REVIEW gREVIEW

Atrial Fibrillation in Africa—An Under-Reported and Unrecognized Risk Factor for Stroke



A Systematic Review

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ABSTRACT

Over three-quarters of deaths from cardiovascular disease and diabetes occur in low- and middle-income countries, which include many African countries. Global studies showed that the prevalence of the cardiac arrhythmia atrial fibrillation (AF) appeared to be lower in Africa. A systematic search of PubMed and African Journals Online was conducted to determine the prevalence of AF and associated stroke risk factors in Africa and to quantify the need for screening. The publications search yielded a total of 840 articles of which 41 were included. AF was often not identified as the disease of primary interest with its own risks. Data on prevalence in the general population was scarce. The prevalence of stroke risk factors showed a large variation between studies, as well as within clustered subpopulations. AF in Africa is under-reported in published reports. The study types and populations are highly heterogeneous, making it difficult to draw a definitive conclusion on AF prevalence.

Noncommunicable diseases cause more than one-half of the total burden of disease in Sub-Saharan Africa (SSA) [1-4]. Cardiovascular diseases constitute a major part of this burden. Strokes and heart attacks are by far the most common cause of cardiovascular death [1,5,6]. For prevention of these conditions, a total-risk approach is recommended by the World Health Organization (WHO) with integrated management of hypertension, diabetes, high cholesterol, and other cardiovascular risk factors. WHO identified counselling and multidrug therapy for people with a high risk of developing heart attacks and stroke as being among the best-buy interventions [7]. Interestingly, targeting of cardiac arrhythmias such as atrial fibrillation (AF) is not mentioned in any of the best-buy strategies, although it is an established risk factor for stroke. AF increases the risk of stroke about 5-fold and it is associated with a 1.5- to 2.0-fold increase in all-cause mortality [8]. A global increase of 42.8% in AF-related deaths has been seen from 2006 to 2016, though a proportion may be attributable to an increased likelihood of reporting AF or atrial flutter as an underlying cause of death [9]. The disease has a high prevalence that can go up to at least 10% in people age 80 years and over in highincome countries [10,11]. The prevalence of AF is not clear in low- and middle-income countries (LMIC), including countries in Africa. It is uncertain if the low incidence and prevalence reported for the African region is due to poor surveillance, under-reporting, or genetic predisposition [12]. Underestimation could have resulted from poor access to health care and weak or nonexistent surveillance systems. We therefore conducted a systematic

review of published reports to determine the prevalence of AF and associated stroke risk factors in Africa and to assess the need for appropriate screening and stroke prevention strategies.

METHODS

Search Strategy

We searched the PubMed database, which includes MEDLINE, for articles published between January 1, 2000, and April 15, 2018, on the epidemiology of AF in Africa. An additional search was performed in the African Journals Online (AJOL) database within the period January 1, 2000 (inserted as "1st of January 2004" as the earliest available date to choose) up to April 15, 2018 [13]. For the initial identification of published studies, we used the following search terms: "atrial fibrillation," "stroke," and "anticoagulation" as Medical Subject Headings, or MeSH, terms. Details of the search are outlined in the Online Appendix.

After the initial search phase in the databases, the secondary inclusion criterion was the term "atrial fibrillation" in the abstract combined with a number that could be the number of people with AF (n or %), the number of AF episodes, or prevalence and/or incidence of AF. If there was no indication for an incidence and/or prevalence number in the abstract, but it did contain the term "atrial fibrillation," the full text would be screened to see whether a number was included in the full-text manuscript. Articles that included the words "atrial fibrillation" in the abstract but did not provide any insight into incidence and/or prevalence were excluded. Articles were only included if

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the research described the African population living in Africa, the data were original (no duplicate publication), the full text was available, and the article was available in English. International articles were excluded if they did not provide country- and/or region-specific numbers for African countries [14]. All the articles were initially screened by 1 reviewer.

Data Extraction

The following data, if available, was extracted from the included publications: country/location; study design; type of population; population size; proportion of female subjects, average age (mean or median); estimates of AF prevalence and/or incidence; method of diagnosis; type of AF (paroxysmal, persistent, permanent); CHADS₂ (Congestive Heart Failure History, Hypertension History, Age ≥75 Years, Diabetes Mellitus History, Previous Stroke or Transient Ischemic Attack Symptoms) or CHA2DS2-VASc (Congestive Heart Failure of Left Ventricular Dysfunction, Hypertension, Age ≥75 [doubled], Diabetes, Thromboembolism or Stroke History [doubled], Vascular Disease, Age 65 to 74 Years, and Sex Category) score; HAS-BLED (Hypertension, Abnormal Renal or Liver Function, Stroke History, Bleeding History, Labile International Normalized Ratio, Elderly (Age >65), Drugs or Alcohol) score; presence of stroke risk factors including diabetes, hypertension, stroke history vascular disease, heart failure, and age ≥75 years old; anticoagulant and/or antiplatelet drug use in AF patients. The stroke risk factors included were based on the items included in the CHA2DS2-VASc score [15].

If specific numbers for the AF cohort were not available, we reported the information available for the whole population. Study country was the country where patient recruitment took place. The study types were coarsely classified as a retrospective cohort study, prospective cohort study, or a case-control study.

Quality Assessment

Study quality was assessed using the checklist of Downs and Black [16]. This checklist is appropriate for both randomized and nonrandomized studies. The checklist item on power (item 27) was not taken into account; the maximum achievable score was therefore 27. Score ranges were given an overall quality level to qualify the available evidence included in the review: excellent (26 to 27); good (20 to 25); fair (15 to 19); and poor (≤14).

RESULTS

The systematic search identified 776 eligible articles from PubMed and 52 articles through the AJOL search. Two articles were indexed in both PubMed and AJOL. Of the identified articles, 115 were excluded because a full text was not available or the article was not in English. A total of 699 abstracts were screened and after exclusion of all articles that did not meet the predefined inclusion criteria, 41

original articles remained. The results are summarized in the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) flowchart (Fig. 1).

Quality Assessment

Overall, 14 studies had the quality score "poor," 26 studies were scored "fair," and only 2 were scored "good." Low scores were partially due to the exploratory, non-randomized nature of the studies. Study quality assessment results with a detailed summary of subscores for the questionnaire items can be found in Online Table 1.

AF Prevalence

The studies identified mainly focused on different patient subgroups and only 3 of the included articles described a population that would be representative for the general population. The results are summarized in Table 1 [17-57]. AF prevalence in the older, general population was 0.30% to 0.70% [17-19]. The study of Dewhurst et al. [17] in a rural area calculated a crude prevalence of 0.67% and an age-adjusted prevalence of 0.64%. Prevalence was different between men (0.31%) and women (0.96%). Also, prevalence increased with age from 0.46% in people age 70 to 74 years up to 1.3% to those age 85 years and older [17]. The age limit for the prevalence number of 0.30% was 50 years and over, and for the higher prevalence of 0.70%, the age limit was 70 years and over [17,18]. In patients with known cardiovascular disease presenting at the hospital, the prevalence varied from 3.8% to 59.0% [20-30]. Patient inclusion for these studies on patients with cardiovascular disease was mainly carried out at a hospital and also described patients that were previously diagnosed with AF. The most frequently used method for AF diagnosis confirmation was a 12-lead electrocardiogram. In heart failure patients, the prevalence was higher and ranged from 15.7% up to 34.0% [31-35]. Four of the studies on heart failure patients were hospital-based and 1 included outpatients [32]. Patients who (recently) underwent cardiac surgery had an AF prevalence of 4.0% to 17.0% [36-40]. Stroke patients, which include patients who had an acute stroke or had a history of stroke, had an AF prevalence of 1.5% to 17.6% [41-47]. Patients with rheumatic heart disease (RHD), had a prevalence of 13.9% to 44.5% [27,48-52]. Patients with RHD represented a more heterogeneous population, including children as well as adults, and with 2 studies also including outpatient data [48,52]. Studies in anticoagulation clinics showed that 25.1% to 65.0% of the patients were using oral anticoagulation for nonvalvular AF as the indication [53-57].

Several studies had information on the type of AF, which is generally categorized as paroxysmal, persistent, or permanent [21,23,24,27,36,44]. Persistent AF was the most found type of AF, ranging from 21.2% to 81.0%. Permanent AF was seen in 5.0% to 58.8% of the patients, and paroxysmal AF accounted for 11.8% to 32.1% of the AF cases (see Table 2) [17-57].

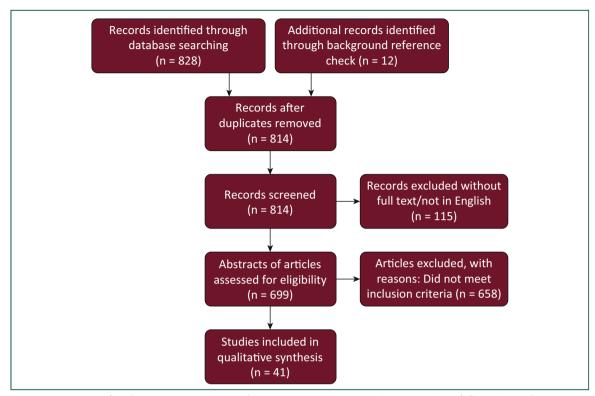


FIGURE 1. PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) flow chart of systematic review article screening and reasons for exclusion.

Stroke Risk Factors

The prevalence and presence of stroke risk factors were deducted from patients' characteristics tables of the included studies. The prevalence of stroke risk factors showed a large variation between studies but also within clustered subpopulations. Studies describing a population with known cardiovascular disease—AF or any other type of cardiovascular disease—included the most comprehensive information on stroke risk factors. Stroke history and the proportion of patients age 75 years and over were the factors that were least reported. The proportion of patients age 75 years and over was the least reported (non-modifiable) factor, reported in only 4 studies, and in general <30% of the patients were age ≥75 years.

Hypertension was the most frequently reported stroke risk factor, with prevalence at least around 60% to 70% for all studies. Prevalence seemed higher in the stroke subpopulation with a range of 31.0% to 93.8%, and it was lowest in RHD patients ranging from 5.2% to 53.3%. Diabetes prevalence ranged from as low as 8% in AF patients without valvular disease in Kenya up to 80.0% in coronary artery bypass surgery patients [27,39]. The definition of vascular disease varied between studies ranging from 6.0% up to 52.2% in post—cardiac surgery patients in South Africa [38]. Some studies only used a history of myocardial infarction, and other studies did not include a

definition for vascular disease. Stroke history was reported in less than one-half of the included studies. For the sub-population of stroke patients, the stroke history was scored prior to the admission. Overall, the reported stroke history showed a large variation ranging between 1.3% and 40.0% for the nonstroke patient subpopulation. Heart failure was present in 16.6% up to 60.3% of the population when excluding the subpopulation with known heart failure.

Indication for Stroke Prevention, Anticoagulant Use, and Aspirin Use

Only 9 studies mentioned the CHADS₂ or CHA₂DS₂-VASc score. The reported CHADS₂ scores ranged from 1.5 to 2.4 and CHA₂DS₂-VASc scores ranged from 3.1 up to 3.9. The use of oral anticoagulation (OAC) and/or aspirin was reported in 10 studies, excluding the studies that focused on international normalized ratio (INR) clinics. All studies conducted in INR clinics described warfarin as the OAC being used [53-57]. Anticoagulants use showed a wide variation from 7.0% up to 81.2% of the patients, which included AF and non-AF patients. Aspirin was used in 14.0% to 94.1% of the patients. AF was the indication for OAC use in 25.1% to 65.0% of the patients that visited an INR clinic. Only 2 studies reported OAC and aspirin use for patients with a CHADS₂ score of 2 or higher, OAC use

TABLE 1. Study characteristics of included articles describing AF prevalence in Africa

					Patients	Female		
Author(s) Ref. #	Year	Study Type	Country	Population	(n)	(%)	Age*	AF (%)
General population								
Dewhurst et al. [17]	2012	Prospective cohort study	Tanzania	People age 70 yrs and over in a rural African community	2,232	56.3	77.8	0.70
Koopman et al. [18]	2014	Prospective cohort study	Ghana	Individuals age 50 yrs and over in a rural area	924	48.1	66.0	0.30
Gray et al. [19]	2016	Prospective cohort study	Tanzania	People age 70 yrs and over	2,232	56.3	77.8	0.60
Patients with known cardiov	ascular dise	ease						
Nqayana et al. [20]	2008	Retrospective cohort study	South Africa	Patients with cardiac disease in pregnancy	95	100.0	_	9.5
Ntep-Gweth et al. [21]	2010	Prospective cohort study	Cameroon	Patients >18 yrs with AF documented on an ECG during the index visit	172	56.4	65.8	100.0
Sliwa et al. [22]	2010	Prospective cohort study	South Africa	Patients presenting to the cardiology unit (Heart of Soweto study)	5,328	61.0	58.8	4.6
Shavadia et al. [23]	2013	Retrospective cohort study	Kenya	Patients with discharge diagnosis of either AF or atrial flutter	162	44.0	67.0	59.0
Jardine et al. [24]	2014	Prospective cohort study	South Africa	Patients with confirmed AF drawn from the private insured sector	302	40.1	67.0	100.0 [†]
Gamra et al. [25]	2014	Cross-sectional observational study	Algeria, Egypt, Morocco, Tunisia	Patients with a history of AF documented by ECG	1,680 [‡]	61.1	64.2	100.0
Akpa and Ofori [26]	2015	Retrospective cohort study	Nigeria	All patients referred to the cardiology unit or seen in the cardiac clinic with ECG evidence of AF	228	42.6	59.8	29.8
Bloomfield et al. [27]	2015	Prospective cohort study	Kenya	Patients age 18 yrs and over with AF based on ruling out significant valvular heart disease	298	53.0	68.0	24.0
Ajayi et al. [28]	2016	Prospective cohort study	Nigeria	Patients with an ECG at the hospital	1,462	67.4 [§]	67.3	3.8
Yameogo et al. [29]	2016	Retrospective cohort	Burkina Faso	Patients at the cardiology department with nonvalvular heart disease	970	44.6	65.5	7.0
Temu et al. [30]	2017	Prospective cohort study	Kenya	AF patients in a teaching/referral hospital: both nvAF as well as vAF	146	55.0	69.4	47.3
Patients with HF								
Stewart et al. [31]	2008	Prospective cohort study	South Africa	De novo presentation of patients with HF and related cardiomyopathies	844	57.0	55.3	53.0
Magaña-Serrano et al.	2011	Prospective cohort study	Tunisia, Algeria	Patients with HF and preserved ejection fraction	648	47.0	63.0	34.0
Damasceno A [33]	2012	Prospective cohort study	Cameroon, Ethiopia, Kenya, Mozambique, Nigeria, Senegal, South Africa, Sudan	Patients admitted to the hospital with diagnosed acute HF	1,006	50.8	52.3	18.3

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NA-1In: -+ -1 [24]	2014	Dunana ati sa nahant	Tanzania	Patients age 18 yrs and over with HF	427	F1 0	FF 0	15.7
Makubi et al. [34]	2014	Prospective cohort study	attending a cardiovascular center		427	51.0	55.0	15.7
Ogah et al. [35]	2015	Prospective cohort study	Cameroon, Ethiopia, Kenya, Mozambique, Nigeria, Senegal, South Africa, Sudan, Uganda	Patients with HF from cardiology units	1,006	50.8	52.4	18.4
Patients undergoing/who u	nderwent ca	ordiac surgery	, ,					
Lorgat et al. [36]	2012	Prospective cohort study	South Africa	Patients who underwent robotically assisted catheter ablation therapy in a hospital	95	25.0	59.4	100.0 [†]
Mazibuko et al. [37]	2012	Retrospective cohort study	South Africa	Pregnant patients with a prosthetic valve prostheses referred to a tertiary hospital	61	100.0	24.0	13.1
Mansoor [38]	2014	Retrospective cohort study	South Africa	Post—cardiac surgery patients	997	44.1	51.9	5.9
Abdel-Salam and Nammas [39]	2016	Prospective cohort study	Egypt	Patients scheduled for CABG with/without valve surgery	740	30.5	56.5	10.4
Bun et al. [40]	2016	Prospective cohort study	Morocco	Patients scheduled for radiofrequency catheter ablation or pacemaker implantation	75	29.3	67.8	4.0
Chtaou et al. [41]	2016	Retrospective cohort study	Morocco	All patients treated with rt-PA at a stroke unit	52	50.5	63.0	17.0
Stroke patients								
Alkali et al. [42]	2013	Prospective cohort study	Nigeria	Patients presenting with acute stroke	272	38.2	55.1	9.2
Walker et al. [43]	2014	Case-control study	Tanzania, Hai	Incident stroke cases, age/sex-matched control subjects	93	46.2	68.2	6.5
Walker et al. [43]	2014	Case-control study	Tanzania, Dar-es- Salaam	Incident stroke cases, age/sex-matched control subjects	39	38.5	61.1	7.7
Germain et al. [44]	2015	Retrospective cohort study	Burkina Faso	Hospitalized patients at the cardiology and neurology departments with ischemic stroke	391	62.3 [§]	63.3	17.6
Gomes et al. [45]	2015	Case-crossover study	Mozambique	Patients admitted to the hospital for newly occurring ischemic and hemorrhagic stroke	593	47.5	58.8	5.6
Adeoye et al. [46]	2017	Prospective cohort study	Nigeria, Ghana	Acute stroke patients who had a 12-lead ECG recording within first 24 h of admission	890	58.3	58.3	4.2
Lekoubou et al. [47]	2017	Retrospective cohort study	Cameroon	Stroke patients that were admitted to the hospital	1,678	50.2	62.0	1.5
Patients with RHD								
Sliwa et al. [48]	2010	Prospective cohort study	South Africa	De novo cases of RHD presenting to the cardiology unit	344	68.0	43.0	34.0
Okello et al. [49]	2013	Prospective cohort study	Uganda	Patients with newly diagnosed RHD	309	63.5	30.0	13.9
								(continued)

TABLE 1. Continued

Author(s) Ref. #	Year	Study Type	Country	Country Population		Female (%)	Age*	AF (%)
Zhang et al. [50]	2013	Prospective cohort study	Uganda	Patients with newly diagnosed RHD	130	71.0	33.0	13.9
Bloomfield et al. [27]	2015	Prospective cohort study	Kenya	Patients age 18 yrs and over with nvAF and vAF	298	78.0	38.0	24.0
Zühlke et al. [51]	2016	Prospective cohort study	LIC: Ethiopia, Kenya, Patients with symptomatic RHD visiting the Malawi, Rwanda, hospital Uganda, and Zambia		1,110	65.8	24.0	18.2
Zühlke et al. [51]	2016	Prospective cohort study	UMIC: Namibia and South Africa	nd Patients with symptomatic RHD visiting the hospital		71.3	39.0	27.5
Okello et al. [52]	2017	Prospective cohort study	Uganda	Patient ages 5-60 yrs with established RHD	449	66.8	31.4	20.3
Patients visiting an INR clinic								
Njovane et al. [53]	2013	Retrospective cohort study	South Africa	Patients using warfarin that attended a primary health care clinic	111	_	_	44.5
Anakwue et al. [54]	2014	Retrospective cohort study	Nigeria	Patients that utilized anticoagulation services in a university hospital	26	53.4	53.8	30.8
Sonuga et al. [55]	2016	Retrospective cohort study	South Africa	Patients attending the INR clinic	136	56.6	64.0	65.0
Schapkaitz and Sithole [56]	2017	Prospective cohort study	South Africa	Patients attending the anticoagulation clinic	147	57.8	56.8	36.1
Ebrahim et al. [57]	2018	Retrospective cohort study	South Africa	Patient visiting the INR clinic	363	65.6	55.0	25.1

AF, atrial fibrillation; CABG, coronary artery bypass graft; ECG, electrocardiography; HF, heart failure; INR, international normalized ratio; LIC, low-income country; nvAF, nonvascular atrial fibrillation; RHD, rheumatic heart disease; rt-PA, recombinant tissue plasminogen activator; UMIC, upper-middle-income country; vAF, valvular atrial fibrillation.

^{*}Italics indicate median age.

[†]Prevalence is 100% because the nominator and denominator were the same due to preselection of AF patients and no numbers of the larger sample population were given.

[‡]Also includes patient from Lebanon, numbers not differentiated for African countries.

[§]Average for AF subcohort.

TABLE 2. Stroke risk factors in patients with AF in Africa

		AE Timo.				04011	Aspirin					
Author [Ref. #]	Year	AF Type: Paroxysmal; Persistent; Permanent	CHADS ₂ * or CHA ₂ DS ₂ -VASc	HAS- BLED	Diabetes (%)	HT (%)	Stroke History (%)	Vascular Disease (%)	HF (%)	Age ≥75 yrs (%)	OAC Use in AF Patients (%)	Use in AF Patient (%)
General population						_						_
Dewhurst et al. [17]	2012	_	_	_	_	_	_	_	_	60.7	_	_
Koopman et al. [18]	2014	_	_	_	_	24.2	1.2 [†]	_	_	_	_	_
Gray et al. [19]	2016	_	_	_	_	73.2	2.9	-	_	_	_	_
Patients with known c	ardiova	scular disease										
Nqayana et al. [20]	2008	_	_	_	_	_	_	13.7^{\dagger}	_	_	_	_
Ntep-Gweth et al. [21]	2010	22.7; 21.5; 55.8	1.9 \pm 1.1	-	10.5	64.5	17.4	6.4	58.1	27.3	30.8	58.7
Sliwa et al. [22]	2010	_	1.5 ± 0.9	_	_	60.0 [‡]	_	6.5 [‡]	56.0 [‡]	_	50.0 [§]	50.0 [§]
Shavadia et al. [23]	2013	_	64.7% ≥2	_	33.0	68.0		19.0	38.0	_	79.8	17.8
Jardine et al. [24]	2014	32.1; 21.2; 46.7	3.08	_	15.6	65.9	_	26.8	32.5	_	75.2	39.4
Gamra et al. [25]		27.9; 21.6; 50.5	_	_	22.7	54.7	_	19.1	16.6	_	52.5	21.9
Akpa and Ofori [26]		_	39.7% CHADS ₂ =1	_	_	58.8 [‡]	_	_	-	_	8.8	94.1
Bloomfield et al.	2015	-	-	_	8.0	67.0	25.0	_	_	-	_	_
Ajayi et al. [28]	2016	_	2.38	_	18.2 [‡]	87.3 [‡]	40.0 [‡]	_	36.3 [‡]	_	_	_
Yameogo et al. [29]		11.8; —; 58.8	3.9 ± 1.6	3.5 ± 1.5		_	_	_	60.3 [‡]	26.0	35.3	76.5
Temu et al. [30]	2017	_	2.2	0.0 _ 1.0	8.7	72.5	26.1	_	49.0	_	81.2	-
Patients with heart fai			2.2		0.7	, 2.3	20.1		13.0		01.2	
Stewart et al. [31]	2008	_	_	_	_	_	_	_	100.0	_	_	_
Magaña-Serrano	2011	_	_	_	42.0	76.0	5.0	50.0	100.0	_	_	_
et al. [32]	2011				42.0	70.0	5.0	30.0	100.0			
Damasceno et al.	2012	_	_	_	11.4	55.5	_	-	100.0	_	7.0	35.0
Makubi et al. [34]	2014	_	_	_	45.0	_	3.7	9.0	100.0	_	_	_
Ogah et al. [35]	2015	_	_	_	11.4	55.5	2.5	9.4	100.0	_	_	_
Patients undergoing/w		lerwent cardiac si	ırgerv		11.7	33.3	2.3	ут	100.0			
Lorgat et al. [36]		14.0; 81.0; 5.0	— —	_	9.0	52.0	24.0#	_	_	_	_	_
Mazibuko et al. [37]		— —	_	_	- -		_	_	_	_	_	_
Mansoor [38]	2012	_	_	_	40.7 [‡]	59.3 [‡]	_	52.2	_	_	_	_
Abdel-Salam and Nammas [39]	2014	_	_	_	80.0	66.8	_	_	_	_	_	_
Bun et al. [40]	2016	_	_	_	_	_	_	_	_	_	_	_
Chtaou et al. [41]	2016	_			12.0	31.0	10.0	6.0				
Stroke patients	2010	_		_	12.0	31.0	10.0	0.0			_	_
Alkali et al. [42]	2013		_	_	25.6	83.7		_				_
		-	_		25.0	05./		_		_		_
Walker et al. [43]	2014	_	_	_	_	_	_	_	_	_	_	_
Walker et al. [43]	2014	— 12 0. E2 1. 34 0	4.72	20 10/ >	— 21 7 [‡]	— ог г <u>†</u>		— 7 2 [‡]			— 70.7	140
Germain et al. [44]		13.0; 52.1; 34.8	4.72	39.1% ≥ 3	21.7 [‡]	85.5 [‡]	100 [‡]	7.2 [‡]	20.3 [‡]	18.8 [‡]	79.7	14.0
Gomes et al. [45]	2015	_	_	_	13.4	86.2	18.9	_	_	_	_	_
Adeoye et al. [46]	2017	_	_	_	32.9	93.8	_	_	_	_	_	_
Lekoubou et al. [47]			_	_	12.8	67.9	14.5	_	_	_	_	_
Patients with rheumat		t disease										
Sliwa et al. [48]	2010	_	_	_	_	_	_	_	18.0	_	_	_
Okello et al. [49]	2013	_	_	_	-	32.7	1.3	_	46.9**	_	_	_
Zhang et al. [50]	2013	_	_	_	_	53.3	_	_	45.9**	_	_	_

TABLE 2. Continued

							Aspirin					
		AF Type: Paroxysmal;					Stroke	Vascular		Age	OAC Use in AF	Use in AF
		•	CHADC *	LLAC	D:-b-+	шт				•		
		Persistent;	CHADS ₂ * or	HAS-	Diabetes		History	Disease		≥75	Patients	Patients
Author [Ref. #]	Year	Permanent	CHA ₂ DS ₂ -VASc	BLED	(%)	(%)	(%)	(%)	HF (%)	yrs (%)	(%)	(%)
Bloomfield et al.	2015	_	_	_	1.0	27.0	23.0	_	_	_	_	_
[27]												
Zühlke et al. [51]	2016	_	_	_	_	5.2	_	43.5	_	_	_	_
Zühlke et al. [51]	2016	_	_	_	-	14.5	_	25.8	_	_	_	-
Okello et al. [52]	2017	_	_	_	_		_		_	_	_	_
Patients visiting an IN	R clinic											
Njovane et al. [53]	2013	_	_	_	16.2	60.1	_	_	_	_	100.0	-
Anakwue et al. [54]	2014	_	_	_	_	_	_	_	_	_	100.0	-
Sonuga et al. [55]	2016	_	_	_	27.2	69.9	12.5	25.7	_	_	100.0	-
Schapkaitz and	2017	_	_	_	_	_	_	_	_	_	100.0	-
Sithole [56]												
Ebrahim et al. [57]	2018			_	_	_	_	_	_	_	100.0	

CHADS₂, Congestive Heart Failure, Hypertension, Age \geq 75, Diabetes, Stroke or Transient Ischemic Symptoms; CHA₂DS₂.VASc, Congestive Heart Failure of Left Ventricular Dysfunction, Hypertension, Age \geq 75, Diabetes, Thromboembolism or Stroke History, Vascular Disease, Age 65 to 74 Years, and Sex Category; HAS-BLED, Hypertension, Abnormal Renal or Liver Function, Stroke History, Bleeding History, Labile International Normalized Ratio, Elderly (Age > 65), Drugs or Alcohol; HT, hypertension; INR, international normalized ratio; OAC, oral anticoagulation.

was 52.5% to 79.8% and aspirin use was 17.8% to 21.9% in these patients [23,25]. Patients with known AF had a mean OAC use between 8.8% and 81.2% and aspirin use ranging from 17.8% up to 94.1% [21-27,29,30].

DISCUSSION

AF is under-reported in published reports from the African region. None of the included studies identify the extent of asymptomatic AF, which is crucial information to determine the overall prevalence and incidence. Underdiagnosis is most probably high and studies that focus on AF screening are lacking. This information is essential to determining the true AF burden and related severe consequences such as stroke. The study types and study populations are highly heterogeneous, making it difficult to draw definitive conclusions on AF prevalence in African populations. The focus of published studies seems to be on AF as a risk factor in other comorbid disease such as stroke, heart failure, or cardiovascular disease in general. AF is often not identified as the disease of primary interest with its own risks and complications. Data on prevalence in the general population was scarce; numbers that approached generalizable prevalence numbers ranged from 0.30% to 0.70% in patients ages 50 or 70 years and over, respectively. Remarkably, none of the studies included in this systematic review were used in the GBD (Global Burden of Disease) 2010 study [12]. The GBD estimates for AF seem not to be representative of the true AF burden in SSA and it is most likely an underestimation because studies with higher estimates and studies describing asymptomatic AF were not taken into account. This also illustrates that evidence on AF in this region is difficult to identify, even with a systematic approach, and search strategies need to be more tailored to the region. Based on published data, prevalence of AF in Africa seems low when compared with in the Western population where prevalence is at least 5.5% in people age 65 years and over [10,58]. AF patients in Africa are in general younger than AF patients in high-income countries [59]. Stroke also tends to occur at an earlier age in the African population when compared with age of occurrence in high-income countries [42-47]. Reasons for an earlier presentation of the disease and its complications are not clear, though it seems that stroke type patterns are different for Africans.

The INTERSTROKE (Global and Regional Effects of Potentially Modifiable Risk Factors Associated With Acute Stroke in 32 Countries) study [60] showed that the

^{*}CHADS2 scores are italic.

[†]Only myocardial infarction scored.

[‡]Percentages for the AF subcohort.

[§]Fifty percent used an anticoagulant or aspirin.

 $^{\|}$ In patients with CHADS $_2 > 2$.

Ten percent of these patients also used warfarin.

[#]Ischemic heart disease

^{**}Pulmonary hypertension.

prevalence of ischemic stroke was lower and the prevalence of hemorrhagic stroke was higher in Africa and other LMIC than in high-income countries. In a SIREN (Stroke investigative research and educational network) study [61], a case-control study carried out in Ghana and Nigeria that aimed to identify and characterize stroke risk factors in SSA, 68% of the cases had ischemic stroke. However, this SIREN study [61] did not include data on AF as an individual factor and was only included in the combined risk factor "cardiac disease," which had an odds ratio of 2.21 (99% confidence interval: 1.38 to 3.54) for ischemic stroke.

A genetic component could play an important role in global and regional differences. The INTERSTROKE study [60] looked at global and regional effects of risk factors associated with acute stroke. AF was a significant predictor for ischemic stroke in Africa (odds ratio: 4.59; 99% confidence interval: 3.66 to 5.75; n = 1,364), and this effect was comparable in Western countries [60]. The population attributable risk of AF in stroke was lower in Africa than in the Western countries: 7.9% versus 17.1% [60]. The ORBIT-AF (Outcomes Registry for Better Informed Treatment for Atrial Fibrillation) study [62] found no difference in outcomes associated with AF, which included stroke, systemic embolism, transient ischemic attack, and various types of hospitalizations. Studies included in this review that described subpopulations of stroke patients showed an AF prevalence ranging from 1.5% to 17.6%, which was much lower than expected based on the knowledge that 20% to 30% of all strokes are due to AF [8].

The higher prevalence of RHD in African countries introduces an increase in complications related to AF, already at a young age [63]. Valvular heart disease can be associated with an increased thromboembolic risk, which probably also adds to the stroke risk in AF patients [64]. In the studies included in this systematic review, we found AF prevalence ranging from 13.9% up to 44.5% in the subpopulations describing RHD. The REMEDY (Global Rheumatic Heart Disease Registry) study [51], a large prospective registry in 12 African countries, India, and Yemen found AF as a comorbidity in 18.2% of the patients for the low-income countries (Ethiopia, Kenya, Malawi, Rwanda, Uganda, and Zambia), 22.6% for the LMIC (Egypt, India, Mozambique, Nigeria, Sudan, and Yemen), and 27.5% in the uppermiddle-income countries (South Africa and Namibia).

Anticoagulant use in the populations studied was low and so was access to surgical interventions, especially in SSA. The included studies suggest that stroke prevention strategies in AF are suboptimal. WHO recommends aspirin as the drug of choice to reduce heart attacks and strokes in LMIC. It is understandable to choose a drug therapy that is affordable in these regions; however, this also implies that we would accept to provide suboptimal therapy. Analyses comparing the use of aspirin to OAC have shown that antiplatelet therapy is around 40.0% to 50.0% less effective in reducing the stroke burden in nonvalvular AF patients, though treatment benefit from vitamin K OAC is highly

dependent on the time in therapeutic range [65,66]. Nonvitamin K antagonist OAC do not require routine monitoring and could therefore be a good alternative to vitamin K OAC therapy, with due consideration of prices that in all probability will drop the next couple of years because of expiring patents and also with possible inclusion on WHO's essential drug list.

AF is not mentioned as a target for stroke prevention in any of the global actions plans or best-buy strategies of WHO. The World Heart Federation has published a very useful roadmap for nonvalvular AF in LMIC. One of the potential solutions includes optimization of AF detection, for example with a screening program by nonphysician health workers [67]. Screening for AF as a target for stroke prevention can be a cost-effective strategy depending on how the screening is arranged, the prevalence of AF and also the initiation of an appropriate stroke prevention strategy [68-77]. Opportunistic screening for AF is recommended in patients >65 years of age and systematic electrocardiogram screening may be considered to detect AF in patients age >75 years or those at high stroke risk according to the European Society of Cardiology guideline and the American Heart Association [8,78]. Systematic screening would be a good approach to determine the true prevalence of AF in the general population in Africa. In a low-resource region, it is very plausible that underdiagnosis is keeping the AF prevalence estimates artificially low. When taking into account that AF patients seem to be younger in Africa, one should consider screening at an earlier age to determine in which subpopulation this intervention would be most beneficial. Research has to be carried out to unravel racial differences in AF prevalence as well as consequential stroke severity and mortality. More attention needs to be paid to diagnosis and timely detection of AF in Africa. A broader prevention approach that also focuses on conditions predisposing to AF such as hypertension, heart failure, valvular heart disease, diabetes mellitus, and obesity should be the future practice.

CONCLUSIONS

AF is under-reported in published reports from the African region. Prevalence of AF seems lower and patients are generally younger compared with numbers found in high-income countries. Reasons for the earlier presentation of the disease and its complications are not clear. Numbers found most likely represent an underestimate because access to health care, use of routine monitoring electrocardiography, and patient surveillance is less common in Africa. Screening for AF would be an important approach to determine the true prevalence in the general population and to identify asymptomatic AF patients at risk for stroke.

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