A Case Report of Neonatal Meningitis and Myocarditis Caused by Enterovirus

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Abstract

Enteroviral infection is very common in newborns. Most newborns recover without serious complications [1]. Invasive neonatal enterovirus infection is rare but often fatal and should be considered in the differential diagnosis with sepsis [2]. We report a case of a neonate who got meningitis and myocarditis caused by enterovirus. A 2-day-old neonate presented with fever. He was diagnosed with enteroviral meningitis. The neonate was hospitalized treated with intravenous immunoglobulin (IVIG) and antibiotics. However, he was diagnosed myocarditis subsequently. Despite those treatment, the newborn ultimately died. We reported a case of myocarditis developing after the diagnosis of enteroviral meningitis. Physicians need to pay attention to clinical conditions of newborns during the treatment of enteroviral infections in newborns.

Keywords: Enterovirus; Meningitis; Neonate; Myocarditis

Introduction

Enteroviruses consist of polioviruses, coxsackieviruses, echoviruses, and new varieties of enteroviruses that have a clearly different genome. Infection with enterovirus in newborns may result in mild to severe and even critical illness [2]. Unspecific febrile illness is the most common presentation of enteroviral infection. More severe infections may cause sepsis-like syndromes, myocarditis, meningoitis, encephalitis, hepatitis, and even death. The mortality in the cases of myocarditis was 44% [3,4]. Newborns are susceptible to get enteroviruses infection via vertical transmission either trans placentally or at the time of delivery. The following case study describes a 38 weeks and 6 days gestational age neonate with meningitis and myocarditis caused by enterovirus.

Case Report

A male neonate was born through meconium contaminated amniotic fluid by cesarean section to a 30-year-old, gravida 2, para 2 mother at 38 weeks and 6 days gestational age and his birth weight was 3480g. The prenatal labs of the mother were all negative, her pregnancy was uncomplicated. The mother had febrile illnesses about 4 days before delivery. After giving birth, the mother had recurrence of fever and was treated with Ceftriaxone; Newborn APGARs were 10 and 10 at 1 and 5 minutes, respectively.

After birth, he was kept in the room for mothers and infants for 1 day. On the 2nd day, the neonate was noted to be drowsy and with a body temperature of 37.8°C. No action has been taken other than encouraging breastfeeding. On the 3rd day, the neonate was grunting, with reduced tone. He was transferred to the NICU and received oxygen. The laboratory tests showed that the leukocyte (WBC) count was 8500/ml (lymphocytes, 25.5%; neutrophils, 65.0%) and the level of C-reactive protein (CRP) was 12.0mg/L. A chest X-ray showed no infiltrates. The TORCH tests were negative. We started with meropenem and vancomycin considering early-onset neonatal sepsis. The fever persisted, and lumbar puncture was performed on the 4th day of hospital. Examination of cerebrospinal fluid showed that the number of WBC was 1840/ml (lymphocytes, 10%; neutrophils, 60%; monocytes 30%), the glucose level was 2.0mmol/L and the study of protein level was 227.1 mg/dl. The PCR of enterovirus was positive but the latex agglutination tests for bacteria were all negative. The neonate was diagnosed with enteroviral meningitis and treated three times with IVIG (100 mg/kg/day). On hospital day 7, the breathing rate was 80 breaths/min and the heart of the newborn was 200 beats/min. EEG showed paroxysmal supraventricular tachycardia and frequent ventricular premature beats. A repeated CSF study showed that the leukocyte count was 180/ml (lymphocytes,
50%; neutrophils, 20%; monocytes 30%), the glucose level was 2.34mmol/L and protein level was 181.7mg/dl. Laboratory tests showed a leucocyte count of 11200/ml (lymphocytes, 37.1% neutrophils, 37.2%) C-reactive protein (CRP) of 18.1mg/L, alanine transaminase level of 560U/L, aspartate transaminase level of 128 U/L, lactate dehydrogenase level of 1102U/L B-type natriuretic peptide (BNP) level of >25000pg/ml and creatine phosphokinase level of 400U/L. Troponin-I and Creatine kinase muscle brain (CK-MB) increased to 6.97ug/l and 275u/L, respectively. His blood gas was normal, but his chest x-ray revealed pneumonia. Echocardiography showed mitral regurgitation, decreased wall motion, small amounts of pericardial effusion with left ventricular ejection fraction of 33.2% and atrial septal defect. Acute respiratory failure and myocarditis was diagnosed. Ventilator care, diuretics, cardiotonic was administered; despite the treatment, symptoms were exacerbated and ultimately fatal. Written informed consent was obtained from the parents of the newborn. This study was approved by the Ethics Committee of Jiaxin Maternity and Child Health Care Hospital.

Discussion

More than 100 serotypes have been identified to date [5]. New types of enteroviruses have a distinctly different genome. Enteroviruses are small RNA viruses belonging to the Picornaviridae family. EVs are known to be the leading cause of febrile and aseptic meningitis in neonates, as well as potentially lethal neonatal infections such as meningencephalitis, hepatitis, and myocarditis. Intrauterine infection is rare, but it can be rapidly fatal [6,7]. 60-70% of newborns diagnosed with enterovirus infection are infected at the time of delivery in the first 10 days of life. The risk factors for critical neonatal disease include the absence of neutralizing antibody, prematurity, maternal illness during the perinatal period, early age, and more virulent viruses infection [8]. The diagnosis of EV infection has been based on positive viral culture, nasopharyngeal swab, stool or cerebrospinal fluid. However, new nucleic acid amplification methods such as PCR for viral RNA identification are more sensitive and have replaced viral culture as standard [9,10]. In our case, enterovirus was detected by RT-PCR in cerebrospinal fluid. Meningitis and myocarditis in neonates can cause significant long-term morbidity [9]. Immunoglobulin has been used in neonates as a therapeutic agent, but clinical efficacy has not been demonstrated. Specific antiviral treatment is not available. Therefore, the treatment is favorable but clinical efficacy has not been demonstrated. Specific antiviral treatment is not available. It is important for physicians to remember that enteroviral infections have a poorer prognosis in neonates than in older children.

Conclusion

Neonatal meningitis and myocarditis caused by enterovirus are rare but often fatal and should be considered in the differential diagnosis in febrile neonates. For clinicians caring for neonates, it is important to consider a general differential diagnosis in neonates with sepsis-like illness. virologic diagnosis can be done quickly by polymerase chain reaction (PCR). Early treatment and prevention measures are the only guarantors of a reduced in morbidity and mortality.

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Conflict of Interest

The authors declare that there is no conflict of interests regarding the publication of this paper.

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