Kinase activity profiling identifies putative downstream targets of cGMP/PKG signaling in inherited retinal neurodegeneration

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Inherited retinal diseases (IRDs) are a group of neurodegenerative disorders that lead to photoreceptor cell death and eventually blindness. IRDs are characterised by a high genetic heterogeneity, making it imperative to design mutation-independent therapies. Mutations in a number of IRD disease genes have been associated with a rise of cyclic 3',5'-guanosine monophosphate (cGMP) levels in photoreceptors. Accordingly, the cGMP-dependent protein kinase (PKG) has emerged as a new potential target for the mutation-independent treatment of IRDs. However, the substrates of PKG and the downstream degenerative pathways triggered by its activity have yet to be determined. Here, we performed kinase activity profiling of different murine organotypic retinal explant cultures (diseased rd1 and wild-type controls) using multiplex peptide microarrays to identify proteins whose phosphorylation was significantly altered by PKG activity. In addition, we tested the downstream effect of a known PKG inhibitor CN03 in these organotypic retina cultures. Among the PKG substrates we identified were potassium channels belonging to the K1 family (KCNQ1, KCNQ2, KCNQ3, KCNQ6), cyclic AMP-responsive element-binding protein 1 (CREB1), DNA topoisomerase 2-α (TOP2A), α-phosphofructo-2-kinase/fructose-2,6-biphosphatase 3 (F263), and the glutamate ionotropic receptor kainate 2 (GRIK2). The retinal expression of these PKG targets was further confirmed by immunofluorescence and could be assigned to various neuronal cell types, including photoreceptors, horizontal cells, and ganglion cells. Taken together, this study confirmed the key role of PKG in photoreceptor cell death and identified new downstream targets of cGMP/PKG signalling that will improve the understanding of the degenerative mechanisms underlying IRDs.

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INTRODUCTION

Inherited retinal degeneration (IRD) relates to a genetically highly heterogeneous group of neurodegenerative diseases causing photoreceptor cell death and eventually blindness [1, 2]. To this day, in almost all cases, these diseases are untreatable. Causative mutations have been identified in over 300 different disease genes (https://sph.uth.edu/retnet; information retrieved October 2021), calling for the development of mutation-independent therapies. Mutations in more than 20 IRD disease genes have been linked to increased levels of cyclic 3',5'-guanosine monophosphate (cGMP) in photoreceptors [3] and are thought to affect at least 30% of IRD patients [4]. A key effector of cGMP-signalling is cGMP-dependent protein kinase (PKG), the overactivation of which may trigger photoreceptor cell death [5, 6].

Mammals possess two different genes encoding for PKG, prkg1 and prkg2 [7]. Splicing of prkg1 leads to two distinct isoforms – PKG1α and PKG1β – which differ in first 80 to 100 amino acids in their N-terminal. The prkg2 gene gives rise to only one isoform called PKG2. PKGs exist as a homodimer and binding of cGMP to one of the four cGMP binding sites induces a conformational change which activates the kinase [7]. Activated PKG phosphorylates numerous cellular proteins at serine/threonine amino acid positions, which in turn regulates numerous cellular pathways. In mammals, PKG1 regulates smooth muscle contraction [8], platelet activation and adhesion [9], cardiac function [10], feedback of the NO-signalling pathways [11], and various processes in the central nervous system, such as hippocampal and cerebellar learning [12]. PKG2 is involved in translocation of CFTR channels in jejunum [13] and regulation of bone growth by activation of kinases such as MAPK3/ERK1 and MAPK1/ERK2 in mechanically stimulated osteoblasts [14].

Intriguingly, PKG also plays an important role in cell death, which has been ascertained, for instance, through studies where PKG activation inhibited tumour progression in colon cancers, breast cancers, ovarian cancers and melanoma [15, 16]. Furthermore, PKG seems to play a central role in photoreceptor degeneration [17–19]. In the rd1 mouse retina, a well characterised model for IRD, photoreceptor cell death is triggered by abnormally high concentrations of retinal cGMP. This event is linked to dysfunction of phosphodiesterase 6 (PDE6) – involved in the regulation of intracellular cGMP levels – caused by a nonsense mutation in the rod Pde6b gene [20]. Increased cGMP signalling has been found in several other models for IRDs [4, 17] and is likely to over-activate PKG [3]. In vitro and in vivo pharmacological inhibition of PKG showed strong photoreceptor protection in rd1 retina as well as in the retina of rd2 and rd10 mouse models [3, 21]. Together, these studies suggest a key role for PKG activity in cGMP-mediated cell
death and highlight PKG as a potential common target for strategies aiming to reduce photoreceptor degeneration. 

The pathways downstream of PKG in degenerating photoreceptors are nonetheless still poorly understood. Increased cGMP/PKG signalling has been associated with increased activity of poly-ADP-ribose-polymerase (PARP), histone deacetylase (HDAC), and calpain proteases, all known to be involved in photoreceptor cell death [17]. However, to date there is no evidence that directly links these events to PKG. Intracellular changes of known PKG targets such as vasodilator-stimulated phosphoprotein (VASP) and cAMP response element-binding protein (CREB) have been observed in dying photoreceptors [19, 22, 23]. While this can be a direct consequence of excessive cGMP/PKG signalling, these targets may also be phosphorylated by or have phosphorylation sites for other kinases, including cAMP-dependent protein kinase (PKA), characterised by substrate motifs similar to those of PKG [24]. A better insight into the downstream effects of PKG and its phosphorylation targets is needed to understand the mechanisms of photoreceptor cell death and to, furthermore, guide the development of both new neuroprotective strategies and biomarker applications.

Using multiplex peptide microarrays, PamChips®, we measured PKG1- and PKG2- mediated phosphorylation of specific peptides by kinases in the lysates of murine retinal explant cultures treated or not with the PKG inhibitor CN03 [21]. We identified several new PKG substrates potentially connected to IRD and confirmed their retinal expression in murine tissue. This study thus provides the groundwork for future studies aimed at the elucidation of cGMP/PKG-dependent cell death pathways.

RESULTS

PKG inhibition reduces photoreceptor cell death in rd1 retinal explants

To investigate potential targets of PKG and their possible role in photoreceptor cell death, we used the PKG inhibitor CN03 on rd1 organotypic retinal explant cultures. CN03 is a cGMP analogue, and as such is able to bind to cGMP binding sites on PKG, without inducing the conformational changes required for kinase activation [25]. This culminates in reversible and competitive inhibition of PKG. In previous studies, CN03 showed marked protection of photoreceptors in retinal explants derived from the rd1 mouse model [21, 26]. We therefore collected wild-type (WT) and rd1 retinal explants treated or not with 50 µM CN03, using a treatment paradigm based on the aforementioned studies. Thus, retinas were explanted at postnatal (P) day 5 when photoreceptor degeneration had not yet started. The CN03 treatment was given at P7 and P9 and cultures were terminated at P11. The latter time-point corresponds to the beginning of rd1 photoreceptor cell death [27], and is thus well suited to assess the protective effects of a given compound and to study events downstream of abnormal cGMP/PKG signalling. We confirmed the protective effects of CN03, by characterising the degree of cell death using the TUNEL assay (Fig. 1). CN03-treated retinas showed marked photoreceptor protection as previously reported [21].

Serine/Threonine Kinase (STK) activity in rd1 retinal explants

To evaluate possible differences in kinase profiles of the murine retinal explant samples (rd1, n = 8; WT, n = 5), we used PamChip® peptide microarray-based Serine/Threonine Kinase (STK) activity assays [24]. The overall STK activity between the samples is represented as heatmap (Fig. S1a). A violin plot was used to visualize the phosphorylation signal intensity of the peptides and its distribution within the same sample groups (Fig. 2a). Increased phosphorylation was observed for 43% of the total 142 peptides in rd1 retinal explants, indicating higher kinase activity in rd1 NT when compared to WT (Fig. 2b). The peptides, SRC8_CHICK_423_435, RBL2_959_971, CDN18_151_163 and RADI_559_569 showed significantly higher phosphorylation (p < 0.05) in rd1 explants than in wild type controls. Using the upstream kinase analysis tool of the BioNavigator® software, the peptides with increased phosphorylation were linked to kinases that are most likely to be responsible for
The phosphorylation of these peptides (See Data Analysis, Materials and Methods Section). The kinase statistics and kinase score were calculated as a metrics for identifying highly active kinases. The kinases that were predicted to be more active in rd1 explants include CaMK4, PKG1, PKG2, PKAα, and Pim1 (Table 1). In order to present relative kinome activity profiles of retinal explants (rd1 vs. WT), kinase score and kinase statistics were used in color coding branches and nodes on the phylogenetic tree of protein kinase families (Fig. 2c).

STK activity in rd1 retinal explants treated with PKG inhibitor CN03
As we found a higher phosphorylation of peptides in rd1 explants, when compared to WT explants, and with a clear role for PKGs observed, we sought to investigate the effect of the PKG inhibitor CN03 on the STK activity. The PKG inhibitor CN03 significantly decreased photoreceptor cell death (cf. Fig. 1). The overall STK activity profiles for treated and untreated rd1 retinal explants are shown in a heatmap (Fig. S1b). The distribution of phosphorylated peptides for both the samples is represented by a violin plot (Fig. 3a). Phosphorylation decreased for approximately 80% of the 142 peptides present on the STK PamChip®. Fourteen peptides were identified whose phosphorylation decreased significantly (p < 0.05) in rd1 CN03 as compared to untreated rd1 (Fig. 3b). Table 2 shows peptides that displayed lower phosphorylation (22 peptides, p < 0.1) in CN03 treated rd1 retina than untreated controls, p-value, names of the proteins they are derived from, with their UniProt IDs, substrate score of PKG1 and PKG2, and localization within the retina.

We subsequently linked the phosphorylated peptides to the putative upstream kinases and found that kinase activity of particularly PKAα, Pim1, PKG1, PKG2, CaMK4 was suggested to be

Fig. 2  Serine/Threonine Kinase (STK) activity in untreated retinal explants. Organotypic retinal explants derived from wild-type (WT) and rd1 mice (WT, n = 5; rd1, n = 8) were maintained in culture medium from P5 till P11. The kinase activity of their lysates was measured on PamChip® Serine/Threonine kinase (STK) arrays. a Violin plot showing the global phosphorylation of the peptides on PamChip® STK array as Log2 signal intensity and their intensity value distribution, when comparing WT to rd1 explants. The thick line connects the average values of each group. b Volcano plot representing Log Fold Change (LFC) and −Log10 p-value for peptide phosphorylation. Red dots indicate significantly changed phosphopeptides (p-value < 0.05), black dots represent peptides with no significant alteration in phosphorylation. c The high-ranking kinases were visualized in a kinome phylogenetic tree, where branch and node color are encoded according to the kinase statistic, with values > 0 (in red) representing higher kinase activity in rd1 retinal explants. The node size is encoded by the kinase score, that ranks kinases based on their significance and specificity in terms of sets of peptides used for the corresponding kinase.
To identify possible associations of the kinases with biological pathways that are activated in retinal degeneration as represented by rd1 explants, we performed an analysis of relevant biochemical pathways using the Kyoto Encyclopedia of Genes and Genomes database (KEGG; Version 2021). Pathway analysis of kinase activity in rd1 vs. WT explants yielded as the major associated pathways Neurotrophin signaling pathway, proteoglycans in cancer, circadian entrainment, insulin resistance, HIF-1 signaling pathway, long-term potentiation, FMAPK signaling (Fig. 4a). After treatment of rd1 explants with CN03, the high scoring pathways i.e., insulin resistance, mTOR, MAPK signaling, long-term potentiation, circadian entrainment, and HIF-1 signaling (Fig. 4b) were the same as found in Fig. 4a.

To identify the putative kinases involved in retinal degeneration, based on the rd1 model and PKG inhibitor CN03 treatment, we compared the kinase list derived from each comparison as represented in a Venn diagram (Fig. 4c). Here, almost 77% of kinases with high activity in rd1 as compared to WT were overlapping with the kinases showing reduced activity in rd1 retinal explants treated with CN03.

### Table 1. Upstream kinase analysis results for rd1 vs. WT and rd1 CN03 vs. rd1.

| Rank | Kinase Name | Kinase Score | Kinase Statistic |
|------|-------------|--------------|-----------------|
| 1.   | CaMK4      | 3.39         | 0.46            |
| 2.   | PKG1       | 3.23         | 0.32            |
| 3.   | PKG2       | 3.16         | 0.30            |
| 4.   | PKAα       | 3.09         | 0.26            |
| 5.   | Pim1       | 2.02         | 0.19            |
| 6.   | p70S6Kβ    | 1.78         | 0.24            |
| 7.   | CaMK2α     | 1.92         | 0.46            |
| 8.   | RSK1/p90RSK | 1.92     | 0.42            |
| 9.   | MSK2       | 1.89         | 0.52            |
| 10.  | PKD1       | 1.95         | 0.43            |
| 11.  | PKCβ       | 1.77         | 0.23            |
| 12.  | Pim2       | 1.70         | 0.21            |
| 13.  | PRKX       | 1.82         | 0.23            |
| 14.  | ROCK2      | 1.69         | -0.37           |
| 15.  | MSK1       | 1.71         | 0.36            |
| 16.  | Pim3       | 1.68         | 0.18            |

| Rank | Kinase Name | Kinase Score | Kinase Statistic |
|------|-------------|--------------|-----------------|
| 37.  | Akt1        | 1.70         | -0.30           |
| 38.  | DCAMKL1     | 1.70         | -0.50           |
| 39.  | PKCβ       | 1.70         | -0.40           |
| 40.  | Akt2        | 1.70         | -0.30           |
| 41.  | COT         | 2.60         | -0.40           |
| 42.  | PKCβ       | 1.60         | -0.40           |
| 43.  | RSKL1       | 1.60         | -0.40           |

The kinase statistic shows the overall change of the peptide set that represents a given kinase. Positive values indicate higher kinase activity in rd1, while negative values indicate lower activity, e.g., in rd1 CN03. The kinase score includes the sum of significance and specificity score (Scores > 1.5 shown in the table).

Reduced by CN03 treatment (Table 1, Fig. 3c). Notably, these were the same kinases that were predicted to be more active in the diseased rd1 explants in comparison to WT explants, indicating specific targeting by CN03 (Fig. 2c).

### Putative biological pathways involved in retinal degeneration

To identify possible associations of the kinases with biological pathways that are activated in retinal degeneration as represented by rd1 explants, we performed an analysis of relevant biochemical pathways using the Kyoto Encyclopedia of Genes and Genomes database (KEGG; Version 2021). Pathway analysis of kinase activity in rd1 vs. WT explants yielded as the major associated pathways Neurotrophin signaling pathway, proteoglycans in cancer, circadian entrainment, insulin resistance, HIF-1 signaling pathway, long-term potentiation, FMAPK signaling (Fig. 4a). After treatment of rd1 explants with CN03, the high scoring pathways i.e., insulin resistance, mTOR, MAPK signaling, long-term potentiation, circadian entrainment, and HIF-1 signaling (Fig. 4b) were the same as found in Fig. 4a.

To identify the putative kinases involved in retinal degeneration, based on the rd1 model and PKG inhibitor CN03 treatment, we compared the kinase list derived from each comparison as represented in a Venn diagram (Fig. 4c). Here, almost 77% of kinases with high activity in rd1 as compared to WT were overlapping with the kinases showing reduced activity in rd1 retinal explants treated with CN03.

### PKG target validation for retinal localization

Next, we tested whether the proteins corresponding to the peptides that were differentially phosphorylated in the three retinal explant groups (i.e., WT, rd1, and rd1/CN03) were present in the retina. While some of these proteins had already previously been shown to be present in mouse retina (Table 2), we performed immunostaining on retinal tissue sections derived from P11 WT mice to assess the retinal expression and cellular localization for six proteins with high PKG preference (Table 2), that might potentially be connected to photoreceptor degeneration.

The analysis of the WT retinal sections immunostained for cyclic AMP-responsive element-binding protein 1 (CREB1) showed abundant expression in the INL as well as the GCL (Fig. 5). The potassium voltage-gated channel subfamily A, member 3 (Kv1.3; KCNA3) was found to be localised in the ONL, possibly in photoreceptor axons, and in two discrete sublamina of the IPL [28], where the synapses between bipolar cell axons and ganglion cell dendrites reside. The potassium voltage-gated channel subfamily A, member 6 (Kv1.6; KCNA6) was prominently expressed in the ganglion cell layer (GCL) and the nerve fibre layer (NFL).
DNA topoisomerase 2-α (TOP2A) expression was restricted to the GCL. 6-phosphofructo-2-kinase/fructose-2,6-biphosphatase 3 (F263) – an enzyme involved in the control of glycolytic flux [29] appeared to be present in the OPL, likely in horizontal cells, as well as in the GCL. Finally, staining with the antibody directed against glutamate ionotropic receptor kainate 2 (GRIK2) confirmed its presence in the INL and GCL, in line with previous findings [30].

Taken together, the immunostaining data as well as the signal intensity in the distribution plots confirmed the retinal expression of several of the discovered PKG targets [24] (Fig. 5a, b). Furthermore, the differences in phosphorylation of these targets between WT, rd1, and rd1/CN03 retinas (Figs. 2 and 3) may suggest a potential role for these PKG substrates in the mechanism leading to photoreceptor cell death.

**DISCUSSION**

Excessive activity of PKG has been directly linked to retinal degeneration and its inhibition has been shown to provide photoreceptor protection in several in vivo IRD models [3, 21]. Yet, at present it is still unclear how PKG exerts its detrimental effects and what are the protein targets that, when phosphorylated by PKG, mediate photoreceptor cell death. Here, we combined PKG inhibitor treatment with multiplex peptide microarray technology and immunohistochemistry to identify novel PKG phosphorylation targets in the retina.

**PKG inhibition mediates photoreceptor neuroprotection**

Over-activation of PKG has been connected to neuronal cell death in different experimental settings and conditions, including in human neuroblastoma derived cell cultures [31], peripheral nerve injury [32], and in photoreceptor degeneration in IRD [6, 33]. For photoreceptor degeneration an excessive accumulation of cGMP was already established in the 1970s and while it had already then become evident that high cGMP-levels were cytotoxic [34, 35], it was unclear how cGMP would exert its negative effects. Research initially focused on a proposed detrimental role of cyclic-nucleotide-gated (CNG) channels [36, 37], however, in recent years it has become increasingly obvious that PKG-signalling plays a major role in photoreceptor degeneration [3, 38]. Our study,
Table 2. Differentially phosphorylated peptides ($p < 0.1$) in rd1 untreated vs. rd1 CN03 treated retinal explant cultures.

| S. No. | ID      | UniProt ID | Protein name                              | p-value | PKG1 score | PKG2 score | Localisation        | Ref. |
|--------|---------|------------|-------------------------------------------|---------|------------|------------|---------------------|------|
| 1.     | IKBA_26_38 | P25963     | NF-kappa-B inhibitor alpha                 | 0.016   | 0          | 0          | ONL, INL, GCL       | [81] |
| 2.     | GBRB2_427_439 | P47870    | Gamma-aminobutyric acid receptor subunit beta-2 precursor | 0.026   | 10         | 6          |                     |      |
| 3.     | PTN12_32_44 | Q05209     | Tyrosine-protein phosphatase non-receptor type 12 | 0.031   | 10         | 5          |                     |      |
| 4.     | KCNA6_504_516 | P17658    | Potassium voltage-gated channel subfamily A member 6 | 0.032   | 9          | 3          | GCL                 | [42] |
| 5.     | NCF1_296_308 | P14598     | Neutrophil cytosol factor 1               | 0.032   | 9          | 7          | GCL                 | [82] |
| 6.     | CREB1_126_138 | P16220    | cAMP response element-binding protein     | 0.034   | 10         | 6          | ONL, INL, GCL       |      |
| 7.     | GRK2_708_720 | Q13002     | Glutamate receptor, ionotropic kainate 2 precursor | 0.035   | 10         | 4          | INL, IPL            | [30] |
| 8.     | NCF1_321_333 | P14598     | Neutrophil cytosol factor 1               | 0.036   | 0          | 0          |                     |      |
| 9.     | EPB42_241_253 | P16452    | Erythrocyte membrane protein band 4.2     | 0.037   | 9          | 7          |                     |      |
| 10.    | VTNC_390_402 | P04004     | Vitronecin precursor                      | 0.037   | 10         | 4          | RPE, ONL, GCL       | [83] |
| 11.    | ADRB2_338_350 | P07550    | Beta-2 adrenergic receptor                | 0.039   | 8          | 4          | Müller cells        | [84] |
| 12.    | BRCA1_1451_1463 | P38398   | Breast cancer type 1 susceptibility protein | 0.039   | 0          | 0          |                     | [85] |
| 13.    | TOP2A_1463_1475 | P11388   | DNA topoisomerase 2-alpha                | 0.039   | 6          | 5          | GCL                 |      |
| 14.    | MIP3_208_220 | P30307     | M-phase inducer phosphatase 3             | 0.041   | 0          | 0          |                     |      |
| 15.    | CDC2_154_169 | P06493     | Cell division control protein 2 homolog   | 0.046   | 0          | 0          |                     |      |
| 16.    | KAP3_107_119 | P31323     | cAMP-dependent protein kinase type II-beta regulatory subunit | 0.046   | 8          | 4          |                     |      |
| 17.    | MYP3_268_280 | Q14896     | Myosin-binding protein C, cardiac-type    | 0.049   | 10         | 4          |                     |      |
| 18.    | RYR1_4317_4329 | P21817   | Ryanodine receptor 1                     | 0.053   | 9          | 5          | ONL, INL, GCL       | [86] |
| 19.    | TY3H_65_77    | P07101     | Tyrosine 3-monoxygenase                   | 0.054   | 9          | 3          |                     |      |
| 20.    | KCNA3_461_473 | P22001    | Potassium voltage-gated channel subfamily A member 3 | 0.055   | 8          | 7          | ONL, IPL            | [42] |
| 21.    | F263_454_466 | Q16875     | 6-phosphofructo-2-kinase/fructose-2,6-biphasate 3 | 0.059   | 10         | 3          | INL, GCL            |      |
| 22.    | VASP_150_162 | P50552     | Vasodilator-stimulated phosphoprotein     | 0.081   | 8          | 4          | ONL                 | [23] |

The table shows the peptide name, name of the protein that the peptide is derived from, its UniProt ID, p-value, PKG1 and PKG2 substrate score, and localization in the retina with reference. The scores for PKG1, PKG2 are as described in [24], and range from 0 (no substrate) to 10 (very good substrate). Colour scheme for the PKG specificity score: 10–8: Good, 7–4: Intermediate, and 3–1: Poor substrate, 0: no substrate [24]. For proteins previously detected in the mouse retina, the reference is added. The expression of proteins printed in bold was studied using immunodetection (cf. Fig. 5).
Fig. 4  Biological pathways involved in retinal degeneration. Key biological pathways with potentially higher activity in rd1 (a) and lower activity in rd1 treated with CN03 (b). Pathways are ranked according to their p-values and colored by their q-values. A q value is the p-value adjusted for multiple testing using the Benjamini-Hochberg procedure [79, 80]. The node size indicates the gene ratio, i.e., the percentage of total genes or proteins in the given KEGG pathways (only input genes or proteins with at least one KEGG pathway were included in the calculation). The overlap between kinases changed in rd1 NT vs. WT and vs. rd1 treated with CN03 respectively (Figs. 2c, 3c), is visualized in a Venn Diagram. Here, increased or reduced activity of a kinase is indicated by arrows pointing up- or down-wards, respectively.

Fig. 5  Presence and localisation of PKG target proteins in the retina. a The panel shows retinal cross-sections derived from P11 WT mice and stained with secondary antibody for negative control (neg. ctrl.), anti-CREB1, anti-KCNA3, anti-KCNA6, anti-TOP2A, anti-F263, or anti-GRIK2. b The localisation of CREB1, KCNA3, KCNA6, TOP2A, F263, and GRIK2 in the retina of WT mice is illustrated by a signal distribution plot along vertical sections across the retina. Antibody labelling obtained on stained retinal tissue sections from three different animals is represented in different shades of green, while the negative control is represented in grey. OS/IS Outer segment/inner segment, ONL Outer nuclear layer, OPL Outer plexiform layer, INL Inner nuclear layer, IPL Inner plexiform layer, GCL Ganglion cell layer, NFL Nerve fiber layer.
In the retina, inhibition of KCNA1 and KCNA3 in vivo was different from CD95 (CD95) our study [43]. It has been shown that within a few minutes of play a role in regulating cell death, making them interesting for surprisingly, these proliferation-related channels also appear to apoptotic volume, also known as cell shrinkage in lymphocytes [45]. In the retina, inhibition of KCNA1 and KCNA3 in vivo was protective for retinal ganglion cells (RGCs) after optic nerve axotomy [46]. The localization of KCNA6 and KCNA3 in the WT mouse retina and their differential phosphorylation by PKG in rd1 and CN03 treated rd1 retinal explant cultures suggest a role in photoreceptor cell death. Past studies on the involvement of K1 channels in cell death, although conflicting, strengthen this idea [43–45]. Establishing whether PKG-mediated phosphorylation of K1 channels involves activation or inhibition of these channels would help shed light on their contribution to photoreceptor death.

Among the various differentially phosphorylated substrates located in the retina of both rd1 and WT, CREB1 is a known target of PKG [47, 48]. CREB1 is a transcription factor implicated in neuronal survival [49] and constitutively expressed in different types of human cancer [50]. As shown by the ex vivo results, CREB1 is widely distributed in the mouse retina. Furthermore, through upstream kinase analysis we predicted the activation of several other kinases such as CaMK4, PKC theta, and PKA, known to target CREB [51–53]. Thus, the abnormal activity of PKG, as well as that of the other kinases, can lead to increased CREB phosphorylation, in contrast to previous studies in which downregulation of CREB1 was associated with photoreceptor degeneration in both rd1 and rd10 mice [22, 54]. The increase in activity of kinases that phosphorylate CREB combined with an increase in CREB peptide phosphorylation in rd1 may due to the activation of parallel signals after the cellular insult: the induction of cell death and the activation of a CREB1-directed survival program [55].

Other PKG targets that we have identified and localized in the retina include vitronectin (VTNC) and F263. VTNC is a cell adhesion protein, upregulated in inflammation and traumatized tissues [56] and a major component of extracellular deposits specific to age-related macular degeneration (AMD) [57]. In the same study, VTNC mRNA expression has been found distributed in the various layers of the human retina. Its high phosphorylation in rd1 explants might be linked to increased inflammation, which is an early phenomenon observed for degenerative retinal disorders such as RP, AMD, and diabetic retinopathy (DR) [58]. If so, VTNC would have the potential to be a predictive biomarker for certain retinal degenerative diseases.

6-phosphofructo-2-kinase/fructose-2,6-biphosphatase 3 or F263 is a pro-glycolytic enzyme. Its activation following excitotoxic stimulation has been associated with neuronal cell death [59]. Furthermore, dysregulation of the HIF-1–F263 pathway has been proposed as crucial for two key aspects of the pathogenesis of DR, namely angiogenesis and neurodegeneration [60]. With ex vivo analysis we localized F263 in GCL and INL while it was not detected in ONL. The identification of F263 as a target of PKG and its high phosphorylation in rd1 compared to WT may suggest a change in retinal energy metabolism during degeneration and may also help to understand the mechanism of aerobic glycolysis in the retina, which remains relatively poorly understood to date [61].

**Novel targets for PKG in the neuroretina**

One of the identified targets in our analysis was KCNA6, which, together with KCNA3 (rank 20 based on p value, Table 2), belongs to the K1 family of voltage-dependent potassium channels. The K1 family mainly consists of channels with a slow and delayed activation. Immunoreactivity studies revealed expression of both KCNA3 and KCNA6 in the mouse retina [42], which is in line with our immunofluorescence results. The K1 family is involved in the regulation of progression through cell cycle checkpoints by defining the membrane potential and ensuring the driving force for calcium and chloride entry. Surprisingly, these proliferation-related channels also appear to play a role in regulating cell death, making them interesting for our study [43]. It has been shown that within a few minutes of apoptosis induction KCNA3 is inhibited by the cluster of differentiation 95 (CD95) via tyrosine phosphorylation [44]. Other studies have shown the activation of KCNA3 at an early stage of apoptosis, as well as its contribution to a decrease in apoptotic volume, also known as cell shrinkage in lymphocytes [45]. In the retina, inhibition of KCNA1 and KCNA3 in vivo was protective for retinal ganglion cells (RGCs) after optic nerve axotomy [46]. The localization of KCNA6 and KCNA3 in the WT mouse retina and their differential phosphorylation by PKG in rd1 and CN03 treated rd1 retinal explant cultures suggest a role in photoreceptor cell death. Past studies on the involvement of K1 channels in cell death, although conflicting, strengthen this idea [43–45]. Establishing whether PKG-mediated phosphorylation of K1 channels involves activation or inhibition of these channels would help shed light on their contribution to photoreceptor death.

Among the various differentially phosphorylated substrates located in the retina of both rd1 and WT, CREB1 is a known target of PKG [47, 48]. CREB1 is a transcription factor implicated in neuronal survival [49] and constitutively expressed in different types of human cancer [50]. As shown by the ex vivo results, CREB1 is widely distributed in the mouse retina. Furthermore, through upstream kinase analysis we predicted the activation of several other kinases such as CaMK4, PKC theta, and PKA, known to target CREB [51–53]. Thus, the abnormal activity of PKG, as well as that of the other kinases, can lead to increased CREB1 phosphorylation, in contrast to previous studies in which downregulation of CREB1 was associated with photoreceptor degeneration in both rd1 and rd10 mice [22, 54]. The increase in activity of kinases that phosphorylate CREB combined with an increase in CREB peptide phosphorylation in rd1 may due to the activation of parallel signals after the cellular insult: the induction of cell death and the activation of a CREB1-directed survival program [55].

Other PKG targets that we have identified and localized in the retina include vitronectin (VTNC) and F263. VTNC is a cell adhesion protein, upregulated in inflammation and traumatized tissues [56] and a major component of extracellular deposits specific to age-related macular degeneration (AMD) [57]. In the same study, VTNC mRNA expression has been found distributed in the various layers of the human retina. Its high phosphorylation in rd1 explants might be linked to increased inflammation, which is an early phenomenon observed for degenerative retinal disorders such as RP, AMD, and diabetic retinopathy (DR) [58]. If so, VTNC would have the potential to be a predictive biomarker for certain retinal degenerative diseases.

6-phosphofructo-2-kinase/fructose-2,6-biphosphatase 3 or F263 is a pro-glycolytic enzyme. Its activation following excitotoxic stimulation has been associated with neuronal cell death [59]. Furthermore, dysregulation of the HIF-1–F263 pathway has been proposed as crucial for two key aspects of the pathogenesis of DR, namely angiogenesis and neurodegeneration [60]. With ex vivo analysis we localized F263 in GCL and INL while it was not detected in ONL. The identification of F263 as a target of PKG and its high phosphorylation in rd1 compared to WT may suggest a change in retinal energy metabolism during degeneration and may also help to understand the mechanism of aerobic glycolysis in the retina, which remains relatively poorly understood to date [61].

**Metabolic pathways in the retina regulated by PKG**

The pathway analysis showed insulin resistance and mTOR as important pathways inhibited by CN03. mTOR is a serine/threonine kinase which regulates protein synthesis, cellular metabolism and autophagy [62]. The mTOR pathway regulates neurogenesis in the eye and is crucial to normal development of retina and the optic nerve [63]. Activated mTOR signaling has been shown to be involved in retinal neurodegenerative diseases such as DR and AMD [64, 65]. Treatment with the mTOR inhibitor rapamycin, improved mitochondrial dysfunction and provided neuroprotection to 661 W cells, a cellular model that shares certain features with photoreceptors [66]. Similarly, inhibition of the mTOR/PARP-1 axis leads to photoreceptor protection against light-induced photoreceptor cell death [62]. However, the targeting of this pathway axis in IRD as treatment strategy is still
contentious as there are also studies that show mTOR stimulation delays cone cell death in IRD [67, 68].

Among the peptides whose phosphorylation was high in untreated rd1 retinal explants and significantly decreased in CN03-treated rd1 retinal explants was F263. The protein F263 is a key regulator of glycolysis [69]. Photoreceptors metabolize glucose through aerobic glycolysis (the ‘Warburg effect’) to satisfy their very high energy demand for the maintenance of the dark current, as well as for the recycling of visual pigments and the renewal of photoreceptor OS [70, 71]. As a consequence of the high energy demand, the retina has an elevated oxygen consumption, probably the highest in the body, and is therefore particularly prone to oxidative stress and reactive oxygen species (ROS)-induced mitochondrial damage [72]. The high phosphorylation of F263 found in the rd1 retina may be linked to an abnormal increase in metabolic activity in the retina resulting in mitochondrial damage. Considering that mitochondrial dysfunction has been associated with RP [72], the identification of F263 as a substrate of PKG and its possible role in the degeneration process could help to clarify the metabolic state of the diseased retina.

CONCLUSION
Using multiplex peptide microarray technology, we provide further evidence for the likely involvement of PKG in degenerating rd1 photoreceptors in vitro and confirm the already known neuroprotective effects of the PKG inhibitor CN03 [21]. Importantly, we identified several novel downstream PKG targets that might play a role in cGMP/PKG-mediated photoreceptor degeneration. This will form the basis for future studies, which may further elucidate the role of PKG target phosphorylation, as well as the beneficial effects of CN03. Our work could likewise be employed to develop novel diagnostic and therapeutic biomarkers for retinal degenerative diseases. For example, the use of phospho-specific antibodies may allow to confirm a reduction of target phosphorylation and in situ studies may allow to localize in which cellular compartment such changes occurred. Finally, our results further connect several metabolic pathways with retinal degeneration, including insulin, mTOR, and HIF-1 signaling, which in future studies may help to understand the complex mechanisms behind photoreceptor cell death.

MATERIALS AND METHODS
Animals
C3H Pde6b<sup>rd1</sup>/rd1 (rd1) and congenic C3H wild-type (WT) mice were housed under standard light conditions, had free access to food and water, and were used irrespective of gender. All procedures were performed in accordance with the law on animal protection issued by the German Federal Government (Tierschutzgesetz) and approved by the institutional animal welfare office of the University of Tübingen.

cGMP analogues synthesis
Synthesis of the cyclic nucleotide analogue CN03 was performed by Biolog Life Science Institute GmbH & Co. KG according to previously described methods [21] (https://patentscope.wipo.int/search/en/detail.jsf?docId=WO2018010965).

Organotypic retinal explant cultures
Preparation of organotypic retinal cultures derived from rd1 (<i>n</i> = 8-10) and C3H (<i>n</i> = 5) animals chosen randomly out of three different litters was performed as described previously [26, 73]. Animals were sacrificed at postnatal day (P)15. Eyes were rapidly enucleated and incubated in R16 serum-free, antibiotic-free culture medium (07491252 A; Gibco) with 0.12% protease K (21935025; ICN Biomedicals Inc.) for 15 min at 37 °C. Subsequently, eyes were incubated in 20% foetal bovine serum (FCS) (F7524; Sigma) in order to block protease K activity. This step was followed by rinsing in R16 medium. Under a laminar-flow hood and sterile conditions, the anterior segment, lens, vitreous, sclera, and choroid were removed from the eyes. The retina with the RPE still attached was cut in four points resembling a four-leaf clover and transferred to a culture membrane insert (3412; Corning Life Sciences) in a six-well culture plate with completed R16 medium with supplements [26]. The retinal explants were incubated at 37 °C in a humidified 5% CO<sub>2</sub> incubator and left undisturbed for 48 h. At P7 and P9 medium was changed <i>i.e.</i> with replacing the full volume of the complete R16 medium 1 ml per dish, with fresh medium. In this context, half of the retinal explants were treated with CN03 at 50 µM (dissolved in water), while the other half was kept as untreated control. For retinal explant lysis, culturing was stopped at P11, retinal explants were snap frozen in liquid nitrogen and stored at −80 °C. For retinal explant cross-sectioning preparation, culturing was stopped at P11 by 45 min fixation in 4% paraformaldehyde (PFA), cryoprotected with graded sucrose solutions containing 10, 20, and 30% sucrose, embedded in optimal cutting temperature compound (Tissue-Tek) and then cut into 12 µm sections.

TUNEL assay
Representative results showing the protective effects of CN03 on photoreceptor cell death at a concentration of 50 µM, were obtained using terminal deoxynucleotidyl transferase dUTP nick end labelling (TUNEL) assay [74] (based on in situ CellDeath Detection Kit, 11683; U.S. Patent no. 5391950, red fluorescence; Sigma-Aldrich) on sections derived from rd1 and C3H retinal explant cultures. DAPI (Vecstahash Antifade Mounting Medium with DAPI; Vector Laboratories) was used as blue fluorescence nuclear counterstain. Images were captured using 7 Z-stacks with maximum intensity projection (MIP) on a Zeiss Axios Imager Z1 ApoTome Microscope MM dual digital camera (Zeiss, Oberkochen, Germany) with a 20x APOCHRO-MAT objective. The excitation (<i>λ</i>exc) and emission (<i>λ</i>em) characteristics of the filter sets used for the fluorophores were as follows: DAPI (<i>λ</i>exc = 369 nm, <i>λ</i>em = 465 nm) and TMR red (<i>λ</i>exc = 562 nm, <i>λ</i>em = 640 nm), Adobe Photoshop (CSSAdobe Systems Incorporated, San Jose, CA) was used for image processing.

Materials for retinal explant lysis
Mammalian protein extraction reagent (M-PER<sup>Pam</sup>®), Halt<sup>TM</sup> protease and phosphatase inhibitor cocktails and the Coomassie Plus (Bradford Assay Kit were purchased from Thermo Fischer Scientific.

Retinal explant lysis
The retinal explant samples were lysed with lysis buffer (MPER with 1:100 phosphatase inhibitor cocktail and protease inhibitor cocktail reagents) for 30 min on ice. The lysate was centrifuged at 16000 x g for 15 min at 4 °C. The supernatant was immediately aliquoted, flash frozen, and stored at −80 °C. The protein content of the lysate was measured using the Bradford Protein Assay [75].

Kinase activity measurements
The kinase activity for the retina lysates was determined on STK PamChip<sup>®</sup> with four arrays, each array comprising of 142 peptides derived from the human phosphoproteome, according to the instructions of the manufacturer (PamGene International B.V., ‘s-Hertogenbosch, North Brabant, The Netherlands). The peptide names consist of the protein they are derived from and the first and last amino acid positions in that protein. The phosphorylated Serine/Threonine amino acid residues are detected by a primary antibody mix, which is then made visible by addition of FITC-conjugated secondary antibody [76]. The assay mix consisted of protein kinase buffer (PamGene International BV, ‘s-Hertogenbosch, North Brabant, The Netherlands), 0.01% BSA, STK primary antibody mix, ATP (400 µM) and retina tissue lysate (0.25 µg protein/array).

Instrumentation for kinase activity measurements
All experiments were performed on PamStation<sup>12</sup>® where up to 12 assays can be performed simultaneously (PamGene International B.V., ‘s-Hertogenbosch, North Brabant, The Netherlands). To prevent unspecific antibody binding, the PamChips<sup>®</sup> were first blocked with 2% BSA, by pumping it up and down 30 times through the arrays. The chips were then washed three times with Protein Kinase Buffer and assay mix was applied. The assay mix was pumped up and down through the arrays for 60 min. Afterwards, the arrays were washed and FITC labelled secondary antibody mix was applied on the arrays. The images of the arrays were recorded at multiple exposure times [77].
Data analysis

The signal intensity of each peptide spot on the array for each time point was quantified by BioNavigator® software version 6.3.67.0 (PamGene International B.V., S’Hertogenbosch, North Brabant, The Netherlands). For each spot, the signal intensity at the different exposure times was combined to a single value by exposure time scaling [77]. The resulting values were log2-transformed and the overall differences in STK profile between rd1 NT vs. WT or rd1 CN03 vs. rd1 NT were visualized as heatmaps and violin plots which were generated in R software (R version 4.0.2, The R Foundation for Statistical Computing). For heatmaps, hierarchical clustering of peptides was performed using the average-linkage method and clustering was shown as dendrograms on the y-axis of heatmaps. Unpaired t-tests were used to determine significant differences (p < 0.05) in phosphorylation intensity between the groups. Results were represented as volcano plots (GraphPad Prism version 9.2.0).

Upstream kinase analysis

Information on kinases that could be responsible for peptide phosphorylation differences between the two sample groups was obtained through the STK Upstream Kinase Analysis tool of BioNavigator® [77]. This software integrates known interactions between kinases and the phosphorylation sites and is provided in databases such as HIPRD, PhosphoELM, Phosphosite-PLUS, Reactome, UniProt to provide a peptide set for each kinase (fingerprint) and predicts the kinases differentially active between the groups. The results of this analysis are described by two parameters: The Kinase Statistic, indicating the size and direction of the change for each kinase and the Kinase Score, ranking the kinases by the likelihood of this kinase being involved.

The Kinase Statistic depicts the overall change of the peptide set that represents a kinase. For instance, a larger positive value indicates a larger kinase activity in either rd1 explants in comparison to WT or CN03 treated explants in comparison to untreated explants. The Kinase Score is the result of two permutation analyses. The Kinase Score is calculated by addition of the Significance Score and the Specificity Score. The Significance Score indicates the significance of the change represented by the Kinase Statistic between two groups (using 500 permutations across sample labels). The Specificity Score indicates the specificity of the Kinase Statistic with respect to the number of peptides used for predicting the corresponding kinase (using 500 permutations across target peptides).

The kinases are ranked by the Kinase Score. The highest-ranking predicted kinases from the significant STK peptide sets are represented on a phylogenetic tree of the human protein kinase family generated in Coral, a web-based application http://phanstiel-lab.med.ucnc.edu/CORAL/ [78].

Pathway analysis

Pathway analysis was performed in Enrichr (https://maayanlab.cloud/Enrichr/), which is a comprehensive resource for curated gene sets and a search engine that accumulates biological knowledge for further biological discoveries [79, 80]. We used differentially phosphorylated peptides and predicted upstream kinases as input list for Enrichr analysis. Visualization was performed in GraphPad Prism (version 9) using known matrices from the analysis.

Histology

For retinal cross-section preparation, the eyes were marked nasally and cornea, iris, lens, and vitreous were carefully removed. The remaining eyecups were fixed in 4% PFA for 2 h at room temperature. Incubation with graded sucrose solutions was performed for cryoprotection. Eyes were embedded in Tissue-Tek and cut into 14 μm sections. Immunostaining was performed on retinal cross-sections derived from 3 different mice by incubating with primary antibody against CREB1 (1:200; Proteintech), KCNMA1 (1:200; Alomone labs), KCNA6 (1:300; Alomone labs), F263 (1:100; Abcam), TOP2A (1:500; Proteintech), GRIK2 (1:100; Invitrogen) diluted in blocking solution at 4 °C overnight. The negative control was obtained by incubating the retinal cross-sections at 4 °C overnight with the blocking solution used for the primary antibody. Alexa Fluor 488 antibody was used as secondary antibody. Sections were mounted using DAPI. Images were captured using 9 Z-stacks with maximum intensity projection (MIP) on a Zeiss Axio Imager Z1 ApoTome Microscope MRm digital camera (Zeiss, Oberkochen, Germany) with a 20x APOCHROMAT objective. The excitation (λex) / emission (λem) characteristics of the filter sets used for the fluorophores were as follows: DAPI (λex = 369 nm, λem = 465 nm) and AF488 (λex = 490 nm, λem = 525 nm). Adobe Photoshop (CS5, Adobe Systems Incorporated, San Jose, CA) was used for image processing.

Histological image analysis

Fluorescence intensity data were generated from images of retinal sections derived from 3 different wild-type (WT) P11 mice, using the image profiling function included in the Zen software (Zeiss). Based on the fluorescence intensity data, a signal intensity distribution plot was generated in which each antibody was compared to its corresponding negative control (Fig. 5B). To account for the high variability inherent to immunostaining, the relative signal intensity was also calculated (Fig. 5S). The bar graphs represent the ratio of the mean signal intensity for a single antibody (antigen) in a given retinal layer divided by the negative control value. The ratios obtained are expressed as mean ± SD. Graphs were prepared using Prism 8 for Windows (GraphPad Software).

DATA AVAILABILITY

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

REFERENCES

1. Galan A, Chiziolini M, Milan E, Sebastiani A, Costaglia I, Carmaggiore F. Good epidemiologic practice in retinits pigmentosa: From phenotyping to biobanking. Curr. Genomics. 2011;12:3-6.
2. Farrar GJ, Carrigan M, Dockery A, Millington-Ward S, Palfi A, Chadderton N, et al. To unravel an excitement of the molecular genetics of inherited retinal degenerations. Hum. Mol. Genet. 2017;26:621–11.
3. Paquet-Durand F, Hauck SM, Van Veen T, Ueffing M, Ekström P. PKG activity causes photoreceptor cell death in two retinitis pigmentosa models. J. Neurochem. 2009;108:796–810.
4. Power M, Das S, Schütze K, Marigo V, Ekström P, Paquet-Durand F. Cellular mechanisms of hereditary photoreceptor degeneration — Focus on cGMP. Retin Eye Res. 2020;74:100772.
5. Paquet-Durand F, Hauck SM, van Veen T, Ueffing M, Ekström P. PKG activity causes photoreceptor cell death in two retinitis pigmentosa models. J. Neurochem. 2009;108:796–810.
6. Koch M, Scheel C, Ma H, Yang F, Stadlmieier M, Glück AF, et al. The cGMP-dependent protein kinase 2 contributes to cone photoreceptor degeneration in the Cnga3-deficient mouse model of achromatopsia. Int J. Mol. Sci. 2021;22:52.
7. Lorenz R, Bertinetti D, Herberg FW. cAMP-dependent protein kinase and cGMP-dependent protein kinase as cyclic nucleotide effectors. In: Non-canonical cyclic nucleotides. 2015:105–22.
8. Tang M, Wang G, Lu P, Karas BH, Aronovitz M, Hesmer EP, et al. Regulator of G-protein signaling-2 mediates vascular smooth muscle relaxation and blood pressure. Nat. Med. 2003;9:1506–12.
9. Arai M, Von Brühl ML, Eisleberger C, Werner M, Konrad I, Kocher T, et al. IRAG mediates NO/cGMP-dependent inhibition of platelet aggregation and thrombus formation. Blood 2007;109:352–9.
10. Prusyazhnya O, Wolhuter K, Switzer C, Santos C, Yang X, Lynham S, et al. Blood pressure-lowering by the antioxidant resveratrol is counterintuitively mediated by oxidation of cGMP-dependent protein kinase. Circulation 2019;140:126–37.
11. Francis SH, Busch JL, Corbin JD. cGMP-dependent protein kinase and cGMP phosphodiesterases in nitric oxide and cGMP action. Pharmacol. Rev. 2010;62:525–63.
12. Feil S, Zimmermann P, Knorr A, Brummer S, Schlossmann J, Hofmann F, et al. Distribution of cGMP-dependent protein kinase type 1 and its isofoms in the mouse brain and retina. Neurosciencen 2005;135:863–8.
13. French PJ, Bijnan J, Edishove M, Vaandrager AB, Scholte BJ, Lohmann SM, et al. Isotype-specific activation of cyclic fibrosis transmembrane conductance regulator-chloride channels by cGMP-dependent protein kinase II. J. Biol. Chem. 1995;270:26626–31.
14. Rangaswami H, Marathe N, Zhuang S, Chen Y, Yeh JC, Frangos JA, et al. Type II cGMP-dependent protein kinase mediates osteoblast mechanotransduction. J. Biol. Chem. 2009;284:14796–808.
15. Tolone A, Belhaj S, Rentsch A, Schwede F, Paquet-Durand F. The CMAP pathway and inherited photoreceptor degeneration: Targets, compounds, and biomarkers. Genes (Basel). 2019;10:1–16.
16. Quadri M, Comitato A, Palazzo E, Tiso N, Rentsch A, Pellacani G, et al. Activation of cGMP-dependent protein kinase restricts melanoma growth and invasion by interfering with the EGF/EGFR Pathway. J. Invest. Dermatol. 2021.
17. Arango-Gonzalez B, Trufunovic D, Sahaboglu A, Kranz K, Michalakis S, Farinelli P, et al. Identification of a common non-apoptotic cell death mechanism in hereditary retinal degeneration. PLoS One. 2014;9:1–11.
18. Faber DB, Park S, Yamashita C. Cyclic GMP-phosphodiesterase of rd retina: Biosynthesis and content. Exp. Eye Res. 1988;46:363–74.
36. Paquet-Durand F, Beck S, Michalakis S, Goldmann T, Huber G, Mühlfriedel R, et al. Cyclic guanosine monophosphate: Elevation in degenerative rod outer segments of the pde6b rd1 mouse retina. Mol. Vis. 2020;20:1–13.

37. Sahabaglu A, Paquet-Durand O, Dietter J, Dengler K, Bernhard-Kurz S, Eckström PAR, et al. Retinitis pigmentosa: Rapid neurodegeneration is governed by slow cell death mechanisms. Cell Death Dis. 2013;4:e488–e488.

38. Celesia GG, DeMarco PJ. Anatomy and physiology of the visual system. J. Clin. Neuroophthalmol. 1994;14:882–92.

39. Lu Y, Zhang L, Zhu R, Zhou H, Fan H, Wang Q, PKFKB3, a key glucose metabolic enzyme regulated by pathogen recognition receptor TLR4 in liver cells. Ther. Adv. Endocrinol. Metab. 2020;20:42018829023474.

40. Vila A, Whitaker CM, O'Brien J. Membrane-associated guanylate kinase scaffolds organize a horizontal cell synaptic complex restricted to invaginating contacts with photoreceptors. J. Comp. Neurol. 2017;525:605–617.

41. Zhuoniero LMT, Adornetto A, Secondo A, Magi S, Dell'Avvessano C, Scorzelli A, et al. Involvement of the nitric oxide/protein kinase G pathway in polychlorinated biphenyl-induced cell death in SH-SY5Y neuroblastoma cells. J. Neurosci. Res. 1998;69:1–21.

42. González-Feroro D, Portillo F, Gómez L, Montero F, Kasparov S, Moreno-López B. Inhibition of resting potassium conductances by long-term activation of the NO/cGMP/protein kinase G pathway: A new mechanism regulating neuronal excitability. J. Neurosci. 2007;27:6302–6.

43. Xu J, Morris L, Thapa A, Ma H, Michalakis S, Biel M, et al. cGMP accumulation causes photoreceptor degeneration in CNG channel deficiency: Evidence of cGMP cytotoxicity independently of enhanced CNG channel function. J. Neurosci. 2013;33:14939–48.

44. Farber DB, Lolley RN. Cyclic guanosine monophosphate: Elevation in degenerating photoreceptors of the C3H mouse retina. Science 1974;186:449–51.

45. Lolley RN, Farber DB, Rayborn ME, Holfiedy JC. Cyclic gmp accumulation causes degeneration of photoreceptor cells: Simulation of an inherited disease. Science 1977;196:664–6.

46. Barobas P, Beck S, Michalakis S, Goldmann T, Huber G, Mühlfriedel R, et al. A key role for cyclic nucleotide gated (CNG) channels in CNG-channel-related retinopathies. PLoS Genet. 2013;12:1–13.

47. Barabás P, Peck CC, Krizaj D. Do calcium channel blockers rescue dying photoreceptors in the rd1 mouse? J. Neurochem. 2002;80:1302–14.

48. Aversano C, Scorziello A, Riva I, Aversano A, Pizzuti A, Sonveaux P. Anticancer targets in the NO/soluble guanylate cyclase/cGMP system on the functions of human T lymphocytes. J. Biol. Chem. 2003;278:14539–14.

49. Zrenner E, et al. Photoreceptor cell death mechanisms in inherited retinal degeneration. Mol. Neurobiol. 2008;38:253–69.

50. Fan B, Li FQ, Zuo L, Li GY. mTOR inhibition attenuates glucose deprivation-induced cell death in rod outer segments of the pde6b rd1 mouse retina. Front. Cell. 2020;8:1571.

51. Sinha D, Valapala M, Shang P, Hose S, Grebe R, Lutty GA, et al. Lysosomes: Regulators of autophagy in the retinal pigmented epithelium. Exp. Eye Res. 2013;144:46–53.

52. Nakahara T, Morita A, Yagasaki R, Mori A, Sakamoto K. Mammalian Target of Rapamycin (mTOR) as a potential therapeutic target in pathological ocular angiogenesis. Biol. Pharm. Bull. 2017;40:2045–9.

53. Fan B, Li FQ, Zou L, Li GY. mTOR inhibition attenuates glucose deprivation-induced cell death in photoreceptors via suppressing a mitochondria-dependent apoptotic pathway. Neurochem Int. 2016;99:178–86.

54. Arroba AI, Campos-Caro A, Aguilar-Diosdado M, Valverde AM. IGF-1, in inflammation and retinal degeneration: A close network. Front. Aging Neurosci. 2018;10:203.

55. Hurley JB. Retina metabolism and metabolism in the pigmented epithelium: A busy intersection. Am. Rev. Respir. Dis. 1991;143:1–9.

56. Rasmussen M, Welinder C, Schwede F, Ekström P. The cGMP system in normal and degenerating mouse neuroretina: New proteins with cGMP interaction potential identified by a proteomics approach. J. Neurosci. 2011;31:21873–85.

57. Porporato PE, Dhup S, Dadhich RK, Copetti T, Sonveaux P. Anticancer targets in the glycocalyx metabolism of tumors: A comprehensive review. Front. Pharmac. 2012;2:49.

58. Zhang R, Shen W, Du J, Gillies MC. Selective knockdown of hexokinase 2 in rods leads to age-related photoreceptor degeneration and retinal metabolic remodelling. Cell Death Dis. 2020;11:1–15.
71. Léveillard T, Sahel JA. Metabolic and redox signaling in the retina. Cell Mol. Life Sci. 2017;74:3649–65.
72. Moreno ML, Mérida S, Bosch-Morell F, Miranda M, Villar VM. Autophagy dysfunction and oxidative stress, two related mechanisms implicated in retinitis pigmentosa. Front. Physiol. 2018;9:1008.
73. Caffé AR, Ahuja P, Holmqvist B, Azadi S, Forsell J, Holmqvist I, et al. Mouse retina explants after long-term culture in serum free medium. J. Chem. Neuroanat. 2002;22:263–73.
74. Loo DT. In situ detection of apoptosis by the TUNEL assay: An overview of techniques. Methods Mol. Biol. 2011;682:3–13.
75. Zor T, Selinger Z. Linearization of the Bradford protein assay increases its sensitivity: Theoretical and experimental studies. Anal. Biochem. 1996;236:902–8.
76. Hilhorst R, Houkes L, Mommersteeg M, Musch J, Van Den Berg A, Ruijtenbeek R. Peptide microarrays for profiling of serine/threonine kinase activity of recombinant kinases and lysates of cells and tissue samples. Methods Mol. Biol. 2013;977:259–71.
77. Chirumamilla CS, Fazil MHUT, Perez-Novo C, et al. Profiling activity of cellular kinases in migrating T-cells. Methods Mol. Biol. 2019;1930:99–113.
78. Metz KS, Deoudes EM, Berginski ME, Jimenez-Ruiz I, Aksoy BA, Hammerbacher J, et al. Coral: Clear and customizable visualization of human kinome data. Cell Syst. 2018;7:347–50.
79. Xie Z, Bailey A, Kuleshov MV, Clarke DJB, Evangelista JE, Jenkins SL, et al. Gene set knowledge discovery with enrichr. Curr. Protoc. 2021;1:e90.
80. Kuleshov MV, Jones MR, Rouillard AD, Fernandez NF, Duan Q, Wang Z, et al. Enrichr: A comprehensive gene set enrichment analysis web server 2016 update. Nucleic Acids Res. 2016;44:W90–7.
81. Zeng HY, Tso MOM, Lai S, Lai H. Activation of nuclear factor-κB during retinal degeneration in rd mice. Mol. Vis. 2008;14:1075.
82. Dvoriantchikova G, Grant J, Santos ARC, Hernandez E, Ivanov D. Neuronal NAD(P)H oxidases contribute to ROS production and mediate RGC death after Ischemia. Invest. Ophthalmo. Mol. Sci. 2018;7:347–350.
83. Anderson DH, Hageman GS, Mullins RF, Neitz M, Neitz J, Ozaki S, et al. Vitronectin receptor function and oxidative stress, two related mechanisms implicated in retinitis pigmentosa. Front. Physiol. 2018;9:1008.
84. Cammalleri M, Dal Monte M, Miranda M, Villar VM. Autophagy dysfunction and oxidative stress, two related mechanisms implicated in retinitis pigmentosa. Front. Physiol. 2018;9:1008.
85. Ganguly A, Shields CL. Differential gene expression profile of retinoblastoma compared to normal retina. Mol. Vis. 2010;16:1292.
86. Huang W, Xing W, Ryskamp DA, Punzo C, Križaj D. Localization and phenotype-specific expression of ryanodine calcium release channels in C57BL6 and DBA/2J mouse strains. Exp. Eye Res. 2011;93:700–9.

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AUTHOR CONTRIBUTIONS
Conceptualization, JG, TT, and FPD; methodology, AR, AT; writing—original draft preparation, AR, AT, RH, TT, JG, and FPD; writing—review and editing, AR, AT, RH, TT, JG, and FPD; supervision, JG, TT, and FPD; funding acquisition, JG and FPD.

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AR, TT, RH, JG are current or former employees of PamGene International B.V., ‘s-Hertogenbosch, The Netherlands. FPD is shareholder of the company Mireca Medicines, Tübingen, Germany, which intends to forward clinical testing of CN03.

ADDITIONAL INFORMATION
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