Otx2 Controls Identity and Fate of Glutamatergic Progenitors of the Thalamus by Repressing GABAergic Differentiation

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GABAergic and glutamatergic neurons modulate inhibitory and excitatory networks in the CNS, and their impairment may cause neurological and psychiatric disorders. Thus, understanding the molecular mechanisms that control neurotransmitter phenotype and identity of excitatory and inhibitory progenitors has considerable relevance. Here, we investigated the consequence of Otx2 (orthodenticle homolog) ablation in glutamatergic progenitors of the dorsal thalamus (referred to as thalamus). We report that Otx2 is cell-autonomously required in these progenitors to repress GABAergic differentiation. Our data indicate that Otx2 may prevent GABAergic fate switch by repressing the basic helix-loop-helix gene Mash1 (mammalianachaete-schute homolog) in progenitors expressing Ngn2 (neurogenin homolog). The lack of Otx2 also resulted in the activation of Pax3 (paired box gene), Pax7, and Lim1 (Lin-11/Isl-1/Mec-3), three genes normally coexpressed with Mash1 and GABAergic markers in the pretectum, thus suggesting that thalamic progenitors lacking Otx2 exhibit marker similarities with those of the pretectum. Furthermore, Otx2 ablation gave rise to a marked increase in proliferating activity of thalamic progenitors and the formation of hyperplastic cell masses. Thus, this study provides evidence for a novel and crucial role of Otx2 in the molecular mechanism by which identity and fate of glutamatergic precursors are established in the thalamus. Our data also support the concept that proper assignment of identity and fate of neuronal precursors occurs through the suppression of alternative differentiation programs.

Key words: Otx2; neuronal differentiation; glutamatergic fate; GABAergic fate; proliferation; thalamus
in sodium citrate buffer (10 mM), pH 6 (Shi et al., 1991, 1997, 2001), and unmasked by four rounds of microwave boiling (4 min/boiling) at 700 W. Only sections from wax-embedded embryos were used. Antigens were described previously (Puelles et al., 2003, 2004) or correspond to PCR antibodies, for their observation and image capture. A modification of the common procedure described above that allowed us to detect four or more antibodies in the cells of the same section. This procedure has been used up to five sequential rounds for a maximum of 10 different primary antibodies, in this study, sections were used for a maximum of two sequential rounds. In this modified procedure, lack of fluorescence indicated that secondary antibodies, but not necessarily also the primary antibodies, had been removed. To rule out this possibility, control slides immunostained with the same primary and secondary antibodies were boiled and hybridized in parallel only with secondary antibodies to check the absence of staining and, thereby, the absence of residual primary antibodies. We also tested that antibodies used in the first round efficiently worked after the last round of immunohistochemistry, thus proving that, during the procedure, antigens were not severely affected. Importantly, we preselected the primary antibodies for their efficiency and used those with the highest antigen affinity, and thereby more difficult to be removed, in the last round of immunohistochemistry [e.g., a-bromodeoxyuridine (aBrDU)].

**Antibodies.** The rabbit antibodies were directed against Otx2 (1:5000; G. Corte, National Institute for Cancer Research, Genova, Italy), Pax3 (1:200; Zymed, San Francisco, CA), GABA (1:500; Sigma-St. Louis, MO), Lim1 (1:200; Chemicon, Temecula, CA), VGLUT2 (1:300; SySy, Goettingen, Germany), and Ph-H3 (the phosphorylated form of histone H3) (1:200; Upstate, Charlotte ville, VA). The mouse antibodies were directed against GFAP (1:200; Chemicon), S-HT (1:100; Chemicon), TH (1:300; Chemicon), neurofilament (1:100; Developmental Studies Hybridoma Bank, University of Iowa, Iowa City, IA), Lim1 (1:100; Developmental Studies Hybridoma Bank), Mash1 (mammalian achaete-schute homolog) (1:10; F. Guillemot, National Institute for Medical Research, London, UK), Ng2 (neurogenin) (1:25; D. Anderson, Howard Hughes Medical Institute, Pasadena, CA), Pax7 (1:100; Developmental Studies Hybridoma Bank), and BrdU (1:100) (Becton Dickinson, Mountain View, CA). The goat antibody was directed against Pax3/7 (1:100; Santa Cruz Biotechnology), the sheep antibody was directed against BrdU (1:200; Abcam, Cambridge, MA), and the guinea pig antibody was directed against VGLUT2 (1:500; Chemicon).

**Analysis of the cell patches: size, location, and expression pattern.** For the anatomic-histological analysis aimed at studying size and location of the patches, we analyzed six conditional mutants at embryonic day 12.5 (E12.5) and E13.5, and six at E16.5. The same number of wild-type and Otx2flox−/− control embryos was analyzed at the same stages. Embryos were wax embedded, sectioned, and Nissl stained. Every fourth serial section for E12.5 and E13.5 embryos and every sixth serial section for E16.5 embryos were selected and analyzed for the size and location of cell patches. Three different sizes were arbitrarily defined: the small size corresponded to the cell patches that were one-half or less of those with intermediate size, which were, in turn, between one-half and one-fourth of those with the large size.

For *in situ* hybridization analysis with radiolabeled probes, a total of 15 conditional mutant embryos at E12.5 and 11 at E16.5 were analyzed together with a corresponding number of Otx1flox+/−; Otx2flox−/− control embryos.

For the analysis of cell patches in terms of expression pattern and relative distribution of Lim1− and Pax3/7−/− Otx2−/− cells, six conditional mutants were studied at E12.5 and E16.5, respectively. After the immunostaining with Otx2, Pax3/7, and Lim1, 115 patches at E12.5 and 105 at E16.5 were scored to inspect whether the same general pattern of expression reported in Figure 4 was detected in all of the patches. For numerical analysis, 80 patches immunostained for Lim1 and Pax3/7 were selected and analyzed at E12.5 and E16.5, respectively. Images of these patches were captured, cells were counted, and the ratio between Lim1− and Pax3/7−/− Otx2−/− cells was determined.

**Cell proliferation experiments and cell counting.** For short-pulse experi-
analyzed. After a BrdU pulse of 60 min, the ratio between the mean of the BrdU− cell number detected in the patches and that of control embryos gives a numerical estimation of the DNA synthesizing activity (S phase). The same procedure was used for Ph-H3− cells to compare the mitotic activity (M phase) of the patches with that of control embryos.

**Results**

*Otx2* inactivation in the thalamus of *Otx1cre/+; Otx2floxflox* mutants

To study the role of *Otx2* in the differentiation of thalamic progenitors, *Otx2* was inactivated by *Otx1*-driven Cre recombinase activity. The generation of the *Otx1cre* and *Otx2floxflox* mouse models has been reported previously (Puelles et al., 2003).

To define the areas of coexpression, we compared the distribution of *Otx1*, *Cre*, and *Otx2* transcripts in the thalamus of wild-type and *Otx1cre/+; Otx2floxflox* mutants. No difference in expression pattern was detected between *Otx1* and *Cre* in *Otx1cre/+* embryos (data not shown). In these embryos, *Otx1*, *Cre*, and *Otx2* were coexpressed in the thalamic neuroepithelium at E9.5, E10.5, and E12.5 (data not shown) (Simeone et al., 1992, 1993; Puelles et al., 2003). In particular, at E12.5, their coexpression in the thalamus was primarily restricted to the ventricular and subventricular zones (Simeone et al., 1993) (data not shown). *Otx2* inactivation in this territory was monitored by analyzing the distribution of the *Otx2* protein and that of the *Otx2* un-floxed transcripts using the *Otx2* antibody and the *Otx2Δ* probe, respectively. The *Otx2Δ* probe detected only *Otx2* un-floxed transcripts and has been described previously (Puelles et al., 2003, 2004). This analysis showed that, in *Otx1cre/+; Otx2floxflox* mutants, *Otx2* was first inactivated at E10.5 in restricted groups of neuronal precursors (Fig. 1A, B, arrows), which, at E12.5, increased in number and showed heterogeneity in size (Fig. 1C,D, arrows and data not shown). Therefore, as reported previously for the midbrain (Puelles et al., 2003), *Otx1*-driven Cre recombinase showed a limited efficiency also in the thalamus, in which this is reflected in a mosaic-like inactivation of *Otx2* first detected at E10.5.

**Anatomical and histological analysis of the thalamus**

*Otx1cre/+; Otx2floxflox* mice died at birth (Puelles et al., 2003). Morphological analysis of the thalamus of conditional mutants showed no obvious impairments before E11 (data not shown), although severe abnormalities characterized by the presence of invaginations and swellings of the neuroepithelium were revealed at E12.5 (Fig. 2B). At E13.5, cell masses of irregular size were numerous, and some of them reached a remarkable size, exhibited a ventricle-like cavity, and resembled patches of proliferating neuroepithelium (Fig. 2D). At late gestation, cell masses were

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**Figure 1.** Inactivation of the *Otx2floxflox* allele by the *Otx1*-driven Cre recombinase. A–D, In situ hybridization with the *Otx2Δ* probe (A, C) and immunodetection of *Otx2* protein (B, D) in *Otx1cre/+; Otx2floxflox* and *Otx2floxflox* embryos show that, at E10.5, the *Otx2floxflox* allele is inactivated in small groups of progenitors (arrows in A, B), which, at E12.5, may increase in size (arrows in C, D). Th, Thalamus.

**Figure 2.** Anatomo-histological abnormalities in the thalamus of conditional mutants. A–F, Nile staining of frontal sections through the thalamus of *Otx1cre/+; Otx2floxflox* (A, C, E, F) and conditional mutants (B, D, F) at E12.5, E13.5, and E16.5. The arrows in B, D, and F point to swellings and/or cell masses within the thalamus. G, Neuropil immunodetection (α2H3) in frontal sections counterstained with Hoechst reveals that, compared with control embryos (G), conditional mutants (H) exhibit heavy abnormalities of axonal routes in the thalamus. Th, Thalamus; S, L, and I, cell masses of small, large, and intermediate size, respectively.
randomly distributed within the mantle zone of the thalamus (Fig. 2F), where the routes of axonal tracts were heavily affected as revealed by neurofilament staining (Fig. 2H). This abnormality may account for the lethality of conditional mutants. None of these impairments was detected in Otx1<sup>+/−</sup>; Otx2<sup>+/−</sup> control embryos (Fig. 2). We also noted that these cell masses exhibited a size variability quite pronounced. To better describe this aspect, we studied in detail E16.5 Otx1<sup>+/−</sup>; Otx2<sup>+/−</sup> mutants. Approximately 20% of cell masses scored at E16.5 belonged to the group with large size (Fig. 2F). In the same embryos, the majority of the cell masses (∼55%) was of intermediate size, which means in our arbitrary scale between one-half and one-fourth of those with large size. The residual patches (∼25%) detected in these embryos showed a small size, corresponding to one-half or less of those classified as intermediate in size (Fig. 2F). In the thalamus of the E16.5 scored embryos, we detected in total 14–28 cell masses. Regarding the location, cell masses detected at E12.5 and E13.5 were identified within the subventricular zone or at the boundary between the subventricular and the mantle zone, whereas those of large size were prevalently detected in the thalamus (Fig. 2B,D). At E16.5, we only sporadically detected cell masses close to the thalamic ventricular neuroepithelium, with the large majority of the patches located in the mantle zone (Fig. 2F). Finally, we also noted that large patches are sometimes generated by the fusion of multiple cell masses. We interpreted this heterogeneity in size and location as a consequence of the extent and stage of Otx2 inactivation. Thus, large cell masses may reflect an early inactivation of Otx2 (e.g., at E10.5) in a relatively large number of progenitors, whereas small cell masses close to the ventricle may be caused by a late inactivation (e.g., E12.5 and/or E13.5) of Otx2 in few progenitors.

**Lack of Otx2 affects cell identity**

To investigate whether the lack of Otx2 generates abnormalities in cell identity, we first analyzed the expression of a number of molecular markers at E12.5 and E16.5. These included components of the Shh pathway that are normally expressed in the proliferating neuroepithelium of the thalamus such as Gli1, Gli2, Gli3, Smo, and Ptc1, and genes transcribed in territories adjacent to the thalamus, such as Pax3, Pax7, and Lim1 in the prethalamus, Dlx1 and Dlx5 in the thalamus, or expressed in the mantle zone of the thalamus such as Gbx2. Moreover, we monitored the expression of Isl1, Foxa2 [also known as Hnf3B (hepaticocyte nuclear factor)], Mx1, three members of the Nkx family such as Nkx2.1, Nkx2.2, and Nkx6.1, two pallial markers such as Tbr1 and Emx1, and genes encoding signaling molecules such as Shh, Fgf8, Wnt1, Wnt3a, Bmp2, and Bmp6. At E12.5 and E16.5 in conditional mutants, no expression was detected in the patches for Dlx1, Dlx5, Tbr1, Emx1, Mx1, Foxa2, Isl1, and Nkx genes or any of the signaling molecules mentioned above (data not shown). For example, Shh was normally expressed in the forebrain and, in particular, its expression in correspondence of the zona limitans intrathalamic (zli) was unaffected (data not shown). However, Gli1, Gli2, Gli3, Ptc1, and Smo were transcribed in both the unaffected neuroepithelium and the patches (Fig. 3A–J), suggesting that the patches contain cells sharing similarity with progenitors normally located in the subventricular and ventricular zones.

Interestingly, abundant expression of Lim1, Pax3, and Pax7 and a complementary loss of Gbx2 was detected in the patches (Fig. 3K–R). The abnormal expression of these genes was detected reproducibly in all of the patches of conditional mutants analyzed and never in Otx1<sup>+/−</sup>; Otx2<sup>+/−</sup> control embryos (Fig. 3K–R).
3K–R). Thus, the lack of Otx2 generates dramatic transcriptional abnormalities in the patches such as the activation of genes not normally transcribed in the thalamus and the repression of Gbx2, normally transcribed in the mantle zone of the thalamus. Noteworthy, Lim1, Pax3, and Pax7 are normally coexpressed in the pretectum (data not shown) (supplemental Fig. S1, available at www.jneurosci.org as supplemental material).

Otx2 is required to suppress the generation of multiple cell types

To assess the molecular identity at the single-cell level and the timing of cell-type generation in the patches, a combined analysis of the distribution of Otx2, Pax3, Pax7, and Lim1 proteins was performed at E10.5, E12.5, E13.5, and E16.5. To this aim, we developed a procedure allowing the comparison of several antibodies on the same section to attribute a complex molecular code, if any, to the same cell (see Materials and Methods). For Pax3 and Pax7, we used an antibody recognizing both proteins because two antibodies specifically directed against Pax3 or Pax7 would not always correspond to the Otx2− progenitors and that Lim1+ cells were always detected at the edge of the Pax3/7−Otx2− cells. Moreover, a numerical comparison between Pax3/7−Otx2− and Lim1+ cells, performed on 80 patches at E12.5 and 80 at E16.5, revealed that the number of Pax3/7−Otx2− cells was between 25 and 45% of the Lim1+ cells at E12.5 and between 10 and 25% at E16.5. Noteworthy, in 10% (n = 8) of the patches analyzed at E12.5 and in 6% (n = 5) of those scored at E16.5, this proportion between Pax3/7−Otx2− and Lim1+ cells changed to ~65% at E12.5 and 40% at E16.5.

In summary, these data indicate that lack of Otx2 in a restricted number of progenitors of the thalamus affects their normal identity, as revealed by the early derepression of Pax3/7, and is subsequently reflected in the appearance of Lim1+ cells.

Proliferation and cell fate analysis

Previous experiments did not prove whether (1) Pax3/7−Otx2− cells corresponded to proliferating progenitors, (2) proliferation in the patches was abnormal, and (3) Lim1+ cells represented the postmitotic progeny of the Pax3/7−Otx2− precursors.

To address these issues, we first studied cell proliferation by providing a short pulse (60 min) of BrdU to pregnant females at E10.5, E12.5, E13.5, and E16.5. In wild-type (data not shown) and Otx1cre/+; Otx2+/− control embryos, BrdU+ cells were detected at E10.5 throughout the entire neuroepithelium and, subsequently, within the ventricular and subventricular zones, in which they gradually decreased in number and became sporadic at E16.5 (Fig. 5A–D). At E10.5, no obvious difference in the number of BrdU+ cells was observed between the Pax3/7−Otx2− and the unaffected neuroepithelium of conditional mutants or control embryos (Fig. 5A, I, M); conversely, at E12.5 and particularly at E13.5 the number and density of BrdU+ cells within the Pax3/7−Otx2− patches was remarkably increased (Fig. 5J, K, N, O, yellow arrow) or with control embryos (Fig. 5B, C). At E16.5, the number of BrdU+ cells diminished but appeared reproducibly higher in the patches of conditional mutants (Fig. 5L, P). Noteworthy, between E12 and E14, the size of BrdU−Pax3/7− cells masses greatly expanded, sometimes reaching the lateralmost edge of the thalamus (Fig. 5J, K).

Next, to assess whether the increased number of precursors in S phase was reflected in a corresponding increase in mitotic activity, we determined the number of Ph-H3+ positive cells. In conditional mutants at E10.5, Ph-H3+ cells were detected in the Pax3/7+ patches (Fig. 5Q, white arrows), and, as for the BrdU+ cells, their number and density were approximately comparable with those of control embryos (Fig. 5E) or the unaffected neuroepithelium of conditional mutants (Fig. 5Q, yellow arrows).
However, at E10.5, the position of Ph-H3+ cells was abnormal in correspondence of Pax3/7+ patches, suggesting an early alteration in the proliferation pattern. At E12.5 and E13.5, the number of Ph-H3+ cells was clearly increased along the ventricle-like cavity of the patches (Fig. 5F,G and compare yellow and white arrows in R,S). At E16.5, the number of Ph-H3+ cells decreased but was still reproducibly higher in the patches (Fig. 5, compare H, T).

Cell counting at E13.5 (see Materials and Methods) showed that, compared with Otx1Cre/+; Otx2+/− control embryos, the number of BrdU+ and Ph-H3+ cells detected in the patches was increased by 1.8- and 1.7-fold, respectively, whereas the ratio between Ph-H3+ and BrdU+ cells in control embryos (0.11) and in the patches (0.12) was comparable. This indicates that cell cycle progression from the S to the M phase is not impaired.

Therefore, these findings collectively support the possibility that Otx2, directly or indirectly, controls the number of proliferating progenitors by regulating their postmitotic transition between E12 and E14. Failure in this control may result in an increase of Otx2+ progenitors reentering the cell cycle and may account for local overgrowth generating neuroepithelial invaginations and the subsequent incorporation of cell masses within the mantle zone.

To assess whether Lim1+ neurons represented the postmitotic progeny of Pax3/7−/−Otx2− progenitors, we first studied whether Lim1+ cells were also BrdU+ in short-pulse experiments. In these experiments, Lim1− cells did not show BrdU staining (data not shown), thus indicating that they correspond to postmitotic neurons. Then, we performed a birthdating experiment by providing a long pulse of BrdU at E13.5 (see Materials and Methods) to pregnant females and analyzing at E16.5 whether postmitotic neurons BrdU+ labeled at E13.3 expressed Lim1. In this experiment, BrdU+ cells appeared at E16.5 homogeneously distributed in a ring-like shape in the patches and, importantly, did not mix with BrdU− cells generated in the unaffected ventricular zone (Fig. 5U). Coimmunostaining experiments showed that a relevant number of Lim1+ postmitotic neurons were also BrdU+ (Fig. 5V,W), indicating that Pax3/7−/−Otx2− progenitors generate Lim1+ postmitotic neuronal cell types. Moreover, because scattered BrdU−Lim1− cells were reproducibly observed (Fig. 5V,W, arrowheads), it is conceivable that these cells represent an additional population of postmitotic neurons unidentified by our markers and generated by the Pax3/7−/−Otx2− progenitors.

Lack of Otx2 induces GABAergic differentiation in glutamatergic progenitors

We next analyzed whether the lack of Otx2 caused abnormalities in terminal differentiation.
Because the thalamus is a relevant source of glutamatergic neurons, we first tested whether this major fate was retained in the patches by studying the expression of the vesicular glutamate transporter VGLUT2 (Cheng et al., 2004; Fremeau et al., 2004; Schuurmans et al., 2004). In contrast to control embryos, VGLUT2 transcripts were not detected in the patches (Fig. 6A, arrow).

Then we studied whether the lack of Otx2 was reflected in a GABAergic fate switch by using specific markers such as glutamic acid decarboxylase Gad1 (encoding GAD67) and GABA. Strikingly, abundant transcription of Gad1 and numerous GABA+ cells were identified in the patches (Fig. 6B, C, arrow). Detailed inspection at the single-cell level, in E13.5 (data not shown) and E16.5 embryos, showed that all of the GABA+ cells were VGLUT2− (Fig. 6D). Moreover, to reinforce this finding and to examine whether GABA+ cells corresponded to postmitotic neurons belonging to the patches, we performed BrdU long-pulse experiments similar to those shown previously for Lim1 (Fig. 5). In this way, we assessed whether at E16.5 BrdU+ cells labeled at E13.3 expressed GABA and/or VGLUT2. This experiment showed that a relevant percentage of BrdU+ neurons were GABA+, although none of them are VGLUT2+ (Fig. 6E). H–M, Combinatorial immunodetection with GABA (H–K), Lim1 (H, I, L, M), and Pax3/7 (J, K) antibodies shows that all of the GABA+ neurons are Lim1+. I, K, and M correspond to the magnification of the area demarcated in H, J, and L. The green color for VGLUT2 (D) is a pseudocolor.

Figure 6. GABAergic fate switch of glutamatergic progenitors. A–C, In situ hybridization (A, B) and immunohistochemistry (C) of E16.5 Otx1cre+/−; Otx2+/− and Otx1cre−/−; Otx2flox−/− embryos with VGLUT2 (A), Gad1 (B) probes, and GABA (C) antibody show that, in the patches, the lack of VGLUT2 expression correlates with the induction of GABAergic markers. D–G, Combinatorial immunodetection with GABA (D, E), VGLUT2 (D, F, G), and BrdU (E, F) antibodies in E16.5 conditional mutants labeled at E13.3 with BrdU shows that many of the BrdU+ neurons are GABA+, although none of them are VGLUT2+. H–M, Combinatorial immunodetection with GABA (H–K), Lim1 (H, I, L, M), and Pax3/7 (J, K) antibodies shows that GABA+ neurons are Lim1+. I, K, and M correspond to the magnification of the area demarcated in H, J, and L. The green color for VGLUT2 (D) is a pseudocolor.
tion of DBH and TH for catecholaminergic fate and 5-HT for serotonergic fate. In these experiments, no expression for these markers was detected.

These data indicate that Otx2 is required in glutamatergic progenitors of the thalamus to suppress the GABAergic differentiation program. In this context, the finding that all of the GABAergic neurons are Lim1⁺ suggests that Lim1 might represent a postmitotic selector of the GABAergic differentiation program. Altogether, our data also indicate that all of the abnormalities so far described in the patches are compatible with a cell-autonomous requirement for Otx2.

**Lack of Otx2 induces Mash1 expression**

In the forebrain, the proneural genes Ngn1, Ngn2, and Mash1 are expressed in different pools of progenitors in which they are required for neuronal specification. In the dorsal telencephalon and thalamus, Ngn2, in particular Ngn2, play a dual role, both being required to activate glutamatergic differentiation of early born neurons and to repress Mash1 expression and GABAergic fate (Fode et al., 2000; Schuurmans et al., 2004). We therefore studied the expression of Mash1 and Ngn2 in the patches.

Robust expression of Mash1 was first detected in a relevant fraction of the Pax3/7⁺ progenitors at E10.5 and, subsequently, maintained until late development (Fig. 7A–C). Importantly, most but not all of the Pax3/7⁺-Otx2⁻ progenitors exhibited Mash1 expression, thus suggesting that the total absence of glutamatergic neurons (Fig. 6) should primarily depend on the lack of Otx2 in all of the progenitors rather than on the activation of Mash1 in only a percentage of them. In this context, the activation of Pax3/7 in all of the Otx2⁻ progenitors strongly suggests that lack of glutamatergic fate may be the consequence of an early and general change in the identity of the mutant progenitors.

Next, we analyzed whether the activation of Mash1 correlated with the suppression of Ngn2. We found that, at E10.5, Ngn2 was absent in the thalamus neuroepithelium of control and conditional mutant embryos showing the same expression pattern (Fig. 7D,G and data not shown). In particular, in correspondence of the patches, Ngn2 was detected in both Mash1⁺ (Fig. 7D) and Pax3/7⁺ (Fig. 7G) progenitors but, subsequently, its expression was gradually reduced and totally lost at E16.5 (Fig. 7E,F,H,I). These findings suggest that, in the absence of Otx2, (1) Ngn2 expression is not affected until E12 and is subsequently lost, and (2) Ngn2 is not sufficient to repress Mash1. In this context, the gradual loss of Ngn2 expression may reflect GABAergic fate switch of Otx2⁻ progenitors.

**Discussion**

**Otx2 controls identity and fate of glutamatergic progenitors in the thalamus**

Otx genes are required for regionalization and patterning of the developing brain and, in particular, play a crucial role in the establishment of the identity and fate of progenitors located in the ventral midbrain (Puelles et al., 2003, 2004).

As for other regulatory factors, Otx2 is expressed in different brain areas generating specific neuronal cell types. Therefore, understanding how the same protein controls identity and fate of different populations of progenitors may provide relevant insights into the mechanisms by which neuronal diversity is generated.

Here we studied the role of Otx2 in the thalamus to determine whether it is required for the differentiation of glutamatergic progenitors and the regulatory steps in which it is involved.

Our results indicate the following: (1) Otx2 is required for glutamatergic differentiation; (2) lack of Otx2 and GABAergic fate switch correlate with the activation of Mash1 in progenitors expressing Ngn2; (3) lack of Otx2 also correlates with the activation of a pretectal-like gene expression profile as revealed by the induction of Pax3, Pax7, and Lim1; and (4) Otx2 has a relevant role in the control of proliferation. Together, these findings indicate that, in the thalamus, Otx2 is a novel and crucial molecular determinant controlling identity, fate, and proliferation of glutamatergic progenitors.

Relevant data have been provided previously on the genetic functions controlling GABAergic and glutamatergic differentiation (Casarosa et al., 1999; Fode et al., 2000; Marin and Rubenstein, 2002; Parras et al., 2002; Cheng et al., 2004; Miyoshi et al., 2004; Schuurmans et al., 2004). Among these, it has been shown...
that, in the telencephalon and thalamus, Ngn2, in particular Ngn2, are required to promote glutamatergic fate and to simultaneously repress Mash1 and GABAergic differentiation (Fode et al., 2000; Schuurmans et al., 2004). Conversely, in the basal forebrain, Ngn2 is unable to repress Mash1 expression (Parras et al., 2002). Our results show that, in early thalamic progenitors (E10.5) lacking Otx2, glutamatergic differentiation is abolished and Mash1 is de-repressed in a relevant subset of Otx2−/−-Pax3/7−/−Ngn2+ progenitors, suggesting that Ngn2 is not sufficient to suppress Mash1 activation in an Otx2−/− cellular context. Therefore, our findings together with those reported previously (Fode et al., 2000; Parras et al., 2002; Schuurmans et al., 2004) suggest that lack of Otx2 or Ngn2 in the thalamus generates in both cases Mash1 de-repression and GABAergic fate switch. Noteworthy, in the forebrain areas in which Otx2 and Ngn2 are coexpressed (e.g., dorsal telencephalon and thalamus), Mash1 transcription and GABAergic fate are repressed, whereas glutamatergic differentiation is activated; the opposite occurs in the areas in which only Otx2 (e.g., pretectum) or none of them (e.g., basal telencephalon and prethalamus) are transcribed. Our findings also reveal that, besides the GABAergic phenotype, Otx2−/− thalamic cell patches exhibit additional similarities in gene expression profile with the pretectum but not with the prethalamus or the basal telencephalon. Indeed, they show expression of Pax3, Pax7, and Lim1 but never that of Dlx1 and Dlx5. Thus, although not conclusive, this observation suggests that lack of Otx2 in the thalamus is reflected in the generation of GABAergic neurons, which, on the basis of their gene expression profile, appear more similar to those of the pretectum than to those of the prethalamus or basal telencephalon.

However, although in the pretectum Otx2 is normally coexpressed with Pax3, Pax7, Lim1, and Mash1, our data suggest that, in the thalamus, Otx2 is required to repress these genes. A likely possibility reconciling this apparent contrast is that activating or repressing properties of Otx2 might depend on the territorial context in which it is expressed. In agreement with this, Otx2 plays different roles in different tissues during embryonic development (Acampora and Simeone, 1999; Acampora et al., 2001; Simeone et al., 2002). In particular, Otx2 is required to control identity and fate of ventral progenitors generating dopaminergic neurons in the midbrain (Puelles et al., 2003, 2004; Vernay et al., 2005; Prakash et al., 2006). In this context, we have shown that Grg4 (groucho-related gene), also expressed in the ventral midbrain, may interact with both Otx1 and Otx2 proteins and convert them from activators to repressors (Puelles et al., 2004). Thus, it is conceivable that Otx2 may convert its transcriptional properties (from activator to repressor and vice versa) by interacting with specific co-repressing or coactivating partner(s) expressed in restricted brain areas.

Here, we studied the role of Otx2 in the thalamus and have shown that its ablation generates the activation of Pax3, Pax7, Lim1, Mash1, and GABAergic fate switch, whereas none of the genes whose expression is affected by the lack of Otx2 in the ventral midbrain (Puelles et al., 2003, 2004) is activated in the Otx2−/− patches. In this context, our preliminary results aimed at detecting the role of Otx2 in the preetectal area, which generates GABAergic neurons and in which Otx2 is coexpressed with Pax3, Pax7, Lim1, and Mash1 (supplemental Fig. S1, available at www.jneurosci.org as supplemental material), show that severe reduction of Otx2 in this region results in an increase of GABAergic neurons, thus suggesting that Otx2 in the preetectum might represent a mild negative modulator of the GABAergic fate. Thus, our data and those reported previously indicate that, depending on the brain territory, Otx2 is required to control identity and fate of different regionally restricted pools of progenitors. In this context, a crucial issue to be addressed will be to dissect in vivo the molecular basis of this regionally restricted differential control exerted by Otx2.

Finally, because the Otx2+ thalamic territory surrounding the Otx2− patches develops normally and Shh expression at the zli is unaffected, it is likely that the abnormalities reported in this study reflect the lack of a cell-autonomous control by Otx2.

**Otx2 controls proliferating activity of thalamic progenitors**

An additional consequence of Otx2 inactivation is the generation of cell masses distributed within the mantle zone of the thalamus at late gestation. Analysis of proliferating activity indicates that the number of Otx2−/− progenitors in S (BrdU+) and M (Ph-H3+) phases is increased by approximately twofold when compared with the Otx2+ progenitors of control embryos. This increase in proliferating activity is particularly evident between E12 and E14 but, at later stages, is less evident but reproducibly higher than in control embryos. In the patches, the ratio between Otx2−/− dividing progenitors (Ph-H3+) and those in S phase (BrdU+) is essentially unaltered, leading us to assume that the progression from S to M phase is unaffected and that size expansion of the patches would be attributable to altered postmitotic transition of Otx2+ progenitors. In this case, a higher proportion of Otx2−/− progenitors should reenter the cell cycle and be responsible for local and vigorous overgrowth. Whether this impairment may be considered as a somatic preneurogenic event remains to be clarified, but certainly the outcome of this strong hyperplasia may be dramatic. However, to our current knowledge, no proliferation-suppressing activity has been documented for Otx2 in humans, although a similar effect on proliferation has already been described in retinal progenitors of Otx1−/−; Otx2−/− double-mutant embryos (Martinez-Morales et al., 2001).

However, it has been reported recently that Otx2 is a medulloblastoma oncogene (Boon et al., 2005; Di et al., 2005). In this tumor, Otx2 is remarkably overexpressed, but, in the thalamus of this conditional mutant, the lack of Otx2 correlates with increased proliferation. This suggests that, depending on the expression level, Otx2 may have different roles in proliferation: its overexpression may result in a tumorigenic event, although its absence may be correlated to hyperplasia. Moreover, these findings suggest that altered proliferation may also be subject to the territory in which Otx2 expression is either lost or enhanced.

Future experiments designed to study the effect of both overexpression and ablation of Otx2 in specific cell types and at different stages should provide new insights on its role in cell proliferation in normal and pathological contexts.

**References**


