SHORT REPORT

Homozygosity mapping and whole-genome sequencing reveals a deep intronic *PROM1* mutation causing cone–rod dystrophy by pseudoexon activation

Anja K Mayer¹, Klaus Rohrschneider², Tim M Strom³, Nicola Glöckle⁴, Susanne Kohl¹, Bernd Wissinger¹ and Nicole Weisschuh^{*,1}

Several genes have been implicated in the autosomal recessive form of cone–rod dystrophy (CRD), but the majority of cases remain unsolved. We identified a homozygous interval comprising two known genes associated with the autosomal recessive form of CRD, namely *RAB28* and *PROM1*, in a consanguineous family with clinical evidence of CRD. Both genes proved to be mutation negative upon sequencing of exons and canonical splice sites but whole-genome sequencing revealed a private variant located deep in intron 18 of *PROM1*. *In silico* and functional analyses of this variant using minigenes as splicing reporters revealed the integration of a pseudoexon in the mutant transcript, thereby leading to a premature termination codon and presumably resulting in a functional null allele. This is the first report of a deep intronic variant that acts as a splicing mutation in *PROM1*. The detection of such variants escapes the exon-focused techniques typically used in genetic analyses. Sequencing the entire genomic regions of known disease genes might identify more causal mutations in the autosomal recessive form of CRD.

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INTRODUCTION

Cone–rod dystrophy (CRD; MIM 120970) has an estimated prevalence of 1:30 000 to 1:40 000 and is characterized by primary loss of cone function and subsequent or simultaneous loss of rod vision.¹ Key symptoms are poor visual acuity, dyschromatopsia and photophobia.

CRDs are genetically heterogeneous and can show different modes of Mendelian inheritance, with autosomal recessive (ar) traits being more common than autosomal dominant and X-linked forms. Currently, 17 genes have been implicated in arCRD, with *ABCA4* (MIM 601691) being the major contributing gene, whereas each of the other known genes, among them *PROM1* (MIM 612657), account for only a minor fraction of cases.² About 75% of arCRD cases apparently do not harbor mutations in these known genes.²

Here we report on a consanguineous family with CRD. The two affected siblings were diagnosed with CRD in the second decade of life, with rapidly deteriorating visual acuities. As the two affected siblings were born out of a first-cousin marriage, we expected the disease-causing mutation to be located in a homozygous interval.³ To that end, genetic analyses started with the identification of these, followed by a panel-based next-generation sequencing approach and finally whole-genome sequencing.

PATIENTS AND METHODS

Members of the family originated from southwest Germany and were recruited in accordance with the principles of the Declaration of

Helsinki, with informed consent and ethical approval. DNA was isolated from venous blood samples using conventional techniques. Using Affymetrix CytoScan HD Arrays (Affymetrix, Santa Clara, CA, USA), regions of homozygosity were calculated with the online tool Homozygosity Mapper (<http://www.homozygositymapper.org/>).⁴ Panel-based sequencing of 105 retinal degeneration-associated genes and whole-genome sequencing, respectively, were performed as described before.^{5,6} In-house automated data analysis pipeline and variant interpretation tools were used for variant calling.

A 2.7-kb fragment comprising *PROM1* exon 18 with 239 bp of upstream sequence, intron 18 and exon 19 with 367 bp of downstream sequence was amplified using a proofreading polymerase and genomic DNA from family member I:2 (see Figure 1) in order to co-amplify both the normal and the mutant allele. Cloning into the exon-trapping vector pSPL3, transfection of human embryonic kidney (HEK) 293T cells, RNA isolation and cDNA synthesis were performed as described previously.⁷ Transfection and downstream analyses of murine 661W photoreceptor-derived cells were performed likewise.

The identified variant was submitted to ClinVar (www.ncbi.nlm.nih.gov/clinvar/) with accession code SCV00022908.

RESULTS

Homozygosity mapping of the two affected siblings revealed a 7-Mb shared homozygous interval on chromosome 4 (rs12512447–rs7699941). The interval comprises 21 HGNC-approved protein-coding genes,

¹Molecular Genetics Laboratory, Centre for Ophthalmology, Institute for Ophthalmic Research, University of Tuebingen, Tuebingen, Germany; ²Department of Ophthalmology, University of Heidelberg, Heidelberg, Germany; ³Institute of Human Genetics, Helmholtz Zentrum Muenchen, Neuherberg, Germany; ⁴CeGaT GmbH, Tuebingen, Germany
*Correspondence: Dr N Weisschuh, Molecular Genetics Laboratory, Centre for Ophthalmology, Institute for Ophthalmic Research, University of Tuebingen, Roentgenweg 11, D-72076 Tuebingen, Germany. Tel: +49 7071 2987618; Fax: +49 7071 295725; E-mail: nicole.weisschuh@uni-tuebingen.de

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Figure 1 Pedigree of the CRD family and *in silico* splicing predictions for the wild-type and mutant genomic sequences of *PROM1*. (a) Pedigree showing segregation of the *PROM1* intronic variant c.2077-521A>G. (b) Splice-site prediction for the wild-type (top) and the mutant allele (bottom). The position of the c.2077-521A>G variant is indicated by the vertical dotted line. Donor site scores from NNSPLICE and HSF are displayed as green vertical bars. The prediction score is provided beside each bar.

among them two genes were known to be mutated in arCRD, namely *RAB28* and *PROM1*. However, subsequent Sanger sequencing for *RAB28* and targeted next-generation sequencing for *PROM1* ruled out variants that may affect function in the exons and the canonical splice sites of the two genes. The targeted next-generation sequencing generated sequence data for additional 104 retinal degeneration-associated genes,⁵ but no putative disease-causing variants were detected in these. Subsequently, we performed whole-genome sequencing of both the affected siblings, generating 159 and 179 Gb of sequence with an average read depth of 44 and 50, respectively. We found no rare and potentially disease-causing exonic variants compatible with a model of autosomal recessive inheritance and shared by both siblings. In a next step, raw read data of the genomic regions of *RAB28* and *PROM1* were manually checked with the Integrative Genomics Viewer.⁸ In comparison with the hg19 human reference sequence, we detected 18 single-nucleotide variants in *RAB28* and 196 in *PROM1*, which were shared by both siblings. After filtering out annotated SNPs (dbSNP 137 and 1000 Genomes), only one private variant remained that

was located deep in intron 18 of *PROM1* (c.2077-521A>G) and segregated with the disease in the expected autosomal recessive mode (Figure 1a). The variant designation is based on the NCBI reference sequence for *PROM1* transcript NM_006017.2 and the genomic reference sequence NG_011696.1. The exon numbering is based on NG_011696.1. Splice-site predictions using the online tools Human Splicing Finder (HSF; <http://www.umd.be/HSF/>) and NNSPLICE (http://www.fruitfly.org/seq_tools/splice.html) showed significantly increased 5' donor splice-site scores for the sequence surrounding the variant (NNSPLICE 0.99 out of 1; HSF 85.4 out of 100; compared with zero for the wild-type sequence in both prediction programs; see Figure 1b).

To test our hypothesis that this deep intronic variant acts as a splicing mutation, cDNA analyses were performed. Owing to the lack of *PROM1* expression in accessible tissues, we made use of a heterologous splicing assay in HEK 293T cells and in murine 661W photoreceptor-derived cells to test mutant and wild-type *PROM1* minigene constructs in direct comparison. RT-PCR of wild-type RNA resulted in a single product of the expected size (Figure 2a), and bidirectional Sanger sequencing confirmed correct splicing of exons 18 and 19 (Figure 2b). In contrast, RT-PCR of mutant RNA revealed a single transcript larger than the expected correctly spliced product (Figure 2a). Direct sequencing revealed an insertion of a 155-bp cryptic exon spliced between exons 18 and 19 (Figure 2b). The aberrant transcript would lead, if translated, to an insertion of 22 novel amino acids followed by a premature termination codon (Figure 2c). This would result in a truncated protein short of 173 amino acids at the C-terminus and thereby lacking the last of five transmembrane domains. However, it is more likely that the mutant transcript undergoes nonsense-mediated mRNA decay.⁹

DISCUSSION

The Human Gene Mutation Database (<http://www.biobase-international.com/product/hgmd>) lists only five *PROM1* mutations in arCRD that all lead to functional null alleles: two nonsense mutations,^{5,10} two mutations at canonical splice sites^{11,12} and one single-nucleotide duplication.^{13,14} To the best of our knowledge, this is the first report of a deep intronic variant in *PROM1* presumably leading to a functional null allele.

It is important to note that the disease-causing mutation in our family would not have been identified using conventional approaches like Sanger sequencing of coding exons or by using exome sequencing. Genomic analyses for other genes involved in retinal degeneration like *ABCA4*,^{15–17} *USH2A*,^{18,19} *CEP290*²⁰ and *OFD1*²¹ have shown that a probably underestimated number of patients harbor deep intronic variants that interfere with splicing. Although the majority of deep intronic variants that act as splicing mutations to date have been identified through the presence of aberrant mRNA transcripts, this approach is only feasible for those genes that are expressed in accessible tissues (ie, lymphocytes, keratinocytes and fibroblasts). Moreover, this approach might be hampered by the degradation of mutant transcripts through nonsense-mediated mRNA decay. These limitations can be overcome by whole-genome sequencing and the use of heterologous splicing assays. Our findings substantiate the necessity of the analysis of regions outside of the coding exons, especially in

Figure 2 *In vitro* splicing assay. As no difference in splicing effect was observed between the two cell lines, only results from HEK 293T cells are shown. (a) RT-PCR revealed an aberrant, larger transcript in HEK 293T cells transfected with the mutant construct compared with the wild-type construct. NTC, nontemplate control. Size standard: NEB 100 bp ladder. Schemes of the amplified products are presented on the right of the agarose gel. White boxes represent pSPL3 exons and gray boxes *PROM1* exons. (b) Sequence analysis showing that the aberrant larger transcript is due to the insertion of a cryptic exon of 155 bp between exons 18 and 19. (c) Consequence of the c.2077-521A>G variant (underlined). The cryptic exon (in capital letters) translates into 22 novel amino acids followed by a premature stop codon (protein translation is shown below DNA sequence in one letter amino-acid code).

families with multiple affecteds and evidence of linkage to known disease-gene loci.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

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