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***Dlx1&2*-Dependent Expression of *Zfhx1b* (*Sip1*, *Zeb2*) Regulates the Fate Switch Between Cortical and Striatal Interneurons**

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Summary

Mammalian pallial (cortical and hippocampal) and striatal interneurons are both generated in the embryonic subpallium, including the medial ganglionic eminence (MGE). Herein we demonstrate that the *Zfhx1b* (*Sip1*, *Zeb2*) zinc finger homeobox gene is required in the MGE, directly downstream of *Dlx1&2*, to generate cortical interneurons that express *Cxcr7*, *MafB* and *cMaf*. In its absence, *Nkx2-1* expression is not repressed, and cells that ordinarily would become cortical interneurons appear to transform towards a subtype of GABAergic striatal interneurons. These results show that *Zfhx1b* is required to generate cortical interneurons, and suggest a mechanism for the epilepsy observed in humans with *Zfhx1b* mutations (Mowat-Wilson syndrome).

Introduction

Cell type specification within the embryonic basal ganglia is regulated at multiple levels. Distinct subdivisions within this region generate distinct neurons. For instance, the lateral ganglionic eminence (LGE) generates striatal projection neurons whereas the medial ganglionic eminence (MGE) generates pallidal projection neurons. Domains within the MGE are biased towards generating different cell types; whereas the rostromedial MGE

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Supplemental Information

Supplemental information for this article includes six Supplemental Figures, two Supplemental Tables and Supplemental Experimental Procedures.

largely produces cortical and striatal interneurons, the caudoventral MGE largely produces pallidal projection neurons (Flandin et al., 2010; Nobrega-Pereira et al., 2010). Distinct MGE-derived cortical interneuron subtypes appear to be generated from the same progenitors, perhaps in a temporal sequence (Brown et al., 2011).

Cortical and striatal interneurons are both generated from the MGE (Marin et al., 2000). The *Nkx2-1* homeobox transcription factor has a central role in specifying their identity. While *Nkx2-1* is initially required for both of these cell types, *Nkx2-1* expression is repressed soon after immature cortical interneurons tangentially migrate from the MGE, while it is maintained in striatal interneurons (Butt et al., 2008; Marin et al., 2000; Nóbrega-Pereira et al., 2008; Sussel et al., 1999).

Forced expression of *Nkx2-1* in cortical interneurons changes their migration so that they settle in the striatum (Nóbrega-Pereira et al., 2008), providing additional evidence that repression of *Nkx2-1* is a key step in generating cortical interneurons. How *Nkx2-1* expression is repressed in these cells is unknown.

Herein, we provide evidence that the *Zfhx1b* (*Sip1*, *Zeb2*) zinc-finger homeobox transcription factor is required to repress *Nkx2-1* in the generation of cortical interneurons. In its absence, we find a decrease in cortical interneurons concomitant with increased striatal *nNOS/NPY/Sst* GABAergic interneurons. We provide evidence that expression of the *cMaf* transcription factor is a highly specific marker of the cortical interneuron lineage, and discovered that its expression is lost in *Zfhx1b* mutants.

Previous analysis of *Zfhx1b* mouse mutants has shed light on its functions in the development of cortical projection neurons (Miquelajauregui et al., 2007; Seuntjens et al., 2009). In humans, mutations of *Zfhx1b* result in Mowat-Wilson syndrome, a developmental disorder characterized by mental retardation, epilepsy and defects of neural crest-derived tissues, including craniofacial and enteric nervous system (Mowat et al., 2003). Our results that demonstrate *Zfhx1b* is required to generate cortical interneurons suggest a mechanism for the epilepsy observed in Mowat-Wilson syndrome.

Results

Conditional Deletion of *Zfhx1b* in the VZ or the SVZ of the Subpallium Using *Nkx2.1-Cre* or *Dlx12b-Cre*

Zfhx1b prenatal expression has been noted in migrating cortical interneurons and the subpallial telencephalon (Batista-Brito et al., 2008; Seuntjens et al., 2003). We found that *Zfhx1b* RNA is expressed in E12.5 MGE-derived cells that are tangentially migrating through the LGE and into the cortex by performing fluorescent in situ hybridization (FISH) on a brain in which MGE-derived cells expressed EGFP (expressed due to *Nkx2.1-Cre* induced recombination of the *CAG:CAT-EGFP* Cre reporter allele) (Figures S1 A-A”).

To determine the role of *Zfhx1b* in the development of the basal ganglia, we used a conditional mutagenesis approach. Using an allele of *Zfhx1b*, in which exon 7 is floxed (Higashi et al., 2002), we removed *Zfhx1b* expression using two different Cre alleles. Deletion of exon 7 creates a frameshift mutation and premature truncation of the protein. Previous analysis failed to detect the truncated mutant protein in *Zfhx1b* mutant tissues, providing evidence that this is a null allele (Higashi et al., 2002).

To remove *Zfhx1b* in the early progenitors of the MGE, we used the *Nkx2.1-Cre* allele (Xu et al., 2008), which drives Cre expression in the ventricular zone (VZ) of the MGE beginning around E9.5 [later it also drives expression in the subventricular and mantle zones

(SVZ and MZ)]. To differentiate between the role of *Zfhx1b* in the VZ and the SVZ/MZ, we used the *Dlx1/2b-Cre* allele (Potter et al., 2008), which drives Cre expression in the SVZ and MZ of the entire subpallium beginning around E10.5. To examine the pattern of recombination, we used an antisense riboprobe designed against *Zfhx1b* exon 7.

By E12.5, Cre activity from both the *Nkx2.1-Cre* and *Dlx1/2b-Cre* alleles removed *Zfhx1b* RNA expression in the expected patterns (Figures 1A-1C). As previously reported, the *Nkx2.1* allele did not express Cre in the dorsal-most portion of the MGE, thus explaining the persistence of *Zfhx1b* in that location (Figure 1B). Of note, in the *Nkx2.1-Cre; Zfhx1b* conditional mutant brains, *Zfhx1b* RNA expression was not observed in the cells that appear to be migrating from the dorsal MGE into the mantle of the LGE, suggesting that the *Zfhx1b*⁺ cells in the mantle of the E12.5 LGE are likely to be MGE-derived cells (e.g. cortical and/or striatal interneurons) (X in Figure 1B). Also, note that *Dlx1/2b-Cre* leads to recombination in the SVZ and MZ of the LGE, MGE and CGE (white arrowhead, Figure 1C and data not shown).

Next, we examined the expression of *Zfhx1b*'s closely related homologue, *Zfhx1a*, in the E12.5 control and mutant telencephalon. Both *Zfhx1a* and *Zfhx1b* are expressed in the subpallial VZ, whereas only *Zfhx1b* is clearly expressed in the SVZ (Figures 1A and S1B). *Zfhx1a*'s expression did not clearly change in the *Nkx2.1-Cre* mediated *Zfhx1b* mutant (Figure S1B-S1D'). Thus, in the *Nkx2.1-Cre* conditional *Zfhx1b* mutant, only the VZ of the MGE continued to strongly express a *Zfhx* homologue.

MGE-Derived Pallial Interneurons Migrate to the Striatum When Deleting *Zfhx1b* in the VZ of the MGE Using *Nkx2.1-Cre*

We analyzed the effect of deleting *Zfhx1b*, using *Nkx2.1-Cre* at multiple developmental stages, including E12.5, E15.5 and P0. To track the fate of *Zfhx1b* mutant cells, we used the *CAG:CAT-EGFP* Cre reporter allele (Kawamoto et al., 2000). Mutant brains had the following genotype: *Nkx2.1-Cre; Zfhx1b*^{F/-}; *CAG:CAT-EGFP*, whereas controls had the following genotype: *Nkx2.1-Cre; Zfhx1b*^{F/+}; *CAG:CAT-EGFP* (on occasion, some were: *Nkx2.1-Cre; Zfhx1b*^{F/+}). At E12.5, while the control brain showed a robust stream of EGFP⁺ cells migrating into the cortex, the mutant's EGFP⁺ MGE derivatives failed to migrate to the cortex, and many were detected in the LGE mantle (Figures 1D-1G').

Next, we analyzed the phenotype using molecular markers of MGE-derived cells including *Nkx2.1* and *Lhx6*. While *Nkx2.1* RNA and protein is expressed throughout the VZ and SVZ of the MGE, its expression thereafter is restricted to specific neuronal lineages. MGE-derived cortical interneurons repress *Nkx2.1* expression as they migrate out of the MGE while most, but not all, classes of striatal interneurons maintain *Nkx2.1* expression. (Flandin et al., 2010; Marin et al., 2000; Nóbrega-Pereira et al., 2008; Sussel et al., 1999). In the mutants, there was a subtle increase in *Nkx2.1* RNA expression in the LGE and CGE (Figures 1H-1J'). This increase was more apparent at higher magnification when analyzing NKX2-1 protein expression (Figures 1G and 1G'), and at later stages (E13.5 and E15.5) (Figures 1 N-P' and 2A-F').

EGFP and NKX2-1 protein expression in control brains co-localized in a subset of cells derived from the MGE. EGFP/NKX2-1⁺ cells were observed in the MGE VZ and SVZ progenitors and a subset of their derived neurons, including the globus pallidus, and striatal interneurons (Xu et al., 2008; Figure 1G solid arrowheads), while interneurons migrating to the cerebral cortex showed little to no NKX2-1 protein expression (Figure 1G open arrowheads). In mutant brains however, most if not all EGFP labeled cells had detectable levels of NKX2-1 protein, with many cells strongly co-expressing NKX2-1 and EGFP in the LGE MZ, and in a region lateral to the globus pallidus (Figure 1G' solid arrowheads). Thus,

Zfhx1b mutants had a defect in their ability to repress *Nkx2-1* RNA and protein expression, concomitant with failure of MGE-derived migration to the cerebral cortex. While *Zfhx1b* was required to repress *Nkx2-1* expression, we did not find evidence that *Nkx2-1* regulated *Zfhx1b* expression; this conclusion was based on in situ hybridization analysis of *Zfhx1b* expression in mice lacking *Nkx2-1* in newly born MGE neurons at E15.5 (*Nkx2-1* conditional mutant with *Dlx5/6-Cre*)(Figure S6).

Lhx6 RNA is expressed in tangentially migrating cells that are immature cortical and striatal interneurons, as well as cell types that remain in the subpallium (Flandin et al., 2011; Lavdas et al., 1999; Liodis et al., 2007; Sussel et al., 1999; Zhao et al., 2008). In the *Zfhx1b* mutant, *Lhx6*⁺ cells failed to be detected in the pallium, whereas they continued to be densely located throughout the MGE, and as a scattered population in the LGE and CGE (Figures 1K-1M'). On the other hand, *Lhx8* and *Gbx2* RNA expression was not appreciably changed in the mutants (Figures S1H-S1J'). Thus, *Zfhx1b* mutants may have a selective defect in cells fated to become pallial interneurons, but not cholinergic striatal interneurons. To explore this hypothesis we studied the phenotype at later developmental stages.

By E15.5, the tangential migration of immature cortical interneurons can be readily visualized by expression of *Lhx6*, *Somatostatin* (*Sst*), and EGFP (in *Nkx2.1-Cre;CAG-EGFP* brains) (Figures 2, S2D-S2F). By contrast, in *Zfhx1b* mutants (*Nkx2.1-Cre*), pallial expression of *Lhx6*, *Sst*, and EGFP was strongly attenuated (Figures 2, S2D'-S2F'). On the other hand, subpallial expression of these markers was increased in two locations: the striatum (asterisks Figures 2A-2C', 2G-2I', and S2D-S2F') and a region contiguous with the caudoventral striatum, which we believe corresponds to the anlage of the central nucleus of the amygdala (labeled e, for ectopia; Figures 2E', 2H', S2D'; note that Figure S2T shows *Dlx5* expression labeling the central nucleus of the amygdala, CeA). The ectopia in these regions also contained increased expression of *Nkx2-1* and *Sox6* (Figures 2D-2F' and 2J-2L'). These genes are normally expressed in the subpallial projection neurons such as the globus pallidus, striatal interneurons and cortical interneurons (*Sox6* only) (Azim et al., 2009; Batista-Brito et al., 2009).

Next, we tested whether the mutant cells that failed to migrate to the pallium had features of the globus pallidus or striatal interneurons. We examined expression of several globus pallidus markers including *Kcnmb4*, *Kctd12*, *Gbx2* and *Lhx8*. Unlike the abnormal expression of *Lhx6*, *Sst*, *Nkx2-1* and *Sox6*, expression of *Kcnmb4*, *Kctd12*, *Gbx2* and *Lhx8* appeared normal in the *Zfhx1b* mutants (Figures 2M-2O', 2S-2U', and S2J-S2O'), providing evidence that the abnormal collections of cells correspond either to abnormally migrated cortical interneurons, or to striatal interneurons, and not globus pallidus neurons. Furthermore, as *Gbx2* and *Lhx8* expression and function are linked to the development of striatal cholinergic interneurons (Chen et al., 2010; Zhao et al., 2003), these results provided evidence that increased striatal *Nkx2-1* expression did not correspond to cells destined to become striatal cholinergic interneurons.

To distinguish whether the abnormal collections of cells in the mutant striatum were cortical or striatal interneurons, we examined expression of *Cxcr7* and *NPY*. At E15.5 *Cxcr7* marked migrating cortical interneurons, and few cells in the striatum (Figures 7J, 7K and 7L), suggesting that it is a relatively specific cortical interneuron marker (Wang et al., 2011). In the mutant, there was a robust reduction of *Cxcr7* expression in the pattern of migrating cortical interneurons, without a substantive increase in striatal expression (Figures 7J', 7K', 7L'); a similar result was seen for *Cux2* (not shown). On the other hand, at E15.5, *NPY* expression strongly marks scattered striatal cells (probably interneurons), and relatively few migrating cortical interneurons (note, most of the cortical expression at this age resembles that of immature projection neurons in the cortical plate). In the mutant, there was a robust

increase in *NPY* expression in the striatum (Figures 2P-2R'), in a pattern closely resembling the pattern of ectopic *Nkx2-1*, *Lhx6* and *Sox6* (Figures 2D-2L'). Thus, we propose that the mutant cortical interneurons are transformed towards GABAergic striatal interneurons.

Deleting *Zfhx1b* in SVZ of the MGE Using *Dlx12b-Cre* Phenocopies Loss of *Zfhx1b* function in the VZ (*Nkx2.1-Cre*)

Towards defining the stage of differentiation when *Zfhx1b* is required for programming interneurons to migrate to the cortex, and not the striatum, we used the *Dlx12b-Cre* allele (Potter et al., 2008). *Dlx12b-Cre* expression begins in subpallial SVZ cells that express the mitotic marker Ki67 (Figures S1T-S1T''), suggesting that Cre recombination occurs in secondary progenitor cells that are mitotically active. Thus, *Dlx12b-Cre* induces recombination beginning in the SVZ of the entire subpallium, whereas *Nkx2.1-Cre* induces recombination in the VZ of the MGE and preoptic area (Figures 1D-1F').

We analyzed the effect of deleting *Zfhx1b* using *Dlx12b-Cre* at E12.5 and E15.5. In general, all of the phenotypes of MGE-derived cells observed with the *Nkx2.1-Cre* were recapitulated with the *Dlx12b-Cre* (Figures 3, S1, S3), including the strong reduction of tangential migration to the cortex, indicated by analysis of Cre-dependent reporter EGFP expression, and *Lhx6*, *Sst*, and *CXCR7* expression. Like *Nkx2.1-Cre* mutants, *Dlx12b-Cre* mutants showed increased numbers of striatal cells that expressed *Lhx6*, *Nkx2-1*, *NPY*, *Sst* and *Sox6* (Figures 3A-3I', 3M-3O' and S3J-S3L'). Furthermore, these mutants did not show an increase in the number of cells that expressed markers of the globus pallidus (*Lhx8*, *Gbx2*, *Kcnmb4*) or striatal cholinergic interneurons (*Lhx8*, *Gbx2*) (Figures 3J-3L', 3P-3R'; and S1Q'-S1S', and data not shown).

Postnatal Analysis of cortical and striatal interneuron phenotypes in *Nkx2.1-Cre;Zfhx1b* mutants—*Nkx2.1-Cre* conditional mutants died between P17 and P21; at P15, mutants weighed ~30% less than their control littermates, a phenotype that was exacerbated by litter size. We did not observe seizures or other neurological/behavioral phenotypes.

We analyzed postnatal day 0 (P0) and P15 *Nkx2.1-Cre;Zfhx1b^{F/-}* conditional mutants to better understand the nature and extent of their cortical and striatal interneuron defects. In the P0 neocortex there was an ~90% reduction in the number of EGFP⁺ Cre-reporter marked cells, as well as a decrease in Calbindin (CB), *Sst*, and *Lhx6* expressing interneurons (Figures 4A-4C'; 4S). Likewise, at P15 there was a >90% reduction in number of neocortical EGFP⁺ cells (Figure 4D-4F'; 4S). Next, we counted the number cortical interneurons in the *Nkx2.1-Cre* lineage that expressed Parvalbumin (PV) or *Sst*, which are the two main MGE-derived subtypes (Rudy et al., 2011). We saw a strong reduction in double labeled neurons, with the numbers of EGFP⁺ interneurons expressing *Sst* or PV reduced by >90% or more (Figure 4S). The expression of cortical Calretinin (CR), which predominantly marks CGE-derived cortical interneurons, showed little to no change in the *Zfhx1b*; *Nkx2.1-Cre* conditional mutant (Figure 4F-F').

In the striatum at P0, as we saw at E15.5, there was an increase in the number of cells expressing EGFP (Cre reporter), *Sst* and *Lhx6* (Figure 4G-4I'); consistent with the hypothesis that *Zfhx1b* mutant cells that were destined to go to the neocortex, instead migrated to the striatum. Additionally, there was a clear increase in the number of striatal cells expressing *nNos*, *NPY* and *Nkx2-1* (Figures 4J-4K', S4B and B'), while we observed no change in *Lhx8*, a marker for striatal cholinergic interneurons (Figures S4A and S4A').

As *NPY*, *Sst*, and *nNos* are also expressed in subsets of cortical interneurons, their increased striatal expression does not provide unequivocal information about whether supernumerary

cells correspond to cortical interneurons that failed to correctly migrate, or to interneurons that changed fate due to the mutation. To this end, we searched for a marker that is expressed in striatal, but not cortical interneurons. Substance P receptor (*TacR1*) is robustly expressed in striatal interneurons (Ardelt et al., 1996). We found that *TacR1* is almost exclusively expressed in striatal and not cortical interneurons at E15.5, P0 and P15 (Figures 4L, 4L', 4Q, 4Q', S4D-S4F', and data not shown). In *Zfhx1b-Nkx2.1-Cre* mutants at P0, there was increased striatal *TacR1* expression (Figure 4Q-4Q'), supporting the idea that at least some of the mutant cells are adopting a striatal interneuron identity.

At P15 the number of mutant cells (EGFP⁺) was roughly the same as in controls, and they were evenly dispersed within the striatum, lacking the cell clusters and ectopia (striatal and caudal amygdala) that were apparent at younger ages (Figure 4M-4M', 4T). The elimination of the excess mutant striatal cells appears to occur through apoptosis, which is robust at P0 (expression of activated cleaved-caspase 6), particularly in the ectopia (Figure S4C-S4C').

Despite the cell death, *Zfhx1b* conditional mutants at P15 continued to have significantly increased numbers of striatal nNOS, *NPY*, *Sst* and *TacR1* expressing cells (183%, 230%, 225%, and 164%; Figures 4N-4Q', 4T). Importantly, total striatal PV⁺ cells were decreased by 58% (Figure 4S). Furthermore, there was no detectable change in *TrkA* expression (Figure 4R-4R', 4T), which marks striatal cholinergic interneurons. We saw very few CR⁺ cells in the control striatum (1-3 cells per section), which did not noticeably change in the *Zfhx1b* conditional mutant (data not shown). Thus, *Nkx2.1-Cre;Zfhx1b* mutants have a selective increase in striatal interneurons expressing nNos, *NPY*, *Sst* and *TacR1*, but have reduced PV interneurons, and no change in cholinergic or CR interneurons.

We also analyzed the gross morphological properties of nNos/*NPY*/*Sst* striatal interneurons in the *Zfhx1b* mutant, and found that, like control brains, *Zfhx1b* conditional mutants had *Sst* processes restricted to the matrixes (Chesselet and Graybiel, 1986), in a lateral to medial gradient (Figure S4G-S4I' - arrowheads mark CB-poor striosomes), suggesting that the overproduced nNos/*NPY*/*Sst* interneurons in the *Zfhx1b* conditional mutant share grossly similar morphological properties with wildtype striatal interneurons.

***Zfhx1b* Expression Is downstream of *Dlx1/2* in the Developing Basal Ganglia**

Dlx1 and *Dlx2* are necessary for subpallial development, including interneuron migration to the cortex (Anderson et al., 1997a; Long et al., 2009a; Long et al., 2009b; Yun et al., 2002a); thus we examined *Zfhx1b* RNA expression in *Dlx1/2* constitutive null mutants using in situ hybridization (Figures 5A-B'). In control brains at E12.5 and E15.5, *Zfhx1b* was expressed in the VZ and SVZ of the subpallium, in addition to its previous described expression in the cortical plate and SVZ (Miquelajauregui et al., 2007; Seuntjens et al., 2009). *Zfhx1b* expression in the subpallial MZ was restricted to dispersed cells in the LGE and to a nucleus forming near the ventral medial ganglionic eminence (MGE) (Figures 1A; 5A, 5B). In *Dlx1/2*^{-/-} mutants, *Zfhx1b* expression was strongly and specifically decreased in the SVZ of the entire subpallium; expression in the subpallial VZ was maintained, albeit perhaps reduced (Figures 5A-B').

Towards defining the mechanisms that regulate *Zfhx1b* expression in the developing subpallium, we identified two regulatory elements near the *Zfhx1b* locus that drive expression in the developing subpallium. These enhancers, here named #649 and #675, were identified by virtue of their extremely strong evolutionary conservation (Figures 5C-G) and their reproducible enhancer activity in the forebrain of mouse embryos in transgenic experiments (Figures 5H-I) (Visel et al., 2007; Visel et al., 2008). The other genes in this region do not have known expression in the developing subpallium. Analysis of enhancer activity at E11.5 in transgenic whole mounts and sections showed that both enhancers drive

LacZ expression in the subpallium, including the SVZ of the MGE (Figures 5H,I). The spatial overlap of enhancer activities and *Zfhx1b* mRNA expression suggests that these two elements are distant-acting transcriptional activators of *Zfhx1b* in the developing subpallium. Computational analysis identified multiple candidate homeobox binding sites (asterisks in Figures 5D,E and highlighted regions in 5F,G).

To test whether *Dlx2* can regulate these candidate *Zfhx1b* enhancers we used a luciferase reporter assay. Co-transfection of a luciferase reporter construct containing enhancers #649 and #675 with a *Dlx2* expression vector in P19 cells showed that DLX2 strongly activates luciferase transcription when these elements are present (Figure 5J).

To determine whether DLX2 directly regulates enhancers 649 or 675, we performed chromatin immunoprecipitation (ChIP)-qPCR of E13.5 basal ganglia using a DLX2 antibody. We found enrichment over several homeodomain-containing regions of enhancers 649 and 675, with a particular domain of #675 (region #3) showing the strongest enrichment as compared to control regions of the genome (Figure 5K). Also, the relative enrichment of the enhancer fragments was eliminated when a DLX2 polypeptide was included in the immunoreaction as a negative control (Figure 5K).

In summary, we have identified two candidate distant-acting gene regulatory elements whose activity patterns suggest that they contribute to *Zfhx1b* expression in the developing subpallium. These enhancer elements are activated by DLX2 in luciferase reporter assays, and are bound by DLX2 *in vivo*, providing strong evidence that subpallial *Zfhx1b* expression directly depends on *Dlx1/2*. Consistent with this, we observed marked alterations of *Zfhx1b* expression in the subpallial SVZ of *Dlx1/2*-deficient mice. Taken together, these results raise the possibility that the loss of *Zfhx1b* expression in the SVZ could contribute to the defects in differentiation and interneuron migration seen in *Dlx1/2* mutants. To investigate this possibility, we compared the phenotypes of the conditional *Zfhx1b* and *Dlx1/2*^{-/-} mutants.

***Dlx1/2*^{-/-} Constitutive and *Zfhx1b* Conditional Mutants Have Similar Changes in Gene Expression Related to Their Defects in Interneuron Migration**

As *Zfhx1b* expression in the subpallial SVZ was greatly reduced in the *Dlx1/2* mutants (Figures 5A-B'), and because interneuron migration to the cortex was greatly reduced in both the *Dlx1/2*^{-/-} and *Zfhx1b* mutants, we hypothesized that loss of *Zfhx1b* may underlie some of the interneuron migration phenotype of the *Dlx1/2*^{-/-} mutants. To evaluate this idea, we compared gene expression phenotypes of the *Dlx1/2*^{-/-} constitutive null mutant with the *Zfhx1b* conditional (*Nkx2.1-Cre*) mutant at E15.5 (Figure 6).

Indeed, changes in *Nkx2-1* and *Sox6* expression were similar in the *Dlx1/2*^{-/-} and *Zfhx1b* mutants. *Nkx2-1* and *Sox6* RNAs are normally expressed in similar patterns at E15.5 (Figures 6A,B,C,G,H and I); in wild type brains, both RNAs are maintained in the MGE, and in putative striatal interneurons and globus pallidus neurons, while they are downregulated in the MGE-derived cortical interneuron lineage (*Nkx2.1*⁻, *Sox6*^{low}). Both *Dlx1/2*^{-/-} and *Zfhx1b* mutants show increased numbers of *Nkx2-1*⁺ and *Sox6*⁺ cells in the striatum/LGE (asterisks in Figure 6; note: the *Dlx1/2*^{-/-} mutant striatum is small due to *Dlx*-function in the LGE) (Anderson et al., 1997b; Yun et al., 2002b).

Lhx6⁺ cells are lacking throughout the cortex and increased in the striatum of both the *Dlx1/2*^{-/-} and *Zfhx1b* mutants (Figures 6D-F'). Likewise, *Sst*⁺ and *NPY*⁺ cortical interneurons were lost, whereas their expression was increased in the striatum (Figures 6J-L''; *Sst* data not shown).

RNA Expression Array Analysis Identifies Candidate Mediators of *Zfhx1b* Function

Towards identifying the molecular mechanisms underlying the *Zfhx1b* mutant phenotype we used an RNA expression microarray analysis. We compared gene expression from the E12.5 MGE of *Nkx2.1-Cre; Zfhx1b^{F/-}* mutants to that of *Nkx2.1-Cre; Zfhx1b^{F/+}* control littermates. Table S1 (see Table S3 for an extended version) lists the most highly up-regulated and down-regulated genes. Overall, a larger number of genes were found to be significantly up-regulated than down-regulated in *Zfhx1b* mutants, which may reflect *Zfhx1b*'s function as a recruiter of repressive transcriptional complexes (van Grunsven et al., 2003; Verschuere et al., 1999; Verstappen et al., 2008). We verified the results for many of the genes by performing in situ RNA hybridization on E12.5 control and mutant brains (Figure S6 and data not shown).

We were most interested in genes that were altered in both the *Nkx2.1-Cre* and *Dlx1/2b-Cre* *Zfhx1b* mutants, given that both mutants showed altered interneuron migration and specification. Six genes fell into this category: *cMaf*, *MafB*, *CXCR7*, *Dlk1*, *Cited1* and *Gpc4* (Figures 7, S6). *Dlk1*, *Cited1* and *Gpc4* were up-regulated in the MGE both mutants (Figure S6). *cMaf*, *MafB*, and *CXCR7* were down-regulated in the MGE and migrating interneurons; later in the paper we focused more on these genes (Figures 7, S6); below we discuss the other genes.

Dlk1 expression was strongly increased in the VZ and SVZ of the MGE in the *Nkx2.1-Cre* mutant, and increased weakly only in the SVZ of the MGE in the *Dlx1/2b-Cre* mutant. (Figures S6J-S6L' and S6HH-S6JJ"). Given that *Zfhx1b*'s function was required in the SVZ of the MGE, the increase in *Dlk1* expression could play a role in the phenotype. *Dlk1* encodes a secreted delta-like ligand (Ferrón et al., 2011; Moon et al., 2002) that could alter Notch signaling. We used electroporation to increase *Dlk1* expression in wild type MGE, but failed to identify a change in interneuron migration (data not shown).

Other genes related to Notch-signaling were also identified in the array analysis, including the *Id2* and *Id4* helix-loop-helix and *Sox6* HMG-box transcription factors (Table S1). Expression of *Id4* was increased in the VZ of the *Nkx2.1-Cre* mutant; however, no change in expression was detected in the *Dlx1/2b-Cre* mutant (Figures S6M-S6O' and S6KK-S6MM"); this implies that *Id4* does not contribute to the interneuron phenotype. *Id2* expression showed a subtle expression increase in the *Nkx2.1-Cre* mutant (not shown); like *Id4*, we did not find a change in its expression in the *Dlx1/2b-Cre* mutant (not shown). *Sox6* expression was also increased based on the array and an increase was seen in both *Nkx2.1-Cre* and *Dlx1/2b-Cre* mutants by in situ hybridization at E12.5 (Table S1 and Figures S1F-S1F' and data not shown), which became more pronounced at E13.5 and E15.5 (Figures 2J-L' and 3G-I'). Of note, *Sox6* represses MGE expression of *Ascl1* (*Mash1*) (Azim et al., 2009), a basic-helix-loop-helix transcription factor whose expression is promoted by Notch-signaling.

Expression of *Cited1*, a p300-binding transcriptional co-activator that promotes signaling in the TGF-beta pathway (Gerstner and Landry, 2007) was increased in the SVZ of the ventral MGE and POA in both the *Nkx2.1-Cre* and *Dlx1/2b-Cre* mutants (Figures S5S-S5U' and S5QQ-S5SS'). This is of interest given that ZFHXB acts as a SMAD-binding transcriptional co-repressor (Vandewalle et al., 2008). Thus, *Cited1* and *Zfhx1b* may function antagonistically in MGE development. *Gpc4* expression in the SVZ of the MGE was increased in both mutants (Figure S5V-S5X' and S5TT-S5WW'). Glypicans (GPC) are extracellular matrix proteins that promote FGF-signaling (Jen et al., 2009).

Expression of genes related to oligodendrogenesis, including *Olig1* and *GPR17* (Chen et al., 2009; Lu et al., 2000) were down-regulated on the array in the *Nkx2.1-Cre* mutant (Table

S1); we failed to detect *GPR17* expression by in situ hybridization and *Olg1* expression was weak. For this reason, we studied *Olg2* expression; its expression was reduced in the SVZ of the MGE at E12.5 (Figures S5P-S5R'). The down-regulation of oligodendrocyte markers may be related to the increase in *ID4* RNA; ID proteins can repress oligodendrogenesis (Wang et al., 2001). By E15.5, we did not detect a change in *Olg2* expression (Figures S2P-S2R'). The *Dlx112b-Cre* mutant did not show changes in *Olg2* expression (Figure S5NN-S5PP') suggesting the *Zfhx1b* function in the VZ, and not SVZ, regulates oligodendrogenesis.

Zfhx1b* Is Required for Expression of Genetic Markers of Cortical Interneurons: *cMaf*, *MafB* and *Cxcr7

The gene expression array showed a ~3-fold reduction in the expression of *cMaf* (*v-Maf*), a leucine zipper-containing transcription factor (Table S1). *cMaf*, and its relative *MafB*, have been reported to be expressed in cortical interneurons (Cobos, 2006; Faux et al., 2009; Zhao et al., 2008). Likewise, the array identified reduced expression of *CXCR7*, whose expression and function are required during cortical interneuron migration (Sánchez-Alcañiz et al., 2011; Wang et al., 2011).

We compared *cMaf*, *MafB* and *Cxcr7* RNA expression at E12.5 and E15.5 and identified some important features (Figures 7, S2A-S2C and S6D-S6F). *cMaf*, *MafB* and *Cxcr7* RNAs were expressed in the SVZ of the dorsal MGE (and not the ventral MGE), and were maintained in cells migrating through the LGE and CGE and then into the cortex at E12.5 and E15.5 (Figure 7, S2A-S2C and S6D-S6F). *cMaf*, *MafB* and *CXCR7* appear to be excellent markers of the cortical interneuron lineage, as we did not detect their expression in other MGE-derived structures, such as the ventral pallidum or globus pallidus at E12.5, E15.5 or P0 (Figures 7, S2A-S2C, S6D-S6F and data not shown). In the E15.5 and P0 striatum, these genes showed little expression (Figure 7), except for *MafB* and *cMaf* in a very small population of cells (data not shown). Thus, unlike other cortical interneuron markers that are also expressed in striatal interneurons (e.g. *Dlx1*, *Lhx6*, *parvalbumin*, *Sst*), *cMaf*, *MafB* and *CXCR7* expression largely mark only cortical interneurons.

We then compared *cMaf*, *MafB* and *Cxcr7* expression in E12.5 and E15.5 control (*Zfhx1b* heterozygotes), *Zfhx1b* conditional mutants (*Nkx2.1-Cre* and *Dlx112b-Cre*) and the *Dlx1/2^{-/-}* constitutive mutant (Figures 7, S2A-S2C'', S3A-S3I'). *cMaf* expression was nearly eliminated in all three mutants. Much of the remaining *cMaf* expression was in scattered blood cells and in the choroid plexus (Figure 7 H' and H''). *MafB* and *Cxcr7* were also greatly reduced in the SVZ of the ganglionic eminences, although they were not as strongly down-regulated as *cMaf* (Figures 7, S2A-S2C', S3D-S3I', S6D-S6I' and S6BB-S6GG'). Therefore, *Zfhx1b* (and *Dlx1&2*) were required for *cMaf*, *MafB* and *Cxcr7* expression, which are highly specific markers of immature migrating cortical interneurons (*cMaf* and *MafB* are specific for MGE-derived interneurons). Thus, the loss of *cMaf*, *MafB* and *Cxcr7* expression in *Zfhx1b* conditional mutants (*Nkx2.1-Cre* and *Dlx112b-Cre*) provides additional evidence that cortical interneurons fail to be specified.

Next, we determined at a cellular resolution when *cMaf* expression begins in developing cortical interneurons (Figures 7M-M''). Given that *cMaf* expression in the dorsal MGE and migrating cortical interneurons is dependent on *Zfhx1b* expression, and that repression of *Nkx2-1* in these regions is also *Zfhx1b* dependent, we analyzed whether or not *cMaf* and *Nkx2-1* are co-expressed in these cells. To this end, we performed a triple-labeling analysis: fluorescent in situ hybridization to detect *cMaf* in combination with immunofluorescence to label NKX2-1 and the EGFP⁺ *Nkx2.1-Cre* lineage (EGFP expression from *CAG-CAT-EGFP*, the Cre reporter allele). We found that *cMaf* RNA was expressed in EGFP⁺ cells derived from the MGE lineage that were migrating through the LGE corridor (LGE-Co) on

route to the cortex (Figures 7M', M'' and M'''). These cells were NKX2-1⁻. On the other hand, NKX2-1 was expressed in *cMaf* cells of the globus pallidus (Figures 7 M, M'' and M'''). Thus, as NKX2-1 expression is repressed in immature cortical interneurons, *cMaf* expression begins.

Discussion

Herein we demonstrate that *Zfhx1b* subpallial expression is directly positively regulated by *Dlx1&2*, and is required in the MGE to generate cortical interneurons that express *Cxcr7*, *MafB* and *cMaf*. In its absence, *Nkx2-1* expression is not repressed, and cells that ordinarily would become cortical interneurons are transformed towards the *NPY/nNos/Sst* subtype of striatal GABAergic interneuron. Furthermore, it is possible that the *Zfhx1b*^{-/-} phenotype is also caused by defects in migration and differentiation that contribute to the formation of subpallial ectopia. However, below we largely concentrate on discussing the evidence that *Zfhx1b* regulates cell-type specification.

Zfhx1b regulates MGE cell-type generation

The MGE generates multiple cell types, including GABAergic interneurons of the cortex and striatum, GABAergic projection neurons of the basal ganglia (e.g. GP), cholinergic neurons of the striatum and basal telencephalon and oligodendrocytes (Flandin et al., 2010; Petryniak et al., 2007; Xu et al., 2008). The MGE generates roughly 60% of all GABAergic cortical interneurons; these express *PV*, *Sst*, *NPY* and *nNos* (Gelman and Marín, 2010; Rudy et al., 2011).

The MGE also generates striatal interneurons. There are three subtypes of GABAergic striatal interneurons: *PV*⁺, *nNos/NPY/Sst*⁺ and *CR*⁺ (Tepper et al., 2010). The striatum also has cholinergic interneurons. The cholinergic population is marked and regulated by *Gbx2*, *Islet1*, and *Lhx8* (Chen et al., 2010; Fragkouli et al., 2009), and the neurotrophin receptor TrkA (Sanchez-Ortiz et al., 2012).

The ventral MGE is not a major source for cortical interneurons based on fate mapping using *Shh-Cre* (Flandin et al., 2010). Thus, the dorsal MGE must be the source of most MGE-derived cortical and striatal GABAergic interneurons. It is poorly understood whether these cell types are generated from distinct subregions, from distinct but intermixed progenitors, or from the same progenitors in a stochastic or temporally modulated program. While the same neuroepithelial progenitor can generate different types of cortical interneurons (*PV* and *Sst*) (Brown et al., 2011), it is not known whether cortical and striatal interneurons are derived from the same progenitor.

Zfhx1b was required to generate GABAergic cells that migrate to the cortex, and to repress the generation GABAergic cells that migrate to the striatum. We suggest that *Zfhx1b* promotes a fate switch between cortical interneurons and *nNos/NPY/Sst* striatal interneurons through repression of *Nkx2-1* expression. Furthermore, *Zfhx1b* mutants have reduced striatal *PV* interneurons (Figure 4M, 4M'); thus, *Zfhx1b* could also control this fate decision. *Zfhx1b* is required in the MGE SVZ, and not the VZ, to promote the specification of pallial interneurons, as we observed largely the same phenotype using *Dlx12b-Cre* (SVZ recombination, Figures 3, S1, S3) and *Nkx2.1-Cre* (VZ recombination; Figures 1, 2, S1, S2).

We identified perhaps the first specific early marker of dorsal MGE-derived cortical interneurons: *cMaf* (Figure 7). *cMaf* and *MafB* expression are dependent on *Zfhx1b* and *Dlx1/2* function (Figure 7, S2A-C', S3D-F', S6D-S6F' and S6BB-S6DD' (Cobos, 2006; Long et al., 2009a; Long et al., 2009b) Notably, neither *cMaf*, nor *MafB* are strongly expressed prenatally in neurons of the striatum (interneurons and medium spiny neurons),

suggesting that prenatally they may be specific markers of the cortical interneuron lineage. Currently, *Maf* function in the brain has only been studied in the hindbrain (Cordes and Barsh, 1994).

***Zfhx1b* connects the *Nkx2-1* and *Dlx* transcription pathways**

There is genetic evidence for at least three parallel (although interacting) transcriptional pathways in the MGE that are required for cortical interneuron development: the 1) *Ascl1* (*Mash1*); 2) *Dlx*; and 3) *Nkx2-1* pathways (Long et al., 2009a,b). The *Nkx2-1* pathway is the core mediator of MGE regional and cell identity (Butt et al., 2007; Sussel et al., 1999); it functions through induction of *Lhx6* and *Lhx8* (Sussel et al., 1999). *Lhx6* is essential for induction of *Mafb* and *Shh* in neurons, maintenance of *Sox6* in interneurons, and the differentiation of Sst and Parvalbumin cortical interneurons (*Lhx6*) (Liodis et al., 2007; Zhao et al., 2008); *Lhx8* is required in cholinergic striatal interneurons (*Lhx8*) (Fragkouli et al., 2009).

Only a subset of MGE neuronal derivatives maintain *Nkx2-1* and *Lhx8* expression, such as the globus pallidus and cholinergic striatal interneurons (Marin et al., 2000), whereas MGE-derived cortical interneurons suppress *Nkx2-1* and *Lhx8* expression (Nóbrega-Pereira et al., 2008). We propose that *Dlx* and *Nkx2-1* pathways interact at this step. We demonstrated that *Dlx1/2* were required for *Zfhx1b* expression in the subpallial SVZ (Figure 4), and that *Zfhx1b* was required for repression of *Nkx2-1*, but not of *Lhx8* (Figures 2D-F', 2M-O', 3A-C' and 3J-L'). Thus, in the absence of *Zfhx1b*, dorsal MGE-derived neurons continued to express *Nkx2-1*, *Sox6* and *Lhx6*, and migrate into the striatum and not the cortex. These cells failed to express markers of cortical interneurons (*Cxcr7*, *cMaf* and *MafB*) (Figures 7 A-L', S5A-S5I'), but highly expressed the striatal GABAergic subtype markers *NPY*, *nNos* and *Sst* (Figures 2P-R' and 3M-O' (Tepper et al., 2010). Additionally, there was increased expression of TacR1, which is robustly expressed in Sst⁺ and ChAT⁺ striatal interneurons, and in very few cortical interneurons (Ardelt et al., 1996; Figure S4D-S4F').

Finally, *Zfhx1b* mutants did not exhibit clear phenotypes of striatal cholinergic interneurons or the GP. Thus, we propose a distinct *Zfhx1b*-independent mechanism for the generation of the GP and cholinergic neurons; the latter depends on the maintenance of *Lhx8*, perhaps in combination with *Islet1* and *Gbx2* (Chen et al., 2010; Fragkouli et al., 2009).

Downstream of *Zfhx1b* in the MGE Cells

It is unclear whether *Zfhx1b* has a common molecular mechanism in all developing cells. *Zfhx1b* mediates some of its functions through interactions with SMAD proteins, and thus participates in TGF-beta signaling (Vandewalle et al., 2008). Expression of the SMAD-binding transcriptional co-activator *Cited1* was increased in *Zfhx1b* mutants (Sup. Figure 6). The link to SMAD signaling is intriguing because SMAD dominant negative expression can inhibit interneuron tangential migration (Maira et al., 2010).

In the pallium, *Zfhx1b* functions in both progenitors and neurons. In hippocampal progenitors, it functions upstream of Wnt signaling to control development of the entire region (Miquelajauregui et al., 2007). In neocortical neurons *Zfhx1b* regulates *neurotrophin-3* and *Fgf9* expression, to control cortical progenitors (Seuntjens et al., 2009). We did not observe similar regulatory changes in the *Zfhx1b* mutant MGE.

While *Zfhx1b* in the MGE SVZ regulates the switch between cortical and striatal interneurons, *Zfhx1b* is also expressed in the VZ of the MGE (Figure 1A). Two genes related to Notch signaling were up-regulated in the mutant MGE VZ, including the secreted delta-like ligand *Dlk1* (Ferrón et al., 2011; Moon et al., 2002) and the HLH transcription

factor *ID4* (Yun et al., 2004) (Figure S6J-S6O'). *Dlk1* up-regulation in the VZ and SVZ could alter the balance of cell fate decisions.

Previous studies suggested that *Nkx2-1* promotes interneuron integration into the striatum via repression of *Npn2*/*Sema3*-dependent repulsion (Marín et al., 2001; Nóbrega-Pereira et al., 2008). We did not detect a change in *Npn2* and *Npn1* RNA expression in migrating immature *Zfhx1b* mutant interneurons at E12.5. On the other hand, van den Berghe et al (in press) present evidence that *Zfhx1b* regulates interneuron migration through the *netrin* receptor *Unc5b*.

Zfhx1b and Human Disease

Mowat-Wilson syndrome (MWS) is caused by a heterozygous mutation or deletion of the *Zfhx1b* (*ZEB2*, *SIP1*), and is characterized by a distinctive facial appearance, intellectual disability, and variable other features including seizures, agenesis of the corpus callosum, and Hirschsprung disease (Mowat et al., 2003). Given *Zfhx1b*'s critical role in cortical interneuron development, we propose that cortical interneuron defects contribute to the seizure phenotype of MWS. Furthermore, since *Dlx1&2* regulate *Zfhx1b* expression in the subpallium, and *Dlx1&2* also regulate craniofacial and enteric nervous system development (Qiu et al., 1995) it will be intriguing whether *Zfhx1b* is also downstream of *Dlx* function during development of these tissues.

Experimental Procedures

See Supplemental Experimental Procedures for detailed description of methods.

Mice

Zfhx1b^{F/F} mice were genotyped according to (Miyoshi et al., 2006). *CAG-CAT-eGFP* mice were genotyped according to (Kawamoto et al., 2000). *Zfhx1b^{F/F}* males were crossed to *Beta-Actin Cre* mice (Lewandoski et al., 1997) to generate the *Zfhx1b* null allele, which was followed by a cross to wildtype mice to eliminate the *Beta-Actin Cre* allele. *Zfhx1b^{+/-}* mice were crossed with *Nkx2.1-Cre II/2b-Cre* mice, and male *Zfhx1b^{+/-}*; *Cre⁺* mice were crossed with female *Zfhx1b^{F/F}* mice with or without the *CAG-CAT-EGFP* allele to generate conditional mutant embryos.

Histochemistry

Embryonic and postnatal brains were prepared and immunostained (Flandin et al., 2010), or assayed by in situ hybridization (Jeong et al., 2008). Protocols can be found on our lab website <http://physio.ucsf.edu/rubenstein/protocols/index.asp>, with modifications for dual immuno/in situ fluorescence analysis described in the Supplementary Experimental Procedures.

Cell culture, transfections and luciferase assays

P19 cells were cultured as described in (Farah et al., 2000). Experimental conditions were tested in triplicate by transfection of cells in 12-well plates using Fugene 6 (Roche). Cotransfection of a Renilla luciferase expression construct was used as a normalization control for a dual-luciferase assay. The following amounts of DNA were used in each well: 80ng *pGL4.73* (Renilla Luciferase, Promega), 240 ng *pCAGGS-empty* or *pCAGGS-Dlx2*, 240ng *pGL4.23-empty* (Luciferase, Promega) or *pGL4.23-enhancer*. Luciferase and Renilla Luciferase quantification was done using a Promega Dual-Luciferase Assay Kit and a microplate luminometer (Veritas). Chi-square test showed that the levels of activation were significant *: $p < 0.05$.

Chromatin immunoprecipitation (ChIP)

ChIP was performed similar to a published method (McKenna et al., 2011) with modifications described in Supplementary Experimental Procedures.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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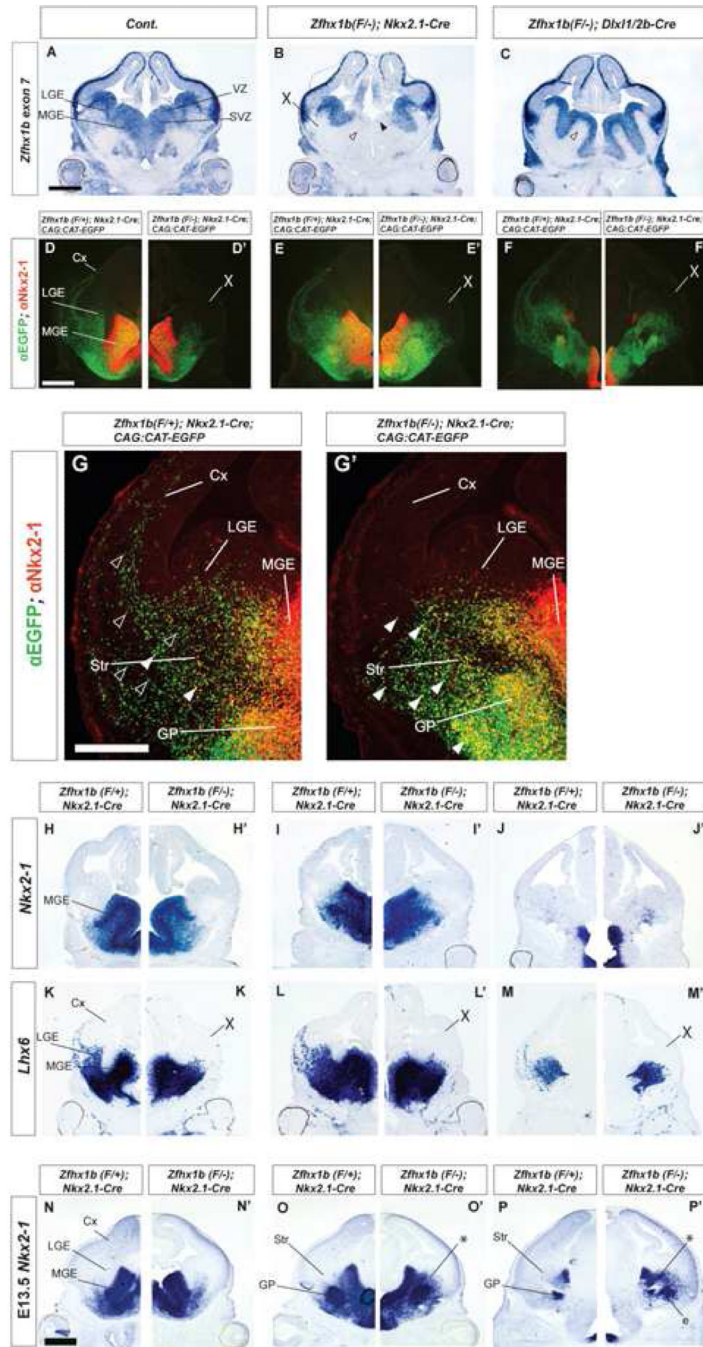


Figure 1.

Zfhx1b expression in the MGE is required for interneuron migration at E12.5. (A-C) *Zfhx1b* RNA expression detected by in situ hybridization in control and conditional *Zfhx1b* mutant telencephalons. (B) *Nkx2.1-Cre*. (C) *Dlx1/2b-Cre*. Black arrowhead in B shows loss of *Zfhx1b* expression in the MGE VZ (except dorsal-most MGE). White arrowheads in B and C show loss of *Zfhx1b* expression in the SVZ of the MGE. X in B shows loss of *Zfhx1b* expression in the SVZ/MZ of the LGE. (D-P') Coronal hemisections of the telencephalon comparing gene expression in three rostral-to-caudal planes of section in control (left side) and *Zfhx1b* *Nkx2.1-Cre* conditional mutants (right side). (D-F') Two color immunofluorescence detection of EGFP (green) and NKX2-1 (red). (G,G') Higher

magnification view of two color immunofluorescence detection of EGFP (green) and Nkx2-1 (red) in control (G) and *Zfhx1b* mutant (G'). Solid white arrowheads show increased numbers of cells that express both EGFP (green) and NKX2-1 (red) in the mutant's LGE/Str. In the wild type cortex, black arrowheads (with white outline) show that cells express EGFP (green) and not NKX2-1. (H-P') In situ hybridization expression analysis at E12.5 of *Nkx2-1* (H-J') and *Lhx6* (K-M'), and at E13.5 of *Nkx2-1* (N-P'). Asterisks in (N-P') show increased numbers of labeled cells in striatum. X in panels (K'-M') notes the loss of labeled cells in the cortex. Abbreviations: Cx: cortex; e: ectopia in region of the ventral striatum and central nucleus of the amygdala; GP: globus pallidus; LGE: lateral ganglionic eminence; MGE: medial ganglionic eminence; MZ: mantle zone; Str: Striatum; SVZ: subventricular zone; Vpd: ventral pallidum; VZ: ventricular zone. Scale bars equal 500 μ m (A and D), and 300 μ m (G).

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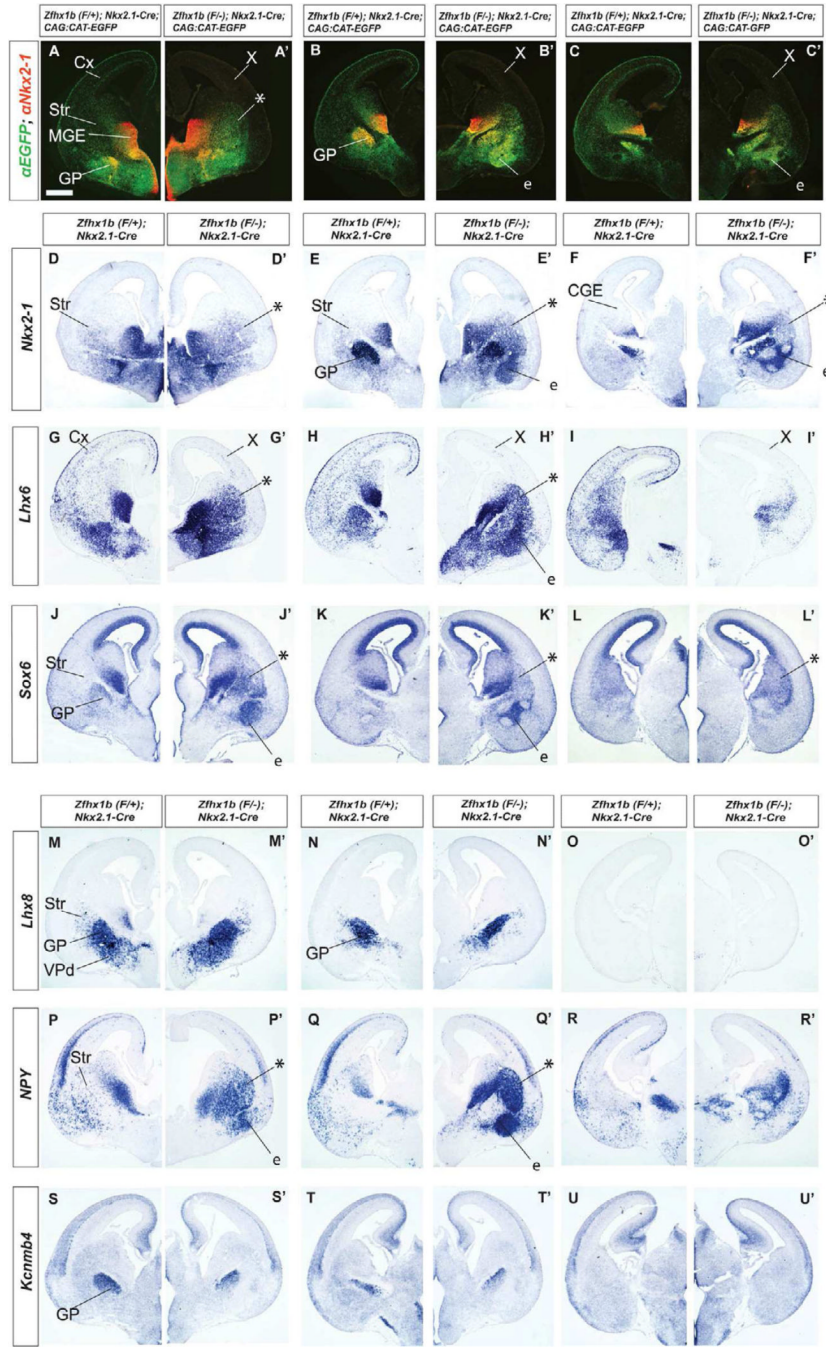


Figure 2. *Zfhx1b* expression in the MGE is required for interneuron migration at E15.5. Coronal hemisections of the telencephalon comparing gene expression in three rostral-to-caudal planes of section in control (left side) and *Zfhx1b*;*Nkx2.1-Cre* conditional mutants (right side). (A-C') Two color immunofluorescence detection of EGFP (green) and NKX2-1 (red). (D-U') In situ hybridization analysis. *Nkx2-1* (D-F'), *Lhx6* (G-I'), *Sox6* (J-L'), *Lhx8* (M-O'), *NPY* (P-R'), *Kcnmb4* (S-U'). Asterisks show increased numbers of labeled cells in the striatum. X shows reduced number of labeled cells in cortex. Abbreviations: CGE: caudal ganglionic eminence; Cx: cortex; e: ectopia in region of the ventral striatum and central nucleus of the amygdala; GP: globus pallidus; LGE: lateral ganglionic eminence; MGE:

medial ganglionic eminence; MZ: mantle zone; Str: striatum; SVZ: subventricular zone; VPd: ventral pallidum; VZ: ventricular zone. Scale bar equals 500 μ m (A).

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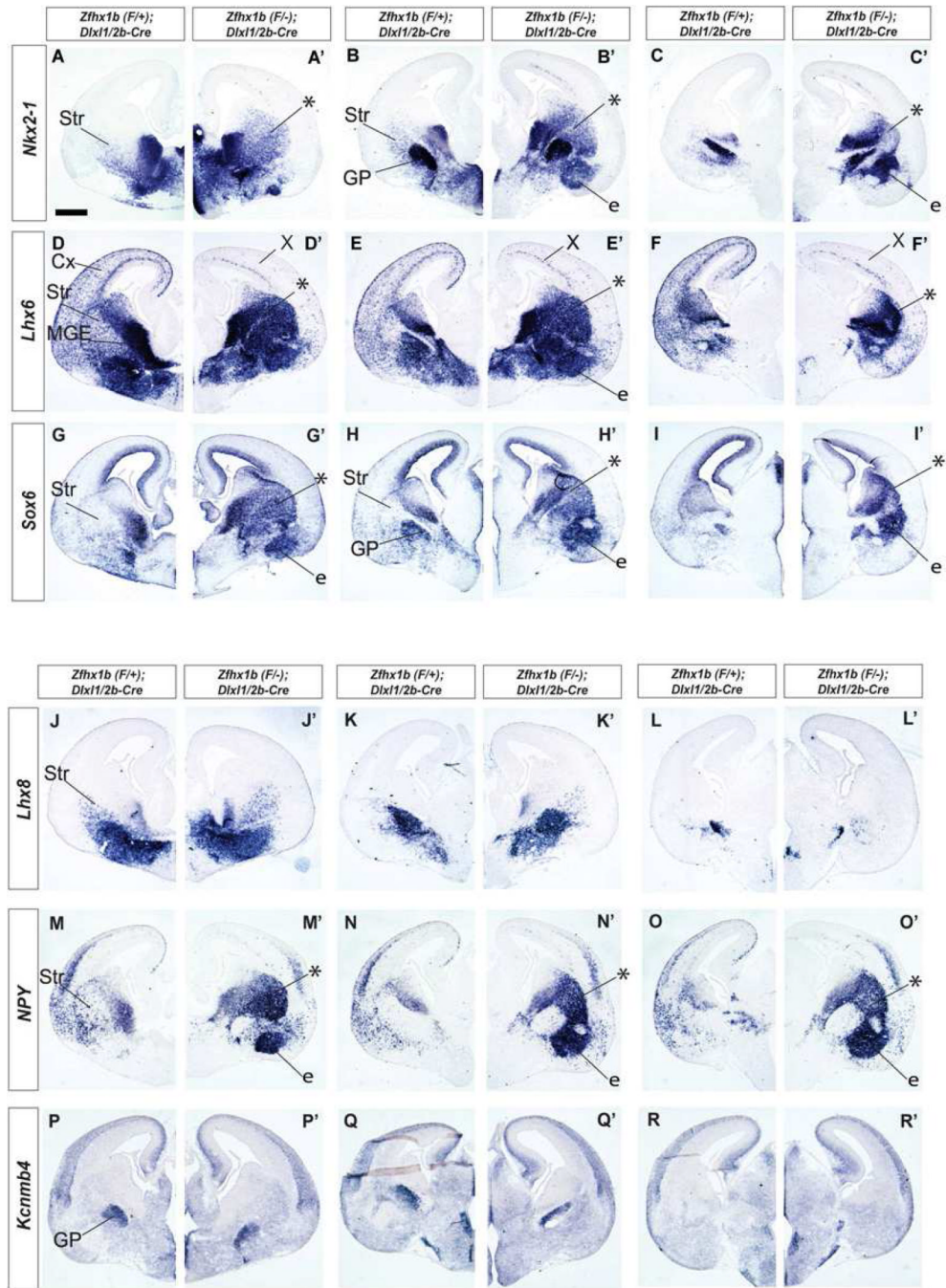


Figure 3.

Zfhx1b expression in the SVZ of the MGE is required for interneuron migration at E15.5. Coronal hemisections of the telencephalon comparing gene expression in three rostral-to-caudal planes of section in control (left side) and *Zfhx1b Dlx1/2b-Cre* conditional mutants (right side). In situ hybridization analysis of: *Nkx2-1* (A-C'), *Lhx6* (D-F'), *Sox6* (G-I'), *Lhx8* (P-R'), *NPY* (M-O'), *Kcnmb4* (P-R'). Asterisks show increased numbers of labeled cells in the striatum. X shows reduced number of *Lhx6*⁺ cells in cortex. Abbreviations: CGE: caudal ganglionic eminence; Cx: cortex; e: ectopia in region of the ventral striatum and central nucleus of the amygdala; GP: globus pallidus; LGE: lateral ganglionic eminence;

MGE: medial ganglionic eminence; MZ: mantle zone; Str: striatum; SVZ: subventricular zone; VPd: ventral pallidum; VZ: ventricular zone. Scale bar equals 500 μ m (A).

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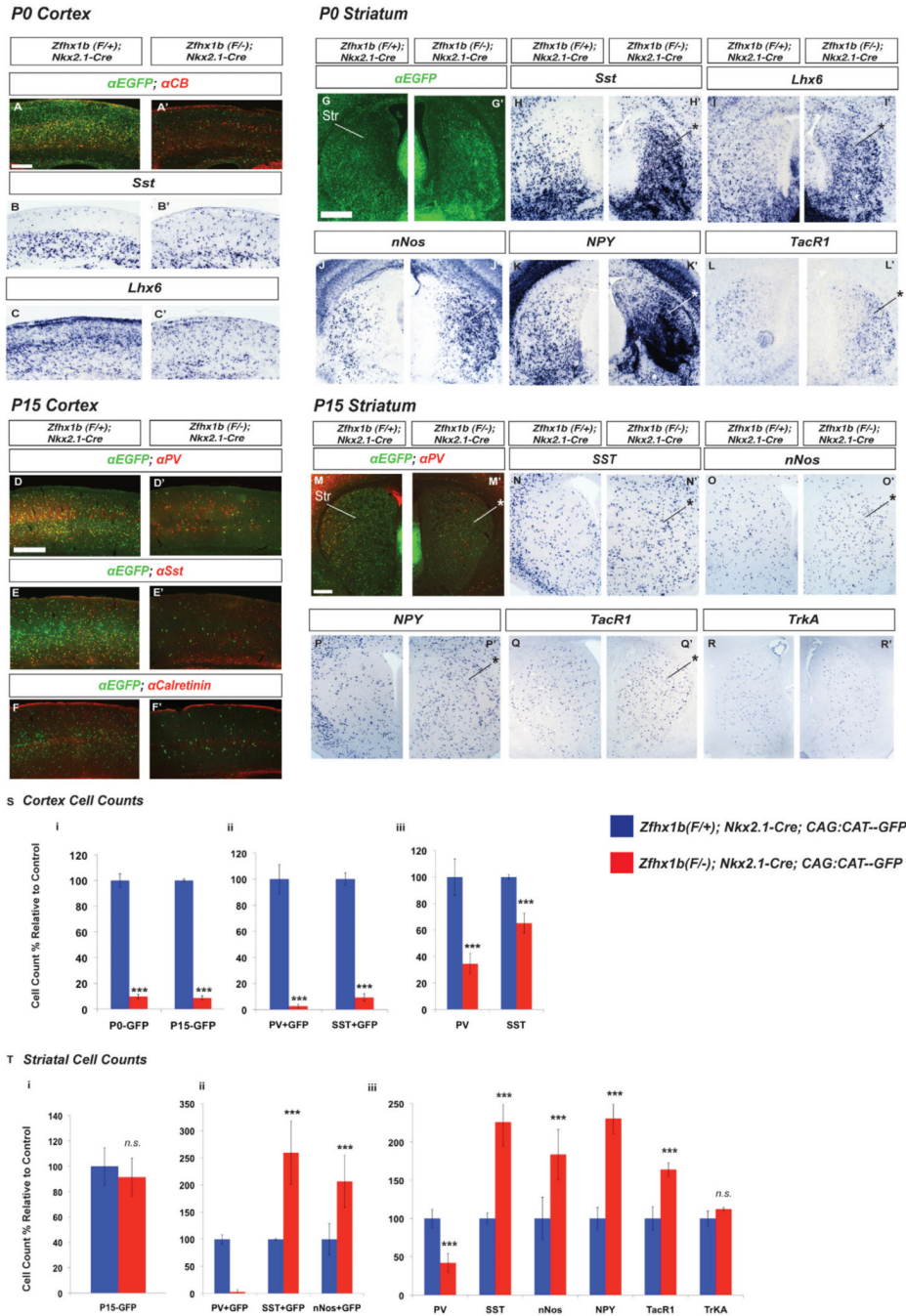


Figure 4. *Zfhx1b* expression in the MGE regulates the numbers and fate of postnatal (P0 and P15) cortical and striatal interneurons. Coronal hemisections showing the neocortex (A-F) and the striatum (G-R), comparing gene expression in control (left side) and *Zfhx1b*;*Nkx2.1-Cre* conditional mutants (right side). (A, A', D-F, G, G', M, M') Two color immunofluorescence with anti-EGFP (Cre reporter; green) and interneuron markers (red); other panels show in situ hybridization results. (S) Cell counts, control relative to mutant, of: total Cre-reporter EGFP⁺ cell numbers in the P0 and P15 cortex (left); of the number of cells that had colocalization of the EGFP Cre-reporter with Sst or PV at P15 (middle); of the total levels of Sst and PV in the P15 cortex (right). (T) Cell counts, control relative to mutant, of: EGFP⁺

Cre-reporter cells in the P15 striatum (left); of EGFP-colocalization with PV, Sst and nNos in the P15 striatum (middle); of the total numbers of the markers PV, Sst, nNos, NPY, TacR1 and TrkA in the P15 striatum (right). Scale bar equals 500 μ m (A,D, G and M). *: p<.05; **: p<.01.

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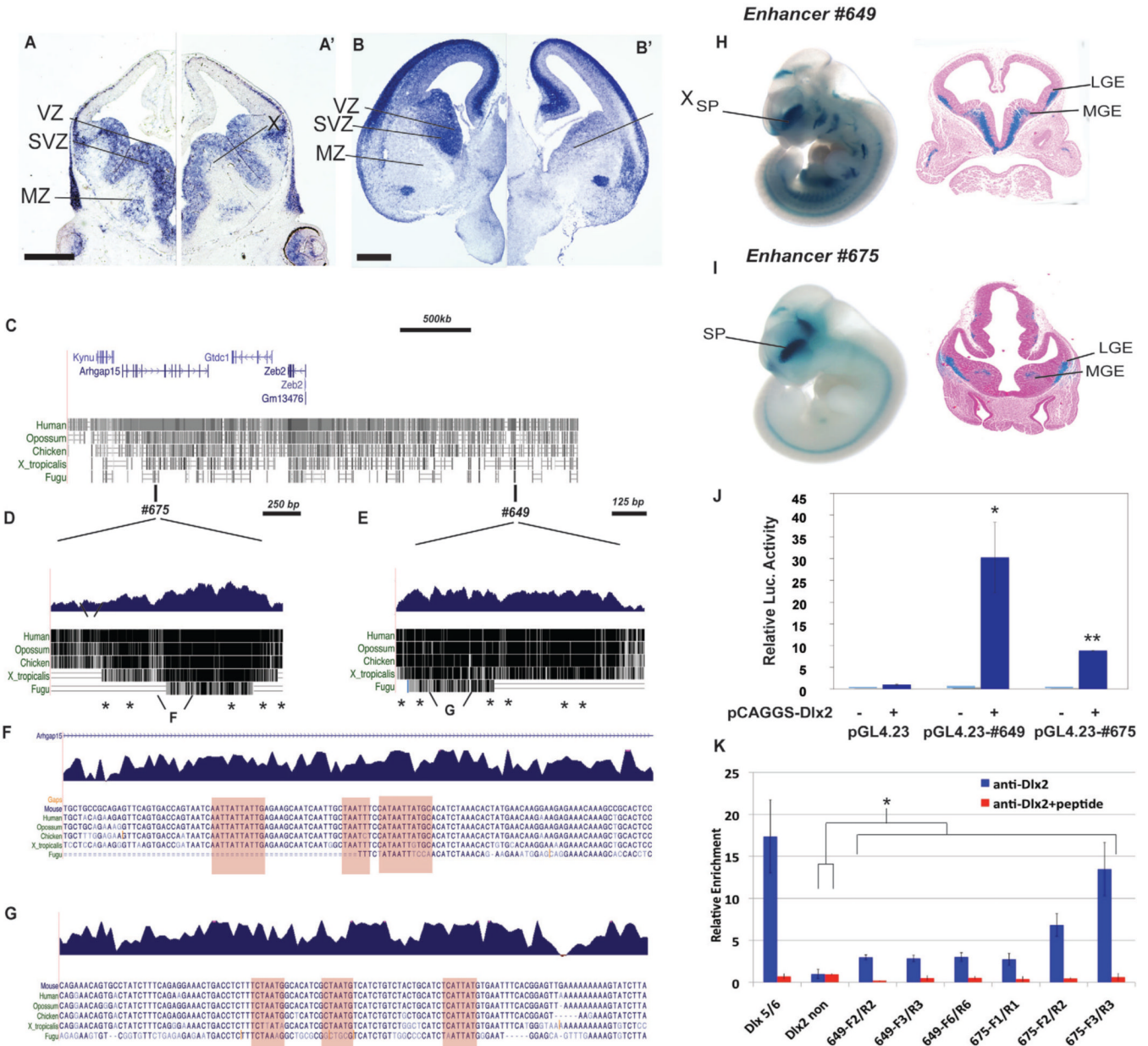


Figure 5. *Dlx1&2* are required for *Zfx1b* expression in the SVZ of the subpallium. (A-B') Coronal hemisections of the telencephalon comparing *Zfx1b* expression in *Dlx1*^{+/-} and *Dlx1*^{2/-} at E12.5 (A-A') and E15.5 (B,B'). Note the greatly reduced *Zfx1b* expression in the SVZ of the LGE and MGE. X notes reduction in *Zfx1b* expression in the SVZ. (C-K) Regulatory elements near *Zfx1b* that drive subpallial expression are bound by DLX2 *in vivo*, and are positively regulated by DLX2. (C) Relative genomic position of two ultraconserved DNA elements near the Human *Zfx1b* (*Zeb2*) locus, #675 (D) and #649 (E) (data from <http://genome.ucsc.edu/>). (D and E) Genomic alignment of enhancers #675 and #649; each contain a number of conserved consensus homeobox sites (asterisks). (F and G) Base-resolution view of regions with homeobox sites within #649 and #675, which are heavily conserved across vertebrate species and are similar to known DLX2 binding sites (Potter et al., 2008). (H and I) Whole mount E11.5 *enhancer-lacZ* transgenic mouse embryos that demonstrated

lacZ expression (X-Gal staining) in the ganglionic eminences (subpallium) (H' and I'). (J) Luciferase assay demonstrating DLX2-dependent transcriptional activation (*pCAGGS* vector) mediated by enhancers #649 and #675 upstream of luciferase (*pGL4.23* vector). (K) Blue bars: DLX2 ChIP qPCR assay (N=3) demonstrates anti-DLX2 binding to chromatin from E13.5 ganglionic eminences to subdomains of enhancers #675 and #649 and the positive control (Dlx5/6 enhancer), and not to the negative control region (a non-conserved domain upstream of *Dlx2*). Red bars: Addition of a DLX2 peptide blocks the anti-DLX2 binding. Abbreviations: Cx: cortex; LGE: lateral ganglionic eminence; Luc. Luciferase; MGE: medial ganglionic eminence; MZ: mantle zone; SP: subpallium; SVZ: subventricular zone, VZ: ventricular zone. Scale bar equals 500 μ m (A and B). ***: $p < .001$; n.s. : not significant.

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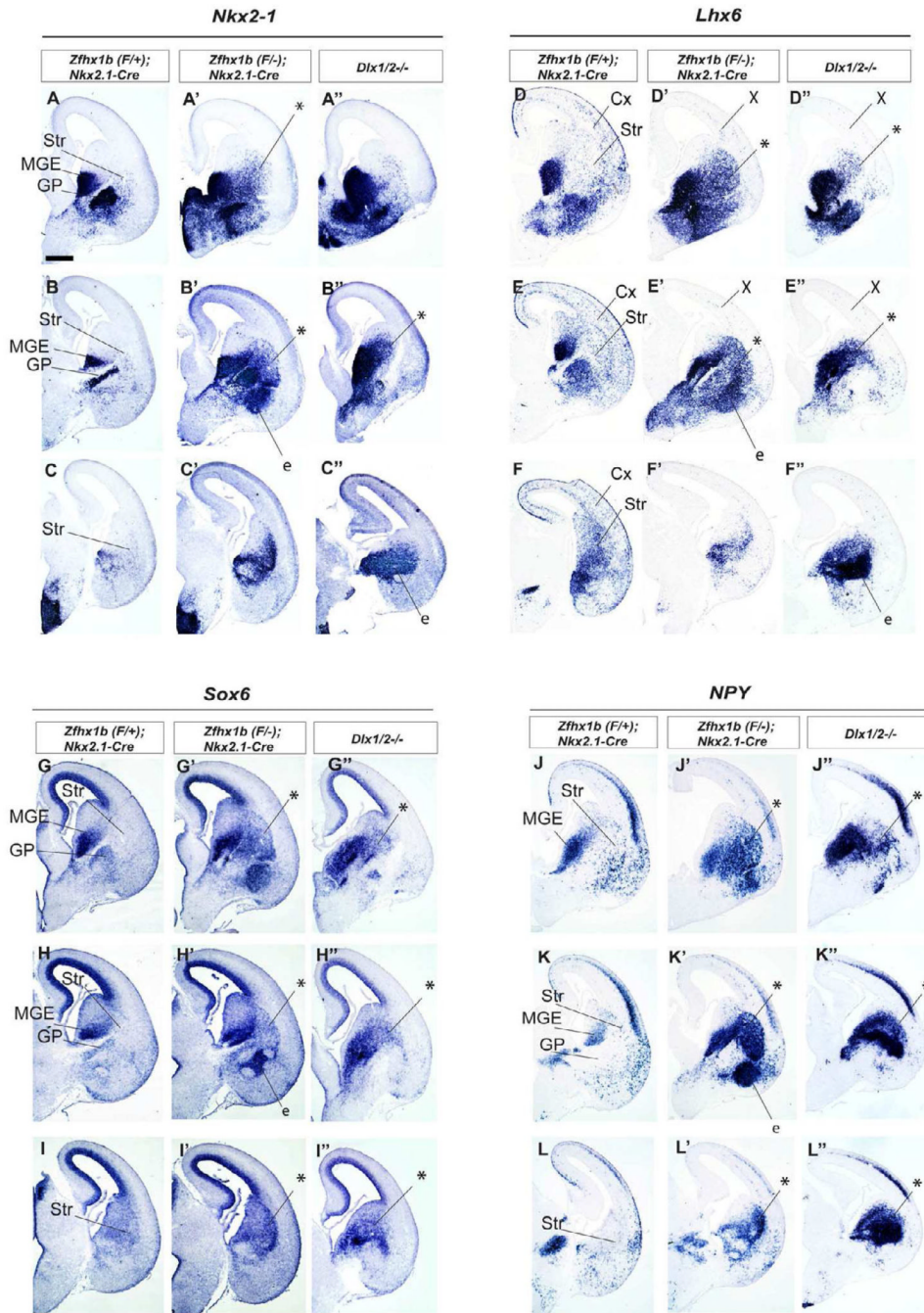


Figure 6. *Zfhx1b*;*Nkx2.1-Cre* and *Dlx1/2^{-/-}* mutants both fail to repress *Nkx2-1* and *Sox6*, lose cortical interneurons, and accumulate MGE cells in their striatum. Coronal hemisections of the E15.5 telencephalon comparing gene expression in three rostral-to-caudal planes of section in controls (left side) and mutants (right side). In situ hybridization analysis of *Nkx2-1* (A-C''), *Lhx6* (D-F''), *Sox6* (G-I''), *NPY* (J-L'') expression was assessed for control, *Zfhx1b*;*Nkx2.1-Cre* mutants and *Dlx1/2^{-/-}* mutants. Asterisks show increased numbers of labeled cells in the striatum. X shows reduced number of *Lhx6⁺* cells in cortex. Abbreviations: CGE: caudal ganglionic eminence; Cx: cortex; e: ectopia in region of the ventral striatum and central nucleus of the amygdala; GP: globus pallidus; LGE: lateral

ganglionic eminence; MGE: medial ganglionic eminence; MZ: mantle zone; Str: striatum; SVZ: subventricular zone; VPd: ventral pallidum; VZ: ventricular zone. Scale bar equals 500 μ m (A).

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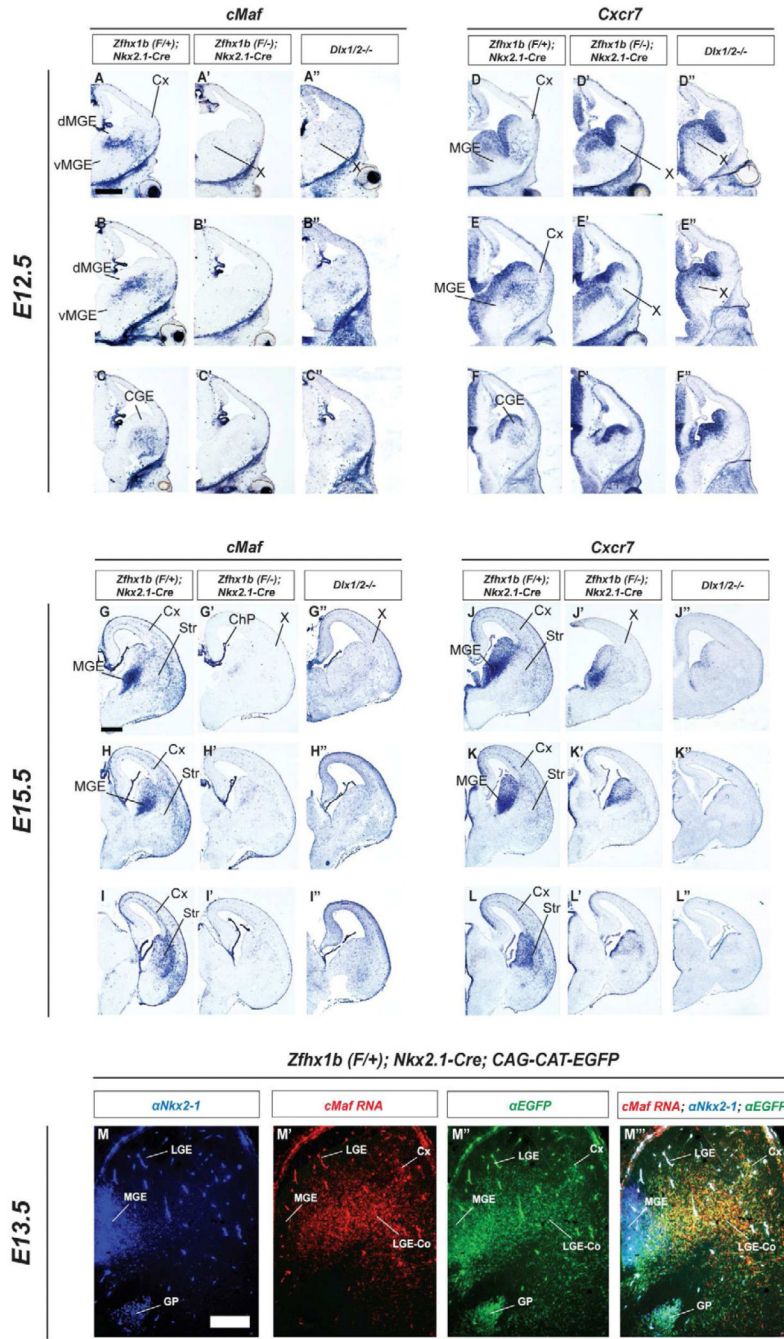


Figure 7. *cMaf* and *CXCR7* are highly specific markers of the cortical interneuron lineage that are lost in *Zfhx1b* conditional mutants and *Dlx1/2*^{-/-} mutants. Coronal hemisections of the E12.5 (A-F'') and E15.5 (G-L'') telencephalon comparing *cMaf* (A-C''; G-I'') and *CXCR7* (D-F''; J-L'') RNA expression by in situ hybridization in three rostral-to-caudal planes of section in control (left panels), *Zfhx1b*;*Nkx2.1-Cre* conditional mutants (middle panels), and *Dlx1/2*^{-/-} mutants (right panels). X shows reduced/absent *cMaf*⁺ or *CXCR7*⁺ cells in cortex or ganglionic eminences. Abbreviations: CGE: caudal ganglionic eminence; ChP: choroid plexus; Cx: cortex; LGE Co: LGE corridor; dMGE: dorsal medial ganglionic eminence;

vMGE: ventral medial ganglionic eminence; Str: striatum. Scale bars are equal to 500 μ m (A) and 200 μ m (M).

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