

Origin of Oligodendrocytes in the Subventricular Zone of the Adult Brain

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Glial fibrillary acidic protein (GFAP)-positive astrocytes (type B cells) in the subventricular zone (SVZ) generate large numbers of new neurons in the adult brain. SVZ stem cells can also generate oligodendrocytes *in vitro*, but it is not known whether these adult primary progenitors generate oligodendrocytes *in vivo*. Myelin repair and oligodendrocyte formation in the adult brain is instead associated with glial-restricted progenitor cells, known as oligodendrocyte progenitor cells (OPCs). Here we show that type B cells also generate a small number of nonmyelinating NG2-positive OPCs and mature myelinating oligodendrocytes. Some type B cells and a small subpopulation of actively dividing type C (transit-amplifying) cells expressed oligodendrocyte lineage transcription factor 2 (Olig2), suggesting that oligodendrocyte differentiation in the SVZ begins early in the lineage. Olig2-positive, polysialylated neural cell adhesion molecule-positive, PDGF receptor α -positive, and β -tubulin-negative cells originating in the SVZ migrated into corpus callosum, striatum, and fimbria fornix to differentiate into the NG2-positive nonmyelinating and mature myelinating oligodendrocytes. Furthermore, primary clonal cultures of type B cells gave rise to oligodendrocytes alone or oligodendrocytes and neurons. Importantly, the number of oligodendrocytes derived from type B cells *in vivo* increased fourfold after a demyelinating lesion in corpus callosum, indicating that SVZ astrocytes participate in myelin repair in the adult brain. Our work identifies SVZ type B cells as progenitors of oligodendrocytes in normal and injured adult brain.

Key words: adult oligodendrogenesis; CNS stem cells; oligodendrocyte progenitors; SVZ astrocyte; myelin repair; Olig2

Introduction

Oligodendrocytes are the myelinating cells of the CNS and are essential for proper brain function. Most oligodendrocytes develop during embryogenesis and early postnatal life from restricted periventricular germinal regions (Warf et al., 1991; Pringle and Richardson, 1993; Timsit et al., 1995; Olivier et al., 2001; Rowitch, 2004; Vallstedt et al., 2004). Forebrain oligodendrocytes originate first from progenitors in the embryonic ventral telencephalon (Spassky et al., 1998; Woodruff et al., 2001). This early wave of oligodendrocytes is transitory and superseded by oligodendrocytes originating later in development from dorsal telencephalic sources (Kessar et al., 2006). From their sites of origins,

oligodendrocyte progenitors (OPCs) migrate to colonize the gray and white brain matter (Thomas et al., 2000; Qi et al., 2002). Many OPCs, characterized by the expression of NG2 chondroitin sulfate, divide to form mature myelinating oligodendrocytes, but others linger as slowly dividing immature cells widely distributed within the adult brain (Polito and Reynolds, 2005). Interestingly, oligodendrocytes continue to be produced in the adult brain (Blakemore, 1972; Prineas and Connell, 1979; Kaplan and Hinds, 1980; McCarthy and Leblond, 1988), and these cells participate in myelin repair (Gensert and Goldman, 1997; Chari and Blakemore, 2002). Here we investigated whether neural stem cells present in the subventricular zone (SVZ) of the adult brain generate new OPCs and mature oligodendrocytes and whether these stem cells contribute to the repair of myelin.

In neonates, retroviral lineage studies have shown that the SVZ is an important source of astrocytes, oligodendrocytes, and neurons (Levison and Goldman, 1993; Luskin, 1993). In the adult, the SVZ continues to generate neurons that reach the olfactory bulb (OB) through the rostral migratory stream (RMS) (Lois and Alvarez-Buylla, 1994). Several lines of evidence indicate that oligodendrocyte progenitors also persist in the adult SVZ. Adult neural stem cells grown from the adult SVZ as neurospheres can generate oligodendrocytes (Morshead et al., 1994; Gritti et al., 1996; Weiss et al., 1996; Doetsch et al., 1999; Pluchino et al., 2003). Importantly, adult SVZ progenitors generate new

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oligodendrocytes *in vivo* after demyelinating lesions of the corpus callosum (CC) (Nait-Oumesmar et al., 1999; Picard-Riera et al., 2002) or after seizures (Parent et al., 2006). The primary progenitors for new oligodendrocytes born in the adult SVZ have not been identified.

Glial fibrillary acidic protein (GFAP)-expressing SVZ astrocytes serve as primary precursors of new neurons (Doetsch et al., 1999). These astrocytes, or SVZ type B cells, generate transit amplifying type C cells that express transcription factors *Dlx2* (distal-less homeobox 2) (Doetsch et al., 2002) and *Mash1* (mammalian achaete-schute homolog 1) (Parras et al., 2004). Here, we show that type B cells at different rostrocaudal levels of the SVZ generate not only neurons but also oligodendrocytes. Type B cells give rise to migrating young oligodendrocytes that differentiate into immature NG2-positive (NG2⁺) cells that continue to divide locally or mature into myelinating cells. Furthermore, primary cultures of single SVZ type B cells gave rise to oligodendrocytes alone or oligodendrocytes and neurons. This work identifies cells and regions in the SVZ giving rise to new oligodendrocytes in the adult brain.

Materials and Methods

Animal care and tissue processing. *Gtv-a* (Holland and Varmus, 1998) or CD-1 (Charles River Laboratories, Wilmington, MA) mice were housed in the University of California, San Francisco Animal Care Facility and were maintained in accordance with the University of California Committee on Animal Research guidelines. For surgical procedures, adult mice were deeply anesthetized with 60 mg/kg body weight of pentobarbital (Nembutal) injected intraperitoneally and maintained with 5% halothane when necessary. For tissue preparation, adult animals received an overdose of Nembutal (200 mg/kg body weight) before transcardial perfusion. For light microscopy, mice were perfused with 3% paraformaldehyde (PFA) in 0.1 M phosphate buffer (PB). Brains were postfixed overnight at 4°C in the same fixative. Whole mounts of the lateral wall of the lateral ventricle were dissected as described previously (Doetsch and Alvarez-Buylla, 1996), or 50- μ m-thick horizontal brain sections were cut with a vibratome. For electron microscopy, adult mice were killed by intracardial perfusion with 2% PFA and 0.8% glutaraldehyde in 0.1 M PB for alkaline phosphatase (AP) staining, 2% PFA and 0.5% glutaraldehyde for immunocytochemistry (ICC), or 2% PFA and 2.5% glutaraldehyde for conventional electron microscopy (EM). Brains were postfixed in the same fixative overnight at 4°C, cut horizontally at 100 μ m, and processed for EM as described previously (Doetsch et al., 1997). Briefly, for AP staining and ICC, tissue samples were washed in maleate buffer and postfixed in 1% osmium tetroxide in 0.1 M PB for 3 h, postfixed in 2% osmium for 2 h, rinsed, dehydrated, and embedded in Araldite. For conventional EM, sections were postfixed in 2% osmium for 2 h instead. Semithin sections, 1.5 μ m thick, were cut with a diamond knife and stained with 1% toluidine blue. For the identification of individual cell types, ultrathin (0.05 μ m) sections were cut with a diamond knife, stained with lead citrate, and examined under a Jeol (Peabody, MA) 100CX electron microscope.

Cell cultures. For coculture experiments, the protocol previously established in the laboratory (Lim and Alvarez-Buylla, 1999) was followed except that DMEM/F-12/B27 medium was used instead of Neurobasal. For single-cell experiments, the GFAP promoter (GFAP_p)-green fluorescent protein (GFP) adenovirus (100 nl) (Doetsch et al., 1999) was first injected *in vivo* into the SVZ of adult CD-1 mice 1.5 d before the dissection. Adult SVZ or striatal cells were then freshly dissociated, purified through a 22% Percoll gradient, and plated on top of a postnatal day 3 (P3) mouse cortical astrocyte monolayer in DMEM/F-12/B27 medium either at 50,000 cells/cm² or as a single cell after selection and isolation using a MoFlo fluorescence-activated cell sorter (DakoCytomation, Carpinteria, CA). Double cells were excluded using pulse width and integrated signal. Cultures were fixed at 1, 3, or 5 d *in vitro* (div) with 3% PFA for 20 min at room temperature and washed three times with PBS.

For cell blots, SVZ cells were dissected from adult CD-1 mice incubated in papain (14 U/ml; Worthington, Lakewood, NJ) for 30 min at

37°C, dissociated to single-cell suspension, and resuspended in DMEM/F-12 for 2 h before fixation with 3% PFA.

Immunocytochemistry. After blocking in PBS containing 10% of normal goat serum for 2 h at room temperature, cells or tissue sections were incubated with primary antibodies overnight at 4°C in the blocking solution. The following primary antibodies were used: mouse monoclonal anti- β -tubulin (Tuj1) (1:500; Berkeley Antibody Company, Berkeley, CA), anti-O₄ (1:50; Chemicon, Temecula, CA), anti-GFAP (1:500; Chemicon), anti-polysialylated neural cell adhesion molecule (PSA-NCAM) (1:400; AbCys, Annecy, France), chicken monoclonal anti-GFP (1:1000; Aves Lab, Tigard, OR), rabbit polyclonal anti-oligodendrocyte lineage transcription factor 2 (Olig2) (1:5000) (Ligon et al., 2004), anti-platelet-derived growth factor receptor α (PDGFR α) (1:50; Santa Cruz Biotechnology, Santa Cruz, CA), for identification of NG2 we used AN2 (1:1000) (Niehaus et al., 1999) and anti-NG2 (Chemicon) anti-myelin basic protein (MBP) (1:100; Chemicon), and anti-CNPase (1:500; Chemicon). For fluorescent ICC, cells, tissue sections, or whole mounts were washed in PBS, incubated with the appropriate secondary antibodies [all cyan at 1:400 (Jackson ImmunoResearch, West Grove, PA); all Alexa at 1:1000 (Invitrogen, Carlsbad, CA)] in blocking solution for 2 h at room temperature [cyanine 2 (Cy2) anti-mouse IgG, Cy3 anti-mouse IgM, Cy3 anti-rabbit IgG, Alexa-488 anti-chicken IgG, or Alexa-564 anti-rabbit IgG], and washed in PBS. For preembedding Olig2 ICC, sections were washed in PB, incubated in blocking solution with a biotinylated anti-rabbit (1:400; Jackson ImmunoResearch) overnight at 4°C, incubated in ABC peroxidase kit (Vector Laboratories, Burlingame, CA) for 2 h, and revealed with 0.02% diaminobenzidine and 0.01% H₂O₂. Controls in which primary antibodies were omitted resulted in no detectable staining.

Retroviral injections. We used a β -actin_p-GFP retrovirus (200 nl) (Carleton et al., 2003) to label all dividing cells in the SVZ and CC. Before injection, β -actin_p-GFP virus was mixed with polybrene (8 μ g/ml; Sigma, St. Louis, MO). Two types of replication competent avian retroviruses (RCASs) (100 nl injection volume) were used to label dividing GFAP-expressing SVZ astrocytes in P60–P120 *Gtv-a* mice (Holland and Varmus, 1998): RCAS-AP (Holland and Varmus, 1998), which encodes AP, and RCAS-GFP (a gift from M. Goetz, GSF, Institut für Stammzellforschung, Neuherberg, Germany), which encodes GFP. RCAS-AP labels thin processes allowing morphological characterization of cells under bright field. This virus, however, labeled fewer cells than the RCAS-GFP, and AP staining is not compatible with most ICC techniques. For ICC costaining, we therefore used the RCAS-GFP retrovirus. A GFAP_p-GFP adenovirus (100 nl) (Doetsch et al., 1999) was used to label SVZ astrocytes for cell sorting and *in vitro* clonal analysis. Injections were made stereotaxically at the following coordinates (anteroposterior relative to bregma, mediolateral, and dorsoventral from surface of the brain): for SVZ, as shown in Table 1; for CC, (1, 1.8, 1.25); or as described in Results. Brains were processed for AP staining as described below. As a control for specificity, a mixture of RCAS-AP and adenovirus β -actin_p-GFP (a gift from Dr. C. Lois, Massachusetts Institute of Technology, Cambridge, MA) (10:1) was coinjected bilaterally into the CC of two adult *Gtv-a* mice.

Demyelinating lesion. Demyelination was induced by injecting lyssolecithin (LPC) into corpus callosum as described previously (Gensert and Goldman, 1997; Nait-Oumesmar et al., 1999). To label the SVZ astrocytes and their progeny, RCAS-GFP was injected into the anterior SVZ of adult P60–P120 GFAP-Tv-a mice. Two days later, 0.5 μ l of 1% LPC (Sigma) dissolved in 0.9% NaCl was ipsilaterally injected into the corpus callosum.

Immunohistochemistry with a rabbit anti-degraded MBP antibody (Chemicon) was performed to detect the demyelinating lesions and verify the extent of demyelination. The control vehicle group received 0.5 μ l of 0.9% NaCl. All animals ($n = 3$ per group) were killed 30 d after lyssolecithin or vehicle injection.

Alkaline phosphatase staining. Endogenous AP activity was inactivated by heating the sections in a 65°C PBS bath for 30 min. The sections were then washed in AP detection buffer (in mM: 100 Tris-HCl, pH 9.5, 100 NaCl, and 5 MgCl₂) for 10 min and incubated in AP buffer containing nitroblue tetrazolium chloride (10 μ l/ml; Boehringer Mannheim,

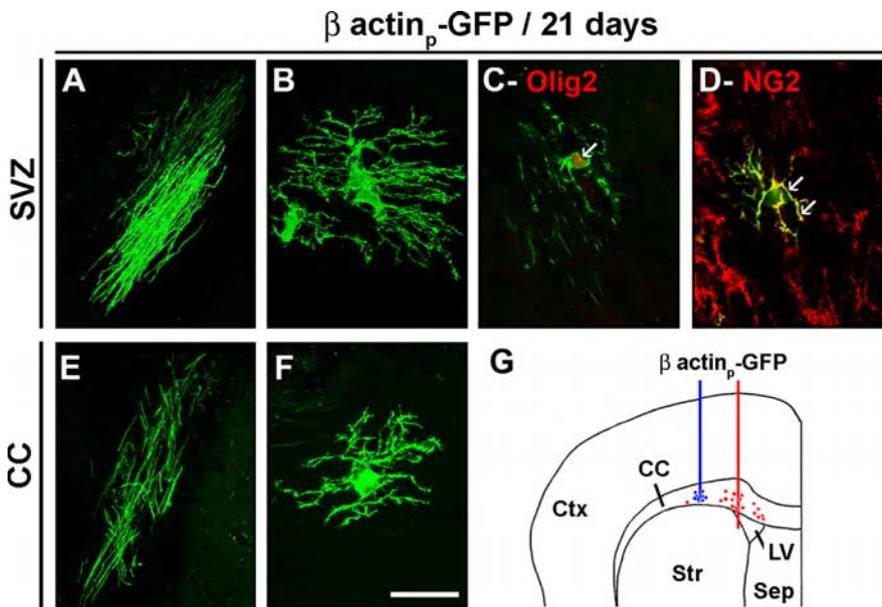


Figure 1. Oligodendrocyte progenitors exist in the SVZ and in the CC of the adult mouse brain. Replication-incompetent murine β -actin_p-GFP virus was injected into the SVZ (**A–D**) or the CC (**E, F**) of adult CD-1 mice. At 21 d after injection, GFP-labeled mature myelinating (**A, E**) and nonmyelinating (**B–D, F**) oligodendrocytes were identified in the CC. **C, D**, Double immunofluorescence with GFP (green) and Olig2 (**C**, red) or NG2 (**D**, red). Note that highly branched GFP⁺ nonmyelinating oligodendrocytes express both markers. **G**, Schematic representation of an adult mouse brain hemisphere showing the location of GFP-labeled cells in the CC after β -actin_p-GFP injection into the SVZ (red) or the CC (blue). Injection tracks for the SVZ (red) and CC (blue) are represented. Note that GFP-labeled cells are widely dispersed in the adult CC after SVZ retroviral injection in contrast to those generated after injection into the CC. Ctx, Cortex; LV, lateral ventricle; Sep, septum; Str, striatum. Scale bar, 60 μ m.

Mannheim, Germany) and 5-bromo-4-chloro-3-indolyl phosphate (2 μ l/ml; Boehringer Mannheim, Germany) for 3–5 h at room temperature. The reaction was stopped in fresh PBS containing 20 mM EDTA.

Results

Adult SVZ and CC contain proliferating oligodendrocyte progenitors

To analyze the progeny of adult SVZ cells, we stereotactically injected β -actin_p-GFP into the SVZ and the CC of the adult mouse brain. Three weeks after injection of this retrovirus into the SVZ, GFP-labeled granular and periglomerular neurons were observed in the OB of the injected hemispheres (data not shown), confirming that SVZ progenitors had been initially infected. GFP-labeled cells with the typical morphology of oligodendrocytes were also found dispersed in the CC. Because GFP labeling fills the cell processes, we were able to identify two subtypes of oligodendrocytes based on morphology: mature myelinating oligodendrocytes (Fig. 1A) (Levison and Goldman, 1993; Butt, 2004) and highly branched cells with the typical morphology of parenchymal OPCs, also known as nonmyelinating oligodendrocytes (Fig. 1B) (Butt, 2004). These GFP⁺ nonmyelinating oligodendrocytes expressed Olig2 (Fig. 1C) and NG2 (Fig. 1D). Proliferating oligodendrocyte progenitors are present in the adult brain parenchyma, including the CC (Polito and Reynolds, 2005). Consistently, injection of retrovirus encoding β -actin_p-GFP into the CC resulted in labeled myelinating (Fig. 1E) and nonmyelinating oligodendrocytes (Fig. 1F) 21 d later. However, in contrast to the SVZ injections, the β -actin_p-GFP retroviral injections into the CC did not label cells that migrate to the OB. Interestingly, GFP⁺ oligodendrocytes derived from the SVZ injections migrated as much as 550 μ m from the injection site. In contrast, labeled oligodendrocytes after CC injections were found no more than 50 μ m from the

injection site (Fig. 1G). Results indicate that dividing cells in both the SVZ and the CC generate oligodendrocytes, but progenitors in the CC give rise to oligodendrocytes that differentiate relatively close to the injection site, whereas oligodendrocytes derived from SVZ progenitors disperse throughout the CC and the neighboring striatum.

SVZ astrocytes give rise to oligodendrocytes in the adult mouse brain

SVZ type B cells, which have characteristics of astrocytes and express GFAP, have been identified as primary precursors of new neurons in the adult mouse brain (Doetsch et al., 1999). We next tested whether such SVZ astrocytes also generate oligodendrocytes *in vivo*. For this, we used transgenic Gtv-a mice expressing the avian leucosis virus receptor subgroup A (Tv-a) under the control of the GFAP promoter (Holland and Varmus, 1998). Infection of P60–P120 Gtv-a mice with RCAS-AP results in specific labeling of dividing GFAP⁺ cells (Holland and Varmus, 1998) and their progeny (Doetsch et al., 1999). We stereotactically injected RCAS-AP and RCAS-GFP retrovirus into the anterior region of the SVZ of adult Gtv-a mice. The

AP encoding RCAS virus allowed morphological identification and Golgi-like reconstruction of labeled cells, whereas RCAS virus encoding GFP allowed double-immunohistochemical labeling. Three weeks after RCAS virus injections into SVZ, AP⁺ or GFP⁺ astrocytes were observed close to the injection site in the SVZ, as well as labeled neurons in the OB (Doetsch et al., 1999). In all of these animals ($n = 18$), we also observed AP⁺ or GFP⁺ cells in the CC and the white matter tracks in the striatum and fimbria fornix that displayed the characteristic morphology of nonmyelinating (Fig. 2A) and myelinating (Fig. 2D, G, I) oligodendrocytes. RCAS-labeled oligodendrocytes were not observed in cortex. We used EM to further identify the oligodendrocytes derived from labeled SVZ astrocytes. AP-labeled cells were first identified as nonmyelinating (Fig. 2A) or myelinating (Fig. 2D) oligodendrocytes in 50- μ m-thick sections using light microscopy. Serial 1.5 μ m plastic sections of these same cells were reembedded and resectioned to 100 nm sections and analyzed by EM. Nonmyelinating oligodendrocytes were relatively large with an electron-lucent nucleus and a light cytoplasm with many polyribosomes (Fig. 2B, C). Myelinating cells had darker cytoplasm with prominent Golgi apparatus and a short endoplasmic reticulum. They also had an electron-dense nucleus with heavy clumps of chromatin that formed a prominent layer beneath the nuclear envelope (Fig. 2E, F). These are typical ultrastructural characteristics of nonmyelinating and myelinating oligodendrocytes (Peters et al., 1991). To confirm the identity of SVZ type B progeny in the CC, we stained tissue sections of brains infected with RCAS-GFP virus with several markers of mature oligodendrocytes. GFP-labeled cells expressed the CNS myelin protein CNPase (Fig. 2G–I) and MBP (Fig. 2J–L).

We next tested whether proliferating astrocytes also serve as

oligodendrocyte progenitors outside the SVZ. We injected RCAS-AP into the CC through the cortex but outside the SVZ ($n = 4$). To confirm the location of the injection, we coinjected an adenovirus encoding GFP (β -actin_p-GFP) that infects all cells. Twenty-one days after injection, GFP⁺ cells were observed in the CC (data not shown), but in none of the hemispheres injected did we observe AP-labeled oligodendrocytes. Similar results were observed after injection of the RCAS virus into different regions, including cortex ($n = 4$), medial and lateral striatum ($n = 4$), and lateral CC ($n = 2$). Occasionally, we found labeled astrocytes close to the injection track, but in none of these mice did we observe labeled oligodendrocytes. These findings indicated that GFAP⁺ oligodendrocyte precursors are in the SVZ.

Oligodendrocytes are generated from SVZ type B cells at different rostrocaudal levels of the adult mouse SVZ

The above experiments targeted the SVZ adjacent to the anterior horn of the lateral ventricle. To investigate whether astrocytes at different rostrocaudal locations in the SVZ generate oligodendrocytes, we used the RCAS-AP retrovirus injected at different rostrocaudal coordinates to target the SVZ, as indicated in Table 1. Animals were killed 21 d after injection; their brains were serially sectioned, processed for AP staining, and AP⁺ myelinating oligodendrocytes as well as differentiated AP⁺ OB neurons were counted. The number of mature oligodendrocytes produced from SVZ astrocytes at different rostrocaudal locations was very similar (Table 1). The number of oligodendrocytes formed was consistently lower compared with the number of neurons that migrate into and differentiate in the OB. This ratio, however, changed depending on the injection site; a proportionally higher number of OB neurons were produced from type B cells labeled in the anterior part of the SVZ compared with caudal SVZ. AP⁺ neurons migrated tangentially to the OB from all the different rostrocaudal SVZ locations tested as described previously (Doetsch and Alvarez-Buylla, 1996). In contrast, AP-labeled oligodendrocytes remained at approximately the same rostrocaudal level of the injection site but migrated laterally or dorsally into the CC, the fornix, and the fiber tracks of the striatum. We did not observe migration of oligodendrocyte precursors into the cortex, the hippocampus (HP), or the OB. These results indicate that both neurons and oligodendrocytes are formed from dividing SVZ astrocytes located at different rostrocaudal levels of the lateral ventricu-

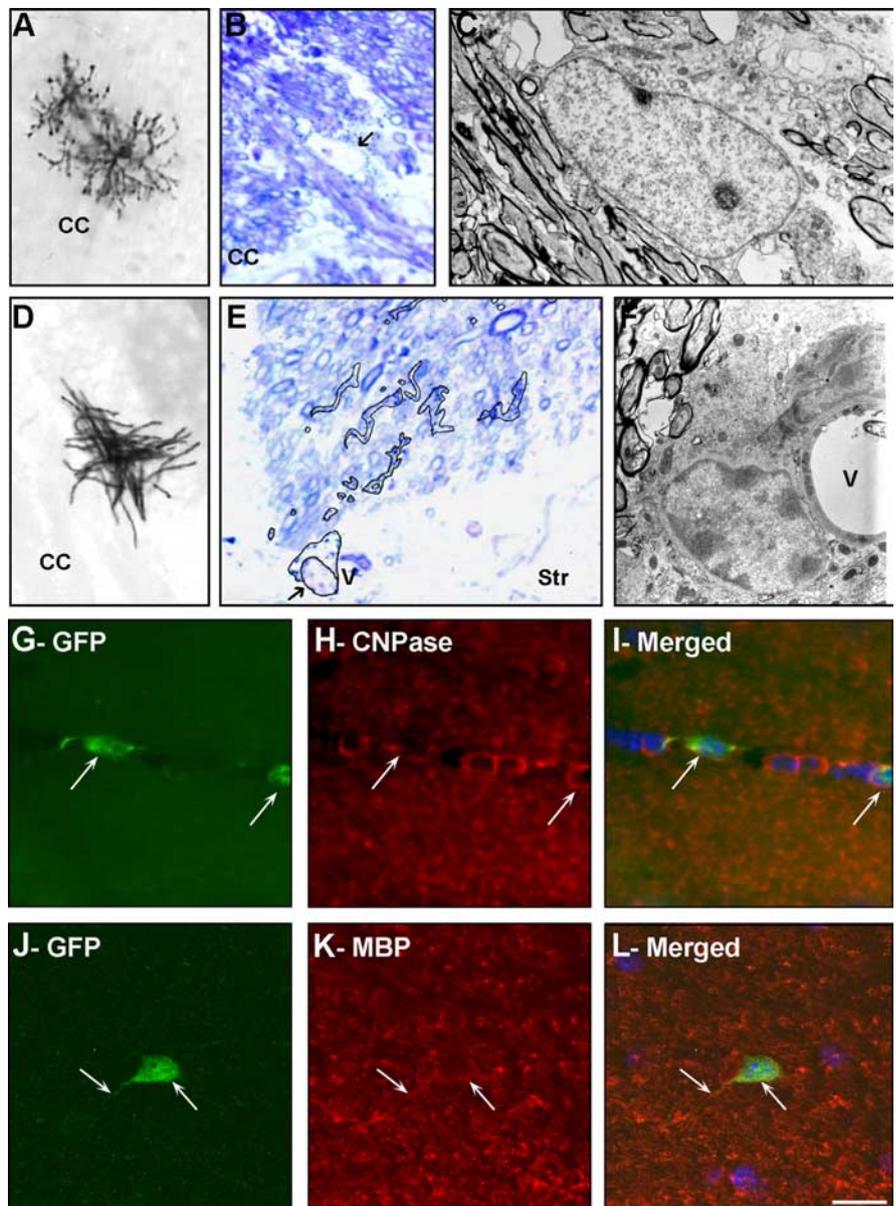
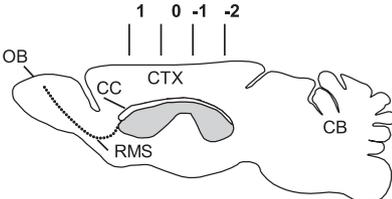


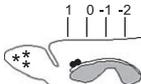
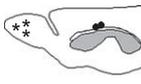
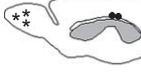
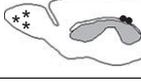
Figure 2. SVZ astrocytes give rise to oligodendrocytes in the adult mouse brain. RCAS-AP (A–F) or RCAS-GFP (G–L) retroviruses were injected into the SVZ of adult Gtv-a mice, and the progeny of the infected SVZ astrocytes was analyzed 21 d after injection using light (A, B, D, E), confocal (G–L), and electron (C, F) microscopy. A–F, Several AP-labeled cells were identified in the CC (A–D) and in the striatum (E, F): highly branched cells presenting typical morphology of nonmyelinating (A–C) and myelinating (D–F) oligodendrocytes. Characteristics of AP⁺ nonmyelinating oligodendrocytes (B, C) and AP⁺ mature myelinating oligodendrocytes (E, F) were determined at the ultrastructural level. B, E, AP-staining in 1.5- μ m-thick semithin sections counterstained with toluidine blue. Arrows indicate the nucleus of the cells analyzed at the electron microscopic level. In E, AP⁺ cell body and processes were traced on top of the picture. C, F, Electron micrograph of AP⁺ cells. Note the different ultrastructure and typical nuclear heterochromatin in myelinating oligodendrocytes compared with that of the nonmyelinating one. G–L, Double immunofluorescence for GFP (G, J) and the oligodendrocyte markers CNPase (H) or MBP (K) of cells in the CC. Merged field shows combined immunostaining for GFP and CNPase (I) or MBP (L). Arrows indicated SVZ-generated cells in the CC-expressing markers of mature oligodendrocytes. Str, Striatum; V, blood vessel. Scale bar: A, D, 80 μ m; B, E, 7 μ m; C, F, 2 μ m; G–L, 20 μ m.

lar wall. The paths of migration for these two cell types are very different. Young neurons migrate tangentially into the OB, whereas oligodendrocytes appear to migrate away from the SVZ and remain at rostrocaudal levels similar to their site of origin.

Identification of migrating oligodendroglial progenitors

In addition to the nonmyelinating and myelinating oligodendrocytes derived from our viral injections described above, we also

Table 1. Cell types produced after RCAS-AP infection at different rostrocaudal SVZ levels


Coordinates, mm			Oligodendrocytes	Neurons	Ratio O/N	Location
A-P	L	D				
1	1	2.3	2	51		
1	1	2.3	6	127	1/27	
1	1	2.3	0	42		
0	1.4	1.6	1	47		
0	1.4	1.6	1	10	1/30	
0	1.4	1.6	2	63		
-1	2.4	1.8	1	8		
-1	2.4	1.8	3	33	1/10	
-1	2.4	1.8	2	19		
-1.5	2.5	1.8	1	4		
-1.5	2.5	1.8	2	3	1/3	
-1.5	2.5	1.8	1	4		
-2	3.3	2	0	1		
-2	3.3	2	2	5	1/3	

This table indicates the number of AP⁺ mature myelinating oligodendrocytes and differentiated neurons found in the adult mouse brain 3 weeks after retroviral infection of astrocytes located at different rostrocaudal levels of the SVZ. The schematic at the top illustrates the lateral wall of the lateral ventricle and its coordinates relative to bregma. Bilateral injections of RCAS-AP were performed in adult Gtv-a mice ($n = 5$ for each coordinate). AP⁺ cells were counted in 1 of every 4 horizontal 50 mm sections, through the whole brain. Ratio O/N indicates the number of oligodendrocytes versus neurons. Note a similar number of oligodendrocytes produced regardless of the original injection coordinates in contrast to neurons. Schematic representation of AP⁺ cell distribution after retroviral infection of SVZ astrocytes at different locations is indicated on the right side of the table. Note that neurons were always found in the OB while oligodendrocytes were located at a rostrocaudal level similar to that of the injection site. CB, Cerebellum; CC, corpus callosum; CTX, cortex; OB, olfactory bulb; RMS, rostral migratory stream.

noticed the presence of elongated cells that resemble migrating cells. We first studied the distribution and antigenic properties of these putative migrating cells in the CC in whole-mount preparations of the SVZ and CC as described previously (Doetsch and Alvarez-Buylla, 1996). For these dissections, paraventricular segments facing the CC were dissected together with the lateral wall of the lateral ventricle. These paraventricular regions correspond to regions of the lateral ventricle that have become obliterated as both walls of the ventricle fuse during development. The most extensive paraventricular region, the subcallosal zone (SCZ), is located between the HP and the CC (Seri et al., 2006). Many elongated cells labeled by PSA-NCAM were present in this region (Fig. 3A). Similar cells were also present close to the SVZ and in more anterior CC. In contrast to the chains of PSA-NCAM⁺ cells described previously in the SVZ and RMS (Doetsch and Alvarez-Buylla, 1996; Peretto et al., 1997), PSA-NCAM⁺ cells in the CC and the SCZ above the HP were not arranged in chains, but the majority appeared to be migrating as individual cells. They had an elongated cell body, with a leading process tipped by a growth cone, and occasionally had a trailing process (Fig. 3B). Unlike PSA-NCAM⁺ young neurons, which also express neuron-specific Tuj1 (Fig. 3C–E) (Doetsch and Alvarez-Buylla, 1996), PSA-NCAM⁺ cells in the CC were negative for Tuj1 (Fig. 3F–H). PSA-NCAM⁺/Tuj1⁻ cells in the CC may correspond to migrating oligodendrocyte precursors (Ben-Hur et al., 1998). Consistently, PSA-NCAM⁺ cells in the CC were also positive for PDGFR α (Fig. 3I–K) and the NG2 chondroitin sulfate proteogly-

can (Fig. 3L–N), which have been shown to be expressed in oligodendrocyte progenitors (Pringle and Richardson, 1993; Nishiyama et al., 1996; Polito and Reynolds, 2005). We next investigated whether these putative migrating cells arise from SVZ astrocytes. RCAS-AP (Fig. 3O) or β -actin_p-GFP (Fig. 3P, Q) retroviruses were injected into the adult SVZ, and animals were killed 1 and 4 d later. In these animals, AP⁺ or GFP⁺ elongated cells were present in the SVZ and the CC. They stained for Olig2 (Fig. 3Q).

It was remarkable how similar the morphology of AP⁺ migrating cells was in the SVZ and CC. We confirmed this observation using EM and preembedding immunocytochemistry for PSA-NCAM. PSA-NCAM⁺ cells first identified in 2- μ m-thick sections (Fig. 3R) were resectioned and analyzed at the electron microscopic level (Fig. 3S). The cells had ultrastructural features of migrating cells: an elongated body and a leading process with a centrosome at its base (Fig. 3S). These elongated cells had a very different ultrastructure from that of astrocytes and oligodendrocytes in the CC. They had scant cytoplasm, numerous ribosomes, and no intermediate filaments. These characteristics are similar to those of young neurons migrating in the RMS and SVZ (Luskin, 1993; Lois et al., 1996; Wichterle et al., 1997). However, unlike young neurons in the SVZ and RMS, putative migrating young oligodendrocytes in the CC were not ensheathed by astrocytes. In contrast, they were tightly associated with myelinated and unmyelinated axons.

A subpopulation of SVZ type B and C cells express Olig2

The above results allowed us to identify oligodendrocytes at different stages of maturation after exiting the SVZ (supplemental data 2, available at www.jneurosci.org as supplemental material). To identify the cells in the SVZ that give rise to these migrating oligodendrocyte progenitors, we used antibodies that label oligodendrocytes at different stages of maturation (Baumann and Pham-Dinh, 2001). Olig2 is a transcription factor necessary for the development of oligodendrocyte lineage, receptor interacting protein (Rip) is expressed by nonmyelinating oligodendrocyte progenitors, and MBP and myelin/oligodendrocyte glycoprotein (MOG) are markers of mature myelinating oligodendrocytes. No labeling was observed in the adult mouse SVZ for Rip, MBP, and MOG (data not shown), indicating that mature oligodendrocytes were not present in this germinal region. As described previously (Hack et al., 2005), Olig2-expressing cells were observed in the SVZ. They were identified as an actively dividing cell population (supplemental data 1, available at www.jneurosci.org as supplemental material) (Hack et al., 2005) often organized as small clusters (Fig. 4A), suggesting that they may correspond to type C cells (Doetsch et al., 1997; Parras et al., 2004). Interestingly, only one or two cells expressed Olig2 within the clusters (Fig. 4A, E). To determine to which cell type Olig2⁺ SVZ cells correspond, we studied 34 Olig2⁺ cells with the electron microscope. Of these

cells, 16 displayed typical characteristics of type C cells (Fig. 4*A,B*): abundant mitochondria, short endoplasmic reticulum and large Golgi apparatus, few microtubules, and intermediate filaments (Fig. 4*B*, inset). Nine of the 34 studied cells had an elongated morphology, and these cells were found to contact myelinated axons (Fig. 4*C,D*). These cells probably correspond to early migrating oligodendrocyte precursors.

Interestingly, the other 15 Olig2-expressing cells analyzed at the electron microscopic level had bundles of intermediate filaments and light cytoplasm typical of SVZ astrocytes (Fig. 4*F*). To confirm this observation at the light microscopic level, we performed double ICC in mouse brain tissue using the anti-Olig2 and the anti-GFAP antibodies. A small fraction of Olig2⁺ nuclei were clearly surrounded by GFAP⁺ staining, suggesting colocalization (Fig. 4*G*). However, this could be attributable to GFAP⁺ processes from neighboring cells wrapping around Olig2⁺ cells. To clearly colocalize the label in individual cells, we freshly dissociated cells from adult SVZ and stained them for Olig2 and GFAP (Fig. 4*H*). We observed that a small fraction ($3.3 \pm 0.4\%$, mean \pm SEM; $n = 4$ independent dissection) of Olig2⁺ cells were also GFAP⁺. In addition, we found that $10.0 \pm 0.7\%$ (mean \pm SEM; $n = 3$ independent dissection) of SVZ Olig2⁺ cells also expressed S100 β (Fig. 4*I*), which is considered a marker for astrocytes by some authors (Reeves et al., 1994; Nishiyama et al., 2002) but has also been suggested to be expressed in cells in the oligodendrocyte lineage (Jennings et al., 2002; Deloume et al., 2004).

Higher number of oligodendrocytes are derived from SVZ type B cells after a demyelinating lesion

Previous studies suggested that progenitor cells in the SVZ contribute to the repair of demyelinating lesions (Nait-Oumesmar et al., 1999; Picard-Riera et al., 2002). We next investigated whether oligodendrocytes derived from SVZ astrocytes contribute to the repair of white matter tracts after a demyelinating lesion. SVZ astrocytes were first labeled by injection of RCAS-GFP into the anterior SVZ of adult GFAP-Tv-a mice. Two days later, lyssolecithin or vehicle was ipsilaterally injected into the CC. Thirty days after demyelination, a 4.3-fold increase in the number of GFP⁺ cells in the corpus callosum of lyssolecithin-injected hemispheres was observed (650 ± 35 cells; $n = 3$) compared with the control vehicle group (149 ± 58 cells; $n = 3$; $p <$

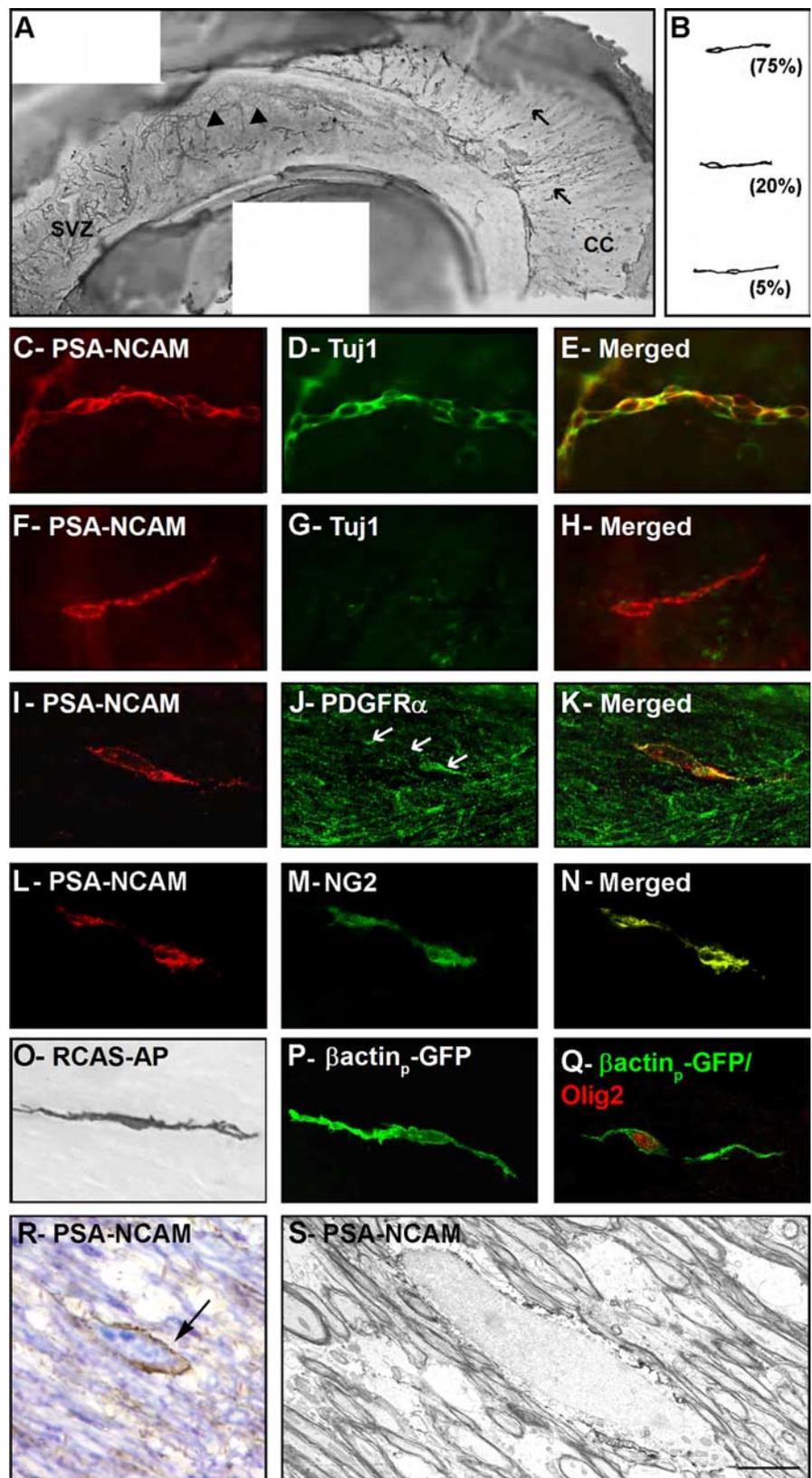


Figure 3. PSA-NCAM-immunopositive cells migrate tangentially within the CC. **A**, Photomontage of PSA-NCAM immunostaining of whole-mount preparation of the lateral wall of the lateral ventricle and the CC. Dorsal is up, and rostral is left. Filled arrowheads show chains of PSA-NCAM⁺ cells migrating on the lateral wall of the lateral ventricle. Arrows show PSA-NCAM⁺ cells migrating within the CC. **B**, Camera lucida drawing of PSA-NCAM⁺ cells observed in the CC. These cells are unipolar, with a short leading process, or bipolar. Percentages of cells displaying each morphology are indicated. **C–N**, Double immunofluorescence for PSA-NCAM (**C, F, I, L**) and Tuj1 (**D, G**), PDGFR α (**J**), or NG2 (**M**) of cells localized on the lateral wall of the lateral ventricle (**C–E**) and on the CC (**F–N**). Merged field shows combined immunostaining for PSA-NCAM and . (Figure legend continues.)

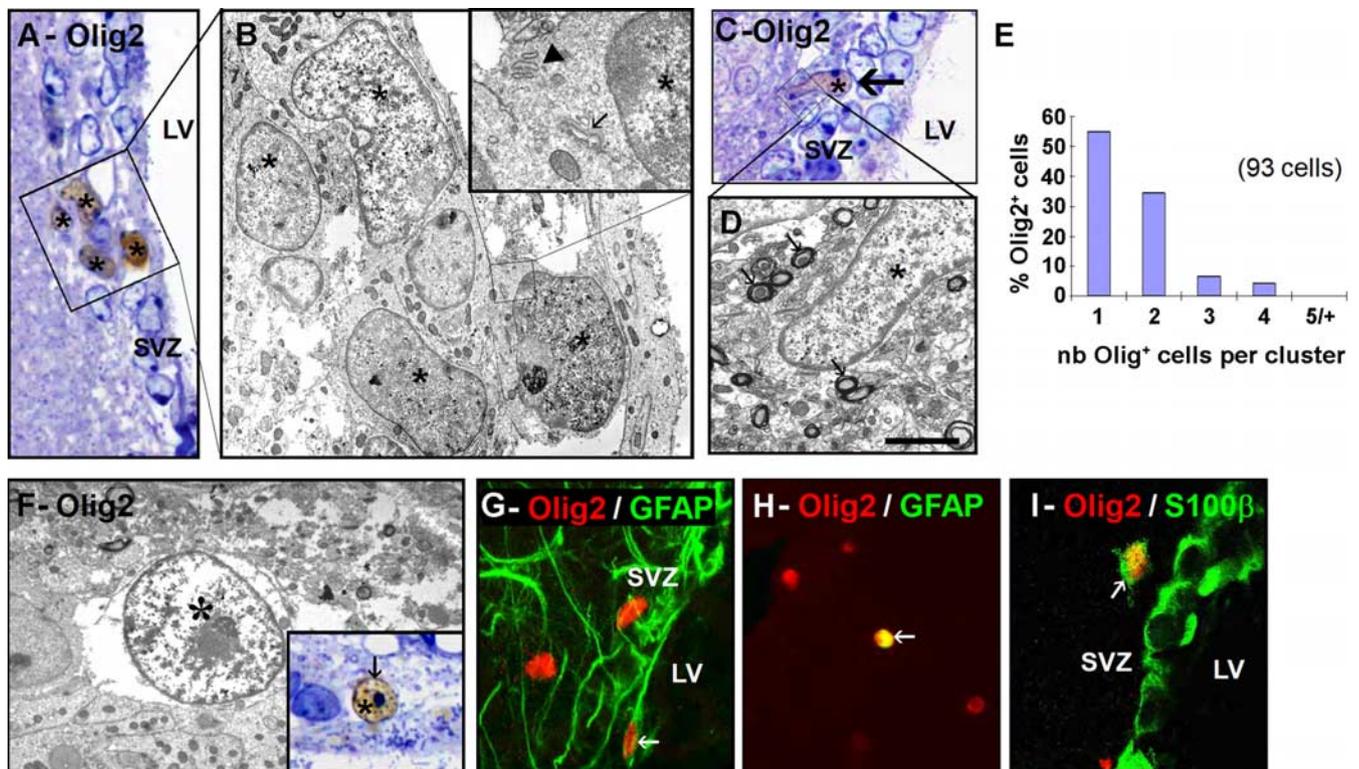


Figure 4. The basic helix–loop–helix transcription factor Olig2 is expressed by a subpopulation of immature transit-amplifying type C cells and by some SVZ astrocytes. Electron microscopic characterization of Olig2⁺ cells in the adult mouse SVZ. **A, C**, Inset in **F**, Olig2 staining in plastic 1.5- μ m-thick semithin section counterstained with toluidine blue. Arrows or asterisks indicate Olig2-labeled cells analyzed at the electron microscopic level. **B, D, F**, Electron micrograph of these cells. Asterisks show the nucleus. **A–D**, Olig2⁺ cells display the ultrastructural characteristics of type C cells: medium dark cytoplasm, numerous mitochondria, large Golgi apparatus (arrows in inset in **B**), and short endoplasmic reticulum (arrowheads in inset in **B**). Note that some Olig2⁺ cells have an elongated morphology (**C**). The cytoplasm of Olig2⁺ cell is often in contact with myelinated axons (arrows in **D**). **E**, Olig2⁺ cells were grouped in small clusters in the SVZ. Ninety-three Olig2⁺ cells were analyzed in 1.5- μ m-thick sections. The diagram indicates the percentage of Olig2⁺ cells that were either alone or associated in small groups of two to five cells. nb, Number of. **F–I**, Olig2⁺ SVZ cells can also present ultrastructural characteristics of type B SVZ astrocytes that appear to be acquiring some characteristics of type C cells. **F**, Electron micrograph of such a cell. **G–I**, This observation was confirmed by performing double immunofluorescence for Olig2 (red) and GFAP (green, **G, H**) or S100 β (green, **I**) performed on coronal sections of adult mouse brain (**G, I**) or on freshly dissociated adult SVZ cells (**H**). Arrows indicate cells expressing both markers. LV, Lateral ventricle. Scale bar: **A, C**, inset in **F**, 10 μ m; **B, D, F**, 3 μ m; inset in **B**, 1 μ m; **G, I**, 20 μ m; **H**, 40 μ m.

0.01) (Fig. 5A, B). Moreover, these GFP⁺ cells appear to be concentrated within and around the demyelination site (Fig. 5B). GFP⁺ cells in the lesion site had the morphology of both mature and immature oligodendrocytes (Fig. 5C). To further characterize these GFP⁺ cells, we coimmunostained these sections with antibodies for NG2, S100 β , and Olig2. Similar proportions of cells stained with these markers were observed in the lesioned and vehicle injected groups: $69.8 \pm 11\%$ S100 β ⁺/GFP⁺ in the lysolecithin versus $62.8 \pm 5\%$ in the control; $42.8 \pm 10\%$ Olig2⁺/GFP⁺ in the lysolecithin versus $35.1 \pm 12\%$ in control; and $48.5 \pm 6\%$ NG2⁺/GFP⁺ in the lysolecithin versus $40.2 \pm 8\%$ in control. We also analyzed an additional control group ($n = 3$) that received RCAS-GFP in the SVZ but in which injection of saline into the corpus callosum was omitted. The number of GFP⁺ cells and proportion of these cells expressing the different oligodendrocyte markers was similar between the saline-injected

and these control animals that only received the virus injection (data not shown). These results suggest that, after the lesion, the increased number of cells derived from SVZ astrocytes had similar characteristics as those normally produced in the control animals. GFP⁺ cells in the CC of lesioned brains expressed CNPase, a marker for CNS myelin protein (Fig. 5D). The precise number of CNPase-positive cells was difficult to quantify because of the diffuse pattern of staining of this marker. However, clear examples of double-labeled cells were observed, indicating that a subpopulation of the cells produced after the lesion mature into myelinating oligodendrocytes.

SVZ astrocytes generate oligodendrocytes and neurons *in vitro*

SVZ cells propagated as neurospheres can differentiate into both oligodendrocytes and neurons *in vitro* (Morshead et al., 1994; Gritti et al., 1996; Weiss et al., 1996; Imura et al., 2003). However, neurospheres are exposed to growth factors, which may modify the normal differentiation program of neural progenitors (Doetsch et al., 2002; Gabay et al., 2003; Santa-Olalla et al., 2003; Hack et al., 2004; Reynolds and Rietze, 2005). To determine whether SVZ progenitors could give rise to oligodendrocytes *in vitro*, we plated freshly dissociated adult SVZ cells onto a

(Figure legend continued.) Tuj1 (**E, H**), PDGFR α (**K**), or NG2 (**N**). PSA-NCAM⁺ cells migrating tangentially as chains expressed the neuronal marker Tuj1 (**C–E**). In contrast, PSA-NCAM⁺ cells present at the surface of the CC did not colabel with Tuj1 (**F–H**) but with the oligodendrocyte markers PDGFR α (**I–K**) and NG2 (**L–N**). **O–Q**, Putative migrating cells located in the CC and labeled after injection of RCAS-AP (**O**) or β -actin₁-GFP (**P, Q**) into the adult mouse SVZ. **Q**, Double immunofluorescence for GFP (green) and Olig2 (red) of a bipolar cell. **R, S**, Electron microscopic characterization of a PSA-NCAM-positive cell in the CC. **R**, PSA-NCAM staining in toluidine blue-stained 1.5- μ m-thick semithin section. Arrow indicates the PSA-NCAM⁺ cell analyzed at the electron microscopic level. **S**, Electron micrograph of this PSA-NCAM⁺ cell. Note that its elongated nucleus contains lax chromatin. Scale bar: **A**, 400 nm; **C–E**, 60 μ m; **F–Q**, 50 μ m; **R**, 5 μ m; **S**, 2 μ m.

confluent monolayer of cortical astrocytes using a modified protocol (see Materials and Methods) used previously to study neuronal differentiation (Lim and Alvarez-Buylla, 1999). We show here that this culture conditions results in robust oligodendrocyte formation from SVZ precursors. One day after plating, surviving SVZ cells were present as single cells. By 3 div, small clusters of three to five cells appeared (Fig. 6A). At this time, double immunofluorescence for Tuj1 and the O₄ antibodies revealed the presence of cells expressing the neuronal or oligodendroglial markers within the same cluster (Fig. 6B–D). Five days after plating, Tuj1⁺ cells (Fig. 6E) and O₄⁺ cells (Fig. 6F) displayed the morphology of neurons and oligodendrocytes, respectively. O₄⁺ cells also expressed Olig2 and NG2 (data not shown). Because oligodendrocyte progenitors are present within the striatum (some of which is dissected with the SVZ), it was possible that our cultures were contaminated with parenchymal OPCs. To control for this possibility, cells from the striatum adjacent to the SVZ of adult animals were plated on cortical astrocytes as described above. Under these culture conditions, neither neurons nor oligodendrocytes were observed in the cultures at 5 div. Finally, oligodendrocytes did not arise from cortical astrocytes without plating SVZ progenitors on top. These observations indicate that SVZ cells are able to generate both neurons and oligodendrocytes in our culture conditions, yet it was possible that oligodendrocytes and neurons were derived from two separate progenitor subpopulations.

To address this question, we examined the potential of single SVZ astrocytes plated onto the monolayer of cortical astrocytes. To isolate SVZ astrocytes, we used a GFAP_p-GFP adenovirus that encodes the GFP marker under the transcriptional control of the GFAP promoter (Doetsch et al., 1999) (Fig. 6G). SVZ astrocytes were first labeled *in vivo* by injecting the adenovirus into the SVZ of adult CD-1 mice. Thirty six hours after injection, GFP⁺ cells were isolated by fluorescence-activated cell sorting (FACS) (Fig. 6H) and plated as single cells on the monolayer of cortical astrocytes. Five days later, the cultures were fixed and immunostained with O₄ and Tuj1 antibodies (Fig. 6I). Twenty-six percent of plated cells gave rise to colonies. Of these, 11.5% generated both neurons and oligodendrocytes. Interestingly, the majority of colonies (80.3%) contained only oligodendrocytes but not neurons. This finding confirms that oligodendrocytes can be derived from GFAP-expressing SVZ cells and that a subpopulation of these cells can generate both neurons and oligodendrocytes.

Discussion

This study shows that adult SVZ astrocytes (type B cells) serve as primary progenitors of new oligodendrocytes in normal and in-

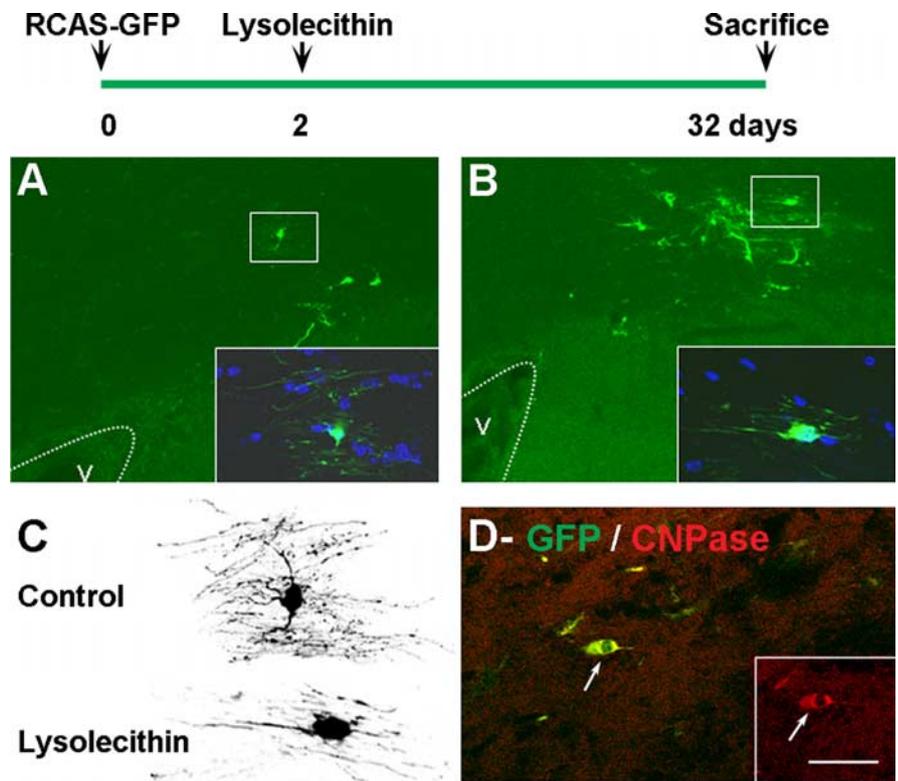


Figure 5. The SVZ astrocytes produce a higher number of oligodendrocytes after a demyelinating lesion. Schematic timeline shows the experimental design. The SVZ astrocytes of GTV-a mice were labeled with RCAS-GFP. Two days later, lyssolecithin or vehicle was injected into the ipsilateral CC. The cellular response to demyelination was evaluated 30 d after lesion. **A, B**, Anti-GFP immunostaining of coronal sections of vehicle-injected (**A**) and lyssolecithin-injected (**B**) mice. In both cases, mature and immature GFP⁺ oligodendrocytes were found in the corpus callosum. Insets show GFP⁺ oligodendrocytes counterstained with the nuclear marker DAPI (4',6'-diamidino-2-phenylindole) under higher magnification. **C**, Z-stack reconstruction of GFP⁺ oligodendrocytes found in corpus callosum in control and lyssolecithin-lesioned brains. **D**, Double immunofluorescence for GFP (green) and the CNS myelin protein CNPase (red) of GFP⁺ cells in the CC of a lesioned brain 30 d after injection of the RCAS-GFP virus. Inset shows the CNPase staining alone in red. Arrows indicate double-labeled cell. V, Lateral ventricle. Scale bar: **A, B**, 160 μ m; insets in **A, B**, 40 μ m; **D**, inset in **D**, 25 μ m.

jured brain. Results suggest that SVZ type B cells, through intermediate Olig2⁺ transit-amplifying cells (type C cells), give rise to young migrating oligodendrocytes that move out of the SVZ into the CC, neighboring striatum, and fimbria fornix to differentiate into nonmyelinating and myelinating oligodendrocytes. Compared with the large number of new neurons born in the SVZ, few oligodendrocytes are normally produced from adult SVZ type B cells. This may be related to the slow turnover of oligodendrocytes in adult brain (McCarthy and Leblond, 1988). Interestingly, however, the number of oligodendrocytes derived from SVZ type B cells was increased after a demyelinating lesion. In these conditions, SVZ-generated oligodendrocytes express markers of myelin, suggesting that they contribute to the remyelination process. This is consistent with the suggestion that, in addition to the parenchymal OPC progenitors (Fancy et al., 2004), SVZ progenitors contribute to the repair of myelinated tracks (Nait-Oumesmar et al., 1999; Picard-Riera et al., 2002; Petratos et al., 2004).

Labeling dividing cells with β -actin_p-GFP retrovirus targeted to either the SVZ or the CC resulted in labeled oligodendrocytes close to the injection site. In contrast, labeling with a retrovirus that infects GFAP-expressing cells results in labeled oligodendrocytes only when the injection is targeted to the SVZ. This suggests that oligodendrocytes are derived from two sources of adult dividing cells, GFAP-expressing SVZ cells and non-GFAP dividing

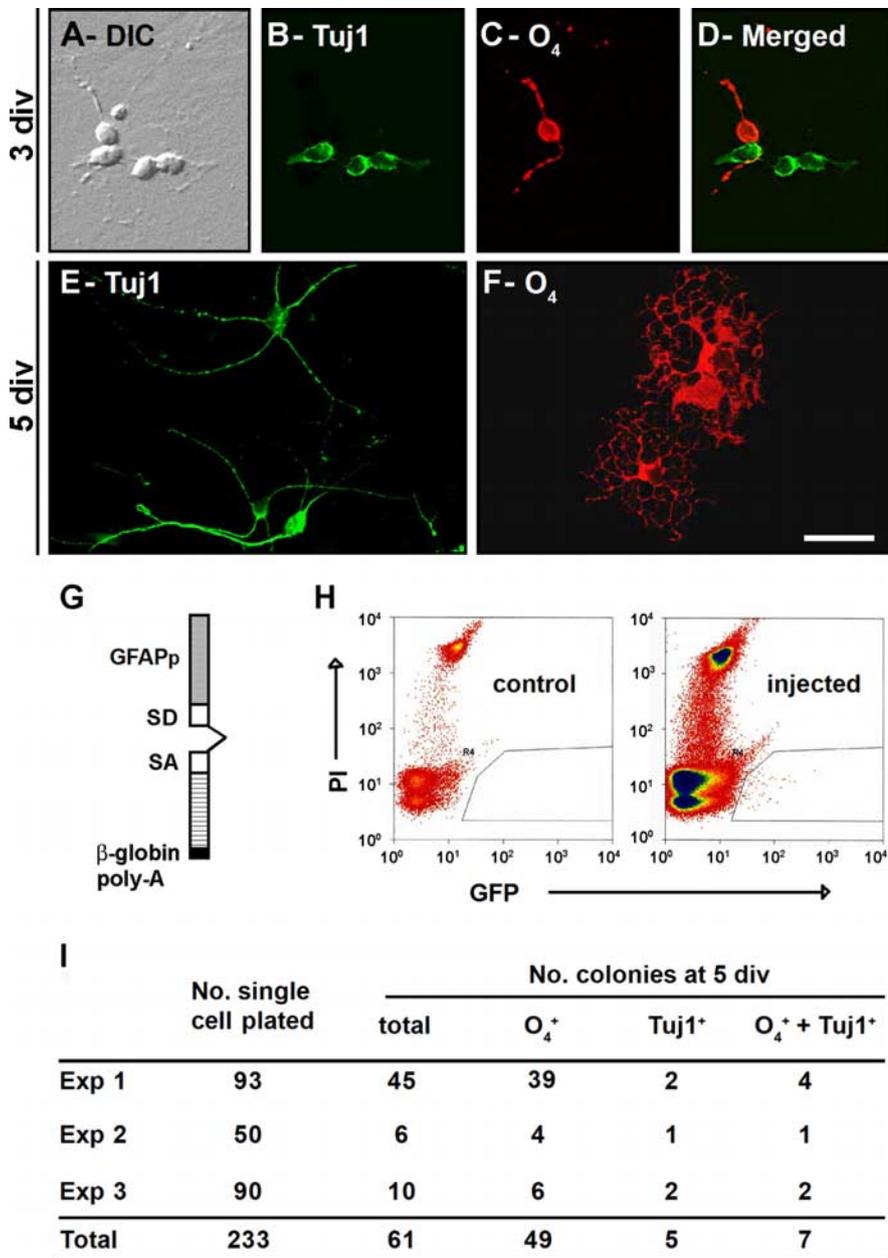


Figure 6. SVZ astrocytes produce oligodendrocytes *in vitro*, and some produce both neurons and oligodendrocytes. **A–F**, Ten thousand SVZ cells isolated from adult CD-1 mice were allowed to differentiate on top of an astrocytic monolayer for 3 div (**A–D**) and 5 div (**E, F**). **A–D**, At 3 div, clusters derived from adult SVZ precursor cells are mixed colonies which contain both O₄⁺ and Tuj1⁺ cells. **E, F**, At 5 div, Tuj1⁺ (**E**) and O₄⁺ (**F**) cells present typical morphologies of differentiated neurons and oligodendrocytes, respectively. **A**, Differential interference contrast (DIC) image of a typical colony. **B, E**, Immunoreactivity for the neuronal marker Tuj1. **C, F**, Immunoreactivity for the oligodendroglial marker O₄. **D**, Merged field showing combined immunostaining for Tuj1 and O₄. **G–I**, Single SVZ astrocytes give rise to both neurons and oligodendrocytes *in vitro*. Adult SVZ astrocytes were specifically labeled *in vivo* or *in vitro* with an adenovirus GFAP_p–GFP. **G**, Schematic showing the GFAP promoter reporter construct used for the adenoviral vector. **H, I**, Single GFP⁺ SVZ astrocytes were purified by FACS, plated on top of an astrocytic monolayer, and grown in culture for 5 div. **H**, Density plots showing gates for selecting SVZ astrocytes [GFP and propidium iodide (PI)]. The right plot corresponds to the sorting of SVZ cells isolated at 1.5 d after injection of the adenovirus GFAP_p–GFP; the left plot is a control non-injected animal. **I**, The number of GFP⁺ cells initially plated is indicated for each experiment (3 independent cultures). The total number of colonies that arose from these cells is presented. These colonies contained only O₄⁺, only Tuj1⁺, or both O₄⁺ and Tuj1⁺ cells. Note that 61 of the 233 single GFP⁺ SVZ cells initially plated gave rise to colonies. Among them, seven generated mixed colonies. Scale bar: **A–F**, 10 μm. SD, Splice donor; SA, splice acceptor.

cells outside the SVZ. Interestingly, the progeny of cells that divide locally within the CC do not appear to migrate far from the site of injection, which is consistent with previous observations showing that the progeny of parenchymal OPCs do not migrate

extensively (Gensert and Goldman, 1997). This is in contrast with the progeny derived from SVZ GFAP-expressing cells that can migrate long distances.

Classical models of cell lineages in the CNS propose that neural stem cells produce two separate pools of progenitors that generate either neurons or astrocytes and oligodendrocytes (Raff et al., 1983; Rao, 1999). However, committed glial precursors, like the optic nerve O-2A progenitor, can be manipulated to also generate neurons (Kondo and Raff, 2000). Recent evidence *in vivo* indicates that neurons and some glial cells are closely related in development and originate from common progenitors in ways that do not fit classical models that separate the neuronal and glial lineages (Williams et al., 1991; Richardson et al., 2000; He et al., 2001; Lu et al., 2002; Zhou and Anderson, 2002; Lui and Rao, 2004; Rowitch, 2004). Consistently, our data suggest that adult SVZ GFAP⁺ cells, which have been shown to function as neuronal precursors, can also give rise to oligodendrocytes. A small subpopulation of single astrocytes isolated *in vitro* gave rise to both neurons and oligodendrocytes. However, we do not know whether single progenitors *in vivo* have this dual potential or whether separate pools of type B cells that generate neurons or oligodendrocytes coexist in the SVZ.

Our results suggest Olig2 expression and function in the adult forebrain is associated to the oligodendrocyte lineage compared with the spinal cord, in which both neuron and oligodendrocyte development depends on Olig2. By using light and electron microscopy, we confirmed that Olig2, a transcription factor that plays an essential role in the generation of oligodendrocytes during embryogenesis (Zhou et al., 2000; Lu et al., 2002), is expressed in a small subpopulation of type C cells (Aguirre et al., 2004; Hack et al., 2004, 2005). In the developing spinal cord of *Olig1/2*^{-/-} compound knock-out mutant mice, the absence of Olig function leads putative motor neuron and oligodendrocyte progenitors to instead generate V2 interneurons and astrocytes (Zhou and Anderson, 2002), demonstrating that Olig activity is essential for the regulation of a binary glial cell fate choice (Rowitch, 2004). Furthermore, Olig2 acts as a repressor of astroglial (Gabay et al., 2003; Fukuda et al., 2004) and neuronal (Hack et al., 2005) differentiation. Intriguingly, we observed the expression of Olig2 in some GFAP-expressing cells, suggesting that Olig2 may begin to be expressed very early in the lineage. It is unlikely that Olig2 function is required for the transition of all SVZ astrocytes to type C cells because no obvious defect in olfac-

pression of Olig2 in some GFAP-expressing cells, suggesting that Olig2 may begin to be expressed very early in the lineage. It is unlikely that Olig2 function is required for the transition of all SVZ astrocytes to type C cells because no obvious defect in olfac-

tory bulb neuron development is observed in 18.5 d postcoitum *Olig2*^{-/-} null animals (K. Ligon and D. Rowitch, unpublished observations). However, our data raise the possibility that *Olig2* expression is critical in a subset of type C cells that develop into oligodendrocytes. This conclusion is consistent with recent work overexpressing *Olig2* in SVZ precursors (Buffo et al., 2005; Hack et al., 2005; Marshall et al., 2005). Although these results suggest that *Olig2* expressing cells in the adult SVZ are progenitors of oligodendrocytes, we cannot exclude that these early progenitors could also generate other lineages. Note that it has recently been suggested that *Olig2* may also be involved in the differentiation of astrocytes (Marshall et al., 2005).

We identified a population of PSA-NCAM⁺/NG2⁺ cells in the CC that express PDGFR α and *Olig2* and have properties of immature migrating cells (Vaughn, 1969; Kakita and Goldman, 1999; Jennings et al., 2002). These young oligodendrocyte progenitors migrate individually or in small clusters but not in chains or ensheathed by glial tubes, as described for SVZ neuroblasts (Lois et al., 1996; Peretto et al., 1997). In contrast, migrating young oligodendrocytes are in intimate contact with myelinated axonal fibers. A similar close interaction between oligodendrocyte progenitors and axonal fibers has been shown in the optic nerve (Fulton et al., 1992). It has been suggested that young migrating oligodendrocytes originating in the neonatal SVZ disperse extensively in both radial and tangential planes (Suzuki and Goldman, 2003). This wide dispersion was not observed in our studies; oligodendrocytes generated in the adult SVZ were specifically found in the CC, the striatum, and fimbria fornix with little evidence of widespread rostrocaudal dispersion, suggesting that migration of oligodendrocyte progenitors born in the adult SVZ is limited to specific locations. A similar conclusion has been reached from developmental studies (Spassky et al., 2002). The affinity for axonal tracks within CC and striatum may explain the differences in dispersion observed between adults and neonates; these fibers may constrain tangential dispersion and explain the limited rostrocaudal distribution we observed.

Cells contained within the radial glia–astrocyte lineage serve as primary progenitors for new neurons in developing and adult brain (Alvarez-Buylla et al., 1990, 2001; Doetsch et al., 1999; Malatesta et al., 2000; Miyata et al., 2001; Noctor et al., 2001; Tamamaki et al., 2001; Merkle et al., 2005). Our results show that adult SVZ astrocytes generate oligodendrocytes. In primary cultures without the addition of growth factors or serum, single GFAP-expressing progenitors generate both oligodendrocytes and neurons. Only a small subpopulation of cells was multipotent in our assay (Fig. 6*J*). This could be attributable to the culture conditions we used, which may favor oligodendrocyte differentiation/survival, or could suggest that subpopulations of SVZ astrocytes are already committed for the generation of specific lineages. Our experimental protocol could also account for this result. For FACS isolation, we had to wait 36 h for accumulation of GFP in SVZ astrocytes after viral injections. By this time, many progenitors may have already moved along a particular lineage. Because our *in vitro* assay favors formation of differentiated clones, it could select for progenitors that are lineage restricted. Even under these stringent conditions, ~11% of the plated cells that survived gave rise to both neurons and oligodendrocytes.

This work highlights the central role that SVZ type B cells, which were taken as fully differentiated astrocytes for many years, play in the generation of not only neurons but also of oligodendrocytes. Our experiments suggest that some SVZ type B cells can generate both lineages *in vitro*, but it remains to be determined *in*

in vivo whether single SVZ type B cells are multipotent. Under normal conditions, SVZ type B cells produce few oligodendrocytes compared with the large number of neurons generated. Interestingly, however, oligodendrogenesis can be significantly upregulated after a demyelinating lesion. It will be important to determine the precise stage in the progression from type B to type C cells at which progenitors become committed to a neuronal or glial lineage and how these decisions are regulated in response to environmental signals. The identification of SVZ type B cells as progenitors for young migrating oligodendrocyte precursors, parenchymal OPCs, and oligodendrocytes will help understand the normal turnover of this essential glial cell in the adult brain.

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