Protection from Alzheimer’s-like disease in the mouse by genetic ablation of inducible nitric oxide synthase

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Brains from subjects who have Alzheimer’s disease (AD) express inducible nitric oxide synthase (iNOS). We tested the hypothesis that iNOS contributes to AD pathogenesis. Immunoreactive iNOS was detected in brains of mice with AD-like disease resulting from transgenic expression of mutant human β-amyloid precursor protein (hAPP) and presenilin-1 (hPS1). We bred hAPP−, hPS1−double transgenic mice to be iNOS+/H11545/H11408/H11545 or iNOS−/−, and compared them with a congenic WT strain. Deficiency of iNOS substantially protected the AD-like mice from premature mortality, cerebral plaque formation, increased β-amyloid levels, protein tyrosine nitration, astrocytosis, and microgliosis. Thus, iNOS seems to be a major instigator of β-amyloid deposition and disease progression. Inhibition of iNOS may be a therapeutic option in AD.

1996, iNOS was identified in AD lesions (10). Collectively, that report and seven confirmatory studies identified iNOS immunoreactivity in neurons (7, 8, 10–12) and astrocytes (7, 12–14) in brains of 75 patients who had AD, but at far lower incidence, extent, and intensity in brains from age-matched controls. Although NOS1 (neuronal NOS) and NOS3 (endothelial NOS) are expressed constitutively in normal brain, widespread expression of iNOS in the central nervous system is pathologic, and has been observed in multiple sclerosis (15), HIV-associated dementia (16), stroke (17), amyotrophic lateral sclerosis (18), and Parkinson’s disease (19). Because of its independence of elevated intracellular Ca2+ (20), iNOS catalyzes a high-output pathway of NO production (21) that is capable of causing neuronal damage and death (16). Multiple mechanisms of NO-dependent cytotoxicity have been identified. For example, inhibition of the mitochondrial electron transport chain by RNI (22) increases mitochondrial production of superoxide anion, and spurs formation of peroxynitrite, which can exacerbate mitochondrial damage (23). RNI can inhibit proteasomal degradation of protein tyrosine residues (4–9), which can report the vicinal production of peroxynitrite from NO and superoxide. However, a functionally important source of AD-associated oxidative or nitrosative brain injury has not been identified that is a plausible target for pharmacologic inhibition.

The online version of this article contains supplemental material.
pathways (24), and perhaps contribute to the markedly decreased proteasome function that was documented in affected regions of brains from patients who had AD (25). Decreased proteasome function is likely to promote further accumulation of Aβ and advanced glycation products, which leads to increased induction of iNOS (13). Based on these observations, we hypothesized that inhibition of iNOS might slow the progression of neuronal damage in individuals in whom levels of intracerebral Aβ are elevated. To test the foregoing hypothesis, we used a genetic approach by breeding disrupted iNOS alleles into mice transgenic for mutant human genes associated with AD.

RESULTS AND DISCUSSION

Construction of strains

First, we backcrossed the original iNOS−/− C57BL/6×129 mice (26) to C57BL/6 for six generations. Descendants of brother–sister matings of the latter mice were crossed with the SJL strain, and their progeny were interbred to derive iNOS−/− C57BL/6×SJL mice. This step was based on evidence that that modifier genes from the SJL background were critical to avoid premature mortality in C57BL/6 mice bearing a mutant (K670N, M671L) human APP (hAPP) transgene (27). We bred the iNOS−/− C57BL/6×SJL mice with a strain called Tg2576, in which a hamster prion protein promoters was critical to avoid premature mortality in C57BL/6 mice bearing a mutant (K670N, M671L) human APP (hAPP) transgene (27).

Comparison of the fate of the three congenic sublines allowed us to test the hypothesis that iNOS exacerbates AD. Mice with the AD-related transgenes and WT iNOS alleles died much earlier than did the WT congenic subline. Deficiency of iNOS exerted a marked protective effect against AD-related mortality, and extended the time at which 75% of the cohort remained alive from 143 d for iNOS+/+ hAPP+/+ hPS1+/+ mice to 315 d for iNOS−/− hAPP+/+ hPS1+/+ mice—a 220% increase (P < 0.0001, logrank test) (Fig. 2 A). Even so, the iNOS−/− hAPP+/+ hPS1+/+ mice still died sooner than did congenic WT mice.

Male (Fig. 2 B) and female (Fig. 2 C) AD-transgenic mice gained weight more slowly than did the WT mice (P < 0.0001, repeated measures ANOVA). However, iNOS alleles had no effect on weight gain, and therefore, iNOS was unlikely to hasten death by impairing feeding or drinking. Moreover, time-lapse video camera recordings every 15 or 30 s over 24-h periods, covering 327 h of life of individual mice, were scored by an observer who was blind to mouse genotype. The video recordings revealed no differences in feeding or other behavior between AD-transgenic mice with and without intact iNOS alleles. Both strains tended to sleep less (4.5 ± 0.9 and 5.4 ± 1.3 episodes per day totaling 519 ± 50 and 581 ± 115 min, respectively) than did WT mice (8.8 ± 1.3 episodes totaling 624 ± 21 min; means ± SD) and both rarely hung from the cage lid, in contrast to WT mice. Necropsies of nine AD-transgenic mice gained weight more slowly than did the WT mice (P < 0.0001, repeated measures ANOVA). However, iNOS alleles had no effect on weight gain, and therefore, iNOS was unlikely to hasten death by impairing feeding or drinking. Moreover, time-lapse video camera recordings every 15 or 30 s over 24-h periods, covering 327 h of life of individually housed mice aged 3–5 mo, were scored by an observer who was blind to mouse genotype. The video recordings revealed no differences in feeding or other behavior between AD-transgenic mice with and without intact iNOS alleles. Both strains tended to sleep less (4.5 ± 0.9 and 5.4 ± 1.3 episodes per day totaling 519 ± 50 and 581 ± 115 min, respectively) than did WT mice (8.8 ± 1.3 episodes totaling 624 ± 21 min; means ± SD) and both rarely hung from the cage lid, in contrast to WT mice. Necropsies of nine AD-transgenic mice with or without intact iNOS alleles revealed a distinct extracerebral pathology in each of four mice that was judged to be incidental (see supplemental material). Thus, the cause of premature mortality in the iNOS−/− hAPP+/+ hPS1+/+ strain was not established.

Impact of iNOS on AD transgene-dependent brain pathology

Cohorts of the three sublines were aged for 20, 40, and 60 wk. Brains from 10 mice per subline and time point were examined for histopathology. No abnormalities were observed in the WT controls. The double-transgenic mice had age-dependent accumulation of plaques in all areas examined: the cingulate, retrosplenial, and motor cortices and the hippocampus. Deficiency of iNOS had

Figure 1. Expression of iNOS in brains of 60- to 64-wk-old mice with or without AD-related transgenes, as judged by immunoblot.

Each lane is from a separate mouse of the genotypes indicated. Tubulin was immunostained as a loading control.
Figure 2. Amelioration of early mortality of AD-transgenic mice by disruption of iNOS alleles without an impact on weight gain. (top, Survival) Individually genotyped mice (iNOS<sup>+/+</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup>, n = 133; iNOS<sup>+/+</sup> hAPP<sup>+/+</sup> hPS1<sup>1/0</sup>; n = 140; iNOS<sup>−/−</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup>, n = 112) were inspected 5 d per week throughout their lifetime, and their mortality was plotted as a function of age. Differences between any two curves were significant (P < 0.0001; logrank test). (middle) Weight gain in males (iNOS<sup>+/+</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup>, n = 43; iNOS<sup>+/+</sup> hAPP<sup>+/+</sup> hPS1<sup>1/0</sup>; n = 67; iNOS<sup>−/−</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup>, n = 48). (bottom) Weight gain in females (iNOS<sup>+/+</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup>, n = 41; iNOS<sup>+/+</sup> hAPP<sup>+/+</sup> hPS1<sup>1/0</sup>, n = 50; iNOS<sup>−/−</sup> hAPP<sup>+/+</sup> hPS1<sup>1/0</sup>, n = 50).

no effect on the accumulation of Aβ during the first 40 wk of life, but markedly inhibited further accumulation thereafter, so that plaque burden was reduced by 61% by week 60 (Fig. 3, A–E). Immunoblot (Fig. 3, F and G) and ELISA (Fig. 3 H) for Aβ were consistent with the immunohistologic findings, in immunoblot, Aβ signal intensity was reduced by 64% in the iNOS-deficient brains (P = 0.012, Student’s t test), whereas in ELISA, Aβ reactivity was diminished by 43% (P = 0.04).

Figure 3. Amelioration of late-stage plaque formation and Aβ deposition in AD-transgenic mice by disruption of iNOS alleles. (A) Representative sections from the neocortex and hippocampus of iNOS<sup>+/+</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup> and iNOS<sup>−/−</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup> mice immunostained with Aβ [1–42] antibody. (B–E) Plaque burden (percentage of area occupied by Aβ [1–42]-immunoreactive plaques) and plaque numbers per unit area as a function of age in cortex (B and C) and hippocampus (D and E). Means ± SEM for 10 mice per strain per time point. Statistically significant differences are marked (P < 0.05; **P < 0.01; ***P < 0.003). No immunoreactivity was detected in mice lacking AD transgenes. Solid green line, iNOS<sup>+/+</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup> mice. Dashed orange line, iNOS<sup>−/−</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup> mice. (F) Representative immunoblot for brain Aβ that was not extractable in physiologic saline or 0.5% Triton X-100, but was soluble in 6% SDS. Lane 1: iNOS<sup>−/−</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup> mice; lane 2: iNOS<sup>−/−</sup> hAPP<sup>+/+</sup> hPS1<sup>1/0</sup> mice; lane 3: iNOS<sup>+/+</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup> mice. (G) Densitometry of four blots like that in F, each for different sets of mice, normalized to β-actin. The x axis sets to 1 the ratio of signal intensity for Aβ to that for β-actin in iNOS<sup>−/−</sup> hAPP<sup>+/+</sup> hPS1<sup>+/0</sup> mice (solid bar marked “+++”); the corresponding ratios for iNOS<sup>−/−</sup> hAPP<sup>+/</sup> hPS1<sup>+/0</sup> mice (hatched bar marked “−−−”) are given as a proportion of the ratio for the “+++” in each of the same four blots (mean ± SEM). (H) ELISA for Aβ in extracts prepared as in F. Aβ burden is indicated in µg/mg brain weight (mean ± SEM, n = 6) for the two strains with AD-related transgenes whose iNOS alleles are indicated as “++/+” (intact iNOS alleles, black bar) or “−−/−−” (disrupted iNOS alleles, gray bar).
Deficiency of iNOS also reduced the extent of protein tyrosine nitration in the brains of the AD-transgenic mice markedly, as shown by immunostaining (Fig. 4 A) and slot-blot (Fig. 4 B) using two different antibodies. Further, iNOS deficiency afforded substantial protection against gliosis, as judged by glial fibrillary acidic protein (GFAP) staining for reactive astrocytes and CD40 staining for activated microglia (Fig. 4, B–E; Fig. S2, available at http://www.jem.org/cgi/content/full/jem.20051529/DC1). Finally, deficiency of iNOS led to a reduction in accumulation of phosphorylated tau protein around plaques as judged by immunostaining (Fig. 4 G) and immunoblot (Fig. 4 H). By densitometric analysis of immunoblots, the ratio of phospho-tau staining/tubulin staining used as a loading control averaged 0.7 ± 0.1 for WT mice, 1.2 ± 0.6 for mice with AD-related transgenes and disrupted iNOS alleles, and 2.7 ± 0.0 for mice with AD-related transgenes and intact iNOS alleles. Thus, iNOS deficiency seemed to reduce tau phosphorylation by 56%.

In summary, iNOS, the catalyst of high-output pathway of NO production (21), is expressed in human AD and in many mouse models of AD. Mice with AD-like disease that were unable to express iNOS genetically lived longer than did their iNOS-expressing counterparts, formed fewer plaques, had lower levels of brain Aβ, suffered less protein tyrosine nitration, accumulated less phosphorylated tau protein, and harbored fewer reactive astrocytes and microglia.

Figure 4. Amelioration of late-stage nitrosative/oxidative injury, astrogliosis, microgliosis, and phospho-tau accumulation in AD-transgenic mice by disruption of iNOS alleles. (A) Nitrotyrosine immunoreactivity in the cingulate cortex of iNOS+/+ hAPP0/0 hPS10/0, iNOS+/+ hAPP+/+ hPS11/0, and iNOS−/− hAPP+/+ hPS11/0 mice. (B) Nitrotyrosine immunoblot. Brain extract proteins (150 μg) were immobilized on a filter with a slot-blot apparatus and were immunoblotted with anti-nitrotyrosine mAb. (C–F) Reduction of GFAP staining indicative of astrocytosis (C and E) and of CD40 staining indicative of microgliosis (D and F) in the cortex (C and D) and hippocampus (E and F) of iNOS−/− hAPP+/+ hPS11/0 mice. Statistically significant differences are marked (***P < 0.001; *P < 0.05). (G) Reduction of phospho-tau immunoreactivity around plaques in iNOS−/− hAPP+/+ hPS11/0 mice. Arrows highlight positive staining. (H) Anti-phospho-tau (p-tau) immunoblot with anti-tubulin as a loading control. Each lane is from a separate mouse of the genotypes indicated. See text for quantitative analysis.
It was shown that fibrillogenic Aβ promotes expression of iNOS (1). However, the striking protective effect of iNOS deficiency on plaque formation and accumulation of Aβ deposits now suggests that iNOS is a major factor that furthers the accumulation of Aβ. Thus, iNOS and fibrillogenic Aβ each seem to promote the other’s accumulation. Selective iNOS inhibitors that can enter the brain should be tested for their ability to slow the progression of AD-like disease in mice.

MATERIALS AND METHODS

Genotyping. Mice were bred and studied according to institutionally approved protocols. The strains that were constructed for this study have been deposited with the Mutant Mouse Regional Resource Centers for distribution to other investigators (http://www.mmrrc.org/catalog/StrainCatalogSearchForm.jsp). All 3,491 mice were genotyped by PCR before adulthood. Those of the three genotypes that were included in this study were genotyped again by PCR; 1 mouse failed to confirm and was excluded from analysis. The specificity of the PCR reactions was confirmed by Southern blot.

For iNOS: 5′-GTGGAACGCTTGGCTGAATTCTGTTAT-3′ (H11032) and 5′-GTGGATAACCCCTCCCCCAGCCTAGACCA-3′ (H11601). For hPS1 transgene, an 80-bp fragment is generated by the following primers: 5′-GGCTTTCTGTCTGTTCTCTC-3′ and 5′-GTCCTTGGGGTCTTCTACCTTTCTC-3′. For disrupted iNOS alleles, the 3′ primer for the mutant allele is homologous to sequences in the neomycin gene and with NOS-A will generate a 268-bp fragment: 5′-ATCAGCCTTTCTCTCTCTCTG-3′ (H11032). For iNOS-B, the common primer 5′-GTGGATAACCCCTCCCCCAGCCTAGACCA-3′ (H11601) and 5′-GTGGTG-3′ (H11032) is generated by the following primers (32): 5′-GGCTTTCTGTCTGTTCTCTC-3′ and 5′-GTCCTTGGGGTCTTCTACCTTTCTC-3′. For WT iNOS alleles, the common 5′ primer is GTCCTTGGCTGAATTCTGTTAT-3′ (H11032). For hAPP transgene, a 300-bp fragment is generated by the following primers (32): 5′-GTGGATAACCCCTCCCCCAGCCTAGACCA-3′ and 5′-GTGGTG-3′ (H11032). For the hPS1 transgene, a 300-bp fragment was restricted with KpnI for iNOS and with BamHI (H11601). The primer pair for the mutant allele is homologous to sequences in the neomycin gene and with NOS-A will generate a 413-bp fragment: 5′-GGCTTTCTGTCTGTTCTCTC-3′ and 5′-GTCCTTGGGGTCTTCTACCTTTCTC-3′. For immunohistology, rabbit anti-iNOS antiserum (anti-iNOS) was restricted with KpnI for iNOS and with BamHI (H11601). The primer pair for WT iNOS (NOS-B) corresponds to bp 337–356 and with NOS-A will generate a 413-bp fragment: 5′-GGCTTTCTGTCTGTTCTCTC-3′ and 5′-GTCCTTGGGGTCTTCTACCTTTCTC-3′. For disrupted iNOS alleles, the 3′ primer for the mutant allele is homologous to sequences in the neomycin gene and with NOS-A will generate a 268-bp fragment: 5′-GGCTTTCTGTCTGTTCTCTC-3′ and 5′-GTCCTTGGGGTCTTCTACCTTTCTC-3′. For iNOS deficiency on plaque formation and accumulation of Aβ deposits now suggests that iNOS is a major factor that furthers the accumulation of Aβ.

Preparation of brains. For immunohistology, mice were anesthetized deeply with pentobarbital and perfused intracardially with ice-cold 0.9% saline followed by ice-cold 4% paraformaldehyde in 0.1 M sodium phosphate buffer, pH 7.2 (PB). Brains were fixed <24 h in 10% neutral buffered formalin, washed in PB, and hemisected sagitally. One hemisphere was cryoprotected in 30% glycerol, 30% ethylene glycol/0.02 M PB and stored at −20°C for preparation of 35 micron frozen sections. The other hemisphere was fixed in 10% neutral buffered formalin for <24 h, transferred to 70% ethanol, and paraffin embedded by standard methods. For Western blots, the perfusion omitted paraformaldehyde and the brains were placed directly in the cryoprotectant solution.

To prepare brain extracts for anti-iNOS and anti-nitrotyrosine immunoblots, 150 ml 0.9% NaCl was used for transcardial perfusion without fixative. Brain quarters were boiled for 10 min in 120 μl sample buffer (200 mM Tris-HCl, pH 6.8, 4% SDS, 4% β-mercaptoethanol, and 20% glycerol), followed by centrifugation at 14,000 g for 30 min, and supernatant was stored at −80°C for immunoblot. Protein concentration was determined by BioRad assay.

Antibodies. For immunohistology, rabbit anti-iNOS antiserum (anti-holoiNOS) was raised against pure, native mouse iNOS (21), and subjected to extensive documentation of monospecificity after ultracentrifugation. These tests included, as a positive control, reactivity with cells in sections of footpads from WT mice infected with Leishmania major (gift from R. Almeida, Federal University of Bahia, Bahia, Brazil) and livers from WT mice infected with Propionibacterium acnes followed 6 d later by bacterial lipopolysaccharide.

As negative controls, we used livers from untreated WT mice and livers from P. acnes- and LPS-treated iNOS knock-out mice. The antiserum was positive with the controls, and negative with the controls, respectively. For immunohistology, the positive samples were negative using nonspecific rabbit IgG in place of anti-iNOS or omitting primary antibody. Moreover, preimmune rabbit serum gave no reaction with sections of brains from AD-transgenic mice expressing iNOS. For further specificity controls, antibody to mouse NOS1 (Santa Cruz K-20) was used to document reaction with NOS1 in cerebellum of WT mice under the same conditions where the anti-iNOS antiserum was negative with cerebellum. Antibody to mouse NOS3 (Santa Cruz N-20) was used to document reaction with NOS3 in heart of WT mice under the same conditions where the anti-iNOS antiserum was negative with heart.

Finally, all results with immunohistochemistry were confirmed by immunoblot. Thus, anti-iNOS antiserum was monospecific for iNOS in mouse, and did not react with mouse NOS1, mouse NOS3, or any other detectable moieties in inflamed organs of iNOS knock-out mice. This anti-iNOS antiserum was furnished to Upstate Biotechnology. Equivalent results were obtained using the original antiserum and samples that we subsequently purchased from Upstate Biotechnology. We have reported similar documentation of specificity for anti-iNOS mAbs 1E8B8 and 5BE36 (Research & Diagnostic Antibodies) (33). For Western blot, we used rabbit anti-iNOS amino terminal domain (Upstate Biotechnology).

Immunohistochemistry. Specimens were provided by M. Sporn (Dartmouth College, Hanover, NH); K. Hsiao (University of Minnesota, Minneapolis, MN); J. Clemens (E. Lilly, Inc., Indianapolis, IN); P. Davies and L. Van Der Ploeg (Merck, Inc., Rahway, NJ); K. Duff and Y. Masugi (New York University, New York, NY); and S. Fu and J. Merrill (Aventis, Inc., Bridgewater, NJ). Staining for iNOS was performed on paraffin-embedded sections. Following antibodies also were used: rabbit anti-GFAP (1:1,000; DakoCytomation), rat anti–mouse CD40 (1:100; Serotech), rabbit antinitrotyrosine (1:50; Upstate Biotechnology), and rabbit polyclonal antibody against tau (phospho T205; 1:100; Abcam Inc.). Antinitrotyrosine slot-blots used mAb 1A6 (0.5 μg/ml) from Upstate Biotechnology.

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Video camera recordings. Mice were housed individually in cages whose water bottles were replaced with gel packs located under the food bin for improved visibility from above. The mice were kept in the standard dark/light cycle, except that a very faint light for the dark cycle was supplied by way of reflection from a 10-W shielded bulb pointed at the wall. Recordings were made with an iSight camera (Apple Computer Inc.) connected to a G4 PowerMacintosh (Apple Computer Inc.). The sensitivity of the camera was increased ~10-fold during the dark cycle. In the first period of recordings, images were taken at 30-s intervals. Subsequently, the frequency was increased to 15-s intervals. Images were collected by way of EvoCam 3.5 software (Evololical Inc., www.evological.com). Recordings were analyzed by one observer, who was blind to the strain of the mice. The observer looked for seizures (none was detected); logged periods of sleep; and characterized motor activity, such as ambulation and hanging from the cage lid. Sleep was defined as an interval of 8 min or longer during which the mouse did not relocate its center of mass and was not engaged in feeding. An interval of sleep was counted as one episode of sleep if it included no more than one excursion lasting <1 min that was preceded and followed by at least 8 min of immobility.

Online supplemental material. Fig. S1 provides immunohistologic evidence for expression of iNOS in brains of mice with or without AD-related transgenes. Fig. S2 illustrates CD40 staining in the cortex. Online supplemental material is available at http://www.jem.org/cgi/content/full/jem.20051529/DC1.

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