Microglial trogocytosis and the complement system regulate axonal pruning \textit{in vivo} 

Tony K.Y. Lim\textsuperscript{1} and Edward S. Ruthazer\textsuperscript{1,2,*}

1. Department of Neurology & Neurosurgery, Montreal Neurological Institute-Hospital, McGill University, Montreal, Quebec, H3A 2B4; Canada
2. Lead Contact

*Correspondence: edward.ruthazer@mcgill.ca
Abstract

Partial phagocytosis—called trogocytosis—of axons by microglia has been documented in \textit{ex vivo} preparations but has not been directly observed \textit{in vivo}. The mechanisms that modulate microglial trogocytosis of axons and its function in neural circuit development remain poorly understood. Here we directly observe axon trogocytosis by microglia \textit{in vivo} in the developing \textit{Xenopus laevis} retinotectal circuit. We show that microglia regulate pruning of retinal ganglion cell axons and are important for proper behavioral response to dark and bright looming stimuli. We identify amphibian regulator of complement activation 3, a homolog of human CD46, as a neuronally-expressed synapse-associated complement inhibitory molecule that inhibits trogocytosis and axonal pruning. Using a membrane-bound complement C3 fusion protein, we demonstrate that enhancing complement activity enhances axonal pruning. Our results support the model that microglia remodel axons via trogocytosis and that neurons can control this process through expression of complement inhibitory proteins.
Introduction

Microglia, the immune cells of the central nervous system (CNS), are vital for the maintenance and development of a healthy brain. Constantly surveilling the brain (Nimmerjahn et al., 2005; Wake et al., 2009), these highly phagocytic cells are thought to contribute to developmental synaptic remodeling by phagocytosing inappropriate or supernumerary synapses, a hypothesis that has derived considerable support from histological and immunohistochemical evidence identifying synaptic components within endocytic compartments in microglia (Paolicelli et al., 2011; Schafer et al., 2012; Stevens et al., 2007; Tremblay et al., 2010). This hypothesis is further supported by numerous studies demonstrating that microglia depletion leads to exuberant axonal outgrowth (Pont-Lezica et al., 2014; Squarzoni et al., 2014), impaired pruning of excess synapses (Ji et al., 2013; Milinkeviciute et al., 2019) and increased spine density (Wallace et al., 2020) during development.

The mechanisms for how microglia shape circuits by engulfing synapses remain poorly understood, and direct evidence of complete elimination of synapses by microglial engulfment remains elusive. Instead of removing entire synapses, microglia have been documented engaging in trogocytosis, or partial elimination, of axons and presynaptic boutons in ex vivo organotypic cultures (Weinhard et al., 2018). However, it remains to be seen whether microglial trogocytosis of axons is a phenomenon that occurs in vivo.

Even if we accept the hypothesis that microglia trogocytose the axonal compartment, many questions remain. What impact does partial elimination of presynaptic structures have on circuit remodeling? It is unclear if this phenomenon is required for proper wiring of neuronal circuits. Does microglial trogocytosis of axons affect axon morphology? While disrupting microglial function enhances axon tract outgrowth (Pont-Lezica et al., 2014; Squarzoni et al., 2014), it is unknown if this result is because of a disruption in microglial trogocytosis, or whether non-phagocytic mechanisms are in play. Is axonal trogocytosis by microglia mechanistically similar to complement-mediated synaptic pruning? There is extensive evidence demonstrating that the complement system regulates synaptic pruning by microglia via the complement protein C3 (Paolicelli et al., 2011; Schafer et al., 2012; Stevens et al., 2007). However, knockout (KO) mice lacking complement receptor type 3 (CR3), the receptor for activated C3, do not exhibit a deficit in microglial trogocytosis (Weinhard et al., 2018), raising the possibility that microglial trogocytosis of axons is mechanistically distinct from complement-mediated synaptic pruning.

In this study, we addressed these questions and directly observed in vivo trogocytosis of retinal ganglion cell (RGC) axons by individual microglial cells in real-time using the developing Xenopus laevis retinotectal circuit. We then developed an assay to monitor microglial trogocytosis of axons among the population of microglia in the optic tectum. To investigate the functional role of microglial trogocytosis, we depleted microglial cells, and found that this enhanced axon arborization and reversed the behavioral responses to dark and bright looming stimuli. We identified amphibian regulator of complement activation 3 (aRCA3) (Oshiumi et al., 2009), a homolog of mammalian CD46, as an endogenously expressed, synapse-associated, complement inhibitory molecule in Xenopus laevis RGC neurons. Overexpression of aRCA3 inhibited trogocytosis and axonal pruning. Conversely, expression of a membrane-bound complement C3 fusion protein in RGCs enhanced axonal pruning. Our findings provide direct in vivo evidence supporting the hypothesis that microglia trogocytose presynaptic axonal elements and supports a model in which microglial trogocytosis regulates axonal pruning to promote...
proper neural wiring during development (Schafer et al., 2012; Weinhard et al., 2018). In this model, neurons exert local control of microglial trogocytosis and axonal pruning by expressing complement regulatory proteins.

**Results**

**Microglia in *Xenopus laevis* tadpoles resemble neonatal mammalian microglia**

Microglia in *Xenopus laevis* tadpoles were labeled with IB4-isoeectin conjugated fluorophores for *in vivo* imaging (Figure S1A). Like microglia in neonatal mammalian models (Dalmau et al., 1997; Smolders et al., 2017), these labeled cells are highly mobile (Movie S1A and Figure S1C), are morphologically dynamic with ameboid-like and primitive ramified-like morphologies (Movie S1B and Figure S1B), and respond to tissue injury by surrounding the damaged zone and removing injured tissue (Movie S2 and Figure S1D).

In developing zebrafish larvae, microglia localize to the cell body layer of the optic tectum and are excluded from the tectal neuropil (Svahn et al., 2013). Conversely, microglia in developing mammalian models are found in neuropil regions (Dalmau et al., 1997; Hoshiko et al., 2012; Tremblay et al., 2010). In the *Xenopus laevis* retinotectal circuit, RGC axons project to the neuropil of the contralateral optic tectum, where they synapse on tectal neurons (Figure 1A). To examine whether Xenopus microglia interact with the tectal neuropil, the neuropil was labeled by bulk electroporation of RGC neurons with a plasmid encoding pH-resistant green fluorescent protein (pHtdGFP). *In vivo* live imaging revealed that, similar to what is observed in mammals, microglia in developing *Xenopus laevis* can be found in both the cell body layer and the neuropil region (Movie S3A). Microglia move in and out of the neuropil region from the cell body layer (Figure 1B and Movie S3B) and move freely through the neuropil (Figure 1C and Movie S3B). Additionally, microglia extend processes into the neuropil to contact axons, with interactions ranging from minutes to hours in duration (Figure 1D and Movie S3C).

**In vivo imaging of RGC neurons reveals microglial trogocytosis of axons and presynaptic structures**

We then sought to examine whether microglia cells engage in trogocytosis of RGC axons in the developing *Xenopus laevis* retinotectal circuit. As endosomal organelles are typically acidified (Casey et al., 2010), pH-stability of dyes and fluorescent proteins is an important consideration when performing live imaging of trogocytosis (Shinoda et al., 2018). To reduce quenching of fluorescence, we used pHtdGFP (pKa = 4.8) which is more pH-stable than EGFP (pKa = 6.15) (Roberts et al., 2016). We expressed pHtdGFP in RGC axons by electroporation in the eye, and labeled microglia with Alexa 594-conjugated IB4-isoeectin. 2-photon live imaging revealed that the amount of green fluorescence associated with individual microglial cells increased following interactions with pHtdGFP-labeled axons (Movies S4A and S4B). In the example shown in Figure 1E, the green fluorescence in the microglial cell increased 3-fold following interaction with a pHtdGFP-labeled axon (Figure 1F). The real-time increase in microglial green fluorescence suggests direct transfer of fluorescent protein from the pHtdGFP-labeled axon and provides direct *in vivo* evidence of microglial trogocytosis of the presynaptic RGC axon.

Because microglia in developing *Xenopus laevis* tadpoles are highly mobile, the possibility that they may leave the imaging field complicates measuring trogocytosis of axons by individual microglia. To quantify microglial trogocytosis, we instead took the approach of measuring microglial green fluorescence across the population of microglia within the optic tectum. A greater number of pHtdGFP-
Figure 1. Microglia surveil the tectal neuropil, contact RGC axons, and increase in green fluorescence following an interaction with pHtdGFP-labeled axons in real-time

(A) A diagram of the developing *Xenopus laevis* retinotectal circuit.

(B) The tectal neuropil does not exclude microglia in *Xenopus laevis*. The yellow dotted line indicates the border of the cell body layer and the neuropil region. Microglia (red) can be observed migrating in and out of the neuropil region from the cell body layer. A single registered optical section is shown.

(C) Microglia surveil the neuropil. A microglial cell is followed over time as it traversed different depths in the tectum.

(D) Microglia extend processes (white arrows) into the neuropil. Contact duration varied between minutes and hours.

(E) A microglial cell (blue arrow) interacts with a pHtdGFP-labeled RGC axon and increases in green fluorescence in real-time. The colocalization of green and red is colorized as white.

(F) The fluorescence change within the interacting microglial cell shown in 1E (blue line), and a non-interacting microglial cell (red line).
labeled axons in the optic tectum is expected to lead to more frequent opportunities for trogocytotic interactions between microglia and pHtdGFP-labeled axons, resulting in greater amounts of green fluorescent material found within the microglial population. Based on this principle, we developed an assay to measure microglial trogocytosis of RGC axons in *Xenopus laevis* tadpoles. At developmental stage 39/40, RGC axons were labeled by retinal electroporation with plasmid encoding pHtdGFP, and microglia were labeled by intraventricular injection of Alexa 594-conjugated IB4-isoelectin (Figure 2A). By two days post-labeling, axons begin expressing pHtdGFP and innervate the optic tectum. This also corresponds to the period when microglia begin to extensively colonize the optic tectum. At day 4 and day 5 post-labeling, the optic tectum was imaged by 2-photon microscopy. The number of pHtdGFP-labeled axons present in the optic tectum was counted and the green fluorescence within the population of microglia in the imaging field was quantified using 3D masking with the IB4-isoelectin channel (Figure 2B). To control for the possibility of RGC apoptosis, data were excluded if axonal blebbing was observed, or if the number of axons decreased from day 4 to day 5. Even when electroporation yields no labeled axons, some basal green fluorescence in microglial cells is still observed (Figure 2C). This is because microglia have high levels of autofluorescent molecules such as lipofuscin, bilirubin, and porphyrins (Mitchell et al., 2010). At day 4, there is a weak positive correlation between microglial green fluorescence and the number of pHtdGFP-labeled axons in the optic tectum, a relationship which is significantly strengthened on day 5 (Figure 2D). The change in green fluorescence associated with microglia from day 4 to day 5 correlates with the number of pHtdGFP-labeled axons in the optic tectum (Figure S2A), suggesting that microglial cells accumulate pHtdGFP from intact axons between day 4 and day 5 post-labeling.

To determine if synaptic material is being trogocytosed by microglia, we generated a synaptophysin-pHtdGFP fusion protein (SYP-pHtdGFP). SYP is a presynaptic vesicle protein (Valtorta et al., 2004), and SYP fusion proteins are commonly used as synaptic vesicle markers (Nakata et al., 1998; Ruthazer et al., 2006). Expressing SYP-pHtdGFP in RGC neurons yielded axons with pHtdGFP puncta along the length of their terminal arbors (Figure 2E). When SYP-pHtdGFP is expressed in RGC axons, on day 4, no correlation is observed between microglial green fluorescence and the number of SYP-pHtdGFP-labeled axons in the optic tectum. However, by day 5, a correlation between microglial green fluorescence and the number of SYP-pHtdGFP-labeled axons is observed (Figure 2F). The change in microglial green fluorescence from day 4 to day 5 is proportionate to the number of SYP-pHtdGFP-labeled axons in the optic tectum (Figure S2B), suggesting that microglial cells trogocytose and accumulate presynaptic elements from RGC axons over the period from day 4 to day 5.

**Depletion of microglial cells by colony stimulating factor 1 receptor (CSF1R) antagonism enhances RGC axon branching and reverses the profile of behavioral responses to dark and bright looming stimuli**

To assess the functional roles of microglial trogocytosis, we first depleted microglia using PLX5622. PLX5622 is an inhibitor of CSF1R, which is a tyrosine kinase receptor essential for microglia survival (Elmore et al., 2014; Erblich et al., 2011). Animals reared in 10 μM PLX5622 had significantly reduced microglia numbers in the optic tectum compared to vehicle-treated animals (Figure 3A and 3B). Morphological analysis of surviving microglia also revealed a reduction in the number of processes per microglial cell, indicating a less ramified morphology (Figure 3C).

Next, we interrogated the effect of microglial depletion on the morphology of single RGC axons. Axons were followed for several days in control and microglia-depleted animals (Figure 3D). Microglial
A

| Day 0 | Day 1 | Day 2 | Day 3 | Day 4 | Day 5 |
|-------|-------|-------|-------|-------|-------|
| Stage 39/40 | Stage 46 | Stage 47 |

RGC electroporation with pHtdGFP and labeling of microglia with IB4-isolectin

Labeled axons innervate optic tectum

Microglia colonize the optic tectum

in vivo imaging

B

Data Acquisition:
2-photon 830 nm

3D masks of microglia

Average microglial green fluorescence

IB4-isolectin Alexa 594
pHtdGFP

C

n = 16 ns
n = 12 ***
**n = 11 ***
**n = 8 ****

Microglial green fluorescence (arbitrary unit)

D

n = 47

Microglial green fluorescence (arbitrary unit)

# pHtdGFP-labeled axons

# pHtdGFP-labeled axons

E

IB4-isolectin Alexa 594
SYP-pHtdGFP

F

n = 57

Microglial green fluorescence (arbitrary unit)

# SYP-pHtdGFP labeled axons

# SYP-pHtdGFP labeled axons
Figure 2. Microglia accumulate green fluorescence label from axons expressing pHtdGFP or SYP-pHtdGFP
(A) Timeline of trogocytosis assay. Axons were labeled with pH-stable GFP (pHtdGFP). 2-photon imaging was performed on day 4 and day 5 post-labeling.
(B) Measurement of green fluorescence signal from microglia. pHtdGFP-labeled axons (green) were imaged concurrently with microglia (magenta). 3D microglia ROIs were automatically generated (magenta outlines). The average microglial green fluorescence was quantified from the population of microglia sampled in the z-stack.
(C) Presence of pHtdGFP-labeled axons in the optic tectum increases microglial green fluorescence between day 4 and day 5. 2-way RM ANOVA interaction F(3,43) = 6.14, p = 0.0014. Sidak’s multiple comparison test ***p < 0.001, ****p < 0.0001.
(D) Microglial green fluorescence weakly correlates with the number of pHtdGFP-labeled axons in the optic tectum on day 4 (n = 47, Pearson r = 0.30, p = 0.038) and moderately correlates on day 5 (n = 47, Pearson r = 0.52, p = 0.0002). The same dataset is analyzed in Figure 2C.
(E) SYP-pHtdGFP fusion protein localizes pHtdGFP to presynaptic puncta.
(F) Microglial green fluorescence weakly correlates with the number of SYP-pHtdGFP-labeled axons in the optic tectum on day 4 (n = 57, Pearson r = 0.063, p = 0.64), and weakly correlates on day 5 (n = 57, Pearson r = 0.34, p = 0.010).
We then sought to delineate the functional effects of microglial depletion on the development of the retinotectal circuit. Previous reports in Xenopus laevis and in zebrafish have shown that the retinotectal circuit is a vital processing and decision-making center for the visual detection of looming objects (Dong et al., 2009; Khakhalin et al., 2014). Therefore, we developed a free-swimming looming stimulus assay in Xenopus tadpoles (Figure 4A) to determine the functional outcomes of disrupting microglial function in this circuit. Tadpole behavioral responses to dark looming stimuli or bright looming stimuli (Figure 4B) were recorded and custom computer vision software was used to track the locomotor response of the tadpole (Figure 4C).
Figure 4D shows contrails from representative animals in response to dark looming stimuli. In vehicle-treated animals, dark looming stimuli evoked stereotypical defensive escape behavior (Movie S5A), whereas microglia-depleted animals were less likely to make defensive responses to dark looming stimuli (Movie S5B). Presentation of dark looming stimuli elicited an increase in velocity in control animals (Figure 4E), with a peak in instantaneous velocity at 1.1 ± 0.1 s (mean ± SD, n = 8) post-stimulus. The distance traveled by control tadpoles over the 3 s immediately following presentation of a dark looming stimulus increased compared to the period before the stimulus was presented (Figure 4F). This increase was absent in microglia-depleted animals. We categorized the tadpole responses to looming stimuli as exhibiting defensive behavior, absence of defensive behavior or undeterminable (excluded from response rate calculation). Microglia-depletion significantly reduced the response rate to dark looming stimuli (Figure 4G).
Figure 4H shows contrails from representative animals in response to bright looming stimuli. Bright looming stimuli rarely evoked defensive escape behavior in control animals (Movie S6A) but often evoked robust responses in microglia-depleted animals (Movie S6B). Presentation of bright looming stimuli elicited an increase in velocity in microglia-depleted animals (Figure 4G), with a peak in instantaneous velocity at 1.9 ± 0.6 s (mean ± SD, n = 8) after the onset of stimulus presentation. The distance traveled by microglia-depleted tadpoles over the 3 s immediately following presentation of a bright looming stimulus increased compared to the period before the stimulus was presented (Figure 4J). This increase was absent in vehicle-treated animals. Furthermore, microglia-depletion significantly increased the response rate to bright looming stimuli (Figure 4K).
In trials where animals exhibited escape behavior to dark or bright looming stimuli, microglial depletion did not significantly alter escape distance (Figure S3A and Figure S3B), maximum escape velocity (Figure S3C and Figure S3D), or escape angle (Figure S3E and Figure S3F), suggesting that microglial depletion did not disrupt motor function.

Figure 3. CSF1R antagonism depletes microglia from the optic tectum and increases axon branch number
(A) Animals were treated with vehicle or 10 μM PLX5622. Brain structures and microglia were labeled using CellTracker Green BODIPY (green) and IB4-isolectin (red), respectively. The white dotted line indicates the border of the optic tectum. Single optical sections are shown.
(B) PX5622 depletes microglia in the optic tectum. Mixed-effects REML model interaction F(3,39) = 14.23, p < 0.0001. Sidak’s multiple comparison post-hoc test **** p < 0.0001. Data are shown as mean ± SEM.
(C) PLX5622 reduces the number of processes per microglia. Mixed-effects REML model main effect F(1,14)=18.42, *** p < 0.001. Data are shown as mean ± SEM.
(D) Monitoring of individual RGC axons in vehicle control and PLX5622-treated animals.
(E) Microglial depletion with PLX5622 did not affect axon arbor length. 2-way RM ANOVA main effect F(1,24)=0.4141, p = 0.53. Data are shown as mean ± SEM.
(F) Microglial depletion with PLX5622 increased axon branch number. 2-way RM ANOVA main effect F(1,24)=5.581, p = 0.027. Data are shown as mean ± SEM.
Figure 4. Microglial depletion reverses the expected behavioral response to both dark and bright looming stimuli

(A) Schematic of a looming behavioral task to assess visuomotor responses in *Xenopus laevis* tadpoles. Stage 47 animals were presented looming stimuli and free-swimming escape responses were recorded.
Bioinformatic identification of amphibian regulator of complement activation 3 (aRCA3), a neuronally-expressed membrane-bound complement inhibitory molecule and homolog of human CD46

Microglial depletion enhanced axonal arborization and perturbed functional development of the retinotectal circuit. As microglia are known to secrete trophic factors and cytokines (Parkhurst et al., 2013; Solek et al., 2018), signaling mechanisms unrelated to trogocytosis may be at work. To better discern a role of trogocytosis in modulating axon morphology, we searched for a molecule that would inhibit axon trogocytosis. Because complement C3 is a known mediator of microglial phagocytosis (Brown and Neher, 2014) and has been proposed to accumulate at synapses and tag them for removal by microglial phagocytosis (Schafer et al., 2012; Stevens et al., 2007), we carried out a bioinformatics screen to identify neuronally expressed proteins that inhibit complement C3.

To search for proteins that inhibit complement C3, we first used the STRING Protein-Protein Association Network to identify proteins which interact with human complement C3 (Figure 5A). To select for complement inhibitory proteins, our search was then refined by focusing on proteins containing CCP (domains abundant in complement control proteins) motifs, which are highly conserved modules abundant in proteins responsible for negative regulation of the complement system (Norman et al., 1991). The top CCP-containing proteins identified by STRING were queried on the Allen Brain Institute human multiple cortical areas RNA-seq dataset (Allen Institute for Brain Science, 2015; Hodge et al., 2019). Of the CCP-containing proteins identified by STRING, only CD46 is highly expressed by human CNS neurons (Figure 5B).

Using NCBI protein-protein BLAST, the *Xenopus laevis* protein with the highest similarity to human CD46 was identified as amphibian Regulator of Complement Activation 3 (aRCA3) (Table 1). aRCA3 is a protein uncharacterized in *Xenopus laevis* but characterized as a membrane-associated complement regulatory protein in *Xenopus tropicalis* (Oshima et al., 2009). Like human CD46, *Xenopus laevis* aRCA3 is a type I transmembrane protein with many extracellular CCP domains and a stop-transfer anchor sequence near the C-terminus of the protein (Figure 5C). As a comparison, the second and third most similar *Xenopus laevis* proteins, complement receptor 2 and complement component 4 binding protein, do not share a protein architecture with CD46. The tertiary structure of *Xenopus laevis* aRCA3...
Table 1. Top 3 most similar proteins to human CD46 in the *Xenopus laevis* genome

| Protein name                                      | Query Coverage | Percent Identity | E-value   |
|--------------------------------------------------|----------------|-----------------|-----------|
| 1. Amphibian Regulator of Complement Activation 3 (aRCA3) | 64%            | 34.77%          | $3 \times 10^{-41}$ |
| 2. Complement Receptor Type 2                     | 70%            | 31.77%          | $4 \times 10^{-35}$ |
| 3. Complement Component 4 Binding Protein          | 76%            | 31.17%          | $4 \times 10^{-35}$ |

resembles that of human CD46 (Movie S7A and Movie S7B). Despite having twice the number of CCP domains as human CD46, aRCA3 is a likely homolog of CD46. The number of CCP domains of CD46 homologs varies across species—for example, the *Gallus gallus* homolog (Cremp) has 6 CCP domains, while homologs in *Equus caballus* (MCP), *Danio rerio* (rca2.1), and *Coturnix japonica* (MCP-like) each have 5 CCP domains.

To investigate whether RGC neurons endogenously express aRCA3, retinal sections were probed by fluorescence RNAscope *in situ* hybridization (Wang et al., 2012). Probes against aRCA3, the positive control transcript RNA polymerase II subunit A (polr2a.L), and the negative control transcript bacterial dihydrodipicolinate reductase (DapB) were hybridized on PFA fixed retinal sections (Figure 5D). As expected, DapB transcripts were not detected in the retina, whereas the housekeeping polr2a transcript was ubiquitously expressed. In contrast, aRCA3 transcripts localize to the ganglion cell layer (Figure 5E), revealing that aRCA3 may be endogenously expressed in RGC neurons.

**aRCA3 associates with synapses, enhances axonal arborization and inhibits microglial trogocytosis**

It is unknown whether aRCA3 localizes to the correct cellular compartments to protect axons from complement attack. To investigate this, we examined the localization of aRCA3 by tagging it with mCherry (Figure 6A). This aRCA3-mCherry fusion protein was expressed together with SYP-pHtdGFP, which localizes to synaptic puncta (Figure 2D). SYP-pHtdGFP and aRCA3-mCherry colocalize when co-expressed in the same axon (Figure 6B). The high degree of colocalization between SYP-pHtdGFP and aRCA3-mCherry suggests that aRCA3 distributes to similar subcellular compartments as the presynaptic marker SYP (Figure 6C).

Next, we examined the effect of aRCA3 overexpression on microglial trogocytosis. pHtdGFP was expressed in RGC neurons with or without overexpression of aRCA3, and microglial trogocytosis was quantified by measuring microglial green fluorescence on day 4 and day 5 post-labeling. Expression of pHtdGFP in RGC neurons resulted in a correlation between microglial green fluorescence and the number of pHtdGFP-labeled axons on day 5 post-labeling (Figure 6D), a relationship that was not observed when aRCA3 was overexpressed (Figure 6E). Overexpression of aRCA3 significantly reduced the correlation coefficient between the change in microglial green fluorescence from day 4 to day 5 and the number of pHtdGFP-labeled axons (Figure S4), suggesting that overexpression of aRCA3 reduces microglial trogocytosis.

We then examined the effect of aRCA3 overexpression on axon morphology in single RGC axons over several days (Figure 6F). Overexpression of aRCA3 resulted in significantly greater axon arbor sizes compared to control axons (Figure 6G). Similarly, axon branch number was greater in axons overexpressing aRCA3 compared to control axons (Figure 6H). These results show that aRCA3...
Figure 5. Identification of a neuronally expressed membrane-bound complement inhibitory protein, amphibian regulator of complement activation 3 (aRCA3), the predicted homolog of human CD46.

(A) Complement C3 was queried on the STRING Protein-Protein Association Network and the top 30 interaction partners are shown in the network diagram. The predicted homolog of CD46 in Xenopus laevis (aRCA3) was identified as LOC108708165.
displayed. Red nodes represent proteins which contain complement control domains that inhibit complement activity. Line thickness indicates the strength of data supporting protein interaction.

(B) Complement inhibitory proteins identified by STRING were screened for neuronal expression in the Allen Brain Map human cortical transcriptomics dataset. Cell type taxonomy and hierarchical clustering was determined according to previous analysis (Hodge et al., 2019). Only CD46 is highly expressed by neurons. Heat map color scale denotes log 2 expression levels as represented by trimmed mean (25%-75%). CPM = counts per million.

(C) Protein architecture of CD46 and the most similar *Xenopus laevis* homolog (aRCA3). Both human CD46 and *Xenopus laevis* aRCA3 are type I transmembrane proteins, with a non-cytoplasmic region that contains many complement control protein modules, and a transmembrane anchor near the C-terminus.

(D) Fluorescent RNAscope *in situ* hybridization on retina sections shows that aRCA3 is endogenously expressed in the retina. Negative control probe DapB was not detected in the retina. Housekeeping gene Polr2a.L was expressed ubiquitously.

(E) High magnification of the highlighted region in Figure 5D. aRCA3 is highly expressed in the GC

overexpression causes axons to have larger more highly branched arbors.

We then examined the effect of aRCA3 overexpression on axon morphology in single RGC axons over several days (Figure 6F). Overexpression of aRCA3 resulted in significantly greater axon arbor sizes compared to control axons (Figure 6G). Similarly, axon branch number was greater in axons overexpressing aRCA3 compared to control axons (Figure 6H). These results show that aRCA3 overexpression causes axons to have larger more highly branched arbors.

While CD46 is best known for its ability to inactivate complement C3 and complement C4, CD46 can also signal through intracellular tyrosine kinase activity under certain conditions (Riley-Vargas et al., 2004). This raises the potential caveat that the increased axonal arborization induced by overexpression of aRCA3 may result from aberrant intracellular signaling. Using a bioinformatic tool, NetPhos3.1 with cutoff scores > 0.6 (Blom et al., 2004), we did not find a predicted tyrosine kinase phosphorylation site on the cytoplasmic region of aRCA3. Nonetheless, it is possible that aRCA3 may exert some of its effects through an intracellular signaling pathway. To provide further functional validation that the complement system affects axonal arborization, we examined the effects of enhancing complement activity on axon morphology.

**Expression of a membrane-bound complement C3 fusion protein reduces RGC axon size and branching**

If aRCA3 affects axon morphology by inhibiting complement activity, enhancing complement activity on single axons should produce effects opposite to that of aRCA3 overexpression. To explore this possibility, we designed an axon surface-localized complement C3 fusion protein to enhance complement activity on individual axons (Figure 7A). Synaptobrevin, also known as vesicle-associated membrane protein 2 (VAMP2), is concentrated in synaptic vesicles, though a significant fraction of VAMP2 is also present on the axon surface (Ahmari et al., 2000; Sankaranarayanan and Ryan, 2000). We cloned *Xenopus laevis* complement C3 and fused it to the extracellular C-terminus of *Xenopus laevis* VAMP2. Thus, expression of this VAMP2-C3 construct in RGC neurons results in axons tagged with extracellular membrane-bound complement C3. While we used complement C3 precursor to generate the VAMP2-C3 fusion protein, complement C3 undergoes spontaneous, low-level activation through the alternative complement pathway (Pangburn et al., 1981).

VAMP2-C3 was co-expressed with pHtdGFP in RGC neurons, and axons were monitored over several days (Figure 7B). To control for the possibility that VAMP2 overexpression may affect axon morphology, we also overexpressed VAMP2 alone in RGC axons. Expression of VAMP2-C3 in RGC
Figure 6. RGC overexpression of synapse-associated aRCA3 inhibits microglial trogocytosis and increases axon arborization

(A) An aRCA3-mCherry fusion protein was co-expressed with SYP-pHtdGFP in RGC axons.

(B) Representative RGC axon expressing aRCA3-mCherry and SYP-pHtdGFP. SYP-pHtdGFP is concentrated at synaptic puncta.

(C) Colocalization analysis of the axon shown in B. Scatterplot of pixel intensities shows a high degree of association between aRCA3-mCherry and SYP-pHtdGFP. After applying Costes threshold, Pearson $r = 0.90$.

(D) Microglial green fluorescence is not significantly correlated with the number of pHtdGFP-labeled axons in the optic tectum on day 4 ($n = 39$, Pearson $r = 0.23$, $p = 0.13$), and moderately correlated on day 5 ($n = 39$, Pearson $r = 0.58$, $p = 0.0001$).

(E) When aRCA3 is overexpressed, microglial green fluorescence is not significantly correlated with the number of pHtdGFP-labeled axons in the optic tectum on either day 4 ($n = 42$, Pearson $r = 0.0082$, $p = 0.96$) or on day 5 ($n = 42$, Pearson $r = -0.018$, $p = 0.96$).

(F) Individual control and aRCA3 overexpressing RGC axons were imaged over several days.

(G) aRCA3 overexpression increases axon arbor length. 2-way RM ANOVA interaction $F(3,102) = 5.43$, $p = 0.0017$. Sidak’s
Discussion

In vivo evidence of microglial trogocytosis of axons during healthy development

Previous experiments in ex vivo slice culture have shown that microglia trogocytose presynaptic elements (Weinhard et al., 2018). Our results now add in vivo support to the hypothesis that microglia engulf synaptic material via trogocytosis.

Numerous past studies have shown that during development microglia have synaptic components within phagocytic compartments (Paolicelli et al., 2011; Schafer et al., 2012; Stevens et al., 2007; Tremblay et al., 2010). However, apoptosis is a prominent feature of neural development (Nijhawan et al., 2000), and it is unclear whether the source of synaptic material within microglia is because of engulfment of apoptotic neuronal components or if it is due to engulfment of synaptic material from live neurons. When investigating trogocytosis, we minimized the effect of apoptosis by excluding measurements when the number of axons between imaging sessions decreased or when axonal blebbing was observed, suggesting that microglia accumulate fluorescent label from living axons.

Expression of complement inhibitory or complement enhancing molecules in neurons regulates axon morphology and trogocytosis by microglia

Activation of complement C3 exposes a reactive thioester bond that covalently attaches to amine or carbohydrate groups on cell surfaces (Sahu et al., 1994). Microglia express CR3, a receptor for complement C3 (Ling et al., 1990), and binding of CR3 to its ligand induces phagocytosis (Newman et al., 1985). Complement C3 localizes to synapses during development and tags synapses for removal by microglia (Stevens et al., 2007). The importance of complement C3 in synaptic pruning has been studied by disrupting the C3 pathway using C3 KO mice (Stevens et al., 2007), CR3 KO mice (Schafer et al., 2012), or recently, with the exogenous neuronal expression of the complement inhibitory proteins Crry (Werneburg et al., 2020) and CD55 (Wang et al., 2020). Here we show that enhancing the C3 pathway with a membrane-bound VAMP2-C3 fusion protein increases axonal pruning at the single axon level. Conversely, overexpression of an endogenous membrane-bound complement inhibitory molecule inhibits axonal pruning and trogocytosis (Figure 8A).

It is hypothesized that neurons endogenously express membrane-bound complement inhibitory molecules to protect synapses from phagocytosis (Stephan et al., 2012; Stevens et al., 2007), and our data support this hypothesis. Recently, sushi repeat protein X-linked 2 (SRPX2) has been identified as a complement C1 tagging (Cong et al., 2020). SRPX2 is a secreted protein, and while secreted complement inhibitory molecules can act via cell-autonomous mechanisms to protect the neuron that produced it, such molecules also have non-autonomous actions due to the ability to diffuse through the extracellular environment. Conversely, a membrane-bound complement inhibitory molecule acts solely through cell-
Figure 7. Expressing VAMP2-C3 fusion protein in RGC neurons reduces axon arbor size and branch number

(A) Complement C3 was fused to the C-terminus of VAMP2 to create a complement enhancing molecule which is targeted to the axon surface and synapses.

(B) Axons from control, VAMP2-C3 expressing, and VAMP2 overexpressing RGC neurons were imaged over several days.

(C) Expression of VAMP2-C3 reduces RGC axon arbor length compared to control and VAMP2 overexpression. 2-way RM ANOVA interaction F(6,75)=3.48, p = 0.0044. Sidak’s multiple comparison test * p < 0.05, ** p < 0.01, *** p < 0.001. Data are shown as mean ± SEM.

(D) Expression of VAMP2-C3 reduces RGC branch number compared to control and VAMP2 overexpression. 2-way RM ANOVA interaction F(6,75)=4.15, p = 0.0012. Sidak’s multiple comparison test * p < 0.05, *** p < 0.001, **** p < 0.0001. Data are shown as mean ± SEM.

autonomous mechanisms and allows for greater spatiotemporal control of local complement protection. While the identity of a membrane-bound complement inhibitory molecule endogenous to mammalian neurons remains elusive, in this study we characterized *Xenopus laevis* aRCA3, an endogenously expressed, synaptic vesicle-associated, complement inhibitory molecule. aRCA3 is the most similar amphibian homolog of human CD46. CD46 is a membrane-bound complement inhibitory molecule that cleaves activated complement C3 and C4 (Barilla-LaBarca et al., 2002). Using the Allen Brain Institute human multiple cortical areas RNA-seq dataset (Allen Institute for Brain Science, 2015), we report that CD46 transcripts are enriched in human neurons. Interestingly, CD46 associates directly with β1-integrins (Lozahic et al., 2000), an adhesion molecule present on neuronal surfaces (Neugebauer and Reichardt, 1991) and enriched in synaptosomes (Chan et al., 2003), suggesting that it too may be localized to axon surfaces and synapses. In our study we show that aRCA3 expression inhibits both axon trogocytosis by microglia and axonal pruning—we speculate that CD46 may perform similar functions in mammals. Excessive synaptic pruning is thought to be one of the underlying causes of schizophrenia (Sellgren CM et al., 2019), and three large-scale genetic susceptibility studies have identified the CD46
gene as a significant Schizophrenia-risk locus (Håvik et al., 2011; Kim et al., 2020; Ripke et al., 2014).

Clearly, the role of CD46 in neurodevelopment warrants further study.

Microglia actively suppress exuberant axonal arborization

Disrupting microglial function by depletion causes exuberant axonal innervation in prenatal models (Pont-Lezica et al., 2014; Squarzoni et al., 2014). However, depleting microglia also increases the number of neural progenitor cells (Cunningham et al., 2013). Therefore, it is unclear whether the exuberant axonal innervation that occurs following microglial depletion results from a deficit of microglial-mediated axonal pruning, or whether it is because of an increase in the overall number of axons. In our experiments, we examined the effect of microglial depletion on the morphology of single axons, showing that microglia depletion significantly increased axon branch number. Thus, our study supports the model that the increase in axonal innervation resulting from microglial depletion is because of increased arborization of individual axons (Figure 8B).

The observations that microglial depletion and overexpression of the complement inhibitor aRCA3 in RGC neurons enhanced axonal arborization suggests that both phenomena may act through a common mechanism. These results support the hypothesis that microglial trogocytosis suppresses both axon branching and axon growth through the removal of extra branches and axonal material. While this explanation is at odds with the observation that microglia-depletion did not increase axon branch size, it is important to note that with aRCA3 overexpression, a single axon gains a relative growth advantage over other axons—this is distinct from the case of microglial depletion, where all axons profit from the same increased growth. If all axons are growing larger at the same pace, competitive mechanisms (Gosse et al., 2008; Ruthazer et al., 2003) might be expected to constrain axon arbor sizes.

It is worth considering that there may be unintended side-effects of microglial depletion with PLX5622 and that many microglial functions beyond trogocytosis of axons may be altered. In our study, we found that CSF1R inhibition reduced process branching of remaining microglial cells, an observation that has also been seen in rodents (Elmore et al., 2014). Decreased process complexity is a sign of microglial activation (Karperien et al., 2013), and microglial activation is associated with microglial release of cytokines and other neuromodulatory molecules (Hanisch, 2002). While we did not study the effect of microglial depletion on cytokine and chemokine production in this study, past studies have shown that cytokines and chemokines in non-disease mouse models are not altered by PLX5622 (Reshef et al., 2017). Moreover, possible confound is that low expression of CSF1R has been reported in some neurons in mice hippocampus (Luo et al., 2013). It is possible that CSF1R is present in Xenopus laevis RGC neurons and we did not exclude the possibility that PLX5622 may be acting directly on RGC axons in our study.

Microglia contribute to proper wiring of the retinotectal circuit in developing Xenopus laevis

A feed-forward network drives visually evoked escape behavior in Xenopus tadpoles (Khakhalin et al., 2014). In the retina, photoreceptors act on bipolar cells, which act on RGCs. Some RGCs are tuned to detect looming (Dunn et al., 2016; Münch et al., 2009), and the population of RGCs that respond to dark looming stimuli are distinct from the population of RGCs that respond to bright looming stimuli (Temizer et al., 2015). In zebrafish, these distinct RGC populations project to different arborization fields in the optic tectum (Robles et al., 2014; Temizer et al., 2015) where the execution of looming computation and behavioral decision-making occurs (Barker and Baier, 2015; Fotowat and Gabbiani,
In highly predated animals such as crabs, zebrafish and mice (Oliva et al., 2007; Temizer et al., 2015; Yilmaz and Meister, 2013), dark looming stimuli—which signal imminent threat such as an oncoming object or predator—reliably induce escape responses, whereas bright looming stimuli—which may occur as the animal exits a tunnel or traverses its environment—are less effective.

Microglial depletion does not appear to alter motor functionality, suggesting that the hindbrain and motor circuitry remain intact. Instead, the effects of microglia depletion on looming-evoked escape behavior must occur further upstream in the retina, the optic tectum, or in the projections from the tectal neurons to the hindbrain. As microglia depletion induces exuberant RGC axonal arborization, one explanation for why microglia depletion disrupts defensive behavior to dark looming stimuli is that the dark looming-sensitive RGC axons cannot effectively wire with their tectal neuron counterparts, resulting in a misclassification of threatening visual stimuli when axonal pruning is disrupted. When microglia are depleted, axons form more errant connections, and dark looming-sensitive RGC axons may instead wire together with tectal neurons which are not associated with threat classification, reducing the probability that dark looming stimuli elicit escape responses. We predict that the reverse is true for bright looming-sensitive RGC axons, whereby microglia depletion results in increased likelihood of errant wiring between bright looming RGC axons and tectal neurons that classify threatening visual stimuli, leading to enhancement of escape behavior to bright looming stimuli (Figure 8C).

**Synaptic pruning and trogocytosis are two sides of the same coin: Axonal pruning**

Classically, synaptic pruning has been described as the engulfment and elimination of synapses, a phenomenon dependent on the complement pathway (Paolicelli et al., 2011; Perry and O’Connor, 2008; Schafer et al., 2012; Stevens et al., 2007). While no deficit in microglial trogocytosis has been observed in ex vivo cultures from CR3 KO mice (Weinhard et al., 2018), one interpretation for this result is that synaptic pruning and trogocytosis are unique phenomena mediated by different mechanisms. However, our study suggests that complement-mediated synaptic pruning and trogocytosis are mechanically related. We find that the complement system influences both axonal trogocytosis and axonal arborization, and it is possible that in CR3 KO mice, compensation through one of the other receptors for activated C3 occurs. For example, microglia express complement receptor type 4 (CR4) (Allen Institute for Brain Science, 2015; Hodge et al., 2019) which can bind activated complement C3 to induce phagocytosis (Ross et al., 1992). Our data supports the hypothesis that neurons control microglial-mediated circuit remodeling through the expression of endogenous membrane-bound complement inhibitory molecules to regulate microglial trogocytosis.
Figure 8. The complement system and microglia regulate axonal pruning at the single axon level

(A) Inhibiting complement activity through overexpression of a membrane-bound complement inhibitory molecule results in impaired axonal pruning; conversely, increasing complement activity through expression of a membrane-bound complement enhancing molecule results in excessive axonal pruning.

(B) During normal development, microglia actively trogocytose and prune axons. Disrupting microglial function via microglial depletion increases axonal arborization.

(C) Proposed model of the role of microglia in the wiring of the *Xenopus laevis* retinotectal circuit. During normal development, errant connections (crossed lines) are removed so that dark looming-sensitive RGC neurons wire primarily with tectal threat detection circuits. When microglia are depleted, errant connections (crossed lines) remain and bright looming-sensitive RGC neurons wire with tectal threat detection circuits.
### Materials and Methods

#### KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Bacterial and Virus Strains** | | |
| DH5α Competent cells | Invitrogen | Cat#18265017 |
| **Chemicals, Peptides, and Recombinant Proteins** | | |
| Isolectin GS-IB4 From *Griffonia simplicifolia*, Alexa Fluor 594 Conjugate | ThermoFisher Scientific | Cat#I21413 |
| CellTracker Green BODIPY | ThermoFisher Scientific | Cat#C2102 |
| RNAscope Probe: *Xenopus laevis* polr2a.L | Advanced Cell Diagnostics | Cat#580841 |
| RNAscope Probe: *Xenopus laevis* aRCA3 | Advanced Cell Diagnostics | Cat#806291 |
| PLX5622 | Plexxikon | N/A |
| Polyethylene glycol 400 | Sigma | SKU#P3265 |
| Poloxamer 407 | Sigma | SKU#16758 |
| D-α-Tocopherol polyethylene glycol 1000 succinate | Sigma | SKU#57668 |
| TRIzol Reagent | Invitrogen | Cat#15596-026 |
| Superscript IV First-Strand synthesis system | Invitrogen | Cat#18091050 |
| Phusion High-Fidelity DNA polymerase | ThermoFisher Scientific | Cat#F530S |
| Sylgard 184 silicone | Paisley | AVDC00184003 |
| Fisher Healthcare Tissue-Plus O.C.T Compound | Fisher Scientific | Cat#23730571 |
| **Critical Commercial Assays** | | |
| RNAscope Multiplex Fluorescent Reagent Kit v2 | Advanced Cell Diagnostics | Cat#323136 |
| **Deposited Data** | | |
| Allen Brain Map Human Multiple Cortical Areas SMART-seq data set | (Allen Institute for Brain Science, 2015) | portal.brain-map.org/atlases-and-data/rnaseq/human-multiple-cortical-areas-smart-seq |
| **Experimental Models: Organisms/Strains** | | |
| Albino *Xenopus laevis* (Xla.NXT-WT: Albino\textsuperscript{NXR}) | National Xenopus Resource | NXR_0082 |
| **Recombinant DNA** | | |
| pCMV eGFP | (Haas et al., 2001) | |
| pFA6a pH-tdGFP | (Roberts et al., 2016) | Addgene Plasmid #74322 |
| pEF1α pHtdGFP | This paper | |
| pEF1α SYN-pHtdGFP | This paper | |
| Construct                   | Source                                      |
|-----------------------------|---------------------------------------------|
| pEF1α aRCA3-Myc-P2A-pHtdGFP | This paper                                  |
| pEF1α aRCA3-mCherry-Myc-P2A-SYN-pHtdGFP | This paper                                      |
| pEF1α pHtdGFP-P2A-VAMP2     | This paper                                  |
| pEF1α pHtdGFP-P2A-Myc-VAMP2-C3 | This paper                                |

### Software and Algorithms

| Name                                | Website/Version                                      |
|-------------------------------------|------------------------------------------------------|
| NCBI protein-protein BLAST         | blast.ncbi.nlm.nih.gov (Altschul et al., 1990)       |
| STRING v11.0 protein-protein association networks | string-db.org (Szklarczyk et al., 2019) |
| SMART protein domain annotation resource 8.0 | smart.embl-heidelberg.de (Letunic and Bork, 2018) |
| Phobius transmembrane topology tool | phobius.sbc.su.se (Käll et al., 2007)               |
| pyMOL 2.4                           | pymol.org/2/ (Schrödinger, LLC.)                    |
| trRosetta                           | yanglab.nankai.edu.cn/trRosetta/ (Yang et al., 2020) |
| ProBLM                               | compbio.clemson.edu/sapp/problm_webserver/ (Kimmett et al., 2014) |
| NetPhos 3.1                          | www.cbs.dtu.dk/services/NetPhos/ (Blom et al., 2004) |
| R 4.0.0                              | www.r-project.org/ (R Core Team, 2018)              |
| RColorBrewer                         | cran.r-project.org/web/packages/RColorBrewer/RColorBrewer.pdf (Neuwirth, 2014) |
| ComplexHeatmap                       | github.com/jokergoo/ComplexHeatmap (Gu et al., 2016) |
| Bioconductor 3.11                    | bioconductor.org (Huber et al., 2015)               |
| Serial Cloner 2.6.1                  | serialbasics.free.fr/Serial_Cloner.html (SerialBasics) |
| Imaris 6                             | imaris.oxinst.com/products/imaris-for-cell-biologists (Oxford Instruments) |
| Fluoview 5.0                         | www.olympus-lifescience.com/en/support/downloads/ (Olympus) |
| ThorImage LS                         | www.thorlabs.com/newgrouppage9.cfm?objectgroup_id=9072#ad-image-0 (Thorlabs) |
| Leica LAS X                          | www.leica-microsystems.com/products/microscope-software/p/leica-las-x-ls/ (Leica) |
| FIJI ImageJ                          | imagej.net/Fiji (Schindelin et al., 2012)          |
| 3D Objects Counter plugin            | imagej.net/3D_Objects_Counter (Bolte and Cordelières, 2006) |
| 3D ROI Manager plugin                | imagejdocu.tudor.lu/plugin/stacks/3d_roi_manager/start (Ollion et al., 2013) |
| TrackMate plugin                     | imagej.net/TrackMate (Tinevez et al., 2017)        |
| Descriptor-based series registration plugin | imagej.net/SPIM_Registration_Method (Preibisch et al., 2010) |
| 3D hybrid median filter plugin | Christopher Philip Mauer and Vytas Bindokas | imagej.nih.gov/ij/plugins/hybrid3dmedian.html |
|-------------------------------|-------------------------------------------|-----------------------------------------------|
| ScatterJ plugin               | (Zeitvogel et al., 2016)                 | savannah.nongnu.org/projects/scatterj         |
| MATLAB                        | MathWorks                                 | www.mathworks.com/products/matlab.html        |
| CANDLE                        | (Coupé et al., 2012)                     | sites.google.com/site/pierrickcoupe/softwares/denoising-for-medical-imaging/multiphoton-filtering |
| Graphpad Prism 8              | GraphPad Software                        | www.graphpad.com/scientific-software/prism/   |
| Fisher r-to-z transformation calculator | VassarStats                | www.vassarstats.net/rdiff.html                |
| Cura 4                        | Ultimaker                                 | ultimaker.com/software/ultimaker-cura         |
| TinkerCAD                     | Autodesk                                  | www.tinkercad.com                             |
| Psychopy 3.0                  | (Peirce et al., 2019)                    | psychopy.org                                  |
| XenLoom (beta): Looming Stimulus Presentation and Tracking of Xenopus laevis tadpoles | This paper | github.com/tonkyylim/XenLoom_beta |
| **Other**                     |                                          |                                               |
| Estink 2000 lumens mini LED projector | Amazon.ca | ASIN#B07F7RT9XZ |
| 96 white 20 lb bond copy paper | Staples.ca | Item#380480 |
| Custom 3D printed mount for projector lens | This paper | www.thingiverse.com/thing:4335379 |
| Custom 3D printed stage for Xenopus laevis behavior | This paper | www.thingiverse.com/thing:4335395 |
| 8-inch soda lime 1500 ml culture dish | Carolina | Item#741006 |
| 60 mm petri dish              | FisherScientific                         | Cat#FB0875713A                                |
| PLA 1.75mm 3D printing filament | iPrint-3D | Transparent purple |
| Logitech C920 webcam           | Logitech                                 | Model# 960-000764                             |
| Webcam scissor arm clamp mount | Amazon.ca | ASIN#B01N77YBLU |
| Anycubic i3 mega 3D printer    | Amazon.ca                                 | ASIN#B07NY5T1LJ                               |
| Grass SD9 Square Pulse Stimulator | Grass-Telefactor | Model SD9 |
| Custom-built 2-photon microscope | (Ruthazer et al., 2006)                |                                               |
| Thorlabs multiphoton imaging system | Thorlabs                        |                                               |
| Ti:Sapphire femtosecond pulsed laser | Spectra-Physics | InSight X3 |

(37/37)
527 **Lead Contact and Materials Availability**

528 Plasmids generated in this study will be made available upon request. Further information and requests
529 for resources and reagents should be directed to the Lead Contact, Edward Ruthazer
530 (edward.ruthazer@mcgill.ca).

531 **Xenopus laevis tadpoles**

532 Adult albino *Xenopus laevis* frogs (NXR_0082) were maintained and bred at 18°C. Female frogs were
533 primed by injection of 50 IU pregnant mare serum gonadotropin (ProSpec-Tany TechnoGene Ltd., Ness-
534 Ziona, Isreal). After 3 days, male and primed female frogs were injected with 150 IU and 400 IU of
535 human chorionic gonadotropin (Sigma-Aldrich, Oakville, CA) into the dorsal lymph sac, respectively. The
536 injected male and female frogs were placed in isolated tanks for mating. Embryos were collected the
537 following day and maintained in Modified Barth’s Saline with HEPES (MBSH) in an incubator at 20°C with
538 LED illumination set to a 12h/12h day-night cycle and staged according to Nieuwkoop and Faber (NF)
539 developmental stages (Nieuwkoop and Faber, 1994). All experiments were conducted according to
540 protocols approved by The Animal Care Committee of the Montreal Neurological Institute and in
541 accordance with Canadian Council on Animal Care guidelines. *Xenopus laevis* sex cannot be determined
542 visually pre-metamorphosis, and thus the sex of experimental animals was unknown.

543 **Labeling of microglia, axons, and brain structures**

544 Stage 39/40 tadpoles were anesthetized with 0.02% MS-222 in 0.1X MBSH. For labeling of microglia,
545 tadpoles received an intracerebroventricular (icv) injection to the 3rd ventricle of 1 mg/ml Alexa 594
546 conjugated IB4-isolectin. A minimum of 48 hours was allowed to pass before live imaging studies were
547 commenced to allow for binding and update of IB4-isolectin by microglial cells. For concurrent labeling
548 of brain structures, 1 mM CellTracker Green BODIPY in 10% DMSO was injected icv after isolectin
549 injection. For labeling of RGC axons, electroporation of plasmid DNA into RGC cells was performed as
550 described previously (Ruthazer et al., 2006, 2013a). In brief, stage 39/40 tadpoles were anesthetized
551 with 0.02% MS-222 in 0.1X MBSH. A glass micropipette was back filled with endotoxin free maxi-prep
552 plasmid solution (2-3 µg/µl) and fast green to visualize the injection. The micropipette was advanced
553 into the eye, and DNA solution pressure injected. The micropipette was then withdrawn, and parallel
554 platinum electrodes were placed to bracket the eye. For bulk labeling of RGCs, a Grass Instruments SD9
555 electrical stimulator was used to apply 4-6 pulses of 2.4 ms duration at 36 V with a 3 µF capacitor was
556 placed in parallel to obtain exponential decay current pulses. For single-cell labeling of RGCs, 4-6 pulses
557 of 1.6 ms duration at 36 V was applied.

558 **Microglial depletion**

559 PLX5622 was dissolved in DMSO at 20 mM, aliquoted, and stored at -20°C. Thawed aliquots of PLX5622
560 were sonicated briefly for 1 minute before dilution in polyethylene glycol (PEG) 400. PEG 400 solution
561 was then further diluted in a solution of 0.1X MBSH containing non-ionic surfactants poloxamer 407 and
562 D-α-tocopherol polyethylene glycol 1000 succinate (TPGS) to form a mixed micelle drug delivery vehicle
563 (Guo et al., 2013). Final vehicle rearing solution consisted of 2.5% PEG 400, 0.04% poloxamer 407, 0.01%
564 TPGS, and 0.05% DMSO in 0.1X MBSH. Initially, animals were reared in 0.1X MBSH. At stage 35-40,
565 animals were transferred to vehicle rearing solution with or without 10 µM PLX5622 for rearing. Rearing
566 solutions were refreshed daily with newly prepared drug and vehicle solutions.
Two-photon live imaging

Two-photon live imaging of axons and microglia cells was performed as described previously (Ruthazer et al., 2013b). Tadpoles were placed in a Sylgard 184 silicone imaging chamber with their dorsal side facing up and were covered with a #1 thickness glass coverslip. 1 μm interval z-stacks of the optic tectum were collected on a custom-built 2-photon microscope (Olympus BX61WI with Olympus FV300 confocal scan head) outfitted with an Olympus 1.0 NA 60x water-immersion objective, or a Thorlabs multiphoton resonant scanner imaging system outfitted with an Olympus 1.0 NA 20x water-immersion objective. Excitation was produced using a Spectra-Physics InSight3X femtosecond pulsed laser. Images were collected at 512x512 pixels with Fluoview 5.0 or ThorImage LS.

For real-time imaging studies, stage 46-48 tadpoles were paralyzed by brief (2-8 min) immersion in freshly thawed 2 mM pancuronium bromide in 0.1X MBSH. Animals were then maintained in 0.1X MBSH for imaging. Z-stacks were collected at 6-minute intervals. For daily imaging studies, stage 43-47 tadpoles were anesthetized by immersion in 0.02% MS-222. Z-stacks were collected, and animals were returned to 0.1X MBSH rearing solution.

When imaging microglia (Alexa 594 conjugated IB4-isolectin) and RGC axons (eGFP or pHtdGFP) concurrently, excitation wavelength was set at 830 nm and a 565 nm emission dichroic was used in conjunction with green (500-550 nm) and red (584-676 nm) filters for fluorescence emission detection on separate photomultiplier tubes. When imaging microglia (Alexa 594 conjugated IB4-isolectin) and brain structures (CellTracker Green BODIPY), imaging was done sequentially. Alexa 594 was first imaged with excitation wavelength set to 810 nm and fluorescence emission detection through the red filter. Imaging of CellTracker Green BODIPY was then performed at excitation wavelength 710 nm and the fluorescence emission detected through the green filter. When imaging RGC axons alone (pHtdGFP), excitation wavelength was set to 910 nm and fluorescence emission detected through the green filter. When imaging aRCA3-mCherry-Myc and SYP-pHtdGFP, images were captured concurrently at wavelength 990 nm through the red and green filters, respectively.

Two-photon laser-induced injury

Laser irradiation injury was carried out by repeatedly scanning an approximately 10x10 μm region under high laser power at wavelength 710 nm for 10 seconds.

Microscopy image processing and analysis

For visualization purposes, the 3D object counter (Bolte and Cordelières, 2006) and 3D ROI manager (Ollion et al., 2013) plugins were used on z-stacks for segmentation and effacement of melanophores on the dorsal dermal surface of the animal before maximum intensity projection. Analysis was performed by an experimenter blind to treatment group.

Axon morphology analysis

Two-photon z-stacks were captured daily from 2 to 5 days post-electroporation. Z-stacks were denoised with CANDLE (Coupé et al., 2012), and were manually traced using Imaris 6 software.

Microglia quantification

The number of microglia and microglia process numbers were counted manually within 150 μm thick z-stacks of the optic tectum.
Microglia mobility

For microglial mobility experiments, microglia in the optic tectum were imaged over 2 hours at 6 minute intervals and manual tracking of microglia was carried out using the TrackMate plugin (Tinevez et al., 2017).

Real-time imaging of trogocytosis of axons by individual microglial cells

The microglia channel was denoised with the 3D hybrid median filter plugin, and the axon channel was denoised with CANDLE (Coupé et al., 2012). Time series were registered by descriptor based series registration (Preibisch et al., 2010). Microglia 3D region of interests (ROIs) were generated using the 3D object counter plugin (Bolte and Cordelières, 2006) using the same threshold across all time points. Green fluorescence intensity within microglia 3D ROIs was measured using the 3D ROI manager plugin (Ollion et al., 2013). Movies were generated using Imaris software or ImageJ.

Trogocytosis assay

Microglia were labeled with Alexa 594-conjugated isoelectin and axons with pHtdGFP or SYP-pHtdGFP as described above. On day 4 and day 5 post-labeling, 150 µM thick z-stacks were collected from the optic tectum. Laser power and photomultiplier tube voltage was kept constant on both days. The number of labeled axons was counted manually. In studies when axons were labeled with pHtdGFP, data were excluded if axonal blebbing was detected, or if the number of labeled axons fell from day 4 to day 5. This data exclusion step was not performed when axons were labeled with SYP-pHtdGFP. Using the red microglia channel, microglia 3D ROIs were automatically generated using the 3D object counter plugin (Bolte and Cordelières, 2006). Segmentation threshold was set to the mean voxel intensity of the red channel plus 2 standard deviations. A minimum threshold size of 1500 voxels was used to exclude background from analysis. The 3D ROI manager plugin (Ollion et al., 2013) was then used to measure the green fluorescence intensity of the microglial cell ROIs within the z-stack. To reduce background signal in the green channel, the mode of the green voxel intensity was calculated from the z-stack and subtracted. Non-microglial ROIs such as melanophores as well as microglia which had overlapping voxels with the axon were excluded manually and the average microglial green fluorescence intensity for the population of microglia in the z-stack was determined.

Colocalization analysis

Two-photon z-stacks were captured on day 3 post-electroporation. aRCA3-mCherry-Myc and SYP-pHtdGFP channels were denoised using the 3D hybrid median filter ImageJ plugin. A 3D mask of the axon was generated by summing the red and green channels and using an intensity threshold for segmentation of the axon. Pearson’s correlation coefficient (Manders et al., 1993) of the masked axon was calculated using Imaris software, with thresholds determined automatically using the method of Costes (Costes et al., 2004).

Looming Stimulus Behavioral Assay

Experimental setup

Stage 47 tadpoles were placed within a closed 60 mm petri dish and allowed to swim freely. The petri dish was placed on the bottom of a large shallow (20 cm diameter, 8 cm deep) glass culture dish filled completely with 0.1X MBSH. The large shallow glass culture dish was placed on a purple 3D printed stage to allow for automated segmentation of albino tadpoles. A webcam was placed above the tadpole to record tadpole behavior, while a 2000 lumens projector customized by 3D printing to shorten the
focal distance, was used to project visual stimuli onto a piece of 96 white 20 lb bond copy paper taped
to the side of the large glass culture dish.

**Stimulus presentation and recording**

Dark or bright looming stimuli were presented using custom code written in Python 3. An expanding
circle was projected onto the culture dish (800x600 pixels or 10.6x8 cm) using the PsychoPy library
(Peirce et al., 2019). For dark looming stimuli, a black (29 cd/m²) expanding circle was shown over a
white (208 cd/m²) background. For bright looming stimuli, a white expanding circle was shown over a
black background. During looming, the diameter of the circle expanded exponentially at 10% per frame
at 60 frames per second, from a diameter of 54 pixels (7 mm) until it encompassed the entire screen 0.5
s later. After another 0.8 s, the screen was reset and a 10 s refractory period commenced. In parallel to
stimulus presentation, 480p video was recorded by webcam using the OpenCV library (Bradski, 2000).
Ten looming stimulus trials per animal were recorded.

**Automated tracking of tadpole behavior**

Custom computer vision tadpole tracking code was written using Python 3 and OpenCV (Bradski, 2000).
Feature detection on the petri dish was used to automatically determine the scale of video data.
Background was subtracted and segmentation of the tadpole was carried out by thresholding. The
resulting mask was fit to an ellipse to extract location and speed, as well as directional data and escape
angle. Location data from 0 – 2 s following the onset of the looming stimulus was summarized and
displayed as a contrail. Instantaneous velocity for the 3 s before and after the looming stimulus was also
extracted and compared by area under curve analysis to obtain distance traveled.

**Discrimination of positive and negative responses to looming stimuli**

An automated python script was used to randomize tadpole videos and a user blinded to treatment
categorized tadpole responses to looming stimuli as defensive escape behavior (positive response),
absence of defensive escape behavior (negative response), or undeterminable (excluded from response
rate calculation). Tadpole responses were categorized as undeterminable if the tadpole was moving
quickly when the looming stimulus was presented. The response rate was calculated as the number of
positive responses divided by the combined number of positive and negative responses.

**Assessment of motor responses**

Positive escape responses were further analyzed to compare motor data. Escape distance and maximum
velocity over 2 seconds from the onset of the loom was calculated. Escape angle was also measured
from heading data. In dark looming stimuli trials, the escape angle was defined as the absolute value of
the change in heading from 0 – 0.6 s. For bright looming stimuli trials, the escape angle was defined as
the absolute value of the change in heading from 0 – 1.2 s. This difference is because dark looming
stimuli evoked escape movement beginning approximately 0.5 s after looming onset, whereas bright
looming stimuli evoked escape movement beginning approximately 1.0 s after looming onset.

**Bioinformatics analysis**

**Identification of complement inhibitory proteins**

To identify candidate complement inhibitory proteins, the STRING database (Szklarczyk et al., 2019) was
used to look for functional interactions between complement C3 and other proteins. Human
complement C3 was queried using a medium confidence (0.4) minimum interaction score, and a limit of
10 and 20 interactors for the 1st and 2nd shell, respectively. Sources was limited to textmining, experiments and databases.

*Expression of complement inhibitory proteins in human cortical cells*

The Allen Brain Map Human Transcriptomics Cell Types Database (Allen Institute for Brain Science, 2015) was used to examine the expression of candidate proteins identified from the STRING query. Cell taxonomy and hierarchical clustering was retained from previous analysis (Hodge et al., 2019). Gene expression of complement inhibitory proteins across distinct brain cell clusters was examined and plotted using R software and the Bioconductor (Huber et al., 2015), ComplexHeatmap (Gu et al., 2016), and ColorBrewer (Neuwirth, 2014) packages.

**Homology search**

The protein sequence of human CD46 (NCBI accession #: NP_002380.3) was queried in the *Xenopus laevis* genome with NCBI protein-protein BLAST (Altschul et al., 1990). The top 3 hits were predicted proteins XP_018102062.1, XP_018100169.1 and XP_018102058.1. The identity of the predicted proteins was obtained by cross-referencing with the *Xenopus tropicalis* genome using BLAST.

**Conserved modular architecture analysis and transmembrane topology**

The SMART protein domain research tool (Letunic and Bork, 2018; Schultz et al., 1998) was used to predict conserved modular architecture and the Phobius transmembrane topology tool (Käll et al., 2004, 2007) was used to predict transmembrane topology.

**Tertiary protein structure prediction**

*De novo* tertiary protein structure was predicted using trRosetta (Yang et al., 2020). The extracellular region, transmembrane region, and intracellular regions of human CD46 and *Xenopus laevis* aRCA3 were modeled separately by trRosetta and joined using pyMOL. The orientation of the protein in the phospholipid bilayer was predicted using the protein bilayer lipid membrane orientation package (Kimmett et al., 2014).

**Kinase prediction**

The NetPhos 3.1 tool (Blom et al., 2004) was used to make predictions of tyrosine phosphorylation sites on the intracellular C-terminus of aRCA3. A cutoff score of 0.6 was used.

**RNAscope in situ hybridization**

**Tissue preparation**

The RNAscope (Wang et al., 2012) fixed-frozen tissue sample preparation and pretreatment protocol provided by the manufacturer was modified for *Xenopus laevis* tadpoles. Stage 46 tadpoles were euthanized in 0.2% MS-222 in 0.1X MBSH. Tadpoles were then fixed in 4% PFA at 4°C for 24 hours on a laboratory rocker. For cryoprotection, tadpoles were then moved sequentially through 10%, 20% and 30% sucrose in 1X PBS, until samples sunk to the bottom of the container. Cryoprotected tadpoles were then embedded in OCT blocks on dry ice. OCT blocks were sectioned at 8 μm thickness on a cryostat and mounted on superfrost plus slides. To enhance tissue adhesion, slides were air dried for 2 hours at -20°C and baked at 60°C for 30 minutes. Slides were then post-fixed by immersion in 4% PFA for 15 minutes, and then dehydrated with 50%, 70% and 100% ethanol. Slides were treated with hydrogen peroxide to block endogenous peroxidases. For target retrieval, using a hot plate and beaker, slides were boiled in RNAscope target retrieval reagent and then treated with RNAscope Protease III.
The RNAscope Multiplex Fluorescent 2.0 Assay was performed according to manufacturer’s protocols using the HybEZ oven. In brief, probes were applied to slides, and 3 amplification steps were carried out. Opal 570 dye was applied to slides along with Hoechst counterstain. Coverslips were mounted with Prolong gold antifade mountant. Slides were imaged with a Leica TCS SP8 confocal.

The following plasmid constructs were constructed and used in this study:

- pEF1α-pHtdGFP
- pEF1α-SYN-pHtdGFP
- pEF1α-aRCA3-Myc-P2A-pHtdGFP
- pEF1α-aRCA3-mCherry-Myc-P2A-SYN-pHtdGFP
- pEF1α-pHtdGFP-P2A-Myc-VAMP2-C3
- pEF1α-pHtdGFP-P2A-VAMP2

Stage 40 Tadpoles were euthanized in 0.2% MS-222 in 0.1X MBSH. Animals were transferred to TRIzol reagent (Invitrogen, Carlsbad, CA) and homogenized by sonication and processed according to manufacturer’s protocols to isolate mRNA. Superscript IV reverse transcriptase (Invitrogen) was then used according to manufacturer’s protocol to generate whole tadpole cDNA.

Cloning and plasmid isolation

Cloning of *Xenopus laevis* cDNA was carried out using primers flanking genes of interest. Primers were generated according to mRNA sequences for aRCA3 (XM_018246573.1), VAMP2.S (NM_001087474.1), and c3.L (XM_018253729.1) predicted by NCBI (NCBI Resource Coordinators, 2018) and Xenbase (Karimi et al., 2018). Synaptophysin was subcloned from mouse synaptophysin (Ruthazer et al., 2006). pHtdGFP was subcloned from a pFA6a plasmid (Roberts et al., 2016). PCR was performed with Phusion High-Fidelity DNA polymerase (Thermo Scientific, Waltham, MA) and DNA fragments were cut with NEB enzymes and ligated into a plasmid with an EF1α promoter, ampicillin resistance, and a P2A self-cleaving construct when appropriate. DH5α bacteria were transformed and single clone colonies containing inserts were isolated. Endotoxin free plasmid preparations of high yield and purity for electroporation was prepared by maxi-prep (Qiagen, Hilden, Germany).

Design of fusion proteins

The SYN-pHtdGFP fusion protein was created by in-frame fusion of pHtdGFP to the C-terminus of SYN between a 3 amino acid linker (QGT). The aRCA3-mCherry-Myc fusion protein was created by an in-frame fusion of mCherry-Myc to the C-terminus of aRCA3 between a 2 amino acid linker (AC). The Myc-VAMP2-C3 fusion protein was created by an in-frame fusion of Myc to the N-terminus of VAMP2 between a 4 amino acid linker (PGKI) and an in-frame fusion of C3 to the C-terminus of VAMP2 between an 18 amino acid linker (ASIKSPVQPLSAHSPVCI). The long linker for VAMP2-C3 was chosen to ensure that the C-terminus transmembrane domain of VAMP2 would not sterically hinder proper folding of C3.

Statistical analysis

Statistical analysis was performed using Graphpad Prism 8 software. All data is presented as mean ± SEM, unless otherwise noted.

Pairwise comparisons

Pairwise comparisons between two groups were tested by unpaired t-test.
Group data

If no datapoints were missing, 2-way repeated measures ANOVA was used to test group data. If an interaction was found, the interaction was reported and Sidak’s multiple comparison test was used to test between groups or time points as appropriate. If no interaction was found, then the main effect was reported. Conversely, if the data contained missing values (due to loss of animals during a time course), the mixed effect restricted maximum likelihood model was used.

Correlation

Correlations were calculated using Pearson’s correlation coefficient. Trendlines were plotted using simple linear regression. Significance testing of the difference between two correlation coefficients was carried out using Fisher’s z transformation.

Data and Code Availability

The documented source code and user guide for the looming stimulus presentation and tadpole tracking software module, XenLoom (Beta), developed in this study is available at https://github.com/tonkyylim/XenLoom_beta.

Acknowledgments

We thank Dr Wayne Sossin, Dr Jean-Francois Cloutier and the MNI Microscopy Core Facility for sharing access to equipment. We also thank Dr Philip Kesner and Anne Schohl for technical advice, Dr Larissa Ferguson for technical assistance, and all members of the Ruthazer lab for helpful discussions. The pFA6a-pHtdGFP plasmid was generously provided by Dr Joerg Stelling. PLX5622 was kindly provided by Plexxikon, Inc. T.K.L. was supported by the CIHR Postdoctoral Fellowship and the McGill Faculty of Medicine McLaughlin Postdoctoral Fellowship. This work was supported by grants to E.S.R. from FRQS and CIHR.

Author Contributions

Conceptualization, E.S.R. and T.K.L.; Methodology, T.K.L. and E.S.R.; Software, T.K.L.; Formal Analysis, T.K.L. and E.S.R.; Investigation, T.K.L.; Writing – Original Draft, T.K.L.; Writing – Review and Editing, E.S.R.; Visualization, T.K.L. and E.S.R.; Resources, E.S.R.; Funding Acquisition, E.S.R.; Supervision, E.S.R.

Declaration of Interests

The authors declare no competing interests.
References

Ahmari, S.E., Buchanan, J., and Smith, S.J. (2000). Assembly of presynaptic active zones from cytoplasmic transport packets. Nat. Neurosci. 3, 445–451.

Allen Institute for Brain Science (2015). Allen Cell Types Database.

Altschul, S.F., Gish, W., Miller, W., Myers, E.W., and Lipman, D.J. (1990). Basic local alignment search tool. J. Mol. Biol. 215, 403–410.

Barilla-LaBarca, M.L., Liszewski, M.K., Lambris, J.D., Hourcade, D., and Atkinson, J.P. (2002). Role of membrane cofactor protein (CD46) in regulation of C4b and C3b deposited on cells. J. Immunol. Baltim. Md 1950 168, 6298–6304.

Barker, A.J., and Baier, H. (2015). Sensorimotor Decision Making in the Zebrafish Tectum. Curr. Biol. 25, 2804–2814.

Blom, N., Sicheritz-Pontén, T., Gupta, R., Gammeltoft, S., and Brunak, S. (2004). Prediction of post-translational glycosylation and phosphorylation of proteins from the amino acid sequence. Proteomics 4, 1633–1649.

Bolte, S., and Cordelières, F.P. (2006). A guided tour into subcellular colocalization analysis in light microscopy. J. Microsc. 224, 213–232.

Bradski, G. (2000). The OpenCV Library. Dr Dobbs J. Softw. Tools.

Brown, G.C., and Neher, J.J. (2014). Microglial phagocytosis of live neurons. Nat. Rev. Neurosci. 15, 209–216.

Casey, J.R., Grinstein, S., and Orlowksi, J. (2010). Sensors and regulators of intracellular pH. Nat. Rev. Mol. Cell Biol. 11, 50–61.

Chan, C.-S., Weeber, E.J., Kurup, S., Sweatt, J.D., and Davis, R.L. (2003). Integrin Requirement for Hippocampal Synaptic Plasticity and Spatial Memory. J. Neurosci. 23, 7107–7116.

Cong, Q., Soteros, B.M., Wollet, M., Kim, J.H., and Sia, G.-M. (2020). The endogenous neuronal complement inhibitor SRPX2 protects against complement-mediated synapse elimination during development. Nat. Neurosci. 1–12.

Costes, S.V., Daelemans, D., Cho, E.H., Dobbin, Z., Pavlakis, G., and Lockett, S. (2004). Automatic and quantitative measurement of protein-protein colocalization in live cells. Biophys. J. 86, 3993–4003.

Coupé, P., Munz, M., Manjón, J.V., Ruthazer, E.S., and Louis Collins, D. (2012). A CANDLE for a deeper in vivo insight. Med. Image Anal. 16, 849–864.

Cunningham, C.L., Martínez-Cerdeño, V., and Noctor, S.C. (2013). Microglia Regulate the Number of Neural Precursor Cells in the Developing Cerebral Cortex. J. Neurosci. 33, 4216–4233.

Dalmau, I., Finsen, B., Tønder, N., Zimmer, J., González, B., and Castellano, B. (1997). Development of microglia in the prenatal rat hippocampus. J. Comp. Neurol. 377, 70–84.

Dong, W., Lee, R.H., Xu, H., Yang, S., Pratt, K.G., Cao, V., Song, Y.-K., Nurmikko, A., and Aizenman, C.D. (2009). Visual Avoidance in Xenopus Tadpoles Is Correlated With the Maturation of Visual Responses in the Optic Tectum. J. Neurophysiol. 101, 803–815.
Dunn, T.W., Gebhardt, C., Naumann, E.A., Riegler, C., Ahrens, M.B., Engert, F., and Del Bene, F. (2016). Neural Circuits Underlying Visually Evoked Escapes in Larval Zebrafish. Neuron 89, 613–628.

Elmore, M.R.P., Najafi, A.R., Koike, M.A., Dagher, N.N., Spangenberg, E.E., Rice, R.A., Kitazawa, M., Matusow, B., Nguyen, H., West, B.L., et al. (2014). Colony-stimulating factor 1 receptor signaling is necessary for microglia viability, unmasking a microglia progenitor cell in the adult brain. Neuron 82, 380–397.

Erblich, B., Zhu, L., Etgen, A.M., Dobrenis, K., and Pollard, J.W. (2011). Absence of colony stimulation factor-1 receptor results in loss of microglia, disrupted brain development and olfactory deficits. PloS One 6, e26317.

Fotowat, H., and Gabbiani, F. (2011). Collision Detection as a Model for Sensory-Motor Integration. Annu. Rev. Neurosci. 34, 1–19.

Gosse, N.J., Nevin, L.M., and Baier, H. (2008). Retinotopic order in the absence of axon competition. Nature 452, 892–895.

Gu, Z., Eils, R., and Schlesner, M. (2016). Complex heatmaps reveal patterns and correlations in multidimensional genomic data. Bioinfor. Oxf. Engl. 32, 2847–2849.

Guo, Y., Luo, J., Tan, S., Otieno, B.O., and Zhang, Z. (2013). The applications of Vitamin E TPGS in drug delivery. Eur. J. Pharm. Sci. 49, 175–186.

Haas, K., Sin, W.C., Javaherian, A., Li, Z., and Cline, H.T. (2001). Single-cell electroporation for gene transfer in vivo. Neuron 29, 583–591.

Hanisch, U.-K. (2002). Microglia as a source and target of cytokines. Glia 40, 140–155.

Håvik, B., Le Hellard, S., Rietschel, M., Lybæk, H., Djurovic, S., Mattheisen, M., Mühleisen, T.W., Degenhardt, F., Priebe, L., Maier, W., et al. (2011). The Complement Control-Related Genes CSMD1 and CSMD2 Associate to Schizophrenia. Biol. Psychiatry 70, 35–42.

Hodge, R.D., Bakken, T.E., Miller, J.A., Smith, K.A., Barkan, E.R., Graybuck, L.T., Close, J.L., Long, B., Johansen, N., Penn, O., et al. (2019). Conserved cell types with divergent features in human versus mouse cortex. Nature 573, 61–68.

Hoshiko, M., Arnoux, I., Avignone, E., Yamamoto, N., and Audinat, E. (2012). Deficiency of the microglial receptor CX3CR1 impairs postnatal functional development of thalamocortical synapses in the barrel cortex. J. Neurosci. Off. J. Soc. Neurosci. 32, 15106–15111.

Huber, W., Carey, V.J., Gentleman, R., Anders, S., Carlson, M., Carvalho, B.S., Bravo, H.C., Davis, S., Gatto, L., Girke, T., et al. (2015). Orchestrating high-throughput genomic analysis with Bioconductor. Nat. Methods 12, 115–121.

Hordijk, W., van der Flier, M.W., van den Berg, R.C., van de Beek, A., van der Mei, D.H., and Pena, M.A. (2012). Microglia actively regulate the number of functional synapses. PloS One 8, e56293.

Käll, L., Krogh, A., and Sonnhammer, E.L.L. (2004). A combined transmembrane topology and signal peptide prediction method. J. Mol. Biol. 338, 1027–1036.

Käll, L., Krogh, A., and Sonnhammer, E.L.L. (2007). Advantages of combined transmembrane topology and signal peptide prediction—Phobius web server. Nucleic Acids Res. 35, W429-432.
Karimi, K., Fortriede, J.D., Lotay, V.S., Burns, K.A., Wang, D.Z., Fisher, M.E., Pells, T.J., James-Zorn, C., Wang, Y., Ponferrada, V.G., et al. (2018). Xenbase: a genomic, epigenomic and transcriptomic model organism database. Nucleic Acids Res. 46, D861–D868.

Karperien, A., Ahammer, H., and Jelinek, H.F. (2013). Quantitating the subtleties of microglial morphology with fractal analysis. Front. Cell. Neurosci. 7.

Khakhalin, A.S., Koren, D., Xu, H., and Aizenman, C.D. (2014). Excitation and inhibition in recurrent networks mediate collision avoidance in Xenopus tadpoles. Eur. J. Neurosci. 40, 2948–2962.

Kim, M., Haney, J.R., Zhang, P., Hernandez, L.M., Wang, L., Perez-Cano, L., and Gandal, M.J. (2020). Network signature of complement component 4 variation in the human brain identifies convergent molecular risk for schizophrenia (Genomics).

Kimmett, T., Smith, N., Witham, S., Petukh, M., Sarkar, S., and Alexov, E. (2014). ProBLM web server: protein and membrane placement and orientation package. Comput. Math. Methods Med. 2014, 838259.

Letunic, I., and Bork, P. (2018). 20 years of the SMART protein domain annotation resource. Nucleic Acids Res. 46, D493–D496.

Ling, E.A., Kaur, C., Yick, T.Y., and Wong, W.C. (1990). Immunocytochemical localization of CR3 complement receptors with OX-42 in amoeboid microglia in postnatal rats. Anat. Embryol. (Berl.) 182, 481–486.

Lozhic, S., Christiansen, D., Manié, S., Gerlier, D., Billard, M., Boucheix, C., and Rubinstein, E. (2000). CD46 (membrane cofactor protein) associates with multiple beta1 integrins and tetraspans. Eur. J. Immunol. 30, 900–907.

Luo, J., Elwood, F., Britschgi, M., Villeda, S., Zhang, H., Ding, Z., Zhu, L., Alabsi, H., Getachew, R., Narasimhan, R., et al. (2013). Colony-stimulating factor 1 receptor (CSF1R) signaling in injured neurons facilitates protection and survival. J. Exp. Med. 210, 157–172.

Manders, E.M.M., Verbeek, F.J., and Aten, J.A. (1993). Measurement of co-localization of objects in dual-colour confocal images. J. Microsc. 169, 375–382.

Milinkeviciute, G., Henningfield, C.M., Muniak, M.A., Chokr, S.M., Green, K.N., and Cramer, K.S. (2019). Microglia Regulate Pruning of Specialized Synapses in the Auditory Brainstem. Front. Neural Circuits 13, 55.

Mitchell, A.J., Pradel, L.C., Chasson, L., Rooijen, N.V., Grau, G.E., Hunt, N.H., and Chimini, G. (2010). Technical Advance: Autofluorescence as a tool for myeloid cell analysis. J. Leukoc. Biol. 88, 597–603.

Münch, T.A., da Silveira, R.A., Siegert, S., Viney, T.J., Awatramani, G.B., and Roska, B. (2009). Approach sensitivity in the retina processed by a multifunctional neural circuit. Nat. Neurosci. 12, 1308–1316.

Nakata, T., Terada, S., and Hirokawa, N. (1998). Visualization of the dynamics of synaptic vesicle and plasma membrane proteins in living axons. J. Cell Biol. 140, 659–674.

NCBI Resource Coordinators (2018). Database resources of the National Center for Biotechnology Information. Nucleic Acids Res. 46, D8–D13.

Neugebauer, K.M., and Reichardt, L.F. (1991). Cell-surface regulation of β1-integrin activity on developing retinal neurons. Nature 350, 68–71.

Neuwirth, E. (2014). Package ‘RColorBrewer.’
Newman, S.L., Becker, S., and Halme, J. (1985). Phagocytosis by receptors for C3b (CR1), iC3b (CR3), and IgG (Fc) on human peritoneal macrophages. J. Leukoc. Biol. 38, 267–278.

Nieuwkoop, P.D., and Faber, J. (1994). Normal table of Xenopus laevis (Daudin): a systematical and chronological survey of the development from the fertilized egg till the end of metamorphosis (New York: Garland Pub).

Nijhawan, D., Honarpour, N., and Wang, X. (2000). Apoptosis in Neural Development and Disease. Annu. Rev. Neurosci. 23, 73–87.

Nimmerjahn, A., Kirchhoff, F., and Helmchen, F. (2005). Resting microglial cells are highly dynamic surveillants of brain parenchyma in vivo. Science 308, 1314–1318.

Norman, D.G., Barlow, P.N., Baron, M., Day, A.J., Sim, R.B., and Campbell, I.D. (1991). Three-dimensional structure of a complement control protein module in solution. J. Mol. Biol. 219, 717–725.

Oliva, D., Medan, V., and Tomsic, D. (2007). Escape behavior and neuronal responses to looming stimuli in the crab Chasmagnathus granulatus (Decapoda: Grapsidae). J. Exp. Biol. 210, 865–880.

Ollion, J., Cochennek, J., Loll, F., Escudé, C., and Boudier, T. (2013). TANGO: a generic tool for high-throughput 3D image analysis for studying nuclear organization. Bioinformatics 29, 1840–1841.

Oshiumi, H., Suzuki, Y., Matsumoto, M., and Seya, T. (2009). Regulator of complement activation (RCA) gene cluster in Xenopus tropicalis. Immunogenetics 61, 371–384.

Pangburn, M.K., Schreiber, R.D., and Müller-Eberhard, H.J. (1981). Formation of the initial C3 convertase of the alternative complement pathway. Acquisition of C3b-like activities by spontaneous hydrolysis of the putative thioester in native C3. J. Exp. Med. 154, 856–867.

Paolicelli, R.C., Bolasco, G., Pagani, F., Maggi, L., Scianni, M., Panzanelli, P., Giustetto, M., Ferreira, T.A., Guiducci, E., Dumas, L., et al. (2011). Synaptic Pruning by Microglia Is Necessary for Normal Brain Development. Science 333, 1456–1458.

Parkhurst, C.N., Yang, G., Ninan, I., Savas, J.N., Yates, J.R., Lafaille, J.J., Hempstead, B.L., Littman, D.R., and Gan, W.-B. (2013). Microglia promote learning-dependent synapse formation through brain-derived neurotrophic factor. Cell 155, 1596–1609.

Peirce, J., Gray, J.R., Simpson, S., MacAskill, M., Höchenberger, R., Sogo, H., Kastman, E., and Lindeløv, J.K. (2019). PsychoPy2: Experiments in behavior made easy. Behav. Res. Methods 51, 195–203.

Perry, V.H., and O’Connor, V. (2008). C1q: the perfect complement for a synaptic feast? Nat. Rev. Neurosci. 9, 807–811.

Pont-Lezica, L., Beumer, W., Colasse, S., Drexhage, H., Versnel, M., and Bessis, A. (2014). Microglia shape corpus callosum axon tract fasciculation: functional impact of prenatal inflammation. Eur. J. Neurosci. 39, 1551–1557.

Preibisch, S., Saalfeld, S., Schindelin, J., and Tomancak, P. (2010). Software for bead-based registration of selective plane illumination microscopy data. Nat. Methods 7, 418–419.

R Core Team (2018). R: A Language and Environment for Statistical Computing (Vienna, Austria: R Foundation for Statistical Computing).

Reshef, R., Kudryavitskaya, E., Shani-Narkiss, H., Isaacson, B., Rimmerman, N., Mizrahi, A., and Yirmiya, R. (2017). The role of microglia and their CX3CR1 signaling in adult neurogenesis in the olfactory bulb. ELife 6, e30809.
Riley-Vargas, R.C., Gill, D.B., Kemper, C., Liszewski, M.K., and Atkinson, J.P. (2004). CD46: expanding beyond complement regulation. Trends Immunol. 25, 496–503.

Ripke, S., Neale, B.M., Corvin, A., Walters, J.T.R., Farh, K.-H., Holmans, P.A., Lee, P., Bulik-Sullivan, B., Collier, D.A., Huang, H., et al. (2014). Biological insights from 108 schizophrenia-associated genetic loci. Nature 511, 421–427.

Roberts, T.M., Rudolf, F., Meyer, A., Pellaux, R., Whitehead, E., Panke, S., and Held, M. (2016). Identification and Characterisation of a pH-stable GFP. Sci. Rep. 6, 1–9.

Robles, E., Laurell, E., and Baier, H. (2014). The Retinal Projectome Reveals Brain-Area-Specific Visual Representations Generated by Ganglion Cell Diversity. Curr. Biol. 24, 2085–2096.

Ross, G.D., Reed, W., Dalzell, J.G., Becker, S.E., and Hogg, N. (1992). Macrophage cytoskeleton association with CR3 and CR4 regulates receptor mobility and phagocytosis of iC3b-opsonized erythrocytes. J. Leukoc. Biol. 51, 109–117.

Ruthazer, E.S., Akerman, C.J., and Cline, H.T. (2003). Control of axon branch dynamics by correlated activity in vivo. Science 301, 66–70.

Ruthazer, E.S., Li, J., and Cline, H.T. (2006). Stabilization of Axon Branch Dynamics by Synaptic Maturation. J. Neurosci. 26, 3594–3603.

Ruthazer, E.S., Schohl, A., Schwartz, N., Tavakoli, A., Tremblay, M., and Cline, H.T. (2013a). Bulk Electroporation of Retinal Ganglion Cells in Live Xenopus Tadpoles. Cold Spring Harb. Protoc. 2013, pdb.prot076471.

Ruthazer, E.S., Schohl, A., Schwartz, N., Tavakoli, A., Tremblay, M., and Cline, H.T. (2013b). In Vivo Time-Lapse Imaging of Neuronal Development in Xenopus. Cold Spring Harb. Protoc. 2013, pdb.top077156.

Sahu, A., Kozel, T.R., and Pangburn, M.K. (1994). Specificity of the thioester-containing reactive site of human C3 and its significance to complement activation. Biochem. J. 302, 429–436.

Sankaranarayanan, S., and Ryan, T.A. (2000). Real-time measurements of vesicle-SNARE recycling in synapses of the central nervous system. Nat. Cell Biol. 2, 197–204.

Schafer, D.P., Lehrman, E.K., Kautzman, A.G., Koyama, R., Mardinly, A.R., Yamasaki, R., Ransohoff, R.M., Greenberg, M.E., Barres, B.A., and Stevens, B. (2012). Microglia Sculpt Postnatal Neural Circuits in an Activity and Complement-Dependent Manner. Neuron 74, 691–705.

Schindelin, J., Arganda-Carreras, I., Frise, E., Kaynig, V., Longair, M., Pietzsch, T., Preibisch, S., Rueden, C., Saalfeld, S., Schmid, B., et al. (2012). Fiji: an open-source platform for biological-image analysis. Nat. Methods 9, 676–682.

Schrödinger, LLC. The PyMOL Molecular Graphics System, Version 2.4.

Schultz, J., Milpetz, F., Bork, P., and Ponting, C.P. (1998). SMART, a simple modular architecture research tool: Identification of signaling domains. Proc. Natl. Acad. Sci. 95, 5857–5864.

Sellgren CM, Gracias J, Watmuff B, Biag JD, Thanos JM, Whittredge PB, Fu T, Worringer K, Brown HE, Wang J, et al. (2019). Increased synapse elimination by microglia in schizophrenia patient-derived models of synaptic pruning. Nat. Neurosci. 22, 374–385.

Shinoda, H., Shannon, M., and Nagai, T. (2018). Fluorescent Proteins for Investigating Biological Events in Acidic Environments. Int. J. Mol. Sci. 19.
Smolders, S.M.-T., Swinnen, N., Kessels, S., Arnauts, K., Smolders, S., Bras, B.L., Rigo, J.-M., Legendre, P., and Brône, B. (2017). Age-specific function of α5β1 integrin in microglial migration during early colonization of the developing mouse cortex. Glia 65, 1072–1088.

Solek, C.M., Farooqi, N., Verly, M., Lim, T.K., and Ruthazer, E.S. (2018). Maternal immune activation in neurodevelopmental disorders. Dev. Dyn. 247, 588–619.

Squarzoni, P., Oller, G., Hoeffel, G., Pont-Lezica, L., Rostaing, P., Low, D., Bessis, A., Ginhoux, F., and Garel, S. (2014). Microglia Modulate Wiring of the Embryonic Forebrain. Cell Rep. 8, 1271–1279.

Stephan, A.H., Barres, B.A., and Stevens, B. (2012). The Complement System: An Unexpected Role in Synaptic Pruning During Development and Disease. Annu. Rev. Neurosci. 35, 369–389.

Stevens, B., Allen, N.J., Vazquez, L.E., Howell, G.R., Christopherson, K.S., Nouri, N., Micheva, K.D., Mehalow, A.K., Huberman, A.D., Stafford, B., et al. (2007). The classical complement cascade mediates CNS synapse elimination. Cell 131, 1164–1178.

Svahn, A.J., Graeber, M.B., Ellett, F., Lieschke, G.J., Rinkwitz, S., Bennett, M.R., and Becker, T.S. (2013). Development of ramified microglia from early macrophages in the zebrafish optic tectum. Dev. Neurobiol. 73, 60–71.

Szklarczyk, D., Gable, A.L., Lyon, D., Junge, A., Wyder, S., Huerta-Cepas, J., Simonovic, M., Doncheva, N.T., Morris, J.H., Bork, P., et al. (2019). STRING v11: protein-protein association networks with increased coverage, supporting functional discovery in genome-wide experimental datasets. Nucleic Acids Res. 47, D607–D613.

Temizer, I., Donovan, J.C., Baier, H., and Semmelhack, J.L. (2015). A Visual Pathway for Looming-Evoked Escape in Larval Zebrafish. Curr. Biol. 25, 1823–1834.

Tinevez, J.-Y., Perry, N., Schindelin, J., Hoopes, G.M., Reynolds, G.D., Laplantine, E., Bednarek, S.Y., Shorte, S.L., and Eliceiri, K.W. (2017). TrackMate: An open and extensible platform for single-particle tracking. Methods 115, 80–90.

Tremblay, M.-È., Lowery, R.L., and Majewska, A.K. (2010). Microglial interactions with synapses are modulated by visual experience. PLoS Biol. 8, e1000527.

Valtorta, F., Pennuto, M., Bonanomi, D., and Benfenati, F. (2004). Synaptophysin: leading actor or walk-on role in synaptic vesicle exocytosis? BioEssays News Rev. Mol. Cell. Dev. Biol. 26, 445–453.

Wake, H., Moorhouse, A.J., Jinno, S., Kohsaka, S., and Nabekura, J. (2009). Resting microglia directly monitor the functional state of synapses in vivo and determine the fate of ischemic terminals. J. Neurosci. Off. J. Soc. Neurosci. 29, 3974–3980.

Wallace, J., Lord, J., Dissing-Olesen, L., Stevens, B., and Murthy, V.N. (2020). Microglial depletion disrupts normal functional development of adult-born neurons in the olfactory bulb. Elife 9, e50531.

Wang, C., Yue, H., Hu, Z., Shen, Y., Ma, J., Li, J., Wang, X.-D., Wang, L., Sun, B., Shi, P., et al. (2020). Microglia mediate forgetting via complement-dependent synaptic elimination. Science 367, 688–694.

Wang, F., Flanagan, J., Su, N., Wang, L.-C., Bui, S., Nielson, A., Wu, X., Vo, H.-T., Ma, X.-J., and Luo, Y. (2012). RNAscope: a novel in situ RNA analysis platform for formalin-fixed, paraffin-embedded tissues. J. Mol. Diagn. JMD 14, 22–29.
Weinhard, L., di Bartolomei, G., Bolasco, G., Machado, P., Schieber, N.L., Neniskyte, U., Exiga, M., Vadiute, A., Raggioli, A., Schertel, A., et al. (2018). Microglia remodel synapses by presynaptic trogocytosis and spine head filopodia induction. Nat. Commun. 9, 1228.

Werneburg, S., Jung, J., Kunjamma, R.B., Ha, S.-K., Luciano, N.J., Willis, C.M., Gao, G., Biscola, N.P., Havton, Leif.A., Crocker, S.J., et al. (2020). Targeted Complement Inhibition at Synapses Prevents Microglial Synaptic Engulfment and Synapse Loss in Demyelinating Disease. Immunity 52, 167-182.e7.

Yang, J., Anishchenko, I., Park, H., Peng, Z., Ovchinnikov, S., and Baker, D. (2020). Improved protein structure prediction using predicted interresidue orientations. Proc. Natl. Acad. Sci. 117, 1496–1503.

Yilmaz, M., and Meister, M. (2013). Rapid innate defensive responses of mice to looming visual stimuli. Curr. Biol. CB 23, 2011–2015.

Zeitvogel, F., Schmid, G., Hao, L., Ingino, P., and Obst, M. (2016). ScatterJ: An ImageJ plugin for the evaluation of analytical microscopy datasets. J. Microsc. 261, 148–156.
Supplemental Figures

Figure S1. IB4-isolectin-conjugated fluorophores label microglial cells in developing *Xenopus laevis* tadpoles, related to Figure 1.

(A) The tadpole brain colorized for identification (yellow = olfactory bulb and forebrain, green = optic tectum, blue = hindbrain). To label microglia, IB4-isolectin conjugated fluorophores are injected into the 3rd ventricle (red).

(B) Dynamic cells with both ameboid-like and primitive ramified-like morphologies are labeled by IB4-isolectin.

(C) The distribution in mobility of IB4-isolectin labeled cells under normal conditions. Average velocity is 1.75 μm/min.

(D) Laser irradiation injury of the neuropil and response by IB4-isolectin labeled cells. Laser irradiation injury induced a region of damaged, autofluorescent, tissue. IB4-isolectin labeled cells mobilize to the injury site and remove the injured tissue by phagocytosis. IB4-isolectin labeled cells are shown in red, and eGFP-labeled RGC axons are shown in green.
Figure S2. Correlation between the change in microglial green fluorescence from day 4 to day 5 and the number of pHtdGFP-labeled and SYP-pHtdGFP-labeled axons, related to Figure 2

(A) The change in microglial green fluorescence from day 4 to day 5 post-labeling correlates with the number of pHtdGFP-labeled axons in the optic tectum. The same dataset is analyzed in Figure 2C and Figure 2D. (n = 47, Pearson $r = 0.48$, $p = 0.0006$).

(B) The change in microglial green fluorescence from day 4 to day 5 post-labeling correlates with the number of SYP-pHtdGFP-labeled axons in the optic tectum. (n = 57, Pearson $r = 0.34$, $p = 0.0092$). The same dataset is analyzed in Figure 2F.
Figure S3. Motor characteristics of escape behavior are not significantly altered by PLX5622-induced microglial depletion, related to Figure 4
A-F analyze all trials where defensive behavior was exhibited to looming stimuli. Data was collected from 8 animals per group.

(A) Microglial depletion did not significantly alter escape distance to dark looming stimuli. Unpaired t-test p = 0.27.
(B) Microglial depletion did not significantly alter escape distance to bright looming stimuli. Unpaired t-test p = 0.26.
(C) Microglial depletion did not significantly alter maximum escape velocity to dark looming stimuli. Unpaired t-test p = 0.32.
(D) Microglial depletion did not significantly alter maximum escape velocity to bright looming stimuli. Unpaired t-test p = 0.25.
(E) Microglial depletion did not significantly alter escape angle to dark looming stimuli. Unpaired t-test p = 0.40.
(F) Microglial depletion did not significantly alter escape angle to bright looming stimuli. Unpaired t-test p = 0.72.
Figure S4. Correlation between the change in microglial green fluorescence from day 4 to day 5 and the number of pHtdGFP-labeled axons with and without aRCA3 overexpression, related to Figure 6

Overexpression of aRCA3 significantly reduced the correlation coefficient between the change in microglial green fluorescence from day 4 to day 5 and the number of pHtdGFP-labeled axons in the optic tectum. Fisher’s z transformation * p < 0.05. Control correlation (n = 39, Pearson r = 0.51, p = 0.0009). aRCA3 overexpression correlation (n = 42, Pearson r = 0.076, p = 0.63). The same datasets are shown in Figure 6D and Figure 6E.
Supplemental Movies

Movie S1. IB4-isolectin labeled cells in *Xenopus laevis* tadpoles are morphologically dynamic and highly mobile, related to Figure 1

(A) IB4-isolectin cells are highly mobile. Timestamp = HH:MM.

(B) IB4-isolectin labeled cells have a dynamic morphology, switching back and forth between ameboid-like and primitive ramified-like morphologies.

Movie S2. Response of IB4-isolectin labeled cells to injury, related to Figure 1

IB4-isolectin labeled cells respond to tissue injury by mobilization to the injury site and phagocytosis of injured tissues. Also shown in Figure S1D. Timestamp = HH:MM.
Movie S3. Microglia surveil the tectal neuropil in developing *Xenopus laevis*, related to Figure 1

(A) Microglia associate with the tectal neuropil in *Xenopus laevis*. Timestamp = HH:MM.

(B) The neuropil does not exclude microglia. Microglia can mobilize into the neuropil region from the cell body layer and can freely move through the neuropil region. Timestamp = HH:MM.

(C) Microglia surveil the tectal neuropil by extending processes into the neuropil from the cell body layer. Timestamp = HH:MM.

Movie S4. *In vivo* real-time trogocytosis in *Xenopus laevis* tadpoles imaged by 2-photon microscopy, related to Figure 1

(A) A microglial cell increases in green fluorescence in real-time after an interaction with a pHtdGFP-labeled axon. The colocalization of green and red is colorized as white. Timestamp = HH:MM.

(B) Another example of a microglial cell increasing in green fluorescence following an interaction with a pHtdGFP-labeled axon. This example is shown in Figure 1E. Timestamp = HH:MM.
Movie S5. Representative responses to dark looming stimuli in control and microglia-depleted animals, related to Figure 4

(A) Representative response to dark looming stimuli in a vehicle control animal.

(B) Representative response to dark looming stimuli in a microglia-depleted animal.

Movie S6. Representative responses to bright looming stimuli in control and microglia-depleted animals, related to Figure 4

(A) Representative response to bright looming stimuli in a vehicle control animal.

(B) Representative response to bright looming stimuli in a microglia-depleted animal.
Movie S7. Predicted tertiary protein structure of CD46 and aRCA3, related to Figure 5

(A) The predicted tertiary protein structure of CD46.

(B) The predicted tertiary protein structure of aRCA3.
A

B

Microglia enter the neuropil region

C

Microglia surveil the neuropil

D

Microglia send processes into the neuropil

brief contacts

prolonged contacts

E

Real-time increase in microglial green fluorescence following interaction with pHtdGFP labelled axon

F

Graph showing microglial green fluorescence over time (mins) with and without interaction with axon.
**A**

|        | Day 0 | Day 1 | Day 2 | Day 3 | Day 4 | Day 5 |
|--------|-------|-------|-------|-------|-------|-------|
| Stage  | 39/40 |       |       |       |       |       |

RGC electroporation with pHtdGFP and labeling of microglia with IB4-isoelectin

**Labeled axons innervate optic tectum**

**Microglia colonize the optic tectum**

*in vivo imaging*

**B**

**Data Acquisition:**
2-photon 830 nm

**IB4-isoelectin Alexa 594**

**pHtdGFP**

**3D masks of microglia**

**Average microglial green fluorescence**

**C**

![Graph](image)

- **n=16** ns
- **n=12** ***
- **n=11** ***
- **n=8** ****

**Microglial green fluorescence**

(adjust units)

**D**

![Graph](image)

- **n=47**

**r = 0.52**

**p = 0.002**

**r = 0.30**

**p = 0.038**

**E**

![IB4-isoelectin Alexa 594 SYP-pHtdGFP](image)

- **20 μm**

**F**

![Graph](image)

- **n=57**

**r = 0.54**

**p = 0.010**

**r = 0.34**

**p = 0.063**

**r = 0.063**

**p = 0.84**
A

Day 2  Day 3  Day 4  Day 5
Vehicle Control

10 μM PLX5622

Day 2  Day 3  Day 4  Day 5
Vehicle Control

10 μM PLX5622

B

Days post-treatment

# microglia

C

Days post-treatment

# of processes

D

Day 2  Day 3  Day 4  Day 5
Vehicle Control

10 μM PLX5622

Day 2  Day 3  Day 4  Day 5
Vehicle Control

10 μM PLX5622

E

Days post-electroporation

Axon arbor length (µm)

F

Days post-electroporation

# of branches
A. Diagram showing the setup of the experiment with a projector (lens removed), webcam, paper screen, large glass bowl, 3D printed lens mount, and 3D printed stage.

B. Graphical representation of the stimuli used in the experiment: Dark looming stimulus and Bright looming stimulus, each with a duration of 0.5 seconds.

C. Time-lapse images of a petri dish with Tadpole showing the response to the stimuli over time (0-3 seconds).

D. Comparison of vehicle control and 10 μM PLX5622 on dark looming stimuli.

E. Graph showing instantaneous velocity over time for vehicle control and 10 μM PLX5622.

F. Histogram showing distance traveled before and after treatment with vehicle and 10 μM PLX5622.

G. Bar graph showing escape probability before and after treatment with vehicle and 10 μM PLX5622.

H. Comparison of vehicle control and 10 μM PLX5622 on bright looming stimuli.

I. Graph showing instantaneous velocity over time for vehicle control and 10 μM PLX5622.

J. Histogram showing distance traveled before and after treatment with vehicle and 10 μM PLX5622.

K. Bar graph showing escape probability before and after treatment with vehicle and 10 μM PLX5622.
A Day 2 Day 3 Day 4 Day 5

Control

B VAMP2

Day 2 Day 3 Day 4 Day 5

pHtdGFP

C 20 μm

D

VAMP2-C3

0 500 1000 1500 2000 2500
Axon length (μm)

Days post-electroporation

VAMP2

0 50 100 150 200
# of branches

Days post-electroporation

VAMP2-C3

pHtdGFP

20 μm

Control (n=7)

VAMP2-C3 (n=9)

VAMP2 (n=13)
A

**Normal axonal pruning**
- Impaired axonal pruning
- Excessive axonal pruning

**Complement activity**

B

**Normal development**
- Microglial-mediated axonal pruning
- Reduced axonal arborization
- Appropriate wiring

**Microglial depletion**
- Impaired microglial-mediated axonal pruning
- Increased axonal arborization
- Inappropriate wiring

C

| Retina                  | Normal development                                      | Microglial depletion                                |
|-------------------------|---------------------------------------------------------|-----------------------------------------------------|
| Dark looming-sensitive RGC neurons | ![Diagram](image1)                                      | ![Diagram](image2)                                  |
| Bright looming-sensitive RGC neurons | ![Diagram](image3)                                     | ![Diagram](image4)                                  |

**Optic tectum**
- Threat detection circuits
- Non-threat detecting circuits

**Escape Behavior**

---

A bioRxiv preprint doi: [https://doi.org/10.1101/2020.06.07.087221](https://doi.org/10.1101/2020.06.07.087221); this version posted August 4, 2020. The copyright holder for this preprint (which was not certified by peer review) is the author/funder. All rights reserved. No reuse allowed without permission.