Chronic Antidepressant Treatment Increases Neurogenesis in Adult Rat Hippocampus

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Recent studies suggest that stress-induced atrophy and loss of hippocampal neurons may contribute to the pathophysiology of depression. The aim of this study was to investigate the effect of antidepressants on hippocampal neurogenesis in the adult rat, using the thymidine analog bromodeoxyuridine (BrdU) as a marker for dividing cells. Our studies demonstrate that chronic antidepressant treatment significantly increases the number of BrdU-labeled cells in the dentate gyrus and hilus of the hippocampus. Administration of several different classes of antidepressant, but not non-antidepressant, agents was found to increase BrdU-labeled cell number, indicating that this is a common and selective action of antidepressants. In addition, upregulation of the number of BrdU-labeled cells is observed after chronic, but not acute, treatment, consistent with the time course for the therapeutic action of antidepressants. Additional studies demonstrated that antidepressant treatment increases the proliferation of hippocampal cells and that these new cells mature and become neurons, as determined by triple labeling for BrdU and neuronal- or glial-specific markers. These findings raise the possibility that increased cell proliferation and increased neuronal number may be a mechanism by which antidepressant treatment overcomes the stress-induced atrophy and loss of hippocampal neurons and may contribute to the therapeutic actions of antidepressant treatment.

Key words: proliferation; granule cell; fluoxetine; tranylcypromine; reboxetine; depression

Depression is a devastating illness that is estimated to affect 12–17% of the population at some point during the lifetime of an individual (Kessler et al., 1994). Antidepressants are commonly prescribed for depression and other affective disorders, although the molecular and cellular mechanisms by which these agents exert their therapeutic effects are not well understood. Preclinical and clinical research has focused on the interactions between stress and depression and their effects on the hippocampus, among other brain regions (Duman et al., 1999; McEwen, 1999). For example, the hippocampus has been shown to undergo morphological changes in response to stress, including atrophy and loss of CA3 pyramidal neurons after exposure to physical or psychosocial stress (Watanabe et al., 1992c; Stein-Behrens et al., 1994; Margo ninos et al., 1996; McEwen, 1999). In addition, brain-imaging studies demonstrate that hippocampal volume is decreased in patients with stress-related psychiatric illnesses, including depression and post-traumatic stress disorder (Sapolsky, 1996; Sheline et al., 1996).

The hippocampus is one of only a few brain regions where production of neurons occurs throughout the lifetime of animals, including humans (Eriksson et al., 1998). Hippocampal neurogenesis can be influenced by several environmental factors and stimuli (Kuhn et al., 1996; Kempermann et al., 1997; Gould et al., 1999a; van Praag et al., 1999b). Importantly, it has been shown that stressful experiences, including both physical and psychosocial stress, suppress the formation of hippocampal granule cells in a number of mammalian species (Gould et al., 1997, 1998; Tanapat et al., 1998). Decreased cell proliferation has also been reported in response to both acute and chronic stress paradigms (Fuchs et al., 1997). It is conceivable that the stress-induced downregulation of granule cell genesis, as well as atrophy and death of CA3 pyramidal neurons, contributes to the reduction in hippocampal volume that is clinically observed (Sapolsky, 1996; Sheline et al., 1996; Duman et al., 1999).

The possibility that antidepressant treatment could oppose or reverse the actions of stress on the morphology and proliferation of hippocampal neurons is suggested by studies demonstrating that antidepressants upregulate the expression of brain-derived neurotrophic factor (BDNF) in the hippocampus (Nibuya et al., 1995). BDNF has been shown to promote the differentiation and survival of neurons during development and in adult brain, as well as in cultured cells (Emberger and Hall, 1995; Palmer et al., 1997; Takahashi et al., 1998). In addition, chronic antidepressant treatment completely blocks the stress-induced downregulation of BDNF expression in the hippocampus, demonstrating that antidepressant treatment can oppose the dystrophic actions of stress (Nibuya et al., 1995). Given the association between depression, stress, and hippocampal neurogenesis, the current series of studies was performed to determine whether antidepressant administration influences hippocampal neurogenesis in the adult rat. After administering different classes of antidepressant drugs or electroconvulsive seizure (ECS), we administered bromodeoxyuridine (BrdU), a thymidine analog that labels dividing cells in S-phase (Takahashi et al., 1992). The effects of antidepressant treatment on proliferation, differentiation, and survival of cells in the dentate gyrus and hilus of the hippocampus were determined using several treatment paradigms.

MATERIALS AND METHODS

Antidepressant treatment. A 175–250 g Sprague Dawley rats (Harlan Sprague Dawley, Indianapolis, IN) were used for all experiments. All animal treatments and maintenance of the rat colony were in accordance with National Institutes of Health laboratory care standards. Animals were housed individually or in pairs and were provided food and water ad libitum. A temperature of 22°C and 12 h light/dark cycle was maintained. Rats were administered either an antidepressant or vehicle. ECS was administered via earclip electrodes (50 mA; 0.3 sec) once daily for 10 d. Earclips were used to control animals, but no electrical current was administered. Drugs and their vehicles were administered intraperitoneally according to standard regimens (Nibuya et al., 1995, 1996): tranylcypromine, 7.5 mg/kg for the first 7 d and then 10 mg/kg for 14 d; reboxetine, 20 mg/kg, 2× per day for 21 d; fluoxetine, 5 mg/kg for 1, 5, 14, or 28 d; haloperidol, 1 mg/kg for the first 7 d and then 2 mg/kg for 7 d; venlafaxine, 1 mg/kg saline for tranylcypromine and reboxetine, 1 ml/kg distilled water for fluoxetine, and 1 ml/kg DSMO for haloperidol (n = 8 for each group).

BrdU labeling. For analysis of BrdU-positive cells, rats were administered BrdU (4 × 75 mg/kg every 2 hr; Sigma, St. Louis, MO) 4 d after the...
last antidepressant or haloperidol treatment (Fig. 1A). The 4-d time point was chosen because a similar paradigm has been used in a previous study of chemical-induced seizures on hippocampal neurogenesis (Parent et al., 1997). Twenty-four hours after the last BrdU injection, rats were killed and transcardially perfused (0.1 M cold PBS for 5 min followed by 4% cold paraformaldehyde for 17 min). For determination of cell phenotype, ECS or fluoxetine (14 d)-treated rats were allowed to survive 28 d after the last BrdU injection (Fig. 1A). To investigate the effect of antidepressant treatment specificity on cell proliferation, ECS- or fluoxetine (14 d)-treated rats were given one injection of BrdU (75 mg/kg) and perfused 2 hr later (Fig. 1A).

To determine the effects of antidepressants on cell survival, BrdU was administered before chronic administration of fluoxetine (Fig. 1B). BrdU (4 × 75 mg/kg every 2 hr) was administered to drug-naïve rats, and 24 hr after the last BrdU injection, rats were started on a chronic regimen of fluoxetine (5 mg/kg for 14 d). Twenty-eight days after the last BrdU injection (~14 d after ECS or fluoxetine treatment, respectively) rats were perfused, and BrdU labeling was examined. After perfusion, all brains were post-fixed overnight in paraformaldehyde (with shaking) at 4°C and stored at 4°C in 30% sucrose. Serial sections of the brains were cut (35 μm sections) through the entire hippocampus (plates 26–40; Paxinos and Watson, 1986) on a freezing microtome, and sections were stored in PBS/NaN3.

Immunohistochemistry. Free-floating sections were used in the determination of BrdU labeling. DNA denaturation was conducted by incubation for 2 hr in 50% formamide/2× SSC at 65°C. Sections were then incubated for 30 min in 2N HCl and then 10 min in boric acid. A filter washing in PBS, sections were incubated for 30 min in 3% H2O2 to eliminate endogenous peroxidases. A filter blocking with 3% normal horse serum (NHS) in 0.01% Triton X-100, cells were incubated with anti-mouse BrdU (1:1000; Boehringer-Mannheim, Indianapolis, IN) followed by amplification with an avidin–biotin complex (Vector Laboratories). Cells were visualized with DAB (Vector Laboratories). Analysis of the number of BrdU-labeled cells demonstrated that chronic antidepressant administration significantly increased the number of BrdU-positive cells in the dentate gyrus (Fig. 2) relative to control. Chronic administration of ECS increased the number of BrdU-labeled cells by ~50%, whereas the chemical antidepressants increased the BrdU labeling by 20–40% (Fig. 3). ECS is clinically the most effective treatment for refractory depression, and this increase in BrdU-positive cells is in agreement with a report by previous investigators (Madsen et al., 2000). The chemical antidepressants tested include a monoamine oxidase inhibitor (tranylcypromine), a serotonin-selective reuptake inhibitor (fluoxetine), and a norepinephrine-selective reuptake inhibitor (reboxetine).

Next, the time course for antidepressant regulation of BrdU labeling was examined. Aministration of fluoxetine for 1 or 5 d did not significantly affect the number of BrdU-positive cells compared with control (Fig. 4). After 14 or 28 d of fluoxetine treatment, a significant increase in BrdU-positive cells was seen compared with vehicle-treated controls and 1 or 5 d of treatment. This indicates that chronic, but not acute, antidepressant treatment increases BrdU labeling in the hippocampus, which is consistent with the time course for the therapeutic action of antidepressants (Duman et al., 1997).

To determine whether upregulation of the number of BrdU-labeled cells is specific to antidepressants, the influence of a non-antidepressant psychotropic drug was examined. Chronic administration of a clinically relevant dose of haloperidol, an antipsychotic agent, did not significantly influence the number of BrdU-labeled cells in dentate gyrus (vehicle, 417 ± 259; haloperidol, 378 ± 238 BrdU-labeled cells; mean ± SEM; n = 6 per group). This indicates that the increase in BrdU-positive cells may be specific to antidepressant treatment.

A modified stereology protocol was used to count the number of BrdU-positive cells throughout the SGZ and hilus (West et al., 1993; Gould et al., 1999a). The SGZ, the border between the granule cell layer and hilus, has been shown to contain the progenitor cells that divide and migrate into the granule cell layer where they mature into neurons or astrocytes (Cameron et al., 1993). The percentage of BrdU-positive cells in the hilus and SGZ was the same in control and antidepressant-treated groups, so the cell counts in the two regions were summed to give the total number of labeled cells per dentate gyrus used in the statistical analysis.

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pressants. In addition, we have found that chronic administration of morphine decreases granule cell proliferation, an effect opposite to that of antidepressant treatment (Eisch et al., 2000).

**Chronic antidepressant treatment increases cell proliferation**

To specifically isolate the effect of antidepressant treatment on cell proliferation, rats were administered ECS (10 d) or fluoxetine (14 d), given a single injection of BrdU, and killed 2 hr later. At this time point, both ECS and fluoxetine treatment significantly increased the number of BrdU-positive cells relative to the respective controls (vehicle, 3610 ± 330; fluoxetine, 4350 ± 420; ECS, 5780 ± 720; $F(2,18) = 6.89; p < 0.05$). These results indicate that antidepressant treatment increases the proliferation of hippocampal cells.

In all of the animals killed at either 2 or 24 hr after BrdU injection, the BrdU-positive cells in the hilus and dentate gyrus were found in clusters (Fig. 5A–C), with irregularly shaped nuclei and diffuse patterns of BrdU staining. This is representative of immature cells undergoing division (Kuhn et al., 1996; Parent et al., 1997). The number of cells per cluster was not affected by ECS or chemical antidepressant treatments. In addition, in all animals killed at 2 hr, a cell undergoing mitosis was seen. The existence of these mitotic figures (Fig. 5B, C) indicates that the BrdU is labeling newly born cells and not labeling cells that are undergoing DNA repair. Taken together, these results demonstrate that chronic antidepressant administration increases the number of BrdU-positive cells in the adult hippocampus and that the increase in labeling can be attributed to an increase in cell proliferation.

In a separate experiment, we examined the influence of antidepressant treatment on cell proliferation when BrdU is administered without a drug washout period. Fluoxetine was administered for 14 d and then BrdU was administered 2 hr after the last fluoxetine treatment. This paradigm also significantly increased the number of BrdU-positive cells (vehicle, 4962 ± 398; fluoxetine, 6740 ± 498; $F(4,35) = 4.35; p < 0.05$). The magnitude of this increase was similar to that observed when BrdU was administered 4 d after the last fluoxetine treatment (Fig. 3). These results indicate that fluoxetine increases proliferation within hours after the last treatment and that this effect is sustained for at least 4 d. Alternatively, this data provides evidence that that the increase in BrdU-labeled cells is not a rebound effect that occurs during the 4 d allowed for the washout period.

To determine whether the antidepressant-induced upregulation of cell proliferation is specific to the hippocampus, another brain region known to contain progenitor cells in adulthood, the subventricular zone of the lateral ventricle (Kuhn et al., 1996), was examined. Chronic ECS or fluoxetine treatment did not influence the number of BrdU-labeled cells per section of this brain region (data not shown; $p > 0.05$). In addition to demonstrating that...
antidepressants specifically increase cell proliferation in the hippocampus, these results also suggest that antidepressant administration does not have a general effect on the amount of BrdU entering the brain or the incorporation of BrdU into the DNA of proliferating cells.

**Antidepressant treatment increases neurogenesis**

Newly born cells in the hippocampus can have several fates: some cells die, whereas others survive and differentiate into mature neurons or glia. To examine the influence of antidepressant treatment on cell fate, the number and phenotype of the BrdU-positive cells was determined by triple immunofluorescent labeling (Fig. 7). Confocal microscopy, using z-plane sections to confirm colocalization for each cell, revealed that the majority of BrdU-positive cells were neuronal (75%) and not glial (13%) in both the control and antidepressant-treated groups. The remaining 12% of cells were not labeled with either a neuronal or glial marker; these cells may represent a phenotype not labeled here, or they may be cells located deeper in the tissue section and therefore not accessible to the antibodies used. They may also represent quiescent undifferentiated cells (Eriksson et al., 1998; van Praag et al., 1999b). These data indicate that the antidepressants do not affect the differentiation of cells into neurons or glia.

In a separate experiment, the influence of antidepressant treatment on the survival of cells that have already been born in the hippocampus was determined. In this experiment, we administered BrdU 1 d before initiating chronic fluoxetine treatment (14 d). Twenty-eight days after the BrdU administration (14 d after the last fluoxetine injection), there was no difference in the number of BrdU-positive cells in the hippocampus (control, 2764 ± 320; fluoxetine, 2808 ± 348 BrdU-labeled cells; mean ± SEM; n = 6). The survival rate of the newly born cells in the hippocampus is approximately the same in both the vehicle and fluoxetine-treated groups (~50% of BrdU-labeled cells in each group survives at this time point). This indicates that chronic antidepressant treatment does not directly affect the rate of maturation and survival of BrdU-positive cells in the hippocampus.

**DISCUSSION**

The results of this study demonstrate that chronic antidepressant administration increases neurogenesis in the dentate gyrus of the adult rat hippocampus. Upregulation of neurogenesis is observed in response to administration of different classes of antidepressants, indicating that increased neurogenesis may be a common action of antidepressant treatment. In addition, increased BrdU labeling is observed after chronic (14 or 28 d), but not short-term (1 or 5 d) antidepressant treatment. These findings indicate that the time course for the upregulation of BrdU labeling is consistent with the time delay required for the therapeutic action of antidepressants (Duman et al., 1997).

In contrast, chronic administration of the non-antidepressant...
psychotropic drug haloperidol, does not increase BrdU labeling in hippocampal granule cells. In addition, we have recently demonstrated that morphine, another non-antidepressant psychotropic drug, decreases BrdU labeling of granule cells (Eisch et al., 2000). These findings indicate that the upregulation of hippocampal BrdU labeling may be pharmacologically selective to the chemical antidepressant drugs. The lack of effect of haloperidol in this study differs from two previous studies, one reporting an increase (Dawirs et al., 1998), and one reporting a decrease (Backhouse et al., 1982) in cell proliferation. However, there are several differences between the current and these previous reports, including dose and time course of drug treatment, species and age of test animals, and the BrdU labeling protocol. One or more of these variables could account for the difference in our results. In the current study, the dose and time of haloperidol treatment were consistent with the therapeutic treatment regimen (Nibuya et al., 1995), in contrast to the two previous studies. Using this relevant treatment paradigm and the same BrdU labeling protocol that was used for the antidepressant studies, haloperidol does not increase BrdU labeling.

Regulation of neurogenesis could occur at several different stages, including cell proliferation, differentiation, and survival. We demonstrate that 2 hr after BrdU administration, antidepressant treatment significantly increased the number of BrdU-labeled cells compared with the saline control group. This indicates that antidepressant treatment increases cell proliferation. To examine the effect of antidepressants on survival of labeled cells, BrdU was administered before the start of antidepressant treatment, and the number of labeled cells was determined 4 weeks later. Under these conditions, the survival of the BrdU-labeled cells can be determined independent of the influence of antidepressant treatment on cell proliferation. In this experiment, the number of BrdU-labeled cells in the treatment group was not different from the control group, indicating that antidepressants do not influence cell survival.

The differentiation of labeled cells was determined 4 weeks after antidepressant or saline treatment by colocalization of neuronal and glial phenotypic markers in BrdU-labeled cells. At this time point, there is a significant increase in the number of BrdU-labeled cells relative to controls. This increase is a result of the upregulation of cell proliferation by antidepressant treatment. The majority (75%) of the surviving BrdU-positive cells express a neuronal marker (i.e., NeuN or NSE) and have physical characteristics of healthy, viable neurons. A much smaller number (13%) of cells express a glial marker (GFAP). The remaining 12% of cells were not labeled with either cell marker and may represent another phenotype or quiescent undifferentiated cells. This ratio of labeled neurons and glia is similar to that reported in previous studies (Eriksson et al., 1998; van Praag et al., 1999b), and was not significantly influenced by antidepressant treatment. This is consistent with our finding that antidepressant treatment increases the proliferation, but not survival, of labeled hippocampal cells. Once the cells are induced to proliferate by antidepressant treatment, their survival and differentiation rates are identical to animals treated with vehicle. The result is a net increase in the number of neurons produced, or neurogenesis, in antidepressant-treated animals compared with vehicle controls.

It is not currently known whether the mature BrdU-positive neurons seen in this study are functional in vivo. However, new neurons in the granule cell layer in hippocampus have been demonstrated to send axons to the CA3 pyramidal cell layer,

Figure 7. Triple labeling confirms that BrdU-positive cells mature into neurons. Rats received BrdU injections 4 d after the last ECS treatment and were killed 4 weeks later. A representative confocal laser-scanning image (66×) of a section from a fluoxetine-treated rat that has been triple-labeled with BrdU (A, green; BrdU-positive cells indicated by arrows), GFAP (B, blue), and NeuN (C, red). The merged image (D) demonstrates cells that are double-labeled in the GCL for BrdU and NeuN but not GFAP.
appropriate projection area for granule cells (Markakis and Gage, 1999). Maturity BrdU-labeled cells in the granule cell layer are also surrounded by synaptic vesicles, indicating that they receive synaptic inputs. Learning and memory tasks that are dependent on the hippocampus result in an upregulation of neurogenesis (Gould et al., 1999a; van Praag et al., 1999b). In addition, increased neurogenesis in response to voluntary running has recently been correlated with an increase in granule cell long-term potentiation, a cellular model of learning and memory (van Praag et al., 1999a). Taken together, these studies demonstrate that newly formed cells differentiate into mature neurons that integrate into the existing hippocampal circuitry and may increase the functional capacity of this brain structure.

The mechanisms underlying the regulation of hippocampal neurogenesis are being actively investigated. Interestingly, both the cAMP cascade and BDNF, which we have found to be upregulated by antidepressant treatment (N ibuya et al., 1995, 1996; Thome et al., 2000), have been shown to play a role in the regulation of neurogenesis. Activation of the cAMP pathway or incubation with BDNF is reported to increase neuronal differentiation and neurite outgrowth of progenitor cells 

**REFERENCES**


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