CASE IN POINT

Disseminated Invasive Candidiasis in an Immunocompetent Host

Dhammika H. Navarathna, DVM, PhD; Eric R. Rachut, MD; Chetan Jinadatha, MD; and Gagan Prakash, MD

Health care providers should consider a nonbacterial source as the causative agent for invasive candidiasis infection in immunocompetent patients.

Candida albicans (C. albicans) is a normal commensal in the human gastrointestinal (GI) tract. In addition to localized infections in healthy human beings, dissemination with fatal outcome can occur in immunocompromised individuals.

Invasive candidiasis (IC) due to C. albicans is the most common nosocomial mycosis in the world and has 2 forms, candidemia and deep-seated tissue candidiasis, which can lead to multisystem organ failure. The deep-seated form may originate from nonhematogenous routes, such as introduction through a peritoneal catheter or ascending infection from cystitis. In addition, about 50% of primary candidemia cases lead to secondary deep-seated candidiasis; however, only about 40% of these cases show positive blood cultures. Since the window of opportunity for a positive culture is narrow, active candidemia may be missed.

Once developed, the prognosis for IC is grim: Mortality is 40% regardless of therapy. IC typically occurs in immunocompromised hosts; IC in immunocompetent persons has rarely been reported. It is challenging to diagnose IC in the immunocompetent patients as 50% to 70% of the general population is naturally colonized by this organism, and when found, it is assumed to be mostly innocuous. Neutrophil-driven cell-mediated immunity associated with IL-1 and IL-17 response prevent fungal growth and dissemination, protecting the immunocompetent host.

We report on a patient who showed no neutropenia or leukocytopenia but developed disseminated candidiasis. This report is one of the rare cases of full-blown disseminated candidiasis with lesions related to C. albicans found in almost all of the important organs.

**CASE PRESENTATION**

A 67-year-old male patient with a history of hypertension, peripheral vascular disease, daily heavy alcohol consumption, and a 50-pack-year history of smoking developed gangrene of the left fifth toe. He underwent vascular surgery consultation with an aortogram/left lower extremity angiography that showed occlusion of the left external iliac artery as well as the left common femoral artery. It was decided to improve inflow in the common iliac artery by placing a bare metal stent and subsequent balloon dilatation before a right to left femoral to femoral artery bypass. The patient tolerated the procedure well and was discharged home.

Two days later, the patient was admitted to a US Department of Veterans Affairs (VA) complexity level 1a hospital with weakness and worsening pain in the left lower extremities. Examination revealed chronic ischemic changes in the feet bilaterally and evidence of dry gangrene in the left fifth toe requiring femoral bypass surgery. But poor nutritional status and cardiac status prevented pursuing a permanent solution.

Following completion of a stress echocardiogram, the patient developed shock with systolic blood pressure of 60 mm Hg, and atrial fibrillation (AF) with rapid ventricular rate (RVR). He was initially treated with IV fluid supplementation, vasopressor therapy, synchronized cardioversion, and IV amiodarone/anticoagulation therapy, due to his persistent AF with RVR. The patient was transferred to a tertiary care center for persistent hypothermia and received treatment with warm saline. After initial recovery with warm saline resuscitation, he had a prolonged, complicated hospital course in...
which he developed progressive respiratory failure requiring intubation and critical care support. He developed a right internal jugular deep venous thrombosis, heparin-induced thrombocytopenia, lower GI bleeding requiring emergent embolization by interventional radiology, inferior vena cava filter placement, renal failure requiring dialysis, small bowel obstruction secondary to right lower quadrant phlegmon and perforation requiring small bowel resection and end ileostomy. His antibiotic regimen included therapy with vancomycin and piperacillin-tazobactam.

He eventually recovered and was extubated and subsequently transferred back to the VA hospital where cefepime was initiated.

**FIGURE 1** *C. albicans* Colorization in Organs Due to Invasive Infection

**A**

Representative section of upper lobe of left lung H&E stain. (A) Acute bronchopneumonia with *C. albicans* yeast and filaments surrounded with inflammatory reaction. (B) A corresponding GMS stain shows yeast and hyphae radiation in right lung tissues. (C) Huge inflammation in kidney matrix shows in a representative H&E stain section of right kidney. (D) GMS stain shows yeast predominant invasively multiplied *C. albicans* inside a glomerulus. (E & F) Representative H&E and GMS section showing *C. albicans* colonization in endocardium.

Abbreviations: *C. albicans*, *Candida albicans*; H&E, hematoxylin and eosin; GMS, Grocott-Gömöri methenamine silver.
because of suspicion of a urinary tract infection and septicemia (urine cultures eventually grew \textit{C} albicans). Over the subsequent 3 days, the patient's renal output and hyperkalemia worsened, he also developed increased anion gap metabolic acidosis and was intubated again and placed on full mechanical ventilatory support. His blood cultures were negative, and sputum cultures revealed normal respiratory flora and 1+ \textit{C} albicans. Infectious diseases consultation recommended an abdominal ultrasound, which revealed nonspecific findings. The antibiotic regimen was changed to daptomycin and piperacillin-tazobactam. A follow-up chest X-ray revealed a developing right lower lobe pneumonia and hilar prominence suggestive of lymphadenopathy. The patient's clinical condition deteriorated, and he subsequently developed cardiac arrest; resuscitation was not successful and he expired.

**Outcome and Follow-up**

An autopsy disclosed the cause of death to be bilateral candida pneumonia, part of a disseminated (invasive) candidiasis, in a patient rendered vulnerable to such infection by peripheral vascular disease and renal insufficiency. Purulent inflammation was noted at the site of disarticulation of the left foot and confluent consolidation of the lower lobes of both lungs as well as focal consolidation of the middle lobe of the right lung. Examination of histologic sections, with staining both by routine method (hematoxylin and eosin) and the Grocott-Gömöri methenamine silver method for fungus, disclosed fungal forms (yeast and filamentous) in most tissues, including the lungs (Figure 1 A and B) and kidneys (Figure 1 C and D). The pulmonary sections in addition to massive inflammation showed macrophages with engulfed yeast (Figure 2 A) and a lymphatic channel, stuffed with yeast in an alveolar septum (Figure 2 B). These findings confirmed the antemortem presence of the fungus and the body's response to it. Inflammation was noted around glomeruli overgrown by candida (Figure 1 C and D); fungi also were seen in capsular regions (not depicted). \textit{C} albicans was present in the myocardium (Figure 1 E and F), brain, thyroid, and adrenal glands (Figure 3); the only organ without \textit{C} albicans was the liver, either because invasion was truly absent here or because sampling had not managed to retrieve it.

Paraffin-embedded blocks of lung tissue, sent to the University of Washington Molecular Diagnosis Microbiology Laboratory for broad-range polymerase chain reaction (PCR) identification, were positive for \textit{C} albicans after extraction of gDNA and conduction of PCR using internal transcribed spacer 1 and 2 specific primers.

**DISCUSSION**

IC is rare among immunocompetent individuals, but \textit{C} albicans can evolve into a fatal disseminated infection. We report an atypical case of IC, with profound pulmonary infection in a patient who died 1 month after hospitalization for lower extremity pain.

Cell-mediated immunity involving neutrophils and macrophages plays a major role in protection against candidiasis, while

---

**FIGURE 2** \textit{Candida albicans} Colonization in the Lung

(A) Histopathology of a dust cell with entrapped \textit{Candida albicans} yeast (arrow); and (B) A lymphatic channel on alveolar septum with multiple yeast cells confirming dissemination of the organism systemically (arrows).
Invasive Candidiasis

cytokines and chemokines involve regulating balanced immunity. A series of recent studies show that alcohol impairs neutrophil-mediated killing and phagocytic-mediated uptake of a pathogen in this process. As the patient chronically misused alcohol, his immune system may have experienced a subclinical immunosuppression, which would have become clinically relevant once \textit{C. albicans} was introduced systemically. Recent studies of bacterial pathogenesis and alcoholism strongly support this hypothesis.

Most patients with the unusual diagnosis of candida pneumonia have had a background of malignancy or immunosuppressive factors (eg, administration of corticosteroids). In a series of 20 cases, 14 had sputum cultures positive for the organism, 6 had positive urine cultures, and 6 had positive blood cultures. Chest radiographs usually showed confluent bronchopneumonia. Five patients were diagnosed antemortem and treated with amphotericin B, but none survived. In the literature a positive blood culture or demonstration of yeast within pulmonary histiocytes has been considered proof of the pathogenicity of the fungus, as opposed to noninvasive colonization of the airways, a common occurrence in patients receiving mechanical ventilation.

As previously discussed, blood cultures are often negative with invasive candidiasis, as the window of opportunity is short and may be missed. As shown in murine models, it is easy to miss a narrow window of candidemia, leading to false-negative blood cultures in clinical practice. Mouse model studies also have found that the window of candidemia is very short in disseminated candidiasis as a lethal IV dose of \textit{C. albicans} disappeared from blood within 48 hours of postinoculation. The biomarker of serum procalcitonin is a great diagnostic resource for the elimination of a likely bacterial sepsis, and conversely, the early suspicion of a fungemia, as serum procalcitonin would typically be elevated in a bacterial but not a fungal sepsicemia. The average cost per test is only about $30, and we recommend testing for serum procalcitonin as well as monitoring of serum lactate levels in cases of nonresponding septicemia.

The \textit{C. albicans} in this case may have been introduced hematogenously from the amputation site or through an ascending cystitis, or possibly have been derived from commensal flora in the GI tract. The iron

\textbf{FIGURE 3 Candida albicans Colonization in Brain, Thyroid, and Adrenal Gland}

(A) Representative Grocott-Gömöri methenamine silver stain sections of brain; (B) thyroid; and (C) adrenal gland to show \textit{Candida albicans} dissemination and colonization in other important organs.
supplementation provided to the patient may have promoted the growth and virulence of the candida; studies have shown that the kidneys assimilate increased levels of iron during disseminated candidiasis thus providing a more favorable site for colonization.\(^\text{17}\)

The presence of *C. albicans* in a single collection of sputum or urine does not ordinarily indicate infection in an immunocompetent individual. Estimation of serum procalcitonin, a biomarker for bacterial infection and sepsis, might be useful if negative, for turning attention to a nonbacterial (such as, candida) source as the causative agent.\(^\text{18}\)

**CONCLUSION**

*C. albicans* can rarely cause disseminated disease in nonimmunocompromised critically ill patients. Low serum procalcitonin levels in a septic patient might indicate nonbacterial cause such as candidiasis. Even with disseminated candidiasis, blood cultures may remain negative.

**Author disclosures**

The authors report no actual or potential conflicts of interest with regard to this article.

**Disclaimer**

The opinions expressed herein are those of the authors and do not necessarily reflect those of *Frontline Medical Communications Inc.*, the US Government, or any of its agencies. This article may discuss unlabeled or investigational use of certain drugs. Please review the complete prescribing information for specific drugs or drug combinations—including indications, contraindications, warnings, and adverse effects—before administering pharmacologic therapy to patients.

**References**


