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Natural disease history and characterization of SUMF1 molecular defects in multiple sulfatase deficiency: a case report

Short title: multiple sulfatase deficiency: a case report

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Abstract

Multiple sulfatase deficiency (MSD) is a very rare lysosomal storage disease(LSD) caused by mutations in the SUMF1 gene. So far, about 143 patients with MSD have been reported in previous studies, although this figure is likely an underestimation due to under-reporting and under-recognition. The present report shows the genetic and clinical aspects of a patient with MSD in comparison to the previously reported patients.

Key words: Lysosomal storage disease - Multiple sulfatase deficiency - SUMF1 gene

Introduction

Considering the number of cases reported with Multiple Sulfatase Deficiency (MSD), caused by mutations in the SUMF1 gene (1-3), it is a very rare Lysosomal Storage Disease (LSD). Previous studies have reported about 143 patients with MSD, which seems to be an underestimation due to under-reporting and under-recognition. This report attempts to clarify the genetic and clinical aspects of a patient with MSD, as compared to the previously reported ones. Prior to the study, an informed consent was obtained from his parents.

Case report

The boy was born in a healthy and Iranian family with first-cousin parents after an uneventful pregnancy at 37 weeks of gestation by cesarean delivery with a seemingly normal Apgar score. He was immediately hospitalized in the neonatal intensive care unit (NICU) and was there for 11 days. Due to apnea coupled with hypotonia and cyanosis, treatment with phenobarbital was administered for him. However, his birth parameters were normal (HC, 33 cm; BL, 49.5 cm; BW, 2450 kg). When he was eight months old, his head suddenly became larger; the diagnosis of communicating hydrocephalus was made for him, and he underwent medication therapy. As a 20-month-old boy, he represented psychomotor delay and developed a tonic seizure. Shortly afterward, he underwent a rapidly progressive loss of muscle tone and psychomotor skills. Social contact eventually deteriorated (no visual contact, no speech, and no parent identification). In the following months, he was admitted repeatedly to our hospital due to recurrent pneumonia, gastroenteritis, and clonic-type seizures. Taking a family history, we found no neurologic problem in the family.

At the last clinical examination, he was 3.5 years old when he presented to our children's hospital emergency department with clonic seizure coupled with apnea. He was bedridden and suffered from global developmental delay (GDD), meaning that he was not able to speak, make eye contact, identify his parents, and sit or walk independently. He weighed 8.5 kg (<1st percentile), his height was 85cm (<1st percentile), and had a head circumflex of 52cm (>97th percentile). The clinical examination revealed hypotonia of the lower and upper limbs, abnormal palmar crease, macrocephalus (head circumflex:52, >95th percentile), hypertrichosis, absent deep tendon reflex, and hepatomegaly (projection of the liver edge 5cm below the costal margin). Also, skeletal deformities (pectus carinatum, kyphosis, and scoliosis) were noted. Morphologically, he presented with hirsutism, broad thumbs, severe muscular atrophy, and delayed growth. His facial features had become slightly coarse and dysmorphic (thick and high arched eyebrows, orbital hypertelorism, long eyelashes, prominent and smooth philtrum, anteverted nostrils, depressed nasal bridge, and posteriorly rotated ears). Furthermore, his neck was considerably short (**figure 1A, B, C, D**). Diffuse ichthyosis of the lower limbs extensor surface and the dorsum of foot was noted. Also, he showed dry, rough, and scaly skin resulting from cradle cap (seborrheic dermatitis). In addition, there was brachydactyly (**figure 2A, B**). The dermatologist confirmed common (vulgar) ichthyosis. The anterior and posterior fontanelles were still open (3cm×2cm and 0.5cm×0.5cm, respectively). The boy had a left undescended testicle and a normal left testicle. The ophthalmologic investigation was normal. **figure 3** shows brain magnetic resonance imaging (MRI) of our patient. Visual examination was normal. The EMG-NCV reported sensory-motor demyelinating

polyneuropathy of both upper and lower limbs and EEG left diffuse slowing pattern (encephalopathic pattern) (**figure 4**). The patient underwent an auditory brainstem response (ABR) test and showed severe hearing loss in both ears. CSF was not sampled. An echocardiogram revealed normal results. Abdominal ultrasound revealed hepatomegaly. X-rays of the thoracolumbar spine, wrist, pelvis and lower extremities were obtained and showed in **figures 5A, B, C, and D**.

Taken together, all of these signs were compatible with disabling and progressive neurological disorder that was ultimately diagnosed as MSD by a genetic test. Instead of enzyme activity testing, the individual's DNA was extracted from a blood sample for whole Exome sequencing(WES). Ultimately, the gene SUMF1 showed a homozygote mutation: c.739G > C, p. G247R in exon 6. The karyotype was normal, 46, XX. The genetic test and clinical findings were compatible with MSD.

Discussion

MSD is an ultra-rare autosomal recessive inborn impairment of metabolism, influencing post-translational activation of human sulfatases by FGE. Due to missense and nonsense mutations in the SUMF1 gene, the catalytic activity of FGE deteriorates, leading to reduced human sulfatase activities (4). Approximately 50 SUMF1 mutations are discovered. The majority of them are missense types that influence residual molecular activity and stability of mutant FGE, that both relate to MSD severity (5, 6). The estimated disorder prevalence is about 1 in 500,000 (7). A genetic test of our patient revealed a mutation in the allele of c.739G > C, p. G247R. This is the most prevalent mutation in MSD and is considered a severe missense variant with severe reduction of stability and activity of human sulfatases (6). The diagnosis is not easy due to the ultra-

rarity of the disorder. In our patient, the signs that led the team to suspect the MSD were the coarse face, the cutaneous features, and particularly the radiological findings consistent with dysostosis multiplex. In previous studies, the most frequently reported sign/symptoms were ichthyosis (71%), organomegaly (57%), dysostosis multiplex (56%), and dysmorphic feature(53%), respectively (6). Our patient had typical manifestations of neonatal MSD, including all these signs. Indeed, the development of these signs is a very helpful indicator in identifying the disorder. Although skeletal (17%) and nutritional and gastrointestinal (14%) problems were the most common signs at birth,(6) in our case, apnea, hypotonia, and cyanosis were detected at birth. It was similar to the case previously described (8). Another Iranian neonate with MSD was also reported who presented the typical clinical symptoms of the disorder (9).

Conclusion

MSD is still an incurable disorder. So, genetic counseling and potential treatment strategies entirely depend on complete functional and molecular analysis of SUMF1 mutations on the biochemical and clinical phenotype.

Conflict of Interest

Authors declared no conflict of interest.

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