

RESEARCH REPORT

Retinal ganglion cell survival after severe optic nerve injury is modulated by crosstalk between Jak/Stat signaling and innate immune responses in the zebrafish retina

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ABSTRACT

Visual information is transmitted from the eye to the brain along the optic nerve, a structure composed of retinal ganglion cell (RGC) axons. The optic nerve is highly vulnerable to damage in neurodegenerative diseases, such as glaucoma, and there are currently no FDA-approved drugs or therapies to protect RGCs from death. Zebrafish possess remarkable neuroprotective and regenerative abilities. Here, utilizing an optic nerve transection (ONT) injury and an RNA-seq-based approach, we identify genes and pathways active in RGCs that may modulate their survival. Through pharmacological perturbation, we demonstrate that Jak/Stat pathway activity is required for RGC survival after ONT. Furthermore, we show that immune responses directly contribute to RGC death after ONT; macrophages/microglia are recruited to the retina and blocking neuroinflammation or depleting these cells after ONT rescues survival of RGCs. Taken together, these data support a model in which crosstalk between macrophages/microglia and RGCs, mediated by Jak/Stat pathway activity, regulates RGC survival after optic nerve injury.

KEY WORDS: Jak/Stat, RGC, Innate immune system, Neuroprotection, Zebrafish

INTRODUCTION

Visual information is transmitted from the eye to the brain along the optic nerve (ON), a structure composed of retinal ganglion cell (RGC) axons. The ON is highly vulnerable to damage and is compromised after acute injury and in neurodegenerative diseases, such as glaucoma. In glaucoma, RGC axons are the initial site of injury; this causes the RGCs to die and ultimately results in irreversible loss of visual function. Neuroprotective strategies for glaucoma treatment seek to maintain the health of RGCs even after axons have been damaged, or to prevent initial damage to the RGC axon itself (Almasieh et al., 2012; Chang and Goldberg, 2012). There has been substantial progress in identifying the molecular and cellular events that lead to RGC death in the glaucomatous eye

(Almasieh et al., 2012; Chang and Goldberg, 2012; Syc-Mazurek and Libby, 2019); however, no FDA-approved therapies currently exist to protect RGCs from death. This highlights the need for new neuroprotective strategies that preserve RGCs during glaucoma or after acute ocular trauma.

Most RGCs die in mammals suffering from glaucoma or acute ocular trauma. For example, in mouse, ~65% of RGCs are lost within 7 days of optic nerve injury (ONI), and >90% by 28 days (Li et al., 2020). Mammals are also unable to regenerate RGCs after ONI, leading to irreparable vision loss. Unlike mammals, zebrafish possess remarkable neuroprotective and regenerative capacity in the central nervous system (Cigliola et al., 2020; Lahne et al., 2020). When the ON is damaged by crush or transection, zebrafish mount a robust regenerative response and regenerate RGC axons, restoring visual connections and function (Diekmann et al., 2015). Moreover, it has been reported that ~75% of zebrafish RGCs are protected from death after ONI, even up to 7-weeks post-injury (Zou et al., 2013), but the mechanisms underlying neuroprotection are unknown. With an interest in developing novel strategies to preserve RGCs during glaucoma and other trauma, we identify potential neuroprotective factors/pathways in zebrafish that mediate RGC survival after ONI.

RESULTS AND DISCUSSION

Zebrafish retain the majority of RGCs after ONT

To enable isolation of RGCs after injury, we first verified that GFP remained expressed in RGCs of adult *isl2b*:GFP fish (Pittman et al., 2008). In retinal cryosections, *isl2b*:GFP cells were detected in the ganglion cell layer (GCL), where they co-labeled with DAPI and the RGC-specific marker zn-8 (Trevarrow et al., 1990); 64.86±8.44% (mean±s.d.) of cells within the GCL were *isl2b*:GFP⁺ (Fig. 1A).

To create an ON injury, we performed ON transection (ONT). Injured retinæ from the left eye (ONT+) and sham surgery retinæ from the right eye (ONT-) were collected at 1, 3, 7 or 14 days post-injury (dpi) (Fig. 1B). We confirmed that there were no differences in cell number between the uninjured left and right eye (left eye: 9844.44±1214.04 cells/mm²; right eye: 10,118.44±1688.71 cells/mm²; *n*=6; *P*=0.6735, Mann-Whitney test). To quantify RGC survival after ONT, we counted *isl2b*:GFP⁺ RGCs from eight regions (four in the peripheral retina and four in the central retina; Fig. 1B), then divided counts from the ONT+ retina by those from the ONT- retina of the same fish. RGC survival was 94.44±5.45% at 1 dpi and 92.52±3.54% at 3 dpi; however, survival decreased to 76.35±2.58% at 7 dpi (*P*=0.0009). At 14 dpi, RGC numbers recovered to 98.71±4.82% (Fig. 1C,D).

isl2b:GFP intensity in ONT+ RGCs declined by 40.43±9.99% at 7 dpi relative to ONT- RGCs (*P*<0.0001), raising the possibility that GFP intensity was falling below the detection threshold after

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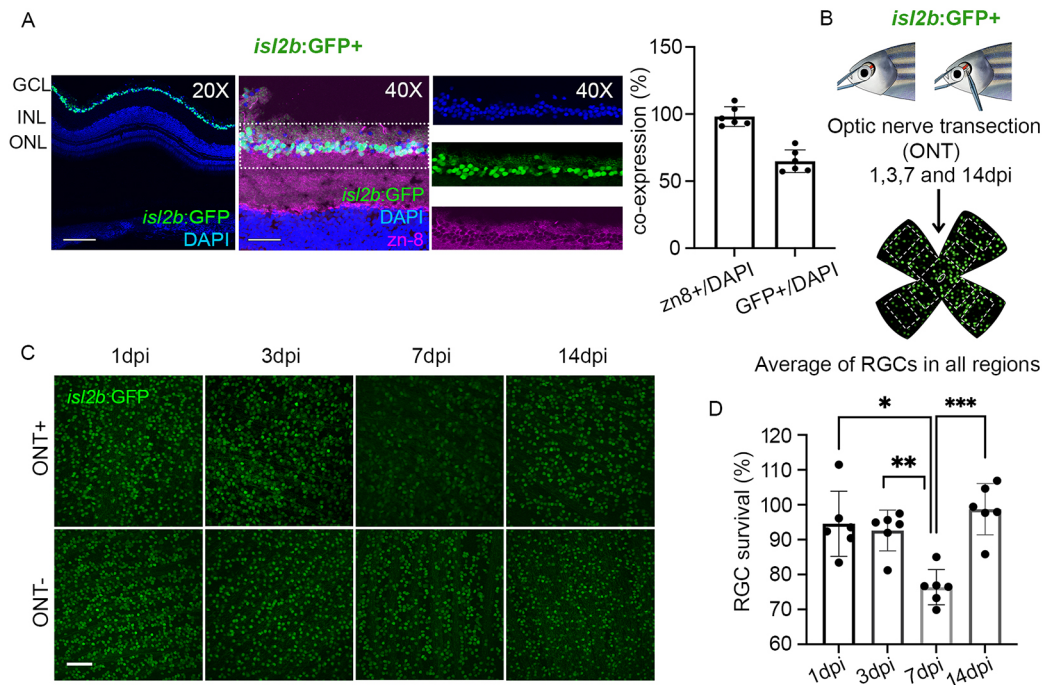


Fig. 1. Zebrafish RGCs are preserved after optic nerve transection. (A) Immunolabeling of RGCs in the GCL with zn-8 (magenta) in the adult *is/2b:GFP+* (green) retinae. Shown are a retinal cross section (20 \times) and high magnification view of the GCL (40 \times). Quantification revealed that ~65% of DAPI (blue)-stained RGCs were *is/2b:GFP+* ($n=xxx$). Scale bars: 100 μm (20 \times); 50 μm (40 \times). (B) Overview of ONT and RGC survival analyses. (C) Images of flat-mount retinae at 1, 3, 7 and 14 dpi. Scale bar: 50 μm . (D) RGC survival percentages at 1, 3, 7 and 14 dpi ($n=6/\text{day}$). Graphs show mean \pm s.d.; * $P<0.05$; ** $P<0.01$; *** $P<0.001$; Kruskal–Wallis ANOVA with Dunn’s multiple comparisons. Data derived from three experiments.

ONT as a result of compromised health rather than death of *is/2b:GFP+* RGCs. Cleaved caspase-3 immunostaining revealed that 12.44 \pm 1.79% (mean \pm s.e.m.) of RGCs in the ONT+ retinae were cleaved caspase-3 $^{+}$ at 7 dpi, compared with 0.06 \pm 0.01% of RGCs in the ONT– retina (Fig. S1). Moreover, no bromodeoxyuridine (BrdU) $^{+}$ cells were detected in the GCL of ONT+ retinae at 7 dpi, indicating that RGCs had not yet regenerated. Taken together, these data indicate that, despite transient reduction in *is/2b:GFP* levels and limited death, most *is/2b:GFP+* RGCs are indeed preserved in zebrafish after ONT.

Identification of neuroprotective factors and pathways after ONT

To identify neuroprotective signals/pathways in RGCs, *is/2b:GFP+* RGCs were isolated by fluorescence-activated cell sorting (FACS) from ONT+ and ONT– retinae at 24 h post-injury (hpi) and analyzed by RNA sequencing (RNA-seq) (Fig. 2A). We identified 308 differentially expressed genes (DEGs) (Fig. 2B), of which 56 were upregulated and 252 were downregulated (Tables S2 and S3). We reasoned that neuroprotective factors would be upregulated upon ONT, and the upregulated DEG group included *stat3*, *irf9*, *sox11b*, *lepr* and *socs3b*, which encode components/regulators of the Jak/Stat signaling pathway (Fig. 2C; Villarino et al., 2017), in addition to other neuroprotective and pro-regenerative genes, such as *gap43* (Chung et al., 2020), *atf3* (Kole et al., 2020) and *atoh7* (Brodie-Kommit et al., 2021). Moreover, the interferon regulatory factor genes *irf9* and *irf1b* (Langevin et al., 2013), and the chemokine receptor *cxcr4b* (García-Cuesta et al., 2019), were also upregulated. Although these genes have been shown to be expressed in rodent RGCs after injury (Dvorianchikova et al., 2012; Hilla et al., 2021), it is also possible that a small number of macrophages and microglia contaminated our sorted *is/2b:GFP+* RGCs. Pathway

enrichment analyses indicated that Jak/Stat signaling was the most highly enriched pathway in RGCs after ONT (Fig. 2D). Furthermore, DEGs associated with the adipocytokine signaling pathway, which regulates Stat3-mediated signals (Kadye et al., 2020), were also enriched in RGCs after ONT (Fig. 2D). Downregulated DEGs revealed that neuroactive ligand receptor interactions, MAPK, and calcium signaling pathways were all significantly changed after ONT (Fig. 2D).

We also investigated gene expression changes at 12 hpi. The same selection criteria as for 24 hpi analyses yielded limited numbers of DEGs at 12 hpi, with five upregulated and three downregulated (Table S4). Relaxing our selection criteria from a false discovery rate (FDR) P -value <0.05 to a P -value <0.05 revealed 49 upregulated and 31 downregulated DEGs (Tables S5 and S6). *stat3* and *socs3b* were amongst the upregulated group, further suggesting activation of the Jak/Stat in RGCs after ONT.

Jak/Stat pathway activity is required for RGC survival after ONT

Jak/Stat activity contributes to RGC survival in multiple injury contexts (Boyd et al., 2003; Huang et al., 2007; Luo et al., 2007) and facilitates ON and retinal regeneration (Elsaedi et al., 2014; Kassen et al., 2009; Leibinger et al., 2013; Mehta et al., 2016; Park et al., 2004; Todd et al., 2016; Zhao et al., 2014). To confirm Jak/Stat pathway activation after ONT, we assessed phosphorylated-Stat3 (pStat3) levels in ONT+ and ONT– retinae (Fig. 3A; Movie 1). pStat3 levels were significantly increased in ONT+ RGCs (Fig. 3B; $P<0.01$), supporting the notion that Stat3 may be neuroprotective after ONT. To determine whether Jak/Stat pathway activity is required for RGC survival after ONT, we performed intravitreal (IV) injections of a pan-Jak inhibitor, Pyridone 6 (P6), or 0.05% DMSO at the time of ONT and again at 1 dpi (Fig. 3C). P6 injection resulted

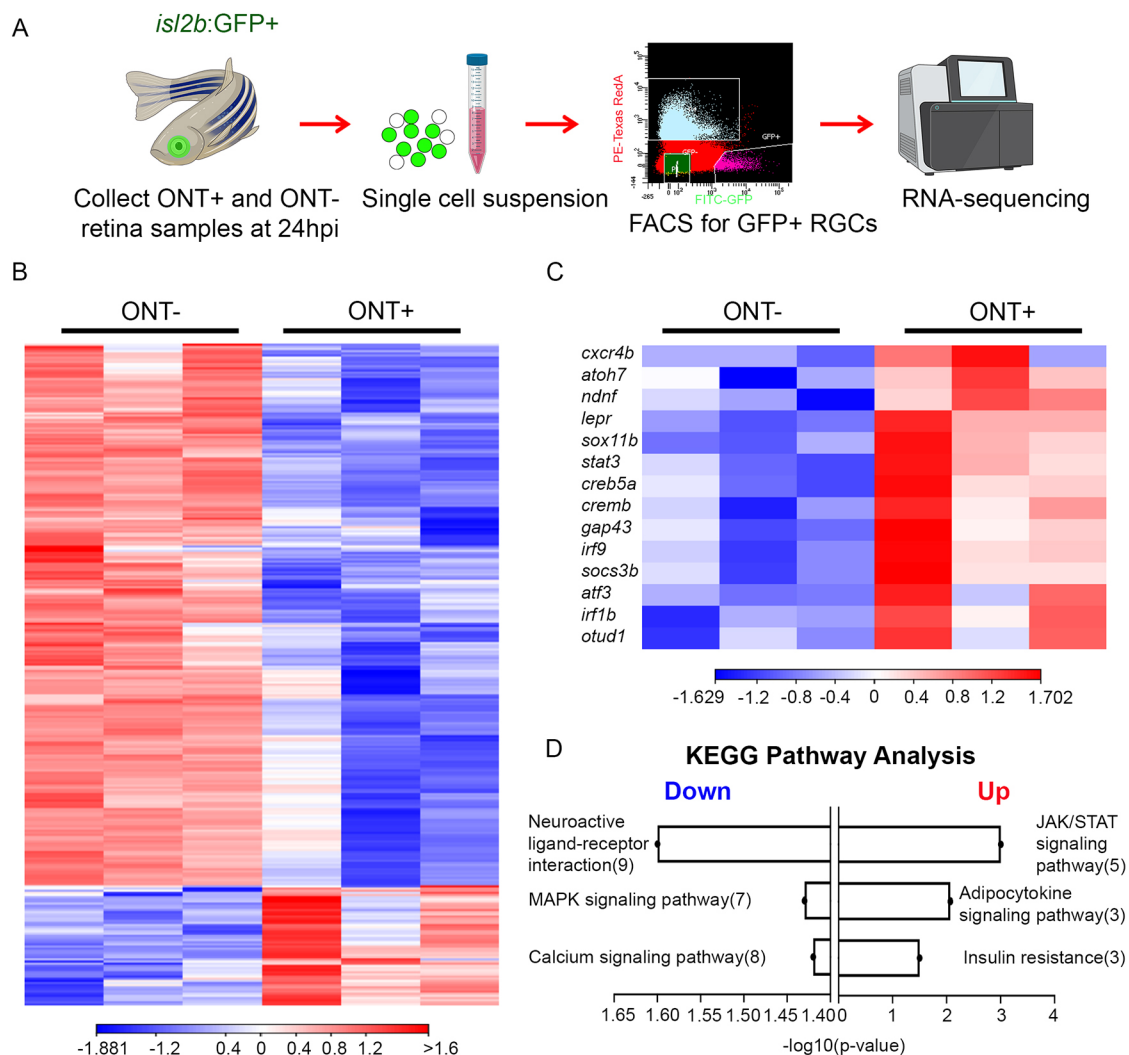


Fig. 2. Identification of DEGs in *isl2b:GFP*⁺ RGCs after ONT. (A) Experimental workflow for FACS-isolation of *isl2b:GFP*⁺ RGCs. An example FACS plot showing a cell-sorting gate is included. Icons were adapted from BioRender.com. Number of experiments=1. (B) Heatmap showing hierarchical clustering of 308 DEGs at 24 hpi from three biological replicates. (C) Heatmap highlighting DEGs of interest based on known neuroprotective and pro-regenerative functions. Heatmap legends show log₂TPM. (D) Pathway enrichment analysis using the KEGG database showing top-3 down- and upregulated pathways after ONT.

in significant reductions to pStat3 levels in ONT+ RGCs at 1 dpi and 7 dpi, supporting the efficacy of P6 in blocking Jak activation in zebrafish (Fig. 3D,E). We next quantified *isl2b:GFP*⁺ RGC survival at 7 dpi when Jak activity was impaired (Fig. 3F,G). DMSO had no effect on RGC survival, and neither did P6 alone ($P=0.4796$; Fig. 3G). However, IV P6 injection in conjunction with ONT resulted in a significant reduction in RGC survival to $39.44\pm 6.99\%$ ($P<0.0001$) and a corresponding increase in cleaved caspase-3 levels in ONT+ RGCs (Fig. S3). These data demonstrate a role for Jak/Stat pathway activity in protecting zebrafish RGCs after ONT.

Innate immune response involvement after ONT

After ONI, the effects of injury-activated immune responses are varied. In some contexts, immune responses stimulate the recruitment and activation of leukocytes, which generate secondary signals to modulate RGC survival and death pathways (Barış and Tezel, 2019; Mac Nair et al., 2016; Nadal-Nicolás et al., 2017; Williams et al., 2017), whereas in others the stimulation of limited neuroinflammation induces pro-survival and regenerative

responses (Kanagaraj et al., 2020; Todd et al., 2019). In zebrafish, leukocyte activity accelerates axonal regeneration after neuronal damage (Tsarouchas et al., 2018), including in RGCs after ON crush (Van Dyck et al., 2021). With this in mind, we next investigated how components of the innate immune system respond to ONT in zebrafish and whether they contribute to RGC vulnerability. We focused on macrophages/microglia, leukocytes that accumulate in the zebrafish retina after a variety of injury types and facilitate repair and regeneration (Leach et al., 2021; White et al., 2017). In zebrafish, the 4C4 antibody recognizes an unidentified protein expressed by macrophages/microglia (Craig et al., 2008). Consistent with other reports (Mitchell et al., 2018), 4C4⁺ macrophages/microglia were located throughout the ONT- retina, including in the GCL (Fig. 4A,B; Movie 2). At 1 dpi, the number of 4C4⁺ cells appeared to increase in the GCL (Fig. 4A,B). Similarly, utilizing *mpeg1:mCherry* transgenic fish (Ellett et al., 2011), we observed an apparent increase in mCherry⁺ macrophages/microglia in the ONT+ GCL at 1 dpi (Fig. 4C). Owing to the morphology and close proximity of macrophages/microglia in the ONT retinae, it was difficult to identify single

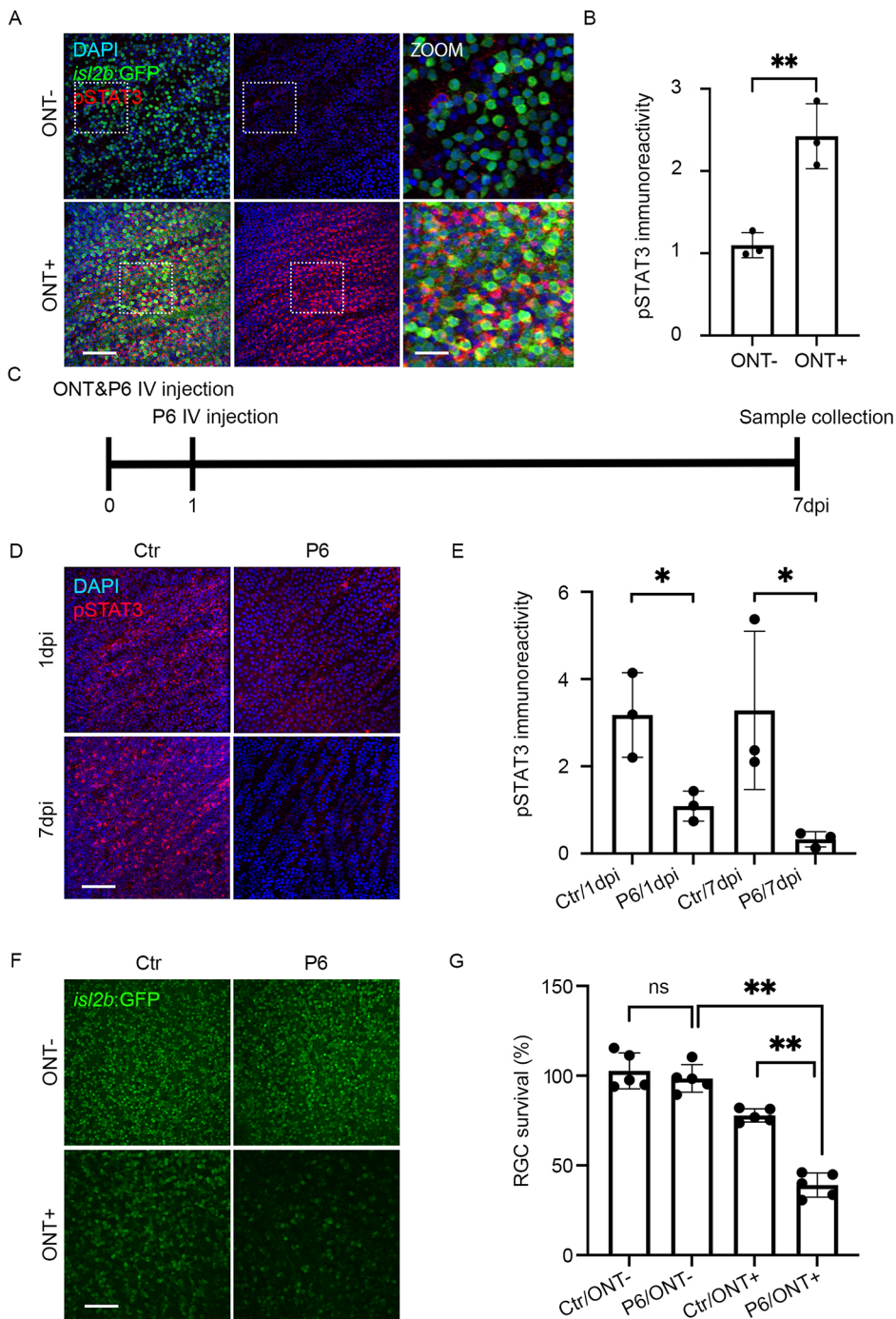


Fig. 3. Jak/Stat pathway activity is required for RGC survival after ONT. (A) pStat3 expression (red) in flat-mount *is/2b:GFP* ONT– and ONT+ retinas at 1 dpi. Nuclei are stained with DAPI (blue). Right-hand panels show 3× magnifications of the boxed regions in the 40× images. (B) Quantification of pStat3 levels at 1 dpi. pStat3 levels in ONT+ RGCs are shown relative to those in ONT– retinas. Shown are mean±s.d. of $n=3$ for each group; ** $P<0.01$; Mann–Whitney test. Data derived from three experiments. (C) Experimental protocol to assess Jak requirement for RGC survival after ONT. (D) pStat3 expression in ONT+ *is/2b:GFP* retinal flat mounts at 1 and 7 dpi with or without IV injection of the Jak inhibitor P6. DMSO was used as control (Ctr). (E) Quantification of pStat3 expression after P6 application at 1 and 7 dpi. $n=3$ /condition. Shown are mean±s.d.; * $P<0.05$, Mann–Whitney test. Data derived from three experiments. (F) Images of P6- or DMSO-injected flat-mount *is/2b:GFP* retinas at 7 dpi. (G) Quantification of RGC survival after P6 injection ($n=5$ /condition). Shown are mean±s.d.; ** $P<0.01$; Kruskal–Wallis ANOVA with Dunn’s multiple comparisons. Scale bars: 50 μm (A, main panels; D,F); 150 μm (A, zoom). Data derived from four experiments.

macrophages/microglia for counting and therefore we quantified the percentage area covered by mCherry⁺ cells within the GCL (Fig. 4D). Compared with ONT– retinas, mCherry⁺ cells covered nearly six times more GCL area after ONT ($P=0.0006$; Fig. 4D). Macrophage/microglia morphologies change upon activation, whereby quiescent cells with a ramified morphology take on a more spherical/amoeboid shape when activated (Karlstetter et al., 2015; Mitchell et al., 2018). Morphological differences were evident in both 4C4⁺ and mCherry⁺ cells within the GCL of ONT+ retinas (Fig. 4B,C), and quantification of mCherry⁺ cell sphericity revealed a significant increase at 1 dpi (Fig. 4E; $P<0.0001$); collectively, these data show that macrophages/microglia accumulate in the GCL and become activated after ONT.

Blocking inflammation or depletion of macrophages/microglia protects RGCs after ONT

To test the hypothesis that macrophage/microglia-mediated inflammation might contribute to RGC death after ONT, we inhibited inflammation after ONT by dexamethasone treatment, a strategy shown to protect RGCs in other injury models (Bollaerts et al., 2019; Dutt et al., 2010; Gallina et al., 2015; Jovanovic et al., 2020), and quantified RGC survival. Experimentally, 2 μl of 10 μM dexamethasone or 0.05% DMSO (vehicle control) was IV injected into *is/2b:GFP* fish at the time of ONT and again at 1 dpi and tissue was collected at 7 dpi (Bollaerts et al., 2019) (Fig. 3C). Dexamethasone efficacy was verified via qRT-PCR for *pxr* (also known as *nr1i2*), a dexamethasone-regulated gene (Fig. S4A;

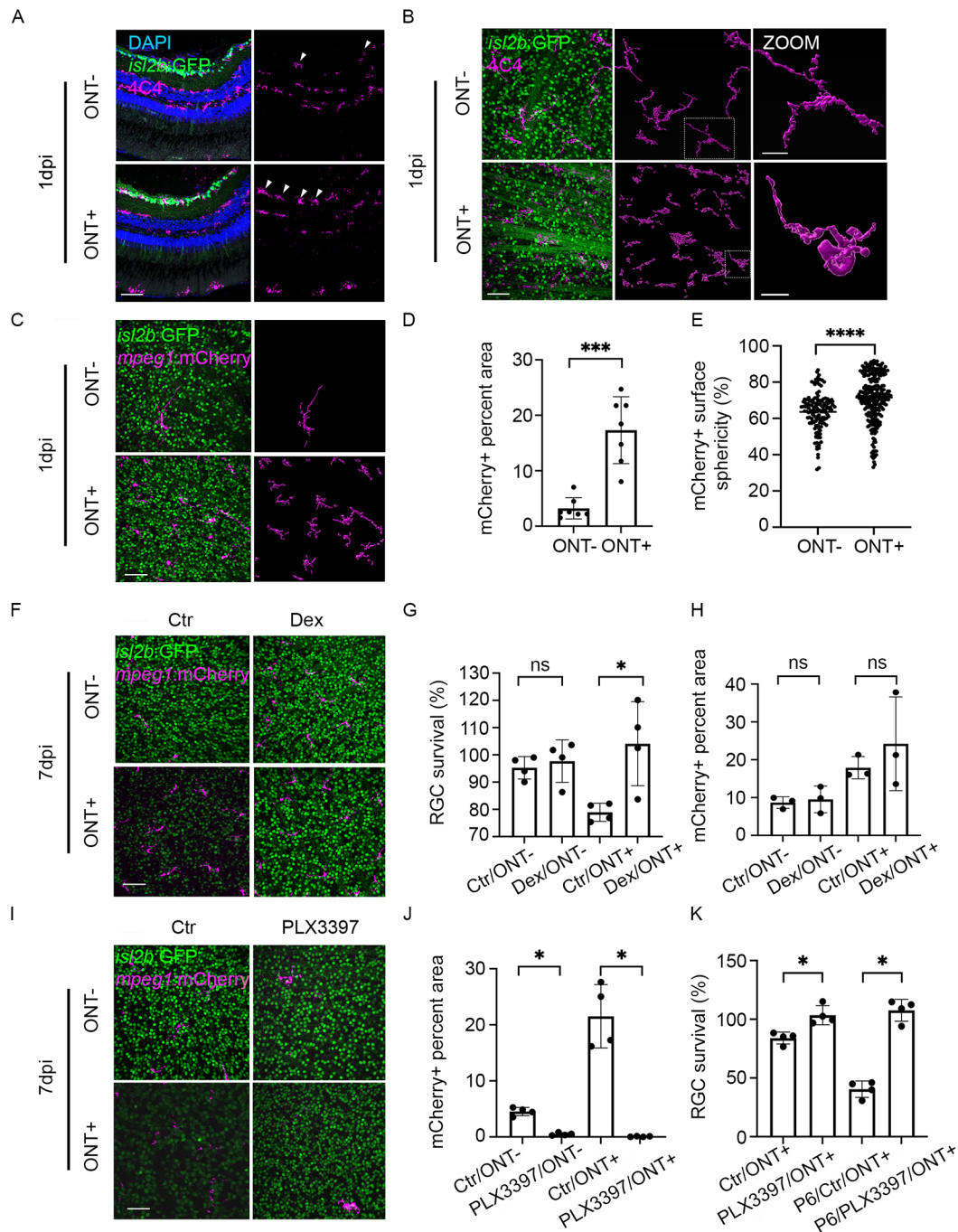


Fig. 4. Macrophages/microglia are recruited to the GCL after ONT and mediate RGC death. (A,B) Immunostaining of 4C4 (magenta) on *isl2b:GFP* retinal cryosections (A) and retinal flat mounts at 1 dpi with surface renderings of 4C4+ macrophages/microglia generated in Imaris (B). (C) Images of retinal flat-mounts from *isl2b:GFP;mpeg1:mCherry* animals at 1 dpi. mCherry+ macrophages/microglia (magenta) surfaces were rendered in Imaris. (D) Quantification of the surface area within the GCL occupied by mCherry+ macrophages/microglia at 1 dpi ($n=7$ /condition). Shown are mean \pm s.d.; *** $P < 0.001$; Mann–Whitney test. Data derived from one experiment. (E) Violin plot showing a significant increase in sphericity of mCherry+ macrophages/microglia in ONT+ retinæ compared with ONT- controls ($n=140$ in ONT- and $n=272$ in ONT+). **** $P < 0.0001$; unpaired *t*-test with Welch’s correction. (F) Flat-mount images of *isl2b:GFP;mpeg1:mCherry* retinæ after IV injection of dexamethasone (Dex) or DMSO (Ctr) at 7 dpi. (G) RGC survival in dexamethasone-treated retinæ increased significantly at 7 dpi compared with control ($n=4$ /condition). Shown are mean \pm s.d.; * $P < 0.05$; Kruskal–Wallis ANOVA test with Dunn’s multiple comparisons. Data derived from three experiments. (H) Quantification of mCherry+ macrophage/microglia coverage of the GCL after ONT and dexamethasone or DMSO injection ($n=3$ /condition). Shown are mean \pm s.d.; Kruskal–Wallis ANOVA test with Dunn’s multiple comparisons. No significant differences were detected. Data derived from three experiments. (I) Flat-mount images of *isl2b:GFP;mpeg1:mCherry* retinæ after PLX3397 treatment or housed in system water (Ctr) at 7 dpi. (J) Quantification of mCherry+ macrophage/microglia coverage of the GCL after ONT and PLX3397 treatment ($n=4$ /condition). Shown are mean \pm s.d.; * $P < 0.05$; Kruskal–Wallis ANOVA test with Dunn’s multiple comparisons. Data derived from three experiments. (K) RGC survival in PLX3397-treated retinæ increased significantly at 7 dpi compared with control. Similarly, RGC survival in PLX3397-treated retinæ increased significantly after P6 addition compared with DMSO controls ($n=4$ /condition). Shown are mean \pm s.d.; * $P < 0.05$; Mann–Whitney test. Data derived from three experiments. Scale bars: 50 μ m. ns, not significant.

Shi et al., 2010). Dexamethasone alone had no effect on *isl2b*:GFP⁺ RGC survival in the ONT⁻ retina (96.56±8.67%, $P=0.5998$), but significantly increased survival of ONT⁺ RGCs at 7 dpi, compared with DMSO controls (104.1±13.35%, $P<0.05$; Fig. 4G). Dexamethasone prevents the recruitment of macrophages/microglia in some contexts (Tsarouchas et al., 2018; White et al., 2017), but not others (Chatzopoulou et al., 2016; Warchol, 1999; Xie et al., 2019; Leach et al., 2021). Quantification of the percentage of the GCL covered by *mpeg1*:mCherry⁺ macrophages/microglia in DMSO- and dexamethasone-injected retinas revealed no significant differences after ONT (Fig. 4H; $P=0.0623$). Quantification of mCherry⁺ cell sphericity, as a proxy for macrophage/microglia activity, revealed no significant differences between dexamethasone-treated retinas and controls at 7 dpi (Fig. S4B). However, dexamethasone suppressed the expression of inflammatory markers, such as *il1b* (Mitchell et al., 2019; Tsarouchas et al., 2018), *il6* and *tnfa* (Bessler et al., 1999), after ONT (Fig. S4A).

Previous reports have identified roles for microglia in contributing to RGC death after a variety of insults (Bosco et al., 2008; Jovanovic et al., 2020; Takeda et al., 2018). However, other studies have shown that microglia are dispensable for RGC survival after ON crush (Hilla et al., 2017), and may instead provide neuroprotective and/or pro-regenerative signals (Bell et al., 2018; Sappington et al., 2006). To test directly the requirement of macrophages/microglia in modulating RGC death in zebrafish after ONT, we depleted macrophages/microglia using PLX3397, a potent inhibitor of the colony-stimulating factor 1 receptor (Csf1r). Csf1r activity is required for macrophage/microglia differentiation (Lin et al., 2008; Sherr et al., 1985) and PLX3397 has been utilized effectively in zebrafish (e.g. Conedera et al., 2019; Leach et al., 2021; Van Dyck et al., 2021). Animals were immersed in 500 nM PLX3397 1 day prior to ONT and retinas were collected at 7 dpi. To validate the efficiency of macrophage/microglia depletion by PLX3397, we quantified the percentage area of the GCL occupied by *mpeg1*:mCherry⁺ macrophages/microglia. PLX3397 significantly reduced the coverage of *mpeg1*:mCherry⁺ cells in both the ONT⁻ and ONT⁺ GCL (Fig. 4I,J). Similar to dexamethasone-mediated RGC protection (Fig. 4F,G), PLX3397-mediated depletion of macrophages/microglia significantly increased RGC survival at 7 dpi (103.49±12.01%, $P<0.05$; Fig. 4I,K) concomitant with decreased cleaved caspase-3 levels in ONT⁺ RGCs (Fig. S3). PLX3397-mediated depletion of macrophages/microglia also rescued RGC survival and decreased cleaved caspase-3 levels at 7 dpi after Jak/Stat pathway inhibition using P6 (107.60±9.32%, $P<0.05$; Fig. 4K; Figs S2 and S3), indicating that Jak/Stat activity is dispensable in the absence of macrophage/microglia recruitment. pStat3 was detected in both RGCs and also some *mpeg1*:mCherry⁺ macrophages/microglia in ONT⁺ retinas (Fig. S5A). Expression of pStat3 was eliminated in both cell types in the P6/PLX3397-treated ONT⁺ retinas (Fig. S5B). However, the observation that pStat3 was detected in both RGCs and macrophages/microglia after ONI indicates that the effects of Jak/Stat pathway inhibition could reflect activities in one or both cell types. Future studies targeting loss- and/or gain-of-function assays to RGCs and/or leukocytes will be needed to parse contributions from each cell type to RGC survival after ONI.

Collectively, these data support a model in which crosstalk between macrophages/microglia and RGCs, mediated by Jak/Stat pathway activity, regulates RGC survival after ONI. As noted above, Stat3 upregulation has been associated with RGC survival in some experimental contexts (Huang et al., 2007; Luo et al., 2007).

Despite this, overall RGC survival is limited under physiological conditions, with over 90% of RGCs lost within 28 days after injury in mice (Li et al., 2020). This is not true for all mammals, however. Indeed, the naked mole-rat retains many RGCs after injury, to at least 28 days (Park et al., 2017). Interestingly, although pStat3 is nearly absent in mouse RGCs, even after injury, it increases significantly in mole-rat RGCs after ON crush, supporting a possible role for Stat3 activity in RGC neuroprotection.

Counter to our expectations, pStat3 localization was predominantly cytoplasmic in ONT⁺ RGCs, rather than nuclear (Fig. 3A; Movie 1). This observation is consistent with cytoplasmic pStat3 localization in the regenerating zebrafish retina (Elsaedi et al., 2014) and in mouse motor neurons responding to cytokine stimulation (Selvaraj et al., 2012). Stat3 possesses transcription-independent functions such as cytoplasmic regulation of autophagy (Shen et al., 2012). Stat3 also localizes to mitochondria after cytokine stimulation in mouse RGCs where it regulates metabolic functions and enhances axon regeneration after ONI (Luo et al., 2016). Thus, it is possible that the function of Stat3 in RGC neuroprotection could be independent of transcription.

Not all zebrafish RGCs survive ONT and cleaved caspase-3⁺ RGCs appeared to be distributed in a non-random pattern in the ONT⁺ retina (Fig. S1). This may indicate that there are RGC subtype(s) that are more susceptible to ONI, similar to observations in mice (Tran et al., 2019). RGC subtypes in zebrafish have been recently characterized (Kölsch et al., 2021) and additional studies will be required to determine whether specific subtypes are lost after ONT, and, if so, whether these subtypes lack the ability to upregulate Jak/Stat activity after injury. Finally, it will be of interest to identify the signals emanating from macrophages/microglia that activate death and/or pro-survival pathways in zebrafish RGCs after injury, as these would also be promising targets around which neuroprotective therapies for glaucoma could be developed (García-Bermúdez et al., 2021; Rashid et al., 2019).

MATERIALS AND METHODS

Animals

Zebrafish (*Danio rerio*) in this study were 3-5 months old and an equal number of males and females were used in all experiments. Transgenic lines used were *isl2b*:GFP (Pittman et al., 2008) and *mpeg1*:mCherry (Ellett et al., 2011; a gift from Dr Neil Hukriede, University of Pittsburgh, USA). Animals were maintained under standard conditions at 28.5°C on a 14 h light/10 h dark cycle. There were no differences in outcomes based on sex of the fish and therefore all data were combined for analyses. All animals were treated in accordance with provisions established by the University of Pittsburgh School of Medicine Institutional Animal Care and Use Committee. Biological replicates are stated in figure legends for each experiment. At least three independent biological replicates were used per experiment.

ONT

ONT was performed as previously described (Elsaedi et al., 2014; Zou et al., 2013). Zebrafish were anesthetized in 0.03% tricaine buffer (MS-222; Fisher Scientific) and placed on a moist tissue paper under a dissecting scope (Leica E65S). ONT surgery was performed on the left eye. After removal of the connective tissue, the eyeball was pulled out from the orbit gently using forceps. The ON and ophthalmic artery, which runs along with the ON, were exposed and the ON was then completely transected using additional forceps, after which the eye was placed back in the orbit. Any animals in which bleeding was observed were euthanized and not used for analysis. The right eye was subjected to a sham surgery as control: connective tissue was removed, the eye was pulled out from the orbit gently, and then placed back in the orbit. Fish were returned to system water in separate tanks to recover.

RGC isolation and FACS

Retinae were harvested from *isl2b*:GFP zebrafish at 12 and 24 h post-injury (hpi) in biological triplicate. Four retinae were collected per sample. For retinal isolations and cell dissociation, animals were euthanized by tricaine overdose and transferred to PBS for enucleation. To achieve a single-cell suspension, the eyeball was rinsed in 1× PBS post-enucleation and incubated in StemPro Accutaset Cell Dissociation Reagent (Thermo Fisher, #A1110501) at 28.5°C for 40 min in a water bath. The cell suspension was then passed through a 70-µm cell strainer (Fisher Scientific) and gently pelleted by centrifugation at 2189 rpm (450 g) for 5 min at 4°C. After two washes in ice-cold 1× PBS, cells were resuspended in ice cold 5% fetal bovine serum in 1× PBS. GFP⁺ cells were sorted using a FACS Aria IIu cell sorter (BD Biosciences) at the Flow Cytometry Core at the University of Pittsburgh School of Medicine Department of Pediatrics. The gate for FACS was set by GFP intensity for wild-type (non-transgenic) and both injured (ONT) and intact (control) *isl2b*:GFP samples. The same gating settings were used for both the ONT and control RGC samples and for all biological replicates.

RNA-seq and bioinformatics analyses

Library preparation, quality control analysis, and next-generation sequencing were performed by the Health Sciences Sequencing Core at Children's Hospital of Pittsburgh as previously described (Leach et al., 2021). cDNA sequencing libraries were prepared using a SmartSeq HT kit (Takara Bio) and Illumina Nextera XT kit (Illumina Inc.) and 2×75 paired-end, 150 cycle sequencing was performed on a NextSeq 500 system (Illumina), aiming for 40 million reads per sample. After sequencing, raw read data were imported to the CLC Genomics Workbench (Qiagen Digital Insights) licensed through the Molecular Biology Information Service of the Health Sciences Library System at the University of Pittsburgh. After mapping trimmed reads to the *Danio rerio* reference genome (assembly GRCz11), DEGs from the 24 hpi time point were identified using the following filter: the maximum of the average group RPKM value >1.5, absolute fold change >2, FDR *P*-value <0.05. Genes with TPM=0 in one or more replicates were excluded. This filtering strategy was also used for DEGs at 12 hpi and a second analysis was performed in which the FDR *P*-value <0.05 was switched to a *P*-value <0.05. Pathway enrichment analyses were performed using DAVID Bioinformatics Resources 6.8 (<https://david.ncifcrf.gov>) and using the KEGG database.

Pharmacological experiments

To assess toxicity and efficacy, the JAK inhibitor Pyridone 6 (P6), or dexamethasone (both Sigma-Aldrich) were IV injected into the intact and injured retina at several different concentrations, as previously described (Elsaedi et al., 2014), and RGC survival was quantified. Similar to previous studies (Bollaerts et al., 2019; Elsaedi et al., 2014), 5 µM P6 and 10 µM dexamethasone were utilized. The first dose of each compound (2 µl) was injected immediately after ONT (0 dpi) and the second dose (2 µl) was injected at 1 dpi. For all control doses, 2 µl of 0.05% DMSO was injected. To deplete macrophages/microglia, fish were immersed in 500 nM PLX3397 (Fisher Scientific) or 0.05% DMSO (control) in system water, as previously described (Kanagaraj et al., 2020). PLX3397 exposure started 1 day before ONT and system water containing PLX3397 was replaced daily during the experiment.

Immunohistochemistry

Immunofluorescence staining on retinal cryosections and flat-mounted retinae were performed as previously described (Uribe and Gross, 2007; Zou et al., 2013) with the addition of an antigen-retrieval step consisting of 100% methanol incubation at -20°C for 30 min for staining pStat3 (MBL International Corporation, D128-3) and cleaved caspase-3 (Abcam, ab13847). For retinal flat-mounts, after euthanasia, fish were decapitated and heads were fixed in 4% paraformaldehyde (PFA) at 4°C overnight. The retina was dissected in ice-cold PBS, washed in PBST (0.1% Triton X-100 in PBS), and then incubated in a 1.5 ml tube on a rotator overnight at 4°C with antibodies against 4C4 (1:200, a kind gift of Dr Peter Hitchcock, University of Michigan School of Medicine, USA; Craig et al., 2008),

pStat3 (1:100, MBL International Corporation), cleaved caspase-3 (1:200, Abcam) and mCherry (1:200, Takara Bio/Clontech Laboratories, 632543). Retinae were then washed in PBST for 3×10 min at room temperature and incubated with goat-anti mouse Cy3 (1:250, Jackson ImmunoResearch Laboratories, 115-165-166) or goat-anti rabbit Alexa 647 (1:500, Cell Signaling Technology, 8940) secondary antibodies for 3 h at room temperature. Samples were then washed with PBST for 3×10 min and carefully cut into four quadrants and mounted on slides with DAPI VECTASHIELD (Vector Laboratories, H-1200). For cryosections, samples were prepared as previously described (Uribe and Gross, 2007); zn-8 (Zebrafish International Resource Center) was used at 1:200 and all other antibodies were used at the same concentrations as for the retinal flat-mount.

BrdU incorporation assays

Adult *isl2b*:GFP⁺ fish were immersed in 10 mM BrdU (Sigma-Aldrich) dissolved in system water from 6 dpi to 7 dpi, and sacrificed at 7 dpi. As a positive control for BrdU incorporation and immunohistochemistry, a needle poke injury was performed following previously published protocols (Fausett and Goldman, 2006) and fish were exposed to BrdU for 24 h prior to being sacrificed. Immunohistochemistry for BrdU proceeded as described above for other antibodies, with the addition of a 10-min incubation of 4 N HCl at 37°C to relax chromatin. Anti-BrdU (Abcam, ab6326) was used at 1:200.

Confocal microscopy, image processing and quantification

For *isl2b*:GFP imaging, retinal flat-mounts were prepared as above, with the head fixed in 4% PFA overnight at 4°C and the retina dissected and mounted on the second day. Images were taken using Olympus Fluoview FV1200 laser scanning microscope and a 40× objective (Olympus Corporation). Images were taken from each of the four quadrants (one peripheral and one central per quadrant). Quantification of RGC numbers was performed using particle analysis in ImageJ after setting up a consistent threshold for all images. RGC survival was calculated as the ratio of *isl2b*:GFP⁺ RGCs in the left (ONT⁺ and/or drug treatment) eye/*isl2b*:GFP⁺ RGCs in the right (ONT⁻ and/or DMSO control) eye of the same fish.

Quantification of macrophages/microglia was performed using Imaris 9.6.0 (Bitplane). Confocal images were first converted into Imaris files, and 3D-rendered surfaces were then created for mCherry or 4C4 using the same algorithm (smoothing=0.4 µm, absolute intensity threshold=1560, objects area>50 µm²) for each dataset. Quantification of total surface area and sphericity was performed using Imaris. Measurements were exported and statistically assessed using Prism 9.0 (GraphPad).

For GFP fluorescence intensity quantification, the GFP signal intensity of 30 randomly selected RGCs in the ONT⁺ and ONT⁻ retina per fish (*n*=6) was measured using Fiji ImageJ and the corrected total cell fluorescence (CTCF) was obtained following the protocol of L. Hammond (The University of Queensland, Australia; <https://theolb.readthedocs.io/en/latest/imaging/measuring-cell-fluorescence-using-imagej.html>). Briefly, using the freehand tool in ImageJ, single RGCs were outlined as a region of interest and GFP intensity was measured as well as the background fluorescence for every retina. CTCF was then calculated as follows: integrated density—(area of selected cell×mean fluorescence of background readings). Relative intensity was calculated as the ratio of the CTCF in ONT⁺ RGCs divided by that of ONT⁻ RGCs.

Localization of pStat3 expression to RGCs was confirmed with surface creation in Imaris 9.6.0. Quantification of pStat3 levels was performed as described (Osborne et al., 2018) using Fiji (ImageJ). Briefly, *isl2b*:GFP labeling was used to identify the location and depth of the GCL. RGC volumes were converted to *z*-projections and background subtraction and speckle removal was performed on all images. Threshold levels for pStat3 were determined from ONT⁻ samples and consistent thresholds were then applied to all images in ONT⁺ samples and integrated density was measured. The expression level of pStat3 was calculated as the ratio of the average integrated density of pStat3 in 1 dpi, 7 dpi, P6/1 dpi or P6/7 dpi retinae to the average integrated density of the contralateral ONT⁻ eye.

Cleaved caspase-3 levels were quantified in two ways: (1) the number of cleaved caspase-3⁺ nuclei relative to the number of RGCs or (2) the ratio of

thresholded area pixels of staining to the number of DAPI+ nuclei. For the latter, cleaved caspase-3 staining volumes were converted to z -projections with a consistent number of z -sections between samples. Background subtraction and speckle removal was performed on all images using Fiji (ImageJ). Threshold levels were determined from control/ONT+ samples and applied consistently for all images. Pixels of cleaved caspase-3 staining areas were then measured and recorded. DAPI+ nuclei counts were performed as described for RGC quantification.

Quantitative real-time PCR (qRT-PCR)

RNA was extracted from adult *isl2b:GFP+* retinæ using Trizol reagent (Fisher Scientific, 15-596-018) and cDNA generated using the iScript cDNA Synthesis Kit (Bio-Rad Laboratories). For each treatment group, three biological replicates were collected with each replicate consisting of three retinæ from three different fish. qRT-PCR was performed on a CFX384 Touch Real-Time PCR Detection System (Bio-Rad Laboratories) using the iTaq Universal SYBR Green Supermix (Bio-Rad Laboratories). Primer sequences are provided in Table S1. Gene expression fold change was calculated using the comparative CT method ($2^{-\Delta\Delta CT}$) and expression in DMSO/ONT- retinæ was normalized to 1 as the control (Livak and Schmittgen, 2001). *gapdh* was used as the housekeeping gene in these experiments (Barber et al., 2005).

Statistics

All statistical analyses were performed using Prism 9.0 (GraphPad). Data are presented as mean \pm s.d., with the exception of cleaved caspase-3 images, which show mean \pm s.e.m. For multiple comparisons, Kruskal–Wallis ANOVA followed by Dunn's multiple comparisons tests between groups were performed. For comparisons between two groups, non-parametric Mann–Whitney tests were performed, with the exception of sphericity comparison, for which an unpaired t -test with Welch's correction was performed. P -values, sample sizes and statistical analyses for each experiment are included in the figure legends.

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Competing interests

The authors declare no competing or financial interests.

Author contributions

Conceptualization: T.K., J.M.G.; Methodology: K.L.L., T.K.; Formal analysis: S.C.; Investigation: S.C.; Writing - original draft: S.C.; Writing - review & editing: J.M.G.; Supervision: T.K., J.M.G.; Project administration: J.M.G.; Funding acquisition: J.M.G.

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Data availability

Raw read and processed RNA-seq data have been deposited in Gene Expression Omnibus under accession number GSE171426.

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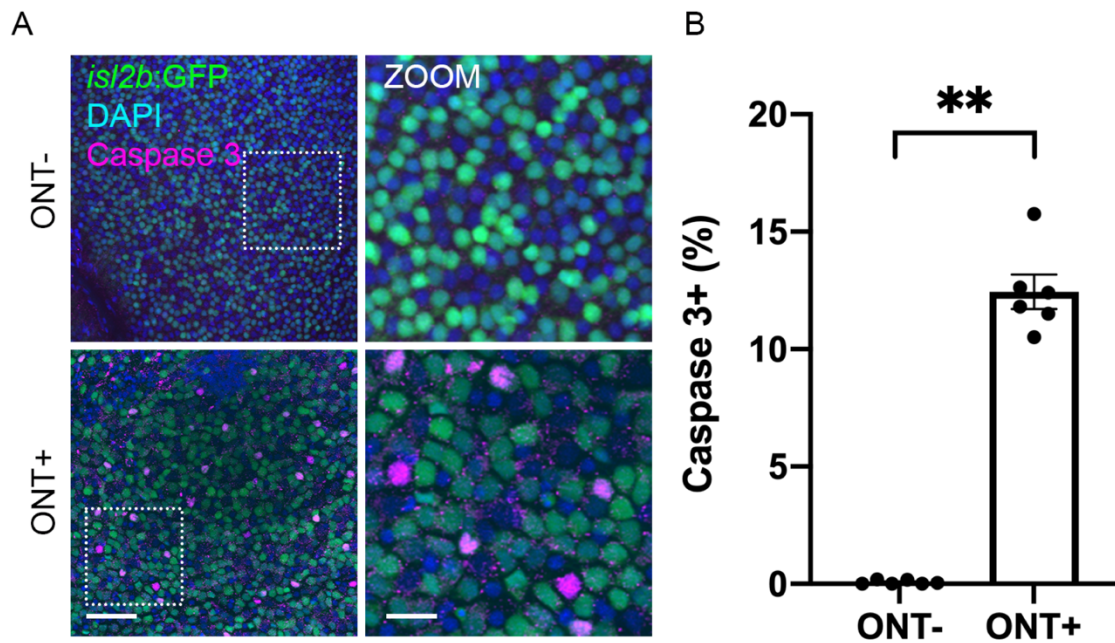


Fig. S1. Cleaved caspase-3 expression in the ganglion cell layer after optic nerve transection.

(A) Cleaved caspase-3 (magenta) expression in *isl2b:GFP*⁺ (green) flat-mount retinæ at 7dpi. DAPI (blue) counterstain for nuclei. **(B)** Quantification of the percentage of cleaved caspase-3⁺ RGCs in the ganglion cell layer after ONT (n=6/condition). Shown are mean±SEM; **p<0.01; Mann-Whitney test. Data derived from 3 experiments. Scale bars = 50µm and 150µm on zoomed images.

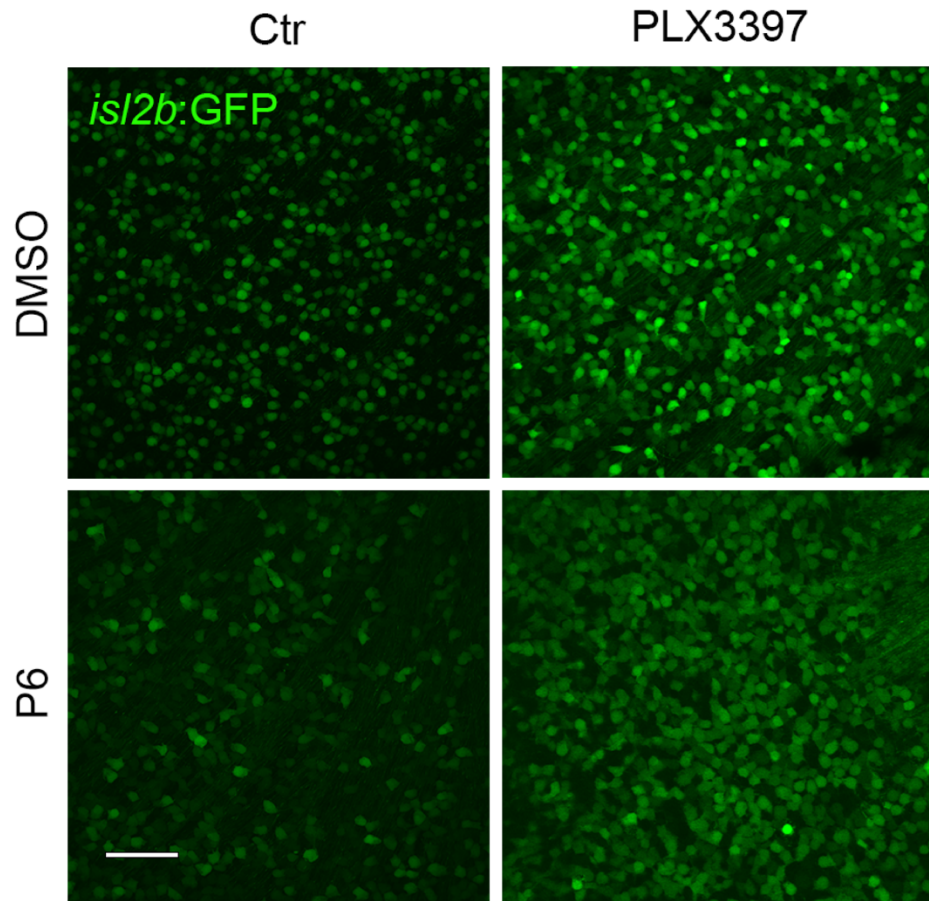


Fig. S2. RGC survival after Jak inhibition and macrophage/microglia depletion.

Flat-mount images of *is/2b:GFP* retinæ after intravitreal injection of P6 or DMSO +/- PLX3397 treatment at 7dpi. Data derived from 3 experiments. Scale bars = 50µm.

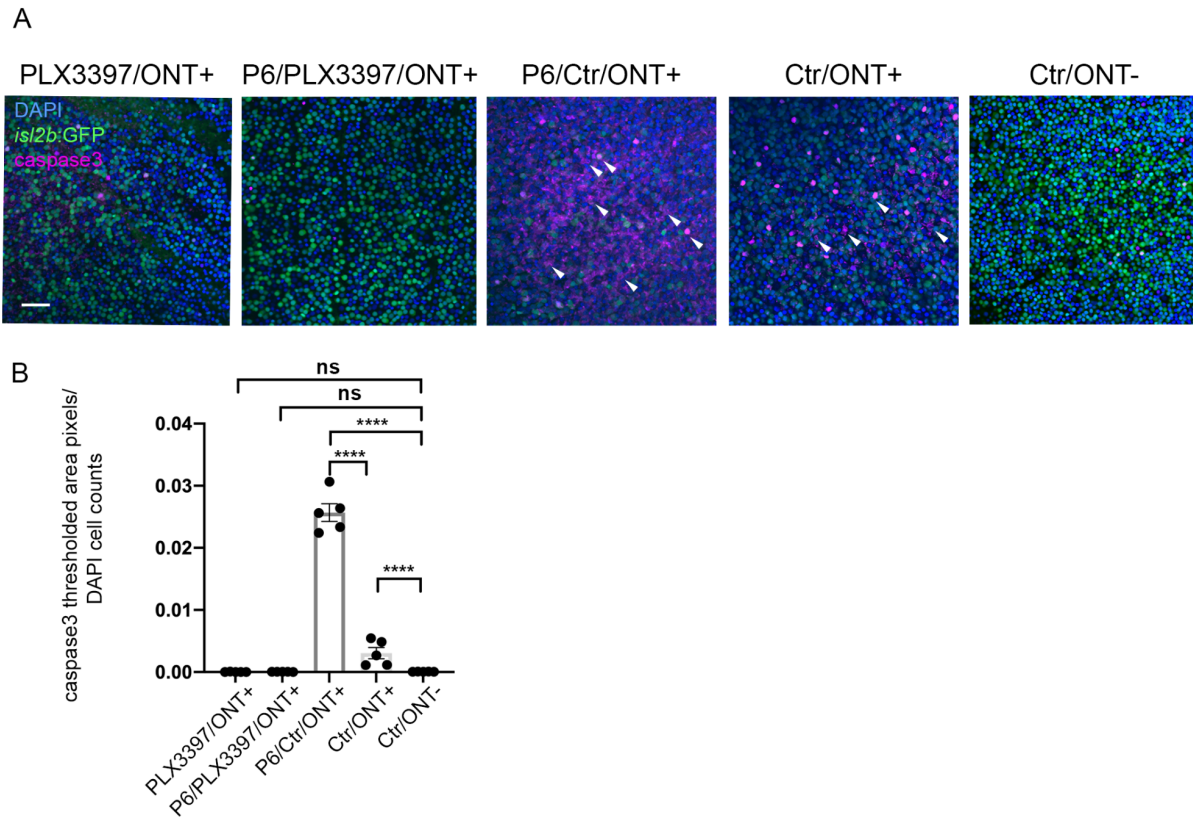


Fig. S3. Cleaved caspase-3 staining after JAK inhibition and macrophage/microglia depletion.

(A) Cleaved caspase-3 (magenta) expression in *is/2b:GFP*⁺ (green) flat-mount retinæ at 7dpi. DAPI (blue) counterstain for nuclei. Scale bar=50 μ m. **(B)** Quantification of the ratio of the thresholded area pixels of cleaved caspase-3⁺ staining to counts of DAPI cells in the ganglion cell layer (n=5/condition). Shown are mean \pm SD; **** p <0.0001; Kruskal Wallis ANOVA with Dunn's multiple comparisons. Data derived from 2 experiments.

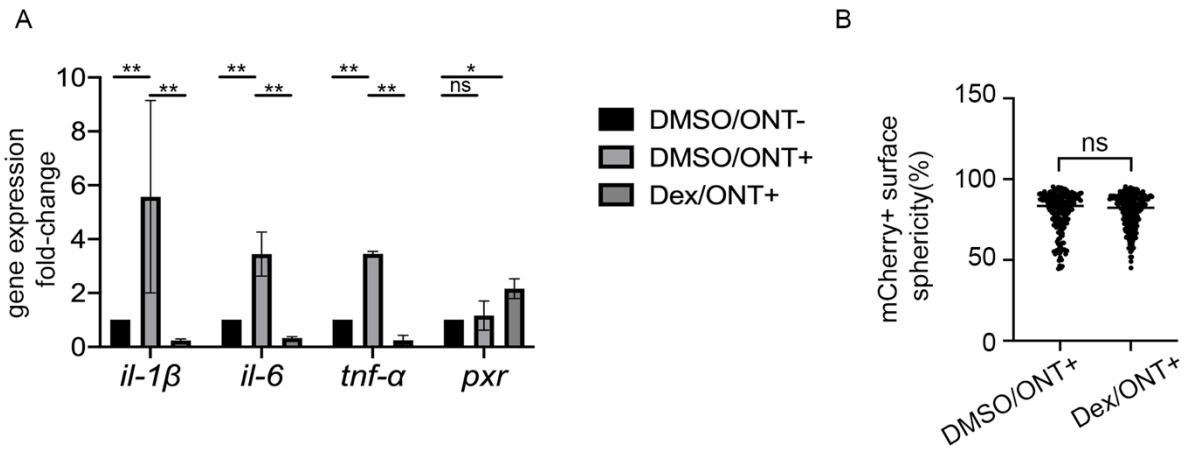


Fig. S4. Quantification of sphericity of macrophages/microglia and qRT-PCR after dexamethasone application.

(A) qRT-PCR for *il1-β*, *il6*, *tnf-α* and a downstream transcriptional target of dexamethasone *pxr* (Pascussi et al. 2000) expression in retinae treated with 10μM dexamethasone or 0.05%DMSO at 7dpi, expression fold change of above genes in DMSO/ONT- retinae were normalized to 1 as the control. *gapdh* were used as the housekeeping gene. Shown are mean±SD; **p<0.01; *p<0.05. Kruskal Wallis ANOVA with Dunn's multiple comparisons. Data derived from 3 experiments. **(B)** Quantification of the sphericity of mCherry+ surfaces after dexamethasone application at 7dpi. Violin plot showing no significant differences in sphericity of mCherry+ macrophages/microglia in Dex/ONT+ retinae compared to DMSO/ONT+ controls (n=180 in Dex/ONT+ and n=198 in DMSO/ONT+); p=0.4235, unpaired t-test with Welch's correction.

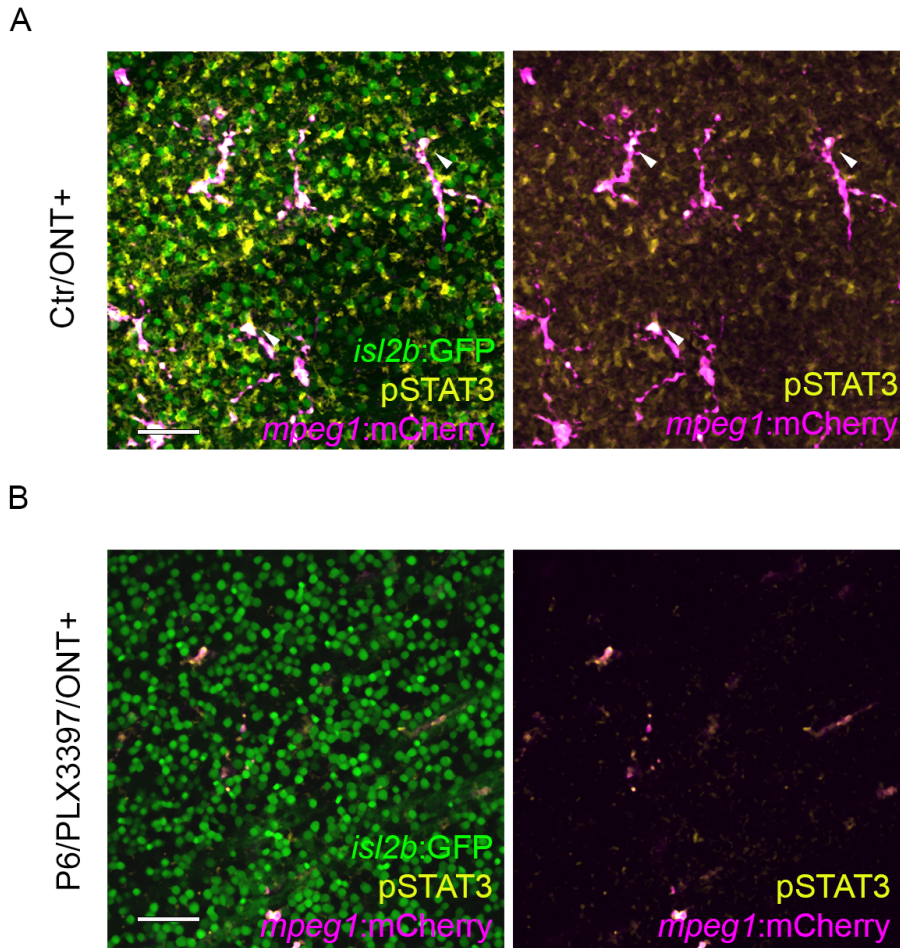


Fig. S5. JAK/STAT pathway activity in macrophages/microglia and after P6/PLX3397 treatment.

(A) Flat-mount images showing pSTAT3 staining (yellow) on *isl2b:GFP*;*mpeg1:mCherry* (*isl2b:GFP* – green; *mpeg1:mCherry* - magenta) retinæ at 7dpi. Image to right has *isl2b:GFP* channel removed. **(B)** Flat-mount images showing pSTAT3 staining on *isl2b:GFP*;*mpeg1:mCherry* (*isl2b:GFP* – green; *mpeg1:mCherry* - magenta) retinæ after P6/PLX3397 treatment or housed in system water and injected with 0.05% DMSO (Ctr) at 7dpi. Image to right has *isl2b:GFP* channel removed. Scale bar=50 μ m.

Table S1. Sequences of q-PCR primers

[Click here to download Table S1](#)

Table S2. 56 up-regulated genes at 24hpi (filtering parameters used: the maximum of the average group RPKM value >1.5, fold change >2, false discovery rate (FDR) p-value <0.05. Genes with TPM=0 in one or more replicates were excluded.

[Click here to download Table S2](#)

Table S3. 252 down-regulated genes at 24hpi (filtering parameters used: the maximum of the average group RPKM value >1.5, fold change >2, false discovery rate (FDR) p-value <0.05. Genes with TPM=0 in one or more replicates were excluded.

[Click here to download Table S3](#)

Table S4. 5 up-regulated genes and 3 down-regulated genes at 12hpi (filtering parameters used: the maximum of the average group RPKM value >1.5, absolute fold change >2, false discovery rate (FDR) p-value <0.05. Genes with TPM=0 in one or more replicates were excluded.)

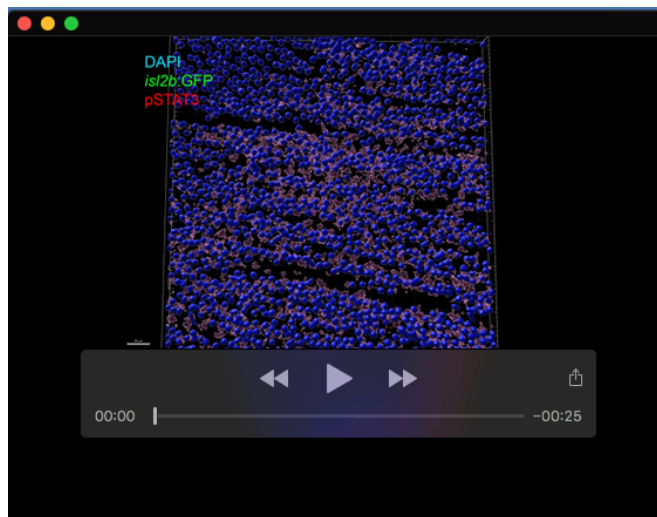
[Click here to download Table S4](#)

Table S5. 49 up-regulated genes at 12hpi (filtering parameters used: the maximum of the average group RPKM value >1.5, fold change >2, p-value <0.05. Genes with TPM=0 in one or more replicates were excluded.

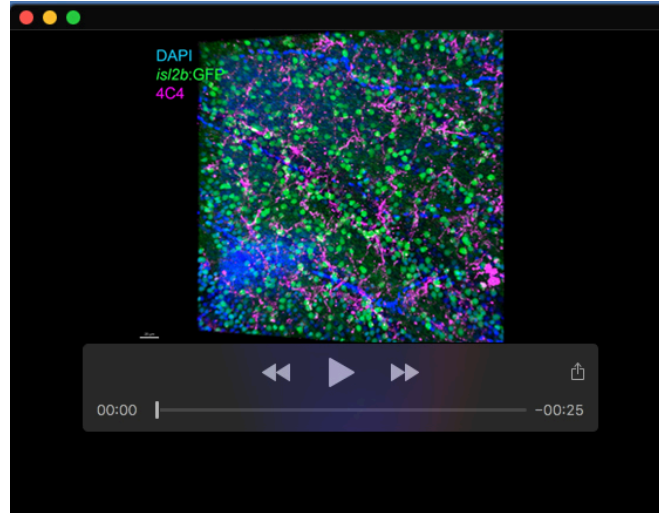
[Click here to download Table S5](#)

Table S6. 31 down-regulated genes at 12hpi (filtering parameters used: the maximum of the average group RPKM value >1.5 , fold change < -2 , p-value <0.05 . Genes with TPM=0 in one or more replicates were excluded.

[Click here to download Table S6](#)



Movie 1. Surfaces of pStat3 staining generated in Imaris. Scale bar = 20 μ m.



Movie 2. 3D image of 4C4 staining generated in Imaris. Scale bar = 20 μ m.