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Hypertension in women, Gambia, Kenya, Mozambique

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Measured and self-reported hypertension among women of reproductive age, Gambia, Kenya, Mozambique

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Abstract

Problem In sub-Saharan Africa, hypertension prevalence is usually estimated from participant recall. We assessed the accuracy of self-reported hypertension in women of reproductive age.

Approach In PRECISE (PREgnancy Care Integrating translational Science, Everywhere), an observational prospective cohort study, we recruited 1825 non-pregnant women of reproductive age, 610 in the Gambia, 609 in Kenya and 606 in Mozambique. We compared self-reported and measured hypertension (systolic blood pressure ≥ 140 mmHg or diastolic blood pressure ≥ 90 mmHg). We adjusted hypertension prevalence for age, body mass index, education, parity, and antihypertensive medicine and oral contraceptive use.

Local setting PRECISE was conducted in both urban and rural hospitals or clinics.

Relevant changes The women were generally in their late twenties and parous. Adjusted measured hypertension prevalence was higher in Mozambique (10.4%; 95% confidence interval, CI: 7.9–12.7) and the Gambia (9.3%; 95% CI: 6.6–12.6) than in Kenya (4.6%; 95% CI: 3.0–6.6). Self-reported hypertension prevalence was highest in the Gambia (12.9%; 95% CI: 10.2–15.9) versus Mozambique (4.2%; 95% CI: 2.8–5.7) or

Kenya (6.7%; 95% CI: 5.0–8.6). Sensitivity of self-reported (versus measured) hypertension was less than 45% in all countries, with specificities more than 89%. Positive likelihood ratios were fair in the Gambia (3.70; 95% CI: 2.47–5.54), and good in Kenya (5.79; 95% CI: 3.36–9.98) and Mozambique (5.18; 95% CI: 2.56–10.46). All negative likelihood ratios were poor (≥ 0.20).

Lessons learnt: Self-reported hypertension is unsuitable for population hypertension estimates among women of reproductive age in these countries.

Introduction

Hypertension is a leading and increasingly prevalent risk factor for cardiovascular disease. Accurate estimates of the prevalence of hypertension are needed for monitoring and evaluation of existing policies and programmes, to address cardiovascular disease. In sub-Saharan Africa, this information is sometimes collected through self-report in surveys. While the reliability of self-reported hypertension varies (e.g. by age, region and sex),¹ the reliability in sub-Saharan Africa is unknown.

We sought to: (i) report the prevalence of hypertension in non-pregnant women of reproductive age, for whom cardiovascular disease prevention has particular potential; (ii) investigate the reliability of self-reported (versus measured) hypertension in such women; and (iii) compare self-reported hypertension prevalence between non-pregnant women of reproductive age and pregnant women in the Gambia, Kenya, Mozambique, within the PRECISE (PREgnancy Care Integrating translational Science, Everywhere), observational prospective cohort.²

Local setting

Coordinated by King's College London, England, PRECISE was conducted in Kenya, Mozambique and the Gambia. PRECISE is an 8.8 million pounds sterling (£) project, with £2.0 million allocated for recruitment of women of reproductive age and completion of the first PRECISE visit.

PRECISE Gambia is led by the Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine at one urban (Farafenni District Hospital; comprehensive maternity services) and two rural sites (Illiasa and Ngeyen Sanjal clinics; essential maternity services) in Farafenni District.

PRECISE Kenya is led by the Aga Khan University in two sites in Kilifi County: one urban (Mariakani Subcounty Hospital; comprehensive maternity services) and one rural (Rabai Subcounty Hospital; essential maternity services).

PRECISE Mozambique is led by the Centro de Investigação de Saúde de Manhiça, in two sites in Maputo Province: one urban (Manhiça District Hospital; comprehensive maternity services) and one rural (Xinavane Rural Hospital; comprehensive maternity services),

Approach

We identified non-pregnant women of reproductive age (15–49 years) from family planning clinics (Mozambique, Kenya), or by random sampling from the local health and demographic surveillance system (the Gambia). Pregnant women of reproductive age were recruited during antenatal care. All provided written, informed consent. The study was approved by the Research Ethics Boards at King's College London, United Kingdom (HR-17/18–7855) and the University of British Columbia (H18–02828), and within each country (2018/REC-74, Kenya; 545/CNBS/18, Mozambique; SCC 1619, the Gambia).

To determine self-reported hypertension, women were asked, “Have you ever been told by a doctor or other health worker that you have high blood pressure?” Then, blood pressure was measured at least twice using a semi-automated oscillometric device,³ after 5 minutes' rest, having removed restrictive arm clothing and checking cuff size and patient position (feet on floor, arm at heart level). For each systolic and diastolic blood pressure measurement, when the first and second readings were within 10 mmHg, we used the average. If the first and second readings were > 10 mmHg different, a third reading was taken and we used the average of the second and third readings.⁴ We defined hypertension as systolic blood pressure \geq 140 mmHg or diastolic blood pressure \geq 90 mmHg.⁵ We referred women with hypertension for relevant care.

For measured and self-reported hypertension, we used multivariable logistic regression to estimate country-specific hypertension prevalence, after adjustment for age, body mass index (BMI), basic education, parity (nulliparous or parous), antihypertensive medicine use (any or none) and oral contraceptives use in the preceding 12 months (before pregnancy in pregnant women of reproductive age). We adjusted country prevalence of hypertension using model-predicted probabilities averaged over adjustment factors. These values are equivalent to the expected prevalence per country, assuming each had the same baseline characteristics for which

adjustment was made. We calculated confidence intervals (CIs) by non-parametric bootstrapping, estimated by the delta method, given sparse data and convergence in some bootstrap samples.

Among the non-pregnant women, we assessed accuracy of self-reported versus measured hypertension by calculating: (i) sensitivity (proportion who self-reported hypertension of women with measured hypertension); (ii) specificity (proportion who self-reported no hypertension, among women with normal measured blood pressure); and (iii) positive and negative likelihood ratios. The positive likelihood ratio indicates how much the odds of measured hypertension increase when hypertension is self-reported, calculated as $(\text{sensitivity})/(1-\text{specificity})$. The negative likelihood ratio indicates how much the odds of measured hypertension decrease when hypertension is self-reported, calculated as $(1-\text{sensitivity})/(\text{specificity})$. A good negative likelihood ratio is < 0.20 and a good positive likelihood ratio is ≥ 5.0 .⁶⁻⁸

Within each country, we compared the prevalence of self-reported hypertension between the non-pregnant and pregnant women by calculating risk difference and 95% bootstrap CIs.

We used R, version 4.2.1 (R Foundation, Vienna, Austria) for analyses.

Relevant changes

From June 2019 to December 2022, we recruited 1825 non-pregnant women of reproductive age and 6770 pregnant women (respectively, 609 and 3450 in Kenya; 606 and 2097 in Mozambique; and 610 and 1223 in the Gambia). There were between-country differences in baseline characteristics for both non-pregnant and pregnant women (Table 1). The non-pregnant women of reproductive age were in their late twenties, while pregnant women were in their early-to-mid-twenties. Non-pregnant and pregnant Gambian women had lower BMI, with a quarter of each group of women being underweight. Most Kenyan and Mozambican women had at least primary education, whereas most Gambian women had no basic education. Few women reported a history of kidney disease or diabetes. Oral contraceptive use in the preceding 12 months was low in the Gambia (3 women; 0.5%) and Kenya (12 women; 2.0%), but more than 10% in Mozambique (98 women; 16.2%).

Among the non-pregnant women of reproductive age, adjusted prevalence rates for measured hypertension were higher in the Gambia (9.3%; 95CI %: 6.6 to 12.6) and Mozambique

(10.4%; 95% CI: 7.9 to 12.7) than Kenya (4.6%; 95% CI: 3.0 to 6.6; Table 2). For self-reported hypertension, adjusted prevalence rates were higher in the Gambia (12.9%; 95% CI: 10.2 to 15.9) than Kenya (6.7%; 95% CI: 5.0 to 8.6) and Mozambique (4.2%; 95% CI: 2.8 to 5.7). Within each country, self-reported hypertension had low sensitivity ($< 45\%$) and high specificity ($\geq 89\%$) compared with measured hypertension. In all countries, the negative likelihood ratios were poor (above 0.65 in all countries), suggesting that self-reported hypertension cannot provide reassurance about the presence of measured hypertension. In Kenya and Mozambique, the positive likelihood ratio indicated an increased likelihood of measured hypertension with self-reported hypertension. The pattern was consistent across baseline characteristics of age, BMI and basic education; due to small sample sizes, we could not examine the effect on sensitivity and specificity of parity, use of antihypertensives or use of oral contraceptives.

Among pregnant women, the adjusted prevalence of self-reported hypertension was lowest in Mozambique (2.7%; 95% CI: 1.9 to 3.5) and highest in the Gambia (12.9%; 95% CI: 10.9 to 15.4), as it was also for the non-pregnant women of reproductive age (Table 2). Adjusted rates of self-reported hypertension did not differ between the non-pregnant women of reproductive age and pregnant women.

Lessons learnt

Our study showed that measured hypertension prevalence among non-pregnant women of reproductive age, after adjusting for participant characteristics, were higher in Mozambique and the Gambia than Kenya, and lower than published estimates, –about 30.0% in Mozambique, 17.0% in the Gambia and 11.0% in Kenya.^{9–11} However, the populations recruited in PRECISE may not be representative of national populations. The women in our study were recruited from selected districts and in different ways. In the Gambia, for example, a quarter of the women were underweight (versus 7.0% at national level),⁹ and $< 1\%$ had known risk factors for hypertension (other noncommunicable diseases). In Kenya, national hypertension prevalence rates are not disaggregated by age, so rates could be higher than in our study. Mozambican women in our study are particularly active, with manual labour jobs (e.g. sugar cane farm work), hence our sample may be different from a national one.^{12,13} As most women were parous and few women were taking antihypertensives or oral contraceptives, we could not adjust for these factors. Therefore, further investigation is needed in nationally representative samples.

We assessed the accuracy of self-reported hypertension versus measured hypertension. Despite the simplicity and widespread use of self-reported hypertension in sub-Saharan Africa, our results show that this method is insensitive for detecting measured hypertension and unsuitable for population hypertension estimates among women of reproductive age (Box 1).

Our findings align with a systematic review that found that self-reporting underestimated hypertension prevalence.¹ While less than half of people with hypertension would have been diagnosed based on self-report (sensitivity: 42.1%; specificity: 89.5%), there was substantial variation in diagnostic accuracy of self-reported (versus measured) hypertension across countries and age groups.¹ Only one study included a sub-Saharan Africa country (Ghana), in which self-report performed poorly (sensitivity: 13.0%; specificity: 97.0%) in a population with mean age of 60 years.¹ Other studies have found reliability of self-reported hypertension to be context-specific.^{6,7}

Our findings have implications for health and demographic surveillance systems, and provide evidence in young African women to support the approach recommended in the Global Monitoring Framework for noncommunicable diseases (STEPwise), to include blood pressure measurement by standardized methods, alongside self-reporting (Box 1).¹⁴ Of note, our blood pressure measurements reflected population screening recommendations, which differ from clinical practice guidelines for diagnosis of hypertension in individuals (that is, repeat blood pressure measurement on more than two separate occasions).⁵

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Competing interests:

None declared.

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Box 1. Summary of main lessons learnt

- We found that self-reported hypertension is specific, but not sensitive, for measured hypertension, making self-report an inadequate screening test.
- Our findings support the World Health Organization's STEPwise approach to noncommunicable disease risk-factor surveillance,¹⁴ which is to include self-reported and biomedical measurements of blood pressure.

Table 1. **Baseline characteristics of the women of reproductive age, by pregnancy status, the Gambia, Kenya and Mozambique, June 2019 to December 2022**

Characteristic	Not pregnant			Pregnant		
	Gambia (n = 610)	Kenya (n = 609)	Mozambique (n = 606)	Gambia (n = 1223)	Kenya (n = 3450)	Mozambique (n = 2097)
Age in years, median (IQR)	28.0 (22.0–36.0)	27.0 (23.0–33.0)	28.0 (23.0–34.8)	26.0 (22.0–31.0)	26.0 (23.0–31.0)	23.0 (19.0–29.0)
Missing	11 (NA)	0 (NA)	0 (NA)	4 (NA)	1 (NA)	0 (NA)
BMI in kg/m², no. (%)						
< 18.5	153/609 (25.1)	53/601 (8.8)	22/597 (3.7)	163/1209 (13.5)	143/3386 (4.2)	39/2070 (1.9)
18.5–24.9	328/609 (53.9)	373/601 (62.0)	357/597 (59.8)	770/1209 (63.7)	1827/3386 (54.0)	1154/2070 (55.7)
25.0–29.9	85/609 (14.0)	120/601 (20.0)	150/597 (25.1)	202/1209 (16.7)	894/3386 (26.4)	652/2070 (31.5)
≥ 30.0	43/609 (7.0)	55/601 (9.2)	68/597 (11.4)	74/1209 (6.1)	522/3386 (15.4)	225/2070 (10.9)
Missing	1 (NA)	8 (NA)	9 (NA)	14 (NA)	64 (NA)	27 (NA)
Religion, no. (%)						
Christian	8/609 (1.3)	379/606 (62.5)	550/605 (90.9)	8/1222 (0.7)	2082/3436 (60.6)	2042/2095 (97.5)
Muslim	601/609 (98.7)	222/606 (36.6)	11/605 (1.8)	1214/1222 (99.3)	1343/3436 (39.1)	27/2095 (1.3)
Traditional, spiritualist, animist	0/609 (0.0)	1/606 (0.2)	35/605 (5.8)	0/1222 (0.0)	4/3436 (0.1)	11/2095 (0.5)
Buddhist	0/609 (0.0)	0/606 (0.0)	0/605 (0.0)	0/1222 (0.0)	0/3436 (0.0)	2/2095 (0.1)
Other	0/609 (0.0)	0/606 (0.0)	5/605 (0.8)	0/1222 (0.0)	1/3436 (0.0)	3/2095 (0.1)
None	0/609 (0.0)	4/606 (0.7)	4/605 (0.7)	0/1222 (0.0)	6/3436 (0.2)	10/2095 (0.5)
Missing	1 (NA)	3 (NA)	1 (NA)	1 (NA)	14 (NA)	2 (NA)
Education^a, no. (%)						
None	373/610 (61.1)	62/606 (10.2)	65/606 (10.7)	781/1222 (63.9)	322/3437 (9.4)	109/2097 (5.2)
Primary	71/610 (11.7)	367/606 (60.6)	266/606 (43.9)	193/1222 (15.8)	1812/3437 (52.7)	704/2097 (33.6)
Secondary	119/610 (19.5)	116/606 (19.1)	271/606 (44.7)	192/1222 (15.7)	895/3437 (26.0)	1258/2097 (60.0)
Higher	47/610 (7.7)	61/606 (10.1)	4/606 (0.7)	56/1222 (4.6)	408/3437 (11.9)	26/2097 (1.2)
Missing	0 (NA)	3 (NA)	0 (NA)	1 (NA)	13 (NA)	0 (NA)
Parity, no. (%)						
Nulliparous	175/610 (28.7)	39/609 (6.4)	60/606 (9.9)	234/1223 (19.1)	990/3450 (28.7)	828/2097 (39.5)
Parous	435/610 (71.3)	570/609 (93.6)	546/606 (90.1)	989/1223 (80.9)	2460/3450 (71.3)	1269/2097 (60.5)
Self-reported hypertension, no. (%)						
No	522/608 (85.9)	551/603 (91.4)	569/606 (93.9)	1063/1220 (87.1)	3202/3429 (93.4)	2044/2095 (97.6)
Yes	83/608 (13.7)	49/603 (8.1)	29/606 (4.8)	153/1220 (12.5)	213/3429 (6.2)	44/2095 (2.1)
Don't know	3/608 (0.4)	3/603 (0.5)	8/606 (1.3)	4/1220 (0.4)	14/3429 (0.4)	7/2095 (0.3)

Missing	2 (NA)	6 (NA)	0 (NA)	3 (NA)	21 (NA)	2 (NA)
Taking	4/83 (4.8)	9/49 (18.4)	1/29 (3.4)	6/153 (3.9)	15/213 (7.0)	0/44 (0.0)
antihypertensive						
medicine, no. (%)						
Chronic kidney disease, no. (%)						
No	605/608 (99.5)	592/603 (98.2)	591/606 (97.5)	1207/1220 (98.9)	3375/3428 (98.5)	2085/2096 (99.5)
Yes	2/608 (0.3)	6/603 (1.0)	1/606 (0.2)	7/1220 (0.6)	29/3428 (0.8)	1/2096 (0.0)
Don't know	1/608 (0.2)	5/603 (0.8)	14/606 (2.3)	6/1220 (0.5)	24/3428 (0.7)	10/2096 (0.5)
Missing	2 (NA)	6 (NA)	0 (NA)	3 (NA)	22 (NA)	1 (NA)
Diabetes, no. (%)						
No	599/608 (98.5)	595/603 (98.6)	599/605 (99.0)	1208/1220 (99.0)	3380/3429 (98.5)	2087/2095 (99.7)
Yes	7/608 (1.2)	4/603 (0.7)	0/605 (0.0)	8/1220 (0.7)	30/3429 (0.9)	3/2095 (0.1)
Do not wish to	2/608 (0.3)	4/603 (0.7)	6/605 (1.0)	4/1220 (0.3)	19/3429 (0.6)	5/2095 (0.2)
answer						
Missing	2 (NA)	6 (NA)	1 (NA)	3 (NA)	21 (NA)	2 (NA)
Oral	3/609 (0.5)	12/609 (2.0)	98/606 (16.2)	11/1222 (0.9)	61/3449 (1.8)	256/2097 (12.2)
contraceptive use						
in previous 12						
months^b, no. (%)						
Missing	1 (NA)	0 (NA)	0 (NA)	1 (NA)	1 (NA)	0 (NA)

BMI: body mass index; IQR: interquartile range; NA: not applicable.

^a We defined basic education as any formal schooling at primary, secondary or post-secondary school-level

^b Among unselected pregnant women, this use was the 12 months before pregnancy.

Table 2. **Measured and self-reported hypertension among women of reproductive age, the Gambia, Kenya and Mozambique, June 2019 to December 2022**

Outcome	Gambia	Kenya	Mozambique
Non-pregnant women	<i>n</i> = 610	<i>n</i> = 609	<i>n</i> = 606
Measured hypertension ^a , % (95% CI)	9.3 (6.6 to 12.6) ^b	4.6 (3.0 to 6.6) ^b	10.4 (7.9 to 12.7)
Self-reported hypertension, % (95% CI) ^a	12.9 (10.2 to 15.9) ^b	6.7 (5.0 to 8.6) ^b	4.2 (2.8 to 5.7)
Pregnant women	<i>n</i> = 1223	<i>n</i> = 3450	<i>n</i> = 2097
Self-reported hypertension ^a , % (95% CI)	12.9 (10.9 to 15.4) ^b	6.0 (5.3 to 6.8)	2.7 (1.9 to 3.5) ^b
Non-pregnant women of reproductive age: self-reported versus measured hypertension			
Sensitivity, % (95% CI)	41 (28 to 56)	38 (21 to 56)	17 (9 to 29)
Specificity, % (95% CI)	89 (86 to 91)	94 (91 to 95)	97 (95 to 98)
Negative likelihood ratio (95% CI)	0.66 (0.53 to 0.83)	0.67 (0.51 to 0.87)	0.86 (0.77 to 0.96)
Positive likelihood ratio (95% CI)	3.70 (2.47 to 5.54)	5.79 (3.36 to 9.98)	5.18 (2.56 to 10.46)
Self-reported hypertension: non-pregnant women of reproductive age versus pregnant women			
Adjusted risk difference, % (95% CI)	0.0 (−3.5 to 3.3)	0.6 (−1.3 to 2.9)	1.4 (−1.6 to 2.9)

CI: confidence interval.

^a We adjusted hypertension prevalence for age, body mass index, basic education, parity, antihypertensive medication and history of taking oral contraceptives in the previous 12 months (and specifically before pregnancy for pregnant women).

^b Data were missing for three women in Kenya and two in the Gambia.