Homozygosity for a Novel ABCA4 Founder Splicing Mutation Is Associated with Progressive and Severe Stargardt-like Disease

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PURPOSE. To clinically characterize and genetically analyze members of six families who reside in the same village and manifest a rare form of retinal degeneration.

METHODS. Ophthalmic evaluation included a full clinical examination, perimetry, color vision testing, and electoretinography. Genomic DNA was screened for ABCA4 mutations with the use of microarray analysis and direct sequencing. RNA analysis was performed with RT-PCR and sequencing.

RESULTS. The authors recruited 15 patients with a unique retinal disease who are members of six highly consanguineous Arab-Muslim families from a single village. During early stages of disease, funduscopic and angiographic findings as well as retinal function resemble those of Stargardt disease. However, later in life, severe, widespread cone-rod degeneration ensues. Marked progressive involvement of the retinal periphery distinguishes this phenotype from classic Stargardt disease. Genetic analysis of ABCA4 revealed two novel deletions, p.Cys1150del and c.4254-15del23. One patient, who was a compound heterozygote, manifested typical Stargardt disease. The remaining 14 patients were homozygote for the c.4254-15del23 intronic deletion and had the progressive form of disease. We identified an identical ABCA4 haplotype in all alleles carrying this mutation, indicating a founder mutation. Detailed RT-PCR analysis in normal retina and lymphoblastoid cells revealed expression of the full-length ABCA4 transcript and three novel transcripts produced by alternative splicing. The full-length ABCA4 transcript, however, could not be detected in lymphoblastoid cells of affected homozygote patients.

CONCLUSIONS. These results expand the genotype-phenotype correlation of ABCA4, showing that homozygosity for the novel c.4254-15del23 splicing mutation is associated with a severe progressive form of disease. (Invest Ophthalmol Vis Sci. 2007;48:4308–4314) DOI:10.1167/iovs.07-0244

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The ABCA4 (ABCR) gene encodes a retina-specific ATP-binding cassette (ABC) transport protein located at the rim of photoreceptor outer segment discs. The ABCA4 protein belongs to the ABC transporters protein family and is involved in retinoid (N-retinylidene-phosphatidylethanolamine) transport across the disc membrane.1,2

Mutations in ABCA4 cause a variety of retinal diseases.3–5 ABCA4 is the only gene known to cause autosomal recessive (AR) Stargardt disease (detection rate, 58%–78%).6,7 Patients with Stargardt disease experience central visual loss in the first two decades of life, often with an initially normal fundus appearance. Later on, Stargardt disease is associated with macular atrophy and yellowish deep retinal flecks, while peripheral vision is largely preserved. Three different subtypes of Stargardt disease have been described based on electroretinal findings,89 with group 3 the most severe phenotype in which peripheral retinal involvement occurs in addition to maculopathy. ABCA4 mutations were also found in a few patients with AR retinitis pigmentosa, a progressive retinal degeneration that initially affects the rod photoreceptors but that ultimately also leads to gradual loss of cones. Mutations in ABCA4 have also been identified in 30% to 50% of patients with AR cone-rod degeneration (CRD),5,10 in which cones are initially affected and rod degeneration ensues. The involvement of ABCA4 in age-related macular degeneration is still controversial, but accumulating data point to involvement in a small percentage of cases.11

We describe here a detailed clinical and genetic analysis of patients from six families residing in the same village who manifest a unique retinal degeneration caused by novel ABCA4 mutations. In addition, we describe the existence of novel ABCA4 transcripts, produced by alternative splicing, in the normal retina.

PATIENTS AND METHODS

Patients

This study, which involved human patients, conformed to the tenets of the Declaration of Helsinki and was approved by the institutional review board. Informed consent was obtained from the subjects after explanation of the nature and possible consequences of the study. DNA was purified from blood samples using the salting-out technique.12

Clinical Evaluation

Full ophthalmologic examination, including assessment of visual acuity (VA), ocular motility, pupillary reaction, biomicroscopic slit-lamp, and dilated fundus examination, was performed in all patients. Subsequently, kinetic perimetry (Goldman visual fields, targets V4e and III4e), color vision testing, full-field electoretinography (ERG), and electrooculography (EOG) were performed.13 Full-field ERGs were recorded using corneal electrodes and a computerized system (UTAS 3000; LKC, Gaithersburg, MD). In the dark-adapted state, a rod response to a dim blue flash and a mixed cone-rod response to a white
ABC4 Mutation Associated with Stargardt-like Disease

ABC4 Haplotype Analysis

DNA amplification was performed by 35 cycles consisting of denaturation at 94°C, annealing at 54°C, and extension at 72°C for 30 seconds each. In addition, an initial denaturation and final extension steps of 5 minutes each were performed. Haplotype analysis included four microsatellite markers (D1S435, D1S188, D1S2719, D1S497) flanking ABC4 and two single nucleotide polymorphisms (SNPs; rs472908 and rs560426 with 5 U NED enzyme Tsp681I at 65°C and Hpy188III at 37°C, respectively) located within ABC4. Restriction products were separated by electrophoresis in 3% agarose gel for 50 minutes.

Mutation Analysis

The single-strand conformation polymorphism and direct sequencing techniques were used to screen all 50 coding exons of ABC4 (RefSeq NM_000350.2). Screening for all known ABC4 sequence changes was performed at Asper Biotech using the ABC4 genotyping microarray (http://www.asperophthalmics.com/ABCRgenetest.htm).

Reverse Transcription and Nested PCR Analysis

We used a previously described protocol to study ABC4 splicing. Total RNA was isolated from Epstein-Barr virus-transformed lymphoblastoid cells (RNeasy; Qiagen, Valencia, CA). RT-PCR was performed (Reverse-iT; Abgene, Epsom, UK) with random decamers, and cDNA techniques were used to screen all 50 coding exons of ABC4 and two single nucleotide polymorphisms (SNPs; rs472908 and rs560426 with 5 U NED enzyme Tsp681I at 65°C and Hpy188III at 37°C, respectively) located within ABC4. Restriction products were separated by electrophoresis in 3% agarose gel for 50 minutes.

RESULTS

Description of Pedigrees

Six reportedly unrelated Arab-Muslim families (Fig. 1A) residing in the same village who share a similar ocular phenotype were studied. Three of the families (MOL0006, MOL0035, and MOL0261) were large and included multiple consanguineous marriages. The inheritance pattern was interpreted as AR in all six families, with a pseudodominant inheritance pattern in MOL0006 because of multiple consanguineous marriages (Fig. 1A).

Clinical Description

The ocular findings in early stages of disease in patients from the six families resemble those of typical Stargardt, mainly involving the macula, with a resultant reduction of central visual field. At the age of 12, his VA dropped to 6/60, and macular changes were evident with atrophy and pigment clumps (Figs. 2I–L). Interestingly, autofluorescence imaging showed a relatively preserved parapapillary ring (Fig. 2K), as previously noted in patients with ABC4 mutations.

Different measures of retinal function showed a similar pattern of progression over time, with initial macular involvement evolving into widespread retinal degeneration. The full-field cone-flicker ERG of MOL0006 II-2 (age 20) was delayed and amplitudes were still within normal limits (Fig. 3A), whereas the response of the foveal cones was severely reduced and delayed (Fig. 3B). Visual field testing of patient MOL0035 II-2 (age 20) showed that in addition to the central scotoma expected in Stargardt disease, there was also some narrowing of the visual fields (Fig. 3C), suggesting peripheral retinal involvement. Patient MOL0261 II-1 (Fig. 1A, age 25) is a representative example of the more advanced stage of disease. Mixed cone-rod and 30Hz cone flicker full-field ERG amplitudes were reduced to approximately 50% of low threshold of normal (Fig. 3D), and cone flicker implicit time was markedly delayed (Fig. 3D, cone flicker waveforms; Fig. 4, other measures of retinal function [diamond correlating to age 25]). Funduscopic findings in this patient (age 29) indeed support widespread retinal involvement.

To better characterize the progress of disease and its association with age, we evaluated retinal function in 14 patients with the same retinal phenotype and genotype. The data presented in Figure 4 show a statistically significant decline in retinal function compared with age in most measures. Cone flicker implicit time showed a trend of reduction with age, which did not achieve statistical significance. In addition, in most patients who could undergo EOG, dysfunction of the retinal pigment epithelium was evident. The only exception to this pattern of progression was patient MOL0006 I-2. At the age of 40, her fundus findings remained localized to the macular area, which had a typical Stargardt-like appearance (Fig. 2C). Full-field ERG function was maintained within normal limits, and EOG findings were at the lower limit of normal (Fig. 4, squares). This exception can be explained by the molecular genetic findings detailed below.

Haplotype and Mutation Analysis

Given the retinal phenotype described and the inheritance pattern of the six families, we considered ABC4 the major candidate gene causative of the disease. Therefore, we performed haplotype analysis of families MOL0006 and MOL0035 using DNA markers located within and flanking the ABC4 gene (Fig. 1B). In family MOL0006, two different haplotypes, 1 and 3, were cosegregated with the disease under the assumption of AR inheritance. Patient MOL0006 I-2 (diagnosed with typical Stargardt) carried both haplotypes, and her unaffected husband carried haplotypes 1 and 2; the latter did not cosegregate with the disease. All affected children, diagnosed with progressive,
severe Stargardt-like disease, were homozygote for haplotype 1. In family MOL0033, two haplotypes, 4 and 6, shared an identical portion within \( \text{ABCA4} \) and cosegregated with the disease. Moreover, haplotype 1, cosegregating in family MOL0006, shared the same \( \text{ABCA4} \) portion with haplotypes 4 and 6.

Based on these data, we assumed that two different \( \text{ABCA4} \) mutations were responsible for the two retinal phenotypes, one of which is shared by haplotypes 1, 4, and 6. To identify these mutations, we screened the DNA of two affected patients for all known \( \text{ABCA4} \) sequence variants by using the Asper biotech \( \text{ABCA4} \) mutation detection microarray. The screen revealed seven sequence changes (c.1269C>T [p.His423His], c.1356+5delG, c.4773+48C>T, c.6069C>T [p.Ile2023Ile], c.6249C>T [p.Ile2083Ile], c.6285T>C [p.Asp2095Asp], and c.6764G>T [p.Ser2255Ile]) that had been previously interpreted as nonpathogenic changes.

We subsequently performed mutation screening of the whole \( \text{ABCA4} \) open-reading-frame and identified two previously reported nonpathogenic changes (c.6282+7G>A and c.302+26A>G) and a novel in-frame deletion (c.3449_3451delGCT [p.Cys1150del]) in exon 25, found heterozygously in patient MOL0006 I-2 (Fig. 5A). The deleted amino acid (Cys1150) is highly conserved, resides within a conserved \( \text{ABCA4} \) region (Fig. 5A, bottom), and is located at the 3'-end of the ABC transporter nucleotide-binding domain. We could not identify this mutation in 190 chromosomes from healthy Arab-Muslim control subjects, nor could we find it in 30 unrelated Arab-Muslim patients with Stargardt disease or CRD. We considered it a pathogenic \( \text{ABCA4} \) mutation. We could not identify a second \( \text{ABCA4} \) mutation in our mutation analysis of the remaining exons, but we were consistently unable to amplify exon 29 by PCR using the DNA of patients with a diagnosis of progressive Stargardt-like disease. We performed a long-range PCR reaction and amplified the region encompassing exon 29. Sequencing analysis of this fragment revealed an intronic deletion of 23 nucleotides overlapping...
with the forward primer of exon 29. The deletion (c.4254-15del23 [IVS28-15del23bp]) was located 15 bp upstream of exon 29 (Fig. 5B), and computer splice-site prediction analysis revealed a high score (0.83 of 1.00) for the wild-type acceptor site; the score for the mutant site was much lower (0.44 of 1.00). The mutation was shared by patients from all six families (Fig. 1) and was absent in 190 Arab-Muslim chromosomes of control subjects. Analysis of two SNP markers within the \( \text{ABCA4} \) gene revealed a shared haplotype (rs472908-A and rs560426-G) in all affected patients who were homozygote for the c.4254-15del23 mutation.

**Transcript Analysis**

Two \( \text{ABCA4} \) transcripts have thus far been reported in normal retinal samples. To explore the possibility that the c.4254-15del23...
15del23 mutation results in aberrant ABCA4 splicing, we isolated total RNA from a normal retina and from Epstein-Barr virus-transformed lymphoblastoid cells of one control subject, a heterozygote individual, and a homozygote patient. Surprisingly, the analysis of ABCA4 exons 27 to 31 in the normal retina yielded four different transcripts (Table 1; Fig. 5C). The
normal ABCA4 transcript, including exons 27 through 31 (ABCA4_v1), had the highest expression level. An aberrant transcript, in which exons 28 and 29 were skipped (ABCA4_v2) causing a frameshift deletion of 224 nucleotides, showed a relatively high expression level (75% of the wild-type expression). Two additional transcripts were in-frame and were expressed at relatively low levels. Exon 29 was skipped in ABCA4_v3 (expression level 9%, comparing to the wild-type transcript), causing a deletion of 99 nucleotides. Exon 29 and the first 114 nucleotides of exon 30 were skipped in ABCA4_v4 (15% of the wild-type expression level). A transcript similar to that of ABCA4_v4 was the first ABCR transcript to be reported.3 It contains exon 29 but not the first 114 nucleotides of exon 30.3,4 Although full-length exons were included or deleted in transcripts ABCA4_v1 to 3, transcript ABCA4_v4 contained only part of exon 30. Sequence analysis of this exon supported previous findings4 that identified a cryptic acceptor splice site located 114 nucleotides within exon 30 (a predicted splice-site score of 0.91). These transcripts were identified in normal retina and in lymphoblastoid cells from the healthy control subject and the heterozygote patient. Detailed RT-PCR analysis of the homozygote patient revealed only the three alternatively spliced isoforms, whereas the full-length ABCA4 transcript could not be detected.

**DISCUSSION**

The present study describes the involvement of two novel ABCA4 mutations in retinal degenerative disease of differing severity. The p.Cys1150del mutation results in a deletion of a highly conserved amino acid located 3' to the ABC transporter nucleotide-binding domain. It was found heterozygously in one patient (MOL0006 I-2) with the diagnosis of Stargardt disease. The founder mutation, c.4254-15del23, causes a splicing defect resulting in the absence of the wild-type ABCA4 transcript. This mutation was found homozygously in 14 patients who manifested progressive Stargardt-like disease and heterozygously in patient MOL0006 I-2.

More than 500 ABCA4 mutations have been identified as causing different retinal diseases, allowing a proposed genotype-phenotype correlation model.4,5,17 The simplified model suggests that patients with two severe mutations have retinitis pigmentosa, patients with one severe and one moderate mutation have CRD, patients with one severe and one mild mutation have Stargardt disease, and patients with only one severe or moderate mutation are at increased risk for age-related macular degeneration. As can be expected, actual genotype-phenotype correlations are more complex, and deviations from this model have been reported.5,17,18 Interestingly, as in the homozygote patients de-

**Table 1. ABCA4 Retinal Transcripts**

<table>
<thead>
<tr>
<th>Transcript</th>
<th>Exons Included</th>
<th>Exons Skipped</th>
<th>Effect on ORF</th>
<th>Relative Expression Level</th>
</tr>
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<tbody>
<tr>
<td>ABCA4_v1</td>
<td>wt (27–31)</td>
<td>None</td>
<td>None</td>
<td>1.00</td>
</tr>
<tr>
<td>ABCA4_v2</td>
<td>27,30,31</td>
<td>28,29</td>
<td>−224 bp frameshift</td>
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<tr>
<td>ABCA4_v3</td>
<td>27,28,30,31</td>
<td>29</td>
<td>−99 bp in-frame</td>
<td>0.09</td>
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<tr>
<td>ABCA4_v4</td>
<td>27,28,30p,31</td>
<td>29,30p</td>
<td>−213 bp in-frame</td>
<td>0.15</td>
</tr>
</tbody>
</table>
scribed here, progression from an initially mild, characteristic phenotype to a severe one, usually CRD, may also occur over time.18–20

The major limitation in obtaining an accurate genotype-phenotype correlation model is the correct interpretation of the effect of a given mutation on protein structure or function. This is particularly difficult with potential splicing mutations, which can lead to mild or severe mutations. Analysis of such mutations is challenging because of technical difficulties in amplifying ABCA4 mRNA from readily available tissues, such as peripheral blood and lymphoblastoid cells.4 In only one study thus far21 has reliable splicing data been obtained through this system, leading to an unexpected result: a frequent base substitution (c.2588G>C) in the first base of exon 17, initially interpreted as a missense mutation (Gly863Ala), created a splicing defect, resulting in a deletion of one amino acid (Gly863). This result emphasizes the importance of accurate mRNA analysis for each suspected splicing mutation before any genotype-phenotype assumptions can be made. Our nested RT-PCR analysis of the retina and lymphoblastoid cells revealed an unexpected result. We found that ABCA4 produced four different transcripts in the normal retina through an alternative-splicing mechanism. The two most common transcripts are the wild-type ABCA4 mRNA and a variant (ABCA4_v2) in which exons 28 and 29 are skipped. This deletion causes a frameshift and is therefore likely to be recognized and degraded by the nonsense mediated decay (NMD) mechanism.21 The other two transcripts (v3 and v4) have in-frame deletions and are expressed at very low levels, probably producing low amounts of nonfunctional ABCA4 proteins. The splicing region of all five ABCA4 transcripts described thus far (the wild-type transcript, a splicing variant with part of exon 30 as reported previously3,4 and three additional splice variants reported here) is located between exons 28 and 30 of the ABCA4 gene. Interestingly, this region contains a number of cryptic splice sites4 and a few splice-site mutations. The novel splicing mutation we identified in this study, c.4254-15del23, prevents the production of the wild-type ABCA4 protein and is likely to be a null mutation. However, we cannot exclude the possibility that small amounts of the wild-type protein are still translated in the retinas of these patients. If such a low level of expression indeed exists, it may explain the delayed appearance of widespread retinal degeneration in our patients compared with other patients who manifest early, full-blown CRD or retinitis pigmentosa because of homozygosity for null ABCA4 mutations.

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References