

Pustular rosacea secondary to *Demodex* infestation shown with reflectance confocal microscopy



Joel Cohen, BS,^a Katharine Hanlon, BFA,^{b,c} Lilia Correa, MD,^{b,c} and James Grichnik, MD, PhD^b

Key words: confocal; *Demodex*; rosacea.

CLINICAL PRESENTATION

The patient, a 75-year-old male with a history of large granular lymphocytosis and common variable immunodeficiency, presented to dermatology with a 2-year history of erythematous, constantly pruritic pustules on his scalp and face that he reports intermittently became warm and inflamed with a pus-filled “whitehead.” (Fig 1) The patient described achieving no symptomatic improvement with topical clobetasol but endorsed some improvement in pruritus with cold compresses. The patient was prescribed topical 1% ivermectin cream, which resulted in some symptomatic improvement at 4-week follow-up.

CONFOCAL MICROSCOPY APPEARANCE

The lesions were examined using the VivaScope 3000 (Caliber Imaging and Diagnostics) reflectance confocal microscope (RCM). Imaging performed in clinic proved follicular infestation with human *Demodex* mites. The handheld confocal exam allowed us to examine several areas on the scalp and forehead, with each area showing many mites in nearly every follicle. In one follicle, 20 individual mites were counted (Figs 2 and 3, Video 1, available on www.jaad.org). Upon follow-up after 4 weeks using ivermectin, the patient returned and was again examined with handheld RCM at several locations, showing no visible *Demodex* mites, and near complete resolution of his skin symptoms (Fig 4).

Abbreviation used:

RCM: reflectance confocal microscope



Fig 1. Clinical photograph of the patient's scalp lesions, pretreatment.

KEY MESSAGE

Human *Demodex* mites (*Demodex folliculorum* and *Demodex brevis*) have become increasingly implicated in cases of folliculitis refractory to standard dermatologic treatment.¹ These species of obligate human ectoparasites are nearly ubiquitous, with infection rates increasing starkly after puberty, as sebaceous units develop, and approaching 100% prevalence by age 70 in the United States.² Demodicosis has been shown to mimic a variety of eruptive pathologies including various subtypes of

From the Morsani College of Medicine, University of South Florida, Tampa, Florida^a; Department of Dermatology & Cutaneous Surgery, University of South Florida, Morsani College of Medicine, Tampa, Florida^b; and Moffitt Cancer Center, Tampa, Florida.^c

Funding sources: None.

IRB approval status: Not applicable.

Patient consent: Consent for the publication of all patient photographs and medical information was provided by the authors at the time of article submission to the journal stating that all patients gave consent for their photographs and medical information to be published in print and online and with the understanding that this information may be publicly available.

Correspondence to: Joel Cohen, BS, Department of Dermatology & Cutaneous Surgery, University of South Florida, 12901 Bruce B. Downs Blvd, MDC 79, Tampa, FL 33612-4742. E-mail: cohen53@usf.edu.

JAAD Case Reports 2023;36:32-3.

2352-5126

© 2023 by the American Academy of Dermatology, Inc. Published by Elsevier, Inc. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

<https://doi.org/10.1016/j.jdcrr.2023.03.022>

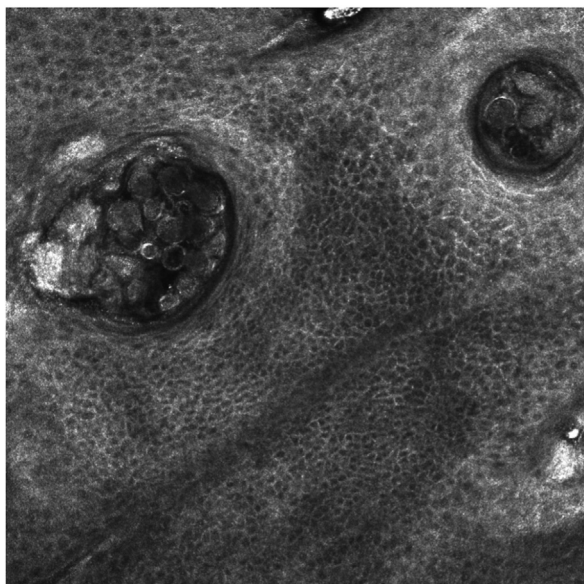


Fig 2. Reflectance confocal microscopy still image of *Demodex* mites within our patient's scalp lesion.

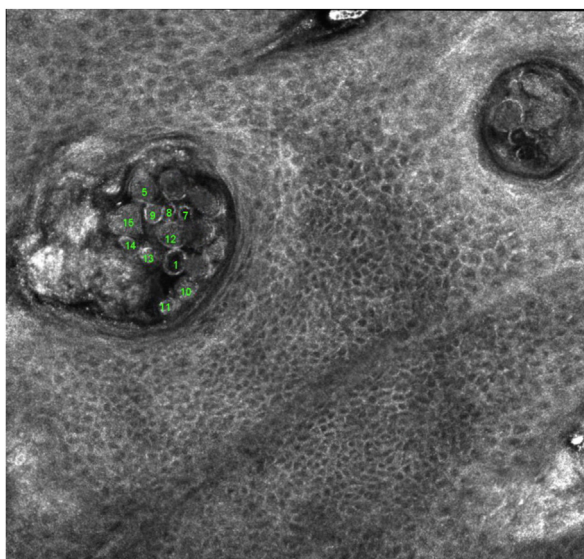


Fig 3. Reflectance confocal microscopy video counting 20 intralesional *Demodex* mites in vivo.

seborrheic, perioral, and contact dermatoses, leading to ineffective treatments often chosen based on a misinterpretation of etiology. Studies have supported the notion that the likelihood of developing a symptomatic infestation increases with increasing *Demodex* density as well as in the context of immunosuppression.³ The mainstay of treatment and symptomatic relief for patients with demodicosis includes the topical application of 1% ivermectin, a medication with a purported dual mechanism of



Fig 4. Clinical photograph of the patient's scalp after 4 weeks of treatment.

action that includes antiinflammatory and antiparasitic properties.⁴

RCM is a powerful noninvasive clinical tool that has been shown to allow in vivo cutaneous imaging and diagnosis and to reduce unnecessary biopsies.⁵ RCM offers dermatologists the ability to identify patterns of cell morphology and potential parasite infestation efficiently and accurately at varying depths of the epidermis to upper dermis. In cases of demodicosis, the benefit of RCM is not limited to diagnosis but can also be used to quickly monitor real-time treatment response at clinical follow-up appointments, using intralesional *Demodex* density and subjective symptomatology as markers of disease severity.

Conflicts of interest

Dr Grichnik notes that he serves as a consultant to Galileo Group and Canfield Scientific; serves on Skin Advisory Board for Regeneron and Dermatology Advisory Council for Melanoma Research Foundation; and receives clinical trial support from Novartis, Eli Lilly, Dermira, Elorac, Boehringer, and Amgen. Dr Correa is a consultant for Accutex Blades and a researcher and consultant for Novartis Pharmaceutical. Dr Cohen and Author Hanlon have no conflicts of interest to declare.

REFERENCES

1. Sattler EC, Hoffmann VS, Ruzicka T, Braunmühl TV, Berking C, et al. Reflectance confocal microscopy for monitoring the density of *Demodex* mites in patients with rosacea before and after treatment. *Br J Dermatol*. 2015;173(1):69-75.
2. Elston CA, Elston DM. *Demodex* mites. *Clin Dermatol*. 2014; 32(6):739-743.
3. Rather PA, Hassan I. Human *demodex* mite: the versatile mite of dermatological importance. *Indian J Dermatol*. 2014;59(1):60-66.
4. Del Rosso JQ. Topical ivermectin: data supporting dual modes of action in rosacea. *J Clin Aesthet Dermatol*. 2017;10(9):39-42.
5. Pellacani G, Farnetani F, Ciardo S, et al. Effect of reflectance confocal microscopy for suspect lesions on diagnostic accuracy in melanoma: a randomized clinical trial. *JAMA Dermatol*. 2022; 158(7):754-761.