Neurons arise in the basal neuroepithelium of the early mammalian telencephalon: A major site of neurogenesis

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Communicated by Kai Simons, Max Planck Institute of Molecular Cell Biology and Genetics, Dresden, Germany, December 23, 2003 (received for review December 15, 2003)

Neurons of the mammalian CNS are thought to originate from progenitors dividing at the apical surface of the neuroepithelium. Here we use mouse embryos expressing GFP from the Tis21 locus, a gene expressed throughout the neural tube in most, if not all, neuron-generating progenitors, to specifically reveal the cell divisions that produce CNS neurons. In addition to the apical, asymmetric divisions of neuroepithelial (NE) cells that generate another NE cell and a neuron, we find, from the onset of neurogenesis, a second population of progenitors that divide in the basal region of the neuroepithelium and generate two neurons. Basal progenitors are most frequent in the telencephalon, where they outnumber the apically dividing neuron-generating NE cells. Our observations reconcile previous data on the origin and lineage of CNS neurons and show that basal, rather than apical, progenitors are the major source of the neurons of the mammalian neocortex.

All neurons generated during the development of the mammalian CNS derive from neuroepithelial (NE) cells (see Supporting Text, which is published as supporting information on the PNAS web site). NE cells undergo three principal kinds of cell division, (i) symmetric, proliferative divisions (two NE cells) (see Supporting Text for terminology), (ii) asymmetric divisions (one NE cell, one neuron) and (iii) symmetric, differentiating divisions (two neurons) (1–6). It remains to be settled whether, at the very onset of neurogenesis, NE cells switch only to asymmetric division, or to both asymmetric and symmetric differentiating divisions.

Real-time imaging studies of cells dividing at the apical surface of the neuroepithelium (7–9) and analyses of intrinsic cell fate determinants with a polarized distribution in mitotic NE cells (7, 10, 11) have provided evidence in support of the widely held view that during early neurogenesis, neurons are generated from NE cells by asymmetric rather than symmetric differentiating divisions (2, 6, 12–14). In contrast, consistent with retroviral lineage tracing studies in vivo (5, 15–18), analysis of the division patterns of isolated NE cells and their progeny has documented the coexistence, in vitro, of asymmetric and symmetric neuron-generating divisions of progenitors at the onset of neurogenesis (19–21). One possible explanation for this apparent discrepancy is the existence of neuronal progenitors other than the canonical NE cell dividing at the apical surface of the neuroepithelium.

One of the problems in studying the divisions of NE cells that generate neurons has been the lack of a marker that allows one to distinguish between proliferating and neuron-generating NE cells and to identify the latter before they enter mitosis. Our group previously reported the first such marker, Tis21, an antiproliferative gene that within the neural tube is selectively expressed in neuroepithelial (NE) cells (22). Moreover, given that >95% of the newborn CNS neurons appear to inherit Tis21 protein from their progenitors (22), Tis21 is presumably expressed in the vast majority, if not all, of neuronal progenitors, an important aspect in view of the possible existence of as yet uncharacterized subpopulations of such cells. Here, we have generated a Tis21-GFP knock-in mouse line and, by using time-lapse multiphoton microscopy, we have analyzed neuron-generating divisions in the embryonic neural tube at the onset of neurogenesis.

Methods

Generation of Tis21-GFP Knock-In Mice. For details about the generation of Tis21-GFP knock-in mice (Tis21+ /m2(Gfp)Whb), see Supporting Text. All data shown in this study are from heterozygous embryos obtained by crossing homozygous males to C57BL/6J females. The day of the vaginal plug was defined as embryonic day (E) 0.5.

Fluorescence Microscopy on Whole-Mounts and Cryosections. Fluorescence microscopy on whole-mounts and cryosections was performed according to standard procedures. Details are given in Supporting Text.

Definitions and Quantitations. “Apical mitoses” were defined as those occurring at the apical (ventricular) surface of the neuroepithelium. “Subapical mitoses” were defined as nonsurface mitoses that were at most three nuclear diameters away from the apical surface of the neuroepithelium. “Basal mitoses” were defined as those occurring in the basal one-third of the ventricular zone (VZ), which is referred to as its “basal portion” or “basal region.”

The abundance of GFP-containing cells in the VZ (Fig. 1 j) was determined by counting the nuclei showing GFP fluorescence and expressing these as percentage of 4’,6-diamidino-2-phenylindole (DAPI)-stained nuclei. For the quantitation of mitotic cells in the VZ and (when present) subventricular zone (SVZ) (Fig. 2 j–m), mitotic cells were identified by phosphohistone H3 staining, corroborated by DAPI staining, and the proportion of mitotic cells showing GFP fluorescence was determined. Basal mitotic GFP-positive cells were also analyzed for βIII-tubulin immunoreactivity.

Multiphoton Time-Lapse Video Microscopy. Brain slices (250–400 μm) from E9.5–E12.5 heterozygous Tis21-GFP knock-in mouse embryos were prepared following established methods (23) and cultured in a POC-Chamber system (Saur, Reutlingen, Germany) in Neurobasal medium (Invitrogen) supplemented with 10% immediately centrifuged mouse serum (Harlan Breeding Laboratories), 1× N2 (Invitrogen), 1× B27 supplement (Invitrogen) and 100 units/ml penicillin/streptomycin. The POC-Chamber system was gassed with 20% O2/5% CO2/75% N2. For details, see Supporting Text.

Time-lapse analysis was usually started ~2 h after the beginning of the culture, after visual verification of the integrity of the slice.

Abbreviations: NE, neuroepithelial; En, embryonic day n; VZ, ventricular zone; SVZ, subventricular zone.

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Images were recorded on an inverted Radiance 2100 multiphoton microscope (Bio-Rad) equipped with Nikon optics (ECLIPSE TE300) and a Mira 900 titanium/sapphire laser (Coherent) tuned to 900 nm, which was pumped by a Verdi 5 W solid-state laser (Coherent). A ×30 water immersion lens, which was heated to 37°C by using an objective heater (Bioptechs) and constantly supplied with immersion water by using a pump, was used to record the GFP fluorescence. Every 5–10 min, 100-μm z stacks, with the individual optical sections being spaced at 3- or 5-μm intervals, were recorded by using a single scanning run at 256 × 256 pixel resolution with a scanning speed of 133–500 lines per second. Images were acquired by using the Bio-Rad software. Three-dimensional reconstructions were computed by using VOL eyCOntly software (Improvision) in either the high-resolution (HR) rendering or 3D-rendering mode (see figure legends).

The amount of GFP fluorescence and nuclear size were quantified from the raw data (Bio-Rad) by using NIH 1.62 software (National Institutes of Health). For details, see Supporting Text.

Results

A Tis21-GFP Knock-In Mouse Reveals Neuron-Generating Progenitors in the Embryonic CNS. We first established a Tis21-GFP knock-in mouse line in which the protein-encoding portion of exon 1 of the Tis21 gene was replaced by GFP carrying a nuclear localization signal (see Fig. 5, which is published as supporting information on the PNAS web site). The appearance of GFP fluorescence in the developing CNS of E9.5–E14.5 heterozygous Tis21-GFP knock-in mouse embryos (Fig. 1 a–d) completely matched the known expression of Tis21 mRNA (22, 24) and correlated precisely with the temporal and spatial gradients of fluorescence. This was corroborated by analysis of GFP fluorescence (the pattern of which was essentially identical to that of GFP immunoreactivity; data not shown) and βIII-tubulin immunoreactivity, a marker of young neurons (25), in sections of the developing CNS. We studied the gradients of neurogenesis by examining, at a magnification allowing an overview of the neural tube wall, either one defined brain region (telencephalon) over time (E9.5–E15.5) (Fig. 1 e–j) or various regions of the CNS along the rostro-caudal axis (telencephalon–spinal cord) at one defined developmental time point (E10.5) (see Fig. 6, which is published as supporting information on the PNAS web site), with essentially the same principal results.

No Tis21-driven GFP expression was detected in the neuroepithelium before the onset of neurogenesis (Fig. 1e). At the onset of neurogenesis (Fig. 1f), a few cells in the VZ (Fig. 1j, E10.5), which lacked βIII-tubulin immunoreactivity and were therefore NE cells, started to express GFP. The fluorescence was confined to the nucleus, as expected from the presence of a nuclear localization signal. Consistent with the notion that Tis21-expressing NE cells generate neurons (22), virtually all of the newborn neurons (as identified by the presence of βIII-tubulin immunoreactivity) in the VZ (Fig. 2h, arrowhead), and still most of the slightly older neurons (“young neurons”) in the adjacent neuron-containing layer (Fig. 1f, asterisk, and Fig. 2d, solid white arrowhead), showed GFP fluorescence (see Table 1, which is published as supporting information on the PNAS web site). Newborn and young neurons are known to lack Tis21 gene expression (22). The presence of GFP fluorescence in these neurons therefore reflected the inheritance, by default, of the GFP protein from NE progenitors exhibiting Tis21-driven GFP expression (see Fig. 4e). This, in turn, implies that most, if not all, neurons are derived from Tis21-GFP-expressing progenitors.

Concomitant with the progress in neurogenesis, the proportion of GFP-expressing NE cells increased (Fig. 1 f–i), with 60% of all cells of the E15.5 telencephalic VZ expressing GFP (Fig. 1j). The GFP positive nuclei appeared to be more abundant and to show brighter fluorescence in the basal than in the apical portion of the VZ. This reflects the fact that (i) the Tis21 gene is transiently expressed in the G1-phase of the cell cycle (22) and, hence, GFP fluorescence peaks when neuron-generating progenitors are in S phase, and (ii) the nuclei of a substantial portion of the neuron-generating progenitors do not migrate to the apical surface in G2 but remain in the basal region of the VZ, as will be described below.

At these later stages of neurogenesis, only few cells in the neuronal layers still showed GFP fluorescence, and there was a clear apical-to-basal decline in both frequency and intensity of these cells (Fig. 1, compare b and i).

Based on the findings that (i) the onset and progression of Tis21-driven GFP expression in NE cells correlates precisely with the onset and progression of neurogenesis and (ii) most, if not all neurons, are derived from Tis21-GFP-expressing progenitors, and in view of the time-lapse observations described below, we conclude that, during early neurogenesis in Tis21-GFP knock-in mouse embryos, divisions of Tis21-GFP-expressing cells in the neuroepithelium generate neurons.

A Mitotic Neuron-Generating Progenitor at the Basal Side of the VZ. We therefore focused on the mitotic cells in the neuroepithelium showing Tis21-driven GFP expression. In the E10.5 hindbrain, consistent with some interphase NE cells expressing GFP (Fig. 2 h, arrow), some of the mitotic cells at the apical surface of the neuroepithelium showed GFP fluorescence (Fig. 2 e–g, open arrows). More apical mitotic cells became GFP-positive as neurogenesis progressed (Fig. 2m, curve A).

Surprisingly, however, even at the onset of neurogenesis, GFP-positive mitotic cells were observed not only at the apical surface of the neuroepithelium, but also in the basal region of the VZ, as illustrated for the E10.5 telencephalon (Fig. 2 a–d, open arrows with asterisk). Like the apical GFP-positive mitotic NE cells, the GFP-positive mitotic cells in the basal portion of the neuroepithelium were undetectable before the onset of neurogenesis. As neurogenesis in the telencephalon progressed, the basal GFP-positive mitotic cells (Fig. 2l, curve B) increased more rapidly in number than the apical ones and outnumbered the latter (Fig. 2l, curve A). The time course of the sum of apical plus basal GFP-positive mitotic cells (Fig. 2l, curve A+B) was consistent with that of the total (mitotic plus interphase) GFP-positive cells in the telencephalic neuroepithelium (Fig. 1j).

Along the rostro-caudal axis of the neural tube, basal GFP-positive mitotic cells were most frequently observed in the regions that are known to form a SVZ at later stages of neurogenesis (26), i.e., the telencephalon (Fig. 2, compare l and m, curves B). Indeed, the basal GFP-positive mitotic cells were found to be abundant in the SVZ of the E13.5 telencephalon (Fig. 3f, filled white arrowheads).

In the telencephalon, almost all (~90%) of the basal mitotic cells observed at the onset of neurogenesis were found to express GFP (Fig. 2j, E10.5), and the same was the case once a SVZ had formed (Fig. 2j, E13.5). The vast majority (~90%) of the basal mitotic GFP-positive cells did not show βIII-tubulin immunoreactivity (Fig. 2d, open arrow with asterisk), which was detected in only 2 of 31 cells analyzed.

We conclude that, from the onset of neurogenesis in the mouse embryonic telencephalon, neurogenic progenitors dividing at the basal side of the VZ coexist with those dividing at the apical side, and eventually outnumber the latter.

Apical Neuron-Generating Progenitors Undergo Asymmetric Division. Next, we studied, during early stages of neurogenesis, the behavior of the GFP-expressing cells dividing at the apical and the basal side of the VZ, and also that of the resulting daughter cells (see Supporting Text for subapical divisions, and Figs. 7d and 8, which are published as supporting information on the PNAS web site). For this purpose, we used time-lapse multiphoton laser scanning microscopy (27) of organotypic slice cultures of E12.5 telencephalon, an established experimental system (28). In the
case of the apically dividing cells, the GFP-labeled nucleus of the mother cell migrated from the basal to the apical side of the neuroepithelium, followed by cell division and migration of the daughter cell nuclei in the basal direction (Figs. 3a and 7a–c, and Movie 1 and Table 2, which are published as supporting information on the PNAS web site). In virtually all cases, the daughter cells behaved differently from each other in several respects.

First, the basally directed migration of the daughter cell nuclei was distinct, with one nucleus (referred to as the leading nucleus, Fig. 3a–d arrowheads) moving ahead of the other (referred to as the trailing nucleus, Fig. 3a arrow; see also Fig. 9, which is published as supporting information on the PNAS web site). Second, the destination of the daughter cells originating from apical divisions differed. Whenever the leading nucleus could be followed for a sufficiently long time, it left the VZ (Fig. 3a, 380 min arrowhead) and apparently entered the adjacent layer containing young neurons (Fig. 7a–c arrowheads). In contrast, the trailing nucleus eventually returned to the apical surface and underwent another mitosis (Fig. 3b arrows). Third, the daughter cells originating from apical divisions were different with regard to the intensity of GFP fluorescence over time. Relative to the leading daughter cell nucleus, the trailing nucleus consistently showed an at least 2-fold increase in GFP fluorescence within the variation of the individual values from the mean.

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first 4–6 h after mitosis (Fig. 4 a and c, apical). Fourth, the increase in GFP fluorescence in the trailing daughter cell nucleus was followed by an increase in the apparent nuclear size as compared to the leading daughter cell nucleus (Fig. 4d, left columns).

We conclude that the apical divisions of Tis21-GFP-expressing NE cells in the telencephalon are asymmetric. One daughter cell becomes a neuron, leaves the VZ, and loses its GFP fluorescence. The other daughter cell remains a neuron-generating NE cell, stays in the VZ, reinduces Tis21-GFP expression, undergoes interkinetic nuclear migration, and divides again at the apical surface.

**Basal Neuron-Generating Progenitors Undergo Symmetric Division.** Unlike daughter cells from apical divisions, the daughter cells from basal divisions behaved similarly to each other. First, in essentially all cases tracked for a sufficiently long time (Fig. 3 c and d, and Movies 2 and 3, which are published as supporting information on the PNAS web site), the sister cells shared the same destination. Both daughter cells, as judged from their GFP-labeled nuclei, migrated basally, left the VZ and entered the adjacent layer containing young neurons (Fig. 3 c 400–510 and d 500–620 min). Second, the basally directed migration of the daughter cell nuclei appeared to occur with similar speed.
GFP (this study) by newborn neurons. P, NE cells undergoing proliferative neuronal progenitors and the passive inheritance of the TIS21 protein (22) and indicate multiple rounds of asymmetric division.

The area of the GFP-stained nucleus over time was set to 100 (black lines), and the total GFP fluorescence in the progenitor nucleus was expressed relative to this (colored curves). (a and b) The curves show the moving average of two consecutive time points each. The curves showing a consistent increase in relative GFP fluorescence (a) correspond, in each case, to the trailing daughter cell that remained in the basal region (b) of the VZ. For each of the time points (10-min intervals), the total GFP fluorescence in the daughter cell with a lesser integral of total GFP fluorescence over time was used to calculate the mean values for (i) the initial GFP fluorescence, (ii) the lag phase before the GFP fluorescence increase, and (iii) the GFP fluorescence increase over time by using linear regression analysis (red line). Similarly, the relative GFP fluorescence data of the three curves shown in a were used to calculate the mean value for the GFP fluorescence over time by using linear regression analysis (blue line). (d) Increase in the size of the nucleus. The area of the GFP-stained nucleus over time was determined in the daughter cells originating from divisions at the apical surface (left columns) and in the basal region of the VZ (right columns). In two of three cases of apical divisions, the area of the nucleus of one of the daughter cells increased, relative to that of the other daughter cell, during the length of the time-lapse observation (left open column, relative area initially after mitosis; left filled column, relative area at the end of the time-lapse). Note that, in each case, this increase in nuclear size occurred in the daughter cell that also showed an increase in relative GFP fluorescence, migrated more slowly in the basal direction, and remained in the VZ. No such size increase was observed for the daughter cells originating from basal divisions (right columns). Data are the mean of two (left) and three (right) observations; bars indicate the variation of the individual values from the mean and the SD, respectively. (e) Scheme summarizing the expression of the TIS21 gene in neuronal progenitors and the passive inheritance of the TIS21 protein (22) and GFP (this study) by newborn neurons. P, NE cells undergoing proliferative division; NG, neuronal progenitors undergoing neuron-generating division; N, newborn neuron; arrows indicate lineage relationship. (f) Putative lineages of basal versus apical Tis21-GFP-expressing neuronal progenitors (green), NE, NE cell; AP, apical progenitor; BP, basal progenitor; N, neuron. Dashed lines indicate multiple rounds of asymmetric division.

Fig. 4. Apical vs. basal neuronal progenitors and their lineages: analysis of Tis21-driven GFP expression and nuclear size of daughter cells. (a–c) Time course of GFP fluorescence. Total GFP fluorescence was determined in the daughter cells originating from divisions at the apical surface (a) and in the basal region (b) of the VZ. For each of the time points (10-min intervals), the total GFP fluorescence in the daughter cell with a lesser integral of total GFP fluorescence over time was set to 100 (black lines), and the total GFP fluorescence in the progenitor nucleus was expressed relative to this (colored curves). (a and b) The curves show the moving average of two consecutive time points each. The curves showing a consistent increase in relative GFP fluorescence (a) correspond, in each case, to the trailing daughter cell that remained in the basal region (b) of the VZ. (c) The relative GFP fluorescence data of the three curves shown in a were used to calculate the mean values for (i) the initial GFP fluorescence, (ii) the lag phase before the GFP fluorescence increase, and (iii) the GFP fluorescence increase over time by using linear regression analysis (red line). Similarly, the relative GFP fluorescence data of the three curves shown in b, which did not show a consistent increase, were used to calculate the mean value for the GFP fluorescence over time by using linear regression analysis (blue line). (d) Increase in the size of the nucleus. The area of the GFP-stained nucleus over time was determined in the daughter cells originating from divisions at the apical surface (left columns) and in the basal region of the VZ (right columns). In two of three cases of apical divisions, the area of the nucleus of one of the daughter cells increased, relative to that of the other daughter cell, during the length of the time-lapse observation (left open column, relative area initially after mitosis; left filled column, relative area at the end of the time-lapse). Note that, in each case, this increase in nuclear size occurred in the daughter cell that also showed an increase in relative GFP fluorescence, migrated more slowly in the basal direction, and remained in the VZ. No such size increase was observed for the daughter cells originating from basal divisions (right columns). Data are the mean of two (left) and three (right) observations; bars indicate the variation of the individual values from the mean and the SD, respectively. (e) Scheme summarizing the expression of the TIS21 gene in neuronal progenitors and the passive inheritance of the TIS21 protein (22) and GFP (this study) by newborn neurons. P, NE cells undergoing proliferative division; NG, neuronal progenitors undergoing neuron-generating division; N, newborn neuron; arrows indicate lineage relationship. (f) Putative lineages of basal versus apical Tis21-GFP-expressing neuronal progenitors (green), NE, NE cell; AP, apical progenitor; BP, basal progenitor; N, neuron. Dashed lines indicate multiple rounds of asymmetric division.

(Table 2). Third and fourth, there was no significant difference between the daughter cells with regard to the intensity of GFP fluorescence over time (Fig. 4 b and e, basal) and their apparent nuclear size (Fig. 4 d, right columns). We conclude that the divisions of Tis21-GFP-expressing progenitors in the basal region of the VZ of the telencephalon (see also Fig. 7 e and f) are symmetric, with both daughters leaving the VZ and becoming neurons.

Discussion

We have taken a comprehensive approach to analyze the generation of neurons in the mammalian CNS. By specifically labeling the vast majority, if not all, of the neuron-generating progenitors within the neural tube in Tis21-GFP knock-in mouse embryos, we found that, at the onset of neurogenesis, neurons are generated by two distinct types of divisions: (i) apical asymmetric divisions of NE cells, which give rise to one neuron and one NE cell that apparently undergoes another neuron-generating division, and (ii) symmetric divisions of progenitors at the basal side of the VZ, which give rise to two neurons (Fig. 4 f).

Two lines of evidence indicate that, in the Tis21-GFP knock-in mouse embryos, we specifically labeled most, if not all, neuron-generating progenitors in the neural tube. The first line of evidence concerns the use of the Tis21 gene to drive GFP expression (Fig. 4 e). GFP fluorescence was found to appear throughout the neural tube in exact correlation with the temporal and spatial gradients of neurogenesis, consistent with the findings that the Tis21 mRNA is specifically expressed in neuron-generating, but not proliferating, NE cells irrespective of any regional patterning (22, 24). The onset of GFP expression was confined to the mitotically active layers, i.e., the VZ and, once present, in addition the SVZ, where fluorescence became detectable in an increasing proportion of cells as more and more neurons were generated during development. GFP fluorescence was also observed in newborn neurons, in which it appeared to decrease with their age. This is in agreement with the previous findings that the Tis21 gene is solely transcribed in neuronal progenitors but not in neurons (22, 24), and with the notion that GFP synthesized in neuronal progenitors will be passively inherited by their daughter cells, i.e., the neurons (Fig. 4 e). The observation that virtually all newborn neurons are GFP-positive (Table 1) therefore implies that most, if not all, neurons must be derived from GFP-expressing progenitors, i.e., the vast majority of neuronal progenitors transcribe the Tis21 gene and hence produce GFP. In other words, Tis21 expression is a “pan-neurogenic” marker.

The second line of evidence concerns how the progeny of GFP-expressing progenitors behave. Whenever the daughter cells of GFP-positive progenitors could be observed for a sufficiently long time, one or both of them left the VZ. Given that the only other cell type produced by NE cells at this early stage of CNS development are neurons (20, 29, 30), which characteristically leave the VZ, it follows that most, if not all, of the GFP-labeled progenitors are neuron-generating. The present approach of exploiting the pan-neurogenic nature of Tis21 gene expression to reveal, specifically and throughout the neural tube, neuron-generating cell divisions explains why a population of basal neuronal progenitors in addition to the canonical apically dividing NE cells was uncovered.

Analysis of the progeny of neuron-generating divisions of NE cells at the apical surface of the VZ (see Supporting Text for relationship to radial glial cells) revealed that, at the onset of neurogenesis, these divisions are asymmetric with regard to daughter cell fate (Fig. 4 f and Fig. 10 d, which is published as supporting information on the PNAS web site). The present demonstration that, at the onset of neurogenesis, the neuron-generating divisions of NE cells at the apical side of the VZ are asymmetric is fully in line with the results of previous studies performed at later stages of neurogenesis (8, 9) and supports the canonical view that neurons arise from asymmetric divisions of NE/radial glial cells at the apical side of the VZ.

In the light of this, the finding of neuron-generating progenitors dividing in the basal part of the VZ (rather than at its apical side) was unexpected. In contrast to the mitotic NE cells at the apical side of the VZ, of which only a minor fraction (~20%) expressed GFP, i.e., turned into neuron-generating progenitors,
nearly all (90%) of the basal mitotic cells were GFP-positive, and a small fraction (<10%) already expressed the neuronal marker βIII-tubulin. The basally dividing neuron-generating progenitors occurred primarily in the neocortex and coexisted with apically dividing neuron-generating NE cells from the onset of neurogenesis. Time-lapse analysis revealed that in contrast to the asymmetric neuronal-generating divisions at the apical side of the VZ, both daughter cells originating from basal neuron-generating divisions migrated out of the VZ and did not show a differential increase in nuclear volume or reinduction of GFP in one of the daughters, consistent with both of them becoming neurons. Hence, in mammals there exists, in particular in the neocortex and from the onset of neurogenesis, a second, hitherto uncharacterized, type of neuronal progenitor, which generates neurons by symmetric divisions at the basal side of the VZ (Figs. 4f and 10c).

These results contribute to resolving the issue as to the origin of CNS neurons. Retroviral lineage studies (5, 15, 16, 18) and clonal analysis of isolated neuronal progenitors (19–21) have shown that neurons can be generated from both symmetric and asymmetric divisions. However, only asymmetric divisions have previously been reported from time-lapse analyses of neuronal-generating divisions of NE/radial glial cells occurring at the apical side of the VZ (7–9). The present description of a second, symmetrically dividing, type of neuronal progenitor in the basal part of the VZ reconciles those seemingly inconsistent previous reports with each other.

Mitotic cells in the basal part of the VZ have been noticed previously, but their role has been unclear. On the one hand, the observed increase in nonapical mitotic cells concomitant with the progression of neurogenesis has led to the suggestion that these cells could contribute to neurogenesis (31, 32). On the other hand, basal mitotic cells may be the progenitors of the SVZ, which constitutes a distinct cell layer at later stages of neurogenesis and which is thought to largely generate glia (26, 29). However, the recent observations that dividing SVZ cells and cortical neurons both express certain genes (33–35) and that mutants affecting SVZ development also affect the generation of upper-layer cortical neurons (34) have been interpreted in favor of a role of dividing SVZ cells also in neurogenesis. This is fully consistent with the direct evidence presented here, showing that the majority of cortical neurons originate from basal progenitors of the VZ and, later, the SVZ.

We believe that (at least some of) the basal Tis21-GFP-expressing neuronal progenitors originate from apically dividing Tis21-GFP-negative NE cells (Fig. 4f). First, the time course of appearance of apical versus basal Tis21-GFP-positive progenitors was distinct, with the basal progenitors being slightly delayed compared to the apical ones. Second, at later stages of neurogenesis in the telencephalon, Tis21-GFP-expressing mitotic basal progenitors outnumber apical ones, and hence at least some of the basal progenitors must initiate GFP expression de novo. In our model, the basal progenitor (Fig. 4f) constitutes an additional mitotic cell intermediate that, like the postmitotic neuron arising from apical progenitors (Fig. 4f), originates from an asymmetric, albeit Tis21-GFP-negative, division.

The basal neuronal progenitor described here has significant implications for mammalian CNS development and, perhaps, evolution. With the progression of neurogenesis in the telenencephalon, the basal neuronal progenitors outnumber the apical ones and constitute the predominant population of neuronal progenitors. Considering that basal progenitors will generate twice as many neurons as apical progenitors (Fig. 4f), the basal progenitors appear to be the major source of cortical neurons. Moreover, by constituting an additional mitotic zone, basal neuronal progenitors provide a means of increasing the number of neurons that can be generated per luminal surface area of neuroepithelium per unit time (Fig. 4f). In this context, it is important to note that basal neuronal progenitors predominate in the region of the neural tube where the greatest number of neurons is generated, the telenencephalon. Interestingly, there has been a substantial increase in the ratio of SVZ to VZ during evolution (34, 36). This most likely reflects the ratio of basal to apical neuronal progenitors as the neocortex enlarged. Accordingly, in humans, progenitors in the SVZ have recently been implicated in neocortical neurogenesis (35).

We thank Drs. E. Fuchs and H. Monyer for support with the ES cell work, Dr. J.-P. Rouault for the Tis21 genomic clone, P. Veh for excellent technical assistance, and Drs. F. Calegari, V. Dubreuil, Y. Kosodo, and especially M. Meredith-Stewart for their helpful comments on the manuscript. W.B.H. was supported by Deutsche Forschungsgemeinschaft Grants SPP 1109, Hu 275/7, SPP 1111, and Hu 275/8 and the Fonds der Chemischen Industrie.