Mutations in \textit{ABCA12} Underlie the Severe Congenital Skin Disease Harlequin Ichthyosis

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Harlequin ichthyosis (HI) is the most severe and frequently lethal form of recessive congenital ichthyosis. Although defects in lipid transport, protein phosphatase activity, and differentiation have been described, the genetic basis underlying the clinical and cellular phenotypes of HI has yet to be determined. By use of single-nucleotide–polymorphism chip technology and homozygosity mapping, a common region of homozygosity was observed in five patients with HI in the chromosomal region 2q35. Sequencing of the \textit{ABCA12} gene, which maps within the minimal region defined by homozygosity mapping, revealed disease-associated mutations, including large intragenic deletions and frameshift deletions in 11 of the 12 screened individuals with HI. Since HI epidermis displays abnormal lamellar granule formation, ABCA12 may play a critical role in the formation of lamellar granules and the discharge of lipids into the intercellular spaces, which would explain the epidermal barrier defect seen in this disorder. This finding paves the way for early prenatal diagnosis. In addition, functional studies of ABCA12 will lead to a better understanding of epidermal differentiation and barrier formation.

Introduction

Harlequin ichthyosis (HI [MIM 242500]) is a very severe and usually lethal skin disorder of unknown cause (Hsu et al. 1989; Moreau et al. 1999; Sarkar et al. 2000). HI is also known as “harlequin fetus” because of the tendency for affected babies to be born prematurely. The neonate is covered in a thick “coat of armor” that severely restricts movement (Hsu et al. 1989; Dahlstrom et al. 1995; Sarkar et al. 2000). The skin dries out to form hard diamond-shaped plaques separated by fissures, resembling “armor plating.” The normal facial features are severely affected, with distortion of the lips (eclabion), eyelids (ectropion), ears, and nostrils. Historically, an HI-affected neonate would usually die within 2 d of birth because of feeding problems, bacterial infection, and/or respiratory distress. However, a number of patients have survived because of the wider availability of neonatal intensive care and likely benefits from oral retinoids (Hsu et al. 1989; Moreau et al. 1999; Elias et al. 2000; Sarkar et al. 2000). In survivors, the disease comes to resemble severe nonbullous congenital ichthyosiform erythroderma (NBCIE) (Haftek et al. 1996; Choate et al. 1998). The skin barrier remains severely compromised, which leads to increased transepidermal water loss, impaired thermal regulation, and increased risk of secondary infection (Moskowitz et al. 2004).

HI occurs in both sexes and in many ethnic groups. Autosomal recessive (AR) inheritance of HI has been...
inferred, but, in a large number of cases, the inheritance pattern cannot be ascertained, and the disorder could be due to a new dominant mutation. It has been subdivided into three types based on the expression of keratin 16 and profilaggrin and the appearance of keratohyalin granules (Dale et al. 1990). Studies have also reported abnormalities in the localization of epidermal lipids and have highlighted the abnormal ultrastructure of epidermal lamellar granules in HI that are associated with the absence of extracellular lamellar structures in all three subtypes (Dale et al. 1990; Akiyama et al. 1996, 1997, 1998). The nuclei of the keratinocytes also show abnormalities, since they appear to become flattened early on in differentiation (in the lower cell layers), but some are retained even as the cornified layer is sloughed off. This parakeratosis along with hyperkeratosis and hypergranulosis (retention of keratohyalin granules in the stratum granulosum) is a characteristic phenotype of the epidermis of neonates with HI (Buxman et al. 1979; Hsu et al. 1989). This may suggest that the process of terminal differentiation is not completed in HI-affected skin. Alterations in calcium-mediated signaling and protein phosphatase activity have also been reported in HI keratinocytes (Kam et al. 1997; Michel et al. 1999). Plausible candidate genes underlying these diverse cellular phenotypes in HI have included filaggrin, claudins, PP2A, and calpain 1 (Kam et al. 1997; Michel et al. 1999).

Mutations in the genes encoding transglutaminase-1 (TGM1 [MIM 190195]), ichthyin, 4 lipoxygenase 3 (ALOXE3 [MIM 607206]), and 12(R)-lipoxygenase (ALOX12B [MIM 603741]) have all been shown to underlie both NBCIE and lamellar ichthyosis (LI) (Huber et al. 1995; Jobard et al. 2002; Lefevre et al. 2004). Recently, missense mutations in the gene encoding the adenosine triphosphate (ATP)-binding cassette (ABC [GenBank accession number NM_173076]) transporter protein ABCA12 (MIM 607800) have also been shown to cause LI (Lefevre et al. 2003). Unlike the other AR forms of ichthyosis, the genetic basis of HI has not been elucidated.

In the present study, we have used SNP-array technology to localize the HI gene, on the basis of the principle of homozygosity mapping. This identified a major locus for HI on chromosome 2q35 and subsequent disruptive mutations in the ABCA12 gene in 11 of the 12 analyzed patients with HI. Most of the mutations identified would result in a severely truncated protein. This genetic study immediately establishes the means for early prenatal DNA diagnosis of HI and provides a molecular clue toward understanding the numerous biological abnormalities of epidermal differentiation in HI-affected skin.

Material and Methods

Clinical

Twelve families, from diverse ethnic backgrounds, with individuals affected with HI were included in the present genetic study—three from the United States, seven from Great Britain, and two from Italy. Ethical approval was obtained from the local research ethics committees. The clinical phenotype was verified as HI by at least two clinicians; the typical clinical presentation at birth is shown in figure 1. A summary of the clinical details is shown in table 1. The control population was of mixed ethnicity and consisted of 60 unrelated individuals.

SNP-Array Analysis

A SNP GeneChip Mapping 10K (Affymetrix) array analysis was performed on affected individuals from six families. Two of these families were consanguineous, with AR inheritance of HI. Patient DNA samples were processed in accordance with the standard GeneChip Mapping 10K Xba Assay protocol. Briefly, 350 ng of DNA was digested with XbaI and ligation to the XbaI adaptor prior to PCR amplification by use of AmpliTaq Gold with Buffer II (Applied Biosystems). For each DNA sample, four 100-μl PCRs were set up to obtain sufficient purified PCR product (20 μg), by use of Ultrafree-MC filtration column (Millipore), for subsequent fragmenta-
<table>
<thead>
<tr>
<th>PATIENT</th>
<th>ETHNICITY</th>
<th>GESTATION (wk)</th>
<th>BIRTH WEIGHT (g)</th>
<th>Retinoid (Dose [mg/kg/d])</th>
<th>Begun at Age (d)</th>
<th>OUTCOME</th>
<th>OTHER PROBLEMS</th>
</tr>
</thead>
<tbody>
<tr>
<td>49</td>
<td>British Bangladeshi</td>
<td>35</td>
<td>2,500</td>
<td>Acitretin (1)</td>
<td>2</td>
<td>SIE, now aged 2.5 years&lt;sup&gt;a&lt;/sup&gt;</td>
<td>Pseudomonal SI, FTT, gastrostomy</td>
</tr>
<tr>
<td>6</td>
<td>British Pakistani</td>
<td>34</td>
<td>1,750</td>
<td>Etretinate (1.15)</td>
<td>2</td>
<td>SIE, now aged 20 years&lt;sup&gt;b&lt;/sup&gt;</td>
<td>Anemia ectropion (surgery)</td>
</tr>
<tr>
<td>7</td>
<td>Bangladeshi</td>
<td>38</td>
<td>2,800</td>
<td>Acitretin NA</td>
<td>NA</td>
<td>SIE, now aged 14 years&lt;sup&gt;c&lt;/sup&gt;</td>
<td>FTT, limb deformity</td>
</tr>
<tr>
<td>55</td>
<td>British Indian</td>
<td>36</td>
<td>2,600</td>
<td>Acitretin (.75)</td>
<td>4</td>
<td>SIE, now aged 8 years&lt;sup&gt;b&lt;/sup&gt;</td>
<td>Neonatal sepsis (SA), hypernatremia, ectropion</td>
</tr>
<tr>
<td>40</td>
<td>Italian</td>
<td>40</td>
<td>3,200</td>
<td>Acitretin (1)</td>
<td>3</td>
<td>SIE, now aged 6 years&lt;sup&gt;a&lt;/sup&gt;</td>
<td>Neonatal sepsis (CA), ectropion surgery, limb strictures</td>
</tr>
<tr>
<td>44</td>
<td>Italian</td>
<td>36</td>
<td>2,000</td>
<td>Acitretin (1)</td>
<td>7</td>
<td>SIE, now aged 6 years&lt;sup&gt;b&lt;/sup&gt;</td>
<td>Nystagmus, PDA, neonatal sepsis</td>
</tr>
<tr>
<td>63</td>
<td>British Pakistani</td>
<td>36</td>
<td>2,600</td>
<td>Acitretin (.5)</td>
<td>4</td>
<td>SIE, now aged 2 years&lt;sup&gt;b&lt;/sup&gt;</td>
<td>Nystagmus, MDD, cataracts, FTT</td>
</tr>
<tr>
<td>50</td>
<td>British white</td>
<td>36</td>
<td>2,300</td>
<td>Acitretin (1)</td>
<td>3</td>
<td>SIE, now aged 4 years&lt;sup&gt;b&lt;/sup&gt;</td>
<td>Neonatal mild hypothermia</td>
</tr>
<tr>
<td>955-01</td>
<td>African-American</td>
<td>36</td>
<td>2,355</td>
<td>Isotretinoin (1)</td>
<td>7</td>
<td>Died at age 6 mo</td>
<td>Septicemia</td>
</tr>
<tr>
<td>915-06</td>
<td>Iranian</td>
<td>36</td>
<td>2,980</td>
<td>Isotretinoin (2)</td>
<td>19</td>
<td>Died at age 4 mo</td>
<td>Septicemia (SA)</td>
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<tr>
<td>928-01</td>
<td>Native American</td>
<td>40</td>
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<td>Not given</td>
<td></td>
<td>Died at age 1.5 d</td>
<td>Respiratory failure</td>
</tr>
<tr>
<td>69</td>
<td>British Pakistani</td>
<td>36</td>
<td>2,820</td>
<td>Not given</td>
<td></td>
<td>Died at age 3 d</td>
<td>Gram-negative sepsis</td>
</tr>
</tbody>
</table>

**NOTE.**—CA = *Candida albicans*; FTT = failure to thrive; MDD = motor developmental delay; NA = information not available; PDA = patent ductus arteriosus; SA = *Staphylococcus aureus*; SI = skin infection; SIE = severe ichthyosiform erythroderma.

<sup>a</sup> Remains on retinoids.
<sup>b</sup> Now off retinoids.
<sup>c</sup> Intermittent courses of retinoids.
tion with DNase I. Fragmentation was visualized by 4% agarose-gel electrophoresis to confirm the production of 50–100-bp PCR fragments prior to 3' labeling with biotin and hybridization to the SNP array. Hybridized arrays were processed with an Affymetrix Fluidics Station 450, and fluorescence signals were detected using the Affymetrix GeneChip Scanner 3000. Raw SNP call data were exported to Microsoft Excel for analysis. The average call rate (expressed as a percentage) were exported to Microsoft Excel for analysis. The average call rate (expressed as a percentage) ± SEM for the six patients was 97.62% ± 0.41%.

Microsatellite and Linkage Analysis

Data from SNP chips revealed an overlapping region of homozygosity, in five of the six affected individuals, at a position on chromosome 2. Microsatellite markers mapping to this region were used to confirm and refine the minimal region of homozygosity. DNA from patient blood samples was first amplified with the microsatellite markers D2S325 (chromosome 2q33.3) and D2S126 (chromosome 2q36.1) from the Applied Biosystems Linkage Mapping Set. Sample DNA was amplified with these primers at 5 μM for 30 cycles at 55°C, by use of Amplitaq Gold PCR Mastermix (Applied Biosystems). D2S128 (chromosome 2q34) and D2S1345 (chromosome 2q34) were selected from the University of California–Santa Cruz (UCSC) Genome Bioinformatics database, and PCR was performed using Bioline Biotaq DNA Polymerase, Bioline PCR buffer, 60 nM MgCl₂, and 20 pM of each primer, with a final reaction volume of 20 μl. Fluorescent-tagged PCR products were analyzed using the ABI 3700 capillary sequencer. Haplotypes were constructed from the microsatellite data of affected individuals and unaffected siblings. Linkage analysis was performed using the Genehunter program (Kruglyak et al. 1996).

Sequence Analysis of ABCA12

Primer design and PCR conditions were the same as those described elsewhere (Lefevre et al. 2003), with the exception that a 15-μl reaction volume was used. The PCR products covering each of the 53 exons and respective intron boundaries were purified using ExoSAP-IT (Amersham Pharmacia Biotech) and were sequenced on either the forward or reverse strand, by use of Ampli-Taq Gold PCR Mastermix (Applied Biosystems). D2S128 (chromosome 2q34) and D2S1345 (chromosome 2q34) were selected from the University of California–Santa Cruz (UCSC) Genome Bioinformatics database, and PCR was performed using Bioline Biotaq DNA Polymerase, Bioline PCR buffer, 60 nM MgCl₂, and 20 pM of each primer, with a final reaction volume of 20 μl. Fluorescent-tagged PCR products were analyzed using the ABI 3700 capillary sequencer. Haplotypes were constructed from the microsatellite data of affected individuals and unaffected siblings. Linkage analysis was performed using the Genehunter program (Kruglyak et al. 1996).

Results

Genetic Mapping

From the SNP-array analysis, only one region of the genome, 2q35, was found to have large blocks of homozygosity in common for five of the six affected individuals assessed by means of this methodology (fig. 2). This common interval was calculated to be a genetic distance of ~1.5 Mb. To support linkage of HI to 2q35, microsatellite markers that map to this region were identified using the UCSC Genome Browser. The microsatellite genotype data are consistent with a recessive mode of inheritance of HI and linkage of the disease gene to 2q35 in the largest consanguineous family with HI (fig. 3A). Linkage analysis was performed for this family, but the size of the family prohibited the calculation of a statistically significant LOD score (data not shown). This highlights the usefulness of the SNP array for homozgyosity mapping in diseases not applicable to classic genetic linkage studies. Affected individuals from families in which the mode of inheritance is unknown were also genotyped with these microsatellite markers. Of the 12 affected individuals analyzed in the present study, 11 were homozygous for one or more of the markers analyzed in the 2q35 region (fig. 3B).

Candidate-Gene Analysis

Six genes were identifiable, from the UCSC Genome Browser, that map within the minimal region, including BARD1, which interacts with the breast-ovarian cancer-associated protein BRCA1; ATIC, in which mutations are the cause of a purine biosynthesis disorder (AICARibosuria [MIM 608688]); Fibronectin 1, which is involved in cell adhesion and migration; the FLJ10116 gene, which encodes a protein of unknown biological function; and the PECA gene, which encodes a peroxisomal trans 2-enoyl CoA reductase. The gene ABCA12, which encodes the ABC transporter protein, also maps within the minimal homozygous region on chromosome 2q35. Missense mutations in this gene have recently been shown to underlie the less severe ichthyotic disorders LI and NBCIE (Lefevre et al. 2003). Since the transport of extracellular epidermal lipids is abnormal in HI (Dale et al. 1990) and a defect in epidermal lipid metabolism has been suggested in the pathogenesis of the disease (Buxman et al. 1979), ABCA12 could be a plausible candidate gene on the basis of chromosome position and function. The 53 coding exons of the ABCA12 gene were individually amplified by PCR and were sequenced for 12 patients. Mutations in ABCA12 were detected in 11
Figure 2. Localization of the HI gene to chromosome 2q35 by use of SNP-array mapping. A, Homozygous SNP profiles of chromosome 2 in six patients with HI. Black bars indicate homozygous SNP call data. Blue bars indicate the minimal common region of homozygous SNPs in patients 1–5, a genetic distance of \(~1.5\) Mb. B, Ideogram of chromosome 2 showing the position of ABCA12 (red bar), located within the minimal common region of homozygous SNPs (blue) identified in panel A. C, The genetic interval of the homozygous SNP “blocks” in each patient with HI, with clear indication that the ABCA12 gene maps within the common homozygous region for five of the six patients.

of the 12 patients (fig. 4). This indicates that ABCA12 mutations are a major genetic cause of HI. The majority of the mutations were predicted to result in a truncated protein; indeed, in one patient (40), the last 25 exons were homozygously deleted. Four patients all harbored the same homozygous frameshift mutation and, on the basis of haplotype analysis (see fig. 3B) and their shared ethnicity, it is likely to be the same founder mutation in at least three of the four patients. In individual 955–01, a homozygous missense substitution (W198C) in exon 6 was identified in addition to the more disruptive homozygous nonsense substitution in exon 44. The latter sequence change is more likely to be associated with HI. Unlike the other individuals with HI in the mutation study who were homozygous for a specific ABCA12 mutation, individual 44 was a compound heterozygote for two different ABCA12 mutations. Both SNP and microsatellite genotypes were heterozygous in the chromo-
Figure 3  Analysis of microsatellite markers flanking the \textit{ABCA12} gene. A, Segregation of microsatellite markers flanking \textit{ABCA12} at chromosome 2q35 in a family with HI. Only affected individual 6 is homozygous at this region of chromosome 2. Note that the majority of siblings with HI are deceased. B, Haplotype analysis of individuals with HI. Microsatellite markers map around the \textit{ABCA12} gene locus (215, 621, 772–215, 828, 657) at chromosome 2q35. Several patients are homozygous for the markers \textit{D2S325} (208, 096, 142–208, 096, 490), \textit{D2S128} (214, 917, 811–214, 918, 065), \textit{D2S1345} (214, 991, 103–214, 991, 425), and \textit{D2S126} (221, 842, 419–221, 842, 782). The genome positions (Mb) for \textit{ABCA12} and microsatellite markers on chromosome 2 are shown in parentheses.

Discussion

The ABC superfamily is a large family of transmembrane proteins involved in transport of substrates across both the plasma membrane and intracellular membranes. Mutations in ABC genes cause diverse diseases, including cystic fibrosis, X-linked adrenoleukodystrophy, Tangier disease, pseudoxanthoma elasticum, and Dubin-Johnson syndrome (Stefkova et al. 2004). In the present article, we report, we believe for the first time, that mutations in the \textit{ABCA12} gene, a member of the ABCA subfamily, cause HI. Genetic studies of this disorder have been hampered by the fact that families with more than one affected child are rare. Families with only one affected individual are unsuitable for genetic linkage studies be-
Figure 4  Mutation analysis of the ABCA12 gene in patients with HL. A, Table summarizing mutation analysis. An asterisk (*) indicates a sequence change to a stop codon. B, i, Chromatogram showing homozygous GC deletion in exon 42 of patient 7. ii, Chromatogram showing homozygous T→C substitution in exon 44 in patient 955-01. iii, Chromatogram showing heterozygous A deletion in exon 33 of patient 44.
cause of (1) the difficulties in determining whether the disease has been inherited in an AR mode or is due to a new dominant mutation and (2) the low statistical power for linkage analysis. An additional complicating factor is the possibility of genetic heterogeneity.

The results of the Affymetrix 10K SNP-array genome scan of five unrelated patients with HI showed suggestive linkage to a region on chromosome 2q35 on the basis of having a large homozygous SNP haplotype “block” in common, a genetic distance of ∼1.5 Mb. Microsatellite markers from this region were identified from the UCSC Genome Browser, and subsequent genotypes were consistent with a locus for HI on 2q35 on the basis of homozygosity at that locus in 11 of the 12 patients with HI described in this study. The ABCA12 gene encoding an ABC transporter protein that is postulated to be involved in the transport of epidermal lipids is localized within the chromosomal region encompassed by these microsatellite markers. This was regarded as an excellent candidate gene to underlie the HI phenotype, since abnormal or absent epidermal lipids in HI skin have been described. Mutation analysis revealed that 11 of the 12 analyzed patients with HI had a disease-associated mutation in the ABCA12 gene. The majority of the mutations were predicted to lead to a severely truncated protein, with the exception of a missense substitution and a splice-site mutation. In contrast, the ABCA12 mutations in LI/NCBIE were all missense and were restricted to within the ATP-binding domain of the protein. Thus, the mutations in HI are likely to disrupt other aspects of ABCA12 function and may provide the molecular reason behind the severity of HI compared with the milder skin conditions LI and NCBIE. Further mutation studies are in progress to identify the molecular spectra of mutations in a larger panel of patients with HI, including those subcategorized into the different types of HI (Dale et al. 1990). The data in the present article reveal that ABCA12 mutations are the major cause of HI.

In normal epidermis, the cytoplasm of cells of the upper spinous and granular cell layers contain small granules, 100–300 nm in size, known as “Odland bodies” (Odland 1960); lamellar granules; lamellar bodies; or membrane-coating granules. These move toward the cell membrane as they enter the granular cell layer. They discharge their lipid components and proteases into the intercellular space and play important roles in barrier function, desquamation, and intercellular cohesion within the stratum corneum (Menon et al. 1992; Elias et al. 2000). Lipid delivery to lamellar bodies has also been shown to be required for subsequent delivery of protein to the organelle (Rassner et al. 1999). The defect in ABCA12 in HI may explain the observed abnormalities in the lamellar granules, possibly associated with severely compromised skin-barrier function. This is analogous to the situation in lung in which surfactant is stored in lamellar bodies in alveolar type II cells. Mutations in ABCA3 cause fatal surfactant deficiency in newborns, with markedly abnormal lamellar bodies observed ultrastructurally (Shulenin et al. 2004). The lamellar bodies in epidermis also contain beta-defensin, a microbicidal peptide that is also found in the intercellular space, which suggests that it might be discharged with lipid components of the lamellar body (Oren et al. 2003). The defect in epidermal barrier and the loss of beta-defensin may also result in increased susceptibility to infection. Additionally, mutations in two other genes of the ABCA family cause disorders of lipid transport. The ABCA1 gene is mutated in Tangier disease and in the less severe, heterozygous, familial hypoalphalipoproteinemia as well as in premature atherosclerosis, the resultant disorder dependent on the site of the mutation (Brooks-Wilson et al. 1999; Rust et al. 1999; Clee et al. 2000). The ABCA4 gene is mutated in patients with Stargardt disease, in some patients with AR retinitis pigmentosa, and in patients with AR cone-rod dystrophy (Stefkova et al. 2004). Our observation that mutations in the ABCA12 gene cause HI is analogous to other systemic disorders of lipid transport. In addition, like ABCA1 and ABCA4, the site and type of mutation in the ABCA12 gene determine the severity of the condition, with the more disruptive genetic lesions, such as deletion or frameshift mutations, causing HI.

Prenatal screening tests for HI historically involved invasive procedures, such as fetoscopically and ultrasonographically guided fetal-skin biopsies that reveal key ultrastructural features, such as multivesicular lamellar granules in affected fetuses at 20–24 wk of gestation (Elias et al. 1980; Suzumori and Kanzaki 1991). Similar morphological changes have been described in amniotic fluid cells at 17 wk of gestation (Akiyama et al. 1994). Other methods that have been used in recent years for the prenatal diagnosis of HI are three-dimensional and four-dimensional real-time sonography (Vohra et al. 2003). A major translational benefit of our present findings is that DNA-based analysis should now be possible, which paves the way for earlier (i.e., first trimester) prenatal testing that is more reliable and conclusive than the other non-DNA–based prenatal screening techniques.

In summary, we have applied the rapid SNP-array genotyping technology to identify, by homozygosity mapping, ABCA12 mutations as the major underlying genetic cause of the devastating skin disease HI. Previously, this disease had proved difficult to map by classic linkage analysis because of the limited size of pedigrees. The association of ABCA12 mutations with HI will facilitate prenatal DNA diagnosis of this life-threatening disorder as well as the possibility of preimplantation genetic testing. In addition, results of the present study suggest a major role for ABCA12 in lipid trans-
port, barrier formation, and keratinocyte differentiation. The increased understanding of the biological role of ABCA12 in HI may lead to the development of targeted therapy to lessen the burden of this disease for both surviving children with HI and their parents.

Acknowledgments

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Electronic-Database Information

The accession number and URLs for data presented herein are as follows:

Ensembl Genome Browser, http://www.ensembl.org/
Online Mendelian Inheritance in Man (OMIM), http://www.ncbi.nlm.nih.gov/Omim/ (for HI, TGM1, ALOXE3, ALOX12B, ABCA12, and AICA-ribosuria)
UCSC Genome Bioinformatics, http://www.genome.ucsc.edu/ (for the Human Genome working draft of chromosome 2)

References


