Keratitis Caused by *Scedosporium apiospermum* Successfully Treated with a Cornea Transplant and Voriconazole

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A case of *Scedosporium apiospermum* keratitis was successfully treated with oral voriconazole and penetrating keratoplasty. Voriconazole levels in the aqueous humor were 53% of the levels in plasma and exceeded the MIC for the isolate by sevenfold.

CASE REPORT

A 47-year-old male presented with a painful corneal ulcer of his left eye, present for 2 weeks. Previously locally applied treatment with chloramphenicol, gentamicin, and tobramycin combined with dexamethasone had been ineffective. At presentation, an infiltrate about 2 mm in diameter was present in the central part of the corneal stroma, surrounded by stromal edema, partially covered with epithelium. The anterior chamber showed severe inflammation and a hypopyon (3 mm). The conjunctiva was heavily injected (Fig. 1a). The patient revealed that he was owner of a worm-breeding station and that he did not use contact lenses.

Corneal scrapings, obtained the same day, showed no bacteria, but possible fungal structures in the Gram stain were seen, which was confirmed by Blankophor P staining. Blankophor P is an optical brightener, similar to calcofluor white, with high affinity for β-glicosidically linked polysaccharides, and is used to trace glucan and chitin in the fungal cell wall (11). The scrapings were inoculated onto blood agar, brain heart infusion agar, and Sabouraud agar and broth and were incubated at 29°C. After 2 days of incubation, fungal colonies were cultured.

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The resected cornea button was cut in three pieces that were fixed in formalin and embedded in paraffin. Subsequently, hematoxylin-and-eosin staining and periodic acid-Schiff (PAS) staining were performed on 4-μm-thick paraffin sections. Microscopic examination revealed desquamation of large parts of the corneal epithelium and an acute inflammatory infiltrate with necrosis in the central anterior part of the corneal stroma, accompanied by multiple fungal elements that were highlighted in the PAS stain (Fig. 2a to c). The hyphae showed septation and branching. Some hyphae were found close to Descemet’s membrane. However, penetration of this membrane was not found, nor did the inflammatory infiltrate or fungal elements reach the peripheral rim of the cornea button.

The aqueous humor stained with Blankophor P showed no fungal structures, and cultures remained sterile. The corneal biopsy, however, showed abundant septate mycelium structures, and the culture was positive with *S. apiospermum*. Cultures for other pathogens, including *Acanthamoeba*, remained negative.

Itraconazole topical therapy was discontinued after 3 days following the corneal transplantation, since white crystals appeared at the anterior stroma at the site of the epithelial defect. These crystals were assumed to be itraconazole precipitates, as this drug is not water soluble. The crystal precipitate disappeared very slowly during the succeeding months. Oral voriconazole therapy (two loading doses of 6 mg/kg of body weight, followed by 4 mg/kg b.i.d.) was started 3 days after surgery and was continued for 3 months. Slow improvement of the eye occurred in the following months. The fungal infection...
did not reactivate in the donor cornea or in the remnants of the recipient cornea.

The in vitro activities of amphotericin B, voriconazole, miconazole, fluconazole, itraconazole, posaconazole, ravuconazole, and terbinafine against the strain were tested with a microdilution technique, according to NCCLS M38-P guidelines (8). After incubation for 70 to 74 h at 35°C, final reading of the microtitration plate was performed visually. The MIC endpoint for amphotericin B was defined as the lowest concentration at which there was 100% inhibition of growth. For voriconazole, miconazole, fluconazole, itraconazole, posaconazole, ravuconazole, and terbinafine, the MIC endpoint was 50% inhibition of growth, compared to that of the drug-free control. The minimum fungicidal concentration (MFC) was determined by plating 50 μl of the contents from each clear well onto Sabouraud dextrose agar plates. The MFC was defined as the lowest concentration at which ≥99% of the inoculum was killed (3). In addition, a two-dimensional checkerboard titration with serial dilutions with combinations of voriconazole and terbinafine was performed to determine if synergy between the two drugs existed. The fractional inhibitory concentration index was calculated with the method described by Walsh et al. (15). The MICs of the drugs, compared with those of other clinical S. apiospermum isolates from our laboratory, are shown in Table 1 (6). All drugs exhibited fungistatic activity against the isolate. There was no synergistic

FIG. 1. Left eye of patient. (a) On presentation. An infiltrate was present in the central part of the cornea and a hypopyon was visible in the anterior chamber. The conjunctiva was heavily injected. (b) Three days after presentation. Shown are increased corneal infiltrate and hypopyon in the anterior chamber, after treatment with amphotericin B and itraconazole eyedrops. Note the severe edema and hyperemia of the conjunctiva.

FIG. 2. (a) Composition of a cross-section of the cornea button; the darker areas in the cornea stroma represent inflammatory infiltrate with necrosis and fungi. Note that the periphery of the cornea does not show this change. (b) Shown is a higher magnification of the area indicated by the rectangle in panel a. It illustrates that some fungi (arrows) and inflammatory cells reach close to Descemet’s membrane (arrowheads); however, penetration of this membrane was not found. (c) Shown is the area indicated by the rectangle in panel b. The fungi appear as septated, branching hyphae with variable diameter. PAS staining is shown in all panels. Original magnifications, ×50 (a), ×100 (b), and ×400 (c).
interaction between voriconazole and terbinafine (fractional inhibitory concentration index, 1.03).

After 12 days of treatment with oral voriconazole, a puncture of the anterior chamber was performed and, at the same time, a blood sample was taken to determine the concentration of voriconazole. A bioassay agar diffusion method was used as described by Perea et al. (10). The assay was performed in duplicate for the aqueous humor and in triplicate with plasma. Inhibition zones were read after incubation for 16 h at 28°C. The calibration curve was plotted on a logarithmic scale to determine the concentrations in the samples. The mean voriconazole concentration in plasma was 3.4 μg/ml and in the aqueous humor was 1.8 μg/ml (53% of the level in plasma).

Environmental sampling was performed in order to trace the source of the infection. Cultures were obtained from materials related to the profession of the patient, including dry and wet feed, peat, cellulose, wheat, soil, and water. Furthermore, eyelid fluid and container were cultured, since the patient indicated that he wiped the snout of the eyelid container manually.

V. volvocis was not cultured from any of the materials.

*S. apiospermum* is a ubiquitous filamentous fungus in nature and can be recovered from soil, decaying matter, and polluted water (9, 16). It may cause localized and disseminated opportunistic infections in immunocompromised patients and in immunocompetent patients after massive exposure and trauma (16).

Keratitis caused by *S. apiospermum* is rare, although several cases have been reported (1, 13, 16). *S. apiospermum* keratitis is often associated with trauma or exposure to vegetation or other organic matter (9, 16). Keratitis caused by simultaneous *Acanthamoeba* and *S. apiospermum* infection, usually in patients with contact lenses, or by contact with polluted water has been reported (4, 12). In the present case, the patient had no history of trauma, but he had contact with many potential organic sources. Unfortunately, we were unable to identify the source of infection.

*Scedosporium* is known to be resistant to many antifungals, and *S. prolificans* is generally more resistant than *S. apiospermum* (2, 6). Our strain was inhibited by a concentration of 1 μg of itraconazole, posaconazole, ravuconazole, miconazole, or voriconazole per ml. The MICs are within the range of those found with other *S. apiospermum* strains previously tested in our laboratory (Table 1) (6) and those found in other studies (1, 5, 9). The antifungals exhibited fungistatic activity, which is in accordance with previous in vitro studies (2, 5). No synergistic activity between voriconazole and terbinafine was detected, although synergistic interaction of this combination has been shown in vitro previously in *S. prolificans* (7).

We found that, after oral administration of 200 mg of voriconazole b.i.d. for 12 days, the concentration in the aqueous humor was more than 50% of the concentration in plasma and almost seven times higher than the MIC for the strain. It has been shown that voriconazole concentrations in cerebrospinal fluid are also 50 to 70% of the levels in plasma (14). Furthermore, treatment with voriconazole was successful in patients with central nervous system aspergillosis and *S. apiospermum* brain abscess (9). In a recent study, voriconazole could be detected in the aqueous humor of rabbit eyes, after administration of 5-μg (50 μg/ml) or 10-μg (100 μg/ml) voriconazole eyedrops b.i.d. for 11 days. There was significant variability among individual animals, but the voriconazole concentration might still be high enough to treat fungal infection (17).

It is not clear yet which therapy is optimal to treat *S. apiospermum* infections. Usually, surgical intervention with adjuvant antifungal therapy is recommended (9). It is possible that transplantation of the cornea alone would have been sufficient to prevent further spread of the infection in our patient, because Descemet’s membrane was not perforated and because the periphery of the cornea was free of inflammation with fungi. However, since miconazole eyedrops were not available and since we were aware of the possible toxic effect of systemically administered miconazole and of the previously reported treatment failure of *S. apiospermum* corneal infections, we decided to administer voriconazole orally.

Voriconazole, together with penetrating keratoplasty, was effective for the treatment of *S. apiospermum* keratitis. High levels of the drug were found in the aqueous humor following oral administration, indicating a possible role for voriconazole in the treatment of fungal eye infections.

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**REFERENCES**