

Role for Neuronal Nitric-Oxide Synthase in Cannabinoid-Induced Neurogenesis

Sun Hee Kim, Seok Joon Won, Xiao Ou Mao, Catherine Ledent, Kunlin Jin, and David A. Greenberg

Buck Institute for Age Research, Novato, California (S.H.K., S.J.W., X.O.M., K.J., D.A.G.); and Université Libre de Bruxelles, Institut de Recherche Interdisciplinaire en Biologie Humaine et Moléculaire, Bruxelles, Belgium (C.L.)

Received May 11, 2006; accepted July 7, 2006

ABSTRACT

Cannabinoids, acting through the CB1 cannabinoid receptor (CB1R), protect the brain against ischemia and related forms of injury. This may involve inhibiting the neurotoxicity of endogenous excitatory amino acids and downstream effectors, such as nitric oxide (NO). Cannabinoids also stimulate neurogenesis in the adult brain through activation of CB1R. Because NO has been implicated in neurogenesis, we investigated whether cannabinoid-induced neurogenesis, like cannabinoid neuroprotection, might be mediated through alterations in NO production. Accordingly, we measured neurogenesis in dentate gyrus (DG) and subventricular zone (SVZ) of CB1R-knockout (KO) and wild-type mice, some of whom were treated with the cannabinoid agonist *R*(+)-Win 55212-2 [(+)-[2,3-dihydro-5-methyl-3-[(morpholinyl)methyl]pyrrolo[1,2,3-de]-1,4-benzoxazin-yl]-(1-naphthalenyl)methanone] or the NO synthase (NOS) inhibitor

7-nitroindazole (7-NI). NOS activity was increased by ~25%, whereas bromodeoxyuridine (BrdU) labeling of newborn cells in DG and SVZ was reduced by ~50% in CB1R-KO compared with wild-type mice. 7-NI increased BrdU labeling in both DG and SVZ and to a greater extent in CB1R-KO than in wild-type mice. In addition, *R*(+)-Win 55212-2 and 7-NI enhanced BrdU incorporation into neuron-enriched cerebral cortical cultures to a similar maximal extent and in nonadditive fashion, consistent with a shared mechanism of action. Double-label confocal microscopy showed coexpression of BrdU and the neuronal lineage marker doublecortin (Dcx) in DG and SVZ of untreated and 7-NI-treated CB1R-KO mice, and 7-NI increased the number of Dcx- and BrdU/Dcx-immunoreactive cells in SVZ and DG. Thus, cannabinoids appear to stimulate adult neurogenesis by opposing the antineurogenic effect of NO.

Cannabinoids, which include naturally occurring plant-derived compounds [e.g., Δ^9 -tetrahydrocannabinol (THC)], endogenous signaling molecules found in animal brains (e.g., anandamide and 2-arachidonoylglycerol), and synthetic drugs [e.g., *R*(+)-Win 55212-2], act on receptors in the brain to modify neuronal function. In addition to their effects on normal physiological functions such as blood pressure, immunity, pain perception, appetite, and cognition, cannabinoids can also regulate the severity of brain injury.

We reported previously that administration of cannabinoids acting on the CB1 cannabinoid receptor (CB1R) reduces neuronal death from cerebral ischemia (Nagayama et

al., 1999). Consistent with this finding, stroke size is increased in mice with genetic deletion of CB1R (Parmentier-Batteur et al., 2002). Although the molecular mechanisms that underlie cannabinoid-induced protection from brain ischemia are not fully understood, one possibility is that cannabinoids act by suppressing excitotoxic signaling cascades, which have been implicated in ischemic brain injury. Supporting evidence includes the ability of cannabinoids to inhibit excitatory neurotransmission (Shen et al., 1996) and to block the neurotoxicity of excitatory amino acids like *N*-methyl-D-aspartate (NMDA) (Shen and Thayer, 1998). A principal signaling module through which NMDA operates to produce its toxic effects is thought to involve the calcium-dependent activation of neuronal nitric-oxide synthase (nNOS) and the resulting production of NO and metabolites such as peroxynitrite (Dawson et al., 1991). It is, therefore, notable that cannabinoids reduce NMDA- and NO-mediated

This work was supported by National Institutes of Health Grant NS39912 (to D.A.G.).

Article, publication date, and citation information can be found at <http://jpet.aspetjournals.org>.
doi:10.1124/jpet.106.107698.

ABBREVIATIONS: THC, Δ^9 -tetrahydrocannabinol; *R*(+)-Win 55212-2, (+)-[2,3-dihydro-5-methyl-3-[(morpholinyl)methyl]pyrrolo[1,2,3-de]-1,4-benzoxazin-yl]-(1-naphthalenyl)methanone; CB1R, CB1 cannabinoid receptor; NMDA, *N*-methyl-D-aspartate; nNOS, neuronal nitric-oxide synthase; NO, nitric oxide; DG, dentate gyrus; SVZ, subventricular zone; KO, knockout; NOS, NO synthase; eNOS, endothelial nitric-oxide synthase; iNOS, inducible nitric-oxide synthase; PBS, phosphate-buffered saline; BrdU, bromodeoxyuridine; 7-NI, 7-nitroindazole; Dcx, doublecortin; i.p., intraperitoneal.

neurotoxicity in retina (El-Remessy et al., 2003) and brain (Kim et al., 2005).

Cannabinoids also stimulate adult neurogenesis or the generation of new neurons from endogenous precursors in the dentate gyrus (DG) and subventricular zone (SVZ) of adult brain (Rueda et al., 2002; Jin et al., 2004c; Jiang et al., 2005). This effect, like protection from cerebral ischemia, appears to arise through activation of CB1R because it is diminished in CB1R-knockout (KO) mice (Jin et al., 2004c). Of interest, signaling by NO has been implicated in neurogenesis as well. Studies with NO donor or NOS inhibitor drugs have shown variable effects (Zhang et al., 2001; Cheng et al., 2003; Packer et al., 2003; Zhu et al., 2003; Matarredona et al., 2004; Moreno-Lopez et al., 2004), probably because they differ in their specificities for NOS isoforms and because nNOS, endothelial NOS (eNOS), and inducible NOS (iNOS) affect neurogenesis differently. This is evident from studies that employ isoform-specific NOS-knockout mice, which show that nNOS activity inhibits neurogenesis (Packer et al., 2003; Sun et al., 2005), whereas iNOS (Zhu et al., 2003) and eNOS (Reif et al., 2004) promote neurogenesis.

In view of these findings, we asked whether in neurogenesis as in stroke, the biological effect of cannabinoids might be mediated through their antagonism of nNOS-derived NO. To address this question, we measured neurogenesis in DG and SVZ of CB1R-KO (Ledent et al., 1999) and wild-type mice, some of whom were given a cannabinoid agonist [*R*(+)-Win 55212-2] or NOS inhibitor (7-nitroindazole) drug. We report that cannabinoids stimulate adult neurogenesis in DG and SVZ, at least partly by relieving the neurogenesis-inhibiting effect of nNOS.

Materials and Methods

Mice. CB1R-KO and wild-type littermates were bred in-house from mice originally obtained from Dr. Catherine Ledent (Ledent et al., 1999). Mice weighing 30–35 g were used for experiments and were euthanized by decapitation. Genotyping was performed as described in a previous publication (Parmentier-Batteur et al., 2002). Experiments were approved by local committee review and conducted according to National Institutes of Health guidelines.

NOS Activity Assay. NOS activity was measured using a commercial nitrate/nitrite colorimetric assay kit (catalog no. 78001; Cayman Chemical, Ann Arbor, MI), according to the manufacturer's instructions. In brief, the cerebral cortex was dissected on ice, homogenized in PBS, and centrifuged at 10,000*g* for 20 min. The supernatant was centrifuged at 100,000*g* for 30 min and then filtered through a 10-kDa molecular mass cutoff filter. Nitrate/nitrite was measured in triplicate in 40- μ l aliquots of the final filtrate on 96-well plastic microtiter plates. Absorbance was read at 570/620 nm using a Cytofluor series 4000 multiwell plate reader (Applied Biosystems, Foster City, CA).

Western Blot Analysis. Cerebral cortex was homogenized in lysis buffer containing 50 mM Tris-HCl, pH 7.5, 1% Nonidet P-40, 150 mM NaCl, 0.5% deoxycholic acid, 0.1% SDS, 100 mg/ml phenylmethylsulfonyl fluoride, and 1 μ g/ml aprotinin. Cell lysates were centrifuged at 12,000*g* for 15 min, and 10 μ g of protein was electrophoresed on 12% SDS-polyacrylamide gels and transferred to polyvinylidene difluoride membranes. Membranes were incubated overnight at 4°C with rabbit polyclonal anti-CB1 (1:1000; Calbiochem, San Diego, CA) or mouse monoclonal anti- β -actin (1:5000; Sigma, St. Louis, MO), then washed with PBS containing 0.1% Tween 20 and reacted with anti-rabbit or anti-mouse secondary antibodies (1:2000; Santa Cruz Biotechnology, Santa Cruz, CA). Immunoreactivity was

detected with luminol reagent (PerkinElmer Life Sciences Inc., Boston, MA).

Injection of Bromodeoxyuridine and 7-NI. Bromodeoxyuridine (BrdU) (50 mg/kg in saline) and 7-nitroindazole (7-NI; 50 mg/kg) or vehicle (dimethyl sulfoxide/1,2-propanediol/distilled water, 1:3:6) were administered i.p. twice daily at 8-h intervals for 3 consecutive days. After 24 h, 50- μ m coronal brain sections were stained with mouse monoclonal anti-BrdU (2 μ g/ml; Roche, Indianapolis, IN), and BrdU-positive cells in DG subgranular zone and SVZ were counted blindly in five sections per animal, spaced 200 μ m apart, under high power on a Nikon E800 microscope with a Magnifier digital camera (Nikon, Melville, NY).

Cell Culture. Cerebral cortical cultures enriched in cells of neuronal lineage were prepared from 16-day Charles River CD1 mouse embryos (Charles River, Wilmington, MA) in neurobasal medium containing 2% B27 supplement, 2 mM glutamate, and 1% each of penicillin and streptomycin. After 4 days, half of the medium was replaced with neurobasal medium containing 2% B27, and experiments were conducted after 5 days in vitro. At this time, cultures were maintained for 24 h in the presence of 50 μ g/ml BrdU and in the absence or presence of 100 nM *R*(+)-Win 55212-2, 100 μ M 7-NI, or both drugs combined.

Immunohistochemistry. Immunohistochemistry was performed on 50- μ m frozen coronal brain sections, corresponding to coronal coordinates interaural 8.7 to 10.2 mm, bregma -0.30 to bregma -1.2 mm (for SVZ), and interaural 4.48 to 5.86 mm, bregma -4.52 to bregma -3.14 (for DG subgranular zone). Sections were cut with a cryostat, fixed with 100% methanol for 10 min, incubated in 2 M HCl at 25°C for 1 h, and rinsed in 0.1 M boric acid, pH 8.5, at room temperature for 10 min. Sections were placed in 1% H₂O₂ for 15 min and then in blocking solution (2% goat serum, 0.3% Triton X-100, and 0.1% bovine serum albumin in PBS) for 2 h at room temperature and incubated with 2 μ g/ml mouse monoclonal anti-BrdU (Roche; 1:200) and goat polyclonal anti-doublecortin (Dcx) (Santa Cruz Biotechnology; 1:100) at 4°C overnight. Sections were washed with PBS, incubated for 2 h at 25°C with biotinylated goat-anti-mouse secondary antibody (Vector Laboratories, Burlingame, CA; 1:200), washed, and placed in avidin-peroxidase conjugate solution (Vector Laboratories) for 1 h. The horseradish peroxidase reaction was detected with 0.05% diaminobenzidine and 0.03% H₂O₂. Processing was stopped with H₂O, and sections were dehydrated through graded alcohols, cleared in xylene, and coverslipped in permanent mounting medium (Vector Laboratories). For confocal microscopy, a Nikon PCM-2000 laser-scanning confocal microscope and Simple PCI imaging software (Compix, Cranberry Township, PA) were used.

Statistics. Data were analyzed using Student's *t* test (single comparisons) or analysis of variance with post hoc Student-Newman-Keuls tests (multiple comparisons). Differences having *p* < 0.05 were considered significant.

Results

If CB1R-mediated signaling provides an important constraint on nNOS, then NOS activity might be increased in CB1R-KO mice. Accordingly, we measured cerebral NOS activity (which includes the nitrate- and nitrite-generating capacity of all NOS isoforms) in CB1R-KO and wild-type mice. Figure 1 shows that NOS activity was approximately 25% higher in knockouts, although expression of nNOS, which is not inducible, was not increased. These results are consistent with an inhibitory effect of CB1Rs on NOS activation.

As noted previously, nNOS inhibits (Packer et al., 2003; Sun et al., 2005), whereas iNOS (Zhu et al., 2003) and eNOS (Reif et al., 2004) promote adult neurogenesis. Because deletion of the CB1R gene enhances nNOS activity, CB1R-KO mice might be expected to show reduced levels of basal neu-

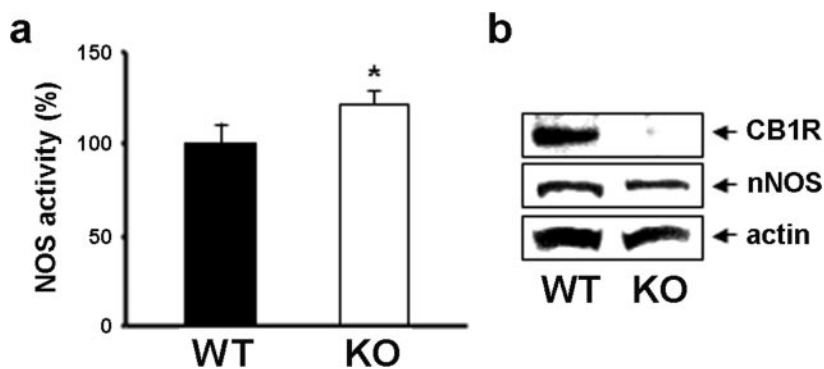


Fig. 1. NOS activity (a) and nNOS expression (b) in wild-type (WT) and CB1R-KO mice. *, $p < 0.05$ compared with WT.

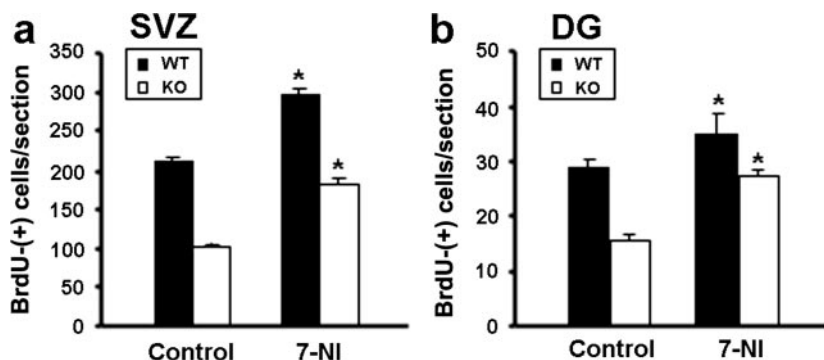


Fig. 2. Effect of in vivo inhibition of NOS with 7-NI on BrdU labeling in SVZ (a) and DG (b) of WT and CB1R-KO mice. 7-NI (50 mg/kg i.p.) or vehicle was administered together with BrdU, twice daily at 8-h intervals for 3 consecutive days. *, $p < 0.05$ compared with same-genotype control.

rogenesis. In addition, basal neurogenesis in these mice might be more readily stimulated by NOS inhibitors. We compared BrdU incorporation, an index of cell proliferation, in DG and SVZ of wild-type versus CB1R-KO mice (Fig. 2). In line with our prior findings (Jin et al., 2004c), BrdU labeling in DG and SVZ of CB1R-KO mice was only approximately 50% of that observed in wild-type mice. Next, we examined the effect of 7-NI, a relatively selective nNOS inhibitor (Southan and Szabo, 1996). 7-NI (50 mg/kg i.p., given twice daily for 3 days) increased BrdU labeling in both DG and SVZ and in both wild-type and CB1R-KO mice. However, the extent of this effect was greater in knockouts. Thus, 7-NI raised BrdU labeling in DG by only approximately 20% in wild-type but by almost 80% in CB1R-KO, mice. In SVZ, the observed increases were approximately 40 and 85%, respectively. Hence, the disinhibition of NOS activity in CB1R-KO mice is accompanied by enhanced sensitivity to the neurogenesis-promoting action of an NOS inhibitor.

We next examined the relative effects of a cannabinoid agonist and 7-NI on BrdU incorporation into cortical cultures enriched in cells of neuronal lineage (Fig. 3). As reported previously (Jin et al., 2002), antibodies against the neuronal lineage markers, β III tubulin and Hu, each stain $\geq 75\%$ of cells in these cultures. Cultures were treated for 24 h with concentrations of *R*(+)-Win 55212-2 (100 nM) or 7-NI (100 μ M) that produced maximal enhancement of BrdU incorporation. The extent of increase in BrdU labeling was similar for *R*(+)-Win 55212-2 and 7-NI, and the combination both drugs produced no further increase. These findings would be expected if *R*(+)-Win 55212-2 and 7-NI acted through the same pathway.

Although our in vivo studies focused on the two principal neuroproliferative zones in the adult brain, and our in vitro studies employed cultures consisting mostly of cells of neuronal lineage, BrdU labels dividing cells irrespective of cell

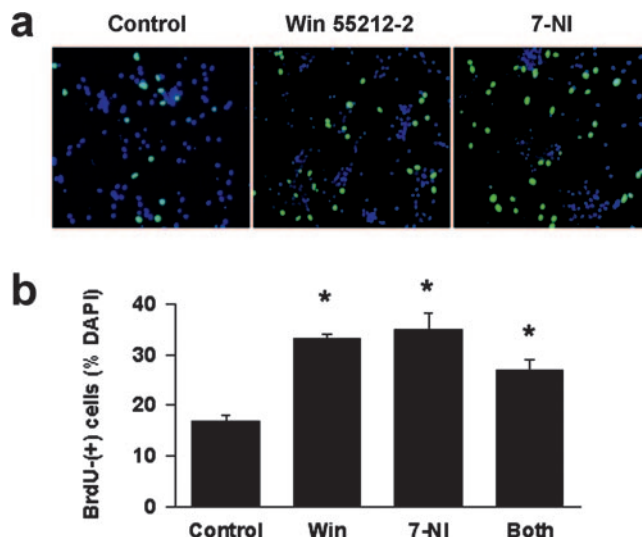


Fig. 3. Effects of the CB1R agonist *R*(+)-Win 55212-2 (100 nM), the NOS inhibitor 7-NI (100 μ M), and both drugs in combination on BrdU incorporation into wild-type cerebral cortical cultures at day 5 in vitro. a, 4,6-diamidino-2-phenylindole, blue; BrdU, green. b, *, $p < 0.05$ compared with control.

type. Consequently, it was important to determine directly whether the BrdU-immunoreactive cells that we analyzed were neuronal. To address this question, brain sections through DG or SVZ of BrdU-treated mice were stained with antibodies against both BrdU and the early neuronal marker Dcx (Francis et al., 1999; Gleason et al., 1999). Sections examined by double-label confocal fluorescence immunohistochemistry showed extensive coexpression of nuclear BrdU and cytoplasmic Dcx in DG and SVZ in both untreated and 7-NI-treated CB1R-KO mice (Fig. 4a). In addition, 7-NI treatment increased the number of Dcx-immunoreactive

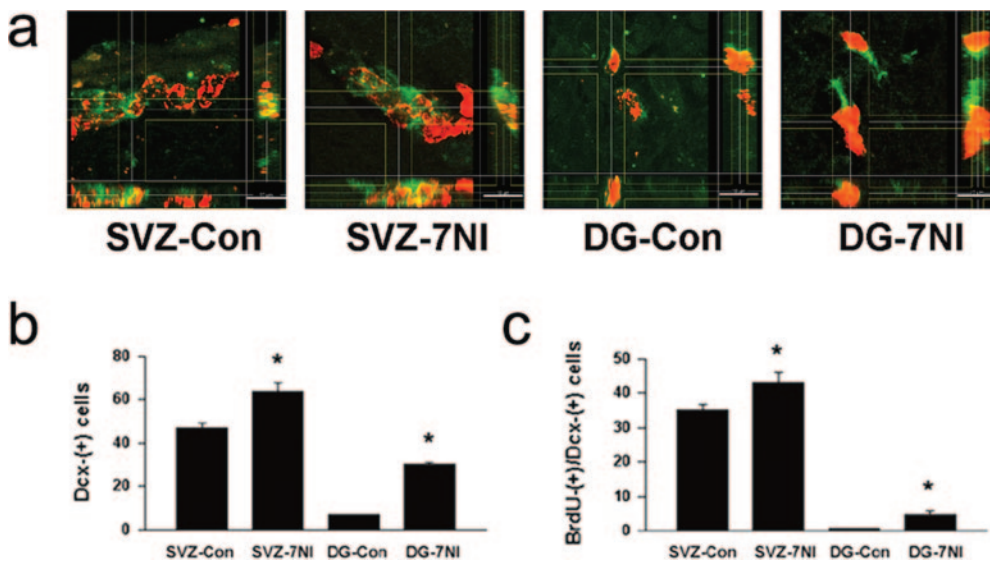


Fig. 4. Effects of in vivo inhibition of NOS with 7-NI on Dcx expression in SVZ and DG of CB1R-KO mice. 7-NI (50 mg/kg i.p.) or vehicle was administered together with BrdU, twice daily at 8-h intervals for 3 consecutive days. a, *xyz*-plane confocal images of sections stained with antibodies against BrdU (red nuclei) and Dcx (green cytoplasm). b, Dcx-immunopositive cell counts. *, $p < 0.05$ compared with controls not given 7-NI. c, Dcx- and BrdU-doubly immunopositive cell counts. *, $p < 0.05$ compared with controls not given 7-NI.

(Fig. 4b) and of BrdU- and Dcx-doubly immunoreactive (Fig. 4c) cells in SVZ and DG, confirming that the number of new neurons was increased.

Discussion

Understanding how adult neurogenesis is regulated is important because of its proposed role in brain repair and recovery from injury. Neurogenesis in DG, SVZ, or both is enhanced in response to diverse cerebral insults, including animal models of stroke (Jin et al., 2001), Alzheimer's disease (Jin et al., 2004a), and Huntington's disease (Jin et al., 2005), as well as Alzheimer's disease (Jin et al., 2004b) and Huntington's disease (Curtis et al., 2003) in patients. These findings have led to the hypothesis that an increase in neurogenesis may be one of the adaptive mechanisms the brain uses to limit injury and resulting functional impairment. If this is the case, even greater benefit might be achieved by therapeutic enhancement of neurogenesis,

A surprisingly broad range of drugs have been found to stimulate neurogenesis in vitro or in the brains of experimental animals in vivo. These appear to include cannabinoids, although some studies have yielded discrepant results, at least partly because some of the drugs employed were not entirely specific for cannabinoid receptors (for discussion, see Jin et al., 2004c). We reported previously (Jin et al., 2004c) that CB1R are expressed on neuronal precursor cells in the adult mouse brain and that the number of BrdU-labeled cells in DG and SVZ was reduced by approximately 50% in CB1R-KO compared with wild-type mice; as in the present study, most BrdU-labeled cells in these regions expressed Dcx, indicative of neuronal lineage. These data argue that CB1R-mediated activity normally exerts a proneurogenic influence. However, we did not examine the signaling mechanisms involved downstream of CB1R. In another study, Jiang et al. (2005) also found colocalization of CB1R with nestin, a marker for neuroepithelial precursors, some of which will undergo neuronal differentiation, in adult rat brain. They showed that the synthetic cannabinoid HU210 stimulated proliferation of neuronal precursors in vitro, and that this effect was mediated through CB1R, guanine nucleotide bind-

ing ($G_{i/o}$) proteins, and extracellular signal-regulated kinase. Furthermore, long-term administration of HU210 to rats increased neurogenesis, measured by colocalization of BrdU with the mature neuronal marker protein NeuN, in DG. Cannabinoids appear to regulate the production of oligodendrocytes as well because CB1R activation promotes the survival of oligodendrocyte precursors from rat brain, via $G_{i/o}$, phosphatidylinositol 3-kinase, and Akt (Molina-Holgado et al., 2002).

In the present study, we found evidence implicating an interaction between cannabinoid and NO signaling in the regulation of adult neurogenesis. Thus, NOS activity was increased in CB1R-KO mice, and the associated reduction in neurogenesis was reversed by the nNOS-preferring NOS inhibitor, 7-NI. Moreover, the cannabinoid agonist *R*(+)-Win 55212-2 and 7-NI increased BrdU labeling in mouse brain cultures to a similar extent, and their effects were not additive. Finally, BrdU was localized to cells of neuronal lineage, defined by Dcx expression, in DG and SVZ.

Cannabinoid and NO signaling pathways have been linked in the past, although not in the setting of neurogenesis. For example, several studies have reported that cannabinoids inhibit iNOS in macrophages or glia (Jeon et al., 1996; Waksman et al., 1999; Esposito et al., 2001; Molina-Holgado et al., 2003). Inhibition of iNOS could not account for cannabinoid-induced enhancement of neurogenesis, however, because iNOS is itself proneurogenic, and neurogenesis is defective in iNOS-KO mice (Zhu et al., 2003). Cannabinoids also inhibit nNOS activity. In cultured cerebellar granule cells, CB1R agonists blocked the stimulation of nNOS by K^+ -induced depolarization (Hillard et al., 1999) by reducing voltage-gated Ca^{2+} influx, upon which nNOS activation depends. Δ^9 -THC protected rat retinal neurons from NMDA-induced, NO- and peroxynitrite-mediated toxicity in vivo, partly through interaction with CB1R (El-Remessy et al., 2003). However, cannabidiol, which does not bind to CB1R, was also protective, and the mechanism employed by Δ^9 -THC was not further defined. Cannabinoids and NO also appear to interact in the regulation of body temperature. The hypothermic effect of Δ^9 -THC was abolished in nNOS-KO mice (Azad et al., 2001), and the combination of a cannabinoid agonist [*R*(+)-Win 55212-2] and an NOS inhibitor

(*N*^ω-nitro-L-arginine methyl ester) reduced temperature in a synergistic fashion (Rawls et al., 2004).

These studies, and especially that of Hillard et al. (1999), suggest a plausible mechanism for the coregulation of adult neurogenesis by cannabinoids and NO. CB1R activation inhibits Ca²⁺ influx through neuronal (N- and P/Q-type) voltage-gated Ca²⁺ channels (Twitchell et al., 1997), which uncouples membrane depolarization from nNOS activation. Ca²⁺ signaling appears to be an important determinant of neuronal maturation (Owens and Kriegstein, 1998). However, the route of Ca²⁺ influx exerts a strong influence on its downstream effects (Bading et al., 1993). Thus, although reduced Ca²⁺ influx through N- and P/Q-type channels may explain how cannabinoids promote neurogenesis, proliferation of neuronal precursors in the adult DG is also enhanced by increased Ca²⁺ influx through L-type voltage-gated or NMDA receptor-gated channels (Deisseroth et al., 2004). Further studies will be required to achieve a fuller understanding of the relationships among cannabinoids, signal transduction, and neurogenesis.

References

- Azad SC, Marsicano G, Eberlein I, Putzke J, Zieglerberger W, Spanagel R, and Lutz B (2001) Differential role of the nitric oxide pathway on D⁹-THC-induced central nervous system effects in the mouse. *Eur J Neurosci* **13**:561–568.
- Bading H, Ginty DD, and Greenberg ME (1993) Regulation of gene expression in hippocampal neurons by distinct calcium signaling pathways. *Science (Wash DC)* **260**:181–186.
- Cheng A, Wang S, Cai J, Rao MS, and Mattson MP (2003) Nitric oxide acts in a positive feedback loop with BDNF to regulate neural progenitor cell proliferation and differentiation in the mammalian brain. *Dev Biol* **258**:319–333.
- Curtis MA, Penney EB, Pearson AG, van Roon-Mom WM, Butterworth NJ, Dragunow M, Connor B, and Faull RL (2003) Increased cell proliferation and neurogenesis in the adult human Huntington's disease brain. *Proc Natl Acad Sci USA* **100**:9023–9027.
- Dawson VL, Dawson TM, London ED, Bredt DS, and Snyder SH (1991) Nitric oxide mediates glutamate neurotoxicity in primary cortical cultures. *Proc Natl Acad Sci USA* **88**:6368–6371.
- Deisseroth K, Singla S, Toda H, Monje M, Palmer TD, and Malenka RC (2004) Excitation-neurogenesis coupling in adult neural stem/progenitor cells. *Neuron* **42**:535–552.
- El-Remessy AB, Khalil IE, Matragoon S, Abou-Mohamed G, Tsai NJ, Roon P, Caldwell RB, Caldwell RW, Green K, and Liou GI (2003) Neuroprotective effect of (–)-Δ⁹-tetrahydrocannabinol and cannabidiol in *N*-methyl-D-aspartate-induced retinal neurotoxicity: involvement of peroxynitrite. *Am J Pathol* **163**:1997–2008.
- Esposito G, Izzo AA, Di Rosa M, and Iuvone T (2001) Selective cannabinoid CB1 receptor-mediated inhibition of inducible nitric oxide synthase protein expression in C6 rat glioma cells. *J Neurochem* **78**:835–841.
- Francis F, Koulakoff A, Boucher D, Chafey P, Schaar B, Vinet MC, Friocourt G, McDonnell N, Reiner O, Kahn A, et al. (1999) Doublecortin is a developmentally regulated, microtubule-associated protein expressed in migrating and differentiating neurons. *Neuron* **23**:247–256.
- Gleeson JG, Lin PT, Flanagan LA, and Walsh CA (1999) Doublecortin is a microtubule-associated protein and is expressed widely by migrating neurons. *Neuron* **23**:257–271.
- Hillard CJ, Muthian S, and Kearn CS (1999) Effects of CB(1) cannabinoid receptor activation on cerebellar granule cell nitric oxide synthase activity. *FEBS Lett* **459**:277–281.
- Jeon YJ, Yang KH, Pulaski JT, and Kaminski NE (1996) Attenuation of inducible nitric oxide synthase gene expression by delta 9-tetrahydrocannabinol is mediated through the inhibition of nuclear factor-kappa B/Rel activation. *Mol Pharmacol* **50**:334–341.
- Jiang W, Zhang Y, Xiao L, Van Cleemput J, Ji SP, Bai G, and Zhang X (2005) Cannabinoids promote embryonic and adult hippocampus neurogenesis and produce anxiolytic- and antidepressant-like effects. *J Clin Invest* **115**:3104–3116.
- Jin K, Galvan V, Xie L, Mao XO, Gorostiza OF, Bredesen DE, and Greenberg DA (2004a) Enhanced neurogenesis in Alzheimer's disease transgenic (PDGF-APP^{Sw}, Ind) mice. *Proc Natl Acad Sci USA* **101**:13363–13367.
- Jin K, LaFevre-Bernt M, Sun Y, Chen S, Gafni J, Crippen D, Logvinova A, Ross CA, Greenberg DA, and Ellerby LM (2005) FGF-2 promotes neurogenesis and neuroprotection and prolongs survival in a transgenic mouse model of Huntington's disease. *Proc Natl Acad Sci USA* **102**:18189–18194.
- Jin K, Mao XO, Sun Y, Xie L, Jin L, Nishi E, Klagsbrun M, and Greenberg DA (2002) Heparin-binding epidermal growth factor-like growth factor: hypoxia-inducible expression in vitro and stimulation of neurogenesis in vitro and in vivo. *J Neurosci* **22**:5365–5373.
- Jin K, Minami M, Lan JQ, Mao XO, Bateur S, Simon RP, and Greenberg DA (2001) Neurogenesis in dentate subgranular zone and rostral subventricular zone after focal cerebral ischemia in the rat. *Proc Natl Acad Sci USA* **98**:4710–4715.
- Jin K, Peel AL, Mao XO, Xie L, Cottrell BA, Henshall DC, and Greenberg DA (2004b) Increased hippocampal neurogenesis in Alzheimer's disease. *Proc Natl Acad Sci USA* **101**:343–347.
- Jin K, Xie L, Kim SH, Parmentier-Batteur S, Sun Y, Mao XO, Childs J, and Greenberg DA (2004c) Defective adult neurogenesis in CB1 cannabinoid receptor knockout mice. *Mol Pharmacol* **66**:204–208.
- Kim SH, Won SJ, Mao XO, Jin K, and Greenberg DA (2005) Molecular mechanisms of cannabinoid protection from neuronal excitotoxicity. *Mol Pharmacol* **69**:691–696.
- Ledent C, Valverde O, Cossu G, Petit F, Aubert J-F, Beslot F, Böhme GA, Imperato A, Pedrazzini T, Roques BP, et al. (1999) Unresponsiveness to cannabinoids and reduced addictive effects of opiates in CB1 receptor knockout mice. *Science (Wash DC)* **283**:401–404.
- Matarredona ER, Murillo-Carretero M, Moreno-Lopez B, and Estrada C (2004) Nitric oxide synthesis inhibition increases proliferation of neural precursors isolated from the postnatal mouse subventricular zone. *Brain Res* **995**:274–284.
- Molina-Holgado E, Vela JM, Arevalo-Martin A, Almazan G, Molina-Holgado F, Borrell J, and Guaza C (2002) Cannabinoids promote oligodendrocyte progenitor survival: involvement of cannabinoid receptors and phosphatidylinositol-3 kinase/Akt signaling. *J Neurosci* **22**:9742–9753.
- Molina-Holgado F, Pinteaux E, Moore JD, Molina-Holgado E, Guaza C, Gibson RM, and Rothwell NJ (2003) Endogenous interleukin-1 receptor antagonist mediates anti-inflammatory and neuroprotective actions of cannabinoids in neurons and glia. *J Neurosci* **23**:6470–6474.
- Moreno-Lopez B, Romero-Grimaldi C, Noval JA, Murillo-Carretero M, Matarredona ER, and Estrada C (2004) Nitric oxide is a physiological inhibitor of neurogenesis in the adult mouse subventricular zone and olfactory bulb. *J Neurosci* **24**:85–95.
- Nagayama T, Sinor AD, Simon RP, Chen J, Graham SH, Jin K, and Greenberg DA (1999) Cannabinoids and neuroprotection in global and focal cerebral ischemia and in neuronal cultures. *J Neurosci* **19**:2987–2995.
- Owens DF and Kriegstein AR (1998) Patterns of intracellular calcium fluctuation in precursor cells of the neocortical ventricular zone. *J Neurosci* **18**:5374–5388.
- Packer MA, Stasiv Y, Benraiss A, Chmielnicki E, Grinberg A, Westphal H, Goldman SA, and Enikolopov G (2003) Nitric oxide negatively regulates mammalian adult neurogenesis. *Proc Natl Acad Sci USA* **100**:9566–9571.
- Parmentier-Batteur S, Jin K, Mao XO, Xie L, and Greenberg DA (2002) Increased severity of stroke in CB1 cannabinoid receptor knock-out mice. *J Neurosci* **22**:9771–9775.
- Rawls SM, Tallarida RJ, Gray AM, Geller EB, and Adler MW (2004) L-NAME (*N*^ω-nitro-L-arginine methyl ester), a nitric-oxide synthase inhibitor, and WIN 55212-2 [4,5-dihydro-2-methyl-4(4-morpholinylmethyl)-1-(1-naphthalenyl-carbonyl)-6H-pyrrolo[3,2,1*ij*]quinolin-6-one], a cannabinoid agonist, interact to evoke synergistic hypothermia. *J Pharmacol Exp Ther* **308**:780–786.
- Reif A, Schmitz A, Fritzen S, Chourbaji S, Bartsch C, Urani A, Wycislo M, Mossner R, Sommer C, Gass P, et al. (2004) Differential effect of endothelial nitric oxide synthase (NOS-III) on the regulation of adult neurogenesis and behaviour. *Eur J Neurosci* **20**:885–895.
- Rueda D, Navarro B, Martinez-Serrano A, Guzman M, and Galve-Roperh I (2002) The endocannabinoid anandamide inhibits neuronal progenitor cell differentiation through attenuation of the Rap1/B-Raf/ERK pathway. *J Biol Chem* **277**:46645–46650.
- Shen M, Piser TM, Seybold VS, and Thayer SA (1996) Cannabinoid receptor agonists inhibit glutamatergic synaptic transmission in rat hippocampal cultures. *J Neurosci* **16**:4322–4334.
- Shen M and Thayer SA (1998) Cannabinoid receptor agonists protect cultured rat hippocampal neurons from excitotoxicity. *Mol Pharmacol* **54**:459–462.
- Southan GJ and Szabo C (1996) Selective pharmacological inhibition of distinct nitric oxide synthase isoforms. *Biochem Pharmacol* **51**:383–394.
- Sun Y, Jin K, Childs JT, Xie L, Mao XO, and Greenberg DA (2005) Neuronal nitric oxide synthase and ischemia-induced neurogenesis. *J Cereb Blood Flow Metab* **25**:485–492.
- Twitchell W, Brown S, and Mackie K (1997) Cannabinoids inhibit N- and P/Q-type calcium channels in cultured rat hippocampal neurons. *J Neurophysiol* **78**:43–50.
- Waksman Y, Olson JM, Carlisle SJ, and Cabral GA (1999) The central cannabinoid receptor (CB1) mediates inhibition of nitric oxide production by rat microglial cells. *J Pharmacol Exp Ther* **288**:1357–1366.
- Zhang R, Zhang L, Zhang Z, Wang Y, Lu M, Lapointe M, and Chopp M (2001) A nitric oxide donor induces neurogenesis and reduces functional deficits after stroke in rats. *Ann Neurol* **50**:602–611.
- Zhu DY, Liu SH, Sun HS, and Lu YM (2003) Expression of inducible nitric oxide synthase after focal cerebral ischemia stimulates neurogenesis in the adult rodent dentate gyrus. *J Neurosci* **23**:223–229.

Address correspondence to: Dr. David A. Greenberg, Buck Institute for Age Research, 8001 Redwood Boulevard, Novato, CA 94945. E-mail: dgreenberg@buckinstitute.org