Case Report

The mitochondrial trifunctional protein (MTP) is a multienzyme complex of the fatty acid beta-oxidation cycle. It is composed of four alpha and four beta subunits as HADHA and HADHB. The three functions are 2-enoyl coenzyme A (CoA) hydratase (LCEH), long-chain 3-hydroxy acyl-coenzyme A dehydrogenase (LCHAD), and 3-ketoacyl CoA thiolase (LCTH). The HADHA gene encodes the LCEH and LCHAD enzymes and the HADHB gene encodes the LCTH enzyme. The cytogenetic location of both genes is at 2p23.3. Mitochondrial trifunctional protein deficiency (MTPD) which is a rare condition that leads to failure of converting certain fats to energy is characterized by decreased activity of all three enzymes.

Signs and symptoms of MTPD may begin during infancy or later in life; those that begin after infancy include hypotonia, muscle pain, rhabdomyolysis, and peripheral neuropathy. We report a Turkish boy diagnosed with MTPD after being investigated for polyneuropathy of unknown origin since infancy.

Case. A 5.5-year-old male patient was admitted to our clinic with complaints of weakness in the arms and legs, physical inactivity compared to his peers, fatigue, weakness and, difficulty in climbing stairs since infancy. Electroneuromyography (ENMG) analysis showed moderate symmetric distal sensorimotor and axonal neuropathy. On the background of chronic polyneuropathy, the patient had acute relapsing episodes with progressively worsening severity in the follow-up period until 12.5 years of age. Whole exome sequencing (WES) was performed in the patient and, revealed that the patient had a homozygous c.1390G>A (p.Gly464Ser) pathogenic variant of the HADHB gene. Although rhabdomyolysis is a well defined accompanying clinical feature of MTPD, it was not present in our patient who only had worsening muscle weakness during attacks.

Conclusion. On the background of chronic polyneuropathy and acute relapsing episodes triggered by fasting or illnesses and rhabdomyolysis physicians should suspect disorders of the fatty acid beta-oxidation cycle.

Key words: polyneuropathy, mitochondrial trifunctional protein deficiency, fatty acid oxidation disorder, HADHB gene.
of MTPD that may begin after infancy include hypotonia, muscle pain, rhabdomyolysis, and peripheral neuropathy. Symptoms can be triggered by periods of fasting or by illnesses such as viral infections.

Patients presenting with the prolonged, progressive course of the disease, which has been associated with myopathy, recurrent rhabdomyolysis, and sensorimotor axonal neuropathy tend to survive into adolescence and adulthood. Here, we report a Turkish patient diagnosed with MTPD who had been under investigation for polyneuropathy of unknown origin since his infancy.

Case Report

A 5.5-year-old male patient was admitted to our clinic with complaints of weakness in the arms and legs, fatigue, physical inactivity compared to peers and, difficulty in climbing stairs since infancy. His prenatal follow-up was normal and he was born at term with a birth weight of 3160 g with an uneventful delivery. He was the third child of consanguineous Turkish parents, and his siblings were healthy. He had walked when he was one year old and started using words at 1.5 years old. However, a waddling gate was first noticed at three years of age.

On admission to our center, growth was found to be normal for age. He had mild kyphoscoliosis. Neurological examination revealed absence of deep tendon reflexes and mild waddling gait. Muscle strength was 4/5 in the proximal and distal assessment of all four limbs. Routine investigations including whole blood count, biochemical parameters, Creatine Kinase (CK) level and vitamin B₁₂ level were within normal ranges. An inherited polyneuropathy was suggested in the patient. Metabolic investigations including serum and urine amino acids, serum ammonia level, lactic acid, and pyruvic acid analysis, tandem mass spectrometry, urine organic acid analysis, and very-long-chain fatty acid analysis were normal. Cranial magnetic resonance imaging (MRI) was normal and cardiology and ophthalmology referrals also found no remarkable findings.

In an electromyography (EMG) analysis performed before admission to our clinic, a sensorimotor neuropathy had been suspected. EMG analysis was repeated in our center and it showed moderate symmetric distal sensorimotor and axonal neuropathy prominent in the lower extremity. Muscle biopsy showed prominent fiber type grouping which was consistent with secondary neurogenic changes. The patient was investigated for Charcot-Marie-Tooth disease (CMT), however, the molecular genetic analysis performed for peripheral myelin protein 22 (PMP22) gene mutation was found non-consistent with the diagnosis. Therefore the possibility of mitochondrial disease was suspected in the patient and multivitamin supplements were initiated, which yielded partial benefit.

At 6.5 years of age, he had complaints of walking deterioration and gait disturbance concurrent with an infection. On neurologic examination ataxia was noted in addition to the absence of deep tendon reflexes. Muscle enzymes including CK and lactate dehydrogenase, liver enzymes, renal functions, and electrolytes were checked for rhabdomyolysis and they were within normal ranges. He was investigated for causes of ataxia. Serum alpha-fetoprotein, vitamin A and vitamin E levels were normal. Hearing assessment, ophthalmologic evaluation, echocardiography, spinal MRI, and molecular genetic analysis performed to evaluate Friedreich ataxia were also normal. In the follow-up period, the gait disturbance complaint regressed spontaneously. Gait and walking disturbance attacks repeated during infectious illnesses at 9 and 11 years of age. In these two attacks, the severity of symptoms had progressed and he could not even stand and walk, even though muscle enzymes continued to be normal (CK levels were 123 and 140 U/L, respectively). Repeated cranial and spinal MRI studies were unremarkable. ENMG performed at 12 years of age showed chronic severe sensorimotor and axonal neuropathy. Although
gate disturbance regressed after the course of each attack, his walking became relatively worse at 12.5 years of age and the family declined to use any vitamin supplements.

Whole exome sequencing (WES) was performed in the patient and, revealed that the patient had a homozygous c.1390G>A (p.Gly464Ser) pathogenic variant of the HADHB gene compatible with MTPD. Parents were detected as heterozygous for the variant. After definitive diagnosis, all routine biochemical and metabolic investigations including serum ammonia level, lactic acid, and pyruvic acid analysis, tandem mass spectrometry, urine organic acid analysis, and ophthalmologic evaluation were performed once again and found normal. Enzyme analysis of the three enzymes forming the MTP could not be performed in the patient due to the fact that these tests can only be performed by very few laboratories related to technical difficulties and financial problems. A low-fat diet with restriction of long-chain fatty acid intake and substitution with medium-chain fatty acids (MCT oil) was commenced. At the last visit, he was 13.5 years old, and he had difficulty standing on his heels. However, his walking and balance were much better under diet and MCT oil treatment.

Informed consent was received from the family.

Evaluation of the Variant:

The c.1390G>A mutation, which was detected as homozygous in the HADHB gene in our patient, is a rare variant that has not been detected in healthy population scans previously performed. In the analysis of this variant with in-silico prediction programs, it was evaluated as a disease-causing variant. This variant, which has not been reported in the literature and online databases, is classified as a “variant of unknown significance” according to the American College of Medical Genetics and Genomics variant classification guidelines. However, since our patient’s clinical findings and clinical progression were compatible with MTPD, this variant was thought to be responsible for the patient’s clinic.

The variant corresponds to the first nucleotide of the exon 16 in the genomic sequence. At the translation level, this variant could create a new possible splice site. The analyses according to “Human Splicing Finder”, the c.1390G>A variant at the beginning of the coding region is estimated to create a new acceptor site (New site: +58.9). If a new acceptor site has occurred, the structure of the protein may have changed.

The evaluation at the level of amino acid exchange; glycine at the position 464, is the first amino acid of the beta-strand of the protein. The variant is replaced by the C terminal of the Thiolase domain (Fig. 1a). The newly formed Serin amino acid has an extra-OH group that may cause the formation of a new hydrogen bond (Fig. 1b, c). A new hydrogen bond may decrease the stability of the protein by changing the interatomic interactions, atomic fluctuation, and local flexibility (Fig. 1d,e,f).

Polyphen-2 score: 1

PROVEAN: -5.403 (cutoff:-2.5)-deleterious

Discussion

Inborn errors of metabolism (IEMs) can present at any age from childhood to adulthood and may affect the peripheral nervous system, usually as part of a diffuse neurological or systemic clinical picture. Inherited metabolic neuropathies comprise a clinically, biochemically, and genetically heterogeneous group of diseases. Rarely, neuropathy can be the unique initial sign of an IEM. Rare causes of metabolic neuropathy include fatty acid beta-oxidation disorders, mitochondrial disorders, porphyria, disorders of the lipid or glycolipid metabolism (eg. Refsum disease, Fabry disease, abetalipoproteinemia, Tangier disease). MTPD is a fatty acid beta-oxidation cycle disorder, and is characterized by a wide clinical spectrum ranging from severe neonatal manifestations including cardiomyopathy, hypoglycemia, metabolic acidosis, skeletal myopathy and neuropathy, liver disease, and death to a mild phenotype with peripheral polyneuropathy.
episodic rhabdomyolysis and pigmentary retinopathy.\textsuperscript{10}

The most common motor symptom of peripheral neuropathy is weakness. It may present as clumsiness, difficulty in climbing stairs, or impaired fine motor skills. These motor symptoms were present in our patient since infancy. Ataxia is another motor symptom associated with neuropathy. Ataxia was a prominent symptom in our patient only during the course of deteriorating attacks that developed in association with an infection. Sensory symptoms, of which none were seen in our patient may include numbness, paresthesia, pain, or burning sensations.

Fu et al.\textsuperscript{11} reported an 8-year-old girl with lower limb weakness since birth who was ultimately diagnosed with MTPD. Blood acylcarnitine analysis revealed slightly increased long-chain 3-OH-acylcarnitine levels; EMG suggested peripheral nerve injury; muscle biopsy confirmed a neurogenic lesion in muscle fibers, as shown by EMG. Analysis of the HADHB gene identified a homozygous missense mutation c.739C > T (p.R247C). They stated that neonatal-onset, as seen in their patient, has not been reported for the neuromyopathic phenotype of mitochondrial trifunctional protein deficiency.\textsuperscript{9} Clinical findings of our patient were similar to their patient, but repeated acylcarnitine analysis including in deterioration attacks were normal in our patient.

Naiki et al.\textsuperscript{12} reported peripheral polyneuropathy, rhabdomyolysis, and infantile-onset hypoparathyroidism in two siblings with HADHB gene mutations. Recently, van Vliet et al.\textsuperscript{13} reported a 20-year-old woman who had axonal motor-sensory polyneuropathy of unknown origin since childhood. She presented with progressive dyspnea, and increased muscle weakness which had been preceded by an infectious disease. Laboratory testing showed rhabdomyolysis and hypocalcemia with low parathyroid levels. Sequence analysis of the HADHB gene showed two heterozygous variants. The authors emphasized the importance of performing metabolic screening when patients are most symptomatic, since normal results could be inconclusive in the absence of metabolic stress.\textsuperscript{11} Serum calcium,
phosphorus, and magnesium levels were normal in our patient, and rhabdomyolysis was not seen, however, he experienced deterioration attacks concurrent with infectious illnesses.

Yamamoto et al.\textsuperscript{14} reported a 45-year-old man presenting with slowly progressive muscle weakness and sensory disturbances in his lower limbs and multiple episodes of exercise-induced severe muscle fatigue and brown urine during his childhood, which had disappeared by age 20. A nerve conduction study showed peripheral axonal neuropathy. Similar to our patient, they considered CMT disease, however, exome sequencing failed to identify a mutation in the known genes associated with CMT. Then, he recurrently developed severe rhabdomyolysis that required hospitalization. They re-examined exome sequencing and found a mutation in \textit{HADHB}. This case of adult-diagnosed MTP deficiency was characterized by slowly progressive peripheral neuropathy masquerading CMT in addition to muscular symptoms.\textsuperscript{14}

No pathogenic variant has been identified in exon 16 of the \textit{HADHB} gene so far, which may be pathologic for the disease. However, it has been reported that mutations detected in the last exons of the gene may be presented with late-onset and milder clinical findings.\textsuperscript{15}

The lack of enzyme level measurement is a limitation of this report. However using online modeling and prediction programs, we have been able to predict that the mutation identified herein can destabilize the structure of the protein. Although acylcarnitine levels were repeatedly normal in our patient, clinical findings may be the result of conformational changes in the protein by the Gly464Ser homozygous variant that may lead to a less functional protein. Nevertheless, functional studies are required to clearly understand the effect of the variants that have not any clinical significance yet.

In conclusion, it is evident that the differential diagnosis of isolated polyneuropathy in childhood may be challenging. Most peripheral nerve disorders in children are hereditary. Characterization of a peripheral nerve disorder involves a careful neurologic examination, family history, electrodiagnostic studies. Although rhabdomyolysis is a well-defined accompanying clinical feature of MTPD, muscle weakness that is getting worse in attacks in the absence of rhabdomyolysis was an interesting aspect of our patient. On the background of chronic polyneuropathy, acute relapsing episodes triggered by fasting or illnesses and rhabdomyolysis should suggest the fatty acid beta-oxidation cycle.

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\textbf{Author contribution}

OU and SK collected the data; OU wrote the manuscript; BC performed genetic analysis.

\textbf{Conflict of interest}

Authors declare no conflict of interest.

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