# Diagnostic and therapeutic challenges in canine sporotrichosis: a case report

Leonardo Sena Santos<sup>1</sup>, Arianne Costa Baquião<sup>2</sup>

<sup>1</sup>Veterinarian.

<sup>2</sup>Universidade Paulista, São Paulo, SP, Brasil

# **ABSTRACT**

#### **OBJECTIVE**

To describe a case of sporotrichosis in a female dog treated at the Veterinary Teaching Hospital in São Paulo, Brazil, with emphasis on clinical signs, diagnosis, treatment, and prognosis.

#### **METHOD**

This study is a longitudinal descriptive case report of a mixed-breed female dog, rescued by her owner, presenting with cutaneous and respiratory manifestations. The animal was attended at the Internal Medicine Department of the Veterinary Hospital of Universidade Paulista, São Paulo, SP, Brazil. Information was obtained through review of the medical records, diagnostic reports, laboratory findings, and clinical observations. The study was approved by the institutional animal ethics committee under protocol number 8295071222.

# **RESULTS**

The patient exhibited non-painful, non-pruritic ulcerative lesions, hyperkeratosis, and alopecia on the rostral region and hind limbs, along with nasal discharge and sialorrhea. Cytopathology, histopathology, radiography, and blood and biochemical analyses were performed but proved inconclusive. Due to lack of improvement after antibiotic and corticosteroid therapy, a fungal culture was conducted, confirming infection by Sporothrix spp. An antifungal susceptibility test revealed intermediate resistance to itraconazole. Nonetheless, itraconazole was selected for treatment at a dosage of 10 mg/kg once daily for 180 days. After this period, both cutaneous and respiratory lesions resolved completely, and a 30-day extension of therapy was recommended to prevent relapse and ensure full therapeutic success.

# **CONCLUSION**

Sporotrichosis should always be considered in the differential diagnosis of ulcerated and non-pruritic skin lesions in dogs; and for diagnosis and implementation of treatment and control measures for this subcutaneous mycosis, fungal culture and antifungal sensitivity tests are essential.

### **KEYWORDS**

Ulcerative lesion; Subcutaneous mycosis; Sporothrix spp.

## Corresponding author:

Arianne Costa Baquião

Faculty Member, School of Veterinary Medicine, Universidade Paulista - UNIP

R. Rua General Leite de Castro, 201 - Jardim Santa Cruz, São Paulo - SP, Brasil

E-mail: arianne.baquiao@hotmail.com

ORCID ID: https://orcid.org/0000-0003-3921-9256

**Copyright:** This is an open-access article distributed under the terms of the Creative Commons.

Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided that the original author and source are credited.

### DOI:



#### INTRODUCTION

Sporotrichosis is a subcutaneous mycosis caused by saprophytic, thermodimorphic fungi, which belong to a phylogenetically related group known as *Sporothrix schenckii* complex. This complex includes four clinically relevant species- *S. schenckii* sensu stricto, *S. brasiliensis*, *S. globosa*, and *S. luriei*- and two others, *S. mexicana* and *S. albicans*, occasionally isolated from humans. Among these, *S. schenckii* sensu stricto and *S. brasiliensis* are considered the most significant species associated with sporotrichosis. While *S. schenckii* sensu stricto has a global distribution and moderate virulence; *S. brasiliensis* emerged in the state of Rio de Janeiro, remains geographically restricted to Brazil and it is considered the most pathogenic species within the complex. 1,2

This mycosis can cause cutaneous, lymphocutaneous, subcutaneous, and systemic lesions; and is primarily transmitted via traumatic inoculation of fungal spores by fomites or scratches, especially from felines.<sup>3</sup>

Sporotrichosis is an anthropozoonosis characterized by insufficient public health policies and limited attention to preventive strategies.<sup>4</sup> Outbreaks have been reported globally, and in Brazil, both human and animal cases have originated primarily in the state of Rio de Janeiro, and subsequent spread to other regions.<sup>5</sup>

Although *Sporothrix schenckii* complex strains affect various mammalian species, particularly cats, sporotrichosis is often underdiagnosed in other animals, as dogs and humans.<sup>6</sup> In dogs, the disease is rare and sporadic, and the cutaneous form is the most frequent. This form presents variable and nonspecific clinical signs, including multiple firm nodules-ulcerated or not-, alopecia and painless, moderated pruritic ulcers, mainly located on the ears, head and body.<sup>3</sup> Disseminated nodular lesions may also appear, often associated to hemorrhagic or hemopurulent scabs and are usually associated to immunosuppression.<sup>7</sup>

Fungal elements are rarely found in dogs and canine diagnosis of sporotrichosis is challenging.<sup>8</sup> Diagnosis is primarily established by isolation and identification of *Sporothrix spp*. in culture, supplemented by cytopathology, histopathology, skin testing, serology, immunohistochemistry, and molecular assays. Fungal culture is considered the gold standard for definitive diagnosic elucidation.<sup>9</sup>

The most common treatment for canine sporotrichosis involves itraconazole<sup>10</sup>, although other antifungal protocols include ketoconazole, potassium iodide, terbinafine, and amphotericin B.<sup>10-13</sup> In Brazil, the first-line treatment is itraconazole at 10 mg/kg, administered until 30 days after complete clinical remission and negative mycological testing.<sup>14-15</sup>

Thus, this study aimed to describe a case of sporotrichosis in a dog and to highlight the key diagnostic and therapeutic aspects observed at a Veterinary Teaching Hospital in the city of São Paulo.

#### **METHODS**

This study is a case report presented in a descriptive and qualitative format. The information was obtained by a review of the medical record, laboratory reports, diagnostic tests, and clinical data provided by the guardian. The study was approved by the Animal Research Ethics Committee of Universidade Paulista (UNIP), under approval number 8295071222.

The data collected were recorded in a descriptive and chronological form, and subsequently analyzed and compared for discussion purposes.

## **RESULTS AND DISCUSSION**

The patient was rescued from the streets and had no previous clinical history available. It presented ulcerated, crusted, non-painful and non-pruritic lesions, as well as hyperkeratosis in the rostral region, accompanied by alopecia on the face, abdominal and pubic regions, and on the thoracic and pelvic limbs, particularly on the caudal surface of the hind limbs. Additionally, the patient exhibited sneezing, serous nasal discharge, ptyalism, and tick infestation (Figure 1).

Figure 1 - Initial lesions in the pubic region, pelvic limbs, and rostral area.







Source: Authors' archive (2025).

Complementary exams were requested, including complete blood count, alkaline phosphatase, alanine aminotransferase (ALT), urea, creatinine, serology for ehrlichiosis, lymph node cytology, and histopathology of the lesion area. Based on the clinical history and physical findings, the initial differential diagnoses included neoplasia and leishmaniasis. However, test results were inconclusive, and treatment was started with topical antiparasitics, oral antibiotic therapy (doxycycline 50 mg every 12 hours for 14 days), topical rifamycin spray on lesions every 12 hours for 14 days, and corticosteroid therapy with prednisone (10 mg every 24 hours until further recommendation).

After 14 days of treatment, with worsening of the cutaneous and respiratory conditions, antibiotic was replaced by amoxicillin asociated to potassium clavulanate (175 mg every 12 hours for 21 days). Mycological culture was requested and, after 14 days, identified Sporothrix spp., along with antifungal susceptibility testing, which showed complete resistance to amphotericin B and fluconazole, intermediate resistance to itraconazole, and sensitivity to clotrimazole, econazole, ketoconazole, miconazole, and nystatin. Treatment was initiated with itraconazole (100 mg every 24 hours for 180 days), administered after meals. The animal was reevaluated monthly for clinical monitoring, and gradual lesion remission was observed, leading to complete clinical resolution (Figure 2). A second fungal culture was performed and returned negative. Therefore, medication was continued for an additional 30 days to prevent recurrence. The guardian was instructed to isolate the animal and use gloves during handling to reduce zoonotic transmission risk.15

Figure 2 - Resolution of skin lesions after treatment.







Source: Authors' archive.

In this report, the initial cutaneous lesions - characterized by ulcerations with hemopurulent and hemorrhagic sacbs - required differential diagnosis with diseases such as leishmaniasis, algomycosis, mycobacteriosis, cryptococcosis, neoplasms, nocardiosis, and sporotrichosis. <sup>15</sup> A conclusive diagnosis was only possible after mycological culture, in agreement with Schubach et al. <sup>3</sup>, who stated that the isolation and identification of *Sporothrix spp*. in culture, based on morphological parameters, remains the gold standard for definitive diagnosis. The diagnostic failure after cytology and histopathology was observed in this case, and it reinforces importance of mycological culture, due the typically low fungal load in canine lesions, which can lead to treatment delays and compromise patient outcomes. <sup>8</sup>

Until diagnosis was confirmed through fungal culture, the patient was treated with corticosteroids (prednisone) and antibacterial agents (doxycycline and later amoxicillin with clavulanate) for 28 days. Corticosteroid therapy was gra-



dually discontinued, and antibiotics replaced by antifungal treatment with itraconazole. The immunosuppressive effect of corticosteroids should be avoided during and after treatment, as clinical recurrence of sporotrichosis has been reported even six months after apparent clinical cure. <sup>16</sup> Moreover, systemic antibiotic therapy should only be prescribed in conjunction with antifungal treatment and in cases of suspected secondary bacterial infection<sup>11</sup>, as inappropriate antibiotic use does not eliminate the fungus and may contribute to antimicrobial resistance- a global public health concern affecting both animal and human populations. <sup>17</sup>

In this study, sporotrichosis treatment was conducted using itraconazole monotherapy. The Sporothrix strain isolated showed intermediate resistance to itraconazole, which led to a prolonged prolonged therapeutic approach with high doses until clinical and laboratory confirmation of cure. <sup>18-21</sup> Galhardo et al. <sup>22</sup> demonstrated that Brazilian Sporothrix strains remain susceptible to itraconazole, while Gremião et al. <sup>23</sup> reported that combination therapy with potassium iodide may be an alternative in cases of therapeutic failure. The drug was not administered concomitantly with H<sub>2</sub> receptor antagonists, which increased its absorption, as described by Reis et al. <sup>24</sup> Renal and hepatic functions were monitored regularly due to potential nephrotoxicity and hepatotoxicity associated with the drug. <sup>12</sup>

Furthermore, after the complete regression of the lesion, a second fungal culture was performed, which was crucial to consolidate the clinical and laboratory cure. This approach was also adopted in our study to ensure complete elimination of the fungus and therapeutic success against Sporothrix spp. infection. <sup>12,15,18,20</sup>

#### **FINAL CONSIDERATIONS**

Although sporotrichosis is not a notifiable disease, its importance should be emphasized to both the medical and veterinary communities due to its zoonotic transmission potential. Common in Brazil, sporotrichosis is an anthropozoonosis with increasing incidence both nationally and internationally. Therefore, it should always be considered in the differential diagnosis of ulcerated and non-pruritic skin lesions in dogs.

For an accurate diagnosis and subsequent implementation of effective treatment and control measures for this subcutaneous mycosis, fungal culture and antifungal sensitivity tests are essential. These tools allow the identification of the etiological agent and the guidance of therapeutic decisions. In these cases, itraconazole has been the antifungal of choice, presenting satisfactory results in clinical management, even in the face of intermediate resistance.

#### **REFERENCES**

- 1. Oliveira MM, Almeida-Paes R, Muniz MM, Gutierrez-Galhardo MC, Zancope-Oliveira RM. Phenotypic and molecular identification of Sporothrix isolates from an epidemic area of sporotrichosis in Brazil. Mycopathologia. 2011 Oct;172(4):257-67. doi: 10.1007/s11046-011-9437-3. DOI: 10.1007/s11046-011-9437-3.
- 2. Chakrabarti A, Bonifaz A, Gutierrez-Galhardo MC, Mochizuki, T, Li S. Global epidemiology of sporotrichosis. Med. Mycol. 2015, 53, 3-14.
  3. Schubach TM, Schubach A, Okamoto T, Barros MB, Figueire-
- 3. Schubach TM, Schubach A, Okamoto T, Barros MB, Figueire-do FB, Cuzzi T, et al. Canine sporotrichosis in Rio de Janeiro, Brazil: clinical presentation, laboratory diagnosis and therapeutic response in 44 cases (1998-2003). Med Mycol. 2006 Feb;44(1):87-92. DOI: 10.1080/13693780500148186.
- 4. Seyedmousavi S, Guillot J, Tolooe A, Verweij PE, Hoog GS. Neglected fungal zoonoses: hidden threats to man and animals. Clin Microbiol Infect. 2015 May;21(5):416-25. DOI: 10.1016/j.cmi.2015.02.031.
- 5. Rodrigues AM, Gonçalves SS, Carvalho JA, Borba-Santos LP, Rozental S, Camargo ZP. Current progress on epidemiology, diagnosis, and treatment of sporotrichosis and their future trends. J Fungi (Basel). 2022 Jul 26;8(8):776. DOI: 10.3390/jof8080776.
- 6. Araújo JMD, Mendes Junior AF, Mothé GB, Tanaka BMBS, Cardoso LF, Bruno NV, et al. Rinite fúngica por Sporothrix spp. em

- cão: Relato de caso. Res., Soc. Dev. 2023 12(4):18412441075. DOI: 10.33448/rsd-v12i4.41075.
- 7. Larsson CE, Lucas R. Tratado de medicina externa: dermatologia veterinária. 2016.
- 8. Cafarchia C, Sasanelli M, Lia RP, Caprariis D, Guillot J, Otranto D. Lymphocutaneous and nasal sporotrichosis in a dog from southern Italy: case report. Mycopathologia. 2007 Feb;163(2):75-9. DOI: 10.1007/s11046-006-0086-x.
- 9. Pereira AS, Gremião IDF, Menezes RC. Sporotrichosis in Animals: Zoonotic Transmission. *In*: Zeppone Carlos I. Sporotrichosis: new developments and future prospects. Switzerland: Ed. Springer International Publishing: Cham; 2015. p. 83-102. 10. Schubach TM, Menezes RC, Wanke B. Sporotrichosis. *In*: Greene CE. Infectious diseases of the dog and cat. Philadelphia: Ed. Saunders Elsevier; 2012 p. 645-650.
- 11. Orofino-Costa R, Freitas DF, Bernardes-Engemann AR, Rodrigues AM, Talhari C, Ferraz CE, et al. Human sporotrichosis: Recommendations from the Brazilian Society of Dermatology for the clinical, diagnostic and therapeutic management. An. Bras. Dermatol. 2022; 97:757-77. DOI: 10.1016/j. abd.2022.07.001.
- 12. Sméra DSR, Sméra FR. Protocolos terapêuticos para o tratamento da esporotricose felina refratária e inovações decorrentes das pesquisas. Revista ft.. 2024 Oct; 28 (139): 28-29. DOI: 10.69849/revistaft/ni10202410270928
- 13. Song Y, Zhong SX, Yao L, Cai Q, Zhou JF, Liu YY, et al. Efficacy and safety of itraconazole pulses vs. continuous regimen in cutaneous sporotrichosis. J Eur Acad Dermatol Venereol. 2011 Mar; 25(3):302-5. DOI: 10.1111/j.1468-3083.2010.03785.x.
- 14. Rosa ĆS, Meinerz ARM, Osório L G, Cleff MB, Meireles MCA. Terapêutica da esporotricose: revisão. Sci. Anim. Health. 2018 May; 5(3):212-28. DOI: 10.15210/sah.v5i3.11337.
- 15. Orofino-Costa R, Macedo PM, Rodrigues AM, Bernardes-Engemann AR. Sporotrichosis: an update on epidemiology, etiopathogenesis, laboratory and clinical therapeutics. An Bras Dermatol. 2017 Sep-Oct;92(5):606-620. DOI: 10.1590/abd1806-4841.2017279.
- 16. Whittemore JC, Webb CB. Successful treatment of nasal sporotrichosis in a dog. Can Vet J. 2007 Apr; 48(4):411-4. 17. Araújo BC, Melo RC, Bortoli MC, Bonfim JRA, Toma TS. Pre-
- 17. Araújo BC, Melo RC, Bortoli MC, Bonfim JRA, Toma TS. Prevenção e controle de resistência aos antimicrobianos na Atenção Primária à Saúde: evidências para políticas [Prevention and control of antimicrobial resistance in Primary Health Care: evidence for policies]. Cien Saude Colet. 2022 Jan;27(1):299-314. DOI: 10.1590/1413-81232022271.22202020.
- 18. Larsson, CE (2011). Esporotricose. Braz. J. Vet. Res. Anim. Sci., São Paulo, 48 (3):250-259. DOI: 10.11606/s1413-95962011000300010.
- 19. Bernardes-Engemann AR, Tomki GF, Rabello VBS, Almeida-Silva F, Freitas DFS, Gutierrez-Galhardo MC, Almeida-Paes R, Zancopé-Oliveira RM. Sporotrichosis Caused by Non-Wild Type Sporothrix brasiliensis Strains. Front Cell Infect Microbiol. 2022 May 27;12:893501. DOI: 10.3389/fcimb.2022.893501.
- 20. Kauffman CA, Bustamante B, Chapman SW, Pappas PG1. Infectious Diseases Society of America. Clinical practice guidelines for the management of sporotrichosis: 2007 update by the Infectious Diseases Society of America. Clin Infect Dis. 2007 Nov 15;45(10):1255-65. DOI: 10.1086/522765.
- 21. Zager VÅ, Santos LA, Malegoni ACS, Roque LZ, Silva TB, Risso FB, et al. (2021) Canine Sporotricosis: Clinic, Epidemiology, Diagnosis and Treatment. OALib. 8: e7406. DOI: 10.4236/oalib.1107406.
- 22. Galhardo MC, Oliveira RM, Valle AC, Paes RA, Silvatavares PM, Monzon A, et al. Molecular epidemiology and antifungal susceptibility patterns of Sporothrix schenckii isolates from a cat-transmitted epidemic of sporotrichosis in Rio de Janeiro, Brazil. Med Mycol. 2008 Mar;46(2):141-51. DOI: 10.1080/13693780701742399.
- 23. Gremião IDF, Martins SRE, Montenegro H, Carneiro AJB, Xavier MO, Farias MR, et al. Guideline for the management of feline sporotrichosis caused by Sporothrix brasiliensis and literature revision. Braz J Microbiol. 2021 Mar;52(1):107-124. DOI: 10.1007/s42770-020-00365-3.
- 24. Reis EG, Gremião ID, Kitada AA, Rocha RF, Castro VS, Barros MB, et al. Potassium iodide capsule treatment of feline sporotrichosis. J Feline Med Surg. 2012 Jun;14(6):399-404. DOI: 10.1177/1098612X12441317.

