Regulation of the p19Arf/p53 pathway by histone acetylation underlies neural stem cell behavior in senescence-prone SAMP8 mice

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Introduction

The ‘stem cell aging’ hypothesis suggests that loss of potential in somatic stem cell pools underlies the progressive decline in tissue renewal observed in multiple systems during aging (van Deursen, 2014). Lifelong generation of olfactory bulb (OB) interneurons takes place in the subependymal zone (SEZ; also known as ventricular/subventricular zone or VS/VZ) of the lateral ventricle wall and is supported by a population of neural stem cells (NSCs, also known as B1 cells). B1 cells divide slowly and can hence retain exogenously administered traceable nucleosides for several weeks, generating transit-amplifying progenitor (TAP) cells, which in turn give rise to neuroblasts that migrate anteriorly to the OB where they differentiate into granular and periglomerular interneurons (Silva-Vargas et al., 2013).

Reduced levels of olfactory neurogenesis that are primarily due to decreased numbers of stem-like cells are found in elderly animals (Maslov et al., 2004; Luo et al., 2006; Molofsky et al., 2006; Shook et al., 2012). Many NSCs remaining in the aged SEZ are primarily quiescent; they are less likely to undergo cellular division or neuronal differentiation than NSCs in the young adult brain, with lower chances of survival (Ahlenius et al., 2009). Those cells remaining in cell cycle, however, are more proliferative than young NSCs, due to increased rates of cell cycle reentry, which can lead to transient expansion of cells before terminal differentiation (Stoll et al., 2011). NSC permanent cell cycle exit results in their differentiation into S100b+ astrocyte-like cells (Raponi et al., 2007; Porlan et al., 2013), although it has been reported that a fraction of B1 cells may also integrate within the agedependymal cell layer and acquire ependymal features (Luo et al., 2008).

Although age-related declines in stem cell function have been associated with altered expression of some growth factors (Tropepe et al., 1997; Enwere et al., 2004), tumor suppressors p16Ink4a and p19Arf (hereafter p16 and p19; encoded by the Cdkn2a locus by the use of two different promoters and alternative reading frames) also appear to play a role (Molofsky et al., 2006; Ahlenius et al., 2009; Mikheev et al., 2009). Cell cycle inhibitor p16 binds to and inhibits CDK4 and CDK6 to retain the growth-suppressive activity of Rb family proteins, whereas p19 binds to and antagonizes the activity of the ubiquitin ligase Mdm2, thereby stabilizing and activating p53 (Gil & Peters, 2006). Histone acetylation controlled by the antagonistic actions of histone acetyltransferases (HATs) and histone deacetylases (HDACs) which either favor or repress transcription (Willis-Martinez et al., 2010) has been involved in the regulation of this locus in other cells (Matheu et al., 2005; Simboeck et al., 2011).

Effects of aging in neural dysfunction have been analyzed in several strains of senescence-accelerated mice (SAM) originated from inbreeding and selection for the early appearance of aging features (Takeda, 2009). Relative to control senescence-resistant SAMR1 mice, young senescence-prone SAMP8 mice exhibit neuronal loss, gliosis, and progressive cognitive deficits (Takeda, 2009; Tomobe & Nomura, 2009). Proliferation in the SEZ and in the dentate gyrus subgranular zone (SGZ), another neurogenic niche, is increased in 2-month-old (2-m) but reduced in 10-m SAMP8 mice (Gang et al., 2011; Díaz-Moreno

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Summary

Brain aging is associated with increased neurodegeneration and reduced neurogenesis. B1/neural stem cells (B1-NSCs) of the mouse subependymal zone (SEZ) support the ongoing production of olfactory bulb interneurons, but their neurogenic potential is progressively reduced as mice age. Although age-related changes in B1-NSCs may result from increased expression of tumor suppressor proteins, accumulation of DNA damage, metabolic alterations, and microenvironmental or systemic changes, the ultimate causes remain unclear. Senescence-accelerated-prone mice (SAMP8) relative to senescence-accelerated-resistant mice (SAMR1) exhibit signs of hastened senescence and can be used as a model for the study of aging. We have found that the B1-NSC compartment is transiently expanded in young SAMP8 relative to SAMR1 mice, resulting in disturbed cytoarchitecture of the SEZ, B1-NSC hyperproliferation, and higher yields of primary neurospheres. These unusual features are, however, accompanied by premature loss of B1-NSCs. Moreover, SAMP8 neurospheres lack self-renewal and enter p53-dependent senescence after only two passages. Interestingly, in vitro senescence of SAMP8 cells could be prevented by inhibition of histone acetyltransferases and mimicked in SAMR1 cells by inhibition of histone deacetylases (HDAC). Our data indicate that expression of the tumor suppressor p19, but not of p16, is increased in SAMP8 neurospheres, as well as in SAMR1 neurospheres upon HDAC inhibition, and suggest that the SAMP8 phenotype may, at least in part, be due to changes in chromatin status. Interestingly, acute HDAC inhibition in vivo resulted in changes in the SEZ of SAMR1 mice that resembled those found in young SAMP8 mice.

Key words: aging; adult neurogenesis; histone deacetylases; histone acetyltransferases; stem cell niche; SAMP8 mice.

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et al., 2013). Moreover, although more neuroblasts are produced in young SAMP8 mice, they are lost at a higher rate due to apoptosis (Gang et al., 2011). These results have been interpreted as a reaction to the neuronal dysfunction/degeneration observed in these animals, but they also raise the possibility that the senescent-prone genetic background has a direct impact on intrinsic properties of NSCs. We have found that increased loss of NSCs in 10-m SAMP8 mice is preceded by their transient overactivation. Despite the higher proliferative activity of young SAMP8 NSCs, they expressed increased levels of p19 and p53 and entered senescence prematurely when cultured in vitro, a deficit that could be restored by abrogation of p53 activity and, interestingly, by inhibition of the activity of HATs. Moreover, we could induce a senescent phenotype, together with increases in p19 and p53, by in vitro treatment with inhibitors of HDACs, whereas in vivo treatment resulted in morphological changes of the SEZ resembling those of young SAMP8 mice.

**Results**

**Increased proliferation and abnormal positioning of B1 NSCs in SAMP8 mice precedes their exhaustion**

A previous analysis had indicated a transient increase in Brdu incorporation in the SEZ of young SAMP8 mice (Díaz-Moreno et al., 2013). To focus specifically on B1 NSCs, we analyzed whole-mount preparations of the lateral ventricle wall by confocal microscopy. The ventricle wall surface is characterized by the repetition of ‘pinwheel’ units, arrangements in which the uniciliated apical thin processes of one or several GFAP+ B1 NSCs are encircled by multiciliated ependymal cells (Mirzadeh et al., 2008). Immunostaining for GFAP, β-catenin to delineate cell membranes, and either γ-tubulin as a marker of cilial basal bodies or acetylated α-tubulin to label cilial cytoskeleton, revealed a normal pinwheel organization in SAMR1 mice (Fig. 1A,B). Compared to other mouse strains, however, 2-m SAMR1 mice had fewer B1 apical surfaces (543 ± 34 per mm² of dorsal SEZ, n = 3, vs. 1,732 ± 53 in C57BL/6 mice, n = 3) and pinwheels (289 ± 30 per mm², n = 4, vs. 952 ± 45 in C57BL/6 mice, n = 3) (for CD1 mice, see also Mirzadeh et al., 2008; Shook et al., 2012), suggesting potential effects of the SAM genetic background in B1-cell population size.

Although all pinwheel components could also be recognized in 2-m SAMP8 mice, the organization was altered (Fig. 1A,B). The number of pinwheels with thin B1-cell processes at their core was remarkably low (59 ± 7 per mm², n = 3). In contrast, we could observe misshapen pinwheels formed by ependymocytes surrounding various GFAP+ cells with large apical areas in the ependymal layer plane, a feature that was also evident in analyses with scanning electron microscopy (Fig. 1A–C). Because cells in these groups were unciliated and GFAP+ but did not express the calcium-binding protein S100β, a marker of ependymal cells and terminally differentiated astrocytes (Raponi et al., 2007) (Fig. 1D), we named them ‘B1-like interposed cells’ (BICs). BICs were very frequent in 2-m SAMP8 mice and neurogenesis, R. Soriano-Cantón et al.

We also detected high numbers of cuboidal GFAP+ cells with multiple cilia integrated within the ependymal layer (Fig. 1G). This type of cell reportedly increases during aging but is rarely found in young C57BL/6 and CD1 mice (Luo et al., 2008; Capilla-Gonzalez et al., 2014). Indeed, we could find only 7 ± 3 cells per mm² (n = 3) in the dorsal SEZ of 2-m C57BL/6 mice, but 49 ± 21 cells per mm² at 12-m and 52 ± 6 at 24-m (n = 2 each). Young SAMR1 and SAMP8 mice had similarly elevated proportions of these cells (Fig. 1G), suggesting that this feature could have been selected through inbreeding.

Notably, the appearance of the SEZ of 10-m SAMP8 mice was similar to that found in SAMR1 controls at 10-m and in normal mouse strains at 12-m. Quantification of pinwheel density revealed a significant reduction in SAMR1 mice between 2 and 10 months, a period at which most age-related dysfunctions develop in SAM strains (Tomobe & Nomura, 2009), similar to reductions observed in C57BL/6 mice. In 10-m SAMP8 mice, however, pinwheels were significantly less frequent (Fig. 1H,I), indicating that BICs were a transient feature and that loss of NSCs is accelerated in SAMP8 mice.

In SEZ sections, we also observed an overall increase in GFAP, both in the number and intensity of positive cells, in 2-m SAMP8 mice (Fig. 2A,B). Moreover, we detected more GFAP+Sox2+ cells (relative to DAPI: 16.4 ± 1.1, n = 4, vs. 10.2 ± 0.3%, n = 3, in SAMR1 mice; P < 0.01) and more GFAP+ cells were also Sox2+ and nestin+ (23.1 ± 1.3 vs. 16.1 ± 1.2% in SAMR1 mice; n = 3, P < 0.05), but fewer were S100β+ (3.3 ± 0.9% vs. 15.9 ± 2.8 in SAMR1 controls; n = 4, P < 0.01; Fig. 2B). These observations indicated the presence of higher numbers of B1 cells in young SAMP8 mice. Similar to observations in whole-mounts, some GFAP+ cells appeared to be integrated within the ependymal layer (Fig. 2C). Indeed, the net increase in GFAP+ cells found in 2-m SAMP8 mice was due to a selective increase in the number of GFAP+S100β-BICs, but not subependymal cells (Fig. 2D). At 10-m, we found similar numbers of GFAP+ cells in both genotypes, but significantly more of them were S100β+ in SAMP8 mice (Fig. 2B), again suggesting that BICs eventually differentiate into S100β+ astrocytes and that NSC senescent features develop faster in SAMP8 mice.

**Increased NSC proliferation in young SAMP8 mice is counteracted by cell death of migrating neuroblasts**

We next investigated whether more B1 cells in 2-m SAMP8 mice resulted in increased neurogenesis. We did not detect changes in the overall proliferation in the SEZ of SAMP8 mice injected with a single pulse of 5-bromo-2′-deoxyuridine (BrdU) one hour before sacrifice (relative to total cells: 8.6 ± 0.6 BrdU+ cells vs. 9.5 ± 0.5% in SAMR1 mice; n = 4). However, the proportions of GFAP+BrdU+ cells (0.38 ± 0.06% vs. a SAMR1 value of 0.22 ± 0.06%; n = 4, P < 0.05) and of GFAP+ Sox2+Ki67+ cells (1.5 ± 0.1 vs. 1.0 ± 0.2%; n = 4, P < 0.01) were significantly increased (Fig. 2E,F). The increase in Ki67+-proliferative GFAP+S100β+ cells in SAMP8 mice was also specifically due to an increase in Ki67+ cells among BICs (relative to total cells: 0.44 ± 0.09 in SAMR1 mice vs. 0.14 ± 0.04% in SAMR1; n = 4, P < 0.05; Fig. 2G,H), suggesting a correlation between abnormal positioning and proliferation.

Activated B1 cells can incorporate and retain BrdU for several weeks, and we could detect more GFAP+ BrdU label retaining cells (BrdU-LRC) in SAMP8 than in SAMR1 mice which had been repeatedly injected with the nucleoside (7 doses in a 12-h period) one month before euthanasia (relative to DAPI: 0.30 ± 0.05 vs. 0.16 ± 0.01%, respectively; n = 3, P < 0.05). Again, GFAP+S100β+ LRCs in contact with the ventricle lumen were more frequent in SAMP8 mice (0.12 ± 0.01 vs. a SAMR1 value of 0.01 ± 0.01; n = 3, P < 0.01; Fig. 2I,J).

Activated B1 cells generate proliferating neuroblasts that can be detected with antibodies to doublecortin (DCX) in the rostral migratory stream (RMS). These neuroblasts differentiate in the OB into periglomerular and granular interneurons that can be identified as they permanently retain BrdU incorporated during the S-phase preceding their cell cycle exit. Despite a higher number of proliferative GFAP+ cells in the SEZ of SAMP8 mice, DCX+ neuroblasts in the RMS (relative to DAPI: 0.30 ± 0.05 vs. 0.21 ± 0.04% in SAMR1 mice, n = 4, P < 0.05; Fig. 2A,B) were significantly more frequent in 2-m SAMP8 mice compared to the SAMR1 controls; n = 4, P < 0.05; Fig. 2C). Moreover, we detected more GFAP+Sox2+ cells (relative to DAPI: 16.4 ± 1.1, n = 4, vs. 10.2 ± 0.3%, n = 3, in SAMR1 mice; P < 0.01) and more GFAP+ cells were also Sox2+ and nestin+ (23.1 ± 1.3 vs. 16.1 ± 1.2% in SAMR1 mice; n = 3, P < 0.05), but fewer were S100β+ (3.3 ± 0.9% vs. 15.9 ± 2.8 in SAMR1 controls; n = 4, P < 0.01; Fig. 2B). These observations indicated the presence of higher numbers of B1 cells in young SAMP8 mice. Similar to observations in whole-mounts, some GFAP+ cells appeared to be integrated within the ependymal layer (Fig. 2C). Indeed, the net increase in GFAP+ cells found in 2-m SAMP8 mice was due to a selective increase in the number of GFAP+S100β-BICs, but not subependymal cells (Fig. 2D). At 10-m, we found similar numbers of GFAP+ cells in both genotypes, but significantly more of them were S100β+ in SAMP8 mice (Fig. 2B), again suggesting that BICs eventually differentiate into S100β+ astrocytes and that NSC senescent features develop faster in SAMP8 mice.

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DAPI-stained nuclei: 16.8 ± 2.6 vs. a SAMR1 value of 18.2 ± 3.0%, n = 3) or BrdU+ newly generated neurons in the OB glomerular layer (relative to DAPI-stained nuclei: 0.28 ± 0.01 vs. a SAMR1 value of 0.27 ± 0.10%, n = 3) were not increased.

To discern among the possible causes that could account for the lack of net effects in neurogenesis, alterations in migration, cell death, and/or proliferation of neuroblasts were investigated. Similar proportions of DCX+ cells were found at different anteroposterior levels of the RMS in SAMR1 and SAMP8 mice (data not shown), indicating that neuroblast migration is not affected. In addition, the proportion of DCX+ cells that incorporated BrdU injected as a one 1-h pulse was similar in SAMP8 and SAMR1 mice (20.6 ± 4.9 vs. 17.3 ± 2.7% respectively, n = 4). The number of TUNEL+DCX+ neuroblasts in the RMS (3110 ± 626 vs. a SAMR1 value of 830 ± 496 cells/mm³, n = 4; P < 0.05) but not of TUNEL+GFAP+ cells in the SEZ (803 ± 464 vs. 839 ± 499 cells/mm³ in SAMR1 mice, n = 4) was increased in SAMP8 mice, indicating that limited survival of newborn cells counterbalances the increased cycling activity of B1 cells in young SAMP8 mice.

Increased cycling activity of young SAMP8 NSCs results in their premature senescence in vitro

In agreement with the presence of more proliferative GFAP+ cells, the SEZ of young SAMP8 mice yielded more primary neurospheres when
individual cells were plated with EGF and FGF (Fig. 3A–C). Compared to SAMR1, SAMP8 primary neurospheres were also significantly larger in diameter (Fig. 3D), had reduced levels of TAP-associated transcription factor Mash1/Ascl mRNA (0.46 ± 0.01 relative to SAMR1 levels, n = 3; P < 0.001), and produced more secondary neurospheres when their cells were dissociated and reseeded (passage P1) at very low density (Fig. 3E, F). These data indicated that the SEZ of young SAMP8 harbors more NSCs capable of forming neurospheres at young ages and further suggested that BICs are B1 cells modified by the SAM genetic background.

SAMP8 secondary neurospheres were, however, smaller in size after 5 days (Fig. 3E,G), incorporated less BrdU (relative to DAPI: 14.7 ± 1.8 vs. 22.2 ± 1.5% in SAMR1 cultures; n = 7, P < 0.01), and could not be subcultured for another passage. They did not appear to undergo apoptosis (percent caspase 3+ cells/DAPI: 1.51 ± 0.14 in SAMP8, n = 4, and 1.56 ± 0.24 in SAMR1 cultures, n = 3), but adopted a senescent phenotype, characterized by cell enlargement, adhesion to the plastic culture plate, cessation of proliferation, and positivity in the histochemical reaction of the senescence-associated β-galactosidase (SAβ-gal) (Fig. 3H,J). Some P2 neurospheres could still be observed, but were noticeably smaller and SAβ-gal+, and had increased levels of S100β mRNA (Fig. 3H–K). The strikingly premature senescence of SAMP8 cells was never seen in C57BL/6 or CD1 cells subcultured for many months (Ferron et al., 2007). Our data indicated that, although SAMP8 SEZs contain elevated numbers of neurosphere-forming cells, their self-renewal is remarkably reduced.

![Fig. 2 Increased number of proliferating NSCs in the SEZ of young P8 mice. (A) GFAP and S100β detection in the SEZs of 2- and 10-m R1 and P8 mice. (B) Percentage of GFAP+/DAPI cells (left) and of GFAP+S100β+/GFAP+ cells (right) in the SEZs of 2- and 10-m R1 and P8 mice. (C) GFAP and S100β in the SEZ of 2-m R1 and P8 mice. (D) Percentage of GFAP+S100β+ BICs and subependymal (SE) cells relative to total cells. (E) GFAP and BrdU in the SEZ of 2-m R1 and P8 mice. (F) GFAP, Sox2, and Ki67 in the SEZ of 2-m R1 and P8 mice. (G) GFAP, S100β, and Ki67 in the SEZ of 2-m R1 and P8 mice. (H) Percentage of Ki67+GFAP+S100β− BICs and subependymal (SE) cells relative to total cells. (I) GFAP, S100β, and LRCs in the SEZ of 2-m R1 and P8 mice. (J) Percentage of GFAP+S100β−BrdU+ label retaining BICs. Data are shown as mean ± SEM of 3 independent mice from each strain (*P < 0.05; **P < 0.01). Dashed white line indicates the lateral ventricle (v) limit. White arrowheads point at single-positive cells and white arrows at doubly positive cells. Asterisks indicate triple-positive cells. Scale bars: 10 μm.](image-url)
Impaired self-renewal of SAMP8 neurosphere-forming cells associates with increased DNA damage and is mediated by p53 and p19

Cellular senescence can be triggered by a number of negative stimuli (telomere erosion, oxidative stress, replicative stress) which activate the DNA damage response (DDR), a signaling pathway in which ATM or ATR kinases block cell cycle progression through stabilization of p53. Activation of certain oncogenes results in derepression of the Ink4a/Arf locus in the absence of DNA damage and can also contribute, through increases in p19, in the stabilization of p53 (Gil & Peters, 2006; van Deursen, 2014). In adult neurosphere cultures, loss of proliferation control, that is, as a result of p21 deletion and/or Sox2 overexpression, leads to DNA damage and senescence mediated by increases in p53 and p19, but not p16 (Marqués-Torrejón et al., 2013). Likewise, we detected increased levels of p19 and p53, but reduced levels of p16 in SAMP8 relative to SAMR1 cultures by immunoblot (Fig. 4A, B), a molecular profile consistent with a DDR.

In SAMP8 neurosphere cultures, we could indeed detect increased proportions of cells with DNA foci immunopositive for the form of histone H2AX phosphorylated in Ser 139 (γ-H2AX), a widespread marker of DNA damage (Fig. 4C). Consistent with this, the levels of p53 phosphorylated in Ser18 (pp53) by ATM (Chao et al., 2006) were also higher (Fig. 4D). Treatment of the cultures with the ATM inhibitor KU55933 or with pifithrin-α (PFTα), a water-soluble inhibitor of p53 activity (Komarov et al., 1999), prevented the senescent phenotype (Fig. 4E). Senescence was also partly prevented by retroviral delivery of a specific shRNA (Marqués-Torrejón et al., 2013) which lowered p53 to 0.68 ± 0.01 (n = 3) of the levels in control cultures (Fig. 4F, G). These data indicated potential activation of the p53 checkpoint in response to DNA damage in SAMP8 cells.

Changes in chromatin acetylation regulate neurosphere proliferation and senescence state through the Arf-p53 axis

Treatment of normal MEFs with trichostatin A (TSA), a natural compound that interferes with the activity of HDACs classes I and II, but not class III (Willis-Martinez et al., 2010), leads to increases in p19, together with reductions in p16, and induces senescence (Matheu et al., 2005; Simboeck et al., 2011). Treatment with HDAC inhibitors also
increases acetylation of other substrates, including p53 which then becomes protected from degradation and increases its activity (Luo et al., 2000). A 3-day treatment of SAMR1 cells with TSA at 25 or 50 nM raised the levels of acetylated p53 on Lys 379 (ac-p53) and induced the accumulation of p19 in a concentration-dependent manner (Fig. 5A). The same results were found with 2 or 4 mM valproic acid (VPA), which also inhibits HDACs by binding to their catalytic center (Göttlicher et al., 2001) (Fig. 5A). Inhibition of the HDAC activity in SAMR1 primary cultures with either TSA or VPA resulted in reduced BrdU incorporation and drove the cells into a SAβ-gal + cultures with either TSA or VPA resulted in reduced BrdU incorporation.

The same results were found with 2 or 4 mM valproic acid (VPA), which also inhibits HDACs by binding to their catalytic center (Göttlicher et al., 2001) (Fig. 5A). Inhibition of the HDAC activity in SAMR1 primary cultures with either TSA or VPA resulted in reduced BrdU incorporation and drove the cells into a SAβ-gal + cultures with either TSA or VPA resulted in reduced BrdU incorporation.

Therefore, decided to explore the effect of HDAC inhibition in NSCs derived from normal mouse strains at passages when DNA damage is not observed (Porlan et al., 2013). Cells in primary cultures of adult C57BL/6 mice also entered senescence upon TSA treatment (not shown) as also reported for cultures derived from postnatal day 8 SEZs (Foti et al., 2013). At passages 4–7, the same treatment did not cause attachment of the cells to the dish, but resulted in reduced formation of neurospheres together with an increment in SAβ-gal + cells (Fig. 5E). TSA treatment also yielded neurospheres with elevated levels of S100β mRNA in which more S100β + cells could be found (Fig. 5E). TSA effects were not observed in p53- or p19-deficient cultures treated in the same way (Fig. 5E), indicating that upregulation of p19 and/or p53 can result in senescence in the absence of DNA damage.

Our results strongly suggested that HDAC activity is required for the maintenance of self-renewal. As HDACs counteract the actions of HATs, we treated SAMP8 neurosphere cultures with 50 μM of the specific inhibitor anacardic acid (AA). The treatment rescued the senescent phenotype of DNA-damaged SAMP8 P2 neurosphere cells (Fig. 5F) without affecting SAMR1 cells (not shown), suggesting that the senescent state observed in SAMP8 cultures could be dependent on the epigenetic deregulation of the Ink4a/Arf locus.

**HDAC inhibition in vivo reproduces the phenotype of young SAMP8 mice**

We next wondered whether epigenetic alterations due to deficient HDAC activity could also result in alterations of the SEZ similar to those observed in young SAMP8 mice. To investigate this possibility, we injected young adult SAMR1 mice with either TSA at 1 mg/kg of body weight or with the DMSO vehicle twice every day during 3 days and analyzed the lateral ventricle wall. The treatment resulted in increased levels of histone 3 acetylated in Lys 9 (ac-H3), as determined by immunohistochemistry and immunoblot (1.5 ± 0.1-fold increase with respect to vehicle injected mice; Fig. 6A, B).

Despite unchanged numbers of pinwheels, a raise in the occurrence of uniliated GFAP + cells integrated within the ependyma, similar to that previously observed in SAMP8 mice, was induced in SAMR1 mice treated with TSA (Fig. 6C, D). Moreover, counts in SEZ sections indicated that the treatment resulted in a higher proportion of GFAP + S100β + cells integrated in the ependymal layer (Fig. 6E). Together with the restoration of the phenotype of SAMP8 neurospheres in vitro by HAT inhibition, this observation suggests that epigenetic changes may, at least partly, underlie the changes observed in the SEZ of SAMP8 mice.

**Discussion**

Aging is a negative regulator of adult neurogenesis, and in turn, decreased neurogenesis is considered a contributor to age-associated cognitive and olfactory declines. Yet, the mechanisms underlying impaired neurogenesis in the aging brain are poorly understood. We have analyzed B1-NSC behavior in SAMP8 mice with the aim of identifying molecular mechanisms that may impinge in their aging process. Our data support a model in which epigenetic derepression of tumor suppressor p19 drives accelerated senescence and loss of subependymal NSCs with a SAMP8 genetic background. Interestingly, these NSCs exhibit a transient increase in proliferative activity before exhaustion in vivo that can be reproduced by HDAC inhibition.

We find that the SEZ of young SAMP8 and SAMR1 mice contains fewer B1-NSCs than other inbred strains, suggesting a significant effect of the SAM genetic background in B1-cell generation and/or survival.
Increased numbers of multiciliated GFAP+ cells in SAMP8 and SAMR1 mice could indicate that B1 cells are not properly maintained in SAM strains, but are more prone to become ependymal cells, a fate that is associated with older ages in other inbred strains (Luo et al., 2008). However, other reports using BrdU retention do not support B1-cell transformation into ependymocytes (Capilla-Gonzalez et al., 2014). Exploring this possibility would require lineage tracing analysis as ependymal cells are no longer detectable by GFAP immunohistochemistry. Nevertheless, SAMP8 mice loose B1 cells more rapidly, resulting in significant differences in the number of pinwheels between SAMR1 and SAMP8 at 10-m.

Accelerated loss of B1 cells in SAMP8 mice is preceded by abnormally increased numbers of multiciliated GFAP+ cells in SAMP8 and SAMR1 mice (Zhou et al., 2011; Foti et al., 2013), further suggesting that the increase in BICs that we observe after TSA treatment and in SAMP8 mice is also likely to be nonproductive.

Expression of the Ink4a/Arf locus is silent during embryogenesis, but is gradually upregulated with aging (Zindy et al., 1997; Krishnamurthy et al., 2004). One or both proteins encoded by this locus are induced during senescence depending on triggers and cellular context, with notable species-specific differences, that is, murine cells being more dependent on p19 than human cells (Bruggeman et al., 2005; van Deursen, 2014). Protein p16 appears to limit NSC potential during physiological aging (Molofsky et al., 2006), but we find increased levels of p19 specifically in SAMP8 neurospheres. Thus, the net effect of bidirectional changes in tumor suppressor protein expression remains to be elucidated. The p19-p53 pathway not only regulates NSC self-renewal and proliferation in the postnatal and adult brains (Bruggeman et al., 2005; Nagao et al., 2008; Ahlenius et al., 2009), but also their responsiveness to gliogenic signals (Nagao et al., 2008). An increase in p19-p53 could couple senescence to a gliogenic program that might promote terminal differentiation of BICs into non-neurogenic S100β+ ependymal cells.
cells. Furthermore, we have recently shown that loss of proliferation control and/or increased replicative stress leads to higher levels of p19, but not p16 and results in cell senescence in neurosphere cultures and higher proportions of S100β+ cells in vivo (Marqués-Torrejón et al., 2013). Although p53 and p21 control at least in part the proliferation of NSCs (Medrano et al., 2009; Marqués-Torrejón et al., 2013; Porlan et al., 2013), we have failed to detect altered expression of p21 in the SEZ from SAMP8 mice (data not shown).

Our results indicate a regulation of the Ink4a/Arf locus by acetylation. HDAC inhibition reduces proliferation of adult hippocampal progenitors and promotes their neuronal differentiation through the upregulation of the neurogenic basic helix-loop-helix transcription factor NeuroD (Hsieh et al., 2004). In these progenitor cells, the orphan nuclear receptor TLX recruits HDACs to repress transcription of the cell cycle inhibitor p21 and of the tumor suppressor PTEN and deregulation of p21 has been shown to be important for the antiproliferative effect of HDAC inhibitors, although p19 was not tested (Sun et al., 2007). The relationship of PcG complexes with Ink4a/Arf expression is also a provocative possibility given the ability of these complexes to create heritable epigenetic marks. PcG gene product Bmi1 deficiency in mice is associated with premature senescence and loss of NSC self-renewal, which can in large part be rescued by repressing the expression of p16 and p19 (Bruggeman et al., 2004, 2005; Molofsky et al., 2005; Nishino et al., 2008).

The underlying genes responsible for accelerated senescence and pathological changes in SAM strains remain largely unidentified. A recent whole-exome sequencing of various SAM strains has revealed the existence of some SAMP-shared and SAMP8-specific single nucleotide variations (Tanisawa et al., 2013). Two of these mutations are found in DNA repair genes Ogg1 and Mbd4 and thus may be directly involved in the susceptibility to diseases via defects in the DDR. A third specific mutation is present in the Aifm3 gene, encoding the apoptosis-inducing factor mitochondrion-associated protein 3, which may contribute to mitochondrial dysfunction in SAMP8 mice as it induces apoptosis in vitro and has an oxidoreductase domain that might play a role in the respiratory chain (Xie et al., 2005).

Our data suggest that the NSC loss that takes place during physiological aging in mice is dramatically accelerated in a SAMP8 background and provides a system in which to analyze the mechanisms underlying exhaustion of stem cell pools in vivo. Moreover, the involvement of p19 in neurogenic senescence suggests the possibility that neurogenic dysfunction in the elderly could be prevented by pharmacological treatment.

**Experimental procedures**

**Animals**

Adult male mice of SAMP8 and SAMR1 (Jackson Laboratories, (Charles River, Barcelona, Spain)), p53 (Marqués-Torrejón et al., 2013) and p19 knockout (Kamijo et al., 1997), C57BL/6, and CD1 strains were handled in accordance with Spanish RD1201/2005 guidelines, following protocols approved by the institutional committees. BrdU (50 mg/kg body weight in saline solution; Sigma) was intraperitoneally injected, either seven times in a 12-h period (one injection every 2 h) 30 days before perfusion or as a single dose 1 h before perfusion. TSA (1 mg/kg body weight; Sigma) or DMSO was administered by intraperitoneal injection, at one dose every 12 h during 3 days, and the SEZs were dissected 1 h after the last injection.

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Histological analyses

Mice were anesthetized (225 mg/kg of ketamine and 3 mg/kg of medetomidine) and transcardially perfused with 4% paraformaldehyde in 0.1 M phosphate buffer pH 7.4 (PB). The brains were dissected, postfixed by immersion in the same fixative overnight (o/n), and serially sectioned into 25-μm coronal sections using a Leica VT 1000S vibratome. For whole-mounts, the SEZs were processed as described (Mirzadeh et al., 2008). Sections and whole-mounts were blocked using 10% FBS and 0.2% Triton X-100 in PB. For BrdU immunodetection, sections were first treated with 2N HCl for 15 min at 37 °C. Samples were incubated in rabbit anti-Ki67 (1:150, Abcam, Cambridge, England), S100β (1:150, Dako), β-catenin (1:100, Cell Signaling), or ac-H3 (1:10 000, Sigma); rat anti-BrdU (1:600, Abcam); chicken anti-GFAP (1:400, Millipore, Madrid, Spain); goat anti-Sox2 (1:50, R&D), DCX (1:100, Santa Cruz); mouse anti-γ-tubulin (1:600, Santa Cruz, Heidelberg, Germany); or acetylated α-tubulin (1:500, Sigma) antibodies in blocking buffer, o/n at 4 °C. After several PB washes, the sections were incubated for 1 h at room temperature (RT) with DyLight 549, DyLight 488, or Cy3-conjugated appropriate secondary antibodies (1:600 to 1:1500; Jackson ImmunoResearch Laboratories, Suffolk, England). Samples were stained with DAPI (Sigma) at 1 μg/mL and mounted with Fluorsave (Calbiochem, Madrid, Spain). Confocal planes along the rostro-caudal axis or z images of the entire dorsal aspect of SEZ whole-mounts were captured using a Fluoview FV10i (Olympus) confocal microscope and analyzed. Apoptotic cells were detected by terminal deoxynucleotidyl transferase-mediated dUTP nick-end labeling (TUNEL) using ApopTag In Situ Apoptosis Detection Kit (Millipore) according to the manufacturer protocol. For scanning electron microscopy, whole-mounts fixed as before were dried with the critical point method and shadowed with gold protocol. For scanning electron microscopy, whole-mounts fixed as before were dried with the critical point method and shadowed with gold protocol. For scanning electron microscopy, whole-mounts fixed as before were dried with the critical point method and shadowed with gold protocol.

NSC culture, transduction, immunocytochemistry, and SA β-gal staining

Methods for adult NSC culture and self-renewal assessment have been described (Ferron et al., 2007). TSA, AA, VPA, KUS5933, and PFTω were from Sigma. Vectors containing pS3 shRNAs, methods used for retrovirus production, and NSC transduction have been described (Marqués-Torrejón et al., 2013). In brief, single cells obtained from passage 1 (P1) neurospheres were infected o/n by incubation with diluted (1:5) supernatants of shRNA retroviruses produced in Plat-E cells; after 5 days, the clones were dissociated and single cells were adhered for 10 min to a Matrigel substrate and stained. The Senescence Cells Histochemo Staining Kit (Sigma) was used following manufacturer’s instructions. Representative images were obtained in a Nikon Eclipse E800 microscope and positivity was quantified using IMAGE 1.49i software (NIH, USA). Briefly, cyan pixels were segmented by color threshold and their total intensity was divided by the number of cells in each image. For immunocytochemistry, incubation with antibodies for mouse anti-β2HAX (1:100, Millipore), rat anti-BrdU (1:600, Abcam), and rabbit anti-S100β (1:1200, Dako) was performed on Matrigel-attached freshly dissociated NSCs as described (Ferron et al., 2007). Secondary antibodies and DAPI counterstaining were as for histological analyses. Pictures of random fields in each condition were taken on the InCell Analyzer 2000 (GE Healthcare, Valencia, Spain), and positive cells were scored using the InCell INVESTIGATOR 1.6 software.

Protein isolation/Western blotting

Neurospheres were homogenized in lysis buffer (20 mM PBS, 150 mM NaCl, 5 mM EDTA, 1% Triton X-100, 2.5 mM sodium orthovanadate, 20 mM sodium fluoride, 2 mM phenyl-methane-sulfonylfluoride and 1X Complete Mini tablet (Roche) for 20 min and centrifuged at 20 000 g for 15 min at 4 °C. Protein concentration was determined by Pierce BCA Protein assay kit (Thermo Scientific, Madrid, Spain). Protein extracts (50–70 μg) were denatured in Laemmli’s sample buffer containing β-mercaptoethanol, sodium dodecyl sulfate (SDS), and bromophenol blue, separated by 10% SDS–polyacrylamide gel electrophoresis and transferred to a nitrocellulose membrane (Sigma). The membranes were blocked 1 h at room temperature with 5% nonfat dry milk in Tris-buffered saline supplied with 0.1% Tween 20 (TBS-T) and incubated with rabbit anti-p53 (1:500, Novocorsa), ac-p53 (1:1000, Cell Signalling), ac-H3 (1:10 000, Sigma), p16 (1:200, Santa Cruz) or pp53 (S18) (1:500, R&D), rat anti-p19 (1:1000, Abcam), or mouse anti-Gapdh (1:1500, Millipore) antibodies in blocking buffer, o/n at 4 °C. After extensive washing with TBS-T, membranes were incubated for 1 h at RT with horseradish peroxidase-conjugated goat antibodies to mouse (1:2000, Dako), rabbit (1:5000, Amersham), or rat (1:1000, Amersham) IgGs in blocking buffer. Peroxidase activity was detected with Western Lightning Plus reagents (Perkin Elmer) according to supplier recommendations.

RNA isolation and qRT–PCR analysis

Total RNA was extracted using TRIzol and cDNA synthesized using PrimeScript RT reagent kit (Takara); cDNA products were amplified on a StepOnePlus Real-Time PCR System (Applied Biosystems) using Premix Ex Taq (Probe qPCR) (Takara). Taqman probes for mouse S100b (Mm00485897_m1) and Gapdh (Mm99999915_g1) were purchased from Applied Biosystems.

Statistics

Data are presented as the mean ± SEM of a number (n) of cultures or independent mice per experimental condition. Statistical significance was determined by two-tailed Student’s t-tests or two-way ANOVA with post hoc analysis by Bonferroni’s test. Significance was set at *P < 0.05, **P < 0.01, or ***P < 0.001. Statistical analyses were carried out using SPSS V. 17 (2008 SPSS Inc., Chicago, IL, USA) or GraphPad Prism (2007 GraphPad Software Inc.).

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Author contributions

R. S.-C. performed most of the experiments; A. P.-V. performed the analyses in whole-mount preparations; J. M. M.-R. performed the analyses with pS3 and p19 mutant cultures; M. A. M.-T. contributed to the analysis of the mouse phenotypes; M. P. contributed to the design of the experiments; I.F. and F. P.-S. designed and supervised the study, secured funding, analyzed the data, and wrote the manuscript. All authors contributed to experimental design, data analysis, discussion, and writing of the manuscript.

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Conflict of interest
None declared.

References

Ahlenius H, Visan V, Kokaia M, Lindvall O, Kokaia Z (2009) Neural stem and progenitor cells retain their potential for proliferation and differentiation into functional neurons despite lower number in aged brain. J. Neurosci. 29, 4408–4419.

Bruggeman SW, Valk-Lingbeek ME, van der Stoep PP, Jacobs JJ, Kieboom K, Tanger E, Hulsman D, Leung C, Arsenijevic Y, Marino S, van Lohuizen M (2005) Ink4a and Arf differentially affect cell proliferation and neural stem cell self-renewal in Bmi-1-deficient mice. Genes Dev. 19, 1438–1443.

Capilla-Gonzalez V, Cebrian-Silla A, Guerrero-Cazes H, Garcia-Verdugo JM, Quiriones-Hinojosa A (2014) Age-related changes in astrocytic andependymal cells of the subventricular zone. Glia 62, 790–803.

Chao C, Herr D, Chun J, Xu Y (2006) Sirt3 and 23 phosphorylation is required for p53-dependent apoptosis and tumor suppression. EMBO J. 25, 2615–2622.

van Deursen JM (2014) The role of senescent cells in aging. Nature 509, 439–446.

Diaz-Moreno M, Hortiguela R, Goncalves A, Garcia-Carpio I, Manich G, Garcia-Bermudez E, Moreno-Estelles M, Eguiluz C, Vilaplana J, Pelegri C, Vilas M, Mira H (2013) Abeta increases neural stem cell activity in senescence-accelerated SAMIP8 mice. Neurobiol. Aging 34, 2623–2638.

Emwere E, Shingo T, Gregg C, Fujikawa H, Ohta S, Weiss S (2004) Aging results in a reduced epidermal growth factor receptor signaling, diminished olfactory neurogenesis, and deficits in fine olfactory discrimination. J. Neurosci. 24, 8355–8365.

Ferron SR, Andreu-Aguiló C, Mira H, Sanchez P, Marques-Troncoso MA, Fariñas I (2007) A combined exin vivo assay to detect effects of exogenously added factors in neural stem cells. Nat. Protoc. 2, 849–859.

Foti SB, Chou A, Moll AD, Roskams AJ (2013) HDAC inhibitors dysregulate neural progenitors and neurogenesis in SAMP8 mouse model of Alzheimer’s disease. Brain Res. 1389, 183–193.

Gil J, Peters G (2006) Regulation of the INK4b-ARF-INK4a tumour suppressor locus: all for one or one for all. EMBO J. 25, 2085–2095.

Haller C, Schmied H, Knappe B, Wiethase M, Dierichs S, Olschowsky S, Heinrich L, Michalik K, Schlesner H (2012) Senescent neural progenitors and neurogenesis during ageing. Nature 484, 448–452.

Kamijo T, Zindy F, Roussel MF, Quelle DE, Downing JR, Ashmun RA, Grosveld G, Krishnamurthy J, Torrice C, Ramsey MR, Kovalev GI, Al-Regaiey K, Su L, Sharpless NE (2010) Ink4a and Arf differentially affect cell proliferation and neural stem cell self-renewal in young but not old mice by reducing p16Ink4a and p19Arf expression. Cell 143, 227–239.

Kamijo T, Zindy F, Roussel MF, Quelle DE, Downing JR, Ashmun RA, Grosveld G, Krishnamurthy J, Torrice C, Ramsey MR, Kovalev GI, Al-Regaiey K, Su L, Sharpless NE (2010) Ink4a and Arf differentially affect cell proliferation and neural stem cell self-renewal in young but not old mice by reducing p16Ink4a and p19Arf expression. Cell 135, 227–239.

Porlan E, Morante-Redolat JM, Marqués-Troncoso MA, Andreu-Aguiló C, Carnero C, Gómez-Ibáñez E, Soto A, Vidal A, Ferrón SR, Fariñas I (2013) Transcriptional repression of Bmp2 by p21(Waf1/Cip1) links quiescence to neural stem cell maintenance. Nat. Neurosci. 16, 1567–1575.

Raponi E, Agenes F, Delphín C, Assand N, Baudier J, Legraverend C, Deloulme JC (2007) S100B expression defines a state in which GFAP-expressing cells lose their neural stem cell potential and acquire a more mature developmental stage. Glia 55, 165–177.

Shook BA, Manz DH, Peters JJ, Kang S, Conover JC (2012) Spatial-temporal changes to the subventricular zone stem cell pool through aging. J. Neurosci. 32, 6947–6956.

Silva-Vargas V, Crouch EE, Doetsch F (2013) Adult neural stem cells and their niche: a dynamic duo during homeostasis, regeneration, and aging.Curr. Opin. Neurobiol. 23, 935–942.

Simboeck E, Ribeiro JD, Teichmann S, Di Croce L (2011) Epigenetics and senescence: learning from the INK4A-ARF locus. Biochem. Pharmacol. 15, 1261–1370.

Stoll EA, Habibi BA, Mikheev AM, Lasiene J, Massey SC, Swanson KR, Rostomily RC, Horner PJ (2011) Increased re-entry into cell cycle mitigates age-related neurogenic decline in the murine subventricular zone. Stem Cells 29, 2005–2017.

Sun G, Yu RT, Evans RM, Shi Y (2007) Orphan nuclear receptor TLX recruits histone deacetylases to repress transcription and regulate neural stem cell proliferation. Proc. Natl Acad. Sci. USA 104, 15282–15287.

Takeda T (2009) Senescence-accelerated mouse (SAM) with special references to neurodegeneration models, SAMP8 and SAMP10 mice. Neurobiol. Aging. 30, 639–659.

Taniwasawa K, Mikami E, Fukui N, Honda Y, Honda S, Ohsawa I, Ito M, Endo S, Ibara K, Ohno K, Kishimoto Y, Ishigami A, Maruyama N, Sawabe M, Iseki H, Okazaki Y, Hasegawa-Iishi S, Takei S, Shimada A, Hosokawa M, Morii M, Higuchi T, Takeda T, Higuchi M, Tanaka M (2013) Exome sequencing of senescence-accelerated mice (SAM) reveals deleterious mutations in degenerative disease-causing genes. BMC Genom. 14, 248.

Tomobe K, Nomura Y (2009) Neurochemistry, neuropathology, and heredity in SAMP8: a mouse model of senescence. Neurochem. Res. 34, 660–669.

Trepope V, Craig CG, Morshed CM, van der Kooy D (1997) Transforming growth factor-alpha-null and senescent mice show decreased neural progenitor cell proliferation in the forebrain subependyma. J. Neurosci. 17, 7890–7895.

Willis-Martinez D, Richards HW, Timchenko NA, Medrano EE (2010) Role of HDAC1 in senescence, aging, and cancer. Exp. Gerontol. 45, 279–285.

Xie Q, Lin T, Zhang Y, Zheng J, Bonanno JA (2005) Molecular cloning and characterization of a human AIF-like gene with ability to induce apoptosis. J. Biol. Chem. 280, 19673–19681.

Zhao Q, Dalgaard CL, Wynder C, Doughty ML (2011) Histone deacetylase inhibitors SAHA and sodium butyrate block G1-to-S cell cycle progression in neurosphere formation by adult subventricular cells. BMC Neurosci. 12, 50.

Zindy F, Quelle DE, Roussel MF, Sherr CJ (1997) Expression of the p16INK4a tumor suppressor versus other INK4 family members during mouse development and aging. Oncogene 15, 203–211.