

## Case Report/Relato de Caso

### Recurrent *Acremonium* infection in a kidney transplant patient treated with voriconazole: a case report

Infecção recidivante por *Acremonium* em paciente transplantado renal tratado com voriconazol: relato de caso

Felipe Francisco Tuon<sup>1</sup>, Carolina Pozzi<sup>2</sup>, Sergio Ricardo Pentead-Filho<sup>1</sup>, Ricardo Benvenuti<sup>2</sup> and Fabiana Loss de Carvalho Contieri<sup>2</sup>

#### ABSTRACT

*Acremonium* infection is rare and associated with immunosuppression. A case of recurrent cutaneous *Acremonium* infection after short term voriconazole use is described. Surgical resection was the definitive therapy. Oral voriconazole was used in the treatment of *Acremonium* infection, but recurrence was associated with short therapy. Prolonged antifungal therapy and surgical resection are discussed for the treatment of localized lesions.

**Key-words:** Voriconazole. *Acremonium*. Fungal infection.

#### RESUMO

Infecção por *Acremonium* é rara e pode estar associada com imunossupressão. Descrevemos um caso de infecção recorrente de pele por *Acremonium* após tratamento breve com voriconazol. Ressecção cirúrgica foi o tratamento definitivo. Terapia prolongada com antifúngicos e ressecção cirúrgica são discutidas para o tratamento de doenças fúngicas localizadas.

**Palavras-chaves:** Voriconazol. *Acremonium*. Infecção fúngica.

#### INTRODUCTION

*Acremonium* is a ubiquitously fungus present in the soil and human infection is extremely uncommon<sup>1,2</sup>. Dermatophytoses, keratitis and mycetomas are the most common clinical presentation. Immunosuppressed patients show a large clinical spectrum, including pneumonia, arthritis, osteomyelitis, endocarditis, peritonitis, meningitis and sepsis<sup>1,3,4</sup>. Immunosuppressed patients can also present localized cutaneous lesions<sup>1</sup>. A case of cutaneous lesion caused by *Acremonium* sp with recurrence after a short course of voriconazole is described.

#### CASE REPORT

A 47 year-old male farmer, living in Mabore (a southern city in Brazil) was admitted to hospital due to skin lesion in the thigh on March 22, 2009. The patient has been a kidney transplant patient since November 7, 2007, using mycophenolate mofetil, tacrolimus and

prednisone. The lesion, measuring 1.5cm in diameter, appeared two months before admission and presented purulent discharge (**Figure 1**). There was no history of trauma, the rest of the physical examination was normal and laboratorial tests were irrelevant. A skin biopsy showed a granulomatous reaction with giant cells, while Grocott stain showed phialide and phialoconidium (**Figure 2**). The culture was positive for *Acremonium* sp, though species identification was not performed.



FIGURE 1 - Cutaneous lesion due to *Acremonium* sp.

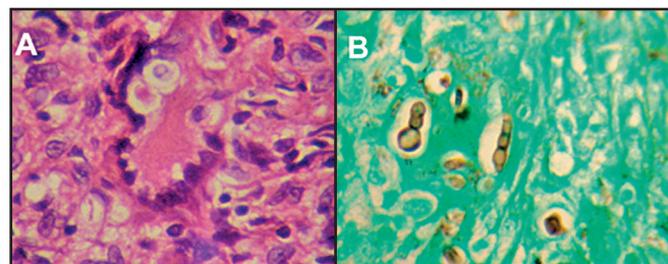


FIGURE 2 - Histological findings verified in a cutaneous lesion caused by *Acremonium* sp. A) Granulomatous infiltrate with giant cells and phagocytosis of yeast-like structures. B) Grocott stain showing phialide and phialoconidium suggestive of *Acremonium* sp.

Immunosuppressive therapy was suspended and voriconazole 200mg b.i.d. was used for 14 days. The lesion improved and drainage was suspended; however, after one month, the lesion returned and a new culture showed *Acremonium* sp with histological analysis showing the same aspect of the first biopsy. Surgical resection of the lesion was performed and immunosuppressive therapy was reintroduced. The lesion cured without recurrence after six months.

1. Division of Infectious and Parasitic Diseases, Hospital Universitário Evangélico de Curitiba, Curitiba, PR. 2. Division of Renal Transplant, Hospital Universitário Evangélico de Curitiba, Curitiba, PR.

**Address to:** Dr. Felipe Francisco Tuon. Infectious and Parasitic Diseases Clinic, Hospital Universitário Evangélico de Curitiba. Alameda Augusto Stelfeld 1908/3º andar, SCIH - Bigorrihlo, 80730-150 Curitiba, PR, Brazil.

Phone: 55 41 3240-5055; Fax: 55 41 3240-5274

e-mail: flptuon@gmail.com

Received in 08/12/2009

Accepted in 21/01/2010

## DISCUSSION

This case suggested that short therapy with voriconazole is inadequate for the treatment of cutaneous lesions caused by certain yeasts occurring on transplant patients. We believe that prolonged therapy should be used for transplant patients. The improvement of the lesion after two weeks of therapy demonstrated the efficacy of voriconazole against *Acremonium*, although no sensitivity test was performed. Nevertheless, short therapy (two weeks) was insufficient for fungus eradication from the skin. The fact that the cutaneous lesion was localized permitted a surgical resection approach.

A previous review summarized clinical experience with infections caused by species of *Acremonium*<sup>5</sup>. Two cases of cutaneous lesions have been published involving transplant patients. The first was described by Strabelli et al in a heart transplant patient<sup>6</sup>. This patient presented a lesion on the knee that was treated with surgery and local therapy. In another case, a subcutaneous infection in a kidney transplant patient was cured with surgical resection of the abscesses and ketoconazole<sup>7</sup>. In some instances, the particular species of *Acremonium* cannot be determined by morphology, as described in several reports and in this case<sup>5</sup>.

The drug of choice for treatment of *Acremonium* infections is amphotericin; however, the species presents low susceptibility to most antifungals, including imidazoles, fluorocytosine and amphotericin B. *In vitro* studies have shown that *Acremonium* can be susceptible to voriconazole, but resistant to other imidazoles<sup>8</sup>. Voriconazole has been used in some cases of *Acremonium* infections with success<sup>9,10</sup>. This patient received voriconazole due to renal graft and risk of renal failure.

In summary, *Acremonium* sp can cause skin lesions in kidney transplant patients. Voriconazole improved the lesion but recurrence occurred with short therapy. Considering the cost of new antifungal drugs, surgical resection is an adequate approach in these cases.

## ACKNOWLEDGMENTS

The authors would like to thank Marcelo Dorneles for the microbiological study.

## REFERENCES

1. Schell WA, Perfect JR. Fatal, disseminated *Acremonium strictum* infection in a neutropenic host. *J Clin Microbiol* 1996; 34:1333-1336.
2. Krcmery Jr V, Kunova E, Jesenska Z, Trupl J, Spanik S, Mardiak J, et al. Invasive mold infections in cancer patients: 5 years' experience with *Aspergillus*, *Mucor*, *Fusarium* and *Acremonium* infections. *Support Care Cancer* 1996; 4:39-45.
3. Guarro J, Gams W, Pujol I, Gene J. *Acremonium* species: new emerging fungal opportunists-*in vitro* antifungal susceptibilities and review. *Clin Infect Dis* 1997; 25:1222-1229.
4. Warris A, Wesenberg F, Gaustad P, Verweij PE, Abrahamsen TG. *Acremonium strictum* fungaemia in a paediatric patient with acute leukaemia. *Scand J Infect Dis* 2000; 32:442-444.
5. Fincher RME, Fisher JF, Lovell RD, Newman CL, Espinel-Ingroff A, Shadomy HJ. Infection due to the fungus *Acremonium* (*Cephalosporium*). *Medicine* 1991; 70:398-409.
6. Strabelli TM, Uip DE, Amato Neto V, Bocchi EA, Higuchi ML, Stolf NA, et al. *Acremonium* infection after heart transplant. *Rev Soc Bras Med Trop* 1990; 23:233.
7. Miro O, Fernando J, Lecha V, Campistol JM. Abscesos subcutaneos por *Acremonium* falciforme en un transplantado renal. *Med Clin (Barc)* 1994; 102:316.
8. Wildfeuer A, Seidl HP, Paule I, Haberreiter A. *In vitro* activity of voriconazole against yeasts, moulds and dermatophytes in comparison with fluconazole, amphotericin B and griseofulvin. *Arzneimittelforschung* 1997; 47:1257-1263.
9. Keynan Y, Sprecher H, Weber G. *Acremonium* vertebral osteomyelitis: molecular diagnosis and response to voriconazole. *Clin Infect Dis* 2007; 45: e5-e6.
10. Mattei D, Mordini N, Lo NC, Gallamini A, Osenda M, Pugno F, et al. Successful treatment of *Acremonium* fungemia with voriconazole. *Mycoses* 2003; 46:511-514.