Germline Mutations in CTNNB1 Associated With Syndromic FEVR or Norrie Disease

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Purpose. Germline and somatic mutations in CTNNB1 have been found in different types of human diseases. This follow-up study aimed to identify causative germline mutations in CTNNB1 and their associated ocular phenotypes through a comparative analysis of whole-exome sequencing data.

Methods. Annotated sequence variations in CTNNB1 were selected from in-house data from whole-exome sequencing of genomic DNA prepared from leucocytes of 3280 unrelated probands with different forms of eye diseases. Potentially pathogenic variants in CTNNB1 were analyzed by multistep bioinformatics analyses. Clinical data from probands with pathogenic variants in CTNNB1 were collected, and potential genotype-phenotype correlations were analyzed.

Results. Eleven rare variants that potentially affect the coding regions of CTNNB1 were detected in 11 of the 3280 samples, and four variants were considered to be potentially pathogenic. All four mutations, namely, c.999delC (p.Tyr333*), c.1104delT (p.His369Thrfs*2), c.1758_1742delinsACA (p.Leu580Thrfs*28), and c.1867C>T (p.Gln623?), were heterozygotes and considered to have a germline origin. Three of the four mutations are de novo mutations, and the status of the remaining mutation is unavailable. All four probands had the same class of closely related ocular diseases: one proband had FEVR, and three probands had Norrie-like retinopathy. The molecular results indicated that three probands showed systemic anomalies, as demonstrated by a follow-up survey, but relevant information for the remaining proband was unavailable.

Conclusions. The data suggest that germline truncating mutations in CTNNB1 cause autosomal dominant syndromic FEVR or Norrie disease. Patients with mutations in CTNNB1, KIF11, or NDP may have similar or overlapping phenotypes, but this phenomenon needs to be studied further.

Keywords: CTNNB1, mutation, FEVR, Norrie

The CTNNB1 gene encodes the beta-catenin protein, which is a key transcriptional co-activator of the Wnt/beta-catenin signaling pathway and is essential for embryonic development, adult tissue homeostasis, stem cell regulation, and tumorigenesis.1 Somatic mutations in CTNNB1 have been reported to be associated with many types of cancer, and germline mutations in the gene have been identified in patients with intellectual disabilities2–8 and autism spectrum disorders.9,10 involving multisissue phenotypes.11–15 Of these patients, 50% display abnormalities in vision, including strabismus,5,15 hyperopia,6 myopia,8 retinal detachment,14 and lens and vitreous opacities.14 Recently, mutations in CTNNB1 have been reported to lead to familial exudative vitreoretinopathy (FEVR) with or without extraocular manifestations.15

FEVR is a genetic disorder characterized by abnormalities in peripheral retinal vascularization and exhibits both clinical and genetic heterogeneity. The phenotypes of FEVR vary widely, from normal visual acuity with only mild non-perfusion zone in the peripheral retina, as demonstrated by fundus fluorescein angiography, to blindness resulting from retinal detachment.16 FEVR mostly occurs alone (nonsyndromic) but also can be accompanied by systemic abnormalities, including mental retardation, microcephaly, and osteoporosis and osteopenia (syndromic).17–19 In addition, FEVR can occur in an autosomal dominant, autosomal recessive, or X-linked manner, and several genes, including FZD4,20 LRP5,18 TSPAN12,21 and NDP,22 which are involved in the Wnt/beta-catenin signaling pathway, and KIF11,17 have been reported to cause FEVR. Mutations in NDP can cause both FEVR and Norrie disease,22 which involves similar vascular dysgenesis of the retina accompanied by microphthalmia, corneal opacity, iris synchia, and mental disorders. Mutations in KIF11 can cause both nonsyndromic and syndromic FEVR,17 involving microcephaly and mental retardation, and mutations in the other three genes cause nonsyndromic FEVR. Three heterozygous mutations in CTNNB1 were recently identified in probands with FEVR: one is associated with syndromic FEVR, and the other two are associated with nonsyndromic FEVR.15

Here, we investigated pathogenic variants in CTNNB1 in 3280 probands with hereditary eye diseases, including 106 probands with FEVR, based on exome sequencing. The results will extend our knowledge of the mutational frequency and spectrum of CTNNB1, as well as the potential genotype-phenotype correlations.
**Table 1.** Four Potential Pathogenic Mutations in *CTNNB1* Identified in Patients

<table>
<thead>
<tr>
<th>Chr Position NM</th>
<th>Amino Acid Change</th>
<th>Status</th>
<th>Patient ID</th>
<th>Allele Frequency in 1000G</th>
<th>ExAC</th>
<th>dbSNP</th>
<th>Novel or Known</th>
</tr>
</thead>
<tbody>
<tr>
<td>chr03 41268761 NM_001904.3 c.999del</td>
<td>p.Tyr333*</td>
<td>Het</td>
<td>QT2648</td>
<td>1 0 0 0</td>
<td>None</td>
<td>Novel</td>
<td></td>
</tr>
<tr>
<td>chr03 41274854 NM_001904.3 c.1104delT</td>
<td>p.His369Thrfs*2</td>
<td>Het</td>
<td>QT1615</td>
<td>1 0 0 0</td>
<td>None</td>
<td>Novel</td>
<td></td>
</tr>
<tr>
<td>chr03 41277269 NM_001904.3 c.1738_1742delinsACA</td>
<td>p.Leu580Thrfs*28</td>
<td>Het</td>
<td>QT409</td>
<td>1 0 0 0</td>
<td>None</td>
<td>Novel</td>
<td></td>
</tr>
<tr>
<td>chr03 41277993 NM_001904.3 c.1867C&gt;T</td>
<td>p.Gln623*</td>
<td>Het</td>
<td>QT1504</td>
<td>1 0 0 0</td>
<td>None</td>
<td>Novel</td>
<td></td>
</tr>
</tbody>
</table>

Hg19 position refers to human GRCh37/hg19 version from the UCSC genome. Chr, chromosome; Het, heterozygous.

**Materials and Methods**

A total of 3280 unrelated probands with different hereditary eye diseases (106 probands with FEVR and 3174 probands with other genetic eye diseases, including retinal degeneration, high myopia, glaucoma, hyperopia, and ocular dysplasia) were recruited to participate in this study. Peripheral blood samples and clinical data were collected from each proband, and genomic DNA was obtained from leukocytes isolated from the peripheral blood samples. In accordance with the tenets of the Declaration of Helsinki, written informed consent was obtained from all the participants or their guardians before the study. This study was approved by the Institutional Review Board of the Zhongshan Ophthalmic Center.

Whole-exome sequencing of the genome of all 3280 probands was performed by Macrogen (http://www.macro gen.com/en/main/index.php). In brief, genomic DNA from each proband was captured using an Agilent SureSelect Human All Exon Enrichment V5 Kit (50M; Agilent, Santa Clara, CA, USA), and the exome-enriched DNA fragments were then sequenced using the Illumina HiSeq4000 system (Illumina, San Diego, CA, USA). The average sequencing depth was at least 125-fold. Variant calling and annotation were performed using SAMtools (http://samtools.sourceforge.net/) and ANNOVAR (http://annovar.openbioinformatics.org/en/latest/), respectively.

The variants in *CTNNB1* were selected and analyzed through a multistep bioinformatics analysis as follows: low-confidence variant positions with coverage less than 10 were excluded; variants with a minor allele frequency ≥0.01, as determined based on the dbSNP database, 1000 Human Genome Project, Exome Variant Server (EVS), and Exome Aggregation Consortium (ExAC; Cambridge, MA, USA; http://exac.broadinstitute.org; accessed October 15, 2018), were excluded; the variants in noncoding regions and synonymous variants without a splice site change, as determined according to the Berkeley Drosophila Genome Project (in the public domain, http://www.fruitfly.org/) or NetGene2 (http://www. cbs.dtu.dk/services/NetGene2/), were excluded; the variants predicted to be benign by two online tools, PolyPhen-2 (http://genetics.bwh.harvard.edu/pph2/) and SIFT (http://sift.jcvi.org), were excluded; and the pathogenic variants identified through a comparative analysis of variants among different groups of diseases were filtered.

A review of the mutations and phenotypes of *CTNNB1* was performed to detect potential genotype-phenotype correlations.

**Results**

A total of 11 rare variants that potentially affect the coding regions of *CTNNB1* were detected in 11 of the 3280 samples, and these consisted of 5 missense variants, 2 splicing variants, 2 nonsense variants, and 2 frameshift variants. The five missense variants were identified in four patients with glaucoma and one patient with high myopia, whereas the two splicing variants were identified in one patient with retinitis pigmentosa and one patient with high myopia (Supplementary Table S1). The two potentially pathogenic frameshift variants (c.1104delT, p.His369Thrfs*2 and c.1738_1742delinsACA, p.Leu580Thrfs*28) were identified in two probands with FEVR or Norrie-like retinopathy, and the two potentially pathogenic nonsense variants (c.999del, p.Tyr333* and c.1867C>T, p.Gln623*) were identified in two probands with Norrie-like retinopathy (Table 1). The four mutations were classified as loss-of-function (LoF) mutations resulting in protein truncation. This type of mutation is extremely rare in *CTNNB1*, as determined by a review of the ExAC database (http://exac.broadinstitute.org/gene/ENSG00000168036; October 15, 2018). Three of the four mutations are de novo mutations, and the status of the remaining mutation is unknown because of the unavailability of genomic DNA from the parents.

The clinical manifestations of the four patients are summarized in Table 2. All four patients were examined before the age of 1 year, and they showed typical temporal dragging of the optic disc or retinal detachment accompanied by a retrolentiform fibrotic mass (Fig. 1, Supplementary Fig. S1). Three of the four patients showed signs of Norrie disease, including microphthalmia, corneal opacity, microcornea, other genetic eye diseases, including retinal degeneration, high myopia, glaucoma, hyperopia, and ocular dysplasia) were recruited to participate in this study. Peripheral blood samples and clinical data were collected from each proband, and genomic DNA was obtained from leukocytes isolated from the peripheral blood samples. In accordance with the tenets of the Declaration of Helsinki, written informed consent was obtained from all the participants or their guardians before the study. This study was approved by the Institutional Review Board of the Zhongshan Ophthalmic Center.

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**Table 2.** Clinical Data of Four Patients With Potential Pathogenic Variants in *CTNNB1*

<table>
<thead>
<tr>
<th>Patient ID</th>
<th>Mutations</th>
<th>Sex</th>
<th>Age at Onset, mo</th>
<th>Best Visual Acuity</th>
<th>Axial Length, mm</th>
<th>Main Phenotype</th>
</tr>
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<tbody>
<tr>
<td>QT2648</td>
<td>c.[999del];[=]</td>
<td>Female</td>
<td>3</td>
<td>NA</td>
<td>NA</td>
<td>14.00</td>
</tr>
<tr>
<td>QT2648</td>
<td>c.[999del];[=]</td>
<td>Female</td>
<td>3</td>
<td>NA</td>
<td>NA</td>
<td>14.00</td>
</tr>
<tr>
<td>QT1615</td>
<td>c.[1104delT];[=]</td>
<td>Female</td>
<td>8</td>
<td>NA</td>
<td>NA</td>
<td>19.00</td>
</tr>
<tr>
<td>QT409</td>
<td>c.[1738_1742delinsACA];[=]</td>
<td>Male</td>
<td>6</td>
<td>NA</td>
<td>NA</td>
<td>17.50</td>
</tr>
<tr>
<td>QT1504</td>
<td>c.[1867C&gt;T];[=]</td>
<td>Male</td>
<td>3</td>
<td>NA</td>
<td>NA</td>
<td>11.80</td>
</tr>
</tbody>
</table>

CAT, cataract; CO, corneal opacity; IPS, iris posterior synechia; MC, microcornea; MD, macular dystrophy; NA, not available; NRO, no response to object; OA, optic atrophy; OD, right eye; OP, occlusion of pupil; OS, left eye; PVF, poor vision fixation; RD, retinal detachment; RFM, retrolentiform fibrotic mass; TD, temporal dragging of optic disc.
posterior synchia of iris, or pupil occlusion (Table 2). After the molecular results were obtained, the four families were re-contacted. Three of the four probands showed systemic anomalies, as determined via telephone surveys, and the remaining proband was lost to follow-up. The proband QT2648, who is currently 15 months of age, had scoliosis and experienced difficulty in controlling her head. She was also unable to speak, and a cranial computed tomography scan taken when she was 5 months of age showed a local osteolysis lesion on the left parietal, as determined by a review of her medical records. Both proband QT1615 and QT1504 are currently 4 years of age and exhibit autism and learning disabilities. QT1615 showed microcephaly, with a head circumference of 40 cm, when she was 9 months of age.

Since mutations in CTNNB1 were identified in patients with intellectual disabilities in 2012, 2 45 mutations have been identified in 56 families with variable diagnoses and phenotypes (Supplementary Table S2).2–15,24–27 The mutation types and locations in the 56 families with or without eye abnormalities are shown in Figure 2.

DISCUSSION
In this study, four truncating mutations in CTNNB1 were identified in four patients with syndromic FEVR or Norrie disease but not in patients with other eye diseases. These results provide additional evidence for an association of truncating mutations in CTNNB1 with FEVR or Norrie-like retinopathy.

Nonsense and frameshift mutations in CTNNB1 are associated with FEVR because of several points. First, the same type of mutations in CTNNB1 were identified in four unrelated patients with the same class of diseases, namely, FEVR or Norrie-like retinopathy. Second, neither nonsense nor frameshift mutations in CTNNB1 were detected in more than 3000 in-house control individuals in our laboratory or more than 60,000 individuals in the ExAC database. Third, three of the four mutations are de novo, and fourth, all four mutations in CTNNB1 identified in the current study are LoF mutations, which is consistent with the previous finding that most (92.7%, 38/41) mutations in the gene are LoF mutations (Supplementary Table S2). Moreover, this type of mutation results in the haploinsufficiency of CTNNB1 by nonsense-mediated mRNA decay and a conditional knockout mouse model of cttnb1 established in previous studies showed specific vascular

unrelated patients with the same class of diseases, namely, FEVR or Norrie-like retinopathy. Second, neither nonsense nor frameshift mutations in CTNNB1 were detected in more than 3000 in-house control individuals in our laboratory or more than 60,000 individuals in the ExAC database. Third, three of the four mutations are de novo, and fourth, all four mutations in CTNNB1 identified in the current study are LoF mutations, which is consistent with the previous finding that most (92.7%, 38/41) mutations in the gene are LoF mutations (Supplementary Table S2). Moreover, this type of mutation results in the haploinsufficiency of CTNNB1 by nonsense-mediated mRNA decay and a conditional knockout mouse model of cttnb1 established in previous studies showed specific vascular
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In conclusion, we identified four heterozygous truncating mutations in CTNNB1 in patients with syndromic FEVR or Norrie disease. Our results provide additional evidence for an association between CTNNB1 haploinsufficiency and autosomal dominant syndromic FEVR or Norrie disease. Furthermore, mutations in CTNNB1, KIF11, or NDF might cause similar or overlapping phenotypes, but the mechanism needs to be studied further.

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References


