RESEARCH COMMUNICATION

COUP-TFI: an intrinsic factor for early regionalization of the neocortex

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Regionalization of the cerebral cortex is thought to involve two phases: an early regionalization phase and a later refinement phase. It has been shown that early regionalization of the neocortex does not require thalamic inputs and is regulated by intrinsic factors. Recently, two such intrinsic factors, *Pax6* and *Emx2*, have been identified. In this study, we identified *COUP-TFI* as a regulatory factor for early neocortical regionalization. The spatial and temporal expression pattern of *COUP-TFI* suggested a role in specification of the neocortex and in maintaining cortical identity. Altered region-specific expression of marker genes in the cortex as well as miswired area-specific connections between the cortex and the thalamus in *COUP-TFI* null mice indicate *COUP-TFI* plays a critical role in regulating early regionalization. Our results substantiate that *COUP-TFI*, an intrinsic factor, may work in concert with *Pax6* and *Emx2* to specify neocortical identity.

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The mammalian cerebral cortex has a very sophisticated architecture. Tangentially, it consists of many discrete areas with distinct connections and cytoarchitecture. However, the mechanism(s) whereby the neocortex is specified and regionalized still are largely unknown. Two popular hypotheses have been proposed: the proto-map hypothesis, which emphasizes genetic influences, proposes that cortical neurons are prespecified when they are born in the ventricular zone (Rakic 1988); in contrast, the protocortex hypothesis favors the role of epigenetic factors, mainly thalamocortical innervation, in cortical regionalization (O'Leary 1989). Transplantation studies have provided several lines of evidence supporting the protocortex hypothesis during postnatal development while other studies, some of which also involve transplantation experiments, have suggested that regional specification is controlled by intrinsic mechanisms (Price and Willshaw 2000). Taking advantage of mice with mutations of the *Gbx-2* and *Mash-1* genes that have defects in thalamocortical axon projections, recent studies have demonstrated that early regionalization of the neocortex does not require afferent input, instead it is controlled by intrinsic mechanisms (Miya-

shita-Lin et al. 1999; Nakagawa et al. 1999). Therefore, specification and regionalization of the neocortex can be divided into at least two steps: (1) an early regionalization phase in which cortical neurons establish their regional identity through regulating gene expression in a cell-autonomous manner and (2) a refinement phase in which extrinsic influences from subcortical areas, including thalamocortical inputs, shape and maintain the cortical subdivisions. Recent studies showed that neocortical regional identity was shifted in *Pax6* and *Emx2* mutant mice, indicating that *Pax6* and *Emx2* are two such intrinsic factors that regulate the regional expression of marker genes and specify cortical identity (Bishop et al. 2000; Mallamaci et al. 2000).

COUP-TFI is an orphan member of the nuclear-receptor superfamily. Mice with a null mutation of the *COUP-TFI* gene exhibit multiple defects in both central and peripheral nervous systems (Qiu et al. 1997; Zhou et al. 1999). In this study, we have attempted to assess the role of *COUP-TFI* in early neocortical regionalization. We demonstrated that *COUP-TFI* is required for the region-specific expression of many marker genes as well as the precise axonal projections between the thalamus and the cortex. Our findings indicate that *COUP-TFI* is a regulatory factor that works in concert with *Pax6* and *Emx2* to specify cortical identity. In addition, *COUP-TFI* may play a role in maintaining such identity.

Results and Discussion

Like other regions of the central nervous system, regionalization of the neocortex occurs very early during corticogenesis. It has been proposed that gradients of inductive signals in progenitor cells initiate early neocortical regionalization (Levitt et al. 1997). It also was suggested that the intrinsic factors controlling early regionalization would be expressed in a gradient or in a region-specific pattern in the developing neocortex (Bishop et al. 2000). Previously, we have shown that *COUP-TFI* is highly expressed in the developing central nervous system (Qiu et al. 1994). To establish *COUP-TFI* as a candidate-intrinsic factor for cortical regionalization, we first investigated the expression pattern of *COUP-TFI* during corticogenesis by section and whole-mount in situ hybridization assays. At E11.5, the onset of cerebral corticogenesis, *COUP-TFI* expression showed a graded pattern in the neocortex with high-lateral to low-medial expression in the frontal view (Fig. 1A), and high-caudal to low-rostral expression in the sagittal view (Fig. 1B). Section in situ hybridization indicated that the high caudolateral *COUP-TFI* expression gradient was maintained in the cortical plate even after birth (Fig. 1C,D). This *COUP-TFI* expression gradient in the superficial layers (layers 2/3) also can be confirmed by whole-mount, in situ hybridization (Fig. 1E,F). Just prior to our submission of this manuscript, the O'Leary laboratory reported a graded expression of *COUP-TFI* by differential display PCR that is consistent with our findings (Liu et al. 2000). This spatial and temporal expression pattern makes *COUP-TFI* a likely candidate for a regionalization regulatory factor.

The above finding prompted us to examine the potential role of *COUP-TFI* in early regionalization by investigating the region-specific expression of marker genes

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Figure 1. Graded expression of *COUP-TFI* during corticogenesis. In-situ hybridization using 35 S-UTP-labeled *COUP-TFI* riboprobe on frontal (A) and sagittal (B) sections of E11.5, coronal (C) section of P1, and sagittal (D) section of P0 mouse brains. Whole-mount in situ hybridization using digoxigenin-labeled *COUP-TFI* riboprobe on P0 mouse brains (E,F). Because of limitation of probe penetration, only expression in superficial layers (layers 2/3) can be detected by whole-mount in situ hybridization. At E11.5, a high-lateral to low-medial and a high-caudal to low-rostral *COUP-TFI* gradient was observed in frontal (A) and sagittal (B) view. This high-caudolateral expression was detected in the cortical plate after birth as indicated by section (C,D) and whole-mount (E,F) in situ hybridization. cp, Cortical plate; ctx, cortex; vz, ventricular zone. Bar: A and B, 200 μ m; C–F, 500 μ m.

including *Id2*, *ROR β* , and *Cadherin 8*, which has been used to assess the regionalization of the neocortex in P3/P4 *COUP-TFI* mutant cortices. *Id2*, a helix-loop-helix transcription factor, was expressed in a region- and lamina-specific manner (Fig. 2A). *Id2* transcripts were detected in the subplate and layers 6, 5, and 2/3, as reported previously (Bulfone et al. 1995). *Id2* transcripts can be detected in the upper part of layer 5 and stop abruptly at a rostral boundary (arrow in Fig. 2A) that has been suggested to be between motor and somatosensory cortices (Rubenstein et al. 1999). *Id2* expression in layers 2/3 was restricted to the rostral part of the cortex with a boundary within the somatosensory area (arrowhead in Fig. 2A). However, the lamina-specific expression pattern was lost in *COUP-TFI* mutants (Fig. 2B), which may be because of the greatly reduced number of layer-4 neurons and subplate neurons in *COUP-TFI* mutants (Zhou et al. 1999) as well as the uniform expression of *Id2* in layer 5. More importantly, the rostral boundary in layer 5 that separates motor and sensory cortices was not apparent (Fig. 2B), and *Id2* also was expressed throughout the rostro-caudal axis in layers 2/3 in *COUP-TFI* mutants, suggesting that *COUP-TFI* is required for regional expression of *Id2* in layers 5 and 2/3 neurons.

ROR β , which encodes an orphan nuclear receptor, is specifically expressed in layer 4 neurons (Park et al. 1997; Schaeren-Wiemers et al. 1997). We have reported previously that *ROR β* expression was greatly reduced at P0 in *COUP-TFI* mutants because of the death of layer 4 neurons (Zhou et al. 1999). Since the purpose of this investigation was to study the regional expression of *ROR β* rather than compare expression levels, we intentionally took images of *ROR β* transcript in P3 *COUP-TFI* mutants at a much longer exposure time (about twice as long for mutants). Consistent with a previous

report (Miyashita-Lin et al. 1999), *ROR β* expression in P3 control-mouse cortex displayed a distinct regional pattern with a caudal boundary within the somatosensory cortex (arrowhead in Fig. 2C) and a region of much reduced expression in a more rostral position (Fig. 2C, between asterisk and arrow). However, in mutant cortex, uniform expression of *ROR β* was detected from rostral to caudal cortex with expanded caudal expression (Fig. 2D) indicating that *COUP-TFI* controls the region-specific expression of *ROR β* .

Figure 2. Changes in region-specific expression of *Id2*, *ROR β* , and *Cadherin 8* in *COUP-TFI* mutants. Section in situ hybridization using *Id2* (A,B), *ROR β* (C,D), *Cadherin 8* (E,F), and *COUP-TFI* (I) riboprobes on sagittal sections of P3 or P4 control (A,C,E,I) and *COUP-TFI* mutant (B,D,F) cortices. Whole-mount in situ hybridization using digoxigenin-labeled *Cadherin 8* riboprobe on P0 control (G) and *COUP-TFI* mutant (H) brains. *Id2* expression in the upper layer of layer 5 stopped abruptly at a rostral boundary (arrow in A) and in the rostral part of layers 2/3 with a boundary in the somatosensory area (arrowhead in A). However, *Id2* expression in P4 *COUP-TFI* mutant cortex lost its lamina-specific pattern and was uniformly expressed in layers 5 and 2/3 (B). *ROR β* expression in layer-4 neurons showed a caudal boundary (arrowhead in C) and a region of much-reduced expression in a more rostral position (between arrow and asterisk in C) in P3 control cortex, while its expression in *COUP-TFI* mutant was more uniform and expanded caudally with a much lower expression level (D). *Cadherin 8* was expressed in the rostral part of the neocortex in the layers 2/3 and 4 (arrow in E) in P3 control cortex (E). However, in P3 *COUP-TFI* mutant, *Cadherin 8* expression in layers 2/3 and 4 lost its rostral restriction (F). This caudal expansion of *Cadherin 8* expression in superficial layers also was observed by whole-mount in situ hybridization (arrowheads in G,H). In-situ hybridization with *COUP-TFI* riboprobe revealed that *COUP-TFI* was expressed throughout the neocortex (I). Specifically, a higher expression in layers 2/3 and 4 in the rostral part (arrow in I) and a high expression in layer 4 with a boundary in the somatosensory cortex (arrowhead in I) were observed. sp, Subplate. Bar, 500 μ m.

Cadherin 8 belongs to the type II class of cadherin, a family of cell adhesion molecules that are important for morphogenesis of the central nervous system. At P3, in situ hybridization assay has revealed that *Cadherin 8* was uniformly expressed in layer 5 while its expression in layers 2/3 and 4 was restricted to the rostral part of the neocortex (arrow in Fig. 2E). However, in P3 mutant cortex, *Cadherin 8* expression in layers 2/3 and 4 lost its rostral restriction and could be detected throughout the rostral-caudal axis, while expression in layer 5 was unchanged (Fig. 2F). The expression of *Cadherin 8* in superficial layers (layers 2/3) also was detected by whole-mount, in situ hybridization (Fig. 2G,H). *Cadherin 8* was expressed in the frontal area of the neocortex with a boundary around the limit of motor cortex (arrowheads in Fig. 2G) in P0 control brains. The rostral *Cadherin 8* expression boundary in superficial layers was shifted caudally, especially in the medial region, in P0 mutant cortices (arrowheads in Fig. 2H). Therefore, the loss of restricted rostral expression of *Cadherin 8* in layers 2/3 and 4 indicated that *COUP-TFI* is required for region-specific expression of *Cadherin 8*. Taken together, the region-specific expression of many genes including *Id2*, *RORβ*, and *Cadherin 8* around P4, when cortical lamination is close to completion, is altered in *COUP-TFI* mutants indicating that normal neocortical regionalization requires *COUP-TFI*. Since expression of these marker genes is regulated by intrinsic mechanisms and is independent of thalamocortical innervation (Miyashita-Lin et al. 1999; Nakagawa et al. 1999), our findings establish *COUP-TFI* as one of the intrinsic regulatory factors that control the specification of the neocortex.

Interestingly, *COUP-TFI* itself also was expressed in a region-specific manner at P4 (Fig. 2I). Although *COUP-TFI* transcripts were detected throughout the cortex, a higher expression in the rostral part of layers 2/3 and 4 (arrow in Fig. 2I) was observed and it correlated with *Cadherin 8* expression in layers 2/3 and 4. In addition, *COUP-TFI* was highly expressed in layer 4 neurons (Fig. 2I) with several boundaries (arrow, asterisk, and arrowhead in Fig. 2I) that resembled the expression pattern of *RORβ* (Fig. 2C). Although the expression pattern of *COUP-TFI* in the postnatal cortex correlates with components of some marker genes, it differs considerably from other expression patterns, suggesting that *COUP-TFI* is important for maintaining region-specific gene expression of some but not all of the markers examined.

Because expression of *COUP-TFI* displayed a region-specific pattern very early (we can detect it at E11.5) during corticogenesis. We also investigated the expression patterns of *Id2*, *RORβ*, and *Cadherin 8* in late embryonic development. We chose E17.5 because this is a time when generation of cortical plate neurons has been completed (around early E17) (Caviness et al. 1995) and massive innervation of the cortical plate by thalamic axons has not yet started. However, at this stage, most cortical plate neurons still are migrating along radial glia, so layer-specific expression cannot be identified. *Id2* expression displayed a high-caudal to low-rostral gradient in control cortex, including the cortical plate and the ventricular zone. Its expression in mutants, though, showed a uniform rostral-to-caudal expression throughout the cortex (Fig. 3A,B). *RORβ* expression in control cortex showed a high-rostral to low-caudal gradient in the cortical plate (layer 4 neurons), which was interrupted in the future somatosensory cortex in control

Figure 3. Altered expression of *Id2*, *RORβ*, *Cadherin 8*, and *LAMP* during embryonic development. Section in situ hybridization using *Id2* (A,B), *RORβ* (C,D), *Cadherin 8* (E,F), and *LAMP* (G,H) riboprobes on sagittal (A–D) and coronal (E–H) sections of E17.5 or E15.5 control (A,C,E,G) and *COUP-TFI* mutant (B,D,F,H) cortices. Expression of *Id2* exhibited a high-caudal to low-rostral gradient at E17.5 in control cortex (A), while it was uniformly expressed in *COUP-TFI* mutant (B). The high-rostral to low-caudal *RORβ* expression gradient in E17.5 control cortex (C) also became more uniform in *COUP-TFI* mutants (D). The high-medial to low-lateral *Cadherin 8* expression (E) in control cortices was switched to high-lateral to low-medial in *COUP-TFI* mutants (F). *LAMP* transcripts were detected in the cortex of control E15.5 mice with a high-lateral to low-medial gradient (G). However, in *COUP-TFI* mutant mice, *LAMP* expression level in the cortex was greatly reduced with a rather uniform expression and an extended medial boundary (H). cp, Cortical plate; PR, the perirhinal cortex; SM, the sensorimotor cortex; sp, subplate; svz, subventricular zone; vz, ventricular zone. Bar, 500 μ m.

E17.5 cortex (Fig. 3C). Mutation of *COUP-TFI* resulted in the uniform expression of *RORβ*, and in it having a more caudal boundary (Fig. 3D). Expression of *Cadherin 8* in control cortical plate showed a high-medial to low-lateral pattern, while its transcripts displayed a high-lateral to low-medial pattern in the mutant cortices (Fig. 3E,F). Taken together, *COUP-TFI* regulates region-specific expression of *Id2*, *RORβ*, and *Cadherin 8* before E17.5. The fact that graded or region-specific expression of these marker genes was lost at E17.5 (Fig. 3), a time before thalamic axons innervate the cortical plate, again confirms that early regionalization of the neocortex is independent of thalamic innervation. Therefore, specification of the neocortex by *COUP-TFI* occurs shortly after the fate of cortical plate neurons is determined and before the innervation of the cortical plate by thalamic axons takes place.

To further demonstrate that *COUP-TFI* regulates region-specific gene expression independent of thalamic influence, we investigated the expression of other earlier markers, which are expressed before the thalamocortical axons ever reach the cortex. The limbic system-associated membrane protein (*LAMP*) is one of the earliest markers of cortical specification (Horton and Levitt 1988). In cerebral cortex, *LAMP* is expressed in early differentiated neurons in the limbic cortical region. At E15.5, a high-lateral to low-medial expression gradient, with the highest signal in the perirhinal cortex and a gradually decreased signal towards the sensorimotor cortex, was detected in control mice as previously reported (Fig. 3G) (Pimenta et al. 1996). However, in *COUP-TFI*

mutant mice, *LAMP* expression level in the cortex was greatly reduced. More importantly, the high-lateral to low-medial *LAMP* expression gradient was not obvious. Instead, *LAMP* was expressed in a rather uniform manner from the perirhinal cortex to the sensorimotor cortex with an extended medial boundary (Fig. 3H). These results clearly demonstrated that *COUP-TFI* is an intrinsic factor for early cortical regionalization, which plays an important role in specification of the cerebral cortex since the onset of the corticogenesis.

During the processing of sensory information, various functionally distinct regions in the neocortex communicate with corresponding areas in the thalamus by way of precise synaptic connections. These connections between the cortex and the thalamus are reciprocal and in a region-specific manner. For example, the somatosensory cortex receives afferent inputs from, and sends efferent projections to, the ventrobasal thalamus (VB) while the visual cortex connects with the lateral geniculate nucleus (LGN). To test the functional consequences of the altered neocortical regionalization in *COUP-TFI* null mice, we investigated the precise connections between the cortex and the thalamus by dye-tracing experiments. Lipophilic DiI and DiA crystals were placed in the somatosensory and visual cortices respectively (see Fig. 4A,A' for positions). Two types of terminal ends were visualized in the terminal fields in the thalamus (Fig. 4B). Those large ends are retrogradely labeled thalamic neurons while those small ends are boutons, the axonal terminals of cortical neurons. Therefore, the labeling we observed in the thalamus actually is the result of the sum of both retrogradely labeled thalamocortical and anterogradely labeled corticothalamic projections. As expected, placing DiI crystals in the somatosensory cortex of the control brains resulted in DiI labeling in the VB (Fig. 4D, red) while placing DiA in the visual cortex labeled the LGN (Fig. 4D, green). When similar experiments were performed in *COUP-TFI* mutant brains, we first observed a greatly reduced DiI labeling in the VB (Fig. 4F, red), which is consistent with our previously reported, misguided thalamocortical projections in *COUP-TFI* mutants. Placing DiI in the mutant somatosensory cortex labeled the VB as it did in the control brains (Fig. 4F, red). However, placing DiA crystals in the presumptive visual cortex position led to DiA labeling in the VB, which normally is connected with the LGN (Fig. 4F, green). Because the development of the thalamus was normal in P0 mutants as indicated by the normal expression of several thalamic-marker genes such as *Gbx2*, *Id4*, and *Lef1* (data not shown), such miswired corticothalamic connections suggested that the cortical area, which usually is within the visual cortex, adopted the identity of the somatosensory cortex in terms of neural connectivity in mice lacking *COUP-TFI*. Thus, these results confirm that *COUP-TFI* is required for the proper regionalization of the cortex.

Recently, it has been reported that the intrinsic factors, *Pax6* and *Emx2* also regulate the region-specific expression of *Cadherin 6* and *Cadherin 8* (Bishop et al. 2000). To elucidate the possible link between *COUP-TFI* and *Pax6/Emx2*, we investigated the graded expression of *Pax6* and *Emx2* in *COUP-TFI* mutants. In a pattern similar to that observed in the control cortices, *Pax6* transcripts were detected in a high-rostral to low-caudal gradient in sagittal sections (Fig. 5A,B) and in a high-

Figure 4. Changes in region-specific connections between the cortex and the thalamus in *COUP-TFI* mutants. (A,A') Shows the positions of DiI and DiA crystals, which were placed in the somatosensory and the visual cortices, respectively. When visualized in the coronal sections of the thalamus using a laser-scanning confocal microscope (B), two types of labeling were observed: the large ends were retrogradely labeled thalamic neurons (arrows in B) and the small ends were boutons of anterogradely labeled axonal terminals of cortical neurons (arrowheads in B). C,D and E,F were conventional microscope images. C and E were bright fields. D and F were overlapping fluorescent images with Texas Red (for DiI labeling, red) and FITC (for DiA labeling, green) filters. Placing DiI and DiA crystals in control somatosensory or visual cortices resulted in labeling in the VB and the LGN, respectively (red and green in D). Consistent with our previous report of defects in thalamocortical projections, insertion of DiI in the somatosensory cortex of *COUP-TFI* mutants resulted in a much less labeling in the thalamus (red in F). Placing DiI in somatosensory cortex led to labeling in the VB as it did in the controls (red in F). However, when DiA crystals were placed in the presumptive visual cortex of *COUP-TFI* mutants, most of labeling was seen in the VB instead of the LGN (green in F). D, Dorsal; L, lateral; LGN, lateral geniculate nucleus; VB, ventrobasal thalamus. Bar: A, 1 mm; B, 50 μ m; C, 200 μ m.

lateral to low-medial gradient in coronal sections (Fig. 5C,D) in E13.5 mutant cortices. The gradient expression of *Emx2*, which normally was in a high-caudal to low-rostral (Fig. 5E,F) and high-medial to low-lateral (Fig. 5G,H) pattern, also was maintained in *COUP-TFI* mutant cortices at E13.5. The unchanged, graded expression of *Pax6* and *Emx2* in *COUP-TFI* mutant cortices indicated that *COUP-TFI*, *Pax6*, and *Emx2* might function independently in regulating early regionalization of the neocortex. In addition, the fact that *COUP-TFI* is expressed not only in the ventricular zone but also in the cortical plate, while *Pax6* and *Emx2* are expressed mainly in the ventricular zone with complementary expression gradients suggests that *COUP-TFI* does not act downstream of *Pax6* or *Emx2* in the cortical neurons. Additionally, the difference in altered expression profiles (e.g., *Cadherin 8*) between *COUP-TFI* null mice and *Pax6* and *Emx2* mutants further suggests that *COUP-TFI* and *Pax6/Emx2* act through different pathways and these intrinsic factors work in concert to establish the identity of each subdivision of the neocortex.

Figure 5. Normal-graded expression of *Pax6* and *Emx2* in the cortices of E13.5 *COUP-TFI* mutants. Section in situ hybridization using *Pax6* (A–D) and *Emx2* (E–H) riboprobes on sagittal (A,B,E,F) and frontal (C,D,G,H) E13.5 control (A,C,E,G) and *COUP-TFI* mutant (B,D,F,H) cortices. *Pax6* was expressed in a high-rostral to low-caudal (A,B) and high-lateral to low-medial (C,D) pattern in control as well as in mutant cortices. Meanwhile, expression of *Emx2* was in a high-caudal to low-rostral (E,F) and high-medial to low-lateral (G,H) pattern in control as well as in mutant cortices. Bar, 200 μ m.

While the graded expression of *Pax6* and *Emx2* is lost around E15.5 and E16.5, respectively (Walther and Gruss 1991; Gulisano et al. 1996), the graded *COUP-TFI* expression was maintained in the neocortex even after birth (Fig. 1C–F). Furthermore, *COUP-TFI* transcript can be detected throughout the cortex during corticogenesis while *Pax6* and *Emx2* are mainly confined to the ventricular zone (Walther and Gruss 1991; Gulisano et al. 1996; Mallamaci et al. 1998). In addition, *COUP-TFI* expression at P4 also was region-specific (Fig. 2I) and certain domains correlated well with those of *Cadherin 8* and *ROR β* . These findings suggest that *COUP-TFI* is not only required for initiating the early regionalization of the neocortex but also may play a role in maintaining such regional identity, while *Pax6* and *Emx2* mainly function in initiation of the early regionalization.

The molecular mechanisms behind how intrinsic factors such as *COUP-TFI*, *Pax6*, and *Emx2* regulate region-specific gene expression still is largely unknown. We do not know if intrinsic factors act directly or indirectly in regulating transcription of marker genes. The complexity and diversity of the expression patterns suggest that early regional specification is achieved through the co-operation of multiple intrinsic factors. As a transcription factor, *COUP-TFI* regulates gene transcription through both transrepression and transactivation (Zhou et al. 2000) mechanisms. It will be interesting to determine how *COUP-TFI* controls the region-specific expression of those marker genes. Moreover, understanding the mechanism whereby the graded expression of *COUP-TFI*, *Pax6*, and *Emx2* is established will greatly enhance our understanding of how neocortical regionalization takes place.

Previously, we reported defects in guidance of thalamocortical projections in *COUP-TFI* mutants (Zhou et al. 1999). We proposed that improper differentiation of subplate neurons in *COUP-TFI* null mice may lead to altered expression of axon guidance molecules by these subplate neurons and, consequently, impaired thalamocortical projections. These guidance molecules can be diffusible molecules, membrane proteins, membrane-

bound molecules, or extracellular-matrix molecules. Increasing evidence suggests that the cortex also may produce such guidance molecules (Price and Willshaw 2000) to guide thalamocortical axons to their proper targets by either promoting or inhibiting outgrowth of these axons. In this study, we demonstrated that the specification of the cortical areas in *COUP-TFI* mutants was compromised and it is possible this defect altered expression of such guidance molecules in the cortex. As a matter of fact, expression of one such molecule, *LAMP*, was changed in *COUP-TFI* mutants. However, in *COUP-TFI* mutants, most thalamocortical axons lost their direction as soon as they left the internal capsule. Because the distance from the internal capsule to the cortex may be beyond the range that diffusible signals can function, and because thalamocortical axons leave the internal capsule before the majority of cortical-plate neurons are generated, defects in the mutant cortex (other than the subplate) are unlikely the main reason for misguidance of thalamocortical axons at the internal capsule.

Material and methods

Mice

COUP-TFI mouse breeding and husbandry was as described (Zhou et al. 1999). For this study, four pair of mice ranging from P3 to P4 were collected for section in situ hybridization. More than 10 pairs of P0 control and mutant mouse brains were used for whole-mount in situ hybridization experiments with *COUP-TFI* and *Cadherin 8* probes. Five pairs of P0 brains were used for DiI/DiA labeling experiments.

Section in situ hybridization

Section in situ hybridization was performed as described previously (Zhou et al. 1999). The cDNA templates were 1.5 kb of *COUP-TFI* cDNA, 1.9 kb of *ROR β* cDNA (Dr. Michael Becker-Andre, Glaxo Wellcome Research and Development S.A., Switzerland), 235 bp of 3'-UTR of *Id2* cDNA (Dr. Mark Israel, University of California, San Francisco), 1.1 kb of *Cadherin 8* cDNA (Dr. Masatoshi Takeichi, Kyoto University, Japan), 260 bp of *Pax6* cDNA (Dr. Peter Gruss, Max Planck Institute for Biophysical Chemistry, Germany), 300 bp of *Emx2* cDNA (Dr. Shinichi Aizawa, Kumamoto University School of Medicine, Japan), and 442 bp *LAMP* cDNA (+50–+491, generated by RT-PCR).

Whole-mount in situ hybridization

Whole-mount in situ hybridization was carried out using digoxigenin-labeled antisense cRNA probes as described previously (Qiu et al. 1997). Due to limitation of probe penetration, only expression in superficial layers (layers 2/3) can be detected by whole-mount in situ hybridization.

DiI and DiA labeling

Brains from newborn mice (P0) were removed and fixed in 4% buffered paraformaldehyde. Crystals of fluorescent dyes DiI and DiA (Molecular Probes) were inserted into the primary somatosensory or visual cortices, respectively (see Fig. 4A and A' for positions). Brains were incubated in fixative at room temperature for 10 wk, embedded in 4% low-melting-point agarose, and sectioned in the coronal plane at 125 μ m using a Leica VT1000S vibratome. Axonal terminals in the thalamus were visualized using a conventional fluorescence microscope (Zeiss Axiophot) and a laser-scanning confocal microscope (Zeiss LSM 510).

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