

NG2-glia as Multipotent Neural Stem Cells: Fact or Fantasy?

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Cycling glial precursors—“NG2-glia”—are abundant in the developing and mature central nervous system (CNS). During development, they generate oligodendrocytes. In culture, they can revert to a multipotent state, suggesting that they might have latent stem cell potential that could be harnessed to treat neurodegenerative disease. This hope has been subdued recently by a series of fate-mapping studies that cast NG2-glia as dedicated oligodendrocyte precursors in the healthy adult CNS—though rare, neuron production in the piriform cortex remains a possibility. Following CNS damage, the repertoire of NG2-glia expands to include Schwann cells and possibly astrocytes—but so far not neurons. This reaffirms the central role of NG2-glia in myelin repair. The realization that oligodendrocyte generation continues throughout normal adulthood has seeded the idea that myelin genesis might also be involved in neural plasticity. We review these developments, highlighting areas of current interest, contention, and speculation.

NG2-glia in Perspective: The Neuron-Glial Dichotomy

The story of NG2-glia begins nearly thirty years ago, with the discovery of a class of glial precursors—“O-2A progenitors”—that could generate oligodendrocytes or type-2 astrocytes in cultures of perinatal rat optic nerve cells (Raff et al., 1983). (Two types of glial fibrillary acidic protein [GFAP]-expressing astrocytes, type-1 and type-2, were recognized in culture; only type-2 astrocytes arose from O-2A progenitors.) O-2A progenitors express a range of defining molecular markers including the nerve/glial antigen-2 (NG2, a proteoglycan core protein) and the platelet-derived growth factor receptor (alpha subunit, PDGFR α). Using these and other markers, the natural history of O-2A progenitors began to be revealed. It was shown that O-2A progenitors first develop in the ventricular germinal zones of the embryonic spinal cord and brain and disseminate through the developing CNS by proliferation and migration, becoming more-or-less uniformly distributed throughout the CNS soon after birth in rodents (reviewed by Miller, 1996; Richardson et al., 2006). After birth, O-2A progenitors associate with axons and generate myelinating oligodendrocytes, which are required for fast and efficient propagation of action potentials.

Cells with an antigenic phenotype that closely resembled perinatal O-2A progenitors were also identified in adult optic nerves (French-Constant and Raff, 1986; Wolswijk and Noble, 1989). These behaved as bipotential oligodendrocyte-astrocyte precursors in culture, just like their perinatal counterparts, but were found to divide, migrate, and differentiate more slowly (Wren et al., 1992). The existence of these “adult O-2A progenitors” was immediately recognized to have important implications for the repair of demyelinating damage such as occurs during multiple sclerosis. Cells that express *Pdgfra* mRNA, presumed to correspond to adult O-2A progenitors, were also visualized throughout the mature brain in situ (Pringle et al.,

1992). These were surprisingly numerous—around 5% of all cells in the CNS (Pringle et al., 1992; Dawson et al., 2003). Using antibodies against NG2 (Stallcup and Beasley, 1987; Diers-Fenger et al., 2001), a continuous network of NG2-immunopositive cells and cell processes was revealed, extending through all parts of the adult brain and spinal cord (Butt et al., 1999; Ong and Levine, 1999; Nishiyama et al., 1999; Chang et al., 2000; Horner et al., 2000; Diers-Fenger et al., 2001; Dawson et al., 2003). The abundance and ubiquitous distribution of these NG2⁺ cells was visually striking—shocking, even—and they came to be regarded as a novel “fifth neural cell type” after neurons, oligodendrocytes, astrocytes and microglia (Nishiyama et al., 1999; Chang et al., 2000; Butt et al., 2002, 2005; Dawson et al., 2003; Peters, 2004). NG2 and PDGFR α are also expressed by pericytes associated with the CNS vasculature (NG2⁺ and PDGFR α ⁺ pericytes appear to be distinct). However, double immunolabeling has shown that PDGFR α ⁺ and NG2⁺ nonvascular cells are essentially one and the same population (e.g., Nishiyama et al., 1996; Diers-Fenger et al., 2001; Dawson et al., 2003; Rivers et al., 2008). Therefore, in this review we refer to the latter as “NG2-glia” to distinguish them from pericytes.

In the meantime, attempts to identify type-2 astrocytes in the developing CNS in vivo had stalled, so a consensus arose that type-2 astrocytes were an artifact of culture. The term “O-2A progenitor” gradually passed out of general use and was replaced by “oligodendrocyte precursor” (OLP) or “oligodendrocyte precursor cell” (OPC) to reflect the then-prevailing view (in the 1990s) that these cells are dedicated mainly or exclusively to oligodendrocyte production during normal development and presumably also in the adult. The nature of type-2 astrocytes and their relationship to real cells in vivo was—and still is—an interesting conundrum. The relationship between OLPs in the perinatal CNS and NG2-glia in the adult was also

not immediately obvious. Although it seemed likely that the adult cells were descended by lineage from their perinatal counterparts this was not formally demonstrated until later, with the advent of genetic fate-mapping approaches in transgenic mice (see below). In this article we refer to both the perinatal and adult cells as NG2-glia.

The sheer number of NG2-glia in the adult brain and their uniform distribution in both gray and white matter seemed counterintuitive. Given their presumed role as oligodendrocyte precursors, should they not be concentrated in white matter where they would presumably be in most demand for myelinating axons? Why should so many precursor cells persist in the mature adult brain in any case? Moreover, the complex process-bearing morphology of NG2-glia *in vivo* seemed more in keeping with differentiated cells than immature precursors. Perhaps NG2-glia served a dual purpose—as a source of oligodendrocytes during development but fulfilling some more homeostatic or “functional” role in the adult (Nishiyama *et al.*, 1999; Butt *et al.*, 2002; Wigley *et al.*, 2007; Nishiyama *et al.*, 2009). Anatomical studies revealed that NG2-glia form close contacts with neurons—with axons at nodes of Ranvier and in close proximity to synapses at neuronal cell bodies (Butt *et al.*, 1999, 2002; Wigley and Butt, 2009). The hypothesis was born that NG2-glia, or a subset of them, might be involved in some aspects of information processing, in partnership with neurons.

This idea took off—and NG2-glia became really “exciting”—when electrophysiologists weighed in. It was already known that NG2-glia express some ion channels and neurotransmitter receptors and that glutamate can influence their proliferation and differentiation in culture (Barres *et al.*, 1990; Patneau *et al.*, 1994; Gallo *et al.*, 1996). However, the first demonstration that NG2-glia in the hippocampus receive long-range synaptic input from neurons *in vivo* sent waves through the research community (Bergles *et al.*, 2000). Synaptic communication between neurons and NG2-glia, both glutamatergic and GABAergic, was subsequently demonstrated in the cerebellum and cerebral cortex, both in gray and white matter (Lin and Bergles, 2002; Chittajallu *et al.*, 2004; Káradóttir *et al.*, 2005; Lin *et al.*, 2005; Salter and Fern, 2005; Paukert and Bergles, 2006; Kukley *et al.*, 2007; Ziskin *et al.*, 2007; Hamilton *et al.*, 2009). Physical synapses were identified between NG2 glia and unmyelinated axons in the corpus callosum (Kukley *et al.*, 2007; Ziskin *et al.*, 2007). Some NG2-glia were found to display spiking sodium currents in response to an initial depolarization (Chittajallu *et al.*, 2004; Káradóttir *et al.*, 2008; Mangin *et al.*, 2008; De Biase *et al.*, 2010). Suddenly, NG2-glia appeared exotic, ambiguous—glial in form (since they do not possess axons) but with some electrical properties akin to neurons. Their chimeric nature also contributed to the idea that NG2-glia, in their “other” role as precursor cells, might be more malleable than previously imagined and perhaps capable of transforming into neurons as well as glia.

NG2-glia as Multipotent Neural Stem Cells?

One study in particular launched the idea of NG2-glia as latent neural stem cells. This was the demonstration that NG2-glia purified from early postnatal (P6) rat optic nerves can apparently be reprogrammed into multipotent stem cells by first treating with

fetal calf serum (FCS) or bone morphogenetic proteins (BMPs) to generate type-2 astrocytes, followed by growth in basic fibroblast growth factor (bFGF) to generate free-floating balls of cells (neurospheres) containing neural stem cells (Kondo and Raff, 2000). Individual cells from these neurospheres could give rise to colonies containing a mixture of neurons, astrocytes, and oligodendrocytes, judged by immunolabeling with cell type-specific antibodies (Kondo and Raff, 2000). This study suggested that NG2-glia are more plastic than previously believed and suggested an explanation for previous reports that newborn rat optic nerve cells can generate neurons in culture, even though optic nerves do not normally contain neurons (Omlin and Waldmeyer, 1989).

These developments took place against a backdrop of exploding interest in stem cells of all sorts and neural stem cells in particular. A landmark series of papers had shown that subependymal astrocytes (“type-B cells”) in the subventricular zone (SVZ) of the adult rodent forebrain are in fact neural stem cells that generate migratory neuroblasts (“type-A cells”) destined for the olfactory bulb (Doetsch *et al.*, 1999). These neuroblasts follow the “rostral migratory stream” (RMS) from the SVZ to the olfactory bulb, where they differentiate into new olfactory interneurons of various types throughout adult life. It seemed (and still seems) possible that the type-2 astrocytes formed by culturing optic nerve NG2-glia with FCS or BMPs (Kondo and Raff, 2000) might be functionally analogous to the subependymal astrocytes (stem cells) of the SVZ. SVZ stem cells, hippocampal stem cells, and cultured type-2 astrocytes all express the glial fibrillary acidic protein (GFAP), for example, and all generate neurosphere-like bodies when cultured in the presence of bFGF. Subsequent reports that SVZ stem cells and their immediate progeny express NG2 and PDGFR α —as also do type-2 astrocytes—lent support to this interpretation (Belachew *et al.*, 2003; Aguirre and Gallo, 2004; Jackson *et al.*, 2006). Taken together, these observations implied that SVZ stem cells, type-2 astrocytes and parenchymal NG2-glia might all be close relatives.

The world of glia was up-ended. No longer were glial cells simply the “support cells of neurons” but rather the precursors of neurons, with neuron-like character of their own. Revolutionary ideas need firm foundations, so several labs geared up to test the differentiation fates of NG2-glia directly *in vivo*, using Cre-lox technology in transgenic mice. The first wave of such studies is now published and the conclusion is clear, if chastening: by far the most common differentiation products of parenchymal NG2-glia are oligodendrocytes, both in the normal and injured adult CNS. Some NG2-expressing cells produce astrocytes during embryonic development but not in the normal adult CNS. In the damaged CNS, the situation is a little more encouraging; following focal demyelination, for example, NG2-glia can generate remyelinating Schwann cells and possibly some astrocytes in addition to oligodendrocytes. However, the notion of NG2-glia as neuronal precursors has taken a significant blow. Although NG2-glia have some limited lineage plasticity—a source of continuing optimism for therapeutic applications—they are, by and large, precursors of myelinating cells. This shifts attention back to the therapeutic potential of NG2-glia in demyelinating conditions such as multiple sclerosis and spinal cord

injury. It also raises a raft of intriguing new questions concerning the role of myelination during normal adulthood.

Cre-lox Fate Mapping: Potential Pitfalls

The general principles of Cre-lox fate mapping are as follows. Mice expressing Cre recombinase under transcriptional control of a gene that is active in NG2-glia (e.g., *Pdgfra*, *NG2*, *Olig2*, *Plp1*) are generated by conventional transgenesis using a plasmid or bacterial artificial chromosome (BAC) or else by homologous recombination in ES cells (knockin). These are crossed with a Cre-conditional reporter line—e.g., *Rosa26-lox-STOP-lox-GFP*, where *Rosa26* is a ubiquitously active promoter, *lox* the recognition site for Cre recombinase, STOP a series of four cleavage/polyadenylation sites (which effectively stop mRNA production) and *GFP* a cassette encoding green fluorescent protein. In double-transgenic offspring (e.g., *NG2-Cre: Rosa26-GFP*), Cre-driven recombination within the reporter transgene activates expression of GFP irreversibly in NG2-expressing cells and all of their descendants, which are identified retrospectively by immunolabeling for GFP together with cell type-specific markers. This version of the technique, using standard Cre, labels NG2-glia as they come into existence during early development and therefore labels all of the progeny of NG2-glia up to the time of analysis. An important modification is to use CreER*, a fusion between Cre and a mutated form of the estrogen receptor (ER*) that no longer binds estrogen at high affinity but can bind 4-hydroxy tamoxifen (4HT), a metabolite of the anti-cancer drug, tamoxifen. After binding 4HT, CreER* translocates from the cytoplasm (where unliganded ER is normally sequestered) to the nucleus, triggering recombination and reporter gene activation. This version of the technique allows NG2-glia to be labeled inducibly (by administering tamoxifen or 4HT to the mice) at a defined stage of development or adulthood, and the course of division and differentiation of the NG2-glia charted subsequently (Figure 1).

While this sounds straightforward, there are pitfalls. First among these is the transcriptional specificity of the Cre transgene, which rarely if ever targets exclusively the precursor cells of interest. Ideally, one should compare and integrate data from independent mouse lines that express Cre (or CreER*) under different transcriptional control—e.g., under either *Pdgfra* or *NG2* control in the present context of NG2-glia. Another technical issue relates to the kinetics of Cre recombination—this depends on the structure of the reporter transgene (e.g., distance between *lox* sites) as well as the level and duration of Cre expression. Often, a larger proportion of the target cell population can be labeled using a “good recombiner” like *Rosa26-YFP* (Srinivas et al., 2001) compared to a “poor recombiner” like *Rosa26-eGFP* (Mao et al., 2001). The commonly used *Z/EG* (Novak et al., 2000) and *Rosa26-LacZ* (Soriano, 1999) lie somewhere between these. There seems to be a threshold of Cre expression, below which very little recombination of the reporter gene occurs; this threshold is lower for a good recombiner than a poor recombiner. This effect can introduce additional cell-type selectivity; for example, if a given mouse line expresses Cre in two types of cell, but more highly in one than the other, then reporter gene activation can effectively be restricted to the more highly-expressing cells. This can be useful

in some circumstances; for example, it is probably the reason that *Olig2-CreER** drives recombination and reporter gene activation mainly in NG2-glia and not in differentiated oligodendrocytes (Dimou et al., 2008). However, in other situations it might introduce unwanted bias, e.g., by subdividing the NG2-glia population in some unpredictable way. In short, Cre-lox fate mapping studies need to be interpreted with care and an open mind, each study considered on its own merits.

It is worth noting here that differences in the tamoxifen induction protocol—whether tamoxifen or 4HT is used, whether it is administered by injection or gavage, or whether it is administered once or several times, for example—can also affect the efficiency of recombination independently of the reporter mouse line employed. While this will result in a greater or lesser fraction of NG2-glia becoming labeled, it will have no effect on the level of expression of the reporter in individual cells because that is determined purely by regulatory elements in the reporter transgene itself. The flip side of this is that the reporter transgene (e.g., *Rosa26*-based) is not necessarily expressed to the same high level in all cell types, which in principle could lead to under-estimation of certain cell types among the labeled progeny of NG2-glia, although there is no evidence that this has been a problem in the studies reviewed below.

NG2-glia Generate Oligodendrocytes but Not Astrocytes during Normal Adulthood

Initially, constitutively active Cre was used in *NG2-Cre* BAC transgenic mice to label the progeny of NG2 glia during brain and spinal cord development (Zhu et al., 2008a, 2008b). As expected, a large proportion of myelinating oligodendrocytes was found among the GFP-labeled progeny of perinatal NG2-glia. In addition, significant populations of GFP-labeled protoplasmic astrocytes were found in the gray matter of the ventral forebrain and spinal cord, though not in white matter. No other labeled cells were identified—in particular, no neurons. A follow-up study from the same group, using *NG2-CreER** instead of *NG2-Cre*, allowed the progeny of NG2-glia to be traced in the postnatal as well as the embryonic brain (Zhu et al., 2011). When tamoxifen was administered at embryonic ages (E16.5), a similar result was obtained as before—NG2-glia generated mainly oligodendrocytes but also some protoplasmic astrocytes in ventral brain territories. Using a reporter line (*Z/EG*) that recombines inefficiently, Zhu et al. (2011) labeled a sparse subset of embryonic NG2-glia that over time generated discrete clusters (presumed clones) of sibling cells. They found that labeled cell clusters contained either astrocytes or oligodendrocyte lineage cells but not both, suggesting that different subsets of NG2-glia in the embryonic CNS are specialized for production of only ventral astrocytes or only oligodendrocytes. When tamoxifen was administered to postnatal mice (P2, P30, or P60) a different result was obtained—this time no astrocytes were found among the progeny of NG2-glia—concurring with previous experience from other labs that had used different CreER* lines (see below). These data imply that there are two distinct subtypes of NG2-glia—“astrogenic” and “oligogenic”—in the early developing CNS, the astrogenic population being depleted during late embryonic development. A feasible explanation might go as follows. Neuroepithelial precursors

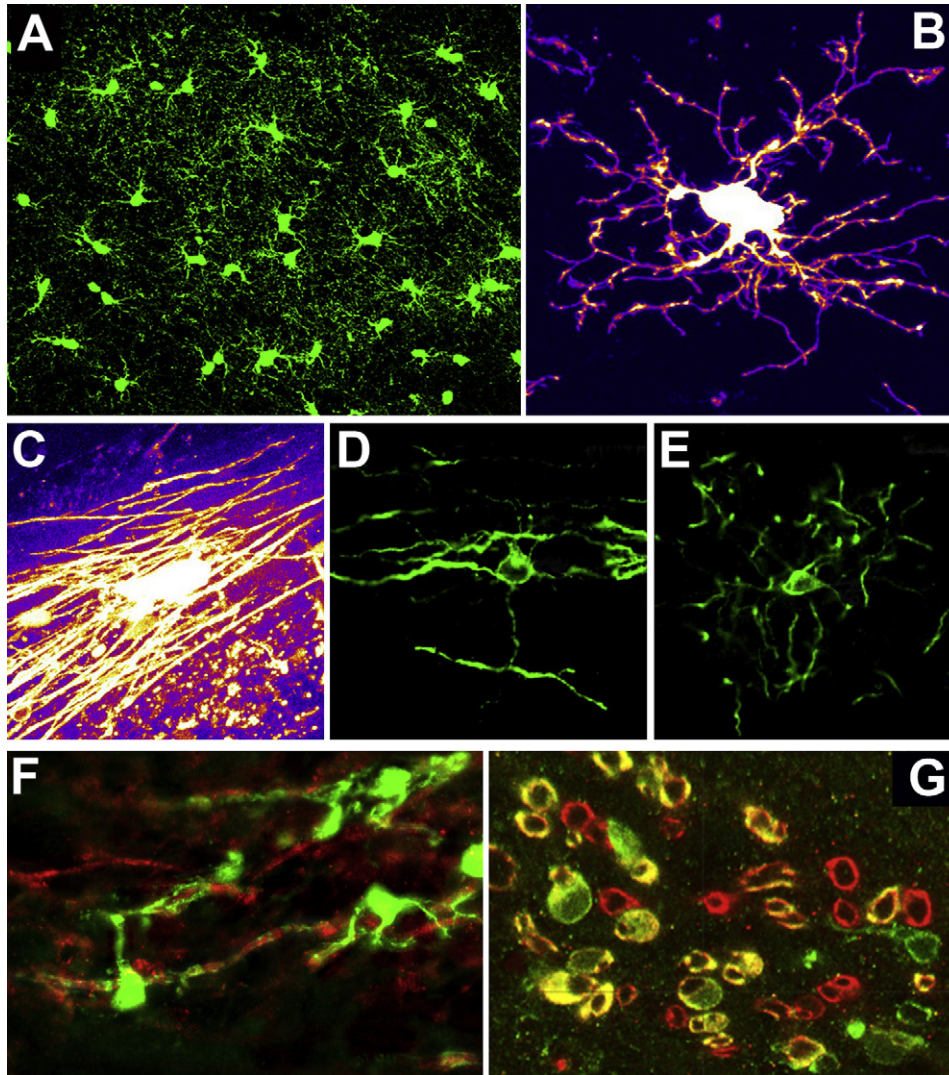


Figure 1. Compendium of Images of NG2-glia and Their Differentiated Progeny in Transgenic Mice

(A) Low-magnification epifluorescence image of the cerebral cortex of a P13 *Sox10-GFP* mouse showing the near-uniform distribution of oligodendrocyte lineage cells, mainly NG2-glia.

(B) A single NG2-gial cell in the cerebellum of a P45 *Pdgfra-CreER* : Rosa26-YFP* mouse, identified by intrinsic fluorescence in a vibratome section, dye-filled through a micropipet with Alexa Fluor 488 to reveal whole cell morphology and imaged by two-photon microscopy (Rivers et al., 2008).

(C) A dye-filled myelinating oligodendrocyte in the corpus callosum of a *Pdgfra-CreER* : Rosa26-YFP* mouse, induced with tamoxifen at P45 and examined 28 days later (P45 + 28) in the two-photon microscope (Rivers et al., 2008). This adult-born oligodendrocyte has >50 internodes. Reproduced from Rivers et al. (2008).

(D) A myelinating oligodendrocyte in the optic nerve of a *Pdgfra-CreER* : Tau-mGFP* mouse at P60 + 28. Membrane-tethered mGFP reveals full cell morphology including the myelin sheaths.

(E) A myelinating oligodendrocyte in the cortical grey matter of a *Pdgfra-CreER* : Tau-mGFP* mouse, showing many short randomly orientated myelin internodes, contrasting with the multiple aligned internodes in white matter.

(F) Remyelinating oligodendrocytes in a *Pdgfra-CreER* : Rosa26-YFP* focally-demyelinated spinal cord (Zawadzka et al., 2010). The mice were induced with tamoxifen on four consecutive days starting at ~P75, lysolecithin (1% v/v) was injected stereotaxically 4 days after the final dose and the animals were analyzed 3 weeks later by epifluorescence microscopy. Oligodendrocyte cell bodies are visualized with anti-GFP (green) and myelin sheaths with anti-PLP (red). Note that cytoplasmic YFP is largely excluded from compact myelin, presumably for steric reasons.

(G) Remyelinating Schwann cells within a focal ethidium bromide (0.1% w/v)-induced lesion in *Pdgfra-CreER* : Rosa26-YFP* spinal cord (Zawadzka et al., 2010). Schwann cells are visualized by immunolabelling for GFP (green) and the Schwann cell-specific protein Periaxin (red).

(radial glia) in the ventral ventricular zone first divide asymmetrically to maintain their own numbers while giving rise to proliferative NG2-glia, which migrate away from the ventricular surface, generating oligodendrocytes during early postnatal development and persisting as oligogenic NG2-glia into adulthood.

Then, just prior to birth, the remaining radial glia transform directly into astrocytes, expressing NG2 transiently as they do so; these astrocytes undergo limited cell division and settle in ventral territories close to their region of origin. Direct trans-differentiation of radial glia is a normal mode of astrocyte

generation in the developing cortex, for example (Mission et al., 1991). The given scenario is consistent with a study using *Olig2-CreER**, in which some astrocytes as well as oligodendrocytes (and motor neurons) were found among the progeny of *Olig2*-expressing neuroepithelial precursors in the embryonic ventral spinal cord (Masahira et al., 2006).

Whatever the precise sequence of events during prenatal gliogenesis, it now seems likely that NG2-glia do not generate astrocytes during normal healthy adulthood. Several Cre-lox studies—using *Pdgfra-CreER** (two independent lines: Rivers et al., 2008; Kang et al., 2010), *NG2-CreER** (Zhu et al., 2011; see above), and *Olig2-CreER** (Dimou et al., 2008) converge on that conclusion. While some reporter-positive astrocytes were observed in the cortical gray matter in the *Olig2-CreER** line, their number did not increase significantly between 8 and 65 days post-tamoxifen administration, indicating that they were not generated continuously from dividing NG2-glia (Dimou et al., 2008). It appears that the *Olig2-CreER** transgene is expressed in some protoplasmic astrocytes in the normal gray matter, resulting in labeling of some of these in addition to NG2-glia. A subsequent study from the same lab (Simon et al., 2011) marked NG2-glia in a different way, by long-term BrdU labeling of 2- to 3-month-old mice, and confirmed that no astrocytes were found among their differentiated progeny.

Neuron Genesis from NG2-glia—Fact or Fantasy?

NG2-glia exposed to appropriate environmental signals in a culture dish appear to revert to a multipotent state, from which they can generate neurons as well as oligodendrocytes and astrocytes (Kondo and Raff, 2000). This sparked the widespread hope that NG2-glia can be a regenerative resource for neurodegenerative diseases that involve neuronal as well as glial loss. A number of studies have encouraged this hope by describing neuronogenic properties of NG2-glia in the normal rodent CNS. For example, NG2-glia in the neocortex and piriform cortex have been reported to express Doublecortin (Dcx), an established marker of migratory neuronal progenitors in the forebrain SVZ/ RMS and hippocampus (Tamura et al., 2007; Guo et al., 2010). Some NG2⁺ cells in the piriform cortex have been found to express Sox2 and Pax6 (Guo et al., 2010), two more neural stem cell markers. Conversely, SVZ and hippocampal stem cells have been reported to express NG2 (Belachew et al., 2003; Aguirre and Gallo, 2004) and PDGFRa (Jackson et al., 2006) and to actively transcribe a *CNP-GFP* transgene (Belachew et al., 2003; Aguirre and Gallo, 2004; Aguirre et al., 2004). However, not all of these observations have survived scrutiny. For example, other labs have failed to confirm NG2 or PDGFRa antibody labeling of SVZ or hippocampal stem cells (Komitova et al., 2009) or to detect NG2 or PDGFRa promoter activity in these stem cell populations in BAC transgenic mice (Rivers et al., 2008; Zhu et al., 2008a; Kang et al., 2010). While antibody-labeling experiments are notoriously difficult and artifact-prone, genetic labeling should be more predictable—so one might imagine—and therefore capable of providing an unequivocal answer to the question “do NG2-glia generate neurons”? However, Cre-lox fate mapping studies have still not completely eliminated the controversy around this question.

Using *Pdgfra-CreER*⁺: Rosa26-YFP* mice, our lab found that although NG2-glia generate predominantly Sox10-positive oligodendrocyte lineage cells during normal adulthood, some Sox10-negative, YFP⁺ cells appeared and accumulated in layers 2 and 3 of the anterior piriform cortex (aPC) (Rivers et al., 2008). The cells acquired NeuN reactivity and morphologically resembled piriform projection neurons. The scale of neuron genesis was small; we estimated that only ~1.4% of the neurons present in layers 2/3 of the aPC were generated in the ~200 days after P45 (Rivers et al., 2008). Two things pointed to their being generated continually from precursor cells: (1) they were not observed until around 1 month after tamoxifen administration, suggesting that they differentiated slowly from NeuN-negative precursors and (2) they steadily increased in number between 28 and 210 days posttamoxifen. It is difficult to imagine how YFP-labeled PC neurons could continue to accumulate months after tamoxifen administration had ceased, unless they were generated from a population of precursor cells that had recombined the YFP reporter gene at the time of tamoxifen addition. They could not have been generated from SVZ stem cells because no YFP⁺ PC neurons were found in *Fgfr3-CreER*⁺: Rosa26-YFP* mice, which marks all GFAP⁺ SVZ stem cells and their neuronal progeny in the olfactory bulb, for example (Rivers et al., 2008; Young et al., 2010). We were unable to colabel the YFP⁺ neurons with BrdU, even after months (100 days) of BrdU exposure via the drinking water, indicating that they might have formed by direct transformation of long-term-quiet precursors. Since we found that ~50% of NG2-glia did not incorporate BrdU over the same time scale (this is controversial), we suggested that the new PC neurons were formed by transdifferentiation of postmitotic NG2-glia (Rivers et al., 2008; Psachoulia et al., 2009). Another possibility is that the new aPC neurons were produced from some other pool of *Pdgfra*-expressing precursors, as yet unidentified. *Pdgfra* is expressed by large numbers of cells outside of the CNS so the new neurons could conceivably originate from precursors that enter the CNS via the circulatory system. Alternatively, they might be generated from *Pdgfra*⁺ perivascular cells within the CNS. (NG2⁺, PDGFRb⁺) perivascular pericytes have been reported to generate neurons and glia in culture in response to bFGF (Dore-Duffy et al., 2006), so it is conceivable that PDGFRa⁺ pericytes might have similar stem cell-like properties.

One other study has reported PC neurons from NG2-glia (Guo et al., 2010). This study used *Pip1-CreER*⁺: Rosa26-YFP* mice to follow fates of NG2-glia in the healthy adult CNS. *Pip1* is expressed in differentiated oligodendrocytes as well as NG2-glia, so Guo et al. (2010) could not address questions about new oligodendrocyte production; however, like Rivers et al. (2008), they did observe YFP-labeling of PC projection neurons. Their labeled neurons first became apparent 17 days posttamoxifen and increased in number for at least 180 days. Control experiments using *GFAP-CreER*⁺: Rosa26-YFP* mice to mark SVZ stem cells ruled out the possibility that the newly-labeled PC neurons were SVZ-derived. Thus, there are strong parallels between the experiments and data of Guo et al. (2010) and our own (Rivers et al., 2008), except that Guo et al. (2010) described their neurons in the posterior piriform cortex (pPC) (Bregma

levels -2.3 mm to -1.1 mm), whereas ours were predominantly in the aPC.

Using their own independently generated line of *Pdgfra-CreER** mice, Kang et al. (2010) failed to detect production of long-term surviving GFP⁺ neurons in the PC or elsewhere in the forebrain. The reason for this difference between their study and ours (Rivers et al., 2008) is not clear. Since the *Pdgfra* BACs used for transgenesis were different (ours contained ~ 55 kb upstream and ~ 74 kb downstream of the *Pdgfra* gene, theirs ~ 70 kb upstream and ~ 40 kb downstream) it is conceivable that they might have inherently different transcriptional specificity—e.g., our BAC but not theirs might contain a regulatory element required for expression of the *Pdgfra-CreER** transgene in a particular set of *Pdgfra*-expressing neuronal precursors. This presupposes that the putative *Pdgfra*⁺ precursors are something other than NG2-glia, since both *Pdgfra-CreER** transgenes are demonstrably expressed in NG2-glia. Alternatively, it might depend on where one looks—we quantified piriform neuron production in the aPC (Bregma levels $+0.22$ mm to $+1.2$ mm) (Rivers et al., 2008 and unpublished), whereas Kang et al. (2010) examined pPC (Bregma -1.9 mm to $+0.5$ mm).

Other groups have reported small numbers of reporter-positive neurons (NeuN⁺) throughout the forebrain—particularly the ventral forebrain—at short times after CreER* induction, but discounted these as probably resulting from sporadic CreER* expression in the neurons themselves (Dimou et al., 2008; Kang et al., 2010; Guo et al., 2010; Zhu et al., 2011). In one study YFP⁺, NeuN⁺ neurons were observed for only a few days following 4HT injection into *NG2-CreER*: Rosa26-YFP* mice (Zhu et al., 2011), suggesting that these neurons were eliminated from the CNS after a short time or else became NeuN negative (or YFP negative). The aPC neurons that we observed (Rivers et al., 2008) are distinct from these and, whatever their origin, cannot easily be explained by sporadic activation of the CreER* transgene in neurons (Kang et al., 2010; Zhu et al., 2011), or by some aspect of the tamoxifen protocol (Simon et al., 2011).

Further experiments will be required to resolve the ambiguity around PC neuron genesis. For example, it will be useful to establish whether there is a real difference between the two *Pdgfra-CreER** lines (Rivers et al., 2008; Kang et al., 2010). If there is, then differences in the transcriptional specificity of the two transgenes might give clues to the origin of the PC neurons observed by Rivers et al. (2008). The transcriptional specificity of the *Pip1* promoter-proximal fragment used in *Pip1-CreER** (Guo et al., 2010) also needs to be examined. This is a fragment of the *Pip1* gene (2.4 kb of 5' sequence plus exon1 and intron1) (Doerflinger et al., 2003), and it cannot be assumed to exactly mimic endogenous *Pip1* expression—which itself is not entirely oligodendrocyte lineage specific, being expressed in a subset of multipotent precursors during development (Delaunay et al., 2009; Guo et al., 2009; Kang et al., 2010) as well as in some differentiated neurons (Nery et al., 2001; Miller et al., 2009). This potentially complicates interpretation of fate mapping studies (Guo et al., 2009, 2010). It is quite important to get to the bottom of these discrepancies because, at the very least, it will refine our understanding of the Cre-lox system and its potential shortcomings.

In any case, there seems to be something interesting going on in the PC. There has been a steady trickle of evidence for neuronal progenitors/immature neurons residing there. For example, cells in the PC have been reported to express Doublecortin, polysialated NCAM, Sox2, and other markers of neural precursor cells (Seki and Arai, 1991; Hayashi et al., 2001; Nacher et al., 2001, 2002; Pekcec et al., 2006; Shapiro et al., 2007; Bullmann et al., 2010; Guo et al., 2010). There have also been reports of continued neuron genesis in the adult rodent and primate PC and its modulation by olfactory stimulation (Bernier et al., 2002; Pekcec et al., 2006; Shapiro et al., 2007; Arisi et al., 2011). Another possibility is that the immature neuron markers might be indicative of neuronal de-differentiation and remodeling in response to changing inputs from the olfactory bulb (OB) (Seki and Arai, 1993; Nacher et al., 2001), since the aPC (otherwise known as the primary olfactory cortex) is the primary target for output neurons (mitral cells) of the OB. The internal OB circuitry is continually changing throughout life, due to the addition of new OB interneurons from the SVZ, so perhaps the whole olfactory system including the aPC is in a constant state of flux.

Do NG2-glia Generate Reactive Astrocytes after Injury?

NG2-glia are known to react to injury by proliferating, upregulating NG2 expression and generating remyelinating oligodendrocytes when required (reviewed by Levine et al., 2001). Since their differentiation potential is known to be influenced by their environment in vitro (Kondo and Raff, 2000), it is possible that they might display a broader range of fates following CNS injury or disease, when their microenvironment is likely to be altered by inflammatory cells and possibly through breach of the blood-brain barrier. Therefore, it is of great interest to discover the fates of NG2-glia in various experimental models of disease or traumatic injury.

There has now been a handful of genetic fate mapping studies of NG2-glia during various experimental pathologies in mice. These include experimental autoimmune encephalomyelitis (EAE) (Tripathi et al., 2010), acute gliotoxin-induced focal demyelination (Zawadzka et al., 2010), spinal cord section (Barnabé-Heider et al., 2010), cortical stab wound (Dimou et al., 2008; Komitova et al., 2011), and a mouse model of inherited amyotrophic lateral sclerosis (ALS; motor neuron disease) (Kang et al., 2010). All these studies found that NG2-glia respond to injury by proliferating and accumulating at the site(s) of damage, and that their major differentiation products are new oligodendrocytes. In addition to oligodendrocytes, modest astrocyte production was also reported in some but not all of these studies—the main source of reactive astrocytes being preexisting astrocytes, not NG2-glia. One study does not conform to this pattern. This is a study of cell generation following a cold-induced injury to the cerebral cortex (Tatsumi et al., 2008), in which the major product of NG2-glia appeared to be protoplasmic “bushy” astrocytes, not oligodendrocytes (see below). NG2-glia derived neurons were not found in any of these studies, however. The main features of all the fate-mapping studies discussed in this review are summarized in Table 1.

Following a cortical (gray matter) stab injury in adult *Olig2-CreER*: Z/EG* mice, Dimou et al. (2008) reported oligodendrocyte

Table 1. Summary of Cre-lox Fate-Mapping Studies Discussed in this Article

CreER*	Inducer	Reporter	Condition	% of NG2-glia Labeled	Cell Types Generated			
					OL	AS	N	Other
Pdgfra ^a	tam	R-YFP	normal	45–50	+++	–	+ (aPC)	
Pdgfra ^b	4HT	Z/EG	normal	40–45	+++	–	–	
		R-YFP	normal	~90	nd	nd	–	
		Z/EG	ALS (SC)	?	+++	–	–	
Pdgfra ^c	tam	R-YFP	EAE (SC)	~30	+++	+/-	–	
Pdgfra ^d	tam	R-YFP	focal demyelination	~40	+++	+/-	–	Schwann cells
NG2 ^e	4HT	Z/EG	normal	<2	+++	–	–	
		R-YFP	normal	~45	+++	–	–	
NG2 ^f	4HT	Z/EG	cortical stab	<2	+++	+/-	nd	
Olig2 ^g	tam	Z/EG+R-GFP	normal	?	+++	+/-	–	
		Z/EG+R-GFP	cortical stab	?	+++	+/-	–	
Olig2 ^h	4HT	R-GAP43-GFP	cryoinjury	?	–	+	–	
Olig2 ⁱ	tam	R-LacZ, R-YFP	axotomy (SC)	~40	+++	–	–	
Plp1 ^j	tam	R-YFP	normal	10	nd	nd	+ (pPC)	

R-YFP, *Rosa26-YFP* etc. (see text for references); SC, spinal cord; ALS, amyotrophic lateral sclerosis; aPC, anterior piriform cortex; pPC posterior piriform cortex; OL, oligodendrocyte lineage; AS, astrocytes; N, neurons; nd, not done. +/- indicates that some astrocytes were found but that these were infrequent (<5% of reporter-positive cells) and/or could be explained by recombination in a subset of astrocytes soon after tamoxifen administration.

^a Rivers et al. (2008).

^b Kang et al. (2010).

^c Tripathi et al. (2010).

^d Zawadzka et al. (2010).

^e Zhu et al. (2011).

^f Komitova et al. (2011).

^g Dimou et al. (2008).

^h Tatsumi et al. (2008).

ⁱ Barnabé-Heider et al. (2010).

^j Guo et al., (2010).

generation but little or no astrocyte production. An accumulation of GFAP⁺ BrdU⁺ reactive astrocytes was found in the vicinity of the lesion, as expected, but these were mostly reporter-negative (i.e., not NG2-glia derived). Very similar results to these were reported following cortical stab wounds in *NG2-CreER*⁺ Rosa26-YFP* mice (Komitova et al., 2011). A subsequent BrdU fate mapping study (Simon et al., 2011) failed to find evidence for any astrocyte production from dividing NG2-glia after cortical stab injury. The emerging consensus from these studies is that the reactive (hypertrophic, strongly GFAP⁺) astrocytes that form the glial “scar” around sites of injury in the cortex are derived predominantly or exclusively from pre-existing astrocytes, not from NG2-glia. This conclusion has been supported by complementary experiments in which astrocytes were labeled specifically by injecting a *GLAST-CreER** lentiviral vector into the cortex of reporter mice, and their fates followed before and after cortical stab injury (Buffo et al., 2008). Before injury, the labeled astrocytes were quiescent (did not incorporate BrdU after a long label) but, after injury, they started dividing and generated many new astrocytes, but not other cell types, at the site of the wound.

This also seems to be what happens after spinal cord injury. Barnabé-Heider et al. (2010) made a transverse cut through the dorsal funiculus of the spinal cord, severing the ascending and descending axon tracts. They observed new oligogenesis but insignificant astrocyte production from NG2-glia (marked using

*Olig2-CreER**), despite a robust astrocytic reaction/gliosis. Most interestingly, they identified two separate components of the astrocytic reaction—a localized accumulation of GFAP⁺ astrocytes at the core of the lesion site in the dorsal funiculus and a more diffuse accumulation/gliosis around the lesion site and throughout the spinal cord at the level of the injury. These two components had different cellular origins; the “core” astrocytes were derived from multipotent stem cells in the ependymal zone surrounding the central canal of the cord (labeled with *FoxJ1-CreER**), whereas the “diffuse” astrocytes were derived from pre-existing astrocytes in the parenchyma (labeled with *Connexin30-CreER**). It is curious that Barnabé-Heider et al. (2010) found that all protoplasmic astrocytes in the spinal cord were Olig2-immunoreactive in their experiments. Despite this, the *Olig2-CreER** transgene did not drive recombination in astrocytes, perhaps because the level of CreER* expression was below threshold (see above, under heading “Cre-lox Fate Mapping: Potential Pitfalls”). While this worked out well for Barnabé-Heider et al. (2010), it does raise the possibility that *Olig2-CreER** might trigger recombination in astrocytes in addition to oligodendrocyte lineage cells in some circumstances.

In marked contrast to the above is a study by Tatsumi et al. (2008), who followed the fates of NG2-glia following freeze-thaw lesions in the cerebral cortex. They used *Olig2-CreER**

mice (the same line used by both Dimou et al. [2008] and Barnabé-Heider et al. [2010]) to mark presumptive NG2-glia prior to injury and reported a robust proliferative response followed by production of “bushy” protoplasmic astrocytes between one and 2 weeks postinjury. Astrocytes appeared to be the major differentiated product of NG2-glia in this injury model; oligodendrocytes were not observed. However, in this study, as in that of Dimou et al. (2008), *Olig2-CreER** triggered recombination in some protoplasmic astrocytes in addition to NG2-glia in the uninjured cortex (~20% of reporter-positive cells were astrocytes at short times after 4HT administration; Tatsumi et al., 2008). This leaves open the possibility that the reactive astrocytes formed after injury were derived from division of pre-existing astrocytes. Tatsumi et al. (2008) discounted this idea because they failed to find BrdU-labeled GFAP⁺ astrocytes shortly after injury but it is possible that there could have been a delayed mitogenic response of astrocytes or slow upregulation of GFAP in previously GFAP-negative astrocytes, either of which might have obscured a transient population of BrdU⁺ astrocytes. Nevertheless, the apparent absence of oligodendrocyte production in the experiments of Tatsumi et al. (2008) marks their study out from the others; perhaps the particular environment of the freeze-thaw injury, as compared to stab injury for example, inhibits NG2-glia from differentiating into oligodendrocytes. It is important to confirm or refute this observation through cryo-lesioning experiments in different *CreER** mouse lines, because it could perhaps provide a link to late-stage multiple sclerosis lesions, in which inhibition of oligodendrocyte differentiation is thought to contribute to remyelination failure.

Other researchers have examined the response of NG2-glia during experimentally induced demyelination. In a gliotoxin-induced focal demyelination model, Zawadzka et al. (2010) found robust remyelination from NG2-glia (labeled using *Pdgfra-CreER*: Rosa26-YFP*)—as expected from previous studies (Redwine and Armstrong, 1998; Watanabe et al., 2002; Dawson et al., 2003; reviewed by Franklin and ffrench-Constant, 2008; Figure 1F). A small proportion of YFP⁺ cells were Aquaporin-4⁺ astrocytes (~3%), but the great majority of reactive astrocytes were derived from *Fgfr3*-expressing cells (ependymal cells and/or preexisting astrocytes) (Young et al., 2010), because they were YFP-labeled in *Fgfr3-CreER*: Rosa26-YFP* mice (Zawadzka et al., 2010). Schwann cells, the myelinating cells of the peripheral nervous system (PNS), are commonly found in remyelinating CNS lesions including some human multiple sclerosis lesions. Often these remyelinating Schwann cells surround blood vessels, which in the past has been taken to suggest that they enter the CNS from the PNS, using the vessels as a migration route. However Zawadzka et al. (2010) found that most remyelinating Schwann cells (Periaxin⁺) in their CNS lesions were YFP⁺ in *Pdgfra-CreER*: Rosa26-YFP* mice, suggesting that they were derived from NG2-glia (Figure 1G). In strong support of this, the CNS-resident Schwann cells were also labeled in *Olig2-Cre: Rosa26-YFP* animals—*Olig2* is not thought to be expressed outside of the CNS. Moreover, almost no CNS Schwann cells, but many Schwann cells in peripheral nerves, were labeled in *Pzero-CreER*: Rosa26-YFP* mice. (*Pzero* is expressed in migrating neural crest and differentiated Schwann cells, but not in the oligodendrocyte lineage.)

Schwann cells were not a minor side product of NG2-glia because 56% of all YFP⁺ cells in ethidium bromide-induced lesions were Periaxin⁺ Schwann cells. (Despite this, most new myelin is oligodendrocyte derived, because Schwann cells each remyelinate only a single internode, whereas oligodendrocytes remyelinate many.) To our knowledge, this is the clearest example to date of lineage plasticity among NG2-glia in vivo. Since both oligodendrocytes and Schwann cells are myelinating cells, relatively subtle reprogramming might be required to cross between them.

In a different demyelinating model—experimental autoimmune encephalomyelitis (EAE), which causes more diffuse and widespread demyelination than gliotoxin injection—Tripathi et al. (2010) found robust production of NG2-glia-derived oligodendrocytes but very few Schwann cells. In EAE, there is a strong inflammatory component to the pathology that is not present in gliotoxin-induced demyelination, suggesting that the local microenvironment in demyelinated lesions exerts a strong influence on the direction of differentiation of NG2-glia. Only a small fraction of YFP⁺ cells (1%–2%) were GFAP⁺ astrocytes in EAE, in keeping with the results from focal demyelination (Zawadzka et al., 2010). A relatively high proportion (~10%) of YFP⁺ cells in this EAE study could not be identified, despite much effort with antibodies against microglia, macrophages, B or T cells, neutrophils, vascular endothelial cells, pericytes, neurons, astrocytes, oligodendrocytes, and Schwann cells.

No astrocyte production from NG2-glia was detected in a mouse model of hereditary ALS (Kang et al., 2010), despite extensive astrocytosis in the ventral horns of the spinal cord where motor neuron degeneration occurs. These authors did, however, observe proliferation and accumulation of NG2-glia and differentiated oligodendrocytes—an unexpected result, since oligodendrocyte involvement in ALS pathology was not previously suspected. Whether reactive NG2-glia are specifically involved in myelin repair in ALS or an incidental byproduct of tissue damage or inflammation is not known.

The gathering consensus seems to be that NG2-glia remain largely committed to the oligodendrocyte lineage in the healthy CNS and in most pathological situations. Exceptions are (1) the still-unresolved question of low-level neuron genesis in the piriform cortex during normal adulthood, (2) robust Schwann cell generation following gliotoxin-induced demyelination, and (3) production of a few GFAP⁺ astrocytes in some but not all injury studies. Overall, lineage flexibility seems to be strongly biased toward myelinating lineages. This injects a healthy dose of realism and tempers our hopes for NG2-glia as a panacea for neurodegenerative disease. It remains possible that NG2-glia might have the potential to generate neurons but do not readily reveal that potential in the environment of the damaged CNS—at least not those conditions that have been examined so far. Pharmacological interventions that can redirect differentiation toward neurons might be found in the future, but it will not be an easy fix. On the other hand, the data reaffirm the central role of NG2-glia in myelin repair, in demyelinating conditions such as multiple sclerosis or spinal cord injury. It has also been useful to learn that the great majority of reactive astrocytes are in most cases not descended from NG2-glia, but from parenchymal astrocytes that re-enter the cell cycle and, in the spinal

cord, from ependymal cells around the central canal. The latter cells represent a relatively unexplored population that is a key target for future investigation. It will be important to discover whether these cells retain, or can be induced to recapitulate, some of the neuronogenic flavor of their forebears in the embryonic neuroepithelium.

Oligodendrocyte Generation and Myelin Dynamics in the Adult

Most newly formed oligodendrocytes in the postnatal forebrain survive long-term and myelinate axons. Myelin formation has been demonstrated by microinjecting live YFP⁺ cells in tissue slices with a fluorescent dye that can spread throughout the cell and expose its full morphology. Like this, newly-formed mature oligodendrocytes that elaborate up to ~50 internodes have been visualized in the adult corpus callosum (Rivers et al., 2008; Figure 1C). Newly formed oligodendrocytes with myelinating morphology were also identified using reporter mice that express a membrane-tethered form of GFP (Kang et al., 2010; Zhu et al., 2011; Figure 1D). That most new oligodendrocytes survive long-term (at least in the corpus callosum) can be inferred by comparing the rate of new cell production (through division of NG2-glia) with the rate of accumulation of differentiated oligodendrocytes, given that the absolute number and density of NG2-glia does not change dramatically during the first year of life (Psachoulia et al., 2009). The cell cycle dynamics of NG2-glia is known from cumulative BrdU labeling experiments (Psachoulia et al., 2009; Simon et al., 2011; see below).

The scale of adult oligogenesis has been something of a surprise. Rivers et al. (2008) calculated that ~29% of all differentiated oligodendrocytes (identified by CC1 immunolabeling) that are present in the corpus callosum of ~8-month-old mice are generated in the 210 days after P45 (*Pdgfra-CreER⁺: Rosa26-YFP*). Zhu et al. (2011) found that ~30% of CC1-positive oligodendrocytes in the corpus callosum of ~4-month-old mice were formed in the 60 days after P60 (*NG2-CreER⁺: Rosa26-YFP*). Comparing these estimates, one might conclude that no more oligodendrocytes are formed after 4 months of age, but Psachoulia et al. (2009) showed clearly that new myelinating oligodendrocytes are still being formed at a low rate even at 8 months. There are a lot of uncertainties in such calculations, (e.g., potential variation in recombination efficiencies from experiment to experiment and at different ages) but, nevertheless, it is clear that oligodendrocyte differentiation continues well into adulthood (Figure 1E), though at a steadily decreasing rate (Rivers et al., 2008; Lasiene et al., 2009; Psachoulia et al., 2009; Kang et al., 2010; Simon et al., 2011; Zhu et al., 2011). NG2-glia in the cortical gray matter also continue to generate oligodendrocytes into adulthood, although the overall rate of oligogenesis in the cortex is considerably less than in the corpus callosum at most ages (Rivers et al., 2008; Kang et al., 2010; Simon et al., 2011; Zhu et al., 2011).

It is not known yet whether adult myelin genesis is required to replace myelin that degenerates through normal “wear and tear” or whether it adds to existing myelin. Only around 30% of axons in the corpus callosum of 8-month-old mice are fully myelinated (Sturrock, 1980), so there is plenty of scope there and in other

major white matter tracts for de novo myelination of previously naked axons. There is evidence from cumulative [³H]-thymidine labeling that oligodendrocytes accumulate modestly in the mouse corpus callosum during the first year, without significant turnover, supporting the idea of de novo myelination (McCarthy and Leblond, 1988). Electron microscopy also shows that the number of myelinated axons in the rodent corpus callosum increases well into adulthood (Nuñez et al., 2000; Yates and Juraska, 2007). If adult-born myelin is laid down exclusively on naked axons, we would expect not to find many new myelinating oligodendrocytes in fiber tracts that consist predominantly of myelinated axons, such as the optic nerve (Honjin et al., 1977); this remains to be tested. On the other hand, myelin turnover is suggested by the observation that average internode length decreases with age, shorter internodes being regarded as a hallmark of remyelination following myelin loss (Lasiene et al., 2009). Perhaps de novo myelination and myelin replacement go on concurrently in different parts of the CNS or within axon tracts, such as the corpus callosum, that contain a mixture of myelinated and unmyelinated axons. If myelin turnover turns out to be commonplace, how neural pathways can cope with continual loss and replacement of oligodendrocytes would need to be understood, because the loss of even one myelin internode has been predicted to cause conduction block (Koles and Rasminsky, 1972; Waxman and Brill, 1978; Smith et al., 1982). Whether action potentials are blocked or delayed will depend on the geometry of the affected fibers, including internode length and axon diameter (e.g., Bostock and Sears, 1976; Waxman and Brill, 1978; Bakiri et al., 2010). Nevertheless, given that one oligodendrocyte usually myelinates many axons, significant problems might be anticipated from oligodendrocyte turnover. Perhaps new internodes can intercalate between existing internodes—i.e., remyelination might initiate at nodes of Ranvier and gradually expand lengthwise, pushing aside the existing internodal sheath(s) while maintaining continuity of myelin. This brief discussion exposes gaps in our knowledge of basic myelin dynamics that need to be filled before we can hope to understand myelin maintenance and plasticity.

Speculations on the Role of Adult Myelination in Learning and Memory

Personal experience tells us that learning a complex motor skill—riding a bicycle, playing a musical instrument, learning a dance step or a sporting activity—requires a great deal of time and practice. On the other hand a motor skill, once learned, is difficult to lose and stays with us throughout our active life. The extended learning experience and long decay time seem consistent with the production and long-term survival of new cells. Could new myelin formation during postnatal life play a part in motor learning? Motor learning is an example of unconscious or “nondeclarative” learning, which includes habituation and classical conditioning (e.g., fear conditioning and Pavlovian conditioning). Nondeclarative learning and memory is an ancient system that is well developed in invertebrate animals—for example, the gill retraction reflex that has been studied in *Aplysia* and other marine molluscs. Studies of this and related phenomena have established that even very small nervous systems have the capacity to learn and remember past

experience and that such memories are an intrinsic part of the circuits involved in the behavioral response, not something that is generated or stored remotely (Carew and Sahley, 1986). The general idea is that, during repetitive use, circuit connectivity becomes strengthened so that the neurons comprising the circuit acquire a lower threshold for firing and become more likely to fire again in future—so-called “Hebbian learning.” How does this happen at a cellular and molecular level? It is well established that repetitive firing can induce changes to the molecular composition of the active synapses and that this can increase the strength of communication between pre- and post-synaptic cells (Milner et al., 1998). Synaptic strengthening is an established mechanism of long-term potentiation (LTP), a cellular correlate of learning and memory in both invertebrates and vertebrates (Bliss and Collingridge, 1993). With the evolution of myelin, vertebrates might have acquired an additional way of modulating circuit activity—by myelinating the interconnecting axons, if previously unmyelinated. New myelination would be expected to increase dramatically the speed of transmission of action potentials and alter the intrinsic circuit properties. Myelination would also provide neurotrophic and physical support to the circuit neurons and make for long-term survival.

There is some evidence that adult myelin genesis might contribute to motor learning in humans. For example, it has been reported that extensive piano practice (Bengtsson et al., 2005) or juggling (Scholz et al., 2009) can cause long-term changes to the structure of white matter tracts, including parts of the corpus callosum, as revealed by magnetic resonance imaging (MRI). It has also been reported that white matter structure is altered in children skilled in abacus use, which involves actual and imagined visuomotor activity (Hu et al., 2011). There is also evidence that training in working memory tasks results in changes in the structure of frontoparietal white matter (Takeuchi et al., 2010; for reviews see Fields, 2008; Ullén, 2009).

For new myelin to be involved in activity-dependent learning, there needs to be a mechanism for regulating oligodendrocyte generation and myelination according to circuit activity. Such a mechanism seems to exist. Recently, Li et al. (2010) showed that electrical stimulation of neurons in the motor cortex led to activity-dependent stimulation of proliferation of NG2-glia in the descending pyramidal (corticospinal) tract. Previously, Barres and Raff (1993) had shown that silencing retinal ganglion neurons, by injecting tetrodotoxin into the developing eye, inhibited proliferation of NG2-glia in the newborn rat optic nerve. Inhibition could be overcome by implanting PDGF-expressing cells next to the nerve, suggesting that electrical activity in retinal ganglion cell axons might normally regulate the supply of mitogens to NG2-glia—possibly by triggering its release from optic nerve astrocytes (Barres and Raff, 1993). This suggests one mechanism by which NG2-glia might sense electrical activity, which, at some threshold, might trigger them to divide and differentiate into myelinating oligodendrocytes. A more direct mechanism involving synaptic communication between axons and NG2-glia can also be envisaged—for example, synaptic signaling might regulate the intrinsic response of NG2-glia to PDGF or other mitogens. Testing these ideas will be an important task for the future.

Does the Decreasing Rate of New Myelin Genesis Contribute to Age-Related Cognitive Decline?

There is an accumulating body of evidence that some measures of general cognitive ability correlate with white matter volume and integrity. For example, cognitive ability and white matter volume increase in parallel into the fourth decade of life and both decline thereafter (Bartzokis et al., 2001; Mabbott et al., 2006; Hasan et al., 2008; Ullén et al., 2008; Zahr et al., 2009; Bartzokis et al., 2010). The reasons behind these age-related changes are unknown but they could conceivably relate to changes in the ability of NG2-glia to proliferate and generate new oligodendrocytes as the brain matures and ages.

We recently measured the cell cycle time (Tc) of NG2-glia in the postnatal mouse brain by cumulative BrdU labeling (Psachoulia et al., 2009) and reported that Tc increases dramatically with age, from ~2 days on postnatal day 6 (P6) to >70 days at P240 (8 months of age) in the cerebral cortex. An age-related increase in the cell cycle time of NG2-glia in the mouse spinal cord has also been reported (Lasiene et al., 2009). The lengthening cell cycle results from the cells' spending more and more time in the early G1 phase of the cycle (Geha et al., 2010; Simon et al., 2011). The decreasing rate of cell division correlates well with the decreasing rate of oligodendrocyte production with age (Psachoulia et al., 2009)—as expected, since new oligodendrocytes must ultimately come from precursor cell divisions. If we assume that oligodendrocytes have a long but finite lifetime *in vivo*, it could be that as the division rate of NG2-glia decelerates and, with it, the rate of oligodendrocyte production, a critical age is reached beyond which the rate of new myelin production does not keep pace with accelerating myelin loss. If so, finding a way to maintain the proliferative rate of NG2-glia in old age might help maintain white matter integrity and slow down age-related mental decline.

Conclusion

Recent experiments indicate that NG2-glia are, first and foremost, oligodendrocyte precursors in the healthy adult CNS. Thus, it is clear that NG2-glia are distinct from neural stem cells that generate hippocampal or olfactory neurons throughout life. Whether they can generate rare neurons in the piriform cortex, as reported by two labs recently, is still unresolved. Following CNS injury, NG2-glia undergo a burst of local proliferation before giving rise to oligodendrocytes and possibly some astrocytes. Following gliotoxin-induced focal demyelination in the spinal cord, they also generate significant numbers of remyelinating Schwann cells. The great majority of reactive astrocytes at sites of damage are not derived from NG2-glia, but from pre-existing astrocytes that re-enter the cell cycle and—in spinal cord—from stem-like cells in the ependymal zone around the central canal. That NG2-glia are mainly precursors of myelinating cells refocuses our attention on their crucial role in remyelination following demyelinating disease or injury. We are also drawn to the role of myelin genesis during normal healthy adulthood, which might play a role in some forms of neural plasticity—motor skills learning, for example. The fact that NG2-glia react rapidly to injury suggests that they might also respond to systemic modulation. Indeed, their cell cycle and/or differentiation rate can be influenced by prolactin levels during pregnancy (Gregg

et al., 2007) or by physical exercise (Simon et al., 2011). Therefore, a key research focus for the future is the potential role of adult myelination in learning and memory and how that might be affected by the environment.

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