

Lentivirus-mediated expression of cDNA and shRNA slows degeneration in retinitis pigmentosa

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Abstract

Mutations in *Pde6b* lead to high levels of signaling molecules cyclic guanosine monophosphate (cGMP) and Ca^{2+} , which ultimately result in photoreceptor cell death in certain forms of retinitis pigmentosa (RP). The level of cGMP, which is controlled by opposing activities of guanylate cyclase (GUCY) and photoreceptor phosphodiesterase-6 (PDE6), regulates the opening of cyclic nucleotide-gated ion channels [CNG] and thereby controls Ca^{2+} influx into the outer segments. Using a lentiviral gene therapy approach, we have previously shown that degeneration can be temporarily slowed either by introducing wild-type PDE6 β or knocking down expression of GUCY2E and CNGB1 in photoreceptors of *Pde6b*^{H620Q}, a mouse model for RP. Rescue was transient with either approach. Therefore, we tested a novel combination therapy using bipartite lentiviral vectors designed to both introduce wild-type PDE6 β expression and knockdown GUCY2E or CNGB1. Immunoblot analysis shows simultaneous increases in PDE6 β and decreases in GUCY2E or CNGB1 in retinas transduced by the vectors, indicating successful transduction. In *Pde6b*^{H620Q} mutants, we observe rescue of photoreceptor function and an increase in photoreceptor rows as compared with untreated controls. However, no evidence of prolonged rescue beyond the limit of the previously tested single therapy was observed.

Keywords: gene therapy, *Pde6b*^{H620Q}, mouse model, PDE6, GUCY2E, CNGB1, bipartite, retinitis pigmentosa, lentiviral vector, shRNA

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Introduction

About 36,000 cases worldwide of simplex and familial retinitis pigmentosa (RP) are caused by loss of function mutations in rod photoreceptor phosphodiesterase-6 (PDE6).^{1–4} Although significant advances in our understanding of RP have been made, the exact interplay between defective PDE6 and the onset of RP pathogenesis remains poorly understood.

Mouse models of human RP have been extensively used to study the mechanisms of onset and progression of the disease.^{5–8} Several *Pde6b* mutant alleles have been identified and studied, including *Pde6b*^{rd1}, *Pde6b*^{rd10} and *Pde6b*^{H620Q}.^{5,6,8} Loss of PDE6 enzyme activity has been shown to result in increased levels of cyclic guanosine monophosphate (cGMP) and Ca^{2+} in *Pde6b*^{rd1} and *Pde6b*^{H620Q} mice.^{5,9–14}

Excessive influx of Ca^{2+} is likely the critical event leading to photoreceptor cell death.^{5,14–16}

Under normal physiological conditions, cGMP levels are regulated by opposing activities of PDE6, which hydrolyzes cGMP into GMP, and guanylate cyclase (GUCY), which produces cGMP from guanosine-5'-triphosphate (GTP). During phototransduction, cGMP controls the excitation state of the photoreceptor through regulation of cGMP-gated (cyclic nucleotide-gated ion channel [CNG]) channels. In the dark, cGMP maintains CNG channels in an open configuration, allowing the influx of Ca^{2+} to the cytoplasm. In the light, PDE6 activation results in the rapid hydrolysis of cGMP and resultant closure of CNG channels. When PDE6 is absent or reduced in mutant photoreceptors, elevated concentrations of cGMP keep CNG channels open, allowing entrance of excess Ca^{2+} , leading to cell death.⁸

Transduction of mutant photoreceptors with gene therapy vectors have been used to delay photoreceptor death.¹⁷ We have previously shown that expression of *Pde6b* cDNA in *Pde6b*^{H620Q} mutant retinas transduced with *Opsin::Pde6b* lentivirus results in improved photoreceptor physiology and increased photoreceptor numbers.⁵ We also took advantage of the known relationships between PDE6, GUCY and CNG to test if reduced expression of GUCY2E or CNCA1 could suppress the degeneration phenotype. Knockdown of *Gucy2e* or *Cnga1* was associated with improved survival and function in *Pde6b*^{H620Q} mutant photoreceptors transduced with shRNA lentiviral vectors.¹⁸

Although our attempts have demonstrated these monopartite vectors can rescue the mutant phenotype, the effect was incomplete and temporary. Here we hypothesized that a combined therapy approach would have an additive or synergistic effect on photoreceptor function and survival. For that purpose, we generated two kinds of bipartite expression vectors to express *Pde6b* cDNA and *Gucy2e* or *Cnga1* shRNAs. After subretinal injection of these lentiviral vectors in *Pde6b*^{H620Q} mice, we analyzed the mutant photoreceptors by electroretinogram (ERG) recording and histology. Our results showed limited functional and morphological rescue, although no synergy was observed.

Materials and methods

Animals

Mice were used in accordance with the Statement for the Use of Animals in Ophthalmic and Vision Research of the Association for Research in Vision and Ophthalmology, as well as the Policy for the Use of Animals in Neuroscience Research of the Society for Neuroscience. *Pde6b*^{H620Q} mice used in this experiment were bred from a colony of mice that has been previously reported.^{5,19} All *Pde6b*^{H620Q} mice analyzed in this study are homozygotes and in the C3H background.

Construction of lentiviral vectors

The viruses used for our experiments were self-inactivating vectors consisting of a 5' long terminal repeat (LTR), a packaging signal ψ , a tRNA primer binding site, a reverse response element and a 3'LTR. This viral vector also contained a central polypurine tract/DNA flap (cPPT) and a Woodchuck hepatitis virus post-transcriptional regulatory element.

Three vectors were assembled: *H1::Gucy2e*, *H1::Cnga1* and *Opsin::Gucy2e* (Figure 1). The vectors were designed to simultaneously express PDE6 β and knockdown GUCY2E or CNCA1. To deliver rod-specific expression of PDE6 β , all vectors contained the mouse rhodopsin 2.0 kb promoter cloned upstream of full-length *Pde6b* cDNA. Both *H1::Gucy* and *H1::Cnga1* vectors drive shRNA expression using the human H1 RNA polymerase III promoter (as per manufacturer's instructions; Biogenova Corporation, Frederick, MD, USA).

The *Opsin::Gucy2e* vector was designed to direct *Gucy2e* shRNA expression in rods, as GUCY2E and GUCY2F are

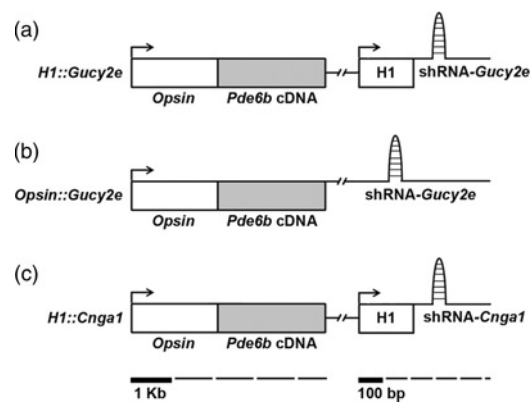


Figure 1 Schematic representation of the lentiviral vectors. The *H1::Gucy2e* (a) construct consists of a 2 kb mouse *Opsin* promoter, a *Pde6b* open reading frame (*Pde6b* cDNA) and the H1 promoter, which drives expression of the shRNA-*Gucy2e*. The *Opsin::Gucy2e* (b) consists of a 2 kb *Opsin* promoter, a *Pde6b* open reading frame and a shRNA-*Gucy2e*. The *H1::Cnga1* lentiviral vector (c) was constructed using 2 kb of the mouse *Opsin* promoter, which directs expression of the *Pde6b* cDNA fragment, and the H1 promoter, which drives expression of the shRNA-*Cnga1*.

required for efficient transport of PDE6 and other proteins to outer segments.²⁰ *Opsin::Gucy2e* contained *Gucy2e* shRNA sequence cloned within the T-cell receptor- β C β 2.1 intron 42, which lies within the same transcription unit of the *Pde6b* cDNA. A unique *MunI* site within the intron was used for insertion of a shRNA oligonucleotide. This configuration is expected to drive expression of both the *Pde6b* cDNA and the pre-*Gucy2e* shRNA within photoreceptors, similar to the expression of pre-miR34 (as per manufacturer's instructions; System Biosciences, Mountain View, CA, USA).^{21,22}

Subretinal transduction of lentivirus

Subretinal injections were described before.^{5,18} In short, *Pde6b*^{H620Q} or C57BL/6J mice at postnatal day (P) 5 were anesthetized in ice and placed under a surgical microscope. The eyelid was opened and 0.8 μ L of virus particles (2×10^7 TU/mL) were injected at the 6 o'clock position 1.5 mm from the limbus. The injection produced a bubble in the subretinal space. Mice received a subretinal injection of lentivirus in the right eye, while the left eye served as a control.

Histochemical analyses

Mice were sacrificed and hematoxylin-eosin (H-E) retinal sections were obtained as described before.^{23,24} Shortly thereafter, animals were sacrificed and eyes were enucleated and fixed in 0.5 $\times$ Karnovsky's fixative (2% paraformaldehyde, 1.25% glutaraldehyde, 0.2 mol/L phosphate-buffered saline). After that they were embedded in paraffin and sectioned every 4 μ m. Sectioning proceeded along the long axis of the segment so that each section contained both upper and lower retina as well as the posterior pole. H-E staining was then conducted. The number and morphology of photoreceptors in injected eyes were compared with control. Quantification of photoreceptor nuclei was conducted on several sections containing the optic nerve.

Immunoblot analysis

C57BL/6J mice were euthanized at eight weeks of age and retinas were homogenized in 10% sodium dodecyl sulfate (SDS) by sonication. To minimize individual differences between subretinal surgeries, retinal extracts were pooled from each virus, and each assay was carried out in duplicate. After denaturation at 100°C for five minutes, total protein was measured by the DC Protein Assay method (Bio-Rad Laboratories, Hercules, CA, USA). Proteins were then separated by SDS polyacrylamide gel electrophoresis. Samples were transferred to nitrocellulose membranes, which were preincubated in blocking solution (3% bovine serum albumin, 150 mmol/L NaCl, 100 mmol/L Tris, 0.5% Tween-20). Membranes were incubated with antibodies against PDE6 β (1:2000; Thermo Scientific, Waltham, MA, USA), GUCY2E (1:2000; kindly provided by Alexander Dizhoor, Pennsylvania College of Optometry, Elkins Park, PA, USA), and CNGA1 (1:12; kindly provided by Robert Molday, The University of British Columbia, Vancouver, BC, Canada). GNAT1 is a rod-specific protein, which served as a control for photoreceptor protein content (Gat1, 1:2000; Santa Cruz Biotechnology, Santa Cruz, CA, USA). Membranes were washed and then incubated with either goat antirabbit conjugated horseradish peroxidase secondary antibodies (1:10,000; Santa Cruz Biotechnology) or goat antimouse IgG-horseradish peroxidase secondary antibodies (1:10,000; Santa Cruz Biotechnology). Antibody complexes were visualized by chemiluminescence detection (Immobilon Western, Millipore Corporation, Billerica, MA, USA) using Kodak BioMax film (Kodak, Rochester, NY, USA). Densitometry analyses were performed using the AlphaImager imaging system (Alpha Innotech, Cell Biosciences, Santa Clara, CA, USA).

Photoreceptor functional analysis

ERGs were performed on mice at P60 and P90, as previously described.^{25–27} ERG b-wave enhancement of maximal response was measured. The enhancement is defined as the difference in maximum ERG responses between transduced and control fellow eyes, measured in microvolts (μ V).

Results

The bipartite lentiviruses used in this study are designed to express both PDE6 β and shRNAs to knockdown either GUCY2E or CNGA1 in *Pde6b*^{H620Q} photoreceptors. To measure the ability of the vectors to alter retinal gene expression, we performed immunoblot analysis on lysates prepared from C57BL6 retinas injected with *H1::Gucy2e*, *Opsin::Gucy2e* or *H1::Cnga1* viral particles (Figure 2). Retinas transduced with *H1::Gucy2e*, *Opsin::Gucy2e* and *H1::Cnga1* showed detectably higher PDE6 β levels than contralateral uninjected retinas (Figures 2a–c). Vectors *H1::Gucy2e* and *H1::Cnga1* also produced reductions in GUCY2E and CNGA1 expression, respectively (Figures 2a and c). Similarly, transduction of the *Opsin::Gucy2e* vector resulted in reduced expression of GUCY2E (Figure 2b). Expression of GNAT1, a rod-specific protein measured as a control, did not vary between controls and virus-transduced

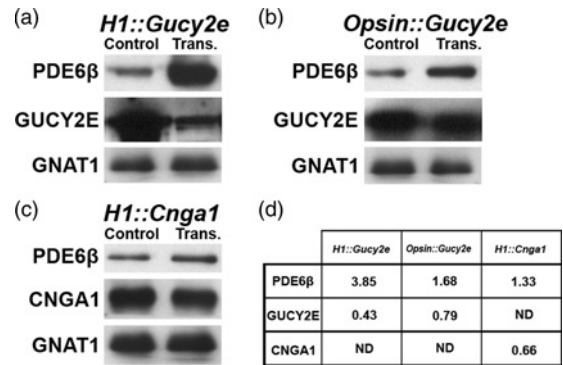


Figure 2 Immunoblotting analysis of retinal lysates from C57BL/6J mice transduced with bipartite vectors. PDE6 β was increased after transduction with either *H1::Gucy2e*, *Opsin::Gucy2e* or *H1::Cnga1* (a–c). GUCY2E is depressed in retinas transduced with *H1::Gucy2e* (a) and *Opsin::Gucy2e* (b) compared with controls. CNGA1 expression is reduced in retinas transduced with *H1::Cnga1* (c) compared with controls. GNAT1 was used as control for photoreceptor protein content and its expression was not reduced by any of the three viruses. Intensities were determined by densitometry to calculate an integral density value for each band. The values were normalized to total protein and GNAT1 expression as a percentage of GNAT1 signal in 20 μ g of control lysate (d). Samples from each experimental group were pooled in order to minimize variation resulting from differences in subretinal surgeries; immunoblot analyses were carried out in duplicate (average values shown). ND, not determined; trans., transduced

eyes (Figure 2). Numerical values were obtained after measuring densitometry of different proteins and each transduced retina was compared with its control (Figure 2d).

To measure the ability of the three vectors to retard *Pde6b*^{H620Q} degeneration, we analyzed the consequences of vector transduction on photoreceptor function and morphology. ERG analysis was performed on injected and control eyes in *Pde6b*^{H620Q} mice at P60 and P90 (Figure 3). In general, maximal ERG responses were higher for transduced than control eyes for all three vectors (Figures 3a–f). In particular, photoreceptor-generated a-waves and inner retina-mediated b-waves were observed at P60 in eyes injected with the bipartite lentiviruses (Figures 3b, d and f) but not in control untreated eyes (Figures 3a, c and e).

To compare the effectiveness of these newly created bipartite vectors to the previously tested monopartite vectors (*Opsin::Pde6b*, shRNA-*Gucy2e*, shRNA-*Cnga1*),^{5,18} we calculated b-wave enhancement values and compared them between all vectors (Figure 3g). We have previously shown that expression of wild-type PDE6 β in *Pde6b*^{H620Q} photoreceptors partially rescues the ERG phenotype (*Opsin::Pde6b*). Similarly, knockdown of CNGA1 by shRNA expression results in partial ERG rescue (shRNA-*Cnga1*). In contrast, shRNA knockdown of GUCY2E does not detectably restore *Pde6b*^{H620Q} ERGs, although photoreceptor cell rescue was observed, likely due to the known effect of *Gucy2e* loss of function on photoreceptor physiology.²⁰ In this study, we found that transduction of the new bipartite vectors resulted in partial *Pde6b*^{H620Q} ERG rescue. *H1::Gucy2e* and *Opsin::Gucy2e* vectors produced higher b-wave enhancement than the shRNA-*Gucy2e* vector, suggesting that co-expression of PDE6 β can mitigate the effect of *Gucy2e* knockdown on phototransduction. However, the apparent rescue of the

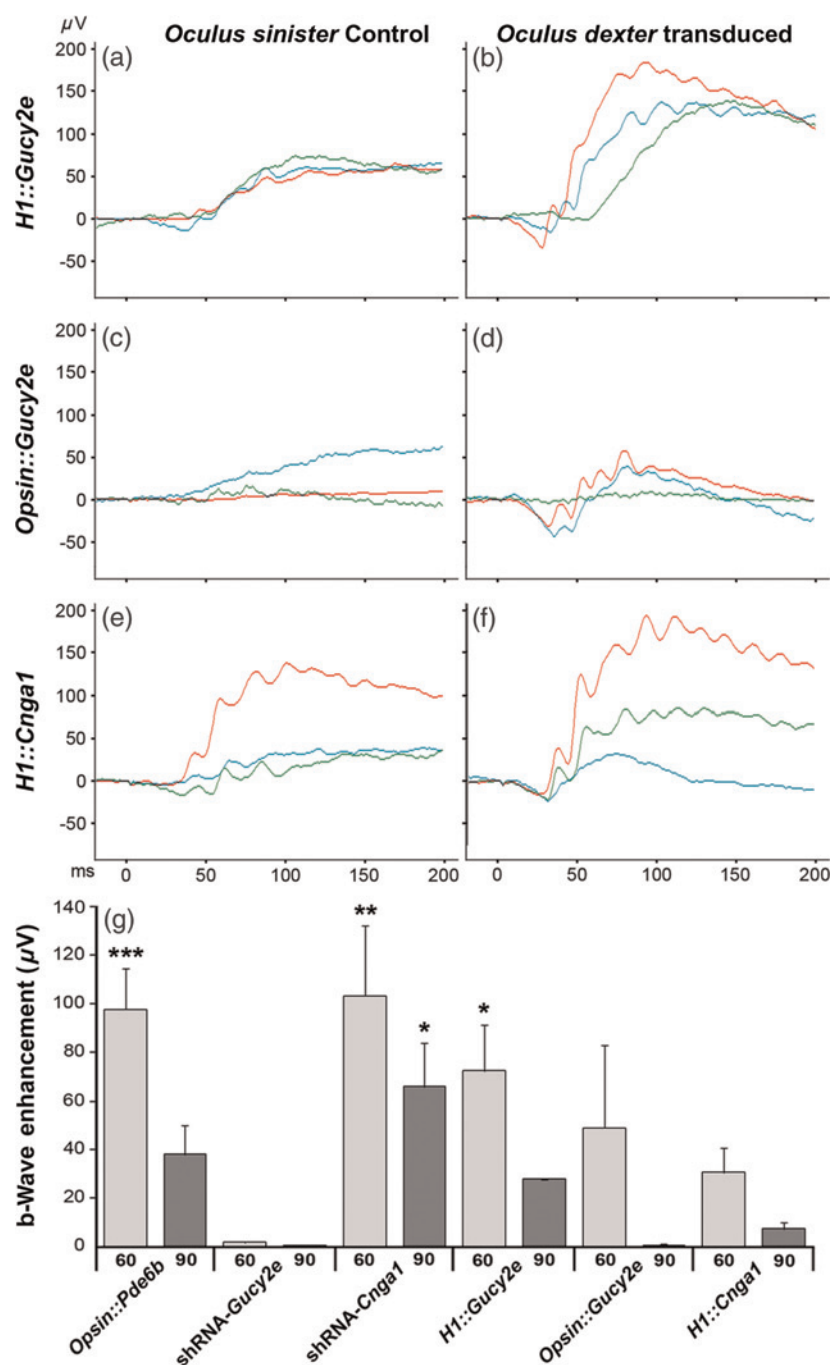


Figure 3 Functional rescue due to bipartite vector transduction. Maximal dark-adapted electroretinograms (ERGs) were performed on both eyes (transduced and control) of *Pde6b*^{H620Q} animals at approximately P60 and P90. In each ERG panel (a–f), the tracings represent different mice from the same treatment group. Compared with untreated controls (a, c, e), the *H1::Gucy2e* (b), *Opsin::Gucy2e* (d) and *H1::Cnga1* (f) vectors improved visual function. Function was measured as both the photoreceptor-mediated a-wave and inner retina-mediated b-wave. ERG traces from three mice are shown for each virus. Enhancement of b-wave was calculated by comparing ERG values from transduced and untreated eyes. Results of the bipartite therapies described in this study (*H1::Gucy2e*, *Opsin::Gucy2e* and *H1::Cnga1*) were compared with previous results of the monopartite therapies (*Opsin::Pde6b*, *shRNA-Gucy2e* and *shRNA-Cnga1*)^{5,18} (g). Significant values: **P* < 0.05, ***P* < 0.01, ****P* < 0.001 (A color version of this figure is available in the online journal)

b-wave was only significant (*P* = 0.0297) when the eye was transduced with the *H1::Gucy2e* vector (b-wave: 132.69 ± 26.02 μV), compared with the non-treated eye (b-wave: 61.17 ± 7.53 μV). *H1::Cnga1* b-wave enhancement was lower than either *Opsin::Pde6b* or *shRNA-Cnga1*, suggesting that simultaneous expression of wild-type PDE6β and knockdown of CNGA1 negatively affect *Pde6b*^{H620Q} mutant photoreceptor function.

To confirm rescue of cell viability, we quantified the number of photoreceptor nuclear rows in the transduced retinas (Figure 4). In untreated control *Pde6b*^{H620Q} retina, the photoreceptor layer contained one row of nuclei at week 8 (Figure 4a). Retinas transduced with bipartite vectors *H1::Gucy2e*, *Opsin::Gucy2e* and *H1::Cnga1* showed greater than one row of photoreceptors in the injected portion of the retina (Figures 4b–d). In some cases the

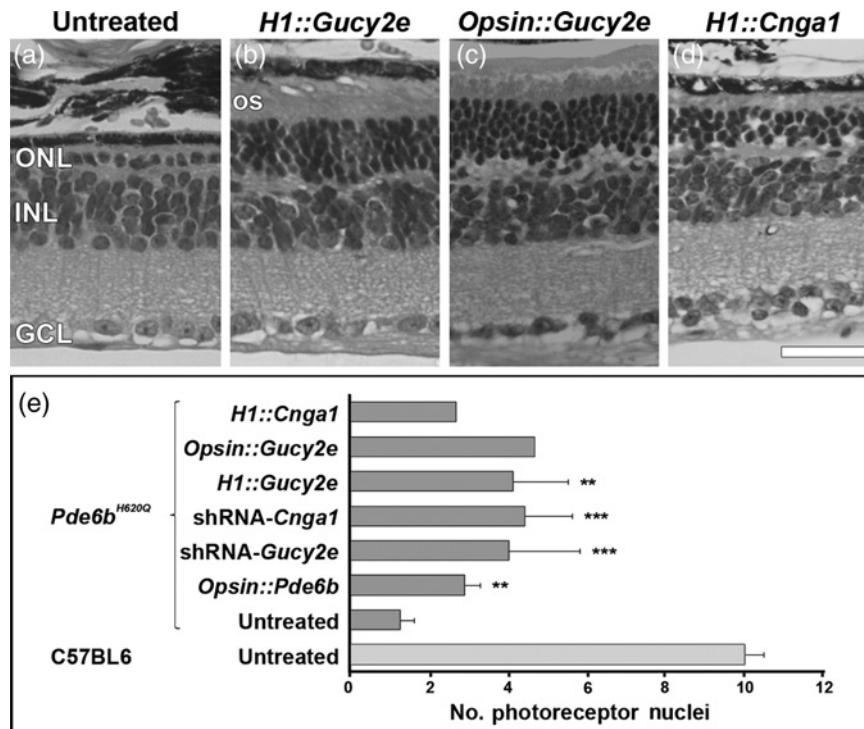


Figure 4 Photoreceptor cell rescue due to bipartite vector transduction. Hematoxylin–eosin labeling in *Pde6b*^{H620Q} retinal sections at P60 stained the different layers of the retina: ONL, INL, GCL (a–d). In non-treated *Pde6b*^{H620Q} retinas, the ONL contained only a single row of photoreceptors (a). Treated retinas with *H1::Gucy* (b), *Opsin::Gucy* (c) and *H1::Cnga1* (d) showed a higher number of photoreceptor rows within the ONL. Outer segments (OS) were also preserved in transduced but not in untreated retinas. Photoreceptor cell rescue in *Pde6b*^{H620Q} using bipartite therapies described in this study (*H1::Gucy2e*, *Opsin::Gucy2e*, and *H1::Cnga1*) was compared with previous data obtained from the use of monopartite therapies (*Opsin::Pde6b*, *shRNA-Gucy2e* and *shRNA-Cnga1*) and from untreated *Pde6b*^{H620Q} and healthy, untreated C57BL6^{5,18} (e). ONL, outer nuclear layer; INL, inner nuclear layer; GCL, ganglion cell layer. Scale bar in (d) is 50 μ m. Bars in (e) represent mean and SEM. Significant values: ** $P < 0.005$, *** $P < 0.001$

outer nuclear layer had around four rows of photoreceptor cells. Photoreceptors in transduced retinas showed outer segments that were not present in *Pde6b*^{H620Q} untreated retinas at this age. The inner nuclear layer and ganglion cell layer remained intact in both treated and untreated retinas. Quantification of the number of nuclear rows for both single and bipartite vectors showed evidence of photoreceptor rescue when compared with untreated control (Figure 4e). Wild-type control mice had approximately 10 cell rows.

Reduction of GUCY2E or CNGA1 expression in conjunction with an increase in PDE6 β retarded photoreceptor cell degeneration. However, this bipartite expression did not provide dramatically greater cell rescue efficacy than conventional monogenic therapies.

Discussion

Defects in the beta (β) subunit of PDE6 result in increased levels of the signaling molecules cGMP and Ca^{2+} in both the *Pde6b*^{rd1} and *Pde6b*^{H620Q} mouse models of RP.^{5,9,14} Such increase of intracellular cGMP and Ca^{2+} levels is toxic to photoreceptors.²⁸ Based on this understanding of the biochemical pathogenesis, we developed a therapeutic strategy using a lentivirus vector to express wild-type PDE6 β in *Pde6b*^{H620Q} mice.⁵ Even though photoreceptor cell count and function showed some rescue, they were

not completely restored to wild-type levels. In our previous study, a control vector (CMV::GFP) was used as a negative control to demonstrate that the rescue resulted from the expression of the specific proteins being introduced, rather than from the presence in the rods of a virally introduced promoter.⁵ As an alternative approach, we studied whether knocking down the expression of GUCY2E would reduce the high cGMP concentration in order to avoid Ca^{2+} overdose. Similarly, we tested whether *Cnga1* knockdown could prevent cation entry; we expected this to have the added benefit of not affecting cGMP levels. Our results of these alternative studies demonstrated that shRNA knockdown of GUCY2E or CNGA1 retards photoreceptor cell degeneration.¹⁸ However, the ability of each of the three monopartite vectors to retard retinal degeneration in *Pde6b*^{H620Q} mice was limited. Our novel bipartite lentiviral vectors were designed to improve photoreceptor function and survival by providing functional PDE6 β and concurrently decreasing abnormal cGMP and Ca^{2+} levels.

Here we have shown that some combined gene therapies can overcome the limitations of single therapies. Two outcomes were used to evaluate the effect of the therapies: cell survival and photoreceptor function. Since both the monopartite and bipartite methods resulted in increased cell survival, we compared the preservation of function by each in order to determine the relative effectiveness of the strategies.

The bipartite *H1::Gucy2e* vector is an improvement over the monopartite *shRNA-Gucy2e* vector. *Gucy2e* knockdown

showed weaker rescue of ERG responses,¹⁸ which is similar to the *Gucy2e* knockout phenotype.²⁰ Although shRNA-*Gucy2e* produces cell survival, these photoreceptors do not express sufficient PDE6 β to elicit a significant response to light. By simultaneously introducing the wild-type *Pde6b* and *Gucy2e* shRNA, we partially corrected this. Because PDE6 β decreases the level of cGMP and GUCY2E ultimately increases the level of cGMP, the two changes induced by this vector should be combined to improve the function more than the monopartite virus.

The bipartite *Opsin::Gucy2e* vector produces a lower ERG rescue than *Opsin::Pde6b* alone. PDE6 β expression in C57BL6 mouse transduced retinas was not significantly higher than endogenous levels. There may not be sufficient PDE6 β to reverse the effect of shRNA-*Gucy2e*. This may be caused by preferential processing of the chimeric *Pde6b/Gucy2e* transcript by the shRNA processing pathway at the expense of *Pde6b* cDNA translation. Future designs for this strategy need to improve *Pde6b* cDNA expression from this vector.

H1::Cnga1 was designed to simultaneously diminish the number of CNG channels, thereby reducing the entry of Ca²⁺ into rods, and express wild-type PDE6 β , increasing cGMP breakdown. Even though the retina showed morphological retention after *H1::Cnga1* transduction, the ERG responses remained much lower than in mice transduced with either *Pde6b* or *Cnga1* monopartite vectors.^{5,18} These data suggest that simultaneous expression of wild-type PDE6 β and CNGA1 knockdown negatively affects *Pde6b*^{H620Q} mutant photoreceptor physiology. The combination of a reduction in CNG channel numbers and an increase in PDE6 β activity may reduce Ca²⁺ to levels insufficient for efficient phototransduction.

Although our lentivirus vectors did produce some rescue, there are several improvements that must be achieved before these treatments can be translated from the lab to the clinic. These include efficient transduction of photoreceptors; early and appropriate expression of wild-type PDE6 β (or shRNAs); prevention of vector silencing; and prevention of host immune responses to transduced cells. For example, we have previously shown low photoreceptor transduction rates using lentiviral vectors expressing LacZ,⁵ in agreement with a report of 5% photoreceptor transduction using other LacZ-expressing viral vectors.²⁹ However, as shRNAs and siRNAs can be transported between cells through gap junctions,^{30,31} the degree of rescue may spread beyond the infected cells. In unpublished serial sections of rescued retinas, a majority of the sections proximal to the injection site contain rescued photoreceptors.

Another question to be addressed is which viral vector system might provide a more effective treatment. Most recently, Jing Pang *et al.*³² used AAV8 (Y733F) to achieve significant long-term rescue of the *Pde6b*^{rd10} mutant phenotype for over six months, which is greater than the rescue achieved with previously tested AAV vectors. The high viral titer and photoreceptor transduction rates of AAV8 vectors may preclude the necessity for a combined therapy approach.

Author contributions: SHT, RJD, C-SL and JT participated in the design of the study; RJD, JT, JS-P and CWH conducted the experiments; JT, JS-P, RJD, KVV, JDS and SHT interpreted the results; and JT, JS-P, RJD, KVV and SHT wrote the manuscript.

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