

Irreversible malarial retinopathy in a returning adult traveler to Marseille, France

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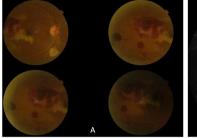
Dear Editor,

Cerebral malaria (CM) is one of the major manifestations of severe malaria, and a leading cause of malaria mortality with a 15–20% fatality rate. The development of CM may occur in 1% of children infected with *Plasmodium falciparum*, and is much more common in children under 5 years living in endemic area, than among non-migrant travelers, adults especially [1]. In France, 2.800 cases of imported malaria were recorded in 2022, of whom 18% were severe, 5% had CM and 0.3% were fatal (French National Reference Center for malaria data). Importantly, neurological signs of CM are not specific and people living in endemic areas often host parasites at subclinical levels. Over there, the definitive diagnosis of CM may significantly be improved through the observation of malaria-specific retinopathy (RP), that can serve as a diagnostic and prognostic criteria for CM [1].

We report a 56-year-old female case of CM with RP. She originated from Cameroon but had been living in France for more than 10 years. She had a history of high blood pressure and chronic alcoholism and had returned in January 2022 from a 2-month stay in Cameroon. She had not taken anti-malarial chemoprophylaxis and had travelled to forest areas. Two weeks after her return to France, she was brought to the emergency department in Marseille, for a febrile loss of consciousness. The clinical examination revealed a Glasgow Coma Scale score of 10/15 and conjunctival jaundice. The diagnosis of severe Plasmodium falciparum malaria infection was confirmed by rapid molecular detection test (LAMP technique) and peripheral blood thin smear finding 4.3% of infected red blood cells. The smear was read at lower magnification and fortuitously showed a microfilaria with a density of 40 microfilariae/ml, further identified as Loa loa by PCR [2]. She was hospitalized in the intensive care unit and treated with intravenous artesunate for 3 days. On the third day, parasitaemia was 0.05% and the patient was transferred to the infectious diseases department starting a 3-day oral relay with dihydroartemisinin-piperaquine. Loiasis was subsequently treated with diethylcarbamazine at increasing doses for a duration of 21 days. Microfilaremia was negative on day 7. The clinical examination found a bilateral visual blurring and persistent temporospatial disorientation. Blood smears were negative 7 and 28 days after the antimalarial treatment. Cerebrospinal fluid (CSF) analysis revealed 6 lymphocyctes/mm³. Brain MRI showed vascular-like leukopathy and diffuse dilation of Virchow Robin's spaces, without significant cortical or hippocampal atrophy. The ophthalmological examination found visual acuity collapsed to 6/10 in the right eye, a calm anterior chamber. Fundoscopy revealed multiple hemorrhages in clear center and cotton wool spots bilaterally; a large area of central retinal ischemia with nasal foveolar sparing and a small retro-hyaloid hemorrhage (right); a patch of central retinal ischemia with nasal foveolar sparing (left) concluding in bilateral occlusive vasculitis (Fig. 1A). Fluorescein angiography confirmed retinal hemorrhage by mask effect and areas of ischemia (Fig. 1B). Although the patient recovered from disorders of consciousness, there was no improvement of retinal lesions at one-year follow-up.

Malaria and loiasis are significant vector-borne diseases that are frequently co-endemic. The overall prevalence of malaria and loiasis co-infection was 1.2% in a recent meta-analysis [3]. The highest prevalence was found in Cameroon (2.7%). Unlike the present case, patients with co-infections usually have lower malarial parasite density, and more asymptomatic clinical status, when compared to malaria mono-infection [3]. Moreover, ocular signs of loiasis are significantly different from malaria. They include painful periorbital swelling, where the worm may present both in the subconjunctival compartment or in the anterior chamber, whereas ophthalmological malaria typically involves the retina. Exceptionally, unilateral retinal artery occlusion has been associated with the persistent presence of *Loa loa* microfilaremia in the bloodstream [4].

Malarial retinopathy is defined by the occlusion of the vessels of the retinal microcirculation secondary to the sequestration of parasitized red blood cells. Retinal hemorrhage correlates with disease severity. Moderate RP may also be found occasionally in patients with uncomplicated *P. falciparum* or *P. vivax* malaria [5]. It was first described in 1878 by Poncet who noted that almost all cases of 'pernicious malaria' were accompanied by retinal hemorrhages. Retinal whitening (macula whitening sparing central fovea and peripheral whitening of the fundus) that we observed in the case presented here, is considered as one of the specific symptoms of CM, and therefore can help differentiate malaria from other causes of coma and fever before microbiological



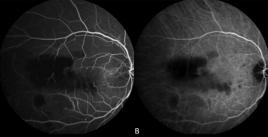


Fig. 1. A. Right fundus. Hemorrhages, Roth spots and retinal bleaching secondary to ischemia by sequestration. B. Fluorescein angiography of the right eye identifying ischemic and hemorrhagic lesions.

confirmation. Data on malarial RP outcomes are scarce. Beare et al. showed in a study in which 116 children with CM were followed up longitudinally by fundus ophthalmoscopy that in those who survived, retinal signs resolved over 1–4 weeks without retinal sequelae [6]. But, in African children survivors of CM, neurological sequelae are more frequent in retinopathy-positive cases, approaching approximately 30% [1]. Few cases of malarial RP reversibility have been reported in adults [7]. Future research is needed to better understand vascular changes in malarial RP, their prognostic significance, and targets for earlier intervention with adjunctive therapies.

Ethical declaration

Written informed consent was obtained from the patient.

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None to declare.

Declaration of competing interest

None to declare.

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