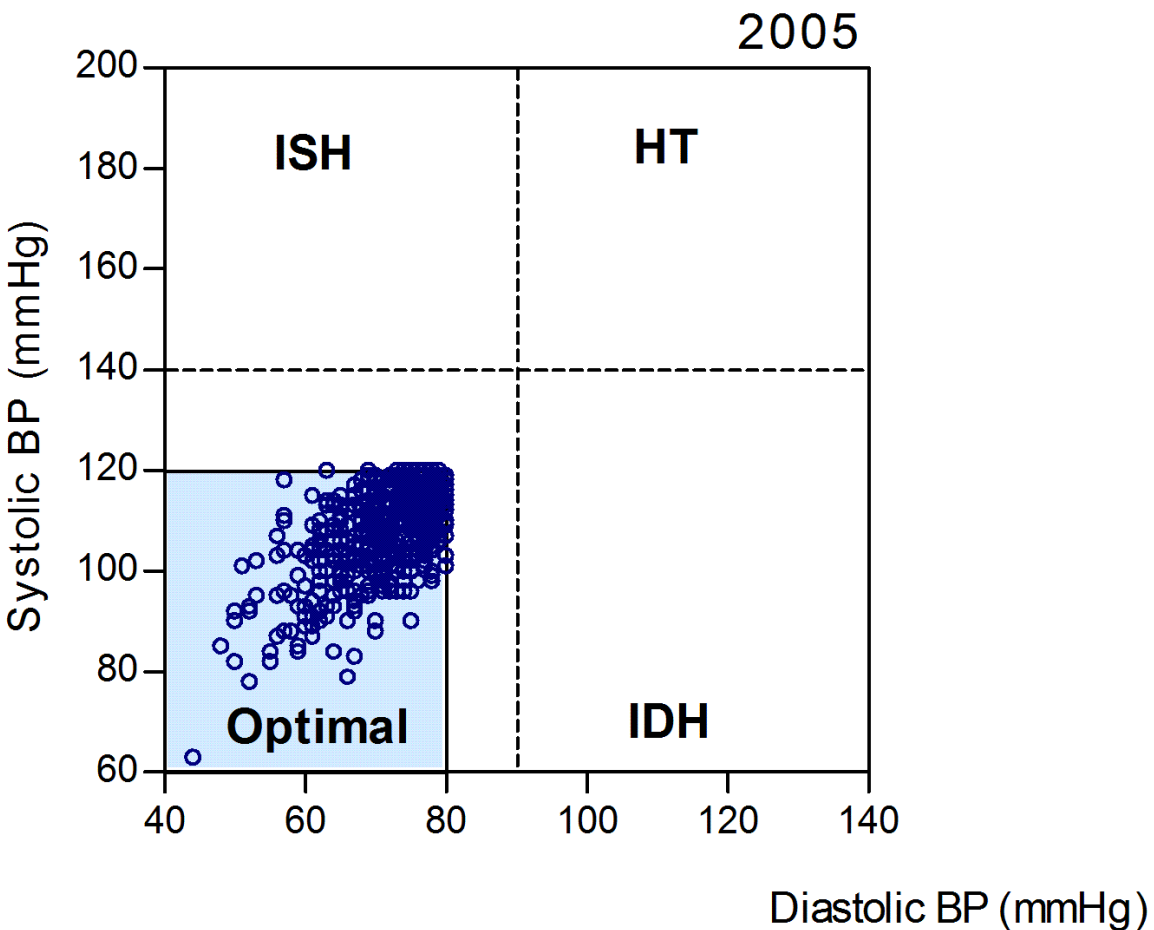


Methods:

Study population:



Participants with Optimal BP at baseline followed for 5 years



Independent relationships with change in SBP:

	Δ SBP ($R^2 = 0.23$)
Age	
Gender (m/f)	
Rural/urban	
WC	$\beta=0.18$; $p=0.006$
HIV	$\beta= -0.23$; $p<0.001$
γ GT	$\beta= 0.13$; $p=0.029$
$\Delta \gamma$ GT	$\beta= 0.17$; $p=0.005$
HbA1c	

Independent relationships Δ SBP or cSBP as dep var with γ GT substituted with self-reported alcohol intake:

	Δ SBP ($R^2 = 0.23$)	cSBP ($R^2 = 0.14$)
Age		
SBP	$\beta = -0.36; p < 0.001$	$\beta = 0.16; p = 0.011$
Rural/urban		$\beta = -0.23; p = 0.008$
WC	$\beta = 0.18; p = 0.005$	
HIV	$\beta = -0.23; p < 0.001$	$\beta = -0.22; p < 0.001$
Alcohol no/yes	$\beta = 0.13; p = 0.029$	$\beta = 0.19; p = 0.004$
HbA1c		$\beta = 0.14; p = 0.025$

SABPA study

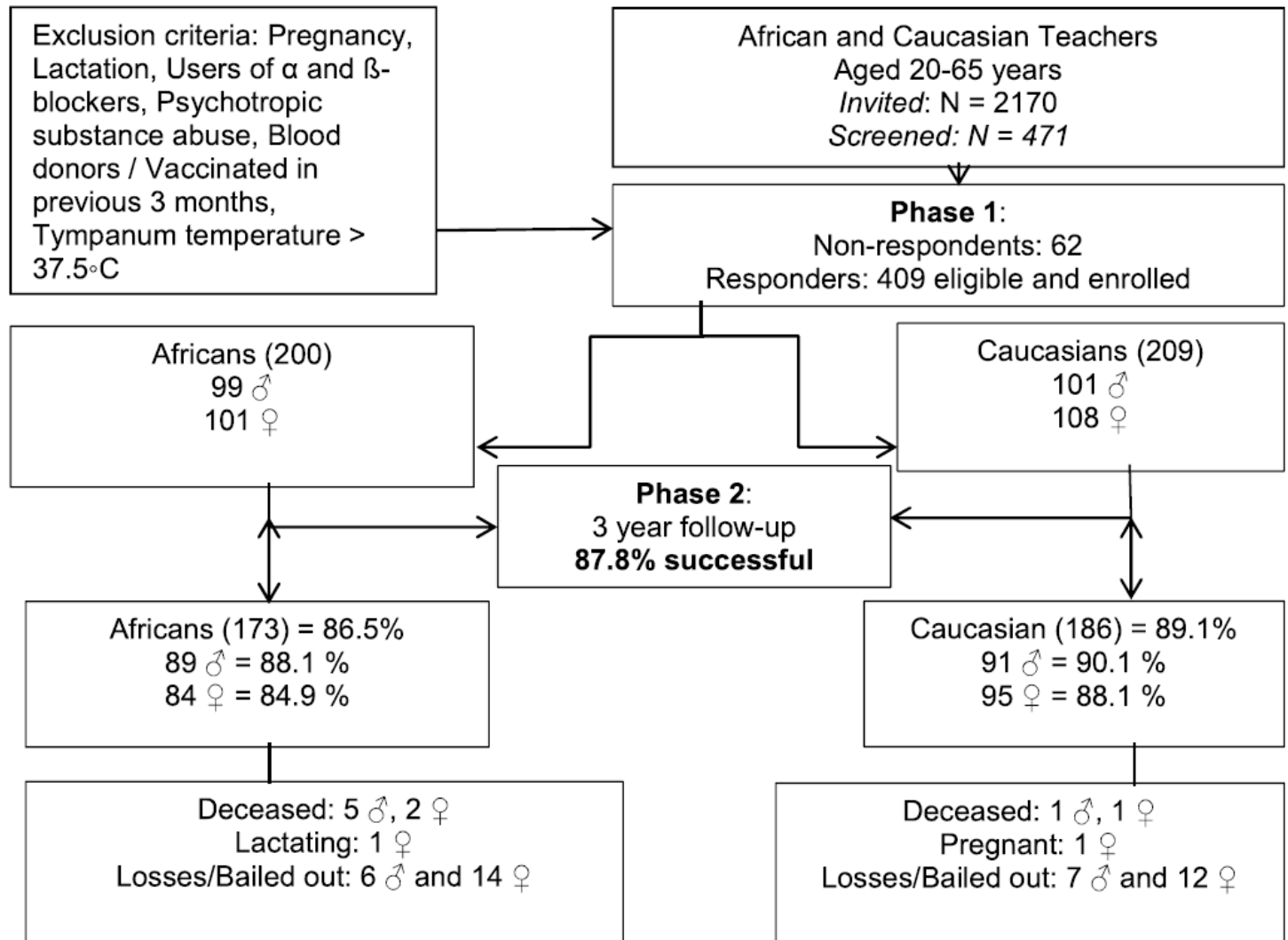


Figure 2. The Sympathetic activity and Ambulatory Blood pressure in Africans (SABPA) prospective cohort study population.

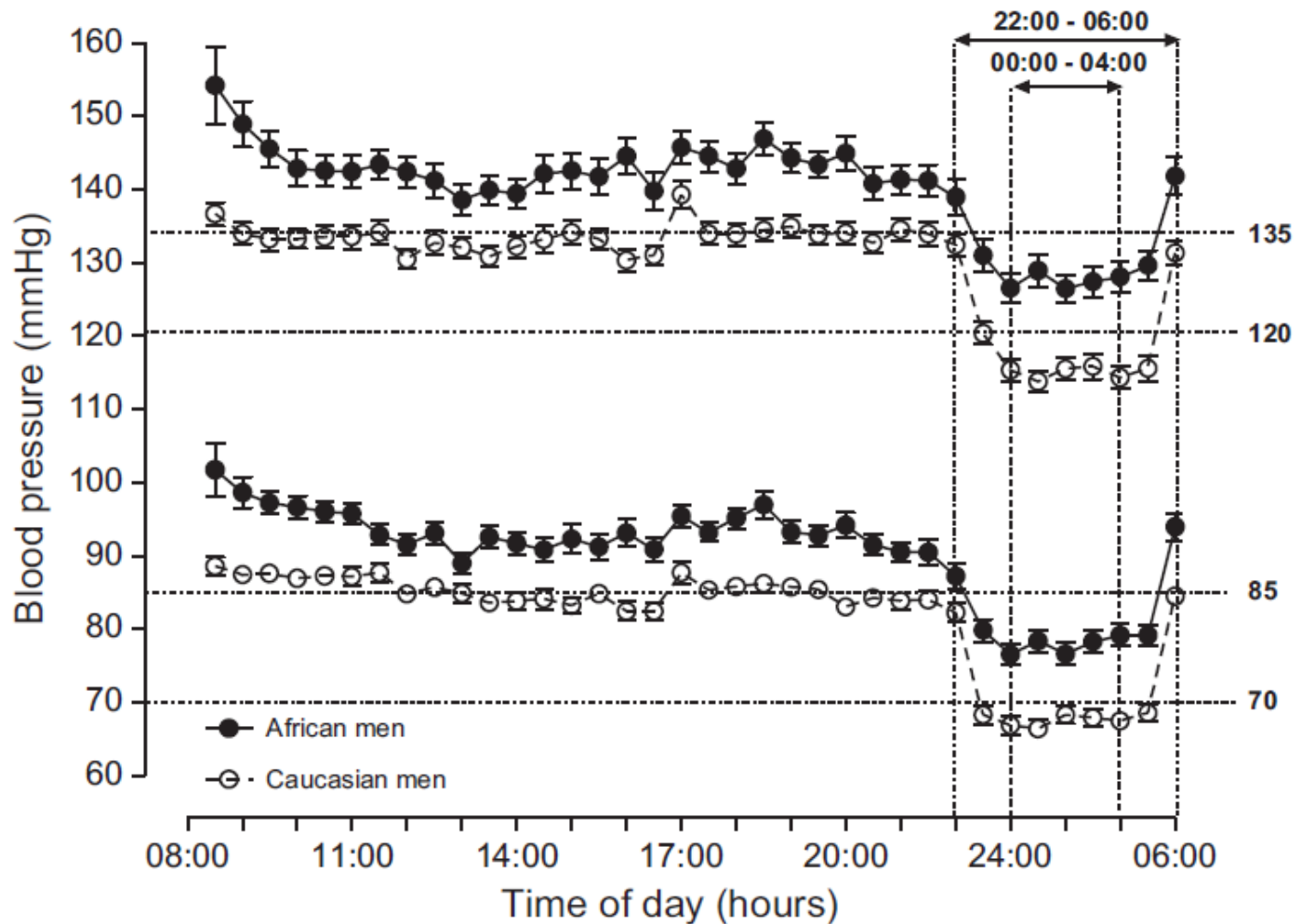


Fig. 1 – Systolic and diastolic ambulatory blood pressure of the African and Caucasian men. Bars indicate standard error.

Sympathetic activity and Ambulatory BP study (SABPA)

Atherosclerosis 238 (2015) 52–54



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journal homepage: www.elsevier.com/locate/atherosclerosis



Progression of cardiovascular risk factors in black Africans: 3 year follow up of the SABPA cohort study



Mark Hamer^{a, b, *}, Roland von Känel^c, Manja Reimann^d, Nico T. Malan^b, Alta E. Schutte^b, Hugo W. Huisman^b, Leone Malan^b

^a Department of Epidemiology and Public Health, University College London, London, UK

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^c Department of Neurology, Inselspital, Bern University Hospital and University of Bern, Switzerland

^d Autonomic and Neuroendocrinological Laboratory Dresden, Department of Neurology, University Hospital Carl Gustav Carus, Dresden University of Technology, Dresden, Germany

Table 1

Comparison between white (reference group) and black participants of CVD risk factor progression over a three-year follow-up period.

Risk factor	Adjusted coefficient (95% CI)	p-value
24 h systolic blood pressure	6.02 (3.51, 8.55)	<0.001
24 h diastolic blood pressure	3.57 (1.96, 5.17)	<0.001
CIMT	-0.048 (-0.033, -0.064)	<0.001
Cross sectional wall area	-1.27 (-0.73, -1.81)	<0.001
Endothelin-1	0.21 (-0.99, 1.41)	0.73
HDL cholesterol	-0.02 (-0.08, 0.04)	0.50
Total cholesterol	0.65 (0.45, 0.86)	<0.001
Total: HDL cholesterol ratio	0.86 (0.64, 1.08)	<0.001
Triglycerides	0.26 (-0.11, 0.17)	0.72
Glycated haemoglobin	0.13 (-0.05, 0.30)	0.16
Glucose	1.22 (0.91, 1.52)	<0.001
Insulin	0.35 (-0.94, 1.65)	0.59
Fibrinogen	0.55 (0.41, 0.69)	<0.001
D-dimer	132.15 (31.36, 232.94)	0.01
Interleukin-6	-1.48 (-4.37, 1.42)	0.32
Tumour necrosis factor- α	-0.57 (-1.00, -0.14)	0.01
BMI	-0.24 (-0.73, 0.25)	0.33
Waist	1.92 (0.24, 3.59)	0.03

Coefficients reflect differences between white (ref) and black Africans, adjusted for age, sex, serum cotinine, physical activity energy expenditure, history of CVD, medication, and baseline levels of the respective risk factor. Where: CIMT, carotid intima media thickness; HDL, high density lipoprotein; BMI, body mass index.

Behavioural and other associated CVD risk factors

- **Obesity & diet (sugar intake)**
- Inflammation
- Alcohol
- Endothelial function/NO bioavailability
- HIV infection

Added sugar intake in South Africa: findings from the Adult Prospective Urban and Rural Epidemiology cohort study¹⁻⁴

Hester H Vorster, Annamarie Kruger, Edelweiss Wentzel-Viljoen, H Salome Kruger, and Barrie M Margetts

Results: Added sugar intake, particularly in rural areas, has increased rapidly in the past 5 y. In rural areas, the proportion of adults who consumed sucrose-sweetened beverages approximately doubled (for men, from 25% to 56%; for women, from 33% to 63%) in the past 5 y. After adjustment, subjects who consumed more added sugars ($\geq 10\%$ energy from added sugars) compared with those who consumed less added sugars had a higher waist circumference [mean difference (95% CI): 1.07 cm (0.35, 1.79 cm)] and body mass index (in kg/m^2) [0.43 (0.12, 0.74)] and lower HDL cholesterol [-0.08 mmol/L (-0.14 , 0.002 mmol/L)].

Conclusions: This cohort showed dramatic increases in added sugars and sucrose-sweetened beverage consumption in both urban and rural areas. Increased consumption was associated with increased NCD risk factors. In addition, the study showed that the nutrition transition has reached a remote rural area in South Africa. Urgent action is needed to address these trends. *Am J Clin Nutr* doi: 10.3945/ajcn.113.069005.

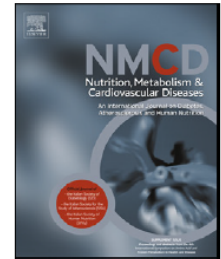


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Nutrition, Metabolism & Cardiovascular Diseases

journal homepage: www.elsevier.com/locate/nmcd



Evaluation of waist-to-height ratio to predict 5 year cardiometabolic risk in sub-Saharan African adults



L.J. Ware ^a, K.L. Rennie ^b, H.S. Kruger ^c, I.M. Kruger ^d, M. Greeff ^d, C.M.T. Fourie ^a,
H.W. Huisman ^a, J.D.W. Scheepers ^a, A.S. Uys ^a, R. Kruger ^a, J.M. Van Rooyen ^a,
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Behavioural and other associated CVD risk factors

- Obesity & diet (sugar intake)
- **Inflammation**
- Alcohol
- Endothelial function/NO bioavailability
- HIV infection

ORIGINAL ARTICLE

Inflammation, obesity and cardiovascular function in African and Caucasian women from South Africa: the POWIRS study

AE Schutte, D van Vuuren, JM van Rooyen, HW Huisman, R Schutte, L Malan
and NT Malan

School for Physiology, Nutrition and Consumer Sciences, North-West University, Potchefstroom, South Africa

□ Inflammation (CRP, IL-6, suPAR) as prognostic markers for mortality

International Journal of Cardiology 184 (2015) 631–636

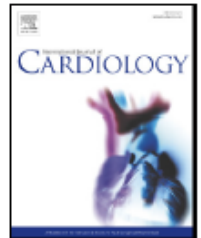


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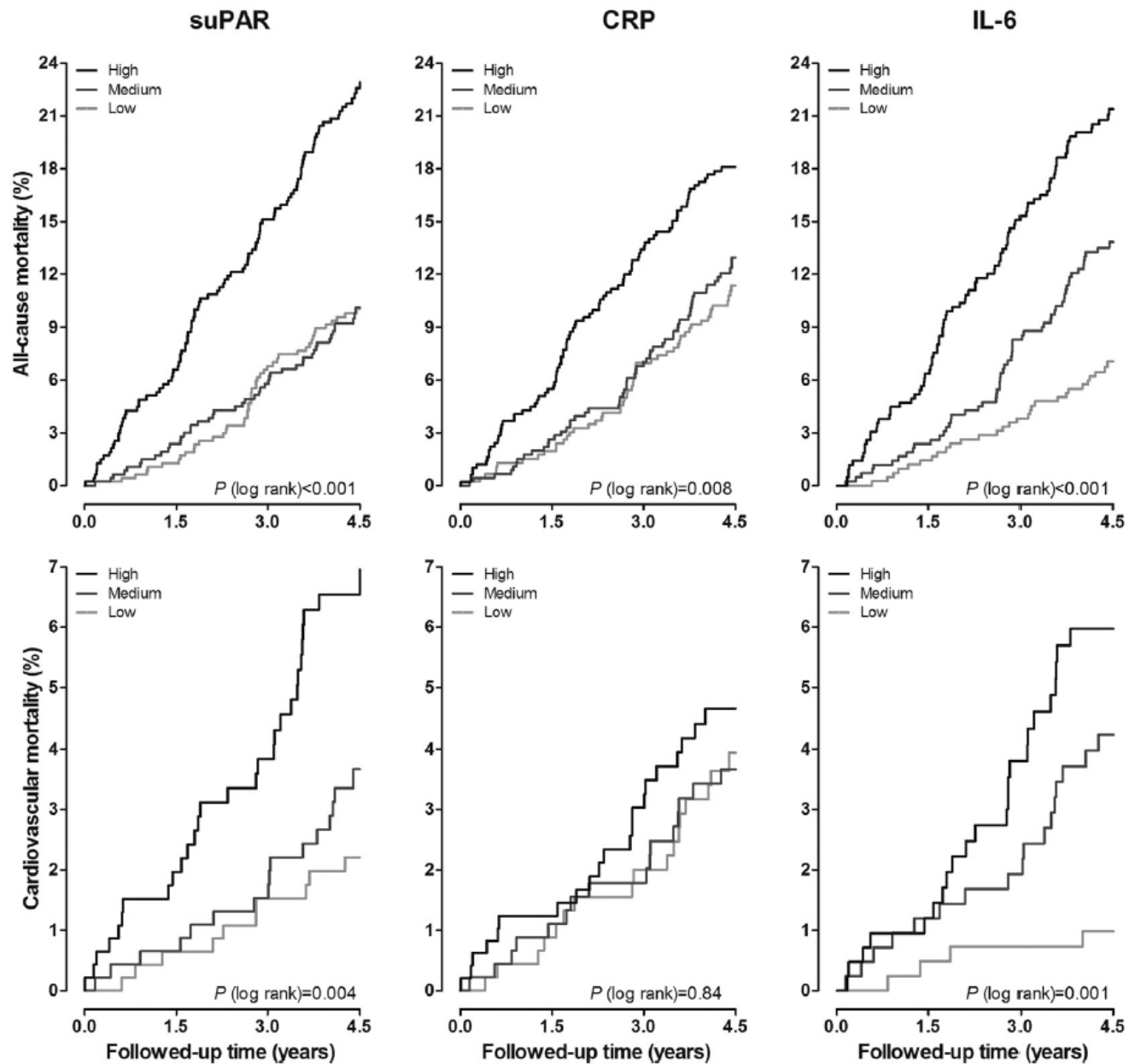
journal homepage: www.elsevier.com/locate/ijcard



Soluble urokinase plasminogen activator receptor as a prognostic marker of all-cause and cardiovascular mortality in a black population[☆]



Shani Botha^{a,*}, Carla M.T. Fourie^a, Rudolph Schutte^{a,b}, Jesper Eugen-Olsen^c, Ronel Pretorius^d, Aletta E. Schutte^{a,b}



Behavioural and other associated CVD risk factors

- Obesity & diet (sugar intake)
- Inflammation
- **Alcohol**
- Endothelial function/NO bioavailability
- HIV infection

Original scientific paper

European Journal of
**Preventive
Cardiology**



Alcohol intake, hypertension development and mortality in black South Africans

**Mandlenkosi C Zatu^{1,2,3}, Johannes M Van Rooyen¹,
Annamarie Kruger⁴ and Aletta E Schutte¹**

European Journal of Preventive
Cardiology

0(00) 1–8

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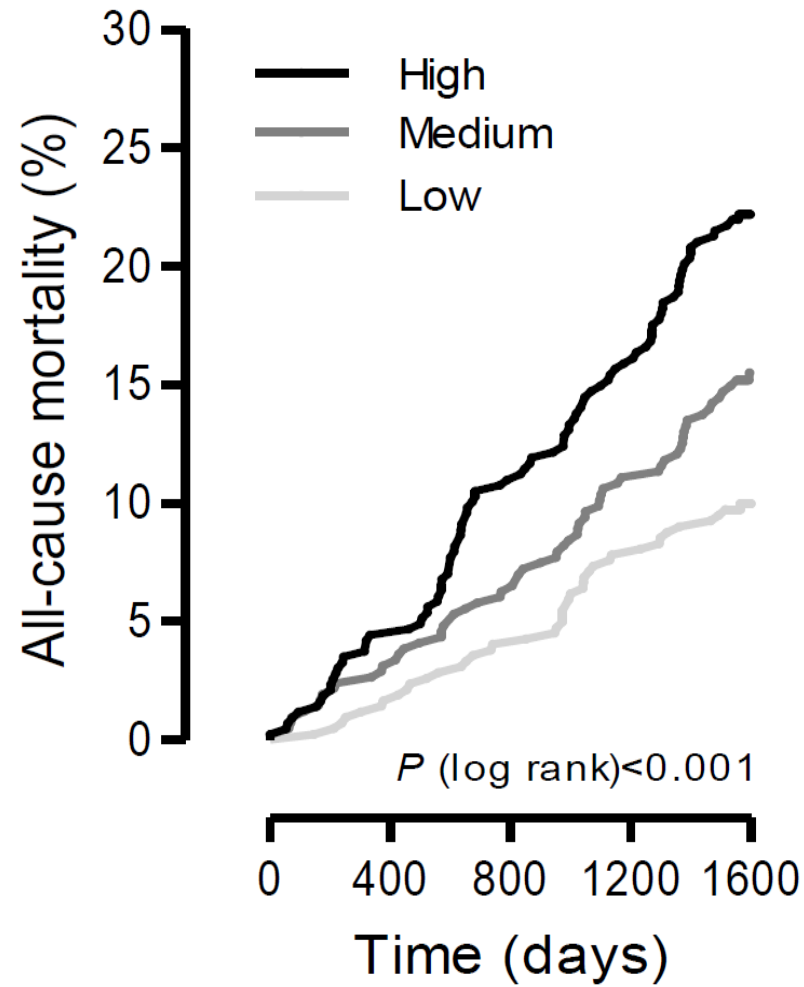
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DOI: 10.1177/2047487314563447

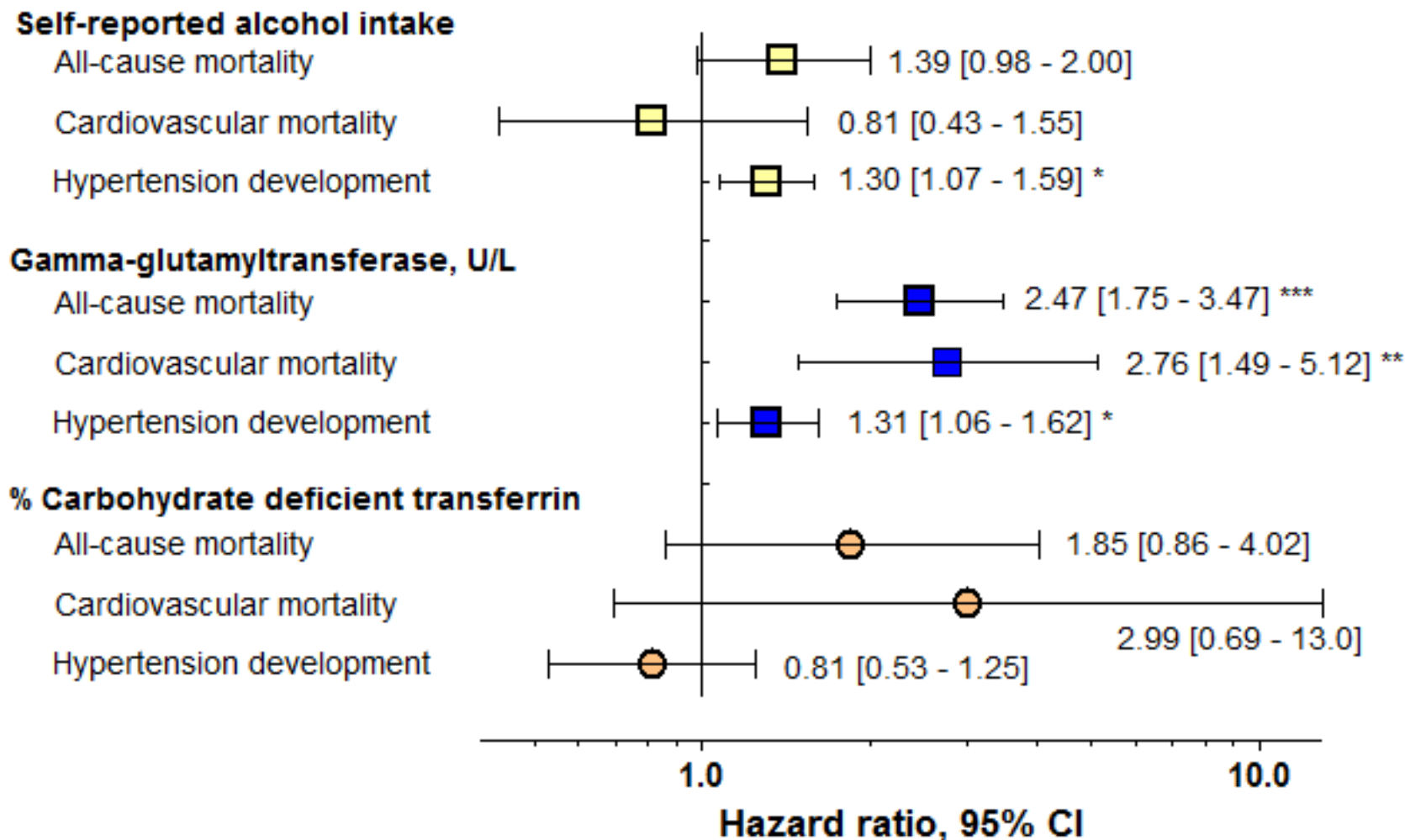
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□ Gamma-glutamyl transferase as predictor of mortality



□ Gamma-glutamyl transferase compared to other alcohol measures as indicator/predictor of mortality?

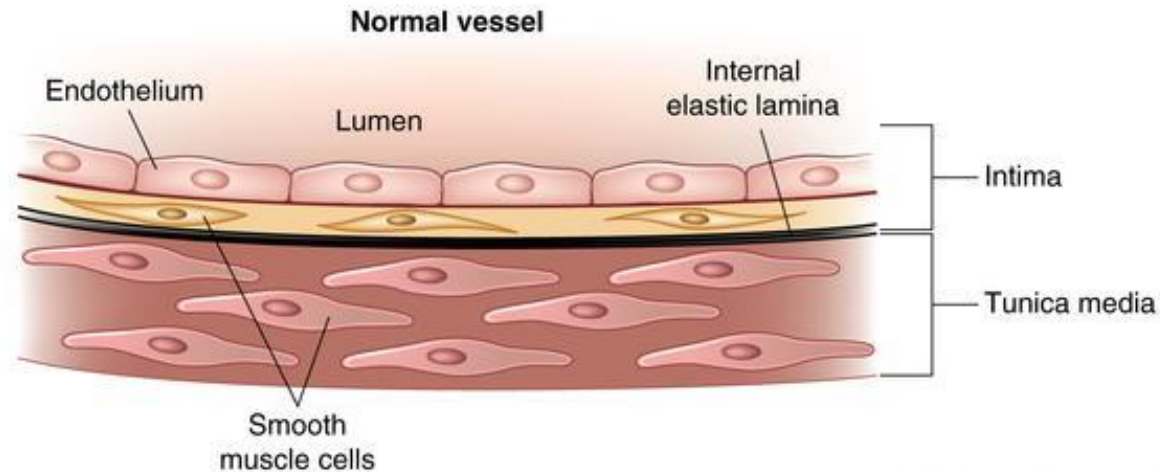


Behavioural and other associated CVD risk factors

- Obesity & diet (sugar intake)
- Inflammation
- Alcohol
- **Endothelial function**
- HIV infection

Endothelial function ...

Microalbuminuria (systemic endothelial dysfunction)



© Current Medicine Group

Urinary Albumin Excretion From Spot Urine Samples Predict All-Cause and Stroke Mortality in Africans

Rudolph Schutte,¹ Roland E. Schmieder,² Hugo W. Huisman,¹ Wayne Smith,¹ Johannes M. van Rooyen,¹ Carla M. T. Fourie,¹ Ruan Kruger,¹ Lisa Uys,¹ Lisa Ware,¹ Catharina M. C. Mels,¹ Minrie Greeff,³ Iolanthé M. Kruger,³ and Aletta E. Schutte¹

□ Microalbuminuria and mortality

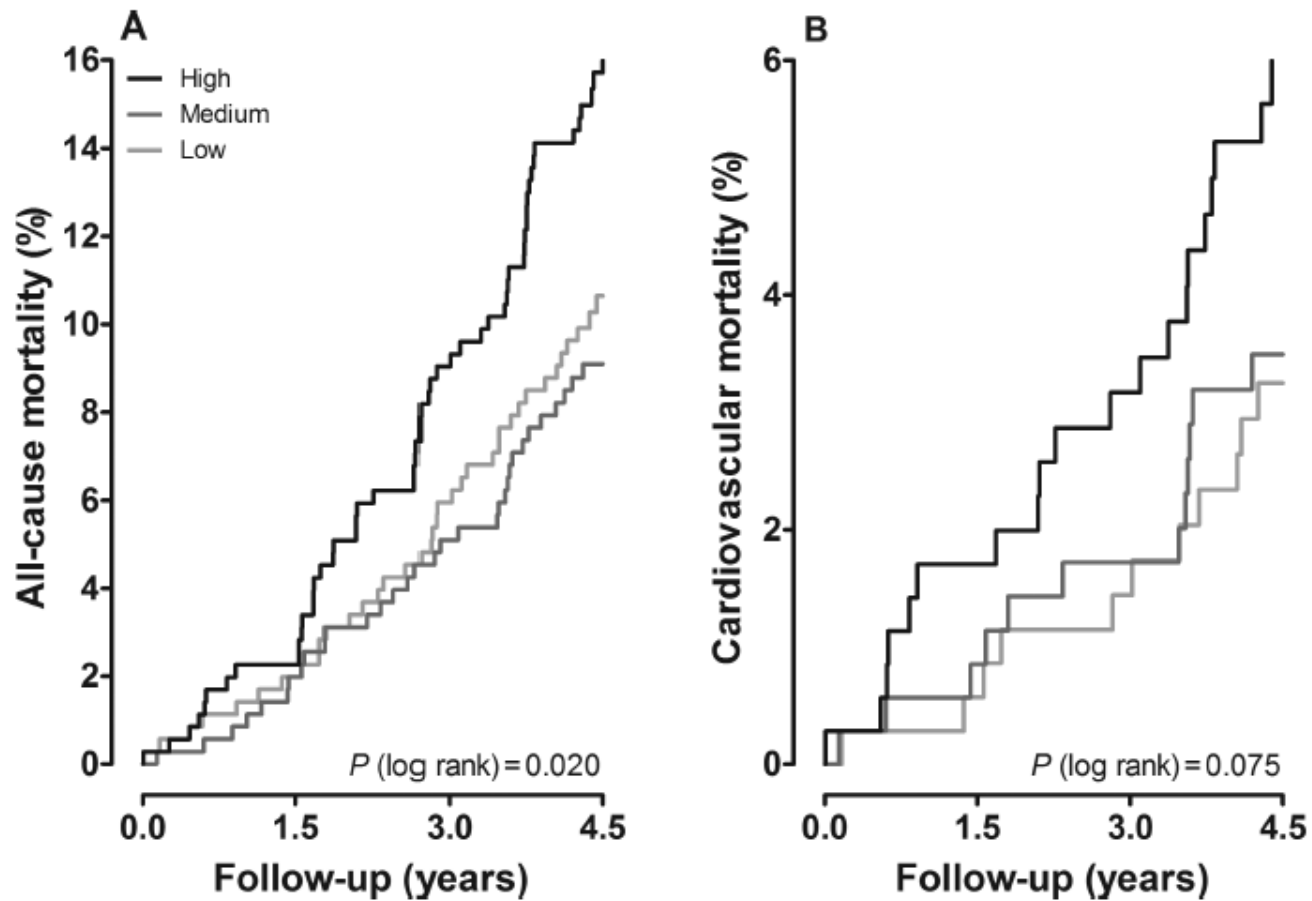
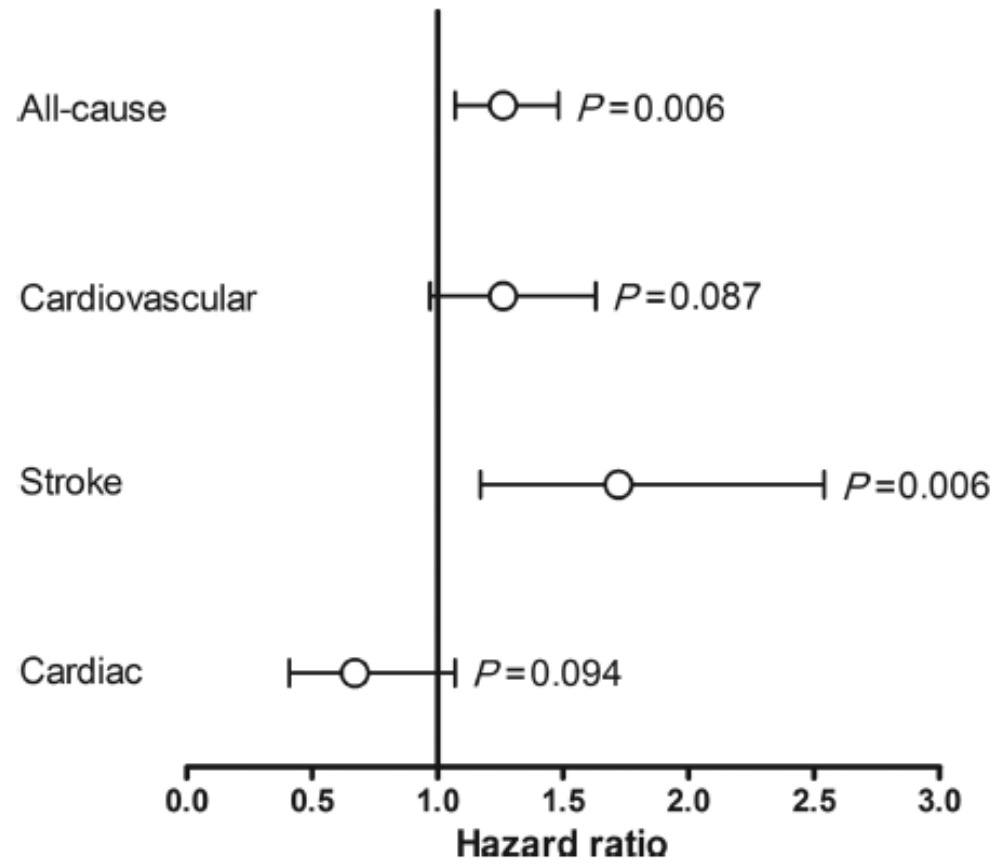


Figure 1. Kaplan-Meier survival function estimates for all-cause (A) and cardiovascular mortality (B) by tertiles of urinary albumin-to-creatinine ratio. *P* values refer to the significance of the log-rank test across three tertiles.

□ Microalbuminuria and mortality



Behavioural and other associated CVD risk factors

- Obesity & diet (sugar intake)
- Inflammation
- Alcohol
- Endothelial function/NO bioavailability
- **HIV infection**

Is HIV-1 infection associated with endothelial dysfunction in a population of African ancestry in South Africa?

C FOURIE, J VAN ROOYEN, M PIETERS, K CONRADIE, T HOEKSTRA, A SCHUTTE

The objective of this study was to assess whether newly identified, never-treated, HIV-1-infected South African participants showed signs of endothelial dysfunction, accelerated atherosclerosis and increased blood coagulation.

We compared 300 newly diagnosed (never antiretroviral-treated) HIV-infected participants to 300 age-, gender-, body mass index- and locality-matched uninfected controls.

□ Crossroads between NCDs and HIV-infection

TABLE 2. ODDS RATIOS OF HIV-INFECTED PARTICIPANTS VS UNINFECTED PARTICIPANTS

	<i>Odds ratios HIV infected vs HIV uninfected</i>	<i>95% CI</i>
HDL-C < 1.36 mmol/l	3.69	2.6–5.2*
TG ≥ 1.0 mmol/l	1.70	1.2–2.3*
TG:HDL ratio ≥ 0.75	3.33	2.4–4.7*
hsCRP ≥ 2.7 mg/l	1.78	1.3–2.5*
hsIL-6 ≥ 4.2 pg/ml	1.67	1.2–2.3*
sICAM-1 ≥ 516 ng/ml	2.04	1.5–2.8*
sVCAM-1 ≥ 693ng/ml	3.92	2.8–5.5*

HDL-C: high-density lipoprotein cholesterol; TG: triglycerides; TG:HDL: triglycerides–high-density lipoprotein ratio; hsCRP: high-sensitivity C-reactive protein; hsIL-6: high-sensitivity interleukin 6; sICAM-1: serum intercellular adhesion molecule-1; sVCAM-1: serum vascular cell adhesion molecule-1. For all variables, the median of total group was used as cut-off value.

*Significant.

□ Crossroads between NCDs and HIV-infection

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Endothelial activation and cardiometabolic profiles of treated and never-treated HIV infected Africans



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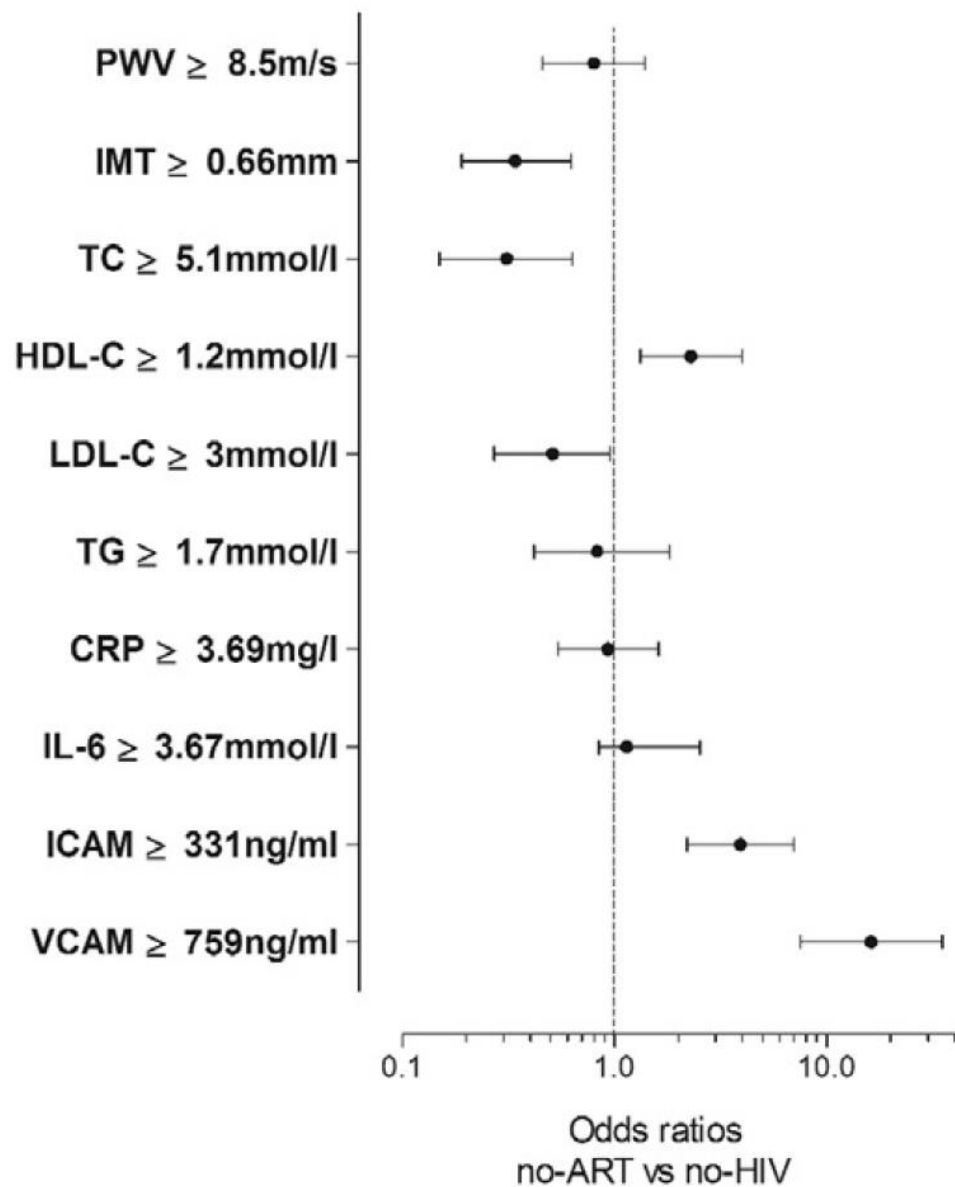


Fig. 2. Odds ratios for increased cardiovascular disease risk

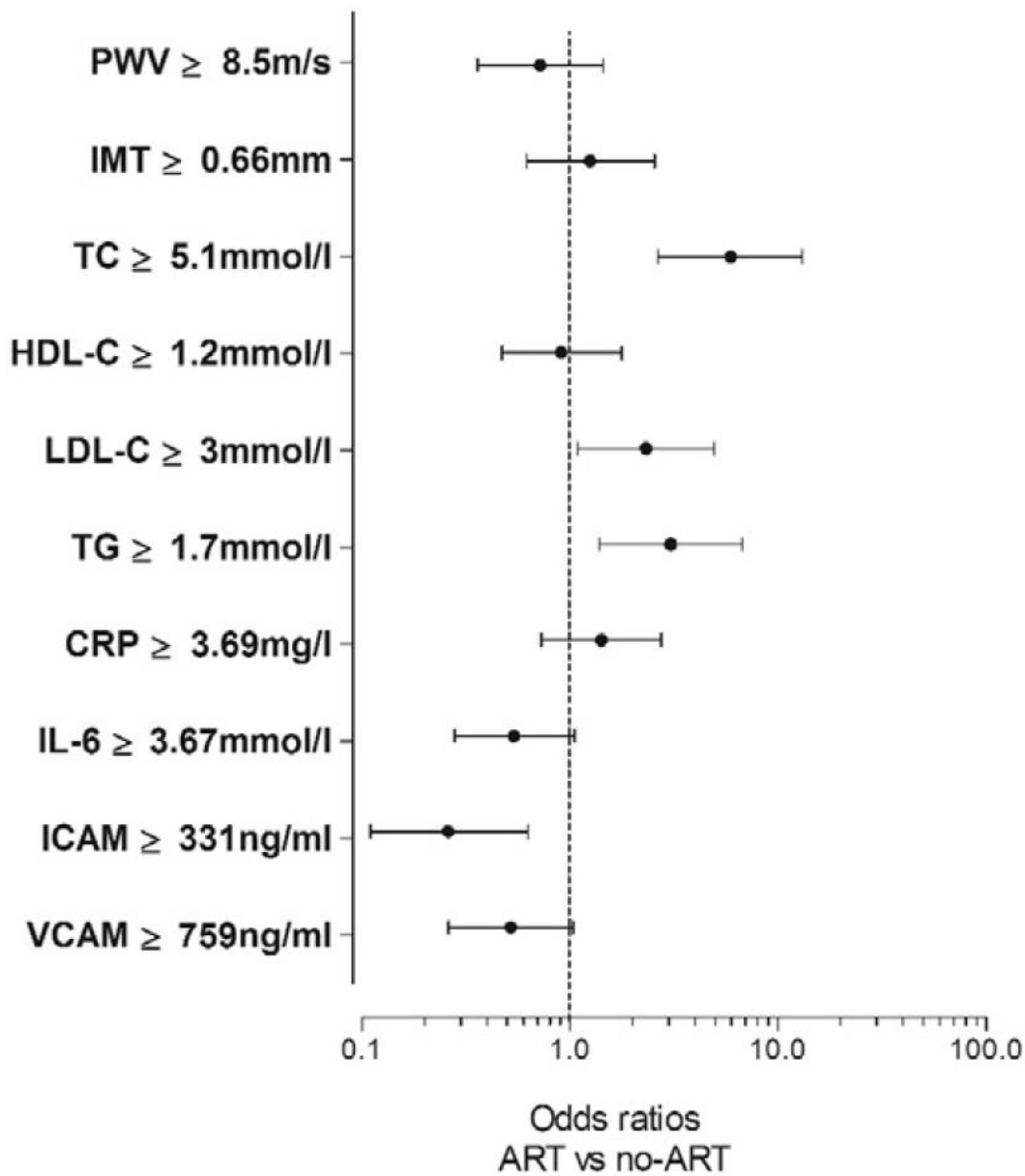


Fig. 2. Odds ratios for increased cardiovascular disease risk

Behavioural and other associated CVD risk factors

- Obesity & diet (sugar intake)
- Inflammation
- Alcohol
- Endothelial function/NO bioavailability
- HIV infection
- **Salt intake?**

□ About salt intake...

Population health in South Africa: a view from the salt mines

www.thelancet.com/lancetgh Vol 1 August 2013

Mandatory regulations passed in March, 2013, to begin in 2016, will affect the salt content of processed food and will be a key weapon in the fight against the rising burden of hypertension.¹



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SALT
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Conclusions:

- Although **CVD is the most important contributor to mortality** world-wide, it has been *different for Africa*.
- But all recent statistics point to CVD overtaking **infectious diseases**, such as HIV, tuberculosis and malaria as the greatest threats of health in Africa.
- Our findings confirm that **CV risk factors** associated with **urbanisation** (obesity, excessive alcohol use, sugar intake, psychological distress, inflammation) indeed **predict hypertension and CV mortality**.
- These risk factors, specifically **blood pressure**, are on an **increasing trajectory** that will be highly challenging to turn around (Westernised environment, poor health systems, low awareness, diagnosis, treatment and control).
- Question remains – how can we **get back the BPs from Donnison's era?**

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