**Title:** Relative preservation of foveal outer retinal structure in infants with *AIPL1* associated Leber’s Congenital Amaurosis: implications for gene therapy

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Leber’s congenital amaurosis (LCA) is a group of heterogeneous disorders characterized by an early-onset rod-cone dystrophy and severe visual loss. The estimated incidence of LCA is 2 - 3 per 100,000 births, making it the commonest cause of inherited blindness in childhood. There have been several human clinical trials that have demonstrated the safety and efficacy of gene therapy in LCA type 2 (caused by RPE65 variants). One of the principal criteria for successful gene therapy of inherited retinal disorders is a degree of cellular preservation of the diseased retina, in order to allow for the potential of functional restoration. In this regard, LCA type 4 is of particular interest. LCA type 4 is caused by variants in the gene Aryl-hydrocarbon-interacting protein-like 1 (AIPL1), which plays a role in molecular chaperoning within photoreceptors. Although AIPL1 sequence variants are associated with a relatively severe LCA phenotype, recent electrophysiological findings suggest that there may be some preservation of retinal function in young children with AIPL1 mutations. Encouragingly, there have now been successful gene replacement interventions in AIPL1 animal models, thus lending support for a human clinical trial to be considered in AIPL1 LCA. Given the very poor prognosis of severe and rapidly progressive visual loss of AIPL1 LCA, and, at least early in the disease, the possibility of preserved retinal function, we aimed to determine if there was an early window of opportunity during which patients might be suitable for a human treatment trial, by means of assessing high resolution retinal imaging and clinical data, in order to identify young patients who might have relatively preserved outer retinal structure that would potentially be amenable to intervention.
Methods

Given that *AIPL1* mutations are estimated to account for only about 5% of LCA, this would equate to an approximate incidence of 1 in a million births. With such a rare disease, it was necessary to draw from an international pool of potential patients. We collected data on molecularly confirmed *AIPL1* patients including demographics, visual acuity, fundus examination findings, optical coherence tomography (OCT) findings and genotype, and analyzed this database to give us an impression of the known pool of *AIPL1* patients and their defining characteristics. In young patients where standard OCT imaging proved difficult or unsatisfactory, attempts were made where possible to obtain high resolution OCT images using hand-held OCT probes.

Results

We compiled data on 42 patients with molecularly confirmed *AIPL1* mutations and a LCA4 phenotype from 18 countries worldwide. The age of these patients ranged from 0.5 to 43 yrs old (median age 8 yrs), with 24 patients (57%) aged less than 10 yrs, and 10 patients (24%) aged less than 5 yrs old. The modal visual acuity was perception of light (PL), which was found in 21 patients (50%), with a range of visual acuities from no perception of light (NPL) to LogMAR 0.90. Posterior pole examination findings were available in 39 patients (93%). There was a normal posterior pole appearance in 7 of these 39 patients (18%) (age range 0.5 - 5 yrs), with a further 18 patients (46%) having retinal pigmentary changes without macular atrophy, and 13 patients (33%) exhibiting macular atrophy. One further patient had an epiretinal membrane (3%). The youngest patient with macular atrophy was 6
The commonest observed sequence variant was p.W278X(c.834G>A), which was found in at least one allele in 26 patients (62%), and in the homozygous state in 15 patients (36%) (Table).

OCT images were obtained for 19 patients (45%). The significant nystagmus and poor vision observed in this phenotype, along with the young age of some of the patients, posed considerable challenges to obtaining high quality foveal OCT scans. Of these 19 patients, six (32%) had poor quality scans or scans that were not foveal. Of the remaining 13 patients in whom good OCT images had been obtained, nine patients (69%) had no evidence of photoreceptor structure at the macula. Four of these 13 patients who had good macular scans, had undergone imaging with hand-held OCT probes. Promisingly, however, three of these 13 patients (23%) did demonstrate significant outer retinal structure, with relative preservation of the inner segment ellipsoid (ISe) layer and outer nuclear layer at the fovea, with one further patient (patient #41, 3 yrs old) demonstrating qualified evidence of a foveal ISe layer (Figure). Three of these four patients (75%) were homozygous for the common AIPL1 mutation p.W278X(c.834G>A). Of note, the three patients with clear evidence of a foveal ISe layer constituted the youngest in the cohort, each being only one year of age or younger. A 4-month follow-up scan carried out on one of the 1-yr-old patients (patient #37) continued to demonstrate evidence of retained outer retinal structure.

**Discussion**

This study shows that there is promising evidence from high resolution OCT imaging that, in the very youngest AIPL1 LCA patients imaged, there is relative preservation of foveal outer
retinal structure. To the best of these authors’ knowledge, the three youngest children in this cohort represent the youngest AIPL1 LCA patients in the literature to date that demonstrate foveal outer retinal structure, which might be amenable to gene therapy. It is also encouraging that the majority of the patients with preserved foveal outer retinal structure in this study harbored the most commonly observed severe sequence variant found in AIPL1 LCA; if these findings can be extrapolated, it may be that the majority of LCA AIPL1 patients still retain preserved foveal outer retinal structure when very young.

Given the very poor visual prognosis of this condition, and the possibility of a therapeutic window which seems to exist both structurally and functionally, along with the success of recent animal and human trials in this and other forms of LCA, a human gene therapy based approach may be worthy of consideration in a small group of selected patients with preserved outer retinal structure in AIPL1 LCA. The results of this study suggest that any intervention may need to be initiated in the first few years of life.

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