A Novel Lysolecithin Model for Visualizing Damage in vivo in the Larval Zebrafish Spinal Cord

Angela D. Morris1 and Sarah Kucenas1,2*

1 Neuroscience Graduate Program, University of Virginia, Charlottesville, VA, United States, 2 Department of Biology, University of Virginia, Charlottesville, VA, United States

Background: Lysolecithin is commonly used to induce demyelinating lesions in the spinal cord and corpus callosum of mammalian models. Although these models and clinical patient samples are used to study neurodegenerative diseases, such as multiple sclerosis (MS), they do not allow for direct visualization of disease-related damage in vivo. To overcome this limitation, we created and characterized a focal lysolecithin injection model in zebrafish that allows us to investigate the temporal dynamics underlying lysolecithin-induced damage in vivo.

Results: We injected lysolecithin into 4–6 days post-fertilization (dpf) zebrafish larval spinal cords and, coupled with in vivo, time-lapse imaging, observed hallmarks consistent with mammalian models of lysolecithin-induced demyelination, including myelinating glial cell loss, myelin perturbations, axonal sparing, and debris clearance.

Conclusion: We have developed and characterized a lysolecithin injection model in zebrafish that allows us to investigate myelin damage in a living, vertebrate organism. This model may be a useful pre-clinical screening tool for investigating the safety and efficacy of novel therapeutic compounds that reduce damage and/or promote repair in neurodegenerative disorders, such as MS.

Keywords: demyelination, oligodendrocyte, zebrafish, myelin, central nervous system, spinal cord

INTRODUCTION

Multiple sclerosis (MS) is a neuroinflammatory disorder characterized by the inappropriate attack of the myelin sheath by the immune system, which results in demyelination, axonal degeneration, and eventually severe disability. MS primarily attacks the central nervous system (CNS) in the form of focal demyelinating lesions found within the brain, spinal cord, and/or optic nerve, and patients battle various neurological challenges that are unique to lesion location. Mammalian models commonly used to study MS, including experimental autoimmune encephalomyelitis (EAE) and drug-induced demyelination, as well as patient data or clinical samples, do not allow researchers to directly visualize processes underlying myelin destruction and repair in vivo.
Lysophosphatidylcholine, commonly referred to as lysolecithin, has been used in mammalian models, including mice, rats, rabbits, cats, and Macaque monkeys, to induce demyelinating lesions in the spinal cord and the corpus callosum (Blakemore et al., 1977; Dousset et al., 1995; Foote and Blakemore, 2005; Girard et al., 2005; Gregg et al., 2007; Blakemore and Franklin, 2008). Injections of 1% lysolecithin are sufficient to induce demyelination in the CNS of these animals, and evidence of remyelination begins between 7 and 10 days post-injection in rodents (dpf) (Jeffery and Blakemore, 1995; Kotter et al., 2001).

Although a generalized demyelination model exists in zebrafish that takes advantage of the ease of drug submersion combined with tissue-specific ablation by the bacterial nitroreductase gene, a major limitation of this model is the inability to create focal lesions, which are a hallmark of MS pathology (Chung et al., 2013; Fang et al., 2014; Karttunen et al., 2017; Karttunen and Lyons, 2019). A previous study has also demonstrated that 1% lysolecithin placed on gelatin foam can successfully induce demyelination when applied directly to the adult zebrafish optic nerve (Münzel et al., 2014). More recently, a model of lysolecithin injection into the trunk of larval zebrafish was reported that utilized in vivo imaging to describe the pro-inflammatory response after demyelination (Cunha et al., 2020). In this manuscript, we have also created and characterized a model that induces focal lesions in the zebrafish spinal cord, allowing us to visualize lysolecithin-induced damage in a living, intact vertebrate model system.

RESULTS

Creating a Focal Lysolecithin Injection Model in Zebrafish

In order to inject into the spinal cord of 4–6 days post-fertilization (dpf) larvae (Figure 1A) to create focal areas of damage, we mounted larvae laterally on 2% agar pads solidified onto 22 mm × 60 mm borosilicate glass coverslips (Figure 1B). Coverslips containing mounted larvae were placed directly onto a Zeiss Axio Observer Z1 microscope stage equipped with an ASI MPPI-3 pressure injection rig and mechanical micromanipulator (Figure 1C). A micropipette needle containing either a 1× phosphate-buffered saline (PBS) control solution or a 0.875% lysolecithin experimental solution was inserted in the dorsal trunk and injected into the larvae via passive diffusion. When the micropipette was not inserted successfully into the spinal cord, the solution was inappropriately injected into the muscle, evidenced by the muscle appearing disturbed upon solution dispersal (Figure 1D). Successful injections resulted in the solution remaining localized within the spinal cord (Figure 1E). To confirm that injection volumes were comparable between the control and experimental groups, we included a fluorescent dye (Dextran-647) with the injection cocktail. In these injections, we observed comparable dispersal between the control (Figure 1F) and experimental solutions (Figure 1G), which on average, was 1–2 somites both anterior and posterior to the injection site. Additionally, we observed a similar survival rate 24 h post-injection for both groups, with the control survival at 72.67% (n = 236) and experimental survival at 69.4% (n = 183). From these studies, we present a targeted method to dispense solutions focally and precisely into the larval zebrafish spinal cord.

Lysolecithin Alters the Number of Sox10+ Cells in the Zebrafish Spinal Cord

Myelination in the zebrafish CNS occurs in an anterior to posterior manner, commencing around 3 dpf (Almeida et al., 2011; Czopka et al., 2013). Therefore, the presence of maturing myelin sheaths between 4 and 6 dpf, coupled with the transparency of transgenic larvae, is ideal for in vivo imaging of the oligodendrocyte lineage cell (OLC) response following exposure to a demyelinating agent, such as lysolecithin. Previous publications have reported a decrease in oligodendrocyte number, demyelination, oligodendrocyte progenitor cell (OPC) proliferation, and OPC migration to the focal injury after injection of lysolecithin into the mammalian spinal cord (Blakemore et al., 1977; Kotter et al., 2001; Blakemore and Franklin, 2008; Lau et al., 2012; Münzel et al., 2014; Azin et al., 2015; Keough et al., 2015). Therefore, we sought to determine if lysolecithin similarly altered OLCs in our model.

To quantify the effects on OLC number in our model, we injected either a lysolecithin cocktail (0.875% lysolecithin + 0.21% Dextran-647 in PBS) or a control solution (PBS + 0.21% Dextran-647) into the spinal cord of 4 dpf Tg(sox10:mrfp) larvae, where sox10 regulatory sequences drive the expression in OLCs. Following injection, we fixed larvae at distinct time points post-injection and performed whole mount immunohistochemistry with an antibody specific to zebrafish Sox10 to count the number of Sox10+ cells in the spinal cord (Binari et al., 2013). As a control, we quantified the number of Sox10− cells anterior to the injection/dispersal region within a region of the spinal cord spanning approximately three motor nerves in width to avoid effects caused by the injection injury itself and observed no difference in Sox10− cell number at 20 hpi (Figures 2A–D, left panels, control average, n = 5, 29.3 Sox10− cells, experimental average, n = 6, 31.3 Sox10− cells, p = 0.3391). We also quantified within the injection/dispersal region to capture the effects from either the lysolecithin or control solution (Figures 2A–E, right panels). The region posterior to the injection site was not quantified to reduce the risk of capturing a decrease in the number of OLCs that might be secondary to Wallerian degeneration (Waller, 1851). At 8 hours post-injection (hpi) within the injection site dispersal region, we observed a significant reduction in the number of Sox10+ cells (Figure 2F) in the lysolecithin-injected group (Figure 2B, right panel) (average 24.94, n = 6) as compared with control larvae (Figure 2A, right panel) (average 30.33, n = 5), demonstrating that lysolecithin is toxic to zebrafish OLCs. When we looked at 20 hpi, we observed an increase in the number of Sox10− cells in lysolecithin-injected larvae (Figure 2D, right panel) (average 32.96, n = 8) as compared with control-injected larvae (Figure 2C, right panel) (average 26.8, n = 5), which
FIGURE 1 | Focal delivery of solutions into the zebrafish spinal cord. (A) Cartoon of a zebrafish larvae identifying the brain and spinal cord (green). The boxed region identifies the injection site/targeted region of interest (ROI). (B) An agar pad (arrow) solidified on a borosilicate glass coverslip is used as a mounting platform for larvae to perform spinal cord injections. (C) An agar pad containing mounted larvae is placed on a microscope stage, equipped with a pressure injection rig and mechanical micromanipulator to hold the injection micropipette. (D) Muscle injections are identified by a wavy appearance following dispersal of the injection solution. Yellow arrows identify the injection site. Dotted lines denote the muscle segment injected. (E) Solutions remain localized within the spinal cord following a successful spinal cord injection. Yellow arrows identify the injection site in the dorsal spinal cord. White arrowheads identify the solution dispersal region. (F) Dispersal region of a control solution containing water and the fluorescent tracer, Dextran-647. Yellow arrows identify the injection site. White arrowheads identify the dispersal region. (G) Dispersal region of a lysolecithin solution including the fluorescent tracer Dextran-647. Yellow arrows identify the injection site. White arrowheads identify the dispersal region. Red dashed lines denote the spinal cord. Scale bar, 50 µm.
FIGURE 2 | Lysolecithin alters the number and behavior of Sox10+ cells in the spinal cord. All images are lateral views of the spinal cord with anterior to the left and dorsal to the top. (A–D) 4 dpf Tg(sox10:mrfp) zebrafish larvae injected with either a control solution [control, (A,C)] or lysolecithin [experimental, (B,D)] containing the fluorescent tracer Dextran-647 to identify the injection site. Injected zebrafish were fixed at 8 hpi (A,B) or 20 hpi (C,D) and labeled with an antibody to Sox10 (green). (E) Quantification of the average number of Sox10+ cells within the injection site and spanning the width of approximately three motor nerves to capture the effects from lysolecithin (experimental) or control solutions. *p = 0.0252 at 8 hpi (n = 5 control; n = 6 experimental) and **p = 0.0033 at 20 hpi (n = 5 control; n = 8 experimental). Statistical significance was measured using an unpaired t-test. (F) Time-lapse imaging of proliferation (red arrowhead) and migration (white arrowhead) of sox10+ cells in a Tg(sox10:nls-eos) zebrafish larva following injection of lysolecithin at 4 dpf. White dashed ellipse denotes the injection site. (G) Time-lapse movie following injection of lysolecithin and photoconversion of the lesion area in a 4 dpf Tg(sox10:nls-eos) zebrafish larva. Red fluorescence denotes the photoconverted cells within the lesion. Green fluorescence denotes the cells outside of the lesion. Arrowhead identifies a sox10+ cell anterior to the injection site that migrates posteriorly into the drug dispersal region. White dashed lines denote the spinal cord. Scale bars, 25 µm.

we hypothesize is due to OPC proliferation, a phenomena reported in mammalian lysolecithin-induced demyelination models (Nait-Oumesmar et al., 1999; Keough et al., 2015; Sahel et al., 2015). To determine if OLCs were proliferating after exposure to lysolecithin and to investigate whether migration also occurred in our model, we injected the lysolecithin cocktail into 4–6 dpf Tg(sox10:nls-eos) (McGraw et al., 2012; Prendergast et al., 2012) larvae and used in vivo, time-lapse imaging to follow sox10+ cells with an imaging interval of 8 min. Following injection of lysolecithin, we observed that sox10+ OLCs near the injection site divide (Figure 2F and Supplementary Movie 1). We also observed that OLCs migrate into the injection lesion from outside of the dispersal region (Figure 2F and Supplementary Movie 1). To investigate if sox10+ cells also migrated from areas significantly anterior to the lesion, we used Tg(sox10:nls-eos) embryos to selectively label OLCs within a two-somite region encompassing the dispersal region. Following injection of lysolecithin at 4 dpf, we photoconverted the Eos protein, changing the green fluorescence to red via UV-light-induced photoconversion (Wiedenmann et al., 2004; Stys et al., 2012). We then used in vivo, time-lapse imaging to track the migration of both green (non-photoconverted) and red (photoconverted) OLCs with an imaging interval of 10 min (Figure 2G). Starting approximately 6.5 hpi, we observed that green sox10+ OLCs from anterior to the injection dispersal region migrate posteriorly into...
the lesion (Figure 2G). Interestingly, we rarely observed that OLCs within the lysolecithin dispersal region migrate during our imaging. This data demonstrates that a subset of sox10+ OLCs found in the lesion originate from areas anterior to the drug dispersal region. Taken together, these results demonstrate that lysolecithin affects sox10+ OLCs in the zebrafish spinal cord.

Lysolecithin Induces Myelin Membrane Changes in Zebrafish Larvae

Because we observed a reduction in Sox10+ OLC number and subsequent OPC proliferation and recruitment in lysolecithin-injected larvae, we next sought to investigate whether lysolecithin-induced myelin membrane changes by performing in vivo, time-lapse imaging in Tg(mbp:egfp-CAAX) (Almeida et al., 2011) larvae, where myelin basic protein (mbp) is labeled with membrane-tethered GFP. mbp is expressed in anterior regions of the zebrafish CNS beginning around 60 hpf (Almeida et al., 2011). At later stages, between 4 and 6 dpf, we observed stable GFP+ membrane sheaths in both the ventral and dorsal spinal cords (Czopka et al., 2011; Kearns et al., 2015; Figures 3A,B). Following injection of the PBS control solution into 4 dpf Tg(mbp:egfp-CAAX) larvae, we observed minimal to no changes to the GFP+ membrane near the injection site throughout the duration of our 16-h time-lapses with an imaging interval of 6 min (n = 6) (Figure 3C). In contrast, we observed distinct changes in GFP+ myelin membrane near the lesion after injection of the lysolecithin solution into the spinal cord of 4 dpf Tg(mbp:egfp-CAAX) larvae (n = 6) (Figure 3D). Specifically, we imaged myelin on somite anterior to the injection site so as to avoid possible damage induced by the injection itself. In these time-lapses, we observed that tight GFP+ membrane sheaths change after lysolecithin injection to take on a more ovoid-like appearance, beginning around 2.5 hpi (Figure 3D). Formation of the ovoid-like/onion bulb structures has previously been reported as evidence of myelin degradation (Acar et al., 2004), and we did not observe the same phenotype in control-injected larvae (Figure 3C).

Because the mbp+ membrane ovoids were obvious by 8 hpi in lysolecithin-injected larvae, we decided to quantify the percentage of zebrafish demonstrating GFP+ membrane changes within the dorsal spinal cord at this time point and observed a significant difference between the control and experimental groups, with approximately 17% of control zebrafish (n = 6) with GFP+ membrane changes within the dispersal area as compared with 60% of zebrafish injected with lysolecithin at 8 hpi (n = 5; Figure 3E). Additionally, we quantified the number of myelin ovoid structures within the control and experimental dispersal regions from 4 to 8 hpi, time points where the GFP+ membrane was dynamically changing in lysolecithin-injected larvae. In the lysolecithin-injected larvae, we observed an increase in the average number of GFP+ membrane ovoids that formed (n = 6 for both groups, at 6 hpi p = 0.0493, Figure 3F).

Because mbp:egfp-CAAX is a stable transgenic line, it is difficult to visualize individual oligodendrocyte membrane changes. Therefore, to investigate individual mbp+ membrane sheath dynamics, we injected a plasmid for mbp:mcherry-CAAX (Mensch et al., 2015) into one-cell embryos to mosaically label mbp+ oligodendrocytes in the spinal cord. At 6 dpf, we then injected either a PBS control solution or lysolecithin posterior to an mbp+ oligodendrocyte, such that the oligodendrocyte would reside anterior to the injection site but would be exposed to the injected solutions (Figure 3G). Using in vivo imaging with an image acquisition interval of 10 min, we observed an mbp+ ovoid form at approximately 8.1 hpi within the lysolecithin-injected group (n = 3; Figure 3G, bottom), but this phenotype was not observed in the control-injected larvae (n = 2; Figure 3G, top). Taken together, our results demonstrate that lysolecithin induces mbp+ oligodendrocyte membrane changes in the zebrafish spinal cord.

Axons Are Indistinguishable Between Lysolecithin- and Control-Injected Larvae

In mammals, previous studies demonstrate that lysolecithin causes minimal damage to axons (Blakemore et al., 1977; Keough et al., 2015). To investigate the effects of lysolecithin on axons in our focal injection model, we injected either a control solution or lysolecithin into the spinal cord of 4 dpf Tg(mbp:egfp-CAAX);Tg(cntn1b:mcherry) (Almeida et al., 2011; Czopka et al., 2013) larvae to label myelin with membrane-tethered GFP and axons with cytosolic mCherry. This transgenic line labels axons that have previously been described as being preferentially myelinated early in development in the zebrafish spinal cord (Czopka et al., 2013), and therefore, it allows us to specifically look at axonal integrity in relation to the lysolecithin-induced demyelination we observe. Using in vivo imaging, we observed that axons were indistinguishable between the control (n = 3; Figure 4A) and the lysolecithin-injected spinal cords (n = 2; Figure 4B) at 8 hpi, a time point in which we previously observed active mbp+ myelin changes via in vivo, time-lapse imaging (Figure 3D). From these data, we conclude that the effects observed on mbp+ membrane sheaths within the lysolecithin group described above were not secondary to axonal degeneration.

Professional Phagocytes Are Recruited to Lysolecithin-Induced Lesions

Macrophages and microglia respond to 1% lysolecithin injections in mammalian models and upon application to the adult zebrafish optic nerve and injection into the zebrafish spinal cord (Kotter et al., 2001; Münzel et al., 2014; Cunha et al., 2020). Therefore, we evaluated if macrophages and microglia were recruited to the focal lesion in our lysolecithin injection model in the larval zebrafish spinal cord.

To investigate professional phagocyte recruitment in our model, we injected either a control solution or lysolecithin into the spinal cord of 4 and 6 dpf Tg(mbp:egfp-CAAX);Tg(mpeg1:mcherry) (Almeida et al., 2011; Ellett et al., 2011) larvae, where mbp+ oligodendrocytes are labeled with membrane GFP, and macrophages/microglia were labeled with cytosolic mCherry. In both 4 and 6 dpf larvae, we observed that mpeg+ cells move quickly throughout the CNS, maneuvering through mbp+ myelin membrane layers. In an in vivo, time-lapse movie with an imaging interval of 6 min of a 4 dpf larva injected...
**FIGURE 3** | mbp+ membrane changes are observed following exposure to lysolecithin. All images are lateral views of the spinal cord with anterior to the left and dorsal to the top. (A,B) Expression of mbp:egfp-CAAX appears as sheath-like structures in the dorsal and ventral spinal cords in 4 (A) and 6 dpf (B) zebrafish larvae. (C) Following injection of a control solution into the spinal cord of a 4 dpf mbp:egfp-CAAX larva, the mbp+ membrane sheaths remain unchanged throughout the duration of the time-lapse movie. (D) Following injection of a lysolecithin (experimental) solution into the spinal cord of a 4 dpf mbp:egfp-CAAX larva, the mbp+ sheaths dynamically change to form ovoids (arrowheads). (E) Quantification of the percentage of zebrafish with mbp:egfp-CAAX membrane changes at 8 hpi. p < 0.0001; n = 6 control; n = 5 experimental. (F) Quantification of the average number of mbp:egfp-CAAX ovoids observed at 4, 6, and 8 hpi demonstrates an increase in the number of ovoids within the lysolecithin (experimental) group (n = 6) when compared with the control group (n = 6). At 6 hpi, *p* = 0.0493. (G) Mosaically labeled oligodendrocytes by injection of mbp:mcherry-CAAX. Larvae injected with a control solution (control, top) or lysolecithin (exp. group, bottom) at 6 dpf. Following injection of a control solution (top), the oligodendrocyte remains relatively unchanged throughout the course of the time-lapse movie. In contrast, an ovoid-like structure is observed within the mbp:mcherry-CAAX+ oligodendrocyte beginning around 8.1 hpi. Red dashed lines denote the spinal cord. Scale bars, 25 µm.
with lysolecithin, we observed that maneuvering mCherry\textsuperscript{+} phagocytes physically displaces mbp\textsuperscript{+} membrane sheaths while traversing through the dispersal region within the spinal cord (Figure 5A and Supplementary Movie 2).

We next quantified the number of mpeg\textsuperscript{+} phagocytes present in the lesion area to determine if larvae injected with lysolecithin had a greater response by professional phagocytes. Following injection of a control solution or lysolecithin into the spinal cord of 4 dpf Tg(mbp:egfp-CAAX);Tg(mpeg1:mcherry) larvae, we saw a modest increase in mpeg\textsuperscript{+} phagocytes within the lysolecithin-injected group from 2 to 6 hpi as compared with control injections (n = 6 control; n = 7 experimental; Figure 5B). Taken together, these results are consistent with the possibility that mpeg\textsuperscript{+} phagocytes are recruited to the lesion area, and this recapitulates the macrophage/microglia recruitment response that has been reported in mammalian lysolecithin models (Kotter et al., 2001; Münzel et al., 2014).

In addition to phagocyte migration toward the lesion, macrophages and microglia proliferate during demyelination and inflammation (Chiang et al., 1996; Remington et al., 2007; Jenkins et al., 2011). To investigate if this occurs in our lysolecithin model, we injected lysolecithin into the spinal cord of 4 dpf Tg(mbp:egfp-CAAX);Tg(mpeg1:mcherry) larvae and performed time-lapse imaging with an imaging interval of 15 min. In these movies, we captured mpeg\textsuperscript{+} cells proliferating within the lysolecithin dispersal region (Figure 6A and Supplementary Movie 3), demonstrating that professional phagocytes migrate into the lesion and proliferate in response to the lysolecithin insult.

Because we observed that mbp\textsuperscript{+} membrane changes within lysolecithin-injected larvae, we next wanted to determine if professional phagocytes were responding to clear oligodendrocyte/myelin debris. To capture the clearance of oligodendrocyte debris in vivo, we injected lysolecithin into 6 dpf Tg(sox10:mrfp; mpeg1:egfp) larvae and performed time-lapse imaging with an imaging interval of 10 min. As described above, mpeg\textsuperscript{+} cells were recruited to the lysolecithin dispersal region. By 3.6 hpi, we observed mpeg\textsuperscript{1+} cells with distinct cytoplasmic vacuoles surrounding sox10\textsuperscript{+} debris (Figure 6B and Supplementary Movie 4). Taken together, our results demonstrate that our focal lysolecithin injection model in zebrafish recapitulates the professional phagocyte response previously reported in lysolecithin-injected mammalian models (Kotter et al., 2001; Kucharova et al., 2011; Rawji and Yong, 2013; Doring et al., 2015).

**Oligodendrocyte Cytoskeleton Dynamics Following Exposure to Lysolecithin**

Actin dynamics are essential for myelin sheath formation (Nawaz et al., 2015; Samanta and Salzer, 2015; Zuchero et al., 2015). Specifically, filamentous actin (F-actin) is integral for initiating myelin wrapping and, upon completion of myelination, is disassembled from the inner tongue/leading edge and becomes localized to the lateral edges of the myelin sheath (Nawaz et al., 2015; Samanta and Salzer, 2015; Zuchero et al., 2015). Interestingly, actin polymerization is involved in peripheral myelin fragmentation and ovoid/onion bulb formation following injury in vitro, and inhibiting actin polymerization prevents myelin ovoid formation (Jung et al., 2011). Furthermore, cytoskeletal plasticity was recently identified as an important component of demyelination, suggesting that mechanisms within the oligodendrocyte cytoskeleton may be actively involved in reacting to demyelinating insults (Locatelli et al., 2015).
To investigate actin dynamics within sox10+ oligodendrocytes in vivo, we imaged with an interval of 10 min, control 6 dpf Tg(sox10:Gal4);Tg(UAS:Lifeact-gfp) larvae (Helker et al., 2013) and observed GFP+ F-actin arranged as sheath-like structures (Figure 7A), reminiscent of the appearance of mbp+ myelin membrane sheaths (Figure 3). To evaluate changes in F-actin, we injected lysolecithin into 6 dpf Tg(sox10:Gal4);Tg(UAS:Lifeact-gfp) larvae, a time point when the GFP+ sheath-like structures were stable in control-injected larvae. At 2 hpi, GFP+ F-actin within sox10+ cells appeared sheath-like. However, by 4 hpi, we observed GFP+ F-actin rearranging from a sheath-like structure to form ovoid-like structures by 8 hpi (Figure 7B). The arrangement of the F-actin from sheath-like into ovoid-like structures is reminiscent of the mbp+ membrane changes described above and occurred at a similar time point post-injection. These results suggest that the dynamic responses from components of the oligodendrocyte cytoskeleton appear to be involved in the lysolecithin-induced membrane changes that we observed post-injection.

DISCUSSION
Toxin-Induced Animal Models for Recapitulating CNS Damage
The use of toxin-induced demyelination models has been imperative for investigating mechanics underlying oligodendrogiopathies as well as processes involved in remyelination (Blakemore et al., 1977; Foote and Blakemore, 2005; Blakemore and Franklin, 2008; Ransohoff, 2012). Although toxin-induced demyelination models lack an ongoing immune response, there is a unique advantage of being able to separate demyelination events from processes underlying remyelination for studying neurodegenerative disorders, such as MS (Miller and Fyffe-Maricich, 2010; Ransohoff, 2012). Because of this
spatiotemporal predictableness, toxin-induced demyelination models can also be useful for screening therapeutic strategies that promote remyelination (Ransohoff, 2012).

Mammalian models commonly used to study MS, including EAE and toxin-induced demyelination models, as well as clinical samples, are not amenable for investigating mechanics and processes underlying myelin destruction and subsequent repair \textit{in vivo}. By developing a focal lysolecithin injection model in zebrafish, we can visualize \textit{mbp} \textsuperscript{+} membrane changes in a live, intact vertebrate model system and investigate mechanics and cellular interactions involved in myelin breakdown and repair. Recently, a paper using injection of lysolecithin into larval zebrafish trunks also described demyelination, and their data is similar to ours (Cunha et al., 2020); in the future, we envision that these focal injection models could be used as a first-pass drug-screening tool to investigate the safety and efficacy of novel compounds that either inhibit lysolecithin-induced damage, or promote its repair, to potentially reveal therapeutic strategies for treating demyelinating disorders, such as MS.

\textbf{Observing \textit{mbp} \textsuperscript{+} Membrane Changes \textit{in vivo}}

Intramyelinic vacuoles and fluid accumulation within the myelin sheath have been reported following lysolecithin-induced demyelination (Triarhou and Herndon, 1985) and in other toxin-induced demyelination models (Blakemore, 1973, 1974). In our focal lysolecithin injection model, we were able to visualize \textit{mbp} \textsuperscript{+} membrane changes and observed that these sheaths change to form ovoid-like vacuoles \textit{in vivo}, in real time. However, future studies are needed to confirm that the ovoids we see in our \textit{in vivo} imaging assays are from intramyelinic edema (Hirano et al., 1965; Hirano and Llena, 2006; Kaufmann et al., 2012).

Additionally, the dynamic rearrangement of F-actin within the oligodendrocyte cytoskeleton suggests that actin polymerization may be involved with the formation of the \textit{mbp} \textsuperscript{+} ovoids that we observed in our \textit{in vivo}, time-lapse imaging. Actin dynamics are essential for myelin sheath formation, and it is reasonable to assume that they play an integral role in kinetics underlying myelin membrane breakdown and repair following a toxin-induced insult. Additionally, after \textit{in vitro} peripheral nerve injuries, actin polymerization is involved in the formation of onion bulbs, and inhibiting actin polymerization prevents their formation (Jung et al., 2011), demonstrating that actin polymerization is actively involved in myelin membrane sheath changes. Mechanisms within the oligodendrocyte cytoskeleton may also be actively involved in reacting to insults as β-actin and β-tubulin gene transcription are upregulated in response to diphtheria-toxin oligodendrocyte death and EAE autoimmune inflammation (Locatelli et al., 2015). To our knowledge, there have been no reports on actin dynamics following lysolecithin-induced damage. Thus, future studies are needed in

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\caption{Professional phagocytes proliferate and clear Sox10\textsuperscript{+} debris. All images are lateral views of the spinal cord with anterior to the left and dorsal to the top. (A) Following injection of lysolecithin into a 4 dpf zebrafish spinal cord, a mCherry\textsuperscript{+} phagocyte recruited to the lysolecithin lesion proliferates (arrowheads). (B) Following injection of lysolecithin into a 6 dpf zebrafish spinal cord, sox10\textsuperscript{+} debris is cleared by an mpeg\textsuperscript{+} cell. Arrowheads identify the sox10\textsuperscript{+} debris within mpeg\textsuperscript{+} vacuoles in single z-planes. Yellow dashed lines denote the spinal cord. Scale bars, 25 µm.}
\end{figure}
FIGURE 7 | Oligodendrocyte cytoskeletal components dynamically change following exposure to lysolecithin. All images are lateral views of the spinal cord with anterior to the left and dorsal to the top. (A) In vivo imaging of Tg(sox10:Gal4; UAS:Lifeact-gfp) larvae at 6 dpf. Arrowheads denote the actin in sheath formation.

(B) Following injection of lysolecithin into a 6 dpf Tg(sox10:Gal4; UAS-Lifeact-gfp) zebrafish, in vivo time-lapse imaging reveals that GFP⁺ F-actin dynamically changes from a sheath-like arrangement (2 hpi) to forming ovoid-like structures (8 hpi). Arrowheads identify the GFP⁺ F-actin changes in a sox10⁺ cell. Scale bars, 25 µm.

our model of demyelination to determine if actin dynamics are actively rearranging the myelin sheath to form these ovoid-like structures, or are simply a passive bystander that dynamically rearranges as a result of the myelin sheath changes following lysolecithin-induced damage.

**Professional Phagocyte Response Following Lysolecithin Insult**

A previous report demonstrates a rapid recruitment of macrophages and microglial cells following injection of lysolecithin into the mouse spinal cord (Ousman and David, 2000). In the study by Ousman and David (2000), phagocytes were observed within the injection region by 6 hpi, and ultrastructural analysis at 4 dpi revealed macrophages containing myelin debris. Microglia/macrophages have also been shown to proliferate extensively in early demyelination (Matsumoto et al., 1992; Schönrock et al., 1998; Raivich and Banati, 2004) as well as in the EAE model (Rinner et al., 1995) and lysolecithin-induced demyelination in primates (Dousset et al., 1995). In the adult zebrafish optic nerve as well as the larval spinal cord, there is an increase in the number of professional phagocytes (Münzel et al., 2014; Cunha et al., 2020).

In our model, we also observe that mpeg1⁺ professional phagocytes respond to the injection region rapidly. The mpeg1⁺ professional phagocytes migrated to the lesion and proliferated, as previously reported in a primate model of lysolecithin-induced demyelination (Dousset et al., 1995). Since lysolecithin acts as a chemotactic factor for peripheral macrophages (Quinn et al., 1988), we expected to see an increase in the number of phagocytes within the lysolecithin lesion. Although there was an increase in the average number of professional phagocytes that...
we observed from 2 to 6 hpi, the results were not statistically significant between the control and lysolecithin-injected larvae, likely because the control larvae also experienced an injury from the injection site.

Phagocytes have been described as having both beneficial and detrimental roles in MS (Huitinga et al., 1990; Bitsch et al., 2000; Hinks and Franklin, 2000; Ousman and David, 2000; Arnett et al., 2011) and \textit{Tg} \textit{Travnickova et al., 2015} by crossing beneficial phagocytic responses to detrimental behaviors \textit{in vivo} offer a unique advantage for investigating the transition from beneficial phagocytic responses to detrimental behaviors \textit{in vivo} as macrophages can be depleted using a nitroreductase genetic model (Pisharath et al., 2007; Curado et al., 2008; Pisharath and Parsons, 2009), allowing for inducible and reversible depletion of \textit{mpeg1}+ cells. \textit{Escherichia coli} nitroreductase can be expressed under the control of the \textit{mpeg1} promoter (Palha et al., 2013; Travnickova et al., 2015) by crossing \textit{Tg} (mpeg1:Gal4FF)$^{25}$ (Ellett et al., 2011) and \textit{Tg}(UAS:nls-mCherry)$^{26}$ (Davison et al., 2007) lines together. To initiate cell depletion, larvae are immersed in the prodrug metronidazole, and nitroreductase converts the metronidazole into a cytotoxin, resulting in cell death. The effects are reversible upon removing the zebrafish from the prodrug solution. This genetic ablation model allows for spatial and temporal control over cell ablation and can be used in our focal lysolecithin injection model to investigate the impact that the absence of \textit{mpeg1}+ cells has on de- and remyelination events in real time, in a living vertebrate organism, feat mammalian systems currently cannot offer. These types of experiments will provide us with a better understanding about the roles and temporal dynamics of professional phagocytes in lysolecithin-induced injury and repair.

In conclusion, the lysolecithin injection model we developed using zebrafish larvae recapitulates phenotypes that have been described in mammalian systems following injection of lysolecithin, with a major advantage of being able to visualize damage in real time, \textit{in vivo}. An important difference is that our study uses larvae and most mammalian studies utilize adults. Future studies investigating the temporal dynamics of juvenile mammalian demyelination would be interesting given the rapid responses we see in our model. Finally, this focal injection model may become a useful drug-screening assay for revealing therapeutic strategies that help to prevent damage in myelinating glia or accelerate and enhance repair kinetics following damage.

**EXPERIMENTAL PROCEDURES**

**Fish Husbandry**

Animal studies were approved by the University of Virginia Institutional Animal Care and Use Committee. Zebrafish strains used in this study included \textit{Tg}(mbp:eGFP-CAAX) (Almeida et al., 2011), \textit{Tg}(sox10:7.2:mrfp)$^{nu234}$ (Kucenas et al., 2008), \textit{Tg}(cntn1b:mCherry) (Czopka et al., 2013), \textit{Tg}(mpeg1:mCherry) (Ellett et al., 2011), \textit{Tg}(mpeg1:eGFP) (Ellett et al., 2011), \textit{Tg}(sox10:nls-eos) (McGraw et al., 2012), \textit{Tg}(sox10:Gal4-VP16) (Chung et al., 2013), and \textit{Tg}(UAS:Lifect-GFP)$^{nu271}$ (Helker et al., 2013). Table 1 identifies the expression of all lines used in this study. Embryos were produced by pairwise matings, raised at 28.5°C in egg water, staged according to hpf or dpf, respectively, and embryos of either sex were used for all experiments (Kimmel et al., 1995). Embryos used for lysolecithin injections, immunohistochemistry, and microscopy were treated with 0.003% phenylthiourea (PTU) in egg water to inhibit pigmentation.

### Table 1

| Transgene name | Description of expression |
|----------------|---------------------------|
| \textit{Tg}(mbp:eGFP-CAAX) | Membrane eGFP in \textit{mbp}+ cells (oligodendrocytes) |
| \textit{Tg}(mpeg1:mCherry) | Cytosolic mCherry in \textit{mpeg1}+ cells (macrophages, microglia) |
| \textit{Tg}(mpeg1:eGFP) | Cytosolic eGFP in \textit{mpeg1}+ cells (macrophages, microglia) |
| \textit{Tg}(sox10:nls-eos)$^{nu18}$ | Nuclear localized Eos in \textit{sox10}+ cells (OPCs, oligodendrocytes) |
| \textit{Tg}(sox10:mRFP)$^{nu234}$ | Membrane RFP in \textit{sox10}+ cells (OPCs, oligodendrocytes) |
| \textit{Tg}(sox10:Gal4) | Gal4 specifically expressed in \textit{sox10}+ cells (OPCs, oligodendrocytes) |
| \textit{Tg}(UAS:Lifect-GFP)$^{nu271}$ | UAS promoter driving Lifect-GFP to label filamentous actin (F-actin) |
| \textit{Tg}(cntn1b:mCherry) | Cytosolic mCherry in mature neurons and axons |

All lines were stable, germline transgenics. Only cell types pertinent to this study are listed for each transgene.
containing either the control or lysolecithin solution, was inserted into the larvae, starting from the dorsal spinal cord and moving ventrally within the spinal cord, between somites 16 and 18. Solutions were dispersed passively into the spinal cord at a pressure range below 16 psi, until the dispersal region spanned the width of one to two somites. Following injections, the glass coverslip containing the mounted larvae was removed from the microscope stage and placed into a Petri dish (Falcon Sterile Petri Dish, 100 mm × 15 mm style, REF 351029) withegg water. The coverslip and larvae were completely submerged in egg water until the agar pad was saturated, resulting in the larvae floating off the agar pad. A pipette pump (Bel-Art, Scienceware Pipette Pump 10 ml Pipettor, Cat. No. 378980000) with a 2 ml glass Pasteur pipet tip (VWR Pasteur Pipet Disposable BBD Borosilicate Glass, 5 3/4 inch size, Cat. No. 53283-916) was used to transfer the larvae into a separate Petri dish with egg water to allow the larvae to recover.

**In vivo Imaging**

All embryos used for live imaging were transferred to egg water containing PTU at 24 hpf to inhibit pigment formation. At specified stages, embryos and larvae were anesthetized using 3-aminobenzoic acid ester (Tricaine), immersed in 0.8% low-melting point agarose on glass-bottomed 35 mm Petri dishes (Electron Microscopy Sciences). Images were captured using either a 40x (numerical aperture 1.2) or a 63x (numerical aperture 1.2) water-immersion objective mounted on a motorized Zeiss Axio Observer Z1 microscope equipped with a Quorum WaveFX-X1 spinning disc confocal system (Quorum Technologies). For time-lapse imaging, z-stacks were collected at specified intervals, and three-dimensional datasets were compiled using MPEG-4 video compression at 10 frames per second (fps) and exported to QuickTime (Apple) to create movies. Image adjustments were limited to contrast enhancement and level settings using MetaMorph software, Adobe Photoshop, and ImageJ.

**Whole Mount Immunohistochemistry**

Larvae were fixed in AB Fix [4% paraformaldehyde (PFA), 0.1% Triton-X 100, 1x PBS] for either 3 h at 23°C or overnight at 4°C, followed by a 5-min wash with PBSTx (1% Triton X-100, 1x PBS), a 5-min wash with DWTx (1% Triton X-100 in distilled water), a 5-min wash with acetone at 23°C, a 10-min wash with acetone at −20°C, and three 5-min washes with PBSTx. Larvae were preblocked in 5% goat serum/PBSTx for at least 1 h and incubated in primary antibody for 1 h at 23°C and overnight at 4°C. The primary antibody used in this study was a rabbit antibody to Sox10 (1:5,000) (Binari et al., 2013). Larvae were washed extensively with 1x PBSTx and stored in 50% glycerol–PBS at 4°C until imaging. Larvae were mounted on their sides in 0.8% low-melting point agarose on glass-bottomed 35 mm Petri dishes and imaged using the confocal microscope described above. Image adjustments were limited to contrast enhancement and level settings using MetaMorph software, Adobe Photoshop, and ImageJ.

**Data Quantification and Statistical Analysis**

All graphically presented data represent the mean of the analyzed data. Statistical analyses and graphing were performed with GraphPad Prism software. The level of significance was determined by using an unpaired t-test or a chi-square test using a confidence interval of 95%. The data in plots and the text are presented as mean ± SEM.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author/s.

**ETHICS STATEMENT**

The animal study was reviewed and approved by the University of Virginia Institutional Animal Care and Use Committee.

**AUTHOR CONTRIBUTIONS**

AM conducted and analyzed all of the experiments. Both authors conceived the study and wrote the manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fcell.2021.654583/full#supplementary-material

**Supplementary Movie 1** | Sox10+ cells proliferate and migrate after injection of lysolecithin into the larval spinal cord. Panels are partially overlapping regions of the spinal cord in lysolecithin-injected (blue region) 4 dpf Tg(sox10:nls-eos) larvae imaged from 2 to 10 hpi. The right edge of the left panel is the left edge of the right panel. OLC proliferation is denoted by pink arrowheads (left panel), and a migratory OLC is denoted by a green dot (both panels). Images were taken every 8 min, and the movie runs at 10 fps.

**Supplementary Movie 2** | mpeg+ phagocytes migrate through myelin layers in lysolecithin-injected spinal cords. Lysolecithin was injected into 4 dpf zebrafish larvae.
REFERENCES

Acar, G., Tanriver, G., Demir, N., Kayisli, U. A., Satir, G. L., Yaba, A., et al. (2004). Ultrastructural and immunohistochemical similarities of two distinct entities: multiple sclerosis and hereditary motor sensory neuropathy. Acta Histochem. 106, 363–371. doi: 10.1016/j.acthis.2004.08.004

Almeida, R. G., Czopka, T., Ffrench-Constant, C., and Lyons, D. A. (2011). Individual axons regulate the myelinating potential of single oligodendrocytes in vivo. Development 138, 4443–4450. doi: 10.1242/dev.071001

Arnett, H. A., Mason, J., Marino, M., Suzuki, K., Matsushima, G. K., and Ting, J. P. (2001). TNF alpha promotes proliferation of oligodendrocyte progenitors and remyelination. Nat. Neurosci. 4, 1116–1122. doi: 10.1038/nn738

Azin, M., Mirnajafi-Zadeh, J., and Javan, M. (2015). Fibroblast growth factor-2 enhanced the recruitment of progenitor cells and myelin repair in experimental demyelination of rat hippocampal formations. Cell J. 17, 540–546. doi: 10.2207/cell.2015.14

Binari, L. A., Lewis, G. M., and Kucenas, S. (2013). Perineurial glia require Notch signaling during motor nerve development but not regeneration. J. Neurosci. 33, 4241–4252. doi: 10.1523/JNEUROSCI.4893-12.2013

Bitsch, A., Kuhlmann, T., Da Costa, C., Bunkowski, S., Polak, T., and Brück, W. (2000). Tumour necrosis factor alpha mRNA expression in early multiple sclerosis lesions: correlation with demyelination activity and oligodendrocyte pathology. Glia 29, 366–375.

Blakemore, W. F. (1973). Demyelination of the superior cerebellar peduncle in old age. J. Neurol. Sci. 193–212. doi: 10.1016/0022-510x(73)90118-4

Blakemore, W. F. (1974). Remyelination of the superior cerebellar peduncle in old mice following demyelination induced by cuprizone. J. Neurol. Sci. 22, 121–126. doi: 10.1016/0022-510x(74)90059-8

Blakemore, W. F., Eames, R. A., Smith, K. J., and McDonald, W. I. (1977). Remyelination of the superior cerebellar peduncle in old age. J. Neurol. Sci. 22, 121–126. doi: 10.1016/0022-510x(77)90179-4

Blakemore, W. F., and Franklin, R. J. (2000). Delayed changes in growth factor gene expression after elimination of macrophages. J. Exp. Med. 193, 1025–1033. doi: 10.1084/jem.20000926

Czopka, T., and Lyons, D. A. (2011). Dissecting mechanisms of myelinated axon formation using zebrafish. Methods Cell Biol. 105, 25–62. doi: 10.1016/b978-0-12-381320-6.00002-3

Davidson, J. M., Akitake, C. M., Goll, M. G., Rhef, J. M., Gosse, N., Baier, H., et al. (2007). Transactivation from Gal4-VP16 transgenic insertions for tissue-specific cell labeling and ablation in zebrafish. Dev. Biol. 304, 811–824. doi: 10.1016/j.ydbio.2007.01.033

Domingues, H. S., Portugal, C. C., Socodato, R., and Relvas, J. B. (2016). Oligodendrocyte, astrocyte, and microglia crosstalk in myelin development, damage, and repair. Front. Cell Dev. Biol. 4:71. doi: 10.3389/fcel.2016.0071

Doring, A., Sloka, S., Lau, L., Mishra, M., van Minnen, J., Zhang, X., et al. (2015). Stimulation of monocytes, macrophages, and microglia by amphotericin B and macrophage colony-stimulating factor promotes remyelination. J. Neurosci. 35, 1136–1148. doi: 10.1523/JNEUROSCI.1797-14.2015

Dousset, V., Brochet, B., Vitali, A., Grossi, C., Benazzouz, A., Boulouerne, A., et al. (1995). Myeloid-derived remyelination in primates: preliminary in vivo study with MR and magnetization transfer. Am. J. Neuroradiol. 16, 225–231.

Ellett, F., Pase, L., Hayman, J. W., Andrianopoulos, A., and Lieschke, G. J. (2011). mpeg1 promoter transgenes directly control macrophage lineage differentiation in zebrafish. Blood 117, e49–e56. doi: 10.1182/blood-2010-10-314120

Fang, Y., Lei, X., Li, X., Chen, Y., Xu, F., Feng, X., et al. (2014). A novel model of demyelination and remyelination in a GFP-transgenic zebrafish. Biol. Open 4, 62–68. doi: 10.1242/bio.20141076

Foa, K. E., and Blakemore, W. F. (2005). Inflammation stimulates remyelination in areas of chronic demyelination. Brain 128(Pt 3), 528–539. doi: 10.1093/brain/awv417

Girard, C., Remelmann, A. P., Dufoir, N., Mallet, J., Bachelin, C., Nait-Oumesmar, B., et al. (2005). Grafts of brain-derived neurotrophic factor and neurotrophin 3-transduced primate Schwann cells lead to functional recovery of the demyelinated mouse spinal cord. J. Neurosci. 25, 7924–7933. doi: 10.1523/JNEUROSCI.4890-04.2005

Gregg, C., Shikar, V., Larsen, P., Mak, G., Chojnacki, A., Yong, V. W., et al. (2007). White matter plasticity and enhanced remyelination in the maternal CNS. J. Neurosci. 27, 1812–1823. doi: 10.1523/JNEUROSCI.4441-06.2007

Helker, C. S., Schuermann, A., Karpanen, T., Zeuschner, D., Belting, H. G., Aifolter, M., et al. (2013). The zebrafish common cardinal veins develop by a novel mechanism: lumen ensheathment. Development 140, 2776–2786. doi: 10.1242/dev.019876

Hill, K. E., Zollinger, L. V., Watt, H. E., Carlson, N. G., and Rose, J. W. (2004). Inducible nitric oxide synthase in chronic active multiple sclerosis plaques: distribution, cellular expression and association with myelin damage. J. Neuroimmunol. 151, 171–179. doi: 10.1016/j.jneuroim.2004.02.005

Hinks, G. L., and Franklin, R. J. (2000). Delayed changes in growth factor gene expression during slow remyelination in the CNS of aged rats. Mol. Cell Neurosci. 16, 542–556. doi: 10.1006/mcne.2000.0897

Hirano, A., and Llena, J. (2006). Fine structure of neuronal and glial processes in neuropathology. J. Neuroimmunol. 151, 171–179. doi: 10.1016/j.jneuroim.2004.02.005

Hirano, A., Zimmermann, H. M., and Levine, S. (1965). Fine structure of cerebral fluid accumulation: VI. Intracellular accumulation of fluid and Cryptococcal polysaccharide in oligodendroglia. Arch. Neurol. 12, 189–196. doi: 10.1001/archnev.1965.00400260079009

Huitinga, I., van Rooijen, N., de Groot, C. J., Uitdehaag, B. M., and Dijkstra, C. D. (1990). Suppression of experimental allergic encephalomyelitis in Lewis rats after elimination of macrophages. J. Exp. Med. 172, 1025–1033. doi: 10.1084/jem.172.4.1025
Kearns, C. A., Ravanelli, A. M., Cooper, K., and Appel, B. (2015). Fbxw7 limits demyelination by inhibiting mTOR signaling. J. Neurosci. 35, 14861–14871. doi: 10.1523/jneurosci.4968-14.2015

Keough, M. B., Jensen, S. K., and Yong, V. W. (2015). Experimental demyelination regenerates CNS correlates with growth of axons in caliber. PLoS One 12:e0178058. doi: 10.1371/journal.pone.0178058

Karttunen, M. J., and Lyons, D. A. (2019). A drug-inducible transgenic zebrafish model for myelinating glial cell ablation. Methods Mol. Biol. 203, 253–310. doi: 10.1007/978-1-4939-9072-6_13

Kamm, W., Bolon, B., Bradley, A., Butt, M., Czasch, S., Garman, R. H., et al. (2012). Proliferative and nonproliferative lesions of the rat and mouse central and peripheral nervous systems. Toxicol. Pathol. 40(4 Suppl), 875–1575. doi: 10.1177/0190223612439125

Kears, C. A., Ravaneilli, A. M., Cooper, K., and Appel, B. (2015). Fbxw7 limits myelination by inhibiting mTOR signaling. J. Neurosci. 35, 14861–14871. doi: 10.1523/jneurosci.4968-14.2015

Kotter, M. R., Zhao, C., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs oligodendrocyte remyelination following lysolecithin-induced demyelination. Glia 55, 204–212. doi: 10.1002/glia.1085

Kotter, M. R., Zhao, C., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Kucenas, S., Takada, N., Park, H.-C. C., Woodruff, E., Broadie, K., and Appel, B. (2008). CNS-derived glia ensheath peripheral nerves and mediate motor root development. Nat. Neurosci. 11, 143–151.

Kucharova, K., Chang, Y., Boor, A., Yong, V. W., and Staluppi, W. B. (2011). Reduced inflammation accompanies diminished myelin damage and repair in the NG2 null mouse spinal cord. J. Neuroinflamm. 8:158. doi: 10.1186/1742-2094-8-158

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.

Lassmann, H., van Rooijen, N., and Franklin, R. J. M. (2005). Macrophage-depletion impairs remyelination of experimental CNS remyelination is associated with a reduced oligodendrocyte progenitor cell response and altered growth factor expression. Neurobiol. Dis. 18, 166–175.
Schönrock, L. M., Kuhlmann, T., Adler, S., Bitsch, A., and Brück, W. (1998). Identification of glial cell proliferation in early multiple sclerosis lesions. *Neuropathol. Appl. Neurobiol.* 24, 320–330. doi: 10.1046/j.1365-2990.1998.00131.x

Stys, P. K., Zamponi, G. W., van Minnen, J., and Geurts, J. J. (2012). Will the real multiple sclerosis please stand up? *Nat. Rev. Neurosci.* 13, 507–514. doi: 10.1038/nrn3275

Travnickova, J., Tran Chau, V., Julien, E., Mateos-Langerak, J., Gonzalez, C., Lelièvre, E., et al. (2015). Primitive macrophages control HSPC mobilization and definitive haematopoiesis. *Nat. Commun.* 6:6227. doi: 10.1038/ncomms7227

Triarhou, L. C., and Herndon, R. M. (1985). Effect of macrophage inactivation on the neuropathology of lysolecithin-induced demyelination. *Br. J. Exp. Pathol.* 66, 293–301.

Waller, A. (1851). Experiments on the section of the Glossopharyngeal and Hypoglossal nerves of the frog, and observations of the alterations produced thereby in the structure of their primitive fibres. *Edinb. Med. Surg. J.* 76, 369–376.

Wiedenmann, J., Ivanchenko, S., Oswald, F., Schmitt, F., Röcker, C., Salih, A., et al. (2004). EosFP, a fluorescent marker protein with UV-inducible green-to-red fluorescence conversion. *Proc. Natl. Acad. Sci. U.S.A.* 101, 15905–15910. doi: 10.1073/pnas.0403668101

Zuchero, J. B., Fu, M. M., Sloan, S. A., Ibrahim, A., Olson, A., Zaremba, A., et al. (2015). CNS myelin wrapping is driven by actin disassembly. *Dev. Cell* 34, 152–167. doi: 10.1016/j.devcel.2015.06.011

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

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