



An Infant with Liver, Kidney, Skin and Musculoskeletal Abnormalities

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Clinical Image

A 21 day-old boy infant was admitted to pediatric intensive care unit with hypotonia, jaundice and abdominal distention. He was born at 37th week of gestation from first cousin Saudi parents via cesarian section with a birth weight of 2200 g. Physical examination revealed the weight, the length was and the head circumference were <3th percentile. Dysmorphic features included flattened nasal bridge, high arched palate, lax skin and micrognathia. There were generalized hypotonia, hepatomegaly, abdominal distention and multiple contractures such as radial deviation of the wrist joint (Figure 1). Laboratory investigations revealed hemoglobin was 16.6 g/dl, leucocytes: 14.9/mm³, platelets: 83,000/mm³, serum total bilirubin level: 13.5 mg/dl, conjugated bilirubin level: 4.3 mg/dl, aspartate aminotransferase (AST): 555 U/L, alkaline phosphates (ALP): 960 U/L. Amino acid analysis of blood and urine, tandem mass were all within normal limits. He was diagnosed as renal tubulopathy based on polyuria with proteinuria: 167 mg/dL, elevated fractional excretion of Na (FENA: 4.89%) and fractional excretion of K (FEK: 66%). Abdominal ultrasound revealed hepatomegaly and bilateral mildly increased renal echogenity. Whole exome sequencing (WES) was performed and showed mutations in the VPS33B gene. Based on all signs and laboratory findings, the baby was diagnosed as arthrogryposis, renal dysfunction, and cholestasis (ARC) syndrome. Arthrogryposis, renal dysfunction, and cholestasis (ARC) syndrome is an autosomal recessive disorder caused by mutations in the VPS33B gene [1]. This syndrome mainly effects liver, kidney, and skin central nervous and musculoskeletal systems. The first component of ARC syndrome is arthrogryposis multiplex congenital [2]. The most common anomalies of musculoskeletal system in ARC syndrome are muscle atrophy, radial deviation of the wrist and flexion contracture of the knee. Osteopenia and pathological fractures can be observed in this syndrome owing to reduced reabsorption of phosphate ions via renal tubules and secondary to hyperparathyroidism [2]. The second component ARC syndrome is cholestatic jaundice and hepatomegaly [3]. These are the most common symptoms in ARC syndrome. Normal GGT levels, mildly elevated AST and ALT levels without biliary obstruction have been described in all patients who have ARC syndrome [3]. Liver histology in the patients with ARC syndrome suggests paucity of bile ducts, lipofuscin deposition and giant cell hepatitis [4]. The third component of ARC syndrome is renal tubular dysfunction which is characterized by multiple features of renal Franconia syndrome including glucosuria, phosphaturia, generalized aminoaciduria and renal tubular acidosis [5]. Renal tubular dysfunction

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Figure 1: Appearance of ARC syndrome. Lax skin, radial deviation and contracture of the wrist joint.

may present in the first few days of life or later around the age of two months. Most of the patients also present symptoms of nephrogenic diabetes insipidus and Fanconi syndrome [5]. Additional clinical symptoms of ARC syndrome principally include ichthyosis, abnormal platelet count and function, secondary infection, and cardiovascular anomalies [6]. The prognosis of the syndrome is very poor and most patients die by the age of 7 months because of recurrent infections, severe dehydration, metabolic acidosis or internal hemorrhaging [7]. No specific treatment for ARC syndrome currently exists; rather, supportive care including fluid infusion, anti-infection, supplement with ursodeoxycholic acid, fat-soluble vitamins, calcium gluconate, L-thyroxine and phosphate is administered to patients for improving the quality of life. Nevertheless, some patients with joint contractures, congenital hip dislocation, and vertical talus are in need of immediate orthopedic intervention due to delayed diagnosis. Unfortunately as curative treatment for ARC syndrome is unavailable, all of the patients die within several months. Therefore extensive research of family history classical clinical presentations and genetic mutational analysis should be performed for diagnosing and to initiate the therapy at early stage.

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