Age-related loss of neural stem cell O-GlcNAc promotes a glial fate switch through STAT3 activation

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Increased neural stem cell (NSC) quiescence is a major determinant of age-related regenerative decline in the adult hippocampus. However, a coextensive model has been proposed in which division-coupled conversion of NSCs into differentiated astrocytes restrict the stem cell pool with age. Here we report that age-related loss of the posttranslational modification, O-linked β-N-acetylglucosamine (O-GlcNAc), in NSCs promotes a glial fate switch. We detect an age-dependent decrease in NSC O-GlcNAc levels coincident with decreased neurogenesis and increased gliogenesis in the mature hippocampus. Mimicking an age-related loss of NSC O-GlcNAcylation in young mice reduces neurogenesis, increases astrocyte differentiation, and impairs associated cognitive function. Using RNA-sequencing of primary NSCs following decreased O-GlcNAcylation, we detected changes in the STAT3 signaling pathway indicative of glial differentiation. Moreover, using O-GlcNAc-specific mass spectrometry analysis of the aging hippocampus, together with an in vitro site-directed mutagenesis approach, we identify loss of STAT3 O-GlcNAcylation at Threonine 717 (T717) as a driver of astrocyte differentiation. Our data identify the posttranslational modification, O-GlcNAc, as a key molecular regulator of regenerative decline underlying an age-related NSC fate switch.

Significance

Depletion of the neural stem cell (NSC) pool is a major driver of age-related regenerative decline in the hippocampus. While increased quiescence is a major contributor to this decline, NSCs can also undergo terminal differentiation into astrocytes, thus restricting the stem cell pool. The mechanisms underlying this fate switch and their relation to age-related regenerative decline have not yet been fully elucidated. In this study, we report an age-related decline in NSC O-GlcNAcylation, coincident with reduced neurogenesis and increased gliogenesis. We identify loss of O-GlcNAcylation at STAT3 T717 in the hippocampus with age, and demonstrate that O-GlcNAcylation of this site is a critical determinant of NSC fate. Our work expands our understanding of how posttranslational modifications influence the aging brain.


The authors declare no competing interest.

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immunohistochemical analysis. We detected a sharp decrease in O-GlcNAc levels in Sox2/GFAP+ NSCs in mature compared to young mice (Fig. 1A and B). Concurrently, this decline was paralleled by a decrease in the number of Nestin+ NSCs, double-cortin (Dcx)/MCM2+ neuroblasts, and Dcx+/MCM2− newly born neurons in the mature hippocampus (Fig. 1C and D), consistent with previous reports (7, 10). While glial differentiation has been reported at old age (3, 27), whether the decline in adult neurogenesis in the mature hippocampus is concurrent with an increase in gliogenesis is unknown. Therefore, we assessed astrocyte differentiation using a long-term 5-bromo-2′-deoxyuridine (BrdU) incorporation paradigm, in which differentiated astrocytes express both BrdU and the glial markers GFAP or S100β. We observed an age-dependent increase in the number of long-term retaining

![Graphs and images showing changes in markers over time](https://www.pnas.org/cgi/doi/10.1073/pnas.2007439117)

**Fig. 1.** Age-related decreased NSC O-GlcNAc levels is coincident with a decline in neurogenesis and an increase in gliogenesis in the mature hippocampus. (A) Representative single z-planes and orthogonal projections of a confocal image z-stack of 2- or 6-mo-old mouse hippocampus, labeled with anti-Sox2, anti-GFAP, anti-O-GlcNAc, and DAPI. (Scale bar, 10 μm.) (B) Quantification of the fold-change of the average O-GlcNAc intensity normalized to area measured in each GFAP+/Sox2+ NSC across all z-stacks per animal (n = 4 animals per group). (C) Representative maximum intensity projections of a confocal image z-stack of 2- or 6-mo-old mouse hippocampus, labeled with DAPI and anti-Nestin (Upper) or DAPI, anti-Dcx, and anti-Mcm2 (Lower). (Scale bar, 100 μm.) (D) Quantification of Nestin+ radial NSCs (Top), Dcx+/MCM2+ proliferating neuroblasts (Middle), and Dcx+/MCM2− immature neurons (Bottom) in 2- or 6-mo-old mice (n = 4 to 5 animals per group). (E) Representative maximum intensity projections of a confocal image z-stack of 2- or 6-mo-old mouse hippocampus, labeled with anti-GFAP and anti-BrdU. Mice were given six daily intraperitoneal injections of BrdU (50 mg/kg) 30 d before killing. Quantification of the percentage of GFAP+/BrdU+ astrocytes out of total BrdU+ cells in the subgranular zone (SGZ) of the DG (n = 4 to 5 animals per group; arrowheads, astrocytes; white dashed line, SGZ). (Scale bar, 10 μm; also applies to F.) (F) Representative maximum-intensity projections of a confocal image z-stack of 2- or 6-mo-old mouse hippocampus, labeled with anti-s100β and anti-BrdU (Left). Mice were treated as in E. Quantification of the percentage of s100β+/BrdU+ astrocytes out of total BrdU+ cells in the SGZ of the DG (n = 4 to 5 animals per group; arrowheads, astrocytes; white dashed line SGZ). Data are represented as mean ± SEM; **P < 0.01, ***P < 0.001, ****P < 0.0001; t test.
BrdU/GFAP<sup>+</sup> and BrdU/S100β<sup>+</sup> astrocytes in the mature compared to the young hippocampus (Fig. 1 E and F), in a timeframe consistent with decreased neurogenesis. These data raise the possibility that decreased NSC O-GlcNAcylation in the mature hippocampus is associated with age-related regenerative decline.

**Decreased NSC O-GlcNAc Levels Impairs Proliferation and Promotes Astrocyte Differentiation In Vitro.** To gain insight into the functional consequence of reduced NSC O-GlcNAcylation, we made use of a primary hippocampal NSC in vitro model. Using RNA-seq, we profiled gene expression in primary NSCs following either pharmacological inhibition of Ogt with OSMI-1 treatment (*SI Appendix*, Fig. S1) or viral-mediated abrogation of Ogt by RNAi (*SI Appendix*, Fig. S2). We found 845 differentially expressed genes following Ogt inhibition (*SI Appendix*, Fig. S2) and 741 differentially expressed genes after Ogt abrogation (*SI Appendix*, Fig. S3A). Gene ontology (GO) analysis of the most differentially down-regulated genes identified biological processes, including cell cycle, mitotic nuclear division, and cell division (Fig. 2B and *SI Appendix*, Fig. S3B). Functionally, we assessed NSC proliferation in vitro following inhibition or abrogation of Ogt and observed a significant decrease in the percentage of 5-Ethynyl-2′-deoxyuridine (EdU)<sup>+</sup> and K<sub>67</sub><sup>+</sup> proliferating cells compared to control conditions (Fig. 2C and *SI Appendix*, Fig. S3C). No differences in cell death were detected by immunocytochemical analysis for cleaved caspase-3 (*SI Appendix*, Fig. S4A). These in vitro data indicate that reducing NSC O-GlcNAcylation negatively regulates NSC proliferation.

Given the relatively low number of significantly up-regulated genes following Ogt inhibition (Fig. 2A), we performed gene list enrichment analysis using Enrichr for chromatin immunoprecipitation (ChIP) enricher analysis (ChEA) and ENCODE transcription factor targets datasets. We identified transcription factors previously associated with NSC function, including STAT3 (Fig. 2D and *SI Appendix*, Fig. S3D), a major initiator of astrocyte differentiation (28, 29). Correspondingly, we surveyed expression of known astrocyte genes in NSCs following Ogt inhibition using our RNA-seq data to annotate several other astrocyte genes, including *Sl100y*, *Aldh1l1*, and *Ssa9* (Fig. 2E). Next, we examined whether reducing NSC O-GlcNAc levels impacted NSC differentiation in vitro. Primary hippocampal NSCs isolated from either postnatal or adult (2 mo) mice were treated with the Ogt inhibitor OSMI-1 or vehicle control and cultured under differentiation conditions. Ogt inhibition resulted in a concomitant decrease in neuronal (Fig. 2F) and increase in astrocyte (Fig. 2G) differentiation compared to vehicle control conditions. To ensure that these changes in cell fate were not due to increased cell death during differentiation, we analyzed cytotoxicity (*SI Appendix*, Fig. S4B) and cell viability (*SI Appendix*, Fig. S4C) throughout the paradigm and observed no difference between Ogt inhibitor-treated and control NSCs. These data suggest that the observed loss of proliferative capacity and neuronal differentiation following decreased NSC O-GlcNAcylation (Fig. 1) may be the result of an altered neuron to glia fate switch.

**Mimicking an Age-Related Decline in NSC O-GlcNAcylation Impairs Adult Neurogenesis and Cognitive Function and Increases Gliogenesis in the Young Hippocampus.** To understand the relationship between O-GlcNAc expression and adult neurogenesis in vivo, we asked whether mimicking an age-related decline in NSC O-GlcNAcylation within the adult hippocampus would impair neurogenesis. To resolve the role of O-GlcNAc in a temporally defined and cell-type-specific manner in the adult brain, we generated male Ogt<sup>fl/fl</sup> mice carrying an inducible NestinCre-ERT<sub>2</sub> gene, in which the X-linked Ogt gene is excised specifically in adult NSCs (Ogt cKO) upon tamoxifen administration (Fig. 3A). Young (2 mo) mice were administered tamoxifen and changes in neurogenesis were assessed in young adult (3 mo) Ogt cKO and littermate control (Ogt<sup>cre/cre</sup>) mice by immunohistochemical analysis. Loss of Ogt expression in adult NSCs resulted in a decrease in the number of Nestin<sup>+</sup> NSCs and Dcx/MCM2<sup>+</sup> neuroblasts in the hippocampus of Ogt cKO compared to littermate control animals (Fig. 3 B and C). We assessed neuronal differentiation and survival using a long-term BrdU incorporation paradigm, in which mature differentiated neurons express both BrdU and the neuronal marker NeuN. We observed a decrease in the number of long-term retaining BrdU/NeuN<sup>+</sup> mature differentiated neurons Ogt cKO compared to control animals (Fig. 3 B and C). To ensure that these differences in neurogenesis were not due to increased cell death, we compared the number of cleaved caspase-3<sup>+</sup> cells and observed no difference in cell death between Ogt cKO and littermate control mice (*SI Appendix*, Fig. S5). To corroborate our genetic studies, we abrogated hippocampal Ogt expression in wild-type animals utilizing an in vivo viral-mediated RNAi approach. Young adult (3 mo) animals were stereotaxically injected with high-titer lentivirus encoding Ogt or control shRNA sequences into the DG of contralateral hippocampi (*SI Appendix*, Fig. S6A). Decreased Ogt in the DG resulted in a significant decrease in the number of Dcx<sup>+</sup>/EdU<sup>+</sup> neuroblasts and long-term retaining BrdU/NeuN<sup>+</sup> neurons compared with the control contralateral DG (*SI Appendix*, Fig. S6B). Collectively, our genetic and in vivo RNAi data indicate that loss of NSC O-GlcNAcylation mirrors an aging condition by negatively regulating neurogenesis in the adult hippocampus.

To investigate whether mimicking an age-related decline in NSC O-GlcNAcylation also impaired cognitive processes associated with adult neurogenesis, we assessed hippocampal-dependent object recognition and associative fear memory using novel object recognition (NOR) and contextual fear-conditioning paradigms, respectively (Fig. 3A). In the testing portion of the NOR task, control animals exhibited a significant preference for the novel object over the familiar object; however, this preference was lost in Ogt cKO mice (Fig. 3D). During contextual fear-conditioning testing, we observed decreased freezing in Ogt cKO mice compared to controls animals (Fig. 3E). No differences were observed in baseline freezing during fear-conditioning training (*SI Appendix*, Fig. S7A) or in cued fear-conditioning testing between genotypes (Fig. 3E). As a control, we profiled general health using an open-field paradigm (*SI Appendix*, Fig. S7B) and observed no differences in overall activity, total distance traveled, or time spent in the periphery or center of the open field, indicative of normal motor and anxiety functions (*SI Appendix*, Fig. S7C–E). These behavioral data demonstrate that decreased adult NSC O-GlcNAcylation impairs hippocampal-dependent learning and memory.

Next, to examine the relationship between O-GlcNAcylation and gliogenesis, we asked whether mimicking an age-related decline in NSC O-GlcNAc levels would promote astrocyte differentiation using a genetic lineage-tracing approach. We generated male Ogt cKO and control mice carrying a loxp-flanked STOP cassette preventing transcription of a CAG-promoter driven tdTomato (tdT) reporter inserted into the Gt(Rosa)<sub>26</sub>Sor locus, in which tdT is expressed in adult NSCs upon tamoxifen administration. Young (2 mo) mice were administered tamoxifen and changes in gliogenesis were assessed in young adult (3 mo) Ogt cKO and littermate control mice by immunohistochemical analysis. Efficacy of our knockout model was confirmed by immunohistochemistry (*SI Appendix*, Fig. S8). Loss of Ogt expression in adult NSCs resulted in an increase in the number of tdT/GFAP<sup>+</sup> and tdT/S100β<sup>+</sup> astrocytes in the hippocampus of Ogt cKO compared to control animals (Fig. 3 F and G). Our in vivo lineage tracing analysis identifies an increase in adult gliogenesis in young mice following decreased NSC O-GlcNAcylation that reflect age-related changes in the mature hippocampus (Fig. 1 E and F).

**O-GlcNAcylation of STAT3 at T717 Is Reduced in the Aging Hippocampus and Promotes a Glate Fase Switch in NSCs.** To gain mechanistic insight into site-specific changes in O-GlcNAcylation potentially regulating
Fig. 2. Decreased NSC O-GlcNAcylation promotes a neuron to glia fate switch in vitro. (A) Heatmap of significantly differentially expressed genes with fold-change $> 2$ and $P < 0.05$ in primary hippocampal NSCs following 24-h 12.5 μM OSMI-1 treatment compared to vehicle control in vitro. (B) Biological process GO terms associated with genes down-regulated following OSMI-1 treatment. (C) Representative field and quantification of primary mouse hippocampal NSCs following 24 h of vehicle (Left) or OSMI (Right) treatment, labeled with DAPI and an 8-h EdU-pulse (Upper), or DAPI and anti-Ki67 (Lower) ($n = 3$ per group). (Scale bars, 10 μm.) (D) Quantification of enriched transcription factor targets as identified by ChEA or ENCODE based on up-regulated genes after 24-h OSMI-1 treatment compared to vehicle control. (E) Fold-change expression (FPKM) of astrocyte-associated genes: S100β, Aldh1l1, and Sox9 in primary hippocampal NSCs following OSMI-1 treatment in vitro ($n = 5$ to 6 per group). (F, G) Differentiation was induced in vehicle or OSMI-1 treated primary postnatal (P1) and adult (2-mo-old) hippocampal NSCs by withdrawing growth factors 7 d prior to fixation. Cells were labeled with DAPI and anti-MAP2. MAP2+ neurons were quantified as a percentage of total DAPI+ cells ($n = 6$ replicates per group). (Scale bar, 10 μm.) (G) Primary postnatal and adult hippocampal NSCs were treated as in F. Cells were labeled with DAPI and anti-GFAP. GFAP+ astrocytes were quantified as a percentage of total DAPI+ cells ($n = 6$ per group). (Scale bar, 10 μm.)
Fig. 3. Decreased NSC O-GlcNAcylation impairs adult neurogenesis and cognitive function while increasing gliogenesis in the young hippocampus. (A) Schematic of experimental paradigm and cognitive testing timeline. Three-month-old Ogt<sup>lox/lox</sup> (control) or NestinCre-ERT2<sup>−/−</sup>; Ogt<sup>lox/lox</sup> (Ogt cKO) littermate mice were administered five daily intraperitoneal tamoxifen injections (75 mg/kg). One month later, hippocampal-dependent learning and memory were assessed by NOR or contextual fear-conditioning paradigms. One week after behavioral assays, mice were administered six daily intraperitoneal injections of BrdU (50 mg/kg). Thirty days later mice were killed, and tissue was analyzed. (B) Representative maximum-intensity projections of a confocal image z-stack of control or Ogt cKO hippocampus, labeled with DAPI and anti-Nestin (Top) or DAPI, Dcx, and anti-Mcm2 (Middle), or anti-NeuN and anti BrdU (Bottom). (Scale bar, 100 μm, Top or 10 μm, Bottom.) (C) Quantification of Nestin<sup>+</sup> radial NSCs (Top), Dcx/Mcm2<sup>+</sup> proliferating neuroblasts (Middle), and NeuN<sup>+</sup>/BrdU<sup>+</sup> newborn mature neurons (Bottom) in control or Ogt cKO littermate mice (n = 10 to 11 animals per group). (D) Object recognition memory was assessed in control and Ogt cKO littermate mice using the NOR task. Quantification of percentage of time spent with the novel object is shown (n = 11 to 12 per group). (E) Associative fear memory was assessed in Ogt cKO and control littermate mice using contextual fear conditioning. Quantification of percentage freezing in response to cue or context 24 h after training is shown (n = 12 per group). (F) Three-month-old NestinCre-ERT2<sup>−/−</sup>; Ai14<sup>lox/lox</sup> (control) or NestinCre-ERT2<sup>−/−</sup>; Ai14<sup>lox/lox</sup>; Ogt<sup>lox/lox</sup> (Ogt cKO) littermate mice were administered five daily intraperitoneal tamoxifen injections (75 mg/kg). One month later, mice were killed and tissue was analyzed. Representative maximum intensity projections of a confocal image z-stack of control or Ogt cKO hippocampus, labeled with endogenous TdTomato and anti-GFAP. Quantification of TdTomato/GFAP<sup>+</sup> astrocytes (n = 3 to 5 per group; arrowheads, astrocytes; white dashed line, SGZ). (Scale bar, 10 μm,) (G) Mice were treated as in F. Representative maximum-intensity projections of a confocal image z-stack of control or Ogt cKO hippocampus, labeled with endogenous TdTomato and anti-s100β. Quantification of TdTomato/s100β<sup>+</sup> astrocytes (n = 3 to 5 per group; arrowheads, astrocytes; white dashed line, SGZ). (Scale bar, 10 μm.) Data are represented as mean ± SEM; *P < 0.05, **P < 0.01, ***P < 0.001; t test (C and E–G); one-sample t test versus 50% (D).
impaired NSC function during aging, we performed comparative
O-GlcNAc-specific proteomics analysis of the hippocampus at
ages before and after changes in adult neurogenesis and gli-
genesis (Fig. 4A). Due to technical limitations in the number
of aged animals necessary to obtain sufficient amounts of primary
NSCs to enrich for dynamic posttranslational modifications, we
opted to subdissect and pool hippocampi from young adult (3 mo),
middle aged (12 mo), and old (24 mo) mice. We isolated total
hippocampal protein, trypsin digested, and labeled the resulting
peptides with isobaric tandem mass tags (TMT) before enriching
for O-GlcNAcylated peptides using lectin weak-affinity chroma-
tography (LWAC). Glycopeptides were analyzed by electron-
transfer/higher-energy collision dissociation (ETdH) mass spec-
trometry. We detected 1,344 O-GlcNAcylation sites across 474
unique proteins (Dataset S1). We next examined mechanistic
synergies between decreased sites of protein O-GlcNAcylation in
the aging hippocampus (Dataset S1) and cell-type-specific mo-
lecular changes elicited in NSCs by decreased O-GlcNAc levels
(Fig. 2 and SI Appendix, Fig. S3), and selectively identified STAT3.
A representative ETdH spectrum used to resolve an unambigu-
ous O-GlcNAcylation site on a peptide from the STAT3 protein
(Fig. 4B), as well as a schematic depicting an age-related decrease
in STAT3 O-GlcNAcylation at T717 and Serine 719 (S719) (Fig. 4C)
(Fig. 4A are provided). Collectively, our proteomic analysis of the
aging hippocampus, together with RNA-sequencing of primary
NSCs, indicate changes in STAT3 signaling downstream of age-
related decreased O-GlcNAcylation.

Currently, the functional consequences of STAT3 O-GlcNAcylation
at S719 are unknown; however, increased STAT3 O-GlcNAcylation
at T717 has been previously shown to inhibit STAT3 activation in
immune cells by preventing STAT3 phosphorylation at Tyrosine
705 (Y705) (30). We therefore investigated whether decreasing
O-GlcNAcylation in primary NSCs could conversely increase
STAT3 activation. We assessed NSC STAT3 phosphorylation at
Y705 following Ogt inhibition with OSMI-1 treatment in vitro by
Western blot (Fig. 4D). While STAT3 expression was not altered,
pharmacological inhibition of Ogt significantly increased phos-
phorylation levels of STAT3 in NSCs (Fig. 4D). Additionally, we
made use of our RNA-sequencing dataset to survey expression of
known STAT3 target genes in NSCs following Ogt inhibition, and
likewise observed an up-regulation of several genes associated
with STAT3 activation (31), including p21, Zeb1, and Ccncl1
(Fig. 4E).

To investigate whether blocking STAT3 activation would miti-
gate the effects of decreasing O-GlcNAcylation on differentiation
in postnatal and adult NSCs, we generated viral expression con-
structs encoding a nonphosphorylatable STAT3 by converting
Tyrosine into Phenylalanine at position 705 (Y705F STAT3). As
a control, primary NSCs were infected with a virus encoding a wild-
type STAT3. When treated with OSMI-I, control NSCs exhibited
decreased neuronal (Fig. 4F) and increased astrocyte (Fig. 4G)
differentiation, corroborating our previous findings (Fig. 2 F and
G). Expression of Y705F STAT3 increased neuronal (Fig. 4F) and
decreased astrocyte differentiation (Fig. 4G), consistent with
previous reports (28, 29). Moreover, the effects of decreasing NSC
O-GlcNAcylation on STAT3 activation (SI Appendix, Fig. S9A)
and neuronal differentiation (Fig. 4F) were mitigated by expres-
sion of the Y705F STAT3 mutant. Next, we investigated whether
mimicking an age-related decline in STAT3 O-GlcNAcylation at
T717 altered NSC differentiation. It should be noted that the T717
site is only present on the predominant, full-length STAT3 α-
isoform, and not on the shorter β-isoform. We generated viral ex-
pression constructs encoding STAT3 with site-directed mutations
converting Threonine to Alanine at position 717 (T717A STAT3).
Expression of O-GlcNAcylation-deficient T717A STAT3 in NSCs
resulted in increased activation of the STAT3 α-isoform (SI Ap-
pendix, Fig. S9B), along with decreased neuronal (Fig. 4H) and
increased astrocyte (Fig. 4J) differentiation compared to wild-type
STAT3 control conditions. These in vitro data indicate that increased
STAT3 activity, stemming from reduced NSC O-GlcNAcylation,
promotes an astrocyte differentiation program in NSCs.

Increased NSC O-GlcNAc Levels Promote Neuronal and Inhibit Astrocyte
Differentiation In Vitro. Given the prolagiogenic effects of reducing
NSC O-GlcNAc levels, we next investigated whether increasing
NSC O-GlcNAc levels would conversely promote neuronal differ-
entiation. We generated a viral expression construct encoding
Ogt under the control of the Nestin promoter and infected pri-
mary postnatal and adult NSCs with this virus or a control
encoding GFP. Ogt overexpression in NSCs resulted in a signifi-
cant increase in Ogt and O-GlcNac levels (SI Appendix, Fig. S10).
Additionally, we observed an increase in neuronal differentiation
(Fig. 4J) and a corresponding decrease in astrocyte differentiation
(Fig. 4K) in NSCs overexpressing Ogt compared to a GFP control.
These data indicate that increased O-GlcNAcylation is sufficient
to promote a neuronal differentiation program and inhibit astro-
cyte differentiation in NSCs.

Discussion
Cumulatively, our data indicate that age-related loss of NSC
protein O-GlcNAcylation regulates a STAT3-mediated neuron to
glia fate switch that contributes to a decline in adult hippocampal
neurogenesis. While increased NSC quiescence has been highlighted
as a major determinant of age-related neurogenic decline in the
hippocampus (7, 32), a complementary model has been proposed in
which division-coupled conversion of NSCs into mature astrocytes
restrict the stem cell pool with age (3). Our data identifying an
O-GlcNAc-mediated glial fate switch in the mature hippocampus
provide evidence for this coextensive cellular mechanism, in which
age-related increased gliogenesis contributes to regenerative decline
and associated cognitive impairments.

Previous reports have observed that the absolute number of
newly generated astrocytes in the adult brain remains relatively
constant with age (3, 33, 34). Given the well-documented age-
related decline of total NSC number, it has been suggested that
maintaining this constant level of gliogenesis from a shrinking
adult NSC pool requires that an increasing proportion of NSCs
adopt a gliogenic fate (3, 35), a phenomenon that we demon-
strate in the mature hippocampus. While adult neurogenesis has
been shown to be a division-coupled process capable of main-
taining the NSC pool (3, 5), gliogenesis is indicative of a terminal
cell-cycle exit and leads to a reduction in total NSC number (3).
This shift toward terminal astrocyte differentiation with age
further exacerbates the age-related decline of the NSC pool. In
fact, mathematical modeling of NSC differentiation estimates that
maintaining constant levels of gliogenesis accounts for nearly half
of the NSCs lost during aging (34). In our study, we demonstrate
that this shift in NSC cell fate occurs alongside a decrease in NSC
O-GlcNAc. Moreover, we find that O-GlcNAcylation of STAT3
T717 declines in the hippocampus with age, and that loss of this
posttranslational modification is sufficient to promote astrocyte
differentiation in NSCs. Ultimately the shrinking of the adult NSC
pool, which stems in part from terminal astrocyte differentiation,
results in decreased neurogenesis during aging, with major func-
tional implications for age-related decreased regenerative capacity
and neurodegenerative disease. Indeed, in elderly humans, main-
taining higher levels of hippocampal neurogenesis is thought to be
neuroprotective, as they are associated with increased cognitive
scores and slower disease progression in the context of Alzheimer’s
disease (13, 14).

The ability to utilize the neurogenic capacity of adult NSCs is
predicated on understanding underlying molecular mechanisms
that can be targeted to reverse the effects of aging in the brain.
Although posttranslational modification of proteins represents a
fundamental mechanism by which information is transmitted, its
role in regulating adult NSC function with age has been largely

Fig. 4. STAT3 O-GlcNAcylation at T717 is reduced in the aging hippocampus and mimicking this decline in NSC promotes a gliogenic fate switch. (A) Schematic of proteomic workflow. The hippocampus was isolated from 3-, 12-, or 24-mo-old mice. Hippocampi were pooled to obtain sufficient protein (n = 5 animals per pool). Proteins were isolated from tissue and digested with trypsin before undergoing isobaric labeling. Enrichment was performed with TMT-labeled peptides for LC-MS/MS and analyzed via electron transfer dissociation mass spectrometry (ETD-MS). (B) Representative spectra of hippocampal proteins isolated from OGT-deficient and wild-type NSCs were infected, treated, and differentiated as in F. Cells were labeled with DAPI and anti-GFAP. GFAP+ astrocytes were quantified as a percentage of total DAPI+ cells (n = 6 replicates per group). (C) Primary postnatal and adult hippocampal NSCs were infected with a lentiviral construct expressing a nonphosphorylatable STAT3 (Y705F) or wild-type STAT3 (Control). Differentiation was induced in vehicle or OSMI-1 treated NSCs by withdrawing growth factors 7 d prior to fixation. Cells were labeled with DAPI and anti-GFAP. GFAP+ astrocytes were quantified as a percentage of total DAPI+ cells (n = 6 replicates per group). (D) Primary postnatal and adult hippocampal NSCs were infected, treated, and differentiated as in H. Cells were labeled with DAPI and anti-GFAP. GFAP+ astrocytes were quantified as a percentage of total DAPI+ cells (n = 6 replicates per group). (E) Primary postnatal and adult hippocampal NSCs were infected with a lentiviral construct expressing Ogt or GFP under the Nestin promoter. Differentiation was by induced withdrawing growth factors 7 d prior to fixation. Cells were labeled with DAPI and anti-GFAP. GFAP+ astrocytes were quantified as a percentage of total DAPI+ cells (n = 6 replicates per group). Data are represented as mean ± SEM; *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001; t test.
overlooked. Previous work has suggested a role for O-GlcNAcylation in mediating proper brain development. Specifically, genetic ablation of the O-GlcNAc cleaving enzyme, Oga, in NSCs during development has been shown to increase NSC proliferation and lead to an accumulation of progenitors and immature neurons in the olfactory bulb (26). In our study, we identify O-GlcNAc as a posttranslational regulator of regenerative decline in the adult brain and posit site-specific ameliorations in O-GlcNAcylation as potential targets to counter age-related regenerative dysfunction.

In particular, age-related loss of STAT3 O-GlcNAcylation at T717 provides a mechanistic link between increased NSC STAT3 signaling and an aberrant astrocyte differentiation program, pointing to STAT3 activation as a driver of age-related neurogenic decline. Interestingly, inhibition of STAT3 signaling in aged muscle stem cells restores regeneration at old age (35). Moreover, we demonstrate that increasing O-GlcNAc in NSCs by overexpressing Ogt is sufficient to promote neuronal differentiation. These data raise the exciting possibility of rejuvenating regenerative capacity in the aging brain by inhibiting STAT3 signaling through an increase in NSC O-GlcNAcylation. Ultimately, our data frame O-GlcNAc as a proneurogenic posttranslational modification, with functional implications for cognitive processes.

From a therapeutic perspective, while decreased O-GlcNAc has been implicated in Alzheimer’s disease pathology and neurodegeneration (18, 19), we recently demonstrated that increasing neuronal O-GlcNAcylation ameliorates age-related cognitive decline (17). Similarly, others have shown that inhibiting STAT3 dephosphorylation in neurons and astrocytes as a focal point in combating age-related functional decline in the aging and diseased brain. Of note, the benefits of interventions that ameliorate cognitive decline during aging and Alzheimer’s disease, such as exercise, have been shown to involve increased adult neurogenesis (37, 38). Given recent observations of the persistence of adult neurogenesis in the aged human brain (13, 14), the promise of restoring cellular and cognitive plasticity in the elderly by targeting adult NSCs is increasingly evident. Here we identify both an age-related NSC fate switch, and the posttranslational modification governing it, as potential cellular and molecular targets by which to maintain regenerative capacity at old age, restore adult neurogenesis and ameliorate cognitive decline.

Methods
Experimental Animal Model and Subject Details. All mouse handling and use was in accordance with institutional and ethical guidelines approved by the University of California, San Francisco Institutional Animal Care and Use Committee. The following mouse lines were used: C57BL/6J young mice (Jackson Laboratory), B6;129-Ogt<sup>Tg126</sup>(J) (The Jackson Laboratory line 004860), C57BL/6-Tg(Nes-creERT2)E8Jsc/Il (Jackson Laboratory line 016261), and B6.Gc-GIRDSAS26ortm1(CAG-tdTomato)Hert<sup>2</sup> (Jackson Laboratory line 007914). All studies were done in male mice. The number of mice used to result in statistically significant differences were calculated using standard power calculations with α = 0.05 and a power of 0.8. We used an online tool (http://www.stat.uiowa.edu/~rlenth/Power/index.html) to calculate power and sample size based on experience with the respective tests, variability of the assays, and interindividual differences within groups. Mice were housed under specific pathogen-free conditions under a 12-h light/dark cycle. All experiments were randomized and blinded by an independent researcher. Researchers remained blinded throughout histological, molecular, and behavioral assessments. Groups were unblinded at the end of each experiment upon statistical analysis.

Immunohistochemistry. Tissue processing and immunohistochemistry was performed on free-floating sections following standard published techniques (39). Briefly, mice were anesthetized with a ketamine (100 mg/kg)-xylazine (10 mg/kg) mixture (Paterson Veterinary, Henry Schein) and transcardially perfused with cold PBS. Brains were removed, fixed in phosphate-buffered 4% paraformaldehyde, pH 7.4, at 4 °C for 48 h followed by cryoprotection in 30% sucrose, and coronally sectioned at 40 μm with a cryostome (Leica SM2010 R). Sections were washed three times in Tris-buffered saline with 0.1% Tween 20 (TBST) and incubated in 3% normal donkey serum (Thermo Fisher Scientific) for 1 h. After overnight incubation in primary antibody (mouse anti-O-GlcNAc [R2L; 1:500], mouse anti-Nestin [clone rat-401; 1:200], goat anti-Actin [C-18; 1:7,500], mouse anti-BM28 [MCM2; 1:250], goat anti-Sox2 [Y-17; 1:200], rabbit anti-GFAP [1:1,000], mouse anti-NeuN [clone A60; 1:1,000], rat anti-Brdu [BU1/75 (ICR1)]; [1:10,000], mouse anti-Ki-67 [clone B56; 1:500], mouse anti-MAP2 [1:500; rabbit anti-s100] [1:500]) at 4 °C. Staining was revealed using fluorescence-conjugated secondary Alexa antibodies (donkey anti-rabbit: Alexa-488 conjugated secondary antibody; donkey anti-rabbit: Alexa-555 conjugated secondary antibody; donkey anti-rat: Alexa-488 conjugated secondary antibody; donkey anti-goat: Alexa-555 conjugated secondary antibody; donkey anti-mouse: Alexa-647 conjugated secondary antibody [1:500]. Antigen retrieval for BrdU labeling required incubation in 3M HCl at 37 °C for 30 min before incubation with primary antibody; Nestin labeling required incubation in Citrate Buffer (Sigma-Aldrich) at 95 °C (three times 5 min) prior to incubation with primary antibody. To estimate the total number of immunopositive cells per DG, confocal stacks of coronal sections of the DG (three to six sections per mouse, 40×/1.3 thick, 360×/um apart) were acquired on a Zeiss LSM 800.

In Vivo O-GlcNAc Quantification in NSCs. Immunohistochemistry and imaging were performed as previously described. To quantify O-GlcNAc levels, NSCs were identified based on GFAP/Sox2<sup>+</sup> staining, localization within the subgranular zone, and radial morphology. As O-GlcNAc staining is nuclear, DAPI was used to identify the nucleus of the analyzed cells. To quantify O-GlcNAc, NSCs in a single cell, intensity in the O-GlcNAc channel was measured within the region defined by DAPI and normalized to the area of that region in a single z-plane. This measurement was repeated over every plane in which that nucleus appeared in order to quantify the entirety of the cell. These values were then summed and normalized to the number of planes quantified for that cell. This process was repeated for every neuron stem cell which was fully included within the section (i.e., not within territory of another NSC). For each individual cell, averages were analyzed in all sections and averaged to determine a value per animal. Four animals were quantified per group. Fiji was used for quantification.

Isolation of Primary NSCs from the Mouse Hippocampus. Primary neuronal stem cell isolation and culture were preformed following previously published techniques (40). Hippocampi were dissected from male postnatal day 1 (P1) or adult (2-mo-old) wild-type C57/BL6 mice and pooled by age for NSC isolation. After removing superficial blood vessels, hippocampi were mechanically dissociated by fine mincing and enzymatically digested for 30 min at 37 °C in DMEM media containing 2.5 U/mL Papain (Worthington Biochemicals), 1 U/mL Dispase II (Boehringer Mannheim), and 250 U/mL Dnase I (Worthington Biochemicals). NSCs were purified using a 65% Percoll gradient and cultured in neural ba M Medium supplemented with 2% B27 containing 1% glutamine, 1% ascorbic acid, 1% glutamine, 1% thymidine, 10 ng/mL EGF, 10 ng/mL FGF) as a monolayer on poly-L-lysine and laminin-coated plates at a density of 10<sup>5</sup> cells/cm<sup>2</sup>.

Viral Plasmids and Viruses. We generated lentiviruses encoding shRNAs targeting Ogt or luciferase using a lentiviral shRNA expression system (pGreenPuro shRNA, System Biosciences) according to the manufacturer’s instructions. The targeted sequences were cloned into the pGreenPuro vector (Ogt, 5′-AGGGAACTAGATAACGCTCTT-3′; luciferase, 5′-GGCACTCTACTCCTTAGAGGGA-3′). Overexpression sequences were generated as follows. The Ogt coding sequence (CDS) and partial 3′ and 5′ UTRs were PCR-amplified form mouse hippocampal cDNA (Ogt forward primer: CACCGT TCGATTTTGTGTGGCCGCC; Ogt reverse primer: TAGGGCAATCTCTCGGTCGCC) and cloned into the pENTR D-TOPO vector (Thermo Fisher Scientific). The CDS was further amplified and cloned into a lentiviral expression plasmid containing a mouse Nestin promoter using the restriction sites NheI and BamH1. The STAT3 CDS was obtained from the ORFeme S collection and sequenced using a sequencing primer. STAT3 mutant constructs were generated using the QuikChange Lightning Site-directed Mutagenesis kit (Agilent) in combination with the following primers: TTTGACAGGTTGCTGCCCGT- CTGTGTGAGACTCTAGCCACTCATCTTTGCTTCTGCTTGCTAGGCAC -ACC (Y705F forward), CTTGCTTCAAGAAGGGCGCTTC (Y705F reverse), AGTGGGATGTTTTCTGAGGCTTAC (T717A forward), ATGTTATGGCTGAGGCTGGTTGTGCACACAG (T717A reverse). The lentiviral Nestin-promoter plasmid was linearized using the restriction enzymes NheI and SalI and the wild-type and mutant STAT3 expression constructs were

White et al.
generated using Gibson assembly (New England Biolabs). All coding plasmid sequences were verified using Sanger sequencing. Endotoxin-free plasmid maxiprep kits were used for viral plasmid preparation. Lentivirus was generated according to University of California, San Francisco viracore protocols. Briefly, Lenti-X 293T cells were transfected with the lentiviral expression plasmid along with Pax2 and VSV-G packaging and envelope plasmids, respectively. Twenty-four hours later, media was supplemented with ViralBoost (Alstem) according to manufacturer’s instructions. Twenty-four hours later, media was removed and filtered through a 0.45-μm filter. The filtered virus-containing media was then supplemented via viral centrifugation at 24,000 × g. The resultant pellet was resuspended in 100 μL PBS. Viral titer was determined using Lenti-X Go Stix Plus (Takara) according to manufacturer’s instructions.

NSC Proliferation Assay. NSCs were seeded in 500 μL growth medium (neurobasal A medium supplemented with 2% B27 without vitamin A, 1% gluta- tamine, 1% penicillin streptomycin, 10 ng/mL EGF, 10 ng/mL BFGF) at a density of 10,000 cells per well on PDL/amin-coated glass coverslips in a 24-well tissue culture plate. Twenty-four hours later, cells were treated with either DMSO or 12.5 μM OSMI-1 (Sigma-Aldrich). Sixteen hours later, cells were treated with 20 μM EdU for 8 h prior to fixing with 4% PFA for 20 min. Acute knockdown of Ogt was induced by treating plated NSCs with shRNA encoding lentiviruses (10 viral particles per cell) for 3 d prior to EdU treatment as described. Cells were washed three times with PBS after fixation, blocked in 3% donkey serum, and incubated in primary antibody at 4 °C for 16 h. After three washes with PBS, EdU incorporation was revealed using a Click-iT EdU Alexa Fluor Imaging Kit (Thermo Fisher Scientific) and nuclei were counterstained with Hoechst 33342 (1:10,000; Thermo Fisher Scientific). EdU+ cells were counted per field-of-view at three randomly determined locations per coverslip. In vitro experiments were conducted in triplicate for each condition and repeated to ensure reproducibility.

NSC Differentiation Assay. NSCs were seeded in 500 μL growth medium and treated with OMSI-1 or infected with lentivirus at a ratio of 10 viral particles per cell. Twenty-four hours (OSMI-1) or 72 h (knockdown and overexpression) after treatment, media was replaced with differentiation medium (neurobasal A medium supplemented with 2% B27 without vitamin A, 1% glutamate, 1% penicillin streptomycin, 10 ng/mL EGF, 10 ng/mL BFGF) and maintained in differentiation medium for 2 d, with the half the media replaced every other day. Cells were fixed with 4% PFA for 20 min. Cells were washed three times with PBS after fixation, blocked in 3% donkey serum, and incubated in primary antibody at 4 °C for 16 h. Staining was revealed using fluorescence-conjugated secondary Alexa antibodies (1:500). MAP2- or GFAP+ cells were counted per field-of-view at three randomly determined locations per coverslip. In vitro experiments were conducted in triplicate for each condition and repeated to ensure reproducibility.

Lactate Dehydrogenase Assay. Primary NSCs were seeded at 3,000 cells per well in a 96-well plate and were treated with OSMI-1 or a vehicle control and differentiated as previously described in NSC Differentiation Assay. At 0, 1, 3, 5, and 7 d after removal of growth factors, cell viability was assayed using Trypan blue. All cells and media were removed from the plate and centrifuged at 400 × g for 5 min. The pellet was resuspended in fresh media and diluted with an equal volume of Trypan blue. Total cells, live cells (Trypan blue excluding), and dead cells (Trypan blue including) were counted using a hemocytometer. Cell viability was determined by dividing the number of live cells by the total number of cells. Counts were performed on six replicates to ensure reproducibility.

Trypan Blue Assay. Primary NSCs were treated with OSMI-1 or a vehicle control and differentiated as previously described in NSC Differentiation Assay. At 1, 3, 5, and 7 d after removal of growth factors, cell viability was assayed using Trypan blue. All cells and media were removed from the plate and centrifuged at 400 × g for 5 min. The pellet was resuspended in fresh media and diluted with an equal volume of Trypan blue. Total cells, live cells (Trypan blue excluding), and dead cells (Trypan blue including) were counted using a hemocytometer. Cell viability was determined by dividing the number of live cells by the total number of cells. Counts were performed on six replicates to ensure reproducibility.

Western Blot Analysis. Primary NSCs were lysed in RIPA lysis buffer (500 mM Tris, pH 7.4, 150 mM NaCl, 0.5% Na deoxycholate, 1% Nonidet P-40, 0.1% SDS, complete protease inhibitors; Roche, Halt Phosphatase Inhibitor Mixture; Thermo Fisher, and PUGNAC; Sigma-Aldrich). Protein lysates were mixed with 4x NuPage LDS loading buffer (Invitrogen) and loaded on a 4 to 12% SDS polyacrylamide gradient gel (Invitrogen) and subsequently transferred onto a nitrocellulose membrane. The blots were blocked in 5% milk in TBST and incubated with primary antibody (mouse anti-β-Actin [1:500; Abcam, ab2739], rabbit anti-O-glcNAc [1:500; Novus, NB1-32791], rabbit anti-Histone H3 [1:1,000; Thermo Fisher Scientific, PAS-16183], rabbit anti-STAT3 [1:1,000; Cell Signaling 12640], rabbit anti-pSTAT3 [Y705; 1:1,000] [Cell Signaling, 9131]) at 4 °C for 1 h. Horseradish peroxidase-conjugated secondary antibodies and an ECL kit (GE Healthcare/Bio-Rad) were used to detect protein signals. Blots were imaged with a ChemiDoc (Bio-Rad) and quantified using ImageJ software (v1.80_9.1). Actin or HH3 bands were used for normalization.

PCR Genotyping. Ogt floxed, Nestin-CreERT2, and TdTomato alleles were genotyped from skin biopsies using PCR. Primers specific for the beta-globin gene were included in the reaction as a control. Ogt floxed forward primer: CACCTCTCCAGCCCGCACAAAACTG. Ogt floxed reverse primer: GAGAGGCGGACGAGAAGGAC. Primers Cre forward: CACCCCTGCTATAGCC. Primers Cre reverse: GAGATGCTCTTACAGGGCGTA. TdTomato mutant forward primer: CTGCCCTGCTATAGCCGATG. TdTomato mutant reverse primer: GGCAATGAAACCGGTATC. TdTomato wild-type forward primer: AAGAGGCTGTCAGCGGAG. TdTomato wild-type reverse primer: CGGAAATACTTGGGGAAGCTG. β-Globin forward primer: CACATGTCGTCACACAGAGATAGGAGGACG. β-Globin reverse primer: CCTTGAGGCTGCCTAGGTTATTGGCAGATG.

Tamoxifen Injections. Animals were injected via intraperitoneal injection with 5mg/kg of tamoxifen or vehicle once every 24 h for a total of five injections per animal. Animals were monitored after recovery and 4 wk allowed to pass after the final injection before any analyses were performed.

BrdU Administration and Quantification. For short-term proliferation studies, mice were intraperitoneally injected with EdU (50-mg/kg body weight, Sigma-Aldrich) daily for 4 d prior to being killed. Ten minutes after the final injection, EdU+ cells were counted per field-of-view at three randomly determined locations per coverslip. In vitro experiments were conducted in triplicate for each condition and repeated to ensure reproducibility.

Contextual Fear Conditioning. Paradigm follows previously published techniques (10). Mice learned to associate the environmental context (fear-conditioning chamber) with an aversive stimulus (mild foot shock; unconditioned stimulus, US) enabling testing for hippocampal-dependent contextual fear conditioning. The mild foot shock was paired with a light and tone cue (conditioned stimulus, CS) in order to also assess amygdala-dependent cued fear conditioning. Conditioned fear was displayed as freezing behavior. Specific training parameters are as follows: Tone duration is 30 s; level is 70 dB, 2 kHz; shock duration is 2 s; intensity is 0.6 mA. On day 1 each mouse was placed in a fear-conditioning chamber and allowed to explore for 2 min before delivery of a 30-s tone (70 dB) ending with a 2-s foot shock (0.6 mA). Two minutes later, a second CS–US pair was delivered. On day 2 each mouse was first place in the fear-conditioning chamber and allowed to explore for 2 min before a 30-s tone (70 dB) ending with a 2-s foot shock (0.6 mA). Mice were placed in a new context containing a different odor, cleaning solution, floor texture, chamber walls, and shape. Animals were allowed to explore for 2 min before being reexposed to the CS. Freezing was analyzed for 1 to 3 min. Freezing was measured using a FreezeScan video tracking system and software (Clever Sys). All behavior is performed double-blinded.

Novel Object Recognition. The NOR task was adapted from a previously described protocol (41). During the habituation phase (day 1), mice could freely explore an empty arena for 10 min. During the training phase (day 2), two identical objects were placed in the habituated arena, and mice could explore the objects for 5 min. For the testing phase (day 3), one object was replaced with a novel object, and mice could explore the objects for 5 min. Time spent exploring each object was quantified using the Smart Video Tracking Software (Panlab; Harvard Apparatus). Two different sets of objects were used. To control for any inherent object preference, half of the mice were exposed to object A as their novel object and half to object B. To control for any potential object-independent location preference, the location of the novel object relative to the trained object was also varied. The objects were chosen based on their ability to capture the animal’s interest, independent of genetic background or age. To determine percent time with novel object, we calculated (time with novel object)/(time with trained object + time with novel object) × 100. In this preference index, 100% indicates full preference for the novel object, and 0% indicates full preference for the trained object. A mouse with a value of 50% would have spent equal time...
Precursor ions were measured from 375 to 1500 m/z. Fusion Lumos (Thermo Scientific) equipped with a NanoAcquity UPLC (Waters) was used to analyze glycopeptides on an Orbitrap Exactive Plus (Thermo Scientific), according to the manufacturer's instructions. Frozen protein pellets were resuspended in 50 mM ammonium bicarbonate containing 6M guanidine hydrochloride, 6X Phosphatase Inhibitor Mixtures II and III (Sigma-Aldrich), and 80 mM PUGNAc (Tocris Bioscience). Protein concentrations were estimated with bicinchoninic acid (BCA) protein assay (Thermofisher Scientific). The protein lysate (1 mg from each sample) was reduced for 1 h at 56 °C with 2.5 mM Tris(2-carboxyethyl)phosphine hydrochloride and subsequently carbamidomethylated using 5 mM iodoacetamide for 45 min at room temperature in the dark. Lysates were diluted to 1M guanidine hydrochloride with 50 mM ammonium bicarbonate, pH 8.0, and equal amounts of each sample were digested overnight at 37 °C with sequencing grade trypsin (Thermofisher Scientific) at an enzyme to substrate ratio of 1:50 (w/wt). Following digestion, samples were acidified with formic acid (FA) (Sigma-Aldrich), desalted using a 360-μg C18 Sep-Pak SPE cartridge (Waters), and dried to completeness using a SpeedVac concentrator before glycopeptide enrichment.

TMT Mass Spectrometry Data Analysis. Data were filtered to only include O-GlcNAcylated peptides unique to a single protein. Quantitation of TMT data was performed by calculating ion peak intensities of reporter ions, which correspond to a 5% local false localization rate. Modified peptides were identified with a protein and peptide false-discovery rate of 1%. O-GlcNAc and O-GalNAc modifications were differentiated based on known protein subcellular localization and HexNAc oxonium ion fragment ratios (44).

RNA Isolation. mRNA of NSCs was isolated by lysis with TRIzol Reagent (Thermo Fisher Scientific), separation with chloroform (0.2 mL per mL TRIzol), and precipitated with isopropyl alcohol according to the manufacturer's instructions.

NSC Library Construction and RNA-Sequencing. After RNA isolation, RNA-sequencing libraries were constructed using the Smart-Seq2 protocol from (46), with modifications. Briefly, 1 ng high-quality RNA was reverse transcribed using SuperScript II (Life Technologies, 18064-014) with a poly-dT anchored oligonucleotide primer, and a template switching oligonucleotide primer that generated homotypic PCR primer binding sites. The cDNA underwent 10 rounds of PCR amplification using KAPA HiFi Hotstart (Kapa Biosystems, KK2601), followed by Ampure bead (Agenencourt) clean-up. The quality of the amplified CDNA was tested using qPCR for GAPDH and nucleic acid quantitation. One nanogram of high-quality amplified cDNA was fragmented with the Trn5 transposase from the Illumina Nextera kit (FC-131-1096) to a median size of ~500 bp. The fragmented library was amplified with indexed Nextera adapters (FC-131-1002) using 12 rounds of PCR. Final libraries were purified with Ampure beads and quantified using a qPCR Library Quantification Kit (Kapa Biosystems, KK4824). Libraries were pooled for sequencing on an Illumina HiSeq 2500 (paired reads 2 × 100 bp).

Bioinformatic Analysis of NSC RNA-Sequencing Data. Alignment of RNA-sequencing reads to the mouse mm10 transcriptome was performed using STAR v2.7.3a (47) following ENCODE standard options, read counts were generated using RSEM v1.3.1, and differential expression analysis was performed in R v3.6.1 using the DESeq2 package v1.38.0 (48) (detailed pipeline v2.0.1 and options available on https://github.com/ermc2cubebioinformatics). Significance was determined using a corrected P < 0.05 and a fold-change > 0.5. GO and ChEA/ENCODE transcription factor enrichment analysis was performed with Enrichr (https://amp.pharm.mssm.edu/Enrichr), Heatmaps were generated using IDeP (http://bioinformatics.dataedd.org). All data available under Gene Expression Omnibus accession no. GSE143388.

Data and Statistical Analysis. Graphed data are expressed as mean ± SEM. Statistical analysis was performed with Prism 6.0 software (GraphPad Software).
Data Availability. RNA-seq data were deposited at GEO (accession no. GSE143388). Mass Spectrometry raw data were deposited at MassIVE (MSV000084825). Mass spectrometry spectra can be viewed at MS-Viewer (qchzi0to8). (49–51).

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