The differentiation of NSCs is regulated by various intrinsic and extrinsic factors. This neuronal replacement therapy is based on the idea that neurological functions lost might be recovered by introducing new cells that can differentiate and integrate appropriately to replace the functions of the lost neurons. However, the underlying molecular mechanisms for their fate specification remain ill defined. Accumulating evidence has confirmed that the differentiation of NSCs is regulated by various intrinsic and extrinsic factors.

The mammalian target of rapamycin (mTOR) is a serine/threonine kinase of the phosphoinositide 3-kinase (PI3K)-related kinases and the catalytic subunit of two distinct complexes called mammalian target of rapamycin complex 1 (mTORC1) and mTORC2. Activation of AKT can be initiated by phosphatidylinositol 3-OH kinase (PI3K), which in turn activates mTORC1 and further downstream targets such as p70S6 kinase. This leads to stimulation of gene transcription, protein synthesis (Parker et al., 2015). mTOR kinase is also a key regulator of the homeostasis of several stem cell pools in which it finely regulates the balance between self-renewal and differentiation in stem cells. Several studies have shown that mTOR activation causes enhanced generation of NSCs/neural progenitor cells, followed by neuronal differentiation (Endo et al., 2009; Marfia et al., 2011), and the inhibition of mTOR abrogates the increase of differentiated neurons (Han et al., 2008). In the CNS the mTOR pathway is also involved in axon regeneration, neuronal activity, and dendritic arborization (Li et al., 2016). Even though the important roles of mTOR in regulating differentiation and development are known, the profound target of mTOR undertaking this process is remained to be addressed.

The transcriptional regulation of cell-type-specific gene expression involves complicated epigenetic events such as DNA demethylation, histone modifications, and chromatin remodeling. DNA methylation/hydroxymethylation influence different stages of NSC differentiation and have been implicated in neural development and differentiation (Kim et al., 2014). The deletion of DNMT1/3b in neuronal progenitor cells results in DNA hypomethylation and triggers astrocyte and early neuronal differentiations, respectively (Fan et al., 2005; Martins-Taylor et al., 2012). In DNMT3a-null mice, impaired post-natal neurogenesis is seen in both the subventricular zone and subgranular zone (Wu et al., 2010). Moreover, DNA methylation can also be catalyzed by TET enzymes such as DNA demethylation, histone modifications, and chromatin remodeling. DNA methylation/hydroxymethylation influence different stages of NSC differentiation and have been implicated in neural development and differentiation (Kim et al., 2014). The deletion of DNMT1/3b in neuronal progenitor cells results in DNA hypomethylation and triggers astrocyte and early neuronal differentiations, respectively (Fan et al., 2005; Martins-Taylor et al., 2012). In DNMT3a-null mice, impaired post-natal neurogenesis is seen in both the subventricular zone and subgranular zone (Wu et al., 2010). Moreover, DNA methylation can also be catalyzed by TET enzymes such as DNA demethylation, histone modifications, and chromatin remodeling. DNA methylation/hydroxymethylation influence different stages of NSC differentiation and have been implicated in neural development and differentiation (Kim et al., 2014). The deletion of DNMT1/3b in neuronal progenitor cells results in DNA hypomethylation and triggers astrocyte and early neuronal differentiations, respectively (Fan et al., 2005; Martins-Taylor et al., 2012). In DNMT3a-null mice, impaired post-natal neurogenesis is seen in both the subventricular zone and subgranular zone (Wu et al., 2010). Moreover, DNA methylation can also be catalyzed by TET enzymes such
as TET1, TET2, and TET3 that revert the methylation status of DNA by successive oxidation of 5-methylcytosine (5-mC) into 5-hydroxymethylcytosine (5-hmC), 5-carboxycytosine, and 5-formylcytosine, which are intermediates of an active DNA demethylation mechanism (Hu et al., 2015). Loss of TET1 results in a 45% decrease of NSCs in the subgranular zone, and the neurosphere derived from TET1−/− mice shows impaired growth function (Zhang et al., 2013). However, overexpression of TET2 and TET3 increases 5-hmC formation and leads to defects in neuronal differentiation (Hahn et al., 2013). Furthermore, DNA methylation/demethylation is found to be coupled with histone modifications during neural development. The upregulation of H3K9 acetylation enhances neural differentiation and activates multiple neurodevelopmental genes (Qiao et al., 2015). However, decreased H3K9 trimethylation severely impairs early neurogenesis and enhances astrocyte formation (Tan et al., 2012). Pro-neural and terminal neuronal genes are poised in stem/progenitor cells by the balance between a repressor histone modification (H3K27me3) and an activator modification (H3K4me3) (McGann et al., 2014). Inhibition of histone deacetylase (HDAC) induces neuronal differentiation of adult NSCs, which is likely mediated through upregulation of neuronal-specific genes, such as neurogenic basic-helix-loop-helix transcription factors NeuroD, Ngn1, and Math1 (Hsieh et al., 2004; Yu et al., 2009).

Valproic acid (VPA) is a short-chain carboxylic acid commonly used for the treatment of epilepsy and bipolar disorders. Its pharmacological effects involve distinct mechanisms that affect the transmission of nerve signals in vitro and in vivo. Several intracellular pathways, such as the mTOR pathway, are affected by VPA. VPA activates mTOR signaling in the prefrontal cortex and hippocampus of autistic model rats, characterized by enhanced phospho-mTOR and phospho-S6 (Qin et al., 2009; Teng et al., 2014). VPA activates the PI3K/AKT/mTOR pathway in muscle, neuron, and induced pluripotent stem cells (Gurpur et al., 2009; Teng et al., 2014). VPA could also suppress the AKT/mTOR pathway in prostate cancer cells and postmortem fusiform gyrus (Nicolini et al., 2015; Xia et al., 2016). Some recent studies have shown that VPA reduces HDAC activity and promotes neuronal differentiation of NSCs (Hsieh et al., 2004). However, the underlying mechanisms are not fully understood.

Although both mTOR signaling and epigenetic regulation are essential for the differentiation of NSCs in developing or adult brains, a direct connection between mTOR signaling and epigenetic modifications remains uncharacterized in NSCs. In this study, we have set out to determine whether such a link exists, and, if so, how it might impact neural differentiation of VPA-induced NSCs.

**RESULTS**

**Activation of mTOR Signaling Is Required for VPA-Induced Neuronal Differentiation of NSCs**

To evaluate the function of mTOR signaling in VPA-induced neural differentiation, we examined the effects of mTOR inhibition and overexpression on neural differentiation, following VPA exposure. We found that VPA treatment caused neuronal differentiation of NSCs (Figure 1A). Interestingly, pretreatment of NSCs with the mTOR-specific inhibitor, rapamycin, remarkably attenuated NSCs from VPA-induced neuronal differentiation (Figure 1A). To further confirm the prodifferentiation role of mTOR, we expressed constitutively active (CA)-mTOR activator Rheb in NSCs and found that mTOR overexpression sensitized NSCs to VPA-induced neuronal differentiation (Figures 1B and 1C). These results suggest that mTOR signaling is critical for VPA-induced neuronal differentiation.

It has been well established that HDAC inhibitors decrease the proliferation of NSCs (Dozawa et al., 2014). We further determined whether the mTOR pathway mediated the effect of VPA on the proliferation of NSCs. We found that rapamycin increased the percentage of bromodeoxyuridine (BrdU)-positive cells and obviously prevented the decrease of VPA-induced proliferation of NSCs. A total of 17% of the cells in VPA-treated cultures were BrdU positive, compared with 47% of the cells in control cultures (Figure 1D). Unexpectedly, rapamycin dramatically enhanced NSCs proliferation (from 17% in VPA-treated cultures to 38% in rapamycin-treated cultures). There was no significant difference in NSC viability among three groups.

**VPA Activates the AKT/mTOR/p70S6K Pathway**

Previous studies have demonstrated that VPA activates the PI3K/AKT/mTOR pathway in muscle (Gurpur et al., 2009). However, the signaling pathway by which VPA induces neuronal differentiation of NSCs is not clear. To determine the mechanism by which VPA promotes neuronal differentiation, the AKT/mTOR/p70S6K pathway was tested. NSCs treated with VPA for 48 or 72 hr had higher levels of activated AKT, mTOR, and p70S6K (Figure 2A). Immunofluorescence observation revealed that VPA activated p70S6K, and that rapamycin attenuated the expression promoted by VPA (Figure 2B), confirming that VPA activates AKT/mTOR/p70S6K in NSCs.

To determine whether this AKT activation and its resulting downstream effects are triggered by VPA at a shorter time after exposure, 1-day-old differentiating cultures were treated with various concentrations of VPA for 1 hr, extracted, and immunoblotted for total and phosphorylated AKT. Treatment of NSCs with VPA activated AKT within 1 hr. Activation of AKT exhibited a dose-dependent response to VPA (Figure 2C).
AKT Activation by VPA Is Dependent on PI3K

Since VPA has HDAC inhibitor activity, kinases expressed or activated as a result of gene modulation may activate AKT independently of PI3K. Alternatively, VPA may activate AKT through the classical PI3K pathway. To differentiate between these possibilities, 1 mmol/L VPA was added on day 2 of differentiation either with or without 100 nmol/L wortmannin, a potent inhibitor of PI3K. Fresh VPA and wortmannin were added every 24 hr thereafter. Microscopic observations revealed that wortmannin attenuated neuronal differentiation of NSCs promoted by VPA (Figure 2D). Immunofluorescence and western blots of extracts made 48 hr after drug exposure demonstrated that wortmannin attenuated the activation of AKT by VPA, confirming that VPA activates AKT through PI3K (Figures 2E and 2F). To exclude the non-specificity of wortmannin on PI3K, the data from knockdown PI3K p110α or p110β were consistent with our above results (Figure S1).
Figure 2. VPA Activates the AKT/mTOR/p70S6K Pathway through a PI3K-Dependent but Not a Histone Acetylation-Dependent Manner in NSCs

(A) VPA (1 mmol/L) was added to duplicate cultures of NSCs for 48 or 72 hr, followed by western blotting using antibodies against the signaling proteins. Higher levels of activated AKT, mTOR, and p70S6K were detected in the VPA-treated NSCs. Representative western blot of three independent experiments.

(B) Immunofluorescence staining for p-p70S6K on NSCs. Higher expression was observed at 1 mmol/L VPA. Scale bar, 50 μm. Representative graph of three independent experiments.

(C) NSCs were exposed to various concentrations of VPA for 1 hr, followed by extraction and western blotting for AKT. VPA activates AKT in a dose-dependent manner, with highest activation seen at 1 mmol/L VPA. Representative western blot of three independent experiments.

(D) NSCs were treated for 48 hr with 1 mmol/L VPA with or without 100 nmol/L wortmannin. Immunofluorescence staining for NeuN showed that the prodifferentiation effect of VPA on NSCs was attenuated by wortmannin. Scale bar, 250 μm, indicating that VPA depends on PI3K for this effect. **p = 0.001 < 0.01 compared with control; ##p = 0.006 < 0.01 compared with VPA. Data are from three independent experiments.

(E) NSCs treated as above were extracted and immunoblotted for phosphor AKT (p-AKT) and total AKT. Ratios of p-AKT to total AKT band intensities were plotted. The AKT-activating effect of VPA was attenuated by wortmannin. *p = 0.02 < 0.05 compared with VPA. Data are from three independent experiments.

(F) Immunofluorescence staining for p-AKT on NSCs. Wortmannin decreased the expression of p-AKT induced by VPA. Scale bar, 250 μm. Representative graph of three independent experiments.

(G) VPA activated AKT/p70S6K signaling and enhanced histone H3 and H4 acetylation. NSCs were stimulated with 1 mM of VPA for various times and the levels of p-AKT, phospho-p70S6K, acetyl-H3, and acetyl-H4 were analyzed by immunoblotting. Both AKT and S6K proteins activation preceded the onset of histone H3 and H4 acetylation. *p = 0.03, 0.01 < 0.05 compared with 12 and 24 hr; †p = 0.007, 0.005, 0.007, 0.01, 0.03 < 0.05 compared with 1, 12, 24, 48, and 72 hr; ‡p = 0.008, 0.02, 0.01, 0.01 < 0.05 compared with 48 and 72 hr. Data are from three independent experiments.

(H) The effects of VPA and TSA on acetyl-H3 accumulation appeared to be additive. TSA alone appeared to elevate levels of phospho-p70S6K, but the increases were not concentration dependent. A higher concentration of TSA (300 nm) attenuated VPA-induced increase in phospho-p70S6K. *p = 0.03, 0.006, 0.02, 0.008 < 0.05 compared with control. Data are from three independent experiments.

(I) NSCs were treated by VPA or VPA plus rapamycin. Lysates from the cells were analyzed by immunoblotting for the levels of HDAC1, HDAC3, and β-actin. Rapamycin (Rapa) increased HDAC1 and HDAC3 levels but did not significantly alter the magnitude of VPA-induced decreases in HDAC1 and HDAC3 levels. **p = 0.006 < 0.01 compared with control; #p = 0.008 < 0.01 compared with control. One-way ANOVA and Student’s t test were used to determine the statistical significance. Data are from three independent experiments.
AKT Activation by VPA Is Independent on Histone Acetylation

VPA rapidly and concentration-dependently induces accumulation of acetylated H3 and H4 (Godino et al., 2015). To exclude the possibility that AKT activation may be triggered by histone modification, we tested the effects of VPA on phosphorylation of AKT, S6K, and acetylation of histone H3 and H4. Our results showed that VPA activated S6K from 12 hr and reached a plateau at 24 hr. VPA activated AKT from 1 hr and reached a plateau at 12 hr, which preceded the increase of S6K phosphorylation. In addition, both AKT and S6K protein activation also preceded the onset of histone H3 and H4 acetylation (Figure 2G). Moreover, trichostatin A (TSA), a potent HDAC inhibitor, concentration-dependent induced accumulation of acetylated H3. TSA at a low concentration induced a significant increase in phospho-S6K levels, yet this increase plateaued at a higher TSA concentration (Figure 2H). These data indicate that TSA produces a limited effect on the mTOR pathway that does not parallel its effect on acetyl-H3 accumulation (Figure 2H). TSA at a low concentration induced additive effects on VPA-induced acetyl-H3 accumulation, but not on VPA-induced decreases in phospho-S6K levels. TSA at a high concentration significantly attenuated VPA-induced increases in phospho-S6K levels. These results suggest that VPA-induced AKT activation is independent on its effect on histone modification.

Although both AKT and S6K protein activation preceded the onset of histone acetylation, to exclude the possibility that AKT/mTOR/p70S6K signaling activation in turn promotes histone acetylation, we investigated the effects of VPA with or without rapamycin on expression of HDACs. Unexpectedly, we observed that rapamycin increased HDAC1 and HDAC3 levels, but did not significantly alter the magnitude of VPA-induced decreases in HDAC1 and HDAC3 levels (Figure 2I). This finding indicates that AKT/mTOR/p70S6K signaling may not direct the histone acetylation.

PI3K/AKT/mTOR Activity Regulated Passive DNA Demethylation during Differentiation Induced by VPA

DNA methylation/demethylation are essential epigenetic modifications controlling gene expression during differentiation and development (Orlanski et al., 2016). Since the inhibitors of histone deacetylation cause epigenetic DNA modifications, we next further evaluated whether PI3K/AKT/mTOR activity regulates VPA-induced neuronal differentiation by repressing DNA demethylation. We found that the mRNA expression profiles of passive DNA demethylation pathway enzymes DNMT1, DNMT3a, but not DNMT3b and DNMT3L, were significantly decreased in NSCs treated with 1 mM VPA for 1 day (Figure 3A). Similar results were obtained from the protein blot experiments (Figure 3B). Surprisingly, PI3K/AKT/mTOR inhibition markedly attenuated the decreases in the passive DNA demethylation pathway enzymes (Figures 3A and 3B). Interestingly, we also found that the expression of the neuron-specific gene Ngn1 was much greater in VPA-treated cells than in the DMSO-treated controls, and that its expression was also affected by PI3K/AKT/mTOR inhibition (Figures 3A and 3B). Our results suggested that PI3K/AKT/mTOR activity may stimulate the activation of Ngn1 through suppressing the expression of DNMT 1 and 3a, which triggered VPA-induced neural differentiation of NSCs.

To clarify whether the increased expression of Ngn1 induced by VPA is a direct result of inhibiting the putative DNMTs, we studied the effects of VPA on the genome-wide DNA demethylation and gene-specific promoter demethylation. In the control group, 5-mC immunostaining (im) was highly intense and distributed throughout the nucleus (Figure 3C). However, in the VPA-treated group, decreased 5-mC-im intensity was seen in the nucleus, but preferentially distributed near the nuclear membrane into differentiation (Figure 3C). We next examined the effect of PI3K/AKT/mTOR inhibition on genome-wide DNA methylation. As expected, rapamycin obviously attenuated the decreased 5-mC-im intensity induced by VPA (Figure 3C), suggesting an important role for mTOR signaling in VPA-induced DNA demethylation, which resulted in Ngn1 expression. This notion was further confirmed by dot blotting for 5-mC (Figure 3D). As Ngn1 expression is known to be repressed by methylation of CpG dinucleotides within its promoter, We next further examined the effect of VPA on gene-specific promoter demethylation; bisulfite sequence analysis revealed that NSCs treated with VPA showed nearly 40% loss of 5-mC in the Ngn1 promoter region between −171 and 140 than that of control NSCs (Figure 3E). Furthermore, rapamycin treatment showed 20% gain of 5-mC in the same region (Figure 3E). DNA methylation levels of the Ngn1 promoter region in rapamycin-treated cells with or without VPA were further verified by EpiMark 5-mC Analysis Kit (Figure 3F). Taken together, our findings indicate that mTOR inhibition mostly blocks the DNA demethylation of neuronal target gene caused by VPA during differentiation.

PI3K/AKT/mTOR Activity Facilitates TET-Mediated DNA Hydroxymethylation during VPA-Induced Differentiation

DNA demethylation involves passive and active demethylation. By converting 5-mC to 5-hmC following the base excision repair pathway, TET family proteins were shown to promote active DNA demethylation in mammalian cells (Kafer et al., 2016). We next focused our investigation on the genes mediating active demethylation. We found upregulation of
TET1, TET2, and TET3 protein levels in VPA-treated cells, this effect can be inhibited by rapamycin treatment (Figure 4A). However, the small difference in TET1, TET2, or TET3 protein expressions between DMSO- and rapamycin-treated cells showed that the effect of VPA on TETs is dependent on the mTOR signaling in NSCs (Figure 4A). To further confirm the expression of the TET family members in every group, we immunohistochemically evaluated the intensity of the TET family. As expected, the immunohistochemical results were consistent with previous TET protein levels (Figure 4B). TET-im intensity was significantly increased in NSCs subjected to VPA when compared with control (Figure 4B). Furthermore, following rapamycin treatment, TET-im intensity was significantly lower in VPA + rapamycin-treated cells than in VPA-treated cells (Figure 4B).

To further verify the effect of TETs on active DNA demethylation, an anti-5-hmC antibody-based dot blot analysis showed an increase of 5-hmC induced by VPA treatment (Figure 4C). Whereas rapamycin and VPA treatment strongly decreased 5-hmC levels (Figure 4C), consistent with our dot blot results we found that 5-hmC covered an increasing amount of the genome, and 5-hmC-im became more intense in VPA-treated cells, a low density for 5-hmC signature was observed in cells treated by VPA + rapamycin (Figure 4D). To determine if 5-hmC has a functional role in regulating neural differentiation, we calculated enrichment of 5-hmC at the Ngn1 promoter and found that 5-hmC becomes especially enriched in VPA-treated cells (Figure 4E). We also observed a significant reduced enrichment of 5-hmC at promoter region of Ngn1 in VPA + rapamycin-treated cells (Figure 4E). These results confirmed that the loss of Ngn1 promoter DNA methylation in NSCs treated with VPA is induced by increased expressions.
of active DNA demethylation pathway enzymes through activating mTOR signaling.

**VPA but Not PI3K/AKT/mTOR Activity Regulates Histone Modifications in the Ngn1 Gene Region**

As mentioned above, VPA elevated the Ngn1 expression in differentiated cells and also regulated the expression of HDACs. To determine whether PI3K/AKT/mTOR signaling mediates VPA-induced Ngn1 expression via regulating the histone modifications, we employed western blot and chromatin immunoprecipitation (ChIP) assays to test histone acetylation and histone modifications at the proximal promoter of the Ngn1 gene after VPA treatment. We detected an abundance of acetylated histone H3 and H4 in extracts of cells treated by VPA (Figure 5A). Interestingly, the level of acetylated histone H3 and H4 remained relatively unchanged in extracts of cells treated by VPA + wortmannin compared with VPA (Figure 5A), suggesting that there is less functional relationship between PI3K/AKT/mTOR signaling and histone acetylation during VPA-induced differentiation. To further define the relationship, we investigated histone modifications at the proximal promoter of the Ngn1 gene after VPA and/or rapamycin treatment. Our data showed that, while trimethylated H3K4 (H3K4me3) significantly increased with VPA treatment at the proximal promoter of the Ngn1 gene, acetylated H3K9 (H3K9Ac) was also obviously enhanced (Figure 5B). The results suggest that the efficient recruiting of H3K4me3 and H3K9Ac at the promoter is a prerequisite for the activation of the Ngn1 gene after VPA treatment. We then analyzed the recruiting of methylated lysine in H3K27 and H3K9 on the promoter region of Ngn1 with ChIP assay. Our observation showed that the cells treated with VPA decreased the trimethylated H3K9 (H3K9me3) and dimethylated H3K9 (H3K9me2) levels in the Ngn1 promoter (Figure 5B). By contrast, no significant differences in histone modifications were found between VPA-treated and VPA + wortmannin-treated cells (Figure 5B). In addition, western blot analysis was in accordance with the above data (Figure 5C). These findings suggest that VPA but not PI3K/AKT/mTOR activity mediates histone modifications in the Ngn1 gene region.
DNMTs Inhibitor Rescues the Differentiation Arrested by PI3K/AKT/mTOR Inhibition or VPA Deprivation

To address whether the elevated expression of DNMT1 and 3a represents a significant cause for the differentiation arrested by loss of PI3K/AKT/mTOR activity or VPA deprivation, we applied the DNMT inhibitor, 5-aza-2'-deoxycytidine (AZA), to the culture medium. AZA is a cytidine analog that blocks actions of DNMTs and causes passive demethylation. Treatment with AZA in the presence of VPA + PI3K/AKT/mTOR inhibition resulted in a 1.7-fold increase in the number of neuronal nuclei (NeuN)-positive cells, compared with VPA + wortmannin treatment control (Figure 6A). Upon removing VPA and wortmannin, we could still observe a significant 2.2-fold increase in NeuN-positive cells from AZA-treated cells compared with control (Figure 6A). Immunohistochemical staining for doublecortin (DCX) and western blot further confirmed these data (Figures 6A and 6B). These results indicated that inhibition of DNMT1 and 3a significantly rescues the trapped differentiation by PI3K/AKT/mTOR inhibition or VPA deprivation.

TET1, TET2, and TET3 Are Necessary for the Neuronal Differentiation of NSCs Induced by VPA

To explore whether the activity of TETs is required for VPA-induced neuronal differentiation, loss of function of TETs was testified. We found that expression of TET1, TET2, and TET3 proteins was efficiently downregulated by shTET1, shTET12, and shTET13 treatments (Figure 7A), along with a reduction of 5-hmC protein as well as an increment of 5-mC protein compared with VPA-treated cells (Figures 7B and 7C). The knockdown of TET1, TET2, and TET3 markedly abolished VPA-mediated increase of DCX + neuron yields (Figure 7D) and Ngn1 protein expressions (Figure 7E). Furthermore, shTET1, shTET12, and shTET13 treatments in VPA-treated cultures led to a robust reduction of 5-hmC enrichment and increase of 5-mC in the Ngn1 gene promoter (Figures 7F and 7G). These findings suggested that VPA-induced neuron differentiation is dependent on TETs-mediated DNA hydroxymethylation/demethylation.

DISCUSSION

Some recent studies have shown that activation of the mTOR pathway participates in the neuronal differentiation regulation of mouse olfactory bulb stem cells and neural progenitor cells (Han et al., 2008; Otaegi et al., 2006). We demonstrated here that VPA activated mTOR signaling and elevated mTOR activity, significantly enhancing VPA-induced neuronal differentiation of NSCs. VPA-induced differentiation was suppressed by the mTOR inhibitor, rapamycin, indicating activation of the mTOR pathway in differentiation induced by VPA. Immunoblotting with antibodies against activated or total AKT, mTOR, and p70S6K showed that VPA activated this pathway, indicating the effects of VPA on NSCs are mediated, at least in part, by activating the AKT/mTOR/S6K pathway.

Wortmannin is a selective and irreversible inhibitor of PI3K and was used to distinguish whether VPA activated...
AKT via the classical PI3K pathway or via an alternate pathway. Wortmannin reduced AKT activation by VPA, indicating that AKT activation by VPA is dependent on PI3K. Moreover, wortmannin inhibited the neuronal differentiation promoted by VPA, confirming that PI3K-mediated AKT activation by VPA is necessary to induce neuronal differentiation in NSCs. Because wortmannin is a non-specific inhibitor of PI3K, we also found that PI3K p110α or PI3K p110β knockdown inhibited AKT activation and attenuated the neuronal differentiation of NSCs with VPA, which was consistent with the results of wortmannin.

However, the mechanism by which VPA activates the PI3K/AKT pathway is currently unknown but it may be mediated through direct effect of VPA on PI3K.

VPA is an HDAC inhibitor and plays a role in modifying chromatin structure and gene expression. The post translational modifications of the amino-terminal tails of the H3 and H4 histones also involve in general changes in condensation of chromatin connected with various tissue-specific genes, which maintain them in either an activated or repressed state. VPA treatment enhances histone acetylation, but also activates mTOR signaling. HDAC1 and HDAC2 knockdown demonstrated that the interaction with mTOR signaling is initiated by histone H3 acetylation (Makarević et al., 2014). The HDAC inhibition represses cancer proliferation via suppressing PI3K/mTOR pathways (Wilson-Edell et al., 2014).

Figure 6. DNMTs Inhibitor Rescues Impeded Differentiation by PI3K/AKT/mTOR Inactivation or VPA Deprivation
(A) Immunofluorescence staining and quantification for NeuN and DCX on NSCs under different treatment conditions. Scale bar, 100 μm. **p = 0.002 < 0.01 compared with control; §§p = 0.004 < 0.01 compared with control; $$p = 0.006 < 0.01 compared with VPA + wortmannin; &p = 0.006 < 0.01 compared with VPA + wortmannin. Data are from three independent experiments.

(B) The expression of Ngn1 protein was analyzed by immunoblotting in different treatment groups. **p = 0.004 < 0.01 compared with control; §§p = 0.006 < 0.01 compared with VPA + wortmannin. Data are from three independent experiments.

Histone acetylation and DNA methylation operate in a concerted manner. In agreement with this hypothesis, HDAC inhibitors such as MS-275 and TSA also reduce DNMT1, 3a and 3b protein levels and enzyme activity in NT-2 precursor cells, tentatively explaining the mechanism of DNA demethylation (Kundakovic et al., 2009). In cultured NSCs, we found no changes in DNMT3b and DNMT3L levels upon VPA administration. VPA or other HDAC inhibitors induce promoter demethylation by activating DNA demethylation mechanisms (Detich et al., 2003; Guidotti et al., 2011). We discovered that PI3K/AKT/mTOR inhibition enhanced DNMT1 and 3a expression and thus enhanced the stimulation of methylation activity specific for neuronal gene silencing during differentiation. Although there is no direct evidence, our finding that PI3K/AKT/mTOR activity stimulated DNMT1 and 3a expression therefore sheds new light on the role played by VPA signaling for both inhibition of proliferation and differentiation of NSCs. Inhibition of HDAC activity...
promotes the ubiquitin-dependent proteasomal degradation of DNMT1 (Du et al., 2010), further suggesting a synergistic promotion of DNA demethylation by suppression of DNMTs activities. Furthermore, we also observed that VPA promotes the DNA hypomethylation in mTOR-dependent manner. The fact that mTOR activation is necessary for other protein kinases to decrease gene-specific DNA methylation suggests that mTOR signaling promotes the ubiquitin-dependent proteasomal degradation of DNMT1 (Du et al., 2010), further suggesting a synergistic promotion of DNA demethylation by suppression of DNMTs activities. Furthermore, we also observed that VPA promotes the DNA hypomethylation in mTOR-dependent manner. The fact that mTOR activation is necessary for other protein kinases to decrease gene-specific DNA methylation suggests that mTOR signaling promotes the ubiquitin-dependent proteasomal degradation of DNMT1 (Du et al., 2010), further suggesting a synergistic promotion of DNA demethylation by suppression of DNMTs activities. Furthermore, we also observed that VPA promotes the DNA hypomethylation in mTOR-dependent manner. The fact that mTOR activation is necessary for other protein kinases to decrease gene-specific DNA methylation suggests that mTOR signaling
may critically link cell-signaling mechanisms with DNA methylation.

VPA is a well-known DNA demethylating agent. It can induce passive DNA demethylation by suppressing DNMT1, DNMT3a, and DNMT3b, which triggers the activation of genes (Dong et al., 2010). The evidence we present here thus supports a model where PI3K/AKT/mTOR activation by VPA orchestrates the epigenetic events during differentiation, allowing complete activation of the neuron-specific gene, Ngn1. We found that hypermethylation of the Ngn1 promoter region, as the result of PI3K/AKT/mTOR inhibition, is tightly correlated with the significantly elevated DNMT1 and 3a expression compared with control cells. Accordingly, we demonstrated that inhibiting DNMT activity by AZA significantly enhanced Ngn1 expression and rescued the differentiation arrested by PI3K/AKT/mTOR inhibition or VPA deprivation. Our data thus strongly argue that PI3K/AKT/mTOR signaling plays a critical function in facilitating the passive demethylation of endogenous global and neuron-specific genes by suppressing the expression of DNMT1 and 3a in cells with VPA.

Active demethylation of DNA is a common mode of epigenetic control of tissue-specific gene expression during development through TET enzymes (Hirabayashi and Gotoh, 2010). DNA 5-hmC, catalyzed by TET enzymes, recently opened up as an important epigenetic signature to control biological activities in developing and adult tissues (Kriaucionis and Heintz, 2009). Our study showed that TET1, TET2, and TET3 were expressed in NSCs. Addition of VPA to the NSC cultures enhanced TET enzyme activity, and led to a robust increase of global and local 5-hmC in the Ngn1 gene promoter region as well as in the number of DCX-positive cells. In NSCs cells, a 5-hmC increase was followed by a substantial decrease of 5-mC, indicating that 5-hmC is an intermediate to be eliminated ultimately by the base excision repair pathway. By contrast, the decreases of 5-mC were shown in the VPA-treated NSCs in our study, which is consistent with the previous report of the 5-mC pattern in adult mouse brain (Dong et al., 2010). However, our TET1, TET2, and TET3 knockdown experiments in the presence of VPA revealed an interaction in the opposite direction. These data suggest that 5-hmC may slowly be converted to cytosine during differentiation. This is in agreement with earlier reports on the redundancy and compensatory actions of 5-hmC or TET family proteins (Etchegaray et al., 2015; He et al., 2015) in accounting for global effects. Moreover, we demonstrated that inhibition of PI3K/AKT/mTOR blocks DNA demethylation of Ngn1 regulatory regions induced by VPA via suppressing TET1, TET2, and TET3, indicating a strong connection between mTOR signaling and TET enzymes during Ngn1 gene activation.

Epigenetic control of long-distance gene expression by histone modifications is a pivotal mechanism determining cell-fate and cell-phenotype maintenance (Rivera and Ren, 2013). Compared with the generally permissive histone acetylation, histone methylations are more diverse in function and their regulation is complex and dynamic. Methylated histones are associated with promoter activation (H3K4me3) and a repressive heterochromatin structure (H3K9me2, K27me3) (Heintzman et al., 2007; Mosammam-parast and Shi, 2010). Enzymes responsible for these modifications are histone methyltransferases and demethylases. We showed that DNA hydroxymethylation promoted by VPA was accompanied with the formation of an open chromatin structure. VPA regulated histone acetylation and methylation, which activates neuron-specific gene expression. Our data showed that VPA exposure caused a reduction in the levels of trimethylated histone H3 on lysine 4 (H3K27 me3), and histone H3-dimethylated lysine 9 (H3K9me2) on the Ngn1 promoter region accompanies this gene repression in differentiating NSCs. Concomitantly, relatively high levels of the trimethylation of lysine 4 (H3K4 me3) and histone H3-acetylated lysine 9 (H3K9Ac) on the Ngn1 promoter region were found. These data suggest that the profoundly increased activity of the Ngn1 gene is associated with transcriptionally active markers (H3K4me3 and H3K9Ac) and two others are associated with repressed markers (H3K9me3 and H3K27me3). A similar conclusion was derived from the observation that the active H3K4me3 and repressive H3K27me3 markers are detected in a large number of developmental regulator genes in mammalian ESCs (Bernstein et al., 2006) and adult stem cells (Cui et al., 2009; Wei et al., 2009). Our present findings also show that mTOR inhibition does not markedly affect the expression of the above H3 modifications during differentiation, indicating that VPA may promote histone modifications through suppressing HDACs activity rather than activating mTOR signaling.

**EXPERIMENTAL PROCEDURES**

Full experimental procedures are provided in the Supplemental Information.

**Cell Culture and Transfection**

Hippocampi were dissected from embryonic day 16 (E16) embryos of adult pregnant Sprague-Dawley rats into Hank's balanced salt solution without calcium or magnesium. The dissociated cells were plated on dishes coated with 50 μg/mL poly-L-ornithine and 10 μg/mL fibronectin (Sigma-Aldrich) in N2 medium and incubated at 37°C in a humidified atmosphere of 5% CO2 and 95% air. Fibroblast growth factor 2 (FGF2, 20 ng/mL, Millipore) was added to expand the population of proliferative progenitors. Cells at 80% confluence were subcultured in N2 medium in the presence of...
Immunofluorescence
Cells plated on chamber slides were fixed with 4% paraformaldehyde, permeabilized with 0.2% Triton X-100, and incubated with the primary antibody at 4°C overnight. Chamber slides were subsequently incubated with fluorescein isothiocyanate- or tetramethylrhodamine-conjugated secondary antibody for 1 hr at room temperature. Coverslips were mounted in Gel Mount (VECTORSHIELD) and imaged with a Nikon E400 fluorescent microscope (Nikon). The name, catalog numbers, and supplier for antibodies are listed in Table S1.

Statistical Analysis
One-way ANOVA followed by Bonferroni’s post hoc test was performed. Student’s t test was used for comparisons of two groups. p Values <0.05 were considered significant.

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Experimental Procedures, one figure, and three tables and can be found with this article online at http://dx.doi.org/10.1016/j.stemcr.2017.04.006.

AUTHOR CONTRIBUTIONS
Conception and Design, A.X., D.L., J.L., and W.J.; Performed Research, X.Z., X.H., Q.L., X.K., Z.O., L.Z., Z.G., and L.X.; Collection and Assembly of Data, M.Z., W.Z., L.Z., and Z.G.; Data Analysis and Interpretation, A.X., L.X., and W.J.; Manuscript Writing, A.X. and M.Z.; Final Approval of Manuscript, all authors.

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