**IFT81 as a Candidate Gene for Nonsyndromic Retinal Degeneration**

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**P**rimary cilia are slender hair-like protuberances present on the cell surface, functioning as sensory antennae to relay extracellular signals to the cell. Mutations in genes encoding for ciliary proteins can cause structural or functional defects in the cilia and have been implicated in a number of developmental and degenerative diseases collectively called ciliopathies. Phenotypes observed in these diseases can range from syndromic multisystem pathologies affecting the central nervous system, heart, kidney, skeletal system, and retina, to nonsyndromic ciliopathies affecting a single organ system.1–8 Genetic studies indicate that the core of the IFT-B complex is functionally conserved in all ciliated organisms from Chlamydomonas reinhardtii to vertebrates.1,3,5 Consistent with these studies, two

**M**ethods. Whole exome sequencing was performed on 50 cone-rod dystrophy (CRD) patients that were previously screened for mutations in known retinal disease genes. The impact of candidate mutation was studied using in vitro cell system and in vivo zebrafish assay to determine the pathogenicity of the variant.

**R**esults. Compound heterozygous mutations in IFT81, including one nonsense (c.1213C>T, p.R405*) and one missense variant (c.1841T>C, p.L614P), were identified in a nonsyndromic CRD proband. Extensive functional analyses of the missense variant in cell culture and zebrafish strongly suggests its pathogenic nature. Loss of IFT81 impairs ciliogenesis and, interestingly, the missense variant displayed significantly reduced rescue of ciliogenesis in the IFT81 knockdown in vitro system. Consistently, dramatic reduction of rescue efficiency of the ift81 mutant zebrafish embryo by mRNA with the missense variant was observed, further supporting its pathogenicity.

**C**onclusions. Consistent with the function of the IFT-B complex in the maintenance of photoreceptor cilium, we report a case of mutations in a core IFT-B protein, IFT81. This represents the first report of mutations in IFT81 as a candidate gene for nonsyndromic retinal dystrophy, hence expanding the phenotype spectrum of IFT-B components.

Keywords: IFT-B complex, IFT81, cilia, cone rod dystrophy

**P**urpose. IFT81, a core component of the IFT-B complex, involved in the bidirectional transport of ciliary proteins, has been recently implicated in syndromic ciliopathies. However, none of the IFT-B core complex proteins have been associated with nonsyndromic retinal dystrophies. Given the importance of ciliary transport in photoreceptor function and structural maintenance, we sought to investigate the impact of IFT (intraflagellar transport) mutations in nonsyndromic retinopathies.

Keywords: IFT-B complex, IFT81, cilia, cone rod dystrophy

**C**onsistent with the function of the IFT-B complex in the maintenance of photoreceptor cilium, we report a case of mutations in a core IFT-B protein, IFT81. This represents the first report of mutations in IFT81 as a candidate gene for nonsyndromic retinal dystrophy, hence expanding the phenotype spectrum of IFT-B components.

Keywords: IFT-B complex, IFT81, cilia, cone rod dystrophy
core subunits of the IFT-B complex (IFT88, and -27) have been previously associated with syndromic ciliopathies,\textsuperscript{14-17} indicating the importance of these proteins in maintaining ciliary function in humans. Furthermore, mutations in peripheral members of the IFT-B complex, such as IFT172, have been previously linked to not only syndromic ciliopathies but also nonsyndromic retinal dystrophies.\textsuperscript{18} Interestingly, however, none of the core complex components have been associated with nonsyndromic retinal pathology to date in humans.

As a member of the IFT-B core complex, IFT81 forms the backbone of the core complex along with IFT72/74. The N-terminal calponin homology domain of IFT81 interacts with the N-terminal domain of IFT72 to form a tubulin binding module required for transportation of tubulin during ciliogenesis.\textsuperscript{8,19,20} Like other IFT-B core members, mutations in \textit{IFT81} have been observed to cause syndromic ciliopathies featuring renal medullary cysts, paraxial polydactyly, early onset rod-cone dystrophy, cerebellar atrophy, and intellectual disability.\textsuperscript{21} However, thus far, the association between mutations in \textit{IFT81} and nonsyndromic human disease has not been reported.

In our study, we investigated the genetics of cone-rod dystrophy (CRD), a progressive inherited retinal disorder by collecting and performing whole-exome sequencing of a cohort of CRD patients whose mutations remain undetermined after known retinal disease-related gene panel screening.\textsuperscript{22} Among them, one proband carrying bi-allelic mutations in \textit{IFT81} was identified. Both in vitro and in vivo functional assays of the putative mutant allele indicate that it negatively impacts protein function and is likely to be pathogenic. Therefore, our finding provides the first report linking the core IFT-B protein, IFT81, to nonsyndromic retinopathy.

**Materials and Methods**

**Subject and Clinical Evaluation**

The proband was diagnosed with CRD and recruited at the Department of Ophthalmology, Peking Union Medical College Hospital (PUMCH). Ophthalmic examinations were performed including best-corrected visual acuity (BCVA) testing, fundus examination, optical coherence tomography (OCT, 3D OCT-2000 Spectral Domain; Topcon, Tokyo, Japan), autofluorescence (AF, Spectralis HRA+OCT; Heidelberg, Germany) and electroretinogram (ERG, RetiPort ERG system; Roland Consult, Wiesbaden, Germany). Informed consent was obtained from the patient for this study. Blood samples were obtained from the patient and her parents. This study adhered to the Declaration of Helsinki and was approved by the Institutional Review Board PUMCH.

**Whole-Exome Sequencing and Bioinformatics Analysis**

Genomic DNA sample (approximately 1 μg) was sheared into 300- to 500-bp-long fragments and repaired with a single adenine base added to the 3′ ends using Klenow exonuclease. Illumina bar-coded adapters were ligated to the ends, and DNA fragments were PCR amplified. The DNA was then captured using the NimblegenSeqCap EZ Human Exome Library v.2.0 following the manufacturer’s protocols for whole-exome sequencing. Captured libraries were sequenced on the Illumina HiSeq 2000 (Illumina, San Diego, CA, USA) to generate 100-bp paired-end reads according to the manufacturer’s protocol.

Reads were mapped to the human reference genome hg19 using the Burrows-Wheeler Aligner.\textsuperscript{23} Base quality recalibration, local realignment, and variant calling were performed as previously described.\textsuperscript{24} Because CRD is a rare Mendelian disease, variants with a normal control population allele frequency higher than 0.5% (for a recessive model) or 0.1% (for a dominant model) in public or internal control databases were excluded.\textsuperscript{24} Databases used for this purpose include the 1000 Genome Database, dbSNP135 (National Center for Biotechnology Information, http://www.ncbi.nlm.nih.gov/SNP in the public domain), the National Heart, Lung, and Blood Institute (NHLBI) Exome Sequencing database (http://evs.gs.washington.edu/EVS, in the public domain), the National Institute of Environmental Health Sciences (NIEHS) Exome Sequencing database (http://evs.gs.washington.edu/niehsExome, in the public domain), and an internal control database of 997 exomes. We also retrieved variant frequency from the Exome Aggregation Consortium (ExAC) database.\textsuperscript{25} Pathogenicity of missense variants was predicted using SIFT,\textsuperscript{26} PolyPhen2,\textsuperscript{27} LRT,\textsuperscript{28} MutationTaster,\textsuperscript{29} and MutationAssessor\textsuperscript{30} as previously described.\textsuperscript{24}

**Sanger Sequencing and Segregation Test**

For each putative mutation, a 500-bp flanking sequence at both sides was obtained from the University of California, Santa Cruz (UCSC) genome browser (hg19 assembly). Primer\textsuperscript{31} was used to design a pair of primers for generating a 400- to 600-bp PCR product containing the variant for Sanger validation. After PCR amplification, the amplicons were sequenced on an ABI 3730xl. Family members of the patient were also Sanger-sequenced to test allele segregation with the disease.

**DNA Constructs**

Human \textit{IFT81} cDNA in pENTR-221 vector was obtained from Ken Scott and subcloned into the p3XFLAG-myc-CMV-14 vector obtained from G. Pazour. The patient-specific missense variant at exon 18 (c.1841T>C, p.L614P) was created using the Agilent’s QuikChange XL-II site-directed mutagenesis kit. GPZ human \textit{IFT81} shRNA (GENE Dharmacon) was obtained through the Baylor College of Medicine CBASS-shRNA library core facility (shRNA1: CATCTATCATTTTCCCTGAA, shRNA2: CCGTTTAGGAAGAACTATAA). The shRNA-mir is coexpressed with Turbo GFP as a bi-cistronic transcript allowing the visual marking of shRNA-mir-expressing cells. To avoid shRNA silencing of the exogenous mRNA, two silent mutations per seed sequence were introduced in \textit{IFT81} wild-type and c.1841T>C mutant vectors (Supplementary Fig. S5).

**In Vivo Zebrafish Functional Experiments**

Capped mRNA from the human cDNA construct of the wild-type and c.1841T>C mutant \textit{IFT81} was synthesized using Invitrogen’s mMESSAGE mMACHINE Kit (T7). RNA was purified using Zymo’s RNA Clean and Concentrator column. Zebrafish rescue experiments were performed on embryos of progeny from \textit{ift81}\textsuperscript{hi409tg/} \textit{ift81}\textsuperscript{c.1841T>C} crosses (parental zebrafish were heterozygous mutants because homozygous mutant is lethal). Embryos were injected at the one-cell stage with 1 nl (50 ng/μL) of human wild-type and c.1841T>C mutant IFT81 mRNA tagged with C-myc and flag. Approximately 100 embryos were injected for each group (with four biological repeats). The phenotype was scored at days post fertilization 3.5. The \textit{P} value was calculated from 2-way ANOVA analysis. Western blotting was performed on day 0 (hours post fertilization 6.0) and day 1 (hours post fertilization 26.0) whole embryo lysate. Anti–C-myc antibody (MMS-150p; Covance, Biologen, San Diego, CA, USA) was used to detect C-myc-IFT81 protein with bactin as a loading control (ab8227; Abcam, Cambridge, MA, USA). Experiments were designed and conducted in adherence to the Institutional Animal Care and Use Committee (IACUC.)
to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research.

Cell Culture, Transfection, and Ciliary Induction

hTERT-RPE1 and HEK 293T cells were cultured in 1:1 DMEM:F12 or DMEM, respectively, supplemented with 10% FBS. For transfection purpose, hTERT-RPE1 cells on coverslips were transfected with DNA constructs using Viafect chemical transfection reagent (Promega, Madison, WI, USA). For rescue experiments, short hairpin RNA (shRNA) and shRNA-resistant IT81 wild-type/mutant cDNA constructs were cotransfected at a 1:1 ratio. Following plasmid transfection, the formation of primary cilia was induced 14 hours after transfection by serum starvation (DMEM-F12 supplemented with 0.3% FBS) for 48 hours. For protein analysis, HEK 293T cells were transfected using MIRUS transit-293T reagents using the manufacturer’s protocol.

Immunofluorescence and Imaging

For immunofluorescence staining, hTERT RPE-1 cells were fixed, permeabilized, and blocked as previously described. Primary antibodies against γ-acetylated tubulin (clone 6-11B-1, 1:500; Sigma-Aldrich Corp., St. Louis, MO, USA), green fluorescent protein (GFP, 1:500; Invitrogen, Carlsbad, CA, USA), and Flag (1:300, F7425; Sigma-Aldrich Corp.) were used along with AlexaFluor 488 anti-mouse, Cy3 anti-rabbit, and Cy5 anti-mouse secondary antibodies, respectively, (Jackson Immunolabs, Bar Harbor, ME, USA). Cells were counterstained with 4',6-diamidino-2-phenylindole (1 μg/mL). Representative images were acquired on an Axio observer Z.1 inverted fluorescence microscope with apotome.2 based optical sectioning and structured illumination, on a 60× EC plan-neofluar objective (Carl Zeiss International, Dublin, CA, USA). For cell counting, image stacks were taken with a z-distance of 0.5 μm and projected as a maximal intensity image using Zeiss Zen 2 core software (Carl Zeiss International). Three biological replicates were scored with 100 cells per replicate for all test panels using images taken on a 40× EC plan-neofluar objective (Carl Zeiss International).

See Supplementary Methods for protein analysis and statistical analysis.

RESULTS

Identification of IFT81 Mutations as a Candidate for CRD

We performed whole-exome sequencing on a group of 50 unsolved CRD patients to discover candidate novel genes underlying the disease. One proband, a 22-year-old female, who

FIGURE 1. Fundus autofluorescence, ERG, and OCT tests display degeneration of macular region and loss of visual response in the proband. (A, B) Left and right fundus autofluorescence images display oval shaped macular hypofluorescence with a hyperfluorescent ring characteristic of CRD. (C) Scotopic and photopic ERGs: both rod and cone responses are significantly reduced, with more severe cone function loss. (D, E) Left and right eye OCT images display thinning of whole retina layers in the macular area observed in both eyes.
displayed a progressive decrease in vision with photophobia since age 12, had impaired color vision and poor visual acuity (left eye, 0.05; right eye, 0.06). The proband lacked any skeleto-developmental defects previously reported including postaxial polydactyly. Ultrasound (B-scan) of the various organs including liver, kidney, and pancreas presented normal structure of the liver and kidneys. Blood routine, liver function, renal function, and blood tests were all in normal functional range (Supplementary Dataset S2: Test report), suggesting a lack of any syndromic defects. Fundus images display oval shaped macular hypofluorescence with a hyperfluorescent ring, which are characteristic pigmented lesions found in CRD. In addition, ERG results displayed reduced responses in both rod and cone cells, with more severe cone function loss (Fig. 1A–C). Finally, we observed thinning of retina layers in the macular area by OCT (Figs. 1D, 1E). Other than retina defects, the patient did not display an additional clinically syndromic phenotype.

Because no mutations were identified during the initial screen of all known retinal disease genes, the proband was subjected to whole-exome sequencing. With mean sequence coverage of 90X, a total of 105,490 variants were initially identified. On filtering as described in the Materials and Methods section, 329 rare variants in 305 genes that affect protein coding remained. Among them, only four genes containing bi-allelic rare variants in their coding region were predicted to be damaging, including FAM186A, PKHD1L1, IRS1, and IFT81 (Supplementary Table S1). FAM186A and PKHD1L1 genes were excluded from the candidate list as homozygous loss of function mutations have been observed in a large number of individuals in the ExAC control cohort.25 Furthermore, IRS1 was excluded from further analysis as IRS1 has been previously linked to type 2 diabetes without retinal degeneration phenotype.33 As a result, IFT81 is the best candidate gene with one nonsense (c.1213C>T, p.R405*) and one missense variant (c.1841T>C, p.L614P) identified in the proband (NM_001143779). Sanger sequencing and segregation analysis of the proband’s family members showed that these two alleles were inherited separately from each parent and hence present in trans in the proband (Fig. 2A). Both variants are extremely rare as they have not been found in the public or in our internal control databases.

Mutant (c.1841T>C) IFT81 Fails to Rescue Ciliogenesis Defect In Vitro

To assess the effect of the missense variance identified in the patient on IFT81 protein function, the hTERT-RPE1 cell was
used as the model system. The hTERT-RPE1 is an immortalized cell line derived from human retinal pigment epithelial cells capable of forming cilia. Consistent with the previous report that IFT81 is required for cilia assembly in the cell, knockdown of endogenous IFT81 by shRNA (sequence: Materials and Methods and Supplementary Materials) leads to ciliogenesis defects. Specifically, on serum starvation, only 21% of shRNA-transfected cells grow cilia compared with 60% in wild-type cells (Fig. 3). To test whether the knockdown phenotype can be rescued by overexpression of IFT81 cDNA, shRNA-resistant (Supplementary Fig. S5) wild-type and mutant IFT81 cDNAs carrying the c.1841T>C mutation were cloned into mamma-

\[ \text{IFT81} \]\( ^{\text{C}} \)-IFT81 could rescue the ciliopathy defects in IFT81 deficient cells. In contrast, in cells cotransfected with mutant IFT81 cDNA along with shRNA, only 28% cells contain cilia, which is not significantly different from shRNA knockdown cells (Fig. 3C). This indicates that the mutant cDNA fails to rescue the ciliogenesis defect in cells with IFT81 knockdown (Fig 3B). To confirm whether this difference in rescue efficiency between wild-type and mutant cDNA is not caused by a difference in protein level, Western blot was carried out. As shown in Supplementary Figure S2C, similar levels of protein are detected for the wild-type and mutant alleles, indicating the mutation does not affect protein translation and stability. In addition, a similar phenotype was observed when two different shRNA targeting endogenous IFT81 (shRNA1, shRNA2, Supplementary Figs. S3A, S3B) were used. In both cases, the mutant flag tagged L614P-IFT81 protein displays localization signal throughout the cellular cytoplasm. Taken together, the (c.1841T>C) IFT81 allele does not rescue the ciliogenesis defect due to endogenous IFT81 knockdown, suggesting it is indeed deleterious to the protein function.

**IFT81 c.1841T>C Mutation Fails to Rescue Ciliary Defects In Vivo**

To further assess in vivo whether the missense (c.1841T>C) mutation is detrimental to IFT81 function, we performed a rescue experiment using ift81 mutant zebrafish. It has been reported that ift81 mutant zebrafish embryos exhibit ciliary defects including spine curvature and kidney cyst formation. To first assess whether this p.L164P missense variant affects IFT81 protein expression, we performed a Western blot on zebrafish embryos injected with wild-type or mutant IFT81 mRNA. Consistent with the result obtained from the cell line as described above, mutant IFT81 is expressed at similar levels to wild type, indicating that the variant does not affect protein level (Fig. 4A). To determine whether c.1841T>C-IFT81 could rescue the ciliopathy defects, we injected wild-type or mutant mRNA into embryos resulting from ift81<sup>1404066</sup>/<sup>1404066</sup> crosses. As expected, these crosses produced about 25% of the ift81<sup>1404066</sup>/<sup>1404066</sup> mutant embryos, which were further assessed for rescue potential of the mutant allele. A significant rescue effect (P = 0.0006) was observed when wild-type IFT81 mRNA was injected, with only 7% fish displaying ciliary defects. In contrast, injection of c.1841T>C-IFT81 mRNA was unable to rescue defects.
the phenotype, as 21% of injected embryos display ciliary defects, similar to the proportion of noninjected control embryos (~25%) (Fig. 4C). In addition, partial rescue in a small proportion of fish injected with wild-type IFT81 mRNA has also been observed (7%). These fish display normal kidney morphology but still exhibit increased spinal curvature in the postvent (Figs. 4B, 4C). Therefore, IFT81 mRNA with a c.1841T>C point mutation has a significantly lower rescue potential than that of the wild-type mRNA (P = 0.0006), consistent with the idea that the allele identified in the proband is likely to be pathogenic (Fig. 4C).

In summary, the p. L614P missense negatively impacts the function of IFT81 in both in vitro and in vivo assays. Together with the observation that the loss-of-function mutation in IFT81 results in a severe syndromic defect in the patient while our proband only exhibits retina defects, the missense variant identified in this study is likely to be a hypomorphic allele, thereby exhibiting pathogenic effects.

**DISCUSSION**

In this report, we identified mutations in IFT81 as candidates for a recessive form of nonsyndromic retinopathy. This is the first report highlighting the importance of a core IFT-B complex protein in retinal ciliopathy. The proband harbors a truncating mutation in exon 12 and a missense mutation in exon 18 acting in trans. Because the allele containing the nonsense truncating (c.1213C>T, p.R405*) mutation is likely subjected to nonsense mediated decay (NMD), the only remaining functional allele contains a nonsynonymous missense mutation leading to the substitution of leucine with a proline residue in the protein. The in silico predictions suggest that this substitution is functionally deleterious, supported by both in vivo and in vitro functional studies. Specifically, the c.1841T>C mutant allele displayed a significantly lower potential for rescuing the ciliary defects in IFT81-knockout hTERT-RPE cells and zebrafish, in comparison to wild type, supporting the functionally deleterious nature of this allele. Together, this provides compelling evidence supporting the pathogenicity of the variant according to American College of Medical Genetics and Genomics (ACMG) guidelines and classifies the variant as likely pathogenic allele class II. Due to limitations of exome sequencing, however, it is possible that larger deletions and structural aberrations in known retinitis pigmentosa genes may be missed. With exception of this caveat, given the functional evidence presented along with a lack of other potentially explanatory mutations, the c.1841T>C mutation in IFT81 is the likely candidate for inherited retinal dystrophy in this proband.

Ciliopathies are known to have characteristically variable phenotypes caused by different mutations in the same gene, depending on the exact nature of the patients’ alleles. For example, it has been reported that mutations in IFT-A core components, specifically, IFT140, IFT144, and IFTB peripheral component IFT172, can cause both syndromic and
nonsyndromic cases of retinal dystrophies. Similarly, mutations in IFT81 have also previously been associated with a syndromic disease featuring severe skeletal anomalies (polydactyly), nephropathies leading to renal abnormalities, intellectual disability, and retinal degeneration.\textsuperscript{21} Retinal pathology was observed in only one of the two patients displaying abnormal ocular movements, hemeralopia, and poor vision with altered scotopic and photopic ERG. The variation between the syndromic phenotype reported by Perrault et al.\textsuperscript{21} and the isolated retinal phenotype observed by our group is likely due to the nature of the identified mutations and their different corresponding impact on IFT81 function (Fig. 2). Due to the importance of IFT81 in maintaining the stability of the IFT-B complex, and by extension ciliary structure and functions, severe mutations leading to complete loss of IFT81 function may not be tolerated in nature and hence potentially lead to severe developmental defects. Indeed, it has been shown that loss of the C-terminal domain renders the IFT81 protein unstable, resulting in no observable assembly of IFT-B components in vitro.\textsuperscript{38} Severe, but not complete, loss-of-function mutations, like the in-frame exon loss and extension of IFT81 C-terminal end by 11 amino acids reported by Perrault et al. can lead to a syndromic phenotype.\textsuperscript{21} Therefore, it is plausible that a milder mutation, such as the missense mutation reported in this study, is likely to be a hypomorphic allele, thereby leading to nonsyndromic ciliary defects.

A simple model explaining the nonsyndromic clinical manifestation is that the retina is more sensitive to the IFT81 activity. The likely cause is that IFT complexes in the retina are involved in massive protein transportation from the protein synthesizing inner segment to the photoreceptive outer segment of the photoreceptor cells. This protein transport process is highly active and is crucial for proper photoreceptor architecture, outer-segment turnover, as well as phototransduction, hence rendering photoreceptors more sensitive to a decrease in function of the IFT complex.

It is possible that IFT81 function specifically in the retina is affected by the c.1841T>C mutation, for example, by altering/weakening its interactions with other retinal specific proteins. A similar case has been previously observed for ciliary proteins involved in massive protein transportation from the protein synthesizing inner segment to the photoreceptive outer segment of the photoreceptor cells. This protein transport process is highly active and is crucial for proper photoreceptor architecture, outer-segment turnover, as well as phototransduction, hence rendering photoreceptors more sensitive to a decrease in function of the IFT complex.

In conclusion, our findings highlight IFT81 as a candidate for inherited retinal dystrophy, thereby implying the importance of a core IFT-B protein, IFT81, in the human retina. This report represents the first link between IFT81 and nonsyndromic retinal dystrophy, expanding the phenotypic spectrum of IFT-B core members as a potential cause of nonsyndromic retinal dystrophy in humans.

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