Full Length Research Paper

Single nucleotide substitution mutations and polymorphisms in ECM1 gene in lipoid proteinosis in siblings of a Pakistani family

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A number of mutations in extracellular matrix protein 1 (ECM1) that is a glycoprotein and expressed in skin and other tissues are reported to cause a rare, autosomal recessive disorder called lipoid proteinosis (LP). The peculiar manifestation of LP is hoarseness of voice caused by laryngeal infiltration in infancy. Skin and mucous membrane changes clinically become apparent, and the disease typically follows a slowly progressive, yet often benign, course. About 300 cases of LP have been reported, but occurrence in siblings is rare. In this study, two siblings (18 and 24-year-old) of a Pakistani family were reported to have LP. This study presents two brothers with scaly itchy lesions on whole body, hoarse voice and macroglossia. Their deceased father had similar clinical manifestations but mother, younger brother and sister were unaffected. Blood from affected and clinically unaffected family members were collected with informed consent. The ECM1 gene containing 10 exons were amplified and sequenced. Both patients showed non-pathogenic missense and silent mutations in exon 6 and 8. In exon 6, a nucleotide C was substituted to T (C®®®®T) at codon 2, in patient 1. This non-pathogenic missense mutation causes appearance of amino acid cysteine instead of arginine that is part of normal ECM1 protein. In patient 2, polymorphism of nucleotide C to T (C/T) was observed observed in exon 6 that may lead to the appearance of cysteine and/or arginine in the resulting gene product. In exon 8, a nucleotide G was substituted to A (G®®®®A) at codon 53, in patient 1. This substitution leads to a silent mutation as serine is coded by both forms of codon. In patient 2, polymorphism of nucleotide G to A (G/A) was observed in exon 8 that do not cause any change in the coded amino acid. These findings represent a set of missense and silent mutations supporting an unusual function of ECM1 protein, broadening the spectrum of disease-linked mutations in rare cases of LP.

Key words: Lipoid proteinosis, extracellular matrix protein 1 (ECM1), missense, silent mutation, single nucleotide polymorphism, exons 6 and 8, genodermatosis.

INTRODUCTION

Lipoid proteinosis (LP) is a rare autosomal recessive genodermatosis which is characterized by deposition of an amorphous hyaline material in skin, mucosa and viscera. LP is also known as hyalinosis cutis et mucosae or Urbach-Wiethe disease (OMIM: 24700) which was first described by Urbach and Weithe (1929). The peculiar manifestation of LP is onset in infancy with hoarseness of voice caused by laryngeal infiltration. Skin and mucous membrane changes clinically become apparent, and the

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Abbreviation: LP, Lipoid proteinosis.
Figure 1. Family pedigree of the affected and unaffected members. Two sons and deceased father are affected, while the mother, son and only daughter have normal phenotype.

disease typically follows a slowly progressive, yet often benign, course. Virtually, any organ may be involved, but visceral involvement rarely leads to clinically significant consequences. The exceptions are involvement of the central nervous system and respiratory tract, which may result in seizures and airway obstruction, respectively. Lifespan is otherwise normal (Horey et al., 2009; Dyer, 2008).

Lipoid proteinosis has been linked to mutations in the gene encoding extracellular matrix protein 1 (*ECM1*). To date, no effective treatment is known (Chan et al., 2007). The *ECM1* protein has important physiological and biological roles in epidermal differentiation, binding of dermal collagens and proteoglycans, and regulation of angiogenesis. The precise function of *ECM1* is still unknown. Many pathogenic mutations have been reported so far, mostly nonsense, missense or splice site mutations with the majority occurring in exon 6 or 7 of the 10-exon in *ECM1* gene encoding a glycoprotein (Salih et al., 2011). LP is histologically characterized by intercellular deposits of periodic acid Schiff (PAS)-positive hyaline material in the skin, mucous membranes and internal organs. Loss-of-function mutations in the gene encoding *ECM1* on chromosome 1q21 7 have been identified in LP; fewer than 20 variants have been reported, that occurred in siblings rarely.

This study reports an unusual finding of non-pathogenic single nucleotide base substitution mutations in exons 6 and 8 in two brothers of a Pakistani family suffering from LP. The results of this study help to understand the spectrum of genetic variations in this rare case of genodermatosis.

**MATERIALS AND METHODS**

Following informed consent and approval from institutional ethical committee two brothers of 18 and 24 years of age suffering from LP, from Karachi, Pakistan were selected for this study. Both patients were presented with scaly itchy lesions on the whole body, hoarse voice and macroglossia. These lesions started at infancy in both brothers. Their hoarseness and low pitch voice was so severe that it was difficult to hear them. On examination, they had eczema, with multiple infections and scars on their faces, chest and back. Patients were found to have characteristic features of LP such as multiple beaded papules along the eyelids, fissured lips and slight macroglossia with a hard woody tongue having a homogenous look. According to their mother, her husband had died almost with similar skin problem and hoarseness, whereas their younger brother and sister were normal (Figure 1).

Routine investigations (Blood CP, SUCE and LFT) together with abdomen USG and CT scan were normal. However, buccal mucosa biopsy with PAS stain was suggestive of this condition. Skin sections stained with haematoxylin and eosin (H&E) showed widespread deposition of hyaline-like material around blood vessels and adnexal epithelia, and thickening of the basement membrane at the dermal-epidermal junction. On periodic acid-Schiff (PAS) staining and PAS-diastase, these sites were PAS positive and diastase resistant (Figure 2).

**Detection of mutation in ECM1 gene**

The DNA of the patients and their clinically unaffected family members were extracted from peripheral blood by the standard phenol-chloroform DNA extraction method (Maniatis et al., 1982). Polymerase chain reaction (PCR) amplification of *ECM1* gene was performed using eight pairs primers (forward and reverse) encompassing all 10 exons and situated in flanking introns as described earlier (Hamada et al., 2003) and briefly given in Table 1.

For PCR amplification, the 50 µl PCR reaction volume was used containing 1xPCR amplification buffer, 250 ng genomic DNA, 0.6 mM each primer, 1.5 mM MgCl2, 0.2 mM each dNTPs and 2.5 U Taq DNA polymerase. The amplification was performed by using standard protocol (Hamada et al., 2003; Hougenhouck-Tulleken et al., 2004). The amplifications were performed at 95°C for 5 min, followed by 35 cycles of 95°C for 45 s, primer-specific annealing
Table 1. Genomic primers used for PCR amplification of ECM1.

<table>
<thead>
<tr>
<th>ECM1 exon</th>
<th>Primer sequence (5’-3’)</th>
<th>PCR product size (bp)</th>
<th>Annealing temperature (°C)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ECM1-1F</td>
<td>AGCTGGGACTGAAGTCATGGC</td>
<td>416</td>
<td>62</td>
</tr>
<tr>
<td>ECM1-1R</td>
<td>TAAAGGCTCCACTGGGCTAG</td>
<td>622</td>
<td>59</td>
</tr>
<tr>
<td>ECM1-2/3F</td>
<td>TCCTACACTCTTGATCTCCA</td>
<td>622</td>
<td>59</td>
</tr>
<tr>
<td>ECM1-4/5F</td>
<td>CAGTGACCCTCCAGTTTCT</td>
<td>484</td>
<td>59</td>
</tr>
<tr>
<td>ECM1-6F</td>
<td>AGCCTTGAGAAGCAGAGGA</td>
<td>671</td>
<td>59</td>
</tr>
<tr>
<td>ECM1-6R</td>
<td>AGTGAACGGGACCTGAGTT</td>
<td>548</td>
<td>59</td>
</tr>
<tr>
<td>ECM1-7F</td>
<td>TTATCTGCTGCCCAGTGTC</td>
<td>408</td>
<td>59</td>
</tr>
<tr>
<td>ECM1-7R</td>
<td>ACATGGATGGATGACTGTC</td>
<td>408</td>
<td>59</td>
</tr>
<tr>
<td>ECM1-8F</td>
<td>CACATCAACAGTTGGCTCT</td>
<td>499</td>
<td>59</td>
</tr>
<tr>
<td>ECM1-8R</td>
<td>GCATCTTCTGGCATCAGAT</td>
<td>548</td>
<td>59</td>
</tr>
<tr>
<td>ECM1-9F</td>
<td>AGTTGCTAGTCTTCCCACA</td>
<td>469</td>
<td>60</td>
</tr>
<tr>
<td>ECM1-9R</td>
<td>AGGCGAGGTAGGATGAGGA</td>
<td>548</td>
<td>60</td>
</tr>
<tr>
<td>ECM1-10F</td>
<td>AATCCAGCTGTGCAAGGAG</td>
<td>469</td>
<td>60</td>
</tr>
<tr>
<td>ECM1-10R</td>
<td>GTAATGAGTTTCAGATGGA</td>
<td>469</td>
<td>62</td>
</tr>
</tbody>
</table>

Figure 3. Amplification of ECM1 exons 1 to 10 resolved on 2.5% agarose gel.

temperature for 45 s, 60°C for 45 s. PCR products (5 µl) were analyzed by 2.5% agarose gel electrophoresis (Figure 3). PCR products were then purified by using commercial kit (QiAquick PCR Purification Kit, Qiagen, Crawley, UK) and sequenced directly using Big Dye® Terminator v3.1 cycle sequencing kit in an ABI 3100 genetic analyzer (Applied Biosystems, Foster City, CA, USA).

RESULTS

Direct sequencing of PCR amplified products from the affected siblings with LP were shown to have one base point mutation by substitution of a nucleotide in exon 6 and 8. In patient 1, a nucleotide C was substituted to T (C→T) as compared to normal control, at codon 2, in exon 6 (Figure 4). This non-pathogenic missense mutation replaces the of amino acid residue arginine (CGC) to cysteine (TGC), unlike the normal ECM1 protein. In patient 2, at the same location, polymorphism of nucleotide C to T (C/T) was observed (Figure 5), which may lead to the appearance of cysteine and/or arginine in the resulting gene product.

The sequence analysis of exon 8 in patient 1 and control
Subjects showed that a nucleotide G was substituted to A (G→A) at codon 53, in exon 8 (Figure 6). This substitution leads to a silent mutation as in both forms of codon code for same amino acid residue serine (TCG → TCA). In patient 2, at the same location of exon 8, polymorphism of nucleotide G to A (G/A) was observed (Figure 7), but this does not cause any change in the coded amino acid.

**DISCUSSION**

A number of recent genetic studies reported the involvement of homozygous frame shift, missense, nonsense and single nucleotide substitution, deletion and insertion mutations in ECM1 gene LP which is located on chromosome 1q21.2 (Salih et al., 2011; Samdani et al., 2010; Kumar et al., 2009; Nasir et al., 2009). About 300 cases of LP have been so far reported. It is found worldwide, but is more common in some countries, such as South Africa, where a founder effect has been demonstrated (Van Hougenhouck-Tulleken et al., 2004). The ECM1 has three known splice variants: ECM1a, ECM1b and ECM1c. To date, 41 distinct germline missense, nonsense, splice site, small and large deletions and insertions, have been reported (Chan et al., 2007). Approximately, 50% of the mutations cluster to
The exact pathogenesis of this disease is unknown. It was considered either as a lysosomal storage disease or disorder of disturbance in collagen synthesis. Recently, loss of function mutations in the gene encoding ECM1 on band 1q21 has been identified as the cause of lipoid proteinosis. The most common location is at exons 6 and 7, patients with exon 6 mutation result in a more severe phenotype, while exon 7 mutations display slightly milder clinical features (Kumar et al., 2009).

This study reports an unusual finding of non-pathogenic single nucleotide base substitution mutations in exons 6 and 8 in two brothers suffering from LP of a Pakistani family. The coding region of ECM1 gene containing 10 exons were amplified and sequenced. Both patients were shown to have one base mutation by substitution of a nucleotide in exons 6 and 8. In patient 1, a nucleotide C was substituted to T (C→T) at codon 2, in exon 6 (Figure 4). This non-pathogenic missense mutation causes conversion of amino acid residue arginine (CGC) to cysteine (TGC) unlike the normal ECM1 protein. In patient 2, at the same location, polymorphism of nucleotide C to T (C/T) was observed (Figure 5), which may lead to the appearance of cysteine and/or arginine in the resulting gene product.

The sequence analysis of exon 8 in patient 1 and control subjects showed that a nucleotide G was substituted to A (G→A) at codon 53, in exon 8 (Figure 6).
This substitution leads to a silent mutation as in both forms of codon code for same amino acid residue, serine (TCG → TCA). In patient 2, at the same location of exon 8, polymorphism of nucleotide G to A (G/A) was observed (Figure 6), but this does not cause any change in the coded amino acid. Earlier, a case of homozygous 62-bp insertion (Nasir et al., 2009) and a case of homozygous frame shift mutation leading to premature termination of ECM1 protein causing LP (Samdani et al., 2010) have been reported in Pakistani families.

This study presents an unusual missense and silent mutations in siblings of Pakistani family suffering from LP. These findings support an unusual function of ECM1 protein and broaden the spectrum of disease-linked mutations in this rare case of genodermatosis in this region. The identification of these mutations in ECM1 gene may be of help in making the involvement of ECM1 gene clear in LP. The results of this study may also be exploited to establish a phenotype-genotype correlation, more precise and accurate prognosis and/or diagnosis, carrier screening for the transfer of LP within families horizontally as well as vertically and also in making the effective strategies for genetic counseling in this regard.

REFERENCES


