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Therapeutic benefits of a component of coffee in a rat model of Alzheimer disease¹

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Abstract

A minor component of coffee unrelated to caffeine, eicosanoyl-5-hydroxytryptamide (EHT) provides protection in a rat model for Alzheimer's disease (AD). In this model, viral expression of the phosphoprotein phosphatase 2A (PP2A) endogenous inhibitor, the I₂^{PP2A} or SET protein in the brains of rats leads to several characteristic features of AD including cognitive impairment, tau hyperphosphorylation, and elevated levels of cytoplasmic β-amyloid protein. Dietary supplementation with EHT for 6–12 months resulted in substantial amelioration of all of these defects. The beneficial effects of EHT could be associated with its ability to increase PP2A activity by inhibiting the demethylation of its catalytic subunit PP2Ac. These findings raise the possibility that EHT may make a substantial contribution to the apparent neuroprotective benefits associated with coffee consumption as evidenced by numerous epidemiological studies indicating the coffee drinkers have substantially lowered risk of developing AD.

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¹This paper is dedicated to Dr. Inge Grundke-Iqbal who supervised most of the immunohistochemical and biochemical studies before she passed away on September 22, 2012.

Disclosure Statement

Conflict of interest disclosure: Signum Biosciences has a U.S. patent on the EHT compound used for this study.

The animal studies conform to NIH guidelines and were approved by our institutional IACUC.

Keywords

Protein phosphatase-2A; Tau; A β ; Hyperphosphorylation of tau; Rat model of sporadic Alzheimer disease; coffee; eicosanoyl-5-hydroxytryptamide; cognitive impairment; methylation of protein phosphatase-2A; adeno-associated virus vector serotype 1; inhibitor-2 of protein phosphatase-2A; SET

1. INTRODUCTION

Alzheimer's disease (AD), the major cause of dementia in middle and old age, is characterized by neurodegeneration that is associated with neurofibrillary tangles and neuritic plaques. A current major goal in medicine is the development of disease modifying therapeutic drugs for AD. The microtubule associated, tau protein is abnormally hyperphosphorylated in AD where it is the principle component of neurofibrillary tangles (Grundke-Iqbal et al. 1986a; Grundke-Iqbal et al. 1986b; Bancher et al. 1989). Similarly, the β -amyloid polypeptide, A β , is the principle component of neuritic plaques (Masters et al. 1985; Wong et al. 1985). Evidence suggests that tangle and plaque precursors, the non-fibril forms of abnormally hyperphosphorylated tau and soluble oligomers of A β are the major cytotoxic species in AD (Iqbal et al. 1986; Grundke-Iqbal et al. 1989; Kopke et al. 1993; Alonso et al. 1994; Klein 2002; Santacruz et al. 2005; Alonso et al. 2010). As much as 40% of the abnormally hyperphosphorylated tau is cytosolic in AD brain (Kopke et al. 1993), and intraneuronal A β accumulation precedes plaque deposition (Bancher et al. 1989; Grundke-Iqbal et al. 1989; Mori et al. 2002; Cataldo et al. 2004), and is correlated with neuronal cell death in AD transgenic mouse models (Oddo et al. 2003; Espana et al. 2010; Gandy et al. 2010).

Phosphoprotein phosphatase 2A (PP2A) accounts for approximately 70% of the total phospho-tau phosphatase activity in healthy human brain (Gong et al. 1993; Gong et al. 1995; Benneceib et al. 2000; Gong et al. 2000; Liu et al. 2005) and also functions to dephosphorylate the β -amyloid precursor protein so as to reduce the formation of the A β (Sontag et al. 2007). In AD, PP2A activity is curtailed so that levels of hyperphosphorylated tau and A β increase, leading to neurodegeneration and dementia (Gong et al. 1993). Two cellular inhibitor proteins, I₁^{PP2A} and I₂^{PP2A}, regulate the activity of PP2A (Li et al. 1995; Li et al. 1996). I₂^{PP2A} or SET inhibits PP2A activity towards hyperphosphorylated tau (Tsuji et al. 2005). I₂^{PP2A} is a 277 amino acid long nuclear protein that is overexpressed and selectively cleaved at N175 into N-terminal (I₂^{N_{TF}}) and C-terminal (I₂^{C_{TF}}) fragments, which are translocated from the neuronal nucleus to the cytoplasm in affected areas of the AD brain (Tanimukai et al. 2005). Both I₂^{N_{TF}} and I₂^{C_{TF}} bind to PP2Ac and inhibit its phosphatase activity towards hyperphosphorylated tau (Arnaud et al. 2011). Transduction of brains of new born rat pups with Adeno Associated Virus serotype 1 (AAV1) encoding I₂^{N_{TF}} and I₂^{C_{TF}} inhibit PP2A activity and cause abnormal hyperphosphorylation and aggregation of tau and accumulation of intraneuronal A β in 13 month old animals (Bolognin et al. 2012). These protein hallmarks of AD are associated with cognitive defects in memory and learning.

From these results, it seems likely that a therapeutic agent that acts to maintain healthy levels of PP2A might provide a disease modifying approach for the treatment of AD. A number of different PP2A-activating compounds and mechanisms have been identified (Vorontsov et al. 2011). An in vitro screen was conducted for natural products that support PP2A activity toward phospho-Tau, and a suitable activity was identified in coffee extracts. The active agent was purified to homogeneity and identified as eicosanoyl-5-hydroxytryptamide (EHT). Synthetic EHT exhibited the same ability to support PP2A activity as EHT isolated from coffee. Dietary supplementation with synthetic EHT exhibited neuroprotective efficacies in mouse models for Parkinson's disease (Lee et al. 2011; Lee et al. 2013). Here we report the beneficial effects of chronic dietary supplementation with EHT on PP2A activity, abnormal tau hyperphosphorylation, accumulation of intraneuronal A β and cognitive performance in a rat model for AD.

2. MATERIALS AND METHODS

2.1 Study design

Adeno-associated virus 1 vector (AAV1) was used to express the I $_2$ ^{PP2A} C-terminal and N-terminal fragments (I $_2$ -N/C) in rat brain to replicate the cleavage of I $_2$ ^{PP2A} found previously in AD brains (Tanimukai et al. 2005). AAV1-I $_2$ -N/C infected rats express the predicted I $_2$ -N/C fragments, are cognitively impaired, show hyperphosphorylation and aggregation of tau, and accumulate intraneuronal A β (Bolognin et al. 2012). As described previously (Bolognin et al. 2012), on the day of birth (p 0.5), Wistar rat pups were transfected by injecting 2 μ l containing 4×10^9 AAV1 genomic equivalents encoding I $_2$ -N/C or, as a control, green fluorescent protein (GFP) into each lateral ventricle of the brain (Figure 1). Successful transfection of rat brains with I $_2$ -N/C was confirmed by RT-PCR. At 21 days of age, the pups were weaned and 15 female AAV1- I $_2$ -N/C and 15 female AAV1-GFP rats were put on 0.1% (w/w) EHT formulated diet (Research Diets; New Brunswick, NJ). As controls, 15 female AAV1- I $_2$ -N/C and 15 female AAV1-GFP pups were put on similar diets lacking EHT. The rats were housed and bred according to the PHS Policy on Human Care and Use of Laboratory animals with 2–3 animals per cage, a 12:12-h light/dark cycle and *ad libitum* access to food and water. Studies on animals were carried out according to protocols approved by the Animal Welfare Committee of the New York State Institute for Basic Research. Rats were subjected to behavioral tests at 6 months of age. Seven animals from each group were perfused after behavioral tests, and the remaining eight animals from each group were perfused after a second set of behavioral tests at 12 months while they were still on EHT or vehicle diet. Animals were anesthetized with sodium pentobarbital (125 mg/kg) and then sacrificed by transcardial perfusion with 0.1 M phosphate buffered saline (PBS). The left hemispheres were dissected into hippocampus, cerebral cortex and subcortical structures and kept at -80°C for biochemical analysis, while for immunohistochemical investigation the right hemisphere was immerse-fixed for 48 h in 4% paraformaldehyde in PBS, then cryoprotected in 30% sucrose and cut in 40 μ m sagittal sections using a freezing-sliding microtome.

2.2 Recombinant plasmid production and vector packing

AAV1- I₂-N/C and AAV1-GFP were generated as described previously (Wang et al. 2010; Bolognin et al. 2012). Briefly the plasmid pEGFP-N3/I₂^{PP2A} (Tsujiro et al. 2005) was used as a template to generate by PCR I₂^{PP2A} C- and N-terminal fragment encoding cDNAs. After verification by DNA sequencing, the cDNA fragments were cloned into the multicloning site of the AAV viral genome containing plasmid pTRUF12. Expression was driven by the CMV promoter/enhancer. Serotype 1 virus was produced (Henckaerts et al. 2009) and titers were calculated from standard curves generated from pTRUF12 as previously described (Zolotukhin et al. 2002).

2.3 Western blots

Rat hippocampus was homogenized to 10% (w/v) final concentration in cold buffer containing 50 mM Tris-HCl (pH 7.4), 8.5% sucrose, 2 mM EDTA, 2 mM EGTA, 10 mM β-mercaptoethanol, 5 mM benzamidine, 0.5 mM AEBSF, 4 μg/ml pepstatin A, 10 μg/ml each of aprotinin and leupeptin, 20 mM β-glycerolphosphate, 100 mM sodium fluoride, 1 mM sodium vanadate and 100 nM okadaic acid. Protein concentrations were determined by the modified Lowry method (Bensadoun and Weinstein 1976). Tissue homogenates were heated in Laemmli's buffer and subjected to SDS-PAGE. Proteins were transferred to PVDF membrane of 0.45 μm pore size and membranes were blocked with 5% non-fat dry milk. The following primary antibodies were used: anti-GAPDH (1:2000; Santa Cruz Biotechnology, Santa Cruz, CA, USA), 92e to total tau (1:5000; (Grundke-Iqbal et al. 1988), phosphospecific tau antibodies: tau pS199, tau pT205, tau pT212, tau pS214, tau pS396 (1:1000, BioSource, Camarillo, CA, USA), R145 to tau pS422 (1:3000; (Tanaka et al. 1998), 12E8 to tau pSer396/Ser404 (1:500, (Seubert et al. 1995), 4D9 to methyl-PP2A (1:200; Princeton University, NJ, USA), 1D6 to unmethylated PP2A (Millipore), and 6A3 to total PP2A (Millipore). Immunoblots were probed with the corresponding anti-mouse or anti-rabbit-HRP secondary antibodies (1:5000; Jackson Immuno Research, West Grove, PA, USA) and detected using enhanced chemiluminescence reagents (Thermo Scientific, Rockford, IL, USA). Multi-Gauge V3 software (Fuji Photo Film, Tokyo, Japan) was used to quantify the density of the protein bands in Western blots. The quantified values were statistically analyzed with the non-parametric t-test.

2.4 Reverse-transcription PCR (RT-PCR) and quantitative PCR (qPCR)

Total RNA was extracted from cerebral cortex, with RNeasy plus mini kit (Qiagen, Valencia, CA, USA) according to manufacturer's instructions. cDNA synthesis was performed using Super Script First-Strand Synthesis System kit (Invitrogen, Carlsbad, CA, USA). rt-PCR amplification was achieved in a thermocycler for 30 cycles: denaturation for 30 s at 95°C, annealing for 30 s at 60°C, polymerization for 30 s at 72°C. The I₂^{PP2A} N-terminal-FLAG primer sequence was the following: forward 5'gcaagaagcgattgaacaca-3' and reverse 5'-gcagtgccctctcatctcc-3'. The amplification products were resolved on 1% agarose gels and quantified using the Molecular Imager System (Bio-Rad, Hercules, CA, USA).

Immunohistochemistry (IHC)—IHC was performed on free-floating cryostat sagittal sections of right brain hemispheres. The following antibodies were used at the indicated

dilution: anti-A β_{1-40} (1:200; Invitrogen, Camarillo, CA, USA), Alexa 555-conjugated goat anti-mouse and goat antirabbit IgG (H+L) (1:500; Molecular Probes, Carlsbad, CA, USA) were used as secondary antibodies. Sections were analyzed using confocal microscope Nikon eclipse 90i (Nikon, Melville, NY, USA). For quantitative analysis the images were taken using 10X objective, six images into 5 μm depth through the z axis were scanned, and horizontal z sections were collected and projected as superimposed stacks. The antibody staining was semi-quantitated by measuring mean fluorescence intensities (MFIs) with NIH Image J software (U.S. National Institutes of Health, Bethesda, MD, USA). MFI per square micrometer area was calculated by dividing the MFI units by the area of outlined regions. The CA3 region from four sections per brain and four animals per group were used for fluorescence intensity and quantification of A β_{1-40} .

2.5 PP2A activity assay

PP2A activity towards phospho-tau was assayed as described previously (Chohan et al. 2006). Briefly, 96-well plates were coated for 8 h at room temperature with 60 μl of 35 mM NaHCO₃ pH 9.5, containing 8.0 $\mu\text{g/ml}$ of a synthetic tau phosphopeptide in which Ser199 was phosphorylated. The coating solution was removed, and the wells were blocked with 150 μl of protein-free blocking buffer (Pierce, Pittsburg, PA) at 4°C overnight and then washed with 50 mM Tris-HCl, pH 7. Phosphatase activity was assessed with 60 $\mu\text{l/well}$ of 0.15 μg tissue extract (prepared with protease but no phosphatase inhibitors) resuspended in reaction buffer (20 mM β -mercaptoethanol, 2 mM EGTA, 2 mM MnCl₂ and 0.01 mg/ml BSA) in the presence or absence of 20 nM okadaic acid (OA) for 60 min at 30°C in a moist chamber. Each well was then incubated overnight at 4°C with 75 μl of a monoclonal antibody, Tau-1 (1:25,000), specific for tau that is unphosphorylated at Ser-198/199/202 (Grundke-Iqbal et al. 1986b). Plates were developed with anti-mouse HRP secondary antibody (1:5,000; Jackson Immuno, West Grove, PA, USA) and tetramethylbenzidine (TMB). Development was monitored in a microtiter plate reader at 650 nm with a 30 min kinetic reading every 2 min. To determine PP2A activity, values in the presence of OA were subtracted from the corresponding values in the absence of OA.

2.6 Behavioral studies

Once a week, the condition of each animal was assessed by measuring body weight, rectal temperature, food consumption, grooming, physical state, and clasp reflex. After 6 and 12 months of treatment, animals were subjected to hippocampal-dependent spatial memory tests using the water maze and object location, respectively. All the behavior procedures on animals were conducted in strict compliance with approved protocols from our institutional Animal Welfare Committee.

2.7 Spatial reference memory evaluation at 6 months

The spatial reference memory task evaluated in a water maze assesses hippocampal-dependent reference memory in rodents, requiring that rats use a spatial navigational strategy to find a fixed submerged escape platform. The hippocampal system processes information about the relationships among distal environmental cues into a spatial map where spatial coordinates of the submerged platform are encoded (Morris et al. 1982). The hippocampus is also crucial for memory storage, consolidation and restitution of the spatial

information (Riedel et al. 1999). The procedure was performed in a 180 cm diameter circular tank. The pool was filled with water ($20^{\circ}\text{C} \pm 1$) made opaque by adding white non-toxic paint. Acquisition was started with the escape platform (14 cm diameter submerged 1 cm below water surface) in the northwest quadrant and each animal was given 90 s to find the platform. If the rat did not find the platform in 90 s it was gently guided to it. At the end of each trial the rat was left on the platform for 20 s, then dried and returned to its home cage until the next trial. Four such acquisition trials, 20 min apart, were given on each day for 3 consecutive days. A test for retention (i.e., a probe trial, PT) was given 24 h after the last day of training. During the PT the rat was allowed to swim in the tank without the escape platform for 60 s. The measures of learning were the time and the distance swum to reach the virtual escape platform. For PT the number of entries in the platform zone was recorded. Rat behavior in the water maze was monitored by a Samsung digital camera (SDC 4304) mounted to the ceiling, and tracked and timed by a SMART (Pan Lab/San Diego Instruments) version 2.0.14 software.

2.8 Object location evaluation

This task was performed after 12 months of treatment and was used to measure hippocampal functioning since this brain structure is critical for associating objects with locations (Malkova and Mishkin 2003). Animals were exposed to two similar objects and they had to identify the spatial location of these two objects in an open-field as novel or familiar, based on the memory of an earlier experience with one of the two different object locations. The familiar location was explored a shorter time than the novel location, because the spatial representation of the former was still available in the memory. The test was developed in the classical open field apparatus (i.e. a PVC square arena, 100×100 cm, with Plexiglas walls, 70 cm high). The open field was placed in a different room from the experimenter. The open field was surmounted by a video camera connected to a computer for tracking. Before the object location test, animals received six sessions of habituation to the arena (2 session/day, 2 h apart; 10 min/session). During habituation sessions, an object was placed in the center of the arena. During the first habituation session, the time of exploration of the object was measured to evaluate neophobia. 24 hours after the last session of habituation, rats performed the object location test consisting of a sample phase and a test phase. During the sample phase, the rat was exposed to two similar objects and was allowed to explore for 5 min. The test phase occurred 1 h after the sample phase. One of the two identical objects was moved to a new location. To analyze cognitive performance, a discrimination index was calculated as follow: $(\text{time exploring the new location} - \text{time exploring familiar location}) \times 100 / \text{time exploring both locations}$. Rat behavior in the open field was monitored by a Samsung Digital Camera (SDC 4304) mounted to the ceiling and tracked and timed by a SMART (Pan Lab/San Diego Instruments) version 2.0.14 software. Time spent close to each object was manually recorded by the experimenter.

2.9 General Behavioral studies: monitoring of animals

During the period of the treatment individual condition of animal was assessed every week by evaluating grooming and physical state, and by measuring body weight, rectal temperature and food consumption.

2.10 Anxiety

Anxiety and exploratory activity was evaluated after 6 months of treatment with EHT by allowing rats to freely explore an open field for 20 min. The testing apparatus was a classic open field (i.e. a PVC square arena of 100 × 100 cm, with 70 cm high plexiglass walls). The open field was placed in a part of the room separated from the experimenter and the control station with a black opaque curtain. Rats were individually submitted to a single 20 min-session. Because for rodents the middle of a non-familiar arena is anxiogenic, anxiety was studied analyzing the time spent in the middle of the arena during the first five minutes of the session. To assess exploratory activity, the total distance the animals covered in the arena was tracked and measured. Data collection was performed using tracking files of the experiment recorded with SMART (Pan Lab/San Diego Instruments) version 2.0.14 software.

2.11 Neurological evaluation

After the first 6-months of treatment, rats were submitted to a battery of behavioral tests to perform a quantitative evaluation of reflexes, muscle strength and motor coordination. Using a scoring system adapted from Korenova and collaborators (2009), we were able to measure the consequences of the neurodegenerative processes on neurological and neuromuscular functions (see Table S1).

The beam-walking test. Three sorts of traversing segments were used (3 cm × 3 cm, 4 cm × 2 cm (traversing segment was 2 cm) and a round beam of 3.5 cm diameter). All had the same length of 200 cm and were placed 75 cm above the floor. Two training and one testing trials were performed. Traversing latency and number of hind-limb slips made during test performance were measured and scored according to the pre-defined rating scale (Table S1). The task was repeated at different sensitivity according to the following scheme:

- beam with a square section of 3 cm × 3 cm (1st day)
- beam with a rectangular cross-section of 4 cm × 2 cm (2nd day)
- beam with a round cross section of 3,5 cm diameter (3rd day)

The prehensile traction test. Rats were allowed to grasp with their forepaws a horizontal steel wire (3 mm in diameter) suspended 75 cm above a padded surface. Latency to fall from the wire was measured. The scoring conditions are described in Table S1.

After 12 months of treatment, neurological functions were evaluated using a test for neuromuscular functions (i.e. the hind-limb extension reflex test), a test for muscle strength (i.e. the prehensile traction test) and a test for motor coordination (i.e., the footprint test).

The hind-limb extension reflex test to measure the consequences of the neurodegenerative processes on neurological and neuromuscular functions. The scoring conditions are described in Table S2.

The prehensile traction test for evaluation of forelimb muscle strength. Rats were allowed to grasp with their forepaws a horizontal steel wire (3 mm in diameter) suspended 75 cm above

a padded surface. Latency to fall from the wire was measured. The scoring conditions are described in Table S3.

The footprint test evaluates motor coordination and synchrony by examining gait during normal walking. For this test, the animal paws are stained with non-toxic acrylic paint (forepaws with red and hind paws with blue). The rat has to walk through a $12 \times 12 \times 50$ cm transparent Plexiglas tunnel over absorbent paper. The following records were made from the walking tracks (see Fig. S1): a) stride length (distance between forepaw-forepaw and hind paw-hind paw); b) gait width (distance between left and right hind paws); c) placement of hind paw relative to fore paw (distance between hind paw-fore paw in each step cycle).

3. RESULTS

3.1 I_{2NTF} and I_{2CTF} genes were effectively expressed in rat brains

Rat pups were injected ICV on the day of birth with AAV1 containing N-terminal and C-terminal fragments of the PP2A inhibitor, I_2^{PP2A} (I_2 -N/C) or as control, AAV1-GFP, and after weaning on day 21, GFP and I_2 -N/C animals were put on either standard rat chow or chow containing 0.1% EHT for up to 1 year (Figure 1). As shown by us previously (Bolognin et al. 2012), considering a FLAG-tag was encoded into the I_2^{PP2A} N-terminal fragment- sequence, we were able to use I_2^{PP2A} N-terminal fragment primers that contain a FLAG sequence not homologous to endogenous I_2^{PP2A} to confirm the successful transduction of the brain with the AAV1- I_2 -N/C. The AAV1-mediated protein expression was less than 5% of the endogenous I_2^{PP2A} level as reported previously (Wang et al. 2010). As expected, FLAG-tagged I_2^{PP2A} N-terminal fragment sequences were not found in GFP control rats.

3.2 EHT prevents I_2 -induced cognitive impairment in rats

The expression in rat brains of the highly active N-terminal and C-terminal fragments of the PP2A inhibitor, I_2^{PP2A} (I_2 -N/C), led to cognitive deficits, and dietary supplementation with EHT relieved these deficiencies (Figures 2 and 3). Hippocampal-dependent cognitive function was assessed at 6 months using the spatial reference memory task in the water-maze (Figure 2). Analysis of the swim speeds of the animals (Figure 2A) revealed that AAV- I_2 -N/C rats swam significantly faster than other groups, independent of EHT supplementation (ANOVA, $p=0.003$; Fisher's post-hoc test, $p<0.049$). Results of the training were therefore analyzed in terms of distance covered to reach the submerged platform rather than time required (Figure 2B). During training in the absence of EHT dietary supplementation, AAV- I_2 -N/C rats displayed delayed performance compared to other groups (Figure 2B, ANOVA, $p=0.038$; Fisher's post-hoc test, $p<0.046$). This finding showed that AAV- I_2 -N/C rats were impaired in the learning of the task compared to AAV-GFP rats, but that treatment with EHT rescued this impairment. During the probe trial, AAV- I_2 -N/C rats also displayed poor performance and visited the platform location significantly less frequently than AAV-GFP rats raised on control diets lacking EHT (Figure 2C, Student t-test, $p=0.023$). This confirmed the impaired ability of AAV- I_2 -N/C rats to encode and memorize spatial information, i.e. the spatial coordinates of the submerged platform. The AAV- I_2 -N/C rats treated with EHT visited the platform location similarly to

the AAV-GFP control animals, confirming that treatment of AAV-I₂-N/C rats with EHT prevented spatial memory impairment.

After 12 months of EHT treatment, hippocampal cognitive function was evaluated using a different test: the object location memory task. During the sample phase of the test, all animals spent similar time exploring the two objects in the arena (Figure 3A, Student t-test, $p > 0.999$). This result shows that the rat's baseline preferences for the objects and locations involved in the test were not significantly affected by I₂-N/C expression or EHT dietary supplementation. During the test phase, control groups and AAV-I₂-N/C rats treated with EHT displayed a discrimination ratio of about 70% showing that these animals tend to explore the new location more than the old. In contrast, AAV-I₂-N/C rats that were not treated with EHT presented a discrimination ratio close to 55%, showing that this group spent similar time in both locations. The performance of this group was statistically different from control and EHT treated groups (Figure 3B, Student t-test, $p < 0.020$). These results confirmed the hippocampal impairment associated with AAV-I₂-N/C expression seen at 6 months in the water-maze task and showed that 12 months of EHT treatment can prevent the impairment of spatial information processing associated with I₂-N/C expression.

3.3 General physical state, body weight and temperature

No alteration in general physical state including grooming and posture, due to either the AAV-I₂NTF/CTF (N/C/vh) infection or the treatment with EHT was detected. Figure 4 represents the follow up of body weight and body temperature during the treatment with EHT (A,C: during the first 6 months; B,D: during the last 6 months). Across months, we observe a general increase of the body weight (Figure 4A and 4C, ANOVAs, $p < 0.001$) and of the body temperature (Figure 4C and 4D, ANOVAs, $p < 0.001$). But, we did not observe any significant effect of AAV-I₂NTF/CTF infection or EHT on these parameters (Figs. 4A and 4C, ANOVAs, $p > 0.141$; Figures 4B and 4D, ANOVAs, $p > 0.263$) during the period of the treatment.

3.4 Evaluation after 6 months of treatment with EHT

Figure 5A represents the neuroscore. Statistical analysis of data obtained from the assessment of neurological examination did not reveal any significant difference among groups (Figure 5A, Kruskal-Wallis test, $p = 0.380$). This result indicates that, at six months of age, AAV-I₂NTF/CTF rats did not present any impairment in neurological function and that treatment with EHT did not induce any neurological side effects.

As shown in Figure 5B, there was a tendency for AAV-I₂NTF/CTF (N/C/vh) rats to visit less the center of the arena than other groups. But statistical analysis did not reveal any significant difference (Figure 5B, ANOVA, $p = 0.736$) among groups. This tendency to explore less the center of the open field suggested that, as the pathology develops in AAV-I₂NTF/CTF rats, anxiety levels would shift toward hyper-anxiety. However, treatment with EHT restored anxiety to normal levels.

During the twenty minutes of free exploration, all groups covered similar distance (Figure 5C, ANOVA, $p = 0.852$), suggesting that all animals displayed similar level of exploration. No effect of the treatment was observed.

3.5 Evaluation after 12 months of treatment with EHT

Figure 6A represents performance of rats in the hind-limb extension reflex test. Statistical analysis did not reveal any difference between groups (Figure 6A, Student t-tests, $p > 0,059$). This result indicates that, at twelve months of age, AAV- $I_2^{NTF/CTF}$ rats did not present any deterioration of peripheral neurological functions and that treatment with EHT did not induce any peripheral neurological side effects.

As shown in Figure 6B, statistical analysis did not revealed any significant difference between groups in the prehensile traction test (Figure 6B, Student t-tests, $p > 0,200$). As represented in the regression chart, correlation analysis of the body weight and the prehensile traction score of the animals did not show any significant effect between body weight and performance in the test (Figure 6C). These results indicate that, neither the AAV- $I_2^{NTF/CTF}$ infection nor the treatment with EHT induced any changes in forelimb muscle strength.

Finally, Figures 6D–G represent analysis of different parameters for the footprint test. Statistical analysis did not reveal any difference between groups in any parameter (Figures 6D–G, Student t-tests, $p > 0,528$). These analyses showed that neither the AAV- $I_2^{NTF/CTF}$ infection nor the treatment with EHT altered motor coordination and synchrony.

3.6 Dietary supplementation with EHT blocks I_2^{PP2A} inhibition of PP2A and prevents tau hyperphosphorylation

I_2 -N/C fragments are known to bind to the PP2A catalytic subunit PP2Ac and inhibit phosphatase activity (Arnaud et al. 2011). We therefore compared the levels of PP2A activity toward phospho-tau in the brains of rats raised with and without dietary EHT supplementation. On normal diets, PP2A activity was significantly lower in rats that express I_2 -N/C than in GFP controls. EHT dietary supplementation completely rescued the I_2 -N/C-induced PP2A deficiency (Fig 7, A, B). Neither I_2 -N/C expression nor EHT treatment caused any significant changes in the level of total PP2Ac protein. These data confirm that expression of I_2 -N/C inhibits PP2A activity, and that treatment with EHT can block I_2 -N/C-induced PP2A inhibition.

The principal form of PP2A that dephosphorylates phospho-tau requires carboxy methylation at its C-terminus (Tolstykh et al. 2000; Wu et al. 2000). Levels of PP2A methylation are controlled by a balance between the activities of two highly conserved, PP2A-specific, enzymes: a methyl transferase, PPMT, that transfers methyl groups from S-adenosylmethionine to the PP2A carboxy terminus; and a methyl esterase, PME, that demethylates PP2A. EHT was initially identified as a component in coffee extracts that inhibited the PP2A demethylation reaction (Lee et al. 2013). Dramatic increases in PP2A demethylation have been observed in brains from AD patients (Sontag et al. 2004). The effect of EHT on levels of PP2A methylation in I_2 -N/C and control rats was therefore investigated, and it was found that EHT treatment significantly reduced the levels of demethylated PP2Ac in I_2 -N/C rats; and a similar trend was seen in GFP controls (Fig 7 C, D). Together these data suggest that the expression of I_2 -N/C decrease PP2A activity, and

that the treatment with EHT, which blocks PP2A demethylation (Lee et al. 2011) can rescue the I₂-N/C inhibition of phosphatase activity.

Since PP2A is the major brain phospho-tau phosphatase (Bennechib et al. 2000; Gong et al. 2000; Liu et al. 2005), I₂-N/C inhibition of PP2A activity would be expected to cause tau hyperphosphorylation. Western blot analysis of tau phosphorylation confirmed this supposition. I₂-N/C expression significantly decreased the total tau level and increased tau hyperphosphorylation compared with GFP control animals at T205, T212, S214, S262/356 and S396, but not at S199, a PP2A non-preferred site (Liu et al. 2005) (Figure 8). Treatment with EHT reduced I₂-N/C-induced tau hyperphosphorylation at most of these sites; there were no significant differences between EHT-treated experimental and GFP control rats. These results indicate that I₂-N/C expression induces abnormal hyperphosphorylation of tau through inhibition of PP2A activity, and that treatment with EHT ameliorates this pathology.

PP2A activity and the levels of tau hyperphosphorylation are known to have a negative correlation at the phosphorylation sites most associated with pathology in AD brain (Liu et al. 2005). As shown in Figures 7 and 8, PP2A activity was significantly decreased in I₂-N/C rats, tau hyperphosphorylation was coordinately increased, and both the changes in PP2A and tau phosphorylation were rescued in EHT-treated animals. To further establish that the hyperphosphorylation of tau that was observed in I₂-N/C rats in the present study was due to the inhibition of PP2A activity, we evaluated this possibility with Spearman non-parametric correlation analysis. We found a significant negative correlation between the PP2A activity and the abnormal hyperphosphorylation of tau at T205 (p=0.0307), T212 (p=0.046), S262/356 (p=0.0002), S396 (p=0.0064) and S396/404 (p=0.0434) (Figure 8). These data strongly support the hypothesis that EHT prevented I₂N/T-induced hyperphosphorylation of tau by maintaining healthy levels of active PP2A in the presence of overexpressed I₂-N/C.

3.7 EHT treatment reduces intraneuronal A β accumulation

Reduced levels of PP2A activity induced by I₂-N/C (Bolognin et al. 2012) and by other means (Sontag et al. 2007) have previously been shown to lead to increases in A β load. The effect of EHT treatment on the accumulation of intraneuronal A β in I₂-N/C rats was therefore investigated. The immunohistochemical analysis revealed a low number of intraneuronal A β -positive cells in rats treated with EHT compared with non-treated rats (Figure 91). These results suggest that I₂-N/C expression promotes the amyloidogenic processing of β APP, and that this effect can be rescued by EHT treatment. Our findings, though preliminary, are in agreement with those of Sontag et al. (2007) who showed that increase in PP2A activity can lead to non-amyloidogenic processing of APP.

4. DISCUSSION

Alzheimer's disease is multifactorial and involves several different etiopathogenic mechanisms (Iqbal et al. 2005b). The familial form, which accounts for less than 1%, and the sporadic form, which represents the remaining over 99% of the cases of AD, are histopathologically identical. While the familial form of AD has been associated with certain mutations in β -APP and presenilins 1 and 2, and the inheritance of the APOE4 allele markedly increases the risk for the disease, the causes of the sporadic form of the disease are

not understood. Nevertheless, the decrease of PP2A activity associated with an increase in the expression, the cleavage and the translocation of I₂^{PP2A} and a decrease in the methylation of PP2Ac reported in AD brain (Gong et al. 1993; Gong et al. 1995; Sontag et al. 2004; Tanimukai et al. 2005; Wang et al. 2010; Bolognin et al. 2012) can lead both to tau and Aβ pathologies. The present study shows that the overexpression and cleavage of I₂^{PP2A} can lead to increases in PP2Ac demethylation, inhibition of PP2A activity, tau hyperphosphorylation, Aβ expression, and cognitive impairment, and that all of these changes are significantly reversed by a minor component of coffee, EHT.

The AAV1-I₂^{N_{TF}-CTF} rat model used in the present study was developed in our lab (Bolognin et al. 2012). Compared to transgenic animals, one of the major advantages of the viral gene transfer technology used to generate the I₂-N/C rat model, is that long-term transgene expression is achieved without affecting the genetic background of the animal (Lawlor et al. 2007). Rats injected with AAV serotype 1 vector encoding the two fragments of I₂^{PP2A} showed a marked reduction of PP2A activity beginning at 4 months of age (Bolognin et al. 2012). The present study extended these findings to 13 months of age, when the cytosolic tau and Aβ pathologies were evident, making this model appropriate to study the effect of changes in PP2A activity on AD-type changes. EHT was administered to rats from the age of 21 days to 13 months. Compared to non-treated animals, a decrease in demethylation of PP2Ac was observed in EHT-treated rats, indicating the efficacy of the compound over chronic long-term use. These results corroborate previous reports showing EHT efficacy in both attenuation of α-synucleinopathies *in vivo* after a 9-months treatment, and inhibition of PP2A demethylation *in vitro* (Lee et al. 2011). Pharmacokinetic studies delivering [³H]-EHT intraperitoneally revealed that this compound crosses the blood brain barrier achieving levels sufficient to inhibit PP2A demethylation, indicating its feasibility for proof of concept studies (Lee et al. 2011). EHT was identified as the major PP2A demethylation inhibitor in coffee (Lee et al. 2011) and was reported to be effective in preclinical studies (Lee et al. 2011; Lee et al. 2013).

Epidemiological studies have established a negative correlation between coffee consumption and the incidence of Parkinson's Disease and AD (Barranco Quintana et al. 2007; Saaksjarvi et al. 2008). In the present study, we observed that rats expressing I₂-N/C displayed increased PP2A demethylation and a decrease in PP2A activity, and that treatment with EHT decreased PP2A demethylation and increased PP2A activity toward tau. Previous studies showed that methylation of PP2Ac affects phosphatase activity in part by facilitating the binding of the regulatory Bα subunits to AC dimers (Tolstykh et al. 2000; Xu et al. 2008). It is not clear, however, that the protective effects of EHT against I₂^{PP2A} inhibition derive entirely from inhibition of PP2A demethylation. The inhibitory effect of EHT on demethylation most likely stems from formation of a complex between EHT and PP2A that precludes the interaction between PME and PP2A that is required for demethylation. Thus, the possibility cannot be excluded that the EHT/PP2A complex may also preclude the formation of inhibitory complexes between PP2A and I₂^{PP2A} in much the same way that it appears to block the interaction between PP2A and PME.

As a consequence of the decrease in PP2A activity in I₂^{N_{TF}-CTF} rats, we observed a clear increase in tau hyperphosphorylation at multiple sites. The association between these events

was confirmed by correlation analysis, showing that decreases in PP2A activity are negatively correlated with the hyperphosphorylation of tau at several sites, particularly the PP2A-dependent sites S262, T212, T205 and S396. S199 and S214 that are not preferred sites for PP2A did not show any correlation. This analysis is consistent with previous reports in AD cases (Liu et al. 2005). PP2A regulates phosphorylation of tau both directly and by regulating the activities of several tau protein kinases (Iqbal et al. 2005a). The effect of I₂-N/C can be either direct by inhibiting the PP2A activity as we previously demonstrated (Arnaud et al. 2011) or through a down regulation of B α subunit PP2A holoenzyme which specifically regulates the phosphorylation of tau (Sontag et al. 2007; Xu et al. 2008).

In addition to tau pathology, we also observed an accumulation of intraneuronal A β in I₂-N/C rats, which was attenuated in animals treated with EHT. These beneficial effects of EHT treatment may target early A β pathological mechanisms through an increase in PP2A activity. PP2A demethylation, for instance, has been reported to be associated with a concomitant decrease in the steady-state release of neuroprotective APP α phosphorylated species and increased secretion of β - and γ -secretase-cleaved APP fragments, inducing a shift in APP processing toward the amyloidogenic pathway (Sontag et al. 2007). Increased APP phosphorylation, either directly through decreased activity of PP2A towards phospho-APP, or indirectly through reduced phosphatase activity towards phospho-JNK, can result in increased A β production (Colombo et al. 2009). Thus, activation of PP2A by small molecules such as EHT, offers a new therapeutic approach for the prevention and treatment of AD and other neurodegenerative disorders, including tauopathies such as frontotemporal dementias (Voronkov et al. 2011).

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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Highlights

- A coffee component, EHT, protects PP2A from demethylation and inactivation.
- EHT inhibits hyperphosphorylation of tau by inhibiting inactivation of PP2A.
- EHT in a rat model of sporadic AD can rescue tau pathology and cognitive impairment.
- EHT can reduce intraneuronal A β accumulation in a rat model of sporadic AD.
- EHT provides a novel therapeutic opportunity for prevention and treatment of AD.

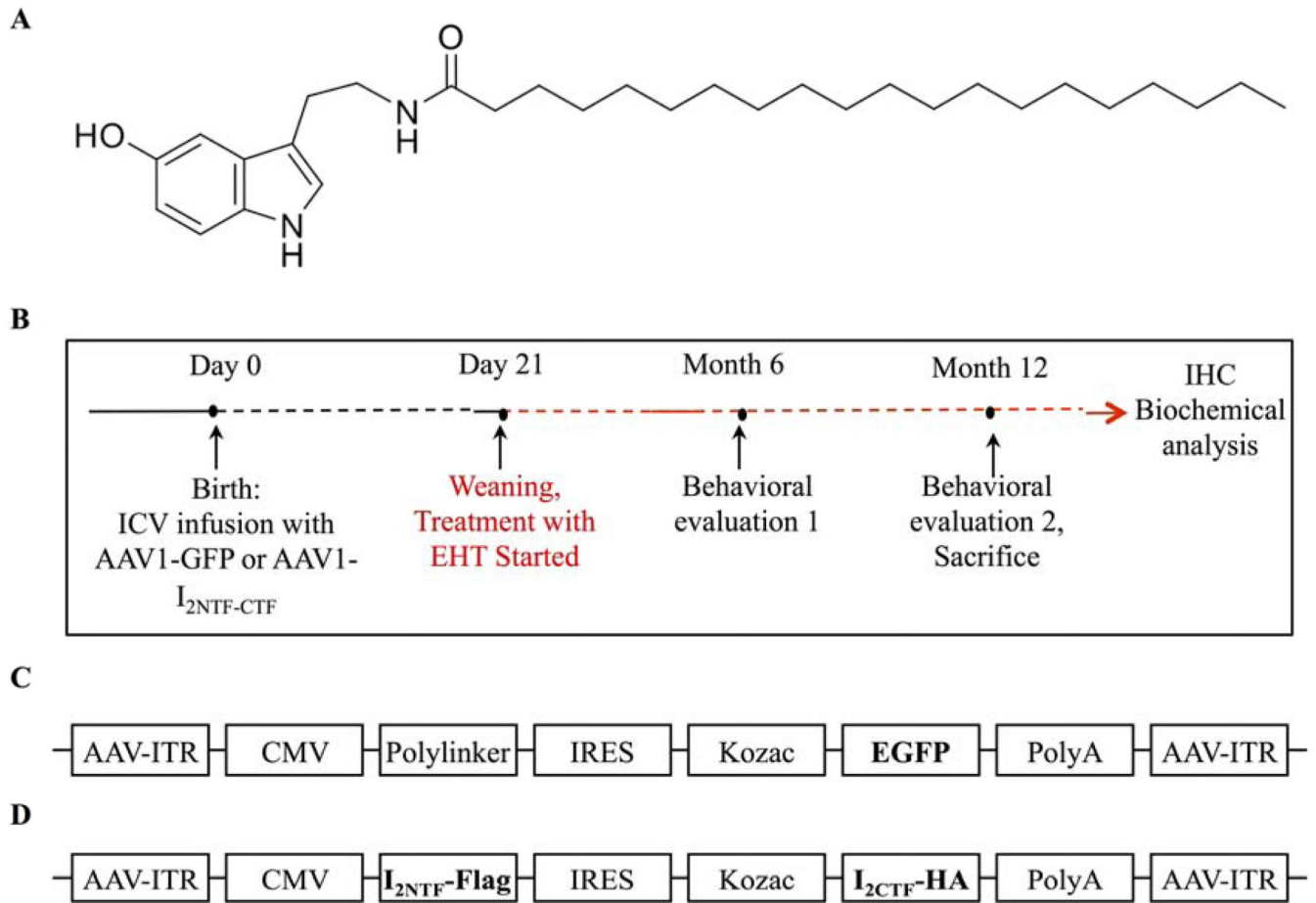


Figure 1.

EHT structure and experimental design and I_2 -N/C transduction in adult rat brains. (A) EHT chemical structure. (B) Schematic representation of the study. Rat pups were injected ICV with AAV on the day of birth. (C, D) Linear maps of the AAV plasmids (based on pTRUF12) containing (C) GFP, or (D) I_2 _{NTF} and I_2 _{CTF} genes inserted between the inverted terminal repeats (ITR). CMV cytomegalovirus promoter, IRES internal ribosomal entry site from poliovirus. After weaning on day 21, GFP and I_2 -N/C animals were put on either standard rat chow or chow containing 0.1% EHT for up to one year.

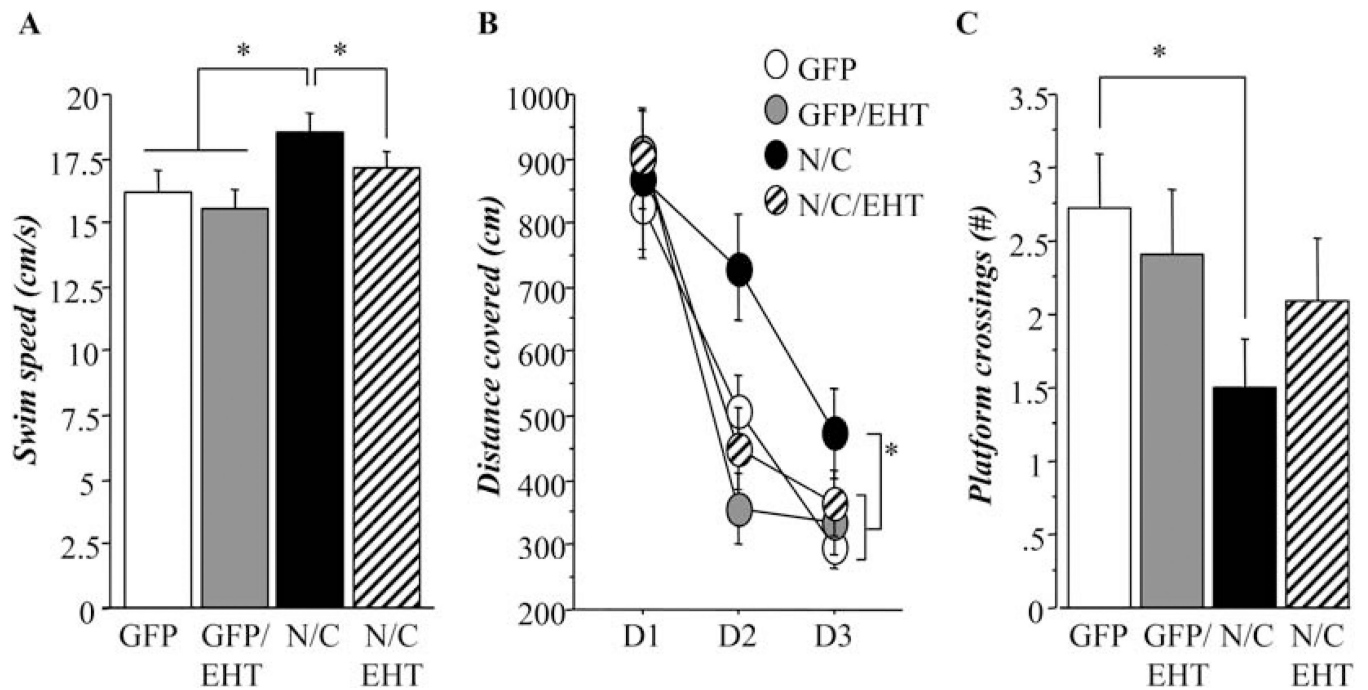


Figure 2.

EHT prevents the reference memory impairment in rats expressing I₂-N/C when tested by the hippocampal-dependent spatial memory water maze test. A) Hippocampal functioning in I₂-N/C rats displayed significantly faster swim speed than other groups (ANOVA, $p < 0.01$; Fisher's post-hoc test, $p < 0.05$), therefore, results of the training phase were analyzed as distance covered to reach the submerged platform. B) Water-maze task: training phase. I₂-N/C rats treated with vehicle displayed delayed training performance compared to other groups (ANOVA, $p = 0.038$; Fisher's post-hoc test, $p < 0.046$) reflecting hippocampal impairment that can be prevented with 6-months treatment with EHT. C) Water-maze task: Probe-trial. I₂-N/C rats treated with vehicle visited less the platform location than other groups (Student t-test, $p = 0.023$), confirming hippocampal impairment and its prevention by 6-months treatment with EHT.

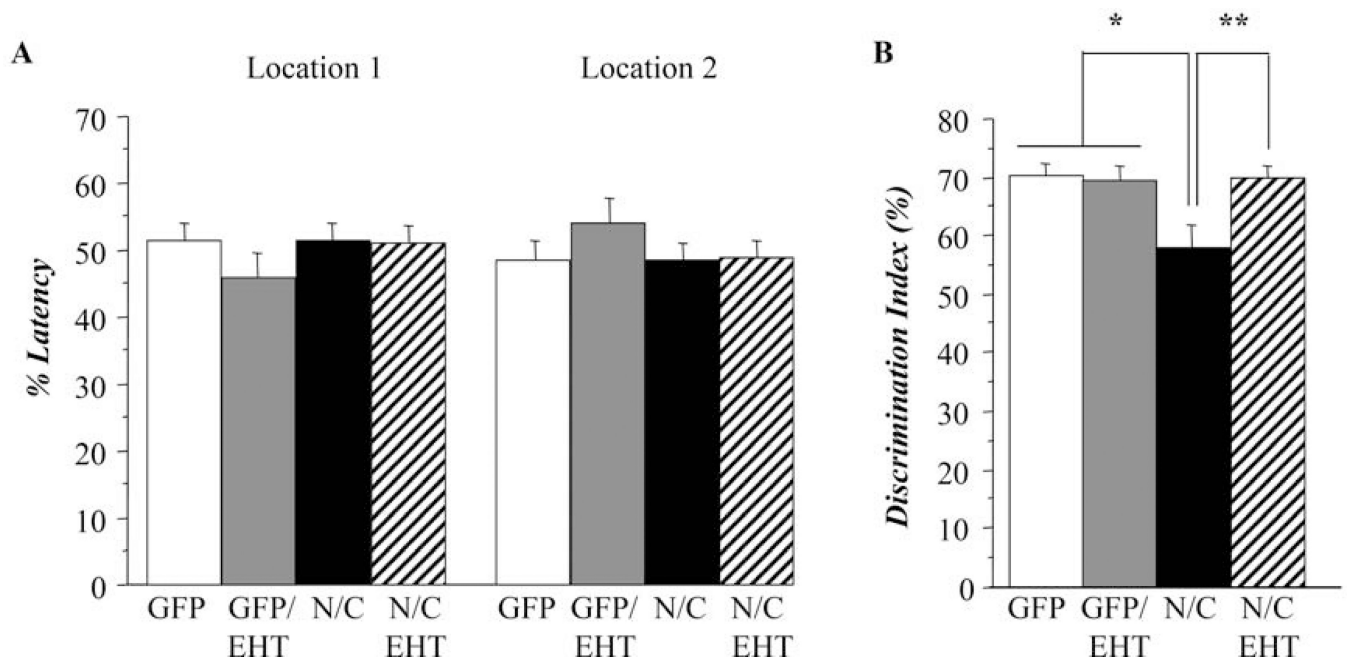


Figure 3.

EHT prevents the cognitive impairment in rats expressing I₂-N/C. Hippocampal functioning was measured to determine cognitive impairment. A) Object location task: sample phase. No differences were found among the groups, indicating no preferences for objects or location (Student t-test, $p > 0.99$). B) Object location task: test phase. I₂-N/C rats spent significantly less time to analyze the object in the new location (Student t-test, $p < 0.02$), indicating cognitive impairment. When treated with EHT for 12 months I₂-N/C rats displayed similar performance as control groups.

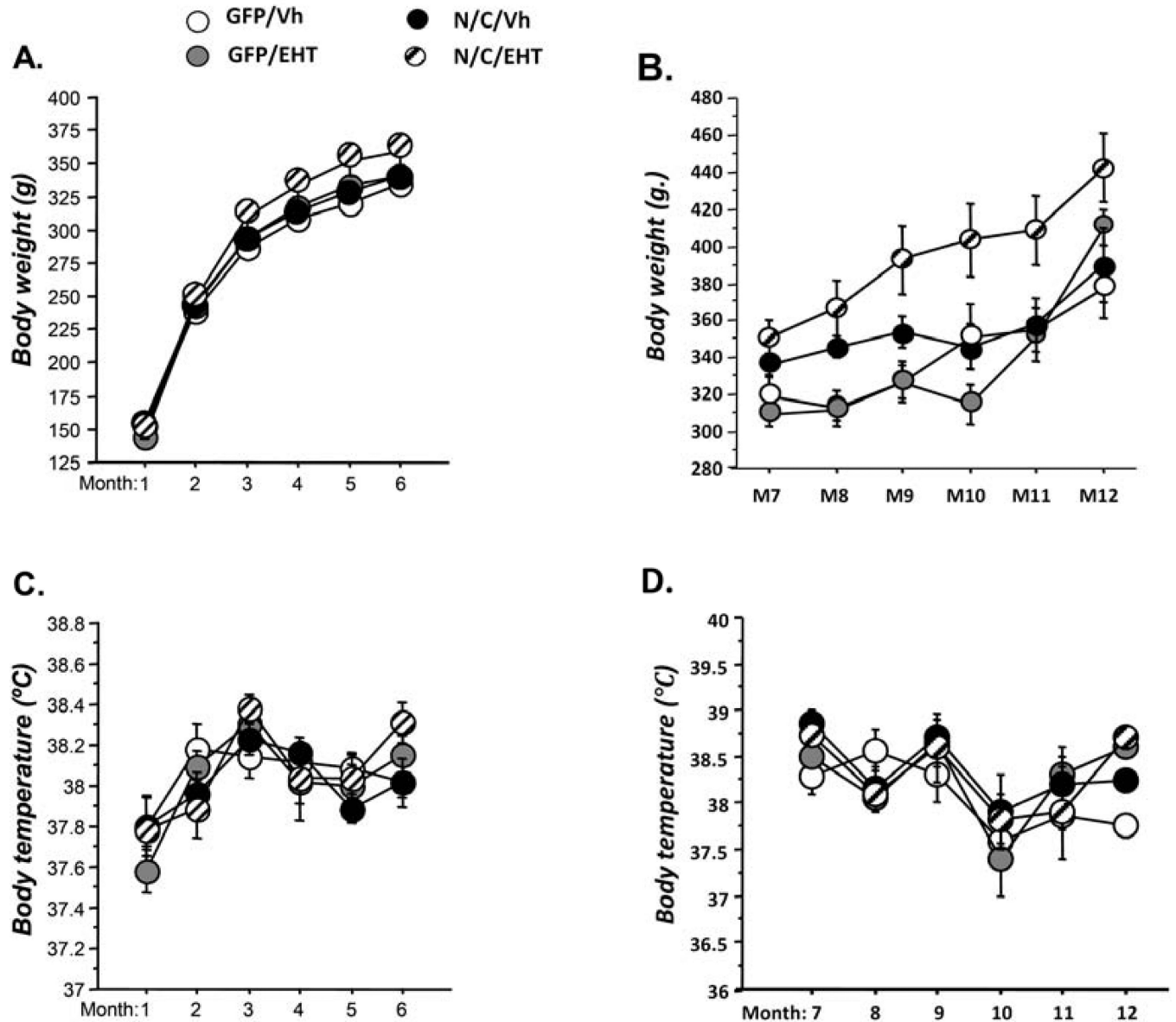


Figure 4. Effect of EHT treatment on body weight and temperature. Follow-up of body weight and body temperature during the treatment with EHT (A-B: during the first 6 months; C-D: during the last 6 months). Across months, we observed a general increase of the body weight (Figure 4A and 4C, ANOVAs, $p < 0.001$) and of the body temperature (Figure 4D and 4E, ANOVAs, $p < 0.001$). But, we did not observe any significant effect of AAV- I_{2NTF}/CTF infection or EHT on these parameters (Figures 4A and 4C, ANOVAs, $p > 0.141$; Figures 4B and 4D, ANOVAs, $p > 0.263$) during the period of the treatment.

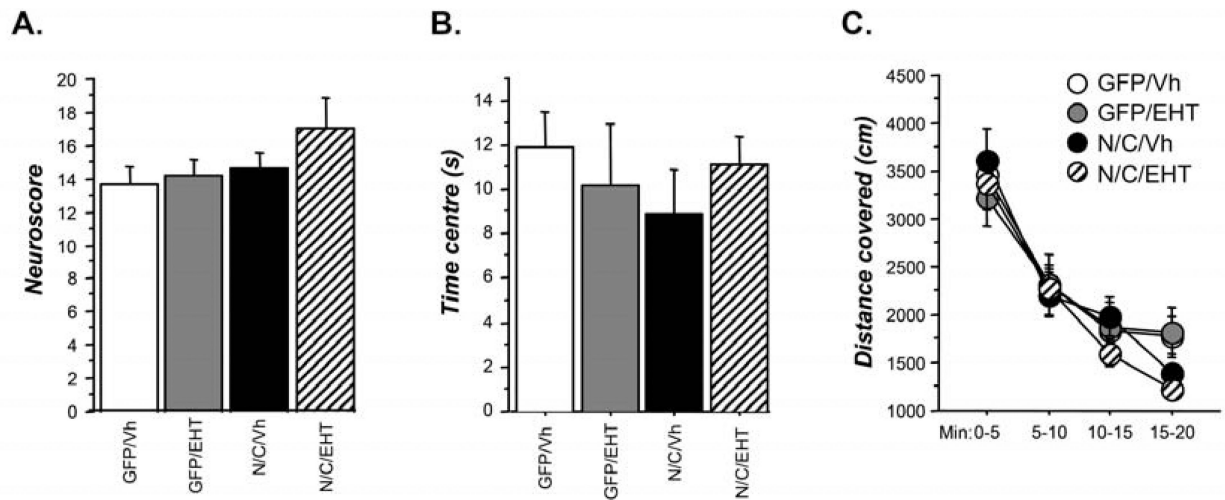
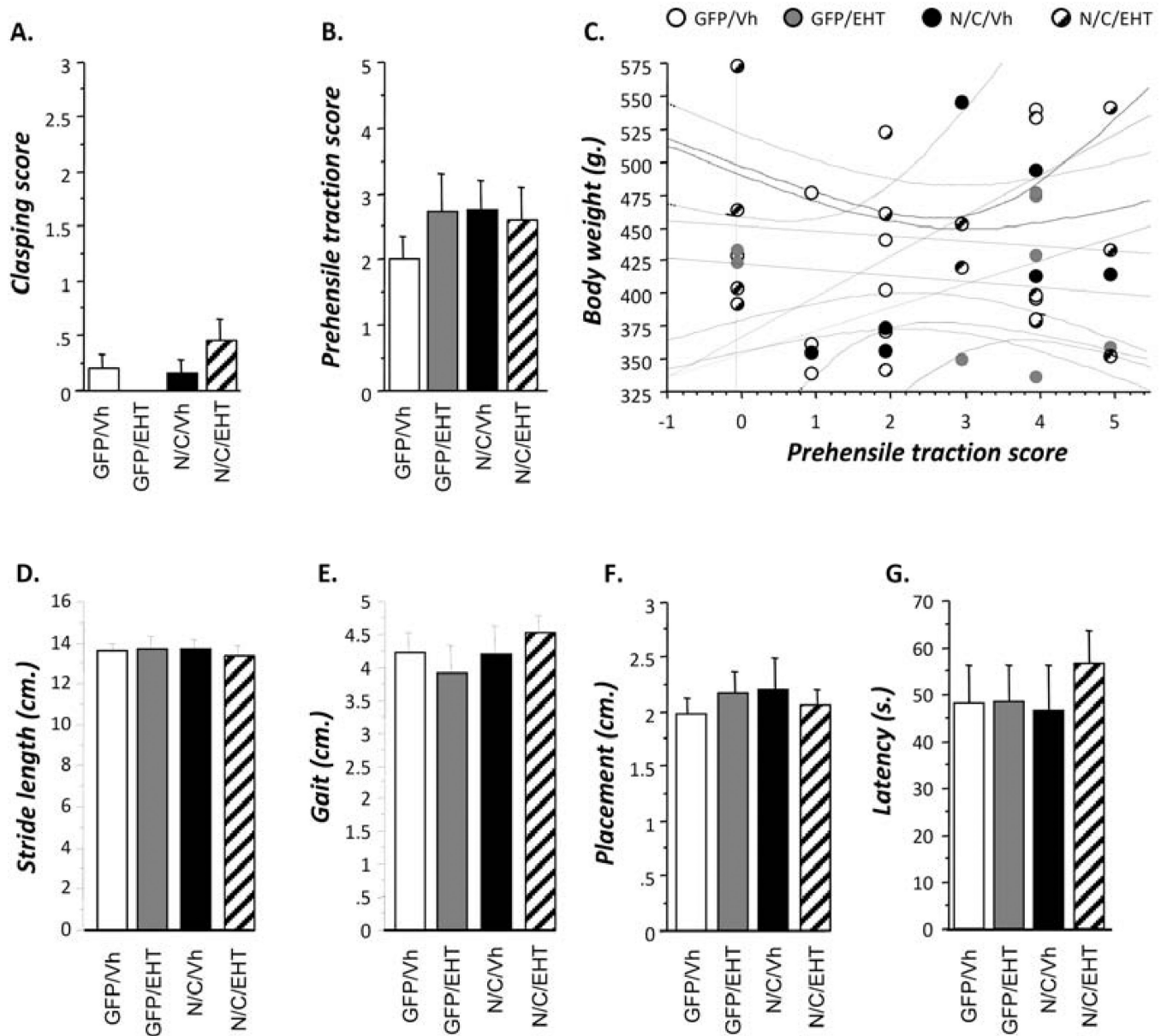


Figure 5. General behavior in 6-month-old rats. A. Assessment of neurological examination (see Table S1) quantitated as neuroscore; B. Anxiety measured by time spent by an animal in the center of the arena; C. Exploratory activity measured by distance covered exploring the arena. EHT treatment had no significant effect on any of the above measures of general behavior.

**Figure 6.**

Lack of any side effects in EHT- and control-treated rats at age of 12 months. Different tasks corresponding to general behavior performed at 12 months of age in the four groups of rats. A. Clasping reflex score; B, C. Prehensile traction score; D-F. Stride length, gait and placement in footprint test. G. Latency to explore novel object in an arena in neophobia test. No significant differences among the four groups of animals were found in any of the above general behavioral tests, indicating no adverse effect of EHT treatment.

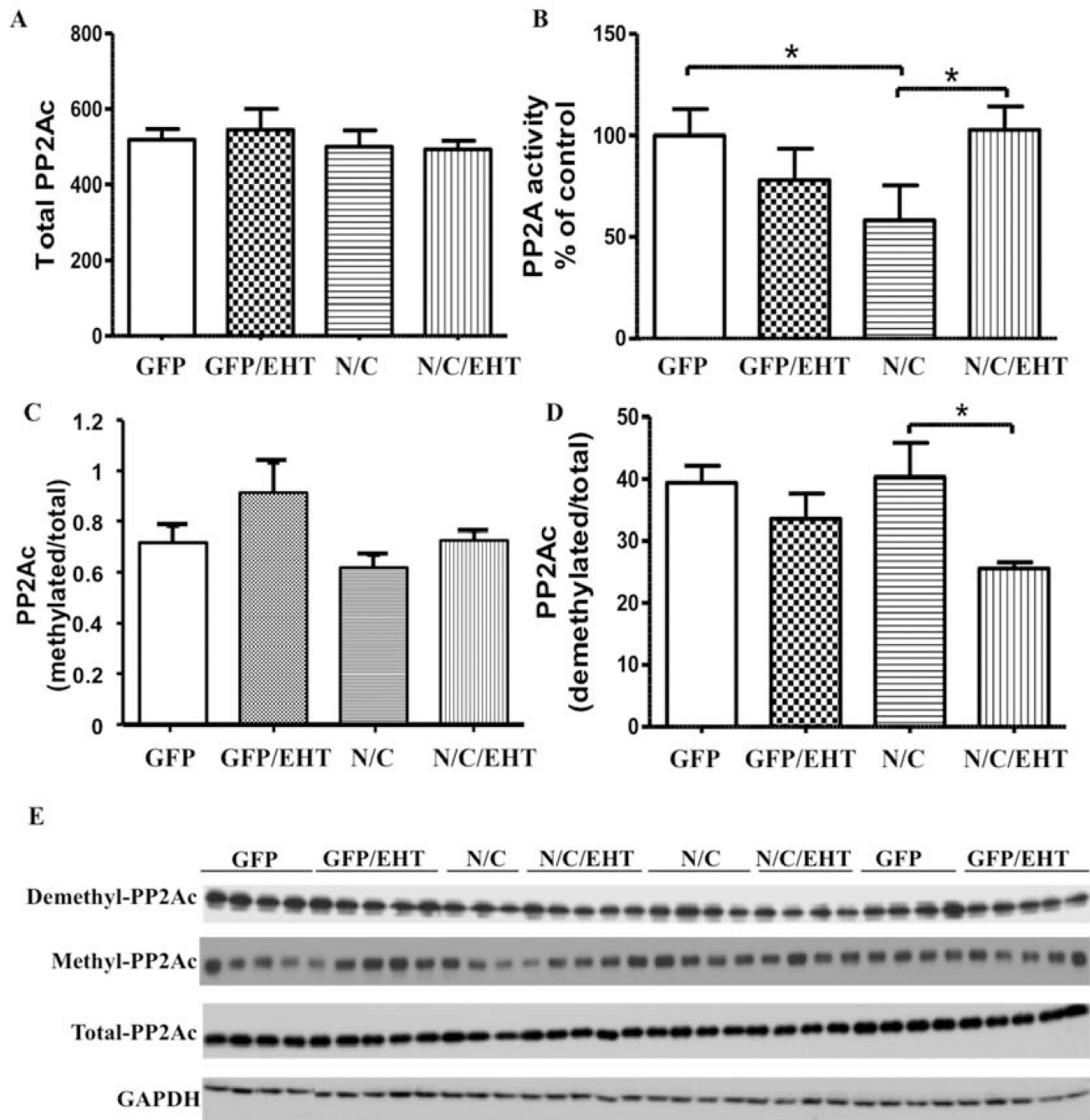
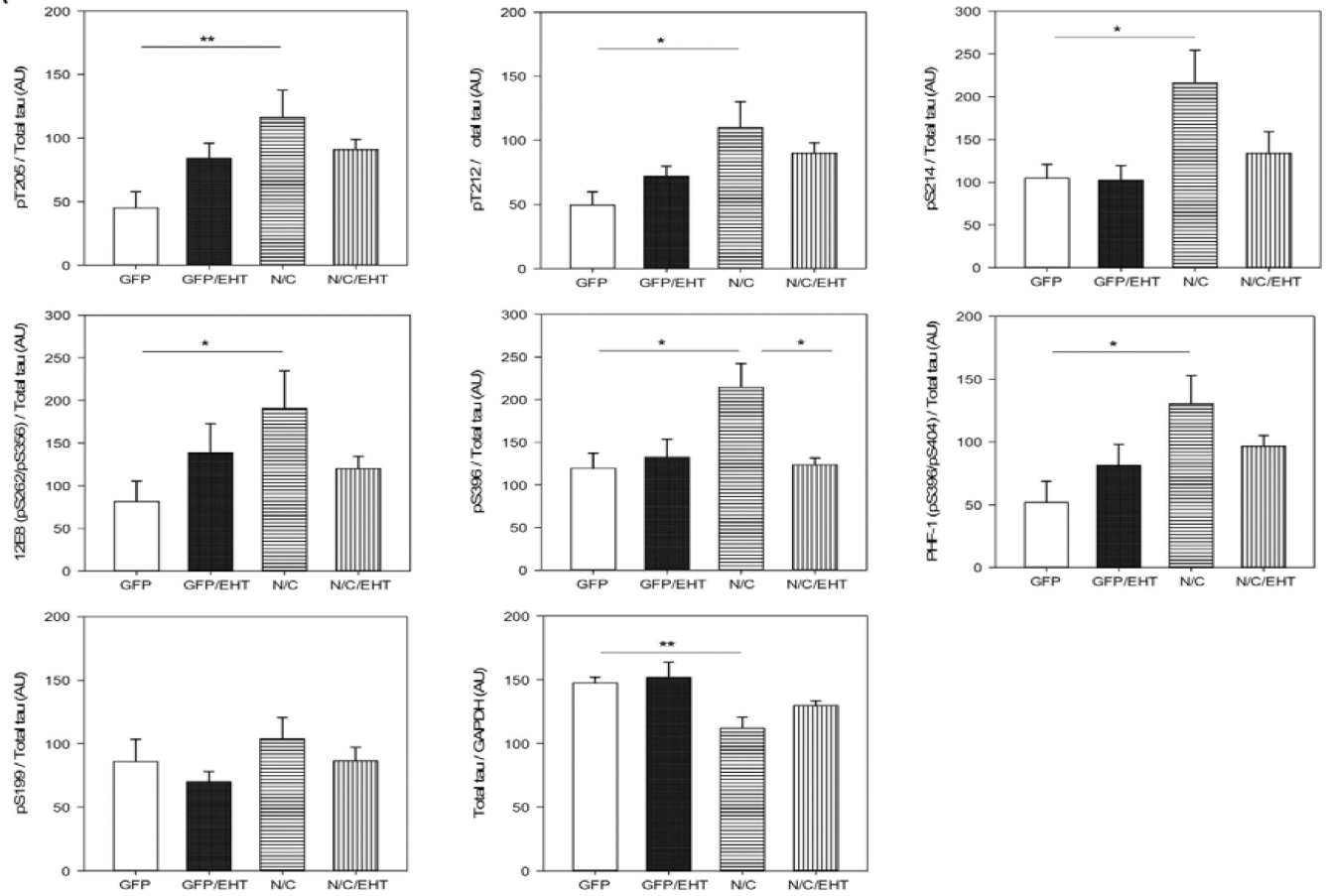


Figure 7.

EHT increases PP2A activity in I₂-N/C rats. A) Western blot quantification of I₂-N/C and GFP rat hippocampi showing the effect of EHT treatment on the level of total PP2Ac. B) PP2A activity was recovered in I₂-N/C rats treated with EHT, evaluated by PP2A phosphatase assay. C) EHT increased PP2Ac methylation and D) reduced PP2Ac demethylation in I₂-N/C rats. E) Representative blots shown for demethylated (1D6); methylated (4D9); total-PP2Ac (6A3) and GAPDH. *p<0.05. The data are shown as mean ± SE, normalized by GAPDH for total PP2A.

A



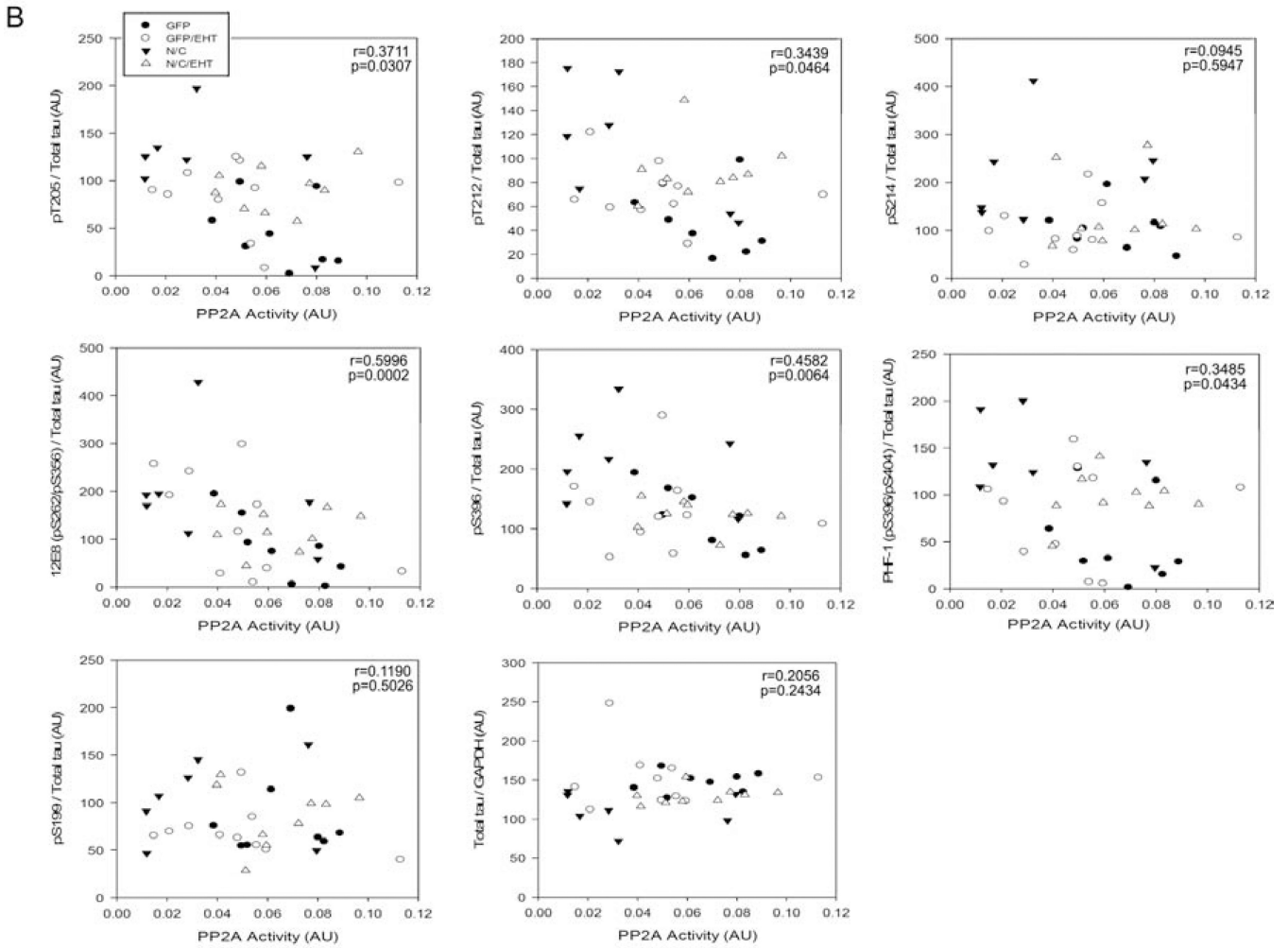


Figure 8. Abnormal hyperphosphorylation of tau in rats expressing I₂-N/C is prevented by EHT treatment. A) Western blot density data are shown as mean ± SE, normalized by GAPDH and for all the phosphorylated sites by total tau. *p<0.05; **p<0.01. B) PP2A activity values from the four groups of rats were correlated with the values corresponding to the densitometry analysis of Western blots developed for various tau phospho-sites. Decrease in hyperphosphorylation of tau at Thr205, Thr212, Ser262/356 (12E8 site), and Ser396/404 (PHF-1 site) directly correlated with increase in PP2A activity.

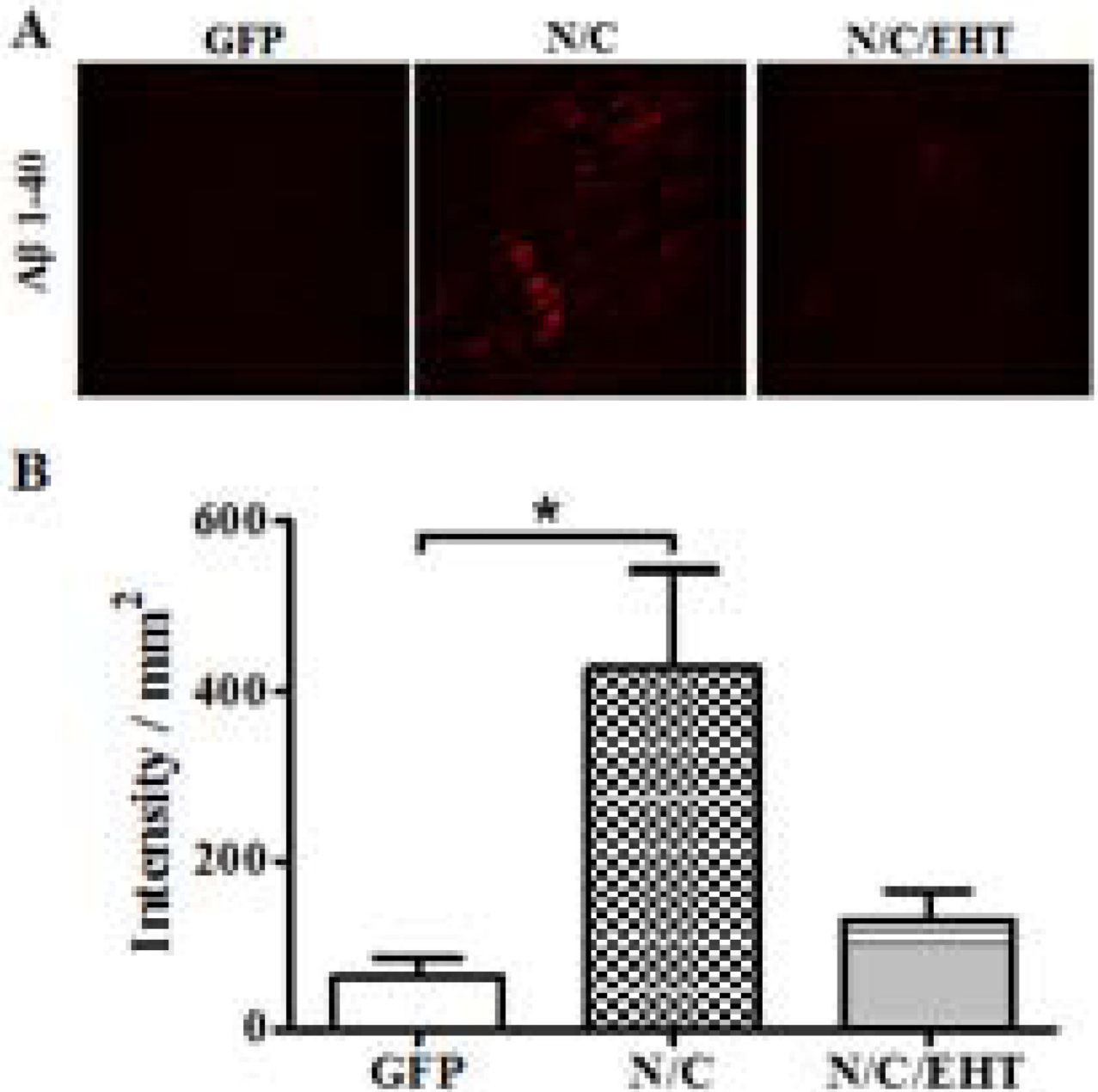


Figure 9. I₂-N/C overexpression increases intraneuronal Aβ 1–40 load, which appears to be attenuated by EHT. A) Representative micrographs of immunohistochemistry (IHC) from cerebral cortex showed Aβ 1–40 load increases in I₂-N/C- (p<0.05; n=3) but not EHT-treated rats. B) IHC quantification of I₂-N/C and GFP rat hippocampi showing the effect of EHT treatment on the level of Aβ 1–40. EHT treatment appears to attenuate the Aβ pathology (p=0.1; n=3 animals/group; p<0.01 for n=12 sections from 3 animals/group).