Early Infantile Galactosialidosis Presenting with an Unusual Renal Involvement

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Abstract

Galactosialidosis is a rare lysosomal storage disease associated with deficiencies of beta-galactosidase and neuraminidase. In this report, we present a 9-month-old early infantile Galactosialidosis infant with renal involvement. In the literature only isolated cases of Galactosialidosis with IgA nephropathy, renal insufficiency and renal transplantation reported. To the best of our knowledge, the patient is the first case reported in the literature in which steroid resistant nephrotic syndrome has been found in a Galactosialidosis patient.

Keywords: Galactosialidosis; Steroid resistant nephrotic syndrome; Early infantile form

Introduction

Galactosialidosis (GS; OMIM 256540) is a rare lysosomal storage disorder caused by deficiency of the protective protein/cathepsin A (PPCA). It is transmitted as an autosomal recessive trait. Protective protein/cathepsin A forms a complex with beta-galactosidase and alpha-neuraminidase and in turn controls the lysosomal compartmentalization, catalytic activation, and stability of the two glycosidases. A mutation in the CTSA gene that encoding PPCA localized on chromosome 20q13.1 were found to be responsible [1].

Galactosialidosis patients present a combination of clinical and biochemical findings of that are typical of those found both in GM1-gangliosidosidosis and sialidosis types 1/II. Three galactosialidosis subtypes are recognized based on age of onset and the severity of clinical manifestations [2]. The early infantile form (type 1) mainly characterized by hydropsfetalis, cherry red spots, hepatosplenomegaly, psychomotor delay, coarse facies, skeletal dysplasia, and early death. Late infantile form (type 2) is characterized by corneal clouding, cardiac involvement, visceromegaly, growth retardation and, rarely, psychomotor retardation. Most patients with the milder juvenile/adult form (type 3) mainly exhibited myoclonus, ataxia, neurological deterioration, mental retardation, angiokeratoma, absence of visceromegaly. Late infantile and juvenile/adult galactosialidosis are slowly progressive diseases that are less severe than the early infantile type. Juvenile/adult form of galactosialidosis is typically associated with a normal life expectancy [3,4].

Nephrotic syndrome is a clinical condition characterized by massive proteinuria, hypoalbuninemia, hypercholesterolemia, and generalized edema. It occurs more commonly in children than in adults, and usually manifests as one of two usually idiopathic diseases: Minimal change nephrotic syndrome or focal segmental glomerulosclerosis. Other histological patterns of nephrotic syndrome seen in children include membranoproliferative glomerulonephritis, and rarely membranous nephropathy and diffuse mesangial hypercellularity. Children having steroid resistant nephrotic syndrome with focal and segmental glomerular sclerosis run a high risk of resistance to immunosuppressive therapy [5,6].

In galactosialidosis patients only a few cases with renal involvement determined. Here, we report a galactosialidosis patient with steroid resistant nephrotic syndrome. The aim of the present report is to both describe the characteristic features of this rare disease, and draw attention to galactosialidosis association with steroid resistant nephrotic syndrome.

Case Report

A 9-month-old male admitted to hospital with abdominal distention and limb edema. He was born at 37 weeks of gestation as a second child of consanguineous parents. Fetal ultrasound and family history was normal. At age of 2 month he had inguinal hernia operation. On physical examination, birth length was 46 cm (<3th percentile), birth weight was 2650 g (10th percentile), and head circumference was 33 cm (3-10th percentile). Blood pressure was 110/60 mmHg and heart rate was 90 beats/min. He didn’t hold his head and eye-tracking had just begun. Muscular hypotonia, coarse face, broad forehead, hypertelorism, bilateral epicanthus, depressed nasal bridge, broad nasal tip, low-set ears, gingival hypertrophy, pectus carinatus, shorth neck, abdomen distention, hepatosplenomegaly, ascites, scrotal edema, thoracolumbar gibbus, and psychomotor retardation detected. Facial dysmorphism at age of 10 month seen in the Figure 1.  

The skeletal survey showed the ovoid spine, beaked vertebra and lumbar scoliosis (Figure 2). The long bones and the pelvis appeared normal. Echocardiography detected mitral insufficiency. Ophthalmological evaluation was normal. Abdominal ultrasound showed bilateral grade 2 echogenicity, enlarged kidneys and ascites.

Nephrological evaluation showed hypoalbuminemia (1.9 gr/dl), massive proteinuria (urine protein/creatinine ratio of 3.5 mg/ml) and hypertriglyceridemia (521 mg/dL) was detected; serum creatinine was 0.7 mg/dl and serum uric acid was 5.4 mg/dl. At age of 3 month he had anuria for 1 month. Nine month old he had proteinuria (urine protein/creatinine ratio of 3.5 mg/mg) and hypertriglyceridemia (521 mg/dL) was detected. The laboratory tests, including hemoglobin, electrolytes, blood glucose, liver function test, and coagulation test were normal.

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stopped and ACE inhibitors were started. The presented patient was diagnosed as steroid-resistant nephrotic syndrome due to massive proteinuria without clinical response after four weeks of prednisone treatment. Cyclosporine could not be started as he had severe infection. Renal biopsy was performed, a sclerotic and nonproliferative glomeruli was seen.

The patient was later admitted to our hospital because of pneumonia at 11 months of age. His psychoomotor development was severe retarded especially concerning motor skills. He developed progressive hepatosplenomegaly, feeding difficulties, and hypoproteinemia. He died because of sepsis and heart failure shortly after then.

**Discussion**

Early infantile galactosialidosis is the most severe form of this disease and is associated with premature mortality. It is not easy to diagnose in the infant period because of early death and necessary of urinary sialyloligosaccharide excretion, and both enzyme activities of beta-galactosialidase and neuraminidase. However the age and presentation as a steroid-resistant nephrotic syndrome corresponds with the most common presentation of the infantile form of sialidosis (nephrosialidosis), including the absence of cherry-red spots or corneal clouding in early infancy, the present case is compatible to early infantile galactosialidosis with both enzyme activities. In the present case, any ophthalmologic findings of galactosialidosis were detected. Although several cases of galactosialidosis reported with no eye abnormalities, galactosialidosis has been usually associated with various ocular anomalies, such as optic nevre atrophy, macular cherry-red spots, cloudy corneae, ocular albinism, and bilateral cataracts [7-9].

Protective protein cathepsin A, is a multifunctional enzyme, with deamidase, esterase and carboxypeptidase activities. In the lysosome, PPCA is essential for the stabilization of a multienzyme complex with lysosomal beta-galactosidase, sialidase/neuraminidase, and galactosaminogluco-6-sulfatase and for activation of neuraminidase. Deficiency of PPCA leads to secondary deficiency of beta-galactosidase and neuraminidase, and causes galactosialidosis. The distended lysosomes primarily contain accumulated sialylated glycoconjugate substrates of neuraminidase. Therefore, galactosialidosis is highly similar to sialidosis which caused by deficiency of neuraminidase [10]. Early and progressive nephrotic syndrome leading to end stage renal failure in the first years of life is one of the components of nephrosialidosis and related to the renal storage of undegraded sialyloligosaccharides [11,12]. Patients with GM1-gangliosidosis also can present extensive renal histological lesions, but impairment of renal function is not typically described. Kidney findings of galactosialidosis in autopsy material of early infantile and young/adult galactosialidosis have been reported [13-15]. These findings include swollen glomerular epithelial cells, tubular epithelial cells, glomerular endothelium, and renal blood vessels with many vacuoles, which are either empty or contain sparse tubular profiles. Proximal tubular epithelial cells contain PAS-positive storage material. Firstly, Sewell et al reported a 4 months old a infantile galactosialidosis patient with large kidneys [16]. Koike et al. described a 31 years old young/adult form of galactosialidosis Japanese patient with IgA nephropathy and normal renal function. They showed the electron microscopy findings of storage material in the kidney which have not been described till now [17]. Shortly after than Kiss et al reported a 9 years old Brazilian girl with galactosialidosis with severe nephrological symptomatology, which led to renal failure and renal transplantation. They hypothesized that end stage renal failure is not a typical finding in galactosialidosis.

Furosemide and albumin infusions in addition to prednisone therapy (2 mg/kg/d) were given. After four-week treatment with prednisone, as massive proteinuria was continued, prednisone was stopped and ACE inhibitors were started. The presented patient was uric acid nitrogen, creatinine, thyroid function tests, plasma and urine aminoacids, urine reducing substances, ammoniac, and lactate levels, were all found to be normal. The peripheral blood smear showed intracytoplasmic vacuolation of lymphocytes. Increased urinary oligosaccharides and sialic acid excretion was detected. We analyzed the activity of beta-galactosidase both in peripheral blood leucocytes and cultured skin fibroblasts. The low beta-galactosidase activities of 20.8 umol/g/h (normal values: 100-400 umol/g/h) and 53.5 umol/g/h (normal controls: 476 and 503 umol/g/h) were detected both in leucocytes and fibroblasts, respectively. The low neuraminidase activity 0.28 umol/g/h (affected control: 0.49 umol/g/h; normal controls: 25 and 34 umol/g/h) was detected in skin fibroblasts.

Figure 1: Frontal view (A) and lateral view. (B) of the patient’s appearance showing coarse face, broad forehead, hypertelorism, bilateral epicanthus, depressed nasal bridge, broad nasal tip, low-set ears, short neck, abdomen distention, and thoracoolumbar gibbus.

Figure 2: (A) Anteroposterior chest image showing lumbar scoliosis. (B) Lateral radiography of the spine showing ovoid vertebra, and beaked vertebra.
and end stage renal failure is secondary to the sialidosis component of the patient [18]. In the present patient, prednisone therapy were given. After four-week treatment with prednisone, as massive proteinuria was continued, prednisone was stopped and ACE inhibitors were started. The patient was diagnosed as steroid-resistant nephrotic syndrome due to massive proteinuria without clinical response after four weeks of prednisone treatment. Cyclosporine could not be started as the patient had severe infection. Renal biopsy was performed, a sclerotic and nonproliferative glomeruli was seen. We speculated that the accumulation of oligosaccharides in the in the organs and the urine may underlie the glomerular visceral epithelial cell dysfunction and proteinuria.

A rare reason of the nonimmun hydrops fetalis is lysosomal storage disorders [19]. In prenatal diagnosis of the lysosomal storage disorders the vacuolated lymphocytes present in the peripheral fetal blood smear may be the first clue to a lysosomal storage disease, but prenatal diagnosis of galactosialidosis is reliably possible by the investigation of the secondary deficiencies of beta-galactosidase and neuraminidase in amniocytes [9]. In the present patient, prenatal diagnosis could not be given to family because of normal fetal ultrasound and no history of recurrent hydropsfetalis.

In conclusion, there were a few cases with galactosialidosis, and only isolated cases of galactosialidosis with IgA nephropathy, renal insufficiency and renal transplantation reported in the literature. Presented case is particularly interesting because of its presentation with an unusual renal involvement. To our knowledge this is the first report of a galactosialidosis patient with steroid resistant nephrotic syndrome. When coarse facies, hepatosplenomegaly, dysostosis multiplex, and early neurological deterioration are detected in an infant, we should be considering galactosialidosis in the differential diagnosis. Also in order to not omit a possible diagnosis of galactosialidosis urinary sialoligo saccharide excretion and both enzyme activities of beta-galactosidase and neuraminidase is necessary. Early diagnosis of this disease will allow for more effective treatment, genetic counseling and prenatal diagnosis in galactosialidosis.

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