Case Presentations

Sepsis with Streptococcus Pneumoniae in a Child with Nephrotic Syndrome

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Abstract

Introduction. Nephrotic syndrome is one of the most frequent glomerular pathologies encountered in pediatric ages, with an incidence of 4.7 cases in 100,000 children. The most frequent infectious complication of nephrotic syndrome is bacterial peritonitis caused most of the times by Streptococcus pneumoniae, a microorganism responsible by a high number of pneumonia, otitis media and sepsis episodes among children.

Case report. We present the case of a 4.7-year-old male patient, diagnosed with nephrotic syndromes with 2 recurrences and transitory hypertension, admitted in our clinic for: diffuse abdominal pain, vomiting, diarrhea, fever and right otalgia. The laboratory tests on the day of admission pointed out elevated inflammatory biomarkers, leukocytosis, anemia and combined dyslipidemia. The urinary exam showed massive proteinuria. The blood culture was positive for Streptococcus Pneumoniae. The abdominal ultrasound revealed mild ascites, the thoracic radiography ruled out a pneumonia, and the ENT exam pointed out erythematous and bulged tympanic membrane. We initiated empirical therapy with Ceftriaxone, steroids and symptomatic drugs. After 3 weeks of hospitalization, the patient is discharged in good general status, with remission of nephrotic syndrome.

Conclusions. Congestive acute otitis media can lead in case of immunosuppressed patients to a severe sepsis with Streptococcus pneumoniae. Thus, anti-pneumococcal vaccination is mandatory, especially in case of high risk patients.

Keywords: nephrotic syndrome, sepsis, otitis media, Streptococcus pneumoniae

Introduction

Nephrotic syndrome (NS) is a frequently encountered pathology in nephrology clinics in both developed and developing countries, representing one of the most frequent glomerular disorders in pediatric ages (1-3). The incidence of NS worldwide is of 4.7 cases in 100,000 children, and the prevalence is estimated at 16/100,000 children below the age of 16 years (2,4). Classically, it is defined as massive proteinuria (>40 mg/m²/h) responsible of hypoalbuminemia (<2.5 g/dl) and generalized edema (1.5-9). Most of the cases of NS that sensitive to corticosteroids, but nonetheless approximately 20% of the cases are refractory to corticosteroids, frequently due to segmental focal glomerulosclerosis. These cases are characterized by the rapid progression towards chronic renal failure requiring dialysis and renal transplantation within the first years after diagnosis (2,4). In idiopathic NS, the most frequent histological finding is minimal changes glomerulonephritis (80% of cases) or segmental focal glomerulosclerosis. Other histological types of NS encountered in children include membranoproliferative glomerulonephritis and less frequently membranous nephropathy (6,4,3).

Many times, severe complications are associated with NS mainly due to the massive losses of plasma proteins (1). The urinary loss of low molecular proteins such as: immunoglobulin G, factor I and factor B can lead to the alteration of opsonization, phagocytosis and the entire process of bacterial elimination resulting to immunosuppression of NS patients (10).
The complications of NS are divided into two main categories: complications due to the disease itself and complications associated to therapy. The first category includes infections (peritonitis, sepsis, cellulitis, and chicken-pox), embolism (venous embolism and pulmonary embolism), hypovolemia (abdominal pain, tachycardia and hypotension), hyperlipemia, chronic renal failure, anemia etc. (1). It is estimated that approximately 1.5% of the cases of NS in children die as a consequence of infections associated to the underlying disease (11). Of this category, the most frequent complication is spontaneous bacterial peritonitis found in 1.5 to 16% of the cases (10). Most of the cases of peritonitis are caused by encapsulated gram positive bacteria, especially Streptococcus Pneumoniae that represent also a major cause of meningitis, pneumonia, acute otitis media and sepsis among children (10,12,13).

Systemic inflammatory response includes at least two of the following criteria: tachypnea, bradycardia, hyperthermia or hypothermia and leukocytosis or leukopenia (14,15).

Sepsis is defined as life-threatening organ dysfunction caused by a dysregulated host response to infection, being one of the most important causes of death worldwide, with an increasing incidence. According to the Society of Critical Care Medicine and the European Society of Intensive Care Medicine in 2016 was established that organ dysfunction can be represented by an increase in the Sequential Organ Failure Assessment (SOFA) which is associated with an in-hospital mortality greater than 10%. Quick SOFA(qSOFA) is less robust than a SOFA score, it does not require laboratory tests and can be assessed quickly and repeatedly and should be used prompt to further investigate for organ dysfunction and to initiate or escalate therapy as appropriate (16-18).

CASE PRESENTATION

Reasons of admission

We present the case of a 4.7-year-old male child admitted to our clinic for the following complaints: diffuse abdominal pain, vomiting, diarrhea, fever, right otalgia. The personal pathological history revealed that the patient was diagnosed with NS with two episodes of relapse and transient hypertension. We mention that the patient had a poor socio-economic status, is unvaccinated, with inappropriate vitamin supplementation, and had extremely low compliance to treatment.

Clinical exam

At the time of admission, the clinical exam revealed the following pathological elements: altered general status, pale and shiny skin, generalized edema with positive dimple sign at the level of pretibial area, right otalgia, tachycardia 146/min, hypertension 115/78 mmHG, tachypnea 30/min, high fever 39,7°C, distended abdomen, abdominal tenderness at palpation, vomiting, diarrhea, weight 17 kg, height 107 cm.

Diagnostic assessment

The laboratory tests on the day of admission pointed out: anemia (hemoglobin of 10.7 g/dl, hematocrit 31%, mean erythrocyte volume 77.1 μL), leukocytosis (leukocytes 21800 μl), neutrophilia (neutrophils 70.6%), thrombocytes 150,000/mm³; elevated inflammatory biomarkers (CPR 58.8 mg/dl, ESR 87 mm/h); hyponatremia (Na 132.2 mmol/l, K 3.97 mmol/l); hypercholesterolemia (412 mg/dl), while the values of triglycerides were within normal ranges. Also, the liver and renal functions proved to be normal. The urinary exam revealed: leukocyturia, hematuria (50/μl), massive proteinuria (500 mg/dl) and the semi quantitative proteinuria was >4500 mg/l. qSOFA score was 3, respiratory rate 30/min, altered general status with Glasgow coma scale 13, hypertension 115/78 mmHG (stage I according to gender and high). A thoracic radiography was performed which ruled out a possible pneumonia. The blood culture was positive for Streptococcus Pneumoniae, and the antibiogram showed susceptibility to: Penicillin, Amoxicillin, Cefotaxime, Ceftriaxone, Imipenem, Levofloxacine, Moxifloxacine, Ofloxacine, Erythromycin, Linezolid, Vancomycine, Tetracycline, Chloramphenicol, Rifampicin and Trimethoprim/Sulfamethoxazole.

The abdominal ultrasound revealed: hepatomegaly, nephromegaly, a fine blade of perirenal fluid and mild ascites. The surgical consult ruled out a spontaneous bacterial peritonitis. The ENT exam pointed out erythematous, bulged tympanic membrane, establishing the diagnosis of acute conge stive otitis media.
Therapeutic assessment

Thereby, following the clinical, paraclinical examination and qSOFA score a diagnosis of sepsis with Streptococcus pneumoniae as a result of acute congestive otitis media, relapsed nephrotic syndrome and anemia were established. Empirical therapy with Ceftriaxone, corticosteroids (methylprednisolone by vein) and symptomatic drugs was initiated, the evolution being favorable. After 3 weeks of hospitalization, the patient was discharged in good general status, in remission (without proteinuria) with the following recommendations: unsalted, hypoglycemic and hypolipemic diet, tapered oral corticosteroids, and adjuvant therapy with potassium supplements, proton pump inhibitors and Ca + Vitamin D3 supplements.

DISCUSSIONS

Among the pediatric population, the most common cause of NS is idiopathic, being diagnosed in 90% of children with aged between 1 to 10 years and in 50% of children diagnosed after above the age of 10 years (7). The incidence of this disease is reported between 2-7/100,000 children bellow 15 years of age, but the prevalence is higher due to the recurrent nature of this pathology (7). The incidence varies according to gender, age, ethnicity and environment, being more common among male patients, with a sex ratio of 1:3 for boys, and it may occur at any age but most frequently with onset between 2 and 7 years (8,18). Similar to the data reported by the literature, the case presented above is a male child whose diagnosis of NS was established at approximately 3 years of age.

Infections are the main complications that threaten the lives of NS patients, due in particular to Streptococcus pneumoniae, the first cause of bacterial peritonitis and sepsis in these patients (9).

However, other microorganisms such as β-hemolytic streptococcus, Haemophilus and Gram negative bacteria are also commonly found (1). Thus, there were reported cases of infectious associations in patients with NS, such as the simultaneous presence of peritonitis, cellulitis and sepsis (10). Moreover, there were described also cases of sepsis with Vibrio cholerae, bacteria that causes mainly diarrhea, but due to severe immunosuppres-
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