Effects of Long-Term Pioglitazone Treatment on Peripheral and Central Markers of Aging

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Effects of Long-Term Pioglitazone Treatment on Peripheral and Central Markers of Aging

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Abstract

Background: Thiazolidinediones (TZDs) activate peroxisome proliferator-activated receptor gamma (PPARγ) and are used clinically to help restore peripheral insulin sensitivity in Type 2 diabetes (T2DM). Interestingly, long-term treatment of mouse models of Alzheimer’s disease (AD) with TZDs also has been shown to reduce several well-established brain biomarkers of AD including inflammation, oxidative stress and Aβ accumulation. While TZD’s actions in AD models help to elucidate the mechanisms underlying their potentially beneficial effects in AD patients, little is known about the functional consequences of TZDs in animal models of normal aging. Because aging is a common risk factor for both AD and T2DM, we investigated whether the TZD, pioglitazone could alter brain aging under non-pathological conditions.

Methods and Findings: We used the F344 rat model of aging, and monitored behavioral, electrophysiological, and molecular variables to assess the effects of pioglitazone (PIO-Actos® a TZD) on several peripheral (blood and liver) and central (hippocampal) biomarkers of aging. Starting at 3 months or 17 months of age, male rats were treated for 4–5 months with either a control or a PIO-containing diet (final dose approximately 2.3 mg/kg body weight/day). A significant reduction in the Ca2+-dependent afterhyperpolarization was seen in the aged animals, with no significant change in long-term potentiation maintenance or learning and memory performance. Blood insulin levels were unchanged with age, but significantly reduced by PIO. Finally, a combination of microarray analyses on hippocampal tissue and serum-based multiplex cytokine assays revealed that age-dependent inflammatory increases were not reversed by PIO.

Conclusions: While current research efforts continue to identify the underlying processes responsible for the progressive decline in cognitive function seen during normal aging, available medical treatments are still very limited. Because TZDs have been shown to have benefits in age-related conditions such as T2DM and AD, our study was aimed at elucidating PIO’s potentially beneficial actions in normal aging. Using a clinically-relevant dose and delivery method, long-term PIO treatment was able to blunt several indices of aging but apparently affected neither age-related cognitive decline nor peripheral/central age-related increases in inflammatory signaling.

Introduction

Adjunct therapy against type 2 diabetes mellitus (T2DM) with thiazolidinediones (TZDs) is on the rise, with increasing numbers of patients prescribed the TZDs rosiglitazone (ROSI, Avandia®) or pioglitazone (PIO, Actos®). These agents are in the top 50 prescribed drugs in North America, and together account for approximately 20 million prescriptions (2008 data, rxlist.com). Results from over two decades of studies have shown that untreated T2DM can negatively impact brain function. Depending on the severity and the duration of the disease, as well as on the age of the individual, the condition is associated with varying degrees of cognitive deficits, motor dysfunction, and depression [1,2,3,4,5,6,7]. While aging worsens the impact of diabetes on cognitive function, it is not clear how diabetes and accompanying peripheral metabolic dysregulation exacerbate this process.

Proposed mechanisms underlying cognitive decline when aging and diabetes coexist include insulin resistance, vascular disease, and inflammation resulting from the release of adipose tissue-derived cytokines. In clinical and animal studies, the brain, and the hippocampus in particular, appear sensitive to peripheral cytokine levels or metabolic stressors [8,9,10,11], with enhanced sensitivity seen in aged animals [8]. Given the role of the hippocampus in memory acquisition, processing and consolidation (reviewed in...
inflammation within the structure likely contributes to memory and cognitive deficits with age and/or AD. Nevertheless, almost nothing is known about the mechanisms through which peripheral metabolic dysregulation as those seen in T2DM impact hippocampal function and cognition.

TZDs are best known for their peripheral actions, where these synthetic PPARγ agonists selectively bind nuclear receptors and enhance lipid accumulation in adipocytes, thereby helping to decrease free fatty acid and lipid levels in plasma [14,15,16,17,18]. This mechanism helps to reestablish insulin sensitivity in T2DM by working on fat, liver, and muscle tissues. Additionally, TZDs are compounds with significant anti-inflammatory actions [19,20]. Recent evidence indicates that TZDs can have beneficial central effects. In particular, ROSI was shown to improve cognition and verbal memory in patients with mild cognitive impairment (MCI) [21]. Further, a recent preliminary study on patients diagnosed with MCI and diabetes also reported improved cognition following 6-months of PIO [22]. Finally, AD patients lacking the ApoE4 allele appear to be selectively sensitive to the beneficial effects of chronic treatment with ROSI [23]. The mechanisms underlying these effects in humans are not clear, but are likely to reflect changes in inflammation, vascular function, insulin and/or glucose levels, energy metabolism or beta amyloid clearance. Whether these improvements are due to changes in the periphery, direct effects in the brain, or some combination, is still unknown. In AD animal models (APPV717I, Tg2576, 3×TG), TZDs have been shown to decrease Aβ deposition [24,25,26,27] (but see [28]), inflammation [24,29,30], and oxidative stress [28].

The predominant information we have regarding the beneficial effects of TZDs in the brain comes from clinical studies in AD patients [21,23]. However, given that most patients with diabetes do not have AD, at least in the earlier stages of the disease, it seems important to determine what the potential effects of these drugs are in the context of normal aging, or in the absence of clinically defined cognitive deficits. Compared to research conducted in AD models, however, little is known about the functional consequences of TZDs on cognition in animal models of normal aging. Therefore, the present studies were undertaken to determine whether PIO, the more brain permeant TZD [26], confers significant benefits within the context of normal brain aging.

Further, because of our prior work identifying new roles of ROSI and PIO in cultured neurons [31], and the work of others suggesting that targets of TZD actions may include Ca2+-mediated pathways in the brain [32,33,34], we investigated select biomarkers of aging including the Ca2+-dependent afterhyperpolarization (AHP), long-term potentiation (LTP), and hippocampal-dependent spatial memory. Other examined variables in the brain and in the periphery included inflammatory cytokine levels, hippocampal gene signatures, and insulin signaling. Our results suggest that at the dose and duration tested, PIO caused expected beneficial effects including reduced peripheral insulin and lipid marker levels, and reduced a central biomarker of aging, namely the AHP. However, other major markers of aging, including increased inflammatory signaling (based on cytokine array measures in the periphery and microarray measures in the hippocampus), impaired cognition, and altered synaptic plasticity, were not altered with PIO treatment.

Methods

Ethics Statement

All procedures were carried out under a University of Kentucky IACUC approved protocol and are in accordance with NIH guidelines for the care and use of laboratory animals.

Animals

Thirty-six male F344/NIA rats were purchased (Harlan, Indianapolis, IN) in 2 groups of 18 with a 6 week stagger between groups. All animals were fed TD94045 diet (Harlan Teklad, Madison, WI) for one week prior to initiation of PIO or control diets. To limit potential cohort effects across two separate animal purchases, treatment groups were balanced across both cohorts. Each group consisted of 8 young (3 months old), and 10 aged (17 months old) animals. Twelve animals from each group were assigned to either the young control (YC, n = 4), young PIO (YP, n = 4), or aged control (AC, n = 4) treatment groups, and the remaining six animals were placed in the aged PIO (AP) group. In all, the study was comprised of 8 YC, 8 YP, 8 AC, and 12 AP animals. Animals were maintained on the diets for 15–20 weeks, and were 7–8 months old, and 21–22 months old at the time of study completion. Five aged animals died in the course of the study. One AP animal had to be euthanized because of an unresolved mandibular/eye infection, and two AP animals stopped eating, lost considerable weight and were euthanized. Based on two gross necropsy reports, another AP animal died of chronic renal failure, and 1 AC animals died of granular lymphocytic leukemia, both major causes of mortality in the aging F344 [35]. The thirty one remaining animals were active, well-groomed and appeared healthy, and were used for behavioral and electrophysiology studies.

Blood collections and analysis

Over the course of the study, three in vivo glucose measures were taken. Animals were placed in a decapacine® restraint (Braintree Scientific, Braintree, MD) while their tails were washed with warm soapy water and dried under a heat lamp. The lateral tail vein was pricked with a 22 gauge needle and a FreeStyle Lite glucometer (Abbott Diabetes Care Inc., Alameda, CA) was used to measure blood glucose levels (mg/dL). Trunk blood from twenty nine animals was collected at the time of hippocampal slice preparation (two samples were lost). Briefly, 2–3 mL was collected in a BD Vacutainer SSD centrifuge tube and allowed to clot at room temperature for one hour. To collect serum, samples were centrifuged at 4000 rpm for 10 min. Half of the serum was sent on dry ice for standard chemistry panel analysis (Comparative Pathology Laboratory, University of California Davis, CA). The remaining serum was frozen (−80°C) and later used to monitor insulin concentrations using the manufacturer’s protocol for an ELISA-based assay (Millipore, Billerica, MA) and our bioassay reader (HTS plus 7000, Perkin Elmer, Wellesley, MA), as well as to monitor for the presence of three proinflammatory cytokines using a Multiplex Bio Assay Analyzer (Millipore).

Insulin Receptor signaling

Liver and brain cortices were used to quantify total and phosphorylated insulin receptor levels according to the manufacturer’s protocol (Calbiochem, San Diego, CA), and using duplicates for each sample. Frozen samples were removed from the −80°C freezer and thawed on ice. After a 2 min homogenization period in PBS followed by centrifugation at 300 rpm for 5 min, the pellet was resuspended in Cell Extraction Buffer (BioSource FNN0011) and left to lyse for 30 min with vortexing every 10 min. At the end of this process the suspension was centrifuged at 14K rpm for 10 min and the protein content in the supernatant was determined using a Bradford assay. Detection of the phosphorylated insulin receptor (IR) was accomplished following the PhosphoDetect ELISA kit protocol (CBA038, Calbiochem). Total IR present in the samples was measured using IR β-subunit ELISA Kit (CBA039, Calbiochem). Briefly,

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samples containing the same concentration of total protein were incubated for 2 h in a 96 wells plate coated with IR β-subunit-specific monoclonal antibody. After washing, an antibody specific for IR phosphorylated at Tyr1162/1163 (CBA039) or specific for IR β-subunit (CBA039) was added (detection antibody). The excess detection antibody was removed after 1 h and a horseradish peroxidase-conjugated antibody (anti-rabbit Ig-HRP) was added to the wells for 30 min. Following a final washing step to remove the excess anti-rabbit Ig-HRP, a substrate was added and absorbance was read at 450 nm.

**Experimental diets**

Pioglitazone (PIO-Actos®) was purchased through our DLAR facility, and was incorporated into standard, color-coded purified rodent diets (TD94045 Harlan Teklad), TD94045 was chosen to approximate the NIH31 diet fed to animals since adulthood (18.8% Kcal from protein, 63.9% Kcal from carbohydrates, and 17.2% Kcal from fat vs. 21% Kcal from protein, 62% Kcal from carbohydrates and 14% Kcal from fat in the NIH31). Because of differences in animal weights and food consumption, two PIO dosages based on animal food consumption and body weight (each measured 3 times a week throughout the course of the study) were chosen: 2.3 mg/Kg/day. Actos® is available in 15–45 mg tablets and in humans, serum concentrations following a single 30 mg oral dose reach approximately 1 μg/mL [36], and following a 10 day treatment with once a day 45 mg oral dosing regimen, peak plasma level was measured at 1.6 μg/mL, as reported in The pharmacological basis of therapeutics [37]. The dose used here (~2.3 mg/Kg/day), is relatively low compared to other reports in animals, and we estimate steady state blood PIO levels at approximately 1.3 μg/mL. This is based on published human clearance values for PIO (1.2 mL/min/Kg) given that the pharmacokinetic properties of PIO in rodents are not available.

**Electrophysiology, AHP and LTP**

Electrophysiological data were recorded between 1.5 and 5 weeks after the end of the Morris water maze training to limit the impact of learning and arousal on transient (about one week [38]) hippocampal excitability changes, and because only a single animal could be monitored daily on the electrophysiology setup. Hippocampal slices slices taken from the medial half of the hippocampus were obtained according to previously published protocols [39], briefly, animals were anesthetized in a CO2 chamber prior to decapitation, hippocampi were removed and transverse slices prepared (350 μm in ice cold low-calcium artificial cerebrospinal fluid (ACSF) composed of (in mM): 128 NaCl, 1.25 KH2PO4, 10 Glucose, 26 NaHCO3, 3 KCl, 0.1 CaCl2, 2 MgCl2) using a Vibratome® (series 3000, TPL, Saint Louis, MO). For AHP experiments, slices were then transferred to a heated (32°C) interface-type chamber, maintained in oxygenated (95% O2, 5% CO2) normal-calcium ACSF containing 2mM CaCl2 and 2mM MgCl2 for least 2 h prior to recording. For LTP experiments, a modified ACSF containing 2.5 mM CaCl2 and 1.3 mM MgCl2 was used.

**AHP experiments**

Each hippocampal slice was placed in a recording chamber (RC22C, Warner Instruments, Co., Hamden, CT) and maintained in a continuous flow of oxygenated ACSF pre-heated at 32°C using a TC2Bip/HPRE2 in line heating system (Cell Micro Controls, Northfolk, VA). This setup was mounted on the stage of a Nikon E600F inverted microscope. As previously described [39], cells were impaled with sharp microelectrodes filled with 2M KMeSO4 and 10mM HEPES, pH 7.4 (tip resistance 100.2±4.7 MΩ), pulled from borosilicate glass capillaries (World Precision Instruments, Sarasota, FL) using a P90 pipette puller (Sutter Instruments, Novato, CA). All experiments were performed in current clamp mode with bridge balance compensation and capacitance neutralization. Signal was digitized at 2 kHz and low-pass filtered at 1 kHz. Recordings of membrane input resistance (IR) were obtained in response to 800 ms, 200 pA hyperpolarizing current injections using an Axoclamp 2B amplifier (Molecular Devices, MDS, Toyota, Canada) while holding the cell at −70 mV. To generate an afterhyperpolarization (AHP) cells were held at −65 mV (baseline) and depolarized with a 100 ms current injection in order to generate three Na+ action potentials. AHPs were elicited every 30 s and at least 6 AHPs were averaged for each cell. The medium AHP (mAHP) was measured as the peak hyperpolarization immediately after the offset of the depolarizing current injection, the slow AHP (sAHP) was measured 800 ms after the end of the current injection. The AHP duration was measured from the end of the depolarizing step until return to baseline. Neurons with input resistance <40 MΩ, holding current >500 pA and action potential height <0 mV, were excluded from this study. Data were acquired using pClamp 8.0 (Molecular Devices) software through a Digidata 1320A A/D converter (Molecular Devices), and quantification of potentials (e.g., amplitude and duration of AHPs) was obtained with Clampfit software (Molecular Devices).

**LTP experiments**

Slices were recorded from within a heated and oxygenated interface-type chamber (32°C) after at least 2 h of recovery. Recording electrodes were 5–10 MΩ (filled with ACSF), and the stimulating electrode was made from twisted insulated stainless steel wire (A-M Systems, Inc. Everett, WA). Stimulation (baseline and LTP) was delivered through a pair of SD9K stimulators (Astro Med Inc., Grass Instr., Warwick, RI). During baseline and after LTP induction, stimulation rate was set to 0.33 Hz. LTP was elicited using a 2 s theta-burst pattern such that eight pulses at 100 Hz (50 ms each) were delivered at 5 Hz [40] in stratum radiatum. Stimulation intensity was set at 33% of the maximum response (determined from an I/O curve prior to LTP induction). This LTP induction protocol was chosen to accentuate the age-dependent decrease in LTP induction and maintenance [13,41,42,43,44,45,46]. For each slice, baseline EPSP slopes averaged across the 20 min prior to LTP induction were used to normalize EPSP slopes after LTP induction. Post-tetanic potentiation (PTP) was derived from EPSP slope measures taken immediately after LTP induction (2 min average) and LTP was derived from EPSP slopes averaged 25–30 min after tetanization (5 min average). A slice was removed from the analysis if the percent change in EPSP slope during the baseline period fluctuated more than 25% (up or down), or if the EPSP was contaminated with a spike following LTP induction.

**Morris Water Maze (MWM)**

The maze (black circular pool, 190 cm in diameter) was placed equidistant (~60 cm) to a continuous wall of black curtains hanging from the ceiling, making the environment relatively neutral. Three high contrast black and white cues (90 cm ×90 cm, representing a circle, triangle and vertical lines), were placed on the curtains. Each day, the animals were placed in one of the four quadrants; this allowed the animal to learn to map the position of the escape platform relative to the cues on the curtain. Pool temperature was maintained at 25–26°C. One quadrant contained
a 15 cm diameter escape platform covered with black neoprene for improved traction. Illumination in the room was set such that the Videomex-V water maze monitoring system (Columbus Instrument, Columbus, OH) could reliably monitor animal movements with no artifacts.

For all training days (days 1–4), three trials were run with animals placed in the pool for 60 s. During the early training days (1–2), animals that did not find the platform within the allotted 60 s were gently guided to the platform. All animals were allowed to stay on the platform for 60 s. Following this 60 s rest period, animals were taken to a drying cage outside the MMW enclosure for 45 s, and then returned to the MMW for a second trial. The intertrial interval was approximately 165 s, with ~60 s of swimming and 105 s of rest. On day 1, three cue trials were run with animals released in the same quadrant for each trial. In these first trials, a hanging white cup was positioned over the platform (~30 cm above the water surface), and the platform was set right at, or slightly above the water level, providing the animals with salient cues for a mean of escape. On the next 3 days of training (days 2–4) animals were placed in a different starting location along the periphery of the maze for each trial (3 trials/day), and the platform was submerged (~2.5 cm below the water surface). Animals were never placed in the pool within the quadrant containing the platform. On the last day (day 5), a single 60 s probe trial was run with the platform removed. Animals were considered visually impaired if they failed to find the platform within the allotted 60 s on 2 out of 3 trials on day 1 (cue learning), and on 3 out of 3 trials for learning days 3 and 4. Using this criterion, 6 aged animals, 3 AC and 3 AP, were excluded from the behavioral analysis.

Microarrays

**Microarray analysis.** During preparation of hippocampal tissue for electrophysiology, dorsal and ventral quarters from both hippocampi were placed in RNase-free Eppendorf tubes on dry ice, and transferred to a −80°C freezer until further use. For each animal (n = 7–8/group), this tissue was treated as a single sample. Each sample underwent RNA extraction, purification, and cDNA labeling separately, as described previously [47,48,49,50], according to standard Affymetrix procedures. Labeled cDNA for each region from each subject was individually hybridized to an Affymetrix rat microarray (RAE230 2.0, 31099 probe sets). All arrays passed standard Affymetrix quality control: GAPDH 3′–5′ ratio 1.07±0.005, RawQ 2.73±0.02, Background noise 79.3±0.6. Scaling factor, based on target intensity of 500, YC: 0.95±0.03, YP: 0.91±0.04, AC: 0.92±0.04, AP: 0.90±0.04; as well as % Present: YC: 69.7±0.44, YP: 70.0±0.33, AC: 69.8±0.31, AP: 70.1±0.35 were not significantly different across treatment groups (two-way ANOVA p>0.4 for main effects of age, drug, and interaction). Visual inspection of residual signal images (Affy PLM [51]) revealed no major image defects.

The MASS probe level algorithm (Gene Expression Console v 1.1, Affymetrix) calculated signal intensity and presence/absence calls. Only unique probe sets/genes with ‘A’ grade annotation and >2 presence calls were retained for further analysis. Values were transferred to Excel (2007, Microsoft), Bioconductor [52], Multi-Experiment Viewer (MEV, [53]) and the DAVID suite of bioinformatic tools [54] for subsequent analysis. All data are MIAME compliant and the raw data has been deposited in a MIAME compliant database (Gene Expression Omnibus - GEO accession #GSE20219).

**Proinflammatory Cytokine Analysis**

Sample sera were analyzed by multiplex bead array using Milliplex rat cytokine kits (RCYTO-80K) according to procedures recommended by the manufacturer (Millipore). Just prior to analysis, frozen sera were thawed and maintained on ice throughout the assay setup. Briefly, all serum samples were diluted 1:5 in sample diluent and were then incubated in duplicate overnight with capture beads specific for IL-1β, IL-6 and TNFα. Beads were subsequently washed and incubated for 2 h with biotin-conjugated detection antibody and then for 30 min with streptavidin-phycocerythrin. Bead fluorescence was then analyzed on a Luminex 100 IS Multiplex Bio-Assay Analyzer. Cytokine concentrations were determined from standard curves of recombinant rat cytokines in which 4-parameter logistic curve fitting analysis was used. All cytokines are reported as pg/ml ± S.D.

**Statistics**

For all electrophysiological measures presented here, outliers were removed based on the 2 SD rule. For main effects of age or treatment on these measures, two-way ANOVA with Bonferroni post-hoc analyses were used. Behavioral and chemical panel analyses also used two-way ANOVA. For genechip analyses, the filtered genes (7922 probe sets) were tested statistically by two-way ANOVA (main effects of age and drug, as well as interaction) using the False Discovery Rate (FDR [55]) to gauge multiple testing error (see Results) and post hoc Fisher’s Protected Least Significant Difference (PLSD) was used for all-pairwise comparisons among genes with significant main effects/interactions. For all statistical analyses, significance was considered present of p values were less or equal to 0.05.

**Results**

**Chemical panel**

Analysis of blood serum obtained at time of hippocampal dissection for each animal, revealed a significant effect of PIO on lipids, including decreased total cholesterol (F1,125 = 4.46, p<0.05) and triglycerides (F1,125 = 15.9, p<0.001). Insulin levels also were significantly reduced by PIO in both age groups (F1,125 = 16.7, p<0.0005), consistent with similar human studies reported in the literature [56,57,58,59,60]. A significant age-dependent increase in HDL was seen (F1,125 = 5.26, p<0.05) but was not sensitive to PIO. In vivo glucose measures (from tail pricks) did not change with age or treatment during the course of the study (Fig. 1C), and analysis of sera obtained at time of dissection showed no glucose level change (see Table 1). Levels of alanine aminotransferase, a marker of liver health, were not affected by age or treatment. Interestingly, the triglyceride (TG) to HDL ratio (TG/HDL) was reduced to the same degree by PIO (~50%) in young and aged animals (YC: 4.1, YP: 2.3, AC: 2.2, AP: 1.2), suggesting that our use of two PIO diets formulated at different drug concentration for the younger and older animals to compensate for different weights, had similar impact on peripheral lipids. Further, because this ratio is considered a surrogate marker for insulin resistance in humans, it seems PIO levels here where within a therapeutically-relevant range, reducing an indirect, yet classic clinical marker of insulin resistance. Therefore, in the F344 rat and at the dose tested, PIO provided significant reductions in lipid profiles and insulin levels, in a manner similar to that seen in clinical studies [56,58,60].

**Organ and animal health**

Animal weights were not different by age or treatment group by the end of the study, indicating younger animals on either the control or the PIO diet gained comparable weight (Fig. 1A). Because of prior reports that TZDs might be associated with adverse cardiovascular outcome [61], we measured heart weights
in all animals. Normalized heart to body weight ratios showed no difference across groups (Fig. 1 B). Upon examination, no gross adiposity or cirrhosis was noted, and no internal organs showed distinguishable signs of pathology in the PIO group. Overall, it seems PIO was well-tolerated, with no observed anomaly detected in the animals’ coat, eyes or skin, as well as internal organs, body, or heart weights.

Dosage

PIO doses calculated from individual body weights and averaged food consumption (taken three times a week across the duration of the study) were 2.6±0.12 mg/Kg/day for the young PIO group and 1.9±0.07 mg/Kg/day for the aged PIO group. This difference was significant (t-test, p<0.05).

Behavioral characterization

Two-way ANOVA on conventional outcome measures associated with the Morris water maze (MWM) including path length and latency to platform, were tested for significance across the 4 days of training. No age or treatment differences were noted during this learning phase apart from a significant decrease in swim speed with age (F_{1,21} = 36.2, p<0.0001). Following the last day of training, a probe test (platform removed) assessed 24 h retention of platform location. Aged animals showed significantly longer path length to platform (F_{1,21} = 5.06, p<0.05) and latency to platform (F_{1,21} = 13.9, p<0.005), likely explained by a decrease in swim speed (F_{1,21} = 5.73, p<0.05). A proximal analysis which is not dependent on animals’ speed or their original distance to the platform at the beginning of each trial [62] also showed that aged animals were swimming, on average, at a cumulative distance farther from the target than younger animals (F_{1,21} = 6.8, p<0.05). A proximity average scalar derived from the cumulative distance data divided by the latency to platform (Fig. 2C) also revealed significant age-dependent impairment on memory recall (F_{1,21} = 6.7, p<0.05). Thus, irrespective of the analysis used, no main effect of treatment was found, suggesting that at the doses tested, PIO could not reverse the age-dependent decrease in memory recall 24 h after the last training day.

LTP

As described previously [13,41,42,43,44,45,46], an age-dependent deficit in long-term potentiation (LTP) maintenance was seen 25–30 min following LTP induction (F_{1,26} = 6.82, p<0.02). Animals treated with PIO did not show signs of improvement on measures of LTP induction or maintenance (F_{1,26} = 0.21, p = 0.67). Similar results were seen on measures of post-tetanic potentiation (PTP) taken immediately following LTP induction, showing a significant main effect of age (F_{1,26} = 5.51, p<0.05), but no main effect of treatment (F_{1,26} = 0.80, p = 0.38). Figure 3 shows group means of normalized EPSP slopes across both age and treatment during 20 min baseline and for 30 minutes

| Table 1. Blood chemistry panel. |
|--------------------------------|
|                        | YOUNG | AGED |
|------------------------|-------|------|
| CHOL (mg/dl)           | CTRL  | PIO  |
| YC                     | 119.4±19.7 | 124.4±10.1 |
| YP                     | 85.4±8.0 * | 108.1±5.9 * |
| AC                     | 141.1±21.4 ** | 88.4±8.7 ** |
| AP                     | 112.1±4.9 | 102.4±6.6 |
| TG (mg/dl)             | CTRL  | PIO  |
| YC                     | 291.2±45.5 | 184.2±44.1 † |
| YP                     | 141.1±21.4 ** | 88.4±8.7 ** |
| AC                     | 70.3±7.8 | 83.2±5.8 † |
| AP                     | 62.5±5.6 | 76.7±3.6 † |
| HDL (mg/dl)            | CTRL  | PIO  |
| YC                     | 7.2±1.8 | 6.1±1.6 |
| YP                     | 4.4±1.5 # | 4.2±1.3 # |
| AC                     | 112.1±4.9 | 102.4±6.6 |
| AP                     | 109.5±8.2 | 101.4±8.1 |
| INSULIN (ng/ml)        | CTRL  | PIO  |
| YC                     | 4.9±1.8 | 51.0±2.7 |
| YP                     | 37.5±3.2 | 47.7±4.6 |

Blood serum markers in control (CTRL) and PIO-treated (PIO) young and aged animals. Abbreviations: Cholesterol: CHOL; Triglycerides: TG; Alanine aminotransferase: ALT; High Density Lipoprotein: HDL. *; **, and † indicate significant PIO effects (two-way ANOVA, p<0.001, and p<0.0005, respectively).

# indicates significant aging effect (two-way ANOVA, p<0.05). Data represent mean ± SEM in 7–8 animals per group.

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following LTP induction. Absolute measures of EPSP amplitudes prior to LTP induction are shown in Figure 3E and reveal a significant age-dependent decrease (\(F(1,26) = 10.97, p < 0.003\)) but no treatment effect (\(F(1,26) = 0.01, p = 0.92\)). This result is consistent with prior evidence that at the CA1 synapse of aged animals, a decrease in the field EPSP is seen, likely mediated by a decrease in the number of functional synaptic contacts [12].

AHP

Decreased excitability mediated by an increase in the sAHP is a reliable Ca\(^{2+}\)-related biomarker of aging in hippocampal CA1 pyramidal neurons [63,64,65,66,67]. Two-way ANOVA on the AHP (Fig. 4) revealed a significant main effect of aging on measures of sAHP (Fig. 4) amplitude (32.5% increase over young, \(F(1,26) = 4.07, p < 0.05\); two-way ANOVA), and duration (26% increase over young, \(F(1,26) = 5.12, p < 0.05\); two-way ANOVA).

Importantly, a main effect of PIO on these AHP measures also was noted with PIO causing a significant reduction in the AHP amplitude (\(F(1,26) = 4.07, p < 0.05\)) and duration (\(F(1,26) = 5.12, p < 0.05\)). PIO had a greater effect on the AHP recorded from the older animals, essentially reducing the AHP to levels seen in young animals (Bonferroni \(p < 0.05\)). No significant difference was found in the mAHP across the different age and treatment groups. No effects of PIO or aging on measures of neuronal health were noted (input resistances in \(\Omega\), were 57.5 \(\pm\) 0.9 for young controls, 58.3 \(\pm\) 0.5 for aged controls, 52.1 \(\pm\) 0.4 for young PIO, and 57.3 \(\pm\) 0.9 for aged PIO), and all cells displayed overshooting action potentials.

IR Signaling

Because PIO reduced insulin levels in the periphery (Table 1), we tested whether we could detect changes in insulin receptor (IR) signaling in the liver and cerebral cortex of each animal. We estimated the degree of IR signaling by normalizing phosphorylated to total IR (ratio) using two separate IR ELISA kits [68]. PIO significantly reduced phosphorylated IR levels in both age groups (\(p < 0.01\); Fig. 5C). Quantitatively similar decreases in total IR also were seen (\(p < 0.05\); Fig. 5A), suggesting no net effect of age or treatment on liver IR signaling (Fig. 5E). Data from animal cortices also showed significant PIO-mediated decrease in phosphorylated IR levels (\(p < 0.05\); Fig. 5D), with no change in total IR (Fig. 5B). As surrogate indication of activated insulin receptor signaling, this reduction in phosphorylated IR in the brain and the periphery likely reflects PIO-mediated reduction in insulin levels. It is not clear, however, that this reflects decreases in insulin receptor signaling as neither ratios of phosphorylated to total IR signals (Figs. 5E and F), nor glucose levels, were significantly altered by age or treatment. Interestingly, while total IR levels were comparable in the liver and brain cortical tissues (Figs. 5A and B), phosphorylated IR levels were approximately 5 fold lower in the brain (Figs. 5C and D), suggesting lesser IR signaling in this tissue. Given that the animals in this study were non-diabetic, were not challenged with high fat diets, and were not pathologically aged, these results indicate that reducing insulin levels in healthy animals seems to have had little impact on insulin signaling.

Gene Microarrays

Two-way ANOVA applied to the 7922 genes in the filtered list (see Methods) yielded three sets of p-values (main effects of aging and drug treatment, and interaction). Because each of these sets of p-values is vulnerable to the error of multiple testing, we plotted p-value frequency histograms (Fig. 6) for each set of p-values, and superimposed the number of findings expected by chance. Interestingly, both drug and interaction terms performed well below chance (at \(p < 0.05\); FDR=2.3), while the main effect of age showed a strong, reliable signature (at \(p < 0.05\); FDR = 0.21), similar to that seen in prior studies [48,49,51,70,71]. Thus, microarray-based transcriptional signatures of hippocampal tissue
Figure 3. Theta-burst induced synaptic potentiation. (A) and (B) Normalized EPSP slopes measured across both age and treatment groups during baseline and following theta burst stimulation (TBS). Representative averaged EPSP traces for each age and treatment group are shown in insets. Post-tetanic potentiation (PTP) of the EPSPS was measured immediately following LTP induction (C). LTP maintenance was measure 25–30 min latter in CTRL and PIO-treated animals (D). (E) EPSP amplitudes taken before TBS reveal a significant effect of aging. No PIO effects were noted, * indicates significant aging effect (two-way ANOVA, p<0.05). Data represent mean ± SEM in 6–9 hippocampal slices from 3–5 animals per group.

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Figure 4. Measures of the Ca\textsuperscript{2+}-dependent slow afterhyperpolarization (sAHP). (A) Representative examples of AHPs recorded in CA1 pyramidal cells from young (left) and aged (right) animals fed either control (CTRL) or PIO-enriched (PIO) diet. Both amplitude (B) and duration (C) of the sAHP were significantly enhanced in the aged group. In aged animals, however, long-termPIO treatment significantly reduced the sAHP (red traces in A). * indicates significant aging and PIO effects (two-way ANOVA, p<0.05). Data represent mean ± SEM in 12–21 recorded neurons from 6–7 animals per group.

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in this study reliably detected aging-related gene signatures comprised most notably of increased inflammatory pathways. Specifically, age-dependent increases in IL-18, IL-33, and IL-10 (receptor binding unit), and an age-dependent decrease in IL-16 were seen (for a complete list of genes significantly different with age, see Table S1). However, no significant ‘PIO’ or ‘age×PIO’ interaction effects were noted, possibly because the drug has no effect on hippocampal transcription, did not penetrate the CNS, or, alternatively, because the study was underpowered for the discovery of those changes.

To test whether the effects of PIO may have been limited to targeted pathways or processes that may have been lost in the overall non-significant PIO or age×PIO interaction effects reported in our gene microarray analysis (two-way ANOVA, Fig. 6), we interrogated the Gene Ontology (GO; www.geneontology.org) database of biological processes. Two highly relevant GO terms related to PIO’s mechanism of action, namely, insulin receptor signaling pathway (GO:0008286) and inflammatory response (GO:0006954) were isolated. Within these pathways, gene symbols associated with GO, annotated to rat, and present on our microarray platform were identified (total genes: insulin receptor signaling = 37; inflammatory signaling = 141). We next determined if any of the genes in these pathways were significantly changed, based on either the main effect of drug, or on the interaction term (p<0.05) from the two-way ANOVA. Insulin signaling genes were significant by neither main effect of drug nor by interaction. Among inflammatory response genes, 3 were significant by the main effect of drug (Ednra- decreased; Serpina3n- decreased; Zfp36- upregulated), and 3 more were significant by interaction (F3 and Mug1- PIO suppresses an age-related increase in expression, Tlr4- PIO suppresses expression in young subjects only). We then evaluated the likelihood that, by

Figure 5. Insulin Receptor signaling. (A) and (B) Total insulin receptor levels (IR) were measured in liver (left) and cortex (right) as a function of age and diet (CTRL vs. PIO). In liver, the significant age-dependent increase in IR levels was significantly reduced by PIO treatment. (C) and (D) Phosphorylated IR levels (pIR) were also measured in liver (left) and cortex (right). While PIO treatment significant reduced pIR levels in both tissues and both age groups, only liver levels appeared to be sensitive to age. (E) and (F) Measures of signaling through IR, based on the ratio of pIR (C and D) to total IR (A and B) revealed no effects of aging or PIO treatment. * indicates significant aging effects (two-way ANOVA, p<0.05). # indicates significant PIO effects (two-way ANOVA, p<0.05). Data represent mean ± SEM in 5–8 samples per group.

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downregulated genes. Although downregulated genes were significantly overrepresented, the magnitude of the effect was relatively small (61 vs. 40 expected by chance). DAVID analysis of aging changes (Table 2) showed downregulated pathways that are often associated with reduced neuronal activity (e.g., Ca ++ binding, receptor signaling, vesicle, and secretion), and reduced glucose-driven energy production (e.g., phosphorylation, ATP synthesis). Upregulated pathways were strongly associated with increased inflammatory signaling (e.g., antigen processing, response to wounding, adaptive immune response, and leukocyte cytotoxicity) and lipid/membrane processes possibly reflecting a metabolic shift towards fatty acid β oxidation and/or compensatory re-myelination (e.g., lysosome, myelination, membrane lipid catabolism, and lipid synthesis). As with the genes themselves, these age-related up- and downregulated processes, particularly increased inflammatory signaling and decreased neuronal activity, are highly similar to findings from previous microarray studies of rodent hippocampal aging [47,48,50,69,70].

Pro-inflammatory Markers

Peripheral inflammation was monitored using a serum-based multiplex bead assay targeted for key cytokines, namely Interleukin (IL)-1β, IL-6 and tumor necrosis alpha (TNF-α). Of the twenty-nine serum samples analyzed, two outliers (one YC and one AP) were removed from the analysis for having signals twice the standard deviation associated with the mean of that group. ANOVA analysis of the peripheral inflammatory markers on the remaining 27 animals (n = 6 young control-YC; n = 8 young PIO-YP; n = 7 aged control-AC; n = 6 aged PIO-AP) showed an age-dependent trend toward larger IL-6 signals, with a near doubling in signal intensity (F(1,24) = 2.7, p = 0.1), however, no PIO effects were detected (p = 0.70). IL-6 signals for each group were (in pg/mL): 122.8 ± 16.7 for the YC, 142.8 ± 26.9 for the YP, 204.9 ± 36.9 for the AC, and 165.5 ± 40.7 for the AP. Serum signals from TNF-α and IL-1β were below detectable levels (<24.4 pg/mL). Compared to the microarray signatures, the results in the periphery showed only modest increases in inflammatory cytokines with age but also reveal that at the concentration tested, PIO did not significantly reduce inflammation in either tissue.

Discussion

We investigated pioglitazone’s potential role in reducing peripheral and central markers in a well-characterized animal model of cognitive decline with aging. We chose the antidiabetic drug because of its favorable permeability into the CNS (~10%, [71]), positive effects in the treatment of type 2 diabetes mellitus (T2DM), and its reported success in the treatment of neurodegenerative disease in several animal models. Choice of animal age, dose, and delivery method were selected to mimic clinical conditions associated with the treatment of late-onset diabetes. In prior studies on the impact of the TZD in the brain, higher doses have been used, and often in more pathological models, including Alzheimer’s, Parkinson’s disease, and spinal cord or brain injury. These models are characterized by pathological processes that are likely present in normal aging, but to a lesser extent. As such, and because of the reported beneficial use of TZDs in alleviating cognitive decline in elderly patients with or without diabetes, we felt it important to test PIO’s role in an animal model of normal aging. We focused on several well-
established brain biomarkers of aging including learning and memory and inflammatory processes.

Several lines of evidence show that specific TZDs can have direct effects on Ca^{2+} homeostasis, particularly L-type voltage-gated Ca^{2+} channels in cardiovascular tissues [72,73,74,75,76,77], as well as in neurons [31,32]. Given that Ca^{2+} dysregulation is considered a hallmark of brain aging [63,78,79,80], is present in animal models of diabetes [81,82,83,84], and appears to contribute to insulin resistance in the periphery [85], we tested animal models of diabetes [81,82,83,84], and appears to be considered a hallmark of brain aging [65,78,79,80], is present in cognitive performance (reviewed in [12,67,78]). In prior studies focusing on whether long-term in vivo PIO can have inhibitory effects on NMDA currents in hippocampal slices will help to address this issue. With respect to the effect of PIO on the AHP, one possibility is that it is mediated by an indirect action on glucocorticoid regulation. In fact, in AD models, TZDs have been shown to improve cognition via alterations in glucocorticoid signaling [26,86], and there is good evidence that the AHP is sensitive to glucocorticoids [87].

We measured insulin levels as well as insulin signaling in the brain and the periphery of young and aged animals. Our results show that insulin and glucose levels were not elevated with age, but insulin levels were significantly reduced by PIO in both age groups (Table 1). Given that our animals were not diabetic, we show that insulin and glucose levels were not elevated with age, but insulin levels were significantly reduced by PIO in both age groups (Table 1). Given that our animals were not diabetic, we predicted that a decrease in insulin levels would result in a concomitant reduction in insulin receptor signaling, at least in the periphery and perhaps also in the brain. ELISA assays on liver (peripheral) and cortical (central) tissues showed that while the insulin receptor levels (Fig. 5). These results suggest that insulin receptor levels exist in a dynamic insulin-sensing equilibrium both in the brain and the periphery. It is not clear, however, whether the few total receptors remaining might be more efficient at translating insulin’s action through differential regulation of downstream targets of the insulin receptor, including IRS-1, Pi3K, Akt, and SH2 [88]. Further, it is also likely that other insulin sensitive tissues (e.g., muscle or fat) could show enhanced sensitivity to PIO’s effects on insulin, resulting in more robust

### Table 2. Pathway analysis for aging-related genes.

| Downregulated | Upregulated |
|---------------|-------------|
| Ca^{2+} ion binding (# 17, p = 0.0035) - Actn1, Ap1gpb1, Calb1, Camk4, Clstn1, Dai1, Dmp1, Gpd2, LOC684520, Ncald, Nell2, Pcdha1, Rnf111, Scg2, Slc24a3, Syp, Tesc | Lysosome (# 24, p = 4.8^{-06}) - Abca2, A2a, Cd74, Ctdsd, Ctns, Dnase2a, Fhbp1, Fucal1, Gm2a, Gusb, Hexa, Hexb, If30, Lamp1, Lamp2, Laptm5, Lgmn, Neul, Nppa, Ppt1, Psen1, Scl1a3a, Tpp1, Trip10 |
| Receptor activity (# 27, p = 0.0008) - Acvr2a, Atm1, Chin1, Dnkl, Epha7, Gabra5, Gabrb3, Gpr176, Htr1b, I22ra2, Insr, LOC683548, Mmd, Nr3c2, Oprml1, Pcdha1, Pcsk5, Ptpro, Ptpru, Ring1, Sla7a1, Sra1, Sistr2, Strap, Tfr, Trpc5 | Tyrosine phosphorylation of Stat3 (# 04, p = 3.7^{-05}) - Celf1, I22ra2, Ppp2ca, Ppp2cb |
| Phosphorylation (# 27, p = 0.0039) - Acvr2a, Atps5a1, Atps5h, Atps6v0a2, Atps6v0b2, Atps6v1c1, Camk4, Cask, Celf1, Dclkl, Epha7, Ikbpkap, I22ra2, Insr, LOC687516, Map2k1, Map2k5, Map3k12, Mark3, Nme1, Pcsk5, Plk2, Ppp2ca, Ppp2cb, Prpfl4b, Uhrmk1 | Vesicle (# 17, p = 0.0145) - A2c3, Agrp1a, Agrp1t1, Ap1gpb1, Capsa2, Chgb, Copbl1, Klf3a, Rab12, Rab3d, Scamp1, Scg2, Syn1, Synp, Syst17, Tfrc, Trnm9 |
| Secretion (# 18, p = 0.0272) - Agrp1a, Agrp2, Copa, Cops1, LOC498335, Nr3c2, Osbpl5, Rab3d, Sar1a, Scamp1, Scg2, Scml1, Sec22a, Snc1, Syn1, Trim9, Yipf5 | ATP synthesis coupled H^{+} transport (# 06, p = 0.0056) - Atp5a1, Atps5h, Atps6v0a1, Atps6v0e2, Atps6v1b2, Atps6v1c1 |

List of numbers indicate that such a number could be found by chance (p = 0.05, two-way ANOVA main effect of age) down- and up-regulated genes were subjected to overrepresentation analysis using DAVID’s clustering function (Methods). Representative pathways from each cluster found to have significantly more genes than expected by chance (p < 0.05, modified Fisher’s exact test) are shown. The pathway (bold) is followed in parentheses by the number of genes found to be significant in that pathway (p) and the likelihood that such a number could be found by chance (p =). This is followed by a list of the gene symbols within that pathway.

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changes in insulin receptor signaling. Because insulin levels were decreased in both young and aged animals treated with PIO, and AHP reductions were only seen in the aged group, we do not believe insulin levels directly influenced the AHP under the condition tested. There is, however, evidence supporting a decrease in insulin sensitivity in the brain of AD patients and in AD models [89,90,91], and we are currently testing this hypothesis in our aging model. However, it is unclear whether the aging F344 rat is a good model of T2DM, and responds to high fat diets with signs of insulin resistance in both the periphery and the brain [92,93]. Here, therefore, we believe that PIO’s effects on insulin signaling were somewhat blunted because animals were healthy and non-diabetic. On the other hand, one would predict that under conditions more representative of human aging, where accumulated exposure to high fat diets and a sedentary life style contribute to T2DM, PIO might significantly reduce insulin levels [56,57,58,59], increase insulin sensitivity, and likely increase insulin signaling.

Prior studies in CNS and peripheral cell types modeling trauma and insult (e.g., LPS, PMA) demonstrate that PPAR-γ agonists play a critical role in reducing inflammatory cytokines (interleukin-1β and TNF-α), including activation of inducible nitric oxide synthase (iNOS) [94,95,96,97,98,99]. Similarly, in animal models of ischemia, stroke, hypertension, stress, and Parkinson’s disease, which are also characterized as pro-inflammatory conditions, PPAR-γ agonists provide significant neuroprotection [100,101,102,103,104,105,106,107,108,109,110,111,112]. In AD animal models also, PPAR-γ agonists appear to reduce baseline inflammatory levels [24,26,113,114]. Only one prior animal study examined the effects of TZDs under non-pathological aging conditions. The authors reported that the increase in pro-inflammatory cytokine levels (IL-1β) in the hippocampus of aged F344 rats was not sensitive to the actions of the TZD rosiglitazone (10 mg/Kg/day) yet the drug caused significant improvement in contextual fear conditioning [115]. Similarly, a prior publication using 20 mg/Kg/day PIO for four months in transgenic mouse models of AD (Tg2576) revealed very little anti-inflammatory effects of PIO (based on microglial activation, soluble Aβ levels, and plaque burden) when compared to ibuprofen treatment [114]. The same group, however, convincingly showed that a 40 mg/Kg/day PIO dose could significantly reduce brain inflammation in the APPV7171 transgenic mouse [24]. Here, therefore, we examined inflammatory cytokines in the serum, and inflammatory signaling in the brain using hippocampal microarray analyses in the context of normal aging, and at low doses of PIO. While a clear inflammatory signature was present in the brain as previously reported in microarray studies of aging [11,47,48,50,116,117,118,119,120], PIO did not significantly reduce inflammatory markers in the hippocampus. In the periphery, no robust age-dependent change in inflammatory cytokine levels was seen (although a trend in increased IL-6 levels was noted in aged animals), precluding an effect of PIO. Importantly, reductions in peripheral insulin and lipids indicate the target therapeutic window for PIO was reached. Under these conditions, thus, it is unclear that PIO was able to reduce central and peripheral inflammatory markers in an animal model of aging, and together, these results suggest that higher PIO doses might be necessary to reduce inflammatory pathways and exert beneficial cognitive effects.

Using an unbiased microarray analysis approach on hippocampal tissue, our study compared the effects of a brain permeant TZD treatment in younger and older animals, and showed that while age-dependent gene signatures were clearly present at both the gene [Fig. 6] and pathway (Table 2) levels, PIO effects on gene expression were virtually absent. Possible reasons for this include: low statistical detection power, insufficient drug exposure, or lack of influence of this treatment regimen on hippocampal transcription. Low statistical power is a possibility; however, the treatment main-effect histogram dips well below chance at small p values (Fig. 6), and it seems unlikely that increasing the number of subjects would allow us to detect significant transcriptional effects of PIO. Regarding drug exposure, PIO reduced peripheral blood chemistry measures significantly and in the direction predicted by prior work (see discussion above and Table 1), suggesting that treatment levels were appropriate. Thus, it seems reasonable to conclude that PIO did not exert a detectable transcriptional effect on hippocampal gene transcription, and that this lack of central influence may be due to either reduced blood-brain barrier penetration or a frank lack of response from hippocampal tissue. Nevertheless, because PIO has established effects involving insulin and inflammatory processes, we also directly investigated genes associated with these processes and tested their sensitivity to PIO in the brain. Although it is not possible to evaluate the biological importance of the 6 genes that were identified without functional genetic manipulation (e.g., knock-in), our resampling analysis revealed that these targeted pathways were not statistically significant. It is interesting to note, however, that of the six genes identified, one of them (Tlr4) recently was found to be sensitive to PIO in monocytes and macrophages [121], reinforcing the role of PIO as an anti-inflammatory agent, and suggesting Tlr1 might be a common target of PIO in peripheral and central tissues.

Compared to prior aging studies, our overlap analysis (Fig. 6 inset) suggests that, irrespective of where along the dorsal-ventral axis it is sampled, the hippocampus shows increased inflammatory markers with age, and validates the use of hippocampal extremities in future microarray studies. Interestingly, upregulated inflammatory categories, historically the most consistent and largest magnitude of the aging brain transcriptional signatures, remain largely unperturbed by PIO administration.

To our knowledge, this is the first study testing long-term PIO treatment in the F344 rat with age, specifically investigating whether a commonly prescribed drug may have off-target cognition enhancing effects in a rat model of aging. The study was designed using a clinically-relevant dose and delivery method. As expected, several signatures of aging were present in older animals, characterized by weak peripheral and robust central inflammatory increases, reduced spatial memory and LTP maintenance, and increased Ca2+-dependent AHPs. Peripheral insulin levels, phosphorylated insulin receptors in the CNS and the periphery, and the AHP were significantly reduced by PIO. While the mechanism through which PIO may mediate its central effects (AHP reduction, reduced phosphorylated insulin receptor) is not clear, it does not appear to occur via a transcriptional process. Given the increased incidence in metabolic syndrome and T2DM seen in the aging population, together with the high numbers of prescriptions written for TZDs, our study has direct therapeutic relevance and suggests future experiments testing the use of these agents at clinically-relevant doses for the treatment of neurological or cognitive conditions are needed. Nevertheless, the results of our study do not preclude the beneficial effects of TZDs in the elderly where metabolic dysregulation and diabetes are often reported.

**Supporting Information**

**Table S1**

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Author Contributions

Conceived and designed the experiments: EMB NMP OT. Performed the experiments: JTP TP JLS MGA KCC OT. Analyzed the data: EMB JTP TP JLS KLA JP KCC OT. Contributed reagents/materials/analysis tools: JCG DAC NMP OT. Wrote the paper: EMB TP NMP OT.

References

1. Gradman TJ, Laws A, Thompson LW, Reaven GM (1993) Verbal learning and/or memory improves with glycemic control in older subjects with non-insulin-dependent diabetes mellitus. J Am Geriatr Soc 41: 1305–1312.
2. Aswa N, Gagnon M, Mesnier C (2004) The relationship between impaired glucose tolerance, type 2 diabetes, and cognitive function. J Clin Exp Neuropsychol 26: 1094–1100.
3. Gregg EW, Yaffe K, Casley JA, Rolka DB, Blackwell TL, et al. (2000) Is diabetes associated with cognitive impairment and cognitive decline among older women? Study of Osteoporotic Fractures Research Group. Arch Intern Med 160: 174–180.
4. Mesnier C (2005) Impact of impaired glucose tolerance and type 2 diabetes on cognitive aging. Neurobiol Aging 26 Suppl 1: 26–30.
5. Reagan LP (2002) Glucose, stress, and hippocampal neuronal vulnerability. Int Rev Neurobiol 51: 289–324.
6. Ryan CM, Geckle M (2000) Why is learning and memory dysfunction in Type 2 diabetes limited to older adults? Diabetes Metab Res Rev 16: 308–315.
7. Zhan WQ, Al-Awadi M (2001) Role of insulin and insulin receptor in learning and memory. Mol Cell Endocrinol 172: 125–134.
8. Barrientos RM, Frank MG, Heim AM, Higgins EA, Watkins LR, et al. (2009) Time course of hippocampal IL-1 beta and memory consolidation impairments in aging rats following peripheral infection. Brain Behav Immun 23: 46–54.
9. Fuhrer MA, Watson GS, Montine TJ, Wang Q, Green PS, et al. (2005) Hyperinsulinemia provokes synchronous increases in central inflammation and beta-amyloid in normal adults. Arch Neurol 62: 1359–1364.
10. Goodpasture JP, Chen J, Abraham J, Richmond AF, Berg BM, et al. (2005) Exaggerated neuroinflammation and sickness behavior in aged mice following activation of the peripheral innate immune system. FASEB J 19: 1239–1331.
11. Prolla TA (2002) DNA microarray analysis of the aging brain. Chem Senses 27: 299–306.
12. Rosenzweig ES, Barnes CA (2003) Impact of aging on hippocampal function: plasticity, network dynamics, and cognition. Prog Neurobiol 69: 143–179.
13. Barnes CA, McNaughton BL (1985) An age comparison of the rates of acquisition and forgetting of spatial information in relation to long-term enhancement of hippocampal synapses. Behav Neurosci 99: 1040–1048.
14. Berger J, Moller DE (2002) The mechanisms of action of PPARs. Annu Rev Med 53: 499–415.
15. Hauner H (2002) The mode of action of thiazolidinediones. Diabetes Metab Res Rev 18 Suppl 2: S10–15.
16. Martin GI, Schoumans K, Lefebvre AM, Stach B, Auswir J (1997) Coordinate regulation of the expression of the fatty acid transport protein and acyl-CoA synthetase genes by PPARalpha and PPARgamma activators. J Biol Chem 272: 20210–20217.
17. Schoumans K, Pieniadz-Ousahre J, Lefebvre AM, Heyman RA, Briggs M, et al. (1996) PPARalpha and PPARgamma activators direct a distinct tissue-specific transcriptional response via a PPRE in the lipoprotein lipase gene. Embio J 15: 5336–5348.
18. Landreth GE, Heneka MT (2001) Anti-inflammatory actions of peroxisome proliferator-activated receptor gamma agonists. J Neuroimmunol 116: 177–179.
19. Hanyu H, Sato T, Kiuchi A, Sakurai H, Iwamoto T (2009) Pioglitazone activated receptor-gamma (PPARgamma) in Alzheimer’s disease: therapeutic implications. CNS Drugs 22: 1–14.
20. Hanyu H, Sato T, Kiuchi A, Sakurai H, Iwamoto T, et al. (2009) Rosiglitazone attenuates learning and memory deficits in Tg2576 Alzheimer mice. Exp Neurol 199: 286–303.
21. Lacombe P, Mathews PM, Schmidt SD, Breidt T, Heneka MT, et al. (2004) Effect of anti-inflammatory agents on transforming growth factor beta over-expressing mouse brains: a model revised. J Neuroinflammation 1: 11.
22. Landreth GE, Heneka MT (2005) Anti-inflammatory actions of peroxisome proliferator-activated receptor gamma agonists in Alzheimer’s disease. Neurobiol Aging 26: 937–944.
23. Feinstein DL (2003) Therapeutic potential of peroxisome proliferator-activated receptor agonists for neurological disease. Diabetes Technol Ther 5: 67–73.
24. Fishel MA, Watson GS, Montine TJ, Wang Q, Green PS, et al. (2005) Peroxisome proliferator-activated receptor-gamma (PPARgamma) is expressed in hippocampal neurons and its activation prevents beta-amyloid neurodegeneration: role of Wnt signaling. Exp Cell Res 304: 91–104.
25. Fishel MA, Watson GS, Montine TJ, Wang Q, Green PS, et al. (2005) Peroxisome proliferator-activated receptor-gamma (PPARgamma) activation protects neurons from NMDA excitotoxicity. Brain Res 1075: 460–469.
26. Godbout JP, Chen J, Abraham J, Richwine AF, Berg BM, et al. (2005) Distant modulation of voltage-gated and ligand-gated Ca2+ currents by PPAR-gamma agonists in cultured hippocampal neurons. J Neurochem 96: 130–141.
27. Godbout JP, Chen J, Abraham J, Richwine AF, Berg BM, et al. (2005) Distant modulation of voltage-gated and ligand-gated Ca2+ currents by PPAR-gamma agonists in cultured hippocampal neurons. J Neurochem 96: 130–141.
73. Asano M, Nakajima T, Iwasawa K, Morita T, Nakamura F, et al. (1999) Trogilitazone inhibits voltage-dependent calcium currents in guinea pig cardiac myocytes. Circulation 99: 2942–2950.

71. Maeshiba Y, Kiyota Y, Yamashita K, Yoshimura Y, Motohashi M, et al. (1997) Effect of rosiglitazone on the uptake of sodium in rat mesenteric arteries. Pharmacology 73: 15–22.

67. Foster TC (2007) Calcium homeostasis and modulation of synaptic plasticity in the aged brain. Aging Cell 6: 319–325.

66. Gant JC, Thibault O (2008) Action potential throughput in aged rat cardiac ventricular myocytes. J Cardiovasc Pharmacol 44: 109–116.

65. Disterhoft JF, Thompson LT, Moyer JR, Jr., Mogul DJ (1996) Calcium-pertussis toxin-sensitive inward rectifier current in brain inhibits inflammatory pain, dorsal horn expression of Fos, and local edema. Neuropharmacology.

63. Moyer JR, Jr., Thompson LT, Black JP, Disterhoft JF (1992) Nimodipine decreases calcium channel function and myogenic tone in rat cardiac myocytes. Pflugers Arch 418: 395–401.

62. Gallagher M, Burwell R, Burchinal M (1993) Severity of spatial learning impairments in patients with type 2 diabetes mellitus: a retrospective review of randomly selected medical records. Clin Ther 24: 374–378.

61. Lipscombe LL, Gomes T, Levesque LE, Hux JE, Juurlink DN, et al. (2007) Rosiglitazone reverses memory decline and hippocampal atrophy in Alzheimer’s disease. Jama 298: 2634–2643.

60. Spanheimer R, Betteridge DJ, Tan MH, Ferrannini E, Charbonnel B (2009) Pioglitazone and cell proliferation in vascular smooth muscle cells. Br J Pharmacol 128: 101–107.

59. Szapary PO, Bloedon LT, Samaha FF, Duffy D, Wolfe ML, et al. (2006) Effects of pioglitazone on inflammatory cytokines in both early and late diabetic patients with metabolic syndrome. Aging Thromb Vasc Biol 26: 182–188.

58. Spanheimer R, Betteridge DJ, Tan MH, Ferrannini E, Charbonnel B (2009) Pioglitazone and cell proliferation in vascular smooth muscle cells. Br J Pharmacol 128: 101–107.

57. Deeg MA, Bueh JB, Goldberg RB, Kendall DM, Zagar AJ, et al. (2007) Insulin enhances mitochondrial Ca++-dependent afterhyperpolarization in CA1 pyramidal cells in an age- and concentration-dependent manner. J Neurophysiol 68: 2100–2109.

56. Gallagher M, Burwell R, Burchinal M (1993) Severity of spatial learning impairment in aging: development of a learning index for performance in the Morris water maze. Behav Neurosci 107: 618–626.

55. Benjamini Y, Drai D, Elmer G, Kafkafi N, Golani I (2001) Controlling the false discovery rate: a practical and powerful approach to multiple testing. J R Stat Soc B 65: 289–300.

54. Bolstad BM, Collin F, Brettschneider J, Simpson K, Cope L, et al. (2005) Bioconductor: open source system for microarray data management and analysis. Biotechniques 39: 374–378.

53. Deew MA, Bueh JB, Goldberg RB, Kendall DM, Zagar AJ, et al. (2007) Insulin enhances mitochondrial Ca++-dependent afterhyperpolarization in CA1 pyramidal cells in an age- and concentration-dependent manner. J Neurophysiol 68: 2100–2109.

52. Kruglikov I, Gryshchenko O, Shutov L, Kostyuk E, Kostyuk P, et al. (2004) Diabetes-induced alterations in ER calcium mobilization in primary and secondary nociceptive neurons. Pfiegers Arch 448: 395–401.

50. Blalock EM, Chen KC, Sharrow K, Herman JP, Porter NM, et al. (2003) Gene expression profiling of Alzheimer’s disease: implications for treatment. CNS Drugs 17: 27–45.

51. Bolstad BM, Collin F, Brettschneider J, Simpson K, Cope L, et al. (2005) Bioconductor: open source system for microarray data management and analysis. Biotechniques 39: 374–378.

49. Ikeda S, Watanabe T (1998) Effects of troglitazone and pioglitazone on the contractility of coronary vascular smooth muscle cells. Eur J Pharmacol 357: 243–250.

48. Miettinen OS, Hyytinen E, Umbach J, Kuusela L, Jaakola M, et al. (2001) Peroxisome proliferator-activated receptor-gamma agonists induce neuroprotection following transient focal ischemia in normotensive, normoglycemic as well as hypertensive and type-2 diabetic rodents. J Neurochem 77: 15–22.

47. Burger C, Lopez MC, Feller JA, Baker HV, Muzyczka N, et al. (2007) Changes in transcription within the CA1 field of the hippocampus are associated with age-related spatial learning impairments. Neurobiol Learn Mem 87: 21–41.

46. Moyer JR, Jr., Thompson LT, Black JP, Disterhoft JF (1992) Nimodipine increases excitability of rabbit CA1 pyramidal neurons in an age- and concentration-dependent manner. J Neurophysiol 59: 413–420.

45. Gant JC, Thibault O (2008) Pioglitazone in Normal Aging.
103. Aoun P, Watson DG, Simpkins JW (2003) Neuroprotective effects of PPARgamma agonists against oxidative insults in HT-22 cells. Eur J Pharmacol 472: 65–71.

104. Garcia-Bueno R, Madrigal JL, Lizasoain I, Moro MA, Lorenzo P, et al. (2005) Peroxisome proliferator-activated receptor gamma activation decreases neuronal inflammation in brain after stress in rats. Biol Psychiatry 57: 885–894.

105. Heneka MT, Klockgether T, Feinstein DL (2000) Peroxisome proliferator-activated receptor-gamma ligands reduce neuronal inducible nitric oxide synthase expression and cell death in vivo. J Neurosci 20: 6862–6867.

106. Hunter RL, Choi DY, Ross SA, Bing G (2008) Protective properties afforded by pioglitazone against intrastriatal LPS in Sprague-Dawley rats. Neurosci Lett 432: 198–201.

107. Romero C, Hurtado O, Mallolas J, Pereira MP, Morales JR, et al. (2007) Ischemic preconditioning reveals that GLT1/EAAT2 glutamate transporter is a novel PPARgamma target gene involved in neuroprotection. J Cereb Blood Flow Metab 27: 1327–1338.

108. Breidert T, Callebert J, Heneka MT, Landreth G, Launay JM, et al. (2002) Protective action of the peroxisome proliferator-activated receptor-gamma agonist pioglitazone in a mouse model of Parkinson’s disease. J Neurochem 82: 615–624.

109. Victor NA, Wanderi EW, Gamboa J, Zhao X, Aronowski J, et al. (2006) Altered PPARgamma expression and activation after transient focal ischemia in rats. Eur J Neurosci 24: 1653–1663.

110. Dehmer T, Heneka MT, Sastre M, Dichgans J, Schulz JB (2004) Protection by pioglitazone in the MPTP model of Parkinson’s disease correlates with I kappa B alpha induction and block of NF kappa B and iNOS activation. J Neurochem 88: 494–503.

111. Kapadia R, Yi JH, Vemuganti R (2002) Mechanisms of anti-inflammatory and neuroprotective actions of PPAR-gamma agonists. Front Biosci 13: 1013–1026.

112. Sundararajan S, Gamboa J, Victor NA, Wanderi EW, Lust WD, et al. (2005) Peroxisome proliferator-activated receptor-gamma ligands reduce inflammation and infarction size in transient focal ischemia. Neuroscience 130: 685–696.

113. Pedersen WA, Flynn ER (2004) Insulin resistance contributes to aberrant stress responses in the Tg2576 mouse model of Alzheimer’s disease. Neurobiol Dis 17: 500–506.

114. Yan Q, Zhang J, Liu H, Babu-Khan S, Vassar R, et al. (2003) Anti-inflammatory drug therapy alters beta-amyloid processing and deposition in an animal model of Alzheimer’s disease. J Neurosci 23: 7504–7509.

115. Gemma C, Stellwagen H, Fister M, Coultrap SJ, Mesches MH, et al. (2004) Rosiglitazone improves contextual fear conditioning in aged rats. Neuroreport 15: 2253–2259.

116. Ayeni KK, Kumar A, Cui L, Jackson TC, Foster TC (2009) Estrogen effects on cognition and hippocampal transcription in middle-aged mice. Neurobiol Aging 30: 932–945.

117. Burger C, Lopez MC, Baker HV, Mandel RJ, Muzyczka N (2000) Genome-wide analysis of aging and learning-related genes in the hippocampal dentate gyrus. Neurobiol Learn Mem 89: 379–396.

118. Parachikova A, Agadjanyan MG, Cribbs DH, Blurot-Jones M, Perreau V, et al. (2007) Inflammatory changes parallel the early stages of Alzheimer disease. Neurobiol Aging 28: 1821–1833.

119. Stranahan AM, Lee K, Becker KG, Zhang Y, Maudley S, et al. (2008) Hippocampal gene expression patterns underlying the enhancement of memory by running in aged mice. Neurobiol Aging 10: 10–12.

120. Wang X, Zaidi A, Pal R, Garrett AS, Braceras R, et al. (2009) Genomic and biochemical approaches in the discovery of mechanisms for selective neuronal vulnerability to oxidative stress. BMC Neurosci 10: 12.

121. Dauz MR, Park S, Devaraj S, Jialal I (2009) Pioglitazone inhibits Toll-like receptor expression and activity in human monocytes and db/db mice. Endocrinology 150: 3457–3464.