

LETTER - RESEARCH

First case of *Trichophyton indotineae* in Brazil: clinical and mycological criteria and genetic identification of terbinafine resistance[☆]

Dear Editor,

Terbinafine-resistant dermatophytes are currently a global health problem, particularly *Trichophyton indotineae*. Terbinafine, a potent antifungal agent against dermatophytes, inhibits the enzyme squalene epoxidase (SQLE), restricting fungal growth by interfering with ergosterol biosynthesis; point mutations in the SQLE genes are the main cause of antifungal resistance.¹ *T. indotineae*, formerly known as *Trichophyton mentagrophytes* variety VIII, frequently shows mutations in the SQLE gene.^{2,3} The present report describes the first case of a patient diagnosed in

Brazil with dermatophytosis caused by terbinafine-resistant *T. indotineae*.

A 40-year-old male individual, without comorbidities, originally from Brazil and living in London, reported pruritic erythematous-desquamative lesions on the lower limbs and buttocks that began in January 2024 (Fig. 1). He reported frequent short trips during the second half of 2023; to Austria, Slovakia, Hungary, and Poland in August, and to Scotland and Turkey in November and December. He sought dermatological care in Piracicaba (state of São Paulo, Brazil), where direct mycological examination revealed the presence of hyaline septate hyphae, and the culture using Sabouraud agar and Mycosel® showed growth of *T. mentagrophytes*. He was diagnosed with dermatophytosis and in March 2024 he was prescribed 500 mg/day of terbinafine for 14 days.

Although there was no clinical improvement, the patient returned to London; in May 2024 he came back presenting lesions in the same locations (Fig. 2). On this occasion,



Figure 1 Clinical aspect of dermatophytosis lesions caused by *Trichophyton indotineae* affecting the posterior region of the lower limbs.

[☆] Study conducted at the Dermatology Clinic, Hospital da Santa Casa de São Paulo, São Paulo, SP, Brazil.



Figure 2 Clinical aspect of dermatophytosis recurrence caused by *Trichophyton indotineae* affecting the anterior surface of the right thigh.

itraconazole 200 mg/day was prescribed for 14 days, with complete clinical remission. However, the patient developed recurrence after treatment was discontinued, and fluconazole 150 mg/day was prescribed for seven days, which proved ineffective. With a new cycle of treatment using itraconazole at the same dosage, the patient showed the same result: good initial response followed by recurrence four days after treatment was discontinued. A skin scraping specimen was collected for new mycological analysis, and treatment with itraconazole was prescribed again. The condition improved, the patient returned to England and once again was lost to dermatological follow-up. In this scenario of (i) disseminated dermatophytosis refractory to terbinafine but susceptible to itraconazole, (ii) microbiological evidence suggestive of the *T. mentagrophytes*/*T.*

interdigitale species complex, and (iii) history of frequent international travel, it was strongly suggested that this was a case of *T. indotineae*. This suspicion was confirmed using the material from the second collection, identifying the isolate as *T. indotineae* resistant to terbinafine and fluconazole through analysis of mycological exams (Fig. 3) associated with DNA sequencing of the internal transcribed spacer (ITS) region of ribosomal DNA. The *SQL*E gene was also amplified and sequenced using the described primers.⁴ The sequences were deposited in GenBank under access numbers PQ634380 (*T. indotineae*) and PQ655447 (*SQL*E). The sequences used are shown in Fig. 4. In addition, an antifungal susceptibility test for terbinafine, fluconazole, and itraconazole was performed using the *in vitro* broth microdilution reference method described by EUCAST (E.DEF 9.4).

The assessed isolate is resistant to terbinafine and fluconazole, with minimum inhibitory concentration (MIC) values of $\geq 16 \mu\text{g/mL}$ (upper limit value) and $8 \mu\text{g/mL}$, respectively; and susceptible to itraconazole (MIC value of $0.064 \mu\text{g/mL}$). Sequencing results revealed two terbinafine resistance mutations (Phe³⁹⁷Leu and Thr⁴¹⁴His).

In the last decade, *T. indotineae* has caused large outbreaks of severe and difficult-to-treat infections worldwide. Lesions may be atypical with multiple morphologies, including concentric erythematous, desquamative, papulosquamous and pustular plaques, in addition to conditions modified by the use of topical corticosteroids.⁵

Cases of terbinafine-resistant *T. indotineae* described are often introduced by immigrants from endemic countries.^{5,6} The high rate of inter-human transmission is a strong contributor to its spread, where familial cases account for about 50% of patients, and sharing of fomites is a common denominator.⁵⁻⁸ However, few cases have been reported to date, mainly due to misidentification and underreporting.⁶ This may be the scenario in Brazil, where terbinafine-resistant dermatophytosis may be overlooked, since the etiological identification of dermatophytes remains a challenge, as DNA sequencing is not routinely used in the diagnosis of superficial mycoses.

The emergence of terbinafine-resistant *T. indotineae* is noteworthy, considering its frequency of up to 75% compared to 44% for *T. rubrum*.^{9,10} This phenomenon may be



Figure 3 Mycological examinations of *Trichophyton indotineae*. (A) Direct microscopic examination (with 10% KOH) under optical microscopy ($\times 400$) showing branched septate hyaline hyphae and arthroconidia. (B) Macromorphology of fungal culture in Sabouraud medium showing velvety white front and light yellow pigment on the back. (C) Micromorphology under optical microscopy ($\times 400$), stained with lactophenol blue, showing the presence of numerous pyriform and clavate microconidia and septate, spindle-shaped macroconidia.

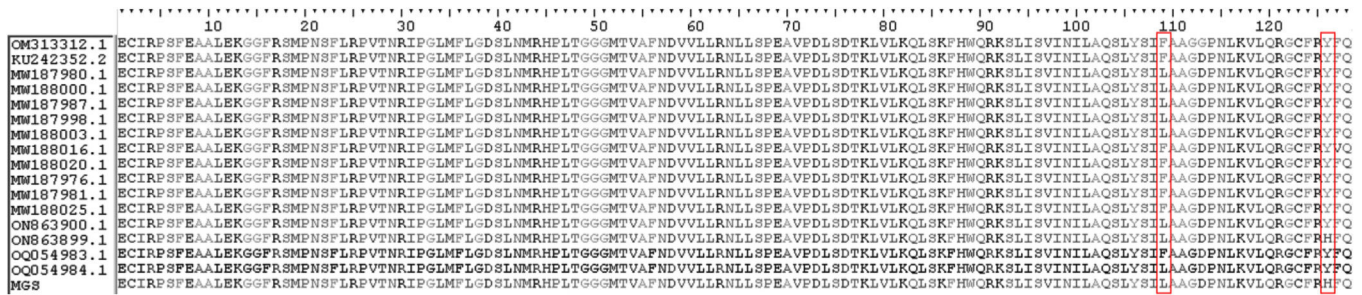


Figure 4 SQLE gene alignment under CLUSTAL multiple sequence alignment in BioEdit. The amino acid sequences of SQLE from the *T. indotineae* isolate IMT-1778 (MGS) were compared with the reference sequence of the *T. mentagrophytes* strain TIMM2789 (GenBank acc. number KU242352.) and the *T. interdigitale* isolate DK-Tinterdig-WT (GenBank acc. number OM313312.1), as well as SQLE sequences of terbinafine-resistant *T. indotineae* strains (GenBank acc. numbers MW187976, MW187980, MW187981, MW187987, MW187998, MW188000, MW188003, MW188016, MW188020, MW188025, ON863900, ON863899, OQ054983 and OQ054984). The amino acid substitutions that were found to be different in IMT-1778-MGS isolate are depicted, and their positions are shown in red boxes.

linked to (i) inappropriate use of antibiotics, antifungals, and corticosteroids; (ii) climate change and indiscriminate use of pesticides; and (iii) the return of intense migratory movements seen after the COVID-19 pandemic.^{7,8}

In summary, the present case is the first report of dermatophytosis caused by *T. indotineae* in Brazil, with the typical evolution of therapeutic resistance to several antifungals and terbinafine resistance associated with mutations in the SQLE gene. Phenotypic and genotypic characterizations were essential for adequate diagnosis and therapeutic choice, but terbinafine resistance complicates treatment options and highlights the need for better surveillance, prevention strategies, and alternative therapeutic approaches.

Financial support

None declared.

Authors' contributions

John Verrinder Veasey: Design and planning of the study; collection of data, or analysis and interpretation of data; drafting and editing of the manuscript or critical review of important intellectual content; effective participation in research orientation; intellectual participation in the propaedeutic and/or therapeutic conduct of the studied cases; critical review of the literature; approval of the final version of the manuscript.

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Conflicts of interest

None declared.

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Received 29 November 2024; accepted 15 January 2025

Available online 24 February 2025

<https://doi.org/10.1016/j.abd.2025.01.001>

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***In vitro* characterization of biofilm produced by *Fusarium oxysporum*, an onychomycosis agent[☆]**

Dear Editor,

Onychomycosis caused by Non-Dermatophyte fungi (NDMs), such as *Fusarium* spp., is more prevalent than previously thought, especially in warmer climates.¹ Furthermore, onychomycosis has currently been attributed to fungi organizing themselves into a biofilm form.² Biofilm is a complex microbial community, highly adhered to the nail and surrounded by a matrix that provides protection and antifungal resistance.^{2,3}

The research group has been studying the genus *Fusarium* spp. as an agent of onychomycosis in immunocompetent hosts. The authors reported its high prevalence in the studied region, established clinical and laboratory criteria for this genus as a causal agent of onychomycosis, and determined the susceptibility profile to the systemic antifungals most commonly used in Brazil.⁴ Later, the authors proved that *Fusarium* spp. uses nail keratin as a single source of nutrients⁵ and began studies on the etiopathogenesis of fusarial onychomycosis based on an *ex vivo* model using sterile human nail fragments.³

More recently the authors reported, for the first time, that *Fusarium oxysporum* is able to form biofilm on the human nail as the only nutritional source.^{6,7} Also, the

authors describe the volatile organic molecule 2-Ethyl-1-Hexanol (2EH) as a quorum-sensing component capable of modulating such biofilm.⁸ These findings were relevant to confirm the etiopathogenesis of fusarial onychomycosis. However, it did not reveal the proper characteristics of the biofilm formed under nutritional support. Thus, the current study aimed to characterize the *in vitro* biofilm formation of *F. oxysporum*, evaluating its natural ability, during 7-days with controlled nutrient availability.

This study was conducted with *F. oxysporum* CMRP2925 isolated from a previously described onychomycosis case.⁴ The isolate was reactivated to confirm its purity and identification, before assays, and was cultured on Sabouraud Dextrose Agar (SDA; DifcoTM, MI, USA) for 7-days at 25 °C. Biofilms were prepared according to Galletti et al.,⁹ with some modifications. A suspension containing 1 × 10⁷ conidia mL⁻¹ was prepared in RPMI Medium 1640 (Gibco, NY, USA), with L-glutamine, sodium bicarbonate, 0.165 M 3-(N-morpholino) propanesulfonic acid (pH 7.2), and 2% glucose. This suspension was placed into 96-well flat-bottomed microtitration plates and incubated at 35 °C in a shaker at 110 rev min⁻¹, for 7-days. Every 24 h, the culture medium was renewed by removing 100 µL of old broth and adding the same volume of fresh RPMI. During the seven days, biofilms were evaluated under different aspects, as previously described.^{6,9} Briefly, cell viability was assessed by counting Colony Forming Units (CFU), quantification of total biomass by Crystal Violet, metabolic activity by the reduction assay of tetrazolium salt, 2,3-(2-methoxy-4-nitro-5-sulphophenyl)-5-([phenylamino]carbonyl)-2H Tetrazolium hydroxide (XTT), characterization of the Extracellular Matrix (ECM), and visualization of biofilm structure by Scanning Electron Microscopy (SEM).

[☆] Study conducted at the Universidade Estadual de Maringá, Maringá, PR, Brazil.