Dynamic Contribution of Nestin-Expressing Stem Cells to Adult Neurogenesis

Diane C. Lagace, Mary C. Whitman, Michele A. Noonan, Jessica L. Ables, Nathan A. DeCarolis, Amy A. Arguello, Michael H. Donovan, Stephanie J. Fischer, Laure A. Farnbauch, Robert D. Beech, Ralph J. DiLeone, Charles A. Greer, Chitra D. Mandyam, and Amelia J. Eisch

Understanding the fate of adult-generated neurons and the mechanisms that influence them requires consistent labeling and tracking of large numbers of stem cells. We generated a nestin-CreERT2/R26R-yellow fluorescent protein (YFP) mouse to inducibly label nestin-expressing stem cells and their progeny in the adult subventricular zone (SVZ) and subgranular zone (SGZ). Several findings show that the estrogen ligand tamoxifen (TAM) specifically induced recombination in stem cells and their progeny in nestin-CreERT2/R26R-YFP mice: 97% of SGZ stem-like cells (GFAP/Sox2 with radial glial morphology) expressed YFP; YFP+ neurospheres could be generated in vitro after recombination in vivo, and maturing YFP+ progeny were increasingly evident in the olfactory bulb (OB) and dentate gyrus (DG) granule cell layer. Revealing an unexpected regional dissimilarity in adult neurogenesis, YFP+ cells accumulated up to 100 d after TAM in the OB, but in the SGZ, YFP+ cells reached a plateau 30 d after TAM. In addition, most SVZ and SGZ YFP+ cells became neurons, underscoring a link between nestin and neuronal fate. Finally, quantification of YFP+ cells in nestin-CreERT2/R26R-YFP mice allowed us to estimate, for example, that stem cells and their progeny contribute to no more than 1% of the adult DG granule cell layer. In addition to revealing the dynamic contribution of nestin-expressing stem cells to adult neurogenesis, this work highlights the utility of the nestin-CreERT2/R26R-YFP mouse for inducible gene ablation in stem cells and their progeny in vivo in the two major regions of adult neurogenesis.

Key words: dentate gyrus; subgranular zone; subventricular zone; rostral migratory stream; olfactory bulb; tamoxifen

Introduction

In the postnatal brain, evidence suggests progeny of glial fibrillary acidic protein (GFAP)- and nestin-expressing stem cells become neurons and astrocytes by progression through distinct stages (Doetsch et al., 1999; Garcia et al., 2004; Kempermann et al., 2004; Ganat et al., 2006). In the subventricular zone (SVZ) and hippocampal subgranular zone (SGZ), the stage at which progeny are restricted to a neuronal fate is unknown. In addition, it is unclear in the adult how many of these cells integrate into the existing neural network. Studies addressing these questions typically use thymidine analogs such as bromodeoxyuridine (BrdU) that target rapidly dividing cells or transgenic reporter mice that medially label discrete stages of adult neurogenesis. In general, these techniques are of limited use for stem cell analysis because their marking (Yamaguchi et al., 2000; Dayer et al., 2003). Although viral-mediated fluorescent labeling can tag stem cells and does not dilute, the necessity of intracranial injection and the restricted spread of virus make it nearly impossible for viruses to consistently label a large number of cells, particularly in the SGZ (van Praag et al., 2002). It is clear that questions of fate restriction and cellular contribution require a new approach to label and track large populations of stem cells and their progeny in the adult brain.

To address this challenge in studying neurogenesis, we and others created conditional and inducible nestin-driven transgenic mice (Beech et al., 2004; Yu et al., 2005; Carlen et al., 2006; Imayoshi et al., 2006; Kuo et al., 2006; Burns et al., 2007). Whereas many of these mouse models are highly effective in labeling and tracking the multipotent lineage of nestin-recombined cells in the embryo, only three of these models have demonstrated efficient labeling within the adult SVZ (Yu et al., 2005; Carlen et al., 2006; Burns et al., 2007), and none have labeled the large pool of adult SGZ stem and progenitor cells.

Here we report the first inducible nestin-CreERT2 mouse that can be used to label, track, and phenotype stem cells and their progeny in the adult SVZ and SGZ. Using this mouse, we explore whether adult-generated neurons are derived from the nestin lineage. In addition, we quantify the diverse composition of la-
beled cells over months following recombination, as well as estimate the total contribution of stem cells and their progeny to adult mice. Our data provide unique insight into the importance of stem cells to neurogenesis in the SVZ and SGZ and underscore the utility of this mouse in gene deletion from stem cells and their progeny in the adult brain.

Materials and Methods

Generation and genotyping of nestin-CreERT2/R26R-yellow fluorescent protein transgenic mice. Animal experiments were performed in accordance with the Guide for the Care and Use of Laboratory Animals and approved by the University of Texas Southwestern Animal Care and Use Committee. The CreERT2 sequence (Indra et al., 1999) (generous gift from P. Chambon, Institute for Genetics and Cellular and Molecular Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the Xh5 plasmid (generous gift from P. Chambon, Institute for Genetics and Cellular and Molecular Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.

Biology, Strasbourg, France) was cloned into the SalI sites of Nestin Xh5 plasmid (generous gift from W. Zhong, Yale University, New Haven, CT). The Xh5 plasmid has been published previously in other mouse models and contains similar elements to Nes/Plac2/3 introns (Zimmerman et al., 1994; Beech et al., 2004). The nestin-CreERT2 founder mice were generated by pronuclear injection of 5aM digest of nestin-CreERT2 into C57Bl/6J fertilized eggs. Five independent lines were transferred to 30% sucrose in 0.1M PBS. Brains were sectioned 30 or 40 μm thick on a freezing microtome in either the coronal or sagittal plane.
Recombination in stem cells and their progeny in the SVZ results in an increasing number of newly formed mature neurons in the RMS and OB

The time course of the appearance of recombinated (YFP+) cells in the SVZ, and subsequently in the OB GCL and glomerular cell layer (GL), revealed a dynamic addition of new cells in the adult (Fig. 2a–c). One day after TAM, nestin-lineage (YFP+) cells were evident in the SVZ, with few in the RMS and none in the OB (Fig. 2a). In agreement with recombination taking place in stem/progenitor cells, YFP+ cells were evident in the SVZ at all time points examined (Fig. 2a), and YFP+ neurospheres could be propagated in vitro from the SVZ of nestin-CreERT2/R26R-YFP mice given TAM in vivo (supplemental Fig. 1a–d, available at www.jneurosci.org as supplemental material). Both the OB GCL and OB GL showed an accumulation of YFP+ cells through 100 d after TAM (Fig. 2b,c) (GCL: F(4,24) = 11.9, p < 0.0001; GL: F(4,25) = 12.2, p < 0.005). The vast majority of YFP+ cells were in the OB GCL rather than in the GL (note difference in y-axis in Fig. 2, b vs c). Within the OB GCL, YFP+ cells were most dense in the deep internal portions of the GCL throughout the longitudinal axis of the OB, consistent with previous reports in adult mice using BrdU to assess OB neurogenesis (Lemasson et al., 2005; Mandairon et al., 2006). Our quantification allowed estimation of the contribution of YFP+ cells to adult OB. Because the adult mouse OB GCL has ~410,000 cells/mm³ (Parrish-Aungst et al., 2007), YFP+ cells represented 0.3, 1.1, and 2.6% of the total OB GCL density at 30, 65, and 100 d after TAM, respectively. In contrast, because each glomerulus is estimated to be surrounded by ~100 periglomerular cells [data not shown and Merson et al. (2006)], YFP+ cells represented 2.7, 3.1, and 4.9% of total periglomerular cell number at 30, 65, and 100 d after TAM, respectively.

Consistent with the significant increase in YFP+ cells over time, phenotypic analysis revealed increasing maturity of YFP+ cells in the RMS/OB. There was a significant difference in the percentage of neuroblasts recombinated [YFP+/DCX+] (Brown et al., 2003) at increasing times after TAM (Fig. 2d) (F(2,14) = 6.1; p < 0.05), as well as among the caudal RMS, RMS in OB, and OB GL (Fig. 2d) (F(2,14) = 28.3; p < 0.0001). Over time, in the caudal RMS, the percentage of YFP+ cells that were neuroblasts increased, whereas in the OB, the percentage of YFP+ cells that...
were neuroblasts decreased. This correlates with the significant increase in the percentage of YFP+ cells in the GCL that did not express DCX between 12 and 30 d or between 30 and 65 d (Fig. 2e) (phenotype by time interaction: \(F(4,18) = 70.8; p < 0.0001\)). These changes are expected because of the differentiation of YFP+ cells and their progeny. Indeed, >75% of YFP+ cells in the OB GCL expressed the mature neuronal marker NeuN at 65 and 100 d after TAM (Fig. 2f) (\(F(2,11) = 10.8; p < 0.005\)), in agreement with BrdU survival studies demonstrating ~80% of OB GCL BrdU+ cells label with NeuN (Mandairon et al., 2006). As expected (Lois and Alvarez-Buylla, 1994), many YFP+ OB GCL cells were also GABAergic (Fig. 2h), and a proportion were also CR+ (Fig. 2i).

Recombination in stem cells and their progeny in the SGZ results in an increasing number of newly formed mature neurons in the hippocampal GCL

At all time points after TAM, YFP+ cells were prominent within the dentate gyrus, with >90% of cells residing within the SGZ.
relative to the other dentate gyrus regions (Fig. 1d,e). There was a significant difference in the number of SGZ YFP+ cells at increasing times after TAM, with significantly more cells between 1 d and all other time points, as well as between 12 and 30 d (Fig. 3a) (F(4,42) = 15.5; p < 0.005). There was no significant change between 30–65 d and 65–100 d after TAM, indicating that the number of YFP+ cells reaches a plateau at 30 d. Because the adult mouse has ~1 million dentate gyrus GCL cells (Abusaad et al., 1999; Harburg et al., 2007), YFP+ cells are estimated to represent 1.0, 0.75, and 0.82% of the total dentate gyrus GCL cells 30, 65, and 100 d after TAM, respectively. The plateau in the SGZ is likely not attributable to inefficient labeling, because at 12 d after TAM, 97.1 ± 1.6% of stem-like SGZ cells were recombined. In this analysis, stem-like cells were identified by their immunoreactivity for both Sox2 (which labels astrocyte stem-like cells, rapidly dividing precursor cells, and mature astroglial cells) and GFAP (which labels astrocyte stem-like cells and mature astroglial cells) in combination with assessment for stem-like radial glial morphology using confocal analysis (Seri et al., 2001; Graham et al., 2003; Pevny and Rao, 2003; Komitova and Eriksson, 2004; Seri et al., 2004; Hattiangady and Shetty, 2007). Moreover, this plateau was not likely caused by TAM- or YFP-induced disruption of gross cellular function, because a neurogenic stimulus, such as free access to a running wheel, significantly increased the number of YFP+ SGZ cells (supplemental Fig. 2a–c, available at www.jneurosci.org as supplemental material) (bregma by treatment interaction: F(11,110) = 2.647; p < 0.01). This suggests that at least some of the recombined progenitor cells or their progeny can respond to a neurogenic stimulus. Considering mice were placed on the running wheels at a time point when YFP+ cells present diverse cellular phenotypes (Fig. 3c), the relatively modest effect of running in our data compared with others likely results from the specific sensitivity of type 2 cells to this neurogenic stimulus (Kronenberg et al., 2003).

As would be expected if labeling SGZ stem cells and their progeny, nestin-lineage YFP+ cells were morphologically heterogeneous after TAM (Figs. 1d,e, 3b,c). Using IHC, morphological analyses, and confocal microscopy, we classified YFP+ cells into nonexclusive phenotypic categories: stem-like (Sox2+/GFAP+), stem/progenitor (nestin+), dividing (Ki67+), immature neurons (DCX+), or postmitotic neurons (DCX+/CR+) (Fig. 3b,c) (Kempermann et al., 2004; Ming and Song, 2005; Hattiangady and Shetty, 2007). There was a significant difference in the proportion of recombined cells in these categories (F(4,60) = 53.18; p < 0.000) that changed after TAM (F(4,60) = 5.87; p < 0.005) (Fig. 3c). Between 1 and 12 d, YFP+ cells matured from being dividing, stem-like, and/or progenitor cells into immature or postmitotic neurons (Fig. 3b,c). In contrast, between 12–30 and 30–65 d, the proportion of YFP+ cells that expressed the different phenotypes remained constant (post hoc >0.05). Together, these data suggest that nestin drives expression in stem-like and progenitor cells in the SGZ. This is supported by the presence of stem-like recombined cells with radial glial morphology in the SGZ at all time points, including 100 d after TAM.

Over time, labeled stem cells and their progeny gave rise to mature dentate gyrus GCL neurons. At 30 d and beyond, the majority of the YFP+ SGZ cells displayed a long process extending up into molecular layer capped by a highly arborized dendritic tree (Fig. 3d). At 65 d and beyond, YFP+ fibers, presumably mossy fibers from YFP+ granule cells, densely innervated CA3. In addition, there was a significant increase in the proportion of recombined cells that expressed NeuN between 30 d and subsequent time points (Fig. 3f) (F(2,8) = 41.4; p < 0.0005). By 65 d after TAM, the percentage of YFP+ cells that expressed NeuN reached a plateau with ~50% of YFP+ cells being neurons at 65 and 100 d (Fig. 3f). YFP+ astrocytes were rare (YFP+/S100B+/GFAP+ and astrocytic morphology), providing additional support that nestin-expressing stem cells give rise to neurons, not...
astrocytes. There was also a significant increase in YFP+ cells that had a mature neuronal morphology in the outer GCL (oGCL) (Fig. 3g) ($F_{1,4,42} = 2.9; p < 0.05$). However, the cells in the oGCL at all time points represented <1% of all recombined cells, supporting that most maturing granule cells do not migrate from the SGZ into the oGCL (Kempermann et al., 2003). Together, these data underscore a link between nestin expression and neuronal fate.

**Discussion**

Multiple lines of evidence support our conclusion that TAM effectively drives recombination in nestin-expressing stem cells in the SVZ and SGZ during young adulthood. In nestin-CreER$^{T2}$/R26R-YFP mice, 96% of YFP+ cells in the SGZ were nestin+ 1 d after TAM, demonstrating that recombination occurs in nestin-expressing stem/progenitor cells. We also find at 12 d after TAM, 97% of stem-like cells (GFAP+/Sox2+/radial glial morphology) were recombined and that even at 100 d after TAM these cells were present. In addition, in vivo recombination led to YFP+ neurospheres in vitro and, as expected from labeled stem cells, maturing YFP+ progeny were increasingly evident after TAM. These data support that this mouse offers a potent tool for gene ablation studies in stem cells and their progeny.

The nestin-CreER$^{T2}$/R26R-YFP mouse allows the first quantifiable assessment of the long-term contribution of nestin-expressing stem cells to adult neurogenesis in both the OB and dentate gyrus. For example, in the SGZ, we estimate the contribution of stem cells is 1%, which is strikingly similar to previous estimates of the contribution of rapidly dividing cells (Doetsch and Hen, 2005) and validates that stem cells give rise to the rapidly dividing progenitor cells in the SGZ. In addition, we reveal intriguing dynamics of YFP+ accumulation in the OB versus SGZ. YFP+ cell density in the OB, which is dependent on the influx of YFP+ RMS neuroblasts and their surviving progeny, increased up to 100 d after TAM. We show the increase in density is attributable to the accumulation of maturing cells in the OB GCL and GL. This finding suggests the addition of new OB cells outpaces new cell turnover and might be expected because the volume of the murine OB increases in the first 2 years of life (Mirich et al., 2002). In contrast to the cell accumulation in the OB, the number of YFP+ cells in the SGZ reached a plateau 30 d after TAM. We were surprised to find a plateau in the number of YFP+ neurons, because we expected an increase based on BrdU-labeling survival studies (Dayer et al., 2003; Kempermann et al., 2003). What could explain the steady state of YFP+ cell number in the SGZ? One possibility is that the proportion of stem, precursor, and/or immature cells decreases with age (Rao et al., 2006; Hattiangady and Shetty, 2007). However, the presence of YFP+ radial glial cells 100 d after TAM and the steady state of the diverse population of recombined cells suggest that, over time, there is not an exhaustion of recombined stem cells. A second possibility is that analysis of longer times after TAM would reveal an eventual reduction in precursor number or activity, leading to a corresponding increase in the proportion of mature neurons. A third possibility is that the YFP transgene is silenced over time, leading to an underestimation of YFP+ cell number in the SGZ and producing the plateau seen 65–100 d after TAM. Although this is important to evaluate directly, this possibility is not supported by the increase in YFP+ cell number in the OB in our mouse, the similar steady-state dynamics seen in the SGZ of another transgenic mouse with an astrocyte-related gene driver of recombination (GLAST::CreER$^{T2}$) (Ninkovic et al., 2007), and the general robustness of genes in the rosa locus. These and other possibilities are important to consider in the ongoing and necessary evaluation of the hypothesis that stem cells and their progeny achieve homeostasis in the adult mouse SGZ.

In addition to revealing the distinct accumulation dynamics in the SVZ and SGZ, our data also address the role of nestin-expressing cells in adult neurogenesis: most progeny of nestin-expressing YFP+ cells are fated to become neurons. These data are in agreement with links between nestin expression and a specific lineage in many non-neuronal and oncogenic stem cells (Wiese et al., 2004). However, they are in contrast to the multilineage role for nestin in the embryo (Beech et al., 2004; Yu et al., 2005; Carlen et al., 2006; Imayoshi et al., 2006; Kuo et al., 2006; Burns et al., 2007), most notably in our nestin-CreER$^{T2}$/R26R-YFP mouse (Battiste et al., 2007). This underscores the importance of the permissive neurogenic microenvironment in determining the ultimate phenotypic fate of the progeny of nestin-expressing cells.

We show that nestin expression can ultimately result in neuronal, but not astrocytic, progeny. It is striking that inducible mouse models using astrocyte-related gene drivers [GFAP, GLAST (Garcia et al., 2006; Mori et al., 2006; Ninkovic et al., 2007)] drive expression in astrocytes throughout the brain, yet also give rise to neurons in the OB and SGZ. Interestingly, type 1 radial glial cells in the SGZ have recently been divided into categories of GFAP+/nestin− and GFAP+/nestin+ (Kempermann et al., 2004; Seki et al., 2007). Thus, the ability of our nestin-CreER$^{T2}$/R26R-YFP mouse to generate YFP+ neurons in the OB and SGZ urges research on factors that guide a GFAP+ or GLAST+ cell to become nestin+, to provide the much-needed information on neuronal fate restriction in the adult brain.

The nestin-CreER$^{T2}$ mouse will clearly be useful in gene deletion or progenitor ablation studies. Although elegant, viral-mediated Cre manipulation does not allow quantification of labeled cells along the longitudinal axis of the SGZ because of limited viral diffusion and often variability in iter between viral preparations (van Praag et al., 2002). Our nestin-CreER$^{T2}$ mouse is extremely consistent between litters, emphasizing its usefulness for gene ablation studies. Similarly, it can be used to inducibly drive cell death of neural stem cells in the adult brain, offering an alternative to the current progenitor ablation strategies (e.g., Garcia et al., 2004). Because the nestin-CreER$^{T2}$ mouse is inducible, it may also allow for future comparison of the present findings in young adulthood with neurogenesis in older mice, because the dynamics of neurogenesis may shift with age.

In conclusion, our data provide novel insights into the dynamic contribution of stem cells and their progeny to OB and dentate gyrus neurogenesis. The data strongly implicate that nestin expression is coincident with neuronal fate restriction. Furthermore, the quantification of the cellular and temporal specificity in the adult SVZ and SGZ validates this model as a valuable tool to guide future research identifying the contribution of stem cells and adult neurogenesis to neural circuitry and function.

**References**


Stereological estimation of the total number of neurons in the murine hippocampus using the optical dissector. J Comp Neurol 408:560–566.


Hattiangady B, Shetty AK (2007) Aging does not alter the number or phenotype of putative stem/progenitor cells in the neurogenic region of the hippocampus. Neurobiol Aging, in press.


