Post-Covid Renal Mucormycosis: A Case Report

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Abstract

The Pandora’s box has been opened in the twenty-first century unleashing “the Coronavirus.” There has been an increase in the incidence of mucormycosis during this coronavirus disease 2019 (COVID-19) pandemic with many case reports of rhino-orbito-cerebral and pulmonary mucormycosis. Diabetes mellitus, COVID, and Mucor fungus together are a dangerous trio associated with high mortality. We report a rare case of renal mucormycosis following COVID pneumonia in a 64-year-old patient with uncontrolled diabetes; with the imaging findings of left emphysematous pyelonephritis and cystitis, nonfunctioning left kidney, and left main renal artery and vein thrombosis. This is the first such case reported to our knowledge. A high degree of clinical suspicion with prompt recognition of imaging findings is essential to improve the chances of survival in such patients.

Keywords

► COVID-19
► diabetes
► renal mucormycosis

Introduction

The coronavirus disease 2019 (COVID-19) pandemic continues despite affecting more than 196 million cases and claiming more than 4.2 million lives globally.1 Adding to the disease burden, we also have incidences of opportunistic fungal infections including aspergillosis, candidiasis, and mucormycosis in patients either after recovery or during the resolving phase of COVID-19 pneumonia.2 There has been a surge of the COVID-19-associated mucormycosis (CAM) especially in India.3 The various presentations of mucormycosis in decreasing order of frequency include oculo-rhino-cerebral, pulmonary, gastrointestinal, cutaneous, and disseminated mucormycosis.4 There has been a case report of fatal primary renal mucormycosis in a post-COVID young male with no prior risk factors, with the imaging finding of renal infarct; the prior history of COVID pneumonia could have been coincidental rather than contributory in that case.5 There has been another case report of renal mucormycosis with a nonfunctioning kidney and main renal artery thrombosis in a post-COVID patient; much similar to our case.6 Till date there have been no reports of emphysematous pyelonephritis with main renal artery and vein thrombosis; an aggressive presentation of renal mucormycosis in a post-COVID individual. Here, we report one such rare case.

Case Report

A 64-year-old male patient presented to the emergency department with complaints of abdominal pain, dysuria, hematuria, and decreased urine output for 1 week. He was a known diabetic and hypertensive on oral hypoglycemic agents and antihypertensives for the past 10 years. He was a known diabetic and hypertensive on oral hypoglycemic agents and antihypertensives for the past 10 years. There were no presenting complaints of fever, breathing difficulty, or loss of consciousness. There was history of fever 1 month back with a positive reverse transcription polymerase chain reaction test for COVID-19; he was treated at an outside
hospital and discharged following which he was apparently asymptomatic till a week back, when he developed the abovementioned symptoms and was referred to our hospital for further management. On examination, the patient was tachypneic and afebrile; blood pressure and oxygen saturation were within normal limits. Diffuse abdominal tenderness more over bilateral loin region was noted.

Laboratory examination revealed normal serum creatinine (0.9 mg/dL initially and an increased total leukocyte count (23,000 cells/mm³) with neutrophilia (94%) and severe lymphopenia (2%). There was elevated erythrocyte sedimentation rate (118 mm/h), C-reactive protein (26.6 mg/dL), D-dimer (1,670 ng/mL), and procalcitonin (6.2 ng/mL) levels pointing toward sepsis. The patient also had poor glycemic control with increased random blood sugar (253 mg/dL), serum ketones (28.0 mg/dL), and glycated hemoglobin levels (11.7%). Urine analysis showed few pus cells and red blood cells on microscopy; no fungal elements were seen. Urine dipstick reflectance photometry detected protein, blood, and glucose but was negative for ketones in urine. There was no growth in aerobic bacterial culture from urine sample.

An unenhanced computed tomography (CT) of the patient’s abdomen and pelvis was performed using 128 slice Multidetector CT-Optima 660 (GE Healthcare, Milwaukee, Wisconsin, United States); following which 100 mL of iodinated contrast Omnipaque (GE Healthcare, Shanghai, China) was administered at a flow rate of 4 mL/s using a pressure injector. Images with a slice thickness of 5 mm and 0.625 mm reconstruction in the nephrogenic, excretory, and delayed phase were obtained at a time delay of 55 seconds, 90 seconds, and 5 minutes, respectively. The study revealed an enlarged left kidney with perinephric fat stranding, thickening of Gerota’s fascia, and multiple air pockets in the left kidney, left upper ureter, urinary bladder, and surrounding soft tissues (►Figs. 1 and 2). There was complete absence of contrast enhancement in the left renal parenchyma (►Fig. 3) with thrombosis of the left main renal artery (►Fig. 4) and thrombosis with air foci within the left main renal vein (►Fig. 5). No evidence of extension of thrombus into the great vessels noted. There were no renal or perinephric abscess/collections. There was no excretion of contrast into
the left pelvicalyceal system (► Fig. 6). Visualized lung bases showed an area of consolidation with central ground-glass opacity (atoll sign) and cavitation in left lower lobe and bilateral minimal pleural effusion (► Fig. 7A and B). A possibility of renal mucormycosis with superadded anaerobic infection was considered.

Cystoscopy was done under general anesthesia which revealed a pus-filled bladder and multiple bladder washes were given. The right ureteric orifice was identified and a double-J stent was placed in the right-side collecting system. The left ureteric orifice could not be identified. The patient was then explored through a left flank cutting incision; left nephrectomy along with removal of left upper ureter and adrenal gland was performed with multiple washes and closed. Gross examination revealed the entirely necrosed left kidney with surrounding fat saponification (► Fig. 8). The histopathological examination (► Figs. 9, 10, 11) showed the necrotic renal parenchyma with neutrophilic infiltration and fungal hyphae with evidence of angioinvasion. The hyphae were aseptate, ribbon-like with a large diameter, irregular width, and were branching at right angles. The left ureter and adrenal gland were not involved.

Despite treatment, the sepsis worsened and the patient continued to deteriorate. The attenders wished to continue his treatment elsewhere and got discharged against medical advice and had succumbed to the disease in a week.

Discussion

Mucormycosis is caused by a group of angioinvasive fungi under the order Mucorales; the most common species involved are the Rhizopus and Mucor. Various risk factors are known to be associated, the most common being
uncontrolled diabetes mellitus; others include hematological malignancy, stem cell or organ transplant recipients, neutropenia, desferrioxamine therapy, and corticosteroid use. An increase in the incidence of CAM is attributed to the widespread use of steroids, hypoxemia, prolonged hospitalization needing intensive care, and previously immunocompromised individuals; but cases have also been reported in immunocompetent individuals with no prior risk factors. The COVID upregulates the glucose receptor protein GRP-78 which helps in the penetration and damage of endothelial cells by Mucor fungi leading to angioinvasion. Glucose, iron, and β-hydroxybutyrate enhance the fungal growth. The COVID-associated inflammation or its treatment modalities might contribute to the increase in these metabolites predisposing the individual to fungal infections. There might exist a deeper pathophysiological link between the two pathogens that waits to be unraveled yet.

Phagocytes (neutrophils) are the first line of body’s defense against fungal pathogens. These white blood cells engulf them in a process called phagocytosis. Sometimes a reverse process occurs, called the “vomocytosis,” in which the pathogen is ejected back out of the cell without the lysis of either the host cell or the pathogen. There has been no study regarding the demonstration of vomocytosis with Mucor fungi yet. There has been a study which showed viral coinfection can enhance the rate of vomocytosis of fungus by macrophages. We hypothesize that COVID coinfection could enhance the vomocytosis of Mucor by macrophages and when at the wrong time and wrong place be catastrophic for the patient. COVID has altered the immune system of our patient predisposing him to fungal infection; the lungs were already affected by the COVID, hence the primary site of inoculation could have been the lung and being an uncontrolled diabetic with ketoacidosis provided a favorable environment for the growth of fungi; the Mucor could have then reached the kidney with the help of phagocytosis and vomocytosis by the neutrophils.

Ultrasonography remains the first line of investigation and can show the enlarged echogenic kidneys with hypoechoic areas of abscesses, perinephric fluid collections, hydronephrosis, and cystitis. CT features include diffuse patchy nephrogram, inhomogeneous enhancement with areas of low attenuation, perinephric fluid collections, and no contrast excretion. There can be multiple hypodense nonenhancing areas (intrarenal segmental infarcts) or even complete absence of contrast enhancement of the kidney. There can be thrombosis of the main renal artery and vein, but are usually spared with thrombosis of intrarenal segmental branches leading to infarction, which can be demonstrated on Doppler sonography. Magnetic resonance imaging study will show hypointense signal on T1-weighted imaging (WI) and hypointense signal on T2WI with diffusion restriction; in contrast to other infections which will be hyperintense on T2WI with diffusion restriction owing to the edema and inflammation. This typical T2 hypointensity with diffusion restriction seen in mucormycosis might be attributed to the completely infarcted tissue with necrotic debris and fungal filled abscesses with high iron (paramagnetic element) content in these tissues and fungi. Other renal fungal infections like candidiasis present as multiple microabscesses, papillary necrosis, and fungal balls within the collecting system whereas renal aspergillosis present as focal mass lesions.

Though the suspicion of mucormycosis can be made with supportive clinical and imaging findings, a definitive diagnosis can be made only with histopathological findings. The treatment of renal mucormycosis is intravenous liposomal amphotericin B and emergency nephrectomy with debridement of nearby involved tissues. Despite treatment, the mortality rate associated with mucormycosis in general are very high averaging around 54% with the highest mortality rate of 96% for disseminated mucormycosis.

Conflict of Interest
None declared.

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References