## Sickle Cell Disease Awareness, Prevention and Management Competence of Student Teachers: Panacea for Strengthening Childhood Education in Nigeria

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Abstract: The study seeks to examine the level of awareness on the prevailing conditions and preventive trends on sickle cell disease among Primary Education Students in Usmanu Danfodiyo University Sokoto. The total population is 755 students, and purposeful sampling technique was used and all the 150 students (UG 3 level) were used as sample for the study. Researcher constructed instrument was used for the study. Samples were subjected to pre-test, awareness/ lecture sessions, and post test. Five research questions and one hypothesis were advanced for the study. Frequency, percentages and t-test were used as statistical instruments. The study found significant difference between pretest and post test scores of students on awareness, knowledge of conditions and preventive measures of sickle cell. The study found difference between pre-test and post test on students decision for spouse compatibility, and knowledge of managing SCD in the class. The study suggests the need for under graduate general studies course content to include lectures on sickle cell conditions, specifically all teacher training professional courses to facilitate wide spread awareness of sickle cell management and preventive measures. The study suggests the need to customize the practice of genotype screening and matching before marriage in various societies, specifically parents, religious leaders and clerics are to sensitize and make gene-type compatibility an important requirement for prospective couples. The study concludes that the best treatment for sickle cell disease is prevention, and this can be achieved with proper and valid laboratory blood screening test results, and matching of spouses before marriage to prevent the high rate of sickle cell birth and the eventual traumatic life they are likely to encounter can be addressed by team work of medical personnel and well trained teachers.

**Keywords**— Sickle Cell, Awareness, Prevention, Students, Competence

#### 1. Introduction

Children with Sickle Cell Disease (SCD) need not only adequate medical treatment, but also proper educational intervention and psycho-social management. This is because a apart from being a fatal disease with high morbidity and mortality rate, it also renders it sufferers handicap educationally, psychologically, socially and generally. Unlike other special need conditions such as seeing hearing, speech, which have special provisions in terms of personnel and facilities, SCD learners attends conventional educational provision for normal children without due regards to their special need conditions. The scenario faced by SCD learners in Nigerian schools is lopsided and need to be strengthen to enable SCD children attain their full educational and general capacity. Specifically, teachers generally are the lead role players in education of a sickle cell learner and should be well informed and equip to function well. To find out awareness knowledge and capacity of student teachers on SCD and its management is the major focus of this study.

Sickle cell disease is caused by transmission of genetic homozygous sickle cell genes(SS) from both parents(Adewoyin,2005). Once transmitted the child acquire an abnormal hemoglobin called hemoglobin S or sickle hemoglobin (Ozenwosu, Chuku, and Ikefuna,2015). Hemoglobin is described by (Adelekin & Ayinmode (2005) as a protein in red blood cells that carries oxygen throughout

the body. In a normal circumstance red blood cells are flexible and round in shape and moves easily through the blood vessel,but in sickle cell patients the sickle hemoglobin cause inability for enough red blood cells to carry adequate oxygen throughout the body causing swelling of limbs, severe pains, infections, crises and a lot of other fatal complications leading to high morbidity and mortality of suffers. This according Cold, to Treadwell, weissman and Vichinly (2008), this account for low rate of survival, and negative implications in their physical, emotional, psycho-social wellbeing. It also affect schooling, educational achievements, work life, marriage and family life. Schooling and education of children with sickle cell condition is of paramount importance and deserve professionally trained personnels who can handle them properly to enable them cope with the rigors of academic demands and educational pursuits in addition to ailments and complications posed on them by Teachers just like doctors and parents are indispensable personnel in management of children with SCD. The relevance of Primary Education students to be properly trained and groomed to enable them organize, plan, and implement primary education for all Nigerian children including children with special need conditions such as the SCD is considered paramount. Arie (2008) conducted a research on the scholastic performance children with sickle cell disease. The sample comprises two groups of African American children with SCD of which 17 are sickle cell

anemia(SS) most severe form, and 15 with Sickle cell disease, a less severe form. They were compared with 34 healthy children with similar demographic characteristics. Their scholastic performance, standardize reading and mathematics achievement test score and school attendance were compared and the result revealed significant difference in favour of healthy children, which suggest that SCD children are at greater educational risk of school failure than their healthy counterparts. The inference derived from this finding showcase the need for teachers to be equipped with some interventions strategies to cater for the educational need of SCD learners.

Researchers like Ameh and Tarfa (2012), Afolayan and jolayemi have attested to the high rate at which the population of SCD are increasing which need to be addressed through advocacy. Education for all and Sustainable Development goals that seek to strengthen educational and life attainments and success for all across the world may not be feasible were some segment of world populations (SCD) are at risk of failure. Available statistics (WHO2005, )shows the prevalence of sickle cell disease shows that 5% of the world's population are carriers of genes responsible for sickle cell disease. Those who inherit only one mutant gene(S hemoglobin) from one parent becomes a carrier, and the presence of single abnormal sickle genes does no harm to the carriers rather it protect them from malaria, but the inheritance of two abnormal genes from two parents( carriers) leads to the condition of sickle cell disease which is irresistible to malaria parasite and malaria is the major cause of ill health condition and death among patients with sickle cell especially children under five( Animasahun, Akitoye and Njokanma (2008). Also Adewoyin (2015) described the scenario of the prevalence of sickle cell disease (SCD) as one of the most common genetic disease globally with highest prevalence in Middle East, Mediterranean region, South Asia, and Sub Saharan Africa especially Nigeria. It is more prevalent in mosquioto endemic areas, rural areas, where awareness, health services and educational attainments is low.

It is generally acknowledged that the SCD is cause by single mutant genes(S gene) donated by each parents (carriers of S genes) resulting in two mutant genes in the offspring (SS genes) leading to sickle cell disease. This means situation can be averted if only one parent has the genes and the other does not have. The case advance by this study is that is a worthy venture to work out modality of spreading the information and raising awareness that this dreadful disease can be averted through simple identification and matching of prospective parents to ensure that two people with S genes do not mate or marry to produce infected SCD child. Thus the study aim to find out the level of awareness and knowledge of sickle cell condition among Primary Education Students (as prospective parents and teachers). As prospective parents since they have not married then equipping them with prior knowledge on selection of compatible genotype spouse might play a great role in their decision on choice of rightful spouse to avoid repeated cycle of generating carriers and SCD children. And as prospective teachers, educating them on the causes, symptoms and educational management of sickle cell disease condition might also help to address some difficulties face by SCD learners and promote their well being, performance and achievements.

Some research studies conducted provide some useful revelations and need for more researches. A cross sectional research conducted by Animasa, Akitoye and Njokanma (2005) on sickle cell anemia awareness among health professional and medical students at the Lagos University Teaching Hospitals. The study revealed that 91.3% had heard about SCD prenatal screening, but only 75% knew that SCD could be prevented by pre-natal screening.48% of them were not aware that prenatal screening is available in Nigeria and 42% would not allow preventive termination of pregnancy for positive screening results. The study shows that respondent do not succumb to the idea of terminating pregnancy which may be due to their belief. The study used health workers and focused on prenatal screening while the current study used student teachers and investigated on premarital screening and decisions. Another study by Abioye-Kuteyi, Oyegbade, Bello and Osakwe (2009), they conducted a survey of 320 Local Government Workers in Ile-Ife, Nigeria, on sickle cell knowledge, pre-marital screening and marital decisions. The findings shows that 69% had poor knowledge about SCD,95% had positive attitude toward premarital screening and 86.7% of the respondents and 74% of their partners had done sickle cell sceening.25% of the married and engaged respondent did not know their status. While 33-66% of the respondents indicated that they would continue with their partners if either or both had hemoglobinopathy. Actually, previous empirical studies conducted made some useful discoveries, but they are not very extensive, therefore, the need for wide spread researches in different communities become relevant, thus the current researcher envisage the need to conduct this study using different environment, approach, research design/instrument and samples.

#### 2. PROBLEM STATEMENT

Sickle cell is one of the predominant genetically acquired killer diseases among human race. Despite the increasing number of educated people in the society and developments associated with it, it is sad to learn that the prevalence of sickle cell disease is also increasing instead of reducing. With a rising population of those affected with sickle cell as Akinyaju and Ann(1989) indicated that about 25% of adults have sickle cell trait and estimated 90.000 out 5.4 million to be SCD and 1.1 million to be carriers. While WHO (2005 gave an estimate of 150,000 birth of sickle cell disorder in Nigeria annually. As at 2016, though no exact estimate but it is evidently becoming a popular disease in most households. This study assume that lack of awareness, knowledge and sensitization about the SCD and improper matching of

marriage partners account for the rise in number of sickle cell birth. Traditionally, there seem to be lukewarm attitude toward blood screening before spouse selection. Traditional and customary arrangements in most society including Sokoto being the area of research does not give prominence ascertain and ensure valid and reliable laboratory genotype screening of prospective couples before entry into marriage union. Degree certificate is high educational attainment which is expected to bring about not only high level man power and resource persons but also people with high level capacity to make wise decision on matters relating to spouse selection and reproduction of healthy children . SCD is assumed to be non-curable but preventable disease. The objective of the study is to investigate the awareness attitude competence of students teachers who are professionally trained to manage children with sickle cell conditions, on the preventive side it is assumed that since most of the student are not married ,then they form the best target group to help in addressing the prevention through their choice of spouse with compatible genotype.

#### 3. METHODOLOGY

The study used three hundred level undergraduate students to find out their awareness and knowledge of the sickle cell condition and their attitude toward prevention and handling of learners with sickle cell disease. The study was quasi experimental with a pretest at the beginning then they were exposed to awareness and lectures and then post test. The population 750 Primary Education students, out of which 150 three hundred level students during the 2015/2016 session offering the courses child psychology and special education respectively were used for the study. The instrument was validated and found reliable with .73 reliablity index. The sensitization given to the respondents was on myth, life stories and updates on sickle cell condition, while lectures were given on the prevention, educational implications and symptoms. management of SCD five research questions and one hypothesis were raised and analysed using frequency count percentages and t-test.

# 4. DATA PRESENTATION Table 1:Showing Frequency Distribution on Biodata of Respondents

Variable	Range	Male percentage	Female Percentage	Total
Sex		100 ( <b>66.67</b> %)	50 (33.33%)	150
Age:	18-30 31-45	92 (61.33%) 08 (5.33%)	42 (28.00%) 08 (5.33%)	150
Marital status	Married Unmarried	08 (5.33%)	10 (6.66%)	150

	92 (61.33%)	40 (26.66%)	
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Table 1 shows the that there are 100 male and 50 females respondent used for this study. The figures are actual number of primary education students in 300 level. The data shows that 92(61.33%) males and 42(28%) females are within the age range of 18 to 30 years and only 8 male and 8 female are above 30 years. And 92 out 100 male students representing (61.33%) ma and 40 out of 50 females students are not married. The number of unmarried students is quite high and could be sensitize on proper matching for compatible partner.

**Research Questions 1**: To what extent are the primary education student teachers aware of some basic information on sickle cell disease.

The level of awareness was measured by number of students that were able to answer questions on basic information considered essential for a potential personnel in education of children. This was ascertain pre-test and post test. Item analysis of their responses were presented in frequency and percentages.

Table 2: Showing Students Level Of Awareness On SCD.

Responses	Pretest		Post-test			
	Aware	Not aware	Aware	Not aware		
Awareness of	58(38.7%	92(62.3%)	134(89.3%)	16(10.7%)		
how SCD is						
transmitted						
Awareness	77(51.3%)	73(48.7%)	121(80.7%)	29(19.3%)		
that it has no						
known cure	25/240/	111/750/	122 (000)	10/10/1		
Awareness	36(24%)	114(76%)	132(88%)	18(12%)		
that it can be						
prevented	25/22 20/	115(76.70()	1.40/02.220/	10(6.70()		
Awareness of	35(23.3%)	115(76.7%)	140(93.33%)	10(6.7%)		
pre-marital						
genotype screening						
Awareness of	17(11.3%)	133(88.7%)	110(73.33%)	40(26.7%)		
prenatal	17(11.570)	133(66.770)	110(75.55%)	40(20.770)		
genotype						
screening						
Awareness of	17(11.3%)	133(88.7%)	112(74.7%)	38(25.3%)		
bone marrow	(	(,,	(,,,,,	23(221272)		
transplant						
Awareness of	00(0%)	16( 100%)	00(00%)	00(100%)		
spouse						
genotype						
before						
marriage						
Awareness of	14(9.3%)	86(90.7)	138(92%)	12(8%)		
need to cross						
match with						
Genotype of						
prospective						
spouse						

Table 2 contain data on the pretest and post test scores on student level of awareness on some basic information about transmission and prevention of SCD. Before the interaction only 58 out 150 shows evidence of awareness of how SCD is transmitted . Similarly, the pretest scores shows that very few students had prior knowledge about how it can be prevented and that it is incurable. The post test result shows increase on number of student that show awareness of basic information on SCD.Out of the 150 respondents only eight of them were married and all of them claimed they don't know the genotype of their spouse.

**Research Question 2:** To what extent do the students teachers in Primary education possess necessary knowledge about sickle cell condition as prospective functional teachers?

Table2: Showing students frequency count on their

knowledge of sickle cell Disease									
Response	Numbe	Pretest		Post test					
S	r								
		know	Don't know	Know	Don't know				
Knowledg	150	32(21.3	118(78.7	125(76.7	25				
e of		%)	%)	%)	(23.3%)				
genotype									
Knowledg e of sickle cell disease condition	150	54(36.%)	96(64%)	145(96.7 %)	5(3.3%)				
Knowledg	150	46(30.7	104	130(86.7	20(30%)				
e of how		%)	(69.3%)	%)					
it is									
acquired									
Knowledg	150	34(22.7	116(77.3	132(88%)	18(12%)				
e of		%)	%)						
causes of									
sickle cell									
condition									
Knowledg	150	36(24%)	114(76%)	140(93.3	10(06.7				
e about					%)				
symptoms									
Knowelge	150	50(33.3)	100(66.7)	137(91.3	13(8.6%)				
about its				%)					
cure									

The data in table 2, shows item analysis of number of students that possess some knowledge on sickle cell disease before and after the treatment. Before the advocacy sessions, the pretest shows that very few students have ideas and knowledge about sickle cell conditions and the post test shows that more students were educated and show evidence of knowlge about sickle cell disease.

Question 3:What factors do the primary education student teachers consider in choice of spouse

Table 4: Factors to consider in selection of spouse

Table 4. Pactors to consider in selection of spouse								
Responses	Pre	test (%)	Post t	est (%)				
Love	46	(30.66%)	15	10%				
Economic status	24	(16%)	11	7.33%				
Educational status	15	(10%)	12	8%				
Health status	20	13.33%)	13	8.66%				

Compatible genotype	10	6.66%)	80	53.33%	
Physical appearance	17	(11.33%)	10	6.66%	
Behaviour and character	18	(12%)	09	%	
Total	150	(100%	150	100	

The table 4 reveals what the participants consider most in spouse selection, the initial response before that interaction shows the highest response of 46 students representing 30.66% of the respondents choose love as what they consider most in decision about spouse , this was followed by economic status and health status . while on the post test the factor with the highest response was genotype compatibility with 80(53.3%) of the respondents, followed by love with only 15 equivalent to 10% and health status with only 13(8.66) respondents.

**Research Question 4:** What is the genotype distribution of respondent and their decision to mary SCD.

Table 5. Student genotype distribution and decision to

marry a person with sickle cell desease.

		Pre-	test	Post-test			
Respo nses	Genot Decision to ype marry SS			Genotype (%)	Decision to marry		
	<b>JP</b> -	Y	es Io	(1-)	Yes	No	
AA genoty pe	68 (45.3% )	24 (16%)	44(29. 3%)	96 (64%)	47(31. 3%)	38(25) 3%)	
AS genoty pe	08 ( 5.3% )	02 (1.3%)	06 (4%)	35(23 3%)	10 (6.7%)	05(3.3 %)	
SC genoty pe	00 (	00 (0 %)	00(0%)	01(0.7%)	00 (00%)	01(.7%	
No idea	74 (49.3% )	38(25. 3%)	36 (24%)	18(12.7%	9(6%)	7(4.7 %)	
Total	150(10 0%	15	0(100%)		150(10	00)	

The table 5 shows genotype indicated by students in pretest and the genotype results obtained laboratory genotype screening for post test .68(45%.33) of the students claim they are AA ,while 74(49.33%) indicate that they don't know their genotype. The post test (laboratory results) shows that 96(64%) respondents have AA, 15(10%) are AS and only one respondent has SC genotype, while 16 (10.66%) students did not present their genotype results. On the decision to choose a person with sickle cell as spouse, a total of 64 respondents (pretest) and 66(post test)declare their willingness to choose person with SCD as marriage partner.

Hypothesis 1: There is no significant difference between student scores before and after exposure to some treatment on classroom management for SCD Learners.

This hypotheses was analyzed using t-test statistics and presented in table 4

Table 6: showing t-test scores of students on Classroom management strategies for SCD

Variabl	N	Df	Mean	Std deviatio	T – cal	p-	Remark	
e			S	deviano	cai	value		
				n				
Pretest	15	29	29.43	12.201	6.34	P=00	Significa	
	0	8			7	0	nt	
Post	15		76.30	9.492				
test	0							

P<.05

The table 6 reveals significant difference between students score in pretest and post test on class room management skill possessed by students. Because the t-caulated value of 6.347 is greater than the p-value at significant level of p<.05. This means that providing students with privilege information on how to handle and manage SCD in classroom increase the capacity as prospective primary school teachers, thus might enhance their teachers to manage SCD learners more effectively.

#### **Major findings**

- 1. There was significant increase in level of awareness on SCD pre-test and post test scores of students teachers in primary education.
- 2. There was significant improvement in level of knowledge of SCD possessed by students teachers in primary education after exposure to sensitization and treatment.
- 3. There was significant change on the need to consider genotype compatibility in choice of spouse.
- 4. The study found fluctuation on student decision on choosing spouse with sickle cell disease.
- 5. The study found significant difference on pre-test and post test scores of students teachers on their capacity to manage and handle children with sickle cell condition in classroom.

#### 5. DISCUSSION OF FINDINGS

Research question one: The study was able to establish (in table 2) that most students teachers are not aware of how SCD is acquired and also lack awareness about different preventive measures. This poses serious risk reproduction of more children with SCD. This may explain the reason for the high increase in the prevalence of the diseases indicated by WHO(2005) and Adewoyin (2015). The inference made by this study is that the students who are being trained to serve in addressing the problem of SCD, are themselves not aware of the basic hints on the transmission and preventions of the disease. Then they are likely to marry blindly without taking precaution on the genetic transmission linkage, so as to avoid incompatible partner. The implications is that of general lack of awareness among people including prospective graduates of primary education who are suppose to serve as personnels in handling childhood education including children with S. The study further revealed that sensitization and advocacy can help to promote awareness and consciousness on SCD so as to address it systematically. Research question two assess the level of knowledge

possessed by respondent and was captured in table 3. The

table provides evidences on lack of necessary knowledge by student teachers in pretest, which was improved upon through subjecting them to advocacy and lessons on sickle cell in post-test result. Actually children with SCD are faced with various problems, teachers who are to handle them need to be conversant with signs, symptoms, complications and the limitations the sickness posed to their educational pursuit which may invariably influence psycho-social, emotional, physical, general wellbeing and life of SCD learners. The issue is though, special education is part of course taught in teacher training programmes in universities, but it was observed that sickle cell disease is not covered in the course content. As such many student are likely to graduate without awareness or knowledge of the disease, its complications and their roles as teachers in helping learners with SCD.

Research question three was on factors that students consider in choice of marital spouse. The study discovered as shown in (Table 4) that respondents do not give prominence to genotype matching of prospective partner in pretest (6.6%) but in the post test 53% of the respondent indicated genotype compatibility as a vital factor to be considered. How the clue deduced from this study is that when students are well informed, they became conscious that they can avoid giving birth to offspring with sickle cell condition even when one is a carrier (AS) or even sufferer (SS). They were also capable of weighing the problem and consequences of wrong matching. This means that adequate sensitization can help to intimate people on the need to be cautious in selection and proper match of marriage partner as a vital factor for reproduction of healthy children, their wellbeing and happiness of the family. The complications and constrain cause by sickle cell disease condition is not limited to the children or sufferer but to a large extent the parents as primary care givers are as well very much affected in so many ways. It may tell very much on job efficiency of teachers who become parents of SCD children. For this reason raising awareness of student teachers early before they become parents is considered necessary.

Research question four tries to find out whether the respondents know and did genotype test. The study was able to establish (Table 5) that most of the respondents do not know or test for their genotype in pretest. But in the post test it was found that 132 out 150 respondent were able to produce their laboratory test results indicating high percentage(64%) of AA genotype and 24% of AS genotype. The students were intimated to go recognized and reliable test and the need to do repeat confirmatory test to ascertain the reliability of the first one. However most of the candidates (pretest) declare that they will not marry an SCD, but in post test more 31% as against the initial 24% show their readiness to marry a spouse with sickle cell disease. The study found that respondent with AA genotype decline to marry SCD, which the only right match for SCD, while ironically those with AS are willing to take SS as spouse. The reason often given by those with AA is that they want to avoid problem of ailments, complication and the demand

that goes with it. While the case of AS respondents declaring to marry another AS pose great danger as it will lead to recycling of more SS genotype, which is detrimental to generation of healthy and viable human race.

One hypothesis was generated and tested using t-test(table 6). It was established by this study that it is not awareness and knowledge of sickle cell disease condition but teachers need to comprehend the peculiarities of SCD learners and the management and intervention strategies to help them overcome the difficulties in educational their pursuits.

### 6 IMPLICATIONS FOR STRENGTHENING CHILDHOOD EDUCATION.

### The findings of this study could serve as a panacea for strengthening childhood education.

Nigeria is among of the countries in the world that declared education as birth right as such is decreed to be Education For All, through the UBE 2004 ACT. However, in reality sickle cell children are not getting adequate and desired service from stake holders. Many researches WHO (2014); Aneke and Okocha(2015); Ezenwosu, Chukwu and Ikefuna, (2015), Alege(2015), had provide a lot of insight in to the increasing trends of carriers and sufferers of SCD with various disposition on the nature of recurrent pain, complication they have to live with, which dully interfere with many aspect of their life including education. The findings of this study suggest that our student teachers are not very much aware of the SCD condition and the predicaments associated with it. And that many of them are not married that is when to get them know that simple blood screening can avert the birth, suffering and demand for special need education for SCD learners. Thus the need to strengthen education of children cannot be fully attained when teachers are inadequately equipped with knowledge and skill to prevent, manage and salvage the suffering of increasing population of SCD learners. Due to the nature of their ailment, they miss school attendance, lessons, assignments, tests and exams very often which are not always compensated for by teachers, putting them at high risk of school failure. They also encounter psycho social and emotional problems in schools which are neglected, thus making them the marginalized or disadvantaged school learners.

- 1.General awareness and sensitization is very necessary using media, schools, hospitals, community meeting points, in public gathering to enable every citin be part of the venture to reduce and address the need of SCD,
- 2. Collobarative teamwork involving all relevant stakeholders such as medical, educational, social welfare, parents, community leaders and civic organization to work as team to address the needs of the SCD patients.
- 3. There is dire need to customize premarital screening in all communities to certified and match prospective couples before marriage to reduce birth of AS and SS traits. Those

- with AA genotype should be encouraged and assisted to marry SS Genotype.
- 4. The Federal and State governments should set up special school for people with SCD and equip the schools with personnel that have stake in their management. It is virtually not easy for classroom teachers to handle both SCD and normal learners in the same class. The demand may be too much for them to cope with.
- 5. All teacher training courses and programme should offer in depth study of sickle cell condition to enable them attend to their needs as teachers. While all students in higher institution should be expose to information about SCD, and be sensitized to screen their blood and make proper choice of reproductive partner.

#### REFERENCES

[1] Abioye-Kuteyi Z.A, Oyegbade, Bello A., I; (2009) Sickle cell knowledge premarital screening and marital decision among local Government working in Ile-Ife Nigeria. *African Journal of* 

Medicine. ncbi.nlm nih.gov retrieved 02-04-2016.

- [2] Adelekin.M.L &Tunde- Ayinmode M.F (2005). Psychosocial impact of sickle cell disease in children seen at University of Ilorin teaching hospital Nigeria: East African Med. Journal 82(2) 73-78. Retrieved 03-09-2016
- [3] Adewoyin A.S (2015-) SCD predominates in sub-sation African & Nigeria.
- [4] Afolayan J.A, Jolayemi F.T, (2011) .Parental Attitude To Children With Sickle Cell Disease In Selected Facilities In Ire-Podan L.G.A of Kwara State. Study On Ethro-Medicine. Krepabishes.com.
- [5] Akinyaju.O.O;& Ann.Y.N.(1989). A Profile of sickle Cell Disease In Nigeria Journal of Academic Science.565 126-136. www.ncbi.nlh.nih.gov/m/pubmed/26729662.
- [6] Ameh S.J, Tarfa F.D, Ebeshi B.U (2012) Traditional health management of sickle cell anemia: Lesson from Nigeria. Anemia vol. 2015 Article ID 791498. 21.
- [7] Aneke.J.C.& Okocha.C.E (2015).Sickle Cell Disease Genetic Counseling and Testing: A Review Archives of medicine and health.4(1) 50-57.
- [8] Aneni E.C, Hamer D.H, Gill C.J (2013) Systematic Review of Current And Emerging Strategies for Reading Morbidity from Malaria in Sickle Cell Diseases. Journal of Tropical Medicine Willing On Line Library.
- [9] Animasahun B.A, Akitoye C.O,& Njokanma. O. (2005).Nigeria Sickle Cell Anemia: Among Health Professional And Medical Students At Lagos University Teaching Hospital. Journal of Medical Info.
- [10] Arie L.N (2008)Scholastic Performance of Children with Sickle Cell Disease. <a href="http://dx.doi.org/10.1300/jo45v05no3-08">http://dx.doi.org/10.1300/jo45v05no3-08</a> .publish on line retrieved 02-09-2016.
- [11] Cold J.L, Mahrer N.E, Treadwell M, Weissman L, & Vichinsly. E. (2008). Psycho social & behavioural out rems in children with SCD and their healthy siblings. Journal of

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behavioural medicine. De 31(6) 506-16. Doi:10../007/0865,-008-9175-2

Epubhttp://dv.dol.org//1b.1155/2015/79/791498

- [12] Ezenwosu.O.U; Chukwu B.F,Ikefuna. A. N.(2015). Knowledge and Awareness of Personal Sickle Cell Genotype among Parent of Children With Sickle Cell Disease in South East. Nigeria. Journal Of Community Genetics6(4)369-374 <a href="http://dx:dol.org/10.1300/j045v05n03-08">http://dx:dol.org/10.1300/j045v05n03-08</a> Phobiah arlne aet 2008 pp123-140.
- [13] William-Smith, M. (2015).Factors that Contributes to the Knowledge ,Health, Belief, Attitudes and Behaviours Regarding Sickle Cell Disease among College Students. Uta-2502d-13155pdf. Retrieved 03-06-2016.