The PPARδ Ligand GW501516 Reduces Growth but Not Apoptosis in Mouse Inner Medullary Collecting Duct Cells

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The collecting duct (CD) expresses considerable amounts of PPARδ. While its role is unknown in the CD, in other renal cells it has been shown to regulate both growth and apoptosis. We thus hypothesized that PPARδ reduces apoptotic responses and stimulates cell growth in the mouse CD, and examined the effect of GW501516, a synthetic PPARδ ligand, on these responses in mouse IMCD-K2 cells. High doses of GW501516 decreased both DNA and protein synthesis in these cells by 80%, but had no overall effect on cell viability. Although anisomycin treatment resulted in an increase of caspase-3 levels of about 2.59-fold of control, GW501516 did not affect anisomycin-induced changes in active caspase-3 levels. These results show that a PPARδ ligand inhibits growth but does not affect anisomycin-apoptosis in a mouse IMCD cell line. This could have therapeutic implications for renal diseases associated with increased CD growth responses.

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1. Introduction

Traditional prostaglandin (PG) signaling in renal epithelial cells occurs through activation of prostanooid receptors [1–3]. In recent years, however, a novel signaling pathway for PGs has been found in which ligand-activated transcription factors called peroxisome proliferator-activated receptors (PPARs) [4] are activated at the nuclear membrane. One PPAR type, PPARδ, also known as PPARβ, is ubiquitously expressed in almost every tissue examined [5]. Both prostanooids PGI2 and PGE2 have been known to activate PPARδ [6, 7]. The main role of PPARδ is in cell survival and this has been well characterized in colon cancer [8–10].

In the kidney, PPARδ is found in every part of the nephron with high levels in the glomerulus, cortical collecting duct, and inner medullary collecting duct (IMCD) and low expression in the outer medullary collecting duct. This PPAR has been implicated in the renal complications of metabolic syndrome [11], but most studies describe a link between PPARδ and survival of renal cells. In a study by Hao et al. [6], PPARδ was shown to increase renal medullary interstitial cell survival during high levels of hypertonic stress, in response to PGI2. PPARδ activation also protected kidneys from renal failure after renal ischemia/reperfusion experiments via the antiapoptotic Akt pathway [12]. Although, most of the studies have suggested an antiapoptotic role of PPARδ, a few papers have shown that PPARδ can increase apoptosis [13]. However, PPARδ is recognized as an important protein in the survival of many renal cell types.

PPARδ has also been shown to promote growth responses in a variety of cells. PGE2, working through PPARδ, can stimulate proliferation of stem cells [14] and overexpression of PPARδ has been shown to reverse inhibitory growth signals in a prostate epithelial cell line [15]. Furthermore, upregulation of PPARδ in smooth muscle cells promotes cell cycle progression [16]. The effect of PPARδ activation on cell growth in the kidney, however, is unknown.

The collecting duct (CD), specifically the IMCD, survives in a harsh environment due to the hypertonic conditions in the interstitium needed to concentrate urine [17, 18]. In most cells, apoptosis would occur at the levels of stress at which they reside, but the CD manages to survive and
resist any apoptotic activity [19]. It is known that cells react to high levels of tonicity by increasing the transport of several osmolytes into the cells aiding in their survival [20]. However, in cells where there is chronic hypertonicity, like the IMCD, the accumulation of these osmolytes may not be sufficient for their continued survival [21]. Cultured IMCD cells have been known to increase levels of cyclooxygenase-2 (COX-2) and stimulate PGE2 release up to 33-fold of control in response to hypertonic stress [22]. It is likely then that PGs may contribute to the inherent CD resistance to stress by targeting PPARδ to promote cell survival.

It would appear in most cell lines that PPARδ has opposing growth and apoptotic effects; however, its role in the CD has not been determined. Our study examines the role of PPARδ activation in the survival and growth of IMCD-K2 cells. The mouse IMCD-K2 cell line is derived from the initial portion of the IMCD from a mouse that is transgenic for Simian Virus 40 (SV40) and was shown to retain many characteristics of the intact IMCD, including amiloride-sensitive sodium absorption stimulated by aldosterone [23]. Therefore, we propose that PPARδ activation will regulate growth and apoptotic responses in the mouse IMCD-K2 cells.
2. Materials and Methods

2.1. IMCD-K2 Cell Culture. Cells were obtained from Dr. Bruce Stanton from Dartmouth Medical College (Hanover, NH, USA). The cells were maintained at 37°C and 5% CO₂ in DMEM/F-12 supplemented with 10% fetal bovine serum, (FBS, Gibco, Carlsbad, Calif, USA), 1% penicillin/streptomycin (Gibco), ITS (5 μg/mL insulin, 5 μg/mL transferrin, 5 ng/mL selenium; Sigma, St. Louis, Mo, USA), 2 mM L-glutamine (Sigma), and 5 μM dexamethasone (Sigma). Cell culture medium was replaced every 48 hours. Twenty-four hours prior to experiments, the culture medium was replaced with serum-free medium (DMEM/F-12).

2.2. Reverse Transcriptase-PCR. RNA was isolated from confluent 100 mm plates of IMCD-K2 cells using 1 mL of the TRIzol reagent (Gibco) according to the manufacturer’s instructions and then treated with DNase I (Invitrogen, Carlsbad, Calif, USA). The RNA was reverse transcribed using MuLV reverse transcriptase and random hexamers.
Serum-starved cells were stimulated for 24 hours with serum-free media (control) or increasing concentrations of GW501516 (10⁻⁸ to 10⁻⁵ M) ³H-leucine was added to IMCD-K2 cells while they were being stimulated and leucine incorporation was measured in counts per minute using a scintillation counter, and expressed as fold control. Values are mean ± S.E.M.; n = 3–5. *P < .05 compared to control.

Figure 6: GW501516 causes a decrease in protein synthesis at high doses. Serum-starved cells were stimulated for 24 hours with serum-free media (control) or increasing concentrations of GW501516 (10⁻⁸ to 10⁻⁵ M) ³H-leucine was added to IMCD-K2 cells while they were being stimulated and leucine incorporation was measured in counts per minute using a scintillation counter, and expressed as fold control. Values are mean ± S.E.M.; n = 3–5. *P < .05 compared to control.

2.3. ³H-cAMP Assay. Since prostanoids can alter cAMP, their effect on cAMP production in IMCD-K2 cells was determined. The cells were serum starved for 24 hours and stimulated with control (serum-free media), 10⁻⁵ M forskolin (Sigma) as a positive control, 10⁻⁶ M ciprofloxin and iloprost (prostacyclin analogs, Schering Ag Berlin, Berlin, Germany), or 10⁻⁷–10⁻⁶ M of prostaglandin E₂ (PGE₂, Cayman, Ann Arbor, Mich, USA) The cells were pretreated for fifteen minutes with 0.5 mM isobutylmethylxanthine (IBMX, Sigma), a cAMP phosphodiesterase inhibitor, and 10⁻⁶ M indomethacin (Sigma), a cyclooxygenase (COX) inhibitor. The stimulation was then stopped with cold 10% trichloroacetic acid for 30 minutes on ice. Four ether extractions were carried out, and a cAMP competitive binding assay was done using the ³H-cAMP DPC radioassay (Intermedico, Markham, ON, Canada) as per manufacturer’s instructions. The amount of ³H-cAMP was measured using a scintillation counter (Beckman Coulter, Mississauga, ON, Canada) and the percent stimulation of cAMP levels over control levels of each sample was calculated.

2.4. Western Blotting. IMCD-K2 lysates were obtained by sonicating samples in RIPA lysis buffer containing 1% Non-ident P-40, 1% sodium deoxycholate, 0.1% sodium dodecyl sulphate (SDS, w/v), 4.5 mM NaCl, 2.5 mM Tris (pH 7.4),
24 hours pretreatment followed by 10−8 M GW501516 for 24 hours pretreatment followed by 10−6 M and 10−8 M GW501516 for 24 hours with (GW-6/A250 and GW-8/A250) or without anisomycin (GW-6 and GW-8), Z-VAD-FMK, a pan-caspase inhibitor with (Z/A250) or without (Z) anisomycin. Values are mean ± S.E.M.

Figure 9: Anisomycin increases caspase activity in IMCD-K2 cells, independent of GW501516. Serum-starved cells were treated with serum-free media (control) for 24 hours, 250 ng/mL anisomycin (A250) for 24 hours, 10−6 M GW501516, and 10−8 M GW501516 for 24 hours pretreatment followed by 10−6 M and 10−8 M GW501516 for 24 hours with (GW-6/A250 and GW-8/A250) or without anisomycin (GW-6 and GW-8), Z-VAD-FMK, a pan-caspase inhibitor with (Z/A250) or without (Z) anisomycin. Values are mean ± S.E.M.; n = 8.

8 μM EDTA, 0.2 mM sodium phosphate (pH 7.2), and fresh 0.5 mM PMSF, 1:100 protease inhibitor cocktail (Sigma), 1 mM sodium pyrophosphate, 10 mM sodium fluoride, and 100 μM sodium orthovanadate. The protein was quantified using the Bradford reagent method (Bio-Rad, Hercules, Calif, USA). The lysates were then resolved by SDS-PAGE on a polyacrylamide gel using a Mini-PROTEAN II apparatus (Bio-Rad) and transferred onto nitrocellulose membranes (Amersham, Amersham, UK). After blocking for 90 minutes in 10% milk/TBS-T (137 mM NaCl, 20 mM Tris base, 0.1% Tween20), the membranes were incubated overnight with the corresponding primary antibody. The membranes were then incubated with their respective secondary antibodies (i.e., anti-rabbit, anti-mouse, and anti-donkey) for 90 minutes and then washed for 60–90 minutes in TBS-T. The bands were then visualized by ECL. Detection of β-actin was used to normalize samples and densitometric analysis of the bands was performed using the Kodak 1D Image Analysis software (Eastman Kodak Company, Rochester, NY, USA).

2.5. Real-Time PCR. To determine the effect of PGE2 on PPARδ levels in IMCD-K2 cells, RNA was isolated and DNase-treated as described above. The cells were treated for 2, 8, or 24 hours with serum-free media as control, 10−5 M indomethacin, or 10−6 M PGE2. A fifteen-minute pretreatment with indomethacin was performed prior to stimulation with PGE2 and forskolin. The mRNA levels of PPARδ were ascertained by real-time PCR using TaqMan One-Step RT-PCR master mix reagents (Applied Biosystems) and an ABI Prism 7000 sequence detection system. Reactions were carried out by using 50 ng of total IMCD-K2 RNA under the following conditions: 48°C for 30 minutes, and 95°C for 10 minutes, and 40 cycles of 95°C for 15 seconds and 60°C for 1 minute. The probe and primers used for mouse PPARδ were forward primer, 5′-GAGCCCCAGTTGGTTTTGC-3′, reverse primer, 5′-TTGAAGGCGCCAGGTCTACT-3′, and probe, FAM-AGTTCAATGCCGCTGGAGCTGATGA-TAMRA (Sigma Genosys, Oakville, ON, Canada). All values were normalized to GAPDH mRNA levels in the same sample, which was determined by the TaqMan Rodent GAPDH control reagent kit (Applied Biosystems).

2.6. 3H-Thymidine Incorporation. To study the effect of the PPAR activation on cell proliferation, DNA synthesis was measured using the incorporation of 3H-thymidine. IMCD-K2 cells were cultured in 24-well plates, grown to ∼50% confluence, and then starved with serum-free media. Afterwards the cells were treated for 24 hours with vehicle-treated control (DMSO), or GW501516 (10−8, 10−7, 10−6, 10−5 M), GW501516 is among the more specific and most commonly used synthetic PPARδ ligands [24]. 3H-thymidine was forward primer, 5′-GAGCCCCAGTTGGTTTTGC-3′, reverse primer, 5′-TTGAAGGCGCCAGGTCTACT-3′, and probe, FAM-AGTTCAATGCCGCTGGAGCTGATGA-TAMRA (Sigma Genosys, Oakville, ON, Canada). All values were normalized to GAPDH mRNA levels in the same sample, which was determined by the TaqMan Rodent GAPDH control reagent kit (Applied Biosystems).
Figure 11: GW501516 pretreatment has no effect on anisomycin-induced apoptosis. Serum-starved cells were stimulated with (a)–(c) serum-free media (control), (d)–(f) $10^{-6}$ M GW501516 (GW-6), (g)–(i) $10^{-8}$ M GW501516 (GW-8), (j)–(l) 250 ng/mL anisomycin (A250) for 24 hours, (m)–(o) $10^{-6}$ M, or (p)–(r) $10^{-8}$ M GW501516 for 3 hours pretreatment followed by $10^{-6}$ M or $10^{-8}$ M GW501516 for 24 hours with A250 (GW-6/A250 and GW-8/A250). The cells were incubated with terminal transferase and biotin-16-dUTP for one hour, followed by two 45-minute incubations with streptavidin-Cy2 and anti-DAPI. The cells were then mounted on slides, visualized with a fluorescent microscope, and captured with a camera. Apoptotic cells are shown in green, while the blue represents DAPI-positive cells.
(Amersham, 0.5 μCi/mL) was added during the final four hours of stimulation. The plates were then washed four times in ice-cold PBS. Next, the cells were permeabilized in 500 μL of 1N NaOH at 37°C for ∼30 minutes, and the amount of 3H-thymidine in counts per minute (cpm) was measured using a scintillation counter. Samples were done in triplicate and thymidine incorporation is expressed as fold control.

2.7. 3H-Leucine Incorporation. To study the effect of the PPAR ligands on cell growth, protein synthesis was measured using the incorporation of 3H-leucine. IMCD-K2 cells were cultured in 24-well plates, grown to ∼50% confluence, and then starved with serum-free media for 24 hours. Next the cells were stimulated for 24 hours with vehicle-treated control (DMSO) and GW501516 (10⁻⁸, 10⁻⁷, 10⁻⁶, and 10⁻⁵ M). 3H-leucine (Amersham, Perkin Elmer, Waltham, Mass, USA, 0.5 μCi/mL) was added to each well during the 24-hour stimulation. The cells were then prepared as described above and the amount of 3H-leucine in counts per minute was measured using a scintillation counter. Samples were done in triplicate and leucine incorporation is expressed as fold control.

2.8. Cell Viability. To evaluate the effect of GW501516 on the viability of IMCD-K2 cells in both the presence and absence of a death response, a cell viability assay was performed. IMCD-K2 cells were grown to 70% confluence in 100 mm plates and starved for 24 hours. Next the cells were treated with 1 μL of ethanol and 1 μL of DMSO in DMEM/F-12, GW501516 (10⁻⁸, 10⁻⁷, 5 × 10⁻⁷, 10⁻⁶, 5 × 10⁻⁶, and 10⁻⁵ M), anisomycin (Sigma; 100, 1000 ng/mL), or cisplatin (Sigma; 50 ng/mL). Some cells were treated with both anisomycin and GW501516 (10⁻⁶ M) following a 24-hour pretreatment with GW501516 (10⁻⁶ M). Anisomycin and cisplatin were used to induce cell death. Following the stimulation, the cells were incubated in 100 μL of a 1x DNA dye-binding solution from the CyQANTA NF kit (Invitrogen). The fluorescence intensity, representing the number of cells, was measured in a FLUOstar Galaxy plate reader (BMG Labtechnologies, Durham, NC, USA). Samples were done in triplicate and fluorescence intensity is expressed as fold control.

2.9. Colorimetric Caspase-3 Activity Assay. To determine if GW501516 is involved in apoptosis, a caspase-3 activity assay was carried out using the colorimetric CaspACE Assay System kit (Promega, Madison, Wis, USA), as per manufacturer’s instructions. IMCD-K2 cells were grown to ∼50% confluence in 100 mm plates and starved for 24 hours. Next the cells were treated with control (1 μL of ethanol and 1 μL of DMSO in 1 mL of DMEM/F-12), 10% FBS, GW501516 (10⁻⁶ M), anisomycin (250 ng/mL), or Z-VAD-FMK (2 × 10⁻⁵ M), a pan-caspase inhibitor supplied in the kit. Some cells were treated with both anisomycin and GW501516 following a 24-hour pretreatment with GW501516. Also, some plates were treated with anisomycin and Z-VAD-FMK following a 24-hour pretreatment with the inhibitor. Following stimulations, the cells were resuspended in 100–200 μL of the caspase lysis buffer subjected to two freeze-thaw cycles (at ∼20°C), and a 15-minute incubation on ice. The lysates were then centrifuged for 20 minutes at 13 100 revolutions per minute and protein was quantified using the Bradford method. 50 μg of each sample was incubated with the colorimetric caspase-3 substrate Ac-DEVD-pNA (2 × 10⁻⁴ M) in a 96-well plate at room temperature overnight. The endogenous active caspase-3 bound to the substrate was determined by the absorbance at 405 nm (FLUOstar Galaxy). Samples were done in duplicate and the caspase activity is expressed as relative absorbance.

2.10. TUNEL Assay. To further evaluate the effect of GW501516 on apoptosis, a terminal deoxynucleotidyl transferase biotin-dUTP nick end labeling (TUNEL) assay was performed. IMCD-K2 cells were grown to ∼60% confluence on glass coverslips in a 24-well plate and treated in duplicate for 24 hours with control (1 μL of ethanol and 1 μL of DMSO in 1 mL of DMEM/F-12), GW501516 (10⁻⁸, 10⁻⁷ M), or anisomycin (250 ng/mL). Some cells were treated with both GW501516 (10⁻⁵, 10⁻⁶ M) and anisomycin, following a 3-hour pretreatment with the respective concentrations of GW501516. After the stimulation, the cells was fixed in 4% paraformaldehyde and 0.2% picric acid in 0.16 M sodium phosphate buffer (pH 6.9). The coverslips were then washed with PBS three times and incubated with 2 μL CoCl₂, 5 μL 5X TdT buffer, 0.1 μL terminal transferase (all taken from the Roche Terminal Transferase recombinant kit), 0.17 μL biotin-16-dUTP (Roche, Mississauga, ON, Canada), and 17.73 μL of H₂O for 1 hour at 37°C in a humid chamber. Next, the cells were washed with 4X SSC (0.6 M NaCl, 76 mM Na Citrate, pH 7), blocked for 30 minutes with 1% milk/4X SSC at room temperature, and incubated with streptavidin-Cy2 (Molecular Probes, Carlsbad, Calif, USA, 1.0 mg/mL) at 1:1000 for 45 minutes at room temperature in the dark. The coverslips were then washed with PBS and incubated with DAPI for 45 minutes at 1:1000. Following three more washes, the coverslips were inverted onto Fluormount mounting media (Electron Microscopy Sciences, Hatfield, Pa, USA) on glass slides. The cells were visualized with the fluorescent microscope Axioskop Mot 2 (Zeiss, Oberkochen, Germany) and captured with the camera AxioCam (Zeiss). For each coverslip, 8 fields of view at 200x magnification were examined using pictures from both the DAPI filter (excitation: ∼360 nm and emission: ∼450 nm) and the green filter (excitation: ∼480 nm and emission: ∼540 nm). The pictures were processed using the Axiovision software (Zeiss). The terminal transferase and biotin-16-dUTP target-free DNA fragments of apoptotic cells produce a fluorescent green signal. The number of apoptotic cells was counted, and from the DAPI filter a percentage of TUNEL-positive cells were determined. A total of 16 fields of view, or two coverslips, were used for each sample.

2.11. Statistics. The GraphPad Prism software (La Jolla, Calif, USA) for Windows (version 4.02) was used to analyze data. Results are expressed as means ± standard error of the mean.
(S.E.M). Either an unpaired t-test or a one-sample t-test (with a hypothetical value of 1) was used to evaluate the statistical significance between data points. A P-value < .05 was considered statistically significant.

3. Results

3.1. EP Receptor Subtypes and PPARδ Expression. PGE2 is known to activate prostanooid receptors, EP1, EP2, EP3, and EP4 as well as PPARδ. Thus, the presence of these receptors in IMCD-K2 cells was determined. PCR was performed and bands corresponding to the prostanooid and PPARδ receptors were visualized as shown in Figures 1(a) and 1(b), respectively. Previous studies in our laboratory demonstrated that at higher stringency the band in the EP2 lane disappears and that the product detected with the IP primers, present at approximately 200 base pair (bp) greater than the predicted size of 407 bp, does not correspond to the IP receptor, as determined by cloning and sequencing. The band seen between 700 and 800 bp in the EP1 lane represents protein kinase N (PKN), as the genes for PKN and EP1 overlap [25]. The multiple bands seen in the EP3 are most likely due to alternative splicing of the EP3 gene [26]. The products in Figure 2(a), prostacyclin (PGI2) analogs cicaprost (CCP) and bands corresponding to the prostanoid and PPAR receptors were visualized as shown in Figures 1(a) and 1(b), respectively. Previous studies in our laboratory demonstrated that at higher stringency the band in the EP2 lane disappears and that the product detected with the IP primers, present at approximately 200 base pair (bp) greater than the predicted size of 407 bp, does not correspond to the IP receptor, as determined by cloning and sequencing. The band seen between 700 and 800 bp in the EP1 lane represents protein kinase N (PKN), as the genes for PKN and EP1 overlap [25]. The multiple bands seen in the EP3 are most likely due to alternative splicing of the EP3 gene [26]. The products were not present when the reverse transcriptase was omitted (negative control; data not shown). These results indicate that IMCD-K2 cells express EP1, EP3, EP4, and PPARδ.

3.2. Effect of PGE2 on cAMP Stimulation. Both the EP3 and EP4 receptors elicit changes in intracellular cAMP; therefore, the effect of different prostanooids on cAMP stimulation was examined. Forskolin was used as a positive control. As shown in Figure 2(a), prostacyclin (PGI2) analogs cicaprost (CCP) and iloprost (ILP) did not alter cAMP levels. PGE2, however, did cause a significant increase in percent stimulation, at about 70% above control, comparable to forskolin. As shown in Figure 2(b), treatment with increasing amounts of PGE2 resulted in a concentration-dependent increase in cAMP, from 14.0 to 85.5%.

3.3. The Effect of PGE2 on PPARδ Expression. Since PGE2 has been known to activate PPARδ to regulate several growth processes [7, 14], it was ascertained that if PGE2 stimulation could affect PPARδ expression in IMCD-K2 cells. Indomethacin, a COX inhibitor, was used to inhibit endogenous PG synthesis. As shown in Figures 3(a) and 3(b), treatment with PGE2, in the presence of indomethacin, for 8 hours resulted in a significant increase, of ~2-fold, in PPARδ protein expression compared to control or indomethacin alone, as measured by Western Blotting. Stimulation for 24 hours also resulted in increased PPARδ protein expression, but this increase was not significant when compared with indomethacin alone. A representative blot for 24 hours is shown in Figure 3(a). However, two-hour stimulation did not alter the protein levels of PPARδ. We also examined the effect of PGE2 on PPARδ mRNA expression by real-time PCR. As shown in Figure 4, PPARδ mRNA remained unchanged with any treatment group or exposure time. Forskolin had no effect on expression either (data not shown) indicating that cAMP does not alter PPARδ protein or RNA expression.

3.4. The Effect of GW501516 on Cell Proliferation/Growth. PPARδ has been known to affect cell proliferation; therefore, we studied the effect of the PPARδ agonist, GW501516, on the proliferation/growth of IMCD-K2 cells. As shown in Figure 5, 10−3 M GW501516 reduced 3H-thymidine incorporation to about 0.05-fold control. PPARδ also regulates cell growth and as shown in Figure 6, 10−5 M GW501516 produced a significant reduction in 3H-leucine incorporation to approximately 0.19 of control. At lower concentrations of GW501516, neither DNA nor protein synthesis was altered.

3.5. The Effect of GW501516 on Cell Viability. To further characterize the effect of GW501516 on IMCD-K2 cells and determine if it has a role in cell survival, a cell viability assay was carried out. As shown in Figure 7, GW501516 (10−8, 10−7, 5 × 10−7, 10−6, 5 × 10−6, and 10−5 M) caused a decrease in cell viability between 0.8 and 0.6-fold of control but none of the values was significant. On its own, GW501516 did not induce death in the IMCD-K2 cells. To examine if the PPARδ agonist could protect cells from an induced death, IMCD-K2 cells were treated with both anisomycin and GW501516. Cisplatin (cis-diaminedichloroplatinum([II])) was used as an additional positive control for cell death. As shown in Figure 8, anisomycin (100 and 1000 ng/mL) caused cell viability to decrease to approximately 0.46- and 0.37-fold of control, respectively. However, 10−6 M GW501516 did not significantly alter the decrease in cell viability seen with the addition of either 100 or 1000 ng/mL of anisomycin.

3.6. The Effect of GW501516 on Caspase Activity. Besides having effects in cell growth, PPARδ has also been linked to changes in apoptosis. Therefore, the antiapoptotic effect of GW501516 was evaluated in IMCD-K2 cells. Caspase-3 activity was examined because it is a downstream effector protease that is common to all apoptotic pathways. As shown in Figure 9, anisomycin treatment resulted in an increase of caspase-3 levels of about 2.59-fold of control. Pre/cotreatment with GW501516 showed a decrease that was 0.79-fold (10−6 M) and 0.81-fold (10−5 M) of anisomycin alone. GW501516 alone showed no considerable change in caspase activity. The caspase inhibitor, Z-VAD completely abolished caspase activity in unstimulated cells, and in anisomycin-treated cells it decreased caspase activity to 0.06-fold of anisomycin alone.

Activation of caspase was also measured by Western Blotting using a cleaved caspase-3 antibody. As shown in Figure 10(a), a 17kDa band corresponding to cleaved caspase-3 was observed for the lysates that had been treated with anisomycin. The intensity of the bands in the lanes representing control and GW501516 alone was too low to be analyzed in the majority of the experiments; therefore, densitometric analysis was not performed for those samples. The levels of active caspase-3 (see Figure 10(b)) increased to 1.48 and 1.18 of anisomycin alone with both cotreatments
(10^-6 and 10^-8 M GW501516, resp.) but the difference was not significant.

3.7. The Effect of GW501516 on Apoptosis. In addition to looking at the effect of GW501516 on caspase activity, TUNEL assays were performed to further evaluate the apoptotic response. As shown in Figure 11, the proportion of TUNEL-positive cells increased while the total number of cells decreases with the addition of 250 ng/mL anisomycin when compared to control or the PPARδ agonist (at both concentrations). Pre/cotreatment of 10^-6 M or 10^-8 M GW501516 with anisomycin was comparable to anisomycin alone.

4. Discussion

4.1. PGE2 and PPARδ. It has been shown previously that prostaglandins, including PGE2, and PPARs influence another one’s activity [27–30]. Although our studies showed that the PGE2 receptors EP1, EP3, and EP4 are present in the IMCD-K2 cells and PGE2 produces an increase in cAMP production, consistent with the expression of EP4, PGE2 had no effect on PPARδ mRNA expression. However, treatment with PGE2 for eight hours did elicit an increase in PPARδ protein levels. There are two likely explanations for this. The first reason is that the PPARδ protein could be stabilized. A previous study in a prostate cancer cell line has shown that PGE2 can stabilize hypoxia-inducible factor 1 alpha (HIF-1α) protein levels, without affecting mRNA levels [31]. This would most likely be due to PGE2 preventing proteolysis of PPARδ. Interestingly, it has been revealed that unlike most nuclear receptors, degradation of PPARδ does not occur upon ligand binding but is in fact inhibited by it [32]. This would indicate that PGE2 could be causing an increase in PPARδ activation. The second possible role of PGE2 is that it could be enhancing the translation of basal PPARδ mRNA levels or stabilizing it. The ability of PGE2 to stabilize mRNA (interleukin 8) has been previously described in a paper by Yu and Chadee [33]. Therefore, from the results reported here and the literature, the effect of PGE2 on PPARδ is most likely a posttranscriptional event. We also observed an increase in PPARδ protein in response to 24 hours indomethacin. It is not clear at this time whether this effect is due to a direct increase in expression, but it has previously been shown that PPAR expression is altered by NSAIDs [34].

4.2. PPARδ Agonist and Cell Proliferation/Growth in IMCD-K2 Cells. It has been previously reported that activation of PPARδ and the use of PPARδ ligands promote cell survival and proliferation. One recent study, using both PPARδ short interfering RNA (siRNA) and a PPAR antagonist, showed that downregulation or blocking the activation of PPARδ, respectively, inhibited the PGE2-induced proliferation of mouse embryonic stem cells [14]. Another study revealed that a PPARδ agonist can stimulate propagation in both human and mouse aortic endothelial cells [35]. However, the effects of PPARδ on renal cell growth responses have not been characterized. Here we show that a highly specific PPARδ ligand, GW501516, decreased DNA and protein synthesis in IMCD-K2 cells, but only at high concentrations. Although most studies indicate that PPARδ activation results in stimulation of cell growth, mostly in human and mouse colon cancer cells [7, 36], there have been previous findings showing that PPARδ agonists, including GW501516, can hinder or have no effect on cell growth in certain human breast cancer cells and melanomas [37, 38]. These differences may be due to species differences or diversity among cell lines.

4.3. PPARδ Agonist and Anisomycin-Induced Apoptosis in IMCD-K2 Cells. Another finding in our study is that treatment with GW501516 does not reverse the reduction in cell survival due to exposure to anisomycin. These results are inconsistent with the literature that demonstrates that activation of PPARδ, or the use of a PPARδ agonist, protects many different cell types, including renal cells [12]. It is possible that the agonist may not actually be activating PPARδ in IMCD-K2 cells, as its efficacy in this cell line has never been tested. In addition, PPARδ activation may not be affecting cell survival if the death response is mostly through necrosis because studies only show a survival role for PPARδ with regard to apoptosis. To evaluate if the agonist is having an effect on PPARδ activation, experiments can be performed with cells transfected with a PPRE-driven reporter plasmid [39]. To completely assess death pathways activated by anisomycin, flow cytometry using the annexin V-FITC and propidium iodide stains should be used [40].

Even though the use of the agonist, in our experiments, elicits no response in overall cell survival in IMCD-K2 cells, the effect on apoptosis had yet to be investigated. When the CD is in a disease state due to urinary tract obstruction, ischemia-reperfusion injury or other insults apoptosis levels tend to increase [41, 42]. Thus, we wished to examine if PPARδ could protect the CD from apoptosis. Using a variety of experiments assessing apoptosis (caspase activity assay, Western blotting, TUNEL), we found no significant change in the levels of apoptosis in cells treated with GW501516 compared to those stimulated with just anisomycin. The variation between our data and the majority of the literature could be due to differences in cell types and the condition of the cells, as most studies have focussed on colorectal cancer tissues [7, 8, 43, 44]. It seems unlikely that the discrepancy in PPARδ effects is due to species differences as most of the cancer studies were done using mouse tissues. However, a few studies clearly indicate that PPARδ may not provide any prevention in certain types of apoptosis, such as in the López et al. [45] study where overexpression of PPARδ did not protect against aspirin-induced apoptosis in Jurkat cells. Thus, it is possible that PPARδ may only protect cells from apoptosis in certain conditions and cell types. From the data we collected, GW501516 has no effect on cell survival or anisomycin-induced apoptosis. The IMCD and possibly the IMCD-K2 cells are resistant to cell death because their environment in vivo is one of hypertension [46]. This inherent resiliency may make them resistant to
different treatments, similar to that previously observed in a mouse IMCD cell line [47].

4.4. Summary. The main objective of this study was to determine whether PPARδ regulates growth and apoptosis in IMCD-K2 cells. PPARδ is highly expressed in the CD, and thus may be responsible for the protection of the CD. We showed that growth responses, including DNA and protein synthesis, in these cells are reduced by PPARδ ligands, but only at high doses. GW501516 had no effect on anisomycin-induced apoptosis in the IMCD-K2 cell line.

References


