Identification of a Novel Arylsulfatase B Gene Mutation in Three Unrelated Iranian Mucopolysaccharidosis Type-VI Patients with Different Phenotype Severity

Nayerossadat Nouri¹, Nargesossadat Nouri², Omid Aryani³, Behnam Kamalidehghan⁴ and Massoud Houshmand*⁵

¹Molecular Genetic Laboratory, Alzahra Hospital, Isfahan University of Medical Sciences, Isfahan, Iran; ²General Tohid Genetic Counseling Center, Isfahan, Iran; ³Dept. of Genetic, Special Medical Center, Tehran, Iran; ⁴Dept. of Pharmacy, Faculty of Medicine, University of Malaya, Kuala Lumpur 50603, Malaysia; ⁵Dept. of Genetic, National Institute for Genetic Engineering and Biotechnology (NIGEB), Tehran, Iran

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ABSTRACT

Background: Mucopolysaccharidosis type-VI (MPS-VI), which is inherited as an autosomal recessive trait, results from the deficiency of N-acetylgalactosamine 4-sulfatase (arylsulfatase B) activity and the lysosomal accumulation of dermatan sulfate. In this study, ARSB mutation analysis was performed on three unrelated patients who were originally from the West Azerbaijan province of Iran. Methods: After PCR and direct DNA sequencing, DNA extraction was performed. Results: Sequencing analysis revealed a novel homozygous missense mutation in the ARSB gene at c.1457A>G [p. D486V] in three unrelated Iranian MPS-VI patients with different phenotype severity. Conclusion: The mutation type in three patients was the same; probably, because of a foundation effect on their population. Iran. Biomed. J. 16 (3): 169-171, 2012

Keywords: Mucopolysaccharidosis type-VI (MPS-VI), Mutation, Iran

INTRODUCTION

Maroteaux-Lamy syndrome (mucopolysaccharidosis type-VI [MPS-VI], MIM# 253200), is a rare, autosomal recessive lysosomal storage disorder caused by a deficiency of arylsulfatase B (ARSB) enzyme that is involved in the degradation of glycosaminoglycans (GAG) dermatan and chondroitin sulfate. The ARSB gene, located on chromosome 5q13-q14, contains 8 exons [1], which produce the arylsulfatase B polypeptide with 533 amino acids [2]. Therefore, ARSB gene mutations lead to lysosomal storage and urinary excretion of these partially degraded substrates. Patients with MPS-VI usually have normal intellectual development, but show many physical symptoms found in Hurler syndrome. A wide variation in the clinical severity is observed: the infantile (severe), juvenile (intermediate) and adult (mild) forms [3]. Although over 130 ARSB mutations have been reported in different countries and different ethnic populations, no ARSB mutation analysis has been reported in the Iranian population. Experimentally, it seems that Maroteaux-Lamy syndrome is the most common type of mucopolysaccharidosis in Iran. In this study, three unrelated Iranian patients with MPS-VI and severe phenotypes were described and homozygous missense mutation in their ARSB gene was also characterized.

MATERIALS AND METHODS

Clinical features of the patients. The three patients were born in first-cousin consanguineous marriages and were originally from the West Azerbaijan province of Iran, but they were completely unrelated. The clinical features of the three patients have been listed in Table 1. The patients were informed of the aim of the study and gave their informed consent to the genetic analysis. The three patients were diagnosed as most likely having the severe form (early onset) of MPS-VI based on the medical interview, physical exam,
Fig. 1. Comparison of missense mutation at exon 8 of ARSB gene at c.1457A>T [p.D486V] in patients with normal sequence of cDNA and protein.
This is the first report of a novel missense mutation in the ARSB gene in three unrelated Iranian patients. Based on the patients' clinical symptoms and their GAG and ASB activities, MPS-IV diagnosis was probable and was therefore necessary to be confirmed by molecular analysis.

Our patients had a missense mutation at exon 8 of the ARSB gene at c.1457A>G [p.D486V]. However, another study indicated that the missense mutation near this location, which was previously reported by Karageorgos et al. [6] at c.1450A>G [p.R484G], causes severe type of MPS-IV. Our novel missense mutation was found to be very near to the mutation in Karageorgos's study and had the same amino acid change. Although all three patients had the same mutation and all were in the same age group, the severity of the disease was different among them. Patient 1 had a more severe form of MPS-VI in comparison with the other two patients, and patient 3 was only presented with growth retardation and ear, nose and throat disorders. However, none of them had central nervous system disorders and mental retardations.

Experimentally, it would seem that Maroteaux-Lamy syndrome is the most common type of mucopolysaccharidosis in Iran; however, there have been no documented reports yet. Our data show that the mutation type in all three unrelated patients was the same. This may be because they were originally from the same region and existence of a foundation effect in their population. In conclusion, more cases from this Iranian province and other locations of the country are needed to identify the most common mutations in the various ethnic groups of the Iranian population.

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REFERENCES