Retinal Cell Biology

Novel Animal Model of Crumbs-Dependent Progressive Retinal Degeneration That Targets Specific Cone Subtypes

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Purpose. Human Crb1 is implicated in some forms of retinal degeneration, suggesting a role in photoreceptor maintenance. Multiple Crumbs (Crb) polarity genes are expressed in vertebrate retina, although their functional roles are not well understood. To gain further insight into Crb and photoreceptor maintenance, we compared retinal cell densities between wild-type and Tg(RH2-2:Cr2b-sfEX/RH2-2:GFP)pt108b transgenic zebrafish, in which the extracellular domain of Crb2b-short form (Crb2b-sfEX) is expressed in the retina as a secreted protein, which disrupts the planar organization of RGB cones (red, green, and blue) by interfering with Crb2a/2b-based cone-cone adhesion.

Methods. We used standard morphometric techniques to assess age-related changes in retinal cell densities in adult zebrafish (3 to 27 months old), and to assess effects of the Crb2b-sfEX transgene on retinal structure and photoreceptor densities. Linear cell densities were measured in all retinal layers in radial sections with JB4-Feulgen histology. Planar (surface) densities of cones were determined in retinal flat-mounts. Cell counts from wild-type and pt108b transgenic fish were compared with both a “photoreceptor maintenance index” and statistical analysis of cell counts.

Results. Age-related changes in retinal cell linear densities and cone photoreceptor planar densities in wild-type adult zebrafish provided a baseline for analysis. Expression of Crb2b-sfEX caused progressive and selective degeneration of RGB cones, but had no effect on ultraviolet-sensitive (UV) cones, and increased numbers of rod photoreceptors.

Conclusions. These differential responses of RGB cones, UV cones, and rods to sustained exposure to Crb2b-sfEX suggest that Crb-based photoreceptor maintenance mechanisms are highly selective.

Keywords: retinal cell density, retinal degeneration, photoreceptor degeneration, retinitis pigmentosa, Leber’s congenital amaurosis, Crumbs

The maintenance of vertebrate retinal photoreceptors is critical for normal vision because photoreceptors normally persist for the entire life span; if they degenerate in mammals, they cannot be replenished. However, understanding mechanisms of photoreceptor maintenance is challenging because photoreceptor degeneration and retinal dystrophy are linked to dozens of genes, including the Crumbs (Crb) family.1–3 For example, pathogenic mutations of human Crumbs 1 (CRB1) cause severe dystrophies, accounting for 5% of retinitis pigmentosa (RP) and 10% of Leber’s congenital amaurosis (LCA).4–9 Despite the relative prevalence of CRB1-based RP and LCA, the molecular etiology of CRB1 in photoreceptor degeneration is not understood.10

Crb proteins are Type I transmembrane proteins that regulate apicobasal epithelial polarity11,12 and cell-cell adhesion.13 In the retina, Crb proteins control epithelial polarity and apical membrane size of photoreceptors in both Drosophila and vertebrates.4,14–16 A single Crb gene exists in the fly, but in mammals, three homologs, Crb1, Crb2, and Crb3, are expressed in photoreceptors and/or Müller glia in the retina.10,17–20 In zebrafish retina, crb1, crb2a, and crb2b gene homologs are expressed, and two transcriptional start sites in the crb2b gene yield two proteins, Crb2b-long form (Crb2b-lf), and Crb2b-short form (Crb2b-sf), that differ in length of their extracellular domain.21–23

To gain insight into potential Crb functions in photoreceptor maintenance, we first analyzed retinal cell densities as a function of age in wild-type zebrafish to provide a baseline for evaluating the effects of genetic abnormalities in Crb on photoreceptor maintenance. To our knowledge, this is the first comprehensive analysis of retinal cell densities in zebrafish, although similar studies have been conducted in other teleost species.24–27 We then compared photoreceptor densities in wild-type with those in the transgenic zebrafish Tg(RH2-
2-Crb2b-sfEX/RH2-2:GFP)pt108b (subsequently referred to as pt108b). In pt108b, blue cones (and less frequently, rods and green cones) secrete Crb2b-sfEX (the extracellular domain of Crb2b-short form), which interrupts Crb2a/2b-mediated adhesion among RGB cones (red, green, and blue cones) by competitively binding to the extracellular domains of endogenous Crb2a and Crb2b. Although the inhibition of Crb-mediated cone-cone adhesion disrupts the planar spatial patterning of cones, and destroys the regular lattice arrangement of the cone mosaic, it is unclear whether or not photoreceptor maintenance is affected. Here we report differential effects on maintenance of rods, RGB cones, and UV cones after interference with Crb-based adhesion.

**Materials and Methods**

**Zebrafish Care**

AB and TL wild-type, and Tg(RH2-2:Crb2b-sfEX/RH2-2:GFP)pt108b transgenic zebrafish were maintained in a 14-hr light and 10-hour dark cycle with standard husbandry procedures. All animal protocols conformed to regulations at the University of Pittsburgh and the University of Michigan as well as to the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research.

**Immunohistochemistry**

To verify photoreceptor subtypes in pt108b fish, we performed immunohistochemistry with the following primary and secondary antibodies: rabbit polyclonal anti-UV opsin for UV cone outer segments (gift from David Hyde, PhD), anti-Zp1 for green/red double cones (ZFIN), Cy3-conjugated goat antimouse IgG (Catalog no. 115-165-166; Jackson Immunoresearch Labs, Jackson, PA, USA), and Cy5-conjugated donkey anti-rabbit IgG (Catalog no. 711-175-152; Jackson Immunoresearch Labs). Images were collected with a Fluoview FV1000 confocal microscope (Olympus, Tokyo, Japan).

**JB4 Plastic Sectioning and Feulgen Staining**

Zebrafish eyes were fixed in 4% paraformaldehyde in PBS at room temperature overnight. The fixed samples were dehydrated in alcohol and embedded in JB4 resin (JB4 embedding kit; Polysciences, Inc., Warrington, PA, USA), following the manufacturer's protocol. The samples were sectioned at 3-μm thickness with a Shandon Finesse microtome (Thermo Electron Co., Waltham, MA, USA). Plastic sections were collected on glass slides, stained with the Feulgen reaction to visualize the cell nuclei, observed under an Olympus UPlanSApo 20/0.80 oil objective (Olympus BX60 microscope), and photographed with a SPOT RT camera, software version 4.6 (Diagnostic Instruments, Inc., Sterling Heights, MI, USA).

**Evaluating Linear Retinal Cell Densities**

We define “linear cell density” as the number of cones of a given subtype counted in a curved segment of retina aligned with the retinal layers in a radial section through the center of the eye. To evaluate cell densities across the retina in Feulgen-stained sections, eyes were sectioned radially in either dorsal-ventral or nasal-temporal (anterior-posterior) planes (Fig. 1A; Supplementary Fig. S1); three consecutive sections through the retinal center were collected and divided into five local regions with an angular subtense of 36 degrees: two peripheral regions, two intermediate regions, and one central region (Fig. 1B). The newly generated tissue in the growing marginal regions (approximately 100 μm) was excluded from quantification (Fig. 1B, black areas). Thus, retinas were systematically sampled at nine local regions representing the midline regions of four retinal quadrants: nasal, dorsal, temporal, and ventral (Fig. 1C).

In each sampling region, to determine the linear cell density, we counted cell nuclei in a retinal segment of 200 μm in length parallel to cellular layers (Fig. 1B). Retinal cells were quantified into six categories according to nuclear morphology and position (Figs. 1D–F): retinal ganglion cells (nuclei localized to the basal layer of the retina), basal inner nuclear layer cells (including amacrine cells, Müller cells, and bipolar cells), horizontal cells (nuclei horizontally elongated and localized to the most apical tier of the inner nuclear layer), rods (nuclei round and more densely stained than cone nuclei, and localized to the basal half of the outer nuclear layer), UV cones (nuclei also rounded but larger and stained lighter than rods, localized apical to rod nuclei and basal to the outer limiting membrane [OLM], and regularly separated from each other), and RGB cones (nuclei elongated in the apicobasal direction, stained lighter than rod nuclei, and localized largely apical to the OLM). Five eyes were sampled for each age and each local region; cell nuclei in three consecutive radial retinal sections were counted and averaged. We performed a paired t-test and ANOVA with post hoc Tukey statistical tests, using JMPPro 12.0.1 (SAS Institute, Inc., Cary, NC, USA) to compare cell densities between different conditions.

**Evaluating Planar Cone Densities by Flat-Mount Imaging**

We define “planar cone density” as the number of cones contained within a unit of surface area in a retinal flat-mount. Retinal flat-mount immunohistochemistry was performed as previously described. In brief, dark-adapted retinas were dissected and flattened with radial relaxing cuts including a large ventral cut, fixed at 4°C overnight in 4% paraformaldehyde in 0.1M phosphate buffer with 5% sucrose, then processed for immunohistochemistry. Primary and secondary antibody incubations were at the following dilutions: mouse anti-ZO-1A-12 (Invitrogen, Camarillo, CA, USA), 1:200; rabbit anti-active Caspase-3 Clone C92–605 (RD Biosciences, San Jose, CA, USA), 1:1000; Dylight 550-conjugated anti-mouse IgG (Thermo Scientific, Waltham, MA, USA), 1:200; and DyLight 650-conjugated anti-rabbit IgG (Thermo Scientific), 1:200. Antigen retrieval in 10-nM sodium citrate with 0.05% Tween20 (pH 6.0) was performed for ZO-1 staining. Images were captured with a Zeiss Axio Image Z1 epifluorescent microscope (Carl Zeiss Microimaging, Inc., Thornwood, NY, USA) with an ApoTome module for optical sectioning using structured illumination. Images were processed for two-dimensional (2D) projection (xy plane) or three-dimensional (3D) “cut views” (xz and yz planes) with AxioVision SE64 or Adobe Photoshop CS6 Extended (Adobe Systems, Inc., San Jose, CA, USA).

**Results**

**Age-Related Changes in Retinal Cell Densities in Wild-Type Zebrafish**

To gain insight into Crb functions in photoreceptor maintenance by analyzing pt108b effects on the retina, we first need...
FIGURE 1. Linear cell densities of six categories of retinal cells were evaluated by JB4-Feulgen histology. (A) Zebrafish eyes were sectioned in either the nasal-temporal (anterior-posterior) or ventral-dorsal axis. (B) Each retinal section was partitioned by a vertical line to exclude the developing marginal region (black) from the differentiated retina, which was further divided into five regions each with angular subtense of 36 degrees. We counted nuclei in 200-μm linear segments (gray bars). (C) Spatial relationships among the nine sampled retinal regions. (D–F) JB4-Feulgen histology illustrates the localization and morphologies of the six retinal cell categories at 1 mpf (D, E; the inset box in D outlines the boundary of E) and 15 mpf (F). The dashed line indicates the location of the OLM. GC, ganglion cell layer; INL, inner nuclear layer; ONL, outer nuclear layer.
to understand the effect of normal aging on retinal cell density. Age-related changes in cell densities can be caused by many factors, including cell proliferation, cell death, and compression or stretching of the retina, all of which have been described in association with growth and differentiation of vertebrate retinas.31–35

Linear densities of most retinal cell categories in the central region declined progressively by approximately 30% to 40% in adult wild-type zebrafish from 3 mpf (months postfertilization) to 27 mpf (Figs. 2A–F). Correlating with a reduction in linear cell densities, the thickness of the ganglion cell layer was reduced from two to three cells thick at 3 mpf to a single cell layer at 27 mpf (Fig. 2G). Similar trends were also found in the intermediate and peripheral regions (Supplementary Figs. S2–S5). In the central region, UV cone linear densities were not significantly changed, although density declined in other retinal regions (Fig. 2E; Supplementary Figs. S2–S5). Although rod linear density did not show a consistent trend, it was slightly higher in some regions at 15 mpf (Fig. 2D; Supplementary Figs. S2, S4), but otherwise relatively constant.

**Topographical Variations of Retinal Cell Densities in Wild-Type Zebrafish**

In addition to age-related changes in linear cell density, we observed topographic variations across the retina, as illustrated in the 3D histogram plots of counts from 15-mpf retinas (Fig. 3). We chose 15 mpf because retinal structure has largely stabilized after the rapid growth in juvenile and young adult stages (3 mpf), but retinal structure has not yet been affected by aging as at 27 mpf (Fig. 2G; Supplementary Fig. S1).

We compared data both within a quadrant radially from central to peripheral, and between quadrants circumferentially across intermediate and peripheral regions. In the radial dimension within a quadrant, linear cell densities tended to decrease from central to intermediate to peripheral in all four quadrants for most retinal cell categories (Supplementary Tables S1, S2), especially ganglion cells (Fig. 3A), horizontal cells (Fig. 5C), UV cones (Fig. 3E), and RGB cones (Fig. 3F), but not basal inner nuclear layer cells (Fig. 3B). In the circumferential dimension (i.e., between quadrants) rod photoreceptor densities vary significantly, and ganglion cells and RGB cones show some variation (Supplementary Table S3). Rod photoreceptor densities in both intermediate and peripheral regions were highest in the dorsal and nasal quadrants (Fig. 3D; Supplementary Tables S1, S5). Conversely, RGB cone density in the ventral intermediate region was higher than in the nasal quadrant (Fig. 3F), and ventral ganglion cells showed a trend toward increased density in the ventral and temporal quadrants compared to dorsal (Fig. 3A; Supplementary Tables S1, S5).

We confirmed that topographical variations in a planar cone density analysis of adult 6-mpf retinal flat-mounts, with immunolabeling of ZO1 to reveal the profiles of photoreceptors and Müller glia at the level of OLM. Cones profiles, identified by their shapes, sizes, and stereotypic pattern, were sampled in approximately the same nine local regions defined for the linear cell density analysis (Figs. 1A–C, 4A). The crystalline lattice cone mosaic pattern characteristic of adult zebrafish retina (Fig. 4D–G) is absent in the central region (Fig. 4H), which represents retina that was present in the larval eye.36 This planar cone density analysis revealed dramatic topographic variations in cone sizes: In particular, in the central region (Fig. 4H) and dorsal quadrant (Figs. 4G, 4G'), UV cones are much larger than RGB cones, and in the ventral quadrant (Figs. 4E, 4E'), all cone profiles are small.

Similar to the linear cone densities, planar cone densities generally decline from central to intermediate to peripheral regions, except in the ventral retina (Figs. 4B, 4C; Supplementary Table S4). Circumpolar comparisons between quadrants confirmed that the highest planar cone density was in the ventral intermediate region (Figs. 4B, 4C; Supplementary Table S4). Although we did not count rod profiles in these flat-mount preparations, the higher density of rods in the dorsal region can be appreciated by the accumulation of small, round rod profiles in the spaces between the cones (Fig. 4G inset; also see Salbreux et al., 2012).36 Thus, the results of the linear and planar cell density counts both show that cones are denser in ventral retina than in dorsal retina, and rods are denser in dorsal retina than in ventral retina.

**Secreted Crb2b-sfES Has Differential Effects on Photoreceptor Maintenance**

In zebrafish, Crb1 is expressed by all cones; Crb2a by cones, rods, and Müller glia; and Crb2b by RGB cones only (Fig. 5A). To gain insight into whether Crb-mediated cone-cone adhesion plays a role in photoreceptor maintenance, we analyzed age-related retinal cell density changes in pt108b transgenic fish, in which Crb2b-sfES is secreted mainly by blue cones, a much more restrictive expression pattern compared with endogenous Crb2a and Crb2b (Fig. 5A; also see Zou et al., 2012).22 We showed previously that expression of Crb2b-sfES gradually disrupts the regular alignment of RGB cones into mirror-image pentamers (G-R-B-R-G) by physically interfering with cone-cone adhesions mediated by Crb2a and Crb2b.22 We reasoned that if Crb2b-sfES disrupts cone adhesion by physically binding to endogenous Crb proteins, Crb2b-sfES may also interfere with signaling pathways that might regulate photoreceptor survival via Crb proteins.10 Comparing flat-mount images of ZO1 cone profiles in 4-mpf young adult pt108b fish with 18-mpf aged adults revealed late-onset disruption of RGB cone spatial organization (Fig. 5B). Importantly, the cone mosaic pattern in 4-mpf pt108b adults is highly organized throughout the retina (Supplementary Fig. S6), with the same topographic variations in cone sizes seen in wild-type adults (Fig. 4), suggesting that Crb2b-sfES interferes with photoreceptor maintenance, but not cone photoreceptor development.

To quantitatively evaluate whether photoreceptor maintenance is affected in the pt108 line, we first examined the histologic characteristics of pt108b retinas in the central and intermediate regions of aged fish (27 mpf). We found no disruptions in overall retinal lamination in pt108b, again consistent with a lack of effect of Crb2b-sfES on retinal development. However, in the aged adult pt108b retinas, very few cone photoreceptor nuclei located apical to the OLM were elongated (Figs. 5C, 5D, 5E), which is a distinctive characteristic of RGB cone nuclei in the wild-type retina (Figs. 5C–E). In their place were round, lightly stained, and sparsely distributed nuclei (Fig. 5C, arrowheads) that resembled UV cones, whose nuclei normally reside below the OLM in wild-type retina (Fig. 5C, arrowhead). Rod nuclei remained small, round, and darkly stained and localized basal to the OLM in pt108b at 27 mpf (Figs. 5C, 5D, 5E), similar to wild-type retinas (Figs. 5C–E).

We verified cone identities with immunohistochemistry in 27-mpf pt108b and found that very few cones expressed arrestin 3a (a regulator of G protein–coupled receptors that is selectively expressed in zebrafish green/red cones and is recognized by the Zpr1 monoclonal antibody), particularly in the central region (Figs. 5D, 5D'). Furthermore, most nuclei of the remaining Zpr1-positive cells had lost their typical elongated morphology and resembled nuclei of UV cones, round and lightly stained (Fig. 5D'; arrow), although most of the cones with round and lightly stained nuclei above the OLM in pt108b retinas were UV opsin-positive (Fig. 5D', arrow-
FIGURE 2. Linear cell density in the central region declines progressively with age, with the exception of rods. (A–F) Six cell categories of linear cell density at 3, 15, and 27 mpf. (A) Ganglion cells, (B) basal inner nuclear cells, (C) horizontal cells, (D) rods, (E) UV cones, and (F) RGB cones. Mean ± 1 SD, n = 5. (G) JB4-Feulgen histology of central retina at 3, 15, and 27 mpf.
These morphologic changes indicate that the standard for distinguishing UV cones from RGB cones by their nuclear morphologies would overestimate UV cones and underestimate RGB cones in *pt108b* retina.

In recognition of these limitations, we quantified differential effects of *pt108b* on photoreceptor maintenance with the following simplified criteria/assumptions: (1) round, lightly stained, and apically localized nuclei were counted as UV cone nuclei, recognizing that some of them may have been deformed RGB cones; (2) elongated, lightly stained, and apically localized nuclei were counted as RGB cone nuclei; and (3) round, darkly stained, and basally localized nuclei were...
counted as rod nuclei. With these standards, the overestimation of UV cones (or underestimation of RGB cones) should be very limited in the central retina of aged *pt108b*, because most apically localized and lightly stained round nuclei were UV cone nuclei (Figs. 5C', 5D'). In central retina (Fig. 6A), cell counts indicated that at both 15- and 27-mpf linear density of UV cones in *pt108b* was unchanged, whereas RGB cones were decreased compared with wild-type (Fig. 6B). Loss of RGB cones in *pt108b* was progressive, declining by approximately 50% from 15 to 27 mpf. Overpopulation of rods was seen at 27 mpf (Fig. 6B).

To examine changes in photoreceptor densities in *pt108b* fish in the context of normal aging, we devised a photoreceptor maintenance index to quantify the effects of Crb2b-sfEX on photoreceptor maintenance (Fig. 6C): a positive value suggests enhanced photoreceptor maintenance/addition, and a negative value suggests a net loss of photoreceptors. (The calculation of these indexes measured changes in photoreceptor cell densities between 15 and 27 mpf ages in *pt108b* normalized to wild-type to eliminate any potential developmental effects on production of photoreceptors, thereby revealing only age-related effects on maintenance during this period.) For the central retinal region, the index values clearly demonstrated that Crb2b-sfEX produced selective loss of RGB cones, almost no effect on UV cones, and an overpopulation of rods (Fig. 6D).

We applied the same index to the remaining retinal regions, but limited the classification of photoreceptors to rods and cones because it was difficult to distinguish UV cones from the residual RGB cone nuclei, which occurred more frequently in the peripheral and intermediate retinal regions. Similarly, across the *pt108b* retina, Crb2b-sfEX caused rod densities to increase and cone densities to decrease (Supplementary Fig. S7).

Taken together, these data show that secretion of an extracellular fragment of Crb2b in the retina differentially affects photoreceptor maintenance, resulting in overpopulation of rods, no effect on UV cones, but selective death of RGB cones.

**Secreted Crb2b-sfEX Changes Planar Cone Patterning**

To determine the effect of secreted Crb2b-sfEX on patterning and morphology in the planar dimension, we examined retinal flat-mounts in aged wild-type and *pt108b* fish. The topographic variations in cone sizes observed in young wild-type adults at 6 mpf (Figs. 4D–H) are still present in older adults at 26 mpf, and the stereotypic lattice mosaic pattern of peripheral and intermediate regions is also retained (Figs. 7A–E). In contrast, in aged *pt108b* retinas (18 and 20 mpf), the mosaic patterning of the surviving cones was disrupted in dorsal and nasal regions (Fig. 7A', 7A''–D', 7D''), as reported previously.\(^\text{22}\)
FIGURE 5. Secrected Crb2b-sfEX affects photoreceptor patterning and maintenance. (A) The expression patterns of Crb genes are highlighted with different colors superimposed on a transmission electron microscopic image of a tangential section through the photoreceptor layer. The Crb2b-sfEX transgene in pt108b is mostly expressed in blue cones; endogenous crb2a is expressed in all photoreceptors and Müller cells; endogenous crb2b is expressed selectively in RGB cones. (B) In flat-mount retinal preparations at 4 mpf, ZO-1 immunolabeled apical profiles of photoreceptors in pt108b retina form a mosaic pattern, but the regular organization is degraded by 18 mpf. (C, C’) JB4-Feulgen staining of retinal cross sections from wild-type...
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The initial increase in rod densities in some retinal regions in adult zebrafish, especially in the pt108b fish, may represent a retinal specialization corresponding to the part of the visual field located above and in front of the animal, which may be important for predator and prey detection. 47-48

**Crb Proteins Function in Photoreceptor Maintenance**

With normal age-related and topographical variations in zebrafish photoreceptor densities as a reference, we found progressive and selective apoptotic death of RGB cones in aged adult pt108b fish, with no effect on UV cones, and overpopulation of rods. In young adult pt108b fish (4 mpf), retinal structure, lamination, and cone mosaic patterning were all normal, but by 15 mpf, disruption of cone photoreceptor morphology and patterning and loss of RGB cones was apparent in retinal sections as well as retinal flat-mounts. Progressive loss of RGB cones continued to 27 mpf, and the anti-active caspase 3 profiles in the photoreceptor layer suggested that cones were dying via apoptosis. Our quantitative analysis of photoreceptor densities was specifically designed to examine age-related rather than developmental effects, and so we excluded the marginal retina, where new cells are being added to the growing retina. Furthermore, the normal appearance of the retina in young adult pt108b fish suggested that photoreceptor differentiation occurred normally. Expression of the Crb2b-sfEX construct was under the control of an opsin promoter, and expression of opsin protein in differentiating photoreceptors is a relatively late developmental event, 36,37 so the initial stages of photoreceptor commitment and differentiation preceded opsin expression. Because production of retinal cells, including photoreceptors, continues in the growing adult retina, newly added cells would be generated in an environment in which Crb2b-sfEX is present. Despite this, the retina appears to develop normally, and the effect on cone photoreceptor maintenance is delayed.

The selective death of RGB cones suggests that Crb2b-sfEX preferentially interferes with function and maintenance of RGB cones. Although the secreted Crb2b-sfEX physically interacts with the broader family of zebrafish Crb proteins (including Crb2a, Crb2b, and Crb1), as suggested previously 22 by in vitro immunoprecipitation analysis (and data not shown), the lack of
FIGURE 6. Secreted Crb2b-sf promotes RGB cone degeneration and rod overpopulation. (A) JB4-Feulgen histology illustrates nuclear morphologies and distributions of photoreceptors in the central area of WT and pt108b retinas at 15 mpf and 27 mpf. (B) Histograms of average linear densities of photoreceptors in central retina of WT and pt108b at 15 and 27 mpf. Mean ± 1 SD, n = 5. (C) Formula for the photoreceptor maintenance index. (D) In the central retina in pt108b, the photoreceptor maintenance indexes show loss of RGB cones, excess rods, and no effect on UV cones.
an effect on UV cones suggests that Crb2b-sfEX interference with Crb2a- and Crb1-mediated mechanisms may not be consequential for photoreceptor maintenance. If this interpretation is correct, the overpopulation of rods may be simply a secondary effect of normal mechanisms of growth-related rod photoreceptor production in the teleost retina. For example, we previously reported overproduction of rods in light-damaged zebrafish retinas with selective loss of UV cones. Nevertheless, we cannot rule out the possibility that in pt108b, Crb2b-sfEX interference with Crb2a, which is expressed by both rods and Müller glia, may directly promote rod overpopulation.

The selective death of RGB cones is observed only in older adults. This slow, progressive degeneration may be the reason why Müller glial cells were not stimulated to regenerate lost RGB cones, as has been reported previously in a zebrafish genetic model of rod degeneration in which loss of rods was neither extensive nor acute. In line with the observation that cones can survive when rods degenerate in zebrafish, the opposite fates of rods and cones in pt108b supports the notion that the maintenance of rods and cones is regulated differently and their survival is not interdependent in zebrafish.

The selective death of cone subtypes has not been observed in genetic mouse models that affect a single Crb gene. Although mutations in the human CRB1 gene can cause severe retinal dystrophies, loss of Crb1 in mice (including the spontaneous rd8 mutation and Crb1 knockout) show a relatively mild phenotype, with focal regions of photoreceptor disorganization, whereas conditional knockout of Crb2 in mice has a more severe and progressive retinal degeneration phenotype that more closely models human disease.

Based on molecular phylogeny of opsins, zebrafish red cones, among the cone subtypes lost in pt108b, are the homolog of mammalian L/M cones; by contrast, the UV cones, persistent in pt108b, are homologous to mammalian S cones. Thus, the pt108b zebrafish line, which targets Crb2b function, provides a useful new tool to investigate the mechanisms of Crb proteins in photoreceptor maintenance and survival.

Although the downstream molecular pathways of Crb2b regulation in zebrafish photoreceptors are yet to be determined, one possible target is the Hippo signaling pathway. Crb is known to activate the Hippo kinase cascade that phosphorylates and inhibits the transcription cofactor YAP/TAZ, which is required for diverse biological functions, including cell proliferation as well as apoptosis, depending
on cellular context.\textsuperscript{53–56} It is tempting to speculate that Crb2b may regulate Hippo signaling to promotes RGB cone mainte-

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