Psoas Abscess Caused by Candida Glabrata: A Case Report

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Abstract

Psoas abscess caused by Candida is an uncommon condition. We report a case of psoas abscess caused by Candida glabrata, which was completely resolved with drainage and oral voriconazole. Because of the nonspecific clinical presentation, the diagnosis of psoas abscess can be a challenge. Prompt suspicion, with early diagnosis and drainage with an appropriate antifungal agent, seems to improve the clinical outcome.

Categories: Infectious Disease

Keywords: candida glabrata, psoas abscess

Introduction

Psoas abscess is an uncommonly encountered clinical entity [1]. It could spread from an adjacent anatomical area, such as a vertebral abscess, gastrointestinal or genitourinary focus, or occasionally seeding from a hematogenous infection [2]. Patients commonly present with nonspecific symptoms such as abdominal pain [3]. A computed tomogram of the abdomen is used to diagnose the condition. The most commonly implicated organisms are *Staphylococcus aureus* and *Mycobacterium tuberculosis (M. tuberculosis)* [4-6]. Psoas abscess caused by Candida species has only rarely been described.

Case Presentation

A 35-year-old female was admitted with fever, dysuria, and suprapubic pain for one week. Her past medical history was significant for diabetes mellitus (DM) and stage 1 B endometrial carcinoma for which she underwent a robotic-assisted laparoscopic hysterectomy, with bilateral salpingo-oophorectomy and lymph node dissection, four months prior to the admission.

Vital signs on admission revealed a temperature of 102.7F, blood pressure 111/70 mm/Hg, heart rate 107/minute, respiratory rate 18/minute, with oxygen saturation of 93% on room air. There was tenderness over the suprapubic and right-sided lumbar area with a full range of motion of the lower extremities.

Laboratory data revealed a white blood cell count of 6X103/µL, hemoglobin 9.7 g/dL (Normal range=12 to 16.3 g/dL), platelet count 365 X103/µL (Normal range=150-450X103/µL), creatinine 0.59 mg/dl (Normal range=0.7-1.3 mg/dl), erythrocyte sedimentation rate 100 mm/hour (Normal range=0-15 mm/hour), and C-reactive protein 27.10 mg/dl(Normal range< 0.29 mg/dl). Urinalysis revealed trace protein, with positive leukocyte esterase, negative nitrite and white blood cells of 240, trace protein. The patient was empirically started on intravenous vancomycin 1 gram every 12 hours and intravenous piperacillin/tazobactam 3.375 grams every eight hours for a presumed urinary tract infection.

The patient underwent computed tomography (CT) of the abdomen/pelvis with contrast, which revealed multiloculated abscess 8 x 9 x 9 cm in the right psoas and quadratus lumborum musculature, however, an underlying necrotic tumor could not be excluded (Figure 1). The patient underwent ultrasound-guided drainage with the placement of a pigtail catheter and the removal of 25 cc of purulent fluid. Gram stain of the fluid showed many white cells and rare yeast. Blood cultures were negative and urine cultures grew Escherichia coli (E. coli) sensitive to ceftriaxone, cefazolin, ceftazidime, nitrofurantoin, and piperacillin/tazobactam. Vancomycin was discontinued, and the patient was continued on piperacillin/tazobactam to complete a seven-day course for a urinary tract infection. Culture of the drained abscess grew Candida glabrata in pure culture. The organism was sensitive to fluconazole with a minimum inhibitory concentration (MIC) of <2 µg, caspofungin MIC < 0.25 µg, and voriconazole MIC <0.12 µg. The patient was started on intravenous fluconazole 400 mg every 24 hours based on susceptibility to fluconazole. Repeat CT of the abdomen/pelvis with contrast was performed two days after starting intravenous fluconazole, which showed an increase in the size of the heterogeneous collection within the right psoas and paravertebral muscles, with its largest component measuring 8 x 9 x 8 cm. This collection extended posterolaterally to involve the right external and internal oblique muscles and more of the right paravertebral musculature. There was overall increased fat stranding within this region. This mass also extended anteriorly with a component just inferior to the lower pole of the right kidney, measuring approximately 3 x 4 x 4 cm; the drainage catheter, with its tip, was seen within posterior right subcutaneous

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Nagarakanti et al. This is an open access article distributed under the terms of the Creative Commons Attribution License CC-BY 4.0., which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited. fat (Figure 2). The patient underwent new ultrasound-guided drainage of the abscess with the removal of 30 cc of purulent fluid and placement of a new pigtail catheter with the removal of the old drain. Intravenous fluconazole was changed to intravenous caspofungin, as the CT abdomen showed an increase in the collection, as susceptibility in-vitro does not always predict the clinical response. The patient continued on intravenous caspofungin for six days. Repeat CT of the abdomen showed an interval decrease in the size of the abscess at the site of the pigtail catheter placement and an interval increase in the size of 4.5 x 2.3 cm abscess in the subcutaneous fat in the right flank superficial to the drainage catheter (Figure 3). The pigtail catheter was removed and intravenous caspofungin was changed to oral voriconazole 200 mg twice daily, and the patient was discharged home. The serum voriconazole level was maintained at 3.1 µg/ml. The patient remained asymptomatic and underwent another CT abdomen four weeks later, which showed complete resolution of the psoas abscess, a necrotic right retroperitoneal mass measuring approximately 4.1 cm, which is inseparable from the right iliopsoas muscle, which is consistent with metastatic disease. There were additional smaller adjacent lymph nodes and no free intraperitoneal air or loculated fluid collections (Figure 4).



FIGURE 1: CT abdomen/pelvis with contrast on admission Multiloculated abscess 8 x 9 x 9 cm in the right psoas and quadratus lumborum musculature

CT: computed tomography



FIGURE 2: CT abdomen and pelvis with contrast after drainage and two days of antifungals

Abscess in the right psoas and paravertebral muscles measuring $8 \times 9 \text{ cm} \times 8 \text{ cm}$ extending posterolaterally to involve the right external and internal oblique muscles

CT: computed tomography



FIGURE 3: CT abdomen and pelvis with contrast after drainage and six days of antifungals

Interval decrease in the size of the abscess at the site of the pigtail catheter placement

CT: computed tomography

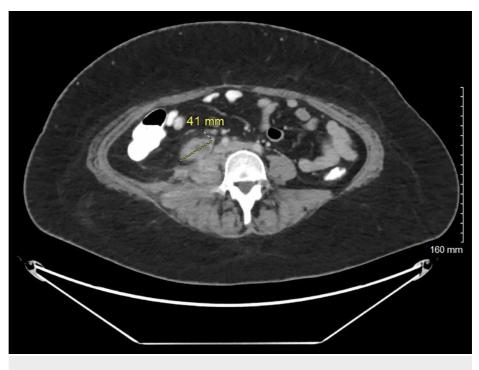


FIGURE 4: CT abdomen/pelvis with contrast at four weeks

Complete resolution of the psoas abscess, a necrotic right retroperitoneal mass measuring approximately 4.1 cm, which is inseparable from the right iliopsoas muscle, which is consistent with metastatic disease

CT: computed tomography

Discussion

Psoas abscess is a rare condition with a reported worldwide incidence of 12 new cases per year [1]. The condition was first described by Mynter in 1881 as psoitis: "a collection of pus in the Psoas muscle" [2]. The psoas muscle courses from the lower part of the thorax to the lower lumbar vertebrae and femur through the retroperitoneum [3]. The muscle has a rich blood supply, making it susceptible to the hematogenous seeding of bacteria, and its anatomical closeness to the gastrointestinal system and spinal column, making it susceptible to infections via direct spread from these sites [4-5].

The most commonly described conditions associated with psoas abscess include vertebral osteomyelitis, epidural abscess, septic arthritis, discitis, sacroiliitis, gastrointestinal diseases, such as Crohn's disease, appendicitis, pancreatitis, infected aortic aneurysms, rectal surgery, and nephrectomy, and immunosuppressive conditions such as malignancy, organ transplantation, human immunodeficiency syndrome (HIV) infection, and tuberculosis of the spine (Pott's disease) [5-7].

Most patients with psoas abscess present with nonspecific symptoms, such as abdominal pain. Patients commonly have an elevated white blood cell (WBC) count and elevated inflammatory markers such as erythrocyte sedimentation rate and C-reactive protein [6]. CT of the abdomen is the most common modality to diagnose psoas abscess. Tabrizian et al. reported that the mean size of the abscess was 6 cm at diagnosis, the abscesses were unilateral in over 50% of the patients and multiple in 25% of the patients [6]. The most common etiology of psoas abscess is *Staphylococcus aureus*, followed by *Streptococci* and *E. coli* [5,8-9]. *M. tuberculosis* was the most common isolate in another psoas abscess case series [7].

Candida sp. is a rare pathogen causing psoas abscesses. *Candida albicans* is reported as a cause of psoas abscess; one case reported a 59-year-old female with diabetes mellitus who presented with a right flank mass of which an aspirate grew *Candida albicans* [10]. *Candida glabrata* as a cause of psoas abscess was reported in two cases, both in the non-English language literature [11-14]. The first case was reported in one out of 38 psoas abscess cases reviewed by Hammani et al. [15]. A second case was described in a 51-year-old diabetic male who was admitted with hip pain and fever, the patient was diagnosed with a psoas abscess with the isolation of *Candida glabrata* from the abscess. The patient improved after percutaneous drainage and oral voriconazole [16].

Our patient's case is similar to the case described by Kim et al. [16], in that it occurred in a patient with underlying DM when both cases improved after percutaneous drainage and voriconazole treatment.

The successful treatment of psoas abscess consists of drainage of the abscess along with systemic antifungal agents [6]. In some cases, surgical drainage is required, and this approach has been shown to have a higher success rate in the treatment than percutaneous drainage [8].

It is not clear why our patient developed this rare condition but as mentioned previously, she had previously described risk factors such as DM as well as endometrial cancer.

Conclusions

To our knowledge, our case is the first case of *Candida glabrata* psoas abscess described in the English literature. Because of the nonspecific clinical presentation, the diagnosis of psoas abscess can be a challenge to the clinician. Early diagnosis with prompt drainage with appropriate antifungal agent seems to improve outcomes.

Additional Information

Disclosures

Human subjects: Consent was obtained by all participants in this study. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

- Garner JP, Meiring PD, Ravi K, Gupta R: Psoas abscess not as rare as we think? . Colorectal Dis. 2007, 9:269-274. 10.1111/j.1463-1318.2006.01135.x
- 2. Mynter H: Acute psoitis. J Buffalo Med Surg. 1881, 21:202-210.
- Cronin CG, Lohan DG, Meehan CP, Delappe E, McLoughlin R, O'Sullivan GJ, McCarthy P: Anatomy, pathology, imaging and intervention of the iliopsoas muscle revisited. Emerg Radiol. 2008, 15:295-310. 10.1007/s10140-008-0703-8
- Leu SY, Leonard MB, Beart RW Jr, Dozois RR: Psoas abscess: changing patterns of diagnosis and etiology. Dis Colon Rectum. 1986, 29:694-698. 10.1007/BF02555310
- Ricci MA, Rose FB, Meyer KK: Pyogenic psoas abscess: worldwide variations in etiology. World J Surg. 1986, 10:834-843. 10.1007/BF01655254
- Tabrizian P, Nguyen SQ, Greenstein A, Rajhbeharrysingh U, Divino CM: Management and treatment of iliopsoas abscess. Arch Surg. 2009, 144:946-949. 10.1001/archsurg.2009.144
- Turunç T, Turunç T, Demiroğlu YZ, Colakoğlu S: Psoas apsesi olan 15 hastanin retrospektif olarak değerlendirilmesi [Article in Turkish]. Mikrobiyol Bul. 2009, 43:121-125.
- Santaella RO, Fishman EK, Lipsett PA: Primary vs secondary iliopsoas abscess. Presentation, microbiology, and treatment. Arch Surg. 1995, 130:1309-1313. 10.1001/archsurg.1995.01430120063009
- Desandre AR, Cottone FJ, Evers ML: Iliopsoas abscess: etiology, diagnosis, and treatment. Am Surg. 1995, 61:1087-1091.
- 10. Fukuhara S, Nishimura K, Yoshimura K, Okuyama A, Yamato M, Kawamori D, Matsuhisa M: A case of psoas abscess caused by Candida albicans [Article in Japanese]. Hinyokika Kiyo. 2003, 49:141-143.
- 11. Thompson WC, Brock JW 3rd: Torulopsis glabrata perinephric abscess: a case report . J Urol. 1983, 130:529-530. 10.1016/s0022-5347(17)51288-4
- 12. Khauli RB, Kalash S, Young JD Jr: Torulopsis glabrata perinephric abscess. J Urol. 1983, 130:968-970. 10.1016/s0022-5347(17)51599-2
- Balkan II, Savas A, Geduk A, Yemisen M, Mete B, Ozaras R: Candida glabrata perinephric abscess. Eurasian J Med. 2011, 43:63-65. 10.5152/eajm.2011.14
- 14. Khemakhem B, Kanoun F, Ben Jemaa M, et al.: Candida glabrata perinephric abscess. A case report [Article in French]. Ann Med Interne. 2001, 152:134-136.
- 15. Hammami BK, Ghorbel H, Abid F, Arab NB, Maâloul I, Hammami A, Jemâa MB: Psoas abscess of the adult: study of 38 cases [Article in French]. Tunis Med. 2007, 85:631-636.
- 16. Kim M, Lee J, Chung CS, Jang JS, Ahn BM, Son W: A case of psoas abscess caused by Candida glabrata . Infect Chemother. 2010, 42:43-47. 10.3947/ic.2010.42.1.43