Abstract

Renal allograft recipients are on life long immunosuppressive medications. This predisposes them to serious infections such as meningitis. However, Escherichia Coli (EC) causing meningitis is uncommon, and extremely rare in adults. There is a dearth of reported cases in the literature. We report a first case of Escherichia Coli meningitis (ECM) in a renal transplant recipient who had undergone craniofacial surgery followed by radiotherapy for an ossifying fibroma associated with Hyperparathyroidism-Jaw Tumour Syndrome (HPT-JTS). Early recognition and choice of appropriate antibiotic therapy gave a successful outcome. Our case demonstrates the risks related to immunosuppression following renal transplantation, such as increased susceptibility to infections and decreased tumour immunosurveillance, which can contribute to potentially life threatening infections. Our case highlights the importance of clinical vigilance necessary in our patient cohort and discusses the pathophysiology of the virulence factors involved in the invasion of blood brain barrier (BBB) leading to ECM.

Keywords: Renal Transplantation; Immunosuppression; Meningitis; Tumour; Immuno Surveillance; Escherichia Coli.

Introduction

Renal Replacement Therapy (RRT) in the form of kidney transplantation is the treatment of choice for end stage kidney disease (ESKD) with better outcomes in comparison to dialysis. However, transplantation gives an additional lifetime risk due to the burden of immunosuppression (IS). Severe life threatening opportunistic infections are common during the immediate and early post transplantation period due to heavy IS burden and decline during the long-term low/maintenance IS period. Meningitis can be a serious life threatening infection in immuno-suppressed patients. ECM is a commonly recognized infection in the neonates, but is an uncommon cause of bacterial meningitis in adults. A PubMed literature search revealed 2 case series: Copenhagen group [1977 – 97; n=20] and a Taiwanese group [1986 – 2003; n=15]. The reported mortality rates varied between the groups with 40% in Copenhagen series and 100% (untreated) versus 27% (treated) in Taiwan series. We describe a case of a rare cause of meningitis in adult renal transplant patient which has not been described before. Our case report will add to the current wealth of available literature about ECM in adults and increase the awareness of being vigilant in patients on long-term IS.

Presentation

Our patient has HPT-JT characterized by the development of multiple parathyroid adenomas and fibro-osseous tumours of the maxilla and the mandible. He developed ESKD secondary to nephrocalcinosis in May 2010. Following a brief period on haemodialysis, he received a living donor kidney transplant in October 2010. He received an initial IS regimen, which consisted of induction therapy with basiliximab and 7 days of glucocorticoids, followed by maintenance therapy with tacrolimus and mycophenolate mofetil (MMF). A protocol biopsy at 3 months post-transplantation was performed in January 2011. This demonstrated Banff 1A/1B cellular rejection in the allograft, which was treated with 3-pulsed doses of methyl-prednisolone (500mg/day) followed by oral prednisolone (20mg/day). Serum creatinine remained stable between 120-135 μmol/L. MMF was withdrawn by May 2011. He remained on maintenance IS with prednisolone (5mg/day) and a dose of tacrolimus with target trough levels of 5-8 μg/L.

His cranio-facial osseous fibroma became aggressive a year after...
transplantation for which he underwent craniofacial surgery in 2011 and again in June 2012. The surgery involved resection of the frontal bone, frontal cranial fossae, medial walls of the orbits, maxillary antra, nasal septum, turbinates and left nasal bone. He had reconstruction with fat to support his frontal lobes. There were no infective complications after his neurosurgical interventions on either occasion. Clinically, he was well with out any evidence of sepsis. A gadolinium-based magnetic resonance imaging in July 2012 prior to the commencement of radiotherapy did not reveal any inflammation or any infection. He required adjuvant radiotherapy due to the recurrence and aggressiveness of the tumour in August 2012. He became acutely unwell following a week after a session of radiotherapy with vomiting, fever, confusion and decreased responsiveness. He became febrile, hypotensive with tachycardia and tachypnoea. He showed signs of agitation and meningism. His Glasgow coma score (GCS) was 10/15 with bilateral up going plantar reflexes and absent focal neurological signs.

Blood tests revealed a white cell count of 29 x10^9/L (neutrophil count of 26) with a C-reactive protein (CRP) of 8mg/L. Renal transplant function remained at its baseline. Blood tacrolimus concentration before this febrile episode was 6 µg/L. Computed tomography scan of his head with contrast enhancement showed changes consistent with previous craniofacial surgery with a residual tumour (Figure 1).

Intravenous hydrocortisone, ceftriaxone and acyclovir with oxygen therapy were administered immediately. Lumbar puncture demonstrated an opening pressure of 28 cm H2O with turbid cerebrospinal fluid (CSF), which contained 1,400 X 10^6/L red blood cells, 44,700 X 10^6/L white blood cells (100% neutrophils) but revealed a negative gram stain. Following lumbar puncture, he received intravenous meropenem, teicoplanin and acyclovir as empirical therapy till CSF culture results were available. Six hours later, he deteriorated and required airway ventilatory support with adjuvant inotropic (noradrenaline) therapy in the intensive care unit.

CSF cultures grew EC sensitive to ceftriaxone, meropenem and gentamicin. Blood cultures from the day of admission also grew EC following 6 to 7 hrs of incubation with the similar antimicrobial sensitivity profile. Serum CRP had rapidly risen to 235. A course of IV meropenem was completed with vital organ support following which he made a full recovery from ECM.

Outcome and Follow-Up

Our patient remained well after successful treatment of this life threatening infection with no neurological sequelae relating to ECM. Serum creatinine remained stable at 116 µmol/l.

Figure 1: This is a contrast-computed tomography of head showing resection of the frontal bone, the frontal cranial fossae, the medial walls of the orbits, the medial walls of the maxillary antra, nasal septum, turbinates and left nasal bone. Fat is grafted to the frontal cranial fossae to support the frontal lobes of the brain and a metal mesh covers the frontal bone defect. Lateral wall of the left maxillary antrum is expanded with disorganised trabeculation and shows bilateral recurrent disease of both orbits consistent with fibrous dysplasia of HPT-JTS.

Discussion

This case is interesting in view of its complexity and shares the challenges one can face in the clinical management of solid organ transplant individuals. Renal transplantation remains the mainstay therapy for ESKD with improved outcomes and better quality of life. They are at an increased risk of infections due to their immunosuppressed state. Risk of infection is higher in the immediate post-transplant period when immunosuppression burden is higher. Infections, though can also occur later during the long-term low immunosuppression maintenance phase. There is often a greater risk of atypical infections in this group of patients. Meningitis is a serious life threatening infection and transplant recipients are at an increased risk of meningitis secondary to atypical organisms as compared to the general population. ECM is a well-known cause of neonatal meningitis but only a few cases of adult ECM have been reported and there has been no such report in a transplanted individual.

Here, we have described for the first time, a case of ECM in an adult renal transplant recipient, highlighting two important risk factors - increased susceptibility to infections and impaired tumour immunosurveillance. It is possible that these two factors have interacted simultaneously in our case. The growth of our patient's ossifying fibroma appears to have accelerated post-transplantation and this may have caused shrinkage of the residual bony tumour with resultant exposure and breach in the bone/soft tissue interface thereby providing a route for EC to cross BBB and infect CSF. Radiotherapy therefore may have exposed our patient to the risk of infection, to which he had become more susceptible to by his immunosuppressive regime. This resulted in a life threatening ECM with out any co-infection with other organisms.

A PubMed search revealed 2 case series of meningitis that have been published: Copenhagen series had 20 consecutive cases from 1977 to 1997 [1] and a Taiwanese group who reported 15 cases from 1986 to 2003 [2]. These series estimate the percentage of all cases of adult bacterial meningitis caused by EC as 1.5% and
and ampicillin. We chose to avoid aminoglycosides to preserve his renal allograft function in the context of severe sepsis. Mero-
penem was chosen as a broad-spectrum beta-lactam bactericidal antimicrobial agent with a very successful outcome. We believe our report to be the first case of ECM in an adult renal transplant recipient with HPT-JTS. Our case also highlights the importance of increased vigilance for unusual and serious causes of infection in transplant recipients.

**Key Learning Points**

- **Immunosuppression** in renal transplant recipients predisposes to higher risk of unusual disseminated infections.
- **Impaired tumour surveillance** and increased susceptibility to infections are two important accepted risk factors in renal transplant recipients.
- E. Coli meningitis is common in neonates, rare in adults and can be life threatening in immunosuppressed individuals.
- **Early detection and prompt consideration** of appropriate antibiotech treatment offers better outcome.

**References**

17. Gomez JB, Maque Y, Moquilaza MA, Anicama WE (2013) E. coli Meningi-
tis Presenting in a Patient with Disseminated Strongyloides stercoralis. Case Rep Infect Dis 2013:424362