Premature neonates are at risk for Elizabethkingia meningoseptica infections, which include meningitis and sepsis.\(^1,2\) Normally found in soil and water, \(E.\) meningoseptica has also been isolated from hospital surfaces and medical devices, such as ventilators. Thus, \(E.\) meningoseptica has increasingly been recognized as the causative organism of nosocomial infections.\(^3\) \(E.\) meningoseptica is a gram-negative, non-fermentative, oxidase-positive, non-motile, aerobic bacillus that tests positive for catalase and urease.\(^4,5\) The risk factors for \(E.\) meningoseptica infection include indwelling central venous catheters, prematurity, immunosuppression, and prolonged and prior exposure to higher antibiotic concentrations.\(^6\) \(E.\) meningoseptica resists \(\beta\)-lactamases antibiotics in two different ways (intrinsic class A extended-spectrum serine-\(\beta\)-lactamases and inherent class B metallo-\(\beta\)-lactamases), which renders it resistant to a broad range of antimicrobials that are routinely used for empiric therapy of infections caused by gram-negative organisms.\(^7\) Therefore, the selection of appropriate antibiotic treatment is clinically difficult. Herein, we describe our experience with two cases of neonatal \(E.\) meningoseptica infection.

CASE REPORTS

Case one
Our first case was a female infant born as one of two twins at 31 weeks gestation with a birth weight of 1.5 kg. No resuscitation was required. She developed respiratory distress and was connected to continuous positive airway pressure. Her chest X-ray showed signs of respiratory distress syndrome, so she was intubated and surfactants were administered. A few hours later, the baby improved and was extubated and shifted to nasal cannula oxygen. No central lines were required. Feeds were started and tolerated. On day six of life, the baby was less active and lethargic. Initial blood tests showed a white cell count of \(18.5 \times 10^3/\mu L\) (reference range (RR): \(6–20 \times 10^3/\mu L\)) with neutrophils \(12.4 \times 10^3/\mu L\) (RR: \(1.0–8.5 \times 10^3/\mu L\)), hemoglobin of 11.8 g/dL (RR: 10.0–14.1 g/dL), and platelet count of \(411 \times 10^3/\mu L\) (RR 150–450 \(\times 10^3/\mu L\)). C-reactive protein (CRP) was 15.7 mg/dL (RR: 0–5 mg/L) and blood cultures were collected and started empirically on ampicillin and gentamycin. Lumbar puncture (LP) was also performed. She showed initial improvement in terms of improved activity,
no fever, and feeding was tolerated, but later her condition deteriorated as she developed convulsions and apnea, which required intubation. Her blood culture initially reported a non-fermentative gram-negative bacteria sensitive to tazocin. The test used was a disc diffusion method according to the clinical and laboratory standard institute (CLSI), and treatment with tazocin was started. The blood culture was sent to a referral laboratory to identify the organism and E. meningoseptica was detected using Vitek®2 (BioMerieux, France). Cerebrospinal fluid (CSF) analysis revealed hypoglycorrhachia (glucose 1.1 mmol/L (2.8–4.4 mmol/L), White blood cells of 2000/mm³ (normal range 0–5), and elevated proteins (280 mg/dL (RR: 15–45 mg/dL)). Blood culture indicated the same organism sensitive to levofloxacin and rifampicin, so these antibiotics were initiated and the infectious diseases team were involved. Head ultrasound (US) scan was normal. The infant’s general condition improved, and she was extubated on day 15 of life. LP was repeated after three weeks of antibiotics and showed no organisms, but the protein was high in the CSF (27.2 mg/dL) with low sugar (1.57 mg/dL). Therefore, antibiotics (levofloxacin and rifampicin) were continued for another two weeks, at which point LP was repeated and showed improvements. The baby received a total of seven weeks of antibiotics and was discharged in stable condition. At follow-up, she had normal growth and development for her age.

**Case two**

Case two was a male infant born at 36 weeks gestation with a birth weight of 2.5 kg. At eight days old, the baby was admitted with poor feeding, fever, and vomiting. Physical examination was unremarkable, and septic work-up was done (complete blood count (CBC), CRP, and blood and urine cultures). The parents refused LP. CBC showed a white cell count (CBC), CRP, and blood and urine cultures). The infant’s general condition improved, and she was extubated on day 15 of life. LP was repeated after three weeks of antibiotics and showed no organisms, but the protein was high in the CSF (27.2 mg/dL) with low sugar (1.57 mg/dL). Therefore, antibiotics (levofloxacin and rifampicin) were continued for another two weeks, at which point LP was repeated and showed improvements. The baby received a total of seven weeks of antibiotics and was discharged in stable condition. At follow-up, she had normal growth and development for her age.

**DISCUSSION**

Neonatal bacterial meningitis is an uncommon but serious infection with high mortality and the morbidity remains high among survivors. The types and distribution of causative pathogens differ according to birth, gestational age, postnatal age, and geographic region with Streptococcus agalactiae being the most common cause of neonatal sepsis and meningitis since the early 1980s. Premature labor is a risk factor for meningitis because most maternal immunoglobulins cross the placenta after 32 weeks gestation, so infants born extremely preterm are at a significantly higher risk for infections. In Oman, prematurity accounts for 63% of deaths in which sepsis due to gram-negative bacteria was a major cause. E. meningoseptica is rare but is associated with high mortality because of its antibiotic resistance and difficult diagnosis. As a primarily opportunistic pathogen, E. meningoseptica mainly infects newborns and immunocompromised hosts from all age groups. Environmental studies have revealed that the organism can survive in chlorinetreated municipal water supplies, often colonizes sink basins and taps, and has become a potential reservoir for infections in hospital settings. As in our cases, premature newborns weighing < 2500 g are at higher risk of E. meningoseptica infection.

The source of an E. meningoseptica outbreak can be detected by obtaining cultures from food and infant formulas, wet areas, dry surfaces, equipment, and the hands of healthcare workers in contact with infected patients. We isolated the organism from a water tank in our institution, which emphasizes the importance of obtaining cultures on a periodic basis. Changing the prescribing policy for empiric antibiotics and protocols for admissions to the neonatal unit, in addition to thorough disinfection
of the unit have been recommended as measures to eradicate E. meningoseptica outbreaks in pediatric wards. Infection control with milder measures has been described in other studies, including alcoholic hand rubs after washing hands, toileting of babies with sterile water instead of tap water; and repairing, cleaning, super chlorinating, and isolating water tanks from all hospital feeder tanks and changing the sink taps. Continuous training should be implemented to reemphasize hand washing and contact precautions for all hospital staff.15,16

The bacterial isolates in our cases were sensitive to ciprofloxacin, piperacillin/tazobactam, and vancomycin. Recent studies have demonstrated the effectiveness of fluoroquinolones, due to their superior pharmacokinetics compared to hydrophilic antimicrobials, such as beta-lactams.17 Sparfloxacin, clinafloxacin, and levofloxacin have shown better activity against E. meningoseptica than ciprofloxacin. Rifampin has been used as part of combination therapy for the treatment of persistent infection. Vancomycin alone or in combination with other agents like rifampin has been successful in the treatment of meningitis in infants.18 Additional studies are required to verify this. Finally, our infection control measures in the special care baby unit were successful as no further E. meningoseptica infections were reported in the last two years.

CONCLUSION
Nosocomial infections caused by E. meningoseptica is an increasing problem in healthcare settings especially, for immunocompromised patients. This is mainly because of its ability to survive in the environment and its antimicrobial resistance nature. It is crucial to increase the capacity of laboratories for the diagnosis of this bacterium and to implement a multidisciplinary approach for the care of infected neonates. Our experience has demonstrated the need for combined antibiotic strategies for better outcomes.

Disclosure
The authors declared no conflicts of interest. Written consent was obtained from the patient.

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