Klebsiella pneumoniae Infection:
A Virulent Cause of Visual Loss

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Abstract
Bacterial endophthalmitis is endogenous in 2–6% of cases and is frequently misdiagnosed initially. Klebsiella pneumoniae is being increasingly recognised as an aggressive causative organism, and it is particularly prevalent in Asian populations. We describe the case of a 71-year-old female of Southeast Asian origin with type 2 diabetes mellitus who presented with visual loss secondary to bacterial endophthalmitis and concomitant cerebral abscesses. Imaging revealed the probable primary source of infection to be a liver abscess. She developed retinal detachment and subsequently underwent an evisceration of her right eye. A Klebsiella spp. was identified from the eye tissue by 16S rRNA amplification. Klebsiella pneumoniae endophthalmitis has a characteristic disease phenotype and a particularly aggressive course with poor visual outcomes observed in most cases. This case highlights the risks of metastatic infection including bacterial endophthalmitis in association with Klebsiella infection.

Introduction
Endophthalmitis, or infection of the vitreous or aqueous humour or both, is a medical emergency. The average annual incidence of endogenous bacterial endophthalmitis is estimated to be 5 per 10,000 hospitalised patients [1], and Klebsiella pneumoniae is being increasingly recognised as a causative organism. It is particularly prevalent amongst Asian populations [2–4], and its association with diabetes mellitus has been well reported. K. pneumoniae endophthalmitis typically occurs as a complication of a primary hepatic infec-
tion [5, 6]. It has a characteristic disease phenotype and a particularly aggressive course, with poor visual outcomes observed in 89% of cases [5].

We have found 1 other case report of *Klebsiella* endophthalmitis as an initial presentation of metastatic infection from a liver abscess in the UK [7].

**Case Presentation**

A 71-year-old female, born in Thailand, who had lived in the UK for more than 20 years, presented to an eye casualty department with a 10-day history of intermittent fever and rigors, accompanied by awareness of floaters and pain in the right eye. She had a background of diet-controlled diabetes, and she had been treated for a presumptive urinary tract infection by her general practitioner 4 days prior to presentation with oral co-amoxiclav.

An initial ophthalmic examination of her right eye revealed a severe anterior uveitis and a hypopyon, accompanied by keratic precipitates and fibrin deposition. Punctate corneal epithelial staining and injected conjunctiva were also noted, and the retina could not be visualised. This was accompanied by increasing visual loss.

Her right eye was almost completely insensitive to light, although direct and consensual pupillary reflexes were preserved. Visual acuity and fields in her left eye were normal, and there were no signs of infection or inflammation. The intraocular pressure in the right and left eye was 32 and 10 mm Hg, respectively. B-scan ultrasonography revealed a retinal traction over the posterior pole, but no haemorrhage or choroidal masses were noted. The patient was started on cyclopentolate 1% eyedrops, betamethasone and neomycin (Betnesol) eye ointment, and dexamethasone 0.1% eyedrops. Her clinical examination was unremarkable; however, blood tests revealed raised inflammatory markers with a white blood cell count of 14.7 × 10⁹/l and C-reactive protein of 128 mg/l.

An unenhanced CT of her head on the day of her initial assessment had shown no focal abnormalities, but a contrast-enhanced CT of her orbits 3 days later demonstrated right-sided proptosis with preseptal and intraconal retro-ocular soft tissue stranding and thickening as well as evidence of associated episcleritis (fig. 1). The patient was commenced on topical dexamethasone, ofloxacin and dorzolamide. Following a transient episode of speech arrest, she was admitted to the stroke unit of another hospital and was found to have a temperature of 38 °C. An MRI of her head showed multiple foci of tiny peripherally enhancing lesions with a slightly T2 hypointense wall, surrounding oedema, and a restricted core on diffusion-weighted imaging. Appearances were in keeping with multiple abscesses, most likely pyogenic in nature (fig. 2). Due to the patient’s clinical condition, she was not transferred back to the eye hospital for further management of her endophthalmitis and did not receive an intravitreal injection of antibiotics. No primary source of infection was evident on clinical examination, and the patient was empirically commenced on intravenous ceftriaxone and metronidazole to likely target the bacterial causes of her brain abscesses.

Five sets of blood cultures and a culture of 3 eye swabs failed to grow a pathogen and an HIV antibody screen, and the culture of methicillin-resistant *Staphylococcus aureus* screening swabs were both negative. A transthoracic echocardiogram showed a thickened mitral valve with no vegetation. A liver ultrasound scan revealed a hepatic abscess (not amenable to drainage) which was characterised further with CT. This confirmed a likely biliary source of sepsis secondary to calculous cholecystitis and obstruction of the common bile duct. The patient had no abdominal symptoms at that time and only had mildly deranged liver function tests (bilirubin 36 µmol/l, alanine transaminase 61 IU/l and alkaline phosphatase 140 IU/l). An endoscopic retrograde cholangiopancreatography (ERCP) was attempted but
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abandoned due to technical difficulties. Our patient's blood glucose level on admission was 12.3 mmol/l. She was known to be homozygous for HbE and did not produce HbA0 and HbA1c; therefore, we were unable to assess her prior glycaemic control.

A week later, the patient had no improvement in vision in her right eye despite being on topical antibiotics, steroids and intravenous antibiotics. However, she was generally feeling better and her inflammatory markers were improving. A slit-lamp examination of the right eye revealed a hazy, opacified cornea, 360° iridocorneal touch, and 4+ white blood cells in the anterior chamber. B-scan ultrasonography of the right eye showed a thickened sclera and retinal detachment. The dense cataract in the right eye caused the fundus view to be obscured. A relative afferent pupillary defect was noted in the right eye, and there was no perception to light in the right eye.

A repeat MRI showed a progression in her perilesional oedema as well as a slight growth of the ring-enhancing lesions (fig. 3). CT orbits demonstrated extensive abnormalities around the right globe, retinal detachments and retro-orbital chemosis. The patient was commenced on Pred Forte 1, atropine 1, dorzolamide 2, Betnesol ointment, cyclopentolate 1 and polyvinyl alcohol 1.4% eyedrops. Intravenous ceftriaxone and metronidazole were continued.

Unfortunately, she went on to develop a perforation in her right cornea, which necessitated the evisceration of her right eye in the following week. This procedure was uncomplicated. Her eye tissue was sent for 16S ribotyping, and *Klebsiella* was identified to species level.

A gradual clinical improvement after her intravenous antibiotic therapy was observed in our patient, with a concurrent reduction in inflammatory markers. She was discharged on an outpatient antibiotic therapy, which was continued for a total of 5 weeks of antibiotics. She was also started on metformin and gliclazide for her diabetes. A repeat MRI scan at 5 months displayed a complete resolution of all ring-enhancing lesions. The patient made a good clinical recovery and returned to living independently with monocular vision. Her left eye remained unaffected and healthy when reviewed at the outpatient clinic 6 months following her discharge. She had a repeat ERCP, and cholecystectomy was recommended.

We would like to thank our patient for consenting to this report of her illness and her clinical course.

**Discussion**

The majority of cases of bacterial endophthalmitis are exogenous, occurring after a trauma or surgery. Endogenous endophthalmitis, resulting from haematogenous spread of an infection, is much less common (2–15% of all cases). Examples of typical sources of infection include endocarditis, complicated urinary tract infections, line-related infections and liver abscesses. Endophthalmitis has occasionally been described in association with a transient bacteraemia, e.g. in injection drug users or following endoscopy [8].

Numerous organisms have been implicated with *S. aureus* and *Streptococcus* spp. being among the most common. Only 5 and 3.6% of cases were attributable to *K. pneumoniae* infections in the UK and the USA, respectively [9, 10]. By contrast, in a case series conducted in Taiwan and Singapore over a period of 4–21 years, the prevalence of *K. pneumoniae* infection in endogenous endophthalmitis was found to be between 54 and 61% [2–4].

Certain capsular types of *Klebsiella* (K1 or K2) have been characterised as having a particular propensity to cause metastatic infection including endophthalmitis [11]. In 3 large
case series, 48–53% of cases with *K. pneumoniae* endophthalmitis were reported to have a liver abscess as a primary infection source [2–4]. Indeed, the incidence of endophthalmitis as a complication of a *K. pneumoniae* liver abscess is cited as being 7–11% [5, 6]. Over 50% of the affected patients also had a diagnosis of diabetes mellitus. In vitro, it has been shown that the neutrophils of diabetic patients have an impaired ability to phagocytose *Klebsiella pneumoniae* species with K1 and K2 surface antigens [11]. This impairment of host immunity may cause increased susceptibility of diabetic patients to *K. pneumoniae* endogenous endophthalmitis, and it also suggests that there may be clinical value in capsular phenotyping. The hypermucoviscous phenotype of *K. pneumoniae* and the magA and rmpA genes overlap with the K1 and K2 serotypes, and are also associated with invasive infection and increased inflammation [12, 13].

Visual outcomes following endogenous bacterial endophthalmitis are usually poor. A study of 22 consecutive patients by Yang et al. [5] found that 89% of patients achieved a visual outcome of light perception at best, with 41% requiring enucleation or evisceration. A case report by Ishii et al. [14] suggests that early vitrectomy and lensectomy may improve the outcome; however, the majority of cases are not identified until the disease is disseminated, and mortality from disseminated extra-ocular infection is 5% [9]. Intravitreal antibiotics should be considered following intravitreal sampling in patients with suspected infectious endophthalmitis [15].

A study by Fung et al. [16] reported a 75% rate of faecal carriage of *K. pneumoniae* in healthy adults in Taiwan. This is greater than the 19.4–38% reported in European studies. It is thought to be a predisposing factor for the higher rates of liver abscess in Asia. In our case, the patient had not visited South East Asia in 5 years, but she may have had enteric carriage. It has also been suggested that host genetic factors may play a role in determining susceptibility to *Klebsiella* infection, and studies in *Drosophila* have identified genes such as phg1 and kil1 essential for the intracellular killing of *K. pneumoniae*, which have known homologues in eukaryotes [17].

*Klebsiella pneumoniae* are commonly resistant to several classes of antibiotics and show an increasing prevalence of carbapenemase-producing strains worldwide. This presents a particular concern and makes choosing the right empirical treatment regimens difficult. Fortunately, our patient responded promptly to a third-generation cephalosporin.

**Conclusion**

This case report was unusual in that the primary source was initially occult, and the patient was a long-standing UK resident. It highlights known risk factors (Southeast Asian origin, diabetes mellitus and liver abscesses) for *K. pneumoniae* infections. Any of these or a history of recent travel to Asia should prompt clinicians to consider this pathogen in a patient with endophthalmitis, even in the absence of positive cultures. K1 and K2 serotypes of *Klebsiella* spp. have been associated with more invasive infections. 16S ribotyping was shown to be a useful method to identify the organism to species level in this case.

**References**

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Fig. 1. Fat stranding and soft tissue swelling overlying the right globe anteriorly, which extends intra-conally and causes a mild degree of proptosis.
Fig. 2. a–c Multiple foci of tiny peripherally enhancing lesions demonstrating a slight T2-hypointense wall and surrounding oedema. The appearances are compatible with multiple microabscesses, most likely of a pyogenic origin.

Fig. 3. A repeat MRI 3 days later showed progression in a perilesional oedema and a mild growth of the ring-enhancing lesions despite antibiotic therapy.