We report 2 new cases of invasive infections caused by *Nannizziopsis* spp. molds in France. Both patients had cerebral abscesses and were immunocompromised. Both patients had recently spent time in Africa.

*Nannizziopsis* spp. molds have been reported in extremely rare cerebral and disseminated infections (1,2), (Table). We describe 2 cases of *Nannizziopsis* infection diagnosed in France during the past 2 years. Both case-patients were immunocompromised and had recently returned from Africa.

The Cases

Case-patient 1 was a 63-year-old woman from France who had been treated for T-cell prolymphocytic leukemia diagnosed in December 2014. She initially received alemtuzumab, switching to bendamustine in March 2015 after tumor progression. That treatment failed, and idelalisib was started on July 11, 2015. The patient was hospitalized 2 days later for acute renal failure with mild fever. She became confused and drowsy, and cerebral computed tomography (CT) scan showed systematized subcortical hypodense areas. Lumbar puncture showed neoplastic cells in the cerebrospinal fluid (CSF) and glucose concentration within reference range. Bacteriological and fungal cultures were sterile. A large volume of ascites fluid remained, despite iterative punctures with negative bacteriological cultures. We initiated intrathecal chemotherapy with methotrexate/cytarabine/methylprednisolone. However, the patient’s condition worsened, with heart failure and loss of consciousness. On July 18, we took new specimens of CSF, bronchial wash, ascites fluid, and blood cultures and sent them for bacteriological investigation. We started empiric treatment with imipenem/aminoglycosides, but the patient died on July 19 of septic shock. No autopsy was performed. Extended-spectrum β-lactamase–producing *Escherichia coli* sensitive to imipenem grew quickly in 1 pair of blood cultures. A second pair was positive 4 days later, with the presence of large septate fungal hyphae and arthroconidia. White and thin cottony mold colonies grew on Sabouraud media incubated at 35°C (online Technical Appendix Figure 1, https://wwwnc.cdc.gov/EID/article/24/3/17-0772-Techapp1.pdf). We performed best model determination and phylogenetic analyses in MEGA6 (http://www.megasoftware.net). We identified *N. obscura* by sequencing the 18S-internal transcribed spacer (ITS) 1–5.8S-ITS2 region (online Technical Appendix Figure 2). The strain had low MICs for antifungals as defined by the European Committee on Antimicrobial Susceptibility Testing (http://www.eucast.org/): amphotericin B 0.06 µg/mL, itraconazole 0.25 µg/mL, voriconazole 0.03 µg/mL, posaconazole 0.06 µg/mL, caspofungin 0.5 µg/mL, and micafungin 0.015 µg/mL. We performed mycologic investigations of CSF and ascites fluid a posteriori on frozen aliquots and conducted PCR assays targeting the ITS region on CSF sampled on July 15 and July 19 and on ascites fluid sampled on July 19. We observed positive amplifications in all samples; subsequent sequencing confirmed the presence of DNA from *N. obscura*. We investigated the origin of the patient’s contamination. She had made several trips to Senegal, the latest in January 2015, during which an ulcerative inflammatory lesion developed on her left little finger. However, Grocott stain and PCR on paraffin-embedded tissue of skin biopsy were negative, and we attributed the lesion to the hematological malignancy.

Case-patient 2 was a 52-year-old woman from France living in Mali, who was hospitalized in Bamako in November 2016 for cough, fever, alteration of general state, and headache. She tested seropositive for HIV (CD4 3/µL; HIV-1 viral load 45.300 copies/mL). Chest radiograph showed bilateral pneumonia, and cerebral CT scan showed a single process on the left temporal lobe. Antiretroviral therapy was initiated with a combination of efavirenz/lamivudine/tenofovir associated with isoniazid, metronidazole, amoxicillin/clavulanate, and trimethoprime/sulfamethoxazole. Because of worsening of her neurologic status, she was repatriated to France. At hospital admission on January 12, 2017, she had a lesion on the left middle fingernail suggestive of onychomycosis, hemiparesis, and paralysis of the right side of the face.
PCR results of the lung lesions and cerebral abscesses; her CD4 cell count was 50/µL and HIV-1 load <40 copies/mL. We switched her antifungal treatment to voriconazole.

**Conclusions**

The molds of the *N. vriesii* complex (*Chrysosporium*-like anamorph, CANV) are members of the *Nannizziopsis* genus (*Oxycyphus*, *Eurotiomycetidae*, *Eurotiomycetes*, *Ascomycota*).
Whole-Genome Analysis of *M. tuberculosis*, Russia

CANV includes the keratinophilic species, which causes skin and fatal disseminated infections in reptiles (3,4). There is no documented evidence of zoophilic species involvement in human infections, but 3 other CANV species have been recovered from human samples (Table). Of the 5 previous cases of *Nannizziopsis* infections, 3 involved HIV patients. One of our patients was seropositive for HIV and the other had T-cell prolymphocytic leukemia, which suggests that lymphopenia could be a key risk factor. All patients with reported *N. obscura* infection had recently traveled in Africa. Results for case-patient 1 showed that the fungus grows in blood cultures and thus has high potential for dissemination. Case-patient 2 had pulmonary lesions, but BAL cultures were rapidly invaded by a *Penicillium* fungus. Although we could not detect *Nannizziopsis* in BAL, the lesion evolved favorably after antifungal therapy. Because *Nannizziopsis* spp. are keratinophilic molds, we looked for cutaneous lesions. Both case-patients had recently developed cutaneous or nail lesions during their time in Africa, but we did not detect *Nannizziopsis* from these lesions. These molds have not been isolated in our laboratory in other kinds of samples (clinical or environmental).

CSF cultures from both our case-patients and ascites cultures from case-patient 1 were negative, but panfungal PCR successfully detected molds (6). Data on biomarkers are scarce. *N. infrequens* cross-reacts with the *Histoplasma* AccuProbe test and *N. hominis* with the *Blastomyces* AccuProbe test (Hologic, San Diego, CA, USA) (5). In case-patient 2, β-D-glucan was positive in CSF and serum but galactomannan antigen was not. *N. obscura* seems to be sensitive to most antifungal agents (2).

These observations show how difficult this infection is to detect, which could explain why so few cases of human infections have been reported. However, the diagnosis of these 2 cases since 2015 suggests that the prevalence of *Nannizziopsis* infections may be underestimated.

Acknowledgments
The authors thank all the members of the Centre National de Référence des Mycoses et Antifongiques (CNRMA, Institut Pasteur de Paris, France) for their invaluable help in the antifungal susceptibility testing.

Figure. Diagnostic testing of a 52-year-old woman from France living in Mali who had *Nannizziopsis* spp. fungal infection. A) Thoracic-abdominal-pelvic scan shows pseudo-nodular lesions in the apex of the right lung, of which one is excavated. B) Cerebral computed tomography scan shows contrast enhancement on several hemispheric nodules on the left and in frontal, parietal, and temporal regions, responsible for large surrounding edema and compression of the left lateral ventricle. The median line is deviated to the right with a subfalcical herniation. C) Hematoxylin-eosin-saffron stain of brain biopsy containing mononuclear inflammatory infiltrates; giant cell granulomas; histiocytes, sometimes with an epithelioid appearance; and neutrophils (original magnification ×200). D) Grocott stain showing thick bulbous mycelial filaments in the cytoplasm of certain giant cells/histiocytes (original magnification ×600). Round shapes correspond to cross-sections of bulbous territories.
About the Author
Dr. Nourrisson is a medical mycologist and parasitologist at the university hospital of Clermont-Ferrand, France. Her research interests include medical fungi epidemiology and digestive protozoa.

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Address for correspondence: Philippe Poirier, Laboratoire de Parasitologie-Mycologie, Centre de Biologie, CHU Gabriel Montpied, 58 rue Montalembert, 63000 Clermont-Ferrand, France; email: ppoirier@chu-clermontferrand.fr

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