N-Wasp Regulates Oligodendrocyte Myelination

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Oligodendrocyte myelination depends on actin cytoskeleton rearrangement. Neural Wiskott-Aldrich syndrome protein (N-Wasp) is an actin nucleation factor that promotes polymerization of branched actin filaments. N-Wasp activity is essential for myelin membrane wrapping by Schwann cells, but its role in oligodendrocytes and CNS myelination remains unknown. Here we report that oligodendrocytes-specific deletion of N-Wasp in mice of both sexes resulted in hypomyelination (i.e., reduced number of myelinated axons and thinner myelin profiles), as well as substantial focal hypermyelination reflected by the formation of remarkably long myelin outffolds. These myelin outffolds surrounded unmyelinated axons, neuronal cell bodies, and other myelin profiles. The latter configuration resulted in pseudo-multimyelin profiles that were often associated with axonal detachment and degeneration throughout the CNS, including in the optic nerve, corpus callosum, and the spinal cord. Furthermore, developmental analysis revealed that myelin abnormalities were already observed during the onset of myelination, suggesting that they are formed by aberrant and misguided elongation of the oligodendrocyte inner lip membrane. Our results demonstrate that N-Wasp is required for the formation of normal myelin in the CNS. They also reveal that N-Wasp plays a distinct role in oligodendrocytes compared with Schwann cells, highlighting a difference in the regulation of actin dynamics during CNS and PNS myelination.

Key words: actin dynamics; axon-glia; myelin; oligodendrocyte; Schwann cell

Significance Statement

Myelin is critical for the normal function of the nervous system by facilitating fast conduction of action potentials. During the process of myelination in the CNS, oligodendrocytes undergo extensive morphological changes that involve cellular process extension and retraction, axonal ensheathment, and myelin membrane wrapping. Here we present evidence that N-Wasp, a protein regulating actin filament assembly through Arp2/3 complex-dependent actin nucleation, plays a critical role in CNS myelination, and its absence leads to several myelin abnormalities. Our data provide an important step into the understanding of the molecular mechanisms underlying CNS myelination.

Introduction

Myelin is a specialized membrane produced by oligodendrocytes and Schwann cells that spiral around axons, thereby enabling fast conduction of action potentials, and providing axonal support (Nave and Werner, 2014; Cohen et al., 2020). During myelination in the CNS, oligodendrocytes undergo extensive morphological changes (Bauer et al., 2009; Seixas et al., 2019; Brown and Macklin, 2020), beginning with the formation of exploratory processes that either make stable axonal contact or retract (Czopka et al., 2013; Almeida and Lyons, 2014). This initial contact is then followed by membrane ensheathment and wrapping, longitudinal extension of the forming myelin unit, and compaction of the myelin membrane layers (Osso and Chan, 2017; Stadelmann et al., 2019). Oligodendrocytes contain two major cytoskeletal systems, microtubules and actin filaments, which regulate process formation and myelination (Bauer et al., 2009; Seixas et al., 2019; Brown and Macklin, 2020). The actin cytoskeleton has a higher turnover and reorganization rate than microtubules, enabling fast reshaping of myelinating oligodendrocytes (Song et al., 2001). Microtubules provide support to the morphology of the cell and allow transport of myelin-specific proteins and mRNA (Lunn et al., 1997; Seiberlich et al., 2015). Moreover, microtubules and actin filaments interact with each other, leading to oligodendrocyte process outgrowth (Song et al., 2001). Previous studies highlight the role of actin dynamics during myelin formation and suggest a two-step model. First,
oligodendrocyte process outgrowth and the initial ensheathment of the axon are driven by Arp2/3 complex-dependent actin polymerization (Zuchero et al., 2015). Subsequently, lateral spreading and growth of the myelin membrane during wrapping and compaction requires F-actin disassembly by ADF/cofilin-1 (Nawaz et al., 2015; Zuchero et al., 2015).

Neural Wiskott-Aldrich syndrome protein (N-Wasp) is a nucleation-promoting factor that drives the generation of branched actin filaments (Alekhina et al., 2017). It regulates cortical actin filament reorganization in response to extracellular stimuli by linking between small GTPases (i.e., Rac and Cdc42) and actin polymerization through the regulation of the Arp2/3 complex (Takenawa and Miki, 2001). N-Wasp and Wasp family verprolin homologous protein-1 (WAVE1) are required for myelination in the CNS and CNS, respectively, and regulate oligodendrocyte differentiation (Kim et al., 2006; Bacon et al., 2007; Jin et al., 2011; Novak et al., 2011). N-Wasp is expressed by both oligodendrocytes and myelinating Schwann cells (Tsuchiya et al., 2006; Novak et al., 2011; Zhang et al., 2014; Marques et al., 2016). It plays a crucial role in Schwann cell membrane wrapping and in longitudinal extension of the myelin unit in the PNS, most likely via regulation of the actin cytoskeleton (Jin et al., 2011; Novak et al., 2011). Schwann cells from N-Wasp-mutant mice formed less radial lamellipodia and shorter axonal processes that lacked defined F-actin-rich cones (Jin et al., 2011). In the absence of N-Wasp, Schwann cells properly sort and ensheathe the axons, but were not capable of proceeding with myelin wrapping (Jin et al., 2011; Novak et al., 2011). In the CNS, N-Wasp was detected with components of the Arp2/3 complex in newly formed oligodendrocytes and in purified myelin fractions (Bacon et al., 2007). Moreover, pharmacological inhibition of N-Wasp prevented process extension and caused filopodium retraction in cultured oligodendrocyte precursor cells. Overall, these data suggest that cellular process extension by oligodendrocyte precursor cells is regulated by N-Wasp and Arp2/3-driven actin polymerization. Here we report that genetic deletion of N-Wasp in oligodendrocytes results in hypomyelination, diverse myelin abnormalities, including outfoldings that enwrap neuronal cell bodies, as well as myelin and axonal degeneration.

Materials and Methods

Mice. Generation of N-Wasp\textsuperscript{lox/lox} (Cotta-de-Almeida et al., 2007) and Cnp-Cre (Lappe-Siefke et al., 2003) mice were previously described and was always kept as heterozygous. Cnp-Cre/N-Wasp\textsuperscript{lox/ls} mice (i.e., homozygous for N-Wasp) were obtained by a conventional breeding scheme. Genotypes were determined by performing PCR on genomic DNA extracted from mice tails. Both male and female animals were used in the study, with no detectable difference in myelin morphology. Rotarod extract was previously described (Zuchero et al., 2015). Both male and female animals were used in the study, with no detectable difference in myelin morphology. Rotarod extract was previously described (Zuchero et al., 2015).

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due to its expression in other cell types, such as neurons and astrocytes. Cnp-Cre/N-Waspflx/flx mice exhibited a severe tremor and a hindlimb paralysis, and had shorter latency to fall from a rotarod compared with WT mice (Fig. 1C). These motor impairments resembled the phenotype of DHH-Cre/N-Waspflx/flx mice (Novak et al., 2011), and are likely attributed the expression of Cnp-Cre in myelinating Schwann cells. In agreement, EM analysis revealed that, similar to DHH-Cre/N-Waspflx/flx animals, sciatic nerves of adult Cnp-Cre/N-Waspflx/flx mice almost completely lacked myelin profiles (Fig. 1J).

In order to evaluate the contribution of N-Wasp to CNS myelination, we first examined sagittal brain sections prepared from 2-month-old mutant mice stained with LFB (Kluver and Barrera, 1953). While the overall brain morphology and the formation of white matter tracks were similar between WT and mutant brains, the latter clearly exhibited hypomyelination in several brain regions, including the corpus callosum and the hippocampal fimbria (Fig. 1D,E). Similarly, we noted a decrease in MBP immunoreactivity throughout the brain (Fig. 1F–I). We next examined optic nerves from 1-month-old WT and Cnp-Cre/N-Waspflx/flx mice by EM. As depicted in Figure 2A, B, Cnp-Cre/N-Waspflx/flx mice displayed pronounced hypomyelination in the optic nerve. These mice exhibited a lower number of myelinated axons (Fig. 2C), thinner myelin profiles (i.e., higher g-ratio; Fig. 2D,E), as well as an increase in the number of abnormal profiles (Fig. 2F; and detailed below) compared with their littermate control. Hypomyelination, increase in g-ratio, and a fourfold increase in the occurrence of additional myelin abnormalities were also detected in Cnp-Cre/N-Waspflx/flx mice at 4 months of age (Fig. 2G–J).

Mice lacking N-Wasp in oligodendrocytes exhibit diverse myelin abnormalities

EM analysis of different brain areas, including the corpus callosum, optic nerve, and the cerebellum, revealed that the most pronounced abnormality resulted from the absence of N-Wasp in oligodendrocytes was the formation of long myelin outfolds (Fig. 3A–F). We noted the presence of long myelin outfolds that extended throughout the neuropil (Fig. 3A), as well as myelin outfolds that surrounded either nonmyelinated (Fig. 3B) or myelinated (Fig. 3C) axons. The association between myelin outfolds and preexisting myelin sheath often resulted in the

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**Figure 1.** Absence of N-Wasp in oligodendrocytes results in hypomyelination. A, RT-PCR analysis of mRNA isolated from optic nerves at 2 months from Cnp-Cre/N-Waspflx/flx mutant and from N-Waspflx/flx (WT) mice. Amplification of N-Wasp and actin was performed using the relevant primers, and products of expected sizes were obtained. B, WB analysis of optic nerves isolated from 2-month-old Cnp-Cre/N-Waspflx/flx mutant and from N-Waspflx/flx (WT) mice using antibodies to N-Wasp and β-tubulin. C, Rotarod test performed with 3-month-old Cnp-Cre/N-Waspflx/flx mutant and from N-Waspflx/flx (WT) mice. The mutant mice had a shorter latency to fall than their littermate control. Error bars indicate SD of n = 5 mice for each genotype (*p < 0.05). D–I, Sagittal sections of brains from 2-month-old N-Waspflx/flx (WT; D,F,H) and Cnp-Cre/N-Waspflx/flx (E,G,I) mice stained with LFB and cresyl violet (D,E), or immunolabeled using antibodies to MBP (F–I). Hypomyelination is detected in the mutant corpus callosum and the fimbria (D–G, arrowheads). D, E, Insets, High-magnification image of the framed area in the corpus callosum. H, I, High-magnification images of the framed regions in F and G showing a decrease in MBP immunoreactivity in the absence of N-Wasp (arrowheads). J, Cnp-Cre/N-Waspflx/flx mice lack myelin profiles in the PNS. EM images of sciatic nerve cross sections from 2-month-old N-Waspflx/flx (WT) and Cnp-Cre/N-Waspflx/flx mutant mice. In mutant nerves, most Schwann cells were arrested at the promyelinating stage, similarly to our DHH-Cre/N-Waspflx/flx mutant nerves (Novak et al., 2011). Scale bars: D–G, 500 μm; Inset, 100 μm; H, I, 50 μm; J, 5 μm.
formation of a pseudo-multimyelin configuration around the axon (Fig. 3D). In these structures, the outer membrane of the myelin outfold contact the outer membrane of exiting myelin (i.e., reverse orientation), and the two inner membranes of the outfolding faced each other (Fig. 3D,M,N). Notably, these are likely distinct from the multimyelin configuration (Fig. 3L) formed by membrane slippage as a result of aberrant axoglial adhesion (Djannatian et al., 2019; Elazar et al., 2019a). In the cerebellum, we observed long myelin outfolds extending from the axon and either partially or completely surrounding neuronal somata in the granular layer (Fig. 3E,F). Throughout the brain and spinal cord, we detected pseudo-multimyelin profiles that were often accompanied by myelin deterioration (emerging cytoplasmic spaces between myelin sheaths), axonal detachment, and degeneration (Fig. 3G–J). We did not detect axonal pathology in non-myelinated fibers, suggesting that the axonal pathology observed in Cnp-Cre/N-Waspflx/flx mice is secondary to myelin disintegration. In support of this conclusion, the appearance of a degenerated axon was frequently associated with the presence of abnormal and degenerated myelin sheath around the axons.

Myelin abnormalities occur already at early developmental stage

We next examined whether the myelin pathology observed in the absence of N-Wasp occurred during the period of active developmental myelination in the optic nerve (Rashband et al., 1999). We performed EM analysis of P10 (Fig. 4A) and P15 (Fig. 4B) optic nerves isolated from WT and Cnp-Cre/N-Waspflx/flx mice to examine the number of myelinated axons, as well as myelin abnormalities. While there was no statistically significant difference in the number of myelinated profiles between the genotypes at P10 (Fig. 4C), the percentage of myelinated axons was significantly lower in the mutant than in WT optic nerves at P15 (Fig. 4E). Nevertheless, the number of myelin outfolds detected in optic nerves of Cnp-Cre/N-Waspflx/flx mutant mice was significantly higher than their age-matched control littersmates at both P10 (Fig. 4D) and P15 (Fig. 4F). Notably, we detected long projecting myelin outfolds (Fig. 4A, right), as well as multilayered and degenerated profiles (Fig. 4A, middle) in the mutant optic nerve already at P10, indicating that the absence of N-Wasp in oligodendrocytes leads to dysregulation of CNS myelination. In addition, while we detected intact axons that were surrounded by abnormal myelin, we rarely detected degenerated axons enwrapped by normal myelin, further supporting the notion that axonal degeneration in these mice is secondary to myelin impairment.

Absence of N-Wasp in oligodendrocytes leads to myelin abnormalities in multiple CNS areas

It was previously reported that deletion of WAVE1, which belongs to the same protein family of N-Wasp, resulted in hypomyelination only in certain regions in the CNS (Kim et al., 2006). To further examine whether deletion of N-Wasp also affects myelination in a regional manner, we performed EM analysis of the corpus callosum and spinal cord of WT and Cnp-Cre/N-Waspflx/flx mice. We detected severe myelin abnormalities, including the formation of outfolds, redundant myelin, and degeneration in both the corpus callosum (Fig. 5A,B) and spinal cord (Fig. 5C,D). Furthermore, in the spinal cord, myelin pathology and degeneration (Fig. 5E,F) were progressive and increased from 1- to 4-month-old animals (Fig. 5G). Together, our results show that, in contrast to the ablation of N-Wasp in Schwann cells, N-Wasp-deficient oligodendrocytes produce abundant
myelin. However, experimental deletion of N-Wasp in oligodendrocytes results in the formation of abnormal myelin membrane extensions and degeneration of myelinated axons, indicating that it is required for the normal CNS myelination.

**Discussion**

During CNS myelination, oligodendrocytes extend cellular processes that contact axons and form myelin internodes by continuously wrapping their membrane around them (Czopka et al., 2013; Snaidero et al., 2014; Stadelmann et al., 2019). These extensive morphological changes are tightly regulated by dynamic reorganization of the cortical actin cytoskeleton (Nawaz et al., 2015; Zuchero et al., 2015; Seixas et al., 2019; Brown and Macklin, 2020; Thomason et al., 2020). N-Wasp affects actin filament polymerization by regulating the nucleation activity of Arp2/3. We found that genetic ablation of N-Wasp in oligodendrocytes results in hypomyelination, extensive formation of myelin outfoldings, and degeneration of myelinated axons throughout the brain and spinal cord, demonstrating that N-Wasp plays a critical role in the formation of normal CNS myelin. Our data support previous studies indicating that the regulation of actin filament dynamics is critical for both the initial extension of cellular process during axonal selection, as well as for membrane ensheathment and wrapping during myelination (Nawaz et al., 2015; Zuchero et al., 2015; Seixas et al., 2019; Brown and Macklin, 2020; Thomason et al., 2020).

*Figure 3.* *Cap-Cre/N-Wasp<sup>flx/flx</sup> mice exhibit severe myelin abnormalities.* **A, B,** EM images of *Cap-Cre/N-Wasp<sup>flx/flx</sup>* corpus callosum at P15. The long extending myelin outfold originated from one axon (labeled a in B) surrounds a nonmyelinated axon (b), and extends an additional outfold (arrowhead) that contact this axon. **C, D,** High-magnification EM images of optic nerve cross sections obtained from 1-month-old *Cap-Cre/N-Wasp<sup>flx/flx</sup>* mutant mice, showing long myelin outfoldings. Myelin outfold cover other myelinated axons. **E, F,** Presence of oligodendrocyte myelin around neuronal cell bodies. EM images of the cerebellum of 4-month-old *Cap-Cre/N-Wasp<sup>flx/flx</sup>* mutant mice. The presence of myelin outfold that either partially (E) or completely (F) surrounding neuronal cell bodies. **G, Arrow indicates the presence of a nearby myelinated axon. High magnification of area indicated by arrowhead in F showing multiple myelin layers around the cell body.** **H-I,** High-magnification EM images of optic nerve cross sections obtained from P10 (G), P15 (H), and P30 (I) *Cap-Cre/N-Wasp<sup>flx/flx</sup>* mutant mice showing different configurations of multmyelinated axons undergoing degeneration and axon (red asterisk) detachment. **J,** EM image of optic nerve cross sections obtained from 4-month-old *Cap-Cre/N-Wasp<sup>flx/flx</sup>* mutant mice showing multiple profiles of focal hypermyelination and degeneration (red asterisk). **K-N,** Schematic model showing normal myelin (K), multmyelin configuration resulting from aberrant adhesion (L), and pseudo-multimyelin configuration formed by wrapping of a myelin outfolding around myelinated axon (M, N). **K,** The myelin sheath (Myelin) and the location of the most inner and outer myelin membranes. Scale bars: **A, B, D-I,** 1 μm; **C,** 0.5 μm; **J,** 2 μm.
The absence of N-Wasp in oligodendrocytes leads to hypomyelination (as reflected by a lower number of myelinated axons and thinner myelin sheath), is also in line with in vitro observations showing that chemical inhibition of N-Wasp in cultured oligodendrocytes by wiskostatin, which prevents activation of the Arp2/3 complex, results in the inhibition of process extension (Bacon et al., 2007). Studies on actin dynamics in developing oligodendrocytes and myelination suggest a role for Arp2/3-dependent actin polymerization during the initial stages of axonal contact and ensheathment, following by actin filament disassembly, which decreases membrane surface tension and promotes radial movement of the membrane during myelin wrapping (Nawaz et al., 2015; Zuchero et al., 2015). In agreement with a spatial role of actin polymerization during the early phase of myelination, N-Wasp along with Arp2/3 complex components were both noted at the edges of cellular extensions in cultured oligodendrocytes (Bacon et al., 2007). In addition, the expression of N-Wasp is higher in premyelinating and newly formed oligodendrocytes than in myelinating oligodendrocytes (Zhang et al., 2014; Seixas et al., 2019). Our findings also support a role for N-Wasp during myelin membrane wrapping, either by inducing actin assembly needed for continuous actin turnover at the edge of the inner lip (Nawaz et al., 2015), or by contributing to the cortical actin scaffold required for a local actin depolymerization-dependent progression of the inner lip (Zuchero et al., 2015). The remarkable similarity between Cnp-Cre/N-Waspflx/flx and Cnp-Cre/N-WaspCre/N-Wasp alleles (Zuchero et al., 2015) strongly suggests that the role N-Wasp plays in myelination is mediated by actin polymerization via the Arp2/3 complex.

One of the most prominent abnormalities detected in Cnp-Cre/N-Waspflx/flx mice is the formation of myelin outfoldings, defined as inappropriate growth of myelin membrane extensions away from the axons. These were observed in the optic nerve, corpus callosum, and spinal cord, and often resulted in the formation of what we term pseudo-multimyelin configuration, in which the outer membrane of the myelin outfold contacted the outer membrane of exiting myelin (i.e., reverse orientation), and the two inner membranes of the outfold were squeezed and faced each other (Fig. 3M,N). Notably, these are distinct from the multimyelin configuration formed by membrane slipping as a result of aberrant axoglial adhesion (Djannatian et al., 2019; Elazar et al., 2019a). In addition to pseudo-multimyelin, we also detected myelin outfoldings around nonmyelinated axons, as well as around neuronal cell bodies, suggesting that their growth is mainly determined by the space constraints within the local environment. The presence of extensive myelin outfoldings detected in the absence of N-Wasp in oligodendrocytes indicates that actin polymerization is required to control membrane growth during myelination. This notion is reinforced by the observation that myelin outfoldings are prominent after genetic deletion of the essential subunit of the Arp2/3 complex ArpC3 (Zuchero et al., 2015). Furthermore, similar myelin outfoldings characterized by abnormal accumulation of cytoplasm in the inner tongue of the myelin sheath were detected after oligodendrocyte-specific ablation of Rho GTPases Cdc42 (Thurnherr et al., 2006), which is a known activator of N-Wasp (Rohatgi et al., 2000). Collectively, our data suggest that membrane growth and correct myelin sheath formation are regulated by actin polymerization via the Cdc42-N-Wasp-Arp2/3 axis. In addition, the misguided movement of the oligodendrocyte leading edge away from the axon after ablation of N-Wasp could result from disrupted axoglial adhesion (Djannatian et al., 2019; Elazar et al., 2019a). N-Wasp is known to regulate cell adhesion (Misra et al., 2007), and the mobility of cell surface receptors (Rey-Suarez et al., 2020), raising the possibility that it may be involved in regulating oligodendrocyte-axon contact. Genetic deletion of N-Wasp in Schwann cells leads to reduced expression of myelin-associated glycoprotein (Jin et al., 2011), an adhesion axoglial molecule that was recently found to cooperate with other axoglial adhesion systems in targeting the inner tongue of oligodendrocytes during myelination (Garcia and Zuchero, 2019). Accordingly, the absence of N-Wasp in oligodendrocytes may not only impair the assembly of actin filaments, but also affect the adherence of the protruding
inner tongue, further leading to the formation of redundant myelin outfoldings. Another intriguing possibility to consider is that contact between the outer membranes of the outfolding and the myelin ensheathing the axon induces a myelin deterioration signal. Such a possibility is supported by observation demonstrating that the spatial segregation of myelin segments in the cortex is mediated by contact-dependent inhibitory signals between oligodendrocytes (Chong et al., 2012). While the exact underlying mechanisms are yet to be explored, our results emphasize the role actin dynamics has in proper myelin wrapping and the control of axonal integrity.

A surprising observation of this study is that the ablation of N-Wasp resulted in two distinct phenotypes in the CNS and PNS. In the PNS, N-Wasp is not required for axonal sorting, a process during which membrane extensions of Schwann cells select larger axons for myelination during development, nor for the initial ensheathment of axons, but is totally necessary for continuous membrane wrapping and the formation of myelin segments (Jin et al., 2011; Novak et al., 2011). In contrast, in the CNS, N-Wasp affects process extension by oligodendrocytes (Bacon et al., 2007), and hence axonal ensheathment and the number of segments formed. In addition, in the absence of N-Wasp in the CNS, membrane wrapping does accrue but often resulted in focal hypermyelination as evident by the formation of aberrant myelin outfoldings. It is interesting to note that, in terms of actin dynamics, the process of axonal sorting by Schwann cells could actually be considered equivalent to oligodendrocytes branching and their initial ensheathment of axons (Feltri et al., 2016). Hence, supported by studies of several other regulators of actin dynamics, such as Myosin II (Wang et al., 2008), Dynamin 2 (Gerber et al., 2019), and Dystonin I (Kornfeld et al., 2016), our findings indicate the existence of fundamental differences in the role actin reorganization plays during CNS and PNS myelination. Alternatively, the relative limited effect N-Wasp ablation has on CNS compared with PNS myelination could result from a possible compensatory mechanism by other actin nucleation-promoting factors of the Arp2/3 complex. N-Wasp belongs to the WASP protein family (i.e., WASP, WAVE, WASH, WHAMM, and JMY), of which several members are expressed in the oligodendrocyte lineage (Cahoy et al., 2008; Zhang et al., 2014; Marques et al., 2016). In line with this possibility, knockdown of JMY (junction-mediating and regulatory protein) in cultured oligodendrocytes impaired the formation of cellular protrusion and branching, and resulted in a reduced ability to contact axons and form myelin internodes when cocultured with neurons (Azevedo et al., 2018). Another candidate protein that may compensate for the
loss of N-Wasp is WAVE1, which like JMY and N-Wasp (Bacon et al., 2007), also regulates process outgrowth and lamellipodia formation in cultured oligodendrocytes (Kim et al., 2006). Furthermore, a general deletion of WAVE1 in mice results in a regional-specific hypomyelination (Kim et al., 2006), which is somewhat complimentary to the one we observed in Cnp-Cre/N-Waspflx/flx mutant mice. Whether WAVE1 provides compensatory function in the absence of N-Wasp will require the use of mice lacking both genes specifically in oligodendrocytes.

References


Czopka T, Ffrench-Constant C, Lyons DA (2013) Individual oligodendrocytes have only a few hours in which to generate new myelin sheaths in vivo. Dev Cell 25:599–609.


Katanov et al. • Role of N-Wasp in CNS Myelination


