

The Thiazolidinedione Pioglitazone Alters Mitochondrial Function in Human Neuronal-Like Cells

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Abbreviations: CO, cytochrome oxidase; DCF, dichlorodihydrofluorescein; DMSO, dimethyl sulfoxide; ETC, electron transport chain; GPx, glutathione peroxidase; HBSS, Hanks buffered salt solution; HRP, horseradish peroxidase; mtDNA, mitochondrial DNA; NRF, normalized relative fluorescence; NRF-1, nuclear regulatory factor 1; NT2, Ntera/D1; PBS, phosphate buffered saline; PPARG, peroxisome proliferator activator receptor γ ; PGC-1 α , peroxisome proliferator-activated receptor γ coactivator 1 α ; PMSF, phenylmethanesulfonyl fluoride; TBS, Tris buffered saline; TTBS, Tween-Tris buffered saline

Abstract

Thiazolidinediones alter cell energy metabolism. They are used to treat or are being considered for the treatment of disorders that feature mitochondrial impairment. Their mitochondrial effects, though, have not been comprehensively studied under chronic exposure conditions. We used the human neuronal-like NT2 cell line to directly assess the chronic effects of a thiazolidinedione drug, pioglitazone, on mitochondria. At micromolar concentrations, pioglitazone increased mitochondrial DNA (mtDNA) content, levels of mtDNA and nuclear-encoded electron transport chain subunit proteins, increased oxygen consumption, and elevated complex I and complex IV Vmax activities. Pioglitazone treatment was also associated with increased cytoplasmic but reduced mitochondrial peroxide levels. Our data suggest pioglitazone induces mitochondrial biogenesis and show pioglitazone reduces mitochondrial oxidative stress in a neuronal-like cell line. For these reasons pioglitazone may prove useful in the treatment of mitochondriopathies.

The thiazolidinedione compound pioglitazone reduces insulin resistance in type II diabetics (Yki-Jarvinen, 2004). It agonizes a transcription activating protein, peroxisome proliferator activator receptor γ (PPARG). Pioglitazone thus mimics the affects of PPARG's physiologic co-activator, peroxisome proliferator-activated receptor γ coactivator 1 α (PGC-1 α). Recent data indicate PGC-1 α also co-activates another transcription activating protein, nuclear respiratory factor 1 (NRF-1) (Wu et al., 1999; Scarpulla, 2002). NRF-1 initiates expression of nuclear-encoded components of the mitochondrial electron transport chain (ETC) and facilitates mitochondrial proliferation. For this reason it seems reasonable to consider the effects of pioglitazone on mitochondria.

Data suggest thiazolidinediones directly affect mitochondrial function (Dello Russo et al., 2003). For example, previous work suggests thiazolidinediones may affect coupling-uncoupling dynamics (Shimokawa et al., 1998; Kelly et al., 1998; Digby et al., 1998). These drugs appear to decrease oxidative phosphorylation coupling, which could in turn increase glucose utilization and influence free radical production/oxidative stress. The effects of thiazolidinediones on oxidative stress are difficult to predict. Uncoupling can be associated with reduced free radical production (Fiskum et al., 2004). On the other hand, the ability of acute thiazolidinedione exposure to increase oxidative stress has been reported (Shishido et al., 2003). To complicate matters, thiazolidinediones may affect mitochondria through more than one mechanism (Feinstein et al., 2005).

Thiazolidinediones are currently under consideration for the treatment of Alzheimer's disease (AD) (Landreth, 2006). The most proposed rationale is that thiazolidinedione anti-inflammatory properties may prove useful for mitigating beta amyloid (A β) induced microglial activation, a process potentially contributing to neurodegeneration. Mitochondrial dysfunction is

also a well-documented feature of AD (Parker et al., 1990; Swerdlow and Kish, 2002, Roses et al, 2007). Whether a primary or secondary phenomenon in AD, mitochondrial dysfunction in AD permits its designation as a mitochondriopathy (Swerdlow, 2007). Because of this, we chose the NT2 neuronal-like cell line for studying how chronic pioglitazone exposure affects mitochondrial function.

Materials and Methods

Cell culture. Human teratocarcinoma Ntera/D1 (NT2) neuronal precursor cells were maintained and expanded in 75 cm² flasks at 37°C and 5% CO₂. Our NT2 cells were originally obtained from Stratagene (La Jolla, CA). The maintenance medium consisted of Optimem (Gibco BRL, Gaithersburg, MD) supplemented with 10% heat inactivated, non-dialyzed fetal bovine serum and 1% penicillin-streptomycin.

Pioglitazone-containing media were prepared through addition of a sterile pioglitazone stock solution. The stock solution consisted of pioglitazone powder (Takeda Pharmaceuticals, Osaka, Japan) dissolved in dimethyl sulfoxide (DMSO). This stock was prepared as a 20 mM solution (15.7 mg in 2 ml of DMSO). Pioglitazone-containing media contained final concentrations of 10 µM, 20 µM, or 40 µM pioglitazone; for some parameters 5 µM pioglitazone was also evaluated. To control for the DMSO vehicle, 1 ml of DMSO was added to 500 ml of the 0 µM medium.

Cells were maintained in their designated medium for at least two weeks prior to any assays. Cells were harvested when flasks reached 90% confluency. We also routinely changed the culture medium one day prior to harvesting. Adherent cells were detached by washing the flask surface with phosphate-buffered saline (PBS) followed by trypsinization. Detached cells were placed in 15 ml (for whole-cell applications) or 50 ml (for applications requiring isolated mitochondria) conical tubes. The larger volume tubes were used for mitochondrial isolation because for this we routinely combined the contents of at least two flasks. After trypsin was removed by a five minute centrifugation at 200g (4°C), the cells were washed twice in PBS and pelleted via centrifugation (5 minutes, 200g, 4°C).

Mitochondrial enrichment. Mitochondrial isolation was performed at 4°C. Washed cell pellets were prepared as described above. These pellets were suspended in isolation buffer (10 mM HEPES, pH 7.3; 1 mM MgCl₂; 0.25 M sucrose) containing fresh protease inhibitors (Protease Inhibitor Cocktail Set I, Calbiochem, San Diego, CA) and 200 μM phenylmethanesulfonyl fluoride (PMSF). The contents of at least two confluent T75 flasks were combined prior to cell rupture, and we used 2 ml of isolation buffer for each harvested flask. Suspended cells were disrupted in a prechilled, 45 ml nitrogen cavitation chamber (Parr Instrument Company, Moline, Ill) at 600 psi for 10 min. Cells were constantly stirred during the pressurization period. The cavitation chamber contained a discharge valve. At the end of the pressurization period the contents were eluted through the discharge valve and collected in a 15 ml conical polypropylene tube. An additional 2 ml of isolation buffer (not supplemented with protease inhibitors or PMSF) was then used to rinse the inside of the cavitation chamber and poured out the open discharge valve into the tube containing the previously eluted pressurized cells. Un-disrupted cells and nuclei were pelleted for removal by centrifugation at 1000g for 10 min. An aliquot of the post-nuclear supernatant, which contained cytoplasmic contents and ruptured membranes, was set aside for glutathione peroxidase (GPx) assays. The rest of the supernatant was centrifuged at 9,000g for 10 min. The resulting crude mitochondrial pellet was suspended in 400 μl of isolation buffer. Protein content of either the 1000g supernatant or mitochondrial fraction was determined using the DC protein assay (BioRad, Hercules, CA).

Cytochrome oxidase, citrate synthase, and complex I V_{max} assays. Cytochrome oxidase and citrate synthase activities were determined on the whole cell pellets described above. These pellets were suspended in Ca⁺²-free, Mg⁺²-free Hank's buffered salt solution (HBSS) (Gibco

BRL) at a concentration of 30×10^6 cells/ml. Protein content of the whole cell suspensions was determined using the DC protein assay. For the cytochrome oxidase assay, a cuvet containing potassium phosphate buffer (20 mM, pH 7.0), dodecyl maltoside (20 μ l from a 10 mg/ml stock solution), and 25 μ g cell protein was warmed to 30°C for two minutes. The reaction was initiated by addition of 25 μ M of reduced cytochrome c, which brought the total cuvet volume to 1 ml. The oxidation of the reduced cytochrome c was followed for 2 minutes at 550 nm on a DU series spectrophotometer (Beckman Coulter, Fullerton, CA). To facilitate calculation of enzyme activity, the maximally oxidized cytochrome c absorbance was determined by adding several grains of potassium ferricyanide to the cuvet. The apparent first order rate constant was calculated and normalized to the amount of protein added to the cuvet ($\text{sec}^{-1}/\text{mg}$ protein). At least 12 independent assays were performed for each condition.

For citrate synthase activity determinations, a cuvet containing 100 mM Tris (pH 8.0), 100 μ M DTNB [5,5'-dithio-bis (2-nitrobenzoic acid)], 0.04% (v/v) Triton X-100 (added as 4 μ l of a 10% Triton solution), and 40 μ g of cell suspension was warmed over two minutes to 30°C. The reaction was initiated by the addition of first 100 μ M oxaloacetate and then 50 μ M acetyl CoA. The total cuvet volume was 1 ml. Citrate synthase catalyzes the release of a thiol from acetyl CoA, which reacts with DTNB to form 5-mercapto-2-nitrobenzoic acid. The extinction coefficient of DTNB reduction is $0.0136 \mu\text{M}^{-1} \text{cm}^{-1}$. The resulting absorbance change was followed in a spectrophotometer for two minutes at 412 nm. This absorbance change was divided by the extinction coefficient, and this value was further divided by the amount of protein in the cuvet. Citrate synthase activities were thus recorded as activity per mg of protein (nmol/min/mg protein). Ten independent assays were performed for each condition.

Complex I activities were determined from crude mitochondrial fractions (prepared as described above). Immediately following suspension of the mitochondrial pellet in 400 μ l of isolation buffer, 100 μ l was taken for protein determination using the DC protein assay. The remaining 300 μ l were transferred to 1 ml polystyrene centrifuge tubes (Fisherbrand, Fisher Scientific, Pittsburgh, PA), and these tubes were tightly capped. The mitochondria in the polystyrene tubes were then subjected to one freeze-thaw cycle, which was accomplished by cooling the suspension to -80°C for 1-2 hours. After thawing, the capped tubes containing the mitochondrial suspensions were sonicated in the cup horn of a Heat Systems-Ultrasonics W-225 sonicator (Farmingdale, NY). During the sonication procedure the cup horn was also filled with an ice water slurry, and the polystyrene tubes were maintained in a horizontal orientation 1 cm above the base of the cup horn. It is important to note when disrupting mitochondria through closed tubes, optimal sonication parameters depend on tube material and the type of sonicator used. For the sonicator and polystyrene tubes we used, we found a 6 minute sonication at 50% duty cycle and output control of 7 provided optimal mitochondrial disruption.

A cuvet containing complex I assay buffer (25 mM KPO_4 , pH 7.4; 0.25 mM potassium EDTA; 1.5 mM KCN) and 10 μ l of a 10 mM β -NADH stock solution was warmed over two minutes to 30°C . The NADH stock solution was prepared using complex I buffer. After this two minute warming period, 100 μ g of sonicated mitochondrial protein was added to the cuvet, which was then warmed for an additional minute at 30°C . The reaction was initiated by addition of coenzyme Q1 (Sigma-Aldrich, St Louis, MO) to a final cuvet concentration of 60 μ M. The coenzyme Q1 was added from a stock solution in which coenzyme Q1 was dissolved in ethanol at a concentration of about 25 mM. The final cuvet volume was 1 ml.

Our complex I assay utilized a spectrophotometer to follow the decrease in NADH absorbance that occurs at 340 nm as NADH is oxidized by complex I to NAD⁺. We followed this absorbance change over 2 minutes at 30⁰C. To account for NADH oxidation by other NADH oxidases, after 2 minutes we added 10 μ M rotenone (10 μ l from a 2 mM stock solution in which the rotenone was dissolved in methanol) to the cuvet and tracked the change in 340 nm absorption for another 2 minutes. The post-rotenone rate was subtracted from the pre-rotenone rate to yield the rotenone-sensitive rate. Typically, the rotenone sensitive rate accounted for 80-90% of the total rate. The rotenone-sensitive rate was divided by 0.00681 μ M⁻¹ cm⁻¹, the combined NADH-Q1 extinction coefficient at 340 nm. This value was then divided by the amount of protein in the cuvet in order to provide the rotenone-sensitive rate as nmol/min/mg protein. At least 15 independent assays were performed for each condition.

Cell oxygen consumption. Whole cells were harvested and washed as described above. The whole cell pellets were then suspended in PBS at a concentration of 30 x 10⁶ cells/ml. 200 μ l of each cell suspension was placed in individual wells of a BD Oxygen Biosensor System plate (BD Biosciences, Franklin Lakes, NJ). Each suspension was run in triplicate. BD Oxygen Biosensor System plates contain a dye that fluoresces with exposure to dissolved oxygen. Plates were scanned using a Wallac Victor2 plate reader (PerkinElmer Life and Analytical Sciences Inc., Wellesly, MA) with an excitation wavelength of 485 nm and an emission wavelength of 630 nm. Readings were taken at 30 minute intervals for a total of two hours. The plate was kept at 37⁰C and 5% CO₂ between readings. Data were normalized according to the manufacturer's normalization protocol (technical bulletin no. 448; BD Biosciences), which converted the data to normalized relative fluorescence (NRF) units. We then divided the calculated NRF units by the

amount of protein added to each well. Our data are thus presented as NRF units per mg protein. The three replicates for a given whole cell suspension were averaged to provide a single data point. The total NRF unit means for the independently analyzed cell suspensions of a particular condition were determined at 1 hour. Over the two hour reading period there was a linear fluorescence increase, and the mean NRF unit slopes for the independently analyzed cell suspensions of a particular condition were calculated. At least 10 independent whole cell suspensions were assayed for each condition.

Oxidative stress determinations. Cytoplasmic peroxide levels were estimated using 2',7'-dichlorodihydrofluorescein diacetate (DCF) (Molecular Probes, Inc., Eugene, OR). For this assay, cells were harvested and washed as described above. The washed whole cell pellets were suspended in a freshly prepared dye loading buffer consisting of HBSS (containing calcium and magnesium but not phenol red), 2 μ M DCF, and 0.1 μ g/ml Hoechst 33342 (Molecular Probes, Inc., Eugene, OR). These suspensions were maintained in the dark for 30 minutes at 37 °C and 5% CO₂. At the end of this dye loading period the cells were pelleted by centrifugation. The cell pellet was suspended in HBSS (containing calcium and magnesium but not phenol red) containing no dyes and the cells were re-pelleted. The washed cells were then suspended in assay buffer (PBS supplemented with 20 mM glucose) at a concentration of 5 x 10⁶ cells/ml. 200 μ l from each cell suspension (1 x 10⁶ cells) were added to individual wells of a clear 96 well plate (Fisherbrand). Each suspension was assayed in triplicate. The plates were analyzed using a Wallac Victor2 fluorescence plate reader at two different settings. The first setting, with excitation at 485 nm and emission at 530 nm, measured DCF fluorescence. The second setting, with excitation at 355 nm and excitation at 460 nm, measured Hoechst fluorescence. Hoechst

stains cell nuclei and its fluorescence provides a linear correlate of the number of cells in a well. The Hoechst readings were done to ensure differences in the number of cells added to each well did not account for differences in DCF fluorescence. After dividing the DCF fluorescence of each well by its Hoechst fluorescence, we averaged the three wells for each sample to obtain a final value. Plates were read at 10 minute intervals over one hour. Between readings plates were kept in the dark at 37°C and 5% CO₂. Fluorescence increase slopes were calculated, which reflect increasing DCF fluorescence as Hoechst fluorescence did not change between readings. Results for each pioglitazone condition are expressed as its fluorescence increase slope relative to that of pioglitazone-untreated cells. Eleven independent measurements were made for each condition.

Mitochondrial peroxide production was assessed using the amplex red (10-acetyl-3,7-dihydroxyphenoxazine) reagent. For this assay we used isolated mitochondria, which were prepared as described above. After suspending crude mitochondria pellets in 400 µl of isolation buffer, we used the BioRad DC protein assay to determine the protein concentration of each suspension. Based on these protein determinations, we added specific amounts of additional isolation buffer (without protease inhibitors or PMSF) to the suspensions of each group so that the final protein concentrations for all the suspensions in a group were equivalent. Stock solutions for glutamate (500 mM in water), malate (500 mM in water), ADP (500 mM in water), amplex red (10 mM in DMSO), and horseradish peroxidase (HRP; 10 U/ml in kit reaction buffer) were made. All stock solutions except the HRP solution were 100x their working concentrations; the HRP stock solution was 50x its working concentration. To prepare one milliliter of working solution, 10 µl each of the glutamate, malate, ADP, and amplex red stocks were added to 20 µl of the HRP stock and 940 µl of isolation buffer (without protease inhibitors

or PMSF). The working solution therefore contained 5 mM glutamate, 5 mM malate, 5 mM ADP, 100 μ M amplex red, and 0.2 U/ml HRP. To initiate the assay, 50 μ l of working solution were added to 50 μ l of mitochondrial suspension in clear 96 well microtiter plates. The plates were analyzed with a Wallac Victor2 fluorescence plate reader using excitation 560 nm and emission 590 nm. Readings were taken at one minute intervals. Thirty readings were taken. Each suspension was measured in triplicate and mean values for each suspension were determined for each time point. Amplex red reacts with H_2O_2 to produce highly fluorescent resorufin. The slope of the resorufin fluorescence increase was calculated for each sample. The slopes for the suspensions containing mitochondria from pioglitazone-treated cells were compared to the slopes of suspensions containing mitochondria from pioglitazone-untreated cells. At least 10 independent assays were performed for each pioglitazone concentration.

The glutathione peroxidase (GPx) reaction follows the oxidation of NADPH to $NADP^+$, which manifests as an absorbance decrease at 340 nm. We performed our measurements with a DU series spectrophotometer (Beckman Coulter). First, NT2 cell post-nuclear fractions were generated as described above with minor modification. Instead of suspending whole cells in 2 ml per flask prior to nitrogen cavitation, whole cells were suspended in 1 ml of isolation buffer per flask, and following nitrogen cavitation the chamber was washed with only 1 ml of isolation buffer. This modification was done in order to concentrate the contents of the post-nuclear supernatant generated by the 1000g spin. We used post-nuclear fractions because GPx is found in both cytoplasm and mitochondria. The protein concentration for each sample was measured using the BioRad DC protein assay. Post-nuclear fractions were then subjected to one freeze-thaw cycle (frozen at $-80^{\circ}C$ for at least two hours). The actual GPx assay was performed using the Total Glutathione Peroxidase Assay Kit (ZeptoMetrix, Buffalo, NY). This kit provides

buffers, NADPH, glutathione, glutathione reductase, and cumene hydroperoxide. Assays were performed according to the manufacturer's instructions, with modifications as specified. Instead of adding 30 μ l of sample, we adjusted our sample volumes so that 100 μ g of post-nuclear, freeze-thawed protein was added to the assay cuvet. This tended to require a 50-60 μ l volume addition. To compensate for this minor increase in sample volume, we reduced the amount of Working Solution A so that the final assay volume in the cuvet was maintained at 960 μ l. Also, a control was run for each sample in which a corresponding volume of de-ionized water was substituted for the actual sample. This provided a background rate that was subtracted from the rate obtained with the sample. The final corrected rate for each sample was divided by the NADPH extinction coefficient ($0.00622 \mu\text{M}^{-1} \text{cm}^{-1}$), and this result was further divided by the amount of protein in the cuvet (0.1 mg). GPx activities from the post-nuclear fractions of pioglitazone-treated cells were compared to those of post-nuclear fractions from pioglitazone-untreated cells. Data are reported as relative activities. Twelve independent measurements were performed for each condition.

Protein electrophoresis and Western blotting. Immunochemical determinations of cytochrome oxidase subunit 1 (CO1) protein levels were performed using isolated mitochondrial protein. Immunochemical determinations of cytochrome oxidase subunit 4 (CO4) protein levels were performed using whole cell protein. Mitochondria and whole cell fractions were prepared as described above and suspended in 200 μ l of PBS and 200 μ l of a 2x Laemmli buffer. The 2x Laemmli buffer was prepared as follows: 10 ml of 1M Tris-HCl (pH 6.8), 20 ml of 20% SDS, 20 ml glycerol, and 40 ml de-ionized H₂O. Samples were heated at 100°C for 5 minutes, and protein concentrations were determined using the BioRad DC assay. DTT and bromophenol

blue were then added to the remaining sample (from a 1 M DTT, 0.1% bromophenol blue stock prepared in 1x Laemmli buffer) in a ratio of 9 parts sample to 1 part DTT/bromophenol blue stock. The sample tube was then heated again at 100°C for 5 minutes. For CO1 determinations, 20 µg of mitochondrial protein were loaded onto 4-20% Tris-HCl polyacrylamide gels (BioRad). For CO4 determinations, 10 µg of whole cell protein were loaded onto 4-20% Tris-HCl polyacrylamide gels. Gels were electrophoresed for 45 minutes at 200 V. Proteins were transferred to polyvinylidene difluoride (PVDF) membranes (BioRad) using 200 mA for 2 hours. The membranes were blocked in Tris-buffered saline (TBS; 50 mM Tris base, 150 mM NaCl, pH 7.5) containing 0.05% Tween-20 and 5% milk for 1 hour at room temperature. Membranes were incubated overnight at 4°C with monoclonal antibodies to CO1 or CO4 (Molecular probes). After stripping, the CO4 blots were later re-stained with a monoclonal antibody to α -tubulin (Calbiochem). After overnight staining with primary antibodies, membranes were washed four times with TBS containing 0.05% Tween-20 (TTBS) and incubated for 1 hour at room temperature with an HRP-conjugated goat anti-mouse antibody (Jackson ImmunoResearch, West Grove, PA). Unbound secondary antibody was removed by four 10 min TTBS washes. The HRP-conjugated secondary antibody was activated with a solution containing an HRP substrate (West Dura Substrate; Pierce, Rockford, IL) and visualized with a Kodak 440 CF Image Station (PerkinElmer Life and Analytical Sciences Inc.). Band densities were determined using Kodak 440CF Image Station software. At least five independent samples were analyzed for each pioglitazone treatment condition, and the mean densities for each condition calculated. Mean densities for the different pioglitazone-treated cell groups were compared to that of the pioglitazone-untreated cell group.

Real time PCR. We used real time PCR to compare relative amounts of mtDNA and nuclear DNA copy numbers within pioglitazone-treated and untreated cells. Amplicons to two different mtDNA regions and one nuclear DNA region were generated for this analysis. The mtDNA amplicons were generated from the CO1 and D-loop segments. The nuclear amplicon was generated through amplification of a β -actin segment. The mtDNA primers were designed to minimize amplification of mtDNA pseudogenes embedded in nuclear DNA

(www.ncbi.nlm.nih.gov/BLAST), and were obtained from Operon (Operon Biotechnologies, Inc., Huntsville, AL). The CO1 primers were as follows: 5'-ATACTACTAACAGACCGCAACCTC-3' (forward primer), and 5'-GAGATTATTCCGAAGCCTGGT-3' (reverse primer). These primers generated a 143 base pair product. The D-loop primers were as follows: 5'-GGAGCACCTATGTGCGCAGTATC-3' (forward primer), and 5'-TGTCTGTGTGGAAAGTGGCTGTG-3' (reverse primer). These primers generated a 106 base pair product. The β -actin primers were as follows: 5'-GAAGGATTCCTATGTGGGCGA-3' (forward primer), and 5'-CAGGGTGAGGATGCCTCTCTT-3'. These primers generated a 103 base pair product.

External standards were prepared as previously described (von Wurmb-Schwark et al., 2002; May-Panloup et al., 2005) with minor modifications. PCR reactions were performed using iProof High Fidelity DNA Polymerase (BioRad) and 100 ng genomic DNA. Thirty cycles of denaturation (30 seconds at 94°C), primer annealing (30 seconds at 58°C), and extension (1 min at 72°C) were performed. Amplicons were purified using QIAquick tubes (Qiagen) and quantified with a DU series spectrophotometer (Beckman Coulter). From each of these amplicons five serial ten-fold dilutions were prepared and stored at -20°C as single use aliquots.

An iCycler iQ Real Time PCR Detection System thermocycler (BioRad) was used to perform real-time PCR amplification using iQ SYBR Green Supermix (BioRad). 100 ng of genomic DNA were used for each reaction. Mitochondrial to nuclear DNA ratios were calculated by dividing the mtDNA quantity for each sample by its corresponding β -actin quantity. The mtDNA : nuclear DNA ratios from pioglitazone-treated cells were compared to that of untreated cells.

Statistical analysis. Data are expressed as mean values \pm standard error of the mean (SEM) for all independent assays of a particular condition. Two-tailed Student's *t*-tests were used to compare group means. *P* values of less than 0.05 were considered significant.

Results

Effects of pioglitazone on cell mitochondrial content and quantity

We used three different approaches to assess whether pioglitazone influences the content or quantity of NT2 cell mitochondria. Our evaluation included real time PCR determination of relative mitochondrial DNA (mtDNA) copy number, immunochemical quantification of both mtDNA and nuclear DNA-encoded mitochondrial proteins, and quantitative measurement of citrate synthase activity (a physiologic correlate of cell mitochondrial mass). The real time PCR data are shown in Figure 1. We independently analyzed mtDNA copy number for two parts of the mtDNA, the CO1 gene and the D-loop. In both instances mtDNA copy number was increased. This was the case for pioglitazone exposures of 10 μ M, 20 μ M, and 40 μ M.

The immunochemical quantification studies used a Western blot approach to measure levels of the cytochrome oxidase CO1 subunit and the cytochrome oxidase CO4 subunit. These data are shown in Figure 2. CO1 is mtDNA encoded, while CO4 is encoded by a nuclear gene, translated in the cytoplasm, and imported into mitochondria. Figure 2 shows levels of both proteins were elevated at sub-toxic pioglitazone concentrations. In each case, with 40 μ M pioglitazone there was increased variance of data and results between concentrations were not significantly different.

Citrate synthase is encoded by a nuclear gene, translated in the cytoplasm, and imported into the mitochondrial matrix. Figure 3 shows the Vmax of the enzyme was clearly increased at 20 μ M pioglitazone. At 40 μ M, the citrate synthase Vmax was actually decreased below baseline.

Effects on cell respiration and electron transport chain function

We used an assay of cell oxygen consumption to assess mitochondrial respiratory rates in pioglitazone-treated cells. Assays were performed using oxygen biosensor plates (BioRad), which facilitate quantitative assessments of cell oxygen consumption. Data were analyzed in terms of oxygen consumption rates and absolute oxygen consumption at 1 hour (Figure 4). For cells maintained in medium containing 20 uM pioglitazone, there was an approximate 20-30% increase in both cell oxygen consumption rates and absolute oxygen consumption. This was not the case with exposure to 40 uM pioglitazone, suggesting this concentration induced functional mitochondrial toxicity.

High cell respiration rates could correlate with increased electron transport chain (ETC) enzyme activities. Figure 5 shows this was indeed the case. We determined complex I V_{max} activity in purified mitochondrial fractions prepared from NT2 cells disrupted through nitrogen cavitation, and complex IV (cytochrome oxidase) V_{max} activity in whole NT2 cells. The results of these spectrophotometric biochemical assays are shown in Figure 5. Complex I activity was increased with 10 uM pioglitazone. With 20 uM pioglitazone there was a trend towards increased complex I activity that was not statistically significant. At 40 uM, there was a suggested trend towards reduced complex I activity. There was a non-significant trend towards increased complex IV activity with 5 uM pioglitazone ($p=0.075$). Complex IV activity was increased with 10 uM and 20 uM pioglitazone exposures, but not with exposure to 40 uM pioglitazone.

Effects on reactive oxygen species production and oxidative stress

We utilized three different approaches to measure peroxide production. First, we used DCF, a peroxide-sensitive fluorescent dye. DCF is localized to cell cytoplasm, and provides sensitive measurements of cytoplasmic peroxide levels. Cells that overproduce peroxide can compensate to some extent by upregulating antioxidant enzyme activities. We felt it was important to determine whether this was indeed happening, and thus we measured GPx activities in our pioglitazone-exposed NT2 cells. Finally, we directly assayed mitochondrial peroxide production using another fluorescent dye, amplex red.

Figure 6 summarizes the results of these assays. The DCF assay demonstrated a pioglitazone dose-dependent increase in cytoplasmic peroxide production. Associated with this increase in cytoplasmic peroxide production was a concomitant upregulation of glutathione peroxidase activity, indicating the observed increase in cytoplasmic peroxide levels was enough to induce a cell compensatory response. Paradoxically, we also found a robust reduction in mitochondrial peroxide levels.

Discussion

Chronic exposure to the thiazolidinedione pioglitazone affects mitochondria. The pioglitazone-induced mtDNA increase we observed is consistent with either more mtDNA per mitochondrion or increased mitochondrial number. Similarly, increased ETC protein and citrate synthase activity (per fixed amounts of total cell protein) could reflect denser packaging of cell mitochondrial protein or more mitochondria per cell. We feel in each case the latter scenario, mitochondrial biogenesis, is more likely. Based also on real time PCR analysis of mtDNA quantity, expression of nuclear-encoded, mitochondria-relevant genes, and expression levels of mtDNA genes other groups previously concluded pioglitazone induces mitochondrial biogenesis in human subcutaneous adipose tissue and cultured preadipocytes (Bogacka et al., 2005; Roses and Saunders, 2006; Roses et al, 2007). If pioglitazone is indeed promoting mitochondrial biogenesis, it is worth considering the fact PPARG and NRF-1 transcription factors share a common co-activator, PGC-1 α (Scarpulla, 2002). NRF-1 promotes mitochondrial biogenesis. Although our data do not address this possibility, it is worth considering whether thiazolidinedione drugs that mimic the effects of PGC-1 α on PPARG also mimic the effects of PGC-1 α on NRF-1.

Although NT2 cell citrate synthase activity was increased with chronic exposure to 20 μ M pioglitazone, with 40 μ M exposure it was decreased from baseline. *In vitro* pioglitazone studies typically utilize 10-20 μ M concentrations. These concentrations exceed clinically achieved serum levels. Budde et al. determined serum pioglitazone concentrations in pioglitazone-treated diabetics. The mean serum C_{max} was 1.329 μ g/ml, or ~3 μ M (10 μ M=3.92 μ g/ml) (Budde et al., 2002; Ito et al., 2003). We suspect decreased citrate synthase activity at 40 μ M pioglitazone reflects toxicity at that concentration. Although our real time PCR data suggest

mitochondrial biogenesis was still promoted at 40 μ M pioglitazone, production of functional mitochondria was reduced. Decreased citrate synthase activity may reflect compromised mitochondrial integrity, impaired mitochondrial import, or an oxidative stress-mediated inhibition of the enzyme. Our oxidative stress determinations argue the latter scenario is less likely, because mitochondrial peroxide was reduced at all concentrations of pioglitazone evaluated. Taken together, our studies suggest at sufficient concentrations pioglitazone induces mitochondrial proliferation via induction of mitochondrial biogenesis, but at clinically toxic concentrations mitochondrial integrity is compromised.

Our oxygen consumption, complex I V_{max} , and complex IV V_{max} data support this view. These parameters were enhanced with chronic 10-20 μ M pioglitazone exposure, but not with 40 μ M exposure. If the observed loss of functional enhancement at the higher concentration is indeed a toxicity effect, elucidating the underlying mechanism is worthwhile. Possibilities include the cytoplasmic oxidative stress induced by pioglitazone, altered stoichiometric relationships between nuclear DNA and mtDNA encoded ETC subunits, or drug-induced expression of nuclear genes that influence mitochondrial function.

To better assess dose-response relationships we tested the effects of 5 μ M pioglitazone on complex IV activity. We cannot conclude chronic 5 μ M pioglitazone exposure has no effect on mitochondrial biogenesis, but can conclude 5 μ M pioglitazone produces at most a less robust response than 10 μ M pioglitazone. We therefore assume more pioglitazone is needed to induce mitochondrial biogenesis than is needed to activate PPARG.

Complex IV activity was increased with both 10 μ M and 20 μ M pioglitazone. Although the complex I V_{max} was also increased at 10 μ M pioglitazone, we did not show this effect with 20 μ M pioglitazone. Measuring complex I activity is more technically difficult than measuring

complex IV activity, and so this may reflect type II error. Alternatively, this finding may be real. If so, several explanations are possible. Subtle increases in mitochondrial fragility may have begun even with 20 μ M pioglitazone, to the point that mitochondrial purification or sonication procedures caused excess disruption of mitochondrial integrity. Complex I consists of 46 bi-genomically encoded protein subunits (as opposed to 13 for complex IV), and so may be more sensitive to minor inequities between mitochondrial and nuclear-encoded ETC subunit gene expression (Carroll et al., 2005). Another potential explanation relates to the fact that radical species can negatively regulate ETC function, and complex I is susceptible to this (Riobo et al., 2001). Although cytoplasmic peroxide levels increased in a dose-dependent fashion, mitochondrial peroxide levels did not. We therefore think this latter possibility should not account for the lack of complex I V_{\max} increase with 20 μ M pioglitazone.

Acute exposure of isolated mitochondria to thiazolidinediones reduces complex I activity and pyruvate-driven state III respiration (Dello Russo et al., 2003; Brunmair et al., 2004; Scatema et al., 2004; Feinstein et al., 2005). Dello Russo et al. also found mitochondrial hyperpolarization occurs in cultured rat astrocytes exposed to pioglitazone for 6 hours (Dello Russo et al., 2003). The data we obtained using a chronic exposure model, in which human NT2 cells were maintained for weeks in pioglitazone, deviate from those of acute exposure models. Mitochondria in dividing cells over time may respond or adapt to thiazolidinediones differently than mitochondria exposed following removal from cells or after brief intracellular exposures. Our data in no way impugn those from acute exposure models, as critical methodologic differences distinguish our studies from acute exposure studies. Different results between these two approaches suggest, though, that mechanisms influencing mitochondrial function in our chronic exposure model ultimately extend beyond those occurring under acute exposure

conditions. A relationship between time-dependent mechanisms may nevertheless exist, since mitochondrial dysfunction is itself associated with mitochondrial biogenesis (Heddi et al., 1999). It is tempting to speculate acute pioglitazone exposure may trigger a secondary, delayed mitochondrial biogenesis.

Observed increases in complex I and IV V_{max} activities could reflect quantitative or qualitative alterations of these enzymes. Our citrate synthase data are relevant in this regard. Dividing complex I and IV activities by citrate synthase activity at least partly corrects ETC V_{max} increases. This suggests increased mitochondrial quantity mostly accounts for ETC activity increases.

Oxidative stress may contribute to thiazolidinedione toxicity (Shishido et al., 2003). In this study increasing pioglitazone concentrations correlated with cytoplasmic hydrogen peroxide levels. This rise in peroxide occurred despite increased GPx activity. The mechanisms underlying thiazolidinedione-related free radical production are unclear. We considered the possibility increased ETC throughput might secondarily increase oxidative stress. After all, running the ETC of hyperpolarized mitochondria promotes superoxide production. However, in our experiments isolated mitochondria showed reduced peroxide levels, suggesting mitochondria are not responsible for the cytoplasmic increase. Alternative sources of cytoplasmic peroxide include peroxisomes, membrane NADH oxidoreductases, and NADPH oxidase.

Reduced mitochondrial peroxide levels could reflect robust upregulation of mitochondrial antioxidant enzymes. Alternatively, a disproportionate increase in complex III subunits could reduce ETC “electron leak” and lower superoxide production. Dissociation of electron transport from hydrogen translocation/ADP phosphorylation (uncoupling) requires consideration (Kelly et

al., 1998; Digby et al., 1998; Shimokawa et al., 1998; Andrews et al., 2005; Douette and Sluse, 2006).

Insulin resistance in type II diabetes is largely determined by muscle mitochondria (Lowell and Shulman, 2005). While pioglitazone reduces insulin resistance, the role mitochondria might play in this has not been widely considered. Our experiments used a neuronal-like cell line rather than muscle, but nevertheless argue pioglitazone may benefit diabetics by affecting mitochondria. Various neurodegenerative diseases are also associated with ETC enzyme activity reductions and increased mitochondria-generated oxidative stress (Swerdlow, 2007). The ability of pioglitazone to increase complex I and IV Vmax activities and reduce mitochondrial peroxide levels in a neuronal-like cell line make it interesting from a neurodegenerative disease-therapeutic perspective.

The use of thiazolidinedione drugs for AD was previously advocated (Heneka et al., 2005; Watson et al., 2005; Risner et al., 2006; Geldmacher et al., 2006, Landreth, 2006). Exploratory analyses from a phase II trial of rosiglitazone in AD suggest rosiglitazone might benefit cognition in those not carrying *APOE4* alleles (Risner et al., 2006). Interestingly, an apolipoprotein E-mitochondrial nexus exists, since apolipoprotein E C-terminal fragments induce mitochondrial dysfunction (Chang et al, 2005). Interactions between rosiglitazone response and *APOE* in the Risner et al. trial could support a hypothesis stating apolipoprotein E fragments may facilitate AD by altering mitochondrial physiology (Roses et al, 2007).

Proposed rationales for trying thiazolidinediones in AD include their effects on insulin, amyloid precursor protein processing, and inflammation. Our data support another rationale, previously proposed by others (Roses et al, 2007), that postulates thiazolidinediones may benefit AD subjects by enhancing mitochondrial function. Our data further suggest thiazolidinediones

might benefit other neurodegenerative diseases with mitochondrial dysfunction (Swerdlow, 2007). Current thiazolidinediones, though, do not robustly pass the blood brain barrier. One study reported cerebrospinal fluid concentrations of pioglitazone were only 18% that of serum (Maeshiba et al., 1997; Heneka et al., 2005). While serum concentrations of pioglitazone and its active metabolites have surpassed 100 μM in animal studies, under clinical conditions serum concentrations are in the 2-3 μM range (Budde et al., 2002; Feinstein et al., 2005). Development of thiazolidinedione mimetics capable of crossing the blood brain barrier seems justified.

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Footnotes

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Figure Legends

Figure 1. Chronic pioglitazone treatment increases NT2 cell mtDNA content. Data reflect relative mtDNA content of pioglitazone-treated cells versus that of cells not treated with pioglitazone. The analysis was done using two different mtDNA primer sets. One primer set was to CO1 of the mtDNA (A), and the other was to the D-loop (B).

Figure 2. Levels of electron transport chain subunits are increased by chronic pioglitazone exposure. The graphs compare densitometry readings from Western blots of mitochondrial protein (for CO1) or whole cell protein (for CO4) prepared from pioglitazone treated and untreated cells. Mean density values from pioglitazone treated cells are reported relative to that of untreated cells. (A) shows pioglitazone treatment increased levels of CO1, an mtDNA-encoded electron transport chain subunit. (B) shows pioglitazone treatment increased levels of CO4, a nuclear-encoded electron transport chain subunit. For the CO4 analysis, tubulin was used as an internal loading control, and tubulin levels were equivalent between pioglitazone treated and untreated cells. (C) shows a representative CO4 experiment with its tubulin control.

Figure 3. Citrate synthase activities from pioglitazone treated and untreated cells. Data were recorded as nmol/min/mg protein, and in the figure are reported as mean activity of pioglitazone treated cells relative to that of pioglitazone untreated cells.

Figure 4. Relative to untreated cells, oxygen consumption in pioglitazone treated cells is increased. Data are reported two different ways. In (A), the oxygen consumption slopes from

pioglitazone treated cells are shown relative to the slope of untreated cells. In (B), the relative amounts of oxygen consumed over 1 hour are shown.

Figure 5. Relative to untreated cells, electron transport chain Vmax activities from pioglitazone treated cells are increased. (A) Complex I activity. (B) Complex IV activity.

Figure 6. Cell peroxide production and oxidative stress levels are affected by chronic pioglitazone exposure. (A) Relative to untreated cells, higher DCF fluorescence in pioglitazone treated cells indicates higher cytoplasmic peroxide levels. (B) Relative to untreated cells, pioglitazone treated cells show increased glutathione peroxidase activity. (C) Using amplex red fluorescence as a measure of peroxide, peroxide levels are lower in mitochondria isolated from pioglitazone treated cells than in mitochondria isolated from untreated cells.

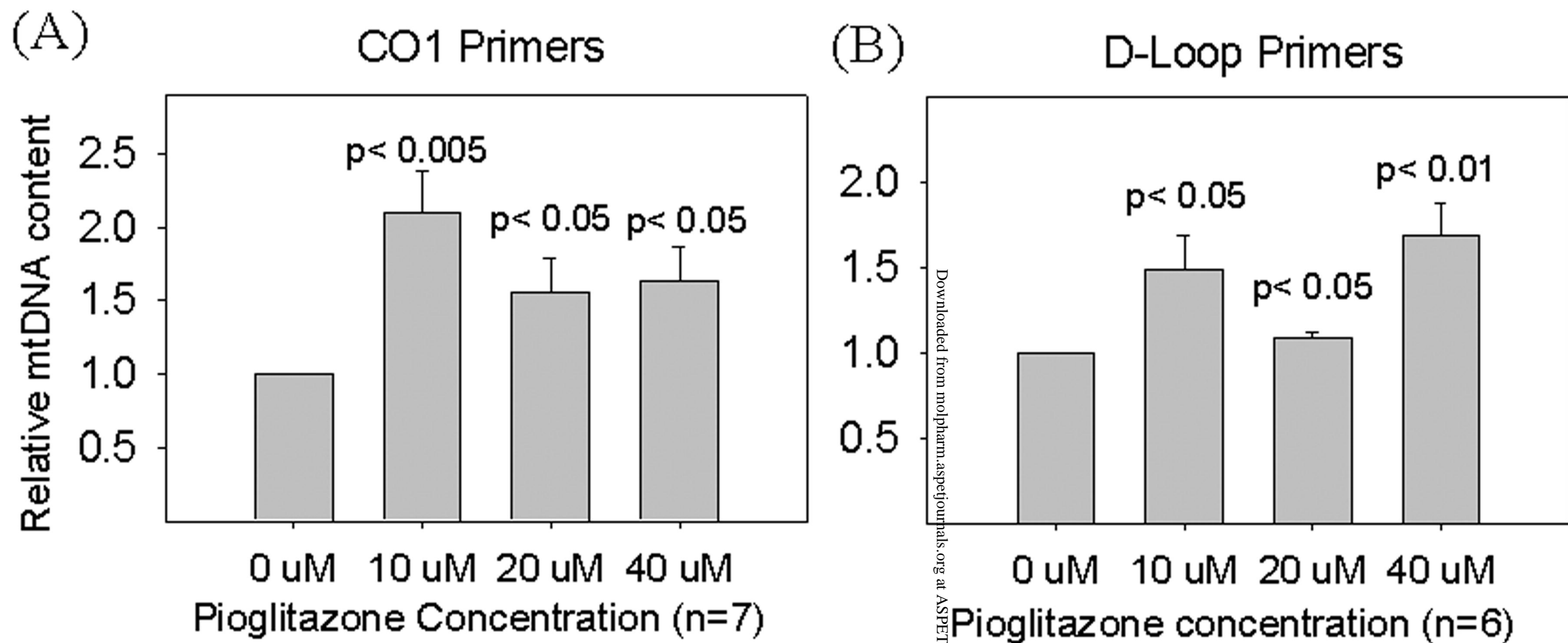


Figure 1

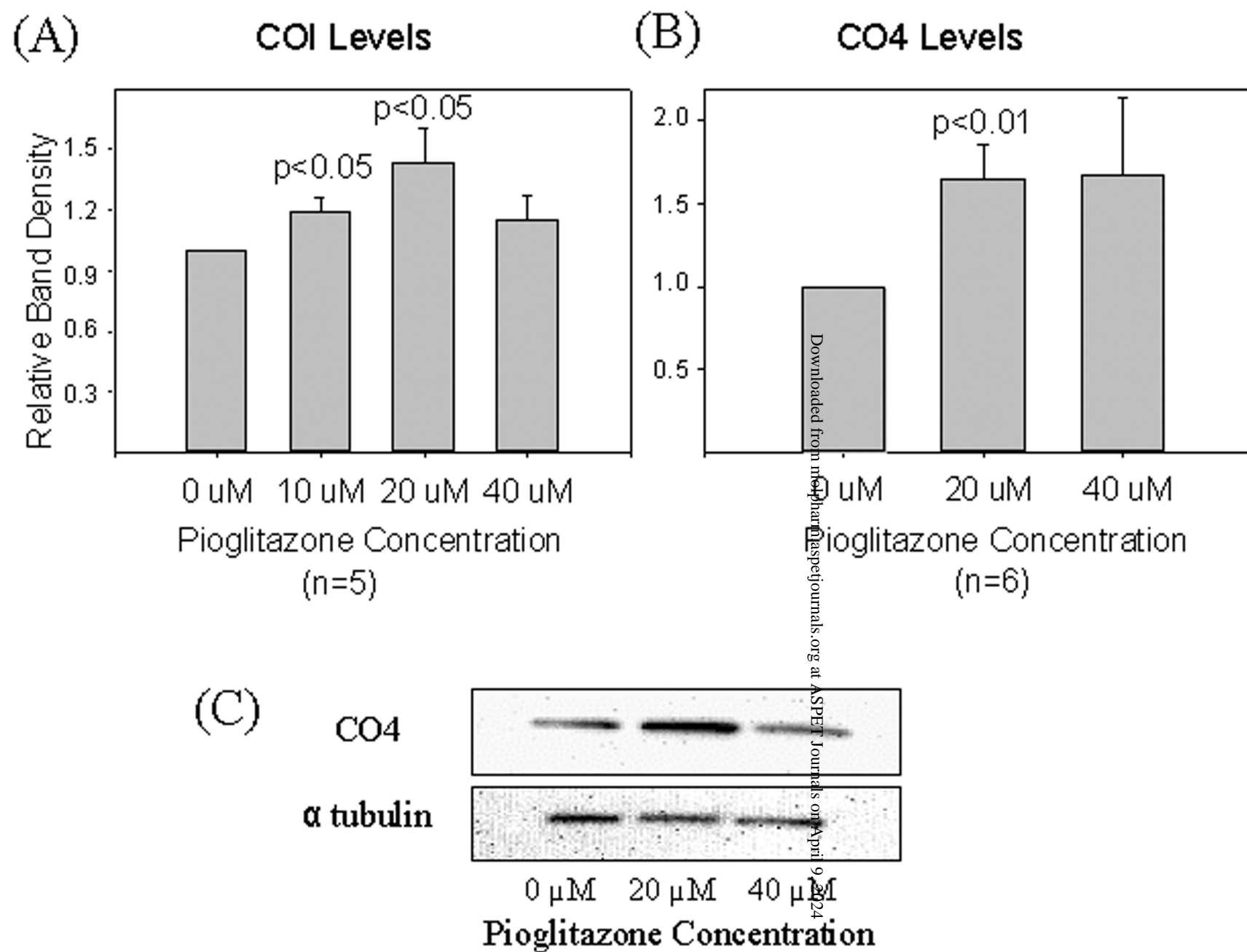


Figure 2

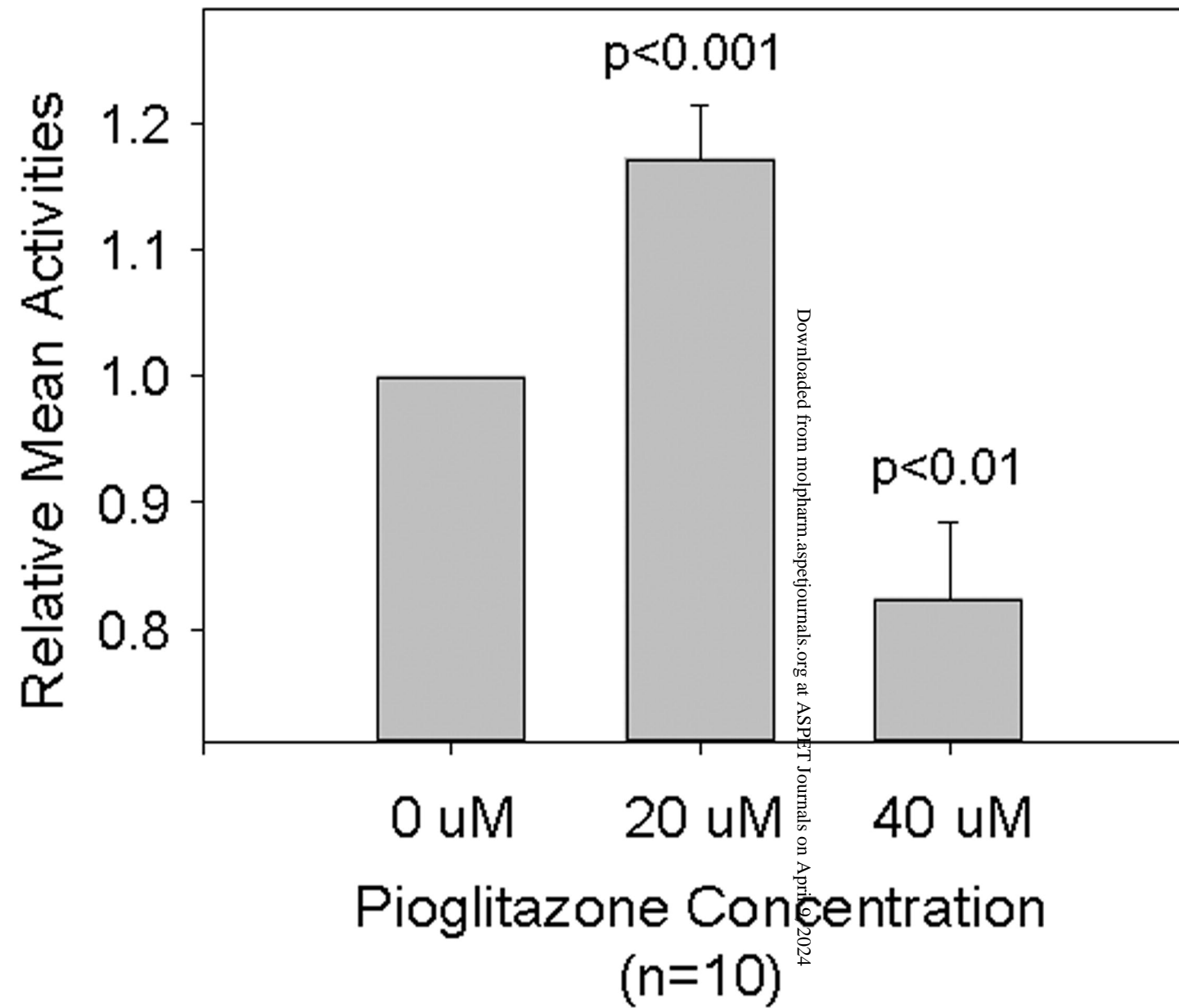


Figure 3

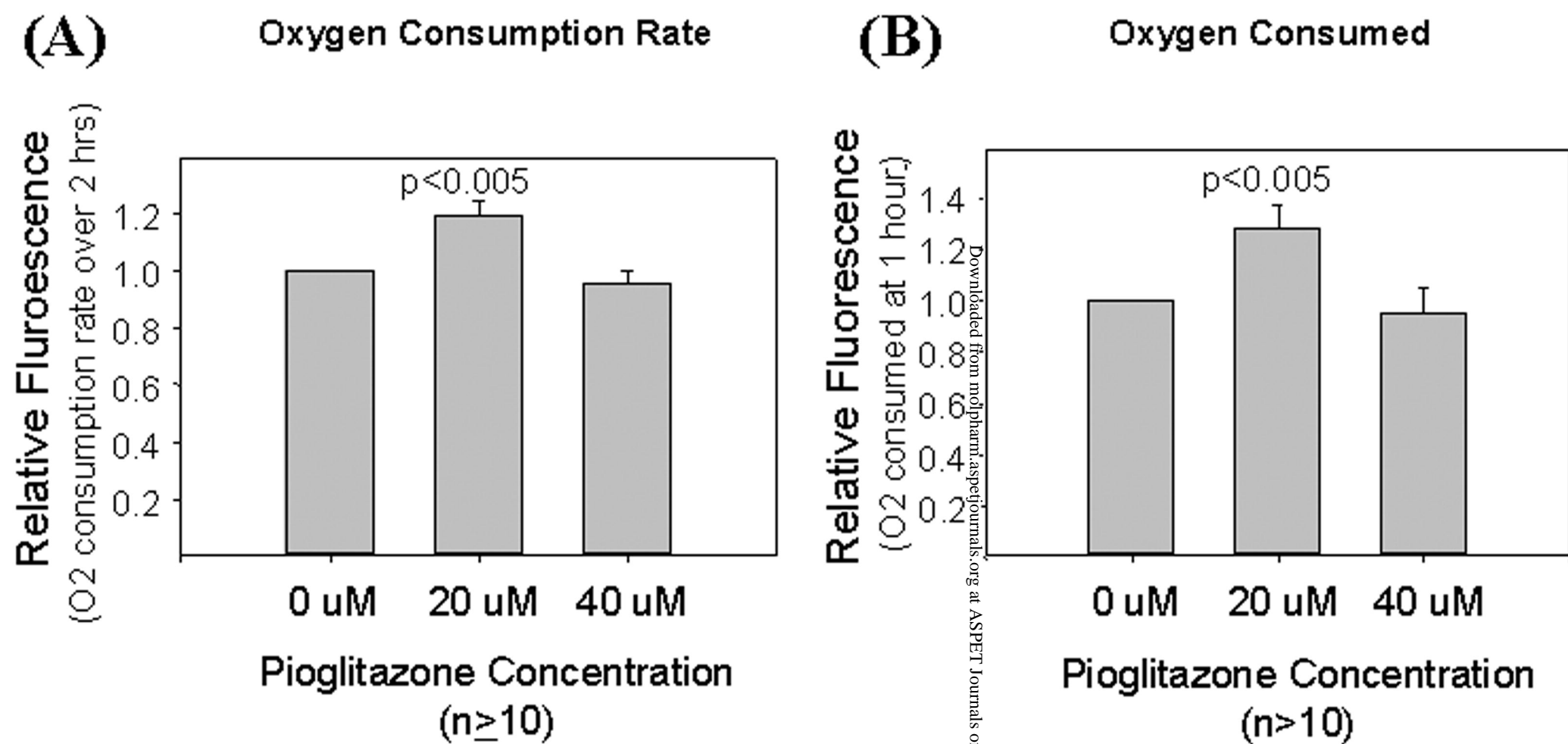
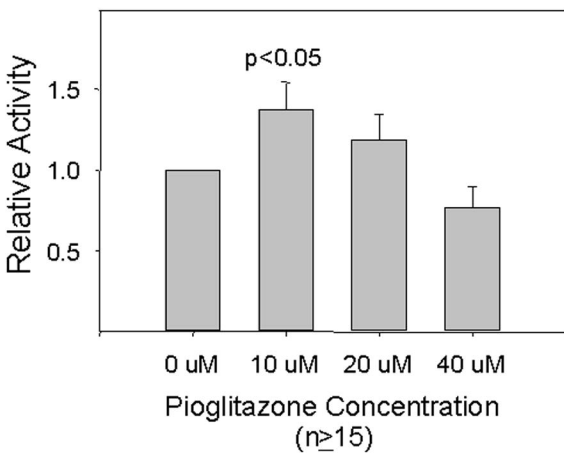


Figure 4

(A) Complex I Activity



(B) Complex IV Activity

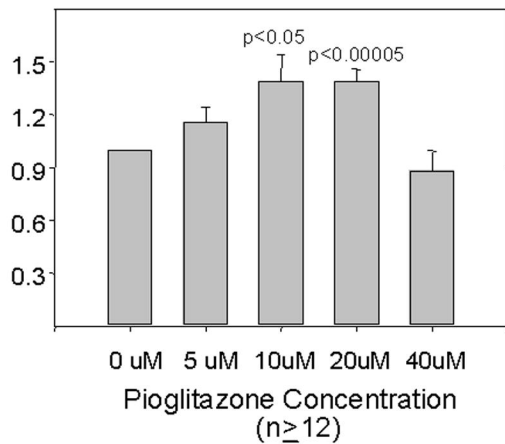


Figure 5

DCF Fluorescence Increase Slope
(Relative Arbitrary Units)

(A) DCF ROS assessment

(B) Glutathione peroxidase activity

(C) Amplex red ROS assessment

Relative GPX activity

Relative Fluorescence

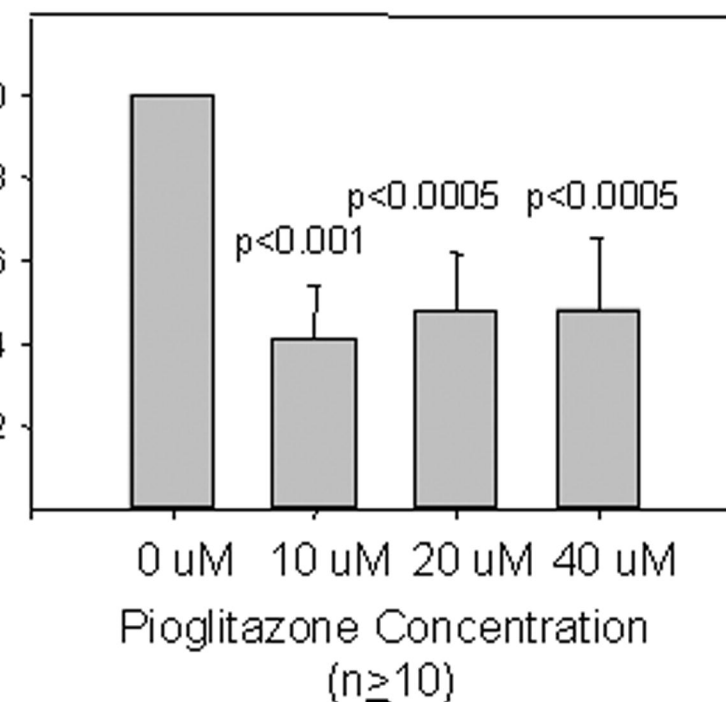
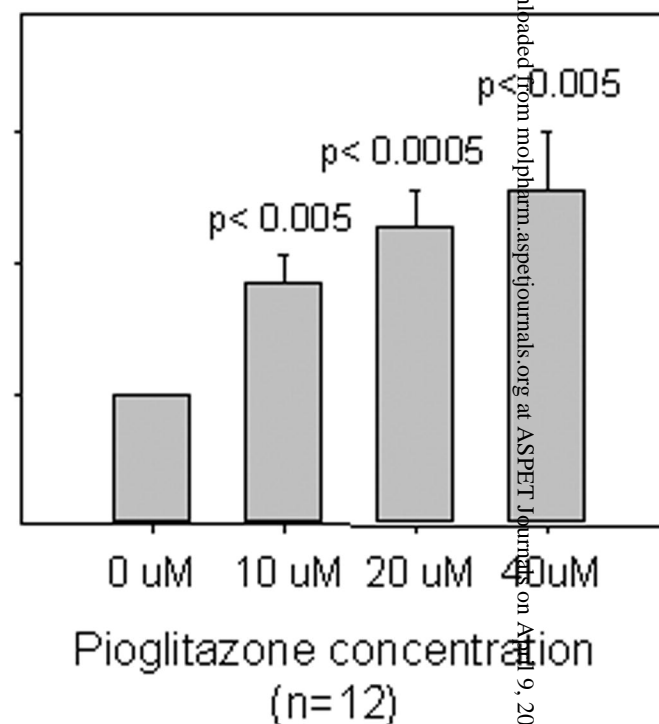
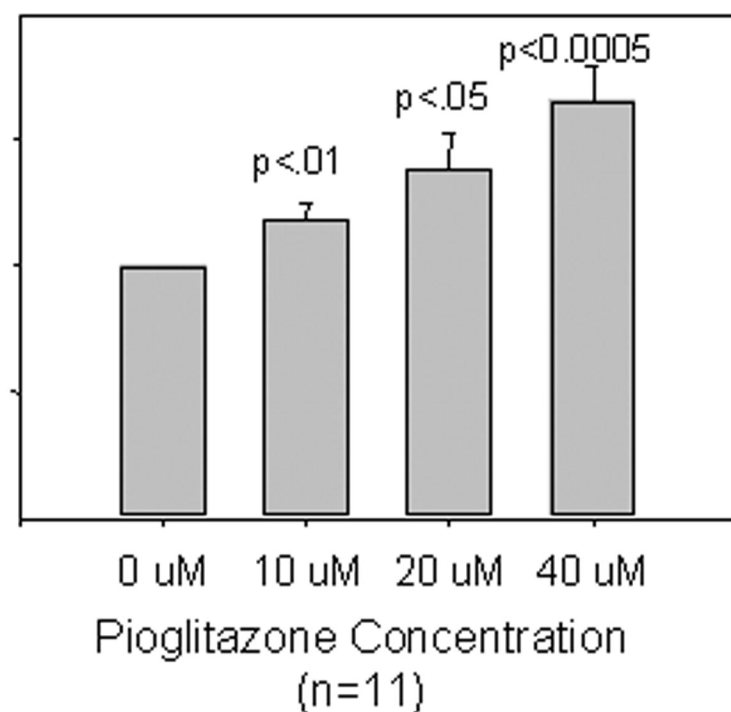


Figure 6