Vaginal mucormycosis: a case report

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Although Zygomycetes cause life-threatening, opportunistic infections in immunocompromised hosts, the first case of vaginitis caused by Mucor species in a healthy woman is reported. Mucor vaginitis, which caused mild symptoms only, was refractory to conventional azole therapy and resistant to flucytosine. Cure was achieved with topical amphotericin B.

Key words: vaginitis, fungal infection, amphotericin B

The vast majority of vaginal fungal infections are caused by Candida species, especially Candida albicans. Recently, attention has focused on an unconfirmed increase in vaginal infections caused by C. glabrata. Most microbiological investigations of large numbers of women with fungal vaginitis have reported infrequent symptomatic vaginitis caused by unusual species of Candida including C. tropicalis, C. parapsilosis, and C. krusei. Rarely, vaginitis caused by Saccharomyces cerevisiae has been reported. Although the Zygomycetes class of microorganisms are recognized as major pathogens in immunocompromised hosts, no mention in the literature has been made of vaginitis caused by these opportunistic pathogens.

CASE HISTORY

A 56-year-old female was referred because of vulvovaginal symptoms that had begun approximately 18 months previously, consisting predominantly of a stinging sensation in the vulva. Other symptoms included vulvovaginal itching, soreness, and discomfort. Because of chronic symptoms and positive 10% KOH microscopy, she had received repeated courses of topical (miconazole, clotrimazole and terconazole) and systemic azoles (fluconazole) as well as topical boric acid without any relief. On one occasion, a fungal culture obtained was positive for Mucor species. She had no abnormal discharge or dyspareunia and her husband was asymptomatic. Prior to the onset of symptoms, she had repeated attacks of bacterial cystitis, treated with antibiotics. Her past medical history was significant only for interstitial cystitis.

Examination demonstrated normal appearance of vulva and vestibule. The vagina contained fairly copious, pasty white discharge, which was not malodorous and was otherwise normal; the cervix was normal. Laboratory tests showed a vaginal pH of 4.2 with a negative amine test. Saline microscopy revealed large numbers of atypical blastoconidia and broad, non-septate hyphae branching at irregular intervals. A 10% KOH examination revealed large numbers of blastoconidia and hyphal elements consistent with Zygomycetes. Definitive diagnosis was established by culture, which showed Mucor indicus species. In vitro susceptibility tests showed that the vaginal isolate was sensitive to amphotericin B and resistant to flucytosine, imidazoles and triazoles. She was placed on topical therapy with 3% amphotericin B cream 5 g daily.
for 3 weeks. After an initial clinical response to amphotericin B, she had a relapse with minimal symptoms and cultures remained positive. She was retreated with 4 additional weeks of topical amphotericin B and then placed on a tapering alternate day dose for 2 months. Her symptoms of stinging and itching resolved and she has remained culture-negative and asymptomatic for a further 12-month follow-up period.

**COMMENTS**

Mucormycosis is a rare but commonly fatal invasive fungal infection caused by one of the ubiquitous saprophytic fungi of the Zygomycetes class. Species of the *Rhizopus*, *Mucor*, and *Absidia* genera of the Mucoraceae family are the organisms most commonly isolated. Mucormycosis is known to affect patients who are immunocompromised and usually neutropenic, and those with uncontrolled diabetes mellitus and ketoadiposis. Other risk factors include iron overload, desferrioxamine administration, wound contamination with soil, colonized adhesive tape, corticosteroid use and severe protein calorie malnutrition. Clinical features depend upon the organ involved and most commonly involve paranasal sinuses and lungs and less commonly the gastrointestinal tract. There are few reports of genitourinary mucormycosis in the literature. Genitourinary involvement is predominantly of the kidneys and has been found with systemic dissemination. Parenteral high-dose amphotericin B is the most effective treatment. No case of vaginal infection by Zygomyceetes has been published to date. The lack of response in the reported patient to treatment with topical azoles is not surprising, since these agents are inactive against *M. indicus* and the Mucoraceae in general. Similarly, flucytosine, which is useful in eliminatingazole-resistant *C. glabrata*, has no activity in vitro against Zygomyceetes. Amphotericin B remains the mainstay of therapy in life-threatening mucormycosis, but has been used extremely infrequently as a topical agent for vaginal mycotic infections and is not available commercially for this indication. A 3% concentration was used somewhat arbitrarily, since this concentration is used to treat oropharyngeal candidiasis due to sensitive *Candida* species. Response in the reported case was not dramatic and required retreatment and extension of the treatment duration. The pathogenesis of this obscure infection remains an enigma and could not be correlated with any personal habits (douching) or other risk factors. This unusual and obstinate case emphasizes the value of obtaining vaginal fungal cultures in treatment of unresponsive cases, as well as the need for full identification of the isolated pathogen and basing therapy on susceptibility testing.

**REFERENCES**


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