Allele-specific editing ameliorates dominant retinitis pigmentosa in a transgenic mouse model

Clarissa Patrizi,1 Manel Llado,2 Daniela Benati,1 Carolina Iodice,2 Elena Marrocco,2 Rosellina Guarascio,3 Enrico M. Surae,2,5 Michael E. Cheetham,3 Alberto Auricchio,2,4,* and Alessandra Recchia1,*

Summary

Retinitis pigmentosa (RP) is a group of progressive retinal degenerations of mostly monogenic inheritance, which cause blindness in about 1:3,500 individuals worldwide. Heterozygous variants in the rhodopsin (RHO) gene are the most common cause of autosomal dominant RP (adRP). Among these, missense variants at C-terminal proline 347, such as p.Pro347Ser, cause severe adRP recurrently in European affected individuals. Here, for the first time, we use CRISPR/Cas9 to selectively target the p.Pro347Ser variant while preserving the wild-type RHO allele in vitro and in a mouse model of adRP. Detailed in vitro, genomic, and biochemical characterization of the rhodopsin C-terminal editing demonstrates a safe downregulation of p.Pro347Ser expression leading to partial recovery of photoreceptor function in a transgenic mouse model treated with adeno-associated viral vectors. This study supports the safety and efficacy of CRISPR/Cas9-mediated allele-specific editing and paves the way for a permanent and precise correction of heterozygous variants in dominantly inherited retinal diseases.

Introduction

Retinitis pigmentosa (RP) is a group of genetically heterogeneous retinal diseases afflicting three million people across the globe with an incidence of 1 in 3,500 live births.1–4 Individuals affected by RP initially manifest night-blindness with gradual constriction of the visual field but sparing of central vision. As the rod loss progresses, secondary death of cones occurs, leading to deterioration of visual acuity and eventual blindness. Part of the difficulty in treating RP is its complex and diverse genetic etiology. To date, more than 3,000 disease-associated variants in approximately 70 disease-causing genes have been causally associated with RP5 and currently more than 150 documented missense/nonsense variants in the rhodopsin (RHO) gene are associated with an autosomal dominant RP (adRP) phenotype (RP4 [MIM: 613731]).6,7 The RHO gene encodes for rhodopsin (RHO), a visual pigment found in rod photoreceptors, responsible for converting photons into chemical signals initiating vision. Rhodopsin is a 348 amino acid G-protein-coupled receptor characterized by an extracellular N-terminal domain needed to stabilize the protein, seven transmembrane-spanning α helices hosting the binding site for the chromophore 11-cis-retinal, and an intracellular C-terminal domain, involved in vectorial transport of rhodopsin to rod outer segments (OSs).8–11 Although approximately half of the RHO-associated adRP cases in the US are caused by the substitution of proline to histidine at position 23 (p.Pro23His)12 in the extracellular N-terminal domain, class I variants clustered in the C-terminal domain13 give rise to a defect in post-Golgi trafficking to the OS and result in a more severe phenotype and worse prognosis for affected individuals.13–15 The large majority of RHO pathogenic variants are inherited in an autosomal dominant manner. In most of these cases, simply adding a normal copy of the gene is not sufficient,16 as the affected gene needs to be inactivated. Downregulation of RHO variants has been attempted in disease models using ribozymes17, RNA interference16,18 and transcriptional repressor by zinc finger proteins.19,20 Most of these approaches do not distinguish the disease-associated alleles from the wild-type (WT), thus achieving bi-allelic suppression that also requires addition of a WT RHO cDNA (“suppression and replacement”). The ability to correct disease-causing variants while sparing the WT allele has been improved greatly by the discovery of CRISPR/Cas9 genome editing.21 Cas9 endonucleases generate double-strand breaks (DSBs) in a specific genomic region that is located adjacent to a protospacer-adjacent motif (PAM) and targeted by a complementary guide RNA (gRNA).22 In the absence of the exogenous template, the Cas9-induced DSBs are repaired through the non-homologous-end-joining (NHEJ) mechanism, leading to the frequent introduction of insertions or deletions in the target site. Thereby, as a valid alternative to the “suppression and replacement” approach that may be potentially used to treat a wide array of dominant diseases but that requires a double intervention, specific inactivation of the altered allele can be pursued for dominant-negative and gain-of-function variants23 that generate a unique PAM site or allow the design of a gRNA that contains the variant in the seed sequence.

1Centre for Regenerative Medicine, Department of Life Sciences, University of Modena and Reggio Emilia, 41125 Modena, Italy; 2Telethon Institute of Genetics and Medicine, 80078 Pozzuoli, Italy; 3UCL Institute of Ophthalmology, London EC1V 9EL, UK; 4Medical Genetics, Department of Advanced Biomedicine, Federico II University, 80125 Naples, Italy; 5Medical Genetics, Department of Translational Medicine, Federico II University, 80125 Naples, Italy
*Correspondence: alessandra.recchia@unimore.it (A.R.), auricchio@tigem.it (A.A.)
© 2021 The Authors. This is an open access article under the CC BY license (http://creativecommons.org/licenses/by/4.0/).
Given the high prevalence of the c.68C>A RHO allele encoding the p.Pro23His variant in the United States, it is not surprising that this has been the primary target of CRISPR/Cas9-mediated gene editing. Indeed, this strategy has already been demonstrated to be effective in recent studies employing the Pro23His knockin mouse model. In these reports, the authors showed a reduced expression of the disease-associated murine transcript triggered by NHEJ repair occurring in the first exon of the gene. The allele-specific inactivation of the murine allele encoding the p.Pro23His variant resulted in a delay of the degenerative retinal process and rescue of retinal functional activity.

A gene editing approach tailored to the C-terminal domain of human rhodopsin and, in particular, to proline 347, the most common residue affected in European individuals, has been neglected so far. Here, for the first time, we employ both Streptococcus pyogenes Cas9 (SpCas9) WT and the high-fidelity variant carrying seven amino acid substitutions, Asn497Ala, Arg661Ala, Gln695Ala, Gln926Ala, Asp1135Val, Arg1335Gln, and Thr1337Arg (hereafter referred to as the fidelity variant carrying seven amino acid substitutions), to generate the p.C347SseR variant in the transgenic Pro347Ser RHO variant. We characterize in detail the c.1039C>T variant in RHO, which leads to the p.C347Ser RHO variant. We characterize in detail the c.1039C>T RHO allele-specific editing and the predicted genomewide-off-target sites by next-generation sequencing (NGS). Considering the role of the RHO C terminus in protein trafficking/folding and the unpredictable editing occurring at the target site, we have performed in-depth biochemical analyses of the most frequent RHO variants generated upon CRISPR/Cas9-mediated editing. Moreover, subretinal delivery of adeno-associated virus (AAV) vector serotype 2/8 (AAV2/8) carrying the WT or VQRHF1 SpCas9 and the target or scramble gRNA demonstrates the therapeutic potential of AAV-Cas9 gene editing to inactivate the human p.Pro347Ser pathogenic variant in the transgenic Pro347Ser mouse model, ameliorating disease progression.

Material and methods

Plasmids
To generate the pCCL.PGK.wtRHO+3’UTR, a 250 bp region of the RHO 3’ UTR was amplified from the genomic DNA of Pro23His transgenic mice and cloned into pCCL.PGK.wtRHO downstream of the stop codon of the wtRHO cDNA. To generate the pCCL.P347S.RHO+3’UTR, the region including the exon 5 of RHO cDNA, carrying the p.Pro347Ser variant, and a 250 bp region of the RHO 3’ UTR amplified from the genomic DNA of Pro347Ser transgenic mouse were cloned into the pCCL-PGK.wtRHO backbone downstream of the exon 4 of RHO cDNA. The effector plasmid SpCas9 gRNA1 was generated by cloning the gRNA1 in the pX330-U6-Chimeric_BB-3Bb-hSpCas9 plasmid (Addgene: 42230) by oligo annealing into BbsI sites. The effector plasmid VQRHF1-SpCas9 gRNAs was obtained by cloning the gRNAs in the pX330-U6-Chimeric_BB-3Bb-hSpCas9 plasmid by oligo annealing into BbsI sites, followed by subcloning of the U6-gRNAs cassette into the MSP2440 plasmid (Addgene: 72250) expressing the VQRHF1-SpCas9. To generate effector plasmids carrying Hygromycin resistance gene, the expression cassette for the Hygromycin resistance gene under the control of herpes simplex virus thymidine kinase (TK) promoter was subcloned in SpCas9_gRNA1 and VQRHF1-SpCas9_gRNAs downstream of the polyA signal of SpCas9 expression cassette, generating SpCas9_gRNA1-TKHygro and VQRHF1-SpCas9_gRNAs-TKHygro. To generate CMV.HA.wtRHO+3’UTR and CMV.HA.P347SRHO+3’UTR plasmids expressing RHO under the cytomegalovirus (CMV) promoter, the hRHO cDNA carrying the WT or p.Pro347Ser variant and the 250 bp region of RHO 3’ UTR were cloned into CMV.HA.RHO plasmid. RHO variant plasmids (CMV.HA.RHO delG/delGG/del12.1/InsT/del9/del12.5) were generated by using the Site-Directed Mutagenesis Kit (SDM) (NEB, New England Biolabs, Ipswich, MA, USA) according to manufacturer’s protocol. Primer pairs (Table S3) were designed for the incorporation of insertions or deletions into CMV.HA.P347S. RHO+3’UTR and pCCL.PGK.P347S.RHO+3’UTR plasmids. The pAAV2.1-U6-gRNA1-RHO-GFP, pAAV2.1-U6-gRNA5.RHO-GFP, and pAAV2.1-U6-scramble-RHO-GFP plasmids were generated by cloning the expression cassette for gRNA1, gRNAs, or scramble gRNA into a pAAV2.1-RHO-GFP plasmid upstream of the RHO promoter by Affib restriction. The pAAV2.1-IRBP.SpCas9-spA plasmid was generated by cloning the interphotoreceptor retinol binding protein (IRBP) promoter into pAAV-pMecp2-SpCas9-spA (Addgene: PX551) using HindIII and AgeI restriction enzymes. The pAAV2.1-IRBP-VQR-HF1.SpCas9 vector was generated by cloning the CMV/VQR-HF1.SpCas9-BGHPA cassette into the AAV2.1 backbone followed by replacing the CMV promoter with IRBP.

Cell culture
HeLa, CHO, HEK293T, and hTERT-RPE cells were obtained from the American Type Culture Collection (ATCC) and were cultured in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal calf serum (FCS), 100 U/mL penicillin, and 100 µg/mL streptomycin (Lonza, Basel, Switzerland). For protein degradation assay, CHO cells transfected with CMV-HA-Pro347Ser, WT, and variant RHO plasmids were treated with 10 µg/mL of cycloheximide (CHX) or MG-132 (50 µM) and analyzed after 0, 3, and 6 h or 0, 4, and 6 h, respectively.

Viral production
Lentiviral vectors (LVs) pseudotyped with the vesicular stomatitis virus G protein were prepared by transient co-transfection of HEK293T cells with transfer vector, pMD1.Lg/pPRE.Int, pMD2.G.YSV-G envelope-encoding packaging plasmid, and pRSV-Rev. AAV vectors were produced by triple transfection of HEK293 cells followed by two rounds of CaCl2 purification. For each viral preparation, physical titers (GC/mL) were determined by averaging the titer achieved by dot-blot analysis and by PCR quantification using TaqMan (Applied Biosystems, Carlsbad, CA, USA). The probes used for dot-blot and PCR analyses were designed to anneal with the IRBP promoter for the pAAV2.1-IRBP-SpCas9-spA vector and the BGHPA region for vectors encoding for gRNA expression cassettes and RHO-GFP. The length of probes varied between 200 and 700 bp.

Transfections of cells, isolation of single-cell clones, and vector copy number determination
Transfection of 2.5 × 10^5 HeLa cells was obtained using Fugene HD transfection reagent (Promega, Madison, WI, USA) following the manufacturer’s instructions. For each transfection reaction, 2 µg of plasmid DNA were mixed to 6 µL Fugene (3:1 ratio).
Transfection of 2.5 × 10^5 HEK293T cells was performed using CaPO4 protocol with 1 μg of SpCas9-gRNA1-TKHygro or VQRHF1-SpCas9_gRNA5-TKHygro. Starting from the day after transfection, cells were treated with 0.2 mg/mL of Hygromycin for 15 days to select antibiotic resistant cells.

Transfection of 1 × 10^5 hTERT-RPE cells was performed with 1 μg of SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5 or respective control plasmids using TransIT-XI (Mirus Bio, Madison, WI, USA) following the manufacturer’s instructions. Transfection of 2.5 × 10^5 CHO cells was obtained using TransIT-XI following the protocol instructions. Each transfection reaction contained 150 ng of plasmid coding for p.Pro347Ser RHO or wt RHO or the selected RHO variants obtained upon gene editing in Pro347Ser RHO HeLa clone. To obtain HeLa clones expressing WT or p.Pro347Ser RHO, HeLa cells were transduced with LVs carrying the WT or p.Pro347Ser RHO expression cassettes. Transduced bulks were limiting diluted to obtain a concentration of 0.3 cells/well and seeded in a 96-well plate. Genomic DNAs (gDNAs) were extracted from single cell clones, and a PCR on the RHO expression cassette was performed as follows: primers PGK_F and hRHO_ex1_R (Table S3); PCR conditions, 30 s at 94°C, 30 s at 58°C, and 30 s at 72°C for 30 cycles. PCR products were separated on 1% TBE (Tris/Borate/EDTA)-agarose gels and stained with ethidium bromide for analysis.

For vector copy number (VCN) determination, qPCR was conducted with 20 ng gDNA using TaqMan Universal PCR Master Mix (Applied Biosystem) and probes specific for human RHO and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) (hRHO, Hs00892431m1; hGAPDH, Hs03929097_g1; Applied Biosystems, Milan, Italy). Reactions were performed at 50°C for 2 min and 95°C for 10 min, followed by 40 cycles at 95°C for 15 s and 60°C for 1 min. Normalization to GAPDH in the same gDNA was performed and the relative copy number was calculated by using the 2−ΔΔCT quantification.

**Semiquantitative and quantitative RT-PCR analyses**

Total RNA from WT or Pro347Ser RHO HeLa clones, mice retinae, and hTERT-RPE cells was isolated with the RNeasy Mini and Micro kits (QIAGEN, Hilden, Germany) according to the manufacturer’s instructions. cDNA was synthesized in a 20 μL reaction by the Superscript III Reverse Transcription kit (Invitrogen). Semiquantitative RT-PCR analysis was performed with the following oligonucleotides: hRHO-Cterm_F and WPRE_R, GAPDH.F and GAPDH.R for mRNA analysis of HeLa cells, hRHO-Cterm_F and hRHO_3’ UTR_RC, Cas9.F and Cas9.R GPPF and GPPR, m.s26rRNA.F and m.s26rRNA.R, and PDE6b.F and PDE6b.R for mRNA analysis of treated retina mice. PCR cycles were as follows: 94°C for 30 s, 58°C for 30 s, 72°C for 30 s, and 72°C for 30 s. TaqMan real-time PCR analysis was performed using the ABI Prism 7900 Sequence Detection System (Applied Biosystems, Monza, Italy) with TaqMan Universal PCR Master Mix and probes specific for human RHO, human and mouse GAPDH, and mouse Pde6g (hRHO, Hs00892431m1; hGAPDH, NM_020463.3; mGAPDH, Mm99999915_g1; mPde6g, Mm00501964_m1; Applied Biosystems, Monza, Italy). Reactions were performed at 50°C for 2 min and 95°C for 10 min, followed by 40 cycles at 95°C for 15 s and 60°C for 1 min. The relative expression of the target genes was normalized to the level of GAPDH housekeeping gene for HeLa clones or Pde6g photoreceptor housekeeping gene in the same cDNA by using the 2−ΔΔCT quantification. The replicated relative quantity (RQ) values for each biological sample were averaged.

**Targeted deep sequencing and off-target analysis**

Genomic DNA was extracted from HeLa clones transfected with SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5, HEK293T cells transfected with SpCas9_gRNA1-TKHygro or VQRHF1-SpCas9_gRNA5-TKHygro and selected with Hygromycin, and mice retinae using QIAamp DNA Mini or Micro kits (QIAGEN) following the manufacturer’s instructions. For NGS analysis, the genomic regions flanking gRNA target sites were amplified by PCR using the Accuprime Taq DNA Polymerase System (Thermo Fisher) and primers in Table S3. PCR products were subjected to library preparations. Briefly, primers hRHO-Cterm_F and WPRE_R and primers hRHO-Cterm_F and hRHO_3’UTR_RC (primers 1rd PCR amplification) were, respectively, used to specifically amplify the p.Pro347Ser-coding RHO cDNA in HeLa stable clones and human p.Pro347Ser-coding RHO gene in transgenic mice. A second amplification with primers (primers 2nd PCR amplification) was required. For off-target analysis in Hygromycin-selected HEK293T cells, individual single pairs of primers were used (primers 1rd PCR amplification). For NGS library preparation, individual barcode was added to each DNA fragment by a limited number (n = 8) of PCR cycles using primers detailed as primers 3rd PCR amplification. Equimolar amounts of library were mixed, diluted, and sequenced with an Illumina MiSeq system by CIBIO Trento. The percentage of indels was quantified by the CRISPResso webtool.

The off-target analysis for each gRNA was performed by using the COSMID webtool. Off-targets analysis was performed by NGS and the percentage of indels was quantified by CRISPResso V2 webtool.

**Immunoblotting analysis**

Cell lysates were extracted with RIPA buffer: 50 mM Tris-HCl pH 8.0, 150 mM w/v NaCl, 1 mM EDTA, 1% w/v NP-40, 0.1% v/v SDS, and 0.05% w/v sodium deoxycholate in the presence of 2% of protease inhibitor cocktail (Roche, Basilea, Switzerland). Sixty μg of protein extracts from HeLa clones and 20 μg of protein extracts from transfected CHO cells were loaded on 12% sodium dodecyl sulfate (SDS)-polyacrylamide gel electrophoresis (PAGE). After electrophoresis, samples were transferred PVDF membranes (GE Healthcare). The membranes were incubated with monoclonal 4D2 primary antibody (1:500, Millipore, Burlington, MA, USA) or monoclonal anti-HA primary antibody (Sigma, 1:1,000) and anti-b-actin antibody (Abcam, Cambridge, UK) for protein loading normalization. Horse-radish-peroxidase-conjugated anti-mouse antibody (diluted 1:10,000 for RHO variant expression analysis, 1:5,000 for rhodopsin expression in HeLa clones) was used for chemiluminescent detection (Pierce). Quantification was performed by densitometry analysis of scanned images using ImageJ software.

**Immunofluorescence and “In-Cell Western” analysis**

For immunofluorescence, cells were fixed using 4% v/v paraformaldehyde (PFA) and permeabilized with 0.01% v/v Triton X-100/PBS, while non-permeabilized cells remained in PBS. Non-specific binding sites were blocked using blocking solution consisting of 3% of bovine serum albumin (BSA) and 10% of normal goat serum in PBS. Non-permeabilized cells were incubated with 4D2 primary antibody (1:500, Millipore) and anti-HA primary antibody (1:1,000, Sigma) in blocking solution; permeabilized cells were incubated with anti-HA primary antibody (1:500, Sigma) and binding immunoglobulin protein (BIP) (1:200, Sigma). As secondary antibodies, Alexafluor488-conjugated goat...
anti-Mouse (1:1,000) and Alexafluor594-conjugated goat anti-Rabbit (1:1,000) were used. After washing, incubation with 6-diamidino-2-phenylindole DAPI (1:5,000) was performed, then slides were mounted with Dako Fluorescent Mounting Medium (Dako Omnis). Immunofluorescence were visualized using Zeiss LSM 700 laser scanning confocal microscope and analyzed with ZEISS ZEN Microscope software.37

To perform "In-Cell Western" analysis, cells were fixed, treated with blocking solution, and stained with anti-HA antibody (1:1,000, Sigma). The secondary antibody IRDye 680RD goat anti-mouse (LI-COR Biosciences, Lincoln, NE, USA) was used. Acquisition was performed using the Odyssey Imager (LI-COR Biosciences).

Cytotoxicity
The Cytotoxicity Detection Kit (LDH, Roche) assay was used for the quantification of cell death based on the measurement of the lactate dehydrogenase (LDH) released from the cytosol of damaged cells into the supernatant. Transfected WT and Pro347Ser RHO HeLa clones were incubated with reaction mixture provided by the kit and prepared following the manufacturer's instructions. Not-transfected HeLa cells were used as "low control" (spontaneous LDH release), while Triton 100X was added to "high control" samples (maximum LDH release). The absorbance was measured at 492 nm by a spectrophotometer (Safire, Tecan, UK). The percentage of sample cytotoxicity was measured following this equation: cytotoxicity (%) = (exp. value – low control)/(high control – low control) × 100.

Animal care
Mice were housed at the TIGEM animal house (Pozzuoli, Italy) and maintained under a 12 h light/dark cycle. Pro347Ser transgenic mice were maintained as F0 by crossing them with themselves and were crossed with C57BL/6J mice purchased from Envigo Italy SRL (Udine, Italy) to generate experimental F1 mice.

Subretinal injection of AAV vectors in Pro347Ser transgenic mice
This study was carried out in accordance with the Association for Research in Vision and Ophthalmology Statement for the Use of Animals in Ophthalmic and Vision Research and with the Italian Ministry of Health regulation for animal procedures (Ministry of Health authorization number 147/2015-FR). Surgery was performed under general anesthesia, and all efforts were made to minimize animal suffering. One-week-old mice were anesthetized with an intraperitoneal injection of 2 mL/100 g of body weight of ketamine/medetomidine, then AAV2/8 vectors were delivered subretinally via a trans-scleral trans-choroidal approach, as described by Liang et al.36 Eyes were injected with 1 µL of vector solution. The AAV2/8 dose (GC/eye) was 1 × 109 of each vector/eye, and thus, co-injection resulted in a maximum of 2 × 109 GC/eye.

Electrophysiological recordings and pupillary light response analysis
The retinal electrophysiological recordings of Pro347Ser mice were performed as previously described.39 Pupillary light responses (PLRs) from Pro347Ser mice were recorded in dark condition using the TRC-501X retinal camera connected to a charge-coupled device NikonD1H digital camera (Topcon Biomedical Systems). Mice were exposed to light stimuli at 1 lux for approximately 10 s and one picture per eye was acquired using the IMAGEnet software (Topcon Biomedical Systems). For each eye, the pupil diameter was normalized to the eye diameter (from temporal to nasal side).

Histological analysis
Mice were sacrificed and eyes were fixed in Davidson’s fixative (deionized water, 10% acetic acid, 20% formalin, 35% ethanol) overnight, followed by dehydration in serial ethanol and embedding in paraffin blocks. Ten-µm thick microsections were cut along the horizontal meridian, progressively distributed on slides. Paraffin-embedded mouse retinae were treated with a citrate buffer for antigen retrieval, incubated with primary antibodies, and developed with 3',3'-Diaminobenzidine (DAB) using a Bond-III Automated IHC Stainer from Leica Biosystems according to the manufacturer’s instructions. The primary antibodies used were as follows: monoclonal ID4 (1:1,000, a gift from Robert Mollay), monoclonal 4D2 (1:30,000, Millipore, Burlington, MA, USA), and monoclonal anti-GFP (1:2,000, Proteintech, Manchester, UK). Images were acquired in bright field using a Zeiss LSM 510 AxioCam microscope.

Statistical analysis
Data were analyzed for statistical significance using two-way ANOVA or Student's t test. All values in each group were expressed as the mean ± SEM. All group comparisons were considered significant at p < 0.05, p < 0.01, and p < 0.001.

Results
CRISPR/Cas9 system specifically edits, in vitro, c.1039C>T RHO encoding the p.Pro347Ser variant
CRISPR/Cas9-mediated inactivation of dominant variants in RHO would in turn minimize and delay photoreceptor degeneration and visual loss in individuals affected by RP. To specifically target the p.Pro347Ser variant, which is caused by a C-to-T transition in RHO exon 5 (c.1039C>T), two gRNAs were designed. Guide RNA1 carries the variant (T) in the seed sequence, as the last nucleotide of the 20 nt protospacer, and guides the SpCas9 to the 5'-CGG-3' PAM sequence. On the reverse complementary strand, gRNA5 guides the high-fidelity VQRHF1SpCas9 variant to the 5'-CGAG-3' PAM sequence that includes the variant (A, in the reverse complementary strand) (Figure 1A).

In the absence of human cell lines constitutively expressing RHO, HeLa cells were engineered with a lentivirus expressing from the phosphoglycerate kinase (PGK) promoter, the WT, or p.Pro347Ser-coding RHO cDNA followed by a 250 bp-long region of the 3' UTR for a more comprehensive analysis of the translated alternatives upon CRISPR-mediated editing. Two clones carrying two copies of WT or p.Pro347Ser-coding RHO cDNA (Figure S1A) that expressed RHO at comparable levels (Figure S1B) were selected and used for further experiments. To assess gRNA specificity and efficiency, WT and Pro347Ser RHO HeLa clones were transfected with effector plasmids expressing the gRNA1 or gRNA5 and the appropriate SpCas9 nuclease (SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5, respectively), or cognate plasmids without gRNAs as negative controls, and analyzed by NGS.
Figure 1. CRISPR/Cas9 targeting of c.1039C>T RHO dominant variant encoding p.Pro347Ser

(A) Schematic representation of human chromosome 3. The picture illustrates two gRNAs (gRNA1 and gRNA5) targeting the variant (T, in bold) in the exon 5 of RHO and the PAM sequences. Capital letters indicate the exon 5, whereas the 3' UTR is in lowercase.

(B) CRISPResso analysis of NGS data obtained on Pro347Ser and WT RHO HeLa clones transfected with effector plasmids (SpCas9_gRNA1 in blue and VQRHF1-SpCas9_gRNA5 in orange). The experiment was performed in triplicate, and the mean is presented.

(C) CRISPResso analysis of indels occurring in c.1039C>T RHO transgene and in the endogenous WT RHO gene after transfection of effector plasmids in Pro347Ser RHO HeLa clone. The experiment was performed in triplicate and is presented as mean ± SEM.

(D) Indels analysis of off-target sites predicted for gRNA1 (top) and for gRNA5 (bottom). The color bars (blue and orange) represent values ascending from bottom to top ranking the indels frequency.

(E and F) CRISPResso graphic representation of indels scored in the target site of Pro347Ser RHO HeLa cells transfected with SpCas9_gRNA1 (E) and VQRHF1-SpCas9_gRNA5 (F). The top sequence is the unmodified reference. The percentage of indel frequency and the number of reads scored are indicated. Red boxes indicate nucleotide insertion and nucleotide in bold indicates substitution.

(G) Type of indels, and their relative percentage, generated in Pro347Ser RHO HeLa clone transfected with SpCas9_gRNA1 and VQRHF1-SpCas9_gRNA5. (H) CRISPResso analysis of indels generated by SpCas9_gRNA1 (top pie chart) and VQRHF1-SpCas9_gRNA5 (bottom pie chart) leading to frameshift or in-frame alterations.
CRISPResso analysis on sequence reads from the Pro347Ser RHO HeLa clone transfected with SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5 scored 65.7% and 28.6% editing, respectively, in a representative experiment (Figures S2A and S2B), while 6% and 1% of reads from transfected WT RHO clone were edited by gRNA1 and gRNAs, respectively (Figures S2C and S2D). NGS analysis on three independent experiments confirmed the efficiency and specificity of gRNA1 (62.0 ± 1.8) and gRNAs (25.4 ± 1.6) in targeting the c.1039C>T RHO variant, with barely detectable editing of WT RHO (gRNA1, 6.2 ± 0.1; gRNAs, 0.9 ± 0.1) (Figure 1B). To further eliminate any bias due to different transfection efficiency between WT and Pro347Ser RHO HeLa clones, allele-specific editing generated by SpCas9_gRNA1 and VQRHF1-SpCas9_gRNA5 was analyzed in the Pro347Ser RHO HeLa clone by NGS targeted to the p.Pro347Ser-coding transgene copies or the endogenous WT RHO alleles. Allele-specific analysis showed that 97.7% and 98.5% of all indels generated in Pro347Ser RHO HeLa clone by SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5, respectively, occurred in the variant transgene (Figure 1C), confirming the highly specific editing of the c.1039C>T RHO variant. While the genomic landscape of the RHO region in HeLa cells could contain epigenetically silenced genomic loci less prone to tether the CRISPR/Cas9 complex compared to viral derived cDNA, the high frequency of indels occurring in the transgene strongly supports the specificity of the designed gRNAs.

The major drawback of the CRISPR/Cas9 system is the possibility to induce genome-wide off-target effects. The COSMID webtool predicted 28 (Table S1) and six putative genome-wide off-targets for gRNA1 and gRNAs, respectively. We investigated, by NGS, the 10 top-ranked off-targets predicted for gRNA1 mapping to intragenic regions and all the potential off-targets predicted for gRNAs (Figure 1D). Briefly, a Hygromycin (Hygro) resistance cassette under the control of TK promoter was cloned in both effector plasmids (SpCas9_gRNA1-TKHygro and VQRHF1-SpCas9_gRNA5-TKHygro) to analyze the off-target sites in Hygro-selected non-clonal HEK293T cells expressing Cas9 nuclease and gRNA. Targeted deep sequencing detected cleavage above background in the intronic sequences of four genes: three predicted for gRNA1 and one for gRNAs (Figure 1D). The CRISPResso analysis of potential splice site modifications predicted no risk of interference with the canonical splicing signals of the hit introns (Figure S3A). Indeed, the expression of the cleaved genes in human immortalized retinal pigment epithelium (hRPE) cells transfected with SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5 was not perturbed by the intronic editing (Figures S3B and S3C).

p.Pro347Ser-specific editing leads to efficient degradation of RHO variants

To better characterize the RHO variants generated upon gene editing, we analyzed the frequency and the type of indels scored by CRISPResso in Pro347Ser RHO HeLa clones transfected with SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5 and we investigated whether small insertions or deletions created by NHEJ repair involving the last codons of c.1039C>T RHO cDNA would generate frameshift changes that in turn would knock out this allele. The analysis, set for a −10 +10 window surrounding the target sites, revealed that the most frequent indels generated by gRNA1 were 1–2 nt deletions occurring around positions −2 to +1 relative to the cleavage site (Figure 1E). Conversely, the insertion of a T nucleotide in the cut site was the top-ranked indel in gRNAs-treated samples (Figure 1F). To refine the analysis further, we calculated the frequency of deletions, substitutions, and insertions in the gRNA1- and gRNAs-treated samples. The results showed that deletions were the most frequent type of modifications for both gRNAs (88.9% and 47.8% of all indels, Figure 1G) but, interestingly, a higher prevalence of substitutions and insertions were scored in the gRNAs (32.8% and 19.4% of all indels)-treated samples compared to gRNA1 (9.9% and 1.9% of all indels)-treated samples (Figure 1G). The analysis of indel distribution revealed extended deletions, up to 80 nt, and insertions, up to 13 nt, in samples treated with SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5 (Figure S4). More importantly, despite the differences in the type of indels and their distribution, 81.5% and 68.9% of all edited sequences lead to frameshift alterations (Figure 1H), suggesting a potentially favorable outcome of p.Pro347Ser RHO knockdown.

Since Cas9-mediated editing is occurring in the RHO C terminus, a comprehensive in vitro study of the localization and degradation of RHO variants generated upon editing was performed. The six most frequent indels identified by CRISPResso upon gRNA1- or gRNAs-mediated editing resulted in shifted reading frame (delG, delGG, and insT, Figure S5) or in-frame deletion of a region including the TAA stop codon and generation of new termination codons downstream of the canonical one (del9, del12.1, del12.5, Figure S5). Cellular localization of these six RHO variants was investigated by immunofluorescence in CHO cells transfected with plasmids expressing the RHO variants fused at the N-terminal region to a human influenza hemagglutinin (HA) tag. All these variants showed localization of RHO at the plasma membrane (Figure 2A), as observed for the WT and p.Pro347Ser rhodopsin (Figure S6), but distinct from p.Pro23His RHO, which is retained in the endoplasmic reticulum (ER). The plasma membrane localization was also quantified by “In-Cell Western” assay on non-permeabilized and permeabilized CHO cells transfected with the selected six RHO variants. The assay confirmed the almost complete localization of all analyzed variants to the plasma membrane, as well as the ER retention of p.Pro23His rhodopsin (Figures S7A and S7B). Persistent expression of rhodopsin variants generated upon editing could impair the therapeutic benefits of this strategy. Therefore, the expression of WT, p.Pro347Ser, and the selected six RHO variants was evaluated by immunoblot analysis (Figures S8A and S8B).
Figure 2. Biochemical characterization of the most frequent RHO variants generated after editing

(A) Immunofluorescence analysis of CHO cells transfected with plasmids coding for RHO variants. Permeabilized cells (left) were stained with anti-BIP and anti-4D2 antibodies, and the scale bar represents 5 μm. Cells not permeabilized (right) were stained with anti-4D2 and anti-HA antibodies, and the scale bar represents 10 μm.

(B) Immunoblot analysis of WT, p.Pro347Ser, and most frequent RHO variants generated after editing expressed in CHO cells transfected with the respective coding plasmids. Anti-HA antibody was used to detect rhodopsin (RHO). The immunoblotting was normalized with an anti-beta-actin antibody.

(C) Densitometric analysis of immunoblots performed on CHO cells transfected with plasmids coding for p.Pro347Ser, del9, del12.1, and del12.5 RHO variants and treated with 10 μg/mL cycloheximide (CHX). The experiment was performed in triplicate and is presented as mean ± SEM. *p value < 0.05.

(D) Densitometric analysis of immunoblots performed on CHO cells transfected with plasmids coding for p.Pro347Ser, del9, del12.1, and del12.5 RHO variants and treated with 50 μM MG-132 proteasome inhibitor. The experiment was performed in triplicate and is presented as mean ± SEM. *p value < 0.05.

in CHO cells 2 days after transfection (Figure 2B and Figure S7C). The rate of RHO variant degradation in transfected CHO cells was measured upon treatment with the translation inhibitor cycloheximide (CHX). Compared to p.Pro347Ser rhodopsin, rhodopsin proteins carrying 9- and 12-nt deletions were rapidly degraded, with between 60%–95% removed after 6 h of translation inhibition (Figure 2C). Treatment with proteasome inhibitor MG-132 at different time points (0, 4, 6 h) revealed that the degradation of del19, del12.1, and del12.5 rhodopsin variants was proteasome mediated (Figure 2D). The degradation of the most frequent variants generated by Cas9 editing demonstrates the desired robust reduction of rhodopsin observed in Pro347Ser RHO HeLa cells transfected with SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5 effector plasmids. Similarly, p.Pro347Ser RHO protein was reduced to 40% upon transfection of effector plasmids in Pro347Ser RHO HeLa cells (Figures 3A and 3B). Furthermore, the transcript coding for p.Pro347Ser RHO was also significantly downregulated after editing as demonstrated by quantitative reverse-transcriptase PCR (qRT-PCR) (Figure 3C). Control plasmids did not influence the expression of p.Pro347Ser RHO and, notably, WT rhodopsin was not perturbed by the treatment with SpCas9_gRNA1 or VQRHF1-SpCas9_gRNA5, as expected by gene editing occurring specifically on the c.1039C>T RHO allele (Figures 3A–3C).

Interestingly, expression of Pro347Ser RHO in the corresponding HeLa clone induced higher cytotoxicity than WT RHO (Figure 3D). Indeed, an LDH cytotoxicity assay showed that the Pro347Ser RHO HeLa clone had lower viability (90%) than the WT RHO HeLa clone (99%). Following treatment with Cas9 and specific gRNA, there was a 50%–60% increase in cell viability compared to negative controls, suggesting that approximately 60% of knockdown of p.Pro347Ser RHO protein was sufficient to significantly reduce the toxic effects of this variant in vitro (Figure 3D).

Allele-specific editing significantly reduces p.Pro347Ser RHO expression in mouse photoreceptors
To translate these in vitro findings to a preclinical model of RP, we treated transgenic mice carrying the human RHO allele with the p.Pro347Ser variant. This model carries the two wild-type murine Rho alleles as well as an undefined number of transgenic alleles with the p.Pro347Ser variant and has a 1:1 ratio of transgene to endogenous opsin...
mRNA, which causes severe retinal degeneration by postnatal day (P) 30, as seen by electroretinogram responses and loss of outer nuclear layer (ONL) thickness.14 To improve the translational potential of our approach, CRISPR/Cas9 components were packaged into AAV2/8 vectors. AAV delivery of the CRISPR/Cas9 system or therapeutic genes has been proven very successful for the treatment of retinal diseases in various preclinical studies.43–45 To restrict the expression of SpCas9 to photoreceptors, we employed the interphotoreceptor retinoid-binding protein (IRBP) promoter, while GFP expression, driven by the RHO promoter, tracked the expression of gRNAs in injected eyes. Briefly, effector vector combinations were AAV expressing WT or VQRHF1 SpCas9 with AAV expressing gRNA1 or gRNAs5, while control AAV vector combinations were SpCas9 with AAV expressing gRNA scramble (Figure 4A). Pro347Ser transgenic mice received a single subretinal injection per eye at P7 of two AAV2/8 vectors carrying either WT or VQRHF1 SpCas9 in combination with either effector or scramble gRNA. Four weeks after injection (P40), molecular analyses of injected retinae showed co-expression of SpCas9, WT or VQRHF1, GFP, and Phosphodiesterase 6b (Pde6b) genes (Figure S8), indicating that the AAV vectors targeted photoreceptors upon subretinal injection. Then, the frequency and type of indels in the target locus and the inactivation of the transcript coding for p.Pro347Ser RHO were evaluated. NGS analysis detected RHO indels up to 14% in 11 retinae derived from SpCas9þgRNA1 treatment and up to 30% in 15 retinae from VQRHF1-SpCas9þgRNA5 treatment (Figure 4B). No RHO indels were detected in the retinae treated with the control vectors (n = 6, data not shown). The most common alterations on target sites were insertions and deletions that lead to frameshift as observed in vitro (Figure 4C), suggesting destabilization of p.Pro347Ser RHO transcripts and protein, which has the potential to provide therapeutic benefit to photoreceptor degeneration. Our strategy specifically targeting the human p.Pro347Ser-coding RHO gene resulted in a significant reduction, from 20% to 60%, of the c.1039C>T mRNA in 11 out of 20 retinae expressing the effector vectors compared to the contralateral retinae injected with control vectors (Figure 4D).
The localization of RHO was investigated at P50 in retinae injected with the effector and control vectors (Figure 4E). In retinae injected with control vectors, total RHO (murine WT RHO + human p.Pro347Ser RHO) was detected using the 4D2 antibody targeting the N terminus of the protein, which would recognize both the WT and C-terminal RHO variant. The predominant localization was observed in the ONL and, to a lesser extent, in the inner segment (IS) and outer segment (OS). By contrast, in retinae injected with effector vectors that downregulate the expression of the dominant human p.Pro347Ser RHO, RHO was more evident in the IS and OS, suggesting
the restoration of a correct localization to the OS. To extend these data, endogenous murine WT rhodopsin was stained using the 1D4 antibody that recognizes an epitope in rhodopsin C-terminal region, which includes the Pro347 residue, and does not react with the Ser347 protein. In retinae transduced with control vectors, the predominant localization of murine RHO was identified in the rod cell body in the ONL, along the IS, and also in the OS. Transduction with effector vectors rescued the localization of endogenous WT rhodopsin primarily to the OS. Staining for GFP in both control and treated eyes identified the transduced portion of the retina (Figure S9).

CRISPR/Cas9-mediated specific RHO allele knockdown in Pro347Ser mice

We then investigated whether c.1039C>T RHO allele-specific knockdown has a therapeutic efficacy benefit on visual function. The retinal function of Pro347Ser transgenic mice treated with AAV effector and control vectors was examined 1 month after injection (P40) by both electroretinography (ERG) and PLRs. ERG analysis showed significant improvement of the b-wave amplitudes at 20 cd.s/m² in retinae treated with either SpCas9 + gRNA1 or VQRHF1-SpCas9 + gRNA5 compared to their corresponding controls (Figure 5A). The improvement in retinal electrical activity was mirrored by post-photoreceptor responses to light stimuli, which result in transient pupil constriction. The eyes injected with SpCas9 + gRNA1 AAVs showed significantly greater pupillary constriction than gRNA scramble-injected eyes (Figure 5B). In contrast, the eyes injected with VQRHF1-SpCas9 + gRNA5 AAVs showed more variation in their pupillary constriction that was not statistically significantly different compared to their control group.

Taken together, these results indicate that a precise editing approach to selectively silence a dominant disease-associated variant in RHO holds promise for slowing down the photoreceptor degeneration observed in dominant RPs.

Discussion

RP causes the progressive death of photoreceptors and eventually blindness. The treatment of RP is challenging, but an early therapeutic intervention aimed at blocking or reducing rod degeneration would be an effective approach to preserve vision in individuals affected by RP. This was the aim of earlier nutrient trials in RP, and results were assessed over several years.46–48

More recently, gene therapy approaches aimed at correcting disease-associated variants identified in genes causing RPs have been reported.4,49,50 In this study, we addressed a CRISPR/Cas9-mediated gene editing approach for a common variant in RHO that causes autosomal dominant RP (adRP) (RP4 [MIM: 613731]). The ClinVar Miner database51 lists 156 variants in RHO, and at least 50 of them are pathogenic or likely pathogenic variants associated with RP4. In this scenario, the knockdown of both alleles followed by gene supplementation represents the most cost-effective approach to pursue. Indeed, the ablate-and-replace strategy could be used for the treatment of all RHO disease-associated variants, thus circumventing the allelic heterogeneity of the disease. However, in the case of editing, this strategy requires bi-allelic events, a
difficult goal to achieve in vivo. Moreover, the threshold level of RHO protein in the cell represents a crucial issue. For rhodopsin, there is a fine balance between insufficiency and toxicity\textsuperscript{2,5} that requires a fine-tuning of the replacement or augmentation of WT rhodopsin. Indeed, the "suppression and replacement" approach involves the risk of converting a dominant condition to a recessive RP should replacement not be as effective as suppression. Moreover, an excess of RHO expressed by an exogenous transgene cassette could exert a detrimental effect for photoreceptor cells, as shown by Mao et al., who described that WT RHO overexpression leads to retinal degeneration in WT mice.\textsuperscript{54}

Conversely, a specific and permanent silencing of the variant allele by CRISPR/Cas9, as proposed in this study, would prevent the pathogenic effects of most dominant variants while preserving the WT allele, which would also not require the ethical issue of disrupting a functional human gene. Indeed, allele-specific knockout would address both dominant-negative and gain-of-function changes, while conventional gene replacement therapy is indicated for haploinsufficiency. Notably, haploinsufficiency has been linked to adRP caused by disease-associated variants in PRPF3 or PRPH2 but not in RHO.\textsuperscript{6,23}

Several studies demonstrated beneficial effects of CRISPR/Cas9 on adRP treatment in preclinical models.\textsuperscript{12,21,55} The majority of the reported studies were designed to correct Pro23His allele, the most frequent disease-associated variant major of the reported studies were designed to correct RHO in WT mice.\textsuperscript{54} That WT RHO overexpression leads to retinal degeneration, as shown by Mao et al., who described that WT RHO overexpression leads to retinal degeneration in WT mice.\textsuperscript{54}

To test this hypothesis, we transferred the CRISPR/Cas9 editing platform to the retina of Pro347Ser transgenic mice using AAV2/8. Although Pro347Ser transgenic mice do not match the copy number or genomic context of the p.Pro347Ser RHO variant in individuals affected by RP, they represent a valuable in vivo model to test the efficacy of gene therapy. AAV2/8 vectors efficiently transfer genes to photoreceptors in the retina.\textsuperscript{31,49,57} and successful applications of CRISPR-AAV vectors in retinal diseases are already reported.\textsuperscript{58–60} Pro347Ser transgenic mice received a single injection of either effector or control AAV2/8 vector combinations. One month later, during rod degeneration, molecular analyses revealed a variable but effective permanent knockdown of human c.1039C>T transcript as a consequence of allele-specific gene editing provided by both SpCas9 variants. The frequency of indels significantly differs from that scored in vitro. This could be due to a lower transduction efficiency with respect to the transfection of HeLa cells, additional copies of the Pro347Ser allele in the transgenic mice or to the requirement of two AAVs to reconstitute the effector SpCas9/gRNA system. The most frequent indels types were insertions and deletions that lead to frameshift as observed in vitro, suggesting a comparable outcome for p.Pro347Ser rhodopsin protein in vivo. Significant evidence of therapeutic benefit was
obtained using both PLRs and ERG. ERG comparisons between treated and control eyes at P40 demonstrated significantly improved responses in treated eyes with respect to the controls.

The beneficial effects observed using ERG were also observed at P50 using histological analyses on treated and control retinae. The mislocalization and retention of rhodopsin in the ONL was partially rescued in retinae injected with effector AAV2/8 vectors, showing an increased localization of WT murine rhodopsin to OS as consequence of downregulation of the dominant effect exerted by human p.Pro347Ser RHO. It is noteworthy that although the ONL structure was not significantly improved by the treatment, a significant functional recovery by ERG and PLR was registered, indicating that the reduction of human RHO transcript coding for p.Pro347Ser was sufficient to improve the function of surviving rods but did not significantly prevent their death.

Overall, our study provides proof of concept for CRISPR/Cas9-mediated allele-specific targeting of a common RHO variant associated with adRP. Whether using the WT or the VQRHF1 SpCas9 variant, which show here similar efficiency, it could be beneficial for 62% of the pathogenic and likely pathogenic variants in RHO, listed in ClinVar Miner, displaying genomic sequences suitable for allele-specific inactivation (Table S2), with no need for RHO supplementation therapy. The translation of this genome editing approach to the clinic will require further pre-clinical testing including using affected-individual-derived cell lines carrying one WT and one Pro347Ser variant RHO allele, potentially through human retinal organoids to model the variant editing in right genomic and cellular context. Moreover, this CRISPR/Cas9 approach would benefit from the development of more efficient clinically relevant non-toxic viral or non-viral delivery systems allowing the targeting and editing of a higher number of photoreceptors. Allele-specific genome editing therefore has the potential to become the therapeutic intervention of choice for precise silencing of genetic variants causing dominant retinal degeneration.

Data and code availability

The NGS datasets supporting the current study have not been deposited in a public repository but are available from the corresponding author (A.R.) on request.

Supplemental information

Supplemental information can be found online at https://doi.org/10.1016/j.ajhg.2021.01.006.

Acknowledgments

The authors are grateful to Valeria Marigo (University of Modena and Reggio Emilia) for helpful discussion throughout the development of this study. The authors wish to thank the LaBSSAH - CI-BIO Next Generation Sequencing Facility of the University of Trento for sequencing samples and Robert Molday (Department of Biochemistry and Molecular Biology, University of British Columbia, Canada) for the gift of the anti-Rho-1D4 antibody and Idil Ahmed (Department of Eye Pathology, UCL Institute of Ophthalmology) for the support in performing the histology. This work was supported by Fondazione Roma, Italy (call for Retinitis Pigmentosa) to A.R.; by the ERC Advanced grant “EYEGET” (694323), the European Union Horizon 2020 “Upgrade” (825825), and the Foundation Fighting Blindness grant TA-GT-0619-0762 to A.A.; and by grants from the Wellcome Trust, United Kingdom (205041) and the Foundation Fighting Blindness USA (PPA-0717-0719-RAD) to M.E.C.

Declaration of interests

C.P., M.L., D.B., A.A., and A.R. are listed as inventors on a patent application related to this work.

Received: August 31, 2020
Accepted: January 8, 2021
Published: January 27, 2021

Web resources

Addgene, https://www.addgene.org
ClinVar Miner, https://clinvarminer.genetics.utah.edu
CRISPPresso web tool, https://crispresso.pinellolab.partners.org/
RetNet - Retinal Information Network, https://sph.uth.edu/retnet

References


