



ELSEVIER

Contents lists available at ScienceDirect

## International Journal of Infectious Diseases

journal homepage: [www.elsevier.com/locate/ijid](http://www.elsevier.com/locate/ijid)

## Medical Imagery

## Motile microfilaria captured by fluorescent microscopy and the unmasking of eosinophilia following treatment

Gerrit Kann<sup>1,2</sup>, Hermann Juling<sup>3</sup>, Valentina Ilievski<sup>3</sup>, Gerrit Burger<sup>1,4,5</sup>, Stephan Göttig<sup>3</sup>, Christoph Stephan<sup>1</sup>, Nils Wetzstein<sup>1,\*</sup><sup>1</sup> Department of Internal Medicine, Infectious Diseases, University Hospital Frankfurt, Goethe University, Frankfurt am Main, Germany<sup>2</sup> Infektiologikum, Frankfurt am Main, Germany<sup>3</sup> Institute of Medical Microbiology and Infection Control, University Hospital Frankfurt, Goethe University, Frankfurt am Main, Germany<sup>4</sup> Institute of Tropical Medicine, University of Tübingen, Tübingen, Germany<sup>5</sup> Centre de Recherches Médicales de Lambaréné (CERMEL), Lambaréné, Gabon

## ARTICLE INFO

## Article history:

Received 6 June 2023

Revised 29 June 2023

Accepted 10 July 2023

## Keywords:

Loa loa

Filariasis

Microfilaremia

Migrant health

Neglected tropical diseases

NTDs

## ABSTRACT

A 24-year-old patient from Cameroon presented to our hospital because of a foreign structure in her left eye. To our knowledge, for the first time, fluorescent microscopy revealed motile microfilariae, and the diagnosis of loiasis was established. Despite substantial microfilaremia, eosinophilia only unmasked after the initiation of antiparasitic therapy.

© 2023 The Author(s). Published by Elsevier Ltd on behalf of International Society for Infectious Diseases.

This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>)

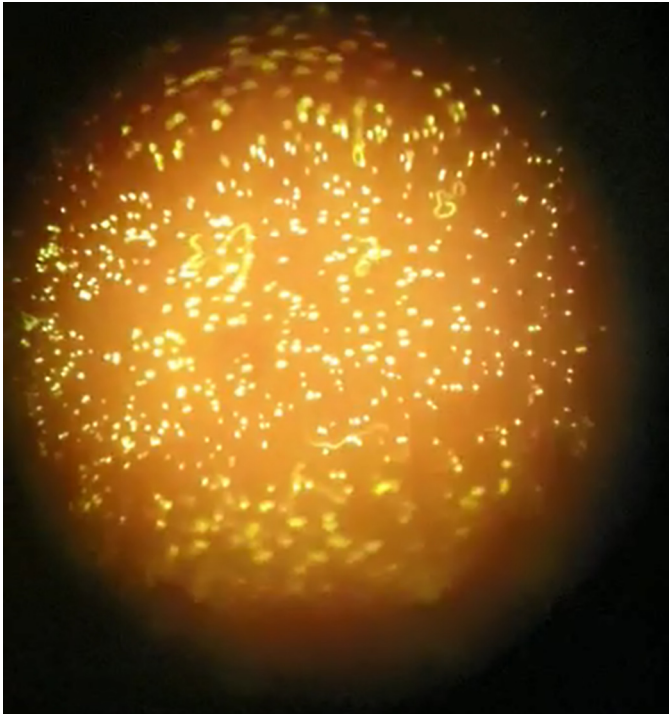
## Explanatory text

A 24-year-old female Cameroonian patient presented to our Infectious Disease Center because of a foreign structure in her left eye (Figure S1). A slightly elevated eosinophil count (0.59/nl) was observed at admission. To rule out parasitic bloodstream infection, fluorescent microscopy was performed and revealed motile structures, identified as live microfilariae (Video 1). This was confirmed in the wet mount microscopy and subsequent examinations of blood smears stained with Giemsa. Diagnosis of microfilaremia loiasis with an initial microfilarial density of 1200/ml was established. After 2 weeks of therapy with albendazole 400

mg qd, an increase in the eosinophil count (Figure S2) was observed [1]. The regimen was changed to diethylcarbamazine on day 32 and continued for another 3 weeks leading to the complete resolution of symptoms and the successive normalization of blood smears. Loiasis is a neglected filarial disease caused by *Loa loa* (the African eyeworm) mainly occurring in Central and Western Africa [2]. In the past, albendazole therapy has been linked to inflammation-mediated pathologies [3]. Interestingly, even in patients with substantial microfilarial counts—especially those from endemic countries—immune reaction in the form of eosinophilia can be unremarkable and may only be unmasked after the initiation of antiparasitic therapy [4,5].

\* Corresponding author.

E-mail address: [nils.wetzstein@kgu.de](mailto:nils.wetzstein@kgu.de) (N. Wetzstein).



**Video 1.** Fluorescent microscopy (quantitative buffy coat [QBC]; malaria tubes, QBC Diagnostics, Philipsburg, USA) and wet mount microscopy of blood smears showing motile microfilariae.

#### Declarations of Competing Interest

Gerrit Kann, Hermann Juling, Valentina Ilievski, Gerrit Burger, Stephan Göttig, and Nils Wetzstein have nothing to disclose. Christoph Stephan has received fees for lectures and/or advisory

board services from Gilead Sciences, Janssen-Cilag, MSD & ViiV Healthcare, and travel grants for scientific conference attendance from Gilead, Janssen, and AbbVie.

#### Funding

This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

#### Author contributions

GK, HJ, VI, and SG performed microbiological diagnostics and filmed the presented videos. NW treated the patient. GK and NW wrote the initial draft of the manuscript. GK, HJ, VI, GB, SG, CS, and NW critically revised the manuscript.

#### Supplementary materials

Supplementary material associated with this article can be found, in the online version, at [doi:10.1016/j.ijid.2023.07.006](https://doi.org/10.1016/j.ijid.2023.07.006).

#### References

- [1] Klion AD, Massougbdji A, Horton J, Ekoué S, Lanmasso T, Ahouissou NL, et al. Albendazole in human loiasis: results of a double-blind, placebo-controlled trial. *J Infect Dis* 1993;**168**:202–6. doi:[10.1093/INFDIS/168.1.202](https://doi.org/10.1093/INFDIS/168.1.202).
- [2] Loiasis Boussinesq M. *Ann Trop Med Parasitol* 2006;**100**:715–31. doi:[10.1179/136485906X112194](https://doi.org/10.1179/136485906X112194).
- [3] Herrick JA, Metenou S, Makiya MA, Taylor-Williams CA, Law MA, Klion AD, et al. Eosinophil-associated processes underlie differences in clinical presentation of loiasis between temporary residents and those indigenous to loa-endemic areas. *Clin Infect Dis* 2015;**60**:55–63. doi:[10.1093/cid/ciu723](https://doi.org/10.1093/cid/ciu723).
- [4] Volpicelli L, De Angelis M, Morano A, Biliotti E, Franchi C, Gabrielli S, Mattiucci S, Di Bonaventura C, Taliani G. Encephalopathy in a patient with loiasis treated with albendazole: A case report. *Parasitol Int* 2020;**75**:102006. doi:[10.1016/j.parint.2019.102006](https://doi.org/10.1016/j.parint.2019.102006).
- [5] Klion AD, Massougbdji A, Sadeler BC, Ottesen EA, Nutman TB. Loiasis in endemic and nonendemic populations: immunologically mediated differences in clinical presentation. *J Infect Dis* 1991;**163**:1318–25. doi:[10.1093/infdis/163.6.1318](https://doi.org/10.1093/infdis/163.6.1318).