



Cat-transmitted sporotrichosis by *Sporothrix brasiliensis:* Focus on its potential transmission routes and epidemiological profile

Fernanda de Andrade Galliano Daros Bastos^{® 1,*}, Marconi Rodrigues de Farias², Isabella Dib Ferreira Gremião³, Francelise Bridi Cavassin^{® 4}, Fabiana dos Santos Monti², Rogério Rodrigues Vilas Boas⁴, Bruno Paulo Rodrigues Lustosa⁵, Emanuel Razzolini⁵, Vânia Aparecida Vicente^{® 5} and Flávio Queiroz-Telles⁶

¹Universidade Federal do Paraná, Curitiba, 80000-000, Brazil

²Pontifícia Universidade Católica do Paraná, Curitiba, 80000-000, Brazil

³Oswaldo Cruz Foundation, Rio de Janeiro, 20000–23799, Brazil

⁴Faculdades Pequeno Príncipe, Curitiba, 80000-000, Brazil

⁵Federal University of Paraná, Basic Pathology Department, Curitiba, 80000-000, Brazil

⁶Federal University of Paraná, Department of Public Health, Curitiba, 80000-000, Brazil

*To whom correspondence should be addressed: Fernanda de Andrade Galliano Daros Bastos, M.Sc., Federal University of Paraná, 181, Gen. Carneiro Street. CEP 80060-90, Curitiba – PR, Brazil. Tel: +55 41996154105; E-mail: fernanda-daros@hotmail.com

Abstract

Cat-transmitted sporotrichosis (CTS) by *Sporothrix brasiliensis* is an important epizoonosis with alarming numbers of cases involving felines, canines, and humans. Considering the increasing incidence of CTS this study sought to elucidate the epidemiological characteristics of cats with sporotrichosis and evaluate the potential transmission routes of *S. brasiliensis* in several biological samples from cats with sporotrichosis. Samples were collected from ulcerated skin lesions, front paws, nasal cavity, and droplets collected from sick cats during sneezing episodes in a veterinarian university hospital, in the city of Curitiba, southern Brazil, between June 2021 and April 2022. A total of 100 cats with sporotrichosis were enrolled in the study. The fungus was isolated in 60% of samples from the nasal cavity and 71% of respiratory droplets. The growth of *S. brasiliensis* on the right and left front paw was observed in 41% of the cats included, and in 38%, there was growth of the fungus even without an apparent lesion on the paw. Of the infected cats, 64% had multifocal lesions throughout the body. The identification of *S. brasiliensis* in samples of exudate, paws, nasal cavity, and sneeze droplets suggests that transmission can occur not only through classic routes, but also through the movement of the infected cat, as well as through respiratory droplets expelled by the cat sneezing or nasal drip. Furthermore, the presence of *S. brasiliensis* on the paws of sick cats indicates the possibility that the fungus is being disseminated in the environment in which the animal lives.

Lay Summary

Cat-transmitted sporotrichosis, caused by *Sporothrix brasiliensis*, is an emerging fungal disease that has become a major public health concern in Brazil. *Sporothrix brasiliensis* has been isolated from several biological samples, suggesting non-classical routes of transmission of sporotrichosis.

Key words: Sporothrix brasiliensis, sporotrichosis, zoonotic infection, transmission route, public health.

Introduction

Sporothrix brasiliensis, a thermo-dimorphic fungus, is the most prevalent agent of epi-zoonotic sporotrichosis in Brazil and its emerging in South American bordering countries.^{1–4} In Brazil, Cat-transmitted sporotrichosis (CTS) affects thousands of humans, domestic cats and dogs.^{5–9}

The most common transmission routes include the traumatic transcutaneous or transmucosal implantation of the yeast cells through scratches and bites.¹⁰ Exudates from feline ulcerative mucocutaneous lesions may harbor a huge yeast burden and be capable of infecting the ocular mucosa even without a history of trauma.^{11,12} Recently, Bastos et al.¹³ identified that respiratory samples from cats with sporotrichosis are also potential sources of transmission of *S. brasiliensis* respiratory droplets created by a sneeze contain viable *Sporothrix* yeast that can infect humans and other animals that have experienced mucocutaneous exposure. Other studies also highlight the isolation of *Sporothrix* spp. on the paws of apparently healthy cats, which may indicate a possible source of transmission from colonized paws.^{14,15}

Since it became a notifiable disease in the state of Paraná (SESA Resolution No. 93/2022), in 2024 the number of confirmed cases, according to epidemiological data from the Paraná State Health Department (SESA), reached 4816, 932, and 50, in cats, humans, and in dogs, respectively. Other regions in Brazil also reported outbreaks like Rio de Janeiro, Rio Grande do Sul, Minas Gerais, Espirito Santo, and São Paulo.^{6,16–20} In recent decades, CTS has also had wide geographic expansion in countries in Latin America with few cases already reported in Europe and North America.^{1,21,22}

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Figure 1. Sequence for collecting biological samples from a cat with sporotrichosis. (A) Exudate from an ulcerated lesion collected by sterile swab; (B) Paw imprint in culture medium/Mycosel agar; (C) Secretion collected by sterile swab introduced into the nasal cavity; and (D) Collection of droplets/respiratory secretions expelled during sneezing.

This study intended to contribute to expanding knowledge about the epidemiological aspects of the disease in felines, which indirectly helps in understanding the spread of human cases in the state. Furthermore, identifying other possible transmission routes through the analysis of biological materials from cats helps us to understand human cases in which there is no inoculation of the fungus through the classic routes of biting and scratching. Thus, the study aimed to elucidate the epidemiological characteristics of infected cats and evaluate the potential transmission routes of *S. brasiliensis* in several biological samples from cats with sporotrichosis in the city of Curitiba, Paraná State, Brazil.

Materials and methods

Study design

This was a prospective study that included cats treated at a veterinary teaching hospital in Curitiba, southern Brazil, between June 2021 and April 2022 with suspected sporotrichosis. The inclusion criteria were domestic cats with clinical and/or epidemiological suspicion of sporotrichosis, presenting at least one skin lesion.

Animal sample collection

For each cat included in the study (n = 100), different biological materials were collected: ulcerated skin lesions (n = 100), right and left front paws (n = 200), nasal cavity (n = 100), and droplets expelled by sneezing (n = 28).

The skin lesions and nasal cavities were collected using a sterile swab (Fig. 1A,C). A direct imprint was made on the Mycosel agar culture medium from the right and left front paws regardless of the presence of lesions (Fig. 1B). For respiratory secretions droplets expelled during sneezing, a My-

cosel agar plate (Becton Dickinson - BD) was placed in front of the animal's nostrils, and a sterile swab was used to stimulate sneezing (Fig. 1D).

Data source and data collection

Epidemiological data were collected through a structured questionnaire, covering detailed information such as gender, age, weight, neighborhood of residence, lifestyle, reproductive status, as well as aspects such as contact with other animals, plants, and soil.

According to the location and number of lesions, clinical forms were classified into fixed forms, lymphocutaneous, disseminated form, and ocular forms. To collect the exudate samples, at least one lesion was required along the length of the animal's body (unifocal) or multiple lesions spread across the body (multifocal). To collect the paws and nasal cavity the animal could have lesions or not (Supplemental Content 1).

Laboratory tests

Cytopathological was performed only on ulcerated skin lesions, while fungal culture was performed on all types of specimens (ulcerated skin lesions, nasal cavity, paws, and sneeze). In cats with multiple skin ulcers, cytopathology and culture were collected from the same lesion and the most affected region. For cytological evaluation, the slides were fixed and stained using a Romanowsky-type stain (Rapid Panothic-Laborclin) to visualize fungal structures. Mycosel agar was utilized for fungal culture. All samples were incubated at 25°C–30°C until 30 days. Once the colonies had grown, they were identified by micromorphology and molecular sequencing. Isolated strains were identified by micromorphology as *Sporothrix* spp. according to the Atlas of clinical fungi.²³

Molecular analysis

Sporothrix isolates were identified at species level by partial Calmodulin sequence. DNA extraction was performed by using a cetyltrimethylammonium bromide protocol,²⁴ followed by PCR amplification using primers CL1 (5'-GAR TWC AAG GAG GCC TTC TC-3') and CL2A (5'-TTT TTG CAT CAT GAG TTG GAC-3'), and by sequenced amplicons using a BigDye Terminator v3.1 Cycle Sequencing Kit (Applied Biosystems). Consensus sequences were produced using BioEdit v 7.2.5, and alignment was performed using MAFFT online interface. Finally, phylogenetic reconstructions were done in MEGA v7.0.26, using maximum likelihood with 1000 bootstrap replicates.

All isolates with molecular characterization were deposited in the Microbiological Collection of Parana Network/Taxonline (https://www.cmrp-taxonline.com/) (Supplemental Content 2).

Statistical analysis

To analyze the epidemiological and demographic profile of the animals, a descriptive statistical analysis was performed. For data related to the results of biological sample collections, Microsoft Excel® 365 and Minitab[®] 19 spreadsheets were used. For inferential analysis, Pearson's χ^2 distribution was applied for association, and the value of P < .05 was considered statistically.

Ethics statement

This study was approved by the Ethics Committee on Research in the Use of Animals of the Pontificia Universidade Católica do Paraná under the registration number 01691/2019.

Results

A total of 100 cats were enrolled in the study with confirmed sporotrichosis by culture. The disease was more prevalent in male cats (66%), aged between 1 and 2 years (49%), and average weight of 3.97 kg (range 1.6–7.6). Most of the cats were not castrated (57%). In 77% of cases, the owners had other animals in the same residence. Of those, 66% were cats, 10% were dogs, and a single guardian had a domestic rabbit (n = 1). In 25% of the cases, the cat had contact with another sick cat in the same residence. Additional information and details are presented in Table 1.

Feline sporotrichosis was reported in 31/75 neighborhoods in the city of Curitiba and 4/29 municipalities in the city's metropolitan region (Fig. 2). In the analysis, 57% of the cats were indoors with outdoor access.

Isolation methods and characterization of the strains

A total of 100 cytological examination and 428 culture specimens were done to confirm the sporotrichosis infection in the cats. Fungal culture demonstrated growth of filamentous colonies, with membranous and white aspects in 56% of collected samples (n = 240) (Fig. 3A). Cytologic analyses showed presence of oval and elongated yeastlike cells in a cigar form that are compatible with *Sporothrix* spp. in 50% of cases (Fig. 3B). Microscopic analysis showed hyaline, thin, septate, and branched hyphae, with unicellular, hyaline or brown coni-

 Table 1. Demographic characteristics of 100 cats with sporotrichosis in Curitiba, Brazil.

Demographic variables	TOTAL N(%)	
Age (years)		
< 1	7 (7%)	
1–2	49 (49%)	
3–4	21 (21%)	
5-6	17 (17%)	
7-8	6 (6%)	
Sex	. ,	
Male	66 (66%)	
Female	34 (34%)	
Neutering		
Yes	38 (38%)	
No	57 (57%)	
Not informed	5 (5%)	
Cat's lifestyle		
Indoor	35 (35%)	
Indoor with outdoor access	57 (57%)	
stray cats	8 (8%)	
Contact with other sick cats		
Yes	25 (725%)	
No	75 (75%)	
Contact with soil and plants		
Yes	86 (86%)	
No	14 (14%)	



Figure 2. Distribution of feline sporotrichosis cases in the city and metropolitan region of Curitiba/Brazil between 2021 and 2022.

dia, globose to ovoid, resembling a daisy flower, compatible with genus *Sporothrix*(Fig. 3C).

From the 28 samples collected from respiratory secretions eliminated by sneezing, 20 (71%) showed growth of *Sporothrix* spp. Regarding culture collected from the nasal cavity, *Sporothrix* spp. was observed in 60% (60/100) of sam-



Figure 3. (A) Macromorphology on Mycosel agar of *Sporothrix* in filamentous phase; (B) Impression smear from skin ulcer showing numerous yeast cells (1000x magnification); and (C) Micromorphology of *Sporothrix* in the filamentous phase (400x increase).

ples, of that, the fungal culture was positive in 68% of cats with sporotrichosis nasal cavity lesion (34/50) and in 52% of samples collected from cats without any nasal abnormities (26/50) (P = .102, Table 2).

Considering the cultures of samples collected from the front paws (right and left), *Sporothrix brasiliensis* growth was observed in 40% (81/200) of them. Of the cats that had no apparent lesions on their paws, the fungal culture was positive in 38% (67/178) of the samples, while cats with paw lesions had a positive culture in 64% (14/22) of the cases (P < .05, Table 2).

Regarding the distribution of skin lesions, 36% of the cats had unifocal lesions and 64% of the cats had multifocal lesions. Fungal culture has higher isolation in cats with multifocal lesions (92%, 59/64) when compared with unifocal lesions (56%, 30/36) (Table 2).

Of the skin lesion samples, 79% (n = 100) have a culture positivity, of that, 96% (n = 48) have both positivity for culture and cytological examination. However, in 62% (n = 31) of the cases, positivity was observed only in the culture samples with negative result for the cytological examination.

Of the 100 animals in the study, 74% (74/100) were analyzed by molecular identification. The phylogenetic reconstruction of the calmodulin gene sequence clustered all isolates, with *S. brasiliensis*, identified the species as the only etiological agent (Fig. 4). Of these, 28.4% were from cat sneezes (21/74), 12.2% were from lesions on the cat's right or left paw (9/74), 17.6% from nasal lesions (n = 13/74), and 41.9% from other sporotrichosis lesions (31/74) (Supplemental Content 2).

Discussion

The present study provides insights into the potential transmission routes of *Sporothrix brasiliensis* to humans, demonstrating that even without apparent skin lesions in some felines, the fungus was isolated and may play an important role in the transmission of sporotrichosis. Furthermore, it describes geographic and epidemiological information about cats treated in Curitiba and nearby.

A higher prevalence was observed in unneutered male cats of reproductive age and with free access to the street. These data corroborate with data already published which suggests that cats with the habit of going outside are more susceptible to becoming infected than cats, that don't use go out to fight for food, territory, or by the female.^{6,25,26} In another study, it was observed that the chances of *Sporothrix* positivity in animals that live partially at home are 3.02 times greater than in cats that do not have access to the street.²⁷

In this study, we observed that 25% of animals may become infected after having contact with another sick cat in the same residence. It is noteworthy from these data that guidance to the owner must be clear and objective, highlighting that the animal diagnosed with sporotrichosis must be isolated from the others until clinical cure, which can vary from weeks to months (median time 4 months).^{28,29} In the study by Paiva et al.,³⁰ the authors observed that the distance between animals is linked to cases of spread of the fungus, so it is essential to keep the cat in a separate environment, to minimize direct contact with other animals and humans.

Almost all infected animals had contact with soil and plants, which could be a potential route of environmental transmission. In the study by Lecca et al.,²⁷ there was no statistically significant association between environmental variables and a positive diagnosis for sporotrichosis. Macêdo-Sales et al.²⁵ also evaluated contact with plants and soil and the results demonstrated that 72% of cats that had contact with soil and/or plants had a positive culture, a lower percentage than cats' healthy people in contact with the environment. Nahal et al.,³¹ 49% of human patients reported trauma involving plants and/or contact with soil, with S. brasiliensis being the predominant species. For the authors, non-zoonotic sporotrichosis is associated with infections acquired after traumatic inoculation with plants and/or contact with soil. Although the possibility of a classic route of transmission from soil and plants should not be ruled out, studies appear to support the hypothesis that direct transmission from infected cats plays a more important role in the transmission and ongoing outbreaks of sporotrichosis in Brazil.^{27,32}

Cats with ulcerated lesions, mainly containing exudate, are more susceptible to transmitting *Sporothrix*, either through direct contact or through wiggling (the act of shaking the body), affecting mostly mucous membranes, including the Table 2. Isolation of Sporothrix spp. from different clinical samples.

Variable (N)		Positive culture (%)	Negative culture (%)	<i>P</i> -value
Droplets from the sneeze (28)		20 (71%)	8 (29%)	
Nasal cavity (100)				.102
	With injury (50)	34 (68%)	16 (32%)	
	Without injury (50)	26 (52%)	24 (48%)	
Paws (200)			× 2	.019*
	With lesions (22)	14 (64%)	8 (36%)	
	Without lesions (178)	67 (38%)	111 (62%)	
Distribution skin lesions (100) Unifoca Multifo				<.001*
	Unifocal (36)	20 (56%)	16 (44%)	
	Multifocal (64)	59 (92%)	5 (8%)	
Culture total (428)		240 (56,1%)	188 (43,9%)	
Cytology results (100)				<.001*
	Positive (50)	48 (96%)	2 (4%)	
	Negative (50)	31 (62%)	19 (38%)	

Legend: $* \chi^2$ test.

ocular region.^{10,33} In this context, this study demonstrated highly positive culture in exudate specimens, which indicates a high risk of transmission occurring through this material. Besides, the main clinical manifestation observed in this study was disseminated forms, with multifocal lesions, therefore has an increased risk of transmitting fungus when compared with cats that present a single lesion (unifocal/fixed cutaneous). It should be emphasized that the care taken when handling an animal that presents one or multiple injuries must be the same, highlighting the use of personal protective equipment by the handler, isolation of the animal, hygiene of the environment, and adequate treatment.

Nasal lesions are frequent manifestations in cats with sporotrichosis, and may have an ulcerated appearance, with the presence of blood, pus, or crust or just a nasal elevation (nasal nodule). In those cats, we observed that *S. brasiliensis* was isolated in nasal swab in most of them, however, it was intriguing, half of the cats that did not present any lesion or nasal elevation had positive culture, indicating a potential risk of transmission, which could be eliminated through respiratory droplets from sneezing or through dripping secretions.¹³

The first study that proposed sneezing as a new transmission route was conducted by the same authors,¹³ however, the fungal species had not been investigated. In this study, we identify that all samples from respiratory secretions expelled by a cat's sneeze are *S. brasiliensis*. This is the main study that reports a possible new route of transmission of sporotrichosis, and with the molecular identification completed, it is highlighted that human cases, especially the ocular forms, deserve attention for diagnosis and appropriate treatment.

Cats with sporotrichosis that did not have lesions suggestive of sporotrichosis on their paws tested positive for *Sporothrix* in 38% of the samples. Souza et al.,¹⁴ the authors isolated *Sporothrix* from the paws of 29.1% of healthy cats that lived with sick cats, demonstrating the importance of clinically healthy cats carrying the fungus on their paws. In another study carried out in Peru, where there are no reports of zoonotic transmission, the authors also isolated *S. schenckii* from the nails of healthy cats (2.38%), demonstrating the importance of investigation in the feline population even in non-endemic areas.¹⁵ Other studies on the investigation of *Sporothrix* in the paws or claws of cats demonstrate a lower frequency of positivity in animals without lesions suggestive of sporotrichosis in this location.^{25,29,34,35} Although there is no proof of transmission from paws colonized by *Sprothrix*, the results found in this study demonstrate the importance of research on the feline population, mainly for guidance on the care of sick cats.

We also evaluated the results of cytopathology and fungal culture. Although culture is the gold standard method, cytopathology is a good diagnostic screening resource to detect Sporothrix yeast cells with sensitivity between 52.6% and 87%.^{36,37} In our study, negative cytopathological results but with positive culture (62%) may be associated with the initiation of treatment with compounded itraconazole in some cats included in the research, which could reduce the sensitivity of the method, however, studies show that compounded itraconazole is not bioequivalent to the reference medicine therefore would not be an effective treatment for feline sporotrichosis.^{38,39} Cytopathology is a quick, simple, and low-cost technique, it is advisable that the veterinarian performs this examination in cases where sporotrichosis is suspected, however, it is important to associate culture when possible because the sensitivity of the method is greater.^{25,36}

According to the molecular analysis, all isolates were identified as *S. brasiliensis*, the agent is commonly associated with cat transmitted sporotrichosis in Brazil as reported previously.³² Those strains with near-complete calmodulin gene sequencing, showed a high degree of similarity with previously sequenced isolates from Rio de Janeiro, Curitiba, and Paraguay.^{2,8,32} Multi-typing analysis based on several isolates from Brazil indicates that zoonotic transmission from *S. brasiliensis* is based on clonal and recombinant *Sporothrix* species,⁴⁰ in addition, molecular clock typing show that multiple zoonotic introductions happened during the last decades, but the epidemiological strains from Curitiba match with the strains from Rio de Janeiro epicenter of the disease.⁴¹

The main limitation of this study was not evaluated if there was transmission of sporotrichosis from cats included and their tutors, although the owners who reported lesions were referred for medical evaluation at a reference center.

The cases of sporothrichosis in felines in the city of Curitiba was highest in the Campo Comprido district, which has a total population of almost 27 000 inhabitants, an area of 8,55 km², and with most of the population living in houses.⁴² A region that also presented a significant number of cases in felines was the Industrial City of Curitiba. According to Cognialli et al.,⁸ this location also has a higher incidence of human cases. This neighborhood borders Campo Comprido and is also mostly made up of houses. The animal with sporotrichosis acts as

	Sporochrix brasiliensis CMRP5497
	Sporounix brasiliensis CMR0611/
	Sporoannx brasiliensis CMRP5501
	Sporounix brasiliensis CMRP6137
	sporodninx brasiliensis CMRPS859
	Sporochnx brasiliensis CMRP5517
	Sporochrix brasiliensis CMRP6128
	Sporochnx brasiliensis CMRP6119
	Sporochnx brasiliensis CMRP5502
	Sporochnx brasiliensis CMRP6140
	Sporochrix brasiliensis CMRP6155
	Sporochnx brasiliensis CMRP6164
	Sporochnx brasiliensis CMRP5506
	Sporochnx brasiliensis CMRP5860
	Sporochnx brasiliensis CMRP6136
	Sporochnx brasiliensis CMRP5518
	Sporochnx brasiliensis CMRP5509
	Sporochnx brasiliensis CMRP5519
	Sporochnx brasiliensis CMRP5857
	Sporochnx brasiliensis CMRP6132
	Sporochnx brasiliensis CMRP6131
	Sporochnx brasiliensis CMRP6122
	Sporochrix brasiliensis CMRP6142
	Sporochnx brasiliensis CMRP6154
	Sporochnx brasiliensis CMRP6138
	Sporochrix brasiliensis CMRP5513
	Sporothmx brasiliensis CMRP5504
	Sporothmx brasiliensis CMRP6157
	Sporochrix brasiliensis CMRP6116
	Sporothmx brasiliensis CMRP6115
	Sporothmx brasiliensis CMRP6139
	Sporochrix brasiliensis CMRP6147
	Sporochrix brasiliensis CMRP5862
	Sporochrix brasiliensis CMIRP5842
	Sporochrix brasiliensis CMRP5508
	Sporochnx brasiliensis CMRP6152
	Sporochnx brasiliensis CMRP6124
83	Sporochnx brasiliensis CMRP5856
	Sporochnx brasiliensis CMRP5498
	Sporochnx brasiliensis CMRP6135
	Sporochnx brasiliensis CMRP5512
	Sporochnx brasiliensis CMRP5496
	Sporochrix brasiliensis CMRP5503
	Sporochnx brasiliensis CMRP5499
	Sporochnx brasiliensis CMRP5851
	Sporochnx brasiliensis CMRP6129
80	Sporochrix brasiliensis CMRP6161
	Sporochnx brasiliensis CMRP6165
	Sporochnx brasiliensis CMRP6156
8	Sporochnx brasiliensis CMRP5511
	Sporochrix brasiliensis CMRP5863
	Sporothrax brasiliansis CMRP5500
	Sporochnx brasiliensis CMRP6121
	Sporothrax brasiliansis CMRPS844
	Sporothrax brasiliensis CMRP6125
	Sporochnx brasiliensis CMRP6130
	91 Sporothrix brasilensis CBS 133003
	Sporochrix brasiliensis CMRP5507
	Sporothrix brasilensis CBS 132985
	Snorothrity brasilensis I CBS 133004
	Snorothriv hrasiliansis I CBS 132994
	Shorothur brasiliansis I CMRP6133
	Sporthur bretlancic CMPD5505
	Sporodula biosilienais CMRP300
	Sportburk brokinensis CMRP6126
	Sporodina biosinensis Chirche 141
	Sporochrix brasiliensis CMRP6120
	Sporoennx brasiliensis CMRP6168
	Sporoennx brasiliensis CMRP6123
98	Sporodnitk brasiliensis CMRP6144
	Sporochrix brasiliensis CMRP6127
	83 Sporochnx brasiliensis CMRP6112
	84 Sporodnix brasiliensis CMRP6149
	Sporochrix brasiliensis CMRP6134
	Sporochrix brasiliensis CMRP6137
	Sporothrix brasillensis CBS 120339 (T)
	Sporochrix brasiliensis CMRP6114
	Sporochrix brasiliensis CMRP6146
	Sporochrix brasiliensis Scaniclass globosa CBS 132923
	os Sporothrix globosa CBS 132924
	Sporothrix globosa CBS 130104
100	Sporothrix globosa CBS 120340 (T)
	Sporothrix luner (CBS 937.72 (T)
	Spore they set and LODD 100075
98	sporounix schencki CBS 1329/5
	Sporoznix Schencki CBS 132976
	- Sporothrixschencki CBS 359.36 (T)
	- Sporothrix schenckii (CBS 359.36 (T) 100 Sporothrix brunneo vib Bicea CBS 10 1570
	- Sporothrix schencki CBS 359.36 (T) 100 Sporothrix brunneo vb Bcee CBS 10.1570 Sporothrix brunneo vb Bcee CBS 793.73 (T)

0.06

Figure 4. Phylogenetic tree of the *Sporothrix* pathogenic clade based on calmodulin near-complete, genes constructed with maximum likelihood implemented in MEGA 70.26. Bootstrap values > 80 from 1000 resampled data sets. *Sporothrix branneviolacea* (CBS 793.73) was taken as outgroup. *Sporothrix brasiliensis* is represented inside the red box, *S. schenckii* is represented inside the yellow box, *S. globosa* is inside the green box, and *S. luriei* is inside the purple box. New strains isolates from the presented study are marked in bold.

a sentinel for possible new human and animal cases, which is why disease prevention and control actions must be linked not only to human cases but also to cats from a one-health perspective.³⁰ These results can assist in prevention and control actions in areas with a greater number of human and feline cases. CTS is an endemic and neglected disease in continuous expansion not only in Brazil, and there are still several gaps on the infection. Therefore, the present study contributes mainly with possible new routes of zoonotic transmission and unpublished data on epidemiology in the city of Curitiba, Brazil.

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Author contributions

Fernanda de Andrade Galliano Daros Bastos (Conceptualization, Data curation, Formal analysis, Funding acquisition, Investigation, Methodology, Project administration, Resources, Supervision, Validation, Visualization, Writingoriginal draft, Writing-review & editing), Marconi Rodrigues de Farias (Data curation, Investigation, Methodology, Resources, Supervision, Validation, Writing-review & editing), Isabella Dib Ferreira Gremião (Data curation, Methodology, Supervision, Validation, Writing-review & editing), Francelise Bridi Cavassin (Data curation, Formal analysis, Methodology, Supervision, Validation, Visualization, Writing-review & editing), Fabiana dos Santos Monti (Data curation, Resources, Supervision), Rogério Rodrigues Vilas Boas (Data curation, Software, Writing-review & editing), Bruno Paulo Rodrigues Lustosa (Data curation, Methodology, Software, Writing-review & editing), Emanuel Razzolini (Data curation, Methodology, Software, Writingreview & editing), Vânia Aparecida Vicente (Data curation, Funding acquisition, Methodology, Resources, Software, Validation, Writing-original draft, Writing-review & editing), and Flávio Queiroz-Telles (Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Project administration, Software, Supervision, Validation, Visualization, Writing-original draft, Writing-review & editing)

Supplementary material

Supplementary material is available at *Medical Mycology* online.

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Declaration of interest

The authors have no relevant financial or non-financial interests to disclose.

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