Abstract
Cryptococcosis is a fungal infection that primarily affects immunocompromised hosts; disseminated infection is uncommon in immunocompetent patients. We describe a case of a previously healthy woman without risk factors, who was admitted to the emergency department with headache, fever and altered mental status. As a result, a cryptococcal disseminated infection was diagnosed.

Keywords: Cryptococcus; Meningitis; Cavitation; Immunocompetent.

Resumen
La criptococosis es una infección por hongos que afecta principalmente a huéspedes inmunodeprimidos; la infección diseminada es infrecuente en pacientes inmunocompetentes. Reportamos el caso de una mujer previamente sana, sin factores de riesgo, ingresada a urgencias por cefalea, fiebre y alteración del estado mental quien posteriormente fue diagnosticada con criptococosis diseminada.

Palabras clave: Cryptococcus; Meningitis; Cavitación; Inmunocompetente.
Introduction
Cryptococcosis is a fungal infection that is found all over the world and is caused primarily by *Cryptococcus neoformans* / *Cryptococcus gattii* (*C. neoformans* / *C. gattii*). *C. neoformans* primarily affects immunocompromised patients, whereas *C. gattii* affects immunocompetent patients exposed to the ecological niche of the fungus\(^1\).

Cryptococcosis is classified as an opportunistic infection because it typically manifests in patients with predisposing factors such as liver disease, kidney disease, sarcoidosis and other immunosuppressant conditions, particularly HIV infection\(^2\).

*C. neoformans* is the leading cause of meningitis in HIV patients, with an estimated incidence of 223,100 cases per year and an 81% mortality rate. Each year, approximately 5,300 HIV-infected individuals in Latin America are affected by cryptococcal meningitis, with a mortality rate ranging from 13 to 71%; Brazil and Colombia have the highest incidence in the region (1,001-2,500 cases), followed by Argentina and Mexico with 501-1,000 cases\(^1\).

Although cryptococcal infections in immunocompetent people are rare, there is growing interest in this patient population. According to reports, between 17 and 22% of diagnosed cases are from persons with "normal" immune function; however, some have been linked to primary or acquired immunodeficiency\(^3,4\). This article emphasizes the importance of considering *Cryptococcus* as an etiological agent in previously healthy individuals who require prompt diagnosis to improve their prognosis\(^3\).

Case description
A previously healthy 19-year-old woman was admitted to the emergency department after complaining of a headache, tinnitus, blurry vision and fever for 4 weeks. Two weeks prior, she had a medical appointment for the same symptoms, which were diagnosed as anxiety disorder; upon her return, additional symptoms of hallucinations and alterations in gait were noted. The patient denied alcohol, -or illicit drugs use-, and refused a previous medical history of unsafe sexual practices, recent travel, blood transfusion, animal contact, or being exposed to the ecological niche of the fungus.

She appeared to be ill. Her body temperature was 99.5°F (37.5°C), blood pressure was 120/60 mm/Hg, heart rate was 85 beats per minute, respiratory rate was 18 breaths per minute, and the pulse oximeter read 97% at room temperature. The patient appeared lethargic and confused. She had a normal cranial nerve examination, no Kerning's or Brudzinski's signs,
no neck stiffness, no motor weakness or cerebellar ataxia, normal and symmetrical tendon reflexes with no pathological reflexes, and no sensory impairment. The rest of the physical examination was normal. During her evaluation in the emergency department, she developed a generalized tonic-clonic seizure less than 1 minute (self-limited).

The white blood cell count was 10 x 10³ /ml (5-12 x 10³), with 81% neutrophils and 8% lymphocytes; hemoglobin level was 11.5 g/dL (normal values: 13-15.7)- and serum glucose level was 69 mg/dL. Other routine laboratory tests were normal. Additional laboratory and imaging studies were performed, including two HIV 1 and 2 tests using fourth generation ELISA serologies and viral load for HIV, both of which were negative.

A chest X-ray revealed a round lung lesion on the right middle lobe that suggested cavitation (figure 1A); chest computed tomography (CT) confirmed a right middle lobe peripheral cavitary lung lesion (figure 1B); and magnetic resonance imaging (MRI) of the brain revealed leptomeningeal enhancement (figures 1C and 1D).

**Figure 1.** X-ray, CT scan and MRI.
A. Chest X-ray: Round lung lesion on the right lung base suggests cavitation
B. Chest CT: Middle lobe peripheral cavitary lung lesion
C,D. Brain MRI: Leptomeningeal enhancement
Lumbar puncture yielded an opening cerebrospinal fluid (CSF) pressure of 13 cm H$_2$O (12-15). The leukocyte count was 68 cells/ml, with 5% polymorphonuclear cells and 95% mononuclear cells, a total protein level of 150 mg/dl, a glucose level of 35 mg/dl (40-70), and a CSF/serum glucose ratio of 0.5. A cerebrospinal fluid polymerase chain cerebrospinal fluid polymerase chain (PCR) for herpes simplex virus type 1 and 2, varicella zoster virus, and cytomegalovirus was negative. A CSF GeneXpert Tb/Rif test was also negative, as were serum and cerebrospinal fluid Cryptococcal antigen latex agglutination tests.

On the India ink stain, encapsulated yeasts were compatible with *Cryptococcus* spp., and *Cryptococcus* neoformans var neoformans was isolated in Sabouraud dextrose agar cerebrospinal fluid culture (figure 2).

A video-assisted thoracoscopic surgery was performed, as well as a right lung biopsy (figure 3). The lung biopsy revealed non-caseous necrosis areas as well as a chronic inflammatory infiltrate containing spherical thick-walled microorganisms. Both PAS and mucicarmine stains confirmed round yeasts compatible with Cryptococcosis. *Cryptococcus neoformans var neoformans* was also isolated from lung tissue culture. A peripheral blood lymphocyte immunophenotype revealed a slight decrease in CD$_4^+$T cells (table 1), while serum immunoglobulins were normal.
Table 1. Immunophenotyping of lymphocyte T in the peripheral blood

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<th>%</th>
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<tr>
<td>CD₃</td>
<td>79.0</td>
<td>50.1 – 79.4</td>
<td>728</td>
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<tr>
<td>CD₄</td>
<td>42.4</td>
<td>23.0 – 44.5</td>
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<td>CD₈</td>
<td>29.9</td>
<td>16.3 – 41.7</td>
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<td>CD₁₆ +CD₅₆</td>
<td>9.1</td>
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Figure 3. Diagnostic testing.
A. Lung H&E stain, 40X. B. Lung H&E stain, 100X. C. Lung PAS stain, 100X. D. Mucicarmine stain
The patient was diagnosed with disseminated cryptococcal disease and was treated for four weeks with liposomal amphotericin B 4 mg/kg IV QD (once a day) and fluconazole 800 mg IV QD, followed by fluconazole 400 mg QD maintenance therapy. She developed new onsets of altered mental status, headaches, meningismus, papilledema, hearing loss, and cranial nerve III and VI palsy shortly after switching to maintenance treatment. The intracranial pressure was noticeably elevated. Despite repeated lumbar punctures showing temporal improvement, a ventricular peritoneal shunt was required due to persistently high intracranial pressure. All her symptoms subsided, and she was discharged from the hospital asymptomatic. She continues the treatment phase with fluconazole 400 mg QD daily for the next 12 months.

Discussion
Disseminated cryptococcosis in immunocompetent individuals is uncommon. The lungs are the primary site of infection, and inoculation is usually via inhalation of yeast spores, with hematogenous spread mainly to the central nervous system\(^{(5)}\). Although the initial symptoms were from central nervous system involvement, a cavitary lesion at the lung level could be seen in this case. This radiological finding is unusual in immunocompetent patients, where non-calcified nodulation is more common in pulmonary cryptococcosis\(^{(6)}\).

The diagnosis of cryptococcal meningitis in immunocompetent hosts is difficult and delays in diagnosis have been shown to have negative consequences for patients with a reported mortality rate of 9-27%. Due to brain-lung involvement, the diagnostic consideration was made in this case during the second admission and cryptococcus was actively searched for during the study. Although the changes in cerebrospinal fluid cytology are indistinguishable from those seen in HIV patients\(^{(8)}\), some authors have suggested ruling out meningeal cryptococcosis when lymphocytosis is found in the cerebrospinal fluid in meningitis\(^{(7)}\).

In the diagnostic approach, the antigen test for cryptococcus has a sensitivity of 97% in cerebrospinal fluid and 87% in serum; false negatives have been described due to prozone effect, as may have occurred in this case; a negative test does not preclude diagnosis of cryptococcosis, particularly if only a single specimen has been tested and the patient exhibits symptoms consistent with cryptococcosis\(^{(7,9)}\). However, it has been shown that the India Ink of the cerebrospinal fluid has a 70–90% sensitivity in cryptococcal meningitis\(^{(5)}\).
The diagnosis of disseminated cryptococcosis is defined by the identification of the fungus in blood cultures or isolated in cultures from two sites. In this case, the fungus was identified in lung tissue and cerebrospinal fluid with the growth of *Cryptococcus neoformans var neoformans*. In addition, the gold standard in lung cryptococcosis demonstrated the presence of the fungus in biopsy\(^{10}\).

As recorded in outbreaks in Canada and the United States\(^{11}\) over the last decade, there has been an increase in cases of *Cryptococcus* infections in individuals without risk factors; this group may have subclinical innate or acquired immunodeficiencies.

*Cryptococcus* infections have been consistently reported in patients with idiopathic CD\(_4\) lymphopenia count below 200-100 cells/μl\(^{12}\). The absolute lymphocyte count in this patient was always abnormal, and a peripheral blood lymphocyte immunophenotype revealed a slight decrease in CD\(_4\)+ T cells; serum immunoglobulins were within normal limits and other acquired immunodeficiencies were ruled out. Other tests for the laboratory diagnosis of primary immunodeficiencies, such as flow cytometric-based assays to test immune cell function (neutrophil oxidative burst, NK cytotoxicity), intracellular cytokine production (TH17), cellular signaling pathways (phosphor-STAT analysis), and protein expression (BTK, Foxp3)\(^{13-15}\), are not available in our institution.

In this group of otherwise normal patients, the outcomes and complications may be more severe, including more likely permanent neurologic sequelae such as stroke, blindness, deafness, and other focal cranial nerve abnormalities\(^{7,13}\). Some studies have suggested that immunocompetent individuals develop post-infectious inflammatory response syndrome (PIIRS), a paradoxical worsening of symptoms caused by immune dysregulation or aberrant immune control, like immune reconstitution syndrome (IRIS) in patients with HIV infection, after the start of antiretroviral therapy\(^{5,14,15}\).

In both cryptococcal meningitis IRIS and PIIRS, there is activation of the dendritic T-cells of the synapse and an increase in IFN-γ and IL-6 cytokines in response to T-cell activation; however, PIIRS lacks effective macrophage activation. This macrophage dissociation -T cell leads to the persistence of tissue antigen, which perpetuates the inflammatory response, resulting in clinical deterioration despite adequate antifungal therapy\(^{5}\).
This case teaches that disseminated cryptococcal infection in immunocompetent patients is uncommon, can manifest in a variety of clinical presentations and innate or acquired immunodeficiencies must be ruled out.

**Conclusion**
Disseminated cryptococcosis in an immunocompetent patient is difficult to diagnose; a high index of suspicion and early treatment are essential for reversing this otherwise fatal condition.

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Authors.

**Conflicts of Interest**
The authors declare that there are no conflicts of interest.

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**Protection of human and animal subjects**
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**Confidentiality of data**
No data that identifies patient are revealed.

**References**


