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Invasive Klebsiella pneumoniae Liver Abscess with Multiple Septic Metastatic Complications – Case Report and Review of Literature

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Authors' contributions

This work was carried out in collaboration between all authors. Author KK did the study design, literature review, data gathering, drafting of manuscript and critical revision. Author EE did the study design, literature review and critical revision. Author HR did study design, literature review, data gathering and critical revision. All authors read and approved the final manuscript.

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Case Study

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ABSTRACT

Klebsiella pneumoniae is a common bacterial pathogen that is known to cause diverse community acquired and nosocomial infections [1,2]. These infections commonly occur in hospitalized individuals with impaired host defenses [3]. However the proportion of community acquired vs nosocomial infection has markedly increased in recent years [4]. A community acquired syndrome of cryptogenic invasive Klebsiella pneumoniae Liver Abscess Syndrome (CIKPLA) has been known to typically affect individuals of Southeast Asian origin, and is complicated by devastating septic spread to other organs [5]. We report a case of community acquired invasive Klebsiella pneumoniae infection with multi organ involvement in an individual of non-Asian origin. The aim of this case report is to alert clinicians about increasing prevalence of CIKPLA syndrome, its different clinical presentations and occurrence in different populations suggesting a need for increased vigilance for early diagnosis and prompt management to prevent disastrous sequelae.

Keywords: Klebsiella pneumonia; liver abscess; septic; CIKPLA; endophthalmitis.

1. CASE REPORT

A 64 year old Sudanese male with a background of hypertension and diabetes mellitus for 13 years presented with history of low grade fever and productive cough of 10 days duration. Two days after the fever, the patient developed pain in the right ankle along with redness and mild pain and swelling in the left calf. He visited a private clinic and received amoxicillin for 5 days without resolution of his symptoms. On the day of admission he developed dull aching, continuous, non- radiating pain on the right side of the abdomen, unrelated to food intake and without alteration in bowel or bladder habits, nausea, vomiting or iaundice along with urinary discomfort. The following day he also developed pain, redness and photophobia along with sticky. mucoid discharge from his right eye.

On physical examination, he had a regular pulse of 80 beats/min, a respiratory rate of 17 breaths/min, temperature of 38.2℃ and a blood pressure of 145/90 mmHg. The other notable physical findings were distention of neck veins. Pitting pedal edema was present bilaterally up to the mid-calf region. There was a red, warm, tender swelling over the right medial malleolus measuring 7 X 5 cm with minimal purulent discharge. Abdominal examination revealed tenderness in the right upper quadrant and a palpable liver edge of 4 cm. Respiratory examination revealed decreased breath sounds and crackles over right basal area. The right eve was chemosed with conjunctival and ciliary injection along with mucoid discharge. The left eye appeared normal. Visual acuity was counting fingers at 1 meter & at 5 meters for right and left eye, respectively. Pupil was sluggishly reactive on the right side and normal on the right. Extra ocular movements were normal bilaterally. Other systems were unremarkable.

His initial laboratory tests showed a platelet count of 132 X 103 U/L with a normal white cell count and hemoglobin. The serum sodium was 125 mmol/L, bicarbonate of 18 mmol/L and blood glucose of 23.42 mmol/L. The remaining electrolytes were within the reference range. Of note was a serum albumin of 26 g/L, ALT of 92 U/L, ALP of 225 with a normal AST, lipase and bilirubin. Chest X-ray revealed evidence of bilateral increased bronchovascular markings with a homogenous opacity on the right side, extending up to the right middle zone with an air-bronchogram suggestive of a parapneumonic

effusion (Fig. 1). Blood cultures were positive for Klebsiella pneumoniae resistant only to ampicillin. Culture from the urine sample and the leg ulcer also grew the same organism. Ultrasound (Fig. 2) abdomen revealed a liver abscess. A CT guided percutaneous drain yielded 250 cc of purulent fluid, with Klebsiella pneumoniae isolated from cultures of the aspirated pus. The patient was initially started on Piperacillin/Tazobactam and urgent ophthalmology consultation was sought. Despite the initial antibiotic treatment, the patient had worsening eye symptoms with tense proptosis and conjunctival chemosis. The patient was diagnosed with endogenous ophthalmitis and total retinal detachment. CT scan showed orbital cellulitis. The antibiotic was changed to Meropenem and IV steroids were added. Unfortunately he developed globe perforation of right eye and underwent evisceration under general anesthesia. The patient's general condition improved after this. He completed the course of antibiotics and was subsequently discharged.

2. DISCUSSION

Klebsiella pneumoniae is an encapsulated Gramnegative bacillus which belongs to the family of Enterobacteriaciae. It is known to cause diverse community-acquired and nosocomial infections [1] and presents with distinct clinical and epidemiological features throughout the world. It is also known to be associated with gastrointestinal infections [6].

Klebsiella pneumoniae has emerged as a major cause of primary or cryptogenic liver abscess, along with an increasingly recognized condition known as Cryptogenic Invasive Klebsiella pneumoniae Liver Abscess Syndrome (CIKPLA). It is commonly found in persons with Diabetes mellitus - possibly due to their impaired phagocytosis mechanism of encapsulated organisms [7,8]. Other than diabetes, invasive Klebsiella pneumoniae infections have been also found in individuals with impaired host defenses (such as chronic alcoholism, malignancy, hepatobiliary disease. chronic obstructive pulmonary disease, glucocorticoid therapy, and renal failure) [4.9-14], suggesting intrinsic virulence factors enabling the organism to escape host immunity [4]. K1 and K2 capsular strains, coded by the rmpA genes have been attributed to the invasive and virulent potential of Klebsiella pneumoniae [15,16], with K1 strains being significantly associated with pyogenic liver abscess and endogenous endophthalmitis [17,18]. However, to date, its exact pathological mechanisms remain unclear [2]. Although the Klebsiella pneumoniae organism was not serotyped for the presence of magA gene and K1 capsular type in our patient, he demonstrated poor prognostic features suggesting a virulent form of strain.

CIKPLA syndrome is frequently associated with one or more complications such as meningitis, endophthalmitis, lung abscess, or fasciitis and has been commonly reported in Taiwanese population [19], with a few reports emerging in United States (of Asian descent) [20]. This syndrome has never been reported in Middle Eastern or Sudanese population.

A few studies have analyzed poor prognostic features associated with CIKPLA syndrome. These include a Glassgow coma scale < 7 - prior to initiation of treatment [21], thrombocytopenia, raised white cell counts and reduced CSF glucose levels [16,22]. Treatment of choice for CIKPLA syndrome includes third generation Cephalosporin, preferably Ceftriaxone.

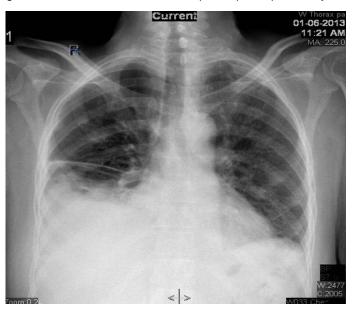


Fig. 1. Chest X-ray showing a heterogeneous opacity involving the mid and lower zone on the right side along with obliteration of the right costophrenic angle and an airbronchogram, suggestive of a parapneumonic effusion



Fig. 2. Ultrasound scan of abdomen showing hepatomegaly (17cm) with multiple echogenic foci likely representing aerobilia probably as a result of gas forming microorganism with abscess, without any obvious fluid

Endogenous endophthalmitis, caused Klebsiella pneumoniae liver abscess is often devastating septic metastatic infection. Considered to be rare previously, it accounts for about 2 - 8% of all endophthalmitis cases [16]. It has mostly been described in association with primary liver abscess in patients from East Asia, particularly Taiwan [8,23-27], and to a much lesser extent in other parts of the world. After diagnosis pyogenic liver abscess, occurrence of ocular symptoms within 48-72 hours suggests the possibility of septic endophthalmitis via hematogenous spread from the liver abscess [24]. Vigorous search for an intraocular infection should be considered in patients presenting with Klebsiella pneumoniae liver abscess.

3. CONCLUSION

We highlight a case of Sudanese male with CIKPLA syndrome demonstrating liver abscess. pneumonia, urosepsis, cellulitis and endophthalmitis with total retinal detachment. The risk factors in our patient included poorly controlled diabetes and chronic heavy alcohol consumption. Despite intensive therapy by an ophthalmologist, visual outcome was poor in our patient. The aim of this case report is to alert clinicians about increasing prevalence of CIKPLA syndrome, its different clinical presentations and occurrence in different populations. We suggest a need for increased vigilance while managing such cases and regularly performing eye examinations at the time of diagnosis so as prevent serious debilitating disease complications.

CONSENT

All authors declare that written informed consent was obtained from the patient (or other approved parties) for publication of this case report and accompanying images.

ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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