



When dyspnea is a Hickam's Dictum

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ABSTRACT

Synchronous opportunistic infections are luckily rare in people living with HIV (PLWH) in the era of highly effective antiretroviral medications. We describe the case of a middle-aged man who presented with diarrhea and shortness of breath and was found to have pneumocystis pneumonia, disseminated histoplasmosis and disseminated mycobacterium avium complex infection along with a new diagnosis of human immunodeficiency virus (HIV) infection. This case highlights that individuals who remain undiagnosed with HIV infection for a long time can still present with concurrent infections and clinicians should remain aware of this.

Case

A 41-year-old male without any notable past medical history presented with 3 weeks of progressive watery diarrhea associated with severe fatigue and one week of non-productive cough and dyspnea. He was afebrile, tachycardic (140 beats/min), hypertensive (162/99 mmHg) and had oxygen saturation of 97 % on room air. He appeared chronically ill but was not in distress. Screening for HIV was reactive and later confirmed positive, and subsequent testing revealed an absolute CD4 count of 28 cells/mm³ and viral load of 721,988 copies/mL. His hospital course was complicated by acute renal failure requiring hemodialysis and acute hypoxic respiratory failure requiring intubation. Imaging revealed extensive bilateral interstitial pulmonary opacities and multiple enlarged retroperitoneal lymph nodes. Bronchoscopy was performed. Direct fluorescent antibody (DFA) smear from bronchoalveolar lavage (BAL) demonstrated a large cyst with honeycomb appearance fluorescing bright green at an excitation wavelength of 490–500 nm (Fig. 1a) and was identified as *Pneumocystis jirovecii*. Bronchoalveolar lavage (BAL) specimen was plated on an inhibitory mold agar plate and grew *Histoplasma capsulatum*, colonies of which appear as a white mold with a cotton consistency, after 15 days of incubation at 25 °C (Fig. 1b). This dimorphic ascomycete also grew in regular blood cultures and AFB blood cultures and was seen as yeast phagocytosed within a segmented neutrophil (Fig. 1c). In addition, AFB blood cultures showed positive staining bacilli on the Kinyoun stain which was identified as *Mycobacterium avium* complex (MAC) (Fig. 1d).

Our patient was initiated on intravenous Liposomal amphotericin B 3 mg/kg/day for disseminated histoplasmosis and was later transitioned

to oral itraconazole 200 mg twice daily. Oral trimethoprim-sulfamethoxazole was initiated for *Pneumocystis jirovecii*. He received oral ethambutol 15 mg/kg/day, oral rifabutin 1500 mg once daily and intravenous amikacin for disseminated MAC infection. Azithromycin was deferred due to concern for prolongation of Qtc with concurrent azole use. Antiretroviral therapy consisting of oral abacavir, lamivudine and dolutegravir was initiated. He had a prolonged hospital stay but recovered completely.

Concurrent infection with *Pneumocystis jirovecii*, *Histoplasma capsulatum* and *Mycobacterium avium* complex is a very rare entity among patients living with HIV in the era of highly effective antiretrovirals [1]. This case highlights the occurrence of these synchronous infections in individuals with Acquired Immune Deficiency Syndrome (AIDS). These coinfections portend significant morbidity and mortality but can be successfully treated with appropriate antimicrobials, antiretroviral therapy, and supportive care [2,3].

CRediT authorship contribution statement

Annette Abraham: Writing – original draft. **Nicholas Moore:** Writing – review & editing. **Sarah E. Sansom:** Conceptualization, Writing – review & editing.

Ethical approval

Not applicable.

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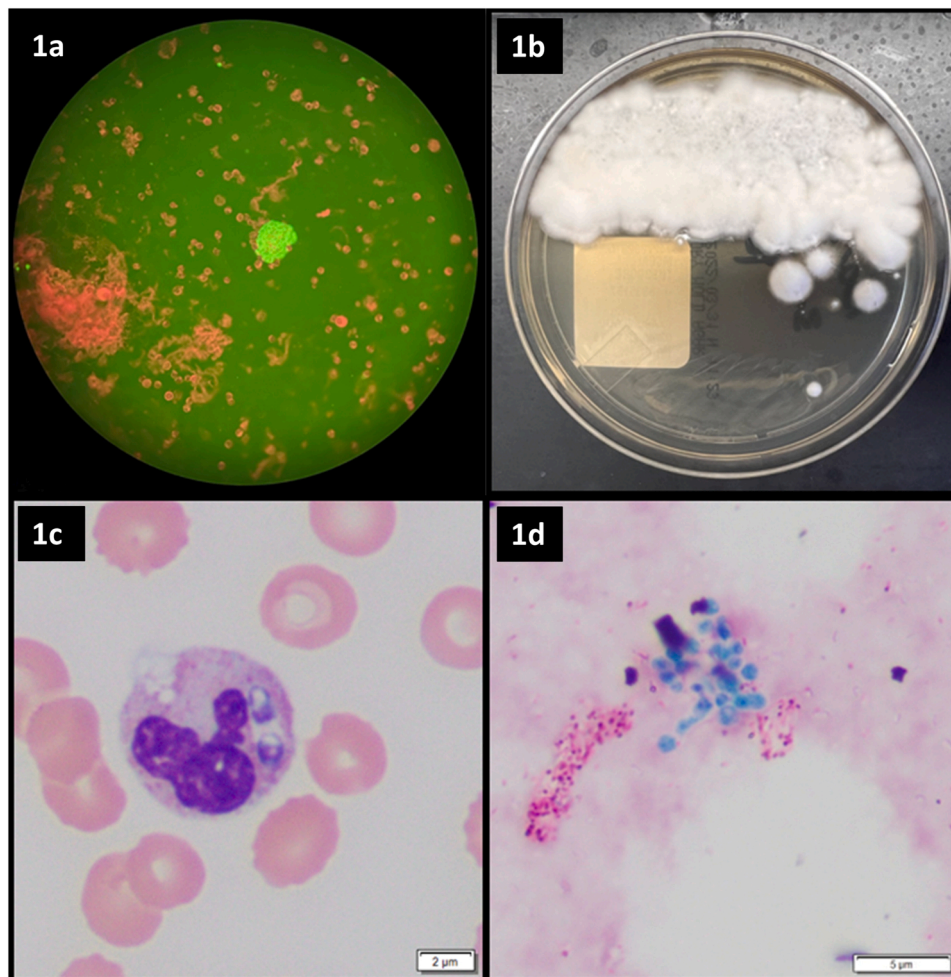


Fig. 1. (1a) Direct fluorescent antibody stain performed from the bronchoalveolar lavage fluid. (1b) Colony morphology of *Histoplasma capsulatum* on Sabouraud dextrose agar isolated from the BAL fluid after 12 days of incubation. (1c) Wright Giemsa stain showing yeast phagocytosed within neutrophils. (1d) Kinyoun stain performed from the Mycobacterium growth indicator tube after incubation onboard the Bactec MGIT system.

Consent

Not applicable

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Declarations of interest

None.

References

- [1] Nevez G, Le Gal S. Pulmonary co-infection with *Pneumocystis jirovecii* and *Histoplasma capsulatum* in AIDS patients is not a rare event. *Int J Infect Dis* 2019;87: 126–7. <https://doi.org/10.1016/j.ijid.2019.08.012>.
- [2] Carreto-Binaghi LE, Morales-Villarreal FR, García-de la Torre G, Vite-Garin T, Ramirez J, Aliouat E, et al. *Histoplasma capsulatum* and *Pneumocystis jirovecii* coinfection in hospitalized HIV and non-HIV patients from a tertiary care hospital in Mexico. *Int J Inf Dis* 2019;86:65–72. <https://doi.org/10.1016/j.ijid.2019.06.010>.
- [3] Dugan J, Grinsztejn E, Glasgow A, John A. Co-infection with *Histoplasma Capsulatum* and *pneumocystis jirovecii* in a patient with liver cirrhosis: a diagnostic dilemma. *Chest* 2020;158:A529–30. <https://doi.org/10.1016/j.chest.2020.08.50>.