

RESEARCH COMMUNICATION

Neurofibromatosis-1 regulation of neural stem cell proliferation and multilineage differentiation operates through distinct RAS effector pathways

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Neurofibromatosis type 1 (NF1) is a common neurodevelopmental disorder caused by impaired function of the neurofibromin RAS regulator. Using a combination of *Nf1* genetically engineered mice and pharmacological/genetic inhibition approaches, we report that neurofibromin differentially controls neural stem cell (NSC) proliferation and multilineage differentiation through the selective use of the PI3K/AKT and RAF/MEK pathways. While PI3K/AKT governs neurofibromin-regulated NSC proliferation, multilineage differentiation is MEK-dependent. Moreover, whereas MEK-regulated multilineage differentiation requires Smad3-induced Jagged-1 expression and Notch activation, MEK/Smad3-regulated Hes1 induction is only responsible for astrocyte and neuronal differentiation. Collectively, these findings establish distinct roles for the RAS effector pathways in regulating brain NSC function.

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Neurofibromatosis type 1 (NF1) is a common neurogenetic disorder in which individuals manifest numerous CNS abnormalities that reflect impaired neuronal and glial cell lineage function. In this regard, 60%–80% of children with NF1 exhibit impairments in learning, attention, and memory (Diggs-Andrews and Gutmann 2013), and 15%–20% of affected children develop low-grade astrocytomas (gliomas) involving the optic pathway and brain stem (Guillamo et al. 2003). The fact that both neuronal and astroglial lineages are impacted raises the possibility that the *NF1* gene product neurofibromin is a critical regulator of neural stem cell (NSC) growth and differentiation. Consistent with this idea, previous reports have revealed that neurofibromin negatively controls NSC proliferation and self-renewal as well as multilineage differentiation (Hegedus et al. 2007; Lee et al. 2010) such that *Nf1* inactivation leads to increased numbers of spinal cord neuroglial progenitors (Bennett et al. 2003) and increased telencephalic NSC proliferation (Dasgupta and Gutmann 2005). Similarly, conditional *Nf1* gene inactivation in BLBP- or GFAP-expressing neuroglial progenitor cells results in increased

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NSC proliferation and glial lineage differentiation in vivo (Hegedus et al. 2007; Wang et al. 2012).

Neurofibromin is widely expressed in the developing brain, where it primarily functions as a negative regulator of RAS activity. Previous studies have demonstrated that loss of neurofibromin expression in NSCs results in increased proliferation and glial differentiation in a RAS- and AKT-dependent fashion (Hegedus et al. 2007; Lee et al. 2010). However, other studies have implicated RAS/ERK signaling as the responsible pathway dictating *Nf1*-deficient neural progenitor cell growth and differentiation in the forebrain (Wang et al. 2012) and cerebellum (Sanchez-Ortiz et al. 2014). Additionally, neurofibromin control of mouse astrocyte and optic glioma growth is mediated by both MEK and AKT effector arms through convergence on the mammalian target of rapamycin (mTOR) complex (Kaul et al. 2015). In contrast, neurofibromin regulation of human and mouse neuronal cyclic AMP homeostasis is RAS-dependent but operates in a MEK/AKT-independent manner through PKC ζ (Anastasaki and Gutmann 2014). Together, these observations argue that neurofibromin regulation of nervous system cell biology may be cell type- or cell function-specific.

To mechanistically define the signaling pathways responsible for brain NSC function, we leveraged *Nf1* genetically engineered mice and converging inhibition strategies to demonstrate that neurofibromin regulation of NSC proliferation and multilineage differentiation involves the engagement of distinct RAS downstream signaling pathways. Here, we establish that neurofibromin control of NSC proliferation is PI3K/AKT-dependent, while MEK/Smad3/Jagged1/Hes1-dependent signaling is required for neurofibromin-regulated NSC glial and neuronal differentiation in vitro and in vivo.

Results and Discussion

To determine which RAS downstream effectors were hyperactivated following *Nf1* inactivation, we focused on third ventricle zone (TVZ) NSCs (Lee et al. 2012). *Nf1*^{−/−} and wild-type TVZ NSC cultures were generated from postnatal day 1 (P1) *Nf1*^{fl/fl} pups following Cre or LacZ gene adenovirus infection, respectively. Following neurofibromin loss, increased ERK (3.5-fold; Thr202/Tyr204) and AKT (1.8-fold and threefold; Ser473 and Thr308) phosphorylation was observed (Fig. 1A).

To identify which RAS effector pathway was responsible for neurofibromin regulation of NSC growth and multilineage differentiation, we used PI3K/AKT and MEK pharmacological inhibitors. While MK2206 treatment inhibited AKT activation (Fig. 1B) and reduced *Nf1*^{−/−} NSC growth (Fig. 1C [direct cell counting], D [percentage of Ki67 $^{+}$ cells]) to wild-type levels, it had no effect on ERK phosphorylation (Fig. 1B). PD0325901 (PD901) treatment inhibited MEK activation but did not decrease *Nf1*^{−/−} NSC proliferation (Fig. 1C) or AKT phosphorylation (Fig.

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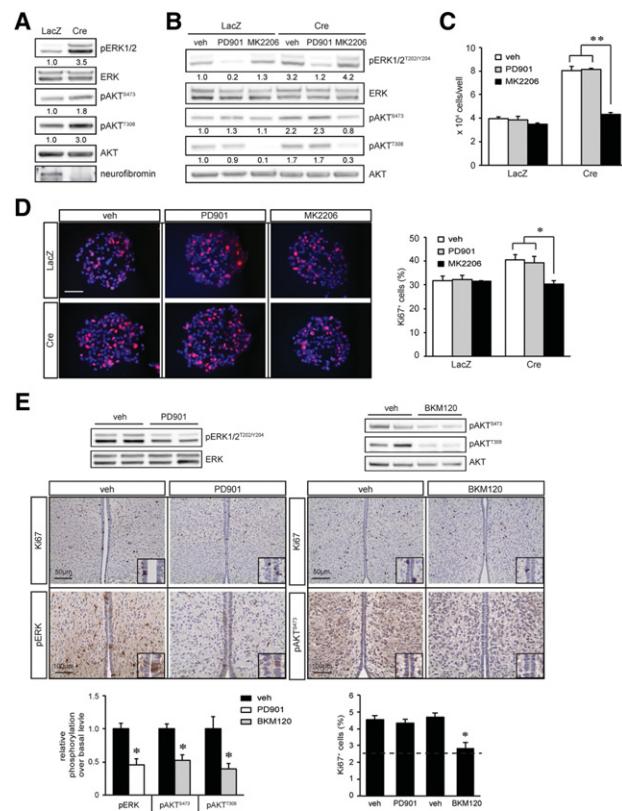


Figure 1. Neurofibromin regulates NSC proliferation in a PI3K/AKT-dependent manner. (A) *Nf1* loss in TVZ NSCs resulted in increased ERK and AKT activation (pERK^{T202/Y204}, pAKT^{S473}, and pAKT^{T308} phosphorylation) relative to wild-type controls. (B) Treatment with the MEK (5 nM PD0325901 [PD901]) or AKT inhibitor (50 nM MK2206) reduced ERK and AKT hyperactivation. Reduced NSC numbers (direct cell counting) (C) and percentage of Ki67⁺ cells (D) were observed only in *Nf1*^{-/-} TVZ NSCs treated with MK2206. (E) NVP-BKM120 (BKM120) but not 5 mg/kg PD901 treatment decreased the percentage of Ki67⁺ cells in the TVZ of P18 *Nf1*^{BLBP} conditional knockout mice in vivo ($n = 4$ per group) to nearly wild-type levels (dotted line). ERK and AKT hyperactivation were decreased following PD901 and BKM120 treatment, respectively, relative to wild-type controls. $n = 4$ per group. (veh) Vehicle. Nuclei were counterstained with DAPI. Error bars denote the mean \pm SD. Bar, 100 μ m. (*) $P < 0.05$; (**) $P < 0.01$.

1B). Similar results were obtained with additional PI3K (NVP-BKM120 [BKM120]) and MEK (UO126) inhibitors (Supplemental Fig. S1A,B). The observation that MEK inhibition had no effect on *Nf1*^{-/-} NSC growth but reduced *Nf1*^{-/-} astrocyte proliferation (Kaul et al. 2015) further underscores the importance of cell type specificity (Lee et al. 2012).

To determine whether differential RAS pathway control of NSC proliferation was also observed in vivo, we used *Nf1*^{BLBP} conditional knockout mice to inactivate *Nf1* gene expression in BLBP⁺ NSCs at embryonic day 9.5 (E9.5) (Hegedus et al. 2007). At P0.5, increased ERK phosphorylation (Thr202/Tyr204), AKT phosphorylation (Ser473 and Thr308), and proliferating cells (percentage of Ki67⁺ cells) were observed in *Nf1*^{BLBP} conditional knockout mice relative to wild-type controls (Supplemental Fig. S1C,D). Following the treatment of pregnant females with either 5 mg/kg PD901 or 30 mg/kg BKM120 from E15 to E18, the percentage of proliferating Ki67⁺ cells within the TVZ was quantified at P0.5. ERK hyperactiva-

tion in the TVZ of *Nf1*^{BLBP} conditional knockout pups was reduced by PD901 treatment; however, there was no change in TVZ cell proliferation (Fig. 1E). In contrast, BKM120 treatment inhibited AKT (Ser473 and Thr308 phosphorylation) hyperactivation and reduced TVZ cell proliferation (Fig. 1E). Collectively, these data demonstrate that neurofibromin regulates NSC proliferation in a PI3K/AKT-dependent manner in vitro and in vivo, consistent with previous findings (Lee et al. 2010).

AKT maintenance of NSC growth has been reported in mice and flies (Lee et al. 2010, 2013; Amiri et al. 2012); however, the role of AKT in regulating NSC multilineage differentiation is less clear (Peltier et al. 2007). Following in vitro differentiation, *Nf1* loss in NSCs resulted in an increase in the percentage of GFAP⁺ and O4⁺ cells (3.4-fold and fourfold, respectively) and a decrease in Tuj1⁺ cells (2.5-fold) relative to wild-type cells (Fig. 2A). While treatment with the AKT inhibitor (MK2206) had no effect on these *Nf1*-deficient NSC differentiation defects, MEK inhibition (PD901) restored *Nf1*^{-/-} NSC astrocyte, oligodendrocyte (Olig2⁺ and O4⁺ cells), and neuron differentiation to near wild-type levels (Fig. 2A; Supplemental Fig. S2A). Similar results were observed using other PI3K (BKM120) and MEK (UO126) inhibitors (Supplemental Fig. S2B).

To determine whether MEK activation was responsible for these multilineage defects in vivo, *Nf1*^{BLBP} conditional knockout pups were treated from P0.5 to P18 (astrocytes and oligodendrocytes) or from E15 to E18 (neurons and Olig2⁺ progenitors) with PD901 or BKM120. While *Nf1* inactivation in BLBP⁺ neural progenitor cells led to increased numbers of GFAP⁺ astrocytes and APC⁺ oligodendrocytes at P18 (Supplemental Fig. S2C), decreased numbers of NeuN⁺ neurons and increased numbers of Olig2⁺ cells were observed at P0.5 (Supplemental Fig. S2E). As observed

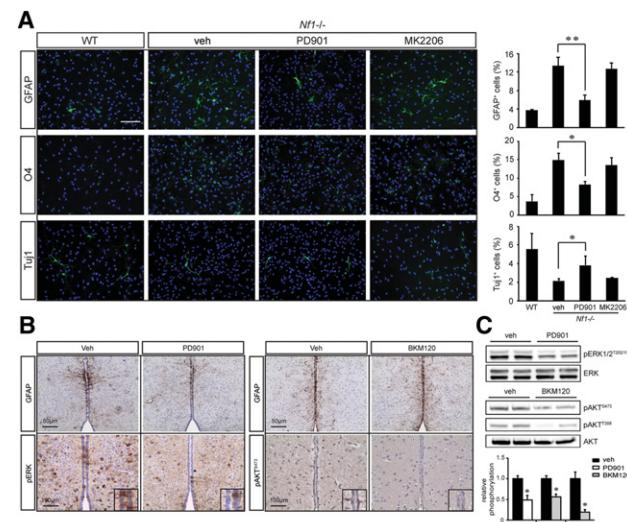


Figure 2. Neurofibromin regulation of NSC multilineage differentiation is MEK-dependent. (A) The astrocyte, oligodendrocyte, and neuronal differentiation defects observed in *Nf1*^{-/-} TVZ NSCs were restored to wild-type (WT) levels following PD901, but not MK2206, treatment. (B) PD901-treated *Nf1*^{BLBP} conditional knockout mice exhibited reduced astrocyte numbers compared with vehicle-treated controls. $n = 4$ per group. No change in astrocyte numbers was observed following BKM120 treatment. $n = 4$ per group. (C) Denitometric analyses of pERK1/2^{T202/Y204}, pAKT^{S473}, and pAKT^{T308} immunoblots relative to total ERK and AKT levels. $n = 4$ per group. Error bars denote the mean \pm SD. (veh) Vehicle. Bar, 100 μ m. (*) $P < 0.05$; (**) $P < 0.01$.

NF1 controls neuroglial growth and differentiation

in vitro, MEK, but not PI3K/AKT, inhibition restored astrocyte [Fig. 2B,C] and oligodendrocyte numbers in *Nf1*^{BLBP} conditional knockout mice to wild-type levels at P18 (Supplemental Fig. S2D) as well as ameliorated the increase in Olig2⁺ cells and decrease in NeuN⁺ cells at P0.5 (Supplemental Fig. S2E). Together with the in vitro results, these data reveal that AKT and MEK independently regulate NSC proliferation and multilineage differentiation, respectively.

The observation that MEK is a central driver of gliogenesis is consistent with prior reports demonstrating that *Mek1/2*-deficient mice exhibit impaired glial cell specification (Li et al. 2012) and that neonatal MEK inhibition rescues the developmental defects in *Nf1*-deficient brains by restoring normal neuron-glial specification (Wang et al. 2012). However, the mechanism responsible for neurofibromin/MEK-driven multilineage differentiation has not been elucidated. Two transcription factors, Erm and Ascl1, can regulate gliogenesis in response to elevated RAS/ERK signaling (Li et al. 2012; Breunig et al. 2015). While *Nf1*^{-/-} NSCs exhibit increased *Erm* expression by quantitative RT-PCR (qRT-PCR) and Western blotting, this was not attenuated following MEK inhibition (PD901) (Supplemental Fig. S3A,B). In addition, no change in Ascl1 protein levels was observed after *Nf1* loss, and nearly 100% of wild-type and *Nf1*^{-/-} NSCs were Ascl1⁺ (Supplemental Fig. S3B,C). Since Erm and Ascl1 function can also be regulated by phosphorylation (Li et al. 2014), these proteins could still play a role in *Nf1*^{-/-} NSC gliogenesis.

Based on increased Jagged-1 expression in *Nf1*-deficient mouse astrocytes (Banerjee et al. 2011) and numerous studies highlighting the critical role of Notch1 signaling in specifying neural cell fate during development (Lutolf et al. 2002; Stump et al. 2002), we examined Jagged1/Notch pathway activation. Following neurofibromin loss in NSCs, there was increased Jagged1 expression and Notch activation (Notch intracellular domain [NICD] expression) in vitro (Fig. 3A) and in vivo (Fig. 3B). However, in contrast to *Nf1*-deficient astrocytes (Banerjee et al. 2011), Jagged1 was not regulated by mTOR activation. While there was a 2.1-fold and 3.6-fold increase in S6 Ser240/244 and Ser235/236 phosphorylation, respectively, mTOR inhibition with rapamycin did not affect Jagged1 or NICD expression in *Nf1*^{-/-} NSCs (Supplemental Fig. S3D). Consistent with the hypothesis that neurofibromin control of Jagged1 in NSCs is AKT/mTOR-independent, MK2206 treatment of *Nf1*^{-/-} NSCs did not reduce Jagged1 or NICD expression (Fig. 3C) or attenuate ERK hyperphosphorylation (Supplemental Fig. S3E). Instead, MEK inhibition (PD901) restored Jagged1 and NICD expression in *Nf1*^{-/-} NSCs to wild-type levels (Fig. 3D) without any change in AKT phosphorylation (Supplemental Fig. S3F). Similar results were observed with additional MEK (UO126) and PI3K (BKM120) inhibitors (Supplemental Fig. S3G). Moreover, Jagged1 and NICD expression in the TVZ of *Nf1*^{BLBP} conditional knockout mice was reduced following PD901 treatment (Fig. 3E) but not by PI3K/AKT (BKM120) inhibition. Collectively, these results establish that neurofibromin regulation of Jagged1/Notch activation is mediated by MEK/ERK signaling in vitro and in vivo.

The importance of Jagged1 to gliogenesis is further supported by studies using conditional Jagged1 deletion in cerebellar neuroepithelial cells (Weller et al. 2006) as well as reports demonstrating that Jagged1-mediated Notch pathway activation promotes astrogliogenesis in vivo (Hu et al. 2013) and inhibits neurogenesis in vitro (Wilhemsson et al. 2012). The ability of activated Notch1 to dictate mul-

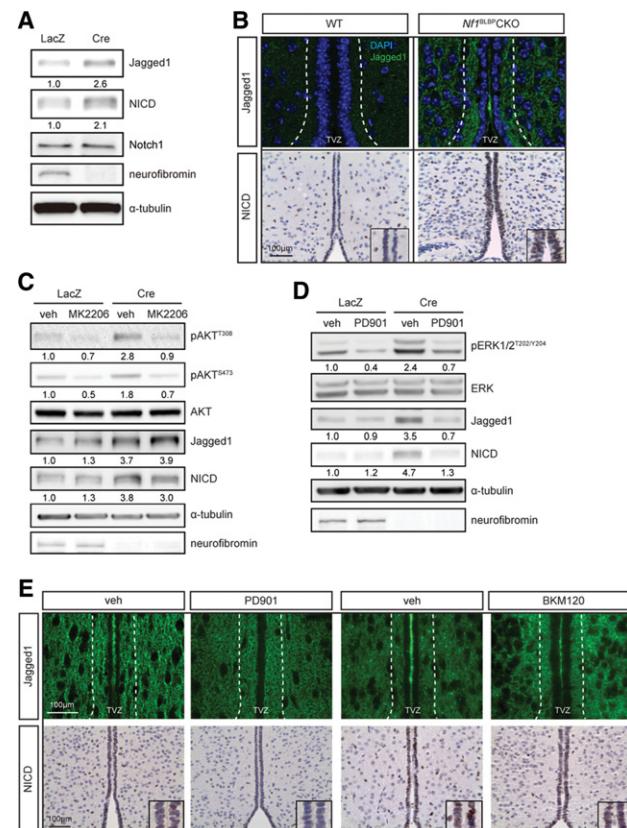


Figure 3. Neurofibromin loss in NSCs results in MEK-dependent Jagged1/Notch activation. Neurofibromin loss resulted in increased Jagged1 and cleaved Notch1 (NICD) expression in vitro (A) and in vivo (B) relative to controls. While MK2206 treatment did not reduce Jagged1 and NICD expression (C), PD901 treatment reduced *Nf1*-deficient NSC Jagged1 and NICD expression to wild-type (WT) levels (D). (E) Increased Jagged1 and NICD expression was observed in the TVZ of *Nf1*^{BLBP} conditional knockout (CKO) mice, which was ameliorated by PD901, but not BKM120, treatment. $n = 4$ per group. (veh) Vehicle.

tilineage differentiation in neural progenitor cells typically involves the Hes1 and Hes5 transcription factors (Furukawa et al. 2000; Hojo et al. 2000). Following *Nf1* loss in NSCs, there was increased Hes1 and Hes5 expression (Fig. 4A), which was reduced by PD901 treatment (Fig. 4B) in vitro. Moreover, ectopic expression of an activated MEK (caMEK), but not an activated AKT (myrAKT), molecule in NSCs increased Jagged1, NICD, Hes1, and Hes5 levels (Supplemental Fig. S4A). Finally, MEK inhibition restored Hes1 and Hes5 expression to wild-type levels in the TVZ of *Nf1*^{BLBP} conditional knockout mice in vivo (Fig. 4C). Together, these findings demonstrate that the Notch1 signaling pathway is activated following neurofibromin loss in a MEK-dependent manner.

Based on conflicting reports regarding Hes5 regulation of gliogenesis (Hojo et al. 2000; Wu et al. 2003) and the nearly exclusive expression of Hes1 within the TVZ, we chose to focus on Hes1. Using two independently generated *Hes1* shRNA constructs to decrease Hes1 expression in NSCs (60% reduction), the increased astrocyte differentiation observed following neurofibromin loss was reduced to near wild-type levels (Fig. 4D). Importantly, Hes1 reduction had no effect on *Nf1*^{-/-} NSC growth (direct cell counting) (Fig. 4E) or proliferation (percentage of Ki67⁺ cells) (Fig. 4F). Moreover, Hes1 knockdown ameliorated

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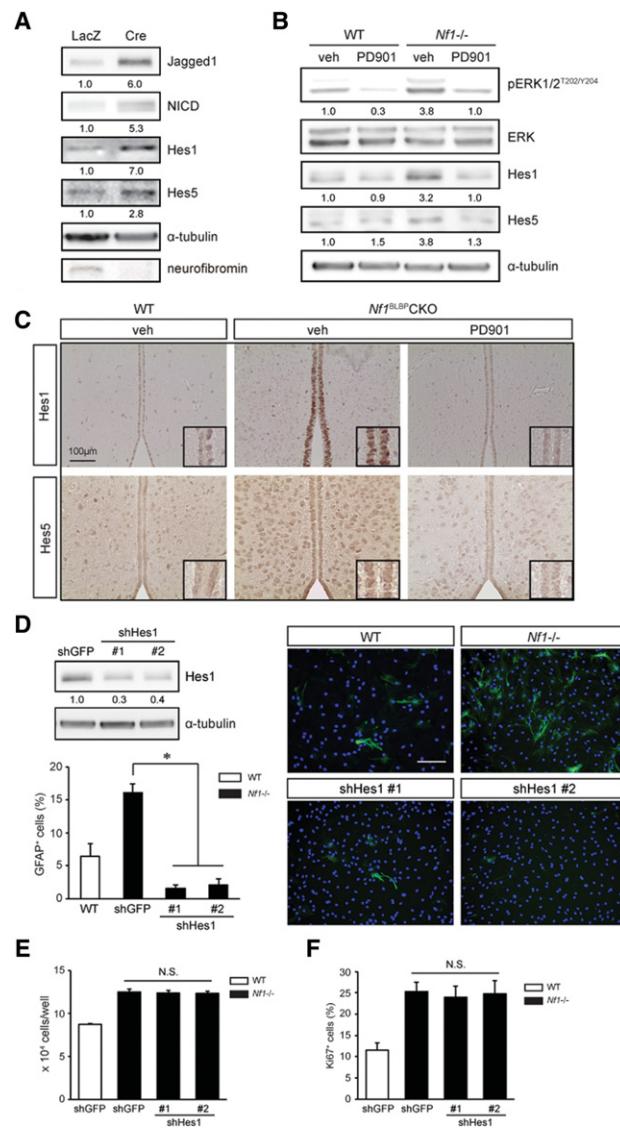


Figure 4. Neurofibromin regulation of NSC astrocyte differentiation is Hes1-dependent. *Nf1* loss results in increased expression of the Hes1 and Hes5 Notch downstream effectors [A], which was restored to wild-type (WT) levels following 5 nM PD901 treatment [B]. [C] *Nf1*^{BLBP} conditional knockout (CKO) mice treated with 5 mg/kg PD901 (P0.5–P18) have reduced Hes1 and Hes5 expression relative to vehicle-treated mice. $n = 4$ per group. [D] Hes1 shRNA knockdown reduced the percentage GFAP⁺ astrocytes following *Nf1*^{-/-} TVZ NSC differentiation. Hes1 knockdown (shHes1) did not reduce *Nf1*^{-/-} TVZ NSC growth (direct cell counting) [E] or proliferation (percentage of Ki67⁺ cells) [F]. (veh) Vehicle. Error bars denote mean \pm SD. Nuclei were counterstained with DAPI. Bar, 100 μ m. (*) $P < 0.01$; (N.S.) not significant.

the decrease in neuronal differentiation in *Nf1*-deficient NSCs but surprisingly had no effect on oligodendrocyte differentiation (percentage of O4⁺ cells) (Supplemental Fig. S4B). These findings demonstrate that neurofibromin regulation of astrocyte and neuronal differentiation is mediated by Hes1 in a reciprocally coordinated fashion, whereas other mechanisms underlie MEK-dependent oligodendrocyte differentiation. In this regard, Hes1 knockdown did not reduce the number of Olig2⁺ cells (Supplemental Fig. S4C). While RAF/MEK signaling is an important determinant of oligodendrocyte differentia-

tion in mice (Galabova-Kovacs et al. 2008) and zebrafish (Shin et al. 2012), other neurofibromin-regulated MEK downstream pathways are likely responsible for governing oligodendrocyte differentiation.

To determine how neurofibromin controls Jagged1 expression, we examined *Jagged1* mRNA expression using real-time qRT-PCR (Fig. 5A). *Jagged1* mRNA expression was regulated in *Nf1*^{-/-} NSCs on the transcriptional level through MEK, since PD901 treatment restored *Jagged1* mRNA to wild-type levels. Several potential regulators of Jagged1 expression have been identified, including β -catenin, YAP, and TGF β /Smad3 (Chen et al. 2010; Zhang et al. 2010; Tschaharganeh et al. 2013). While we observed no changes in β -catenin activation or YAP expression following neurofibromin loss, the increased Smad3 expression observed in *Nf1*-deficient NSCs was reduced to wild-type levels following PD901 treatment (Fig. 5B,C).

We next used genetic and pharmacologic approaches to reduce Smad3 function. Following Smad3 knockdown (using two different shRNA constructs), Jagged1, NICD, and

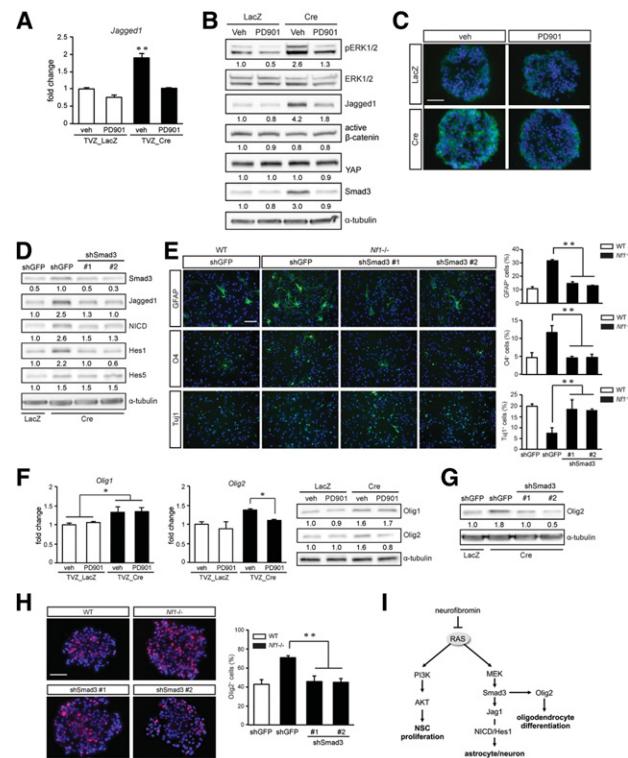


Figure 5. Neurofibromin/Jagged1 regulation of astrocyte differentiation requires MEK-mediated Smad3 expression. Increased *Jagged1* transcription [A] and protein levels [B] in *Nf1*^{-/-} NSCs were reduced to wild-type levels following 5 nM PD901 treatment. PD901 treatment of *Nf1*^{-/-} NSCs had no effect on β -catenin activity or YAP expression but reduced Smad3 expression by Western blotting [B] and immunocytochemistry [C]. [D] Smad3 knockdown restored Jagged1, NICD, and Hes1, but not Hes5, expression to wild-type levels. [E] The astrocyte, oligodendrocyte, and neuronal differentiation defects observed in *Nf1*^{-/-} TVZ NSCs were restored to near wild-type levels following Smad3 knockdown. [F] MEK inhibition (5 nM PD901) reduced Olig2, but not Olig1, expression by qRT-PCR and Western blotting. Smad3 knockdown reduced Olig2 expression by Western blotting [G] as well as the percentage of Olig2⁺ cells within the *Nf1*^{-/-} neurospheres [H]. [I] Proposed model of neurofibromin/RAS regulation of NSC growth and multilineage differentiation. (veh) Vehicle. Error bars denote mean \pm SD. Bar, 100 μ m. (*) $P < 0.05$; (***) $P < 0.01$.

Hes1 expression in *Nf1*^{-/-} NSCs was restored to wild-type levels (Fig. 5D). Similar to PD901 treatment, Smad3 knockdown of *Nf1*^{-/-} NSCs restored astrocyte, oligodendrocyte, and neuronal differentiation to wild-type levels (Fig. 5E). It should be noted that Smad3 knockdown did not change Hes5 expression (Fig. 5D), arguing against Hes5 as a mediator of neurofibromin-controlled multilineage differentiation. Similar results were observed in *Nf1*-deficient NSCs treated with the SIS3 Smad3 pharmacological inhibitor (Supplemental Fig. S5A,B; Jinnin et al. 2006). Collectively, these results support a mechanism by which neurofibromin/MEK control of astrocyte and neuron differentiation operates in a Smad3/Jagged1/Hes1-dependent manner.

Neurofibromin regulation of Smad3 function could operate at the level of transcription, protein degradation, or phosphorylation (Massague et al. 2005). While neurofibromin loss results in increased Smad3 levels, subcellular fractionation revealed an enrichment of Smad3 in the nucleus of *Nf1*^{-/-} NSCs (34%) relative to wild-type NSCs (8%) (Supplemental Fig. S5C). However, the mechanism underlying this increase in Smad3 expression was not the result of increased Smad3 RNA levels (qRT-PCR) (Supplemental Fig. S5D) or degradation mediated by increased SCF/ROC1 and GSK3- β binding (Fukuchi et al. 2001; Guo et al. 2008). Whereas Smad3 physically interacted with GSK3- β , but not with SCF/ROC1 (Supplemental Fig. S5E), neurofibromin loss or MEK inhibition (PD901 treatment) had no effect on Smad3 and GSK3- β binding, as assessed by immunoprecipitation. Finally, MEK-dependent Smad3 regulation was not dependent on TGF β -induced phosphorylation, as Smad3-Ser423/425 phosphorylation was similar in wild-type and *Nf1*^{-/-} NSCs (Supplemental Fig. S5F) and did not involve phosphorylation at the best-characterized ERK phosphorylation site (Ser208) (Supplemental Fig. S5F). Future studies will be required to identify the mechanism responsible for neurofibromin regulation of Smad3 levels.

Taken together, our findings establish that neurofibromin control of NSC function involves the selective use of distinct RAS effector pathways. In this regard, RAS activation is critical for both neurofibromin-regulated NSC proliferation and multilineage differentiation such that inhibition using the nonselective RAS inhibitor (lovastatin) (Li et al. 2005) restored both *Nf1*-deficient NSC growth and multilineage differentiation to wild-type levels (Supplemental Fig. S5G-I). However, whereas neurofibromin control of neuron and astrocyte differentiation requires Smad3 regulation of Hes1, the mechanism underlying Smad3-mediated oligodendrocyte differentiation likely involves other transcription factors. In this manner, Olig-1 and Olig-2 have been identified as essential factors for specifying oligodendrogenesis (Lu et al. 2000; Takebayashi et al. 2002). Consistent with recent findings demonstrating that Olig1 is not essential for oligodendrocyte development in mice (Paes de Faria et al. 2014), MEK inhibition (PD901) reduced the increased Olig2, but not Olig1, mRNA and protein expression in *Nf1*-deficient NSCs (Fig. 5F). Moreover, the elevated Olig2 expression in *Nf1*-deficient NSCs was decreased following Smad3 genetic (Fig. 5G,H) or pharmacologic (Supplemental Fig. S5J) inhibition. These findings suggest a model in which neurofibromin control of NSC multilineage differentiation involves distinct transcriptional programs: Neuron and astrocyte differentiation requires MEK/Smad3-dependent Hes1 induction, whereas MEK/Smad3-dependent oligodendrocyte differentiation involves Olig2 function (Fig. 5I).

Coupled with observations that other RAS downstream pathways have cell type-specific functions [neurons vs. astrocytes] relevant to brain cell function (Hegedus et al. 2007; Anastasaki and Gutmann 2014), the observations reported here establish that differential use of distinct RAS effector signaling pathways can govern separable cellular functions even within the same cell type, further underscoring the importance of cellular context in interpreting the impact of genetic mutations on brain function.

Materials and methods

Mice

BLBP-Cre; *Nf1*^{flx/wt} transgenic mice were crossed with *Nf1*^{flx/flx} mice to generate BLBP-Cre; *Nf1*^{flx/flx} (conditional knockout) mice. *Nf1*^{flx/flx} mice were used as wild-type controls. All strains were maintained on a C57BL/6 background and used in accordance with an approved animal studies protocol at Washington University.

Primary NSC analysis

TVZ NSCs were established from the TVZ of P1 *Nf1*^{flx/flx} mouse pups and analyzed as previously described (Lee et al. 2010). Retroviral and lentivirus (Supplemental Table 1) transduction was performed for overexpression and knockdown studies, respectively. All experiments were performed at least three times using primary NSCs generated from independent litters.

Western blotting

Western blotting was performed as previously reported (Lee et al. 2010) using primary antibodies (Supplemental Table 2) and was quantified by densitometry using a chemiluminescence imaging system (UVP). Each experiment was performed at least three times, and representative blots are presented.

Immunostaining

Paraffin or frozen sections were processed (Dasgupta and Gutmann 2005) prior to staining with the appropriate antibodies. The percentage of Ki67 $^{+}$ cells lining the TVZ was quantified as previously reported (Lee et al. 2012).

Pharmacologic inhibition studies

Neurospheres were trypsinized into single cells, and 5×10^5 cells per well were plated onto ultralow-binding 60-mm plates. Cells were treated with specific inhibitors for 4–5 d. PD901 (5 mg/kg/day; Selleck), BKM120 (30 mg/kg; Selleck), or matched vehicle (0.5% hydroxypropyl methylcellulose with 0.2% Tween 80 [Sigma-Aldrich] or 10/90 [v/v] N-methyl-2-pyrrolidone [NMO]/PEG300, respectively) was injected intraperitoneally into pregnant females from E15 to E18. Postnatal PD901 or NVP-BKM120 administration to lactating females (P0.5–P18) was achieved by oral gavage, and the mice were perfused at P18.

Real-time qRT-PCR

Real-time qRT-PCR was performed as previously described (Yeh et al. 2009) with specific primers (Supplemental Table 3), and $\Delta\Delta CT$ values were calculated using *H3f3a* as an internal control.

Statistical analysis

Each experiment was performed with samples from at least three independent groups. Statistical significance was set at $P < 0.05$ using the Student's *t*-test.

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