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Chloroquine retinopathy: pattern of presentation in Ibadan, Sub-Sahara Africa

Abstract

Background Self-medication with chloroquine is common in Ibadan, Sub-Sahara Africa. Retinopathy from chloroquine is not uncommon. The aim was to determine the pattern of presentation. *Methodology* Cases of Chloroquine

retinouology Clases of Chilologune retinopathy seen at the Retina and Vitreous Unit of the University College Hospital, Ibadan between 2008 and 2014 were reviewed. Information on age, sex, duration of chloroquine use, and visual loss were retrieved. Visual acuity at presentation, anterior, and posterior segment findings were documented. The results were analyzed using proportions and percentages.

Results Fourteen cases were seen during the study period. Mean age was 50.7 years. Male to female ratio was 3.5:1. Average duration of visual loss before presentation was 2.7 years. Average duration of self-medication with chloroquine was 5.3 years. Presenting visual acuity showed 2(14%) cases of bilateral blindness(VA < 3/60 in both eyes); 5(35.7%) cases of uniocular blindness; three cases of bilateral low vision(VA worse than 6/18 but better than 3/60). Anterior segment examination showed abnormal sluggish pupillary reaction in those with severe affectation. Dilated fundoscopy showed features ranging from mild macular pigmentary changes and bulls eye maculopathy to overt extensive retinal degeneration involving the posterior pole, attenuation of retinal vessels, optic atrophy, and beaten bronze appearance of atrophic maculopathy.

Conclusion Chloroquine retinopathy is not uncommon in Ibadan, Sub-Sahara Africa. Bulls eye maculopathy, extensive retinal, and macular degeneration with optic atrophy are the main presentations. Public health education is imperative. TS Oluleye, Y Babalola and M Ijaduola

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Introduction

Cambaiaggi and Hobbs et al^{1,2} first described the effect of chloroquine on the eve. A total dose between 100 and 300 g is needed to cause toxicity, which is equivalent to about 200 mg per day for 3 years.³ Cloroquine is cheap and readily available as over the counter medication in Nigeria. Nigeria has more reported cases of malaria and deaths due to malaria than any other country in the world.⁴ Most patients would have used chloroquine for fever before hospital presentation. Most non-skilled workers use chloroquine as prophylaxis for malaria in Nigeria. A study showed that its self-medication is so common that it is a common cause of heart block in Nigerians.⁵ Chloroquine deposits in the retinal pigment epithelium leading to its degeneration and that of the neurosensory retina leading to irreversible visual loss. In Nigeria, hospital presentation is usually a last resort for most people hence the late presentation of chloroquine retinopathy. The aim of this study was to describe the pattern of presentation of Nigerian patients with chloroquine retinopathy seen at the Retina and Vitreous Unit of the Department of Ophthalmology, University College Hospital, Ibadan between 2008 and 2014.

Materials and methods

The retinal register that contain retina unit cases seen in the department of ophthalmology, University College Hospital, Ibadan was reviewed and all cases of chloroquine retinopathy retrieved and studied to determine the pattern of presentation. Fourteen cases were seen during the period under study. Information on age, sex, duration of chloroquine use, and

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Received: 14 April 2015 Accepted in revised form: 14 July 2015 Published online: 2 October 2015 visual loss were retrieved. Visual acuity at presentation, anterior, and posterior segment findings were also documented. The results were analyzed using proportions and percentages.

Results

Fourteen cases were seen during the study period. Mean age was 50.7 years. Male to female ratio was 3.5:1. Average duration of visual loss before presentation was 2.7 years.

Table 1 Presenting features of patients with chloroquine retinopathy in Ibadan, Sub-Sahara Africa

S/No	Age	Sex	Duration of CQ use (years)	Duration of VL (years)	VAR	VAL	Anterior. seg signs	Fundoscopy
1	45	F	7	5	CF	6/6	RAPD R	Bulls' eye macula
2	52	Μ	8	2	6/60	CF	Sluggish pupils	Bulls' eye macula
3	48	F	5	1	6/36	6/60	Sluggish pupils	Bulls' eye macula
4	43	F	10	5	6/24	6/60	Sluggish pupils	Atrophic macula
5	44	F	<5	1	6/5	6/5	Nil	Early pigmentary changes
6	36	Μ	<5	1	6/5	6/5	Nil	Early pigmentary changes
7	49	Μ	3	5	6/60	6/60	Sluggish pupils	Atrophic macula
8	74	Μ	>5	>5	6/9	CF	Sluggish left pupil	Bull's eye R, atrophy Left
9	50	М	>5	>5	CF	CF	Sluggish pupils	Extensive posterior pole degeneration. Attenuated vessels
10	59	Μ	5	1	6/60	6/60	Sluggish pupils	Posterior pole degeneration
11	62	Μ	>5	1	6/24	6/18	_	Bulls eye macula
12	43	М	>5	2	6/18	CF	Sluggish left pupil, RAPD	Extensive posterior pole degeneration, pigments, demarcation line
13	50	М	>5	3	LP	LP	APD	Extensive retina degeneration, atrophic macula, pseudo RP
14	56	М	2	1	HM	6/12	Right RAPD	Extensive posterior pole degeneration with macular atrophy

Abbreviations: APD, afferent pupillary defect; CQ, chloroquine; CF, counting fingers; HM, hand movement; LP, light perception; RAPD, Relative afferent pupillary defect; RP, retinitis pigmentosa; VAL, visual acuity in left eye; VAR, visual acuity in right eye; VL, visual loss.



Figure 1 Bull's eye maculopathy: patient 3 (a) and patient 4 (b) note a ring of hypopigmentation surrounding the fovea signifying RPE window defect degeneration. The upper pictures showed generalized degenerative mottling of the posterior pole. The retina in (b) also showed vascular attenuation with some disc pallor.

Average duration of self-medication with chloroquine was 5.3 years. Presenting visual acuity showed 2(14%) cases of bilateral blindness (VA < 3/60 in both eyes); 5(35.7%) cases of uniocular blindness; 3 cases of bilateral low vision (VA worse than 6/18 but better than 3/60). Anterior segment examination showed abnormal sluggish pupillary reaction in those with severe affectation. Dilated fundoscopy showed features ranging from mild macular pigmentary changes and bulls eye maculopathy to overt extensive retinal degeneration involving the posterior pole, attenuation of retinal vessels, optic atrophy, and beaten bronze appearance of atrophic maculopathy (Table 1 and Figures 1-4). A striking picture noted included a sharp demarcation between the degenerated posterior pole and the healthy retinal and the retinal pigment epithelium in the periphery (Figure 3).

Discussion

66

The study showed that self-medication with chloroquine was the major cause of retinopathy in the patients

presented. Self-medication of chloroquine is a problem in Nigeria.⁶ Males are more affected in this study. In Calabar, south eastern Nigeria, males are the major abusers of chloroquine.⁶ The drug is sold over the counter to treat cases of fever through self-medication. In Ghana, a neighboring country, chloroquine is the most commonly kept antimalarial medication in the homes.⁷

In this study, about one-third of the eyes were blind at presentation with the rest at risk of severe visual loss. Anterior segment finding of sluggishly reacting pupils and afferent nerve defect is a reflection of the posterior segment involvement.

Bull's eye manifestation comprising of an annular parafoveal chorioretinal atrophy have been described and is pathognomonic of chloroquine retinopathy.⁸ The chloroquine accumulates in melanin of the retinal pigment epithelium. However in Nigerians as seen in this study, in addition to the bull's eye maculopathy, most patients with chloroquine retinopathy present with diffuse chorioretinal atrophy involving a large area of the posterior pole with a sharp demarcation between the



Figure 2 Extensive posterior pole degeneration, pigmentary retinopathy with attenuated arterioles and optic disc atrophy in patient 12.



Figure 3 Sharp demarcation between degenerated retina in posterior pole and normal peripheral retina in patient 12.



Figure 4 (Patient 13) advanced chloroquine retinopathy with marked retinal degeneration, optic atrophy, attenuation of both arterioles and veins, with beaten bronze appearance of the atrophic macula. Note bone spickle pigmentation in the periphery (arrow).

posterior pole and the retinal periphery. This suggests the affinity of chloroquine for the retina in the posterior pole of the eye. Extensive retinal degeneration is explained by the late presentation seen in the patients.

Blindness from Chloroquine is irreversible and unfortunately visual loss will progress despite cessation of therapy. Three patients with retinopathy are told to stop using chloroquine and use alternative antimalarials. Most of these patients with low vision will benefit from low vision aids.

Conclusion and recommendation

Chloroquine retinopathy is not uncommon in Ibadan, Sub-Sahara Africa. Bulls eye maculopathy and extensive retinal and macular degeneration are the main presentation. It is recommended that enabling law should be made by government to withdraw chloroquine from the market as is done in Malawi.⁹ Public health education is imperative in Nigeria and Sub-Sahara Africa against self-medication with chloroquine. Currently there are alternative medications to chloroquine for malaria treatment and should be considered.

Summary

What was known before

• Chloroquine retinopathy reported in patients with longterm treatment with chloroquine for rheumatologic conditions.

What this study adds

• Abuse of chloroquine in Sub-Sahara Africa resulting in retinopathy.

Conflict of interest

The authors declare no conflict of interest.

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