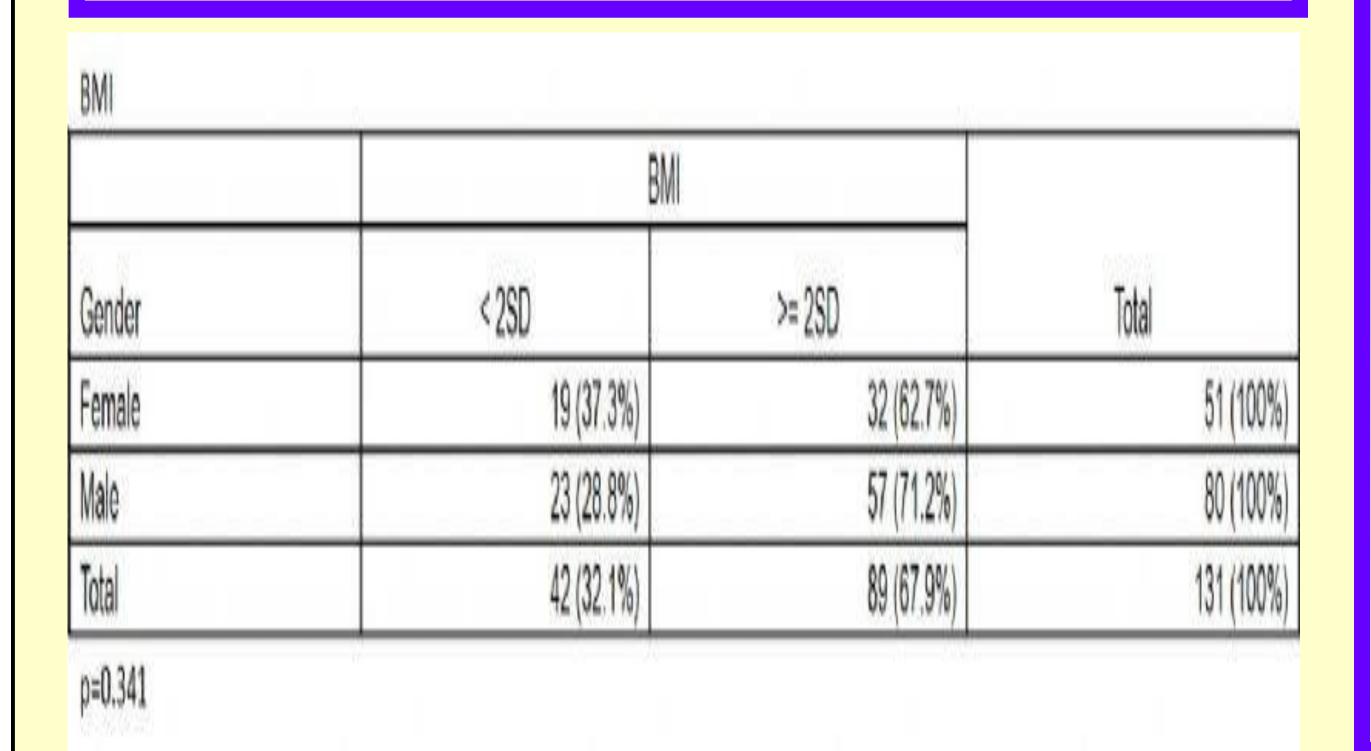
# EFFECT OF SICKLE CELL DISEASE ON GROWTH AND PUBERTY

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## **OBJECTIVES**

•To describe growth and pubertal development in children and adolescents with Sickle cell disease.



## METHODS

•This was a cross-sectional descriptive study involving 142 children with confirmed sickle cell disease who were being followed up at Kenyatta National Hospital's Paediatric Hematology clinic.

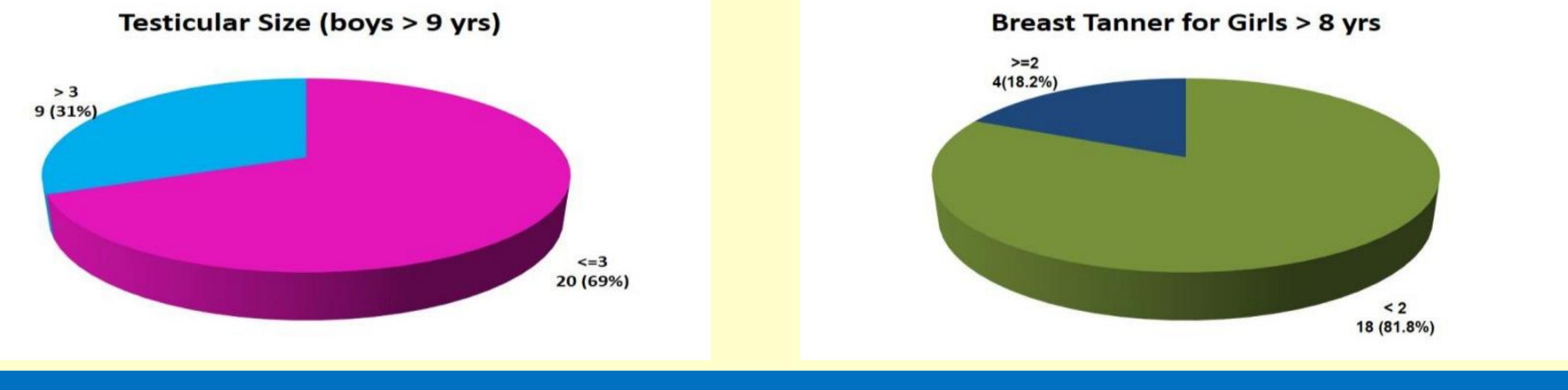
•Demographic information including age, gender, tribe and socioeconomic data were obtained by standard questionnaire while the clinical information including height, weight and BMI were obtained by actual measurements and plotted on the CDC charts for age and

gender.

•Pubertal status in girls was determined by assessing breast tanner stage while in boys it was assessed by measuring the testicular volume. Presence of pubic and axillary hair was assessed in both.

	Characteristic	Frequency	Percentage
	Sex(n=142)		000 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100 - 100
	Female	56	39.4
-	Male	86	60.6
	Age Male (n=86)		
	<=9yrs	53	61.6%
	> 9yrs	33	38.4%
	Age Female (n=56)		
•	<=8yrs	33	58.9%
	> 8yrs	23	41.1%
	Father's ethnic group(n=139)		
	Luo	91	65.5
-	Luhya	35	25.2
	Other	13	9.4
	Mother's ethnic group(n=140)		100 BOR 100
-	Luo	89	63.6
	Luhya	39	27.9
	Other	12	8.6
	Highest level of education in household		
	(n=139)	37	26.6
•	Primary		
	High school	4	2.9
	Secondary	57	41.0
-	College	24	17.3
-	University	17	12.2
	Family income per month (n=138)		
	< 5000	5	3.6
	> 20000	99	717





#### RESULTS

•The median age of patients was 7.0 years (IQR 5.5,9.5 years). Of these patients 86 (60.6%) were male and 56 (39.4%) were female with a male to female ratio of 43:28. Those with underweight were 16.3% male and 16.1% female (<-2 SD) while 11.1% male and 13.5% females were stunted (<-2SD).

•Twenty girls (96.4%) and all the boys in their respective age groups had no axillary hair while 19 of the girls (85%) and 27 of the boys (93%) had no pubic hair.

### CONCLUSIONS

### Reference

•A large proportion of these children were either underweight, stunted or had low BMI. There is need for longitudinal studies to determine whether this growth failure is due to the sickle cell condition or other environmental factors.

•Most of the children with sickle cell disease had not initiated puberty at their expected respective ages. Follow up needs be done on these patients to determine at what age they will go into puberty and what effects this condition has on their fertility rate.

1...Smiley D et al Therapy insight: metabolic and endocrine disorders in sickle cell disease...<u>Nat Clin Pract Endocrinol Metab.</u>
2008 Feb;4(2):102-9.



Growth and syndromes (to include Turner syndrome)

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