Arterial Blood Pressure in Children with Sickle Cell Anaemia and Controls: A Cross Sectional Study

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ABSTRACT

Background: Sickle cell anaemia (SCA) is a very common disorder among indigenous people of Borno and Yobe States in the North-eastern region of Nigeria. It is characterized by recurrent episodes of severe bone pain from occlusion of blood vessels by sickled red cells. Hypertension in children with sickle cell anaemia is rare and when present may be secondary to other disease process and more rarely may be essential hypertension. However, when hypertension occurs in SCA patients, it accelerates the progression of complications such as cerebrovascular accident and sickle cell nephropathy, hence the need to study the pattern of blood pressure in these patients. Early detection of hypertension can help in forestalling its progression and preventing its devastating complications such as cerebrovascular accident. **Objectives:** To compare the arterial blood pressures of children with sickle cell anaemia in steady state with those of age and sex-matched healthy controls. Methods: The study design was cross-sectional comparative. Minimum sample size of 54 was determined using Taylor's formula. Subjects were recruited systematically by enrolling every other patient as they present at the Paediatric Haematology clinic. The controls were apparently healthy age and sex-matched haemoglobin AA children. Anthropometric parameters and the blood pressure were measured and recorded appropriately. Blood pressure was classified using published normative data. Results: A total of 54 children with sickle cell anaemia in steady state and 159 normal children as controls were enrolled. The ages of both the SCA and controls ranged from 3 to 14 years, with median age of 8 years. Of the 54 SCA children 28 (52%) were males while 26 (48%) were females. Mean packed cell volume of the SCA children was 22 ± 3.5 . Although 8 (14.8%) of the SCA children were hypertensive, there was no significant difference in the mean systolic blood pressure (96mmHg) of children with sickle cell anaemia compared to controls (99mmHg) P = 0.078 while, the mean diastolic blood pressure (65mmHg) of the control was significantly higher than that of SCA cohort (59) $P \le 0.0001$. Conclusion: Hypertension is not rare among children with sickle cell anaemia. This stresses the need regular BP recording during follow-up in order to prevent devastating events such as cerebro-vascular accident.

KEYWORDS: Arterial blood pressure, hypertension, sickle cell anaemia, children, North-Eastern Nigeria.

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Introduction

Sickle cell disease (SCA) is the most common inherited blood disorder of great public health importance worldwide. Sub-Saharan Africa accounts for three-quarters of the more than 300,000 SCA children born globally every year¹. It is a multisystemic disease affecting almost all organs of the body through the combined effects of chronic infections, anaemia, recurrent chronic hypoxia and repeated infarction². This

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pathologic processes affect end organ functions with ultimate multi-organ failure³. Hypertension in children with SCA is uncommon and may be secondary to another disease process or more rarely may be essential hypertension⁴. Secondary hypertension is generally more common in children than in adults. Common causes of secondary hypertension in children include predominantly renal disease, endocrine disorders and coarctation of aorta that is amenable to surgery⁵. However, some children and adolescents may have essential hypertension in which a cause of the disease is not identifiable. Essential hypertension in children has been demonstrated to correlate positively family history with of hypertension, obesity and low birth weight.1,6,7 Several researchers have documented that systemic blood pressure is generally lower in individuals with SCA compared to age and sex matched controls with normal haemoglobin.^{4,8} They attributed these findings to the prevailing chronic hypoxic state in SCA leading to vaso dilatation and thus reduced peripheral resistance. Other factors attributable to low systemic blood pressure in children with SCA include salt loosing sickle cell nephropathy and the relatively large plasma volume resulting from low packed cell volume.9 With improved care and close routine follow up of children with SCA particularly in tertiary centres like ours, in addition to several strategies aimed at control of malaria and other infectious diseases; which are recognized important causes of mortality particularly in children with SCA, majority of these children are surviving into adolescent and adulthood. These young adults are likely to harbour several organs dysfunction owing to the underlying pathologic processes in SCA. Kidneys, the commonest cause of

secondary hypertension in children, are one of the earliest organs affected in children with Both structural and SCA. functional abnormalities of the kidneys are recognized progressively beyond the age of five.¹⁰ The abundance of data on blood pressure pattern among these children and young adults with SCA documented low systemic blood pressure and thus, interval blood pressure check or cardiology referral is not part of the standard care protocol for SCA children and adolescents.

We thus, aim to determine the blood pressure pattern among children and adolescents with SCA and compare same with similar children without SCA in the North-eastern region of may allow Nigeria. This for early identification and prompt management of blood pressure abnormalities in children with SCA, thus, reducing the long term complications and improve their quality of quality of life.

Materials and Methods

This cross-sectional comparative study was conducted at the Paediatric Haematology Clinic and General Paediatric Outpatient Clinic of the University of Maiduguri Teaching Hospital (UMTH), over a period of six months, from January to June, 2016. The study population were known SCA children in steady state as defined by Akinola et al,¹¹ on folic acid and proguanil aged between 3 and 14 years on routine follow up visit. Participants were recruited systematically by choosing alternate patients as they presented at the Clinic after meeting the inclusion criteria.

The controls were age and sex matched apparently healthy children with haemoglobin AA on follow up after recovery from minor ailments at the General Paediatric Outpatient clinic.

Ethical considerations

Approval was sought and obtained from the Hospital Research and Ethics Committee of the UMTH, written consent was obtained from the care givers after adequate education. Verbal assent was also obtained from children 7 years and older with unlimited liberty to deny consent or opt out of the study without any stage at any negative consequence. Those detected with hypertension were referred to Paediatric cardiology clinic for further evaluation and subsequent co-management.

Sample size: This was determined using Taylor's formula,¹² and P was taken from a previous study by Garlick et al from South-Western Nigeria who reported prevalence of 3%.13 Thus, 54 SCA children (cases) and 159 HbAA children (controls) were recruited.

Inclusion criteria: Children aged 3 - 14 years with SCA form the cases, apparently healthy non-SCA children age and sex matched form the control after consenting. Children below the age 3 years were not included because the study require measuring height of the subjects which children within this age group cannot cooperative to obtain accurate measurements. Even though length may be used for children in this age category, the procedure is cumbersome. Also, BP can vary by 40 - 50 mmHg in infants and young children following crying. Excluded were children whose parents/caregivers did not give consent to participate in the study. Information obtained from the cases include Biodata, history of previous episodes of crises and previous history of blood transfusion. The standing height in (cm) and weight in were measured using Wunder's (kg) stadiometer fitted with a weighing scale. The body mass index (BMI) was calculated and classified using World Health Organization (WHO) growth charts.

Blood pressure of each subject was measured with Accoson's mercury in glass sphygmomanometer using age appropriate cuffs (one which bladder covers two-third of the subject's right arm in supine position). Systolic blood pressure was taken at the point at which the first Korotkoff sound was heard and the diastolic blood pressure at the point at which the Korotkoff sound became muffled.¹⁴ Systolic blood pressure (SBP) and diastolic blood pressure (DBP) were defined as normal between the 5th and the 90th percentile for age, sex and height. Hypertension was defined as BP greater or equal to the 95th percentile for age, sex and height percentile. Blood pressure was classified using published normative data (US).15

Packed cell volume was determined for all the participants using capillary blood collected from a finger prick into a heparinized capillary tube after cleansing with alcohol. The PCV was read with the Hawksley haematocrit reader after centrifuging the capillary tubes.

Data obtained were analysed using Statistical Package for Social Sciences (SPSS) version 16 of 2008 (SPSS, Chicago, Illinois, USA). Means, standard deviation, frequencies and percentages, tables, graphs, and charts were used to present data. Paired t-test for comparing the means of two samples was used to test for significance of the differences between means of BP of SCA subjects and their controls, Chi-Square was used to compare categorical variables. Relationship were also explored using correlation and p value of 0.05 or less was considered significant at 95% confidence interval.

Results

A total of 54 children with SCA in steady state and 159 controls with haemoglobin AA were enrolled during the study period. The age and sex distribution of the study population is shown in table I.

Age group (years)	SCA n (%)		SCA n (%) CONTROL n (%)		TOTAL
	Male	Female	Male	Female	
<5	6(21.4)	4 (15.4)	14(19.2)	6(7.0)	30 (14.1)
5-10	10(35.7)	14(53.8)	27 (37.0)	48(55.8)	99 (46.5)
>10	12(42.9)	8(30.8)	32(45.2)	32 (34.5)	84 (39.4)
Total	28 (52)	26 (48)	73 (45.9)	86(54.1)	213 (100)

Table 1: Age and sex distribution of the study population

SCA=Sickle cell anaemia

Of the 54 SCA children 28 (52%) were males while 26 (48%) were females giving a Male: Female ratio of approximately 1.1:1. There was no significant difference in ages of males and females $X^2 = 3.437$ and p=0.329 in the SCA group.

The frequency distribution of the PCV of the SCA patients is as shown in figure 1, majority (53.7%) of the SCA patients had PCV between 21% and 25% while only 2 had PCV of >30%



Fig. 1: packed cell volume distribution of children with SCA

The mean blood pressure of the subjects and controls are shown in table III. Althou gh both the mean systolic and diastolic blood pressure of the control group were higher than those of the SCA group, only the diastolic blood pressure was significant (P= 0.0001). Eight

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(14.8%) of SCA children and none of the controls had blood pressures within the hypertensive range. The mean weight and height were also significantly higher in the controls than the children with SCA (P= 0.0243 and 0.0433) respectively.

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Parameters	SCA		Control	
		Mean(SD)	Mean(SD)	P-value
Weight(kg)		21.98(7.39)	25.90(11.93)	0.0243*
Height(cm)		120.80(18.51)	127.29(20.83)	0.0433*
BMI(kg/m²)		14.91(3.12)	15.07(2.87)	0.7296
SBP(mmHg)		95.56(10.55)	99.11(13.38)	0.0780
DBP(mmHg)		59.48(8.39)	64.96(8.57)	0.0001*

Table 2: Mean b	lood pressure a	and anthropometric	data of study	population
			j	

SCA=Sickle cell anaemia, SD= Standard deviation, BMI= Body mass index, SBP= Systolic blood pressure, DBP= Diastolic blood pressure. *p=<0.05

Only one child with SCA was overweight, and none was obese. While most 148 (84.9%) of the controls were underweight and none was overweight. The BMI of the SCA children and controls is shown in table III.

BMI	Frequency (%)	Haem AA
	SCA	Controls
Underweight	49(90.7)	148(84.9)
Normal	4(7.4)	11(15.1)
Overweight	1(1.9)	0(0.0)

Table 3. Distribution	of BMI	of children with	SCA and controls.
Table 5. Distribution	OI DIVII	of children with	och ana controlo.

BMI= Body mass index (Kg/m^2)

The SCA children with hypertension were significantly younger and shorter than those of them with normal blood pressure (p = 0.0112 and 0.0199) respectively. There was no

significant difference in BMI between the hypertensive and normotensive SCA children Table IV.

Table 4: Association of clinical	and laboratory	parameters	with the	e mean	Systolic	blood
pressure in SCA Patients						

Factors	Mean (SD)	Mean (SD)	t-value	p-value	
	Hypertensive	Normotensive			
Frequency∞	8 (14.8%)	46 (85.2%)			
Age in years	5.38 (1.061)	8.80 (3.625)	2.6301	0.0112*	
PCV	22.00 (3.5857)	22.00 (3.4189)	0.0000	1.0000	
Weight in kg	17.88 (3.3568)	22.696 (7.6837)	1.7351	0.0886	
Height in cm	106.88 (5.939)	123.20 (18.916)	2.4028	0.0199*	
BMI(kg/m2)	15.91 (4.2786)	14.74 (2.9024)	0.9763	0.3334	
PCV= Packed cell volume, BMI= Body mass index. ∞ = summary, *p value <0.05					

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Discussion

In this study, we found a significantly lower mean diastolic blood pressure in SCA children compared to the control group. This agrees with the findings of previous studies; Ernst *et al*^{,8} Sergeant *et al*¹⁶ and Pegelow *et al*⁴. Several reasons have been advanced on why the SCA children may have lower blood pressures. Firstly, the chronic hypoxia secondary to the chronic anaemic state in subjects with SCA, causes vasodilatation leading to lower peripheral resistance and hence lower systemic blood pressure.17Furthermore, SCA children are known to have higher serum concentration of prostaglandin which is a potent platelets aggregation inhibitor¹⁸⁻²⁰ and thus lowers Other peripheral resistance. factors attributable to low systemic blood pressure in children with SCA include salt losing sickle cell nephropathy,9 renal tubular damage with concentrating defects and increased renal tubular sodium and water excretion thus promoting lower arterial pressure.²¹⁻²³

In contrast to the prevailing overall lower mean blood pressure among the SCA children, 14.8% of them had elevated blood pressures. This finding is also in agreement with earlier reports of hypertension occurring in children with SCA.²⁴⁻²⁷This high prevalence of hypertension stresses the need for routine measurement of blood pressure in children with SCA. These SCA children with hypertension were significantly younger when compared to those with normal blood pressure. Similar finding was reported from a study that used 24 hours ambulatory BP monitoring.²⁸ Thus, it is unlikely that the finding from our study is due to 'white-coat hypertension'. Higher PCV has been associated with higher BP in SCA from previous studies.^{29,30}However, there is no significant difference in PCV between hypertensive and normotensive groups in our study. The reason for high preponderance of hypertension in the younger age group may not be immediately clear, but may suggest that early phases of SCA nephropathy may be characterized by hypertension.^{25,26}

Anthropometric parameters in this study are similar to those described in most literature that patients with SCA have a growth delay which starts at the age of two years, and this affects weight more than the height and progresses through the adolescent years.^{24,31} Some reports demonstrated an association between elevated blood pressures and BMI.^{24,32-34}

However, we did not find such association in our studies probably because BMI was low in virtually all groups as a lot of the enrolees are internally displaced from their homes due to the insurgency in the North-eastern part of Nigeria.

Other possible explanation for those findings is the fact that SCA patients have changes in plasma renin, endothelin and nitric oxide metabolites due to chronic vaso-occlusion with those changes affecting the balance between vaso constriction and vasodilatation, which is not usually seen in severely undernourished children.^{24,26}

Conclusion

In general, children with sickle cell anaemia have lower blood pressure when compared to those with HB AA. Hypertension may be more prevalent among the younger children with SCA. However, the identification of few SCA patients with hypertension stress the need to routinely measure the BP of children with SCA on follow up. Multi-centre studies with large sample size is desirable in Nigeria to evaluate BP in children and adolescents with SCA for detection of possible causes of hypertension and effective management to

avoid devastating events such as Cerebrovascular accident.

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References

- Grosse SD, Isaac O, Hani KA, Djesika DA, Frederic BP, Thomas NW. Sickle cell disease in Africa: A neglected cause of early childhood mortality. Am. J. Prev. Med 2011; 41(6): 398-405
- Olanrewaju DM, Complication of sickle cell anaemia-A Review. Nig Med Pract 8. 1988; 16(3): 107-111
- George IO, Opara PI. Sickle cell anaemia: A survey of associated morbidities in Nigerian children. Afr J Haematol Oncol. 2011; 2(2): 187-190
- **4.** Pegelow CH, Colangelo L, Steinberg M, Wright EC, and Smith Jand Phillips G *et al.* Natural history of blood pressure in sickle cell disease: Risks for stroke and death associated with relative hypertension in sickle cell anaemia. Am. J. Med 1997; 102: 171-177
- Falkner B. Hypertension in children and adolescents: Epidemiology and natural history. Paediatr Nephrol. 2010; 25(7): 1219-1224
- 6. Iampolsky MN, Fabiola IS, de Souza, Roseli OS, Sarni. Influence of body mass index and abdominal circumference on children's systemic blood pressure. Rev Paul Pediatr. 2010; 28(2): 181-187

Authorship contributions

Conceived and designed the experiments: Farouk AG and Ibrahim HU.

Performed the experiments: Farouk AG, Ibrahim BA and Ibrahim HU. Analyzed the data: Farouk AG, Elechi HA, Ibrahim BA and Idrisa JA.

Contributed materials/analysis tools: Farouk AG, Mustapha MG, Elechi HA, Idrisa JA and Ibrahim BA. Wrote the paper: Farouk AG, Mustapha MG, Ibrahim BA and Elechi HA. Conflict of interest: None declared

- 7. Silva Ana Carolina Pio da, Alberto Augusto Alves Rosa. Blood pressure and obesity of children and adolescents' association with body mass index and waist circumference. Arch latinoamNutr Caracas 2006; 56(3): 244-250
- 8. Ernst AA, Weiss SJ, and Johnson WD. Blood pressure in acute vaso-occlusive crises of sickle cell disease. South Med J. 2000; 93: 590-592
- **9.** Radel EG, Kochen JA, Finberg L. Hyponatraemia in sickle cell disease. A renal salt loosing state. J Paediatr. 1976; 88: 800-805
- **10.** Pham PT, Pham PC, Wilkinson AH, Lew SQ. Renal abnormalities in sickle cell disease. Kidney Int 2000; 57: 1-8
- **11.** Akinola NO, Stevens SM, Franklin IM, Nash GB, Stuart J. Subclinical ischaemic episodes during the steady state of sickle cell anaemia. J Clin Pathol 1992; 45: 902-906
- **12.** Araoye MO. Subjects selection. In: Araoye MO, ed. Research methodology with statistics for health and social sciences. Ilorin: Nathedex, 2003; 115-129
- **13.** Garlick JP, Barnicot NA. Blood group and haemoglobin variant in Nigerian (Yoruba) children. Ann Hum Gen 1957; 21: 420-425.

- 14. Bernstein D. Disease of the peripheral vascular system. In: Behrman RE, Kliegman RM, Jenson HB. Eds. Nelson Textbook of Paediatrics. Philadelphia: WB Saunders Company, 2004: 1591-1598
- **15.** National High Blood Pressure Education Program Working Group on High Blood Pressure in Children and Adolescents. The fourth report on the diagnosis evaluation, and treatment of high blood pressure in children and adolescents. Paediatrics 2004; 114(2): 555-576
- **16.** Sergeant GR, Sergeant BE. Clinical features. Sickle cell disease. Third edition. Oxford University Press, 200: 194-208
- 17. Grell GAC, Alleyne GAO and Sergeant GR. Blood pressure in adults with homozygous sickle cell disease. Lancet 1981; 2: 1166. Ref.: https://tinyurl.com/y5jl3w5s (https://tinyurl.com/y5jl3w5s)
- 18. Johnson CS and Giorgio AJ. Arterial blood pressure in adult with sickle cell disease. Arch Int Med 1981; 141: 891-893
- **19.** de Jong PE, Landman H and Van Eps LWS. Blood pressure in sickle cell disease. Arch Int Med 1982; 142: 1239
- **20.** Karayaylali I, Onal M, Yildizer K, Seyrek N, Paydas S, *et al.* Low blood pressure, decrease incidence of hypertension and renal, cardiac, and autonomic nervous system functions in patient with sickle cell syndromes. Nephron. 2002; 91: 535-537
- **21.** de la Sierra A. Segura J. Banegas JR, Gorostidi M, de la Cruz JJ, Armario P *et al* Clinical features of 8295 patients with resistant hypertension classified on the basis of ambulatory blood pressure monitoring. Hypertension 2011; 57(5): 898-902
- **22.** The fourth report on the diagnosis, evaluation and treatment of high blood pressure in children and adolescents.

Paediatrics 2004; 114(2Suppl 4th Report): 555-576

- 23. Elebute O. The effect of age and sickle cell disease on renal function in Nigerian Children. West Afr Med 1973; 22: 93
- **24.** Ho CH, Joao TAC, Josefina APB. Blood pressure in children with sickle cell disease. Rev Paul Pediatr. 2012; 30(1): 87-92
- **25.** Shatat IF, Jackson SM, Blue AE, Johnson MA, Orak JK, Kalpatthi R. Masked hypertension is prevalent in children with sickle cell disease: A Midwest pediatric nephrology consortium study. Pediatr Nephrol. 2012; 28: 115-120
- **26.** George IO, Tabansi PN and Onyearugha. Hypertension in Children with Sickle Cell Disease: A Comparative Study from Port Harcourt, Nigeria. International Blood Research & Reviews 2015; 3(3): 130-134
- 27. Johnson CS. Arterial blood pressure and hyper viscosity in sickle cell disease. Hematol Oncol Clin North Am. 2005; 19: 827-837
- 28. Moodalbail DG, Falkner B, Keith SW, Mathias RS, Araya CE et.al. Ambulatory hypertension in a pediatric cohort of sickle cell disease. Journal of American Society of Hypertension. 2008; 1 – 9.
- **29.** Wolf RB, Saville BR, Roberts DO, Fissell RB, Kassim AA et al. Factors associated with growth and blood pressure pattern in children with sickle cell anemia: Silent Cerebral Infarct Multi-Center Clinical Trial cohort. Am. J. Hematol. 2015; 90: 2-7.
- **30.** Rodgers GP, Walker EC, Podgor MJ. Is "relative" hypertension a risk factor for vasoocclusive complications in sickle cell disease? Am J Med Sci. 1993; 305: 150–156
- **31.** Zago MA, Kerbauy J, Souza HM, Figueiredo MS, Costa FF, Cruz SM *et al.* Growth and sexual maturation of Brazilian patients with sickle cell diseases. Trop Geogr Med 1992; 44: 317-321

- **32.** Sarni RO. Souza FI, Pitta TS, Fernandez AP, Hix S, Fonseca FA. Low birth weight: Influence on blood pressure, body composition and anthropometric indexes. Arc Med ABC. 2005; 30: 76-82
- **33.** Sorof JM, Lai D, Turner J, Poffenbarger T, Portman RJ. Overweight, ethnicity and the

prevalence of hypertension in school-aged children. Pediatrics. 2004; 113: 475-482

34. 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 Hypertens 1989; 2: 2-8

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