Pattern of Presentation of Sickle Cell Retinopathy in Ilorin

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Purpose: This study aimed at determining the pattern of clinical presentation of sickle cell retinopathy in the eye clinic at the University of Ilorin Teaching Hospital, Ilorin.

Methods: A retrospective review of the medical records of twenty-three sickle cell patients who were seen over a period of 5 years. Necessary information was extracted from the records.

Results: A total of 23 patients were seen. The majority of the participants were males (65.2%) and 87% were of the hemoglobin SC genotype. The mean age was 33.4 ± 8.7 years with a range of 18 to 60 years. About 78% were below the age of 35 years, however, most (60.9%) were within the age range of 26-35 years (table 1). Of all participants, 56.5% had tertiary education and 21.7% had secondary education. Varying grades of proliferative sickle cell retinopathy (PSCR) were present in all the patients. Less than half of the patients presented within three months (43.5%) of onset of symptoms. Blurring of vision (60.9%) and seeing of floaters (39.1%) were the commonest symptoms. About half (47.8%) of the presenting eyes had a visual acuity (VA) less than 3/60 at presentation and 70% of presenting eyes had grade 4 or 5 PSCR. Grade 3 PSCR was present in 39.1% of fellow eyes. The patients with tertiary education were more likely to present early (p=0.048). One

Table 1: Age and gender of patients

		Age category (Years)				Total (%)
		15 -25	26-35	36-45	>45	
Sex	Female	2	4	1	1	8 (34.8)
	Male	2	10	3	0	15 (65.2)
Tota	I	4	14	4	1	23 (100)

patient was bilaterally blind from PSCR. Loss of vision was due to PSCR in 95.6% of cases.

Discussion: HbSC genotype accounts for 87% of patients with retinopathy as seen in this study and this is similar to previous reports. 1-3 Most of the patients were between 26 and 35 years of age in this study. This is in agreement with previous studies. 1-3 Almost half of the patients seen in this study were blind at presentation as was reported by Oluleye.1 Late presentation of these cases has been reported in previous studies 1,4 and this was a similar finding in our study.

Conclusion: Most patients with SCR in this study presented within the age range of 26-35 years with stage 4 or 5 disease. Awareness on screening needs to be created among the patients and the physicians on the preventable nature of vision-threatening sickle cell retinopathy.

References

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