Mutation Spectrum of the \textit{LRP5}, \textit{NDP}, and \textit{TSPAN12} Genes in Chinese Patients With Familial Exudative Vitreoretinopathy

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\textbf{Purpose.} \textit{LRP5}, \textit{NDP}, and \textit{TSPAN12} are known to be associated with familial exudative vitreoretinopathy (FEVR). In this study, a comprehensive mutation screening for the three genes was performed in patients with a clinical diagnosis of FEVR in Han Chinese.

\textbf{Methods.} Genomic DNA and clinical data were collected from 100 probands and their family members. Sanger sequencing was performed to screen for \textit{LRP5}, \textit{NDP}, and \textit{TSPAN12} mutations and phenotype-genotype correlation was analyzed.

\textbf{Results.} There were 23 causative mutations identified in 23 unrelated probands (10/23 in \textit{LRP5}, 8/23 in \textit{TSPAN12}, and 5/23 in \textit{NDP}). Apart from \textit{NDP} mutations, only two \textit{LRP5} mutations inherited in an autosomal recessive manner. Among the 23 causative mutations, 13 were novel variants (4/10 in \textit{LRP5}, 6/8 in \textit{TSPAN12}, and 3/5 in \textit{NDP}). According to the modified classification system, statistical significance was observed in the distribution of mutated genes ($P = 0.049$). None of the causative mutations was found in group I FEVR. Proband with \textit{LRP5} or \textit{NDP} mutations were mainly categorized into group III and IV, \textit{TSPAN12} mutations were mainly observed in probands with group IV and V FEVR.

\textbf{Conclusions.} The detection rate for mutations in the three known genes was 23%. Mutations in \textit{LRP5} and \textit{TSPAN12} were more frequent, accounting for 10% and 8%, respectively. The \textit{NDP} mutations were only identified in 6% in this cohort. There were 13 novel variants found, which provided a deeper understanding of this disease. Potential phenotype-genotype correlation was observed in the modified system. \textit{TSPAN12} mutations might lead to the most severe phenotype.

Keywords: familial exudative vitreoretinopathy, mutational analysis, phenotype-genotype correlation

\section*{Familial exudative vitreoretinopathy (FEVR) is a genetic disorder affecting the development of retinal vasculature. The primary feature in FEVR is referred to as a premature arrest of vascularization in the peripheral retina. The subsequent ischemia and hypoxia can induce neovascularization and variable secondary complications, including vascular leakage, fibrovascular proliferation, traction of the posterior pole structure, macular dragging, retinal fold, retinal tears, and partial or total retinal detachment. The severity of clinical manifestation ranges from the absence of visual impairment to total blindness.

FEVR is genetically heterogeneous and has been reported to have autosomal dominant (adFEVR; Mendelian Inheritance in Man [MIM# 133780]), autosomal recessive (arFEVR; MIM# 133780), and X-linked recessive (XL-FEVR; MIM# 305390) inheritances. To date, mutations in the \textit{FZD4} (MIM# 604579)\textsuperscript{1} have been shown to contribute to the dominant FEVR. The \textit{TSPAN12} (MIM# 613138) and \textit{LRP5} (MIM# 605056) mutations are believed to be inherited as dominant or recessive trait\textsuperscript{2,5} while mutations in the \textit{NDP} (MIM# 300658) gene have been found in X-linked FEVR.\textsuperscript{6} Moreover, mutations in the \textit{ZNF408} (MIM# 616468),\textsuperscript{7} \textit{KIF11},\textsuperscript{8} \textit{RCBTB1},\textsuperscript{9} and \textit{CTNB1} (MIM# 617572)\textsuperscript{10} have been linked to FEVR patients in recent studies.

The proteins, encoded by the \textit{FZD4}, \textit{LRP5}, \textit{NDP}, and \textit{TSPAN12} genes, are involved in the Wnt/Norrin-$\beta$-catenin signaling pathways that are critically required for retinal angiogenesis during eye development.\textsuperscript{11,12} As an atypical Wnt ligand, Norrin activates the canonical Wnt pathway through its specific binding to the Frizzled4 receptor and Lrp5/6 coreceptor,\textsuperscript{13} which is selectively enhanced by Tetraspanin 12.\textsuperscript{12,13} It produces a downstream signal that stabilizes the cytosolic $\beta$-catenin. The increasing $\beta$-catenin translocates into the nucleus and interacts with the lymphoid enhancer factor/T-cell factor (LEF/TCF) transcription factor, which induces the transcription of target genes.\textsuperscript{14}

In this study, sequence analysis of \textit{LRP5}, \textit{NDP}, and \textit{TSPAN12} was performed in 100 unrelated Chinese individuals with clinical diagnosis of FEVR and their families. A comprehensive mutation spectrum for \textit{LRP5}, \textit{NDP}, and \textit{TSPAN12} in China was determined.

\section*{Methods}

\section*{Patients}

Written informed consent in accordance with the guidelines of the Declaration of Helsinki was obtained from the participating individuals or their guardians before the clinical data and DNA
samples were collected. All procedures were approved by the internal review board of the Zhongshan Ophthalmic Center, Sun Yat-sen University. There were 100 probands recruited and had a final diagnosis of FEVR. Patients with a gestational age of less than 38 weeks or a neonatal birth weight of less than 2000 g were excluded, to eliminate the possible presence of retinopathy of prematurity. The clinical diagnosis of FEVR was described in a previous study and "FZD4" mutations were found in 21/100 individuals.15 There were 70 ethnically matched individuals without history of ocular diseases recruited as controls. Comprehensive fundus examinations were performed, and no signs of retinal diseases were observed.

Genetic Analysis

Tissue samples (blood or buccal swabs) were obtained from each participant for genomic DNA extraction using the standard protocols of the QIAGEN QIAamp DNA Blood Kit (Qiagen, Hilden, Germany) and the Epicenter Master Pure Complete DNA and RNA Purification Kit (illumina, madison, wi, USA). For mutational analysis, the coding exons of "LRP5", "NDP", and "TSPAN12", and their flanking intronic sequences were amplified with PCR. The resulting amplicons were analyzed with Sanger sequencing. Direct sequencing was performed with an ABI BigDye Terminator Cycle Sequencing Kit using an ABI PRISM 3100 Genetic Analyzer (Applied Biosystems, Foster City, CA, USA). The reference sequences of "LRP5" (GenBank NG_015853.1 for gDNA, NM_001291902.1 for mRNA), "NDP" (GenBank NG_009852.1 for gDNA, NM_000266.3 for mRNA), and "TSPAN12" (GenBank NG_023203.1 for gDNA, NM_012538.5 for mRNA) were used for the identification of variants.

In Silico Analyses

Protein alignments for novel missense mutations were performed using the online COBALT program. Four online algorithms were used to estimate the pathogenicity of the missense changes, including SIFT, PolyPhen-2, Align GVGD, and Mutation Taster. For splice site mutations, another four online programs were used to predict their potential outcome, including Human Splicing Finder, NNSPLICE, FSPlice, and NetGene2. For secondary structural analysis of mutant protein, PsiPred v.3.3 was used.

Cloning Sequencing

Heterozygous frameshift mutations were confirmed by cloning sequencing. The target fragments covering the mutation sites were amplified with PCR. The PCR products were gel purified by using the Universal DNA Purification Kit (TIANGEN, Beijing, China) and cloned into pMD19-T (Takara BIO, Shiga, Japan) according to the manufacturer's instructions. Escherichia coli DH5α was transformed with the resulting plasmids and cultured in Luria-Bertani (LB) agar plates with ampicillin at 37°C for 12 to 14 hours. A single colony was placed in 5 mL of liquid LB medium with ampicillin and grown overnight in the 37°C incubator shaker for 12 to 14 hours. The plasmids were isolated from the suspension with the TIANprep Mini Plasmid Kit (TIANGEN), and Sanger sequencing was used to confirm the mutant and wild-type alleles.

Statistics

All analyses were performed using SPSS software version 16.0 (IBM Corp., Armonk, NY, USA). The independent samples t-test was used for normally distributed continuous variables, while the Kruskal–Wallis test was used for the nonnormally distributed continuous variables. A P value of less than 0.05 was considered statistically significant.

Results

In this study, 23 potentially pathogenic mutations were identified in 23 unrelated FEVR probands, including one patient harboring compound heterozygous mutations in "LRP5" (c.[1058G>A];[92-2A>C]) and two unrelated individuals carried the same reported "LRP5" mutation (c.4488+1T>G). Apart from "NDP" mutations, only two "LRP5" mutations inherited as an autosomal recessive trait. The remaining nucleotide changes in "LRP5" and "TSPAN12" were heterozygous and inherited as autosomal dominant pattern (Table 1). Of the 23 probands with "LRP5", "NDP", and "TSPAN12" mutations, nine had a family history of FEVR and 14 were isolated cases (Fig. 1).

Among these 23 variants, 13 of them were novel (4/10 in "LRP5", 6/8 in "TSPAN12", and 5/5 in "NDP"). None of these novel mutations was detected in 70 ethnically matched control individuals. These variants were not found in current online databases of human sequence variants, including the Ensembl, dbSNP ExAC databases, gnomeAD, and The Human Gene Mutation Database (HGMD). Chromatograms of these novel variants and clone sequencing of heterozygous insertion and deletion mutations are shown in Figure 2.

Apart from these diseases causing mutations, 20 synonymous variants in "LRP5" and "TSPAN12" were excluded from this study (data not shown). Six known variants in "LRP5" were considered as benign because the minor allele frequency was higher than 0.1% in the East Asian population (Supplementary Table S1). Besides that, three variants were considered as uncertain significance (Supplementary Table S2). Novel variant c.286-1G>T was located in the splice acceptor site of exon 4 in "TSPAN12". It was predicted to damage the natural acceptor splice site by three algorithms (NNSPLICE, FSPlice, and NetGene2). The 4-year-old male proband showed a retinal fold in his right eye, however, the same variant was found in his healthy father. The c.1512G>T (p.[W504C]) variant in "LRP5" previously identified in an osteoporosis-pseudoglioma syndrome (OPPG) family with compound heterozygous mutations (p.[W504C];[W478R]) and members with a single p.(W504C) variants only exhibited low bone mineral density.16 The heterozygous p.(W504C) was detected in our index patient with an ectopic macula and her healthy mother. Additionally, the splice site variant c.3637+4C>T (rs746451473), located in the splice donor site of exon 16 of "LRP5", was found in a sporadic male patient. Three of four algorithms considered it as benign.

Mutations Detected in "LRP5"

Three novel heterozygous missense mutations were found. The c.1318A>T (p.[I440F]) was found in a 9-month-old girl with retinal fold in both eyes, her mother and elder sister carried the same mutation and were diagnosed as FEVR. The isoleucine at codon 440 was conserved among various species (Fig. 3A) and three variants were considered as benign because the minor allele frequency was higher than 0.1% in the East Asian population (Supplementary Table S1). Besides that, three variants were considered as uncertain significance (Supplementary Table S2). Novel variant c.286-1G>T was located in the splice acceptor site of exon 4 in "TSPAN12". It was predicted to damage the natural acceptor splice site by three algorithms (NNSPLICE, FSPlice, and NetGene2). The 4-year-old male proband showed a retinal fold in his right eye, however, the same variant was found in his healthy father. The c.1512G>T (p.[W504C]) variant in "LRP5" previously identified in an osteoporosis-pseudoglioma syndrome (OPPG) family with compound heterozygous mutations (p.[W504C];[W478R]) and members with a single p.(W504C) variants only exhibited low bone mineral density.16 The heterozygous p.(W504C) was detected in our index patient with an ectopic macula and her healthy mother. Additionally, the splice site variant c.3637+4C>T (rs746451473), located in the splice donor site of exon 16 of "LRP5", was found in a sporadic male patient. Three of four algorithms considered it as benign.

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**WE**, without effect; LNSS, loss of the natural splice site; NPSS, new predicted splice site; Hetero, heterozygous; Homo, homozygous; Hemi, hemizygous; NA, not available; D, deleterious; T, tolerated.

* One patient harbored compound heterozygous mutations in LRP5.
† Two unrelated probands carried the same mutation.

Table 1. Causative Mutations Identified in 100 Proband`s With Familial Exudative Vitreoretinopathy

The hydrophilic cysteine was replaced by hydrophobic phenylalanine. The cysteine at codon 913 was highly conserved from human to various species (Fig. 3C) and the substitution was predicted as pathogenic by all algorithms.

The heterozygous change c.1582G>A (p.(R353Q)) was found in a 7-month-old boy with a retinal fold in the left eye and in his affected father with peripheral vascular deficiencies detected by fundus fluorescein angiography. The c.4733C>T (p.(T1578M)) variant was identified in a 16-year-old boy with a retinal fold and his relatives were not available for both clinical and genetic examinations; however, no family history was reported. The hydrophilic threonine was highly conserved and was substituted with hydrophobic methionine. All four algorithms predicted these as deleterious substitutions. The allele frequency of these heterozygous variants was below 0.1% and they have never been reported in FEVR patients before.

Two *LRP5* mutations inherited an autosomal recessive pattern. The homozygous mutation c.1183G>T (p.(R395W)) was found in a 3-year-old boy. A falciform retinal fold was observed in his right eye. His healthy parents carried the same mutation in heterozygous state. The conserved arginine at codon 395 was replaced by hydrophobic tryptophan. Align GVGD and Mutation Taster predicted it as pathogenic. The compound heterozygous mutation (c.[1058G>A];[92-2A>C]) was found in a 4-month-old boy, who visited our clinic due to no response to light since birth. Bilateral retrolenticular fibrotic mass and total retinal detachment was observed. His healthy father carried the heterozygous c.92-2A>C mutation while healthy mother carried the heterozygous c.1058G>A mutation. The c.1058G>A (p.[R353Q]) was previously reported in patients with OPPG.17 No skeletal problem was reported in this family. The novel c.92-2A>C mutation located in the splice acceptor site of exon 2 and the most common outcome for a splice acceptor site mutation was a deletion of the following exon. Four algorithms predicted the loss of natural splice site.

Two recurrent mutations were found in our study. The splice site mutation c.4488+2T>G was found in two unrelated probands with retinal folds and segregated among their families. The c.4087G>C (p.[D1363H]) mutation previously reported in a Chinese family,18 was identified in a 3-year-old boy with falciform retinal fold in the right eye. His healthy parents did not carry the same mutation.

Mutations Detected in TSPAN12

Three novel frameshift mutations were found. The single base pair insertion c.438_439insT caused a frameshifting resulting in 11 incorrect amino acids after codon 146 and a premature termination at codon 158 (p.[T147Yfs*12]). The proband was a 1-month-old girl with total retinal detachment and massive vitreous proliferation in her left eye and macular ectopic in her right eye. Her father with bilateral retinal folds carried the same
FIGURE 1. Schematic pedigrees of the families with causative mutations. Arrows indicate proband; open symbols, clinically unaffected; solid symbols, clinically affected; +, clinically evaluated; +/-, homozygous mutation; +/-, heterozygous mutation; --, wild type.
A novel mutation. A 1-bp deletion c.655delC was identified in a singleton. This novel mutation caused a frameshift resulting in four incorrect amino acids after codon 218, followed by premature termination at codon 223 (p.[Q219Nfs*5]). The female patient developed total retinal detachment in her left eye during early childhood, which resulted in no light perception. She visited our clinic at the age of 44 with decreased vision in the right eye, and a retinal fold with remarkable temporal exudates was observed. Her brother and father were reported to have the congenital cataract from childhood.

A novel stop loss mutation, c.916_918+3delTAAAAA, was identified in a 4-year-old boy with dragged macular in his right eye. His mutation-carrying mother manifested bilateral exotropia with characteristic peripheral avascular retina. The 6-bp deletion removed the termination codon (TAA) at the end of exon 8 and the extra three bases in the following intron (AAA). It was predicted to abolish the original stop codon 306 (*306E).
and resulted in an elongated protein with 34 additional amino acids at the C-terminus (p.*306Eext*35). To establish whether the stop loss mutation could affect the structure of Tetraspanin 12, PsiPred v.3.3 was used to analyze the secondary structure of wild-type and mutant protein. According to the prediction, two extra α-helices and one new β-sheet would be generated, hypothetically, the additional amino acids would alter the structure of Tetraspanin 12 (Fig. 4).

Two novel splice acceptor mutations were found. The c.150-1G>A mutation, located in the splice acceptor site of exon 4, was identified in a 10-year-old boy with bilateral retinal folds and his asymptomatic father. Four algorithms revealed the creation of a novel splice site, which would result in a one nucleotide deletion in the mRNA, and thereby would lead to a shift in the open reading frame and protein truncation. The proband showed asymmetric manifestation with worse visual acuity in his right eye due to retinal fold involving the macula. The second splice acceptor mutation, c.469-1G>A, was identified at the end of intron 6. All four online tools predicted the loss of natural splice site. The proband was a 5-year-old boy with bilateral retinal folds. His healthy parents did not carry the same mutation.

The novel c.2T>C (p.[M1T]) was identified in a 17-year-old male patient. The proband was noticed to have FEVR associated rhegmatogenous retinal detachment in his right eye, which pars plana vitrectomy was performed. Falciform retinal fold was observed in his left eye. His asymptomatic father exhibited with typical FEVR features of bilateral peripheral avascularization; however, no DNA sample was available for mutational analysis. The methionine at codon 1 was highly conserved from human to other species (Fig. 3D), and this mutation led to the replacement of a hydrophobic residue by a hydrophilic threonine residue. This change was predicted to be deleterious in all four online programs.

The missense mutation c.464G>C (p.[R155T]), was found in a patient with asymmetric manifestation. This 28-year-old male patient suffered total blindness in the right eye with retinal detachment during early childhood, while his left eye was asymptomatic. Peripheral retinal capillary nonperfusion was observed by fluorescein angiography. P.(R155T) was never reported in patients with FEVR.

Figure 3. Amino acid conservation of novel missense variations. (A) P.(I440F) in LRP5, (B) p.(V648I) in LRP5, (C) p.(C913F) in LRP5, (D) p.(M1T) in TSPAN12, (E) p.(H68P) in NDP, and (F) p.(H94L) in NDP.

Figure 4. The predicted secondary structure of wild type and elongated mutant TSPAN12 protein (c.-916+918+3delTAAAAA p.*306Eext*35)). (A) The predicted secondary structure of wild type TSPAN12 protein from codon 281 to 305. (B) The predicted secondary structure of mutant TSPAN12 protein from codon 281 to 339, two extra α-helices and one new β-sheet were generated (highlighted in red line).
reported as the causative mutation in FEVR and the minor allele frequency of this change was lower than 0.1%.

A recurrent mutation c.285+1G>A, located in the splice donor site of intron 5, was identified in a full-term infant, and cosegregated among his family. The boy visited our clinic 1 week after birth and fundus screening showed bilateral retinal detachment with pronounced peripheral exudation and hemorrhage. After the intravitreal injection of Ranibizumab in both eyes, the subretinal exudate and hemorrhage were absorbed, however a falciform retinal fold was presented in the left eye. This mutation was previously reported in a Nigerian female with autosomal recessive FEVR, who carried two heterozygous TSPAN12 mutations, c.[146C>T];c.285+1G>A.3

**Mutations Detected in NDP**

Two novel missense mutations were detected. The c.203A>C (p.[H68P]) mutation was found in a 35-year-old male with ectopic macular in the right eye. The hydrophilic histidine at codon 68 was conserved across majority of the species (Fig. 3E). This mutation changed the hydrophobic alkaline histidine into the hydrophobic proline residue. No family member was available for fundus examinations and molecular analysis. The c.281A>T (p.[H94L]) was identified in a 3-month-old male infant with bilateral retinal fold. The characteristic features of FEVR, including peripheral avascular zone and retinal exudates were observed in both eyes. No family member had complained about ocular symptoms and his healthy mother was heterozygous for this mutation. The histidine at codon 94 was highly conserved from human to various species (Fig. 3F), and this mutation led to the replacement of a hydrophilic alkaline residue by a hydrophobic leucine residue. Both p.[H68P] and p.[H94L] were predicted to be deleterious by two algorithms (Align GVGD and Mutation Taster).

The novel c.334delG mutation was found in a 6-month-old boy, who exhibited bilateral retinal detachment. The single base pair deletion was predicted to cause a frameshift after codon 112 and generate a prolonged protein at the C-terminus (p.[G113Afs*149]). His elder brother with tractional retinal detachment in both eyes carried the same mutation. His mother carried the same mutation in the heterozygous state, including 9 cases in group III and 29 in group IV. None of the probands with asymptomatic FEVR, in whom the diagnosis of FEVR was only confirmed by fundus fluorescein angiography, were categorized into group I. Patients with vitreous hemorrhage due to retinal neovascularization were categorized into group II. Patients staged into group I or II visited our clinic for the first time mainly in adolescent or adulthood. Patients with an ectopic macula were categorized into group III, while patients with retinal folding were categorized into group IV. Patients with severe total tractional retinal detachment at least in one eye were categorized into group V.

Combined with FZD4 mutations in these probands we reported before, most of the probands (86.4%, 38/44) with pathogenic variants were categorized into group III and IV, including 9 cases in group III and 29 in group IV. None of the FZD4, LRP5, TSPAN12, and NDP mutation was found in probands with group I FEVR, and only one proband was categorized into group II (1/21 in FZD4 mutations). Five patients were staged into group V (5/44). Among the five probands, three of them carried the TSPAN12 mutation (3/5) and one of them harbored the NDP variants (1/5), while only one patient with compound heterozygous mutation in LRP5 exhibited extremely severe FEVR (1/5). According to the modified system, statistical significance was observed in the distribution of four genes ($P = 0.049$) (Table 3).

**Clinical Finding in Patients With Causative Mutations**

There were 100 unrelated Chinese individuals with the clinical diagnosis of FEVR enrolled in this study. Potentially pathogenic mutations were found in 44 probands, including 10 patients with LRP5 mutations, 8 with TSPAN12, 5 with NDP mutations, and 21 patients with causative FZD4 mutations in our previous study.15 There were 44 probands with causative mutations staged according to the Trese’s staging system.24 In Trese’s system, $35 \ (35/44)$ patients with causative mutations in these four genes were staged into severe FEVR (stage 3–5), including 15 (15/21) in FZD4, 10 (10/10) in LRP5, 7 (7/8) with TSPAN12, and 3 (3/5) with NDP mutations. However, statistical significance was not established in this classification system ($P = 0.411$ (Table 2).

After analyzing the clinical features of the 100 probands, a modified approach based on the Trese’s staging system was addressed. Probands with asymptomatic FEVR, in whom the diagnosis of FEVR was only confirmed by fundus fluorescein angiography, were categorized into group I. Patients with vitreous hemorrhage due to retinal neovascularization were categorized into group II. Patients staged into group I or II visited our clinic for the first time mainly in adolescent or adulthood. Patients with an ectopic macula were categorized into group III, while patients with retinal folding were categorized into group IV. Patients with severe total tractional retinal detachment at least in one eye were categorized into group V.

**Discussion**

After screening the coding and flanking regions of $LRP5$, $NDP$, and $TSPAN12$ for mutations, 23 pathogenic mutations were found in 25 unrelated FEV probands, considering our previous study that 14 causative mutations in FZD4 was found in 21 unrelated individuals,15 17 causative mutations in FZD4, LRP5, NDP, and TSPAN12 were identified in 44 unrelated probands. The detective rate of four known genes was $44\% \ (44/100)$. FZD4 mutations accounted for the largest proportion (21/44), followed by LRP5 mutations (10/44), TSPAN12 mutations (8/44), and NDP mutations (5/44) (Fig. 5).

This study was one of the largest cohorts of FEVR in Chinese populations and provided a comprehensive mutation-
TABLE 3. Phenotype-Genotype Analysis in Modified Classification System

<table>
<thead>
<tr>
<th>Group</th>
<th>N</th>
<th>FZD4</th>
<th>LRP5</th>
<th>TSPAN12</th>
<th>NDP</th>
<th>N With Causative Mutation (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td>10</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0 (0)</td>
</tr>
<tr>
<td>II</td>
<td>21</td>
<td>5</td>
<td>1</td>
<td>1</td>
<td>2</td>
<td>1 (25.0)</td>
</tr>
<tr>
<td>III</td>
<td>48</td>
<td>15</td>
<td>8</td>
<td>4</td>
<td>2</td>
<td>29 (60.4)</td>
</tr>
<tr>
<td>IV</td>
<td>17</td>
<td>0</td>
<td>1</td>
<td>5</td>
<td>1</td>
<td>5 (29.4)</td>
</tr>
<tr>
<td>V</td>
<td>102</td>
<td>21</td>
<td>10</td>
<td>8</td>
<td>5</td>
<td>44 (44)</td>
</tr>
</tbody>
</table>

*p Patients were staged in the basis of the highest stage of FEVR in either eye.

al screening of four confirmed genes that caused FEVR. Compared with the previous study of a large cohort of 98 FEVR patients in Caucasian persons, the detection rates of causative mutations in Wnt/Norrin signal pathway were similar with the previous study, accounting for 46% (42/92). However, the involvement of each gene was different. LRP5 was the most frequently mutated gene in Caucasian population (19.6%), and FZD4 mutations accounted for 15.2%. The reported mutation frequencies in the LRP5, TSPAN12, and NDP gene varied in different populations. The frequency of LRP5 mutations ranged from 10% to 22.2%, while the frequency of NDP mutations ranges from 6.5% to 12.5%. The frequency of TSPAN12 mutations identified 2% to 10% in patients with autosomal dominant or autosomal recessive FEVR, while the frequency of NDP mutations ranges from 6.5% to 12.5%. The reported mutation frequencies in the LRP5, TSPAN12, and NDP gene varied in different populations. The frequency of LRP5 mutations ranged from 10% to 22.2%, while the frequency of NDP mutations ranges from 6.5% to 12.5%.

In summary, this study provided a comprehensive understanding of the mutation spectrum of FEVR-associated genes of FEVR in Wnt/Norrin pathway. Mutation screening of known genes would provide invaluable information for diagnosis and genetic counseling, especially for patients without family history or with atypical presentations. Further analysis of novel causative genes may explain some mild clinical findings such as group I FEVR patients, and definitely will contribute to our understanding of the molecular mechanisms of retinal angiogenesis, which is crucial for FEVR and other retinal vascular disease.

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