Identification of Two Novel BCKDHB Mutations in Korean Siblings with Maple Syrup Urine Disease Showing Mild Clinical Presentation

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Case Report
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Maple syrup urine disease (MSUD; OMIM#248600) is a rare autosomal recessive aminoacidopathy caused by a defect in the branched-chain a-keto acid dehydrogenase complex (BCKDH), a multi-subunit enzyme found in mitochondria that is involved in the metabolism of the branched chain amino acids (BCAAs) leucine, isoleucine, and valine. Impaired activity of the BCKDH complex causes accumulation of these three amino acids and their keto acid derivatives, leading to metabolic encephalopathy and progressive neurodegeneration in untreated patients [1]. It has an estimated prevalence of 1 in 185,000 worldwide [1, 2] and 1 in 230,000 in Korea [3].

Disease-causing mutations have been identified in the BCKDHA, BCKDHB, or DBT genes, which encode different subunits of the BCKDH complex. Although encephalopathy and progressive neurodegeneration are its major manifestations, the severity of the disease may range from the severe classic type to milder intermediate variants. We report two Korean siblings with the milder intermediate MSUD who were diagnosed with MSUD by a combination of newborn screening tests using tandem mass spectrometry and family genetic screening for MSUD. At diagnosis, the patients’ plasma levels were elevated for leucine, isoleucine, valine, and alloisoleucine, and branched-chain a-keto acids and branched-chain a-hydroxy acids were detected in their urine. BCKDHA, BCKDHB, and DBT analysis was performed, and two novel mutations were identified in BCKDHB. Our patients were thought to have the milder intermediate variant of MSUD, rather than the classic form. Although MSUD is a typical metabolic disease with poor prognosis, better outcomes can be expected if early diagnosis and prompt management are provided, particularly for milder forms of the disease.

Key words: Maple syrup urine disease, BCKDHB, Novel mutation, Korean.
creasing disease severity in this order. Among these, the classic form represents 75–80% of all MSUD patients [1,4].

Early diagnosis and BCAA-restricted dietary treatment promote normal intellectual development and prevent neurological complications [5,6]. Therefore, in 2006, MSUD was added to the nationwide newborn screening (NBS) program for all newborns in Korea [3]. However, MSUD is quite rare, and less than 10 MSUD patients have been reported to date in Korea. Moreover, only 3 of these patients have received molecular confirmation of the diagnosis, and all of them had the classic form of the disease [7].

In this study, we present two siblings with MSUD arising from two novel BCKDHB mutations who exhibited mild clinical courses and were thought to have a variant form of MSUD, which would be the first description of this disease type in Korea.

Case

1. Patient 1 (younger brother)

A 2-month-old Korean boy visited the Seoul National University Children's Hospital (Seoul, Korea) because of recurrent abnormal results of NBS for inherited metabolic diseases. He was born as a dizygotic twin with a birth weight of 2.80 kg at 37 weeks of gestation and was the third child born to healthy and nonconsanguineous parents. At the age of 3 days, an NBS performed using liquid chromatography/tandem mass spectrometry (LC-MS/MS) yielded an abnormal result. Therefore, a second-tier test was performed on day 20 after birth, and the result also revealed abnormally high levels of leucine (550.5 μmol/L) and valine (382.8 μmol/L). An amino acid analysis of the serum performed at the age of 34 days showed elevated concentrations of isoleucine (299.9 μmol/L), leucine (749.9 μmol/L), and valine (605.1 μmol/L). Alloisoleucine was also detected at the level of 58.2 μmol/L. His urine organic acid analysis revealed elevated branched chain α-keto acid and branched chain α-hydroxy acid levels. MSUD was diagnosed biochemically, and a low-BCAA formula and thiamine supplementation were started on day 34 after birth.

At the age of 2 months, the initial examination in our hospital showed that the patient's height and weight were 59.6 cm (+0.75 standard deviation [SD]) and 5.1 kg (–1.0 SD). His head circumference was 39.4 cm (+0.73 SD). He did not show any pathological signs or symptoms for encephalopathy, and he could establish eye contact with his mother and exhibited social smiling. The follow-up serum amino acid analysis showed decreased levels of BCAAs (isoleucine, 188 μmol/L; leucine, 371 μmol/L; and valine, 256 μmol/L) and alloisoleucine (23 μmol/L) (Table 1). The plasma alloisoleucine-to-isoleucine ratio was 0.19. The patient had normal levels of aspartate aminotransferase, alanine aminotransferase, glucose, and electrolytes. Ketone was not detected in the serum or urine. Brain and abdominal ultrasonography showed no abnormalities.

The patient is currently 9 months of age with a BCAA-restricted diet (isoleucine 40 mg/kg, leucine 60 mg/kg, and valine 45 mg/kg, total protein 2.5–3.0 g/kg, and total calorie 120–130 kcal/kg per day) and thiamine supplementation, and shows a normal growth rate with regard to height (74.0 cm, +0.66 SD) and weight (9.0 kg, +0.0 SD), as well as normal developmental milestones, without any signs or symptoms of neurological deterioration. During the follow-up period, he had no acute episodes of metabolic derangement, and the levels of BCAAs had been normalized. Considering the clinical manifestations of the patient, he was thought to have the mild intermediate variant of MSUD, rather than the severe classic subtype.

2. Patient 2 (elder brother)

The elder brother of patient 1 was brought to the outpatient clinic of our institution at the age of 4.5 years for family screening of MSUD. Regarding his history, he was born with a birth weight of 3.65 kg at 40 weeks of gestation. An NBS that was performed using LC-MS/MS yielded normal results at that time. Although he showed

Table 1. Results of plasma amino acid analyses at diagnosis in two maple syrup urine disease siblings

<table>
<thead>
<tr>
<th>Amino acids</th>
<th>Patient 1 (µM)</th>
<th>Patient 2 (µM)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alanine</td>
<td>301 (143-439)</td>
<td>150 (120-600)</td>
</tr>
<tr>
<td>Arginine</td>
<td>45 (12-133)</td>
<td>45 (12-112)</td>
</tr>
<tr>
<td>Citrulline</td>
<td>34 (3-35)</td>
<td>21 (8-47)</td>
</tr>
<tr>
<td>Glutamic acid</td>
<td>85 (10-133)</td>
<td>43 (14-78)</td>
</tr>
<tr>
<td>Glutamine</td>
<td>229 (246-1,162)</td>
<td>118 (333-809)</td>
</tr>
<tr>
<td>Glycine</td>
<td>183 (81-436)</td>
<td>135 (107-343)</td>
</tr>
<tr>
<td>Alloisoleucine</td>
<td>58.2 (ND)</td>
<td>59.0 (ND)</td>
</tr>
<tr>
<td>Isoleucine</td>
<td>299 (31-86)</td>
<td>159 (6-122)</td>
</tr>
<tr>
<td>Leucine</td>
<td>749 (47-155)</td>
<td>458 (30-246)</td>
</tr>
<tr>
<td>Lysine</td>
<td>159 (66-270)</td>
<td>49 (66-270)</td>
</tr>
<tr>
<td>Ornithine</td>
<td>151 (22-103)</td>
<td>34 (20-136)</td>
</tr>
<tr>
<td>Phenylalanine</td>
<td>64 (22-108)</td>
<td>38 (26-98)</td>
</tr>
<tr>
<td>Proline</td>
<td>167 (52-298)</td>
<td>150 (40-332)</td>
</tr>
<tr>
<td>Serine</td>
<td>77 (71-186)</td>
<td>101 (70-194)</td>
</tr>
<tr>
<td>Threonine</td>
<td>168 (24-174)</td>
<td>55 (40-204)</td>
</tr>
<tr>
<td>Tyrosine</td>
<td>64 (22-108)</td>
<td>33 (19-119)</td>
</tr>
<tr>
<td>Valine</td>
<td>605 (64-294)</td>
<td>482 (132-480)</td>
</tr>
</tbody>
</table>

Values are presented as data (reference range). Bold letters indicate abnormality.

Patient 1, younger brother; patient 2, elder brother. ND, not detected.
a mild delay in language development, his growth velocity with regard to height and weight and his motor development were normal. When he was 3.5 years of age, he was admitted to a regional hospital for 3 days because of severe dehydration and acute deterioration of his mental status associated with viral acute gastroenteritis. The levels of ketone in his urine were elevated, and intravenous rehydration with dextrose solution was required to recover his general and mental condition, although specific metabolic studies were not performed at that hospital.

At the age of 4.5 years, his height and weight were 104.5 cm (+0.16 SD) and 16.0 kg (−0.42 SD), respectively. His head circumference was 51.0 cm (+0.20 SD), and there were no abnormal findings upon physical examination. Serum amino acid analysis revealed elevated levels of BCAAs (isoleucine, 159 μmol/L; leucine, 458 μmol/L; and valine, 482 μmol/L) and alloisoleucine (59 μmol/L) (Table 1). His plasma alloisoleucine-to-isoleucine ratio was 0.37. Branched chain α-keto acids and branched chain α-hydroxy acids were also detected in the urine. The levels of ketones and electrolytes in the serum were normal. MSUD was confirmed, and a BCAA-restricted diet (isoleucine 400 mg/day, leucine 600 mg/day, and valine 450 mg/day; total protein 35 g/day, total calorie 2,000 kcal/day) and thiamine supplementation were started. He was also thought to have the mild intermediate variant of MSUD.

3. Molecular genetic analysis

The parents of these patients provided formal informed consent to perform peripheral blood sampling for genetic analyses. All coding exons and exon–intron boundaries of the BCKDHA, BCKDHB, and DBT genes were screened. No mutations were identified in the BCKDHA and DBT genes. The analysis of the BCKDHB gene revealed the presence of two novel mutations in the siblings. They were heterozygous for two missense mutations: c.508C>T (p.R170C) in exon 5 and c.673C>G (p.L225V) in exon 6 (Fig. 1). The mother and the father harbored the c.508C>T (p.R170C) and c.673C>G (p.L225V) mutations, respectively. Moreover, the remaining two siblings were carriers of one of the mutations identified here. All three in silico prediction algorithms predicted that the novel variants might affect protein function and might be pathological mutations (Table 2) [8-10].

Discussion

Traditionally, MSUD patients can be classified into different clinical phenotypes. The classic form, which is most severe with less than 2% of BCKDH activity, manifests itself within the neonatal period as poor feeding, seizures, and coma, and represents 75–80%
of all patients. The remaining 20–25% of patients have milder variant forms, with later or episodic onset of symptoms and signs and BCKDH activity ranging from 2% to 40%. There is a rare form that exhibits thiamine responsiveness where pharmacological doses of thiamine lead to normalization of the BCAA levels [1,11].

MSUD is diagnosed based on the presence of typical clinical features and elevated levels of BCAA and alloisoleucine in the plasma and of branched-chain hydroxy acid and keto acid in the urine. In particular, a level of more than 5 μmol/L alloisoleucine in the plasma is a distinctive metabolic feature that is present in all forms of MSUD, whereas the plasma levels of BCAAs may be variable and are incomplete for the prognosis [12]. BCKDH enzyme activity, which can be measured using a variety of cell types, including skin fibroblasts, lymphocytes, or liver cells, is also important for diagnosis. However, previous studies have reported that enzyme activity measurements have variable accuracy and may not be clinically useful [13]. The plasma alloisoleucine-to-leucine ratio can be used as an indirect indicator of disease severity because invasive procedures, such as skin biopsy, are not required for this test, and this ratio is inversely correlated with the BCKDH enzyme activity measured in fibroblasts [14]. The siblings described here showed ratios of 0.19 and 0.37, respectively. These values are lower than those observed in patients with severe classic forms (0.59-0.72) and imply that our patients have milder variants [14].

Molecular genetic testing for the BCKDHA, BCKDHB, and DBT genes has been used preferentially for diagnostic confirmation of MSUD. Differences in enzymatic activity and biochemical derangements among mutations in these 3 genes have not been found, and no strict genotype–phenotype correlations have been defined for MSUD [15,16].

In this study, we identified two novel missense mutation variants in BCKDHB, p.R170C and p.L225V. This gene codes for the E1β subunit that has transketolase activity. Although there is no known mutational hot spot, arginine 170 and leucine 225 are amino acids with a unique pattern in the pyrimidine-binding domain that are highly conserved across species. We were not able to perform a functional analysis to measure the residual enzyme activity in vitro. However, the novel variants identified in this study were thought to be pathological mutations that caused the loss of enzymatic activity because these variants were not found in 50 healthy Korean controls, and all three in silico algorithms used in this study predicted that the novel variants might affect protein structure and function (Table 1).

Since the benefit of NBS for MSUD was demonstrated for the first time in 1991 [17], early initiation of treatment after an early diagnosis via NBS has evidently resulted in fewer and less severe clinical symptoms and has yielded better neurodevelopmental outcomes [18]. The nationwide NBS program, including screening for MSUD, has been applied for all newborns in Korea since 2006. However, milder variant forms of MSUD, including the intermediate or intermittent type, can be missed during NBS. There are some reports of patients with variant forms that were not detected by NBS [19,20], and our patient 2 is an example of such a missed case. Second-tier testing, including alloisoleucine and complementary application of the ratios of leucine to other amino acids, has been attempted and may improve NBS sensitivity for MSUD [5,15].

In summary, we report the identification of the first Korean family with the variant form of MSUD resulting from two novel BCKDHB mutations. Although MSUD is a rare disease, and most of the patients with the classic form can be detected by NBS currently, milder variants of MSUD might escape NBS detection as was the case in patient 2. Thus, a high index of suspicion and repeated biochemical tests are necessary for the early detection and prevention of irreversible neurological complications, particularly in children showing delayed neurodevelopment with a history of unexplained ketonuria. Considering that MSUD is transmitted as an autosomal recessive condition and that interfamilial variability of the clinical manifestations is suggested, as observed in the siblings reported here, family screening using molecular genetic analysis should be performed in the patients’ siblings, even if they show a normal result upon NBS.

Acknowledgements

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