Dawn and dusk peaks of outer segment phagocytosis, and visual cycle function require Rab28

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Abstract
RAB28 is a farnesylated, ciliary G-protein. Patient variants in RAB28 are causative of autosomal recessive cone-rod dystrophy (CRD), an inherited human blindness. In rodent and zebrafish models, the absence of Rab28 results in diminished dawn, photoreceptor, outer segment phagocytosis (OSP). Here, we demonstrate that Rab28 is also required for dusk peaks of OSP, but not for basal OSP levels. This study further elucidated the molecular mechanisms by which Rab28 controls OSP and inherited blindness. Proteomic profiling identified factors whose expression in the eye or whose expression at dawn and dusk peaks of OSP is dysregulated by loss of Rab28. Notably, transgenic overexpression of Rab28, solely in zebrafish cones, rescues the OSP defect in rab28 KO fish, suggesting rab28 gene replacement in cone photoreceptors is sufficient to regulate Rab28-OSP. Rab28 loss also perturbs function of the visual cycle as retinoid levels of 11-cRAL, 11cRP, and atRP are significantly reduced in larval and adult rab28 KO retinas (p < .05). These data give further understanding on the molecular mechanisms of RAB28-associated CRD, highlighting roles of Rab28 in both peaks of OSP, in vitamin A metabolism and in retinoid recycling.

Abbreviations: cpd, cycles per degree; CRD, cone-rod dystrophy; CT, circadian time; dpf, days post fertilization; IFT, intraflagellar transport; mpf, months post fertilization; OKR, optokinetic response; OS, outer segment; OSP, outer segment phagocytosis; ZT, Zeitgeber time.

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1 | INTRODUCTION

Visual impairment resulting from photoreceptor dystrophy is a common phenotype of inherited ciliopathies, a group of disorders characterized by ciliary dysfunction. The photoreceptor outer segment (OS) is a highly modified primary cilium. As such, the OS is an intricate membranous organelle which detects light photons and converts them into neurotransmitter signals via phototransduction. The molecular machinery required for normal ciliary formation, transport, and signaling is also required for photoreceptor OS development and function.

Rod OS discs appear to develop from nascent, evaginated, ciliary ectosomes whose shedding is blocked by peripherin. While rod OS are comprised of enclosed discs, cone OS develop differently, forming open ended evaginations called lamellae. Both discs and lamellae are established via actin polymerization regulated by Arp2/3 and steadily shift toward the OS tip. This continual process works in synergy with the highly regulated process of outer segment phagocytosis (OSP). As new OS discs and lamellae are continuously formed from evaginated ectosomes produced at the OS base, the oldest discs and lamellae are removed from the OS tip, and those are phagocytosed by the supporting retinal pigment epithelium (RPE). The photoreceptor outer segment (POS) is exposed to high levels of light; thus, phagocytosis of the oldest OS tips by the RPE prevents the accumulation of photo-oxidative compounds and maintains photoreceptor health and viability.

Outer segment phagocytosis deficits are directly linked to progressive forms of inherited blindness. Defective OSP may also contribute to age-related macular degeneration. However, a large proportion of the molecular mechanisms instigating OSP in photoreceptors remain to be identified and characterized, leaving a knowledge gap in retinal biology and how perturbed OSP leads to retinal or macular degeneration. We consider OSP as a process encompassing demarcation of the OS area to be shed, overlapping with phagocytosis (consisting of three distinct phases, recognition/binding, internalization, and digestion). Furthermore, OSP by the RPE is a putative alternative source of essential retinoids to integrate into the visual cycle, and support vision. Approximately 7%–10% of distal OS are phagocytosed daily, providing a significant source of retinal following opsin proteolysis. Discovery of ABCA4 expression in the RPE cells, in addition to photoreceptor disc membranes, provided strong evidence of an ABCA4 role to recycle this pool of free retinaldehyde via the visual cycle to prevent formation of toxic dimers (bisretinoids).

Previous studies also postulate that bisretinoids associated with the visual cycle could be transformed following phagocytosis in the acidic RPE phagolysosomes to A2E, the major lipofuscin fluorophore following phagocytosis in the acidic RPE phagolysosomes.

In 2012, the small ciliary GTPase Rab28 was identified as the first Rab GTPase associated with human cone-rod dystrophy (CRD). Rab28 null and hypomorphic alleles cause autosomal recessive cone-rod dystrophy (arCRD) with eight pathogenic alleles attributed to date.In Caenorhabditis elegans, we have shown that Rab28 is an intraflagellar transport (IFT) and BBSome-associated ciliary protein, which regulates extracellular vesicle biogenesis in a subset of ciliated neurons. In addition, we employed zebrafish knockout and transgenic reporter models to probe Rab28's localization and, we previously described the localization, GTP/GDP nucleotide regulation, and a protein interaction of Rab28 in cone photoreceptors. Significantly, the rab28 KO zebrafish model shows a dawn OSP phenotype similar to the mouse rab28 KO model; however, no retinal degeneration was observed in the zebrafish model.

This study further investigated the role of Rab28 in the zebrafish retina, primarily focusing on its role in OSP. As the molecular mechanisms of cone OSP are insufficiently understood our aim was to elucidate regulators of OSP in zebrafish, a cone-rich retina model, similar to humans. At 15 days post fertilization (dpf), we investigated if rab28 KO zebrafish display reduced peak RPE phagosomes at both morning (ZT 4) and evening (ZT 17) OSP peaks, as described in Lewis et al. We hypothesized that overexpression of Rab28 specifically in cone photoreceptors could rescue the OSP defect in rab28 KO fish, demonstrating a cell autonomous role. We undertook proteomic profiling by mass spectrometry to identify novel regulators of OSP regulators and factors dysregulated at OSP peaks by the loss of Rab28. Finally, we determined if visual cycle retinoid levels are significantly reduced in larval and adult rab28 KO eyes.

2 | METHODS

2.1 | Zebrafish strains and maintenance

Zebrafish larvae up to 131 hpf were grown in 10 cm dishes of E2 embryo medium (0.137 M NaCl, 5.4 mM...
KCl, 5.5 mM Na3HPO4, 0.44 mM KH2PO4, 1.3 mM CaCl2, 1.0 mM MgSO4, and 4.2 mM NaHCO3, conductivity ~1400–1600 µS, pH~7) maintained at 27°C in an incubator on a 14 h/10 h light-dark cycle, unless otherwise stated.30

Adult zebrafish were reared in 3 L/10 L tanks in system water and maintained at a temperature of 28°C on a 14 h/10 h light-dark cycle unless otherwise stated. Up to 28 dpf, the adolescent zebrafish were fed a specialized diet (Special Diet Services) and were steadily transferred to a main diet of brine shrimp (Artemia sp.) and pellet food.

Zebrafish strains used in this study were as follows:
- WT (Tübingen).
- rab28ede7
- rab28ede7Tg[gnat2:eGFP-rab28WT]
- rab28ede7Tg[gnat2:eGFP-rab28Q72L]
- rab28ede7Tg[gnat2:eGFP-rab28T26N]27

### 2.2 Ethics statement

All experiments with zebrafish were performed according to ethical exemptions and approvals granted by the UCD Animal Research Ethics Committee (AREC-Kennedy 20-15, 20-16) and authorizations from the Health Products Regulatory Authority (Project authorization AE18982/P062 and AE18982/P187).

### 2.3 Zebrafish behavioral assays

The optokinetic response (OKR) assay to test visual acuity (VA) and contrast sensitivity (CS) of 5 dpf larvae, was initiated by placing larvae in 9% methylcellulose in a 55-mm Petri dish. The dish was placed inside a rotating drum with a black-and-white striped pattern with 20%–100% contrast on the inside and 0.02–0.2 cycles per degree (cpd) grating spatial frequency, unless otherwise stated.31 For VA assays, cpd grating spatial frequency was varied, whereas contrast remained at 100%. For CS assays, the cpd was consistently 0.02 cpd, with alterations made to contrast within the ranges highlighted above. The drum was rotated at 18 rpm for 30 s in a clockwise and 30 s in a counterclockwise direction. Using a stereoscope, eye movements (saccades) per minute were recorded manually. All visual behavioral measurements were performed in larvae at 5 dpf, prior to 131 hpf, between 10 a.m. (ZT 2.5) and 3 p.m. (ZT 7.5).

### 2.4 Electroretinogram

Electroretinogram (ERG) recordings were performed on juvenile zebrafish eyes 2 months post fertilization (mpf) as previously described.32 In short, zebrafish were dark-adapted for at least 30 min prior to testing, followed by euthanization. The eyes were stimulated with six light pulses (100 ms) with an intensity of 500, 2300, 6000, 6000, 2300, and 500 lux. Electrical signals were captured using an electrode that was inserted within the eyeball and subsequently amplified. The recorded maximum b wave amplitudes per zebrafish were averaged per light intensity and normalized against wild type. All statistical analyses were performed using PRISM software (v9.0.0). Average scores were calculated, followed by an unpaired student’s t test.

### 2.5 Transmission electron microscopy

Zebrafish were humanely euthanized with tricaine methanesulfonate and stored in a 2% PFA and 2.5% glutaraldehyde in 0.1 M Sorenson phosphate buffer pH 7.3 fixative solution at 4°C for a minimum of 2 days. Sample processing involved post-fixation in 1% osmium tetroxide and subsequent dehydration using a 4-day gradient of ascending series of ethanol concentrations prior to Epon 812 resin for a minimum of 24 h. 80 nm sections were cut on a Leica EM UC6 microtome and mounted on copper grids and post-stained with 2% uranyl acetate and 3% lead citrate. To ensure consistency between samples for imaging and quantitative analysis, the optic nerve was used as a reference point for sectioning. Imaging was performed on an FEI Tecnai 12 electron microscope. For phagosome analysis by TEM, phagosomes were manually counted, and the density was calculated as phagosomes per micron of RPE as previously described.27

### 2.6 Retinoid analysis

For retinoid analysis carried out on adult zebrafish, two fish per strain were euthanized in tricaine (MS-222) prior to enucleation on ice under dim red light (four eyes per genotype, per replicate). Samples were snap frozen in liquid nitrogen and stored at −80°C. Analysis of larval retinoid levels were carried out on 100 larval heads per strain as previously described.33 Subsequent to euthanasia, larval heads were collected under dim red light on dry ice and snap frozen in liquid nitrogen and stored at −80°C. For normal-phase high-performance liquid chromatography (HPLC) analysis of retinoids, all retinoid extractions were carried out under dim red light in a dark room. Four zebrafish eyes (adult) or 100 heads (larval) were stored at −80°C, thawed on ice, and homogenized in 500 µl of phosphate-buffered saline (PBS) using a glass tissue grinder (Kontes). Protein concentration was...
determined using 50 μl of sample and a Micro BCA protein assay kit (Pierce). To the remaining 450 μl of homogenate, 25 μl of 5% SDS (~0.2% SDS final concentration) and 50 μl of brine were added, and samples were briefly mixed. Hydroxylamine hydrochloride (500 μl of 1.0 m solution in pH 7.0 PBS) was added to generate retinal oxime, and samples were vortexed and incubated at room temperature for 15 min. The aqueous phase was quenched and diluted using 2 ml of cold methanol. The samples were twice extracted by the addition of 2-ml aliquots of hexane, brief vortexing, and centrifugation at 3000 g for 5 min to separate the phases. Pooled hexane extracts were added to 13 × 100-mm borosilicate test tubes and evaporated to dryness under a stream of nitrogen. Dried samples were dissolved in 100 μl of hexane and analyzed by normal-phase HPLC using an Agilent 1100 series chromatograph equipped with a Supelcosil LC-Si column (4.6 × 250 mm, 5 μm) using a 0.2%–10% dioxane gradient in hexane at a flow rate of 2 ml per min. The eluted mobile phase was analyzed using a photodiode-array detector. Spectra (210–450 nm) were acquired for all eluted peaks. The identity of eluted peaks was established by comparison with spectra and elution times of known authentic retinoid standards. Retinoid amounts were quantitated by comparing their respective peak areas to calibration curves established with retinoid standards.

### 2.7 | Bisretinoid analysis

For bisretinoid analysis carried out on adult zebrafish, three fish per strain were euthanized in tricaine (MS-222) prior to eye dissection on ice under dim red light. Samples were snap frozen in liquid nitrogen and stored at −80°C. Animals had been dark adapted for 12 h prior to euthanasia. Bisretinoids were extracted from zebrafish eyes under red dim light. Briefly, six zebrafish eyes per each biological sample were washed with PBS and homogenized in 1 ml PBS. Chloroform/methanol (4 ml, 2:1, vol/vol) was added, and the samples were extracted with the addition of 4 ml of chloroform and 3 ml of dH2O, followed by centrifugation at 1000 g for 10 min. Extraction was repeated with the addition of 4 ml of chloroform. Organic phases were pooled, filtered, dried under a stream of argon, and redissolved in 100 μl of 2-propanol. Bisretinoid extracts were analyzed by normal-phase HPLC with a silica column (Zorbax-Sil 5 μm, 250 × 4.6 mm; Agilent Technologies) as previously described in Radu et al (IOVS 2008).34 The identity of each bisretinoid peak was confirmed by online spectral analysis. A2PE data were presented as means with standard deviation of four biological samples for each genotype.

### 2.8 | Statistical analysis

Statistical analysis of all data was performed using Graphpad Prism v7, unless otherwise stated. All data values shown represent the mean for the number of replicates (n) indicated in each figure legend. Error bars represent standard errors of the mean (SEM) or standard deviations (SD), as indicated. Student’s t test or an ANOVA was used to determine the statistical significance between groups, where indicated. Statistical analysis of all data for Figures 6 and 7 was performed using R, unless otherwise stated. The boxplots in Figure 6(D–F) illustrate the minimum and maximum values (ends of the whiskers), interquartile range (length of the box), and median (line through the box) for the three replicates at each time-point ZT 4, ZT 9, and ZT 17. A one-way repeated-measures ANOVA was used to determine if there was a statistically significant difference in the mean across time (i.e., for ZT 4, ZT 9, and ZT 17). Then a student’s t test was used to perform pairwise comparisons to examine if there is a statistically significant difference in the mean for the following pairs: ZT 4 relative to ZT 9; ZT 9 relative to ZT 17; and ZT 4 relative to ZT 17. Using the results from the t test Figure 6(G) shows the number of proteins that are upregulated or downregulated at a specific time-point in comparison with all other time points. Figure 6(H) shows the mean of the seven proteins that were downregulated at time-point 9 at each time-point ZT 4, ZT 9, and ZT 17; the difference in the mean and its upper and lower confidence interval for ZT 4 relative to ZT 9 and ZT 9 relative to ZT 17.

For proteomic profiling of OSP regulators as described in Figure 7, a two-way repeated-measures ANOVA was carried out to examine the effect of group type (Mutant or Sibling) over time (ZT 4, ZT 9, and ZT 17) on log-transformed LFQ intensity. There was a statistically significant interaction between the effects of group and time on LFQ intensity for the proteins listed in Figure 7(C). This implies that the trend of the log-transformed LFQ intensity over time was significantly different if the individuals were rab28 mutant versus sibling for these proteins. Some examples of proteins with significantly different trends over time are illustrated in Figure 7(E–I). Subsequently, pairwise t tests were used to determine if there was a statistically significant difference in the mean of the log-transformed LFQ intensity for rab28 mutant versus sibling at each time point. Figure 7(D) provides the list of proteins that show a significant difference in LFQ intensity for at least one time-point. All p-values were adjusted using the Bonferroni multiple testing correction method. If the p < .05, it is highlighted in the Table in Figure 7(D). Normality checks and Levene’s test were carried out, and the assumptions were met.
2.9 Proteomic analysis

Two animals were euthanized per genotype at each time point with three biological replicates per time point (four eyes per genotype, per replicate, per time point). Eyes were dissected on ice and snap frozen in liquid nitrogen with 1× protease inhibitor (Roche) in milliQ H₂O. Protein concentrations were determined using the BCA protein assay kit (ThermoFisher Scientific). Samples were stored at −80°C until protein preparation processing using PreOmics iST Sample Preparation Kit (PREOMICS) as per manual (https://www.preomics.com/resources). Samples were analyzed on a Thermo Scientific Q-Exactive mass spectrometer connected to a Dionex Ultimate 3000 (RSLCNano) chromatography system. Peptides were separated on C18 home-made column (C18RP Reposil-Pur, 100 × 0.075 mm × 3 μm) over 150 min at a flow rate of 250 nl/min with a linear gradient of increasing ACN from 1% to 27%. The mass spectrometer was operated in data-dependent mode; a high resolution (70 000) MS scan (300–1600 m/z) was performed to select the 12 most intense ions and fragmented using high-energy C-trap dissociation for MS/MS analysis.

2.10 Data processing and bioinformatics

Raw data from the Q-Exactive were processed using MaxQuant35,36 (version 1.6.4.0) incorporating the Andromeda search engine (ANDROMEDA). To identify peptides and proteins, MS/MS spectra were matched against Uniprot Danio rerio database (2018_05) containing 46 933 entries. All searches were performed using the default setting of MaxQuant, with trypsin as specified enzyme allowing two missed cleavages and a false discovery rate of 1% on the peptide and protein level. The database searches were performed with carbamidomethyl (C) as fixed modification and acetylation (protein N terminus) and oxidation (M) as variable modifications. For the generation of label-free quantitative (LFQ) ion intensities for protein profiles, signals of corresponding peptides in different nano-HPLC MS/MS runs were matched by MaxQuant in a maximum time window of 1 min. The Perseus computational platform (version 1.6.2.3) was used to process MaxQuant results. Data were log transformed. To examine changes in protein expression using volcano plots, data were log transformed, and missing values were imputed with values from a normal distribution. Data were normalized using z-score and visualized using heat maps. Pathway enrichment analysis was performed using the ClueGo (v2.3.5)39 and Cluepedia (v1.3.5) plugins40 in Cytoscape (v3.4.0)41 with the Danio rerio (7955) marker set. The KEGG functional pathway databases, consisting of 6004 genes, as well as the Gene Ontology biological processes database with 17 136 genes were used GO term levels (min = 3; max = 8) and GO term restriction (min#genes = 3, min% = 1%) were set, and terms were grouped using a kappa score threshold of 0.4. The classification was performed by the right-sided hypergeometric enrichment test, and its probability value was corrected by the Benjamini–Hochberg (adjusted % term p < .05).

3 RESULTS

3.1 Proteome profiling reveals cellular pathways regulated by Rab28

To elucidate the biological effects of Rab28 loss on the zebrafish retina, an unbiased proteomic profiling approach was undertaken, comparing eyes of a rab28 KO model to wild-type siblings. To capture changes in protein expression at peaks and troughs of OSP, biological triplicates were collected at ZT 4, ZT 9, and ZT 17 (Figure 1A). From 2 mpf eyes, a total of 4112 proteins were identified in rab28 KO and wild-type sibling samples by mass spectrometry. 498, 310, and 207 proteins were significantly altered (−log p-value ≥1.3) in rab28 KO eyes at ZT 4, ZT 9, and ZT 17, respectively. The data presented in heat maps (Figure 1B–D) show 325, 208, and 71 upregulated proteins and 173, 102, and 136 downregulated proteins at ZT 4, ZT 9, and ZT 17, respectively. Eyes were grouped using a k-score threshold of 0.4. The classification was performed by the right-sided hypergeometric enrichment test, and its probability value was corrected by the Benjamini–Hochberg (adjusted % term p < .05).
**FIGURE 1**  
*rab28* knockout zebrafish have altered proteome profiles  
(A) Time course highlighting sample collection time (red x)  
in relation to phagosome level fluctuation. (B–D) Heat maps representing  
the differentially expressed proteins between 2 months post  
fertilization wild-type siblings and *rab28*−/− at ZT 4, ZT 9 and ZT 17  
respectively, illustrating proteins significantly upregulated (in red)  
or downregulated (in blue), given a cut-off of −log *p*-value ≥1.3.  
Student’s *t* test was used for statistical analysis (*N* = 3). (E–G) Volcano  
plot representing differentially expressed proteins between wild-type  
siblings and *rab28*−/− given a fold change cut-off of ±1.2 and −log *p*-value  
≥1.3. Student’s *t* test was used for statistical analysis. (H,I) Venn diagram  
representing common and mutually exclusive proteins upregulated  
(H) and downregulated (I) between *rab28* KO and wild-type siblings at ZT 4, ZT 9, and ZT 17.
significantly, differentially expressed proteins identified phototransduction (e.g., short-wave opsins), phagocytosis (e.g., Tubb1, Tubb2, Atp6v1) and oxidative phosphorylation pathways (e.g., Ndufs6, Ndufb7) altered at ZT 4, ZT 9, and ZT 17 (Figure 2A–C).

Further interrogation of our proteomic data shows significantly reduced expression of core ciliary transport proteins, for example, Bbs1, Bbs4, and Bbs9 in rab28 KO eyes (Figure 2D). This is in agreement with our previous research which showed a link between Rab28 and the BBSome complex, an octameric protein complex necessary for protein trafficking into and out of the cilium.25 We also observed increased Pde6d levels in the rab28 KO at ZT 9, which is interesting as Pde6d is the molecular chaperone known to shuttle lipidated Rab28 into the cilium by a trafficking pathway known as lipidated protein intraflagellar targeting (LIFT).25–27

Previously, we reported that rab28 KO larvae did not show an impaired visual phenotype up to 15 dpf using the OKR assay under standard conditions.27 However, as the proteomic analysis revealed reduced levels of phototransduction and visual cycle proteins (e.g., Short-wavelength opsin 1 and 2, Rhodopsin, and Cellular retinaldehyde-binding protein 1) in 2 mpf rab28 KO eyes, we re-investigated vision using more sensitive visual behavior methods (Figure 2G,H).31 These assays probed for defects in CS or VA (Figure 2G,H). However, in rab28 KO larvae at 5 dpf, no significant differences in either the VA or CS were observed compared with control siblings. ERG data collected at 2 mpf also did not reveal a significant difference in the normalized b-wave amplitude of the rab28 KO compared with the age-matched wild-type sibling control, suggesting there is no reduction in retinal function in rab28 KO at juvenile stages (Figure 2I). The absence of an impaired visual/retinal phenotype at the larval-juvenile stages indicates that Rab28 function is compensated for by other proteins or that impaired vision is a mid-late-onset phenotype.

Our proteomic analysis of the rab28 KO model revealed a significant reduction in the levels of Abca4b (Figure 2F), a fish parologue of the gene encoding ABCA4; a member of the ATP-binding cassette transporter gene sub-family A (ABC1). ABCA4 defects are associated with recessive Stargardt’s disease.43 In retinal photoreceptors, ABCA4 actively flips N-retinylidene-phosphatidylethanolamine from the lumen to the cytoplasmic leaflet of OS membranes, thereby facilitating the removal of potentially toxic retinoid compounds from photoreceptors.44,45 Recently, ABCA4 was shown to be expressed in the RPE, aiding in recycling of free retinaldehyde released during proteolysis of opsins following OSP, thereby attenuating formation of bisretinoids in the RPE.13

### 3.2 Rab28 loss results in reduced RPE phagosomes at both zebrafish OSP peaks

In the context of OSP, we previously reported 55% lower levels of phagosomes in the RPE of rab28 KO at ZT 4,27 the dawn OSP peak.29 This concurred with similar reports for a Rab28 KO mouse model.26 However, different species display one, two, or potentially even three OSP peaks.29,46,47 The zebrafish retina has two OSP peaks; a dawn peak at ZT 4 and a dusk peak at ZT 17, comprising of phagocytosed cone and rod photoreceptor OS tips at each time-point.29 From the proteomic data, KEGG pathway analysis revealed the phagosome pathway to be altered in the rab28 KO at ZT 4 (Figure 2A). Although individual phagocytosis-associated proteins were altered at ZT 4, ZT 9, and ZT 17, the term phagocytosis did not reappear in the KEGG pathway analysis at time points other than ZT 4. Thus, to directly assess the integrity of OSP in the rab28 KO throughout the day, samples were collected at discrete time points in accordance with the OSP flux reported.29 As in Carter et al., TEM was performed on the retinae of 15 dpf rab28 KO and siblings but extended to ZT 0, ZT 4, ZT 12, and ZT 17 and the number of RPE phagosomes counted. As expected, the phagosome number oscillated between time points (Figure 3B). Importantly, the TEM analysis ascertained that loss of Rab28 also disrupts the dusk OSP peak reducing the peak number of overall RPE phagosomes by ~50% compared with the sibling controls. TEM of additional time points confirmed the rab28 KO line does not display phase-shifted peaks at ZT 0 or ZT 12 (Figure 3B–N). Notably, there is a basal level of RPE phagosomes (46% of the peak phagosome number, when comparing ZT 4 vs. ZT 0) observed at each time-point in both wild-type siblings and rab28 KO (Figure 3B). This reveals that Rab28 is required for the burst of RPE phagosomes at dawn and dusk peaks, but not for basal levels of OSP.

### 3.3 Reduced RPE phagosomes caused by loss of Rab28 is rescued using cone-specific eGFP:Rab28

In the retina, Rab28 is expressed in the RPE, and rod and cone photoreceptors.26,48 We hypothesized that the reduced RPE phagosomes at OSP peaks in rab28 KO zebrafish resulted from the absence of Rab28 in cones. Thus, a Tol2 Rab28-eGFP transgene, expressing EGFP-tagged zebrafish Rab28, regulated by a cone-specific gnat2 promoter,49 was crossed into the rab28 KO line. TEM sections were collected from transgenic animals at the dawn OSP peak, and the number of RPE phagosomes quantified (Figure 4). This revealed that gene replacement to
selectively express wild-type zebrafish Rab28 in cone photoreceptors is sufficient to restore the dawn OSP peak in rab28 KOs to wild-type levels (Figure 4B,E), suggesting zebrafish Rab28 plays a more important role in cone photoreceptors compared with other retinal cells.

GTPase switching between the GTP, or GDP-bound conformations of Rabs is the major mechanism by which most Rabs are thought to be activated/inactivated.50 We previously reported that both GTP and GDP-binding variants of zebrafish Rab28 localize to the photoreceptor

![KEGG pathway analysis of rab28 KO vs wildtype sibling at ZT 4](image)

![KEGG pathway analysis of rab28 KO vs wildtype sibling at ZT 9](image)

![KEGG pathway analysis of rab28 KO vs wildtype sibling at ZT 17](image)

| UNIPROT | GENE | Log Student's T-test p-value | Student's T-test Difference | Zeitgeber Time |
|---------|------|-----------------------------|-----------------------------|---------------|
| B3DFT2  | bbs1 | 3.61                        | 1.59                        | 17            |
| F18BR9  | bbs4 | 1.37                        | 1.85                        | 4             |
| E7F6E0  | bbs9 | 2.87                        | 0.29                        | 4             |
| F10Z5A  | pde6d| 1.50                        | -0.32                       | 9             |

rab28 KO showed altered expression of core ciliary transport proteins

| UNIPROT | GENE | Log Student's T-test p-value | Student's T-test Difference | Zeitgeber Time |
|---------|------|-----------------------------|-----------------------------|---------------|
| F1QVE1  | abcd3| 2.38                        | 0.5                         | 4             |
| F1QVD3  | rbgla| 2.52                        | 0.30                        | 9             |

rab28 KO showed reduced levels of visual cycle proteins

![Contrast sensitivity assay of 5 dpf rab28 KO](image)

![Visual Acuity of rab28 KO at 5 dpf](image)

![ERG responses of 2 mpf rab28 KO](image)
OS, albeit to varying degrees.\textsuperscript{27} While Rab28 localizes to the cone OS independent of nucleotide binding, it is unknown if the role of Rab28 in OSP is influenced by nucleotide binding. This question was investigated with three eGFP-Rab28 variants expressed in zebrafish cones in the \textit{rab28} KO background. Surprisingly, both GTP-preferring and GDP-preferring transgenic models restored normal peak levels of RPE phagosomes in the knockout background (Figure 4B,E–G). Notably, the Rab28\textsuperscript{T26N} variant was recently reported as pathogenic in CRD patients.\textsuperscript{19} A slight reduction in RPE phagosomes was observed in the eGFP-Rab28\textsuperscript{T26N} retinas, however, this was not statistically significant (Figure 4B,F). We also observed an increase in the amount of RPE phagosomes with eGFP-Rab28\textsuperscript{Y72L} in comparison with the wild-type variant, albeit again this difference did not show statistical significance (Figure 4B,G).

3.4 | Investigation into light regulation of zebrafish OSP

To inform future studies of OSP regulation in zebrafish, the issue of whether the peaks show a circadian regulation, as in mice and rats,\textsuperscript{46,51,52} or a light-dark transition regulation as in frogs\textsuperscript{47} needed to be resolved. Thus, the effect of a dark-dark light cycle on RPE phagosomes was quantified by TEM at ZT 4, ZT 9, and ZT 17 (Figure 5). With dark:dark environmental conditions, the number of RPE phagosomes at circadian time 4 (CT4) (0.12 phagosomes/\(\mu\)m RPE) and CT 17 (0.10 phagosomes/\(\mu\)m RPE) was significantly (\(p = .005, p \leq .0001\) respectively) reduced (Figure 5F,H,I,J) compared with both peak numbers (0.18 and 0.25 phagosomes/\(\mu\)m RPE, respectively) at ZT4 and ZT 17, under a normal light dark cycle (Figure 5B,D,I,J) suggesting an absence of circadian regulation. Indeed, at both CT 4 and CT 17, in the absence of light, the number of RPE phagosomes was equivalent to the basal number (0.14 phagosomes/\(\mu\)m RPE) present at OSP trough time ZT 9 (Figure 5C,I,J). Notably, RPE phagosomes were not eliminated in the dark:dark samples but maintained at a level equivalent to the basal level at an OSP trough in light-dark reared samples (Figure 5I,J). In summary, exposure to light:dark transition was necessary to fully initiate the burst of OSP at the expected dawn and dusk peak time in the zebrafish retina.

3.5 | Proteomic analysis of wild-type zebrafish retinae reveals potential OSP regulators

Significant progress has been made in identifying factors that mediate phagocytosis of rod OS in the RPE.\textsuperscript{53–55} However, our understanding of the factors (e.g., in photoreceptors) that initiate OSP is limited, as is whether OSP is regulated by the same factors at dawn and dusk peaks. To probe these research questions, proteins with significant changes in expression between peaks and troughs of OSP were identified from proteomic profiling of wild-type zebrafish eyes collected at ZT 4, ZT 9 and ZT 17 (Figure 6A,B). Cross comparisons of all three time points identified 676 proteins with significant changes between peak-to-trough or peak-to-peak (Figure 6B). Gene ontology analysis identified the cellular processes of intracellular transport and cellular protein localization (e.g., Rab2a, Rab3ab, and Dkc10) as differentially expressed at the dawn and dusk OSP peaks compared with the OSP trough (Figure 6C). To assess the quality of this novel list of differentially expressed proteins during OSP, we examined the profile of canonical OSP proteins (Figure 6D–F). Encouragingly, Integrin alpha-V (Itgav) and ATP binding cassette subfamily F member 1 (Abcf1) peaked at the dusk OSP peak ZT 17. Lysosomal-associated membrane protein 1b (Lamp1b) stayed constant throughout the day, despite high variability between biological replicates at ZT 17 (Figure 6E). Using a more stringent analysis of changes in the pattern of expression, 52 proteins were significantly differentially expressed at the dawn and/or dusk OSP peaks compared with the OSP trough (Figure 6G,H, Table S1). Two and sixteen proteins were uniquely upregulated at ZT 4 and ZT 17, respectively. Twelve and fifteen proteins...
were uniquely downregulated at ZT 4 and ZT 17, respectively. No proteins were significantly downregulated at both OSP peaks compared with the OSP trough. Notably, seven proteins were significantly upregulated at both the dawn and dusk OSP peaks compared with the OSP trough (ZT 4 and ZT 17 vs. ZT 9) (Figure 6G,H). These proteins
include Jacalin 8, Sorting nexin 6, Dyskerin, Calpain 2, N-myc downstream-regulated 1b, Exportin 1 and an unannotated 6.8 kDa mitochondrial proteolipid like protein (Figure 6H). Details on proteins significantly altered at solely one OSP peak are listed in Table S1. Most (7 of 9) of the proteins upregulated at ZT 4 are also upregulated at ZT 17, whereas only 7 of the 23 proteins upregulated at ZT 17 are also upregulated at ZT 4. There is no overlap between the proteins downregulated at ZT 4 and ZT 17.

Further investigation is required to elucidate the precise roles, if any, of these proteins in OSP regulation, and at which stage of the OSP process they are involved; demarcation, recognition, binding, engulfment, or degradation. In summary, our analysis has uncovered novel proteins potentially involved in regulating OSP in vivo. In particular, as the zebrafish is a cone-dominant model, some of these candidate proteins may be specifically important for cone photoreceptor OSP.
Zebrafish outer segment phagocytosis (OSP) is regulated by external light sources. (A,E) Time course highlighting sample collection time (red x), and light-dark cycle. Samples were collected at ZT 4, ZT 9, and ZT 17 from 16 days post fertilization (dpf) wild-type zebrafish retinae. At each time point, one cohort had undergone a regular 14:10 light:dark cycle. The other had been raised in dark from ZT 10 the day prior to collection and were euthanized under red light. (B–D, G,H) Representative TEM images of wild-type retinae at 16 dpf, with phagosomes indicated (pink arrows). Scale bar; 5 μm. (I) Scatter plots of phagosome/μm of retinal pigment epithelium (RPE) in dark-reared and control retina at ZT 4 and ZT 17 (OSP peaks) and ZT 9 (OSP trough). Fish raised in the dark for have ~35%–55% reductions in the number of RPE phagosomes containing engulfed OS material in comparison with controls at ZT 4 and ZT 17 respectively. Statistical analysis carried out by student’s t test between individual groups. RPE of >570 μm was measured per animal. N = 3 animals (6 retinae) per group per time point. (J) Trend line demonstrating the level of RPE phagosomes present in the dark-raised wild-type eyes (black) compared with controls exposed to the regular light-dark cycle (orange) at defined time points throughout the day. A two-way ANOVA, followed by Turkey’s multiple comparison testing revealed no significant difference between CT 4 CT 9 or CT 17 (black), but a significant difference between ZT 4 versus ZT 9 (p = .0229), ZT 9 versus ZT 17 (p ≤ .0001), and ZT 4 versus ZT 17 (p = .003) (orange)
FIGURE 6 Investigation into internal regulators of zebrafish outer segment phagocytosis (OSP). (A) Time course highlighting sample collection time (red x) in relation to phagosome-level fluctuation. (B) Heat map representing differentially expressed proteins between 2 months post fertilization wild-type siblings at ZT 4, ZT 9, and ZT 17, respectively, illustrating proteins significantly upregulated (in red) or downregulated (in green), given a cut-off of $-\log p$-value $\geq 1.3$. Student’s t-test was used for statistical analysis. (C) Gene ontology analysis of significantly downregulated and upregulated proteins at OSP peaks (ZT 4 and ZT 17 grouped) compared with an OSP trough (ZT 9). (D–F) Box and whisker plots showing the normalized expression profile of known OSP-associated proteins Itgav, Lamp1b, and Abcf1 at ZT 4, ZT 9, and ZT 17. (G) Table containing the number of proteins that are specifically upregulated or downregulated at ZT 4, ZT 17, or ZT 9. (H) Table documenting the mean of the seven proteins specifically downregulated at the OSP trough (ZT 9) compared with OSP peaks (ZT 4 and ZT-17); the difference in the mean and the upper and lower confidence intervals for ZT 4 relative to ZT 9 or ZT 9 relative to ZT 17.
Identification of potential OSP regulators with modified expression profiles in rab28 KO retinae

Given the list of 52 proteins identified as significantly altered at one/both OSP peaks, we took the opportunity to discriminate how Rab28 could regulate OSP, by determining which of these 52 proteins display significantly altered expression patterns in the rab28 KO model (Figure 7A,B). For a substantial cohort of proteins, two-way ANOVA revealed the linear trend of the log-transformed LFQ intensity over time was significantly

\[ F = \frac{\text{sum of squares between groups}}{\text{degrees of freedom between groups}} \]

\[ p = \text{probability of observing the data if the null hypothesis is true} \]
different between rab28 mutant versus sibling. Of the original 52 candidate OSP proteins, 21 showed a significant profile difference in the rab28 KO eyes compared with sibling controls (Figure 7C). Several were previously associated with OSP, or general phagocytic processes including Rab7a and Tuba1a. A deeper analysis was undertaken to determine if there was a statistically significant difference in the mean of the log-transformed LFQ intensity for rab28 mutant versus sibling at distinct time points (Figure 7D). In line with Rab28’s role in OSP, the 21 candidates were predominantly proteins showing significantly altered levels at a single OSP peak time point, either ZT 4 or ZT 17. The profiles of Acat2, Septin 3, Lsm4, Calr3b, Crub2b, Ncaldb, and Camk2b1 showed a significant simple main effect alteration at ZT 4 (e.g., Figure 7E). In contrast, Calb2a, Nplc4, Gpr37l1b, Rab7a, Cryba1l1, and Tuba1a showed a significant simple main effect alteration in the rab28 KO at ZT 17 (e.g., Figure 7F). Only one protein profile, Dnml11, was significantly different at both OSP peaks in the rab28 KO compared with wild-type controls (Figure 7H).

Additionally, Rpe2 (p = .016) and Polr2c (p = .04) displayed a significant difference in the mean log-transformed LFQ intensity for rab28 mutant versus sibling, but there is no significant association with time (ZT 4, ZT 9, and ZT 17; see Figure 7I for an illustration).

3.7 rab28 gene knockout leads to age-dependent changes of visual cycle retinoid and bisretinoid levels

Defective phagocytosis of photoreceptor OS by the RPE is linked to diminished health and function of the eye in some inherited retinal dystrophies and in age-related blindness. Therefore, we investigated if there were functional consequences in retinal homeostasis due to the loss of OSP peaks in rab28 KOs. The light-sensitive chromophore 11-cis-retinaldehyde (11cRAL) is needed for phototransduction and is recycled from its all-trans-retinaldehyde (atRAL) photoisomer via the RPE visual cycle. OSP by the RPE generates a pool of retinaldehyde to shuttle either into the visual cycle as atRAL or being directly recycled as 11cRAL to photoreceptors.

Hence, we hypothesized that Rab28 loss would affect levels of visual cycle retinoids and bisretinoids. Eyes were collected under dark-adapted conditions from rab28+/+ and rab28 KO eyes at larval (15 dpf) and adult (8 mpf) stages, and the retinoid profile was evaluated by HPLC.

At 15 dpf, a significant reduction in the rab28 KO levels of retinyl esters; 11-cis-retinyl palmitate (11cRP) (p = .0127) and all-trans-retinyl palmitate (atRP) (p = .0306) was observed (Figure 8A). At 8 mpf, a significant reduction in the visual chromophore 11cRAL (p = .0110), atRP (p = .0313), and total cumulative retinoids was observed (Figure 8B). From these data, we infer a direct or indirect role of Rab28 in the visual cycle, with the reduction in 11cRAL potentially leading to a visual deficit at a later time point. At the 8 mpf time point, we noticed a trend towards increasing levels of atRAL, a precursor of bisretinoid biogenesis and a marked reduction in total retinoid content.

Thus, we investigate whether rab28 KO zebrafish show changes in vitamin A dimerization processes which materialize as accumulation of bisretinoid-lipofuscin in RPE cells. Bisretinoids are a family of photoreactive vitamin A aldehyde-derived adducts that form nonenzymatically as a by-product of the visual cycle and implicated in disease processes that threaten vision and retinal health. We quantified bisretinoids by normal-phase HPLC in dark-adapted rab28+/+ and rab28 KO eyes at 7.5 and 12 mpf (Figure 8C). Levels of A2E, a major bisretinoid found in the lipofuscin granules, were similar in the zebrafish eye irrespective of
genotype and age (data not shown). However, the levels of the A2PE bisretinoid, an immediate precursor of A2E, were significantly increased in eyes of 12 mpf rab28 KO compared with wild-type zebrafish eyes (Figure 8C). There was no difference in A2PE levels at 7.5 mpf, suggesting the rate of bisretinoid by-product formation is augmented by age.

**DISCUSSION**

Since linkage of human RAB28 mutations to inherited blindness and cone-rod dystrophy, studies have focused on Rab28 function in the retina. However, there remains a vast gap in understanding of the protein networks regulated by Rab28. Here, we endeavored to bridge that void,
to shed more light on Rab28 function and Rab28-CRD disease mechanisms. This study uncovered that Rab28 is required from the larval zebrafish stage for the burst of RPE phagocytosis, of the photoreceptor outer segments at both dawn and dusk peaks. We further investigated the regulation of OSP in the zebrafish model and the role of Rab28. Using a gene complementation approach, we uncovered that Rab28 plays an important role in the cone photoreceptors, as cone-specific expression of Rab28 was sufficient to restore the OSP defect in our knockout model. Proteomic analysis by mass spectrometry demonstrates loss of Rab28 effects the expression profile of several known regulators of OSP and identified potential novel regulators of the OSP process. We determined that rab28 KO can result in long-term physiological effects on the retina, as retinoid and bisretinoid analysis on the rab28 KO revealed altered levels of visual cycle components, as well as an increase in the toxic vitamin A compound A2PE.

Our proteomic analysis demonstrates that when Rab28 expression is ablated at adult stages, expression levels of several protein families are significantly reduced, including those of ciliary transport proteins, the visual cycle and phototransduction (Figure 2). In agreement, previous studies in ciliated C. elegans cells show that prenylated Rab28 is an IFT cargo, associating with the molecular transport trains in a GTP-dependent manner.24 Furthermore, Rab28 was dependent on the BBSome for localization to the ciliary membrane, and on the molecular chaperone Pde6d for transport into the cilium.24,25 The interaction of Rab28 and Pde6d has been confirmed in mouse and zebrafish retinas.26,27 Indeed, Ying et al26 suggested that visual impairment in Pde6d mouse knockouts relates to Rab28’s absence from photoreceptor OSs. The rodent study saw no changes in the expression levels of Pde6d in the Rab28 KO’s. Here, we have observed a small, but significant increase in the levels of Pde6d in the rab28 KO model at 2 mpf at ZT 9, which could potentially be an attempt at compensation for reduced Rab28 levels.

In relation to ciliary transport, we also observed a significant reduction in the levels of BBSome proteins Bbs1, Bbs4, and Bbs9 at ZT 4 or ZT 17, in zebrafish rab28 KO eyes (Figure 2D). This suggests that Rab28 loss has a consequential effect on the BBSome, a finding not previously reported at the protein level. Previous work shows that the C. elegans orthologue of BBS-8 is required for ciliary localization of Rab28.25 Based on available evidence, Rab28 is postulated not to be a member of the core BBSome complex but rather associated with the BBSome to undergo IFT within the cilium.24 As such, further investigation in needed into how loss of an IFT and LIFT cargo could reduce protein levels of trafficking machinery.

At 2 months of age, many components of cone and rod phototransduction were significantly reduced in Rab28 KO eyes which could lead to impaired vision. Previously, Carter et al27 used the standard OKR and visual motor response assays to assess the visual behavior of the rab28 KO model up to 21 dpf, which showed no reduction in visual function compared with siblings. Thus, we adapted our OKR assay, allowing investigation of more subtle defects in VA or CS.31 However, in 5 dpf rab28 KO larvae, no deficit was observed in VA or CS (Figure 2H—I). We conclude that rab28 KO larvae do not have an early-onset vision impairment and that assays in adult rab28 KOs are most likely to uncover impaired vision consistent with reduced phototransduction proteins at 2 months.

As mouse and zebrafish Rab28 KOs display diminished dawn OSP peaks, we expected proteomic analysis to reveal significant changes in phagocytosis related proteins. However, from KEGG pathway analysis the phagocytosis pathway as a whole was only downregulated at ZT 4, but with individual OSP associated proteins significantly altered at the latter time points (Figure 2A–C). To independently expand on the findings from Carter et al27, we quantified the number of RPE phagosomes present at consecutive time points in 15 dpf rab28 KO retinæ. This revealed that OSP peaks were not time shifted in the rab28 KO and confirmed that Rab28 is required for the peak levels of OSP at dawn and dusk. Strikingly, this broader temporal analysis of OSP peaks and troughs, revealed significant basal levels of RPE phagosomes throughout the day. Notably, the basal levels of OSP were not altered in rab28 KO, suggesting that Rab28 is required for the burst of OSP that occurs at dawn and dusk peaks (Figure 3). Similar results are observed in the avb5 and anxa5 mouse knockout models, wherein absence of either protein causes a complete lack of the characteristic burst of phagocytosis.8,59 In these models, there is a higher basal level of phagocytosis throughout the day which may compensate for the reduced peaks.8,59 This may happen in rab28 KO models and could explain why rapid retinal degeneration is not observed. Likewise, the avb5 KO shows no defect in visual behavior until beyond 4 months and in the RPE of older avb5 mice accumulation of autofluorescent lipids occurs.8 Thus, chronic defects in OSP may be needed before visual impairment presents.

Here, we demonstrate that Rab28 expressed in cone photoreceptors is sufficient to rescue the defect in peak OSP observed in rab28 KO. This indicates that Rab28 in cone photoreceptors has a cell-autonomous role in regulating OSP peaks. Previous studies suggest that OSP requires a synergistic relationship between photoreceptors and the RPE, but that RPE proteins were the key regulators for efficient OS tip engulfment and removal.6,11 However, our data suggest that Rab28 expressed in the cone photoreceptors is necessary to enable peaks of OSP. Similarly, genetic
mutants of kif17 show reduced levels of RPE phagosomes throughout the day. Kif17 shows rhythmic expression, localizing to the cone OS tip at both OSP peaks, suggesting that cone expressed Kif17 also regulates OSP. Clearly, OSP is not merely a passive process for photoreceptors, but one which requires an interplay between RPE- and photoreceptor-specific regulators.

When wild-type eGFP-tagged Rab28 was returned solely to cone photoreceptors, this was sufficient to restore peak RPE phagosomes, to wild-type sibling levels (Figure 4). To ask if this effect was dependent on Rab nucleotide binding, we expressed dominant negative and dominant active transgene variants in the analyzed may be too early to see a phenotype, as RAB28-CRD is a progressive inherited retinal dystrophy. From a therapeutic perspective, rescue of the peak OSP defect by cone-specific expression of Rab28, suggests that gene therapy for RAB28-associated CRD may be successful by Rab28 gene restoration to cones.

Having confirmed a role for cone photoreceptor Rab28 in regulating dawn and dusk peaks of OSP, we decided to investigate the fundamental mechanisms regulating peaks of OSP in vivo. This also holds biomedical significance as defective OSP may be part of the disease pathway in Rab28-linked CRD and in other retinal and macular degenerations. Environmental cues are major regulators of OSP in other species with light-dark transitions regulating OSP peaks in frog and Arvicandus ansorgi, whereas internal circadian rhythms can drive OSP peaks in rat and mouse models. In zebrafish, we observed a significant decrease in the number of RPE phagosomes present in the retinae of animals reared under a dark-dark environment for <24 h (Figure 5), suggesting that the burst of OSP in zebrafish is not under circadian regulation.

We then undertook to identify novel molecular regulators of OSP, involved in all stages of the OSP process. Unbiased proteomic analysis allowed us identify 676 proteins differentially expressed in wild-type sibling eyes across the three time points analyzed, ZT 4, ZT 9, and ZT 17 including Itgav, Abcf1, and Lamp1b, proteins previously linked to OSP regulation. Analyzing protein expression profiles upregulated/ down regulated at ZT 4 and ZT 17 (OSP peaks) compared with ZT 9 (OSP trough) identified 52 proteins following multiple comparison testing (Figure 6). Only seven proteins are upregulated at both OSP peaks, suggesting that different regulators are involved in dawn and dusk peaks. Importantly, this list included proteins with known involvement in OSP including Rab7a, Tuba1a, and Rp2. KEGG pathway analysis uncovered processes expected to be affected during OSP including, protein localization, vesicle docking, organelle organization, and intracellular transport. Notably, several proteins on the list are not annotated, for example, Uniprot entries B3DFK4, A0A0R4ITV0, B8A688. Therefore, this list of proteins differentially expressed during OSP provides several avenues to elucidate novel regulators of OSP. One interesting protein upregulated solely at the ZT 4 peak is the lens protein, Cryba1/βA3/ A1-crystallin. This protein is expressed in the RPE, where it is involved in acidification of lysosomes by vacuolar ATPase modulation. A significant interaction between Rab28 and Cryba1/βA3/ A1-crystallin was previously reported, including in the T26N, or GDP-preferring variant. Using this list of proteins upregulated at one or both peaks of OSP in zebrafish, we returned to elucidating how Rab28 controls OSP by identifying which of them lose differential expression in Rab28 KO eyes. (Figure 7). Of the 52 candidate proteins, 21 showed a statistically significant relationship between group (rab28 mutant vs. sibling) and time (ZT 4, ZT 9, and ZT 17) (Figure 7C). For an additional two proteins the difference in the group (rab28 mutant vs. sibling) is associated with significantly different LFQ intensity only.

Interestingly, only Park7 was dysregulated at both peak times (ZT 4 and ZT 17) in the rab28 KO (Figure 7C,D,H). Park7, or DJ-1, is a redox-sensitive molecular chaperone linked with Parkinson’s disease and also involved in regulating the RPE’s response to oxidative stress. No previous studies have postulated a role for Park7 in OSP, but it is well known that phagocytosis itself increases oxidative stress leading to increase levels of reactive oxygen species. Oxidative stress is also inducible as a result of imbalance between the OS renewal and the phagocytic activity which scavenge the OS tips, which may account for an altered expression pattern in our rab28 KO model. Notably, we also saw distinct differences in the profiles of Rab7a and Rp2, two proteins previously associated with OSP (Figure 7). Rab7a is known to be involved in phagosome maturation in the
RPE, and we hypothesize that lower levels of Rab7a may be due to reduced levels of phagosomes produced in the KO retina.\textsuperscript{63} Rp2 is also of interest as human mutations lead to CRD.\textsuperscript{64} Knockout of Rp2 leads to defects in OS renewal, and it has been postulated that this could be in part due to a defect in OSP.\textsuperscript{60}

Diminished levels of peak OSP, as mentioned earlier, may contribute to later-onset impaired vision. Therefore, to determine if the loss of OSP peaks in rab28 KOs has earlier functional consequences in the retina, we investigated defects in the visual cycle. Several studies link OSP to the visual cycle, as engulfment of OS tips by the RPE is another method of retinoid recycling and distinct to the canonical and non-canonical pathways that shuttle retinoids from photoreceptors to the RPE or Muller glia.\textsuperscript{12,13}

From our proteomics analysis of 2 mpf eyes, the retinyl flippase Abca4b was downregulated in the rab28 KO at both ZT 4 and ZT 9 (Figure 2F). Investigation of retinoid levels unveiled reduced retinyl esters atRP and 11cRP from 15 days old in rab28 KO (Figure 8A). Adult rab28 KO eyes show a progressive defect in the visual cycle, as levels of the visual chromophore 11cRAL become significantly reduced from 15 dpf to 8 mpf (Figure 8B). Bisretinoid analysis revealed an age-dependent increase in A2PE levels at 12 months old, confirming defective retinoid cycling in rab28 KO (Figure 8C). A2PE is a by-product of the visual cycle, resulting from two molecules of all-trans-retinal and phosphatidylethanolamine. A2PE can be formed in both photoreceptors and RPE cells, unlike A2E a bisretinoid that results after hydrolysis of A2PE in the acidic environment of the RPE lysosomal compartment only.\textsuperscript{16} We hypothesize that A2PE is elevated within photoreceptor OS due to the defective OSP phenotype and lower levels of Abca4b in rab28 KO. Accumulation and formation of A2PE may contribute to photoreceptor cell dysfunction.\textsuperscript{65}

In summary, this study further highlighted the role of Rab28 in OSP, showing its requirement for both the dawn and dusk peak of OSP. Rab28 dysregulation impacts vitamin A dynamics in the eye, by perturbing the recycling of essential retinoids and favoring formation and accumulation of toxic byproducts. Our investigation has also uncovered potential novel regulators of OSP, providing a resource for future studies of OSP, and in particular cone photoreceptor OSP.

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DISCLOSURES

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

AUTHOR CONTRIBUTIONS

Ailis L. Moran, Stephen P. Carter, and Breandán N. Kennedy: conceptualization; Ailis L. Moran: data curation; Ailis L. Moran and Breandán N. Kennedy: formal analysis; Ailis L. Moran, Joanna J. Kaylor, Zhichun Jiang, Sanne Broekman, Eugene T. Dillon, Alicia Gómez Sánchez, Michelle Carey, and Breandán N. Kennedy: validation. Ailis L. Moran, Joanna J. Kaylor, Zhichun Jiang, Sanne Broekman, and Alicia Gómez Sánchez: investigation. Ailís L. Moran and Breandán N. Kennedy: methodology; Ailís L. Moran and Breandán N. Kennedy: writing—original draft. Breandán N. Kennedy: resources; Breandán N. Kennedy: supervision; Breandán N. Kennedy: project administration; Ailís L. Moran collected all samples for proteomics, TEM, phagosome analysis, retinoid analysis, bisretinoid analysis and generated the transgenic models in the knockout background. Alicia Gómez Sánchez performed visual function analysis. Sanne Broekman performed ERG analysis. Stephen P. Carter created the original genetic and transgenic models. Joanna J. Kaylor performed retinoid analysis. Zhichun Jiang performed bisretinoid analysis. Eugene T. Dillon performed mass spectrometry analysis. Michelle Carey and Sajal K. Minhas performed statistical analysis of proteomics. Ailís L. Moran, Erwin van Wijk, Roxana A. Radu, Oliver E. Blacque, Gabriel H. Travis, and Breandán N. Kennedy interpreted the results and provided significant intellectual input.

DATA AVAILABILITY STATEMENT

Raw data supporting the findings of this study are available from the corresponding author on request.

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**SUPPORTING INFORMATION**

Additional supporting information may be found in the online version of the article at the publisher’s website.

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