U.S. Case Report of Cerebral Phaeohyphomycosis Caused by *Ramichloridium obovoideum* (*R. mackenziei*): Criteria for Identification, Therapy, and Review of Other Known Dematiaceous Neurotropic Taxa

DEANNA A. SUTTON,1,* MALCOLM SLIFKIN,2 ROBERT YAKULIS,2 and MICHAEL G. RINALDI1,3

Fungus Testing Laboratory, Department of Pathology, University of Texas Health Science Center at San Antonio3 and Audie L. Murphy Division, South Texas Veterans Health Care System,3 San Antonio, Texas 78284, and Department of Laboratory Medicine, Allegheny General Hospital, Pittsburgh, Pennsylvania 15212-47722

Received 21 August 1997/Returned for modification 24 October 1997/Accepted 25 November 1997

We report a case of cerebral phaeohyphomycosis in a 36-year-old male caused by the neurotropic fungus *Ramichloridium obovoideum* (Matsushima) de Hoog 1977 (*Ramichloridium mackenziei* Campbell et Al-Hedaithy 1993). This man resided in the Middle East, where the fungus appears to be endemic and, possibly, geographically restricted, since all previous reports of brain abscesses due to this organism have been for patients indigenous to this area. As a servant of the Saudi Arabian royal family, he appeared in the United States seeking treatment for chronic weight loss, fatigue, decreased memory, and a more recent 2-week history of right-hand weakness which worsened to involve the entire right upper extremity. On the day prior to his admission, he had a focal motor seizure with rotation of the head and eyes to the right, followed by secondary generalization. A computerized tomogram showed a ring-enhancing hypodense lesion in the left parietal subcortical region with associated edema and mass effect. Diagnosis of a fungal etiology was made following a parietal craniotomy and excisional biopsy by observation of septate, dematiaceous hyphal elements 2 to 3 μm in width on hematoxylin-and-eosin-stained sections from within areas of inflammation and necrosis. Culture of the excised material grew out a dematiaceous mould which was subsequently identified as *R. obovoideum*. At two months postsurgery and with a regimen of 200 mg of itraconazole twice a day, the patient was doing well and returned to Saudi Arabia. His condition subsequently deteriorated, however, and following a 7-month course of itraconazole, he expired. We use this case to alert clinicians and personnel in clinical mycology laboratories of the pathogenicity of this organism and its potential occurrence in patients with central nervous system signs and symptoms who have resided in the Middle East and to review and/or compare *R. obovoideum* with other neurotropic, dematiaceous taxa and similar nonneurotropic, dematiaceous species.

The numbers and types of saprobic, opportunistic, and dematiaceous moulds that have been documented as etiologic agents of phaeohyphomycosis continue to escalate. Clinical presentations may include superficial, cutaneous, subcutaneous, and systemic disease. Dematiaceous fungi that occur in compromised hosts may go on to disseminate hematogenously, inciting disease in various organs. Some that disseminate appear neurotropic, i.e., have a predilection for central nervous system (CNS) tissue, where they may localize, causing brain lesions and/or abscesses. These neurotrophic organisms are frequently thermotolerant or thermophilic, growing at 40°C or higher. Dematiaceous filamentous taxa known to be neurotropic include the hyphomycetous fungi Cladosphialophora bantiana (Saccardo) de Hoog, Kwon-Chung and McGinnis (4, 12, 20, 24, 25, 30, 44, 52), *Exophiala dermatitidis* (Kano) de Hoog (Wangiella dermatitidis McGinnis) (21, 23, 24, 27, 35, 46, 52), and *Ochroconis gallopavum* (Wangiella dermatitidis McGinnis) (21, 23, 24, 27, 35, 46, 52), as well as the dematiaceous ascomycete Chaetomium atrobrunneum Ames (1, 39). *Bipolaris spicifera* (Bainier) Subramanian (5, 24, 53) and *Bipolaris hawaiiensis* (M. B. Ellis) Uchida et Arakaki (24, 31, 41) may also invade the CNS via paranasal sinus extension. Additional dematiaceous agents incriminated include *Rhinocladiella atrovires* Nannfeldt (10, 24), *Cuvularia pallescens* Boedijn (16, 24, 26), and *Fonseccae pedrosii* (Brumpt) Negroni (2, 17, 18, 24, 52). Most of these species have a rather widespread distribution and are encountered in various ecological niches. Based on cases previously reported as either *Ramichloridium obovoideum* or *Ramichloridium mackenziei*, however, this etiologic agent appears to be geographically restricted to the Middle East (7, 22, 28, 32). Nonneurotropic species sharing macroscopic and microscopic morphologies similar to those of *R. obovoideum* include *Ramichloridium schulzeri* Saccardo de Hoog (8, 9, 13, 14, 40), *Rhinocladiella aquaspersa* (Borelli) Schell et al. (8, 42), and *Veronaea botryosa* Ciferri et Montemartini (3, 9, 35). A comparison of culture characteristics, microscopic morphologies, and in vitro susceptibilities may aid in differential diagnosis for patients with CNS signs and/or symptoms of a presumed, dematiaceous fungal etiology.

**MATERIALS AND METHODS**

**Case report.** The patient was a 36-year-old male servant of the Saudi Arabian royal family who presented at the Allegheny General Hospital in Pittsburgh, Pa., in May of 1995 with a 2-week history of right-hand weakness which worsened to involve the entire right upper extremity. On the day prior to admission, he had a focal motor seizure with rotation of the head and eyes to the right, followed by secondary generalization.
Hodgkin's disease diagnosed by biopsy of the left axillary lymph node in 1983. The patient was treated with alternating cycles of MOPP (nitrogen mustard, vincristine, procarbazine, and prednisone) and ABVD (doxorubicin, bleomycin, vinblastine, and dacarbazine) until complete remission was achieved. In 1986, the patient suffered a relapse which required six cycles of MOPP and ABVD. The patient relapsed again, was treated with CMVP-16 (cyclophosphamide, methotrexate, etoposide), and had been in complete remission since 1987. Past medical history was also significant for schistosomiasis as a child and chronic hepatitis B with portal hypertension, hepatomegaly, and splenomegaly. Since 1993, the patient had noticed a 10- to 12-kg weight loss, was chronically fatigued and apathetic, and had a decreased memory.

Upon admission on 6 May 1995, the patient was fully conscious, had a bitten tongue, normal fundi, and normal visual fields. There was increased weakness in the right upper extremity which was more pronounced distally than proximally, hyperactive deep tendon reflexes, and normal sensation. During his hospital stay, there was deterioration of normal lower-extremity strength over a period of 2 weeks. Except for hepatosplenomegaly, the remainder of the physical examination was within normal limits. The clinical impression was of upper neuron weakness of the right upper and lower extremities with right focal motor seizure with secondary generalization. A space-occupying lesion in the left midrolandic gyrus was suspected.

An electroencephalogram showed a normal pattern of waking and sleeping. A preoperative computerized tomogram with and without contrast showed a ring-enhancing hypodense lesion in the left parietal subcortical region with associated edema and mass effect, but no significant midline shift (Fig. 1). Within the areas of inflammation and tissue necrosis were occasional brown pigmented hypal forms approximately 2 to 3 μm in width, with the presence of septa in some areas. The fungal elements were also readily apparent in paraffin-embedded Congo red tissue sections (Fig. 3). Although the hypal form predominated, there were also moniliform elements (hypal elements with swellings at regular intervals) measuring up to 4 μm in diameter at their widest point. These were most easily visualized in the Gomori methenamine silver-stained sections (Fig. 4). Following histopathological confirmation of a dematiaceous mould as the etiologic agent, a regimen of 200 mg of itraconazole (ITRA) twice a day (BID) was begun.

The fungal culture was referred to the Fungus Testing Laboratory, Department of Pathology, University of Texas Health Science Center at San Antonio (accession no. 95-1147) for identification and susceptibility studies. In July 1995, 2 months postsurgery, the patient was doing well and the weakness, involving his right arm and leg had improved considerably. He returned to Saudi Arabia, continuing his regimen of 200 mg of ITRA BID. However, his condition subsequently deteriorated, requiring rehospitalization. He expired on 1 January 1996, after having received approximately 8 months of itraconazole. The patient did not garden and had no occupational exposure to the soil. Interestingly, he did walk barefoot in the desert. He had formerly been employed as a telephone operator before his most recent employment as a servant for the royal family.

Histopathology. Approximately 2 by 2 cm of white and brown tissue was submitted for frozen sectioning and hematoxylin and eosin staining showed a severe mixed acute and granulomatous infiltrate with large areas of tissue necrosis (Fig. 2). Within the areas of inflammation and tissue necrosis were occasional brown pigmented hypal forms approximately 2 to 3 μm in width, with the presence of septa in some areas. The fungal elements were also readily apparent in paraffin-embedded Congo red tissue sections (Fig. 3). Although the hypal form predominated, there were also moniliform elements (hypal elements with swellings at regular intervals) measuring up to 4 μm in diameter at their widest point. These were most easily visualized in the Gomori methenamine silver-stained sections (Fig. 4).

Mycology. Brain tissue from the excisional biopsy was submitted to the mycology laboratory for fungal cultures. The specimen was plated on Sabhi and brain heart infusion agars with 5% sheep blood (Becton Dickinson Microbiology
Systems, Cockeysville, Md.), with incubation at 30°C. These primary plates had been sealed with Shrink Seal (Scientific Device Laboratory, Inc., Glenview, Ill.) to maintain adequate humidity.

The isolate was referred to the Fungus Testing Laboratory, University of Texas Health Science Center at San Antonio, San Antonio, Tex., for identification and susceptibility testing. The isolate was plated onto potato flakes agar (PFA), which was prepared in-house (37), and Mycobiotic Agar (Remel, Lenexa, Kans.) at 25°C (ambient air, alternating daylight and darkness), at 35°C, and at 42°C (ambient air in the dark). Subcultures at 25 and 35°C were held for 4 weeks, while those at 42°C were maintained for 2 weeks. Slide cultures, maintained for 7 days at 25°C, were prepared on PFA blocks and examined in lactophenol cotton blue (Poly Scientific Research and Development Corp., Bay Shore, N.Y.) mounts.

Antifungal susceptibility testing. The case isolate was tested to determine its susceptibilities to antifungal agents. Tests were performed by previously described macrodilution methods (34, 38). Briefly, the case isolate and the Pae- cilomyces control strain (UTHSC 90-459) were grown on PFA, which was prepared in-house, for 14 days at 25°C to induce conidial formation. Mature PFA cultures were overlaid with sterile, distilled water, and suspensions were made by gently scraping the colonies with the tip of a Pasteur pipette. Heavy hyphal fragments were allowed to settle, and the upper, homogeneous conidial suspensions were removed. Conidial suspensions were adjusted spectrophotometrically to 95% transmittance at 530 nm and were further diluted 1:10 in medium. Final drug concentration ranges were as follows: for amphotericin B (AMB; E.R. Squibb & Sons, Princeton, N.J.), 0.03 to 16 µg/ml; for 5-fluorocytosine (5-FC; Hoffmann-La Roche Inc., Nutley, N.J.), 0.125 to 64 µg/ml; for fluconazole (FLU; Pfizer, Inc., New York, N.Y.), 0.125 to 64 µg/ml; and for ITRA (Janssen Pharmaceutica, Titusville, N.J.), 0.015 to 8 µg/ml. AMB was tested in Antibiotic Medium 3 (Difco, Detroit, Mich.); other antifungal agents were tested in RPMI-1640 with L-glutamine and morpholinepropanesulfonic acid (MOPS) buffer at a concentration of 165 mM and without sodium bicarbonate (American Bioanalytics, Inc., Niagara Falls, N.Y.). Previously prepared, frozen drug tubes containing 0.1 ml of drug were allowed to thaw and were inoculated with 0.9 ml of the conidial medium suspension. A drug-free growth control tube was included with the case isolate and control organism. The tubes were incubated at 35°C, and MICs were read at the first 24-h interval when growth was observed in the drug-free growth control. MICs were defined in terms of the first tube that gave a score of 0 (optically clear) for AMB and a score of 2 (reduction in turbidity of ≥80% in contrast to the drug-free control tube) for 5-FC, FLU, and ITRA.

RESULTS

Mycology. At the Allegheny General Hospital, the mould was first detected on Sabhi and brain heart infusion agars as small, gray, lanose colonies at 4 days of incubation at 30°C. After 7 days of incubation, colonies were 8 to 10 mm, lanose, and gray black. The reverse color on Sabhi agar was gray black, with the tops of the colonies on both media being black, woolly, and domed. The microscopic morphology, by tape and teased mounts in lactophenol cotton blue (Becton Dickinson Microbiology Systems), was described as similar to that seen in *F. pedrosoi*. Often two smooth-walled and lightly pigmented conidia were seen projecting from both sides of the relatively thick-walled rachis, giving a “Mickey Mouse” appearance (Fig. 5A). Cultures in the Fungus Testing Laboratory on PFA at

![FIG. 2. Brain lesions showing mixed acute and granulomatous inflammation with hematoxylin and eosin staining. Magnification, ×300.](http://jcm.asm.org/)

![FIG. 3. Paraffin-embedded Congo red tissue sections containing hyphal elements of *R. obovoidum*. Magnification, ×630.](http://jcm.asm.org/)

![FIG. 4. Gomori methenamine silver stain showing moniliform hyphal elements of *R. obovoidum*. Note the moniliform (bead-like) hyphae characteristic of agents of phaeohyphomycosis. Magnification, ×1,200.](http://jcm.asm.org/)
25°C revealed colonies that were dark brown to black with a jet-black reverse, woolly, and slow growing, reaching only 2 to 4 mm in 10 days. Six weeks of incubation at 25°C produced colonies with a high-domed central area and a submerged margin throughout. Growth was better at 35 than 25°C with diameters reaching 12 to 20 mm in 17 days. Growth also occurred at 42°C and on media containing cycloheximide, although measurements were not taken. Hyphae were septate and brown, measuring 1.3 to 2.0 μm in diameter. Conidiophores arose at more or less right angles to the vegetative hyphae and were not strongly differentiated from it (micronematous). The conidiogenous areas elongated sympodially, becoming flexous (Fig. 5B). The conidal rachis, 3 to 12 μm long by 3 to 5 μm in diameter, was somewhat larger than the lower part of the conidiogenous cells. Relatively few conidia per fertile axis were present. They were pale brown, ellipsoid to ovoid, single celled, smooth, 5.85 to 8.7 μm long by 2.7 to 4.8 μm wide, with a protuberant hilum or a flat secession scar up to 1.3 μm wide (Fig. 5C).

**DISCUSSION**

**Case report.** Although the patient’s past medical history was significant for stage IIIb mixed-cellularity Hodgkin’s disease (in complete remission since 1987), chronic hepatitis B with portal hypertension, hepatomegaly, and splenomegaly, and shistosomiasis as a child, it is difficult to establish whether these conditions were contributory to acquisition of the fungus. The patient denied any occupational exposure to the soil, but he did walk barefoot in the desert. Cerebral phaeohyphomycosis due to other neurotropic agents, particularly *C. bantiana*, is also frequently seen in patients without obvious (detectable), immunosuppressive, coincidental patterns of preexisting illness and without an unambiguous occupational predisposition, al-
though culturally proven cases do have a male-to-female ratio of about 3:1 (12, 24). One exception appears to be the relationship between preexisting *Nocardia asteroides* and phaeohyphomycotic brain abscesses due to *C. bantiana* (24, 30, 32, 43). In the 8 cases of *R. obovoideum* referenced here, in which the sex of the patient was known, the male-to-female ratio was 1:1.

**Therapy.** Cerebral phaeohyphomycosis is one of the most serious clinical manifestations caused by dematiaceous fungi and has a high degree of morbidity and mortality, requiring early and aggressive therapy. The patient in this case report was admitted on 6 May 1995, was diagnosed on 18 May 1995, and remained on this regimen until 26 December 1995, 6 days prior to expiring on 1 January 1996. Although the patient was diagnosed rather quickly after admission, he had experienced a weight loss of 10 to 12 kg, was chronically fatigued and apathetic, and had had a decrease in memory since 1993. Seventeen cases of intracranial mycoses requiring neurosurgical intervention were reviewed by Jamjoom et al. (22). Citing a 41% mortality, they indicated that reasons for a fatal outcome included (i) failure to consider a fungal etiology, (ii) failure to obtain an early tissue diagnosis because of late referral, and (iii) failure to respond to antifungal therapy. Naim-ur-Rahman et al. reported three cases with a 100% mortality due to rupture of recurring abscesses into the ventricles despite adequate antifungal chemotherapy and multiple surgical procedures (33). The fatal outcome in this patient may have been attributed, in part, to his seeking medical attention late in the course of the disease to possible failure of ITRA therapy. However, he did appear to receive an adequate dosing regimen for an isolate demonstrating in vitro susceptibility to this agent (MICs of less than 0.015 µg/ml) (45). Although levels in serum documenting adequate absorption of the drug were not measured, the patient did initially respond to the drug, permitting his return home to Saudi Arabia. ITRA, like other azole antifungals, is fungistatic rather than fungicidal and may have merely suppressed the organism. Factors in addition to in vitro susceptibility which may have influenced the outcome, as outlined by Rex et al., include the pharmacokinetics of the drug, general host factors, site of infection, and virulence of the pathogen (36). In vivo-in vitro drug studies with other neurotropic agents, *C. bantiana*, *E. dermatitidis*, and *O. gallopava* (*Dactylaria constricta* var. *gallopava*), prior to the introduction of ITRA, suggested that the greatest protection in experimentally infected mice was offered by 5-FC followed by AMB, FLU, and ketoconazole and that in vitro susceptibility testing had no predictive value (11). In vivo animal studies with *R. obovoideum* versus ITRA were not available at the time of the patient’s therapy.

A review by Dixon et al. of 26 cases of CNS fungal infections due to *C. bantiana* suggested that the neurosurgical resection of the lesion, with or without antifungal therapy, was the most important determinant for cure and survival (12). A resectable lesion was defined as a discrete mass demarcated by a peripheral gliotic capsule surrounding necrotic infected debris, in contrast to abscesses having poorly delimited margins and satellite abscesses. Complete resection of the lesion in this case was not thought possible.

**Antifungal susceptibility.** While standardization in antifungal susceptibility testing has been attained for the major, clinically significant yeasts with publication of the National Committee for Clinical Laboratory Standards document no. M27-A (34), standardization of susceptibility testing for filamentous fungi is only commencing (15). Parameters previously defined for yeast testing, with some modification, also appear to be useful for mould testing. The case isolate in the present study was
<table>
<thead>
<tr>
<th>Species</th>
<th>Macroscopic morphology</th>
<th>Microscopic morphology</th>
<th>Ecology</th>
<th>Physiology</th>
<th>% Susceptible in vitro (n)</th>
<th>Comment(s)</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>R. obovoideum</td>
<td>Black, woolly, domed colony; slow growth</td>
<td>Sympodial; few conidia per fertile axis; brown conidia; 4.7–9.6 by 2.7–6.0</td>
<td>Middle East; soil? plants?</td>
<td>40°C, 1; cyclo., V 100 (1) (AMB)</td>
<td>Domed colony occurs at maturity 7, 22, 28, 32, 50</td>
<td>100 (1) (5-FC) 100 (1) (FLU) 100 (1) (ITRA)</td>
<td></td>
</tr>
<tr>
<td>C. (Xylohypha) bantiana</td>
<td>Olivaceous to black; woolly; moderate growth</td>
<td>Long, poorly branched, non-fragile chains of conidia; hila absent</td>
<td>Wide distribution in soil and plants</td>
<td>40°C, 1; nitrate, 1, 83 (6) (AMB)</td>
<td>Other names include Cladosporium trichoides</td>
<td>12, 20, 24, 25, 30, 44, 50, 52</td>
<td>100 (4) (5-FC) 100 (1) (FLU) 100 (6) (ITRA) 100 (1) (MON)</td>
</tr>
<tr>
<td>E. (Wangiella) dermatitidis</td>
<td>Olivaceous to black; mucoid; moderate growth</td>
<td>Black yeast synanamorph present; ellipsoidal conidia; 2.5–4 by 2–3</td>
<td>Wide distribution in soil, plants, and water</td>
<td>40°C, 1; nitrate, 2</td>
<td>Both annellides and phialides may be present; colony maybe more filamentous on Sabouraud dextrose agar</td>
<td>21, 23, 24, 27, 35, 46, 50, 52</td>
<td></td>
</tr>
<tr>
<td>O. gallopavum, (Dactylaria gallopava)</td>
<td>Reddish-brown, velvety, maroon diffusing pigment; slow growth</td>
<td>Two-celled, clavate conidia; 11–18 by 2.5–4.5; on denticles</td>
<td>Wide distribution in thermal areas</td>
<td>42°C, 1; cyclo., 2; urease, 1</td>
<td>100 (2) (AMB) 100 (1) (5-FC) 100 (2) (ITRA)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>C. atrobrunneum</td>
<td>Gray to black with production of perithecia; moderate growth</td>
<td>Perithecia diam, 70–150; sub-globose, dark brown; few straight setae; brown asco-spores; fusoidal; 9.3–10.8 by 4.9–6.2</td>
<td>Wide distribution in soil, plants, and air</td>
<td>40°C, 1</td>
<td>None available Fatal in a leukemic patient 1, 39</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**TABLE 2. Dematiaceous neurotropic taxa**

<table>
<thead>
<tr>
<th>References</th>
<th>Comment(s)</th>
<th>Species</th>
</tr>
</thead>
</table>

*This list is not all inclusive.*

*Additional dematiaceous agents reported include R. atrovirens, C. pallescens, and F. pedrosoi. In some instances, C. bantiana may have been reported as F. pedrosoi.*

*PFA was prepared in-house for 10 days at 25°C.*

*Dimensions are given in micrometers.*

*1, growth; V, variable growth; 2, no growth; cyclo, cycloheximide.*

*en, number of isolates tested by the National Committee for Clinical Laboratory Standards Reference method for broth dilution antifungal susceptibility testing of yeasts (proposed standard M27-A, modified to accommodate mould testing). Mould susceptibility testing is currently nonstandardized, and breakpoints have not been established. Percent susceptibility and percent resistance, respectively, are based upon the following MICs (in micrograms per milliliter): AMB, #1 and $2; 5-FC, #16 and $32; FLU, #32 and $64; ITRA, #0.5 and $1.0; and miconazole (MON), #8 and $16.*
tested by a modified M27-A method. Although standardized antifungal susceptibility testing now permits large surveys of yeast isolates for correlating MIC data with clinical response, such correlations for infrequently encountered moulds are not possible, still making empiric therapy necessary. Clinical outcomes with *R. obovoideum* have been dismal, regardless of antifungal therapy, and appear to lack correlation with in vitro data (7).

**Taxonomy and identifying features.** The genus *Ramiichloridium*, causing leaf-spots on banana, was first described by Stahel (49). However, the description was invalidly published, lacking a Latin description. In 1977, de Hoog reviewed a number of dematiaceous hyphomycetes that produce holoblastic conidia from a sympodial proliferating axis, including *Ramichloridium* and *Rhinoocladiella* (8). These genera were separated mainly on the basis of (i) macronematous conidiophores in *Ramichloridium* versus micronematous conidiophores in *Rhinoocladiella* and (ii) human pathogenicity, lacking in *Ramichloridium* but present in *Rhinoocladiella* species. A third, less stable characteristic mentioned was the presence of a yellow or an orange diffusable pigment, often present in *Ramichloridium* but lacking in *Rhinoocladiella*. *R. obovoideum* (Matsushima) de Hoog comb. nov. was included in Stahel’s review and was originally described as *R. obovoideum*. In the late 1980s, case reports incriminating *R. obovoideum* as an agent of cerebral phaeohyphomycosis emerged. In 1993 Campbell and Al-Hedaithy published a review of eight cases due to an organism they named *R. mackenziei* (Table 1) (7). Characteristics which deviated from the previously-described genus *Ramichloridium* included (i) a lack of yellow or orange diffusible pigment, (ii) conidiophores not obviously differentiated from the vegetative hyphae, and (iii) the unambiguous human pathogenicity of the organism. Despite these deviations, they felt the most appropriate genus to accommodate these fungi was *Ramichloridium*. Because Matsushima’s isolate was not available for comparison with the case isolates and conidia from the brain abscess isolates were smaller than those for *R. obovoideum* (4.7 to 9.0 by 2.7 to 4.0 μm versus 8.8 to 12 by 3.8 to 5 μm), the new name *R. mackenziei* was proposed in honor of D. W. R. Mackenzie. Our case isolate also had conidial dimensions smaller than those originally described. Until further isolates are reported for comparison and current and future isolates are evaluated at the molecular level, it is difficult to ascertain whether the size of the conidia is a stable characteristic, a view held by Campbell and Al-Hedaithy (7), or whether this variability in conidial dimensions is within the limits for the species (8, 29).

Other dematiaceous hyphomycetes with potential or genuine cerebral pathogenicity include *C. bantiana* (also named Xylohypha bantiana), *Cladosporium bantianum*, *Cladosporium trichoides*, *Cladosporium trichoides* var. *chlamydosporum*, and *Xylohypha emmonsii*, *E. (Wangella) dermatitidis*, and *O. gallopavum* (*Dactylaria gallopava* and *Dactylaria constricta* var. *gallopava*). The darkly pigmented ascomycete *C. atrobrunneum* is an additional taxon causing cerebral phaeohyphomycosis. Characteristics of these five dematiaceous neurotropic taxa, including their in vitro susceptibility data, are outlined in Table 2. A closely related species, *Chaetomium strumarium*, although not dematiaceous, is also an agent of brain abscesses (1). Additional organisms which have been reported to invade the CNS include the graminicolous (living on grass) species *B. spicifera* (5, 24, 53), *B. hawaiensis* (31, 41), and *C. pallescens* (16, 26). *R. atrovirens* was also incriminated in CNS involvement in a human immunodeficiency virus-positive male drug abuser (10).

Nonneurotropic dematiaceous fungi sharing macroscopic and microscopic morphologies similar to those of *R. obovoideum* include *R. Schulzeri*, *R. aquaspersa*, and *V. botryosa*. *R. Schulzeri*, which was reported in a case of “golden tongue” in a leukemic patient, is also sometimes seen from respiratory isolates. It more closely matches the original description of the genus *Ramichloridium* in that the conidiophores are markedly differentiated from the vegetative hyphae, there are numerous conidia per fertile axis (Fig. 6), and a distinct yellow diffusible pigment is usually present. *R. aquaspersa*, an agent of human chromoblastomycosis, bears closely packed hyaline-to-subhyaline and ellipsoid conidia on crowded denticles. *V. botryosa*, an agent of subcutaneous phaeohyphomycosis in China, produces predominately two-celled, crowded conidia on a sympodially proliferating fertile axis.

The numbers and types of opportunistic dematiaceous fungi continue to escalate. While most have occurred in immunocompromised hosts, patients without obvious detectable immunosuppression may also be at risk for disseminated disease. *R. obovoideum* (*R. mackenziei*) should be considered as a possible etiologic agent in cerebral phaeohyphomycosis, particularly in patients indigenous to Middle Eastern countries. It is also prudent for laboratory workers studying this organism to bear in mind its likely neurotropic nature. Because of its potential respiratory acquisition and until the route of infection is continued, all studies with this mould, as well as with all other filamentous fungi, should be accomplished with a biological safety hood.

**REFERENCES**


