AAV-CRB2 protects against vision loss in an inducible CRB1 retinitis pigmentosa mouse model

Thilo M. Buck,1 Rogier M. Vos,2 C. Henrique Alves,1,3 and Jan Wijnholds1,2

1Department of Ophthalmology, Leiden University Medical Center (LUMC), 2333 ZC Leiden, the Netherlands; 2Netherlands Institute of Neuroscience, Royal Netherlands Academy of Arts and Sciences (KNAW), 1105 BA Amsterdam, the Netherlands

Loss of Crumbs homolog 1 (CRB1) or CRB2 proteins in Müller cells or photoreceptors in the mouse retina results in a CRB dose-dependent retinal phenotype. In this study, we present a novel Müller cell-specific Crb1KO Crb2LowMGC retinitis pigmentosa mouse model (complete loss of CRB1 and reduced levels of CRB2 specifically in Müller cells). The Crb double mutant mice showed deficits in electroretinography, optokinetic head tracking, and retinal morphology. Exposure of retinas to low levels of DL-α-aminoadipate acid induced gliosis and retinal disorganization in Crb1KO Crb2LowMGC retinas but not in wild-type or Crb1-deficient retinas. Crb1KO Crb2LowMGC mice showed a substantial decrease in inner/outer photoreceptor segment length and optokinetic head-tracking response. Intravitreal application of rAAV vectors expressing human CRB2 (hCRB2) in Müller cells of Crb1KO Crb2LowMGC mice subsequently exposed to low levels of DL-α-aminoadipate acid prevented loss of vision, whereas recombinant adeno-associated viral (rAAV) vectors expressing human CRB1 (hCRB1) did not. Both rAAV vectors partially protected the morphology of the retina. The results suggest that hCRB expression in Müller cells is vital for control of retinal cell adhesion at the outer limiting membrane, and that the rAAV-cytomegalovirus (CMV)-hCRB vector is more potent than rAAV-minimal CMV (CMVmin)-hCRB1 in protection against loss of vision.

INTRODUCTION
Mutations in the Crumbs homolog 1 (CRB1) gene are associated with retinitis pigmentosa (RP), Leber congenital amaurosis (LCA), and cone-rod dystrophies and are sporadically found in foveal retinoschisis and macular dystrophy.1-3 The human and nonhuman primate retina express and localize CRB1 and CRB2 proteins in Müller glial cells (MGCs) and photoreceptor cells (PRCs) at the outer limiting membrane (OLM).1-6 The mouse retina also expresses and localizes the CRB2 protein at the OLM in MGCs and PRCs. However, whereas the mouse retina does express and localize the CRB1 protein at the OLM in MGCs and retinal progenitor cells, the mouse retina does not express the CRB1 protein in PRCs. The human and nonhuman primate retinas express a CRB1 protein of 1,406 aa, whereas the mouse retina expresses a CRB1 protein of 1,405 aa.7,8 Loss of the CRB1 or the CRB2 protein in the retina results in loss of adhesion between MGCs, between PRCs, and among MGCs and PRCs.5,6,10-12 No therapy is available for the treatment of CRB1-related retinal dystrophies. Recombinant adeno-associated viral (rAAV) vector-mediated gene supplementation may provide a lasting therapy to CRB1 RP patients. We previously showed that CRB2 can rescue retinas lacking CRB1 or CRB2 proteins from retinal degeneration in two fast-progression RP mouse models by increasing the levels of CRB2 into both MGCs and PRCs.13 However, rescue at mid-stage retinal disease could not be achieved with CRB1 cDNA supplementation, or by supplementation of CRB2 cDNA only in PRCs or only in MGCs. In the present study, we developed a sensitive RP MGC-specific mouse model to test for protection at early stage retinal disease by rAAV-human CRB (hCRB) gene therapy vectors.

We also investigated the development of the mouse retinal phenotype to explore the window of opportunity for rAAV-hCRB gene therapy. Previously, we analyzed Crb-related retinal degeneration mouse models. (1) The knockout (KO) of the Crb1 gene (Crb1KO) ablates the expression of the CRB1 protein in MGCs and retinal precursor cells, and it resulted in slow progression of retinal disorganization and degeneration from postnatal day 14 (P14) on.9,10,14 (2) In Crb2KO mice, retinal degeneration occurs at foci in the inferior temporal quadrant of the retina. (3) Cell-type specific ablation of CRB1 in MGCs, or of CRB2 in MGCs, suggested that CRB proteins execute important overlapping roles in MGCs. Loss of CRB1 protein in mouse MGCs, or the loss of CRB2 in MGCs, results in disruptions at the OLM, protrusion of rows of photoreceptor nuclei into the photoreceptor inner and outer segment layers, and ingresses of rows of photoreceptor nuclei into the outer plexiform layer (OPL). These retinas mimic RP in which the retinal degeneration process remains slow during the period of 1 year.9,6,11,15,16 (4) Most importantly, the complete loss of CRB2 in MGCs in Crb1KO mice worsened the retinal
phenotype from a RP-like to LCA-like phenotype. These \( Cbh^{1KO}Crb^{2\text{LMGC}} \) retinas lacking CRB1 and CRB2 specifically in MGCS show in addition to protrusion of photoreceptor nuclei into the segment layers, also an intertwining of photoreceptors with inner retinal cells. The \( Cbh^{1KO} \) RP and \( Cbh^{1KO}Crb^{2\text{LMGC}} \) retinas are not suitable for testing gene therapy vectors since the onset of retinal degeneration is too slow, whereas the \( Cbh^{1KO}Crb^{2\text{LMGC}} \) MGC-KO LCA model is not suitable to test gene therapy vectors because the onset of retinal degeneration occurs during retinal development and is too fast. In this study, we analyzed a novel mouse model by reducing endogenous mouse CRB2 (mCRB2) expression in MGCS from one instead of two \( Cbh^{1KO} \) alleles in \( Cbh^{1KO}Crb^{2\text{LowMGC}} \). Compared to littermate control \( Cbh^{1KO}Crb^{2\text{Flox/Flox}} \) (floxed/wild-type \( Cbh^{2} \)) the \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) showed a worsened retinal phenotype; therefore, we used these mice to test rAAV human \( CRB1 \) (hCRB1) and human CRB2 (hCRB2) gene therapy vectors that specifically target the MGCS.

Many underlying diseases show a nominal phenotype until a stressor triggers an escalation. DL-\( \alpha \)-amino adipic acid (DL-AAA) or cysteine/glutamate-specific antiporter inhibitor, decreasing the reserve pool of cysteine and glutathione (GSH) in MGCS. Low doses of DL-AAA disrupt the distal Müller glial sealing at the OLM by downregulating the adherens junction-associated protein zonula occludens-1 (ZO-1). The decrease in adhesion mediated by DL-AAA intravitreal injection is linked to photoreceptor nuclei protrusions in control mice and in the S334ter-line-3 rat model of RP. First, a decrease of glial fibrillary acidic protein (GFAP) was found 3 days after intravitreal injection of DL-AAA in the S334ter-line-3 rat model of RP, followed by an upregulation of GFAP 2 weeks later. Subretinal injection of DL-AAA to nonhuman primates caused a reduction in photoreceptor nuclei and a decrease in the electroretinography (ERG) response. The long-term effects of low doses of DL-AAA on vision-guided behavior in RP models have not been thoroughly investigated.

Here, we studied the effects of DL-AAA on retinas with decreased levels of CRB2 in MGCS lacking CRB1 (\( Cbh^{1KO}Crb^{2\text{LowMGC}} \)) compared to retinas with normal levels of CRB2 in MGCS lacking CRB1 (\( Cbh^{1KO}Crb^{2\text{Flox/Flox}} \)). We challenged the following mice on a 99.9% C57BL/6J-OlaHsd genetic background to DL-AAA: \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) KO MGCs, and \( Cbh^{1KO} \) KO mice not expressing Cre recombinase, and \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) mice that express Cre recombinase specifically in MGCS to ablate one allele of \( Cbh^{2} \). Our data suggest that \( Cbh^{1KO}Crb^{2\text{Flox/Flox}} \) and \( Cbh^{2\text{Flox/Flox}} \) retinas are less sensitive to DL-AAA than are \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) retinas, suggesting that raising the levels of CRB2 by rAAV gene therapy targeting \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) MGCS might prevent the adverse effects of the glial toxin DL-AAA.

In summary, we demonstrate that (1) low levels of CRB2 in \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) MGCS lacking CRB1, with normal levels of CRB2 expressed at the OLM in photoreceptors (see the cartoon in Figure S1 on CRB protein expression in the mouse model), result in a RP retinal phenotype with foci of retinal disorganization mostly in the inferior retina. Interestingly, our previous studies showed that complete loss of CRB2 in MGCS lacking CRB1 resulted in an LCA retinal phenotype throughout the entire retina. Furthermore, we show that (2) reduction of CRB2 protein levels worsens the retinal phenotype in the inferior quadrants as found in \( Cbh^{1KO} \) mice, (3) \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) RP mice are more susceptible to stress on MGCS than are \( Cbh^{1KO}Crb^{2\text{Flox/Flox}} \) and \( Cbh^{2\text{Flox/Flox}} \) mice, and (4) rAAV-hCRB2 therapy protects \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) retinas against loss of vision due to exposure to DL-AAA.

RESULTS

Reduction of CRB2 and loss of CRB1 in MGCS leads to ERG and OKT deficits

We expressed Cre under control of the Pdgfrα promoter in MGCS to ablate CRB2 expression of one floxed allele (\( Cbh^{1KO}Crb^{2\text{LowMGC}} \) = \( Cbh^{1KO}Crb^{2\text{Flox/Flox}} \) PdgfrαCre\(^{TG}\); Figure S1). High levels of Cre recombinase expression in photoreceptors and other neuronal cells can cause toxicity, impairing neuronal function. We assessed whether Cre expression in \( Cbh^{1KO} \) MGCS \( (Cbh^{1KO}Crb^{2\text{Flox/Flox}}) \) has an impact on retinal morphology, retinal transmission (ERG responses), or vision-guided optokinetic head-tracking thresholds (optokineti

Next, we measured flash ERGs and OKT responses in 1-, 3-, 6-, 9-, and 12-month-old \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) mice and age-matched littermate controls (\( Cbh^{1KO}Crb^{2\text{Flox/Flox}} \); Figures 1A–1J; Figure S3). One-month-old \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) mice showed normal ERG responses (scotopic, photopic, flicker) and OKT thresholds (visual acuity [VA] thresholds; contrast sensitivity thresholds) compared to age-matched littermate controls (Figures S3A, S3B, and S3H). Three-month-old dark-adapted \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) mice showed a reduced a-wave and b-wave response (Figures 1A–1D) and a reduced ERG flicker response (0.5 Hz; Figure S3M) compared to \( Cbh^{1KO}Crb^{2\text{Flox/Flox}} \) mice, suggesting a reduced rod photoreceptor retinal function. The ERG dark-adapted a-wave and b-wave response worsened over time in \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) and \( Cbh^{1KO}Crb^{2\text{Flox/Flox}} \) mice (6–9-, 12-month-old mice; Figures 1A and 1E–1G; Figure S3). The a-wave and b-wave amplitudes were proportionally lower (Figures 1H and 1I), but the b-wave/a-wave ratio of the scotopic ERG was not affected (Figure 1J), indicating that the overall retinal transmission (b-wave) and the photoreceptor response (a-wave) were impeded.

Vision-guided OKT responses (Figure 1K) were assessed on visual acuity (spatial frequency) and contrast sensitivity measurements. The spatial frequency threshold (visual acuity) was lower in 12-month-old \( Cbh^{1KO}Crb^{2\text{LowMGC}} \) littermates (Figure 1L). The contrast sensitivity threshold was markedly lower already at 6 and 12 months at a wide range of spatial frequencies measured (Figure 1M). The effects on contrast sensitivity threshold differences were detected with a higher statistical significance level at the spatial frequencies 0.064 and 0.092 cycles/degree (c/d; Figure 1M),
Figure 1. Decreased retinal function and vision-guided behavior in Crb1KO/Crb2LowMGC compared to Crb1KO/Crb2Flox/WT age-matched littermates. Crb1KO/Crb2LowMGC measurements are indicated in red (experimental group), and Crb1KO/Crb2Flox/WT age-matched littermates with a similar genetic background are indicated in black (control group). (A) Electroretinographic analysis of retinal function: scotopic single-flash intensity series (–4, –3, –2, –1, 0, 1, 1.5, and 1.9 log cd·s/m² light). (legend continued on next page)
suggesting that these spatial frequencies are most informative for hCRB gene therapy studies.

Reduction of CRB2 in MGCs results in a more severe CRB1 phenotype in the inferior part of the retina

We analyzed the morphological phenotype on mouse eyes on plastic sections. We included a WT-like mouse with a similar genetic background (Crb2^{Flox/WT}) because ectopic cell counts and photoreceptor inner/outer segment (IS/OS) length quantification compared to Crb1^{KO}Crb2^{Flox/WT} mice on plastic sections had not been done previously on mice with a 99.9% C57BL/6JOLAhsD genetic background. At 3 months of age, in Crb1^{KO}Crb2^{LowMGC} retinas compared to littermate control Crb1^{KO}Crb2^{Flox/WT} retinas, disorganization of the retinal layering at foci was detected in the two inferior quadrants of the retina (Figures 2A–2E; symbols: arrows, protrusions; open triangles, loss of photoreceptor IS/OS; asterisks, neovascularization). These disorganizations at foci included protrusions of photoreceptor nuclei into the photoreceptor segment layers, ingestion of photoreceptor nuclei into the OPL, disruptions at the OLM, and intermingling PRCs with inner retinal cells. Interestingly, outside the foci of retinal disorganization the layering of the retina remained intact. At 12 months of age the severity and number of retinal disorganizations at foci increased. Retinal disorganization could also be observed in the littermate control Crb1^{KO}Crb2^{Flox/WT} retinas, but outside of the affected foci the retinal layering remained intact (Figures 2F–2J). In the superior retina of the Crb1^{KO}Crb2^{LowMGC} mice, but not of the Crb1^{KO}Crb2^{Flox/WT} mice, at 3 months of age, sporadic protrusions of photoreceptor nuclei at foci could also be detected (Figure 2D), and such protrusions were observed at 12 months of age in Crb1^{KO}Crb2^{LowMGC} retinas as well as in littermate control Crb1^{KO}Crb2^{Flox/WT} retinas (Figures 2I and 2G). At 12 months of age, we observed that four out of five Crb1^{KO}Crb2^{LowMGC} retinas, and one out of five in the littermate Crb1^{KO}Crb2^{Flox/WT} control retinas, developed focal neovascularization in the inferior quadrants (Figure 2J, asterisks).

We generated retinal spidergiders for retinal thickness, outer nuclear layer (ONL) thickness, inner nuclear layer (INL) thickness, number of rows of photoreceptor nuclei in the ONL, photoreceptor IS/OS length, ectopic cells in the subretinal space, and ectopic cells in the OPL. In 12-month-old mice, the retinal thickness, the number of rows of photoreceptor nuclei in the ONL, and the ONL thickness were decreased compared to Crb1^{KO}Crb2^{Flox/WT} mice (Figures 2L, 2M, and 2O). No major difference was found in the INL thickness in 12-month-old mice (Figure 1N). Crb1^{KO}Crb2^{LowMGC} mice displayed many displaced retinal cells (at 1, 3, and 6 months of age; Figures 2E and 2P–2R). Most of the ectopic nuclei were found in the OPL and some at the subretinal space (compare Figure 2S and Figure 2Q). The photoreceptor IS/OS length in the inferior quadrants of Crb1^{KO}Crb2^{LowMGC} retinas was shorter compared to Crb1^{KO}Crb2^{Flox/WT} retinas at 3 months, and both mouse lines had similar but shorter IS/OS in the inferior quadrants compared to the superior quadrants at 6 months of age (Figures 2T and 2U).

We also assessed the morphology on protein expression by immunohistochemistry in 3-month-old mice for gliosis, Müller glial microvilli, IS/OS of photoreceptors, synapses at the OPL and IPL, and OLM disruptions (Figure 3). More GFAP-positive stress fibers extending from the inner limiting membrane (ILM) to the OLM were observed in the inferior and superior quadrants of Crb1^{KO}Crb2^{LowMGC} compared to Crb1^{KO}Crb2^{Flox/WT} mouse littermate retinas (Figures 3A–3E). When further looking at Müller glial morphology, also shortened and collapsed microvilli (CD44+) were observed in Crb1^{KO}Crb2^{LowMGC} compared to Crb1^{KO}Crb2^{Flox/WT} retinas (Figures 3F–3J). This matches our previous transmission electron microscopy (TEM) observation that the CRB levels from photoreceptors at the OLM and here in MGCs are important for MGC microvilli extensions.11,12 Also, the outer segments of photoreceptors (labeled with cone arrestin for cones and rhodopsin for rods) were lost at foci in the inferior and partially in the superior quadrants of Crb1^{KO}Crb2^{LowMGC} compared to Crb1^{KO}Crb2^{Flox/WT} retinas (Figures 3K–3O). Similarly, the internalization of rhodopsin around the cell nucleus was found in photoreceptor ingresses (Figures 3L, 3M, and 3O, asterisks). The IPL synapses (sublamina-a OFF-bipolar cell synapses; sublamina-b ON-bipolar cell synapses) were relatively unaffected as shown by the synaptic marker plasma membrane calcium ATPase 1 (PMCA1), but the OPL synapses (protein kinase Cz [PKCz]/PMCA1 markers) were disrupted in the superior and inferior quadrants of the retina of Crb1^{KO}Crb2^{LowMGC} compared to Crb1^{KO}Crb2^{Flox/WT} retinas (Figures 3P–3T, asterisks indicate OPL disruption).27 We also found many more OLM disruptions and a lower CRB2 expression at the OLM in Crb1^{KO}Crb2^{LowMGC} compared to Crb1^{KO}Crb2^{Flox/WT} and WT-like (Crb2^{Flox/Fl}) retinas (Figures 3U–3Y, asterisks indicate OLM disruptions).

We previously hypothesized that the loss of mCRB2 protein expression at the OLM determines the retinal phenotype in Crb1^{KO} mice,
but it was not clear how much mCRB2 is contributed by MGCs and PRCs. In this study, we semi-quantified mCRB2 protein expression (on fluorescence intensity) at the OLM and the number of OLM breaks in 3-month-old Crb2ΔRods (WT-like), Crb2ΔRods (ablation of Crb2 in rods), Crb1KO/Crb2LowMGC, Crb1KO/Crb2LowMGC, and Crb1KO/Crb2LowMGC mice. We validated our previous results, indicating that the OLM mCRB2 protein expression in the inferior retina compared to the superior retina was similar to the previous mixed genetic background of Crb1KO mice. The Crb2 protein expression was decreased at the OLM by 49% (21 ± 4 AU; n = 3 mice), 34% ± 10% in Crb1KO/Crb2LowMGC (n = 5 mice), and 62% ± 12% in Crb1KO/Crb2LowMGC (n = 3 mice) compared to Crb1KO/Crb2LowMGC retinas (n = 7), suggesting that MGCs contribute about half of the total CRB2 protein levels to the OLM (Figure 3Z). CRB2 protein expression between the peripheral (27 ± 6 AU [arbitrary fluorescence unit] and 15 ± 4 AU; n = 5 mice) and central retina (21 ± 1 AU and 14 ± 5 AU) was not statistically different in Crb1KO/Crb2LowMGC mice (p = 0.08) or Crb1KO/Crb2LowMGC mice (p = 0.30; data not shown).

The adherens junctions and the subapical region are located at the OLM. We and others have previously investigated the recruitment of adherens junction markers (e.g., catenins or catenins) by the Crumbs complex at the subapical region. The Crumbs complex consists of the CRB protein family (Crb1 and Crb2), the PALS1 (MPP5-PATJ-MUPT1 protein complex, and the PAR6-PAR3-aPKC-CDC42 protein complex. Disruption of the Crumbs complex leads to loss of polarity and loss of adhesion in many Crb mouse models. Semi-quantification of p120-catenin, an adherens junction marker, showed a 28% ± 3% (SEM) decrease in Crb2ΔRods compared to WT and a 45% ± 7% reduction in Crb1KO/Crb2ΔRods compared to Crb1KO/Crb2ΔRods mice (Figure 3AA). No statistical difference in p120-catenin expression was found between Crb1KO/Crb2ΔRods and Crb1KO/Crb2LowMGC mice (Figure 3AA).

Furthermore, the level of CRB1 and CRB2 proteins at the OLM determined the number of OLM breaks. More OLM breaks were found in mice with less CRB protein expression at the OLM (Figure 3BB). We further validated our previous results that the loss of CRB1 or CRB2 reduces the p120-catenin protein localization at the OLM, subsequently destabilizing the OLM and reducing adhesion between MGCs and photoreceptors, causing OLM breaks, and finally facilitating photoreceptor loss in the form of protrusion of photoreceptor nuclei through the OLM into the layer of the inner segments.

**Figure 2. Removal of CRB1 and low levels of CRB2 in MGCs lead to abnormal layering in the inferior quadrants** (A–K) Toluidine-stained light microscopy of retinal sections from control (Crb2ΔRods/Flox, Crb1KO/Crb2ΔRods/Flox, and Crb1KO/Crb2LowMGC) mice at 3 and 12 months of age. Representative morphological changes: protrusions (red arrows downward), ingrowths, neovascularization (asterisks), and loss of inner/outer segments of photoreceptors (red triangles). (K–Q) Representative 12-month-old retina of a Crb1KO/Crb2LowMGC mouse indicates photoreceptor layer presence in periphery and superior quadrants of the retina. (L–U) Spidergrams of the retina of wild-type (WT; Crb2ΔRods/Flox, Crb1KO/Crb2LowMGC, and Crb1KO/Crb2LowMGC) mice. (L and M) Decrease of retinal thickness (outer limiting membrane (OLM) to the inner limiting membrane) and outer nuclear layer (ONL) thickness at 1 mm distance on the inferior retina in 1-, 3-, 6-, 9-, and 12-month-old mice. (N and O) Inner nuclear layer (INL) thickness and the number of rows of photoreceptor nuclei in the ONL of 12-month-old mice. (P–S) Total number of ectopic cells at 1, 3, and 6 months of age. (S) Most ectopic cells are in the outer plexiform layer (OPL) at 3 months of age. (T and U) Inner/outer segment length of photoreceptors in the periphery (2.0 mm) and central (1.0 mm) from the optic nerve head (ONH). Scale bars, 50 μm. Data are presented as mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001.

**Exposure to DL-AAA causes lasting retinal damage and worsened sight** A low dose of 100μg of DL-AAA injected intravitreally in WT mice was shown by others to disrupt the OLM, causing photoreceptor nuclei protrusions into the photoreceptor segment layers that were resolved during 48 h. In this study, we examined whether retinas with reduced levels of CRB2 proteins in Crb1KO/Crb2LowMGC MGCs are more sensitive to DL-AAA than are retinas expressing normal levels of CRB2 proteins in Crb1KO/Crb2LowMGC MGCs. We challenged 2-month-old WT-like (Crb2ΔRods/Flox) and the RP (Crb1KO/Crb2ΔRods/Flox and Crb1KO/Crb2LowMGC mice with different doses of DL-AAA (100, 150, or 200 μg) and analyzed the retinal morphology, the retinal transmission (ERG), and vision-guided behavior (OKT) 1 month after intravitreal injection (3-month-old mice; Figure 4A). No effect was seen on OKT and ERG when we injected PBS as a control (Figures 4B–I). An overall dosage effect of DL-AAA on retinal transmission (ERG) and vision-guided behavior (OKT) was found for all mouse lines (Figures 4B–I). 100 μg of DL-AAA markedly decreased the retinal transmission for the scotopic a-wave and b-wave (Figures 4C–E) and the OKT thresholds (Figures 4F–I) in the Crb1KO/Crb2LowMGC mice but not in the Crb2ΔRods/Flox and Crb1KO/Crb2LowMGC mice.

No effect was seen on morphology when we injected PBS as a control (Figures 4K–N). Very little effect on morphology was found at 100 μg of DL-AAA except for the Crb1KO/Crb2LowMGC retinas (Figures 4O–S). The Crb1KO/Crb2LowMGC retinas had an overall worsened retinal phenotype on the superior as well as the inferior retina as indicated by the ingestions (asterisks) and protrusions (arrow) in the retina (Figures 4R and 4S). 150 and 200 μg of DL-AAA caused irreversible retinal damage on morphology in all mouse lines (Figures 4T–4Z, 4AA, 4BB, and 4CC). The decrease in ERG/OKT and worse retinal morphology indicates that Crb1KO/Crb2LowMGC retina might be more sensitive to OLM disruptions induced by DL-AAA. In the next section we investigated whether the increased sensitivity could be alleviated by hCRB1 or hCRB2 gene supplementation therapy.
Figure 3. \(Crb1^{KO}Crb2^{LowOGC}\) retinas compared to \(Crb1^{KO}Crb2^{Flox/WT}\) retinas show more disruptions at the OLM. Immunohistochemistry of 3-month-old mice. A representative image is shown of three to six retinas per group analyzed. (A–Y) Sections were stained for: (A–E) glutamine synthetase (green) for MGCs and glial fibrillary acidic protein (GFAP; red) for MGC stress fibers; (F–J) CD44 (red) for Müller glial microvilli processes; (K–O) cone arrestin (CAR; red) for cone photoreceptor segments and rhodopsin for rod photoreceptor outer segments (green); (P–T) microtubule-associated protein 2 (MAP2) (green) for ganglion cells.

(legend continued on next page)
rAAV-hCRB protects retinal morphology, OKT, and ERG response in the DL-AAA-challenged Crb1KO-Crb2lowMGC RP mouse model

We previously determined the tropism and potency of the ShH107445F capsid (and the cell-specific expression of the cytomegalovirus [CMV] and minimal CMV [CMVmin] promoter) and the hCRB1 codon-optimized or the hCRB2 codon-optimized cDNA delivery to MGCs by intravitreal delivery of rAAVs in WT mice. We previously showed that the ShH107445F capsid can effectively infect and efficiently express GFP in more than 40% of all mouse MGCs by intravitreal injection.36,37 We also previously showed that subretinally injected rAAV serotype 2 (rAAV2)/9.CMV.min.hCRB1ico.spA or intravitreal rAAV2/ShH107445F.CMV.min.hCRB1ico.spA can express hCRB1 protein at the OLM in Crb1KO mice. Finally, we previously demonstrated that we can express hCRB2 protein at the OLM by intravitreal delivery of rAAV2/ShH107445F.CMV.hCRB2ico.spA in Crb2 conditional KO (cKO) retinas.13

rAAV2/ShH107445F.CMV.min.hCRB1ico.spA (rAAV-hCRB1) or rAAV2/ShH107445F.CMV.hCRB2ico.spA (rAAV-hCRB2) was injected intravitreally into one eye of Crb1KO-Crb2lowMGC mice at P21. Then, 2-month-old retinas were challenged by intravitreal injection of 100 μg of DL-AAA in both eyes. ERG, OKT, spectral domain optical coherence tomography (SD-OCT), and retinal morphology were assessed in 3-month-old mice (Figure 5A).

The overall retinal morphology improved with both rAAV vectors (Figures 5B–5G). Retinas receiving the rAAV-hCRB treatment had more rows of photoreceptor nuclei in the ONL (Figure 5H). The expression of hCRB in mouse MGCs protected against the protrusion of photoreceptor nuclei into the subretinal space and OPL (Figure 5I). In the central-inferior retina of Crb1KO-Crb2lowMGC mice, the photoreceptor IS/OS length was 33 ± 6 μm (see also Figure 2P), 21 ± 6 μm in 100-μg DL-AAA-treated mice, 41.7 ± 5 μm in 100-μg DL-AAA+rAAV-hCRB1-treated mice, and 40 ± 22 μm in 100-μg AAA+rAAV-hCRB2-treated mice (Figure 5J), indicating a significant protection against loss of photoreceptor segment lengths in the central area for rAAV-hCRB-injected eyes. We then investigated the retinal morphology by SD-OCT (Figures 5K–5M). We found retinal degeneration in the central inferior quadrants at around 0.5–0.9 mm from the optic nerve head (ONH) on the volume intensity projection (VIP) in the ONL comparable to changes found on plastic sections (Figures 5H and 5J). More extensive disruptions in the OPL/ONL/OLM of control mice (100 μg of DL-AAA only) were found in the inferior and superior retinal quadrants (red arrows in Figures 5K–5M). Interestingly, some mice injected with rAAV-hCRB1 showed many vitreous-infiltrating cells (asterisks in Figure 5L).

We measured ERG and OKT visual acuity and contrast sensitivity in these mice. We analyzed the differences between the rAAV-hCRB-treated eye against the eye not receiving the rAAV treatment (control eye) because the variation between mice was considerable, and comparison on an individual mouse reduces variation and permits a pairwise comparison, reducing the numbers of mice needed to show effects (see Figures 6 and 54 for absolute values). The retinal function measured by scotopic and photopic ERG was significantly higher in rAAV-hCRB2-treated eyes, but no significant changes were found in rAAV-hCRB1-treated eyes compared to control eyes (Figures 6A–6C; see also Figure 54). The visual acuity improved for rAAV-hCRB2-treated eyes but not for rAAV-hCRB1-treated eyes (Figure 6D). The contrast sensitivity threshold (spatial frequency: 0.032, 0.064, and 0.092) was significantly increased upon delivering hCRB2 cDNA but not for hCRB1 cDNA to MGCs (Figures 6E–6G; Figures S4N and S4O). The tracking at 0.092 cycles/degree was significantly worse for rAAV-hCRB1-injected eyes compared to control eyes (Figure 6G; Figures S4N and S4O).

rAAV-hCRB therapy reduces gliosis and protects Müller cell microvilli and photoreceptor IS/OS

MGCs extend from the endfeet at the ILM up to the apical vili above the OLM and support/maintain retinal homeostasis and retinal integrity. Long Müller glial stress fibers (GFAP-positive) extended through the retina in the inferior-central retinal areas of Crb1KO-Crb2lowMGC mice injected with 100 μg of DL-AAA (Figure 7A, arrows indicate stress fibers). However, eyes treated by rAAV-hCRB showed shortened and less GFAP-positive stress fibers extending into the ONL (Figures 7B and 7C, arrows indicate stress fibers). Similarly, shortened and thickened Müller microvilli were observed in the retina of eyes injected with DL-AAA without rAAV-hCRB treatment. However, retinas with rAAV-hCRB treatment displayed less severely affected Müller microvilli in the inferior retinal quadrants (Figures 7E and 7F).

Crb1KO-Crb2lowMGC retinas exposed to DL-AAA-mediated loss of adhesion had long stretches of no inner/outter photoreceptor segments in the inferior retinal quadrants stained by rhodopsin and cone arrestin (Figure 7G, arrows). Also, more ectopic nuclear rhodopsin expression in photoreceptor ingresses was observed (Figure 7G, asterisks indicate ingresses). The most affected inferior-central retinal quadrants frequently had some outer photoreceptor segments left in rAAV-hCRB-treated eyes compared to the control eyes (Figures 7G and 7I).

Retinas of Crb1KO-Crb2LowMGC mice injected with DL-AAA had more ectopic synapses and loss of horseshoe-shaped synapses (Figure 7G).

and synapses in the inner plexiform layer (IPL),26 PMAco1 (red) for pre-synapses of photoreceptors at the OPL and lamina a/b in the IPL,27 430 Molecular Therapy: Methods & Clinical Development Vol. 20 March 2021 and PKCz (light blue) for bipolar cells and bipolar post-synapses at the OPL; and (U–Y) subapical region marker CRB2 (red) and the adherens junction marker p120-catenin (green), (Z and BB) The decrease in CRB proteins and p120-catenin proteins at the OLM increases breaks at the OLM in 3-month-old WT, Crb2Flox/Wt, Crb1KO-Crb2Flox/Wt, Crb1KO-Crb2LowMGC and Crb1KO-Crb2A2lowMice. (AA) Normalized CRB2 protein expression to Crb1Flox mouse (fluorescence). (AB) Normalized p120-catenin protein expression to Crb1Flox mouse (fluorescence). (BB) OLM breaks per 100 μm retinal length. GCL, ganglion cell layer. Scale bars, 50 μm; inserts, 50 μm. Data are presented as mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001.
Retinas treated with rAAV-hCRB showed smaller disruptions of horseshoe-shaped synapses at the OPL (Figures 5G–5L; see also Figures 5K–5M, arrows). Also, DL-AAA-induced retinal stress may induce synaptic changes in the IPL but no differences between the previously described conditions without DL-AAA (Figures 3P–3T), with 100 μg of DL-AAA, or DL-AAA with rAAV-hCRB treatment were observed (Figures 7)–7L, asterisks indicate photoreceptor [synaptic] ingressions.

### Neovascularization, activated microglial cells, ectopic hCRB1 expression, and ciliary body changes related to rAAV-hCRB1 intravitreal injections

rAAV-hCRB1 treatment produced hCRB1 protein at the OLM (Figures S5A–S5E), but it was also found in the ciliary body (Figure S5D). Similarly, CRB2 was found at the OLM in rAAV-hCRB2 of non-treated retinas (Figure S5G). The benefit of the rAAV-hCRB2 over rAAV-hCRB1 therapy on ERG retinal transmission and OKT behavior outcome measures (see Figure 6 and Figure S4) compared to similar benefits on retinal morphology (Figure 5) prompted us to further investigate whether one of these vectors increases neovascularization events or microglial activation.

First, we characterized the background of neovascularization and microglial activation in the inferior-central quadrants of 3-month-old Crb2Flox/Flox, Crb1KO/Crb2Flox/WT, and Crb1KO/Crb2LowMGC mice injected with DL-AAA (100 μg) against the noninjected control eyes (Figures S5H–S5M). A strong increase in neovascularization events was detected in the degenerate inferior retinal quadrants of Crb1KO/Crb2LowMGC mice injected with DL-AAA (100 μg) upon immunohistochemical staining for plasmalemma vesicle-associated protein (PLVAP; a marker for early vascular leakage) or activated microglial cells (CD11b-positive microglial cell dendrites and migration to the ONL and GCL) (Figure S5M). Neovascularization was not observed in the RPE, ONL, or INL when rAAV-hCRB2 treatment group. However, we observed in the treatment groups of rAAV-hCRB2 compared to rAAV-hCRB1 fewer GFAP-positive stress fibers (Figures 6B and 6C) and fewer neovascularization events in the GCL/NFL (Figures S5H–S5O). Additionally, the rAAV-hCRB1 treatment group showed consistent neovascularization at the ciliary body that was not found in control eyes (100 μg of DL-AAA injected) or in rAAV-hCRB2-treated eyes. Neovascularization at the ciliary body was previously observed after 4.5 months after intravitreal injection of rAAV-hCRB1 but not by 150 μg of DL-AAA injected in the rAAV-hCRB1 treatment group. However, we did not detect major protein contaminants in two independently produced rAAV batches used in the study (Figure S5T).

In summary, we observed more neovascularization events with immunohistochemistry in the ONL/ONL/NFL when 100 μg of DL-AAA was injected intravitreally in Crb1KO/Crb2LowMGC mice compared to noninjected Crb1KO/Crb2LowMGC mice (Figures S5L and S5M). Fewer neovascularization events in the ONL/ONL/NFL were observed with immunohistochemistry and SD-OCT in the rAAV-hCRB2 treatment group or rAAV-hCRB1 treatment group compared to the control eyes. However, we observed in the treatment groups of rAAV-hCRB2 compared to rAAV-hCRB1 fewer GFAP-positive stress fibers (Figures 6B and 6C) and fewer neovascularization events in the GCL/NFL (Figures S5H–S5O). Additionally, the rAAV-hCRB1 treatment group showed consistent neovascularization at the ciliary body that was not found in control eyes (100 μg of DL-AAA injected) or in rAAV-hCRB2-treated eyes. Neovascularization at the ciliary body was previously observed after 4.5 months after intravitreal injection of rAAV-hCRB1 but not by 150 μg of DL-AAA injected in the rAAV-hCRB1 treatment group. However, we did not detect major protein contaminants in two independently produced rAAV batches used in the study (Figure S5T).

### DISCUSSION

In this study, we show that (1) in Crb1KO/Crb2LowMGC mice, a minimum of half of the endogenous mCRB2 levels in MGCs lacking...
Figure 5. rAAV-hCRB2 protects against retinal disorganization and degeneration in the AAA-challenged Crb1<sup>−/−</sup>Crb2<sup>LowMGC</sup> mice

(A) Retinitis pigmentosa model: rAAV-hCRB was injected in one eye versus noninjected eye (control) at P21/P22. Both eyes were then injected with a MGC-specific OLM stressor (100 μg of DL-AAA) at the 2-month time point. The potential rAAV therapy protective effect was measured at 3 months. (B–G) Toluidine-stained light microscopy of retinal sections from 3-month-old Crb1<sup>−/−</sup>Crb2<sup>LowMGC</sup> mice that were treated with (B–D) rAAV2/ShH10 Y445F.CMVmin.hCRB1 or treated with (E–G) rAAV2/ ShH10Y445F.CMV.hCRB2. (H–J) Spidergrams of the retina of Crb1<sup>−/−</sup>Crb2<sup>LowMGC</sup> mice (100 μg of AAA injected, n = 9; rAAV-hCRB1 injected, n = 4; and rAAV-hCRB2 injected, n = 8). (H–J) rAAV-hCRB treated retinas had more rows of photoreceptor nuclei in the ONL (H), fewer ectopic photoreceptor cell nuclei in the subretinal space and (legend continued on next page)
CRB1 (with normal levels of CRB2 expressed at the OLM in photoreceptors) results in a RP retinal phenotype. Interestingly, in previous studies we showed that complete loss of CRB2 in MGCs lacking CRB1 resulted in an LCA retinal phenotype.\(^1\) (2) Mice with reduced levels of CRB2 protein in MGCs lacking CRB1 in MGCs (Crb1\(^{KO}\) Crb2\(^{LowMGC}\)) showed increased sensitivity to OLM disruptions upon exposure to DL-AAA. (3) rAAV-hCRB2 therapy to MGCs protects against OLM disruptions, a decrease of ERG responses, and a loss of OKT contrast sensitivity in Crb1\(^{KO}\) Crb2\(^{LowMGC}\) mice.

**CRB1**-related RP patients have a nonfunctional or less functional CRB1 protein in MGCs and photoreceptors.\(^4\) We have shown that we can model the **CRB** phenotype in human retinal organoids in vitro\(^6\) and in mice in vivo. In the present study, we further explored the effect of ablating **CRB** in late-born retinal cells, such as MGCs. Previously, we showed that the full ablation of Crb1 and Crb2 in MGCs (Crb1\(^{KO}\) Crb2\(^{AbMGc}\)) caused a severe LCA-like retinal phenotype with no ERG response already in 1-month-old mice.\(^2\) Previously, we showed that full ablation of Crb2 specifically in MGCs (Crb2\(^{AbMGC}\)) causes a slow progressing RP-like phenotype with sporadic disruptions at the OLM with protrusion of photoreceptor nuclei into the photoreceptor segment layer, similar to previous observations in retinas lacking CRB1, without effects on the ERG response.\(^3\) In this study, we show that Crb1\(^{KO}\) Crb2\(^{LowMGC}\) mice develop a more severe RP-like phenotype than do Crb1\(^{KO}\) or Crb2\(^{AbMGC}\) mice with clear effects on ERG and OKT responses and retinal morphology, suggesting similar functions of CRB1 and CRB2 proteins in MGCs.

Mice are housed under standard low light conditions that do not resemble the retinal stress that RP patients undergo in regular life. We have shown previously that light exposure can worsen the **CRB1** RP mouse phenotype.\(^8\)\(^,\)\(^12\) Bright light exposure causes prolonged inflammation, neovascularization, and retinal damage in RP mouse models. Blue light exposure induces retinal degeneration, oxidative stress, and neuroinflammatory activity similar to dry age-related macular degeneration (AMD). In this study, we explored intravitreal DL-AAA injections that act on glial cells such as MGCs, causing disruptions at foci at the OLM. Since exposure of DL-AAA to the mouse retina, or loss of CRB1 or loss of CRB2 proteins in the mouse retina, causes disruptions at foci at the OLM, we examined whether mice with decreased levels of CRB proteins in MGCs are more sensitive to DL-AAA exposure than retinas with higher levels of CRB proteins in MGCs. The OLM disruptions caused by low doses of DL-AAA are reversible in WT mice.\(^7\) The phenotype described indicated some similarities such as photoreceptor nuclei protrusions into the photoreceptor segment layer that we previously observed in Crb1 RP mouse models. However, it was not known how DL-AAA could affect the visual tracking thresholds (OKT), the retinal function (ERG), quantitative retinal morphology, or mouse models with reduced cell adhesion. West et al.\(^{17}\) showed that low levels of DL-AAA (100 \(\mu\)g) did not affect the gross retinal morphology in WT mouse retinas. In this study, we first demonstrated that mice expressing normal levels of endogenous CRB2 proteins in MGCs (Crb2\(^{LowMGC}\) or Crb1\(^{KO}\) Crb2\(^{LowMGC}\)) are less sensitive to exposure of 100 \(\mu\)g of DL-AAA than are mice with reduced levels of endogenous CRB proteins in MGCs (Crb1\(^{KO}\) Crb2\(^{LowMGC}\)). Subsequently, we demonstrated that increasing the levels of recombinant hCRB2 in MGCs in Crb1\(^{KO}\) Crb2\(^{LowMGC}\) retinas prior to exposure to DL-AAA decreased the sensitivity to DL-AAA exposure. We applied the rAAV-hCRB gene therapy vectors at P21, and it could have been more effective to apply rAAV-hCRB at earlier time points. In this study, we show that rAAV-hCRB2 applied at P21 to Crb1\(^{KO}\) Crb2\(^{LowMGC}\) MGCs at early stage retinal disease effectively prevents against the adverse effects of exposure of DL-AAA. The Crb1\(^{KO}\) Crb2\(^{LowMGC}\) mice pre-treated intravitreally with rAAV-hCRB2 and subsequently exposed to DL-AAA showed significantly fewer disruptions at the OLM, fewer protrusions of photoreceptor nuclei in the photoreceptor segment layers, less loss of photoreceptors, an improved contrast sensitivity as measured by OKT, and an improved retinal function as measured by ERG.

Intravitreal injection of rAAV vectors is more efficient in supplementing cDNA to MGCs compared to subretinal injections. For example, we have shown that rAAV2/ShH10\(^{14455}\) can effectively infect and efficiently express GFP in more than 40% of all mouse MGCs.\(^{36,37}\) Yet, intravitreal injections increase the risk of alternate rAAV vector biodistribution and ectopic vector expression.\(^{13}\) Others have indicated a transient inflammation of the aqueous and the vitreous body by empty rAAV capsids and rAAV supplementation vectors at high doses.\(^{42}\) In this study, we re-assessed the effect of rAAV-hCRB1 and rAAV-hCRB2 to the vitreous body, ciliary body, and the neuroretina. We previously found that rAAV-hCRB1 applied at P14 causes ectopic CRB1 expression within the epithelium of the ciliary body and the iris epithelia, affecting the corneal thickness, the eyeball perimeter, and CD11b- and CD3-positive infiltrating cells to the ciliary body.\(^{13}\) We found similar ectopic rAAV-hCRB1 expression at 3 months of age in the Crb1\(^{KO}\) Crb2\(^{LowMGC}\) mouse model injected at P21 with rAAV-hCRB1 and at 2 months of age with DL-AAA injected. Interestingly, cDNA supplementation of hCRB1 to MGCs (by rAAV capsids) improved the retinal morphology, such as a decrease in ectopic cells in the OPL/subretinal space and the maintenance of the number of rows of photoreceptor nuclei in ONL in the central retina.

We hypothesize that hCRB1 applied to Müller cells can protect against loss of OLM integrity, but OKT visual behavioral and ERG
electrical transmission studies suggest that the neural network is not sufficiently restored, whereas morphological studies of the treated eyes suggest that there are adverse effects upon ectopic expression of CRB1. Potentially, the neovascularization events seen in the GCL/NFL of the inferior mouse retinas and the poor ERG/OKT responses measured in the rAAV-hCRB1-treated eyes, but not in the rAAV-hCRB2-treated eyes, may be linked to differences of hCRB2 over hCRB1 (over-)expression in (1) MGCs, (2) astrocytes, (3) protein-protein interactions, or (4) immunogenic properties of hCRB1 protein in a Crb1 KO mouse. We discuss these points below.

(1) MGCs contribute to the maintenance and rigidity of the retinal layers (i.e., tensile strength), regulate blood flow and maintain the retina-blood barrier (e.g., release vascular endothelial growth factor [VEGF]), ensheathe synapses in IPL and the OPL, guide light rays to the segment layer, and remove waste products.43–45 Mouse Crb1 (mCrb1) and Crb2 (mCrb2) expression levels in MGCs are linked to the intermediate capillary plexus development by suppressing the angiogenic growth factor VEGFA46 and promoting MMP-3 expression (extracellular matrix remodeling).47 hCRB1 expression protected MGCs less than did hCRB2 from gliosis. (2) Astrocytes wrap around retinal endothelial cells and pericytes of the superficial capillary plexus at the GCL/NFL. The astrocytic foot processes may release angiogenic growth factors (e.g., VEGF) during retinal development and disease, altering the expression of tight junction proteins in retinal endothelial cells (blood-retinal barrier, ZO-1/occludin/VE-cadherin proteins).48–50 VEGF suppression speeds up programmed capillary regression during development.51 The AAV6 capsid variant ShH10Y445F used in our study efficiently infects astrocytes.52 We found only neovascularization events in the GCL/NFL with rAAV-hCRB1 that may be linked to differences of hCRB1 over hCRB2 expression in DL-AAA-stressed astrocytes. (3) The protective function of hCRB2 over hCRB1 protein may also be explained by unknown differences in the intracellular protein function (both have short intracellular FERM/PDZ/ERLI domains9,14) or differences in hCRB homodimerization/heteromerization of extracellular hCRB/mCRB protein at the OLM. However, both vectors...
rescued the retinal morphology at the OLM where we would expect that the homodimerization/heteromerization takes place. (4) The Crb1KO/Crb2LowMGC mice are naive for full-length mCRB1 protein but not for mCRB2 (expressed in, e.g., RPE cells, the retina, choroid, lung tissue, and the brain5,53–55), and the expression of hCRB1 on a Crb1KO background may induce an immune reaction, whereas due to endogenous mCRB2 expression the retinal expression of hCRB2 is less likely to induce an immune reaction, especially in a CRB1-RP model with a leaky blood-retina barrier (partially aggravated by DL-AAA), allowing more infiltration of immune cells. More research is needed to delineate the effects of hCRB1 overexpression in MGCs or astrocytes on retinal vasculature in the healthy, degenerate, and Crb1KO retina.

Nevertheless, the additive effect of DL-AAA on Crb1KO/Crb2LowMGC MGCs may model the exudative vasculopathy seen in RP-CRB1 patients’ and help to better understand what therapies on MGC-regulated vasculopathy may be beneficial for RP-CRB1 patients. Coats-like exudative vasculopathy (also called idiopathic retinal telangiectasia) is strongly associated with the RP-CRB1 phenotype seen in clinics.1 We observed (early) vascular leakage (PLVAP) in the inner retina in this study, but also previous studies found neovascularization (VEGF, von Willebrand factor [vWF], Fluorescein angiography-cSLO) in RP-CRB1 mouse and rat models.5,8,12,15,28,47,56 Subretinal injection of a high dose of DL-AAA induces MGC injuries that can develop into vascular telangiectasia and hemorrhages of the inner retinal vasculature in rats, rabbits, and nonhuman primates20,57,58. Thus, a (pan-) CRB1 therapy may need to address not only the cell-cell adhesion at the OLM but also regulation of glial cells on retinal vasculature signaling and overall retinal maintenance.

We show that rAAV-hCRB2 does not cause adverse changes at the ciliary body or the GCL/NFL. Furthermore, rAAV-hCRB2 protected against loss of retinal and visual function and retinal morphology. Our study suggests that expression of hCRB2 does not cause adverse effects in mouse retinas and that it can significantly increase retinal adhesion in a new cell-adhesion CRB1 RP mouse model. The study further strengthens the hypothesis that rAAV-hCRB2 retinal gene therapy might be of benefit for RP patients with mutations in the CRB1 gene.

MATERIALS AND METHODS

Mice

Procedures concerning animals were performed according to the Dutch Central Commission Animal Experimentation (CCD) license.
no. AVD1160020172924, the working protocols (OZP nos. PE.18.016.002, PE.18.016.006, PE.18.016.007, and PE.18.016.010) approved by the Local Ethics Committee (Instentie voor Dierenwelzijn [Ivd]) of the Leiden University Medical Center, and the Association for Research in Vision and Ophthalmology (ARVO) statement for the use of animals in ophthalmic and vision research. All mice were used or maintained on a 99.9% C57BL/6J/OlaHsd genetic background with a 12-h light/12-h dark cycle (standard low light housing condition ~10–20 lux) and supplied with food and water ad libitum. All experiments were carried out in male and female mice. All mouse strains below were confirmed to be Crb1Δα-negative, Nnt(exon 7–11)WT/WT, Mmnn1(exon 8)N−/−, α-synuclein(exon 6)−/−, and PdcdgWT/WT, thus similar to the genetic background of C57BL/6J/OlaHsd mice.

Crb1KO/Crb2Δfloxed/Flox mice were crossed with a Crb1KO/Pdgfrα-CreTyg/+ to produce Crb1Δ/−/Crb2Δfloxed/Flox/Pdgfrα-CreTyg/+ mice (Crb1KO/Crb2LowMGC) and Crb1Δ/−/Crb2Δfloxed/Flox mice. The Crb1KO/Crb2LowMGC mice lack mCRB1 in radial glial progenitor cells and MGCs and reduced mCRB2 protein expression in MGCs during early retinal development.2 The Crb1Δ/−/Crb2Δfloxed/Flox/Pdgfrα-CreTyg/+ mice contain a Pdgfrα-Cre transgene (C57BL/6-Tg(Pdgfrα-Cre)1Clef/J) driving the Cre gene specifically in MGCs.5,58 We determined the Cre mosaicism in a reporter mouse line (ROSA26R-TdTomato) and found that 95% of all MGCs excised membrane-targeted enhanced green fluorescent protein (mGFP) by Pdgfrα-Cre-mediated recombination.5 mCRB2 protein is expressed by MGCs and PRCs at the OLM. The Cre mice used express the Cre recombinase specifically in MGCs, generating hemizygote Crb2 MGCs. Assuming that 49% ± 3% (SEM) of mCRB2 is localized at the OLM within photoreceptors. (Figure 3Z), we estimate that 51% of mCRB2 is localized at the OLM in control Crb1KO/Crb2Δfloxed/Flox/WT. Since the total levels of mCRB2 at the OLM of Crb1KO/Crb2ΔLowMGC retinas are reduced by 34% ± 10% (SEM) compared to the reference littermate control Crb1KO/Crb2Δfloxed/Flox/WT (Crb1KO) retinas (Figure 3Z), we estimate that the levels at the OLM of mCRB2 in Crb1KO/Crb2Δfloxed/Flox MGCs dropped from 100% to 33% ([51% − 34%]/51% × 100 = 33%) specifically in Crb1KO/Crb2ΔLowMGC MGCs. The mCRB2ΔRods and Crb1KO/Crb2ΔΔRods were previously described.11 The mCRB2ΔRods contained a Rho-iCre transgene ablating the Crb2 gene in developing rod photoreceptors. Chromosomal DNA isolation and genotyping were performed as previously described.3 All mice were euthanized using CO2/O2. The experimental and control mice were collected at the same time at 5–8 h within the light cycle to have comparable lengths of the IS/OS of photoreceptors.

**ERG**

Dark and light-adapted ERGs were performed under dim red light using an Espion E2 (Diagnosys, Lowell, MA, USA). ERGs were performed on 1-, 3-, 6-, 9-, and 12-month-old Crb1KO/Crb2ΔLowMGC and Crb1KO mice. The ERG values of the right and left eye were averaged for the analysis in Figures 1 and S2. One eye was used for analysis for other experiments (treatment versus control eye). Mice were anesthetized using 100 mg/kg ketamine and 10 mg/kg xylazine intraperitoneally, and the pupils were dilated using atropine drops (5 mg/mL). ERGs were recorded as previously described.16 Scotopic recordings were obtained at −4, −3, −2, −1, 0, 1, 1.5, and 1.9 log cd · s/m^2 light intensity. Flicker recordings were obtained at 0.5 log cd·s/m^2 fixed light intensity at the frequencies 0.5, 1, 2, 3, 5, 7, 10, 12, 15, 18, 20, and 30 Hz. Photopic recordings were obtained at 30 cd/m^2 background light at −2, −1, 0, 1, 1.5, and 1.9 log cd · s/m^2 light intensity. The ERG tests were performed consecutively as follows: (1) scotopic, (2) flicker, (3) 10-min light exposure (30 cd · s/m^2 light intensity), and (4) photopic.

**SD-OCT imaging**

Mice were anesthetized using 100 mg/kg ketamine and 10 mg/kg xylazine intraperitoneally, and the pupils were dilated using atropine drops (5 mg/mL). The mouse was placed on a rodent alignment system with a bit bar (AIM-RAS, Biopoint, USA). The optic nerve was aligned to the center of the image on a SD-OCT imaging device with a mouse retina lens with a 50° field of view (Biopoint Envisu R2210 VHR, Biopoint, USA). The eyes were kept moisturized with eye drops (Styntace Ultra, Alcon) and an eye gel (Vidiscis carbogen, Bausch & Lomb). The following protocols (a-scan × b-scan × frame) for both eyes were run: (1) linear b-scan, 1.0 mm, 1,000 × 2 × 24 (fast fundus); (2) rectangular, 1.8 × 1.8 mm, 1,000 × 100 × 6 × 1 (high-resolution volume); (3) rectangular, 1.8 × 1.8 mm, 1,000 × 400 × 4 × 1 (square pixel volume); and (4) radial, 1.8 × 1.8 mm, 1,000 × 100 × 6 × 1 (high-resolution radial volume). The frames were averaged on the InVivoVue Reader software (Biopoint, USA) and analyzed on Diver 3.4.4 software (Biopoint, USA) for the VIP for the ONL.

**Optokinetic head-tracking response**

Optokinetic head-tracking response (OKT) was measured as previously described.11,23,24 The testing was performed in awake and non-restrained mice. Mice were placed on a pedestal surrounded by four displays that create a visual drum for the mice. The grating was set at 12 degrees/s (spatial frequency). The tracking was recorded in a clockwise (CW) or counterclockwise (CCW) direction. The drum rotation was random from trial to trial, and the experimenter made a forced-choice decision between CW and CCW rotation. The maximum spatial frequency capable of driving head tracking was determined first (visual acuity threshold). The contrast sensitivity was measured at 0.032, 0.064, 0.092, 0.103, 0.192, and 0.272 cycles/degree (spatial frequency). The recording was done twice per mouse. Mice were measured at 1, 3, 6, 9, and 12 months of age at random for the blinded experimenter. The eyes of mice measured in DLAAS, PBS, and rAAV-hCRB injection experiments were recorded separately (treated versus nontreated eye).

**Morphological and immunohistochemical analysis**

Eyes were collected at the time points of 1, 3, 6, 9, and 12 months of age. Mice injected with DL-AAA, PBS, and/or rAAV-hCRB were collected at 3 months of age. Mouse eyes were compared to (age-matched) littermates or the nontreated eye of the same animal. The eyes were marked on the superior side with a dye for superior-inferior orientation.6 For morphological analysis, eyes were enucleated and...
fixed at room temperature with 4% paraformaldehyde in PBS for 25 min. Then, the eyes were dehydrated for 30 min in 30%, 50%, 70%, 2× 90%, and 2× 100% ethanol, 50:50 ethanol/Technovit 7100 (Kulzer, Wehrheim, Germany), and finally in 100% Technovit 7100 overnight at 4°C. The eyes were sectioned (2 μm), stained with 1% toluidine blue, and mounted with Entellan. The sections were imaged on ×51 native resolution (bright field) on a slide scanner (3DHISTECH Pannoramic 250). For immunohistochemistry, we dehydrated the eyes for 30 min in 15% sucrose in PBS, followed by 30% sucrose in PBS (30 min). A detailed immunohistochemistry protocol is described in Buck et al. Cryosections (7 μm) were rehydrated in PBS, blocked (1 h), stained with the secondary antibody overnight at 4°C, washed in PBS three times (10 min), stained with the secondary antibody (Alexa Fluor 488, Alexa Fluor 555, Alexa Fluor 647, Cy3, or Cy5 for 1 h), washed three times in PBS (10 min), and mounted with Vectashield HardSet mounting medium containing DAPI (Vector Laboratories). A Leica TCS SP8 confocal microscope was used for image acquisition. Image analysis was done in Leica X, Fiji ImageJ, and Adobe Photoshop CC2018.

Antibodies

The following primary antibodies were used: glutamine synthetase (1:250; BD Biosciences), rhodopsin (1:500; Millipore), cone arrestin (1:500; Millipore), PKCζ (1:250; BD Biosciences), MPP4 AK4 (1:200; homemade1), CRB1 AK2 (pH 1.5) (1:200; homemade3), CRB2 (1:200), p120-catenin (1:100; BD Biosciences), GFAP (1:200; Dako), CD11b (1:100; BD Biosciences); PLVP (1:200; BD Pharmingen), and VE-cadherin (1:100; BD Biosciences).

DL-AAA

DL-AAA preparations were prepared on the day of injections. DL-AAA in PBS was dissolved/deprotonated by dropwise addition of 10 M NaOH. Then, the pH was raised to 7.3 by adding hydrochloric acid (37% w/v) or fuming. The volume was adjusted with PBS to 100, 150, or 200 μg/mL. The final solution was filter-sterilized (0.22 μm). A volume of 1 μL of DL-AAA or PBS was injected invitroretinally in one eye corresponding to 0 (PBS), 100, 150, or 200 μg of DL-AAA.

Generation and purification of rAAV vectors

The pAAV2-AmpR-inverted terminal repeat (ITR)-CMVmin-hCRB1co-sPA-ITR or pAAV-AmpR-ITR-CMV-hCRB2co-Spa-ITR plasmids consist of the flanking ITRs of AAV2, the minimal CMV promoter (for hCRB1 expression), the full-length CMV promoter (for hCRB2 expression), the human codon-optimized CRB1 or CRB2 cDNA, and a 48-bp synthetic polyadenylation (spA) sequence. The plasmid DNA was produced in Sure-2 cells and extracted on an anion exchange column (endotoxin range, 1–10 endotoxin units [EU]/μg). Endotoxin levels were not measured, and potential endotoxins were not removed from plasmid preps. The hCRB1 and hCRB2 coding sequences used in our rAAV-vectors are highly similar to the mCRB1 or mCRB2 coding sequence proteins. hCRB1/mCRB1 (1,406 and 1,405 aa; UniProtKB: P82279 and Q8VHS2, respectively) and hCRB2/mCRB2 (1,285 and 1,282 aa; UniProtKB: Q5JF48 and Q80Y88, respectively) proteins expressed at the OLM are similar in size and have similar domains (signal peptide, EGF-like, LamG, transmembrane, FERM, PDZ ERLI). The pAAV-hCRB plasmid, pHPlorer, and pXX2-ShH10F were co-transfected in 10× 15-cm dishes of 80% confluent HEK293T cells to generate rAAV2/ShH10F445F.CMV.hCRB2co.sPA or rAAV2/ShH10F445F.CMV.min.hCRB1co.sPA viral particles. After benzonase treatment, the lysates were ultracentrifuged onto an iodixanol density gradient. The purified rAAV stock was filter-sterilized and then concentrated on an ampiclon spin column (100,000 nominal molecular weight limit [NMWL]). All viral titers were determined by quantitative PCR. The final rAAV preparation was stored in 0.001% Pluronic F-68 in PBS at –80°C. No major contaminants were found for the rAAV-ShH10F445F.CMV.min.hCRB1co when a 1010 viral genome (vg) rAAV sample was denatured by lithium dodecyl sulfate (LDS) sample buffer, dithiothreitol (DTT); reducing agent), and heat (96°C; 5 min) and loaded on a SDS-PAGE gel (NuPAGE Bis-Tris mini gel) and protein-stained (Pierce silver stain kit; see Figure S5T).

rAAV and DL-AAA injection

Mice were anesthetized with 100 mg/kg ketamine and 5 mg/mL xylazine intraperitoneally, and the pupils were dilated using atropine drops (5 mg/mL). The pain was blocked locally by topical application of lidocaine (10 mg/mL) on a cotton swap to the eye and surrounding tissue. rAAV-hCRB1 or rAAV-hCRB2 was injected invitroretinally in one eye (around 50/50 right versus left eye) at P21-old mice (1 μL; 1010 vg). AAA-DL or PBS was injected invitroretinally in one eye in 2-month-old mice (1 μL). A Hamilton 10-μL needle was used for AAA, PBS, and rAAV-hCRB injections. The injected and noninjected eyes were washed with hypropemollose (3 mg/mL) drops, covered with chloramphenicol (Teva, 10 mg/g), and placed on a heating mat for recovery.

Quantification and measurement of photoreceptors

The thickness of the retina (ILM to OLM) was measured on plastic sections at 0.5, 1.0, 1.5, and 2.0 mm distance to the ONH, as previously described. Three measurements on three different sections were averaged per mouse. Bright-field images were taken on a slide scanner (3DHISTECH Pannoramic 250) at ×51 native resolution.

Normalized CRB2 protein and p120-catenin quantification and OLM breaks

We stained against nuclei (DAPI), CRB2 (second antibody: anti-rabbit Cy3) and p120-catenin (second antibody: anti-mouse Alexa Fluor 488) using a master mix at the same time with one CrbiKO.Crb2Flox/WT and one CrbiKO.Crb2lowMGC eye on each glass slide (total of six slides; total animals: n = 6 CrbiKO mice, n = 6 CrbiKO.Crb2lowMGC mice). Three sections per eye were imaged with the same laser and gain settings (12 images per mouse; >1 mm retinal length analyzed for periphery and central area). Two images in the periphery and two images in the central area (inferior and superior; four images per section) were made. Confocal microscopy was done in one run (total of 144 images, 12 animals). An additional group of CrbiKO.Crb2Flox/Flox, CrbiKO, Crb2KO, Crb2Flox/Flox, Crb1KO, and additional CrbiKO.Crb2Flox/WT retinas were similarly analyzed in one
immunohistochemical and microscopic section but normalized to the previous Crb1KO/Crb2Flox/WT fluorescence intensity values. The OLM area was defined as an area of ~2.5 μm above and 2.5 μm below the OLM. The fluorescence intensity was measured on grayscale images. The mean gray intensity per pixel over the OLM area was calculated. The mean gray intensity per pixel over the OLM area was calculated.

Supplemental Information can be found online at https://doi.org/10.1016/j.omtm.2020.12.012.

Acknowledgments
The authors thank Ilse Voshart, Ariadna Rocha Sierra, Martijn Koomen, Charlotte A. Andriessen, and Annelies Boonzaaijer-van der Laan for technical assistance, and all Wijnholds Lab members for advice on the manuscript. This work was supported by the Foundation Fighting Blindness (TA-GT-0715-0665-LUMC, to J.W.), the Netherlands Organisation for Health Research and Development (ZonMW grant 43200004, to J.W.), and by the Dutch blindness funds (Uitzicht 2013–13, to J.W. and Uitzicht 2018–6, to C.H.A. and J.W.); Rotterdamse Stichting Blindenbelangen, MaculaFonds, Stichting Blindenhulp, Landelijke Stichting voor Blinden en Slechtzienden, Algemene Nederlandse Vereniging ter Voorkoming van Blindheid, and Stichting Blinden-Penning.

Author Contributions
Conceptualization, T.M.B. and J.W.; methodology, T.M.B and J.W.; investigation, T.M.B. and R.M.V.; formal analysis, T.M.B. and J.W.; writing – original draft, T.M.B.; writing – review & editing, T.M.B., C.H.A., and J.W.; funding acquisition, C.H.A. and J.W.; resources, R.M.V. and J.W.; supervision, J.W.

Declaration of Interests
The authors declare no competing interests. The LUMC is the holder of patent number PCT/NL2014/050549, which describes the potential clinical use of CRB2. J.W. is listed as co-inventor of this patent and is an employee of the LUMC.

References
1. Talib, M., van Schooneveld, M.J., van Genderen, M.M., Wijnholds, J., Florijn, R.J., Ten Brink, J.B., Schalij-Delis, N.E., Dagnelle, G., Cremers, F.P.M., Wolterbeek, R., et al. (2017). Genotypic and phenotypic characteristics of CRB1-associated retinal dystrophies: a long-term follow-up study. Ophthalmology 124, 884–895.
2. Vincent, A., Ng, J., Gerth-Kahler, C., Tavares, E., Maynes, J.T., Wright, T., Tiwari, A., Tumber, A., Li, S., Hansen, J.V.M., et al. (2016). Biallelic mutations in CRB1 underlie autosomal recessive familial foveal retinoschisis. Invest. Ophthalmol. Vis. Sci. 57, 2637–2646.
3. Tsang, S.H., Burke, T., Oll, M., Yzer, S., Lee, W., Xie, Y.A., and Allikmets, R. (2014). Whole exome sequencing identifies CRB1 defect in an unusual maculopathy phenotype. Ophthalmology 121, 1773–1782.
4. Quinn, P.M., Buck, T.M., Mulder, A.A., Onohin, C., Alves, C.H., Vos, R.M., Bialecka, M., van Herwaarden, T., van Dijk, E.H.C., Talib, M., et al. (2019). Human iPSC-derived retinas recapitulate the fetal CRB1 CRB2 complex formation and demonstrate that photoreceptors and Müller glia are targets of AA.V5. Stem Cell Reports 12, 906–919.
5. Quinn, P.M., Mulder, A.A., Henrique Alves, C., Dejours, M., de Vries, S.L., Klooster, J., Dalkara, D., Koster, A.J., Jost, C.R., and Wijnholds, J. (2019). Loss of CRB2 in Müller glial cells modifies a CRB1-associated retinitis pigmentosa phenotype into a Leber congenital amaurosis phenotype. Hum. Mol. Genet. 28, 105–123.
6. Pellissier, L.P., Lunding, D.M.S., Tanimoto, N., Klooster, J., Vos, R.M., Richard, F., Sothilingam, V., Garcia Garrido, M., Le Bivic, A., Seeliger, M.W., and Wijnholds, J. (2014). CRB2 acts as a modifying factor of CRB1-related retinal dystrophies in mice. Hum. Mol. Genet. 23, 3759–3771.
7. van Rossum, A.G.S.H., Aartsen, W.M., Meuleman, J., Klooster, J., Malysheva, A., Versteeg, I., Arsanto, J.-P.P., Le Bivic, A., and Wijnholds, J. (2006). PalS1/Mpp5 is required for correct localization of Crb1 at the subapical region in polarized Müller glia cells. Hum. Mol. Genet. 15, 2659–2672.
8. van de Pavert, S.A., Kandtzhivazhieva, A., Malysheva, A., Meuleman, J., Versteeg, I., Levet, C., Klooster, J., Geiger, S., Seeliger, M.W., Rashbash, P., et al. (2004). Crumbs homologue 1 is required for maintenance of photoreceptor cell polarization and adhesion during light exposure. J. Cell Sci. 117, 4169–4177.
9. Quinn, P.M., Pellissier, L.P., and Wijnholds, J. (2017). The CRB1 complex: following the trail of Crumbs to a feasible gene therapy strategy. Front. Neurosci. 11, 175.
10. van de Pavert, S.A., Meuleman, J., Malysheva, A., Aartsen, W.M., Versteeg, I., Tonagel, F., Kamphuis, W., McCabe, C.J., Seeliger, M.W., and Wijnholds, J. (2007). A single amino acid substitution (Cys249Trp) in Crb1 causes retinal degeneration and deregulates expression of pituitary tumor transforming gene Pttg1. J. Neurosci. 27, 564–573.
11. Alves, C.H., Boon, N., Mulder, A.A., Koster, A.J., Jost, C.R., and Wijnholds, J. (2019). CRB2 loss in rod photoreceptors is associated with progressive loss of retinal contrast sensitivity. Int. J. Mol. Sci. 20, 4609.
12. van de Pavert, S.A., Sanz, A.S., Aartsen, W.M., Vos, R.M., Versteeg, I., Beck, S.C., Klooster, J., Seeliger, M.W., and Wijnholds, J. (2007). Single amino acid substitution (Cys249Trp) in Crb1 causes retinal degeneration and deregulates expression of pituitary tumor transforming gene Pttg1. J. Neurosci. 27, 564–573.
13. Pellissier, L.P., Quinn, P.M., Alves, C.H., Vos, R.M., Klooster, J., Flannery, J.G., Heimel, J.A., and Wijnholds, J. (2015). Gene therapy into photoreceptors and Müller glial cells restores retinal structure and function in CRB1 retinitis pigmentosa mouse models. Hum. Mol. Genet. 24, 3104–3118.
14. Boon, N., Wijnholds, J., and Pellissier, L.P. (2020). Research models and gene augmentation therapy for CRB1 retinal dystrophies. Front. Neurosci. 14, 860.
15. Alves, C.H., Sanz, A.S., Park, B., Pellissier, L.P., Tanimoto, N., Beck, S.C., Huber, G., Murtaza, M., Richard, F., Sridevi Gurubaran, I., et al. (2013). Loss of CRB2 in the mouse retina mimics human retinitis pigmentosa due to mutations in the CRB1 gene. Hum. Mol. Genet. 22, 35–50.
16. Quinn, P.M., Alves, C.H., Klooster, J., and Wijnholds, J. (2018). CRB2 in immature photoreceptors determines the superior-inferior symmetry of the developing retina to maintain retinal structure and function. Hum. Mol. Genet. 27, 3137–3153.

17. West, E.L., Pearson, R.A., Tschernert, M., Sowden, J.C., MacLaren, R.E., and Ali, R.R. (2008). Pharmacological disruption of the outer limiting membrane leads to increased retinal integration of transplanted photoreceptor precursors. Exp. Eye Res. 86, 601–611.

18. Kato, S., Ishiha, S., Sugawara, K., and Kawatsura, K. (1993). Cystine/glutamate antiporter expression in retinal Müller glial cells: implications for α-α-aminoacidotripeptide toxicity. Neuroscience 57, 473–482.

19. Yu, W.Q., Eom, Y.S., Shin, J.A., Nair, D., Gryzawcz, S.X.Z., Gryzawcz, N.M., Craft, C.M., and Lee, E.J. (2016). Reshaping the cone-mosaic in a rat model of retinitis pigmentosa: modulatory role of ZO-1 expression in DL alpha-aminoacidic acid reshap. PLoS ONE 11, e0151668.

20. Shen, W., Zhang, J., Chung, S.H., Hu, Y., Ma, Z., and Gilles, M.C. (2011). Submacular –α-α-aminoacidic acid eradicates primate photoreceptors but does not affect pigment or the retinal vasculature. Invest. Ophthalmol. Vis. Sci. 52, 119–127.

21. Jimeno, D., Feiner, L., Lillo, C., Teofilo, K., Goldstein, L.S.B., Pierce, E.A., and Williams, D.S. (2006). Analysis of kinesin-2 function in photoreceptor cells using synchronized Cre-loxP knockout of Kif3a with RHO-Cre. Invest. Ophthalmol. Vis. Sci. 47, 5039–5046.

22. Murray, S.A., Eppig, J.T., Smalley, D., Simpson, E.M., and Rosenthal, N. (2012). Cell-specific expression of plasma membrane calcium ATPase isoforms in retinal neurons. J. Comp. Neurol. 451, 1–21.

23. Prusky, G.T., West, P.W.R., and Douglas, R.M. (2000). Behavioral assessment of visual acuity in mice and rats. Vision Res. 40, 2201–2209.

24. Prusky, G.T., Alam, N.M., Beekman, S., and Douglas, R.M. (2004). Rapid quantification of adult and developing mouse spatial vision using a virtual optomotor system. Invest. Ophthalmol. Vis. Sci. 45, 4611–4616.

25. Prusky, G.T., and Douglas, R.M. (2004). Characterization of mouse cortical spatial vision. Vision Res. 44, 3411–3418.

26. de Souza, C.F., Ninison-Smith, L., Christie, D.L., Polkinghorne, P., McGhee, C., Kalloniatis, M., and Acosta, M.L. (2016). Macromolecular markers in normal human retina and applications to human retinal disease. Exp. Eye Res. 150, 135–148.

27. Krizaj, D., Demarco, S.J., Johnson, J., Strehler, E.E., and Copenhagen, D.R. (2002). Cell-specific expression of plasma membrane calcium ATPase isoforms in retinal neurons. J. Comp. Neurol. 451, 1–21.

28. Klaassen, I., Van Noorden, C.J.F., and Schlingemann, R.O. (2013). Molecular basis of the inner blood-retinal barrier and its breakdown in diabetic macular edema and other pathological conditions. Prog. Retin. Eye Res. 31, 89–99.

29. Newman, E.A. (2015). Glial cell regulation of neuronal activity and blood flow in the retina by release of gliotransmitters. Philos. Trans. R. Soc. Lond. B Biol. Sci. 370, 1–9.

30. Witmer, A.N., Newmark, J.A., Turunen, H.T., Farivar, T., Liu, J., Song, C., Ye, G.J., Pennock, S., Gaskin, C., Knop, D.R., and Shearnman, M.S. (2020). Inflammatory response to intravitreal injection of adeno-associated virus vector: relative contribution of genome and capsid. Hum. Gene Ther. 31, 80–89.

31. Newman, E., and Reichenbach, A. (1996). The Müller cell: a functional element of the retina. Trends Neurosci. 19, 307–312.

32. Franz, K., Grosse, J., Skatchkov, S.N., Schinkinger, S., Foja, C., Schild, D., Uckermann, O., Travis, K., Reichenbach, A., and Guck, J. (2007). Müller cells are living optical fibers in the vertebrate retina. Proc. Natl. Acad. Sci. USA 104, 8287–8292.

33. Newman, E.A. (2015). Glial cell regulation of neuronal activity and blood flow in the retina by release of gliotransmitters. Philos. Trans. R. Soc. Lond. B Biol. Sci. 370, 1–9.

34. Witmer, A.N., Blazquez, H.G., Weich, H.A., Altalto, K., Vrensen, G.F.J., and Schlingemann, R.O. (2002). Altered expression patterns of VEGF receptors in human diabetic retina and in experimental VEGF-induced retinopathy in monkey. Invest. Ophthalmol. Vis. Sci. 43, 849–857.

35. Son, S., Cho, M., and Lee, J. (2020). Crumbs proteins regulate lateral retinal vascular development required for vision. Biochem. Biophys. Res. Commun. 521, 939–946.

36. Kim, J.H., Kim, J.H., Yu, Y.S., Kim, D.H., and Kim, K.W. (2009). Recruitment of pericytes and astrocytes is closely related to the formation of tight junction in developing retinal vessels. J. Neurosci. Res. 87, 653–659.

37. Gardner, T.W., Lietz, E., Khan, S.A., Barber, A.J., Bonsall, D.J., Lesher, T., Rice, K., and Brennan, W.A. Jr. (1997). Astrocytes increase barrier properties and ZO-1 expression in retinal vascular endothelial cells. Invest. Ophthalmol. Vis. Sci. 38, 2423–2427.

38. Witmer, A.N., Vrensen, G.F.J.M., Van Noorden, C.J.F., and Schlingemann, R.O. (2003). Vascular endothelial growth factors and angiogenesis in eye disease. Prog. Retin. Eye Res. 22, 1–29.

39. Meeson, A.P., Argilla, M., Ko, K., Witte, L., and Lang, R.A. (1999). VEGF deprivation-induced apoptosis is a component of programmed capillary regression. Development 126, 1407–1415.

40. Koerber, J.T., Klimczak, R., Jang, J.H.H., Dalkara, D., Flannery, J.G., and Schaffer, D.V. (2009). Molecular evolution of adeno-associated virus for enhanced glial gene delivery. Mol. Ther. 17, 2088–2095.

41. Xiao, Z., Patrakka, J., Nukui, M., Chi, L., Niu, D., Betsholtz, C., Pikkarainen, T., Vainio, S., and Tryggvason, K. (2011). Deficiency in Crumbs homolog 2 (Crb2) affects gastrulation and results in embryonic lethality in mice. Dev. Dyn. 240, 2646–2656.

42. van den Hurk, J.A., Rashbash, P., Roepman, R., Davis, J., Voesenek, K.E., Arends, M.L., Zonneveld, M.N., van Roeke, M.H., Cameron, K., Rohrschneider, K., et al. (2005). Characterization of the Crumbs homolog 2 (CRR2) gene and analysis of its role in retinitis pigmentosa and Leber congenital amaurosis. Mol. Vis. 11, 263–273.
55. Dolón, J.F., Paniagua, A.E., Valle, V., Segurado, A., Arévalo, R., Velasco, A., and Lillo, C. (2018). Expression and localization of the polarity protein CRB2 in adult mouse brain: a comparison with the CRB1<sup>rd8</sup> mutant mouse model. Sci. Rep. 8, 11652.

56. Zhao, M., Andrieu-Soler, C., Kowalczuk, L., Paz Cortés, M., Berdugo, M., Dernigoghossian, M., Halili, F., Jeanny, J.C., Goldenberg, B., Savoldelli, M., et al. (2015). A new CRB1 rat mutation links Müller glial cells to retinal telangiectasia. J. Neurosci. 35, 6093–6106.

57. Shen, W., Li, S., Chung, S.H., and Gillies, M.C. (2010). Retinal vascular changes after glial disruption in rats. J. Neurosci. Res. 88, 1485–1499.

58. Li, Y., Busoy, J.M., Zaman, B.A.A., Tan, Q.S.W., Tan, G.S.W., Barathi, V.A., Cheung, N., Wei, J.Y., Hunziker, W., Hong, W., et al. (2018). A novel model of persistent retinal neovascularization for the development of sustained anti-VEGF therapies. Exp. Eye Res. 174, 98–106.

59. Roesch, K., Jadhav, A.P., Trimarchi, J.M., Stadler, M.B., Roska, B., Sun, B.B., and Cepko, C.L. (2008). The transcriptome of retinal Müller glial cells. J. Comp. Neurol. 509, 225–238.

60. Alves, C.H., and Wijnholds, J. (2018). AAV gene augmentation therapy for CRB1-associated retinitis pigmentosa. In Methods in Molecular Biology, C.J.F. Boon and J. Wijnholds, eds. (Humana Press), pp. 135–151.

61. Buck, T.M., Pellissier, L.P., Vos, R.M., van Dijk, E.H.C.C., Boon, C.J.F., and Wijnholds, J. (2018). AAV serotype testing on cultured human donor retinal explants. In Methods in Molecular Biology, C.J.F. Boon and J. Wijnholds, eds. (Humana Press), pp. 275–288.

62. Heynen, S.R., Meneau, I., Caprara, C., Samardzija, M., Imsand, C., Levine, E.M., and Grimm, C. (2013). CDC42 is required for tissue lamination and cell survival in the mouse retina. PLoS ONE 8, e53806.