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Research Article

Burden of Hypertension and Abnormal Glomerular Permeability in Hypertensive School Children

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Abstract

Background: Childhood hypertension has been associated with target-organ damage in young adults. It is often asymptomatic in both children and adolescents; when persistent, and long-standing, it could be a significant risk factor for kidney damage and increased glomerular permeability.

Objectives: Burden of hypertension and its impact on glomerular permeability were prospectively determined in randomly recruited primary school children.

Patients and Methods: Blood pressure (BP) measurement was performed by the auscultation method, and abnormal glomerular permeability was assessed by dipstick testing of urine for persistent proteinuria and/or hematuria for \geq three months in hypertensive children.

Results: Of 1,335 pupils aged 10.0 \pm 2.4 (6.0 - 14.0) years, 33 (2.5%) were hypertensive. Overall mean systolic/diastolic BP was 125.6 \pm 6.5/81.7 \pm 3.3 (range: 114.0 - 140.0/80.0 - 90.0) mmHg. Nine (27.3%) had combined systolic and diastolic hypertension, 126.7 \pm 5.7/80.0 - 80.0 \pm 0.0 (120.0 - 130.0/80.0 - 80.0) mmHg. Isolated systolic hypertension, 125.4 \pm 6.7 (114.0 - 140.0) mmHg, was present in 14 (42.4%), whereas 10 (30.3%) had isolated diastolic hypertension, 82.0 \pm 3.5 (80.0 - 90.0) mmHg. Mean systolic and diastolic BP were 131.0 \pm 3.3 (130.0 - 140.0) mmHg and 86.5 \pm 4.43 (80.0 - 90.0) mmHg, respectively. According to the dipstick test, none of the hypertensive pupils showed urinalysis evidence of proteinuria and/or hematuria after three months of testing.

Conclusions: Although the burden of hypertension was 2.5%, the dipstick method did not detect any hypertension-related abnormal glomerular permeability in the school children.

Keywords: Hypertension, Glomerular Permeability, Proteinuria, School Children

1. Background

Hypertension (HTN) prevalence in the normal pediatric and adolescent population ranges from 3.2% to 4.7% (1-4). Childhood HTN has been linked to target-organ damage in young adults (5). HTN is often asymptomatic in both children and adolescents; when persistent and longstanding, it could be a significant risk factor for kidney damage and increased glomerular permeability (6, 7). Proteinuria and/or urinary red blood cells (RBCs) are important urinalysis manifestations of kidney damage and increased glomerular permeability. Such evidence of bilateral kidney damage is an indication of chronic kidney disease if protracted for \geq three months and may progress to end-stage renal disease (ESRD) if not recognized and treated early (6). Renal replacement therapy, the standard treatment for ESRD, is rarely available and affordable for many patients in developing countries (8-11). It was postulated that hypertension-related kidney damage is caused by glomerular ischemia and hypoperfusion secondary to

progressive narrowing of preglomerular arteries and arterioles (12). Animal models of HTN showed that glomerular capillary HTN and hyperperfusion are the principal factors causing glomerular damage and progressive loss of kidney function (13). As a compensatory mechanism, direct transmission of high systemic blood pressure (BP) to the glomerular capillary network is physiologically blocked by an increase in afferent arteriolar resistance (14). The failure of this mechanism leads to increased glomerular capillary pressure with resultant high glomerular filtration rate, increased transglomerular passage of proteins, and mesangial influx of proteins and other macromolecules. The ultimate renal histopathology is glomerular sclerosis secondary to the activation of mesangial cells and upregulation of proinflammatory cytokines and growth factors (14).

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2. Objectives

This study prospectively determined the burden of hypertension and its impact on glomerular permeability in randomly recruited primary school children.

3. Patients and Methods

3.1. Study Area and Population

This study was conducted between June 3 and November 15, 2013, among primary-school pupils, in Ile-Ife, State of Osun, Nigeria. The study location is semiurban and has two local government areas (LGAs), namely Ife central (population, n = 167,254) and Ife East (population, n = 188,087). These LGAs cover land-mass areas of 111 km² and 172 km², respectively.

3.2. Ethics

The study was approved by the ethics and research committee of our institution and the local inspectorates of education in the two LGAs where it was carried out. Informed consent was obtained from the parent(s)/guardian(s) of each pupil. The study conformed to the Helsinki declaration of 1975 on research involving human subjects, as revised in 2000.

3.3. Sampling Technique and Research Protocol

A multistage random sampling technique was employed to select 12 schools from a list of schools in both LGAs. Six primary schools, consisting of three private and three government-owned schools, were selected from each LGA. The school register was consulted for enrollment figures by age and gender in each of the selected schools. A composite register was drawn for subjects aged 6 - 14 years whose parent(s)/guardian(s) had consented. The subjects were categorized according to their classes; a table of random numbers was used to select the pupils. Overall, a total of 1,335 pupils were recruited and studied. The research protocol was administered on each pupil for biodata and anthropometric and clinical information. A general physical examination was carried out on each pupil before the midday break (1100 hours) in a classroom.

3.4. Blood Pressure Measurement

Following the recommendation of the task force on high BP in children and adolescents (7), resting BP was determined by the auscultation method in the right arm after a 10-minute resting period using the mercury gravity sphygmomanometer with bladder cuff sizes 17.0 - 19.0 cm (length) by 7.5 - 9.0 cm (width). Onset of the first tapping sound (Korotkoff sound 1) indicated the systolic BP (SBP), whereas the point of complete disappearance of the sound (Korotkoff sound 5) indicated the diastolic BP (DBP). For each pupil, the average of two measurements was taken after an initial BP trial run to allay fear and anxiety. Pupils with high BP had their BPs checked monthly for three months in their schools; those with persistently elevated BP were thereafter referred to our pediatric nephrology clinic for further assessment and treatment. HTN was defined as SBP and/or DBP \geq 95th percentile for age, gender, and height on three different occasions (7). SBP and/or $DBP \ge 95$ th but not > 99th percentile by >5 mmHg was regarded as stage I HTN, whereas stage II HTN was defined as SBP and/or DBP > 99th percentile by more than 5 mmHg for age, gender, and height (7). BP percentile charts based on age, gender, and height developed by the task force on the diagnosis, evaluation, and treatment of high BP in children and adolescents (7) was used to determine whether a pupil was hypertensive; the same chart was used for staging of hypertension.

3.5. Anthropometric Measurement

Weight was measured using the Seca weighing machine (Seca gmbh & Co.kg, Germany). The scale was standardized daily and measured to the nearest 0.5 kg. Each child was weighed barefoot in light clothing. Height was measured with a Leicester height stadiometer (Marsden weighing machine group, UK) mounted on a vertical wall with the pupil barefoot and with head held in the Frankfort plane. The height was taken to the nearest 0.1 cm. Body mass index (BMI) was calculated as weight divided by the height squared (kg/m²). The International Obesity Task Force cutoff points of > 25 kg/m² and > 30 kg/m² for overweight and obesity, respectively, were used (15), whereas underweight and normal weight were defined using the BMI-for-age profiles developed by the National Center for Health Statistics for boys and girls aged 2 - 20 years (16).

3.6. Urinalysis

The presence of protein and/or red blood cells (RBCs) in the urine was assessed using the UriScreen Combi 10 dipstick (Yercon Diagnostic Co., Ltd., Chang Chu, China). Freshly voided urine samples were collected into plane universal bottles prior to the pupils' midday break. The test strip was dipped into the fresh urine for approximately 1 sec and then drawn across the edge of the container to remove excess urine. After 30 seconds, the test strip was compared with the color scale, and the result was immediately recorded. Color changes after 2 minutes were disregarded. Urine protein level was assessed as negative, trace, 1 + (30 mg/dL), 2 + (100 mg/dL), and 3 + (500 mg/dL). In this study, significant proteinuria by dipstick testing was defined as urinary protein \geq 30 mg/dL (\geq 1+). A dipstick value \geq ca. 5 - 10 erythrocytes/ μ L was taken as significant hematuria. Like their BP, the pupils' urine was examined monthly for significant proteinuria and/or RBCs for three months. According to the national kidney foundation-kidney/disease outcome quality initiative (6), persistent significant proteinuria and/or urinary RBCs determined by the dipstick test for three months or more is regarded as evidence of abnormal glomerular permeability.

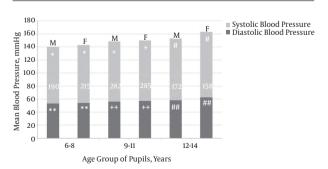
3.7. Data Analysis

Statistical analysis was performed using SPSS 16 for Windows, evaluation version (2006 SPSS Inc.). The comparative statistics used were Student's t-test, and Pearson's correlation (r). Other data were presented as numbers and percentages. A P value < 0.05 was regarded as statistically significant.

4. Results

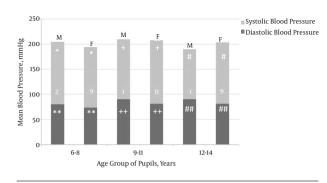
The study population included more females (n = 687; 51.5%) than males (n = 648; 48.5%). Of 1,335 (1.9%) pupils, 25 had prehypertension, and 33 (2.5%) pupils aged 6 - 14 (10.0 \pm 2.4) years were hypertensive; the majority of them were females (29/33; 88.0%), resulting in a male-to-female ratio of 1:7. The mean SBP/DBP in the hypertensive pupils was 125.6 \pm 6.5/81.7 \pm 3.3 mmHg (range: 114.0 - 140.0/80.0 - 90.0; > 95th to > 99th/ > 95th to 99th percentile). There was a weak but significant correlation between SBP (r = + 0.334, P = 0.01), DBP (r = + 0.278, P = 0.01), and age. Gender comparisons of mean SBP and DBP by age group in normotensive and hypertensive school pupils are respectively, summarized in Figures 1 and 2. Table 1 shows the number of pupils per HTN stage by age group and gender, and Table 2 summarizes the pattern and stages of HTN in the pupils.

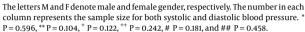
Figure 1. Gender Comparisons of Mean Systolic and Diastolic Blood Pressure in Normotensive School Pupils



The letters M and F denote male and female gender, respectively. The number in each column represents the sample size for both systolic and diastolic blood pressure. * P = 0.545, ** P = 0.218, * P = 0.545, ** P = 0.292, # P = 0.000, and ## P = 0.000.

Figure 2. Gender Comparisons of Mean Systolic and Diastolic Blood Pressure in Hypertensive School Pupils





The overall prevalence of overweight and obesity among the 1,335 pupils was 0.4% and 1.4%, respectively, but none of the hypertensive pupils was obese. HTN was, however, detected in 25 (75.8%) normal weight and seven (21.2%) underweight pupils and in one (3.0%) overweight pupil. The mean heights for the hypertensive male and female subjects were 132.4 \pm 6.91 cm and 138.80 \pm 10.44 cm, respectively (P = 0.04), and the mean BMIs were 16.12 \pm 2.11 kg/m² and 15.91 \pm 1.76 kg/m², respectively (P = 0.72). Overall, SBP(r=+0.213; P=0.01) and DBP(r=+0.148; P=0.01) correlated weakly with BMI. SBP (r = +0.434; P = 0.01) and DBP (r = +0.389; P = 0.01), however, correlated fairly well with height. Analyses for gender difference showed poor correlations between SBP, DBP, mean arterial pressure (MAP), and BMI as well as height (Table 3).

Urinalysis by dipstick revealed no evidence of abnormal glomerular permeability, because none of the hypertensive pupils had significant proteinuria and/or urinary RBCs. Trace proteinuria, however, was found in three of the 33 (9.1%) hypertensive pupils.

5. Discussion

The 2.5% prevalence of HTN among school children in this study, determined based on the fourth task-force criterion (7), is similar to the 2.2% prevalence reported among Swiss school children by Chiolero et al. (17) but is 1.3 times lower than the 3.2% prevalence reported four years earlier by Adegoke et al. (1) in the same locality. The disparity is probably due to the higher upper-age limit of 18 (range: 6 -18) years in the earlier study compared to 14 (range: 6 - 14) years in the current study. In both studies, HTN was diagnosed mostly for the first time in adolescent pupils, thus underlining the need for early screening of school children Table 1. Number of Pupils Per Hypertension Stage by Age Group and Gender

Blood Pressure (BP) by Age Group	Systolic and Diastolic Hypertension Stage	Gender		Total
		Male	Female	-
6 - 8, y				
Systolic BP, mmHg	Ι	0	6	6
Diastolic BP, mmHg	Ι	1	1	2
Systolic BP, mmHg	П	1	2	3
Diastolic BP, mmHg	П	0	0	0
9 - 11, y				
Systolic BP, mmHg	Ι	0	4	4
Diastolic BP,mmHg	Ι	0	0	0
Systolic BP, mmHg	П	0	5	5
Diastolic BP, mmHg	П	1	2	3
12 - 14, y				
Systolic BP, mmHg	I	0	3	3
Diastolic BP, mmHg	Ι	1	4	5
Systolic BP, mmHg	П	0	2	2
Diastolic BP, mmHg	П	0	0	0

Table 2. Pattern and Stages of Hypertension in the School Children

Type and Stage of Hypertension	No. (%)	Mean Blood Pressure (BP), Mmhg (Range)	BP Percentile
Isolated systolic hypertension	14 (42.4)	$125.0\pm 6.7(114.0$ - 140.0)	> 95th to 99th
Isolated diastolic hypertension	10 (30.3)	$82.0\pm3.5(80.0$ - 90.0)	95th to 99th
Combined systolic and diastolic hypertension	9 (27.3)	126.7 \pm 5.7/80.0 \pm 0.0 (120.0 - 130.0/80.0 - 80.0)	> 95th to > 99th/99th
Stage I hypertension	20 (60)		
Systolic BP		121.0 \pm 4.5 (114.0 - 130.0)	95th to 99th
Diastolic BP		80.5 ± 1.2 (80.0 - 84.0)	95th to < 99th
Stage II hypertension	13 (40)		
Systolic BP		131.0 \pm 3.3 (130.0 - 140.0)	99th to > 99th
Diastolic BP		86.5 ± 4.4 (80.0 - 110.0)	99th to > 99th

for hypertension, because target-organ damage in adulthood had been traced to childhood HTN (5, 18).

Isolated HTN was more frequent (72.7%) than combined systolic and diastolic HTN (27.3%) in this study. Furthermore, isolated systolic HTN (SHTN) prevalence (42.4%) was higher than isolated diastolic HTN (DHTN; 30.3%), confirming earlier reports that isolated HTN is not rare (19, 20). Studies by Rosner et al. (19) (SHTN, 4.4% vs. DHTN, 3.2%) and Sorof et al. (20) (SHTN, 47% vs. DHTN, 17%) revealed that isolated SHTN is more common than isolated DHTN. The pathophysiology of isolated HTN in children is not yet clear; however, studies have shown that isolated SHTN is more frequently linked with end-organ damage than is isolated DHTN (21, 22). As shown in Figure 1, the mean BP of the normotensive school children tended to increase with increasing age, especially as the children approached pubertal age. This is similar to findings in earlier studies (1, 4, 23-26). This pattern, which can be imputed to normal physiological hormonal changes that attend puberty and the increase of peripheral arterial resistance and cardiac output with age (27), was not replicated in the hypertensive children. In the latter, BP rise was haphazard and failed to correlate with rising age (Figure 2). This suggests that hypertension is associated with a defective normal BP regulatory Table 3. Correlation of Blood Pressure With Body Mass Index and Height by Gender in Hypertensive and Normotensive Pupils

Blood Pressure (BP)		Correlation Coefficient, R				
	Body M	Body Mass Index				
	Male	Female	Male	Female		
Hypertensive pupils ^a						
Systolic BP	- 0.015	- 0.165	+ 0.366	+0.008		
Diastolic BP	- 0.035	- 0.068	+ 0.603	+ 0.274		
MAP	- 0.035	- 0.203	+ 0.673	+ 0.315		
Normotensive pupils ^b						
Systolic BP	+ 0.207	+ 0.249	+ 0.380	+ 0.496		
Diastolic BP	+ 0.133	+ 0.178	+ 0.319	+ 0.452		
MAP	+ 0.171	+ 0.220	+ 0.970	+ 0.503		

 $^{{}^{}a}P = 0.05.$

mechanism.

The mean hypertension age was 10 years, and the majority of the hypertensive pupils had stage I HTN (60.0%). Whether they had stage I or II HTN, the female children tended to be more hypertensive. This may also, in part, be due to earlier puberty onset in females than in males. However, the very high prevalence of stage II HTN (40.0%) in this study indicates the possibility of a secondary etiology for hypertension in a good number of the pupils. In one study, 35% - 50% of hypertensive adolescents were obese, with a positive correlation established between obesity and HTN as early as 5 years of age. The study argued that obesity causes and sustains childhood essential hypertension (28). In another study, HTN prevalence was found to increase progressively with increasing BMI, and approximately 30% of overweight school children had HTN (BMI > 95th percentile) (29). Although these findings agree with those from Port Harcourt (18), a highly Westernized cosmopolitan city in Nigeria where the prevalence of overweight and that of obesity among school children were 5.7% and 5.9%, respectively, they are at variance with the findings from this study. The prevalence of overweight and that of obesity were 0.4% and 1.4%, respectively, but none of the hypertensive pupils was obese; BMI correlated poorly and inversely with BP in the hypertensive pupils in this study. In fact, the majority of the hypertensive pupils had either normal or low BMI. This is consistent with a US study that revealed high BP at low BMI among Black children, indicating a weaker effect of BMI on BP levels in Blacks (19). It is not clear why the majority of the hypertensive pupils in this study had either normal or low BMI. Perhaps an investigation of the etiology of HTN would have made the reasons for this obvious. Environmental factors, sleep disorders, low birth weight/reduced nephron number, positive family history of HTN/cardiovascular disease, and genetic disorders are a few of the factors that have been associated with HTN (30). Some congenital anomalies of the kidneys and urinary tract and even renal artery stenosis and endocrine disorders could have been missed in the children, because the study was not designed to establish the etiology of HTN. Although all of these may constitute limitations of this study, they point to future avenues of research on HTN in school children. Future studies should, therefore, focus not only on HTN prevalence but also on HTN etiology so that management plans can be more robust and definitive.

Evidence for abnormal glomerular permeability could not be established in this study, because none of the hypertensive pupils, including those with stage II HTN, had significant proteinuria and/or urinary RBCs. This may indicate that HTN was not long-standing at the time of the study or that the method used was not sensitive enough to detect abnormal glomerular permeability in HTN. Palatini et al. (31) showed that young adults with stage I HTN and hyperfiltration developed microalbuminuria three times more often than normally filtrating patients after eight years of follow-up, supporting the argument that the impact of increased BP on renal structures depends not only on high BP levels, but also on how long the BP has been elevated (14).

Trace proteinuria, a risk factor for ESRD (32), was found in 9.1% of the hypertensive pupils. The pupils with trace proteinuria who were regarded as normal in this study may have had microalbuminuria, but the convention of regarding trace proteinuria as negative has the tendency to exclude a large number of people with microalbumin-

 $^{{}^{}b}P = 0.01.$

uria (33). Trace proteinuria may serve as an important marker of microalbuminuria in both the general population and those at high risk of cardiovascular disease (33). The dipstick test of \geq +1 that is considered positive evidence of proteinuria has been reported as an unsuitable test in the general population because of its low sensitivity for urinary abnormalities (34, 35). Trace proteinuria should therefore be an indication for exclusion of microalbuminuria in the hypertensive person. The dipstick test showing trace proteinuria may therefore be a useful tool for that purpose (36-38).

5.1. Conclusion

The burden of HTN was 2.5%, with high BP commonly found at normal BMI. Young females were more frequently hypertensive than male school children, indicating the need for routine BP checks in school-aged female children. Abnormal glomerular permeability was rare in the young hypertensive school children, suggesting either recent hypertension or an insufficiently sensitive evaluation method. Testing for microalbuminuria and a urine microscopy examination for RBCs and other urine sediments might be more informative and predictive with respect to abnormal glomerular permeability in hypertension.

Footnotes

Authors' Contribution: The first and last authors conceived and designed the study and were responsible for data analysis, interpretation, drafting, and final manuscript approval. The first author collected the data. The second and third authors were involved in data analysis, interpretation, drafting, and final manuscript approval.

Conflict of Interests: None to declare. Funding/Support: None.

References

- 1. Adegoke SA, Elusiyan JBE, Olowu WA, Adeodu OO. Relationship between body mass index and blood pressure among Nigerian children aged 6-18 years. *Niger Endocrine Pract.* 2009;**3**:35–43.
- McNiece KL, Poffenbarger TS, Turner JL, Franco KD, Sorof JM, Portman RJ. Prevalence of hypertension and pre-hypertension among adolescents. J Pediatr. 2007;150(6):640–4. doi: 10.1016/j.jpeds.2007.01.052. [PubMed: 17517252].
- Hansen ML, Gunn PW, Kaelber DC. Underdiagnosis of hypertension in children and adolescents. JAMA. 2007;298(8):874–9. doi: 10.1001/jama.298.8.874. [PubMed: 17712071].
- Okoh BA, Alikor EA, Akani N. Prevalence of hypertension in primary school-children in Port Harcourt, Nigeria. *Paediatr Int Child Health*. 2012;**32**(4):208–12. doi: 10.1179/2046905512Y.0000000039. [PubMed: 23164295].

- Lurbe E, Torro I, Alvarez V, Nawrot T, Paya R, Redon J, et al. Prevalence, persistence, and clinical significance of masked hypertension in youth. *Hypertension*. 2005;45(4):493–8. doi: 10.1161/01.HYP.0000160320.39303.ab. [PubMed: 15767467].
- National Kidney F. K/DOQI clinical practice guidelines for chronic kidney disease: evaluation, classification, and stratification. *Am J Kidney Dis.* 2002;39(2 Suppl 1):S1–266. [PubMed: 11904577].
- National High Blood Pressure Education Program Working Group on High Blood Pressure in C. The fourth report on the diagnosis, evaluation, and treatment of high blood pressure in children and adolescents. *Pediatrics*. 2004;114(2 Suppl 4th Report):555-61. [PubMed: 15286277].
- Olowu WA. Renal failure in Nigerian children: factors limiting access to dialysis. *Pediatr Nephrol.* 2003;18(12):1249–54. doi: 10.1007/s00467-003-1255-5. [PubMed: 14586684].
- Esezobor CI, Oniyangi O, Eke F. Paediatric dialysis services in Nigeria: availability, distribution and challenges. West Afr J Med. 2012;31(3):181– 5. [PubMed: 23310939].
- Balaka B, Douti K, Gnazingbe E, Bakonde B, Agbere AD, Kessie K. Etiologies et pronostic de l'insuffisance renale de l'enfant a l'hopital universitaire de Lome. J Rech Sci Univ Lome (Togo). 2012;14:11-8.
- Aloni MN, Nsibu CN, Meeko-Mimaniye M, Ekulu PM, Bodi JM. Acute renal failure in Congolese children: a tertiary institution experience. *Acta Paediatr.* 2012;101(11):514–8. doi: 10.1111/j.1651-2227.2012.02827.x. [PubMed: 22931368].
- 12. Zucchelli P, Zuccala A. Primary hypertension-how does it cause renal failure?. *Nephrol Dial Transplant*. 1994;**9**(3):223–5. [PubMed: 8052423].
- Ofstad J, Horvei G, Kvam FI, Morkrid L, Sekse I, Svarstad E, et al. Glomerular hemodynamics in progressive renal disease. *Kidney Int Suppl.* 1992;36:S8-14. [PubMed: 1614073].
- Lurbe E, Alvarez V, Redon J. Predictors of progression in hypertensive renal disease in children. J Clin Hypertens (Greenwich). 2004;6(4):186– 91. [PubMed: 15073472].
- Cole TJ, Bellizzi MC, Flegal KM, Dietz WH. Establishing a standard definition for child overweight and obesity worldwide: international survey. *BMJ*. 2000;**320**(7244):1240–3. [PubMed: 10797032].
- Lahti-Koski M, Gill T. Obesity in childhood and adolescence. Pediatr Adolesc Med. 2004;9:8–10.
- Chiolero A, Cachat F, Burnier M, Paccaud F, Bovet P. Prevalence of hypertension in school children based on repeated measurements and association with overweight. *J Hypertens*. 2007;25(11):2209–17. doi: 10.1097/HJH.0b013e3282ef48b2. [PubMed: 17921814].
- Ejike CE, Ugwu C. Hyperbolic Relationship between blood pressure and body mass index in a Nigerian Adolescent Population. Webmed Cent Hypertens. 2010;1:WMC00797.
- Rosner B, Prineas R, Daniels SR, Loggie J. Blood pressure differences between blacks and whites in relation to body size among US children and adolescents. *Am J Epidemiol.* 2000;**151**(10):1007-19. [PubMed: 10853640].
- Sorof JM, Urbina EM, Cunningham RJ, Hogg RJ, Moxey-Mims M, Eissa MA, et al. Screening for eligibility in the study of antihypertensive medication in children: experience from the Ziac Pediatric Hypertension Study. *Am J Hypertens*. 2001;**14**(8 Pt 1):783–7. [PubMed: 11497194].
- Malcolm DD, Burns TL, Mahoney LT, Lauer RM. Factors affecting left ventricular mass in childhood: the Muscatine Study. *Pediatrics*. 1993;92(5):703–9. [PubMed: 8414859].
- Daniels SD, Meyer RA, Loggie JM. Determinants of cardiac involvement in children and adolescents with essential hypertension. *Circulation*. 1990;82(4):1243–8. [PubMed: 2401062].
- 23. Ayoola EA. Prevalence of adolescent hypertension in Nigeria. *NigJ Paediatr.* 1979;**6**:18–26.
- Akor F, Okolo SN, Okolo AA. Blood pressure and anthropometric measurements in healthy primary school entrants in Jos, Nigeria. SAJCH. 2010;4(2):42-5.
- 25. Antia-Obong OE, Antia-Obong IE. Arterial blood pressure of Nigeria urban-rural school children. *Nig J Paediatr.* 1991;**18**:3–11.

- Asani MO, Bode-Thomas F. Blood pressure pattern and its correlates among primary school children in Jos, Nigeria. *Highland Med Res J.* 2006;3(2):51-61.
- 27. Curran JS, Barness LA. Obesity. In: Behrman RE, Kliegman RM, Jenson HB, editors. Nelson Textbook of Pediatrics. Philadelphia: WB Saunders Company; 2000. pp. 172–6.
- Gutin B, Basch C, Shea S, Contento I, DeLozier M, Rips J, et al. Blood pressure, fitness, and fatness in 5- and 6-year-old children. *JAMA*. 1990;**264**(9):1123-7. [PubMed: 2384936].
- Sorof J, Daniels S. Obesity hypertension in children: a problem of epidemic proportions. *Hypertension*. 2002;40(4):441-7. [PubMed: 12364344].
- Awazu M. Epidemiology of Hypertension. In: Avner ED, Harmon WE, Niaudet P, Yoshikawa N, editors. Pediatric Nephrology. Berlin Heidelberg: Springer-Verlag; 2009. pp. 1459–84.
- Palatini P, Mormino P, Dorigatti F, Santonastaso M, Mos L, De Toni R, et al. Glomerular hyperfiltration predicts the development of microalbuminuria in stage 1 hypertension: the HARVEST. *Kidney Int.* 2006;**70**(3):578–84. doi: 10.1038/sj.ki.5001603. [PubMed: 16788693].
- Iseki K, Ikemiya Y, Iseki C, Takishita S. Proteinuria and the risk of developing end-stage renal disease. *Kidney Int.* 2003;63(4):1468–74. doi: 10.1046/j.1523-1755.2003.00868.x. [PubMed: 12631363].

- Konta T, Hao Z, Takasaki S, Abiko H, Ishikawa M, Takahashi T, et al. Clinical utility of trace proteinuria for microalbuminuria screening in the general population. *Clin Exp Nephrol.* 2007;11(1):51–5. doi: 10.1007/s10157-006-0458-z. [PubMed: 17384998].
- Boulware LE, Jaar BG, Tarver-Carr ME, Brancati FL, Powe NR. Screening for proteinuria in US adults: a cost-effectiveness analysis. *JAMA*. 2003;290(23):3101–14. doi: 10.1001/jama.290.23.3101. [PubMed: 14679273].
- Meyer NL, Mercer BM, Friedman SA, Sibai BM. Urinary dipstick protein: a poor predictor of absent or severe proteinuria. *Am J Obstet Gynecol.* 1994;**170**(1 Pt 1):137–41. [PubMed: 8296815].
- Soonthornpun S, Thammakumpee N, Thamprasit A, Rattarasarn C, Leelawattana R, Setasuban W. The utility of conventional dipsticks for urinary protein for screening of microalbuminuria in diabetic patients. J Med Assoc Thai. 2000;83(7):797-803. [PubMed: 10932516].
- Sam R, Shaykh MS, Pegoraro AA, Khalili V, Hristea I, Singh AK, et al. The significance of trace proteinuria. *Am J Nephrol.* 2003;23(6):438–41. [PubMed: 14583662].
- Zeller A, Sigle JP, Battegay E, Martina B. Value of a standard urinary dipstick test for detecting microalbuminuria in patients with newly diagnosed hypertension. *Swiss Med Wkly.* 2005;135(3-4):57–61. [PubMed: 15729608].