

Palmitoylethanolamide Protects Against the Amyloid- β 25-35-Induced Learning and Memory Impairment in Mice, an Experimental Model of Alzheimer Disease

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Alzheimer disease (AD) is the most common form of neurodegenerative dementia. Amyloid- β deposition, neurofibrillary tangle formation, and neuro-inflammation are the major pathogenic mechanisms that in concert lead to memory dysfunction and decline of cognition. Palmitoylethanolamide (PEA) is the naturally occurring lipid amide between palmitic acid and ethanolamine. Despite its clear role in inflammation and pain control, only limited *in vitro* evidence exist about a role for PEA in neurodegenerative diseases. Here we describe the neuroprotective activities of PEA in mice injected intracerebroventricularly with amyloid- β 25–35 (Ab25–35) peptide (9 nmol). We used spatial and non-spatial memory tasks to evaluate learning and memory dysfunctions. Ab25–35 injection significantly impaired spontaneous alternation performances, water maze spatial reference and working-like memory, and novel object recognition test. PEA was administered once a day (3–30 mg/kg, subcutaneously), starting 3 h after Ab25–35, for 1 or 2 weeks. PEA reduced (10 mg/kg) or prevented (30 mg/kg) behavioral impairments induced by Ab25–35 injection. PEA failed to rescue memory deficits induced by Ab25–35 injection in peroxisome proliferator-activated receptor- α (PPAR- α) null mice. GW7647 (2-(4-(2-(1-cyclohexanebutyl)-3-cyclohexylureido)ethyl)phenylthio)-2-methylpropionic acid; 5 mg/kg per day), a synthetic PPAR- α agonist, mimicked the effect of PEA. Acute treatment with PEA was ineffective. According with the neuroprotective profile of PEA observed during behavioral studies, experimental molecular and biochemical markers induced by Ab25–35 injection, such as lipid peroxidation, protein nitrosylation, inducible nitric oxide synthase induction, and caspase3 activation, were reduced by PEA treatment. These data disclose novel findings about the therapeutic potential of PEA, unrevealing a previously unknown therapeutic possibility to treat memory deficits associated with AD.

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INTRODUCTION

Alzheimer disease (AD) is the most common form of neurodegenerative dementias. AD has a complex and progressive pathological phenotype characterized by hypometabolism and impaired synaptic function, followed by pathological burden (Zawia *et al*, 2009). Amyloid- β deposition, neurofibrillary tangle formation, and neuro-inflammation are the major pathogenic mechanisms that act in concert to produce memory dysfunction and decline of cognition. To date, there is no specific treatment for AD, and the progressive and multifaceted degenerative pheno-

type suggests that new successful treatment strategies need to be equally multi-faceted and stage-specific.

Palmitoylethanolamide (PEA), the naturally occurring amide between ethanolamine and palmitic acid, is an endogenous lipid and belongs to the family of the fatty acid ethanolamides (FAEs). PEA is abundant in mammalian brain. As in the case of other FAEs, the production of PEA occurs through an on-demand synthesis within lipid bilayer (Cadas *et al*, 1996). PEA displays a considerable pharmacological potential when used as a drug (Lo Verme *et al*, 2005a; LoVerme *et al*, 2005b), or when its endogenous levels are enhanced by the inhibition of its catabolism (Solorzano *et al*, 2009).

The first evidence of the potential benefit of FAEs was found in the early 1940s, when Coburn and Moore (1943) reported the antipyretic properties of the dried chicken egg yolk in children with rheumatic fever. Ten years later, the lipid fraction from egg yolk was identified as the component responsible for this effect (Coburn *et al*, 1954), and it was

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also shown to be effective in reducing experimental allergy (Long and Martin, 1956). PEA was then identified to be the active component of this lipid fraction (Kuehl *et al*, 1957), and later, it was also found to be present in mammalian tissue (Bachur *et al*, 1965). After the discovery of its anti-inflammatory properties, PEA has been clinically tested and used to treat respiratory symptoms caused by the influenza virus (Masek *et al*, 1974).

The potential applications of this lipid amid remained overlooked; however, until the characterization of its anti-inflammatory (Mazzari *et al*, 1996), analgesic (Calignano *et al*, 1998, 2001), and anticonvulsant (Lambert *et al*, 2001) properties, and the identification of peroxisome proliferator-activated receptor- α (PPAR- α) as its primary intracellular molecular target (Lo Verme *et al*, 2005a; LoVerme *et al*, 2006; D'Agostino *et al*, 2007; Melis *et al*, 2008; Scuderi *et al*, 2011). PPARs are ligand-activated transcription factors belonging to the nuclear receptor superfamily. Three different subtypes have been described, namely PPAR- α , - β/δ and, - γ , all of which display distinct physiological functions dependent on their differential ligand-activation profiles and tissue distribution (Kliwer *et al*, 1994; Forman *et al*, 1996). Both PPAR- α and - γ subtypes regulate *in vivo* and *in vitro* inflammatory responses (Devchand *et al*, 1996; Delerive *et al*, 2001).

Both PEA and PPAR- α are present in the central nervous system (CNS), although their functions in this district, such as their possible neuroprotective potential, are largely unknown. We have reported that some of the analgesic properties of PPAR- α agonists (including PEA) have a likely centrally mediated component, and that the central PPAR- α activation may mediate a central tissue reprogramming away from central inflammation and sensitization (D'Agostino *et al*, 2007; D'Agostino *et al*, 2009; Sasso *et al*, 2011). Considering the anti-inflammatory properties of PEA, and the existence of a PEA/PPAR- α signaling at CNS level, we hypothesized that PEA may exert neuroprotective effect in an experimental model of neurodegenerative dementia. The aim of the present study was, therefore, to evaluate the effect of PEA treatment in an experimental model of AD, such as the intracerebroventricular injection of aggregated amyloid- β fragment 25–35 (Ab25–35) in mice.

MATERIALS AND METHODS

Animals

All procedures met the European guidelines for the care and use of laboratory animals (86/609/ECC and 2010/63/UE), and those of the Italian Ministry of Health (DL 116/92). Male wild-type (WT) and PPAR- α $-/-$ (B6.129S4-SvJae-Pparatm1Gonz) mice previously backcrossed to C57BL6 mice for 10 generations, were bred in our animal facility, where a colony was established and maintained by heterozygous crossing. Mice were genotyped as described on the supplier webpage (<http://jaxmice.jax.org>), with minor modifications. DNA was extracted from tails using the RedExtract kit (Sigma-Aldrich, Milan, Italy). All animals were maintained on a 12-h light/12-h dark cycle with free access to water and standard laboratory chow.

Chemicals

PEA and GW7647 (2-(4-(2-(1-cyclohexanebutyl)-3-cyclohexylureido)ethyl)phenylthio)-2-methylpropionic acid) were from Tocris (Bristol, UK). PEA and GW7647 were dissolved in PEG and Tween 80 2:1 (Sigma-Aldrich), and kept over night under gentle agitation with a micro stirring bar. Before injection, sterile saline was added so that the final concentrations of PEG and Tween 80 were 20 and 10% v/v, respectively.

Drugs were injected subcutaneously (s.c.) in a volume of 100 μ l per 30 g of body weight. The Ab25–35 and scrambled Ab25–35 (ScAb) peptide were from NeoMPS (Strasbourg, France) or Genepep (Montpellier, France). They were dissolved in sterile bi-distilled water at a concentration of 3 mg/ml, and stored at -20°C until use. To prepare 'aged' Ab25–35 peptide, peptides were aggregated by incubation at 3 mg/ml in sterile bi-distilled water at 37°C for 4 days. Peptides were administered intracerebroventricularly (i.c.v.) in a final volume of 3 μ l per mouse, as previously described (Maurice *et al*, 1996).

Study Design

In Figure 1 are reported the experimental setting used during this study. Amyloid aggregates were generated and i.c.v. injected as reported above. To evaluate its neuroprotective profile, PEA was first administered once a day, with treatment starting 3 h after the amyloid injection. After 7 days of treatment, mice were tested for spontaneous alternation in the Y-maze test. The day after (8 days after amyloid), they were submitted to the Morris water-maze task for 5 days. After a day of resting (day 13), the same mice were submitted (day 14) to a different water-maze procedure to specifically evaluate working memory (Figure 1 flow chart A). A different cohort of mice was treated, as reported above, to evaluate the effect of PEA on novel object recognition (NOR) test, and GW7647 on NOR and Y-maze test. To evaluate acute anti-amnesic effect, a different group of mice was treated i.c.v. with amyloid, as reported above, and then treated with a single dose of PEA 30 min before being tested (Figure 1 flow chart B). For molecular and biochemical analysis, mice were daily treated with PEA and euthanized after 5 days (Figure 1 flow chart C).

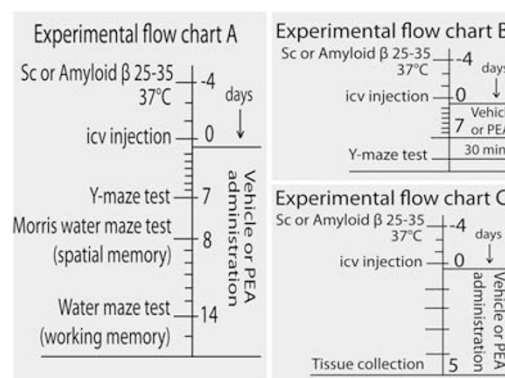


Figure 1 Experimental flow charts. (A) Experimental design for behavioral studies with daily treatment. (B) Experimental design for behavioral studies with acute treatment. (C) Experimental design for molecular, and biochemical and molecular analysis.

Spontaneous Alternation Performances

Each mouse, naïve to the apparatus, was placed at the end of one arm in a Y-maze (three arms, 40 cm long, 120° separate) and allowed to move freely through the maze during a single 5-min session. The series of arm entries, including possible returns into the same arm, was recorded visually. An alternation was defined as entries into all three arms on consecutive trials. The number of the total possible alternations was therefore the total number of arm entries minus two, and the percentage of alternation was calculated as (actual alternations/total alternations) × 100.

Spatial Learning and Memory Tests

The water maze was a circular pool (diameter 170 cm, height 60 cm). The water temperature, 23 ± 1 °C, light intensity, external cues in the room, and water opacity were rigorously reproduced. A transparent Plexiglas non-slippery platform (diameter 10 cm) was immersed under the water surface (1.5 cm) during acquisition trials. Swimming was recorded using a camera capture, and analyzed using a videotrack software (Any-maze, Stoelting, Wood Dale, IL, USA). The software divided the pool into four quadrants. Training consisted of four swims per day for 5 days, with about a 15-min inter-trial time. Start positions were pseudo-randomly selected, and each animal was allowed a 90-s swim to find the platform. Once the mouse reached the platform, it was left 20 s on the platform. The latency, expressed as mean ± SEM, was calculated for each training day.

A probe test was performed 1 h after the last swim on day 5. The platform was removed and each animal was allowed a free 60-s swim. The start position for each mouse corresponded to one of two positions remote from the platform location in counterbalanced order. The platform quadrant was termed the training quadrant, and the percentage of time spent in the training quadrant was determined.

Water-Maze Working Memory

A procedure for specifically assessing the short-term memory component was then performed for 3 days. The platform location changed every day, but not among trials. Each animal was allowed four trials per day, with a 2-min inter-trial time interval. Data represent the mean performance over days for each trial. The first trial is an informative sample trial, in which the animal is allowed to swim to the platform in its new location. During each other trial, the mouse involves its working memory component to increase its ability to reach the platform location (Yamada *et al*, 1999; Maurice *et al*, 2008). The decrease in latency between trials 1 and 4 could therefore be calculated to have a direct measure of the working memory performance.

NOR Test

The NOR test consisted of two sessions: a training session followed by a retention trial 24 h later. Mice were habituated to the testing arena for 2 consecutive days before the test. During the training session, two different objects (A and B) were placed in the testing arena. Each animal was allowed to explore the objects for 5 min. The mouse was considered to be

exploring the object when the head of the animal was facing the object, or the animal was touching or sniffing the object. The total time spent exploring each object was recorded by a trained observer blind to treatments, and expressed as percentage of total exploration time. In the retention session, one identical and one novel object (A and C) were used. A mouse was allowed to explore the objects for 5 min, and the time spent exploring each object was recorded. Exploration time was normalized as percentage of total exploration time. Preference for the novel object was considered as successful retention of memory for the familiar object.

Rotarod Test

Mice were habituated to the testing room for 30 min. Rotarod maximal r.p.m. was 40.0, and the acceleration time was 200 s. Mice were given six trials for 2 consecutive days.

Preparation of Total Tissue Protein Extracts and Western Blot Analysis

Hippocampi were homogenized on ice-cold lysis buffer (10 mM Tris-HCl, 20 mM, pH 7.5, 10 mM NaF, 150 mM NaCl, 1% Nonidet P-40, 1 mM phenylmethylsulphonyl fluoride, 1 mM Na₃VO₄, leupeptin, and trypsin inhibitor 10 µg/ml; 0.25 ml/50 mg tissue). After 1 h, tissue lysates were centrifuged at 14 000 r.p.m. for 15 min at 4 °C, and the supernatant was stored at -80 °C until use. Protein content was measured using bovine serum albumin (Sigma-Aldrich) as a standard. Protein lysates (70 µg) were separated on SDS-polyacrylamide gels, and membranes were incubated with anti-inducible nitric oxide synthase (iNOS; cat. no. 610432, BD Biosciences, from Becton Dickinson, Buccinasco, Italy), and anti-cleaved-caspase-3 (fragment p17) or -uncleaved-caspase-3 (fragment p30; cat. no. 9662, Cell Signaling, from Euroclone, Milan, Italy). Signals were visualized using ImageQuant 400 equipped with Quantity One Software 4.6.3 (GE Healthcare, Milan, Italy).

Lipid Peroxidation Measures

Mice were killed by decapitation, and brains were rapidly removed, weighed, and kept in liquid nitrogen until assayed. After thawing, brains were homogenized in cold methanol (1:10, w/v), centrifuged at 1000 g for 5 min, and the supernatant collected. Homogenate was added to a solution containing FeSO₄ 1 mM, H₂SO₄ 0.25 M, and xylanol orange 1 mM, and incubated for 30 min at room temperature. Absorbance was measured at 580 nm (A₅₈₀₁), and 10 ml of cumene hydroperoxide (CHP) 1 mM was added to the sample and incubated for 30 min at room temperature, to determine the maximal oxidation level. Absorbance was measured at 580 nm (A₅₈₀₂). The level of lipid peroxidation was determined as CHP equivalents according to: CHP equiv. = A₅₈₀₁/A₅₈₀₂ × (CHP (nmol)) × dilution, and expressed as CHP equiv. per wet tissue weight and than as percentage of control (Maurice *et al*, 1996).

Statistical Analysis

Data are expressed as mean ± SEM. Two-groups comparisons were made using the Student's *t*-test. When more than two

groups, data were analyzed using one-way ANOVA followed by the Dunnett's *post-hoc* multiple comparison test. Water-maze escape latencies were analyzed over trials using repeated measures ANOVA. For the probe test of Morris water maze, the time spent in the training quadrant was analyzed *vs* the chance level (15 s); for water-maze working memory, improvement between trial 1–4 was analyzed *vs* the chance level of 0 (no improvement); and for NOR, preference for the new object during the retention trail was analyzed *vs* the chance level of 50% using Wilcoxon signed-rank test. The level of statistical significance was $P < 0.05$.

RESULTS

PEA Counteracts the Ab25–35-Induced Learning Deficits

The possible neuroprotective effects of PEA were first evaluated on the appearance of Ab25–35-induced learning deficits. As previously reported (Maurice *et al*, 1996), acute i.c.v. injection of Ab25–35 induced, 7 days after, a marked impairment of spontaneous alternation performances compared with ScAb25–35. PEA was administered at 3, 10 or 30 mg/kg s.c. once per day, with treatment starting 3 h after the i.c.v. administration of the Ab25–35 peptide. Seven days after, PEA treatment resulted in a dose-dependent attenuation of the Ab25–35-induced spontaneous alternation impairments ($F_{(4,39)} = 5957$, $P < 0.0001$; Figure 2a); whereas no significant differences were found in the number of arm entries (Figure 2b). *Post-hoc* analysis revealed that the dose of 30 mg/kg was the most active, producing a complete prevention of the appearance of Ab25–35-induced spontaneous alternation deficits (Figure 2a).

Spatial and Reference Working Memory

The effect of Ab25–35 peptide injection on spatial and reference working memory was analyzed using the Morris

water-maze procedure. Animals received ScAb (9 nmol) or Ab25–35 (9 nmol), and were submitted 8 days after to a reference memory procedure. They had to learn the location of an invisible platform set at a fixed position, by performing four swimming per day for 5 consecutive days.

The acquisition profiles of ScAb-treated mice showed a significant diminution of swimming latencies, which is indicative of learning abilities. By contrast, the latencies showed by Ab25–35-treated mice were significantly higher for all training trials, as compared with ScAb-treated control group. PEA treatment attenuated this learning impairment in a dose-dependent manner (Figure 3a; effect of treatment $F_{(3,132)} = 6.88$, $p = 0.001$). Although the 3-mg/kg dose was ineffective (data not shown), the dose of 30 mg/kg almost completely normalized the learning profile of mice (Figures 3a and b). Supporting a protective role for PEA against Ab25–35 memory impairment, the probe test revealed that ScAb- and PEA (30 mg/kg)-treated mice explored preferentially the training quadrant at a similar extent (Figure 3c; $P < 0.05$). Although the analysis of swimming speed revealed no significant difference among groups (Figure 3d), we found that the Ab25–35-treated mice displayed a significant increase in tigmotactic behavior during the first trail of the first day (when mice were naïve to the apparatus) that is indicative of higher anxiety. PEA treatment normalized this anxiety behavior (Figure 3e; $F_{(3,23)} = 4.155$, $p = 0.0288$).

Animals were then submitted to the working memory procedure. During 3 days, they had to learn the location of a platform changing every day, by performing four swimming with a short inter-trial interval of 2 min. Animals learned the new platform location on sample trial 1 and then gradually improved their competence based on their working memory ability. Results showed that ScAb-treated WT mice were able to learn the variable platform location, as latencies significantly decreased over trials (Figure 3g). The Ab25–35-treatment impaired this acquisition. On trial 4, a significant difference in latency was measured between the ScAb- and Ab25–35 (9 nmol)-treated groups. Improvement was also analyzed in terms of difference between trial 1 and trial 4 (Figure 3h). The ScAb-treated group showed a latency decrease significantly higher than zero. PEA treatment showed a normalization trend respect to the impairment induced by Ab25–35, with a learning profile comparable to that one of ScAb25–35-treated mice (Figures 3g and h).

PEA Does not Exert Acute Anti-Amnesic Effect Against Ab25–35-Induced Memory Impairments

Behavioral deficits induced by Ab25–35 seem to be related to alterations in cholinergic and glutamatergic corticolimbic systems (Maurice *et al*, 1996; Olariu *et al*, 2001); therefore, drugs boosting these systems may exert acute anti-amnesic properties, in a way independent of their action on neurodegeneration progression. To test this possibility, we performed an additional experiment, and a possible acute anti-amnesic effect of PEA was examined in mice centrally injected with ScAb or Ab25–35 peptide 7 days before. The spatial working memory was examined in the Y-maze test. Mice received PEA (30 mg/kg) 30 min before the test session. In mice treated with Ab25–35, the alternation performance was about 55%, and PEA treatment did not

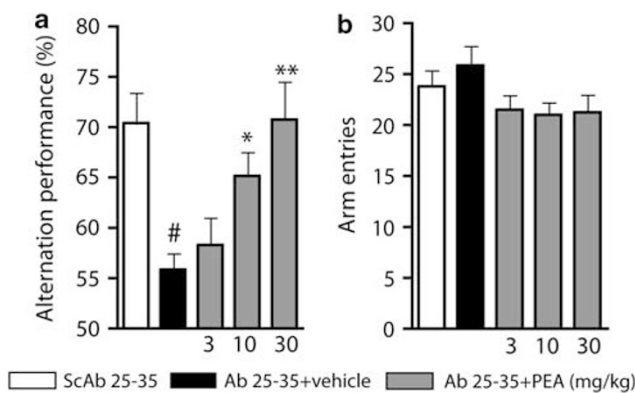


Figure 2 Neuroprotective effect of palmitoylethanolamide (PEA) on amyloid- β 25–35 (Ab25–35)-induced spontaneous alternation deficits in mice. (a) Alternation performances and (b) total numbers of arm entries. Mice were injected intracerebroventricularly (i.c.v.) with scrambled Ab25–35 (ScAb25–35) or Ab25–35 peptide (9 nmol). Mice (8–10 per group) received a daily subcutaneous (s.c.) injection of vehicle or PEA (3, 10, and 30 mg/kg). After 7 days, mice were examined for spontaneous alternation in the Y-maze apparatus. [#] $P < 0.05$ vs (ScAb25–35 + vehicle)-treated group; ^{*} $P < 0.05$, ^{**} $P < 0.01$ vs (Ab25–35 + vehicle)-treated group; Dunnett's test.

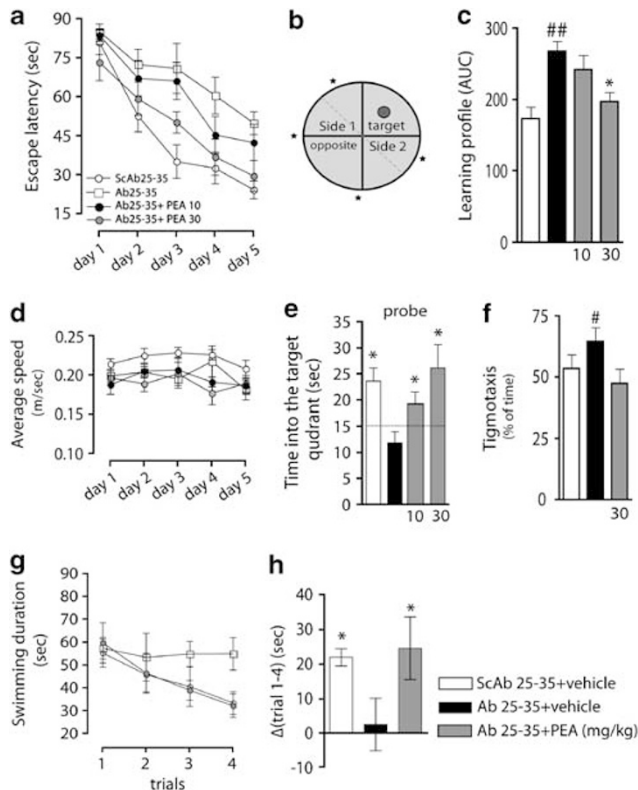


Figure 3 Neuroprotective effect of palmitoylethanolamide (PEA) on amyloid- β 25–35 (Ab25–35)-induced spatial learning and memory deficits in mice. Animals received a single intracerebroventricularly (i.c.v.) administration of scrambled Ab25–35 (ScAb25–35) or Ab25–35 (9 nmol), and they were daily treated with a subcutaneous (s.c.) injection of vehicle or PEA (10 and 30 mg/kg). After 8 days, mice were tested in the Morris Water Maze task. (a) Acquisition profiles: mice were submitted to acquisition of an invisible platform placed in a fixed location (target quadrant) with four swims per day during 5 days. (b) Apparatus drawing. (c) Acquisition profiles visualized as area under the curve (AUC) indicative of learning abilities. (d) Graph showing mice speed during the execution of the task. (e) Probe test sessions: 2 h after the last swim on day 5, the platform was removed, and the time spent in the target quadrant analyzed; $*P < 0.05$ vs a change level of 15 s (Wilcoxon signed-rank test). (f) Tigmotaxis during the first swim of the first day indicative of an anxiety-related behavior; $\#P < 0.05$ vs ScAb25–35 group. On day 14, mice were submitted to acquisition of a platform location changing daily (working memory procedure). (g) Acquisition profiles and (h) improvements between trial 1 and 4; $*P < 0.05$ vs a change of zero (Wilcoxon signed-rank test).

produce any improvement (Figures 4a and b). Central administration of ScAb or Ab25–35 peptide, or the subsequent i.p. treatment with PEA did not change the total number of arm entries (Figure 4b), speed (Figure 4c), or distance traveled during test execution (Figure 4d).

Neuroprotective Effects of PEA Against Ab25–35-Induced Toxicity are Dependent on PPAR- α

Previous studies have shown that PEA and other lipid amides, such as oleoylethanolamide, activate the anti-inflammatory nuclear receptor PPAR- α in a range of concentrations that normally occurs in mammalian tissue (Fu *et al*, 2003; Lo Verme *et al*, 2005a). To assess whether the absence of functional PPAR- α could affect the protective effect of PEA against Ab25–35-induced learning and

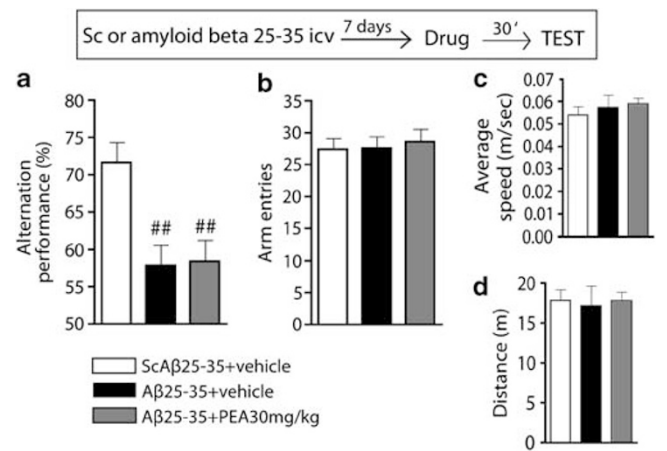


Figure 4 Anti-amnesic effect of acute palmitoylethanolamide (PEA) on amyloid- β 25–35 (Ab25–35)-induced spontaneous alternation deficits in mice. (a) Alternation performances, (b) total numbers of arm entries, (c) average speed during testing, and (d) distance traveled in the apparatus. Mice were injected intracerebroventricularly (i.c.v.) with scrambled Ab25–35 (ScAb25–35) or Ab25–35 peptide (9 nmol). After 7 days, mice (nine per group) received a single intraperitoneal injection of vehicle or PEA (30 mg/kg), 30 min before being examined for spontaneous alternation in the Y-maze apparatus. $\#\#P < 0.01$ vs (ScAb25–35 + vehicle)-treated group; Dunnett's test.

memory impairments, we tested the most effective dose, 30 mg/kg, in PPAR- α knock-out mice. As the role of PPAR- α on brain functions is largely unknown, we first verified if the genetic ablation of PPAR- α could produce phenotypic traits relevant to the present study. As shown in Figure 5, PPAR- α knock-out male mice and their WT littermate controls (8–9 weeks of age) did not differ when tested in the Y-maze apparatus (Figures 5a and b) or Morris water-maze task (Figures 5c and d). In a different group of PPAR- α knock-out and WT mice, we therefore tested the neuroprotective effect of PEA (30 mg/kg per day) against Ab25–35-induced learning and memory impairments. PEA did not protect PPAR- α knock-out mice against Ab25–35-induced learning and memory impairments. PEA failed to normalize spontaneous alternation performances of PPAR- α knock-out mice after 7 days of treatment (Figures 6a and b). To further validate the role of PPAR- α receptor, we also tested a synthetic and selective PPAR- α agonist, GW7647. GW7647 (5 mg/kg per day) reduced the spontaneous alternation deficit caused by Ab25–35 (Figure 6c). As expected, GW7647 was ineffective when tested in PPAR- α knock-out mice (data not shown). To extend our observations to a non-spatial form of cognition, we tested a different cohort of mice in the NOR test. Both PEA (30 mg/kg per day) and GW7647 (5 mg/kg per day) reduced the impairment induced by Ab25–35 in the NOR test evaluated after a retention interval of 24 h (Figure 6d). Increased exploration of the novel object was interpreted as successful retention of memory for the familiar object. Ab25–35-treated mice did not show a preference for the novel object after 24 h, whereas ScAb25–35, and PEA- or GW-treated mice had a percentage of novel object exploration significantly above the chance level of 50% (Figure 6d). We finally tested the same cohort of mice for motor coordination and motor learning by means of repeated rotarod testing. Mice were

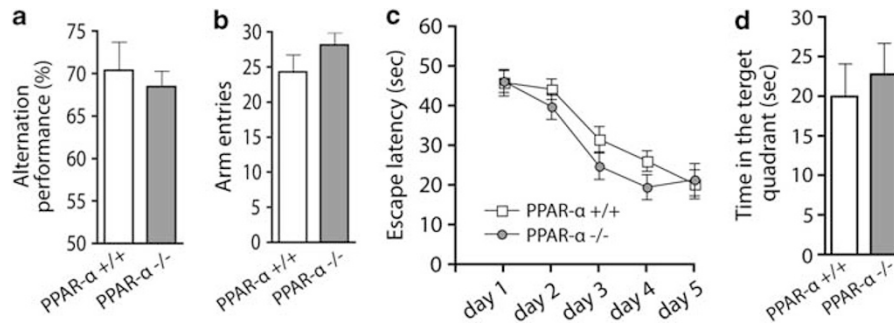


Figure 5 Memory profiles of peroxisome proliferator-activated receptor- α (PPAR- α) knock-out mice. Effect of the absence of functional PPAR- α on (a) spontaneous alternation performances, and (b) number of arm entries in the Y-maze test. (c) Spatial learning profile, and (d) spatial memory recall in the Morris Water Maze task. Wild-type (WT; +/+; $n=8$) and PPAR- α knock-out (-/-; $n=9$) littermate male mice were tested at 8–9 weeks of age. No significant differences were found.

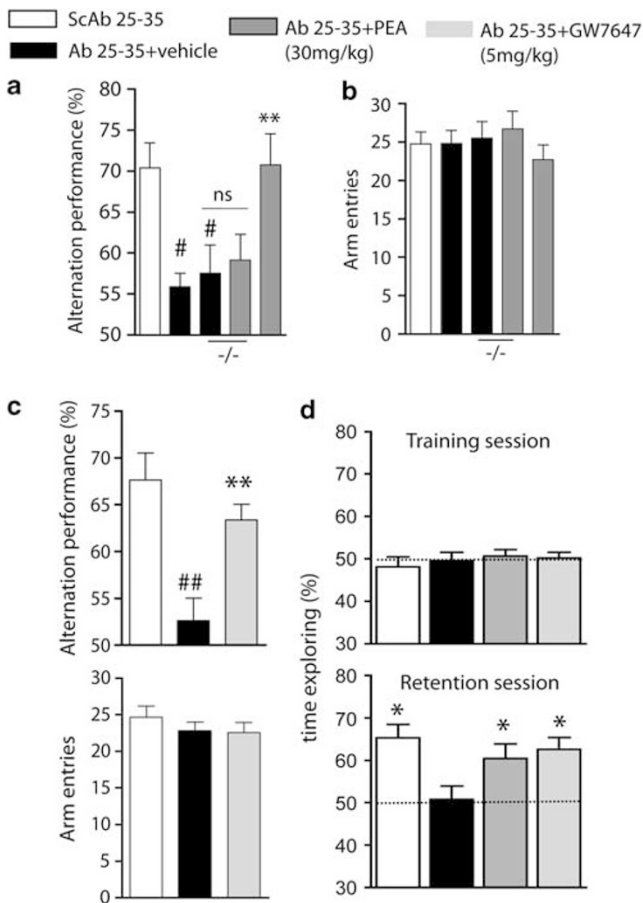


Figure 6 Neuroprotective effects of palmitoylethanolamide (PEA) against amyloid- β 25–35 (Ab25–35)-induced toxicity are dependent on peroxisome proliferator-activated receptor- α (PPAR- α). (a) Alternation performances and (b) total numbers of arm entries. PPAR- α wild-type (WT; +/+) and knock-out (-/-) mice were injected intracerebroventricularly (i.c.v.) with scrambled Ab25–35 (ScAb25–35) or Ab25–35 peptide (9 nmol). Mice (8–10 per group) received a daily subcutaneous (s.c.) injection of vehicle or PEA (30 mg/kg). (c) GW7647 (2-(4-(2-(1-cyclohexanebutyl)-3-cyclohexylureido)ethyl)phenylthio)-2-methylpropionic acid; 5 mg/kg per day, s.c.), a synthetic and selective PPAR- α agonist, mimics the effect of PEA. After 7 days, mice were examined for spontaneous alternation in the Y-maze apparatus. # $P<0.05$ or ## $P<0.01$ vs (ScAb25–35 + vehicle)-treated group; *** $P<0.01$ vs (Ab25–35 + vehicle)-treated group; Dunnett's test. (d) Novel object recognition (NOR) test. Both PEA and GW7647 counteract the impaired novel object recognition induced by Ab25–35 peptide (9 nmol). * $P<0.05$ vs a chance level of 50% (Wilcoxon signed-rank test).

tested for 2 consecutive days with six trails per day. Neither amyloid fragment nor pharmacological treatments impaired motor coordination or motor learning. Independently of treatment, all mice had increasing time performance over testing (data not shown).

PEA Reduces Amyloid-Induced Oxidative Stress and Apoptotic Markers

Biochemical parameters of amyloid toxicity were also analyzed in the hippocampus extracts to validate the neuroprotective activity of PEA. Amyloid peptides, and particularly Ab25–35, induce a massive oxidative stress in forebrain structures. We therefore analyzed some representative biochemical and molecular markers with the aim to support the above reported behavioral observations. As previously reported, Ab25–35 significantly increases the levels of lipid peroxidation, protein nitrosylation, induces iNOS expression, and results in cell death through caspase-dependent apoptosis pathways (Figure 7). PEA (30 mg/kg) significantly reduced the Ab25–35-induced lipid peroxidation increase (Ab 25–35 vs ScAb25–35, $P<0.01$; Ab25–35 vs Ab25–35 + PEA, $P<0.05$; Figure 7a). PEA (30 mg/kg) significantly reduced the Ab25–35-induced iNOS induction (iNOS/ β -actin ratio \pm SEM ($n=4$), ScAb25–35: 1.055 ± 0.074 , Ab25–35: $1.763 \pm 0.12^{##}$, Ab25–35 + PEA: $1.368 \pm 0.17^*$ ($^{##}P<0.01$ vs ScAb25–35; $^*P<0.05$ vs Ab25–35, Dunnett's test); Figure 7c). PEA (30 mg/kg) attenuated ($P=0.055$) the Ab25–35-induced activation of caspase3 (cleaved caspase3/caspase3 ratio \pm SEM ($n=4$), ScAb25–35: 0.435 ± 0.029 , Ab25–35: $0.630 \pm 0.031^{\#}$, Ab25–35 + PEA: 0.515 ± 0.037 ($^{\#}P<0.05$ vs ScAb25–35, Dunnett's test); Figure 7d). Overall, this set of data adds further support to the neuroprotective effect of PEA observed during behavioral studies.

DISCUSSION

The present study discloses a previously unreported therapeutic effect of PEA in an experimental model of AD.

There are two major hallmarks of AD: neurofibrillary tangles and amyloid aggregates. In rodent brain, Ab25–35 induces, after acute injection or chronic infusion, biochemical changes, morphological alterations, and behavioral impairments reminiscent of AD physiopathology

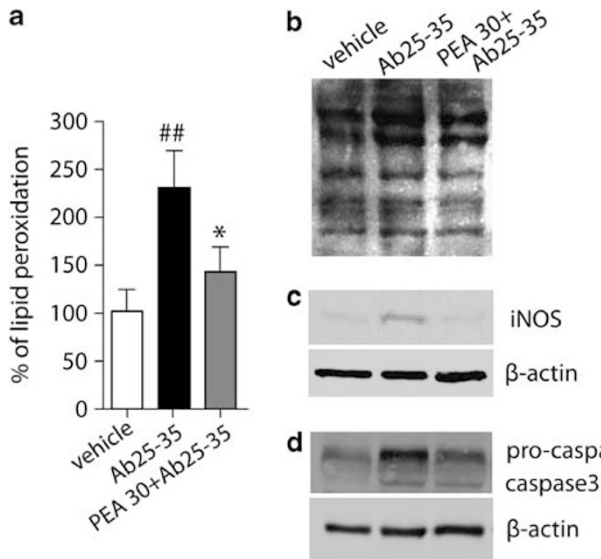


Figure 7 Neuroprotective effects of palmitoylethanolamide (PEA, 30 mg/kg) measured using biochemical markers in the hippocampus in scrambled amyloid- β 25–35 (ScAb25–35)- and Ab25–35-injected mice, daily treated with vehicle or PEA (30 mg/kg, subcutaneously (s.c.)): (a) lipid peroxidation levels, (b) protein nitrosylation levels; (c) inducible nitric oxide synthase (iNOS) expression; (d) caspase-3 expression. Mice were administered s.c. with PEA or vehicle 3 h after the intracerebroventricularly (i.c.v.) injection of ScAb or Ab25–35 peptide (9 nmol) and once a day for 5 days.

(Maurice *et al*, 1996; Delobette *et al*, 1997). By using Y-maze, water maze, and object recognition tasks, we have shown that a daily treatment with PEA reduces learning and memory deficits caused by the central injection of the pre-aggregated Ab25–35. The protective profile of PEA is dose-dependent, and the dose of 30 mg/kg fully prevents the working memory-like impairment evaluated as Y-maze alternation performances. The same dose almost completely rescues spatial learning and memory deficits induced by Ab25–35 when tested in the water-maze task. The restoration of these two aspects of memory performances is of particular interest, as they are highly representative of common clinical features of AD. Patients with AD show decline in immediate memory span and in working memory as the disease progresses, and they often experience difficulties with spatial orientation in everyday activities (Hort *et al*, 2007).

Among the different stages of the disease, neuroinflammation and oxidative stress have an important role in pathogenesis. Inflammatory mediators, such as TNF α , IL-1, and IL-6, have negative effects on hippocampal long-term potentiation that is considered a cellular model of learning and memory (Bellinger *et al*, 1993; Stellwagen and Malenka, 2006; Schmid *et al*, 2009; Tsai *et al*, 2010). Although there exists a vast literature on the anti-inflammatory properties of PEA, our work is, to the best of our knowledge, the first *in vivo* study that shows the ability of this endogenous acylethanolamide to rescue experimentally induced memory deficit. Previous studies have shown that PEA and other fatty acid acyl ethanolamines activate the anti-inflammatory nuclear receptor PPAR- α in a range of concentrations that occur normally in tissues (Fu *et al*, 2003; Lo Verme *et al*, 2005a). The expression of this nuclear receptor is necessary

for exogenous PEA to exert its anti-inflammatory (Lo Verme *et al*, 2005a) and antioxidant effects (Raso *et al*, 2011). Our study showed the obligatory role of PPAR- α for the neuroprotective effect of PEA against amyloid toxicity. As expected, our work also showed that a synthetic PPAR- α agonist mimics the pro-cognitive effect of PEA. These data add further support to the notion that this nuclear receptor is the primary molecular target of PEA, and extend our previous finding about the physio-pathological role of PEA/PPAR- α signaling in the CNS (D'Agostino *et al*, 2007, 2009). With reference to AD, our behavioral observations in PPAR- α knock-out mice are also supported by recent *in vitro* evidence showing that PEA reduces amyloid-induced astrogliosis through a PPAR- α -dependent mechanisms (Scuderi *et al*, 2011). As part of its transcriptional activity, PPAR- α also induces the expression of genes involved in mitochondrial and peroxisomal fatty acid β -oxidation (Berger and Moller, 2002; Willson *et al*, 2000). Peroxisomes have a crucial role for ROS and lipid metabolism, and their importance in brain physiopathology is well established. We hypothesize that PEA acting at PPAR- α receptor could increase the number of peroxisomes and/or the activity of the peroxisomal matrix protein catalase counteracting the redox perturbation following the amyloid excess. Recent studies showed that the PPAR- α agonist fenofibrate is able to promote the survival of cortical neurons in the presence of the NO donors *in vitro* (Gray *et al*, 2011). Moreover, the expression of neuronal peroxisomal and peroxisome-related proteins (eg, PPAR- α) has been reported to change after a challenge with Ab *in vitro* (Cimini *et al*, 2009), supporting the notion that peroxisomal and peroxisome-related proteins may constitute a cellular defense against Ab-mediated oxidative injury. According with this scenario, we demonstrated that behavioral effects exerted by PEA are corroborated by molecular and biochemical evidences, in that PEA reduced the increase of lipid peroxidation, protein nitration, iNOS expression, and the induction of pro-apoptotic pathways induced by Ab25–35.

Although the molecular mechanism underlying the neuroprotective effect of PEA needs further investigations, we provided evidence suggesting that PEA acting at PPAR- α rescues behavioral impairments that can mimic some traits of AD. Considering that PEA is currently used to treat chronic pain, our findings can be worthy of clinical consideration, and they can have immediate utility, at least in the early stages of this pathology.

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DISCLOSURE

The authors declare no conflict of interests.

Authors contribution

GD designed experiments, carried out behavioral studies, analyzed data, and wrote the manuscript. RR, CA, CC

performed research. RR and RM participated in the design of the study. AC conceived the study, participated in its design and coordination, provided financial sources, and wrote the manuscript.

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