

## RESEARCH COMMUNICATION

# The T-box transcription factor Eomes/Tbr2 regulates neurogenesis in the cortical subventricular zone

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**The embryonic subventricular zone (SVZ) is a critical site for generating cortical projection neurons; however, molecular mechanisms regulating neurogenesis specifically in the SVZ are largely unknown. The transcription factor Eomes/Tbr2 is transiently expressed in cortical SVZ progenitor cells. Here we demonstrate that conditional inactivation of Tbr2 during early brain development causes microcephaly and severe behavioral deficits. In Tbr2 mutants the number of SVZ progenitor cells is reduced and the differentiation of upper cortical layer neurons is disturbed. Neurogenesis in the adult dentate gyrus but not the subependymal zone is abolished. These studies establish Tbr2 as a key regulator of neurogenesis in the SVZ.**

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In mammals highly diverse populations of neurons are organized into the complex structure of the six-layered neocortex. Projection neurons populating each layer are generated sequentially in proliferative zones adjacent to the ventricle in an “inside-out” fashion (Rakic 1974). In early neurogenesis, radial glial cells (RG) in the ventricular zone (VZ) generate mainly lower-layer pyramidal cortical neurons through asymmetric divisions directly at the apical surface of the ventricle. As neurogenesis proceeds, RGs additionally generate intermediate progenitor cells (IPCs) that undergo symmetrical divisions predominantly in the subventricular zone (SVZ) (Haubensak et al. 2004; Noctor et al. 2004), giving rise to neurons

destined for upper cortical layers (Wu et al. 2005). Hence, the early embryonic SVZ contains at least two distinct neural cell populations: migrating neurons and IPCs (Tarabykin et al. 2001). The substantial enlargement of the SVZ and upper cortical layers in mammalian phylogeny suggests that IPC-derived neurons have markedly contributed to the evolutionary expansion of the cortex (Smart et al. 2002; Kriegstein et al. 2006). This view is supported by the absence of SVZ mitoses in the developing reptilian and avian dorsal cortex, which lack equivalent cortical layers II–IV (Cheung et al. 2007). The molecular mechanisms that specifically regulate proliferation and differentiation of IPCs are largely unknown. Few genes display predominant SVZ expression, notably subventricular-expressed transcript 1 (Svet1) (Tarabykin et al. 2001), the transcription factors cut-like 2 (Cux2) (Zimmer et al. 2004), and the T-box transcription factor Tbr2 (Englund et al. 2005). Expression of Svet1 and Cux2 appears to persist in IPC-derived upper cortical layer neurons, suggesting they represent SVZ markers for upper-layer progenitors (Molyneaux et al. 2007). This is consistent with recent studies demonstrating that Cux2 knockout mice display increased IPC proliferation and upper cortical layer thickness (Cubelos et al. 2007).

In contrast, Tbr2 is only transiently expressed during SVZ neurogenesis. Tbr2 expression initiates between embryonic day 10 (E10) and E12 in a few differentiated preplate neurons including Cajal-Retzius cells. Highest Tbr2 expression is detected between E12 and E16 in IPCs coincident with the peak of murine cortical neurogenesis and decays after E17 (Englund et al. 2005; Molyneaux et al. 2007). In humans a homozygous chromosomal translocation that disrupts Tbr2 expression is associated with microcephaly, polymicrogyria, corpus callosum (CC) agenesis, cognitive deficits, and severe motor delay with hypotonia (Baala et al. 2007). In mice Tbr2 is required for development of the trophoblast and during germ layer formation, leading to embryonic death before onset of neurogenesis (Russ et al. 2000; Arnold et al. 2008). To study Tbr2 function during brain development we conditionally inactivated Tbr2 in the CNS.

## Results and Discussion

Animals lacking Tbr2 expression in the CNS were generated by crossing Tbr2/Eomes<sup>CA/+</sup>;Sox1 Cre to Tbr2/Eomes<sup>CA/CA</sup> mice (Arnold et al. 2008). Sox1 Cre mice (Takashima et al. 2007) strongly express Cre throughout the neural tube from E9.5 before the onset of Tbr2 expression in the CNS at E10.5 (Bulfone et al. 1999). As a control, Sox1 Cre mice crossed to the R26R reporter strain activate LacZ expression throughout the neural tube at E10.5 (Supplemental Fig. 1A). Excision of the conditional allele in E13.5 brains was confirmed by RT-PCR and Western blots (Supplemental Fig. 1B,C).

Mutant animals (Tbr2/Eomes<sup>CA/CA</sup>;Sox1 Cre) were viable and fertile but displayed severe behavioral abnormalities. Mutants were readily identifiable due to their enhanced aggressiveness upon handling. Breeding of mutant females ( $n = 4$ , eight litters) to wild-type males and vice versa ( $n = 2$ , four litters) gave viable offspring; however, pups were invariably subject to infanticide at birth. If mutant males were removed, offspring survived

[Keywords: Eomesodermin; Tbr2; neuronal progenitor cells; SVZ; brain size]

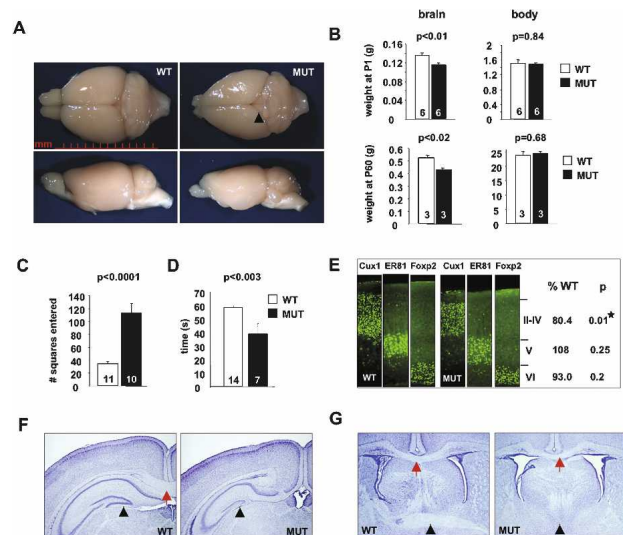
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**Figure 1.** CNS-specific *Tbr2* inactivation causes microcephaly and behavioral abnormalities. (A) Dorsal and lateral views show a significantly reduced size of the cortex (note the more exposed tectum, arrowhead) and OB in adult mutant brains. (B) Brain, but not body, weight is significantly decreased at P1 and P60. (C) Mutants display substantial hyperactivity in the open field. (D) Hang wire exam reveals reduced grip strength. (E) Layer-specific immunostaining shows significantly reduced upper cortical layers, however normal layering is maintained ( $n = 3$  per group). (F) The CC is thinner and smaller in the rostro-caudal dimension (red arrow, also cf. in G) and the DG lacks the lower blade (arrowhead). (G) Coronal sections show normal ventricles and absence of the anterior commissure (arrowheads). ( $n =$  animal number in bars, mean  $\pm$  SEM,  $t$ -test.)

normally. Additionally mutants showed a remarkable hyperactivity. Behavioral examination in the open field test showed their substantially increased exploratory behavior (Fig. 1C). Similar to hypotonia in *Tbr2*-deficient humans, mutant mice display significantly reduced grip strength (Fig. 1D).

Gross examination of mutant brains showed a markedly reduced brain size (Fig. 1A). In contrast, mutant body weight was normal (Fig. 1B). The size of the cortex and the olfactory bulbs (OBs) was markedly reduced. These differences were detectable at birth, suggesting *Tbr2* requirements during embryonic neurogenesis (Fig. 1B).

Histological examination revealed several morphological defects. Mutants displayed a reduction in cortical thickness. Analysis with layer-specific markers *Cux1* (layers II–IV), *ER81* (V), and *Foxp2* (VI) demonstrates that upper-layer thickness is reduced (Fig. 1E). Consistent with reduced upper cortical layers as the principal source of callosal projection neurons (Jensen and Altman 1982), the CC was smaller (Fig. 1F,G). Furthermore, the contralateral projecting fibers from olfactory nuclei and temporal lobes (Zhou et al. 2008), which form the anterior commissure, were absent (Fig. 1G). In the hippocampal dentate gyrus (DG) the suprapyramidal but not the infrapyramidal blade developed (Figs. 1G, 5 [below]). In addition, the mutant OB lacked the characteristic mitral cell layer (Supplemental Fig. 2).

To further explore these abnormalities, we analyzed proliferation, apoptosis, and migration during cortical development. Proliferation of VZ and SVZ cortical progenitors was evaluated by phosphohistone H3 (pH3) immunoreactivity. The number of dividing cells at the ven-

tricular surface was normal. However, a significant decrease of nonventricular surface (nVS) proliferation suggested mutants display reduced SVZ neurogenesis (Fig. 2A). To test whether the reduced number of nVS mitosis is due to prolonged cell cycle of SVZ progenitors or reduced production of IPCs in the VZ, we performed BrdU, Ki67 double-labeling (Chenn and Walsh 2002). Following a 30-min BrdU pulse to label S-phase cells, we performed double-immunostaining for pan-proliferation marker Ki67 and BrdU, consistent with a reduced number of abventricular pH3<sup>+</sup> cells, we found significantly reduced numbers of Ki67<sup>+</sup> cells in the mutant SVZ (Fig. 2B). However, there were no significant differences in the ratio of BrdU<sup>+</sup>, Ki67<sup>+</sup> double-labeled cells over total Ki67<sup>+</sup> cells, indicating that SVZ precursors cycle at similar rates (Fig. 2B). To test whether *Tbr2* might be required for the generation of IPCs we examined cell cycle re-entry at E16 after pulse labeling with BrdU 24 h earlier. Cells remaining in the cell cycle were identified as BrdU<sup>+</sup> and Ki67<sup>+</sup> double-labeled (see the Materials and Methods). The ratio of Ki67<sup>+</sup>, BrdU<sup>+</sup> double-labeled cells over total BrdU<sup>+</sup> cells in mutants was significantly decreased (Fig. 2C), indicating reduced cell cycle commitment.

To evaluate whether increased apoptosis contributes to the reduced brain size, caspase3 activity was analyzed at E15. There was no evidence for enhanced cell death in the mutant cortex (Fig. 2D).

Next we explored potential migration deficits. We found normal Reelin staining, suggesting normal positioning and function of Cajal-Retzius cells; furthermore, Calretinin staining showed normal preplate splitting. Expression of *mDAB1* and *p35*, genes with key roles in neuronal migration (Ayala et al. 2007), did not reveal abnormalities (Supplemental Fig. 4). Thus, correct laminar position of cortical neurons and normal expression of developmental markers suggest that *Tbr2* inactivation does not overtly affect cortical neuronal migration, consistent with the observed down-regulation of *Tbr2* in migrating neurons (Englund et al. 2005).

To explore whether *Tbr2* deficiency impacts more specifically on neurogenesis in the cortical SVZ, we examined the expression of *Pax6*, marking RGs in the VZ, and the SVZ marker genes *Svet1* and *Cux2*. *Pax6* expression in the cortical VZ was unaffected, suggesting normal development of VZ progenitors (Fig. 3A). Likewise, expression of *Pax6* at later stages and additional VZ progenitor markers (*Hes5*, *Sox2*) was normal (Supplemental Fig. 3A). In contrast, expression of *Svet1* and *Cux2* showed a severe and disproportional reduction with respect to the ~25% decrease of proliferating cells in the mutant SVZ (Fig. 3A). This does not simply reflect a developmental delay since similar observations were made at later stages (Supplemental Fig. 3B).

Consistent with decreased IPC proliferation, *NeuroD* and *Tbr1* staining revealed reduced numbers of newly formed neurons (Fig. 3B).

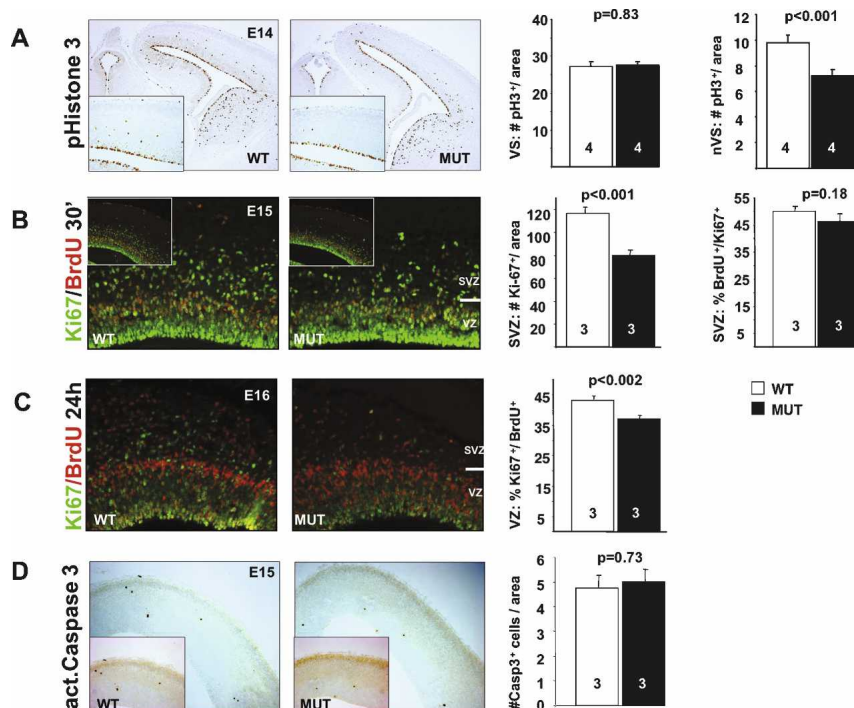
To analyze neurogenesis independently from marker expression, we performed birthdating experiments by injecting BrdU at E16 and analyzing the number of BrdU<sup>+</sup> neurons at postnatal day 10 (P10), when neuronal migration is largely completed. The number of BrdU<sup>+</sup> cells was significantly reduced in mutant upper cortical layers, but as judged by correct laminar positions of BrdU<sup>+</sup> cells, neuronal migration is unaffected (Fig. 3C).

Does transient *Tbr2* expression in IPCs also direct the differentiation of their neuronal progeny? Expression of

Satb2, known to be required for specification of upper-layer neurons (Britanova et al. 2008), is substantially reduced in mutants (Fig. 4A). Intriguingly, expression of Brn2 (Sugitani et al. 2002) was markedly reduced specifically in upper cortical layers II/III, while expression in layer V was unperturbed (Fig. 4B). Similarly, expression of lower cortical layer markers Ctip2, ER81 (layer V), Otx1 (layers V and VI), and Nurr1 (layer VIb) (Fig. 4C,D; data not shown) did not reveal differences. As judged by expression of the pan-neuronal markers NeuN (Fig. 4E) and MAP-2 (Supplemental Fig. 5), loss of Tbr2 does not simply cause a general delay in neuronal differentiation. Normal immunostaining with SMI-32, MAP-2, and Golgi impregnation shows that the reduced cortical size is not associated with gross abnormalities in dendritic arborization (Supplemental Fig. 5). The smaller size of the CC, delayed differentiation of upper cortical layer neurons, and CC agenesis in Tbr2-deficient humans led us to investigate potential abnormalities of interhemispheric connections. DiI tracing experiments reveal Probst bundles in the mutants (Fig. 4F). Thus, a fraction of callosal axons are misrouted. Collectively, our experiments demonstrate that transient Tbr2 expression is required for normal differentiation of upper cortical layer neurons.

The neuronal differentiation defects prompted us to explore the characteristic somatotopic organization in the Barrel cortex. Using either serotonin transporter (5HTT/SERT) immunohistochemistry or DiI tracing of thalamic afferents, we detected normal thalamic projection into supragranular layers. However, the typical input-related patterning was strongly diminished (Fig. 4G,H). Bisbenzamide staining also revealed the loss of cytoarchitectonic barrel pattern (Supplemental Fig. 4). Thus, Tbr2 is essential for normal development of peripheral input-related pattern formation and cytoarchitectonic differentiation in layer IV of the somatosensory cortex.

The sequential generation of supra- and infrapyramidal granule neurons in the DG is reminiscent of the neurogenic phases of lower and upper cortical layer neurons in the VZ and SVZ, respectively. Neurons of the suprapyramidal blade are primarily derived from early cell divisions in the dentate neuroepithelium near the ventricle. In contrast, the infrapyramidal blade originates mainly postnatally during a subsequent wave of neurogenesis in the dentate migration stream and hilus (Altman and Bayer 1990). To investigate neurogenesis in the DG, we performed Tbr2, Ki67 double-staining in wild types at P5. Tbr2 shows strongest expression in the dentate migration stream and hilus, and ~66 ± 7% (mean ± SD) of Tbr2<sup>+</sup> cells ( $n = 600$ ) expressed Ki67 (Fig. 5A). Conversely, among the population of Ki67<sup>+</sup> cells



**Figure 2.** Decreased proliferation in the mutant SVZ. (A) pH3 staining reveals similar numbers of mitoses on the VS but significantly reduced numbers of mitosis in nVS areas. (B) Ki67, BrdU double-staining following 30-min BrdU pulse labeling shows reduced numbers of Ki67<sup>+</sup> SVZ progenitors. Cell cycle length of mutant SVZ progenitors is unchanged as indicated by similar ratios of Ki67<sup>+</sup>, BrdU<sup>+</sup> double-positive cells over total Ki67<sup>+</sup> cells. (C) Ki67, BrdU double-staining following BrdU pulse labeling 24 h earlier reveals increase rate of cell cycle exit in Tbr2 mutants, indicated by the reduced ratio of Ki67<sup>+</sup>, BrdU<sup>+</sup> double-positive cells over total BrdU<sup>+</sup> cells. (D) Activated caspase3 staining of the E15 cortex demonstrates apoptosis is unaffected.

( $n = 606$ ), ~52 ± 14% were Tbr2<sup>+</sup>. In the early postnatal DG substantial gliogenesis occurs concomitantly with the generation of granule neurons (Yuasa 2001). This potentially masks effects of Tbr2 deficiency on proliferation of neuronal precursors in mutants; indeed Ki67 staining was indistinguishable between genotypes (Fig. 5B). Interestingly, immunostaining for activated caspase3 revealed significantly enhanced cell death in the dentate migration stream (Fig. 5C) that partly explains the substantial reduction of neurogenesis in the mutant dentate migration stream and the hilus as revealed by NeuroD immunostaining (Fig. 5D).

In the adult brain, the DG and the periventricular subependymal zone are the principal sites of neurogenesis. In the mutant DG, the number of Ki67<sup>+</sup> (Fig. 5E) and BrdU<sup>+</sup> (data not shown) cells was strongly reduced and we never observed newly formed neurons as revealed by Doublecortin (DCX) (Fig. 5F) or Calretinin (data not shown) staining (Kempermann et al. 2004). Interestingly, while neurogenesis in the DG was completely absent, neurogenesis in the subependymal zone of adult mutants appeared normal as revealed by DCX staining in the rostral migratory stream (Fig. 5G). These findings suggest that Tbr2 is developmentally required for establishment of neuronal progenitors in the adult DG but not for the adult subependymal zone.

Finally, we explored Tbr2 contributions to the generation of OB mitral cells. The typical layer of mitral cells,



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normally seen in Nissl staining, was undetectable (Supplemental Fig. 2A). Reelin staining revealed decreased numbers of mitral cells that failed to organize in a single layer and rather appear ectopically scattered (Supplemental Fig. 2B). In DiI tracing experiments, back-labeling of mitral cells from the lateral olfactory tract revealed fewer cells that were not confined to the mitral cell layer (Supplemental Fig. 2C). Anterograde labeling from the OB showed reduced fibers and more diffuse projections into the lateral olfactory tract (Supplemental Fig. 2D).

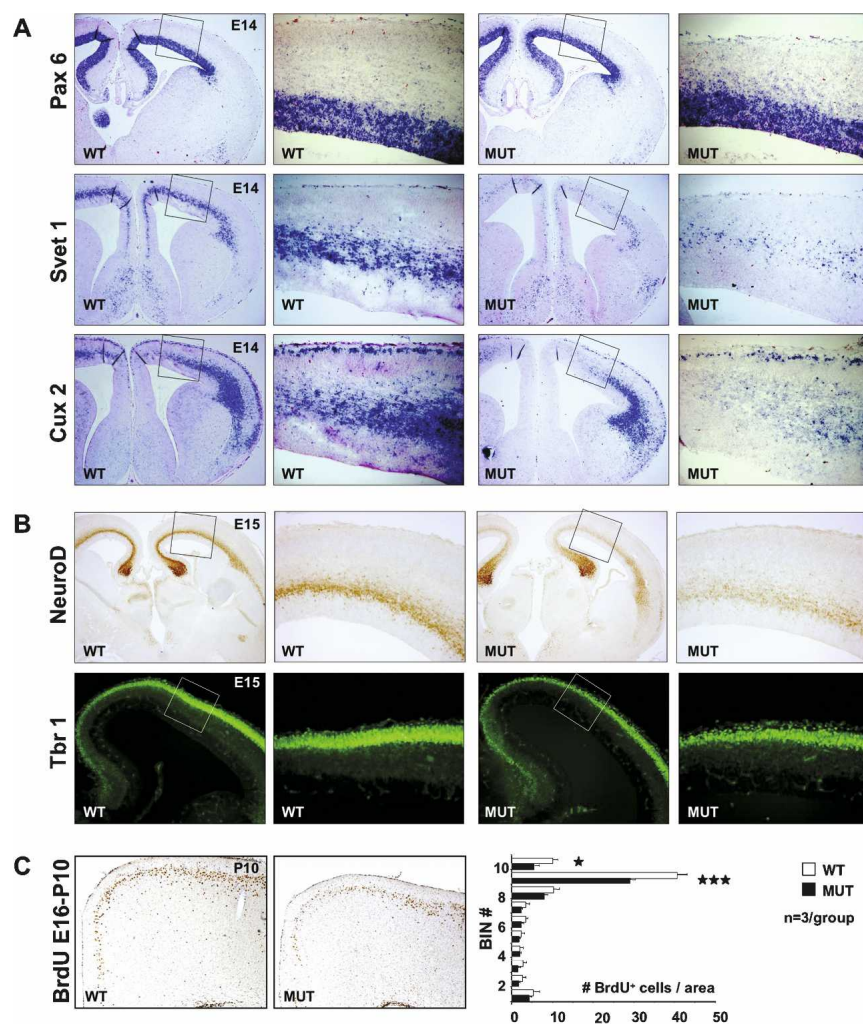
In summary, we show several critical functions for Tbr2 in cortical neurogenesis. Loss of Tbr2 leads to a reduced number of proliferating cells in the SVZ. Tbr2 is expressed at early stages of IPC development concomitant with progressive down-regulation of Pax6, thus marking the transition from RG to IPC (Englund et al. 2005). The generation of Tbr2<sup>+</sup> progenitors and upper cortical layer neurons is reduced in Pax6-deficient mice

(Tarabykin et al. 2001; Quinn et al. 2007). Similarly, the transcription factor *tailless* (*tlx*, alias *Nr2e1*) is expressed in cortical progenitors slightly earlier than Tbr2, and *tlx* mutant mice also show a reduced size of the SVZ and specific reduction of upper cortical layers. As for Tbr2 mutants, they also display enhanced aggressiveness and infanticide (Monaghan et al. 1997; Land and Monaghan 2003). Similar to Tbr2-deficient mice, neither of these mutants shows complete loss of SVZ progenitors, suggesting heterogeneous molecular mechanisms for IPC formation. The reduced *Svet1* and *Cux2* expression levels in Tbr2 mutants are more severe than the reduction of SVZ progenitor numbers. Similarly, the decreased *NeuroD* expression is more pronounced than the reduction of neuronal numbers deduced from cortical thickness measurements. This suggests that loss of Tbr2 affects the specification of IPCs and the differentiation of their neuronal progeny.

Despite reduced SVZ progenitor numbers, the majority of upper cortical layer neurons form in the absence of Tbr2. Furthermore, the expression of specific neuronal markers (*Satb2* and *Brn2*) appears more affected than others (*Cux1* and *NeuN*). Thus, subprograms of neuronal specification are likely to be differentially regulated by Tbr2. In this context most CC fibers developed and crossed the midline but only a fraction of callosal projections displays pathfinding defects. This is in contrast to the observed CC agenesis in humans (Baala et al. 2007) and might indicate species-specific differences.

In the mutant OB mitral projection neurons are reduced and their organization and projection disturbed, similar to Tbr1 mutant mice (Bulfone et al. 1998), suggesting a lineage relationship with Tbr2. Despite the close similarity of Tbr1 and Tbr2 T-box DNA-binding domains, both family members seem to have evolved for distinct functions in brain development. Indeed the decreased reelin expression, defective preplate splitting, and cortical migration deficits characteristic of Tbr1 mutants are not observed in Tbr2 mutants (Hevner et al. 2001).

Loss of Tbr2 function severely compromises DG development. Enhanced apoptosis in the dentate migration stream contributes to the premature exhaustion of DG neurogenesis and profoundly affects the late developing lower blade. Similar to the neocortex, the substantially reduced *NeuroD* expression in the developing mutant DG probably reflects Tbr2 requirements during early neuronal differentiation. The reduced numbers of adult DG progenitors implicate Tbr2 functions in both developing and adult progenitor populations. During devel-



**Figure 3.** Reduced expression of SVZ markers and decreased neuronal output. (A) In situ hybridization for the VZ marker Pax6 is largely unaffected, whereas expression of the SVZ markers *Svet1* and *Cux2* is markedly reduced in Tbr2 mutants. (B) Immunostaining for *NeuroD* and *Tbr1* reveals reduced numbers of cortical neurons generated in the mutant cortex. (C) Birthdating experiment of P10 cortical sections showing BrdU-labeled neurons at E16. Generation of upper cortical layer neurons is significantly ([\*]  $P < 0.05$ ; [\*\*\*]  $P < 0.005$ ) reduced in Tbr2 mutants.

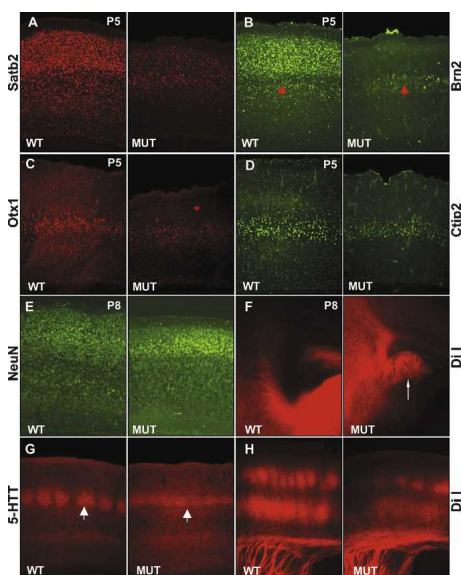
opment Tbr2 is required to establish the pool of adult DG progenitors. Additionally, in the adult DG, Tbr2 expression in IPCs potentially regulates cell cycle re-entry (Hodge et al. 2008). This dual requirement is reminiscent of *tlx* mutants, which display underdeveloped infrapyramidal blades (Monaghan et al. 1997), while *tlx* is independently required for adult DG neurogenesis (Shi et al. 2004). Further studies in inducible conditional mutant mice are necessary to rigorously discern Tbr2 functions in DG neurogenesis in the developing and adult brain.

## Materials and methods

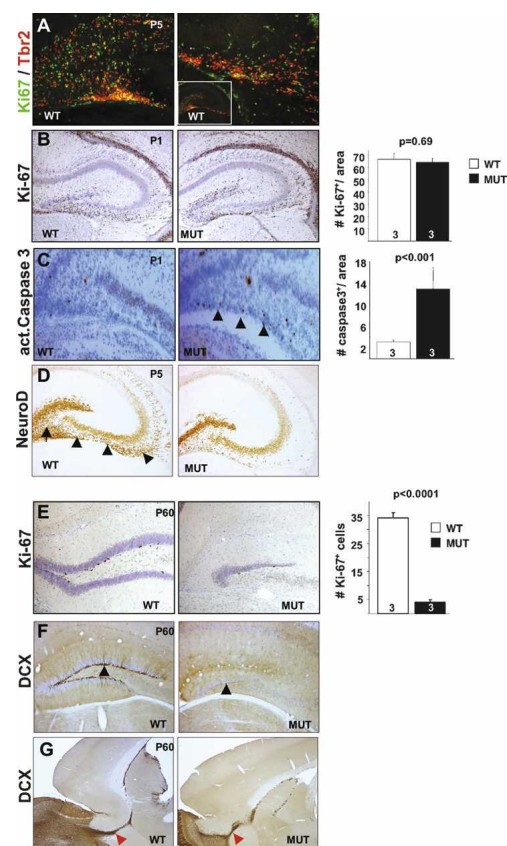
For cell cycle length studies, E15 mice were injected with 50 mg/kg BrdU i.p. and brains were recovered 30 min later. Randomly selected Ki67<sup>+</sup> cells ( $n = 1500$  per genotype) in the SVZ were identified; subsequently, BrdU labeling was determined by switching to the corresponding fluorescence channel (Chenn and Walsh 2002). SVZ precursor cells were identified as those set apart from the characteristic dense layer of BrdU<sup>+</sup> precursors in the upper VZ as described (Cubelos et al. 2007).

To study cell cycle re-entry, E15 mice were injected with 50 mg/kg BrdU i.p. and brains were recovered 24 h later. The interval allows for VZ-derived cells to become Tbr2<sup>+</sup> IPCs (Siegenthaler et al. 2008). Randomly selected BrdU<sup>+</sup> cells ( $n = 1500$  per genotype) in the VZ and VZ/SVZ interface were analysed for Ki67 expression. The SVZ was excluded from the counting frame to minimize contributions of IPC-derived BrdU<sup>+</sup> cells, which were in S phase at the time of injection. The boundary was defined by a characteristic band of strongly BrdU<sup>+</sup> cells due to the 2-h interval of maximum bioavailability of a 50 mg/kg BrdU pulse (Takahashi et al. 1992) and the typical ellipsoid shape and vertical orientation of VZ nuclei (Takahashi et al. 1995; Calegari et al. 2005).

Additional Materials and Methods are provided as Supplemental Material.



**Figure 4.** Tbr2 is required for differentiation and functional organization of upper cortical layer neurons. (A,B) Expression of *Satb2* (A) and *Brn2* (B) in upper cortical layers is reduced, whereas *Brn2* in layer V is maintained (red arrows). (C,D) Expression of lower cortical layer markers *Otx1* (layers V, VI) (C) and *Ctip2* (layer V) (D) appears unaffected. (E) The reduced expression of upper-layer markers is not due to a general differentiation defect as indicated by normal expression of *NeuN*. (F) *DiI* tract tracing of the interhemispheric connections reveals the development of Probst bundles indicative of misrouted callosal axons (arrows). (G,H) 5HTT staining (G) and *DiI* tracing (H) of thalamo-cortical afferents demonstrate disturbed somatotopic organization of the Barrel cortex (arrows depict an example for the characteristic clustering of thalamo-cortical afferents in layer IV).



**Figure 5.** Tbr2 is required for DG development. (A) Tbr2, Ki67 double-labeling at P5 shows Tbr2 expression in progenitors of the dentate migratory stream. Most Tbr2<sup>+</sup> cells are proliferating; however, a substantial fraction of proliferating cells is Tbr2<sup>-</sup>. (B) Overall, the number of Ki67<sup>+</sup> cells in the developing mutant DG is indistinguishable from wild-type littermates. (C) Activated caspase3 staining shows substantially increased apoptosis (arrowheads) in the dentate migration stream and hilus. (D) Neurogenesis in the dentate migration stream is strongly reduced as demonstrated by *NeuroD* expression (arrow, hilus; arrowheads, migratory stream). (E) The adult mutant DG shows strongly reduced number of Ki67<sup>+</sup> progenitors. (F) *DCX* staining demonstrates the absence of neurogenesis. (G) In contrast, neurogenesis in the adult SVZ is unaffected. ( $n =$  animal number in bars, mean  $\pm$  SEM, *t*-test.)

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