

Prevalence and associated factors of hypokalemia in hypertension: the perspective in a low to middle-income setting

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Background: Despite being a common feature that can be present either as a complication of treatment or a sign of primary aldosteronism, the true prevalence of hypokalemia has not been enough documented in a low to middle-income setting. This study sought to determine the prevalence and associated factors of hypokalemia at the outpatient department of the Douala General Hospital (DGH), Cameroon.

Methods: Records obtained from the hypertension registry of the DGH were reviewed and those with confirmed white coat hypertension were excluded from the study. Univariate and multivariate logistic regression was used to determine factors associated with hypokalemia in hypertension.

Results: A total of 687 records (62.1% females, mean age 57.5±12.9 years) were included in the study. One hundred and eighty (26.2%) records had hypokalemia. Participants on diuretics were more likely to develop hypokalemia, and those receiving indapamide or chlorthalidone (OR =5.7, P<0.001 and OR =4.8, P=0.02 respectively) were more likely to have hypokalemia than those on hydrochlorothiazide (OR =1.9, P=0.01).

Conclusions: Hypokalemia was very frequent amongst our patients with hypertension but not systematically well investigated. Diuretics prescription was the main associated factor. These findings have implications for the investigation and management of individuals with hypertension in our setting.

Keywords: Electrolytes; hypertension; hypokalemia; potassium; prevalence

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Introduction

Hypertension is a major cardiovascular risk factor accounting for about 50% of cardiovascular deaths globally (1,2). Compared to high-income countries, the rate of hypertension has been on the rise in low to middle-income settings (3). Effective treatment of this condition warrants adequate lifestyle modifications as well as pharmacologic

management (4,5). Pharmacologic management of hypertension has well-established adverse effects of which electrolyte disturbance are amongst. Hypokalemia defined as serum potassium less than 3.5 mmol/L (6-8) is a well-established adverse effect of pharmacologic management of diuretic therapy (9,10) which remains the cornerstone in the management of hypertension (4,11). Moreover,

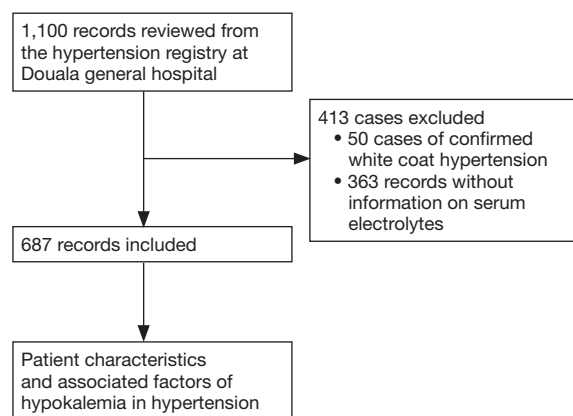


Figure 1 Flow chart representing recruitment of participants (1,100 records were reviewed and 687 were included in the study).

hypokalemia in hypertension can as well be present in specific etiologies of hypertension with the most common being primary aldosteronism (12-14), with recent endocrine society guidelines recommending screening for primary aldosteronism in any case of hypertension presenting with spontaneous or diuretic-induced hypokalemia (12,13).

Several studies have demonstrated the relationship between potassium and blood pressure (BP) (7,15-19) and although serum potassium is not an absolute predictor of total body potassium, studies have determined a U-shaped relationship between cardiovascular outcomes and serum potassium (8,20-22). Compared to Caucasian populations, studies have reported salt sensitivity in the black populations, these studies further demonstrated blunting of the sodium effect on BP once potassium intake is increased (23). Furthermore, data reports low urinary potassium excretion in blacks when compared to Caucasians regardless of potassium intake (24,25).

To our knowledge, most studies reporting the prevalence of hypokalemia in hypertension were carried out in high-income countries, with these studies revealing the prevalence of hypokalemia in hypertension between 6–21% (26,27). Thus, we carried a study to measure the prevalence of hypokalemia in hypertension as well as the factors associated with hypokalemia in hypertension in a low to middle-income setting.

Methods

Study design and setting

This was a 5-year retrospective (January 2014–January 2019)

observational study conducted at the outpatient department of Douala General Hospital (DGH) which is a tertiary hospital located in the urban city of Douala. Data were obtained from the DGH hypertension registry, an ongoing registry created by cardiologists of the DGH since 2010.

Records with confirmed hypertension by an attending cardiologist (based on office BP $\geq 140/90$ mmHg or 24-hour ambulatory BP $\geq 130/80$ mmHg) (28) were included, while records with confirmed white coat hypertension (records with office BP $\geq 140/90$ mmHg with 24-hour ambulatory BP $< 130/80$ mmHg) (28), and records with no information on serum electrolyte were excluded from the study.

Data collection

Administrative authorization was obtained from the central administration of the DGH and ethical clearance from the Faculty of Health Sciences University of Bamenda, Cameroon.

Blood pressure: BP was obtained from patient records which were measured following the European Society of Cardiology (ESC) 2013 guidelines (28) and the average BP for each patient record was obtained by summing the second and third BP reading in each record and dividing by two.

Serum potassium and other laboratory investigations were obtained from patient records, information concerning usage of antihypertensive medications was obtained during the first consultation and those reported to be newly diagnosed with hypertension were considered not to be on any medications. Hypokalemia was considered as serum potassium < 3.5 mmol/L. Mild, moderate and severe hypokalemia was defined as serum potassium between 3–3.4, 2.5–2.9 and < 2.5 mmol/L respectively. *Figure 1* summarizes the steps for the recruitment of study participants.

Statistical analyses

Data obtained from patient records were analyzed using SPSS version 25 for windows, patients were classified into three groups based on serum potassium. Categorical variables were presented as frequencies and percentages while continuous variables were presented as mean and standard deviation. Categorical variables were compared using the Chi-squared (χ^2) test and Fischer exact test whereas one-way analysis of variance was used to compare continuous variables, Brown-Forsythe correction was used to adjust P values for continuous variables which violated the Levene's test for homogeneity of variance. Univariate and

Multivariate logistic regression was used to determine the associated factors of hypokalemia in hypertension with the calculation of odds ratio (OR) and 95% confidence intervals (CIs). A P value of less than 0.05 was considered significant.

Results

Patient characteristics

In total, 687 participants were included in the study. The demographic and clinical characteristics of these participants are summarized in *Table 1* based on three levels of serum potassium. Overall 62% [427] of the participants were females with a mean age of 57.5 ± 12.9 years. The mean systolic BP (SBP) and diastolic BP (DBP) were 163.8 ± 24.0 and 95.8 ± 16.0 mmHg respectively with 65% of participants reported having either grade 1 or grade 2 hypertension based on the ESC classification. Also, 74% of participants reported being on BP-lowering medications. The mean serum potassium was 3.9 ± 0.55 mmol/L with 71% of participants reported to have serum potassium between 3.5–5.0 mmol/L.

Prevalence of hypokalemia in hypertension

Of the 687 records reviewed, 180 had serum potassium less than 3.5 mmol/L thus the prevalence of 26.2% (95% CI, 22.9–29.7%). Out of these 180 cases, 82.8% were cases of mild hypokalemia, 16.7% had moderate hypokalemia and 0.6% were cases of severe hypokalemia.

Factors associated with hypokalemia

On univariate analysis, use of calcium channel blockers (CCBs) (OR = 1.57, $P < 0.05$) was associated with hypokalemia, also use of diuretics was associated with hypokalemia with use of indapamide (OR = 5.7, $P < 0.001$) associated more with hypokalemia than hydrochlorothiazide (OR = 1.6, $P = 0.01$). Female sex had a decreased likelihood to present with hypokalemia (OR = 0.93, $P = 0.70$) but this was however not statistically significant. All variables used for univariate analysis were entered into the multivariate model, the use of diuretics was significantly associated with hypokalemia, with participants using thiazide-like diuretics (indapamide and chlorthalidone) presenting higher chances of hypokalemia than those using of thiazide-type diuretics (hydrochlorothiazide). *Table 2* summarizes the results obtained from univariate and multivariate logistic regression.

Discussion

In this study, we demonstrated a high prevalence of hypokalemia (26.2%) in a black African hypertensive population, with use of diuretics independently associated with hypokalemia. Furthermore, participants receiving thiazide-like (indapamide and chlorthalidone) diuretics were more likely to present with hypokalemia than those on thiazide-type (hydrochlorothiazide) diuretics. To our knowledge, this is the first study reporting the prevalence of hypokalemia in hypertension in the general hypertensive population in a sub-Saharan African setting.

Prevalence of hypokalemia in hypertension

Most studies reporting the prevalence of hypokalemia in hypertension are focused on the diagnosis of primary aldosteronism with most of the participants presenting with specific characteristics like resistant hypertension or severe hypertension (26,27). This study included all categories of patients thus representing a general hypertensive cohort. The prevalence of hypokalemia in this study was 26.2%, similar to that observed by Douma *et al.* (26). Although the Douma series included only participants with resistant hypertension nevertheless, Monticone and colleagues reported a lower prevalence (6.3%) (27). Notably, this was a study to report the prevalence of primary aldosteronism and antihypertensive medications in most participants were withheld for 4–6 weeks prior to measurement of serum electrolytes and aldosterone renin ratio.

Factors associated with hypokalemia

Diuretics are strongly recommended by studies and hypertension guidelines as first-line agents in the treatment of hypertension, especially in blacks (4,5,29). In addition, studies have established superiority of thiazide-like diuretics (chlorthalidone and indapamide) over thiazide-type diuretics (hydrochlorothiazide) with regards to BP-lowering and cardiovascular risk protection, with similar effects on metabolic profile (serum electrolytes and serum glucose) (11,30,31). Nonetheless, thiazide diuretics are associated with hypokalemia (9). This study revealed diuretics were independently associated with hypokalemia in hypertension and on further analyses, chlorthalidone and indapamide were more associated with hypokalemia than hydrochlorothiazide. These results are in agreement with findings of a systematic review conducted by Dorsch *et al.*

Table 1 Patient characteristics by serum potassium level

Variables	K <3.5 mmol/L, n=180	K 3.5–5 mmol/L, n=488	K >5 mmol/L, n=19	P value
Female	114 (63.3)	302 (61.9)	11 (57.9)	0.88
Age (years), (SD)	57.67 (12.63)	57.45 (13.02)	58.42 (13.50)	0.94
Risk factors				
Diabetes mellitus	19 (10.6)	53 (10.9)	2 (10.5)	0.99
Heart failure	7 (3.9)	13 (2.7)	0	0.53
Tobacco	4 (2.2)	14 (2.9)	1 (5.3)	0.72
Signs and symptoms				
Palpitations	23 (12.8)	78 (15.9)	1 (5.3)	0.29
Clinical parameters				
Weight (kg)	80.06 (14.20)	80.54 (15.02)	77.42 (15.12)	0.64
Heart rate (bpm)	79.39 (14.36)	79.58 (14.49)	77.21 (17.43)	0.78
SBP (mmHg)	168.08 (25.63)	163.94 (22.99)	185.05 (24.83)	<0.001
DBP (mmHg)	95.19 (16.42)	95.84 (15.78)	100.13 (16.21)	0.44
Laboratory parameters				
Sodium (mmol/L)	137.58 (6.55)	139.13 (6.91)	142.11 (11.34)	0.004
Creatinine (mg/dL)	1.02 (0.37)	1.15 (1.0)	2.16 (2.48)	0.05
eGFR (mL/min/1.73 m ²)	87.56 (26.63)	88.90 (29.60)	72.61 (36.16)	0.06
LDL (g/L)	1.28 (0.43)	1.36 (0.46)	1.23 (0.62)	0.32
Total cholesterol (g/L)	1.96 (0.49)	2.10 (0.50)	1.95 (0.79)	0.02
Serum glucose (mg/dL)	99.61 (25.38)	99.36 (37.49)	105.43 (25.75)	0.81
Medications				
On blood pressure lowering medications	155 (86.1)	342 (70.1)	14 (73.9)	<0.001
Beta blockers	14 (7.7)	46 (9.4)	1 (5.3)	0.32
CCBs	91 (50.6)	193 (39.5)	7 (36.8)	0.77
ACE inhibitors	77 (42.8)	189 (38.7)	8 (42.1)	0.51
ARBs	20 (11.1)	38 (7.8)	0	0.33
Diuretics	128 (71.1)	227 (46.5)	6 (31.6)	<0.001
ACEi + diuretics	71 (39.4)	96 (19.8)	2 (10.5)	<0.001

ACEi, angiotensin converting enzyme inhibitor; ARBs, angiotensin receptor blockers; bpm, beats per minute; DBP, diastolic blood pressure; eGFR, estimated glomerular filtration rate; K, serum potassium; LDL, low density lipoproteins; SBP, systolic blood pressure; CCBs, calcium channel blockers.

where chlorthalidone was associated with more hypokalemia than hydrochlorothiazide (11). However, they are somehow different from those observed in other systematic reviews, which included studies conducted almost exclusively in Caucasians and where there was no difference between

thiazide-like and thiazide-type diuretics on serum potassium levels (11,30). These contrasting findings can be explained by racial or nutritional differences. It shall be noted that some studies have shown that urinary potassium excretion is less in blacks than Caucasians even when identical amounts

Table 2 Factors associated with hypokalemia in hypertension (univariate and multivariate logistic regression)

Variables	Univariate analysis		Multivariate analysis	
	OR (95% CI)	P value	aOR (95% CI)	P value
Female sex	0.93 (0.66–1.33)	0.70	1.11 (0.77–1.62)	0.57
Age (years)	1.00 (0.98–1.01)	0.87	0.99 (0.98–1.01)	0.92
Hydrochlorothiazide	1.6 (1.22–2.84)	0.004	1.92 (1.15–3.22)	0.01
Chlorthalidone	4.38 (1.29–14.87)	0.02	4.82 (1.32–17.52)	0.02
Indapamide	5.70 (3.59–9.07)	<0.001	5.89 (3.52–9.87)	<0.001
Furosemide	1.75 (0.46–6.68)	0.41	2.17 (0.54–8.78)	0.27
CCBs (yes)	1.57 (1.12–2.21)	0.01	1.16 (0.79–1.70)	0.44
Beta blockers (yes)	0.83 (0.44–1.54)	0.55	0.79 (0.39–1.57)	0.50
ARBs (yes)	1.54 (0.87–2.73)	0.14	1.66 (0.85–3.27)	0.14
ACE inhibitors (yes)	1.18 (0.83–1.66)	0.34	0.85 (0.53–1.34)	0.48
Fasting plasma glycemia >100 (mg/dL)	1.36 (0.95–1.95)	0.09	1.26 (0.86–1.85)	0.24
eGFR <60 (mL/min/1.73 m ²)	0.93 (0.57–1.53)	0.77	0.84 (0.49–1.44)	0.52

ACE, angiotensin converting enzyme; ARBs, angiotensin receptor blockers; aOR, adjusted odds ratio; CCBs, calcium channel blockers; CI, confidence interval; eGFR, estimated glomerular filtration rate; OR, odds ratio.

of potassium are provided in the diet (24,25). However, because the urinary potassium excretion also depends on sodium intake and diet, we cannot rule out the possibility that a low salt diet which is largely prescribed by physicians in our setting might have influenced plasma potassium levels in our population. Also, potassium depletion has been reported to increase mean arterial BP (32), through pressor effects and alteration of vascular resistance, in addition, it is as well reported that potassium depletion increases urinary calcium excretion which has been stipulated by several studies to have positive pressor effects. Finally, that a large proportion of our population was on angiotensin converting enzyme (ACE) inhibitors or angiotensin receptor blockers (ARBs) could have resulted in less potassium excretion due to a synergistic effect of these medications with diuretics. This has been particularly advocated and demonstrated in populations with low-renin hypertension (33,34). Patients with essential hypertension especially those with resistant hypertension are generally reported to have high plasma renin (35) which can, in turn, result in negative potassium balance by subsequent activation of the renin aldosterone angiotensin pathway. However, studies have reported black patients often present with low renin hypertension (36). It is worth noting that hypertension in blacks is salt-sensitive,

this effect is however blunted by increased potassium intake (23,37). Also, non-pharmacologic treatment of hypertension encourages patients to adopt the DASH (dietary approach to stop hypertension) diet which has been reported to effectively decrease BP, this diet is also noted for its high potassium and calcium content (38).

Limitations of the study

The present study has some limitations that must be underscored. First, its retrospective design precluded any standardizing methods of measuring serum potassium and the collection of data in all eligible patients as well as reporting the proportion of patients who were on potassium supplements which could have an impact on the prevalence of hypokalemia. Second, in the absence of randomization, there was no head to head comparison of diuretics in this study. Third, diuretics does not only reduced potassium, but also increased uric acid and increase total cholesterol and triglycerides but we did not assess other metabolic changes.

Despite these short comes, our study is one of the rare studies evaluating the prevalence and associated factors of hypokalemia in a general hypertensive cohort in a low to

middle-income setting, our findings provide baseline values for planning future research on hypertension in our setting.

Conclusions

This study revealed 1 out of 4 patients with hypokalemia, and though not randomized patients on thiazide-like diuretics (indapamide and chlorthalidone), were more likely to present with hypokalemia than those on thiazide-type diuretics (hydrochlorothiazide). More attention should be paid on serum electrolytes of patients with hypertension and consideration should be given to promoting the intake of potassium-rich foods, as this may balance the consequences of hypokalemia.

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Footnote

Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at <http://dx.doi.org/10.21037/jxym.2020.03.02>). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The protocol of this study was approved by the institutional review board of the faculty of health sciences university of Bamenda (2019/0045H/Uba/IRB), and the study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). Individual informed consent was waived due to the retrospective nature of this study.

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