A Well-Preserved Culprit
(See page 704 for the Photo Quiz.)

Diagnosis: reactivated, disseminated histoplasmosis.

Histological examination of laryngeal biopsy specimens showed the presence of budding yeast cells (Figures 1 and 2). Based on this observation, laryngeal histoplasmosis with underlying iatrogenic hypercortisolism, steroid-induced hypertension, and diabetes mellitus was suspected. Histopathological reexamination of the adrenal glands, which had been removed in 1996, also raised suspicion of adrenal histoplasmosis instead of the previously suspected culture-negative tuberculosis. The presence of *Histoplasma capsulatum* in both the laryngeal and adrenal gland tissues was demonstrated by broad-spectrum polymerase chain reaction (PCR) targeting the fungal internal transcribed spacer (ITS) region, followed by sequence homology analysis of the PCR product [1]. *H. capsulatum* genotyping using 4 targets of sufficient intra-species variability (arf, h-anti, ole, and tub1) [2, 3] provided proof that reactivation, not reinfection, took place. Interestingly, tests for specific antibodies remained negative twice, but *H. capsulatum* antigen detection from a urine sample obtained after 2 weeks of antifungal therapy was positive. After 6 weeks of incubation, *H. capsulatum* was also cultivated from the patient’s urine (Figure 3).

Treatment with itraconazole was initiated according to the current Infectious Diseases Society of America guidelines [4]. The hoarseness disappeared almost completely, but hospitalization became necessary due to progressive asthenia. Cranial computed tomography (CT) scan was not indicative of cerebral histoplasmosis, but CT scans of the chest and abdomen suggested bilateral pulmonary involvement. The patient, who always had sufficient capacity of discernment, chose to discontinue any treatment, including hydrocortisone substitution, and died from hypocortisolism 2 months after the diagnosis of disseminated histoplasmosis was made. The development of diabetes mellitus and immunological changes as a result of advancing age may have contributed to the reemergence of the pathogen 15 years after the initial manifestation.

Only <1% of individuals infected with *H. capsulatum* develop symptomatic disease. The severity of symptoms depends on the degree of exposure and the immune status of...
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H. capsulatum is highly prevalent in the central and southeastern parts of the United States as well as in Central and South America and parts of Africa. However, our case shows that, due to increasing international travel activity, infection needs to be ruled out in a wide variety of clinical scenarios, not only within endemic regions. In addition to traditional diagnostic techniques (histopathology, serology, and culture), broad-spectrum PCR targeting the fungal ITS region is a valuable tool for the detection of this potentially fatal pathogen, not least in settings where reliable histopathological identification may be difficult to obtain.

the host, and can range anywhere from mild pulmonary affection to severe disseminated disease. Disseminated disease mostly occurs in patients with impaired cell-mediated immunity [5]. In this context, it is remarkable that invasive fungal infections are not consistently recognized in patients treated with immunosuppressants like corticosteroids and tumor necrosis factor–α blockers [6, 7].

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Notes

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Figure 3. White to buff-brown mould form of H. capsulatum grown at 25°C on Sabouraud’s dextrose agar. Large numbers of infectious conidia may be released by the mere lifting of the culture plate lid.