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The Epidemiology of Cardiovascular Diseases in Sub-Saharan Africa: The Global Burden of Diseases, Injuries and Risk Factors 2010 Study

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Abstract

The epidemiology of cardiovascular diseases in sub-Saharan Africa is unique among world regions, with about half of cardiovascular diseases (CVDs) due to causes other than atherosclerosis. CVD epidemiology data are sparse and of uneven quality in sub-Saharan Africa. Using the available data, the Global Burden of Diseases, Risk Factors, and Injuries (GBD) 2010 Study estimated CVD mortality and burden of disease in sub-Saharan Africa in 1990 and 2010. The leading CVD cause of death and disability in 2010 in sub-Saharan Africa was stroke; the largest relative increases in CVD burden between 1990 and 2010 were in atrial fibrillation and peripheral arterial disease. CVD deaths constituted only 8.8% of all deaths and 3.5% of all disability-adjusted life years (DALYs) in sub-Saharan Africa, less than a quarter of the proportion of deaths and burden attributed to CVD in high income regions. However, CVD deaths in sub-Saharan Africa occur at younger ages on average than in the rest of the world. It remains uncertain if increased urbanization and life expectancy in some parts of sub-Saharan African nations will transition the region to higher CVD burden in future years.

Keywords

Cardiovascular disease; Sub-Saharan Africa; Mortality; Burden of disease; Global health

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The major CVDs vary in terms of underlying pathology (atherosclerosis, hypertensive target organ damage, infection, inflammation, anatomic deformities), interaction with other organ systems (pulmonary, immune, endocrine, neurologic, hematologic), and segment of the population affected. The prevalence pattern of the CVDs is distinctly different in sub-Saharan Africa compared with the rest of the world: infectious and inflammatory causes are relatively more common and atherosclerosis relatively less common. The past pattern of sub-Saharan African CVD epidemiology is not necessarily permanent, however. As parts of Sub-Saharan Africa urbanize and life expectancy increases, a larger share of the population may develop higher CVD rates, along with persistence of the lower CVD rates in poorer rural areas.¹ CVD epidemiology studies have been scarce in the region, and complete vital registration of deaths by cause and burden of disease assessments have been completed by only a handful of sub-Saharan African nations.² We therefore have an incomplete picture of CVD epidemiology in sub-Saharan Africa. In this review, we present the data on sub-Saharan Africa based on the Global Burden of Diseases, Injuries, and Risk Factor (GBD 2010) Study and discuss the strengths and limitations of study findings for sub-Saharan Africa.

The objective of the GBD 2010 Study was to estimate the epidemiology of 291 major diseases and injuries in 21 epidemiologic regions, including four sub-Saharan Africa regions (sub-Saharan Africa East, West, Central and South; Table 1).³ The GBD 2010 included substantially more source data on fatal and nonfatal CVD outcomes and more advanced estimation methods than prior versions of the GBD.⁴ Despite these advances, the problem of scarcity and uneven quality of data on CVD in Sub-Saharan Africa persists and is discussed in the section on study limitations.

Methods

The core summary measurement of population health in the GBD 2010 Study was disability-adjusted life years (DALYs) in the years 1990 and 2010. DALYs represent the “health gap” between a population's actual health and an ideal standard. DALYs are composed of years of life lost (YLL) to premature deaths and years lived with non-fatal disease disability (YLD).³ Ten major CVD cause categories were defined based on International Classification of Disease (ICD) classifications: stroke, ischemic heart disease, cardiomyopathy, rheumatic heart disease, hypertensive heart disease, endocarditis, atrial fibrillation, aortic aneurysm, peripheral arterial disease and “other cardiovascular and circulatory”. The last “other” category included cardiopulmonary disease (ICD-10 I27, I28), non-rheumatic valvular disease (I34, I35, I36, I37), disorders of the arteries, capillaries or veins (I72, I77, I78, I83, I84, I87, I88, I89), venous embolism and thrombosis (I82), hypotension (I95), post-procedural disorders (I97), and cardiovascular disorders in syphilis and other diseases (I98). For mortality and YLL estimates, the GBD 2010 Study assembled a global cause-of-death data base. Mortality data were mapped to GBD cause categories and distribution of deaths to CVD and other causes and regional mortality and temporal trends were estimated using the cause of death ensemble modeling (CODEm) approach.⁵ The cause of death correct (CoDCorrect) algorithm estimated total CVDs as a proportion of all deaths, after which the component CVD causes were fit into the total CVD mortality envelope. Non-fatal CVD prevalence was estimated from data gathered in systematic reviews of epidemiologic data

using a Bayesian meta-regression method (DisMod-MR).⁶ Disability from each case of nonfatal CVDs and other diseases or injuries was estimated in a household survey of lay people in Bangladesh, Indonesia, Peru, Tanzania, and the U.S.A., and an international web-based survey of health professionals.⁷ Because heart failure is not defined as an underlying cause of death in the International Classification of Diseases (ICD), separate methods were developed in order to distribute heart failure deaths to upstream CVDs and other causes of heart failure.⁸

Cause-of-death data from sub-Saharan Africa were usually not from standard vital registration, but were predominantly gathered from verbal autopsy studies, police reports, sibling histories, and burial and mortuary reports.⁵ With the exception of a few higher quality studies,^{9–11} most data on CVD in sub-Saharan Africa are from small community surveys and hospital-based registries.¹² The GBD dealt with the problem of absent or low quality epidemiologic data from sub-Saharan Africa by 1) incorporating covariates (CVD risk factors, national income, differences in measurement method) and “borrowing strength” from nearby regions and years of observation in CODEm and DisMod-MR models, and 2) using standard assumptions about the relationship between disease-specific incidence, prevalence, case fatality, and mortality in DisMod-MR models.^{3,6,8}

Results

Compared with all other world regions, the sub-Saharan Africa region had the smallest proportion of disease burden attributed to CVD in 2010: 8.8% of total deaths, 3.9 % of years of life lost, and 3.5% of DALYs (“tree plot” visualization of cause proportions, by region, available at <http://viz.healthmetricsandevaluation.org/gbd-compare/>). The corresponding CVD proportions for combined high-income regions were 35.7%, 27.2%, and 16.4%. Within sub-Saharan Africa, the Southern region had the highest proportional CVD burden and the Western region the lowest. The overall all-ages age-standardized sub-Saharan Africa 2010 CVD mortality rate was 892 deaths per 100,000 persons, and combined fatal and non-fatal burden of disease was 18,825 DALYs per 100,000 persons (Table 2; also by sub-Saharan Africa region, with uncertainty intervals, Table 1). Cerebrovascular disease had the highest death and disability rate of the top 10 CVDs in sub-Saharan Africa. Stroke mortality rates in 2010 ranged from 85 per 100,000 persons (95% CI 72–101) in Western sub-Saharan Africa to 126 per 100,000 person-years (95% CI 105–166) in Central sub-Saharan Africa. Age-adjusted DALYs ranged from 1665 per 100,000 persons (95% CI 1377–1968) in Western sub-Saharan Africa to 2434 per 100,000 persons (95% CI 2040–3041) in Central sub-Saharan Africa. Hemorrhagic and other non-ischemic stroke burden dominated over ischemic stroke burden across all regions of sub-Saharan Africa. Hemorrhagic and other non-ischemic stroke was the cause of 55% of stroke deaths and 64% of stroke DALYs.

In sub-Saharan Africa there were more women living with peripheral arterial disease in 2010 than there were men (9.85 versus 4.39 million). Compared to Europe and North America where the bulk of people with peripheral arterial disease is above 55 years, most PAD cases in sub-Saharan Africa were noted amongst younger people (<55 years). For atrial fibrillation, 2010 DALYs were higher for men compared to women (Table 2). However, mortality associated with atrial fibrillation was higher in women compared to men (Table 2).

All non-atherosclerotic CVDs (all CVDs excepting ischemic heart disease, thrombotic ischemic stroke, and peripheral arterial disease) were responsible for about half of all CVD mortality and DALYs in sub-Saharan Africa.

Rank of CVDs by proportion of CVD DALYs changed little in sub-Saharan Africa between 1990 and 2010, though absolute DALYs increased for all CVDs (Fig 1). Of the 20-year, 34% increase in DALYs in sub-Saharan Africa between 1990 and 2010, 65% of the DALY increase was due to population growth, 4% due to aging of the population, and the increase was attenuated by a 35% decrease in age-standardized CVD DALYs per capita population (decrease in disease rates). While absolute DALYs changed little between 1990 and 2010 for causes like endocarditis, DALYs about doubled for CVDs more common in older adults, like peripheral arterial disease and atrial fibrillation. While age-standardized DALY rates decreased for most CVDs in sub-Saharan Africa, age-standardized rates actually increased 16% for atrial fibrillation and 27% for peripheral arterial disease.

Because adults in sub-Saharan Africa are more likely to die from a non-CVD cause compared with the rest of the world, sub-Saharan Africa has the lowest years of life lost due to CVD (Table 3). At the same time, for persons dying of CVD, mean age of death in sub-Saharan Africa is the lowest in the world [64.9 years (95% uncertainty interval, 64.4–65.4)].

Discussion

The epidemiology of CVDs in sub-Saharan Africa is distinct because atherosclerosis causes only about half of CVDs. The GBD 2010 Study estimated that absolute CVD burden has increased in sub-Saharan Africa since 1990, with the largest relative increases in burden being atrial fibrillation and peripheral arterial disease—CVDs occurring most often in the elderly. Cerebrovascular disease, particularly hemorrhagic and other non-ischemic stroke, was the dominant source of CVD burden in the region. Because of the high burden of human immune deficiency virus (HIV), malaria, neonatal, and other non-CVD causes in sub-Saharan Africa, relatively few lives are lost due to CVD. On the other hand, the average age at CVD death in sub-Saharan Africa is the youngest in the world (Table 3).

Consistent with conclusions of systematic review studies, cerebrovascular disease, and particularly hemorrhagic and other non-ischemic stroke, emerged as the leading cause of CVD burden in sub-Saharan Africa.^{12,13} Our data on the relatively high rates of stroke mortality in sub-Saharan Africa compared with developed countries can be attributed to the higher stroke severity (for example, hemorrhagic strokes are clinically more severe than non-hemorrhagic strokes and have significantly greater case-fatality)¹⁴ and less than optimal acute stroke management in the region. The proportionally greater burden of hemorrhagic and other non-ischemic stroke compared with ischemic stroke burden is likely to be attributed to the relatively low prevalence of atherosclerosis (major cause of ischemic stroke) and high prevalence of hypertension (major cause of hemorrhagic stroke) in sub-Saharan Africa, as it was suggested for countries at the initial stages of epidemiological transition from communicable to non-communicable disorders.¹⁵ While there was no substantial variation in stroke burden across Eastern, Southern and Western regions of sub-Saharan Africa, stroke burden in Central sub-Saharan Africa was higher than in other

regions of sub-Saharan Africa. This observation may be related to the more advanced stage of epidemiological transition currently observed in Central sub-Saharan Africa compared with other regions of sub-Saharan Africa. However, further good quality stroke epidemiological studies are required in sub-Saharan Africa to reliably address causes of the very high stroke burden in the region and develop culturally appropriate strategies for stroke prevention.

Among the CVDs, the largest relative increase in burden of disease for sub-Saharan Africa between 1990 and 2010 was observed for atrial fibrillation. Since atrial fibrillation is far more common in men, the 2010 DALYs were predictably higher in men compared to women. However, the higher mortality associated with atrial fibrillation in women is an unexpected finding and clearly needs further detailed investigation. Given the established association between atrial fibrillation and stroke, it is possible that the former is contributing to continued high burden of the latter. Atrial fibrillation burden was not assessed in the first GBD project. However an overall increase in both atrial fibrillation incidence and prevalence at a global level has recently been reported (1990–2010).¹⁶ While our findings in sub-Saharan Africa do not provide a direct explanation for increase in atrial fibrillation burden, these results do allow for some speculation in this regard. Even though rheumatic heart disease has been associated with occurrence of atrial fibrillation, the significant decrease in the burden of this condition would suggest that factors other than rheumatic heart disease are responsible for higher atrial fibrillation burden. Ischemic heart disease, hypertension, and cardiomyopathies saw an increased burden in sub-Saharan Africa and these are all conditions associated with higher occurrence of atrial fibrillation. However, there are likely to be multiple factors affecting the rising burden of atrial fibrillation and these could be region-specific. Especially given the etiologic role of atrial fibrillation for stroke, significant additional surveillance is needed and more detailed analysis of the mechanisms of atrial fibrillation is warranted.

Peripheral arterial disease also stood out as one of the CVDs with the largest relative increase in sub-Saharan Africa since 1990. The observation that the prevalence of peripheral arterial disease was higher among adults younger than 55 years old is concerning. It suggests exposure to peripheral arterial disease risk factors – e.g., tobacco smoke exposure, diabetes, hypertension, or hypercholesterolemia – at relatively young ages.¹⁷ As peripheral arterial disease may be caused by atrial fibrillation – a condition that also increased in prevalence – atrial fibrillation should be considered as a factor leading to peripheral arterial disease in sub-Saharan Africa.

Though the GBD 2010 estimated for sub-Saharan Africa the world's lowest ischemic heart disease death rates and burden per capita, ischemic heart disease ranked second on the sub-Saharan Africa CVD rank list. The rarity of ischemic heart disease in the region has been documented only in a handful of community surveys and hospital registries.^{11,18} Lack of data has led to very different conjectures about the impact of urbanization and lifestyle changes on ischemic heart disease in sub-Saharan African populations.^{1,19} The example of Japan during the 1960's to 1990's should caution us that lifestyle changes and adverse risk factor trends, such as increases in population mean total cholesterol, do not necessarily spur increased ischemic heart disease mortality.²⁰

Heart failure is an important cause of death and disability in sub-Saharan Africa, and unlike in the high-income countries of North America and Europe, it is most often caused by hypertensive heart disease, rheumatic heart disease, or cardiomyopathy and rarely by ischemic heart disease.^{9,21–24} Heart failure is conspicuously absent among the major CVDs reported on here. Because the GBD cause list is based on ICD classifications, and in the ICD heart failure is not classified as an underlying cause of death and disability, in this analysis most of heart failure burden was assigned to its preventable root CVD causes (e.g., ischemic heart disease, rheumatic heart disease and other valvular heart disease, hypertensive heart disease, and cardiomyopathies). The shortcoming of this emphasis on primary heart failure prevention is that the potential benefits of treating heart failure cases and alleviating prevalent heart failure disability burden are not explicitly estimated, to some degree taking heart failure off the public health agenda for sub-Saharan Africa. It is likely that if heart failure underlying causes were reclassified into heart failure burden, heart failure would rise to be among the major CVDs in sub-Saharan Africa. GBD heart failure burden estimates are needed both for sub-Saharan Africa and globally and will be forthcoming.

Disease burden estimates are important for health system planners in the sub-Saharan Africa region. CVD epidemiology may be changing rapidly in the region, especially in areas where life expectancy is increasing (partly through improved neonatal and communicable disease prevention and treatment) and economic development and urbanization are progressing rapidly. The GBD made estimates for sub-Saharan Africa by “borrowing strength” from cause of death and epidemiology study data within the region and, in a geographically nested fashion, from other world regions. However, the problem of scarce and low-quality CVD epidemiologic data remains an endemic problem for sub-Saharan Africa. Even use of standard measurement instruments developed in other regions may risk erroneous estimates. For example, in the GBD ischemic heart disease analysis, World Health Survey self-reported angina prevalence was unexpectedly high in sub-Saharan Africa nations, perhaps because the Rose angina questionnaire is not equally valid in all populations.⁸ In 1998, responding to the original GBD (1990) Study, Cooper et al. argued that given lack of good quality vital registration and other surveillance studies, disease burden estimates for sub-Saharan Africa were overly uncertain and that the “complete” appearance of GBD estimates might lead to the misconception that more and better research is not needed.²⁵ We argue with equal force that the best possible epidemiology and burden estimates are needed for decision-making in sub-Saharan Africa now, and that more and better epidemiologic surveillance is desperately needed.

In conclusion, the GBD 2010 Study used hierarchical models and meta-regression in an effort to overcome the limitation of sparse data and estimate mortality and burden of disease for sub-Saharan Africa. Our estimates suggest that non-atherosclerotic CVDs are responsible for about half of the CVD burden in the region. Though CVD is responsible for a smaller share of deaths and DALYs compared with other diseases in sub-Saharan Africa, the average age at CVD death is the youngest in the world. Two important policy lessons emerge from the GBD 2010 Study's analysis of CVD in sub-Saharan Africa. First, given the predominance of cerebrovascular disease and hypertensive heart disease burden, hypertension control needs to be a high priority. Second, sparse data led to a high degree of uncertainty about the CVD estimates, leading us to point to the dire need for more support

for good quality, population-based epidemiologic research. The recent establishment of a number of high quality local and regional CVD epidemiology studies is an encouraging sign for sub-Saharan Africa CVD epidemiology and burden of disease estimates in the future.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Abbreviations and Acronyms

CVD	cardiovascular disease
GBD 2010	Global Burden of Disease 2010 Study
ICD	International Classification of Diseases
YLD	years lived with disability
YLL	years of life lost due to disability

1990		2010	
5,943,526 (39.5%)	1 Stroke	1 Stroke	7,840,477 (38.8%)
4,163,062 (27.6%)	2 Ischemic heart disease	2 Ischemic heart disease	5,780,399 (28.6%)
1,387,868 (9.2%)	3 Other cardio & circulatory	3 Other cardio & circulatory	1,829,864 (9.1%)
1,147,212 (7.6%)	4 Cardiomyopathy	4 Cardiomyopathy	1,580,805 (7.8%)
1,130,529 (7.5%)	5 Rheumatic heart disease	5 Hypertensive heart disease	1,414,406 (7.0%)
893,598 (5.9%)	6 Hypertensive heart disease	6 Rheumatic heart disease	1,187,594 (5.9%)
195,117 (1.3%)	7 Endocarditis	7 Endocarditis	210,552 (1.0%)
100,952 (0.7%)	8 Atrial fibrillation	8 Atrial fibrillation	193,005 (1.0%)
76,629 (0.5%)	9 Aortic aneurysm	9 Aortic aneurysm	120,453 (0.6%)
27,031 (0.2%)	10 Peripheral vascular disease	10 Peripheral vascular disease	55,219 (0.3%)

Fig 1. Leading causes of cardiovascular disease burden, sub-Saharan Africa, 1990 and 2010

Table 1

Sub-Saharan countries and GBD regions.

Central: Angola, Central African Republic, Congo, Democratic Republic of Congo, Equatorial Guinea, Gabon
East: Burundi, Comoros, Djibouti, Eritrea, Ethiopia, Kenya, Madagascar, Malawi, Mauritius, Mozambique, Rwanda, Seychelles, Somalia, Sudan, Tanzania, Uganda, Zambia
South Botswana, Lesotho, Namibia, South Africa, Swaziland, Zimbabwe
West: Benin, Burkina Faso, Cameroon, Cape Verde, Chad, Cote d'Ivoire, Gambia, Ghana, Guinea, Guinea-Bisub, Liberia, Mali, Mauritania, Niger, Nigeria, Sao Tome and Principe, Senegal, Sierra Leone, Togo

Table 2
Age-standardized mortality and DALYs per 100,000 people, 2010, all sub-Saharan African regions combined, the GBD 2010 Study.

Cause	All Sub-Saharan Africa Regions Combined					
	Males		Females		Both Sexes	
	DALYs	Deaths	DALYs	Deaths	DALYs	Deaths
All cardiovascular and circulatory diseases	20,270.10	926.28	17,563.61	858.36	18,824.98	891.53
Rheumatic heart disease	731.48	24.21	883.81	26.28	807.43	25.30
Ischemic heart disease	6859.08	311.88	4689.42	240.01	5700.08	272.80
Cerebrovascular disease	7955.06	401.89	7203.56	393.29	7568.48	399.03
Ischemic stroke	2810.70	172.24	2682.43	181.49	2749.08	178.43
Hemorrhagic and other non-ischemic stroke	5144.35	229.65	4521.14	211.79	4819.40	220.60
Hypertensive heart disease	1272.43	60.24	2025.35	105.09	1682.73	85.69
Cardiomyopathy and myocarditis	1420.36	54.34	968.39	37.60	1177.39	45.09
Atrial fibrillation and flutter	256.40	3.14	184.57	3.56	215.83	3.36
Aortic aneurysm	172.88	9.23	98.61	5.38	131.18	6.98
Peripheral arterial disease	105.04	3.82	56.97	1.58	77.91	2.52
Endocarditis	105.52	2.89	101.47	2.61	103.03	2.74
Other cardiovascular and circulatory diseases	1391.84	54.61	1351.45	42.94	1360.91	48.00

Table 3

Average age at CVD death and years of life lost (YLL) due to CVD death per capita, 2010, the GBD 2010 Study.

Super Region	Age at Death	YLL Per Capita
Sub-Saharan Africa	64.9 (64.4–65.4)	0.0216 (0.0191–0.0245)
East Asia/Pacific	72.6 (72.2–72.9)	0.0401 (0.0366–0.0421)
Eastern Europe/Central Asia	75.5 (75.3–75.7)	0.1084 (0.1050–0.1112)
High Income	81.2 (81.0–81.6)	0.0368 (0.0354–0.0396)
Latin America/Caribbean	73.6 (73.3–73.9)	0.0294 (0.0279–0.0315)
North Africa/Middle East	68.6 (68.3–68.9)	0.0442 (0.0406–0.0461)
South Asia	67.6 (66.9–68.2)	0.0352 (0.0299–0.0385)