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Development/Plasticity/Repair

Nuclear Factor I Coordinates Multiple Phases of Cerebellar Granule Cell Development via Regulation of Cell Adhesion Molecules

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A central question is how various stages of neuronal development are integrated as a differentiation program. Here we show that the nuclear factor I (NFI) family of transcriptional regulators is expressed and functions throughout the postmitotic development of cerebellar granule neurons (CGNs). Expression of an NFI dominant repressor in CGN cultures blocked axon outgrowth and dendrite formation and decreased CGN migration. Inhibition of NFI transactivation also disrupted extension and fasciculation of parallel fibers as well as CGN migration to the internal granule cell layer in cerebellar slices. In postnatal day 17 Nfia-deficient mice, parallel fibers were greatly diminished and disoriented, CGN dendrite formation was dramatically impaired, and migration from the external germinal layer (EGL) was retarded. Axonal marker expression also was disrupted within the EGL of embryonic day 18 Nfib-null mice. NFI regulation of axon extension was observed under conditions of homotypic cell contact, implicating cell surface proteins as downstream mediators of its actions in CGNs. Consistent with this, the cell adhesion molecules ephrin B1 and N-cadherin were identified as NFI gene targets in CGNs using inhibitor and Nfi mutant analysis as well as chromatin immunoprecipitation. Functional inhibition of ephrin B1 or N-cadherin interfered with CGN axon extension and guidance, migration, and dendritogenesis in cell culture as well as in situ. These studies define NFI as a key regulator of postmitotic CGN development, in particular of axon formation, dendritogenesis, and migratory behavior. Furthermore, they reveal how a single transcription factor family can control and integrate multiple aspects of neuronal differentiation through the regulation of cell adhesion molecules.

Key words: transcription; neuron; differentiation; axon; migration; dendrite

Introduction

During nervous system development, immature neurons are born in spatially and temporally distinct patterns and undergo specialized differentiation programs that give rise to unique populations of mature neurons. This maturation involves a series of morphological stages in which neurons migrate and elaborate axons and dendrites, resulting in appropriate synaptic interconnections. The coordination of these various events as a developmental program is a fundamental question.

Cerebellar granule neurons (CGNs) provide an excellent model for studying various aspects of neuronal development. In the postnatal cerebellum, granule neuron progenitors proliferate in the external germinal layer (EGL) and then form immature neurons within the premigratory zone (PMZ). Within the PMZ, CGNs elaborate bipolar axons that form fascicles of parallel fibers and migrate tangentially. CGNs then extend a third, radial process and migrate through the molecular layer (ML) along Bergmann glia to form the internal granule cell layer (IGL). Postmigratory CGNs complete their differentiation in the IGL by forming dendrites and synaptic connections with mossy fibers and Golgi type II neurons. A number of transcription factors regulate the lineage specification, spatial organization, and development of CGNs, including Math1 (mouse atonal homolog 1) (Ben-Arie et al., 1997; Jensen et al., 2004), En-2 (engrailed protein 2) (Millen et al., 1995), RU49/Zipro1 (zinc finger proliferation 1) (Yang et al., 1999), Pax-6 (paired box gene 6) (Yamasaki et al., 2001; Swanson et al., 2005), Notch-2 (Solecki et al., 2001), Barhl1 (Bar-H-like 1)
(Li et al., 2004), and Zic1 and Zic2 (zinc finger protein of the cerebellum 1 and 2) (Aruga et al., 1998, 2002). These trans regulators have been implicated in specific stages of CGN development, including progenitor migration from the rhombic lip (Gazit et al., 2004), proliferation and onset of differentiation within the EGL (Aruga et al., 1998, 2002; Yang et al., 1999; Solecik et al., 2001), axon formation (Yamasaki et al., 2001), and radial migration (Li et al., 2004). How various transcriptional mechanisms and downstream genes are integrated to elaborate successive stages of CGN development remains poorly understood.

Nuclear factor I (NFI) consists of a family of four genes (Nfia, Nfib, Nfjc, and Nfix) that each gives rise to multiple isoforms via alternative splicing (Gronostajski, 2000). This factor regulates target genes as homodimers and/or heterodimers and has been directly implicated in developmental and cell-specific gene regulation (Chaudhry et al., 1997; Gronostajski, 2000), including nervous system function and development. Mice lacking either Nfia or Nfib exhibit marked neurological defects, including agenesis of the corpus callosum, disruption of midline glia, tremor, and hydrocephalus (das Neves et al., 1999; Shu et al., 2003; Steele Perkins et al., 2005). Nfib-null mice also are defective in development of the basilar pons and hippocampus (Steele Perkins et al., 2005). However, specific roles and direct target genes within neurons for this family of trans regulators have been elusive.

Recently, NFI was shown to be an important transcriptional regulator of a CGN-specific gene. The α6 subunit of the GABA<sub>A</sub> receptor (Gabra6) is highly enriched in CGNs (Zheng et al., 1993), and its gene transcription is directly regulated by NFI (Wang et al., 2004). Although NFI is present in numerous neuronal subtypes as well as glia (Chaudhry et al., 1997), all four NFI genes are expressed at elevated levels in CGNs relative to other cerebellar cell types as well as cortical neurons (Wang et al., 2004). These findings suggested that the NFI family has additional important functions in CGNs besides Gabra6 regulation. Here, we demonstrate that NFI is a key regulator of postmitotic CGN development, controlling axon formation, migration, and dendritogenesis. Furthermore, we show that NFI exerts these multifaceted actions via the regulation of cell adhesion molecules.

**Materials and Methods**

**Recombinant proteins, blocking peptides, and antibody.** Mouse ephrin-B1/Fc and rat EphB1/Fc chimeric proteins were obtained from R & D Systems (Minneapolis, MN). N-cadherin antagonist peptides (HAVDI and INPISSGQ) have been described previously (Williams et al., 2000a,b). N-Acetyl-CHGVC-NH₂ was used as a negative control peptide. The GC-4 N-cadherin blocking antibody was purchased from Sigma (St. Louis, MO).

**Plasmids and cell lines.** Self-inactivating lentivirus expressing NFI dominant repressor (NFI/EnR) or the Drosophila engrailed repressor domain (EnR), both containing a C-terminal hemagglutinin (HA) tag, have been described previously (Wang et al., 2004). HA-tagged NFI/EnR and EnR fragments were released from their lentiviral expression vectors and inserted into the multiple cloning site of pMSCV2.2 internal ribosomal entry site–green fluorescent protein (GFP) to generate NFI/EnR and EnR-expressing retroviruses. Human embryonic kidney 293T cells were obtained from the American Type Culture Collection (Rockville, MD).

**Production of vesicular stomatitis virus glycoprotein-pseudotyped lentiviral and retroviral particles.** Vesicular stomatitis virus glycoprotein (VSVG)-pseudotyped lentiviruses were generated by transient cotransfection of the vector construct (15 μg), the packaging construct pCMV ΔR8.91 (10 μg), and the pMD.G VSV.G viral envelope expression vector (5 μg) into 293T cells using the Calcium Phosphate ProFection Mammalian Transfection System (Promega, Madison, WI) (Wang et al., 2004). Recombinant VSVG-pseudotyped retroviruses were generated by tran-

**Animals.** CD1 mice were used for the preparation of CGNs and for immunohistochemical studies. Nfia<sup>−/−</sup> mice and control Nfia<sup>+/+</sup> littermates were on a C57BL/6NTac background (Shu et al., 2003). Nfib<sup>−/−</sup> mice and control littermates (Steele Perkins et al., 2005) were on a C57BL/6J background, and Nfic-null mice were backcrossed onto a C57BL/6 background. All protocols used for mouse studies were approved by the Institutional Animal Care and Use Committee at the University of Massachusetts Medical School and were in full compliance with the National Institutes of Health Guide and Use of Laboratory Animals.

**Cell and tissue culture studies.** Mouse CGNs were prepared from 6-d-old pups according to previously described procedures (Wang et al., 2004). Briefly, dissected cerebella were digested with 1% trypsin and 1 mg/ml DNase (Sigma) in calcium–magnesium-free PBS, pH 7.4, at room temperature for 3 min. CGNs were prepared by mechanical trituration and enriched by Percoll (Sigma) gradient centrifugation. The granule cell fraction was further purified by prepping the cells on poly-<i>γ</i>-lysine-coated Petri dishes at 33°C for 1 h. For assays of neurites and dendrites in dissociated cultures, cells were plated at a density of 5 × 10⁵ cells/cm² onto poly-<i>γ</i>-lysine/laminin-treated chamber slides in Neurobasal medium containing B-27 serum-free supplement, 2 mM l-glutamine, 100 U/ml penicillin, and 100 μg/ml streptomycin (pen-strep) (Invitrogen, Grand Island, NY) and 0.45% D-glucose. Reaggregate cultures were prepared as described by Gao et al. (1991). Briefly, reaggregates were formed by incubating purified CGN progenitors for ~2 h on uncoated tissue culture dishes in MEM (Sigma) containing 10% fetal bovine serum, 0.45% D-glucose, pen-strep, and 2 mM l-glutamine at a cell density of 4 × 10⁶/ml. Reaggregates were then washed and cultured in Neurobasal/B27 medium on poly-<i>γ</i>-lysine/laminin-treated chamber slides at 37°C/5% CO₂.

**Migration assays.** Migration assays were performed using polycarbonate membrane filters (pore size, 5 μm) inserted into 24-well Boyden chamber-type Transwells (Costar, Cambridge, MA). The undersurfaces of the membrane inserts were coated with laminin at 20 μg/ml in PBS for 2 h at 37°C. Granule cells (10⁶) were added in Neurobasal/B27 medium to the upper chamber and allowed to migrate into the coated undersurface at 37°C for 16 h. Cells remaining on the upper membrane surface were wiped off, and the inserts were fixed with 4% paraformaldehyde and stained with bisbenzimide. For viral studies, cells were transduced with lentivirus during regreaggregate formation in suspension cultures. The next day, cells were enzymatically dissociated and added to Transwell plates as above. Cerebellar slice cultures were performed as described previously (Wang et al., 2004) using either rat or mouse cerebellar slices. Briefly, 200 μm sections were cut from postnatal day 7 (P7) mouse cerebella, embedded in 5% agarose, and incubated in MEM with 10% FBS, 2 mM l-glutamine, pen-strep, and 0.45% D-glucose at 35°C. Slices were then placed on Transwell membrane inserts (Corning) and cultured in Neurobasal/B27 medium at 37°C/5% CO₂. GFP-positive (GFP<sup>+</sup>) cells were analyzed solely within the cerebellar vermis.

**Cell cultures (both dissociated and reaggregate) were infected with lentivirus at a multiplicity of infection of 2–4 as described previously (Wang et al., 2005), resulting in >90% infection of CGNs. For slice experiments, the retroviral titers was reduced to 4 × 10⁶ infectious units per well to permit visualization of individual GFP<sup>+</sup> cells.**

**Bromodeoxyuridine incorporation assays.** CGN reaggregates were allowed to form in the presence of lentiviral expression vectors for 20 h. They were then labeled with 10 μM bromodeoxyuridine (BrDU) (Roche Applied Science) for an additional 24 h. Cells were then dissociated into single cells, allowed to attach to chamber slides treated with poly-<i>γ</i>-lysine and laminin, and then fixed with 4% paraformaldehyde. DNA denaturation was accomplished using 2N HCl for 30 min at 37°C, followed by neutralization with two changes of 0.1 M sodium borate buffer, pH 8.5, for 10 min each. BrDU-labeled cells were detected using anti-BrDU antibody (1:5000, Mab 3518; Chemicon, Temecula, CA).

**Isolation of RNA and reverse transcription-PCR.** Total RNA was extracted from tissues and cells using TRI reagent (Sigma). First-strand
cDNAs were synthesized with oligo-dT_{12-18} primers or random hexamers using the SuperScript reverse transcription (RT)-PCR system (Invitrogen). Transcripts were assayed by semiquantitative RT-PCR using the following primers: ephrin B1 (forward primer, TTGTTGCGATAGTTGTCGC; reverse primer, GCTGTTGCAAATCTCGC) and N-cadherin (forward primer, TTGACTTTGAAAGAAAAGG; reverse primer, TGATGCTGCCATCAGC). 18S ribosomal RNA was assayed for normalization purposes. Other primer sequences are available on request.

**Request for reagents.**

Di labeling of parallel fibers and granule cells. P17 cerebella were fixed with 4% paraformaldehyde, embedded in 5% agarose, and cut into 400 μm coronal sections throughout the vermis using a vibratome. Labeling with the lipophilic dye DiI was performed as described previously (Soha et al., 1997). Briefly, small DiI crystals were implanted into the ML at the midline, and sections were incubated in PBS at 37°C for 4 d. Sections (80 μm) were then prepared and analyzed for parallel fiber extension in the ML and the presence of CGN dendrites within the IGL using fluorescein microscopy.

**Immunofluorescence.** Cerebella from wild-type P6 mice and heads of P17 Nfia and Nfic knock-out mice and embryonic day 18 (E18) Nfib-null mice were fixed in 4% paraformaldehyde in phosphate buffer, pH 7.4, imbedded with 30% sucrose, and cryosectioned at 15 or 50 μm thickness as described previously (Wang et al., 2004). Primary CGNs or slice cultures were fixed with 4% paraformaldehyde for 30 min at room temperature. Immunofluorescence was performed as in the previous studies (Wang et al., 2004). Samples were pretreated with 10% normal goat serum (In-vitrogen), 0.05% Tween 20, and 0.3% Triton-X-100 in PBS at room temperature for 1 h, followed by incubation with primary antibodies at 4°C overnight and then with secondary antibodies at room temperature for 1 h. The following antibodies were used: anti-NFIA (1:10,000), anti-NFIB (1:1,000), and anti-NFIX (1:1,000) (Active Motif, Carlsbad, CA); anti-NFI-C (1:500; Dr. N. Tanese, New York University, New York, NY); anti-HA (1:200; Bethesda Laboratory, Montgomery, TX) and for colabeling studies (1:400; Cell Signaling Technology, Danvers, MA); anti-GFP monoclonal antibody, anti-neuronal-specific nuclear protein (NeuN), and anti-neurogenic differentiation (NeuroD) (each at 1:1000; Chemicon); anti-β tubulin isotype III monoclonal antibody (1:1000; Sigma); anti-Math1 (1:500; Dr. J. Johnson, University of Texas Southwestern, Dallas, TX); anti-cyclin D2 (1:200; Santa Cruz Biotechnology, Santa Cruz, CA); pan-axonal neurofilament monoclonal antibody (SM-132, 1:1000); anti-ephrin B1 polyclonal antibody (1:200; Santa Cruz Biotechnology), anti-mouse N-cadherin (1:50; Developmental Studies Hybridoma Bank, University of Iowa, Iowa City, IA); anti-S100β (1:200; Sigma); and anti-calbindin-D28k monoclonal antibody (1:1000; Sigma). Secondary antibodies were conjugated to cyanine 3, AlexaFluor 594 (Invitrogen), F78, or FITC. For studies of knock-out mice, sections were stained with 1 μg/ml bisbenzimide (Sigma) after treatment with secondary antibodies.

**Chromatin immunoprecipitation assays.** Chromatin immunoprecipitation (ChIP) assays were performed as described previously (Wang et al., 2004). Briefly, 6 d in vitro (DIV) CGN cultures or nuclei isolated from P15 mouse cerebella were crosslinked with 1% formaldehyde for 5 min at 37°C. Immunostaining confirmed that >85% of nuclei from P15 cerebella were positive for NeuN, a granule neuron marker in the mouse cerebellum (Weyer and Schilling, 2003). Cells or nuclei were then collected and lysed in SDS buffer (1% SDS, 10 mM EDTA, 50 mM Tris-HCl, pH 8.1, 1 mM phenylmethylsulfonyl fluoride, and 1 μg/ml protease inhibitor mixture mix). Chromatin was sonicated to an average length of 600 bp, and NFI antibody was added to the lysate. After overnight incubation at 4°C, immune complexes were collected with protein A-Sepharose (GE Healthcare, Little Chalfont, UK). The precipitated chromatin was treated with proteinase K and DNase-free RNase A, followed by overnight incubation to reverse the crosslinking. DNA was extracted and used as template for PCR assay of the mouse ephrin B1 and N-cadherin promoters. PCR primer sequences were as follows: ephrin B1 (forward, 5'-TGG-TAGCTCCTATTATGTG-3'; reverse, 5'-TGGAGGTTCTCAGAAGGTC-3'); N-cadherin (forward, 5'-TGGCTGGCTGTGTTTCTC-3'; reverse, 5'-GCTAAATTGGCATCAGTGA-3').

**Image analysis and statistical tests.** Images of immunostaining or cell cultures were obtained by computer-assisted microscopy using Nikon (Tokyo, Japan) Optiphot or Zeiss (Oberkochen, Germany) Axioverflow inverted microscopes. Images were obtained and analyzed using Axio Vision 4.4 (Zeiss) software. For reaggregate studies, only cellular clusters having similar sizes were compared. For granule cell migration, CGN cell bodies were identified by staining either for HA (for lentiviral experiments) or using bisbenzimide, and their distances from the reaggregate perimeter were determined as described previously (Hirotsune et al., 1998). Neurites were identified by β-III tubulin staining. To analyze neurite length, reaggregates were visually divided into quadrants, and the median neurite length for each quadrant was determined by measuring the longest and shortest neurites from the edge of the cell body cluster. The four quadrant values were then averaged to obtain the overall median neurite length for each reaggregate. Neurite density was determined by measuring the number of neurites extending beyond the edge of the cluster divided by the diameter of each reaggregate. In antibody-treated reaggregate cultures, outgrowth of their nonradial neurites was quantified by measuring the area occupied by all neurites and then normalizing to the area of each cell cluster. A minimum of 20 reaggregates were examined for each experimental treatment. Measurements of axon and dendrite length in dissociated CGN cultures were performed on captured images as described previously (Ronn et al., 2000) from 200–300 neurons per experiment. Dendrite density was defined as the average number of dendrites per cell.

**Results**

**NFI is expressed in CGNs throughout their postmitotic differentiation.**

Previous studies demonstrated that NFI proteins are highly enriched in mature CGNs within the adult mouse cerebellum (Wang et al., 2004). To address their potential function during CGN development, we examined NFI protein expression in the early postnatal cerebellum. Immunohistochemical analysis revealed that NFIa, NIFB, and NFIX are readily detected within the IGL of P6 mice (Fig. 1). NFIc immunostaining was too weak to be reliably discerned (data not shown). NFIa, NIFB, and NFIX were localized to the nuclei of granule cells, consistent with a
transcriptional role for these proteins. Furthermore, nuclear staining for these three family members was detected in fusiform cells within the ML, indicative of migrating postmitotic CGNs that have departed the EGL en route to the IGL (Fig. 1). Within the EGL, intense nuclear localization was observed in its deeper aspect, the PMZ, in which immature postmitotic granule neurons first appear. Staining within the outer proliferative region of the EGL was generally faint and nonspecific (Fig. 1), similar to a negative control (data not shown). Thus, NFI proteins are upregulated in CGNs as they become postmitotic and subsequently undergo radial migration and final maturation within the IGL.

**NFI transactivation is required for neurite outgrowth by CGNs**

The expression pattern for NFI proteins suggested that they might function during several stages of postmitotic CGN development, including axon formation and migration. Reaggregate cultures of CGNs were used to examine this question because they permit assessment of both of these processes. Purified CGN progenitors were allowed to form reaggregates in suspension cultures for ~20 h, followed by plating on a laminin-coated surface. After attachment, reaggregates initially extend axon-like neurites, followed by subsequent migration of cell nuclei along these extended processes (Gao et al., 1991; Bix and Clark, 1998). To probe NFI function, CGN progenitors were infected during reaggregation with lentiviral vectors that expressed either a dominant repressor form of NFI (NFI/EnR) or the Drosophila engrailed repressor domain alone (EnR) (Wang et al., 2004). As reported previously (Wang et al., 2005), lentiviral vectors quantitatively transduce cultured CGNs with high efficiency with a rapid onset of protein expression (within 6 h) without affecting cell survival or differentiation.

The NFI dominant repressor dramatically impaired neurite extension from CGN reaggregates relative to EnR-transduced controls (Fig. 2A). Both reaggregate neurite density and neurite length were reduced threefold by NFI/EnR ($p < 0.001$ for each), with neurites often highly fasciculated and extending in an aberrant nonradial manner (Fig. 2A). Immunostaining for phosphorylated neurofilament confirmed that neurites extending from reaggregate cultures were axons (data not shown). Furthermore, CGN migration was greatly inhibited in NFI/EnR- or EnR-expressing reaggregates (Fig. 2B). Very few migrating cells were observed, and these were present at relatively short distances. In contrast, EnR-expressing CGNs migrated considerable distances from reaggregates (Fig. 2B), identical to mock-infected cultures (data not shown).

CGN progenitors continue to proliferate in reaggregate suspensions (Gao et al., 1991). Because lentiviral expression occurred during re-aggregate formation in the above experiments, it was possible that the NFI dominant repressor affected CGN progenitor cell division and thereby indirectly altered their subsequent differentiation. However, addition of BrdU to reaggregates revealed that the NFI dominant repressor did not significantly affect the number of CGN progenitors undergoing S phase, and there also was no effect of NFI/EnR on the percentage of surviving cells (data not shown). Thus, repression of NFI transactivation in CGNs reflects specific interference with the differentiation process per se. This is consistent with the postmitotic up-regulation of NFI protein observed in these cells (Fig. 1). Furthermore, NFI/EnR did not suppress expression of NeuroD or NeuN (data not shown), both of which are differentiation markers for granule neurons within the cerebellum (Miyata et al., 1999; Weyer and Schilling, 2003). Thus, onset of CGN differentiation was not generally affected.

CGNs elaborate neurites in dissociated cell cultures as part of a cell-intrinsic program (Powell et al., 1997). Interestingly, axons showed no significant differences in length or general morphology when dissociated CGNs were transduced with NFI/EnR or EnR proteins (Fig. 2C and data not shown). One difference between reaggregate and dissociated cell experiments was that, in the former case, CGN progenitors were transduced with lentivirus for 20 h before plating on laminin, whereas dissociated cultures were infected during the plating step. It was therefore possible that the differential effects of NFI disruption between the two cultures reflected a more rapid and efficient repression of NFI function within reaggregates. To address this, reaggregates were transduced with NFI/EnR- or EnR-expressing lentiviruses in suspension cultures for 20 h, and then they were disaggregated into individual cells before plating on laminin. No major differences in either the number of axon-bearing cells or axon length were observed under these circumstances (data not shown). Thus, NFI regulation of axon outgrowth by CGNs was only detected under conditions of homotypic cell contact present in reaggregate cultures.

**NFI controls dendrite formation in maturing CGNs**

In addition to its importance for earlier stages of CGN differentiation within the PMZ/ML, NFI also is required for the expression of the Gabra6 gene within the IGL (Wang et al., 2004). We therefore investigated whether NFI regulated dendritogenesis, which is also a late differentiation event occurring in the IGL. Dendrite formation in NFI/EnR-treated reaggregate cultures could not be discerned on individual CGN cell bodies within reaggregates. Dissociated CGNs were therefore transduced on 1 DIV with either NFI/EnR or EnR lentiviruses, and dendrite formation was examined on 4 DIV using mitogen-activated protein 2 (MAP2) as a marker. Suppression of NFI transactivation caused a marked reduction in dendritogenesis by CGNs (Fig. 2D). The NFI dominant repressor greatly reduced both dendrite length (4.0-fold; $p < 0.001$) as well as the number of dendritic processes per cell (2.4-fold; $p < 0.001$). Thus, in contrast to its effects on axon formation, NFI regulation of dendritogenesis occurred in dissociated cultures.
Disruption of NFI function alters differentiation of CGNs in situ

We extended the above studies by examining the role of NFI in regulating CGN development within their native cellular environment using retroviral infection of cerebellar slices. Bicistronic vectors expressing NFI/EnR or EnR proteins together with GFP were used to assess cell body migration and parallel fiber formation within transduced neurons. Although a high percentage of EnR-transduced CGNs emigrated from the EGL to the ML and IGL, most NFI/EnR-expressing CGNs remained within the EGL region, and only a small fraction reached the IGL (Fig. 3A). Thus, departure of CGNs from the EGL was markedly inhibited by disruption of NFI transactivation.

In coronal sections infected with the control EnR virus, GFP-expressing CGNs extended parallel processes within the EGL/ML region (Fig. 3B). Axons of CGNs expressing the NFI dominant repressor were reduced in length 3.2-fold ($p < 0.001$). Furthermore, these processes were defasciculated, with an ~10-fold increase in fibers oriented toward the EGL and/or IGL (Fig. 3B). Thus, consistent with cell culture studies, NFI regulates the extension and orientation of parallel fibers as well as events required for CGN migration to the IGL in situ. Because very few CGNs reached the deep IGL in cerebellar slice cultures, it was not possible to reliably examine the effects of NFI inhibition on dendrite formation in this preparation.

NFI regulates CGN migration

The NFI dominant repressor markedly inhibited migration of CGNs in both reaggregate and slice cultures (Figs. 2B, 3A). However, these experiments did not address whether NFI has a direct role in this process. For example, migration from reaggregates occurs along extended axons, and NFI/EnR severely inhibited axon formation in these cultures. Similarly, emigration of CGNs from the EGL to the IGL within cerebellar slices likely depends on progression of earlier events within the EGL/PMZ, including parallel fiber extension, which is disrupted in slice cultures by the NFI dominant repressor. To address the role of NFI in CGN migration per se, we performed modified Boyden assays in which dissociated CGNs were cultured on membranes coated on the opposing side with laminin. Expression of NFI/EnR inhibited CGN migration by 50% relative to EnR-transduced control cells ($p < 0.01$).

Thus, NFI function also is necessary for CGN migratory behavior in addition to axon and dendrite formation.

CGN development is altered at multiple steps in Nfia knock-out mice

Based on the above findings, we examined whether CGN differentiation was altered in the cerebella of mice lacking NFI family members. Knock-out mice have been previously generated for Nfia, Nfib, and Nfic (das Neves et al., 1999; Grunder et al., 2002; Wang et al., 2007).

Figure 2. NFI regulates process formation and migration. A, The NFI dominant repressor impairs neurite outgrowth from CGN reaggregates. CGNs were infected with lentivirus expressing HA-tagged NFI/EnR or EnR during cell reaggregation and examined 20 h after plating for neurite extension using β-III tubulin immunostaining. B, NFI is required for migration of CGNs from reaggregates. Top, Reaggregates were stained for HA to identify transduced CGN nuclei. Bottom, The distribution of HA-negative neurons migrating from reaggregates. C, The NFI dominant repressor does not significantly affect neurite outgrowth in fully dissociated CGN cultures. Dissociated CGNs were infected with NFI/EnR or EnR virus on 0 DIV and stained for β-III tubulin on 2 DIV. D, NFI is required for dendrite formation in dissociated CGN cultures. CGNs were transduced with NFI/EnR or EnR lentiviruses and then stained for MAP2. Both dendrite length and number per cell were significantly reduced by the NFI dominant repressor relative to control. * $p < 0.001$, significantly different from control. Scale bars: A–C, 100 μm; D, 50 μm.
Figure 3. NFI transactivation is required for parallel fiber extension and CGN migration in situ. A, The dominant repressor impairs CGN migration from the EGL to the IGL. Top, Representative images of GFP + CGNs infected with NFI/EnR or EnR retroviruses in sagittal cerebellar slices. Bottom, The percentage of GFP + cells present within the EGL, ML, and IGL expressed as the mean. *p < 0.05, **p < 0.001, significantly different from EnR control. B, Parallel fiber extension and orientation are markedly altered by inhibition of NFI. Top, GFP + immunostaining of parallel fibers within the EGL/ML region of coronal cerebellar slices transduced with NFI/EnR or EnR retroviruses. Essentially all GFP + cells detected within this region contained bipolar processes typical of forming parallel fibers. Note that labeled cells lying below the indicated EGL region derive from an apposing EGL. Bottom, The angle of divergence (θ) from the parallel orientation, expressed in degrees, for GFP + axons. *p < 0.001, significantly different from control. Scale bars, 50 μm.

Steele Perkins et al., 2003, 2005). Nfib-null mice die at birth as a result of lung defects (Grunder et al., 2002; Steele Perkins et al., 2005), whereas Nfia-null mice can survive to approximately P20 (das Neves et al., 1999; Wang et al., 2004). Examination of cerebella from P17 Nfic-null mice showed no obvious phenotypes (data not shown), likely reflecting lower expression of this family member in the CNS (Chaudhry et al., 1997). Because postmitotic differentiation of CGNs is primarily postnatal, we initially examined Nfia mice to examine possible alterations in CGN maturation. In particular, we focused on P17 mice in which CGNs elaborate dendrites and have completed parallel fiber extension and radial migration. Analyses were limited to the cerebellar vermis throughout. Grossly, P17 Nfia−/− mice exhibited a deformed foliation pattern relative to wild-type animals (Fig. 4A). Fissures were generally reduced in depth, and, in many instances, lobules appeared either stunted or poorly defined. For example, anterior lobules (I–V) were markedly underdeveloped, and segregation of lobules VI and VII appeared abnormal (Fig. 4A). Altered foliation has been associated previously with defects in postmitotic CGN development, including altered migration (Schwartz et al., 1997; Li et al., 2004).

Dil labeling (Soha et al., 1997) was used to assess both parallel fiber development and CGN dendrites in Nfia-null cerebellum. Within the ML of wild-type P17 cerebella, CGN axons exhibited a typical bipolar and parallel phenotype and extended over a considerable distance (Fig. 4B). However, ML axons were extremely short throughout both anterior and posterior aspects of the vermis in Nfia-null mice (Fig. 4B). Those processes that were sufficiently extended were typically misoriented, frequently extending at nearly a 90° angle toward the meninges or the IGL (Fig. 4B). Furthermore, dendrite formation was dramatically impaired on CGNs throughout the IGL of more anterior lobules in P17 Nfia−/− vermes relative to wild-type (Fig. 4C). Dendrites were generally not evident on these mutant CGNs and those few detected were extremely short, whereas CGNs in wild-type mice possessed typical claw-like dendrites.

Migration of CGNs from the EGL to the IGL is normally completed in mice by P17, and no migrating cells were identified in the ML of wild-type mice using NeuN and NeuroD as CGN markers (Fig. 4D and data not shown). However, although many CGNs completed migration to the IGL in P17 Nfia-null mice, numerous CGN nuclei were observed in the ML (Fig. 4D and data not shown). These exhibited fusiform shapes typical of migrating CGNs, indicating retarded migration in Nfia knock-out mice. Furthermore, a residual layer of NeuroD + and NeuN + cells (on average, two to three cells thick) was still present in the EGL/PMZ of mutants at this age, which was absent in +/+ mice (Fig. 4D and data not shown). No Math1 + or cyclin D2 + cells were detected within the residual EGL (data not shown), indicating the absence of CGN progenitors in this region. Thus, loss of Nfia disrupts parallel fiber extension and orientation, dendritogenesis, and departure of postmitotic CGNs from the EGL/PMZ.

The number and organization of calbindin + Purkinje neurons were similar in P17 wild-type and Nfia knock-out mice (Fig. 4E), indicating that alterations in CGNs were not attributable to loss of or gross changes in Purkinje cells. Subtle decreases in the length of Purkinje dendritic branches were apparent in the upper margins in Nfia-null mice relative to wild type (Fig. 4E). This may reflect reduced maturation of parallel fibers within this region of the ML and the presence of a residual EGL containing CGNs. Finally, S100 immunostaining revealed that the density and distribution of Bergmann glia also were unaltered in Nfia-null mice (data not shown).

CGN axon formation is disrupted in Nfib-null mice

Perinatal death of Nfib knock-out mice precluded an extensive analysis of CGN differentiation in these animals. However, E18
**Figure 4. Abnormal cerebellar development in Nfi-null mice.**

A, Bisbenzimide staining of sagittal sections through the vermis of P17 mice showing altered foliation in Nfi mutant mice. Top, Nfia+/+. Bottom, Nfia−/−. Lobules are labeled with roman numerals. B, DiI-labeled parallel fibers within cerebellar coronal sections from wild-type and Nfia-null animals. Left, Arrowheads indicate parallel fibers within the ML extending away from the DiI crystal inserted at the midline. Axon extension is reduced throughout the cerebellar vermis of Nfia-null mice compared with wild-type mice. Right, Higher magnification of the ML showing dramatically disoriented fibers in Nfia-null mice (data not shown). These findings indicated that ephrin B1 and N-cadherin lie downstream of NFI in differentiating granule neurons.

Based on these results, we determined the expression of ephrin B1 and N-cadherin in CGNs within the developing mouse cerebellum. In P6 mice, ephrin B1 was predominantly localized to the PMZ and ML and also was present in the IGL (Fig. 6B). N-cadherin was detected throughout the cerebellum of P6 mice, including the EGL, ML, and IGL, as well as white matter (Fig. 6B). These expression patterns mirror those reported previously for the two proteins in the developing rat cerebellum (Doherty et al., 1991; Moreno-Flores et al., 2002) and are consistent with their functioning throughout different stages of CGN maturation.

We next examined whether the expression of ephrin B1 and N-cadherin in the cerebellum was affected by disruption of Nfi−/− mice were previously reported to have foliation defects within the forming cerebellum (Steele Perkins et al., 2005), possibly reflecting altered early development of granule neurons. We therefore examined whether parallel fiber formation was disturbed in Nfib-null mice using the axonal marker phospho-neurofilament. Immunostaining was greatly reduced within the EGL/PMZ region in Nfib-deficient mice (Fig. 5). This did not reflect a general disruption of CGN differentiation, because NeuroD protein was readily detected in the EGL/PMZ region of both Nfib+/+ and Nfib−/− mice (data not shown). Thus, loss of NFIB expression alters axon formation by CGNs in the early cerebellum. Together, knock-out mouse results indicated that NFI proteins, in particular NFIA and NFIB, are important for normal development of CGNs, including parallel fiber formation within the EGL/PMZ and dendritic formation in the IGL.

NFI regulates the expression of cell adhesion molecules in postmitotic CGNs

The fact that NFI regulation of axon formation occurred in CGN reaggregates suggested that its actions might be mediated by cell adhesion molecules, at least in part. We therefore examined the expression of several cell adhesion and guidance molecules implicated in axonogenesis and/or migration after treatment of CGNs with the NFI dominant repressor. The mRNAs for two of these, N-cadherin and ephrin B1, were markedly downregulated (approximately threefold) in CGN cultures by NFI/EnR, whereas those for several others (ephrin B2, EphB1, slit2, integrin β1, and L1) were unaffected (Fig. 6A and data not shown). These findings indicated that ephrin B1 and N-cadherin lie downstream of NFI in differentiating granule neurons.

Based on these results, we determined the expression of ephrin B1 and N-cadherin in CGNs within the developing mouse cerebellum. In P6 mice, ephrin B1 was predominantly localized to the PMZ and ML and also was present in the IGL (Fig. 6B). N-cadherin was detected throughout the cerebellum of P6 mice, including the EGL, ML, and IGL, as well as white matter (Fig. 6B). These expression patterns mirror those reported previously for the two proteins in the developing rat cerebellum (Doherty et al., 1991; Moreno-Flores et al., 2002) and are consistent with their functioning throughout different stages of CGN maturation.

We next examined whether the expression of ephrin B1 and N-cadherin in the cerebellum was affected by disruption of Nfib genes. Immunostaining for both proteins was reduced in the ML and IGL of P17 cerebella from Nfib−/− mice relative to wild-type controls (Fig. 6C). Cerebellar expression of another cell adhesion molecule, sialylated-NCAM, was unaltered in Nfib knock-out mice (data not shown). Western blotting confirmed that both
The preceding cell culture studies strongly implicated ephrin B and N-cadherin as critical mediators of NFI actions in axon formation, migration, and dendritogenesis in CGNs. We therefore examined whether ephrin B1-Fc had any effect on CGN migration under these conditions (data not shown). Because migration assays used dissociated CGNs, it was possible that endogenous ephrin B function was masked by reduced cell–cell contact and ephrin–Eph trans interactions. We therefore examined whether ephrin B1-Fc affected CGN migration in the presence of homotypic cell contacts using reaggregate cultures. Under these conditions, soluble ephrin B1-Fc caused a partial reduction of CGN migration (−20%; p < 0.05) compared with NFI/EnR, which inhibited this process by 50% (data not shown). Thus, both N-cadherin and ephrin B contribute to NFI regulation of CGN migration.

Finally, cell adhesion molecule inhibitors also suppressed dendritic formation by dissociated CGN cultures (Fig. 8G,H), with N-cadherin peptide antagonists being more effective. Both the number of dendritic processes as well as their lengths were inhibited. Soluble EphB1–Fc had no effect on dendrite extension (Fig. 8G).

**N-cadherin and ephrin B function in situ**

The preceding cell culture studies strongly implicated ephrin B and N-cadherin as critical mediators of NFI actions in axon formation, migration, and dendritogenesis in CGNs. We therefore addressed whether they regulated these events *in situ* using cerebellar slices infected with GFP-expressing retrovirus. Both ephrin B1-Fc and N-cadherin peptides inhibited CGN egress from the EGL to the IGL in sagittal slices (Fig. 9A,B). In coronal slices, soluble ephrin B1–Fc repressed extension of newly forming axons.
Figure 6. Expression of cell adhesion molecules is regulated by NFI in CGNs. A, Ephrin B1 and N-cadherin transcripts are specifically reduced by the NFI dominant repressor in CGNs. Semiquantitative RT-PCR analysis of cell adhesion or guidance molecule mRNAs in CGN cultures transduced with NFI/EnR or EnR lentiviruses. Samples were analyzed using threefold serial dilutions. 18S ribosomal RNA was assayed as a loading control. B, Immunostaining for ephrin B1 (top) and N-cadherin (bottom) in the vermis of P6 mouse cerebella. oEGL, Outer EGL region. C, Cell adhesion molecule immunostaining is reduced in Nfia mutant mice. Sagittal sections through the vermis of P17 Nfia+/−/− and Nfia−/−/− mice were stained with antibodies against ephrin B1 (top) and N-cadherin (bottom). D, Western blot analysis of ephrin B1 and N-cadherin in mouse cerebella showing reduced expression of these proteins in Nfia-deficient mice. E, Ephrin B1 immunostaining (top) is reduced in E18 Nfib−/− mouse cerebellum relative to Nfib+/−/+ controls. The arrowhead indicates labeled cells within the EGL of wild-type mice. Bottom, Bisbenzimide staining of the same sections. Scale bars, 50 μm.

Figure 7. In vivo binding of NFI to cell adhesion molecule promoters in postmitotic CGNs. A, Locations of consensus NFI binding sites within the mouse ephrin B1 and N-cadherin promoters. The approximate positions of PCR primers within the 5′ regions for ChIP assays are indicated by arrows. B, ChIP assay showing that NFI proteins are bound in vivo to the ephrin B1 and N-cadherin promoter regions in 6 DIV CGN cell cultures. C, NFI binds to the ephrin B1 and N-cadherin promoters in intact mouse cerebellar tissue. ChIP assay using nuclei prepared from P15 cerebella, >85% of which derive from CGNs based on NeuN and NeuroD immunostaining. Mouse brain factor 2 genomic sequences were used as a negative control. Pre, Preimmune serum.

(Fig. 9C), decreasing their average length approximately fourfold (p < 0.001). Similarly, N-cadherin antagonist peptides inhibited CGN axon formation (Fig. 9D) approximately threefold (p < 0.001). Furthermore, both inhibitors altered the parallel orientation of newly extending axons (Fig. 9C,D). Antagonism of N-cadherin caused a more dramatic defasciculation that was similar in extent to that observed with the NFI dominant repressor (Fig. 3A).

Dendrite formation by CGNs within the IGL of cerebellar slices also was disrupted by these antagonists (Fig. 9E,F). Both N-cadherin peptides and ephrin B1-Fc reduced the number of primary dendritic arborizations approximately threefold (p < 0.01), and these processes tended to be thinner.

Discussion

Previous studies involving knock-out mice have directly implicated NFIA and NFIB in CNS development, in particular in the formation of midline glia (das Neves et al., 1999; Shu et al., 2003; Steele Perkins et al., 2005). Recently, NFIA also was found to be required for gliogenesis within the developing chick spinal cord (Deneen et al., 2006). NFIB also has been directly implicated in hippocampal and pons development (Steele Perkins et al., 2005). However, knowledge of the broader
functions and targets for NFI proteins in developing neurons has remained limited. Our cell culture, in situ and Nfi knock-out studies indicate that NFI family members are central transcriptional regulators of CGN development, their function being important for several major aspects of their postmitotic maturation: axon formation in the PMZ, migration, and dendrite formation within the IGL. Furthermore, we demonstrated previously that NFI directly regulates expression of the extrasynaptic neurotransmitter receptor subunit Gabra6, present in the IGL (Wang et al., 2004). Thus, a single transcription factor family can act during postmitotic neuronal differentiation during multiple maturation stages.

Another finding is that cell adhesion molecules are an important class of NFI targets and downstream mediators in maturing CGNs. This observation helps to explain how NFI regulates diverse aspects of CGN development. N-cadherin and ephrin B1 appear to have broad functions as cell adhesion molecules in developing neurons (Barami et al., 1994; Riehl et al., 1996; Gao et al., 1999; Palmer and Klein, 2003; Zhu and Luo, 2004; Halloran and Wolman, 2006). In CGN cultures, both proteins can promote axon formation when used as substrata for outgrowth, and ephrin B1 stimulates dendrite extension as a surface ligand (Doherty et al., 1991; Moreno-Flores et al., 2002). Also, ephrin B reverse signaling interferes in vivo with the chemoattractive effect of stromal cell-derived factor-1 (SDF-1) (Lu et al., 2004).

The present inhibitor studies extend these previous findings, indicating direct involvement of both endogenous ephrin B and N-cadherin specifically in parallel fiber extension, dendritogenesis, and migratory behavior by CGNs both in vitro and in situ. The expression of ephrin B1 within the EGL/PMZ, ML, and IGL of the mouse cerebellum is consistent with these diverse functions. Similarly, N-cadherin has been associated with growth cones derived from extending axons of CGNs (Nakai and Kamiguchi, 2002) as well as with synaptic glomeruli within the granule cell layer of the mouse cerebellum (Fannon and Colman, 1996).

Local cellular interactions are critical for different stages of CGN development. For example, homotypic cell–cell contacts are important for proliferation of cerebellar progenitors (Gao et al., 1991) and for early differentiation and migration from the EGL (Gao and Hatten, 1993; Lu et al., 2004). Furthermore, newly forming axons within the PMZ form fascicles of parallel fibers that involve axonal contacts between neighboring CGNs (Altmann, 1972; Berglund et al., 1999). Cell surface proteins play a central role in these cell contact-dependent events by promoting intercellular signaling and cell adhesion (Berglund et al., 1999; Solecki et al., 2001; Lu et al., 2004).

NFI regulation of axon formation was observed under conditions of homotypic cell contact, consistent with a role for cell adhesion molecules in mediating this process. Inhibitor studies using cerebellar slices indicated that NFI influences parallel fiber orientation/fasciculation by promoting the expression of ephrin B1 and N-cadherin expression, at least in part. This likely involves trans interactions between N-cadherin monomers or ephrin B1/Eph proteins present on neighboring CGN axons. Inhibition of N-cadherin produced a more dramatic disorganization of parallel fiber orientation, suggesting a more primary role as an NFI mediator of axon guidance. Contactin-1 also was previously implicated in parallel fiber fasciculation (Berglund et al., 1999). Thus, multiple cell adhesion molecules appear to cooperate in controlling the orientation of parallel fibers. In addition to fasciculation, our results indicate that the ability of CGNs to extend parallel fibers per se also is dependent on homotypic axon guidance. Contactin-1 also was previously implicated in parallel fiber fasciculation (Berglund et al., 1999).

In contrast to the effects on axon outgrowth, NFI, ephrin B, and N-cadherin regulate dendrite formation in dissociated CGN
transduced CGNs within the IGL of cerebellar slices. Cultures were treated with either ephrinB1-Fc (these cell adhesion molecules. For example, c

orientation for GFP

Figure 9. Cell adhesion molecule inhibitors disrupt differentiation of CGNs in situ. A, B, CGN radial migration is inhibited by cell adhesion molecule antagonists. Cerebellar slices were transduced with GFP retroviruses and treated with 50 μg/ml ephrinB1-Fc, 400 μg/ml N-cadherin inhibitor peptides, or control reagents for 60 h. Distributions of viral-transduced GFP + cells within different cerebellar layers are shown. Both inhibitors significantly reduced CGN migration relative to controls. *p < 0.05; **p < 0.01. C–D, Top, Antagonists inhibit parallel fiber extension and induce axon misorientation. GFP + parallel fibers in the PMZ/ML region of coronal slices treated with 50 μg/ml ephrinB1-Fc or 400 μg/ml N-cadherin inhibitor peptides for 48 h. Labeled cells lying outside the indicated layers derive from an adjacent EGL (see legend to Fig. 3B). Bottom, The orientation of GFP + axons was significantly altered by the soluble inhibitors relative to controls. *p < 0.01; **p < 0.001. θ, Angle of divergence from the parallel orientation for GFP + axons, expressed in degrees. E, F, Antagonists inhibit extension of GFP-labeled dendrites from retroviral-transduced CGNs within the IGL of cerebellar slices. Cultures were treated with either ephrinB1-Fc (E) or N-cadherin antagonist peptides (F) or control reagents for 60 h.

cultures. This could reflect homotypic cellular interactions operating under these conditions (e.g., cell contacts involving extended axons) or, alternatively, cell autonomous functions of these cell adhesion molecules. For example, cis interactions have been observed between ephrin ligands and Eph receptors as well as between N-cadherin monomers (Takeda et al., 1999; Shan et al., 2000; Yin et al., 2004; Marquardt et al., 2005) (see also below).

In addition to process formation, NFI also regulates CGN migration through ephrin B1 and N-cadherin. Ephrin B reverse signaling was previously implicated in CGN egress from the EGL via its antagonism of SDF-1 (Lu et al., 2004), which is a CGN chemoattractant released by the meninges that prevents their premature migration from the EGL (Ma et al., 1999). In addition to this permissive function, our findings indicate that ephrin B2 can also directly promote CGN migration to some extent under conditions of homotypic cell contact, such as those present within the EGL. Inhibitor studies also indicated that N-cadherin has a prominent role in mediating NFI regulation of CGN migration. Facilitation of CGN migratory behavior by endogenous N-cadherin has not been reported previously. One distinction between these two mediators is the apparent requirement for trans signaling in the case of ephrin B.

Studies of NFI-regulated axon extension, migration, and dendritogenesis in cell culture were performed on a laminin surface and thus involved integrin β1-dependent mechanisms. Integrin β1 mRNA is expressed throughout CGN development within the postnatal mouse cerebellum (Blaess et al., 2004). Integrin β1 has been implicated in several aspects of CGN development, including progenitor proliferation (Blaess et al., 2004), axon extension (Mercado et al., 2004), and migration (Fishman and Hatten, 1993; Thelen et al., 2002; Lauro et al., 2006). Interestingly, both ephrins and N-cadherin can interact in cis with integrin β1 pathways during axon extension (Arregui et al., 2000; Nakamoto et al., 2004) as well as cell migration (Li et al., 2000; Huynh-Do et al., 2002). Thus, NFI may promote axon formation, migration, and dendritogenesis through signaling interactions between ephrin B1/N-cadherin and integrin β1. Significantly, cell-autonomous interactions between integrin β1 and N-cadherin were shown recently to regulate dendritic architecture in retinal ganglion cells (Marrs et al., 2006). Similar cell-autonomous, cis interactions may account for the stimulation of dendrite formation in dissociated cultures by NFI, N-cadherin, and ephrin B.

Gene knock-out mice indicate that both NFIA and NFIB are important for proper differentiation of CGNs. Furthermore, downregulation of N-cadherin and/or ephrin B1 in Nfia-null mice suggests that loss of both proteins contributes to these alterations in CGN differentiation. Nfia-deficient mice exhibited disrupted parallel fiber formation and orientation as well as CGN dendritogenesis, consistent with cell culture and in situ studies using the NFI dominant repressor. Radial migration also was incomplete by P17 in these mice, which may reflect inhibition of migratory behavior and/or earlier differentiation events mediated by ephrin B1/N-cadherin such as parallel fiber extension. Alterations in CGN proliferation, growth arrest, or onset of early differentiation attributable to Nfia deficiency also cannot be ruled out. However, this is not supported by the lack of effect of Nfia/EnR on BrdU incorporation in CGN cell cultures and by the unaltered expression of the early differentiation markers NeuroD and NeuN in Nfia/EnR-expressing CGNs and in Nfia-null mice. There was also no evidence of residual CGN progenitors in P17 Nfia-null mice. It is possible that alterations in other cell types also may contribute to the cerebellar phenotype in Nfia gene knock-out mice (Mathis et al., 2003). For example, immunostaining for both N-cadherin and ephrin B1 were broadly diminished in the cerebellum of Nfia-null mice. Finally, the disruption of axonal marker expression in the EGL of Nfib-null mice indicates that this family member also is critical for normal parallel fiber formation in the early forming cerebellum.

In addition to its role in CGN differentiation, loss of NFIA results in reduced formation of midline glia in the mouse fore-
brain (Shu et al., 2003). Furthermore, this factor inhibits neurogenesis via Hes5 (hair and enhancer of split 5) in ventricular zone progenitors within the embryonic chick spinal cord and promotes their gliogenesis and also directs subsequent differentiation of spinal astrocytic precursors (Deneen et al., 2006). Thus, NFIA and its family members appear to be versatile regulators in promoting the differentiation of distinct neural cell types depend-

References


