Molecular findings of Colombian patients with type VI mucopolysaccharidosis (Maroteaux–Lamy syndrome)

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ABSTRACT

Introduction: Maroteaux–Lamy syndrome, or mucopolysaccharidosis (MPS) type VI, is an autosomal recessive lysosomal storage disease caused by a deficiency of the enzyme arylsulfatase B (ARSB), required to degrade dermatan sulfate. The onset and progression of the disease vary, producing a spectrum of clinical presentation. So far, 133 mutations have been reported. The aim of this study is to determine the mutations in the ARSB gene that are responsible for this disease in Colombian patients.

Results: Fourteen patients with clinical manifestations and biochemical diagnosis of MPS VI were studied, including two siblings. The 8 exons of the gene were directly sequenced from patients’ DNA, and 14 mutations were found. 57% of these mutations had not been previously reported (p.H111P, p.C121R, p.G446S, p.*534W, p.S334I, p.H147P, c.900T–G, and c.1531_1553del) and 43% had been previously reported (p.G144R, p.W322*, p.G302R, p.C447F, p.L128del, and c.1143-1G–C). Of the previously reported mutations, 80% have been associated with severe phenotypes and 20% with intermediate-severe phenotypes. Bioinformatic predictions indicate that the new mutations reported in this paper are also highly deleterious.

Conclusions: Most of the Colombian patients in this study had private mutations.

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1. Introduction

Mucopolysaccharidoses are a group of congenital metabolic disorders caused by the deficiency of a specific lysosomal enzyme that affects normal catabolism of glycosaminoglycans (GAGs). Accumulation of GAGs in different organs and tissues leads to the complex signs and symptoms of these multisystemic diseases (Giugliani et al., 2010; Neufeld and Muenzer, 2001). Mucopolysaccharidosis type VI (MPS VI; MIM no. 253200) or Maroteaux–Lamy syndrome is a rare genetic disease with recessive autosomal inheritance caused by deficiency of the N-acetylgalactosamine-4-sulfatase enzyme, also known as arylsulfatase B (ARSB). This enzyme is required to degrade dermatan sulfate and chondroitin sulfate [review in 3]. The disease is characterized by progressive, systemic clinical manifestations that cause significant functional impairment. The rapidly progressive form is typically characterized by a slowing of the growth rate, skeletal and joint deformities, coarse facies, and obstruction and recurrent infections of the upper airway. Later on, patients require wheelchair support or are bed-ridden due to bone deformities, cardiopulmonary disease, blindness, or compression of the spinal canal. Patients with the rapidly progressive disease die in adolescence or by age 20, while individuals with slowly progressive forms have a life expectancy of approximately 40–50 years. Although cognitive impairment is not usually described, physical and functional limitations affect psychomotor development and learning (Giugliani et al., 2007). A global incidence of MPS VI between 1 in 248,000 and 1 in 300,000 live births is estimated (El Dib and Pastores, 2009), but population data indicate that incidence may be higher in Brazil. In a high-risk screening of a Brazilian population with MPS diagnoses, 16% were identified as MPS VI (Giugliani et al., 2007). In the city of Monte Santo, State of Bahia, northeastern Brazil, actual prevalence is estimated to be much higher than the prevalence reported in the literature; a founder effect of the p.H178L mutation has been described in a cluster of patients in this area (Costa-Motta et al., 2011). In 2012, 27 MPS VI cases were identified in Colombia, of which 10 occurred in native groups (Rosselli et al., 2012). Additionally, a series of 20 ceramic artifacts from the Tumaco-La Tolita culture, more than 200 years old, were found. It is possible that these artifacts represent individuals with Maroteaux–Lamy syndrome. The artifacts depict phenotypic characteristics, such as skeletal dysplasia, macrocephaly, coarse features, wide mouths, prominent chests, kyphosis, and scoliosis. These ceramics are likely evidence of this disease occurring in pre-Colombian populations in Colombia (Pachajoa and Rodriguez, 2014). Patients with MPS VI...
have widely variable multisystemic symptoms, with typically chronic and progressive courses, that mainly affect the cardiorespiratory and skeletal systems, comeas, skin, liver, spleen, meninges, and brain (Azevedo et al., 2004). Although the systemic involvement is very similar to the clinical profile of MPS I, intelligence quotient is not affected by MPS VI because there is no accumulation of heparan sulfate, which is predominantly responsible for neurological damage (Neufeld and Muenzer, 2001).

The ARSB gene, located on chromosome 5 (5q13-q14), is made up of 8 exons and synthesizes a 2228-bp mRNA that encodes a precursor protein of 533 amino acids (Valayannopoulos et al., 2010; Litjens et al., 1989). As of October 2015, 165 mutations were reported by The Human Gene Mutation Database (HGMD Professional 2015.3) (http://www.hgmd.cf.ac.uk/) including missense mutations (102), nonsense mutations (9), splice site mutations (18), small deletions (3), small insertions (1), insertion–deletion (indel) mutations, and large deletions (2). In South America, various studies have been conducted on patients with MPS VI, including the first molecular study of South American patients (12 Brazilian and 1 Chilean) by Petry et al. in 2005 that identified 7 new mutations (Petry et al., 2005). Other studies involving South American patients include Karageorgos et al. study in 2007 (Karageorgos et al., 2007) and Garrido et al. study in 2007 (4 Argentinian patients) (Garrido et al., 2007). As mentioned earlier, some studies have been conducted on South American population, but to date, no molecular study has been conducted on Colombian patients.

A correlation between excrated urinary GAGs and phenotype was found by Swiedler et al. in 2005 (Giugliani et al., 2010; Swiedler et al., 2005), but no direct correlation has been established between genotype and phenotype thus far (Giugliani et al., 2010; Litjens et al., 1996). Identification of the genotype may be important for predicting phenotype and, in some MPS cases, making treatment decisions; it is also useful for providing family genetic counseling on reproductive risks, prenatal diagnosis, and prevention of genetic diseases (Giugliani et al., 2010).

The aim of this study is to identify the molecular alterations responsible for Maroteaux–Lamy syndrome in Colombian patients.

2. Material and methods

2.1. Patients

Fourteen patients with MPS VI or Maroteaux–Lamy syndrome, biochemically confirmed by enzyme activity analysis, entered the study after signing an informed consent. All patients came from different areas of Colombia; 50% (7 patients) from Bogotá and 50% (7 patients) from other regions of the country (Cartagena, Ipiales, Funza, and Medellin). In regards to gender distribution, 50% were women and 50% were men. Of the 14 patients, 2 were siblings (patients 4 and 5) and the rest of the patients were not related. This study complied with the provisions established by resolution No. 008430 of 1993 of Colombia Ministry of Health, and it was conducted under the Ethical Principles for Medical Research Involving Humans established by the Declaration of Helsinki. Finally, it was approved by the Ethics Committee of Universidad El Bosque and Pontificia Universidad Javeriana.

2.2. Genomic DNA extraction, polymerase chain reaction (PCR), and sequencing

Genomic DNA was obtained from peripheral blood in EDTA tubes using the salting-out method of extraction (Miller et al., 1988). Each of the 8 exons, including their adjacent intronic regions (approximately 30 bp 5′ and 3′ of each exon), was amplified by PCR. Primers were based on those reported by Garrido et al. (Garrido et al., 2007) and were verified by the Primer 3 program (http://primer3.sourceforge.net). The primer sequences and annealing temperatures are shown in Table 1. The PCR consisted of 40 ng of template DNA, 1.25 U of Taq DNA polymerase (Bioline Ltd., London, UK), 1× buffer, 1 mM MgCl₂, 0.08 mM dNTPs, 0.2 mM primers, and 4% DMSO in a final volume of 25 μl. The reactions underwent initial denaturation for 5 min at 95 °C and 35 cycles of denaturation at 95 °C for 30 seconds, annealing at 59–62 °C (depending of the primer) for 30 seconds, elongation at 72 °C for 30 seconds, and a final extension step at 72 °C for 5 min. The PCR products were purified using the EXOSAP-IT® enzymatic column method, and sequencing was performed using an ABI PRISM 3730XL Analyzer® (96 capillary type).

2.3. Bioinformatic tools

The program Sequencher 5.2.4 (Gene Codes Corporation) was used for sequence analysis. Upon discovery of any discrepancy between the reference sequence (NC_007089.1) and a patient’s sequence, a search was conducted at the National Center for Biotechnology Information (NCBI) database (http://www.ncbi.nlm.nih.gov/gene/411) to determine whether the variant had been previously reported. If it proved to be a single nucleotide polymorphism (SNP), the allelic frequency of this change was searched in the 1000 Genomes database (http://www.1000genomes.org/). Previous reporting of the mutation was corroborated in the database of specific MPS6 mutations (http://mpsi6-database.org/) and the human gene mutations database (http://www.hgmd.cf.ac.uk/ac/index.php). If the variant had not been reported, bioinformatic predictions of the defects in the protein variant were performed using the following bioinformatic tools: PolyPhen-2 (http://genetics.bwh.harvard.edu/pph2/) (Adzhubei et al., 2010), MutationTaster (http://www.mutationtaster.org/) (Schwarz et al., 2014), MuStab (http://bioinfo.ggc.org/mustab/) (Teng et al., 2010), SNPs&GO (http://snps-and-go.biocomp.unibo.it/snps-and-go/) (Calabrese et al., 2009), and Provean (http://provean.jcvi.org/index. php) (Choi et al., 2012). Novel mutated amino acids were located on a 3D ARSB protein graphic using modeling of the enzyme arylsulfatase B (1FSU PDB) described by Bond et al. (1997) for Pymol 1.7.4 free accesses software (http://www.pymol.org/) (The PyMOL Molecular Graphics System, Version 1.7.4 Schrödinger, LLC). Adaptive Poisson–Boltzmann Solver (APBS) and PDB2PQR packages were used to calculate the electrostatic potentials of all the protein atoms (Baker et al., 2001; Dolinsky et al., 2004). The APBS default parameters were set. The thermodynamic stability changes of mutations were computed using the force-field FoldX (http://foldx.crg.es/) (Schymkowitz et al., 2005). The MutationTaster was used to compare the Arylsulfatase B amino acids with different species with mutated proteins, and the conservation scores were calculated by Consurf (http://consurf.tau.ac.il/) (Ashkenazy et al., 2010).
Novel missense mutations p.H111P, p.C121R, p.H147P, p.D300E, p.S334I, and p.G446S had an abnormal electrostatic surface potential as it is shown in Fig. 2. Since electrostatic interaction plays a significant role in diffusion-influenced reactions (Zhou, 1993), it was possibly affected in the mutant protein. These mutations also had thermodynamic stability changes in comparison with the wild type protein (Table 3).

Most of the affected novel mutations conserved amino acid residues. Fig. 3 shows that histidine at position 111, cysteine at position 121, aspartic acid at position 300, glycine at position 446, serine at position 334, histidine at position 147, and leucine at position 129 are highly conserved in arylsulfatase B of different species. The conservation score for these amino acids was 9 (1 less conserved and 9 fully conserved) except for histidine at position 147 whose score was 7.

4. Discussion

Of the previously described mutations, the c.430G>A (p.G144R) mutation was reported (Isbrandt et al., 1994) in a patient with a severe phenotype in the United States of America. c.1143-1G>C (IVS5-1G>C), an intronic mutation (Garrido et al., 2007), alters the splicing acceptor site, causing the loss of exon 6 and a truncated protein of 391 amino acids. It was the most common mutation in a study of Argentinian and Spanish patients, representing 21.9% (7/32) of the mutant alleles and the patients with this mutation had a severe phenotype. The nonsense mutation c.966G>A (p.W322*) was also reported in the Argentinian

Table 2

| Patient | Mutation – Allele 1 | Location | Mutation – Allele 2 | Location | Origin |
|---------|---------------------|----------|---------------------|----------|--------|
| 1       | c.430G>A (p.G144R) | Exon 2   | c.361T>C (p.C121R) | Exon 2   | Bogotá |
| 2       | c.332A>C (p.H111P) | Exon 2   | c.1143-1G>C (IVS5-1g-c) | Intron 5 | Bogotá |
| 3       | c.332A>C (p.H111P) | Exon 2   | c.332A>C (p.H111P) | Exon 2   | Bogotá |
| 4*      | c.332A>C (p.H111P) | Exon 2   | c.1143-1G>C (IVS5-1g-c) | Intron 5 | Funza  |
| 5*      | c.332A>C (p.H111P) | Exon 2   | c.1143-1G>C (IVS5-1g-c) | Intron 5 | Funza  |
| 6       | c.384_386delCCT (p.L129del) | Exon 2 | c.966G>A (p.W322*) | Exon 5   | Bogotá |
| 7       | c.504G>A (p.G302R) | Exon 5   | c.900T>G (p.D300E) | Exon 5   | Bogotá |
| 8       | c.900T>G (p.D300E) | Exon 5   | c.1336G>A (p.G446S) | Exon 7   | Medellín |
| 9       | c.1336G>A (p.G446S) | Exon 7   | c.1601A>G (p.S334I) | Exon 8   | Cartagena |
| 10      | c.440A>C (p.H147P) | Exon 2   | c.1531_1553del (p.V513*) | Exon 8   | Cartagena |
| 11      | c.440A>C (p.H147P) | Exon 2   | c.1340G>T (p.C447F) | Exon 8   | Ipiales |
| 12      | c.1001G>T (p.S334I) | Exon 5   | c.1304G>T (p.C447F) | Exon 5   | Ipiales |
| 13      | c.1001G>T (p.S334I) | Exon 5   | c.1001G>T (p.S334I) | Exon 5   | Bogotá |
| 14      | c.904G>A (p.G302R) | Exon 5   | c.904G>A (p.G302R) | Exon 5   | Bogotá |

* This two patients are siblings.
NA: The programs could not perform the prediction of the effect due to the type of mutation. RI: Reliability Index; PC: Prediction Confidence.

Table 3
Predictions of the effect of novel mutations in Colombian MPSVI patients, all bioinformatics tools showed that the variants are deleterious and causing disease mutations. c.1531_1553del could not be assessed by these bioinformatics tools because the prediction programs are not designed for this type of mutations; this variant is a frameshift mutation that causes a premature stop codon. The FoldX for the wild type is −47.39 kcal/mol.

| Mutation          | PolyPhen-2   | Mutation taster | Mustab           | SNP&GO       | Provean | FoldX   |
|-------------------|--------------|-----------------|------------------|--------------|---------|---------|
| c.332A>C          | Deleterious  | Disease causing | Decrease in protein stability | Disease causing | RI: 6   | Deleterious |
| (p.H111P)         | Score: 1.00  | Probability: 0.99 | PC: 94.6%        | Probability: 0.97 | −55.83 kcal/mol |
| c.361T>C          | Deleterious  | Disease causing | Decrease in protein stability | Disease causing | RI: 5   | Deleterious |
| (p.C121R)         | Score: 0.99  | Probability: 0.99 | PC: 83.6%        | Probability: 0.92 | −39.63 kcal/mol |
| c.900T>G          | Deleterious  | Disease causing | Increased in protein stability | Disease causing | RI: 10  | Deleterious |
| (p.D300E)         | Score: 1.00  | Probability: 0.99 | PC: 26.8%        | Probability: 0.93 | −33.35 kcal/mol |
| c.1336G>A         | Deleterious  | Disease causing | Decrease in protein stability | Disease causing | RI: 3   | Deleterious |
| (p.G446S)         | Score: 0.96  | Probability: 0.99 | PC: 78.4%        | Probability: 0.64 | −49.44 kcal/mol |
| c.1601A>G         | NA           | Disease causing | Probability: 0.99 | NA           | NA      | NA      |
| (p.G534W)         |              |                 |                 |              |         |         |
| c.1531_1553del    | NA           | NA              | NA               | NA           | NA      | NA      |
| c.1001G>T         | Deleterious  | Disease causing | Decrease in protein stability | Disease causing | RI: 8   | Deleterious |
| (p.S334I)         | Score: 1.00  | Probability: 0.99 | PC: 78.3%        | Probability: 0.92 | −48.8 kcal/mol |
| c.440A>C          | Deleterious  | Disease causing | Decrease in protein stability | Disease causing | RI: 9   | Deleterious |
| (p.H147P)         | Score: 1.00  | Probability: 0.99 | PC: 81.6%        | Probability: 0.94 | −44.18 kcal/mol |
|                   |              |                 |                 |              |         |         |

of the 8 new mutations reported in this study (Table 3), 6 were missense mutations. For the c.332A>C (p.H111P) mutation, bioinformatic programs predicted that it was a deleterious and disease-causing mutation. Amino acid 111 is conserved in the arylsulfatase B of different vertebrate species (Fig. 3). The positive charge is missing with the loss of histidine, and this change could result in alteration of contacts. In addition, this mutation could alter the electrostatic potential of the protein, affecting its functioning and interactions. The bioinformatic programs predicted that c.361T>C (p.C121R) was a deleterious and disease-causing mutation; amino acid 121 is conserved in all sulfatases, and there is a disulfide bond at this position. In the literature, a similar variant was reported by Wei-Dong et al. in 1992. They described a mutation in a disulfide bond (p.C117R) (Jin et al., 1992) and the loss of protein stability and enzymatic activity. Bioinformatic programs also showed that c.900T>G (p.D300E) was a deleterious and disease-causing mutation because the aspartic acid at position 300 is one of the 10 residues involved in the active site. The p.D300E mutation was reported by Karageorgos but one caused by a different mutation c.900T>A, which results in a severe phenotype (Karageorgos et al., 2007). c.1336G>A (p.G446S) was found to be a deleterious and disease-causing mutation, according to the bioinformatic programs; the amino acid 446 is conserved in all of the sulfatases. Regardless, both amino acids are neutral, glycine is apolar, and serine is polar, which disrupts the three-dimensional structure of the protein; a similar change in the protein was also reported (Karageorgos et al., 2007) for the p.G446R mutation that causes a severe phenotype. The bioinformatic programs predicted that the c.440A>C (p.H147P) was a deleterious and disease-causing mutation. Histidine 147 is conserved in all of the sulfatases, and this amino acid is in the active site of the ARSB enzyme. Exchanging histidine for proline makes it lose its positive charge, which affects the active site. Finally, c.1001G>T (p.S334I) was found to be also a deleterious and disease-causing mutation according to the bioinformatic programs; serine in the 334 position is conserved in all of the sulfatases. Since serine is

and Spanish patients study (Garrido et al., 2007). This mutation causes a truncation of the ARSB enzyme, eliminating the 212 carboxy-terminal residues; it was reported in a heterozygous patient with a severe-intermediate phenotype. The missense mutation c.904G>A (p.G446S) was described in an Italian patient with a severe phenotype in a homozygous form (Villani et al., 1998), and the amino acid at position 302 may be particularly important for this polypeptide as it is fully conserved in all of the sulfatases of eukaryotic lineage. The c.1340G>T (p.C447F) mutation results in the substitution of cysteine for phenylalanine (Karageorgos et al., 2007); since amino acid 447 is a potential glycosylation site, this change affects its affinity for the mannos-6-phosphate receptor and enzyme uptake.

Fig. 2. Electrostatic surface potential of the mutant proteins, products of the novel missense mutations. The color scale ranges from −5 kT/e (red) to 5 kT/e (blue). A) Wild type protein; B) p.H111P; C) p.C121R; D) p.H147P; E) p.D300E; F) p.S334I; G) p.G446S. There are changes in the electrostatic surface potential in all mutant proteins.
polar and isoleucine is apolar, the substitution of a polar amino acid for a hydrophobic amino acid could affect the residue contacts.

4.1. Stop-loss mutation

According to MutationTaster, the c.1601A > N (p.*534W) mutation causes disease and a change in the amino acid sequence, elongating the protein and therefore resulting in a protein with 584 amino acids (the normal protein has 533). The same effect on the length of protein was described by Arl et al. (Arlt et al., 1994) who reported the p.*533Q mutation that results in a protein with 584 amino acids. Although most precursors of higher size are proteolytically degraded before reaching the trans-Golgi network (Arlt et al., 1994; Lippincott-Schwartz et al., 1988), some proteins are able to avoid this mechanism. Changes in the kinetic and conformational structure could increase the enzymatic activity; this last effect could partially compensate the first effect and be associated to an intermediate phenotype, but an altered three-dimensional structure of the mutant polypeptide causing a higher susceptibility to proteinase. The phenotype described for this mutation is intermediate (Arlt et al., 1994).

4.2. Deletions

According to MutationTaster, c.384_386delCCT causes disease, and a leucine is lost at position 129 (p.L129del), without altering the reading frame (non-frameshift). It is the same c.382_384delCTC mutation that has been described by Fernández-Marmiesse et al (Fernández-Marmiesse et al., 2014) which produces the same change in the electropherogram and in the protein. This loss is located near the active site, and such deletion mutations cause structural changes (Jurecka et al., 2012). The c.1531_1553del(CCCGTGTACCTTCCCTGCACAGGA) is a 23 bp deletion. This deletion changes the reading frame (frameshift), causing a premature stop codon that results in a 513-amino acid protein (truncated protein). A similar protein abnormality was reported by Voskoboeva et al. in 2000; they described the p.Y513* (c.1539C > G) nonsense mutation in Russian patients, which results in a truncated protein of 513 amino acids (Voskoboeva and Krasnopol’skaia, 2000).

The most common SNPs were rs1065757 and rs25415. Two patients carried the minor allele of 5 SNPs and another 2 carried the minor allele of 4 SNPs. This finding and the high frequency of some novel mutations suggest that a haplotype analysis should be performed. Although this study is not sufficient to determine the founder effect of new mutations found in the Colombian population, it constitutes the first step in understanding the population genetics of Colombian patients with MPS VI. A genotype–phenotype correlation analysis was not performed.

5. Conclusions

Molecular analysis allowed the identification of mutations in Colombian patients. Some of the mutations in our patients were identified as private as described in previous studies. This is the first molecular study of patients with MPS VI in the Colombian population, and 9 of the mutations had not been previously reported. As described in the literature, most mutations have been found in exons 1, 2, 5, and 8.
in this study, most mutations were found in exons 2 and 5 (65%). Unlike prior reports, no mutations were found in exon 1. Of the 8 new mutations, 3 of them were found in more than one Colombian patient; p.H111P was found in 4 patients from Bogota, p.H147P in 2 patients from Cartagena, and p.S334I in 2 patients from Ipiales (Fig. 4A). The other 5 mutations were only present in one patient each one (Fig. 4B). Of the mutations that had not been previously reported, bioinformatics predicted that they caused significant changes to the enzyme, and most of these mutations were located near the active site of ARSB (Fig. 5). Additional studies to analyze the possible founder effect of these mutations are required. Furthermore, we will continue to study patients diagnosed with MPS VI through in silico, in vitro, and ancestry studies as well as genotype-phenotype correlation studies.

Declaration of interest

The authors declare that they have no competing interests. The translation of this article was sponsored by Biomarin Colombia LTDA.

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