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Susanne F. Koch, …, Chyuan-Sheng Lin, Stephen H. Tsang

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Hereditary retinal degenerative diseases, such as retinitis pigmentosa (RP), are characterized by the progressive loss of rod photoreceptors followed by loss of cones. While retinal gene therapy clinical trials demonstrated temporary improvement in visual function, this approach has yet to achieve sustained functional and anatomical rescue after disease onset in patients. The lack of sustained benefit could be due to insufficient transduction efficiency of viral vectors (“too little”) and/or because the disease is too advanced (“too late”) at the time therapy is initiated. Here, we tested the latter hypothesis and developed a mouse RP model that permits restoration of the mutant gene in all diseased photoreceptor cells, thereby ensuring sufficient transduction efficiency. We then treated mice at early, mid, or late disease stages. At all 3 time points, degeneration was halted and function was rescued for at least 1 year. Not only do our results demonstrate that gene therapy effectively preserves function after the onset of degeneration, our study also demonstrates that there is a broad therapeutic time window. Moreover, these results suggest that RP patients are treatable, despite most being diagnosed after substantial photoreceptor loss, and that gene therapy research must focus on improving transduction efficiency to maximize clinical impact.

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Halting progressive neurodegeneration in advanced retinitis pigmentosa

Susanne F. Koch,1,2 Yi-Ting Tsai,1,2,3 Jimmy K. Duong,4 Wen-Hsuan Wu,1,2 Chun-Wei Hsu,1,2 Wei-Pu Wu,1,2 Luis Bonet-Ponce,5 Chyuan-Sheng Lin,6 and Stephen H. Tsang1,2,3,6

1Barbara and Donald Jonas Laboratory of Stem Cells and Regenerative Medicine and Bernard and Shirlee Brown Glaucoma Laboratory, Departments of Pathology and Cell Biology, Columbia University, New York, New York, USA. 2Edward S. Harkness Eye Institute, New York Presbyterian Hospital, New York, New York, USA. 3Institute of Human Nutrition, College of Physicians and Surgeons, Columbia University, New York, New York, USA. 4Department of Biostatistics, Mailman School of Public Health, Columbia University Medical Center, New York, New York, USA. 5Department of Physiology, School of Medicine and Dentistry, Catholic University of Valencia, Valencia, Spain. 6Herbert Irving Comprehensive Cancer Center, Columbia University Medical Center, New York, New York, USA.

Hereditary retinal degenerative diseases, such as retinitis pigmentosa (RP), are characterized by the progressive loss of rod photoreceptors followed by loss of cones. While retinal gene therapy clinical trials demonstrated temporary improvement in visual function, this approach has yet to achieve sustained functional and anatomical rescue after disease onset in patients. The lack of sustained benefit could be due to insufficient transduction efficiency of viral vectors (“too little”) and/or because the disease is too advanced (“too late”) at the time therapy is initiated. Here, we tested the latter hypothesis and developed a mouse RP model that permits restoration of the mutant gene in all diseased photoreceptor cells, thereby ensuring sufficient transduction efficiency. We then treated mice at early, mid, or late disease stages. At all 3 time points, degeneration was halted and function was rescued for at least 1 year. Not only do our results demonstrate that gene therapy effectively preserves function after the onset of degeneration, our study also demonstrates that there is a broad therapeutic time window. Moreover, these results suggest that RP patients are treatable, despite most being diagnosed after substantial photoreceptor loss, and that gene therapy research must focus on improving transduction efficiency to maximize clinical impact.

Introduction

Retinitis pigmentosa (RP), a progressive neurodegenerative disease, is the most common cause of hereditary blindness in developed countries (1, 2). In the early stage of the disease, peripheral rod photoreceptor degeneration is underway, and patients experience night blindness; in addition, some cones exhibit structural pathologies (3). Rod degeneration continues in mid and late stages, causing tunnel vision. In the late stage, macular cone loss becomes significant enough that patients lose daylight vision and are often deemed legally blind (1).

There is currently no treatment for RP (indeed, for any retinal degenerative disease) that can halt the loss of both photoreceptors and function. Of those therapies on the horizon, gene therapy may be the most promising (4–6). Therapeutic development is negatively impacted by the fact that when the efficacy of a potential gene therapy is low, it is difficult to control for the various possible reasons. Failure might result if the means of delivering the therapeutic to its target cells is insufficient, or if a therapeutic is administered too late to halt the degeneration cascade. In the current study, we address these 2 potential limitations by first optimizing the means of delivery, and then by testing a gene therapeutic delivered under optimized conditions at multiple disease stages.

Currently, therapeutic genes are administered by subretinal injection. However, there are 3 significant problems with this approach. First, the therapeutic generally reaches only a fraction of the diseased cells. Second, injections are inherently highly variable, leading to large deviations in response and dictating the use of large numbers of animals to compensate for that deviation. Third, injections are invasive and can lead to side effects (e.g., trauma). Thus, a method that does not involve subretinal injection would greatly facilitate the study of disease mechanisms and the development of therapeutics.

A second major issue that impacts the development and testing of any type of therapeutic is the timing of treatment. In preclinical studies, gene therapeutics are almost always delivered before the onset of cell degeneration. However, most, if not all, neural degenerative diseases are diagnosed after the onset of degeneration (7). Therefore, a therapeutic must be able to demonstrate efficacy after disease onset—even at mid- or late-stage disease. However, it has been suggested that treatment at later stages might be hindered by a “point of no return,” beyond which photoreceptor cell death is unpreventable (8). An important research priority, therefore, is to determine the therapeutic time window in order to understand whether mid- or late-stage treatment is even feasible.

In the current study, we generated an RP mouse model in which mutant rod-specific cyclic GMP (cGMP) phosphodiesterase 6b (PDE6b) (Online Mendelian Inheritance in Man [OMIM] #180072) can be restored to WT levels in all rod photoreceptors. Mutations in the gene encoding PDE6B are responsible for 36,000 new cases of RP worldwide each year (1, 9–11). Using a tool
to achieve optimal gene delivery, we then systematically mapped the treatment time window for our RP gene therapeutic relative to disease progression. We demonstrate stable, sustained rescue of photoreceptor structure and visual function, even when the therapy was administered at mid-to-late disease stages.

Results
In the first part of the Results, we describe an RP mouse that allows the delivery of a therapeutic gene to all rod photoreceptors and to do so without subretinal injections. In the second part, we use this mouse model to determine whether RP can be treated well after the onset of photoreceptor cell loss.

An inducible rod-specific Cre driver. To generate a preclinical model for RP, in which the therapeutic gene can be delivered uniformly to all rods, we first generated a Pde6gCreERT2 mouse and then crossed it with our Pde6bStop Pde6bH62Q0 mice to derive Pde6b Stop Pde6bH62Q0 Pde6gCreERT2 mice (referred to herein as mutant). In the Pde6b Stop Pde6bH62Q0 mice, 1 allele carries a point mutation (Pde6bH62Q0) that dramatically reduces expression of PDE6b (12, 13). This hypomorphic rod-specific mutation is not as severe as a null mutation (e.g., Pde6brd1) (14) and, therefore, is a better model of human RP, which is usually caused by a missense mutation. The second allele carries a floxed stop cassette (Pde6bstop) that prevents gene expression in the absence of Cre recombinase activity. In addition to a stop codon, the stop cassette contains 3 polyadenylation signals (3).

To generate the Pde6gCreERT2 mouse, we inserted tamoxifen-inducible CreERT2 into the rod-specific Pde6g locus (Figure 1A). Insertion of CreERT2 was confirmed at the genomic DNA level by Southern blot analysis (Figure 1B). To confirm the cell specificity of the Pde6gCreERT2 recombinase, we crossed these mice with the ROSA26-lacZ reporter mouse line. ROSA26-lacZ reporter mice contain a loxP-flanked stop cassette preventing expression of the downstream lacZ gene; when crossed with a Cre Tg strain, the stop sequence is removed, and lacZ is expressed in cells where Cre is expressed (15). After i.p. injection of tamoxifen, the ROSA26-lacZ Pde6gCreERT2 double-Tg mice showed X-gal staining in photoreceptor cells, but not in cells in the inner nuclear layer (INL) or ganglion cell layer (GCL) (Figure 1C). The subcellular pattern of X-gal staining in photoreceptors — in the inner segment (IS) and synaptic terminals (STs) — has been previously described (16). These data validate functional Cre recombinase expression in photoreceptors.

To demonstrate that Pde6gCreERT2 can remove the stop cassette in the Pde6bStop Pde6bH62Q0 mice, we treated mutant mice with tamoxifen and then performed PCR analysis on DNA isolated from the retinal outer nuclear layer (ONL). A 362-bp band was found in DNA amplified from treated mutant mice, but not in DNA from untreated mice (Figure 1D).

Figure 1. Inducible rod-specific Pde6gCreERT2 driver line. (A) Targeting strategy for the generation of Pde6gCreERT2 mice. Exons 2–4 are represented by black boxes. Exons 2 and 3 were replaced by CreERT2. The targeting vector contains a neo/tk cassette (Neo). After homologous recombination, CreERT2 is placed under Pde6g locus control. (B) Southern blot analysis of EcoRI- and EcoRV-digested genomic DNA from 8 ES clones hybridized to the probe shown in A. The 8.5- and 6.6-kb bands represent WT and targeted Pde6g alleles, respectively. (C) X-gal–stained retinal section from a 4-week-old ROSA26-lacZ Pde6gCreERT2 mouse injected with tamoxifen at 2 weeks of age. Blue, lacZ reporter gene signal. Scale bar: 50 μm. RPE, retinal pigment epithelium. (D) PCR analysis of ONL DNA isolated from 4-week-old retinae identified a 284-bp fragment for WT, 284- and 415-bp fragments for an untreated mutant, and 284-, 415-, and 362-bp fragments for a mutant treated with tamoxifen at 2 weeks of age. (E) Retinal sections from 4-week-old mice immunostained for PDE6b; the treated mutant was tamoxifen injected at 2 weeks of age. Scale bar: 30 μm.
amplified from the ONL of untreated mutant or WT retinae (Figure 1D). A faint 415-bp nonrecombined product was also noted; this is because, in addition to rods, the ONL also contains cone nuclei, in which Pde6gCreERT2 recombinase is inactive. To analyze expression of PDE6b protein after tamoxifen-induced Pde6gCreERT2-mediated recombination, retinal sections were immunolabeled with an Ab against PDE6b. While immunolabeling was absent from mutant retinae, the outer segment (OS) of treated and WT retinae exhibited intense levels of staining (Figure 1E). These data confirm the efficient, near-complete tissue-specific Pde6gCreERT2-mediated recombination.

These reporter gene and target gene experiments demonstrate the high efficiency of our inducible rod-specific Cre driver. With this tool, we can restore PDE6b expression in all mutant rods to levels that are close to those of WT, and we can do so at specific time points. In the following experiments, in addition to our mutant mice, we used Pde6b+Pde6bH620Q, Pde6gCreERT2 mice as a WT (positive) control.

Photoreceptor degeneration in Pde6bH620Q, Pde6bH620Q, Pde6gCreERT2 mutant mice. To test our gene therapy after the onset of photoreceptor degeneration at early, mid, and late disease stages, we first needed to gain a basic understanding of the disease progression timeline in our mutant. To do this, we performed a quantitative histological analysis of the mutant and WT photoreceptor layer, known as the ONL, from mice that were 2, 4, or 8 weeks of age. We measured ONL thickness, cone numbers, and cone and rod photoreceptor OS lengths.

At 2 weeks of age, OS and ONL morphology in the mutant retinae (Supplemental Figure 1, A and D; supplemental material available online with this article; doi:10.1172/JCI82462DS1) were indistinguishable from that of age-matched WT control retinae (not shown). It is important to note that in 2-week-old retinae, developing retinal OS maturation and programmed photoreceptor cell death are ongoing (17, 18); as a result, the 2-week-old mutant rod OS was reduced, and the ONL was thicker than in WT adult retinae (Supplemental Figure 1, G and H). Cone OS lengths and cone numbers were not quantified at 2 weeks, since expression of the cone marker was low (Supplemental Figure 1D). At 4 weeks of age, rod OS length and ONL thickness were reduced by roughly 50% and 30%, respectively (<0.015 and <0.036) compared with WT adult retinae (Supplemental Figure 1, B and G). In contrast, cone OS length and cone numbers were not significantly changed (P = 0.65 and P = 0.37, respectively) (Supplemental Figure 1, E, I, and J). At 8 weeks of age, OS length and ONL thickness were dramatically decreased — by about 70% — compared with WT adult retinae (P < 0.009 and P < 0.001, respectively) (Supplemental Figure 1, C, G, and H). In addition, cone OS length and cone numbers were dramatically decreased — by roughly 65% and 30%, respectively (P < 0.003 and P < 0.023) (Supplemental Figure 1, F, I, and J). Given this progressive decrease in ONL thickness and OS length, we defined the 2-, 4-, and 8-week-old mutant retinae as being in early, mid, and late stages of RP disease.

Restoration of PDE6b expression and retinal function over a broad time window. To determine the therapeutic time window for photoreceptor rescue, we injected tamoxifen into 2-, 4-, and 8-week-old mice (i.e., at early, mid, and late disease stages, respectively). To determine whether PDE6b expression was rescued by the treatment, retinae were subjected to immunoblot analysis (Figure 2). PDE6b expression (i.e., a 90-kDa band) was observed in mutant mice treated at all 3 stages, but the level of expression was highest when tamoxifen was administered at 2 weeks of age. This progressive decrease in expression of rod-specific PDE6b correlates temporally with the decrease noted in rod OS length and ONL thickness (Supplemental Figure 1, A–C).

To assess the degree to which retinal function was rescued, we performed electroretinographic (ERG) analysis in 18-week-old mice. Under dark-adapted conditions with dim flashes, to derive rod responses, WT retinae exhibited a rod-specific b-wave (positive deflection), and mutant retinae showed no response; tamoxifen treatment rescued the rod-specific b-wave, with earlier treatment eliciting a higher response (Figure 3A). Under dark-adapted conditions with bright flashes, to derive mixed rod and cone responses, WT retinae exhibited an a-wave (negative deflection), followed by a b-wave. Mutant retinae lacked the a-wave and exhibited a strongly reduced b-wave; tamoxifen treatment led to nearly complete restoration of the a- and b-waves at the 2- and 4-week treatment time points and partial restoration at the 8-week time point (Figure 3A). Under light-adapted conditions with very bright flashes, to derive cone responses, WT retinae exhibited a cone-specific b-wave, which was strongly reduced in mutant mice; treatment at every stage rescued the cone responses (Figure 3A).

We quantified the amplitudes of the a- and b-waves, comparing treated and untreated mutants (Figure 3, B–E). Treatment at 2 and 4 weeks (i.e., early- and mid-stage disease, respectively) significantly increased the rod b-wave amplitude (P < 0.001). Treatment at 8 weeks (late-stage disease) appeared to rescue rod ERG responses, however, the effect was not statistically significant (P = 0.070) (Figure 3B). This is probably because approximately 70% of rods had died by 8 weeks (Supplemental Figure 1). The b-wave amplitudes in light-adapted ERG responses were significantly greater in treated mice at 2, 4, and 8 weeks of age (P < 0.003, P < 0.001, and P < 0.011, respectively); this indicates that treatment prevented the progressive degeneration of cones (Figure 3C). Treated mice also exhibited significantly greater mixed rod-cone a-wave amplitudes at all 3 treatment time points (2 and 4 weeks, P < 0.001; 8 weeks, P < 0.049) (Figure 3D). Similarly, the
The rhodopsin-positive rod OS manifests structurally as shortening, followed by loss of the rod itself (19). In WT retinae, intense rhodopsin staining was observed in the OS (Figure 4A), which was almost completely absent in mutant retinae (Figure 4B). OS length was comparable to that of WT retinae at 2 weeks (Figure 4C) and was also rescued at 4 and 8 weeks (Figure 4, D and E). This gene therapy prevented any further decrease in OS length.

This rod cell loss manifests structurally as decreased thickness of the ONL. We therefore stained sections of 20-week-old retinae with Hoechst dye, which labels all nuclei. Retinae treated at 2 and 4 weeks had thicker ONLs compared with those of untreated mutants (Figure 4). Retinae treated at 8 weeks had ONLs that were also thicker than those of untreated retinae, but the difference at this late stage was smaller, because the majority of rods had died by 8 weeks (Supplemental Figure 1). In RP, rod degeneration leads to the secondarily degeneration of cones. To specifically examine the response of cones to our treatment, retinal sections were immunostained for cone arrestin. In WT retinae, immunostaining was apparent in cone pedicles, axons, and in the cytoplasm of cone ISs and OSs (Figure 4F). In mutant retinae, cone arrestin immunostaining suggested that both the number of cones and the length of their OSs were reduced (Figure 4G). Tamoxifen injection at 2, 4, or 8 weeks of age preserved cone numbers and structure (Figure 4, H–J).

We next quantified the observed changes in OS length and ONL thickness. Rod OS length and ONL thickness were significantly greater in retinae of mice treated at either 2 or 4 weeks compared with retinae of untreated mutants (P < 0.001); on the other hand, there was no significant difference between mice treated at 8 weeks and untreated mutants (P = 0.7 and P = 0.2, respectively) (Figure 4, K and L). On the other hand, cone OS length was significantly greater in mice treated at 2 or 4 weeks (P < 0.001), or at 8 weeks (P < 0.05) (Figure 4M). Thus, while we were able to rescue the remaining photoreceptors even at late-stage disease, earlier treatment (when there are more photoreceptors) was associated with greater functional and structural rescue (Figures 3 and 4).
Absence of reactive gliosis after gene therapy. Reactive gliosis is an indicator of damage to the CNS, including damage caused by significant levels of cell degeneration. In retina, gliosis is indicated by upregulation of glial fibrillary acidic protein (GFAP) in Müller glial cells. To determine whether our gene therapy reduces retinal damage, we performed GFAP immunolabeling on sections of 20-week-old retinae. In WT retina, GFAP immunolabeling was apparent in the endfeet of Müller glia (Figure 5A). In untreated mutant retina, GFAP immunolabeling was increased, with intensely labeled stress fibers extending to the ONL (Figure 5B); this activation of Müller glia was apparent in 4-week-old untreated mutant retinae (but not in 2-week-old retinae) and was even more severe by 8 weeks (Supplemental Figure 2). In retinae of mutant mice treated at 2, 4, and 8 weeks of age, the GFAP immunolabeling pattern was comparable to that seen in WT retinae (Figure 5, C–E). Thus, treatment prevented Müller glial cell activation and infiltration at an early disease stage and reversed it at mid and late stages.

Long-term effects of treatment. Figures 2–5 demonstrate functional and structural rescue of diseased rods in our PDE6b-deficient RP mouse model. In addition to this direct rescue by gene therapy, data in Figures 3 and 4 provide evidence for indirect, sec-
secondary rescue of cones. To analyze whether these rescue effects are stable, and thus long lasting, we treated mice at 8 weeks of age and analyzed retinal function at 10 and 32 weeks of age and retinal structure at 10, 20, and 32 weeks of age.

In retinae of treated mutant mice, ERG responses at 10 and 32 weeks of age were not significantly different ($P = 0.8$, $P = 0.6$, and $P = 0.1$ for rod, cone, and mixed responses, respectively) (Figure 6). In contrast, in untreated mutant mice, cone and mixed b-wave amplitudes decreased significantly between 10 and 32 weeks ($P < 0.001$). Rod b-wave and mixed a-wave amplitudes (i.e., rod-based ERG measures) were quite low, and the modest decrease between 10 and 32 weeks was not significant ($P = 0.2$ and $P = 0.3$, respectively) (Figure 6). We also examined treatment stability by immunolabeling rod OSs with rhodopsin Ab (Figure 7, A–F) and cones with cone arrestin Ab (Figure 7, G–L); a nuclear stain was used to label the ONL. Over the 22-week study, mutant retinae exhibited rod OS shortening, progressive and dramatic loss of cones, and thinning of the ONL. In treated retinae, on the other hand, these structures did not appear to change. These experiments demonstrate that the gene therapy at a relatively late disease stage prevented degeneration of the remaining rods and cones for up to 6 months.

We have quantitated ONL thickness in WT and untreated mutant mice at 2, 4, and 8 weeks of age (Supplemental Figure 1), as well as in WT, untreated, and treated mutant mice at 20 weeks of age (Figure 4L). We next examined ONL thickness in WT, untreated mutant, and treated mutant mice at 1 year and then combined all of these data (Figure 8). In WT mice (Figure 8, blue line), the decrease in ONL thickness was due to normal programmed cell death of rods, which occurs from days 5 to 24 (17). In untreated mutant mice, ONL thickness decreased over a longer time frame (Figure 8, black line), reflecting the combination of normal developmental cell death and progressive degeneration of mutant photoreceptor cells, which are entirely gone by 1 year. Treatment at 2 weeks (Figure 8, yellow line) prevented the onset of degeneration caused by the mutation. Treatment at 4 and 8 weeks (Figure 8, green and red lines, respectively) prevented further degeneration. Earlier intervention was associated with the rescue of more photoreceptors.
In recent retinal gene therapy clinical trials, some of the research subjects showed functional improvement (23–30). However, several follow-up studies clearly showed that photoreceptor loss had not been halted or even slowed (31–34). It was suggested that there is a threshold of accumulated changes as a result of the mutation, after which photoreceptor death is inevitable (31); this has been called the “point of no return” (8). If such a threshold (i.e., point of no return) exists, our mouse data demonstrate that it would have to occur at very terminal disease stages — after our late stage (i.e., 8 weeks of age). Instead, our data suggest that the failure to rescue photoreceptors in humans, rather than being the result of a point of no return, is due to transduction efficiencies that are therapeutically insufficient. Insufficient transduction would allow abnormalities in untreated mutant cells to continue to stress both treated and untreated photoreceptors. This same logic could be used to explain the failure to halt degeneration in some previous animal studies (13, 35). In support of this idea are gene therapy studies demonstrating a delay in photoreceptor degeneration that was limited to the transduced area (36–38).

Although RP is typically diagnosed in young adulthood, photoreceptor degeneration has been detected in RP patients as young as 6 years of age (1). Therefore, for RP gene therapy, it is especially important to be able to demonstrate long-term efficacy — that is, stability that will last over the many remaining decades of a patient’s life. In addition, any therapy that establishes long-term rescue of cones could have enormous clinical impact, since humans spend most of their waking hours in light levels that are within the dynamic range of cones (i.e., cone loss is blinding). In the current study, we were able to demonstrate long-term stability of both functional and structural rescue (Figures 6–8). In fact, ours is the first study, to our knowledge, to demonstrate a stable long-term halt of rod and cone degeneration. There are probably multiple mechanisms underlying this non–cell-autonomous rescue of cones. First, our direct rescue of rods reduced the level of cell death in the RP retina, which may lead to an increase in positive trophic factors (39) and a decrease in toxic factors. For example, we demonstrated that gene therapy reverses Müller glial cell activation (Figure 5 and Supplemental Figure 2), which, by itself, is likely to exert a positive impact on cell survival (40).

Our findings in Pde6b-mutant mice are likely to be generalizable to humans, because our RP mouse model recapitulates much of human RP disease progression. Specifically, disease progression in both our mutant and RP patients is characterized by early loss of rods, followed by secondary loss of cones (1). In both our mutant and human RP patients, the structural and functional phenotypes manifest as changes in ONL thickness and ERG recordings (41). To develop an effective RP gene therapy for humans, our work points to the importance of improving delivery of gene therapeutics and delivering those therapeutics at the earliest point possible. To improve vector delivery (i.e., to increase both the area treated as well as the percentage of target cells treated), it is important to evaluate newer adeno-associated virus (AAV) vector variants and promoters, generated to enhance transduction efficiency and specificity (42, 43), and to test single versus repeated administration of these AAVs. In addition, a new AAV variant has been evolved to efficiently target cells in the retina from an intravitreal injection (44).
Our work strongly suggests that RP is treatable with gene therapy, even at late disease stages. On the other hand, at very advanced stages — when few or no photoreceptors remain — gene therapy is not an option, and other therapies (e.g., stem cell-based transplantations, ref. 45; optogenetics, refs. 46, 47; or retinal prostheses, ref. 48) could be considered. Further, our work suggests that to minimize cell loss and maximize visual function, it is critical to intervene with an effective gene therapy (such as ours) when photoreceptor loss is still modest. This can be accomplished with better diagnostic tools and education. The combination of improved delivery of gene therapies and earlier diagnosis promises to effectively halt the disease in its earliest stages.

**Methods**

**Animals.** ROSA26lacZ reporter mice (B6.129S4-Gt(Rosa)26Sor<sup>tm1Sor</sup>/J) were purchased from The Jackson Laboratory. Pde6b<sup>H620Q</sup> mice were re-derived via oviduct transfer from morulae provided by the European Mouse Mutant Archive (EMMA) (12, 13). Pde6b<sup>H620Q</sup> mice were generated in the Tsang laboratory, as previously described (3). Pde6g<sup>CreERT2</sup> mice were generated as described below. All mice were maintained in the Columbia University Pathogen-free Eye Institute Annex Animal Care Services Facility under a 12-hour light/12-hour dark cycle.

**Generation of Pde6g<sup>CreERT2</sup> driver mice.** KV1 embryonic stem (ES) cells were resuspended in saline at a density of 4 × 10<sup>7</sup> cells/ml and infected with a linearized targeting vector by electroporation (Bio-Rad electroporator, single pulse, 220 V and 960 μF) (3, 49, 50).

**Blastocysts.** Blastocysts were harvested from pregnant C57BL/6J mice at 3.5 days postcoitus by insertion of a syringe-attached needle into the cervix. Blastocysts were thawed, passaged once, washed with PBS, dissociated with 0.1 M PB, incubated for 5 minutes in 5 μg/ml Hoechst 33342 (catalog H1399; Sigma-Aldrich) was then injected at a concentration of 100 μg/ml, and then injected at a concentration of 100 μg/g BW on 3 consecutive days (1 injection/day), beginning at the time points indicated in the Results section.

**PCR and recombinant assay.** Tamoxifen (100 mg/ml in ethanol; catalog T5648; Sigma-Aldrich) was thoroughly mixed at 42°C, diluted with corn oil to a final concentration of 10 mg/ml, and then injected at a concentration of 100 μg/g BW on 3 consecutive days (1 injection/day), beginning at the time points indicated in the Results section.

**To assay for recombination, ONL DNA was extracted from 10-μm frozen sections using a 22-gauge needle and a surgical microscope (m690; Leica). PCR was performed as previously described (3). To target the Pde6b DNA sequence, we used 3 primers: forward primer, 5′-TGTCCTGTGGTCTGC-3′; reverse primer, 5′-TGGCGATGGAGTGCCTGA-3′; and internal primer, 5′-GTCCGTGACCGAGCTG-3′. The PCR-amplified vector sequence yielded 3 fragments: 284 bp for H620Q/WT, 415 bp for stop, and 362 bp after recombination.**

**Hc and morphometry.** Mice were euthanized according to established IACUC guidelines and previously described procedures (36). At that point, eyes were enucleated and placed in 4% paraformaldehyde for 5 minutes. Retinae were then dissected from the eyecup, fixed in 4% paraformaldehyde for 45 minutes, washed in 0.1 M phosphate buffer (PB), cryoprotected in 30% sucrose overnight, and embedded. Retinae were then sectioned vertically at 10 μm using a Leica cryostat. Sections were washed with 0.1 M PB and incubated overnight at 4°C in primary Ab in 0.1 M PB containing 5% ChemiBLOCKER (catalog 2170; EMD Millipore) and 0.3% Triton X-100. The following Abs were used: rabbit anti-cone arrestin (1:5,000; catalog AB15282; EMD Millipore); mouse anti-rhodopsin (1:500; catalog sc-57432; Santa Cruz Biotechnology Inc.); and Cy3-coupled anti-GFAP (1:800; catalog C9205; Sigma-Aldrich). Subsequently, sections were washed in 0.1 M PB, and goat anti-mouse Alexa 555-conjugated secondary Ab (1:500; catalog A-21425; Molecular Probes, Life Technologies) or goat anti-rabbit Alexa 488-conjugated secondary Ab (1:500; catalog A-11070; Molecular Probes, Life Technologies) was applied for 2 hours at room temperature. Sections were then washed with 0.1 M PB, incubated for 5 minutes in 5 μg/ml Hoechst 33342 (catalog H1399; Molecular Probes, Life Technologies), and analyzed using a confocal microscope (Nikon A1).

**Quantitative analyses of ONL thickness, OS lengths, and cone numbers were performed on sections containing the optic nerve (5 sections per eye). Measurements were performed 250 μm from the optic nerve using ImageJ software (NIH). Cones were counted in 200 × 200 μm fields.**

**LacZ staining.** ROSA26lacZ mice were bred with Pde6g<sup>CreERT2</sup> mice. Tamoxifen was injected into the 2-week-old offspring. Two weeks later, retinae were sectioned, stained with X-gal to detect β-gal activity (LacZ Tissue Staining Kit; InvivoGen), and analyzed using a Leica DM 5000B microscope.

**Immunoblot analysis.** Retinae were collected from 20-week-old mice, homogenized (51), and protein concentrations measured using the BCA protein assay (Thermo Scientific). Proteins were separated by SDS-PAGE (4%–15%) (Bio-Rad) and transferred to nitrocellulose (Bio-Rad). After blocking, membranes were incubated in rabbit anti-PDE6b Ab (1:1,000; catalog PA1-722; Thermo Scientific) overnight at 4°C, washed, and then incubated in goat anti-rabbit IgG-HRP secondary Ab (1:2,000; catalog sc-2004; Santa Cruz Biotechnology Inc.) for 1 hour at room temperature. β-actin loading control was detected using mouse anti-β-actin Ab (1:1,000; catalog ab125248; Abcam).
Membranes were visualized by chemiluminescence detection (EMD Millipore) using Biomax film (Kodak).

**ERG.** Mice were dark adapted overnight, and recordings were carried out under dim red light illumination. Mice were first anesthetized by i.p. injection of 0.1 ml/10 g BW of anesthetic solution (1 ml of 100 mg/ml ketamine and 0.1 ml of 20 mg/ml xylazine in 8.9 ml PBS). Body temperature was maintained at 37°C using a heating pad. Mouse pupils were dilated by topical administration of 2.5% phenylephrine hydrochloride and 1% tropicamide. Handmade electrodes were placed on the corneas, and gonioscopic prism solution (Alcon) was applied to each eye. ERG responses were recorded simultaneously from both eyes using an Espion Visual Electrophysiological System (Diagnosys). Rod and mixed rod-cone responses were recorded in dark-adapted mice using pulses of 0.001 and 3 cd/m² (white 6500 K), respectively. Then, mice were light adapted for photopic responses for at least 10 minutes in the Ganzfeld dome. Recordings were carried out under rod-suppressing continuous background illumination of 30 cd/m² (white 6500 K), and cone responses were recorded using pulses of 30 cd/m² (xenon).

**Statistics.** All data were analyzed using Stata 12 and R 3.1.1. Two-tailed t-tests were used to analyze the data, and P values of less than 0.05 were considered significant. To compare ERG data between groups at a single time point, we first averaged the ERG amplitude data for the 2 eyes of each mouse and then averaged the data for all of the mice within a single group (e.g., untreated mutants). To compare the mean ERG amplitudes between groups, a linear regression model was fit (where the outcome was the measurement and the predictors in the model were the group indicators), and pairwise comparisons between the groups were performed.

To compare ERG data between 2 time points (e.g., between weeks 10 and 32) for a single group, we carried out 2 distinct statistical analyses—one for mutants and a second for treated mutants. This is because while treated mutant mice were subjected to repeated ERG recordings, untreated mutant mice were subjected to a single ERG recording, at 1 time point. For treated mutant mice, we calculated the change over time for each eye and then calculated the average change per mouse; we then calculated the average for all mice within this group. In this way, we were able to study the within-subject changes for treated mutant mice. In untreated mutants, we first calculated the average ERG amplitude for the 2 eyes of each mouse, at the single time point, and then calculated the average for all mice within the mutant group. To compare mean outcomes between time points, we performed a 2-sample t test.

For analyses of OS lengths and ONL thickness, multiple sections were taken from a retina at 1 time point. For each measurement (e.g., rod OS length), values from multiple sections were averaged. Then, a linear regression model was fit, where the outcome was a structural measurement and the predictors were the group indicators, and pairwise comparisons between groups were performed.

**Study approval.** All experiments were approved by the IACUC of Columbia University. 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 and the Policy for the Use of Animals in Neuroscience Research of the Society for Neuroscience.

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Address correspondence to: Stephen H. Tsang, 635 W. 165th Street, 5th Floor New York, New York 10032, USA. Phone: 212.342.1189; E-mail: sht2@columbia.edu.