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Gene co-regulation by *Fezf2* selects neurotransmitter identity and connectivity of corticospinal neurons

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Abstract

The neocortex contains an unparalleled diversity of neuronal subtypes, each defined by distinct traits that are developmentally acquired under the control of subtype-specific and pan-neuronal genes. The regulatory logic that orchestrates the expression of these unique combinations of genes is unknown for any class of cortical neuron. Here, we report that *Fezf2* is a selector gene able to regulate the expression of gene sets that collectively define mouse corticospinal motor neurons

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AUTHOR CONTRIBUTIONS

S.L. and P.A. conceived the work, designed the experiments, analyzed the data and wrote the manuscript. S.L. performed the majority of the experiments. B.J.M. contributed to experimental design and performed microarray analysis. E.Z. performed the ChIP-seq experiments. L.A.G. analyzed the RNA-seq data. H.-H.C. performed the *in vitro* differentiation experiment and assisted in manuscript preparation. W.Y. performed the electrophoretic mobility shift assay experiment. A.M. assisted with FACS purification and the microarray experiments. E.T. performed the HARDI analysis. S.M. analyzed the ChIP-seq data. D.K.G. and J.L.R. supervised the bioinformatics analyses. P.A. supervised all aspects of the project.

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Note: Any Supplementary Information and Source Data files are available in the online version of the paper.

(CSMN). We find that *Fezf2* directly induces the glutamatergic identity of CSMN via activation of *Vglut1* (*Sc117a7*) and inhibits a GABAergic fate by repressing transcription of *Gad1*. In addition, we identify the axon guidance receptor *Ephb1* as a target of *Fezf2* necessary to execute the ipsilateral extension of the corticospinal tract. Our data indicate that co-regulated expression of neuron subtype-specific and pan-neuronal gene batteries by a single transcription factor is one component of the regulatory logic responsible for the establishment of CSMN identity.

During development, neurons of the CNS acquire distinct molecular, anatomical and physiological properties. Development of these traits is dictated by pan-neuronal and lineage-specific molecular cues, which in combination define neuronal subtype-specific identity and set the basis for neuronal diversity.

Among other regions of the mammalian CNS, the cerebral cortex stands as a prime example of how expanded neuronal diversity served the evolution of complex behavior. Here an extraordinary number of neuronal subtypes integrate into local and long-distance circuitry subserving sensory, motor and high-level cognitive functions¹. The majority of cortical neurons are excitatory projection neurons. Their classification is largely based on axonal projection targets, but it is clear that multiple traits—molecular identity, primary and collateral axonal connections, somatodendritic morphology and electrophysiological activity—must be taken into account to accurately distinguish individual projection neuron subtypes². The molecular regulatory logic that orchestrates acquisition of multiple class-specific features is not understood for any neuronal subtype of the cerebral cortex.

CSMN are one specialized class of cortical excitatory neurons, which connect layer Vb of the cortex to the spinal cord. At the molecular level, CSMN are the most extensively characterized class of cortical projection neurons^{3,4}. A master transcription factor—forebrain expressed zinc finger 2 (*Fezf2*)—has been identified that is necessary for the fate specification of CSMN^{5–7}. In the absence of *Fezf2*, CSMN fail to generate. In agreement, CSMN-specific genes are not expressed in layer Vb of the *Fezf2*^{-/-} null cortex, a deficiency accompanied by changes in dendritic morphology and a lack of axonal projections to the spinal cord^{5–7}.

Conversely, *Fezf2* alone can cell-autonomously instruct the acquisition of CSMN-specific features when expressed in diverse, permissive cellular contexts *in vivo*. Ectopic overexpression of *Fezf2* in progenitors and postmitotic neurons of different lineages is sufficient to induce the generation of neurons that express multiple CSMN genes and extend axons to subcerebral targets^{5,8–10}. Beyond acquisition of lineage-specific CSMN features, neurons expressing *Fezf2* also acquire necessary cortical pan-projection neuron features. They become glutamatergic and develop distinctive pyramidal morphology⁸. These data indicate that *Fezf2* is sufficient to instruct the acquisition of multiple aspects of CSMN identity.

Here we demonstrate that *Fezf2* is a selector gene for an individual class of neurons of the neocortex: corticospinal motor neurons. We found that activation and repression of large batteries of genes necessary to orchestrate the acquisition of CSMN defining traits was directly co-regulated downstream of *Fezf2*. *Fezf2* was sufficient to induce vesicular

glutamate transporter 1 (*Vglut1*) and to repress glutamate decarboxylase 1 (*Gad1*) expression to instruct the acquisition of glutamatergic identity. In addition, we uncovered a new role for EphB1 as a functionally relevant effector of *Fezf2*, required to execute the ipsilateral extension of the corticospinal tract through the ventral forebrain. Our data indicate a direct role for *Fezf2* as a selector gene that coordinates the expression of pan-neuronal and class-specific genes required for the developmental acquisition of CSMN identity.

RESULTS

Fezf2 induces signature genes of corticospinal neurons

To understand the molecular logic underlying the acquisition of CSMN traits upon *Fezf2* expression, we compared the *in vivo* gene expression of FACS-purified cortical progenitors that ectopically expressed *Fezf2* to control progenitors. We used *in utero* electroporation to deliver *Fezf2^{GFP}* or *Ctrl^{GFP}* expression vectors to neocortical progenitors at embryonic day (E) 14.5, when they primarily generate callosal projection neurons (CPN) of the upper layers (Supplementary Fig. 1a). Overexpression of *Fezf2* in these progenitors is sufficient to instruct a fate switch resulting in the generation of CSMN and other subtypes of corticofugal projection neurons^{5,7,11}. *Fezf2^{GFP}*- and *Ctrl^{GFP}*-electroporated progenitors were FACS-purified 24 and 48 h after surgery (Supplementary Fig. 1b,c) and acutely profiled by microarrays.

A large set of genes was induced by *Fezf2* as early as 24 h (263 genes; $P < 0.001$, fold change > 1.5), while others were induced not earlier than 48 h (441 genes; $P < 0.001$, fold change > 1.5) after electroporation. In addition, we identified a smaller set of genes repressed by *Fezf2* (90 genes at 24 h; 89 genes at 48 h; $P < 0.001$, fold change > 1.5) (Supplementary Table 1).

To investigate whether genes induced by *Fezf2* mark developing CSMN, we performed *in situ* hybridization for selected candidates (Supplementary Table 1) at multiple stages of embryonic and postnatal development of the cerebral cortex (E15.5, E18.5, postnatal day (P) 3, P7 and P14; Fig. 1 and Supplementary Fig. 2). Remarkably, we identified a series of *Fezf2* target genes (for example, *Ldb2*, *Adcyap1*, *Akap12*, *Ephb1*, *Rgs16*, *Rgs8*, *Acvr1c*, *Pappa2*, *Kif26a*) that labeled developing CSMN (and related subcerebral projection neurons, ScPN) in the early cortical plate (Fig. 1 and Supplementary Fig. 2), whereas other targets (for example, *Kif26a*, *Tshz2*, *Tmem163*, *Cntn6*, *C1ql3*, *Parm1*) were specific to later stages of CSMN development (Fig. 1 and Supplementary Fig. 2).

To precisely define the neuron subtype-specific expression of the identified transcripts, we performed colocalization analyses of all the selected *Fezf2*-induced genes with CTIP2, a well-established marker of CSMN and other ScPN³. As expected, all genes tested showed various degrees of colocalization with CTIP2 (Fig. 2 and Supplementary Fig. 3a). Among the embryonic CSMN marker genes, *Adcyap1* showed restricted expression both in cortical progenitors at E13.5, the peak of CSMN neurogenesis, and subsequently in developing layer V (Fig. 2a). *Adcyap1* expression molecularly defined a subpopulation within the broader group of CTIP2⁺ neurons (Fig. 2b). At postnatal stages, *Kif26a* expression similarly defined

a subset of CTIP2⁺ neurons (Fig. 2c), whereas *Tmem163* labeled most of the CTIP2⁺ population in the cortex (Fig. 2d).

To investigate whether *Fezf2* is required for the expression of the identified target genes, we performed *in situ* hybridization for 13 genes on *Fezf2*^{-/-} cortex. We found that, in the absence of *Fezf2*, the expression of all genes tested was abrogated or decreased in layer Vb. Notably, the effect of *Fezf2* loss on expression was selective for layer Vb, as these genes were maintained in other layers (Supplementary Fig. 3b). These data indicate that *Fezf2* induces numerous early and late CSMN genes and that *Fezf2* is required for the expression of its targets in layer Vb.

Fezf2 induces a transcriptome shift toward nascent CSMN

To understand how early during development *Fezf2* governs expression of CSMN genes, we interrogated our target gene list against available RNA sequencing data from E14.5 cortical plate, subventricular zone (SVZ) and ventricular zone (VZ)¹². At E14.5 the cortical plate is mostly populated by early postmitotic corticofugal neurons (including nascent CSMN). In contrast, the SVZ and the intermediate zone (SVZ/IZ) primarily contain basal progenitors of upper layer CPN¹³⁻¹⁵ and early postmitotic upper layer CPN¹⁶. Finally, the VZ is mostly populated by apical progenitors generating upper layer CPN¹⁵. We found that *Fezf2* overexpression upregulated 186 genes enriched in the cortical plate versus SVZ/IZ and VZ (Supplementary Fig. 4 and Supplementary Table 2).

We next compared the *Fezf2*-induced genes to the available transcriptome of purified CSMN³. Strikingly, within 48 h, *Fezf2* overexpression had already induced 30 genes known to be restricted to early postnatal CSMN (Supplementary Table 3).

Next we investigated whether *Fezf2* repressed genes of alternative neuronal fates. We found that 73 genes repressed by *Fezf2* (at both 24 and 48 h) were preferentially expressed in the E14.5 VZ and SVZ/IZ, which mainly contain progenitors of upper layer projection neurons (Supplementary Fig. 5 and Supplementary Table 2). In agreement, comparative analysis with the available transcriptome of purified CPN¹⁷ showed that *Fezf2* repressed 17 genes including *Cux1*, *Zfx4*, *Fzd1* and *Btd11*. These included early markers of the CPN lineage, such as *Cux2*, shown to mark progenitors fated to an upper layer identity¹⁵. The data indicate that, when expressed in progenitors of upper layer CPN, *Fezf2* induces a battery of transcripts that collectively define developing CSMN and represses genes of developing CPN.

Fezf2 associates with the proximal promoters of CSMN genes

To elucidate the strategy used by *Fezf2* to instruct CSMN identity, we performed a genome-wide DNA binding analysis for *Fezf2* in cortical progenitors, using chromatin immunoprecipitation followed by deep sequencing (ChIP-seq)¹⁸ in combination with RNA-seq. Cortical neural progenitors were isolated at E14.5 and expanded *in vitro* for one passage as neurospheres. The neurospheres were infected with retroviruses encoding epitope-tagged *Fezf2* (3xFlag-*Fezf2*) or control (3xFlag) and harvested 48 h after infection for anti-Flag

ChIP (Supplementary Fig. 6a). The 3xFlag-tagged Fezf2 was confirmed to be functional by *in utero* electroporation (Supplementary Fig. 6b–e).

Two replicates of the ChIP-seq data sets were analyzed independently using the GEM (genome-wide event finding and motif discovery) integrative computational method¹⁹. In the two replicates, GEM analysis predicted 15,665 binding events with significant enrichment in 3xFlag-Fezf2 over control ($P < 10^{-3}$; Supplementary Table 4). Of these, 81.8% fell within 5 kb of the transcription start site (TSS) of 12,860 annotated genes (Fig. 3a and Supplementary Table 4). Genes bound by Fezf2 contained over 63% of known CSMN genes³ (Supplementary Table 4), including those expressed at early (Fig. 3b–c), middle (Fig. 3d) and late stages of CSMN maturation (Fig. 3e–i).

To assess the relationship between DNA binding and transcriptional regulation by Fezf2, we performed comparative RNA-seq analysis of 3xFlag-Fezf2 and 3xFlag neurospheres 48 h after infection (Supplementary Fig. 6a). We identified 1,295 transcripts differentially expressed upon Fezf2 induction, including 747 upregulated and 548 downregulated genes (false discovery rate = 10%; Supplementary Table 5). The fragments per kilobase RNA per million mapped reads (FPKM) values of *Fezf2* indicate absence of transcription (FPKM = 0.42) in control progenitors and high levels upon *Fezf2* overexpression (FPKM = 3,142, data not shown). Confirming Fezf2 overexpression *in vivo*, 224 of the genes induced by Fezf2 in neurospheres were preferentially expressed in the cortical plate and were enriched for postmitotic neuronal genes (Supplementary Fig. 7a and Supplementary Table 5). In contrast, 155 genes downregulated by Fezf2 were preferentially expressed in VZ and SVZ/IZ and were enriched for cell-cycle genes (Supplementary Fig. 7b and Supplementary Table 5). Genes upregulated by Fezf2 in neurospheres also showed enrichment for CSMN versus CPN signature genes by gene score enrichment analysis (GSEA) (Fig. 3j and Supplementary Table 4). Collectively, this indicates that Fezf2 induces similar transcriptional changes in isolated cortical progenitors to the ones observed *in vivo*.

We found that 64.1% of the genes induced and 77.4% of those downregulated by Fezf2 in cortical neurospheres were promoter-bound (Supplementary Fig. 8a,b). Similar percentages were observed when using transcripts regulated by Fezf2 *in vivo* (Supplementary Fig. 8c,d). These represent a significantly greater proportion of genes than expected by chance (Online Methods, Supplementary Fig. 8c–e and Supplementary Table 6). GSEA analysis of genes that were both transcriptionally regulated and promoter-bound by Fezf2, against sets of CSMN and CPN signature genes (Supplementary Table 4), showed that only CSMN signature genes were significantly enriched in the Fezf2-upregulated targets (Fig. 3j).

GEM motif-finding analysis identified a GC-rich sequence (CGCCGC) that is enriched at Fezf2-bound sites (Supplementary Fig. 9a). Since Fezf2-bound promoters are slightly more GC-rich than unbound ones (64.4% versus 49.6% GC content; Supplementary Fig. 9b), it is difficult to conclude whether Fezf2 directly binds to this motif. Nevertheless, this sequence was present in 42.9% of Fezf2-bound TSSs compared to 11.2% of unbound TSSs. In addition, electrophoretic mobility shift assay on selected target promoters containing this sequence (namely, *Ascl1* and *Ephb1*) showed binding by Fezf2 (Supplementary Fig. 9c,d). Previously reported consensus sequences predicted either by SELEX (systematic evolution

of ligands by exponential enrichment) in zebrafish²⁰ or using protein-DNA arrays in the human protein-DNA interactome (hPDI)²¹ were present in 15.4% for hPDI and 3.9% for SELEX of all Fezf2-bound sites, respectively, while the GEM-defined motif was present in 48.6% of these sites (Supplementary Fig. 9b).

Together, these data indicate that Fezf2 activates and represses a broad program of neuron subtype-specific genes by binding to proximal promoters.

Fezf2 selects glutamatergic and represses GABAergic fate

To functionally test whether genes regulated by Fezf2 control the acquisition of CSMN traits, we concentrated on two defining features of CSMN: glutamatergic identity and axonal extension to the spinal cord. Glutamate signaling is a terminal, necessary feature of all cortical projection neurons. It is unknown whether fate-specifying transcription factors for cortical neuron classes also instruct the establishment of this pan-neuronal trait. We found that Fezf2 bound to the promoter of *Vglut1*, the vesicular glutamate transporter used by most cortical projection neurons, including CSMN²², and was sufficient to induce its expression in cortical neurospheres (Fig. 4a,b). The binding appeared specific, as *Vglut2* (*Sc117a6*), used only by glutamatergic neurons in cortical layer IV and outside of the cortex²², was neither bound nor transcriptionally regulated. Concomitantly, Fezf2 bound to the promoter of *Gad1* (*Gad67*), necessary for establishment of GABAergic identity, but repressed its expression (Fig. 4a,b). To test the extent to which Fezf2 can instruct neurotransmitter choice, we established doxycycline-inducible *Fezf2-IRES-GFP* (iFIG) and control *GFP* (iGFP) mouse embryonic stem cell lines and induced neural differentiation using the Serum-free embryoid body quick (SFEBq) protocol²³ (Fig. 4c,d and Supplementary Fig. 10). GFP⁺ cells were FACS-sorted and profiled by RNA-seq. Confirming our results in primary cortical progenitors, *Fezf2* overexpression induced *Vglut1*, had no effect on *Vglut2* and inhibited the expression of *Gad1* (Fig. 4e). Together, these data indicate that Fezf2 is sufficient to select the appropriate effector genes to establish the neurotransmitter identity of CSMN, while repressing an alternative GABAergic fate.

Fezf2 directly controls expression of *Ephb1* in CSMN

Axonal connectivity to the spinal cord is a defining class-specific trait of CSMN. Directed axonal extension begins soon after CSMN fate specification and is strictly guided under the control of multiple axon guidance molecules (reviewed in ref. 24). Gene Ontology analysis of the Fezf2-induced genes showed a significant enrichment in axon guidance molecules. Fezf2 associated with the proximal promoters of 78% of these genes (Supplementary Table 4), including neuropilin 2 (*Nrp2*) [and *Robo1*, which are known to affect the extension and guidance of the corticospinal tract (Supplementary Fig. 11a). Among these, we selected for further analysis *Ephb1* (Fig. 1), encoding a tyrosine kinase receptor that mediates axon guidance and is critical for midline repulsion decisions^{25–27} and whose function in CSMN remains unknown.

We first profiled *Ephb1* expression in the developing cortex by *in situ* hybridization. *Ephb1* mRNA peaked in the cortical plate at E15.5, when CSMN axonal extension begins. *Ephb1* expression remained restricted to developing layer V, and at lower levels in layer VI, until

approximately E18.5 (Fig. 5a). Postnatally, *Ephb1* levels drastically decreased, and by P14 expression was weak and detectable only in layer VI (Fig. 5a) in both TBR1⁺ projection neurons²⁸ and APC⁺ oligodendrocytes (Fig. 5b)²⁹.

To define the neuron subtype-specific expression of *Ephb1* within layer V, we used β -galactosidase (β -gal) as a proxy for endogenous *Ephb1* expression in mice in which a *lacZ* cassette has been knocked into the *Ephb1* locus (see Online Methods; Supplementary Fig. 11b). Virtually all of the β -gal⁺ neurons were CTIP2⁺ and negative for SATB2, a marker for CPN^{30,31} (Fig. 5c). In addition, we retrogradely labeled ScPN and CPN by injecting FluoroGold into the pons and the contralateral hemisphere, respectively. We found that β -gal was expressed in FluoroGold-labeled ScPN and excluded from CPN labeled from contralateral cortex (Fig. 5c). These data demonstrate that, in the developing cortex, *Ephb1* is specifically expressed in neuronal subtypes that extend ipsilateral axonal projections and is excluded from neuronal subtypes that project through the cortical midline.

To investigate whether *Fezf2* is required for *Ephb1* expression, we performed *in situ* hybridization on cortical sections from *Fezf2*^{-/-} mice at E18.5. In the absence of *Fezf2*, *Ephb1* was no longer expressed in ScPN, and expression was maintained only at very reduced levels in layer VI (Fig. 5d). In agreement, our ChIP-seq data showed that *Fezf2* associates with the proximal promoter of the *Ephb1* gene, within a 310-base-pair region at the TSS (Fig. 5d and Supplementary Fig. 9d). *Ephb1* expression was unchanged in the striatum of *Fezf2*^{-/-} mice, a region that does not endogenously express *Fezf2* (Fig. 5d). These data demonstrate that *Fezf2* is required for the expression of *Ephb1* in ScPN.

Abnormal anterior commissure axon crossing in *Ephb1*^{-/-} mice

To decipher the role of EphB1 during CSMN development, we examined *Ephb1*^{-/-} mice. Brains of *Ephb1*^{-/-} and wild-type littermates (P28) were processed by immunohistochemistry for myelin basic protein to visualize myelinated axons. Mutant mice displayed clear abnormalities of major axon tracts (Fig. 6). The internal capsule comprised smaller than wild type and partly unfasciculated axon bundles (Fig. 6a,b). These abnormalities were accompanied by an expansion of the external capsule (Fig. 6a,c,d; arrows) and the anterior commissure (Fig. 6a,d; arrowheads). In addition, ectopic axon bundles extended within the internal capsule toward the external capsule and continued to grow ventrally (Fig. 6a,d).

We hypothesized that, in the absence of EphB1, the trajectories of subcerebral axons are rerouted, abnormally crossing to contralateral targets via the anterior commissure. To test whether the ectopic axon tracts were of neocortical origin, we anterogradely traced neocortical output projections with the lipophilic tracer DiI injected in the sensorimotor cortex of *Ephb1*^{-/-} and wild-type littermates at P2. DiI labeled axons were found in the corpus callosum and internal capsule in both wild-type and *Ephb1*^{-/-} mice. However, only in the mutant mice did we observe distinct DiI-labeled axons crossing at the anterior commissure and extending ventrally and rostrally (Fig. 6e). In wild-type cortex, only neurons located in the most lateral cortical areas, the posterior perirhinal cortex, and the entorhinal cortex project contralaterally through the anterior commissure³². In the *Ephb1*^{-/-}

mice, however, neurons located in dorsal areas projected ectopically through this commissure.

To confirm and visualize the axonal projection routes in the absence of EphB1, we used high-angular-resolution diffusion imaging (HARDI) tractography to render a three-dimensional image of major axon tracts in mutant and wild-type brains^{33,34}. We found that axon fibers in the dorsal areas of the neocortex merged with the anterior commissure in the *Ephb1*^{-/-} but not in wild-type brains (Fig. 6f). To identify a potential source of EphB1 ligands, we performed *in situ* hybridization for the ephrin genes *Efnb1*, *Efnb2* and *Efnb3*. We found that only *Efnb3* was expressed at the ventral forebrain midline and thus might mediate EphB1-dependent repulsion of ipsilateral descending tracts (Supplementary Fig. 12).

Reduced corticospinal tract in *Ephb1* null mutants

To determine whether the corticospinal tract is specifically affected by EphB1 loss, we used the *Fezf2::PLAP* (placental alkaline phosphatase) reporter line⁷ to genetically trace CSMN axons in control (*Ephb1*^{+/-};*Fezf2*^{PLAP/+}) and EphB1-deficient (*Ephb1*^{-/-};*Fezf2*^{PLAP/+}) mice at P7. In control mice, PLAP-positive axons were exclusively observed extending along ipsilateral, corticofugal trajectories (Fig. 7a). In contrast, in EphB1-deficient mice, a large proportion of PLAP-positive axons extended inappropriately to contralateral targets, crossing the midline via an enlarged anterior commissure (Fig. 7b). Most notably, this resulted in a drastic reduction of subcerebral axons reaching the cerebral peduncle (Fig. 7b) and of corticospinal axons in the dorsal funiculus of the spinal cord (Fig. 7b). These results demonstrate that, in the absence of EphB1, the ipsilateral axonal trajectories of the corticospinal tract are compromised and a distinct proportion of axons take contralateral, crossed routes through the anterior commissure.

It is notable that *Fezf2*^{-/-} mutants also show loss of axonal projections to the cerebral peduncle and the spinal cord (Fig. 7c and refs. 5–7), accompanied by an expansion of the anterior commissure (Fig. 7c and ref. 7). This phenotype is reminiscent of that observed in the EphB1-deficient mice (Fig. 7b and Supplementary Fig. 13). Beyond connectivity, in the absence of *Fezf2* the global fate specification of CSMN (molecular identity, electrophysiological properties and connectivity to the spinal cord) is abolished^{5–7}. In contrast, we found that loss of EphB1 did not affect the fate specification of projection neuron classes, which showed normal expression of the canonical projection neuron subtype-specific markers CTIP2, TBR1, SATB2 and CUX1 (Supplementary Fig. 13). The data indicate that EphB1 is an effector molecule directly downstream of *Fezf2* that executes a specific modular aspect of the global *Fezf2*^{-/-} phenotype.

DISCUSSION

Deciphering the molecular logic that orchestrates the acquisition of neuronal subtype-specific features in the mammalian cerebral cortex is fundamental to understanding how its neuronal diversity has evolved. These molecular rules are not yet defined for any subtype of cortical neuron. Here we report that acquisition of defining features of corticospinal motor

neurons is at least partly achieved by co-regulated expression of batteries of CSMN effector molecules directly downstream of a single selector gene, *Fezf2*.

Precise control over activation and repression of transcription factors responsible for acquisition of class-specific traits in projection neurons is critical to resolve neuronal fates during cortical development. *Tbr1* and *Sox5* act as repressors of *Fezf2* to negatively regulate the extension of the corticospinal tract and to promote connectivity to the thalamus^{35–38}. Similarly, reciprocal repression of *Fezf2*, *Satb2* and *Ctip2* is employed to choose subtype-specific long distance connectivity^{39,40}. These studies motivate a more global analysis of the molecular rules beyond single genes that govern the acquisition of lineage-specific traits.

We found that expression of *Fezf2* in cortical progenitors fated to generate upper layer neurons resulted in both the induction of a large series of genes expressed in nascent CSMN and the repression of molecular programs of upper layer neurons. Given that this transcriptional regulation largely occurred via binding of *Fezf2* to the promoters of both activated and repressed genes, the data indicate that lineage bifurcation decisions can be co-regulated by the same transcription factor from very early stages of differentiation. Our data also provide a molecular explanation for why, during development, cortical progenitors tune *Fezf2* expression to the lower levels observed in late-stage cortical progenitors^{5,7,41}. Maintenance of *Fezf2* expression after the completion of neurogenesis of deep layer neurons may be molecularly incompatible with the initiation of upper layer neurogenesis.

Much work is still required to understand the nuances of how *Fezf2* can act as an activator for some target genes and as a repressor for others. It is likely that factors such as post-translational modification of *Fezf2*, the choice of interacting cofactors and the presence of neighboring binding sites for different proteins at different loci critically contribute to these distinct transcriptional outcomes⁴².

Transcription factors endowed with properties of classic selector genes have been identified in different tissues as genes uniquely able to control ‘master’ molecular switches of distinct cellular fates. Elegant work in *Drosophila* has highlighted core properties of selector genes that clarify the logic guiding their activity. Multiple target genes are often regulated downstream of a single selector gene, and targets often include both activated and repressed genes⁴³. Our evidence that *Fezf2* acts as both an activator of CSMN genes and a repressor of genes for alternative fates further supports the idea that *Fezf2* is a selector gene. To our knowledge, this is the first transcription factor endowed with these properties for any class of cortical neurons.

During development of CSMN, expression of lineage-specific and pan-neuronal genes alike display temporally dynamic regulation. It is intriguing that within 48 h of induction, *Fezf2* is already associated with the promoters of CSMN genes whose expression is normally temporally spaced during CSMN differentiation. The data suggest that, in cortical progenitors, *Fezf2* might be ‘priming’ the transcriptional territory of CSMN by also occupying genomic loci that are transcriptionally active in these neurons at later times. This implies that temporal resolution of gene expression over the several weeks of CSMN development likely relies on cofactors that operate by interaction with *Fezf2* or work on the

genomic loci that are primed by early *Fezf2* binding. The fact that *Fezf2* expression is maintained at all stages of CSMN differentiation *in vivo*⁵⁻⁷ supports the view that active binding by *Fezf2* to CSMN loci may be required to poise them for transcriptional regulation at later stages of differentiation.

Precedents exist for the use of early priming of genomic loci in progenitor cells to enable the execution of specific differentiation programs at later stages of development. In *Caenorhabditis elegans*, asymmetric distribution of gene expression in bilaterally symmetrical ASE gustatory neurons relies on early priming of the *Isy-6* microRNA locus in progenitors of only the left neurons⁴⁴. *MyoD* has also been shown to associate with enhancers of both early and late myogenic genes⁴⁵.

In his first definition of selectors, Antonio García-Bellido postulated that these genes would regulate effector molecules necessary for the acquisition of specific cellular features⁴⁶. Here we identify EphB1 as an effector molecule of *Fezf2*. In turn, EphB1 executes CSMN axonal repulsion from the midline in the ventral telencephalon, enabling the ipsilateral extension of the corticospinal tract. Our data are in agreement with work demonstrating a key role for EphB1 in mediating midline repulsion decisions. In the spinal cord, EphB1 instructs the ventral trajectory of motor neuron axons by mediating their repulsion from the mesenchyme of the dorsal limb²⁵. Similarly, in the optic chiasm, EphB1 mediates the repulsion of retinal ganglion cell (RGC) axons from the midline⁴⁷. Notably, in the retina the transcription factor *Zic2* is also upstream of genes critical for both axonal guidance decisions (EphB1) and neurotransmitter identity (*Sert*)⁴⁸.

The regulation of axon guidance molecules in different classes of cortical projection neurons has just begun to be explored⁴⁰. Whether expression of lineage-specific axon guidance molecules is directly controlled by the same transcription factors that govern neuron subtype fate specification remains unknown. It is striking that the phenotypic abnormalities observed in the absence of EphB1 mimic the axon guidance defects observed in the *Fezf2*^{-/-} mutants, without affecting the overall fate specification of CSMN. While the possibility that expression of EphB1 in thalamus or striatum contributes to this phenotype⁴⁹ cannot be ruled out, the fact that *Fezf2* is not expressed in these regions and yet *Fezf2*^{-/-} mice present similar axon guidance defects to *Ephb1* mutants supports a cell-autonomous role of EphB1 in CSMN. Our data point to a direct regulation of key CSMN axon guidance receptors by the same selector gene that governs global programs of fate specification of this neuron subtype.

Not all selector genes have equal properties. Pioneering work in *C. elegans* has identified ones termed terminal selector genes sufficient to instruct specific neurotransmitter fates⁵⁰. Our data indicate that *Fezf2* controls the acquisition of glutamatergic identity by direct binding to the promoter of *Vglut1* and other genes involved in the synthesis, transport and signaling of glutamate. This is in agreement with the fact that overexpression of *Fezf2* in progenitors of GABAergic medium spiny neurons results in a switch to a glutamatergic fate *in vivo*⁸. In addition, like selector genes in *C. elegans*, *Fezf2* expression is maintained in CSMN throughout life, suggesting a requirement for *Fezf2* to both establish and maintain

CSMN traits. In the future, conditional loss of *Fezf2* in adult CSMN will test this possibility and define the extent to which *Fezf2* fulfills the definition of a canonical terminal selector.

An understanding of the molecular regulatory architecture that shapes the identity of CSMN and other classes of cortical neurons is fundamental to gaining insights into how neuronal diversity develops in the cerebral cortex and will furthermore inspire tools to reprogram class-specific neuronal identity in the context of neurodegenerative disease.

ONLINE METHODS

Mice

Fezf2^{-/-} mice were generated by Hirata *et al.*⁴¹ (*Fezf2* GenBank accession code: AB042399). *Ephb1*^{-/-} mice (*Ephb1* GenBank accession code: NM_173447) were generated by Deltagen, Inc. (Menlo Park, CA). The line was generated by targeting the *Ephb1* coding region with a cassette containing a 4.5-kb homology sequence upstream of exon 8, a Lac0-SA-IRES-lacZ-WT Neo/Kan cassette and a 2.2-kb homology sequence downstream of exon 8. The insertion of the Lac0-SA-IRES-lacZ-WT Neo/Kan cassette resulted in a 100-bp deletion of genomic DNA. Sequences of PCR primers employed to genotype the *Ephb1*^{-/-} line are as follows: forward primer, ACGTGGGAGGACTCTAATCCTCTTC; reverse primer (wild-type allele), TCTAGGTTGCTGGCTACAGGACTTG; reverse primer (Neo), GGGCCAGCTCATTCCTCCCATCAT. *Fezf2::PLAP* (*Fezf2*^{PLAP/+}) mice were generated by Chen *et al.*⁷ and crossed with the *Ephb1*^{+/-} mutant line to obtain *Ephb1*^{-/-};*Fezf2*^{PLAP/+} mutants. Timed-pregnant CD-1 mice were obtained from Charles River Laboratories (Wilmington, MA) for *in utero* electroporation. The day of the vaginal plug was designated embryonic day 0.5 (E0.5). The day of birth was designated postnatal day 0 (P0). All mice were maintained in standard housing conditions on a 12-h light/dark cycle with food and water ad libitum. No more than four adult animals were housed per cage. All mouse studies were approved by the Massachusetts General Hospital and Harvard University Institutional Animal Care and Use Committee and were performed in accordance with institutional and federal guidelines.

In utero electroporation

The *Fezf2*^{GFP} and *Ctrl*^{GFP} construct and conditions for *in vivo* electroporation were described previously^{5,51}. The *3xFlag-Fezf2* open reading frame was cloned into the *Ctrl*^{GFP} construct driven by the CAG promoter. Briefly, 800 nl of purified DNA (2 µg/µl) mixed with 0.005% fast green in sterile PBS was injected *in utero* into the lateral ventricle of CD1 embryos at E14.5 under ultrasound guidance (Vevo 770, VisualSonics). Five 40-V pulses of 50 ms were delivered at 1-s intervals in an appropriate orientation across the embryonic head using 1-cm-diameter platinum electrodes placed outside the uterus, using a CUY21EDIT square wave electroporator (Nepa Gene). Injected embryos were collected for FACS purification either 24 h or 48 h after electroporation.

FACS purification

Pregnant dams were deeply anesthetized 24 h or 48 h after surgery. Their embryos were removed and sensorimotor cortex was microdissected, as previously described⁵², from the

electroporated cerebral hemisphere using a fluorescence dissecting microscope to precisely visualize the labeled regions. *Fezf2^{GFP}* and *Ctrl^{GFP}* electroporated cells were purified by FACS directly into RNAlater⁵³ and RNA was extracted as previously described⁵⁴. To control for biological sample variability, we processed multiple independently collected samples for both *Fezf2^{GFP}* and *Ctrl^{GFP}* electroporated cells at each time point of analysis (true biological replicates) for a total of fourteen samples derived from different litters, FACS purifications and microarray hybridizations (three each of *Fezf2^{GFP}* and *Ctrl^{GFP}* at 24 h and four each at 48 h). *Fezf2^{GFP}*- and *Ctrl^{GFP}*-labeled cortical tissues were enzymatically digested in dissociation medium (20 mM glucose, 0.8 mM kynurenic acid, 0.05 mM APV, penicillin-streptomycin (50 U/ml and 0.05 mg/ml, respectively), 0.09 M Na₂SO₄, 0.03 M K₂SO₄ and 0.014 M MgCl₂) containing L-cysteine HCl (0.016 µg/µl) and papain (10 unit/ml; Worthington Biochemical Corp., NJ) at 37 °C for 30 min. Papain digestion was stopped with ovomucoid (10 mg/ml) and BSA (10 mg/ml) in dissociation medium at room temperature. Progenitors were mechanically dissociated to obtain a single-cell suspension by gentle trituration in iced OptiMEM containing glucose (20 mM), kynurenic acid (0.4 mM) and APV (0.025 mM). All chemicals were purchased from Sigma, and all media were purchased from GIBCO-BRL. GFP-positive cells were purified using a BD FACS Vantage SEM DiVa cell sorter. Cells were gated on the basis of on fluorescence, forward scatter and side scatter to select the appropriate population. FACS-purified progenitors and neurons from both time points were collected and stored in RNAlater before RNA extraction⁵³.

Microarray and analysis

Approximately 10,000 to 30,000 FACS-sorted progenitors were used for each biological replicate. RNA was extracted using the StrataPrep Total RNA Micro Kit (Stratagene), and RNA quality was assessed using a Bioanalyzer (Agilent Technologies). RNA was amplified per Affymetrix small sample protocol using two consecutive rounds of linear *in vitro* transcription to obtain 15–20 µg of amplified and labeled cRNA for each hybridization. Microarray hybridization was performed with an Affymetrix Mouse Genome 430 2.0 Array according to standard Affymetrix protocol⁵⁵. For microarray experiments, 3 or 4 replicates were used at each age, which allows the identification of differential gene expression with low false discovery rates³.

Using the Rosetta Resolver software (Rosetta Biosoftware), data from individual microarrays were normalized with the trend function and replicates were combined. Statistical significance of gene expression differences between *Fezf2^{GFP}* and *Ctrl^{GFP}* experiments was determined by pairwise comparisons at each time point using the Rosetta Resolver error modeling method⁵⁶ with *P*-value limit of <0.001 and fold change of >1.5. Similar results were obtained by normalizing with GCRMA in Bioconductor⁵⁷ or with the Bioconductor implementation of MAS 5.0 and performing pairwise comparisons using significance analysis of microarrays (SAM)⁵⁸.

For comparison with RNA-seq expression data from Ayoub *et al.*¹², BAM files were downloaded from NCBI (accession code GSE30765), differential expression was determined with Cuffdiff, and analysis was performed with the Cumberbund package in the

R language and environment for statistical computing and graphics⁵⁹. From a list of genes with significant differential expression between VZ, SVZ and cortical plate ($P < 0.001$) in at least one sample, those with significant up- or downregulation induced by *Fezf2* were identified and subsequently used for cluster analysis with Jensen-Shannon divergence as the distance metric in Cumberbund.

Western blot

Brains were dissected from P1 *Ephb1* knockout, heterozygous and wild-type mice and lysed in RIPA buffer (150 mM NaCl, 1% NP40, 0.5% deoxycholate, 0.1% SDS, 1 mM EDTA, 50 mM Tris-HCl, pH 8) with proteinase inhibitors (S8830, Sigma-Aldrich). Mouse ES cells were treated with 2 $\mu\text{g}/\text{ml}$ doxycycline for 48 h and harvested in RIPA buffer. After quantification by Bio-Rad protein assay (Bio-Rad), proteins were resolved in 4–12% SDS-PAGE gels and transferred to a nitrocellulose membrane. The western blot was performed using goat anti-EphB1 (M19, Santa Cruz, 1:250), rabbit anti-Fezf2/Fezf1 (F441, IBL, 1:500), rabbit anti- β -tubulin (9F3, Cell Signaling, 1:500) and mouse anti- α -tubulin (T9026, Sigma, 1:5,000). Anti-goat, anti-rabbit and anti-mouse HRP-conjugated secondary antibodies (HRP-linked anti-rabbit IgG, Cell Signaling Technology 7074, HRP-conjugated goat anti-mouse IgG, Cell Signaling Technology 7076, HRP-conjugated anti-goat IgG, Santa Cruz sc-2020, all at 1:5000), followed by ECL (Amersham), were used to visualize proteins of interest on the membrane.

Retrograde and anterograde labeling

Pups were anesthetized by hypothermia at P2 and injected with FluoroGold (for retrograde labeling) in the contralateral cortex or the pons, or with DiI (for anterograde labeling) in the motor cortex as previously described^{17,60}. The site of DiI injection was identified by ultrasound guidance. Pups were returned to the care of their mother and deeply anesthetized at P4 and P7 in the retrograde tracer studies, and at P4 for the anterograde tracer study, before perfusion and collection of the brain for immunocytochemistry.

High-angular-resolution diffusion imaging (HARDI)

Four-week-old (P28) wild-type and *Ephb1*^{-/-} littermates were perfused with saline followed by 4% paraformaldehyde, postfixed for 24 h, and subsequently fixed for an additional week in 4% paraformaldehyde solution containing 1 mM gadolinium (Gd-DTPA). For image acquisition, the brains were placed in Fomblin. Brains were scanned on a 4.7-T Bruker Biospec MR system. The pulse sequence used for image acquisition was a 3D diffusion-weighted spin-echo echo-planar imaging sequence, time repetition (TR)/time echo (TE) 1,000/45.47 ms, with an imaging matrix of $96 \times 96 \times 128$ pixels. Spatial resolution was $125 \times 125 \times 125 \mu\text{m}$. Sixty diffusion-weighted measurements ($b = 4,000 \text{ s}/\text{mm}^2$) and one non-diffusion-weighted measurement ($b = 0$) were acquired with $\delta = 12.0 \text{ ms}$, $\tau = 24.2 \text{ ms}$. Total acquisition time was approximately 2 h for each imaging session. The HARDI method was used to reconstruct the orientation distribution function in each voxel and resulting tractography pathways were reconstructed using a streamline algorithm for diffusion tractography^{33,61} Diffusion Toolkit and TrackVis (<http://trackvis.org/>) were used to reconstruct and visualize axonal pathways. Trajectories were propagated by consistently

pursuing the orientation vector of least curvature, and tracking was terminated when the angle between two consecutive orientation vectors was greater than the given threshold of 45° for each specimen. Brain mask volumes created by MRICro (<http://www.mccauslandcenter.sc.edu/mricro/mricro/mricro.html>) were used to terminate tractography structures instead of the FA (Fractional Anisotropy) threshold as previously reported^{61,62}. Commissural pathways were isolated as fibers passing through the manually drawn region of interest that included the entire anterior commissure at the midline, ensuring that other fiber pathways were not contributing, as previously described⁶¹.

Immunocytochemistry, PLAP staining and *in situ* hybridization

Brains for immunocytochemistry were processed as previously described^{55,63}. Primary antibodies and dilutions used were as follows: rabbit anti-TBR1 antibody, 1:2,500 (gift from R. Hevner); rat anti-CTIP2 antibody, 1:1,000 (Abcam ab18465); rabbit anti-GFP, 1:500 (Invitrogen A11122); chicken anti-β-galactosidase, 1:500 (ICL lab CGAL-45A); mouse anti-SATB2, 1:50 (Abcam ab51502); rat anti-MBP, 1:500 (Millipore MAB386); and rabbit anti-CUX1, 1:100 (Santa Cruz CDP M-222). Goat anti-rabbit IgG Alexa Fluor 488, 546, and 647 (Life Technologies A11070, A-11071, A21246), Goat anti-chicken IgG Alexa Fluor 488, 546, 647 (Life Technologies A11039, A11040, A21449), Goat anti-rat 488, 546, 647 (Life Technologies A11006, A11081, A21247), Goat anti-mouse IgG Alexa Fluor 488, 546, 647 (Life Technologies A11017, A11018, A21237) secondary antibodies were diluted 1:750. PLAP activity was detected with alkaline phosphatase staining buffer (0.1 mg/ml 5-bromo-4-chloro-3-indolyl phosphate; 1 mg/ml nitro blue tetrazolium in 100 mM Tris-HCl, pH 9.5; and 100 mM NaCl) as previously described⁷. *In situ* hybridization and combined *in situ* hybridization with immunohistochemistry were performed following published methods and using 40-μm-thick sections cut on a vibrating microtome and mounted on Superfrost slides (Fisher)⁶⁴. Riboprobes were generated as previously described⁵⁴. The cDNA template clone for *Ldb2* was provided by J. Macklis; all other riboprobes were generated from cDNA template clones using the primers listed below. Tissue sections were imaged using a Nikon 90i fluorescence microscope equipped with a Retiga Exi camera (Q-Imaging) and analyzed with Volocity image analysis software v4.0.1 (Improvision). All primary data from immunohistochemistry and *in situ* hybridization experiments were repeated at least three times and analyzed by one investigator, then confirmed by a second, independent investigator who was blinded to genotype and experimental conditions.

In situ hybridization riboprobe templates

cDNA templates for riboprobes were obtained by RT-PCR with the following primer pairs:

Gene name	Forward primer	Reverse primer	Product length
<i>Acvr1c</i>	AGGACTTGCCCTCGAAAAGTGA	GAAAGCAGAAGCGGACATTC	610
<i>Adcyap1</i>	AATGACTTGGGGAATTGCTG	GATGAACAGCACTGGAGAA	542
<i>Akap12</i>	GTTGGGACCTTGAGACAAA	TGGCACACTCATCTGTCCAT	691
<i>C1ql3</i>	AGTGTGGTCCCTCACCTGGA	CAAGAACCAAAGCTGACACG	507
<i>Cntn6</i>	GTATCTGTCCGAGAAGGTCA	GTGGACCTTGGACACTTCTA	238

Gene name	Forward primer	Reverse primer	Product length
<i>Ephb1</i>	CACATCCATCTCCCTTTGCT	TCCAGAAACCCTTTCCCTCT	618
<i>Efnb1</i>	ACAAGCCACACCAGGAAATC	TGGGGGCAGTAGTTGTTCTC	640
<i>Efnb2</i>	ACCACACAGCTATGCAGCAG	CGAACTCCACGCTTCTGGT	675
<i>Efnb3</i>	GCAGTGTGGACATGATGGAC	GCACACTAAAGAGCGGGAAG	675
<i>Kif26a</i>	TCCTCAGCTCCAGACTCCAT	GCGACAGTCTTTCCATCTCC	851
<i>Pappa2</i>	CCATTGTTCCACAAATGCTG	CTCGCTCCACATGTTGCTTA	900
<i>Parm1</i>	TGCACTGACCAAGCCAGATA	ACAAACAGCAAGGCAGTGAA	700
<i>Rgs16</i>	AAAAGGCTGTGTGTGTGGAAC	CTATCACTTCTGAGTCTTACCG	752
<i>Rgs8</i>	TGAGGTCATGTTTGGGTTCA	CAGGCTCTACGGACTTCTGG	741
<i>Tmem163</i>	AGGGTCTCTGCTTGACAGGA	CCCTACATGTTGGCACACAC	650
<i>Tshz2</i>	GGCGAAGAGGACACAGACTC	AAGGAGCGCTGTGCGATAAAA	710
<i>Tmem117</i>	AACTATGCCACAACGGTGCT	TTGTAGACAGTGGGCTGTGC	754

Neurosphere isolation and viral infection

Primary cortical progenitors of the dorsal telencephalon were isolated from E14.5 brains and grown as neurospheres in the presence of growth factors as previously reported⁶⁵.

Neurospheres were expanded for a maximum of one passage and then plated as a monolayer on dishes coated with poly-D-lysine (VWR) and laminin (BD). E14.5 pregnant dams were deeply anesthetized and the embryos removed. The presumptive somatosensory area was microdissected in cold Hanks' medium and mechanically triturated to obtain a single-cell suspension. Cells were spun at 5,000g for 5 min at room temperature, and the pellet was resuspended in neurosphere medium, containing 100 U/ml penicillin/streptomycin, 200 mM L-glutamine, N2 and B27 supplements (Gibco), as well as 20 ng/ml epidermal growth factor (EGF; Sigma) and 20 ng/ml β -fibroblast growth factor (FGF; Millipore) in order to promote the survival of neural stem cells. After quantification of cell density, neural stem cells were plated in 200-ml tissue culture flasks (untreated or low-adherence, Corning) at 200,000 cells/flask and incubated at 37 °C with 5% CO₂. The medium was changed every 3–4 d and the cells were expanded as neurospheres for one passage. Subsequently, cells were plated as a monolayer on dishes previously coated with poly-D-lysine (VWR) and laminin (BD).

3xFlag-Fezf2 and *3xFlag* were cloned into pRETROX-IRES-ZsGreen1 retroviral vectors (Clontech Laboratories, Inc.) to generate high-titer, replication-incompetent, VSVg-coated retroviral particles that were packaged in 293T cells (MOI = 5). 12–18 h after plating, neural stem cells were infected with retroviruses. 16–203h after infection, cells were switched into fresh medium. Cells were collected 48 h after infection for chromatin immunoprecipitation (ChIP, $n = 2$) and for RNA-seq ($n = 2$). Sample sizes for ChIP-seq and RNA-seq experiments on neural progenitors complied with the ChIP-seq ENCODE guidelines⁶⁶.

Chromatin immunoprecipitation (ChIP)

ChIP analysis was performed on 1.5×10^8 neural stem cells per condition (*3xFlag-Fezf2*^{ZsGreen1} and *3xFlag*^{ZsGreen1}, 2 replicates each) as previously described⁶⁷. Briefly, cells were dissociated with 0.25% trypsin (Invitrogen), washed with PBS and chemically cross-linked with formaldehyde solution (1%). Cells were disrupted in lysis buffer (50 mM Tris,

pH 8.0; 10 mM EDTA; 1% SDS; and protease inhibitors) for 20 min on ice and sonicated with a Bioruptor to shear the DNA into 200–700 bp fragments. Dynal magnetic beads (sheep anti-mouse M-28, Invitrogen) were preblocked with 5 μ g of anti-Flag M2 antibody per reaction (Sigma-Aldrich) rotating overnight at 4 °C. Beads were then washed and resuspended in fresh 0.5% BSA in PBS. Bead-antibody complexes were incubated at 4 °C overnight with rotation. Beads were resuspended twice in low-salt immune complex wash buffer (0.1% SDS; 1% Triton X-100; 2 mM EDTA; 20 mM Tris-HCl, pH 8.1; and 150 mM NaCl), incubated for 5 min at 4 °C with rotation and resuspended in LiCl immune complex wash buffer (0.25 M LiCl; 1% NP40; 1% deoxycholate; 1 mM EDTA; and 10 mM Tris-HCl, pH 8.1) twice and incubated for 5 min at 4 °C with rotation. Beads were then rinsed with ice-cold 10mM Tris-HCl; 1mM EDTA, pH8.0 and DNA was eluted in 100 μ l of elution buffer (50mM Tris-HCl; 10mM EDTA; 1%SDS, pH 8.0) at 65 °C for 15 min. Reverse crosslinking was performed by incubating the CHIP DNA overnight at 65 °C. DNA was purified on QIAquick PCR purification columns (Qiagen) for use as template for Solexa library construction. To prepare an input control, whole-cell-extract DNA (reserved from the sonication step) was also treated by reverse crosslinking and purified.

ChIP-seq data analysis

For both ChIP-seq replicates, high-throughput sequencing reads were aligned to the mouse genome (version mm9) using Bowtie version 0.12.5 with options “-q --best --strata -m 1 -p 4 --chunkmbs 1024,” and only uniquely mapping reads were retained for further analysis. GEM was used to detect binding events, using the options “--top 2000 --k_min 7 --k_max 12 --a 20 --q 3 --mrc 1 - constant_model_range --d_l 1000 --d_r 1000 --v 2 --nf --refine_pwm --gc0 0.42.” Reported peaks contain a ChIP-seq enrichment level that is significantly greater than 1.5 times the scaled read count from the corresponding region in the control experiment ($P < 10^{-6}$, binomial test, adjusted for multiple testing using Benjamini & Hochberg’s method). GPS (genome positioning system) was also used to confirm binding locations, using the options “--top 2000 --a 20 --q 3 --mrc 1 --constant_model_range.” Regions of ChIP enrichment (1 Kbp) are highlighted in red or blue, centered on the GEM-predicted binding event locations. *De novo* motif discovery was also performed using GEM, which jointly estimates binding locations and primary DNA motif patterns. Motif frequencies at binding events and gene promoter regions were defined by scanning 100-bp windows around GPS binding locations or annotated TSSs for matches to each motif. Motif scoring thresholds were based on a false discovery rate of 10^{-3} , defined using a third-order Markov model of the mouse genome. Comparison between peaks and gene features was performed using the Ensembl v62 mouse genome annotation. Genes were defined as bound if a Fezf2 binding event was located within 5 kb of a gene’s TSS. A chi-squared test was used to assess the significance of associations between Fezf2 binding and groups of differentially regulated genes *in vivo*.

RNA-Seq library preparation

Total RNA was isolated using QIAzol (Qiagen)/chloroform extraction followed by spin-column purification (RNeasy mini kit, Qiagen) according to the manufacturer’s instructions. RNA concentration and purity were measured with Nanodrop (Thermo Fisher), and RNA integrity was assessed on a Bioanalyzer (Agilent) using the RNA 6000 Nano Total RNA kit

(Agilent). High-quality RNA samples (RNA integrity number ≥ 8) were used for library preparation. Poly(A)⁺ RNA-seq libraries were constructed using the TruSeq RNA Sample Preparation Kit (Illumina) following the manufacturer's protocol. 200 ng of total RNA from neurospheres was used as input for the TruSeq libraries. Before sequencing, libraries were run on a Bioanalyzer DNA7500 chip to assess purity, fragment size and concentration. Libraries free of adaptor dimers and with a peak region (220–500 bp) area $\geq 80\%$ of the total area were sequenced. Individually barcoded samples were pooled and sequenced on the Illumina HiSeq 2500 platform.

RNA-Seq analysis

Paired-end 101-bp reads were aligned to the mouse (mm9) reference genome assembly using Tophat2 (refs. 68,69) with default options. Aligned reads and the UCSC reference transcriptome .gtf file (UCSC genome browser; knownGene track) were used as input for Cuffdiff2 (ref. 70) for expression quantification in fragments per kilobase RNA per million mapped reads (FPKM) and differential testing between conditions using default options ($P < 0.001$). Cummebund v2.1 (<http://compbio.mit.edu/cummebund/>) was then used to process, index and visualize the output of the Cuffdiff2 analyses.

To test for enrichment for CSMN signature genes after *Fezf2* overexpression in cortical progenitors, we performed a preranked gene set enrichment analysis (GSEA). Briefly, CSMN signature genes were selected from Arlotta *et al.*⁵⁴ as the intersection of genes flagged as 'present' (Affymetrix; RMA) with a fold-change of CPN/CSMN $\geq 1/3$ from the P3 and P6 time point comparisons. Inversely, CPN signature genes were created from the intersection of genes with a fold-change of CPN/CSMN ≥ 3 from the same two time points. Additional gene sets consisting of the subset of each gene list that was also selected as *Fezf2*-bound from our CHIP-Seq study were created.

We next created a ranked gene list of all expressed genes (FPKM ≥ 1) from our *3xFlag-Fezf2^{ZsGreen1}* versus *3xFlag^{ZsGreen1}* cortical progenitor RNA-Seq assay. Expressed genes were rank-ordered using the value of the Cuffdiff2 test statistic. All derived gene sets were tested for enrichment or depletion after *Fezf2* overexpression using the preranked GSEA approach⁷¹. CSMN signature and CSMN signature plus *Fezf2*-bound gene sets were determined to be significant with a nominal P value ≤ 0.01 .

Gene ontology enrichment analysis was performed on *Fezf2*-induced genes with specific expression in the cortical plate (as determined by Ayoub *et al.*¹²) and *Fezf2*-repressed genes with specific expression in either the VZ or SVZ/IZ using DAVID⁷². Genes within each gene set were selected using a threshold specificity score $S \geq 0.65$ (described below) for any given condition. Gene sets were tested for significant enrichment using the GO Biological Process FAT collection of gene sets.

We defined the specificity score S_{gi} for a gene (g) in a condition (i) as follows:

$$S_{gi} = 1 - \sqrt{\text{JSD}(p_g, q_i)}$$

where JSD is the Jensen-Shannon divergence between p_g , the expression profile of a gene g across all conditions expressed as a probability distribution, and q_i , the unit vector of perfect expression in condition i . Where applicable, genes are specifically assigned to the condition with the highest S score above a threshold value. The threshold for specificity was selected empirically to be 0.65 through examination of the distribution of S scores across all genes.

Hypergeometric and bootstrap analyses

To determine enrichment for Fezf2-bound genes within the set of significantly differentially regulated genes in the E14.5 cortical neurospheres, we employed a hypergeometric test (phyper from the stats package in R) and a bootstrap method. For the bootstrap method, gene sets of comparable size were sampled (without replacement) from the universe of genes 10,000 times, and P values were estimated as the fraction of iterations where a greater than or equal number of Fezf2-bound genes were selected.

Derivation and differentiation of inducible Fezf2 and GFP mouse ES lines

Mouse *Fezf2-IRES-GFP* was PCR-cloned from the *Fezf2^{GFP}* construct⁵ into P2lox-40 using the Gateway system (Life Technologies). The control GFP construct was a gift from E. Mazzoni⁷³. The plasmid constructs were nucleofected into the A2loxCRE mouse ES line for inducible cassette exchange (ICE) as described in Iacovino *et al.*⁷⁴ The cells were selected on DR4 mouse embryo fibroblasts (GlobalStem) with 300 $\mu\text{g/ml}$ G418. Single inducible *Fezf2-IRES-GFP* (iFIG) clones and inducible GFP (iGFP) clones were manually picked and expanded in standard ES+LIF medium without doxycycline (15% FBS, 1 \times Glutamax, 1 \times non-essential amino acids, 1 \times penicillin/streptomycin and 1,000 U/ml mLIF (ESGROW, Millipore) in Knockout DMEM). All media were obtained from GIBCO BRL. The iFIG and iGFP clones were differentiated *in vitro* into cortical neurons using the SFEBq protocol as described in Eiraku *et al.*⁷⁵ Expression of *Fezf2* and *GFP* was induced at day 10 *in vitro* with 2 $\mu\text{g/ml}$ doxycycline. The doxycycline was maintained in the culture until day 18, and the SFEBqs were dissociated and FACS-sorted for GFP⁺PI⁻ populations, which were then analyzed by RNA-seq.

Electrophoretic mobility shift assay (EMSA)

GST-tagged truncated mouse Fezf2 (GST-tFezf2, AA291-455) protein was expressed in Rosetta DE3 *E. coli* (Millipore) and purified on GST-agarose with standard methods. Given that the full-length Fezf2 aggregates in solution, we chose a truncated version of Fezf2 (tFezf2) containing five of the six zinc finger motifs in the C-terminal domain of Fezf2, highly conserved across all vertebrates. IRDye end-labeled oligonucleotide probes (Integrated DNA Technologies) were designed against proximal promoter regions under ChIP-seq peaks of the *Ascl1* and *Ephb1* genes. Negative controls for each gene were chosen from neighboring sequences not containing ChIP-seq peaks. The probe sequences are as follows (with GEM motifs underlined):

Ephb1 positive probe:

GCCGAGCCCCAGCGGAGACGCGCCGCGTCCCAGGGCGCCGCTGCGCTCCCGGC
GGGTGGCTAGCCACCCGCCGGGAGCGCAGCGGGCGCCCTGGGACGCGGCGCGTC
TCCGCTGGGGCTCGGC.

Ephb1 negative probe:

TGTCAGAAGACAGTCATAATGGGAAGACATGAAGAAACAGCCGAAAGTCTGAC
AACTTGTTAAACAAGTTGTCAGACTTTCGGCTGTTTCTTCATGTCTTCCATTATG
ACTGTCTTCTGACA.

Ascl1 positive probe:

GCTCTGAGCTGCCGCGGCCGCCGCCGCTGCCGCCGCCGCCGCCGGTTCGCAAAGAA
GCAGGCGCCTGCTTCTTTGCGACCGCGGCGGGCGGCAGCGGGCGGCGGCCGC
GGCAGCTCAGAGC.

Ascl1 negative probe:

CCTAGTGGTTATTTTATTGCTGTAAAATAAATTTAACCCTTTCCTTACCAAGCTG
CTTTTAAAAGCAGCTTGGTAAGGAAAGGGTTAAATTTATTTTACAGCAATAAAA
TAACCACTAGG.

EMSA was performed using the Odyssey EMSA Kit (LI-COR Biosciences, USA), following the manufacturer's protocols. Briefly, 100 nM IRDye end-labeled probe was incubated with 0, 10 or 20 µg GST-tFezf2 at room temperature for 30 min in 20 µl reaction buffer (1× binding buffer, 0.2 mM DTT, 0.25% Tween 20, 0.05 µg/µl poly(dI.dC) and 5 mM MgCl₂). The reactions were resolved on 4–20% Mini-PROTEAN TBE precast gel (Bio-Rad) and visualized by the Odyssey CLx infrared imaging system (LI-COR Bioscience, USA).

Statistics

Bar and line graphs represent mean values of all replicates and error bars represent s.e.m. except for RNA-seq results, in which the error bars represent 95% confidence intervals for the Cuffdiff2 model expression estimate. Animals were assigned to groups on the basis of genotype. Age-matched littermates were used as controls in all experiments. No randomization was used. For all animal experiments, data distribution was assumed to be normal, but this was not formally tested. For ISH and immunofluorescence studies, data collection and analysis were performed blinded to the conditions of the experiment. No animals were excluded from analyses. Except for RNA-seq and ChIP-seq sample sizes (explained above), no statistical methods were used to predetermine sample sizes. Our sample sizes are similar to those reported in previous publications³.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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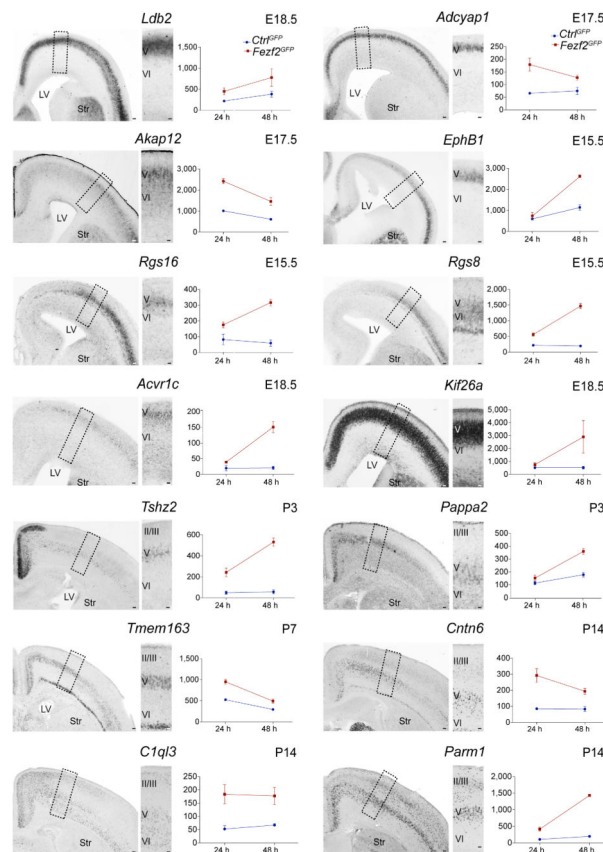


Figure 1.

Fezf2 overexpression in cortical progenitors induces genes that label corticospinal motor neurons. Left, *in situ* hybridizations on coronal sections of the cerebral cortex at different embryonic (E15.5, E17.5 and E18.5) and postnatal stages (P3, P7 and P14) (insets enlarged from boxed areas). Right, expression levels (normalized microarray intensity, see Methods) for each selected gene in *Ctrl*^{GFP} (blue) and *Fezf2*^{GFP} (red) *in utero* electroporated cortical progenitors that were collected at 24 h ($n = 3$ litters per condition) and 48 h ($n = 4$ litters per condition). Vertical axes are normalized intensities (arbitrary units). Error bars indicate s.e.m. LV, lateral ventricle; Str, striatum. Scale bars, 100 μ m; 50 μ m in insets. Source data are shown in Supplementary Table 1.

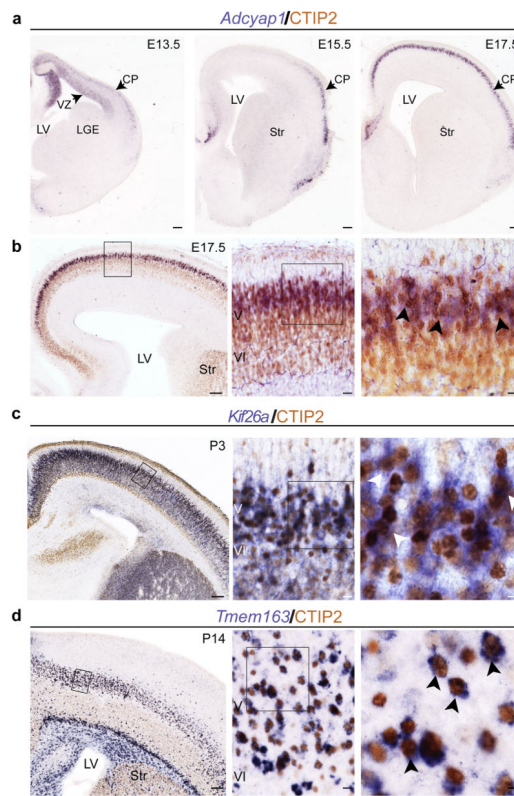


Figure 2. *Fezf2*-induced genes identify native CSMN and label subsets of the broad CTIP2-positive population in layer V. (a) *In situ* hybridization showing expression of *Adcyap1* in E13.5 cortical progenitors (left) and young postmitotic subcerebral neurons in developing cortical plate (middle and right). (b–d) Immunocytochemistry for CTIP2 combined with *in situ* hybridization for *Adcyap1*, *Kif26a* and *Tmem163* (boxed area enlarged in panel to the right). Examples of double-positive cells (arrowheads) are indicated in the right column. CP, cortical plate; LGE, lateral ganglionic eminence; LV, lateral ventricle; Str, striatum. Scale bars, 100 μm (a; b–d left panels), 20 μm (b–d, middle panels), 10 μm (b–d, right panels). The complete gene list is given in Supplementary Table 3.

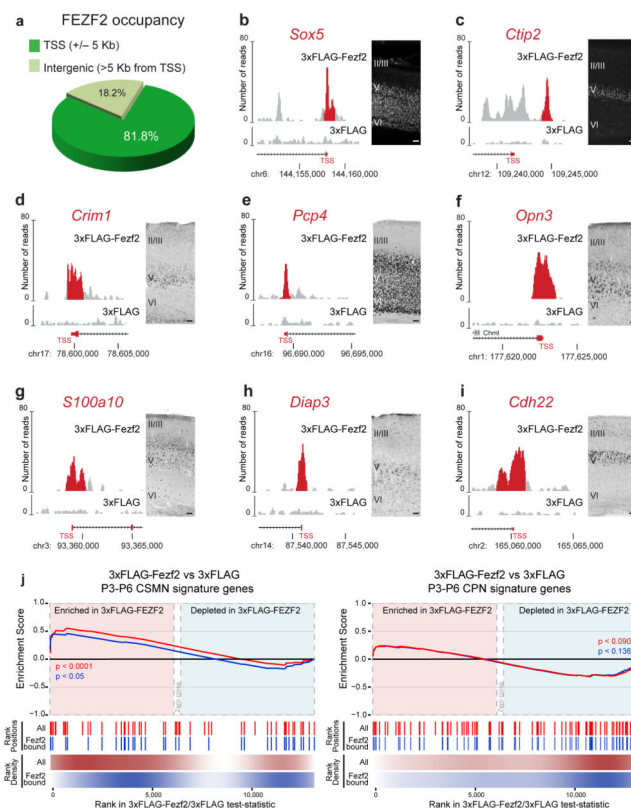
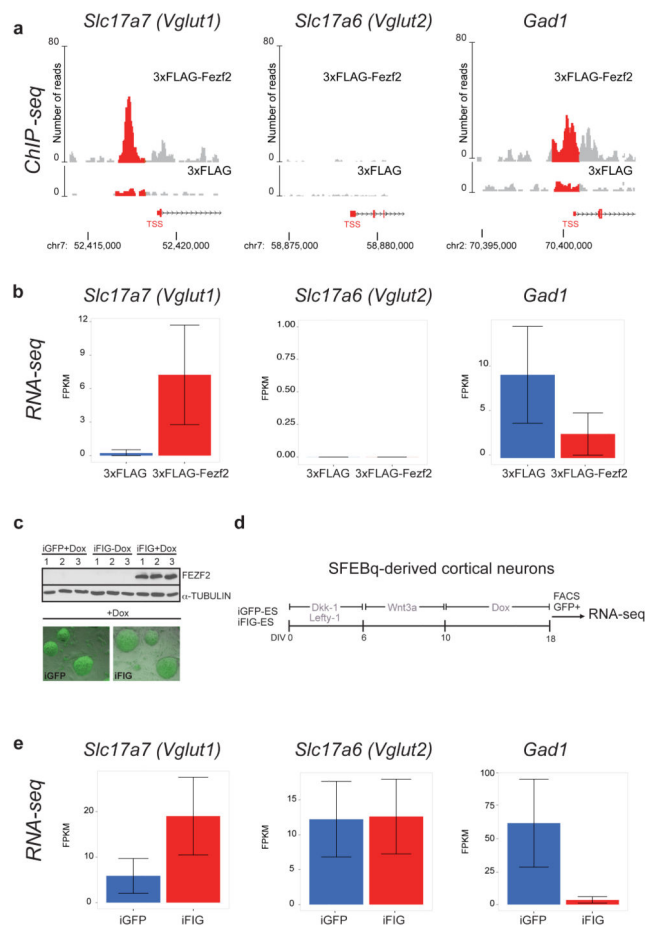
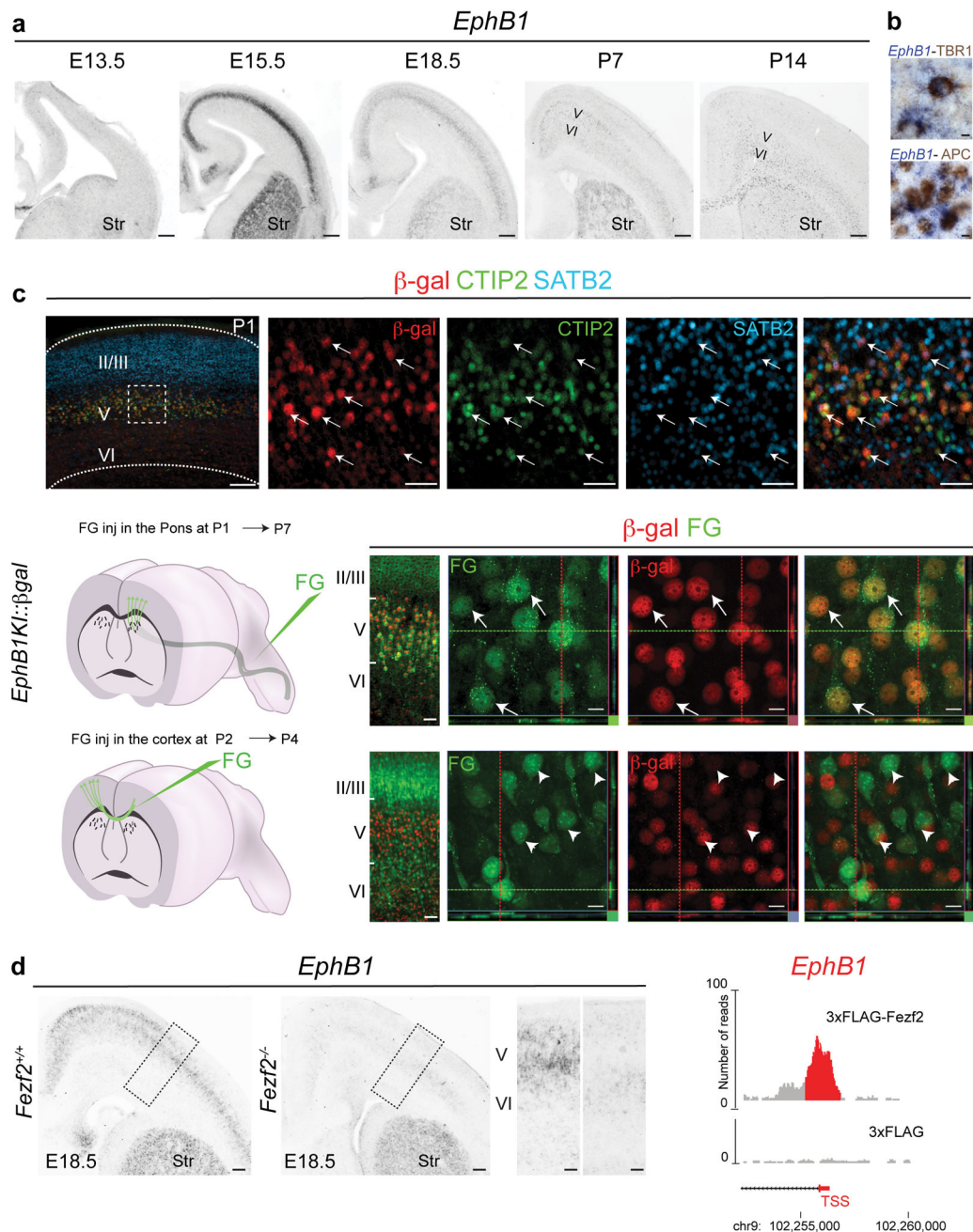


Figure 3.

Genome-wide binding analysis for *Fezf2* shows preferential association with proximal promoter regions of CSMN genes. (a) *Fezf2* binding events preferentially occur in proximity to promoter regions, within 5 kb of the TSS of annotated genes. (b–i) Examples of 3xFlag-*Fezf2* peaks at proximal promoters for early (*Sox5* and *Ctip2*; b,c), middle (*Crim1* (d) and late (*Pcp4*, *Opn3*, *Diap3*, *S100a10* and *Cdh22*; e–i) CSMN genes. *In situ* hybridizations are shown for *Crim1* (P21), *Pcp4* (P21), *Opn3* (P21), *S100a10* (P21), *Diap3* (P14) and *Cdh22* (P21). Immunohistochemistry results are shown for SOX5 (E18.5) and CTIP2 (P1). Scale bars, 100 μ m (b–i). (j) The “zero cross” area represents genes where the Cuffdiff2 test statistic is equal to 0 and no appropriate rank information is available. The color scales used for the rank density represent kernel density estimate of gene rank positions (white = 0, red/blue=max density). GSEA for CSMN and CPN signature gene sets. Signature gene sets (in red) and the corresponding subsets bound by *Fezf2* (in blue) were assessed for enrichment in *Fezf2*-overexpressing neurospheres. Both CSMN signature genes and the subset bound by *Fezf2* were significantly enriched in *Fezf2*-overexpressing neurospheres. Neither set of CPN signature genes showed significant enrichment. Source data are shown in Supplementary Table 4.

**Figure 4.**

Fezf2 promotes glutamatergic and inhibits GABAergic neurotransmitter pathways. **(a)** ChIP-seq trace shows that 3xFlag-*Fezf2* binds specifically to the promoters of *Vglut1* (*Slc17a7*) and *Gad1* but not *Vglut2* (*Slc17a6*). **(b)** RNA-seq analysis shows the effect of 3xFlag-*Fezf2* overexpression on these three genes in neural stem cells *in vitro*. **(c)** Inducible expression of single-copy *Fezf2-IRES-GFP* is shown by immunoblotting and fluorescence microscopy. Uncropped original immunoblots are shown in Supplementary Figure 10. **(d)** Schematic representation of directed differentiation protocol used to instruct embryonic stem cell (ES) differentiation into mixed populations of cortical neurons²³. **(e)** RNA-seq analysis showing the effect of *Fezf2* expression on *Vglut1*, *Vglut2* and *Gad1* in ES-derived neurons. Clones used for RNA-seq were from the differentiation of $n = 2$ independently generated iGFP and $n = 2$ iFIG lines. The error bars represent 95% confidence intervals for the Cuffdiff2 model expression estimate as defined in Methods and described in Trapnell et al., 2013.

**Figure 5.**

Fezf2 controls *Ephb1* selective expression in CSMN by direct association with the *Ephb1* promoter. (a) *In situ* hybridization for *Ephb1* in the forebrain at different stages of embryonic and postnatal development shows highest expression in developing cortical plate at E15.5, the time of initial CSMN axonal extension, and its restricted expression in developing layer V, until approximately E18.5. (b) *In situ* hybridization for *Ephb1* and immunocytochemistry for TBR1 and APC show maintained expression of *Ephb1* at postnatal stages in corticothalamic neurons (TBR1-positive in layer VI) and oligodendrocytes (APC-positive). (c) β -galactosidase immunocytochemistry in *Ephb1*

heterozygous mice at P1 shows that, within layer V, expression colocalizes with CTIP2, and not with SATB2 (arrows upper panel). The dotted rectangle area in the left panel is shown in high magnification in the four panels on the right. Retrograde labeling of ScPN from the pons of P1 *Ephb1* heterozygous mice shows colocalization of FluoroGold with β -galactosidase in layer Vb (arrows). Retrograde labeling of CPN in P2 *Ephb1* heterozygous mice shows no colocalization of FluoroGold with β -galactosidase in callosal neurons of layer Va (arrowheads). Confocal images of layer V were combined to produce three-dimensional reconstructions (Lower right panels). Sidebars represent projections along the x-z axes (right) and the y-z axes (below). **(d)** Left, *in situ* hybridization for *Ephb1* on E18.5 wild-type (left inset) and *Fezf2*^{-/-} (right inset) littermates shows that *Ephb1* levels specifically decrease in layer V of the mutant cortices. Right, DNA regions spanning 5' UTR and first exon of the *Ephb1* gene show enrichment of 3xFlag-Fezf2 binding ($q = 10^{-15}$) compared to control. Str, striatum. Scale bars, 100 μ m (**a**; **c**, far left panels), 20 μ m (**b**), 50 μ m (**c**, top right panels; **d**), 20 μ m (**c**, bottom right panels).

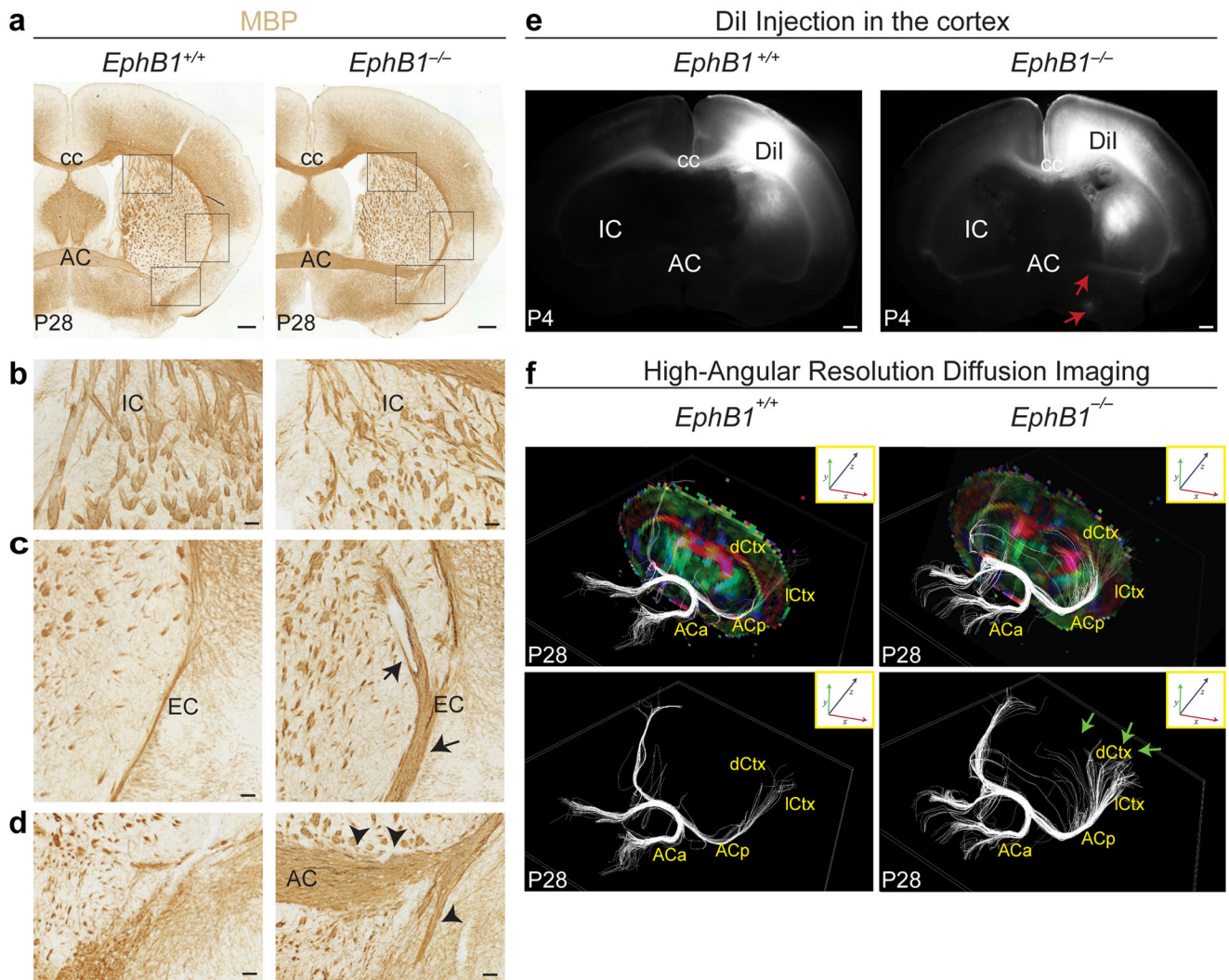


Figure 6. Cortical neurons aberrantly project through the anterior commissure in absence of EphB1. (a–d) Immunocytochemistry for myelin basic protein (MBP) on coronal sections of P28 wild-type and *Ephb1*^{-/-} littermates shows internal capsule reduction and defasciculation, accompanied by an expansion of both the external capsule (arrows) and the anterior commissure (arrowheads). (e) DiI anterograde injections in deep layers of the somatosensory cortex of P2 wild-type ($n = 3$) and *Ephb1*^{-/-} ($n = 3$) pups show axons ectopically crossing at the anterior commissure and extending ventrally and rostrally (red arrows) in the *Ephb1* mutants but not in wild-type animals. (f) Color-coded three-dimensional reconstructions based on HARDI of P28 wild-type and *Ephb1*^{-/-} littermates show distinct axon fibers originating in dorsal areas of the neocortex merging abnormally with the anterior commissure in the *Ephb1*^{-/-} (top right) compared to wild-type (top left) brains. Red represents the anterior–posterior direction; green represents the medial–lateral direction; blue represents the dorsal–ventral direction. Green arrows indicate axon tracts penetrating into the cortex. AC, anterior commissure; ACa, anterior part of the anterior commissure; ACp, posterior part of the anterior commissure; cc, corpus callosum; dCtx,

dorsal cortex; lCtx, lateral cortex; EC, external capsule; IC, internal capsule. Scale bars: 50 μm (a), 20 μm (b–d) 100 μm (e).

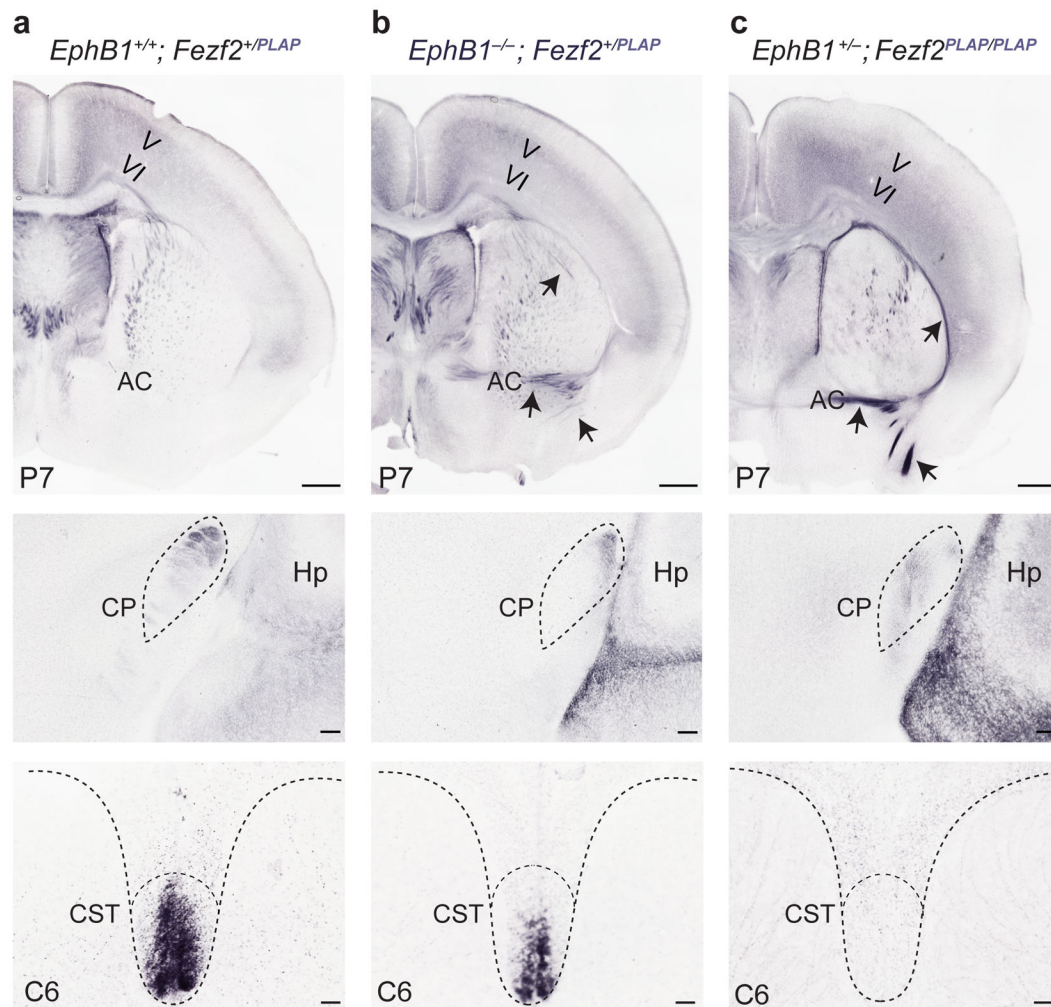


Figure 7.

Ephb1^{-/-} mice recapitulate the axon guidance phenotype observed in *Fezf2* null mutants. (a–c) PLAP-positive axons inappropriately project through the anterior commissure in *Ephb1*^{-/-} mutants (b, arrows), mimicking the *Fezf2*^{PLAP/PLAP} axonal phenotype (c, arrows). Enlargement of the anterior commissure in *Ephb1*^{-/-} mice is accompanied by a reduction of PLAP-positive axons found in the cerebral peduncle (b, middle panel) and the cervical spinal cord (b, bottom), canonical targets of the corticospinal tract in wild-type mice (a, middle and bottom). No axons are found in the cerebral peduncle or spinal cord of the *Fezf2*^{PLAP/PLAP} mice (c, middle and bottom). AC, anterior commissure; Cp, cerebral peduncle; Hp, hippocampus; CST, corticospinal tract. C6, cervical vertebra. Scale bars: 100 μm (a–c, top panels), 50 μm (a–c, middle and bottom).