Congenital Hereditary Endothelial Dystrophy Caused by SLC4A11 Mutations Progresses to Harboyan Syndrome

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Purpose: Homozygous mutations in SLC4A11 cause 2 rare recessive conditions: congenital hereditary endothelial dystrophy (CHED), affecting the cornea alone, and Harboyan syndrome consisting of corneal dystrophy and sensorineural hearing loss. In addition, adult-onset Fuchs endothelial corneal dystrophy (FECD) is associated with dominant mutations in SLC4A11. In this report, we investigate whether patients with CHED go on to develop hearing loss and whether their parents, who are carriers of an SLC4A11 mutation, show signs of having FECD.

Methods: Patients with CHED were screened for mutations in the SLC4A11 gene and underwent audiometric testing. The patients and their parents underwent a clinical examination and specular microscopy.

Results: Molecular analyses confirmed SLC4A11 mutations in 4 affected individuals from 3 families. All the patients were found to have varying degrees of sensorineural hearing loss at a higher frequency range. Guttate lesions were seen in 2 of the 4 parents who were available for examination.

Conclusions: Our observations suggest that CHED caused by homozygous SLC4A11 mutations progresses to Harboyan syndrome, but the severity of this may vary considerably. Patients with CHED should therefore be monitored for progressive hearing loss. We could not determine conclusively whether the parents of the patients with CHED were at increased risk of developing late-onset FECD.

Key Words: corneal endothelial dystrophy, Harboyan syndrome, sensorineural hearing loss, SLC4A11

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Loss of corneal clarity accounts for 2% of blind registrations in children below the age of 16 years in the United Kingdom. This is frequently because of inherited corneal endothelial dystrophies, which include posterior polymorphous corneal dystrophy and congenital hereditary endothelial dystrophy (CHED). Fuchs endothelial corneal dystrophy (FECD), the most common endothelial dystrophy, tends to occur as a late-onset disease. Although clinically distinct, these disorders share many signs and pathological features including altered morphology and loss of endothelial cells, secretion of an abnormal collagenous layer posterior to the basement membrane, and corneal edema.

CHED may be inherited in a dominant (CHED1, MIM 121700) or a recessive manner (CHED2, MIM 217700). CHED2 is more common and is characterized by a more severe phenotype than CHED1. Clinical signs include bilateral cloudy corneas at birth or in the early neonatal period, and sometimes, the condition is associated with congenital glaucoma. Because the severity of corneal opacity in CHED varies considerably, observation is an appropriate management option for patients with mild to moderate corneal clouding. Penetrating keratoplasty is the mainstay of surgical management, although successful Descemet stripping endothelial keratoplasty has recently been reported. Mutations in SLC4A11 (MIM 610206) on chromosome 20p13 cause CHED2. Harboyan syndrome (MIM 217400) is characterized by CHED in association with sensorineural hearing loss. The hearing deficit in Harboyan syndrome is not present at birth but is typically progressive with onset around the age of 10 to 15 years. Recesive mutations in SLC4A11 also account for this phenotype. More than 60 different homozygous or compound heterozygous mutations in SLC4A11 have been reported to cause either CHED or Harboyan syndrome with little evidence to support a genetic difference between these phenotypes.
FECD generally presents in the fifth or sixth decade. Corneal examination reveals endothelial guttae before the patient becomes symptomatic, but as the disease progresses, corneal edema develops. The patient may complain of glare, diminished vision, and discomfort, which is typically worse on awakening because of overnight hypoxia resulting in increased epithelial and stromal edema. FECD accounts for 22% of corneal transplants in the United Kingdom. It is usually sporadic, but familial forms with dominant inheritance have been documented. To date, dominant mutations in 4 genes, SLC4A11, COL8A2 (MIM *120252), ZEB1/TCF8 (MIM *189909), and LOXHD1 (MIM *613072) and 3 loci on chromosomes 5 (FECD5, MIM %613269), 9 (FECD7, MIM %613271), and 13 (FECD2, MIM %610158) have been reported. In addition, an X-linked locus has been described.

To determine whether recessively inherited CHED, Harboyan syndrome, and late-onset FECD may all coexist as allelic disorders within 1 family, we revisited 2 previously reported CHED families and also analyzed an unrelated CHED patient and 1 of his parents who was available for the examination.

MATERIALS AND METHODS

Recruitment and Clinical Assessment

Patients and their families were recruited to the study after their informed consent was obtained using a process that was approved by the local Research Ethics committee. The study adhered to the tenets of the Declaration of Helsinki. Three families (Fig. 1A), 2 of Pakistani origin living in the United Kingdom and the other of Mexican origin, had in total 4 affected members that had been diagnosed with CHED, by an experienced ophthalmologist. A full medical history was taken from all the family members, who were also subject to a detailed ophthalmic examination. In addition, patients with CHED underwent audiometry screening using either the Madsen Aurical audiometer (GN Otometrics A/S, Taastrup, Denmark) or an AD229 diagnostic audiometer (Interacoustics A/S, Assens, Denmark) according to the manufacturers’ instructions. The affected cases and their parents were also examined using specular microscopy (Tomey EM-3000 specular microscope, Tomey GmbH, Erlangen-Tennenlohe, Germany).

Molecular Analysis

Peripheral blood samples were collected from the affected patients and unaffected relatives. Genomic DNA was extracted from blood according to standard procedures and used as a template in the polymerase chain reaction to screen for mutations in the 19 exons of SLC4A11 as described previously.7

RESULTS

Ophthalmic History and Examination

Clinical examination of the patients with endothelial dystrophy suggested a diagnosis of CHED, whereas their family history suggested recessive inheritance (Fig. 1 and Supplementary Figure S1; http://links.lww.com/ICO/A190).

Mutation Screening

Sequencing of SLC4A11 confirmed the presence of homozygous mutations in all the families and the diagnosis of CHED. The patients from families A and B were all homozygous for the previously reported mutation c.1391G->A, p.G464D in exon 11, whereas the patient from family C was homozygous for a novel mutation c.397T->C, p.F133L in exon 4 (Supplementary Figure S2; http://links.lww.com/ICO/A190).

FIGURE 1. Family structure and ophthalmic history of patients with CHED. A, Families A, B, and C are shown. The proband is indicated by an arrow. B, Age at the examination and clinical findings are shown for each patient. Any ophthalmic interventions are also highlighted. IOP, intraocular pressure; OD, right eye; ONH, optic nerve head; OS, left eye; VA, visual acuity.
Audiometric Testing

Audiometric examination of patient II.1 from family A at 12 years of age revealed bilateral mid- to high-frequency sensorineural hearing loss consistent with a diagnosis of Harboyan syndrome (Fig. 2A). Audiometric examination was repeated at age 21, when some deterioration of hearing was measured. His younger affected sibling, whose corneal opacity was more marked at presentation, had subjective hearing problems at the age of 21, expressing a difficulty in hearing others’ conversations when studying at college. The audiometry results

FIGURE 2. Hearing tests. Audiometry of the CHED2-affected patients from families A, B, and C. The graphs show the frequency in hertz (x axis) and the hearing level in decibels (y axis). The test was performed on the patient at 12 years (family A, II.1), 21 years (family A, II.2), 15 years (family B, II.1), and 12 years (family C, II.1), respectively. Note the reduced sensorineural hearing loss in the range 30 to 50 dB at the higher frequencies.
showed a midfrequency bilateral sensorineural hearing loss consistent with these symptoms (Fig. 2A). The findings of the audiometry test in the patient from family B at age 11 were unremarkable. This was repeated at age 15 when a bilateral high-frequency hearing loss was found (Fig. 2B). Audiometric testing in the patient from family C at the age of 12 disclosed a bilateral sensorineural hearing loss in the range of 30 to 60 dB, mainly affecting the higher frequencies (Fig. 2C).

Specular Microscopy Examination
For each family, the parents, who were related by consanguinity, did not manifest obvious visual problems. However, a corneal examination of the parents in family A using a specular microscope showed guttae in the 44-year-old father, whereas the 46-year-old mother’s scan showed moderate pleomorphism with a normal cell count (Fig. 3). A specular microscopic examination of the mother at age 38 in family B showed unremarkable images with a normal endothelial cell count (data not shown). The father in family B aged 40 was unavailable for examination. The specular microscopic examination of the mother in family C, aged 37 years, revealed guttae but a normal cell count (Fig. 3). The father, aged 42 years, was unavailable for clinical examination.

DISCUSSION
Given that CHED, Harboyan syndrome, and some FECD are caused by mutations in SLC4A11, it is perhaps surprising that all these conditions have never been described within the same family. In this article, we report that all 4 patients originally diagnosed with CHED2 were subsequently found to have varying degrees of sensorineural hearing loss at the higher frequency range after audiometric examination, suggesting that CHED2 invariably progresses to Harboyan syndrome. We also investigated whether parents of these patients, who are themselves heterozygous SLC4A11 mutation carriers, have early signs of FECD and found that 2 out of the 4 parents who were available to be examined had guttae in their endothelium. However, the relatively young age of the parents makes it difficult to reach a conclusion regarding their risk of developing FECD. Although based on a small number and therefore speculative, these observations are nevertheless valuable given that CHED is a rare condition. Our findings, together with data from other similar studies, will lead to a better understanding of this rare condition. Interestingly, this pattern of inheritance for a mutation in SLC4A11 is somewhat similar to that recently reported for LOXHD1, in which dominant mutations cause FECD, whereas recessive mutations cause deafness.

The basis for the phenotypic heterogeneity seen in homozygous SLC4A11 mutation carriers with nonsyndromic CHED2 or Harboyan syndrome has been the subject of much speculation. The onset of progressive hearing loss in Harboyan syndrome has been described previously in children as young as 2 years and in adults as old as 33 years. However, auditory abnormalities have not been directly tested for in most cases of isolated CHED2, suggesting that Harboyan syndrome may have gone undetected. Because the proportion of null and missense SLC4A11 mutations identified in both conditions is similar, with no obvious clustering and the coexistence of the conditions within a family has been reported, there is little evidence to support a genetic basis for the difference between them. Instead, our longitudinal study suggests that CHED2 cases eventually experience some degree of sensorineural hearing loss and that variable age of onset of these symptoms

FIGURE 3. Corneal endothelium analysis. A, Specular microscopy images of the father (I.1) and mother (I.2) of family A and also the mother (I.2) of family C are shown. B, The table highlights the values for cell density (per square millimeter), coefficient of variation, and percentage hexagonality for the specular microscopic images. As a guideline, cell density of 2500 cells per square millimeter at middle age is within the normal range, and this value decreases with age, a coefficient of variation >0.4 and <50% hexagonality are indicative of an abnormal endothelium.
may be related to some unknown differences in the expression of genetic modifiers or exposure to environmental triggers.

The auditory phenotype seen in Harboyan syndrome is consistent with the observation that SLC4A11 is not only expressed in corneal endothelial cells but it is also expressed in fibrocytes of the stria vascularis in the inner ear, causing mutants but not those implicated in FECD. This is explained by the presence of the FECD mutant protein reduces the risk of developing late-onset FECD.

To conclude, we report here that CHED2 progresses to Harboyan syndrome and that both conditions are the same disease at different stages of development. We advise that patients with CHED2 should be monitored for progressive hearing loss and that their parents might be at an increased risk of developing late-onset FECD.

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