Bacterial meningitis in a known case of steroid-resistant nephrotic syndrome

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ABSTRACT

Nephrotic syndrome (NS) is a kidney disorder in which, excess protein is being excreted through urine. While most cases of idiopathic nephrotic syndrome respond to steroid therapy and experience a limited number of relapses prior to complete remission, some cases suffer from frequent relapses and become steroid-dependent or are primarily steroid-resistant. Treatment options are limited to immunosuppressive drugs with significant side effect profiles. Infections in nephrotic syndrome are common and can be severe. Meningitis is one of the complications associated with nephrotic syndrome. Here, we report a child with steroid-resistant nephrotic syndrome (SRNS) with meningitis. Early detection and appropriate treatment of these complications can be lifesaving.

Key words: Seizures, Streptococcal pneumonia, Cyclophosphamide, Steroid

CASE PRESENTATION

An 8-year-old female child was brought to our casualty with complaints of convulsions aborted by self at home and by medication in the casualty, followed by post-ictal drowsiness and bowel and bladder incontinence. There was no history of aura or fever, neither any history suggestive of cranial nerve involvement. There was no significant family history. Birth history and developmental history were normal. Child was immunized till 5 years of age as per the National Immunization schedule. No optional vaccines were given. On examination, the child was unconscious, Glasgow Coma Scale (GCS) was 8/15 (E2 V2 M4) with tachycardia and hypertension. Generalized edema was present. Patient was maintaining saturation on room air. Weight, height, and BMI of the child were normal. Multiple striae were present on the arm, shoulder, abdomen, and thigh. Moon facies was present. Rest head-to-toe examination was normal. The abdomen was distended with smiling umbilicus. There was no tenderneess or organomegaly. Fluid thrill was present. Upon examination of the Central Nervous System (CNS), the child had altered sensorium, deep tendon reflexes were brisk and signs of meningeal irritation (neck Stiffness, Kernig’s, and Brudzinski’s signs) were present. Rest CNS examination was normal. Investigations (Table 1) were sent which showed increased total leukocyte count (TLC), raised C-reactive protein (CRP), hypokalemia, hypoalbuminemia, and raised albumin: creatinine ratio. Urine routine showed proteinuria. There was no evidence of hematuria. Examination of the cerebrospinal fluid (CSF) revealed 20 cells/mm³, 20% polymorphs, 8.3 mg/dL protein and 108 mg/dL sugar. CSF culture showed growth of Streptococcus pneumoniae.

Initially, the child was on tablet prednisolone at 60 mg/m² and in August 2020, she was started on injection of cyclophosphamide as she was diagnosed as steroid-resistant nephrotic syndrome. Due to non-compliance and improper follow-up, the child developed relapse and parents started oral steroids on their own. There was no significant family history. Birth history and developmental history were normal. Child was immunized till 5 years of age as per the National Immunization schedule. No optional vaccines were given. On examination, the child was unconscious, Glasgow Coma Scale (GCS) was 8/15 (E2 V2 M4) with tachycardia and hypertension. Generalized edema was present. Patient was maintaining saturation on room air. Weight, height, and BMI of the child were normal. Multiple striae were present on the arm, shoulder, abdomen, and thigh. Moon facies was present. Rest head-to-toe examination was normal. The abdomen was distended with smiling umbilicus. There was no tenderneess or organomegaly. Fluid thrill was present. Upon examination of the Central Nervous System (CNS), the child had altered sensorium, deep tendon reflexes were brisk and signs of meningeal irritation (neck Stiffness, Kernig’s, and Brudzinski’s signs) were present. Rest CNS examination was normal. Investigations (Table 1) were sent which showed increased total leukocyte count (TLC), raised C-reactive protein (CRP), hypokalemia, hypoalbuminemia, and raised albumin: creatinine ratio. Urine routine showed proteinuria. There was no evidence of hematuria. Examination of the cerebrospinal fluid (CSF) revealed 20 cells/mm³, 20% polymorphs, 8.3 mg/dL protein and 108 mg/dL sugar. CSF culture showed growth of Streptococcus pneumoniae.

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Magnetic resonance imaging (MRI) was suggestive of post-ictal encephalitis. After establishing the diagnosis, supportive treatment including injection of ceftriaxone, amikacin, fosphenytoin, and levetiracetam, intravenous (IV) fluids, and antihypertensive were started. There were no signs of raised inflammation of connective tissue (ICT). Proteinuria and edema were gradually resolved. Patient was stabilized and injection of cyclophosphamide and oral prednisolone was started at 60 mg/m²/day. The child responded with the given treatment. A final diagnosis of steroid-resistant nephrotic syndrome (SRNS) with meningitis was made. Repeat imaging of the brain showed resolution of brain parenchymal involvement. Patient has been asked to follow-up for further cyclophosphamide therapy.

DISCUSSION

The hallmark of nephrotic syndrome is massive proteinuria, leading to decreased circulating albumin levels which was seen in our patient. Edema occurs due to decrease in plasma oncotic pressure, as a consequence of low serum albumin levels, causing extravasation of fluid into the interstitial space.

Patients with nephrotic syndrome, particularly children, are at increased risk of developing serious bacterial infections, including pneumonia, empyema, and peritonitis [7]. Sepsis, meningitis, and cellulitis are other serious infections that can occur in children with nephrotic syndrome [8]. The increased risk for infection is related to renal losses of IgG. Loss of opsonizing factors may specifically increase the susceptibility to encapsulated bacterial infection, in particular to pneumococcal infections that are potentially lethal [9,10]. The predominant bacterial causes are *Streptococcus pneumoniae* and Gram-negative enteric organisms such as *Escherichia coli* [4]. Patients with NS are frequently in an immunosuppressive state because of urinary loss of immunoglobulins and the use of immunosuppressive drugs. Therefore, they are at a higher risk of suffering from infections followed by high morbidity and mortality. These infections are also the main reasons for hospitalization along with delayed response to steroid or discontinuation of steroid therapy in some cases. Our patient suffered from bacterial meningitis which was secondary to immunosuppressant. Early diagnosis and initiation of treatment are essential to prevent morbidity and mortality.

CONCLUSION

It can be concluded that meningitis and is a rare complication of NS, and an early detection and management of the same can be life saving.

REFERENCES


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