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Retinoic acid receptor γ is a therapeutically targetable driver of growth and survival in prostate cancer

Kevin Petrie1 | Zuzanna Urban-Wójciuk2 | Yordan Sbirkov3 | Amy Graham4 | Annika Hamann5 | Geoffrey Brown6

Abstract

Background: Prostate cancer (PC) tissue contains all-trans retinoic acid (ATRA) at a very low level ($10^{-9}$ M), at least an order of magnitude lower than in adjacent normal healthy prostate cells or benign prostate hyperplasia. When this is coupled with deregulated expression of the intracellular lipid-binding proteins FABP5 and CRABP2 that is frequently found in PC, this is likely to result in the preferential delivery of ATRA to oncogenic PPARβ/δ rather than retinoic acid receptors (RARs). There are three isotypes of RARs (RARα, RARβ, and RARγ) and recent studies have revealed discrete physiological roles. For example, RARα and RARγ promote differentiation and self-renewal, respectively, which are critical for proper hematopoiesis.

Aims: We have previously shown that ATRA stimulates transactivation of RARγ at sub-nanomolar concentrations (EC50 0.24 nM), whereas an 80-fold higher concentration was required for RARα-mediated transactivation (EC50 19.3 nM). Additionally, we have shown that RAR pan-antagonists inhibit the growth of PC cells (at 16-34 nM). These findings, together with the low level of ATRA in PC, led us to hypothesize that RARγ plays a role in PC pathogenesis and that RARγ-selective antagonism may be an effective treatment.

Methods and results: We found that concentrations of $10^{-9}$ M and below of ATRA promoted survival/proliferation and opposed adipogenic differentiation of human PC cell lines by a mechanism that involves RARγ. We also found that a RARγ-selective antagonist (AGN205728) potently induced mitochondria-dependent, but caspase-independent, cell death in PC cell lines. Furthermore, AGN205728 demonstrated synergism in killing PC cells in combination with cytotoxic chemotherapeutic agents.

Conclusion: We suggest that the use of RARγ-selective antagonists may be effective in PC (and potentially other cancers), either as a single agent or in combination with cytotoxic chemotherapy.

Keywords
prostate cancer, RARγ antagonists, retinoids
1 | INTRODUCTION

Prostate cancer (PC) is one of the most commonly diagnosed non-cutaneous malignancies in elderly males living in the industrialized countries of Western Europe and North America and current estimates suggest that approximately 1 in 10 men over the age 65 years will go on to develop an aggressive metastatic form of the disease.\(^1\) Thus due to an increasingly aging population, the incidence of PC is predicted to greatly increase over the next 30 years.\(^2\) Currently, chemo- or radiotherapy, often coupled with androgen deprivation therapy (ADT), is the gold-standard treatment for most PC patients.\(^3,4\) However, a highly aggressive metastatic hormone-independent form of the disease (castration or hormone resistant prostate cancer (CRPC)) develops in a significant number of patients and the treatment options for these patients remain limited. Even with the recent introduction of improved therapeutics, the average survival time of CRPC tumors and approaches that do not incorporate pan-agonism of RARs, ideally utilizing isotype-specific agonists and/or antagonists may prove useful. Consistent with this notion, we have previously shown that, in contrast to ATRA, pan-RAR antagonists induced growth arrest and apoptosis in PC cell lines and primary patient samples, while normal prostate epithelial cells displayed reduced sensitivity.\(^22,23\) In this study, we therefore sought to investigate how ATRA concentration affects the growth and survival characteristics of PC cell lines and whether this would afford the opportunity to develop new therapeutic approaches.

All-trans retinoic acid (ATRA) plays crucial roles in early development, organogenesis and tissue homeostasis by regulating the balance between cell proliferation, differentiation and apoptosis\(^5\). ATRA serves as an activating ligand for a host of obligate dimers between one of the retinoid acid receptor (RAR\(\alpha\), RAR\(\beta\), RAR\(\gamma\)), and one of the retinoid X receptor (RXR\(\alpha\), RXR\(\beta\), RXR\(\gamma\)) family members. In the absence of ligand, gene expression is repressed, whereas ATRA binding leads to coregulator exchange and gene activation.\(^6\) Individual RAR isotypes have different patterns of temporal and spatial expression during prostate development, suggesting that they play discrete roles during prostate development.\(^7\) ATRA plays a role in the development of early prostate precursor cells derived from the murine fetal urogenital sinus,\(^8\) with signaling by RAR\(\alpha\) or RAR\(\gamma\)-implicated in the promotion of differentiation vs survival/proliferation, respectively.\(^9\) In adults, an adequate supply of ATRA, possibly acting via RXR\(\gamma\), is required for the proliferation and survival of prostate stem cells,\(^9\) as well as the maintenance of proper prostate functionality.\(^7,8,10,11\) Moreover, both ATRA-deprivation and genetic ablation of RAR\(\gamma\) expression block development of the murine prostate, providing further evidence of an important role for RAR\(\gamma\).\(^10\) PC tissue has been observed to contain ATRA at very low levels (in the region of \(10^{-9}\) M), which is at least an order of magnitude lower than in adjacent normal healthy prostate cells or benign prostate hyperplasia.\(^12\) Any effects arising from low concentrations of ATRA are predicted to be compounded by over- and under-expression, respectively, of the intracellular lipid-binding proteins FABP5 and CRABP2, which occurs frequently in PC.\(^13-15\) In addition to functioning as a pan-RAR agonist, ATRA also activates the "orphan" nuclear receptor PPAR\(\gamma\), which promotes proliferation and survival. Whether ATRA is diverted to RARs or oncogenic PPAR\(\gamma\) is dependent on the FABP5:CRABP2 ratio, with FABP5 delivering ATRA to PPAR\(\gamma\) and CRABP2 delivering ATRA to RARs. It is therefore likely that this is likely to divert available ATRA to oncogenic PPAR\(\gamma\) at the expense of retinoic acid receptors (RARs). It is therefore predicted that PC cells become adapted to survive and proliferate in a low-concentration ATRA environment, with ATRA diverted to oncogenic PPAR\(\gamma\) at the expense of retinoic acid receptors (RARs).

Pharmacological concentrations (\(10^{-6}\) M) of ATRA have demonstrated clinical efficacy in a sub-type of acute myeloid leukemia (AML) referred to as acute promyelocytic leukemia (APL), that is generally associated with the t(15;17)(q24;q21) reciprocal translocation between the PML and RARA genes.\(^16\) This is primarily due to the fact that APL is associated with few cytogenetic defects apart from t (15;17) and the success of ATRA (in combination with arsenic trioxide) in this case is due to its direct targeting of the PML-RAR\(\alpha\) fusion oncoprotein.\(^14,17\) This idea is supported by the failure of ATRA to yield positive clinical outcomes in other cancers.\(^18-21\) This may also be due, at least in part, to the fact that ATRA elicits pleiotropic effects—most likely as a result of multiple RAR isotype-specific actions.\(^22,23\) Thus, ATRA may be of limited therapeutic usefulness in complex tumors and approaches that do not incorporate pan-agonism of RARs, ideally utilizing isotype-specific agonists and/or antagonists may prove useful. Consistent with this notion, we have previously shown that, in contrast to ATRA, pan-RAR antagonists induced growth arrest and apoptosis in PC cell lines and primary patient samples, while normal prostate epithelial cells displayed reduced sensitivity.\(^22,23\) In this study, we therefore sought to investigate how ATRA concentration affects the growth and survival characteristics of PC cell lines and whether this would afford the opportunity to develop new therapeutic approaches.

2 | METHODS

2.1 | Compounds

The synthetic retinoids AGN195183, AGN205327, AGN194310, AGN196996, and AGN205728 (Figure 1 and Table 1) were synthesized at the Shanghai Institute of Materia Medica. Their synthesis, development, and specificities have been described previously in Hughes et al and references within.\(^26\) All-trans-retinoic acid (Cat. # 0695); the pan-RAR agonist TTNPB (4-[[E]-2-(5,6,7,8-Tetrahydro-5,5,8,8-tetramethyl-2-naphthalenyl)-1-propenyl]benzoic acid, Cat. # 2020); the RARA- and CRABP2 delivering ATRA to RARs. It is therefore likely that this is likely to divert available ATRA to oncogenic PPAR\(\gamma\) at the expense of retinoic acid receptors (RARs).

However, a highly aggressive metastatic hormone-independent form of the disease (castration or hormone resistant prostate cancer (CRPC)) develops in a significant number of patients and the treatment options for these patients remain limited. Even with the recent introduction of improved therapeutics, the average survival time of CRPC patients is still often less than 36 months,\(^3,4\) necessitating the development of new and better treatment approaches.

All-trans retinoic acid (ATRA) plays crucial roles in early development, organogenesis and tissue homeostasis by regulating the balance between cell proliferation, differentiation and apoptosis. ATRA serves as an activating ligand for a host of obligate dimers between one of the retinoid acid receptor (RAR\(\alpha\), RAR\(\beta\), RAR\(\gamma\)), and one of the retinoid X receptor (RXR\(\alpha\), RXR\(\beta\), RXR\(\gamma\)) family members. In the absence of ligand, gene expression is repressed, whereas ATRA binding leads to coregulator exchange and gene activation. Individual RAR isotypes have different patterns of temporal and spatial expression during prostate development, suggesting that they play discrete roles during prostate development. ATRA plays a role in the development of early prostate precursor cells derived from the murine fetal urogenital sinus, with signaling by RAR\(\alpha\) or RAR\(\gamma\)-implicated in the promotion of differentiation vs survival/proliferation, respectively. In adults, an adequate supply of ATRA, possibly acting via RXR\(\gamma\), is required for the proliferation and survival of prostate stem cells, as well as the maintenance of proper prostate functionality. Moreover, both ATRA-deprivation and genetic ablation of RAR\(\gamma\) expression block development of the murine prostate, providing further evidence of an important role for RAR\(\gamma\). PC tissue has been observed to contain ATRA at very low levels (in the region of \(10^{-9}\) M), which is at least an order of magnitude lower than in adjacent normal healthy prostate cells or benign prostate hyperplasia. Any effects arising from low concentrations of ATRA are predicted to be compounded by over- and under-expression, respectively, of the intracellular lipid-binding proteins FABP5 and CRABP2, which occurs frequently in PC. In addition to functioning as a pan-RAR agonist, ATRA also activates the "orphan" nuclear receptor PPAR\(\gamma\), which promotes proliferation and survival. Whether ATRA is diverted to RARs or oncogenic PPAR\(\gamma\) is dependent on the FABP5:CRABP2 ratio, with FABP5 delivering ATRA to PPAR\(\gamma\) and CRABP2 delivering ATRA to RARs. It is therefore likely that this is likely to divert available ATRA to oncogenic PPAR\(\gamma\) at the expense of retinoic acid receptors (RARs). It is therefore predicted that PC cells become adapted to survive and proliferate in a low-concentration ATRA environment, with ATRA diverted to oncogenic PPAR\(\gamma\) at the expense of retinoic acid receptors (RARs).

2.2 | Cell lines and cell culture

RWPE-1 (ATCC# CRL-11609, RRID:CVCL_3791); LNCaP (ATCC# CRL-1740, RRID:CVCL_1379); DU 145 (ATCC HTB-81, RRID:
CVCL_0105) and PC-3 (ATCC# CRL-1435, RRID:CVCL_0035) cells were purchased specifically for this study from the American Type Culture Collection (ATCC, Manassas, VA). The LNCaP,31 DU 14532 and PC-333 cell lines were derived from metastatic prostate carcinoma. The poorly invasive, androgen-dependent LNCaP cell line expresses RARA and RARG mRNA, and RARα and RARγ, but not RARβ protein.34,35 There are conflicting data regarding expression of RARB mRNA in LNCaP cells.34-37 DU145 and PC3 cells exhibit androgen-independent growth and express RARA and RARG, but RARG mRNA, and also RARα and RARγ, but not RARβ.34,35 Human prostate epithelial non-neoplastic RWPE-1 cells38 express RARα and RARγ, but not RARβ.37 Cell cultures were maintained under serum-free conditions using the Prostate Epithelial Cell Growth Medium (PrEGM) BulletKit (Cat. # CC-3166, Lonza) minus retinoic acid in the presence of antibiotics (Lonza). For experiments, cells were seeded in RPMI-1640 supplemented with x1 ITS+1 liquid media (containing insulin, transferrin, selenium dioxide, linoleic acid, and bovine serum albumin, Merck) in the presence of antibiotics (100 U mL⁻¹ penicillin and 100 μg mL⁻¹ streptomycin Lonza).

2.3 | Analysis of cell growth

Cell proliferation was determined either by direct cell count or indirectly using the ViaLight Plus Cell Proliferation and Cytotoxicity BioAssay Kit (Lonza) using a Berthold LB953 luminometer as previously...
Adipocyte differentiation was induced in serum-free adapted LNCaP, described. Briefly, cells were seeded (400 cells per well) in RPMI-1640 + ITS\(^+\) culture medium in 96-well microtitre plates and treated with retinoids both immediately and at day 2 by replacing the medium. The number of viable cells was assessed at day 5 by measuring cellular ATP levels.

2.4 | Analysis of colony forming potential

Single cell suspensions of serum-free adapted PC cells were prepared by trypsinization of 80% confluent plates using 0.25% (v/v) trypsin in phosphate-buffered saline for 5 minutes at 37°C. Single cell suspensions containing <90% viable cells as measured by trypan blue exclusion were discarded. Cells were seeded at a density of 1000 cells per 65-mm diameter plastic cell culture dish. Cells were incubated for 21 days at 37°C in 5% CO\(_2\) in 4 mL of RPMI 1640 plus ITS\(^+\). Culture medium was replaced every 5 days, and after 21 days cells were fixed using 1% (v/v) formaldehyde in 0.9% (v/v) NaCl and stained with 1% (v/v) methylene blue in 0.9% (v/v) NaCl. Colony-forming efficiency was determined by counting the number of colonies that formed: (number of colonies formed/number of cells inoculated) \(\times\) 100. Determination of clone morphology (holoclone, meroclone, or paraclone) was performed as previously described.

2.5 | Analysis of transactivation

Transactivation assays using CV-1 kidney fibroblast cells have been described previously. Briefly, CV-1 kidney fibroblast cells were first transiently transfected with a mammalian plasmid vector (ERE-Tk-Luc) that expresses Firefly luciferase under the transcriptional regulation of the Xenopus vitellogenin estrogen response element (ERE), together with the pGL4.70[hRLuc] Renilla luciferase control vector (Promega). Cells were also co-transfected with expression vectors encoding a fusion protein comprising the estrogen receptor DNA-binding domain and the ligand-binding domains of either RAR\(\alpha\), RAR\(\gamma\), or RAR\(\gamma\), expressing a fusion protein comprising the estrogen receptor DNA-binding domain and the ligand-binding domains of either RAR\(\alpha\), RAR\(\gamma\), or RAR\(\gamma\). Cells were transfected using Lipofectamine (Thermo) according to the manufacturer’s instructions. Twenty-four hours post-transfection, cells were treated with increasing concentrations of ATRA (10\(^{-10}\)–10\(^{-6}\)) for 16 hours in culture medium-containing 2.5% charcoal-stripped FBS (Merck). Transactivation was measured using the Dual-Luciferase Reporter Assay System (Promega). Firefly luciferase reporter activity was normalized against Renilla luciferase activity, and transactivation data are presented as the percentage of the maximal response produced by 1 \(\mu\)M ATRA.

2.6 | Analysis of adipogenic differentiation

Adipocyte differentiation was induced in serum-free adapted LNCaP, PC-3 or DU145 as described by Zhou et al. Briefly, cells were plated at a density of 3 \(\times\) 10\(^5\) cells/dish in 60 mm cell culture dishes in RPMI 1640 plus ITS\(^+\). In order to induce adipogenic differentiation, this growth medium was supplemented with dexamethasone (10\(^{-7}\) M), indomethacin (5 \(\times\) 10\(^{-5}\) M), and the phosphodiesterase inhibitor 3-isobutyl-L-methyl-xanthine (IBMX, 5 \(\times\) 10\(^{-5}\) M). Adipogenic differentiation medium was refreshed every 3 days and adipogenesis was assessed by Oil red O staining on day 14 and day 21. Cells were washed with ice cold PBS and fixed in 5% formaldehyde (v/v in PBS) for 1 hour. Cells were washed and incubated for 15 minutes with 1 mL of Oil red O (0.6% [w/v] in 60% isopropanol/ PBS 1:1 v/v). After washing, cell-bound Oil red O was extracted in 1 mL isopropanol and absorbance was measured at 510 nm. Values were normalized using the ViaLight Plus Cell Proliferation and Cytotoxicity BioAssay Kit (Lonza).

2.7 | Antibody transfection

Serum-free grown LNCaP cells were transfected with antibodies directed against either RAR\(\alpha\) or RAR\(\gamma\) (Chariot, Active Motif) according to the manufacturer’s recommendations. In brief, 50 \(\mu\)g of antibody and 1 \(\mu\)g of β-galactosidase were resuspended in 100 \(\mu\)L x2 sterile phosphate-buffered saline (PBS) and mixed with freshly sonicated Chariot transfection reagent (6 \(\mu\)L of reagent re-constituted in 100 \(\mu\)L sterile H\(_2\)O). This mixture was incubated at room temperature for 30 minutes to form the transfection complex reagent. LNCaP cells (5 \(\times\) 10\(^5\)) were washed twice with pre-warmed serum-free RPMI 1640 and the final cell pellet was mixed with the antibody/Chariot mixture and incubated at 37°C in a humidified atmosphere containing 5% CO\(_2\). After 30 minutes a further 400 \(\mu\)L of serum-free RPMI 1640 were added to the cells to achieve the final transfection concentration. The LNCaP cells were incubated at 37°C for a further 5 hours to allow internalization of the antibody. Control cells were treated with the Chariot reagent alone (6 \(\mu\)g in 200 \(\mu\)L of sterile PBS). After antibody transfection, the cells were washed twice in RPMI 1640 + ITS\(^+\), and re-suspended in a volume of 1 mL. A 100-\(\mu\)L aliquot of cells was removed, fixed and stained for β-galactosidase expression to assess transfection efficiency. Cells were grown for a further 12 days at which time colonies were fixed with 2% buffered formaldehyde, washed and stained with 1.25% (w/v) crystal violet stain. Polyclonal antiserum to RAR\(\alpha\) was a generous gift from Dr P. Chambon (Institute for Genetics and Cellular and Molecular Biology, Strasbourg, France), and the antibody to RAR\(\gamma\) (Cat. # sc-550) was purchased from Santa Cruz Biotechnology Inc.

2.8 | Cell cycle analysis

Cell cycle status was measured by staining harvested cells with propidium iodide (PI, Molecular Probes) according to the manufacturer’s instructions. The distribution of cells between phases of the cell cycle was determined using a FACS Calibur flow cytometer and the CellFIT cell-cycle analysis software (Becton-Dickinson). The cell
cycle distribution is shown as the percentage of cells containing 2n (G1 phase), 4n (G2 and M phases) and 4n > 3 > 2n DNA amount (S phase) judged by propidium iodide staining. The apoptotic population is defined by the percentage of cells with DNA content lower than 2n (sub-G1 phase).

2.9 | Analysis of apoptosis

Apoptosis-associated DNA fragmentation was measured using the TACS 2 TdT Fluorescein in situ apoptosis detection kit (Trevigen). Briefly, serum-free adapted PC cells were seeded at a density of 5 × 10⁵ cells per 75 cm² flask and treated with retinoids of vehicle control. Treated cells (both adherent cells and those in suspension) were trypsinized, cytopun onto microscope slides and analyzed according to the manufacturer’s instructions. Induction of apoptosis was assessed using fluorescence microscopy and quantified by counting five separate fields of view (minimum of 500 cells).

Activation of caspase was measured essentially as described by Keedwell et al.²⁵ The pan-caspase inhibitor Z-VAD-FMK (R&D Systems) was used to investigate the involvement of caspases in AGN205728-induced apoptosis. A 20 mM stock solution was prepared in DMSO. Z-VAD-FMK was added at 50 μM to bulk cultures of cells 1 hour before adding AGN205728, and cells were monitored using the TUNEL assay. Fluorometric assays for caspase activity were conducted in 96-well microtiter plates. In total, 50 μL of assay buffer (20 mM HEPES, pH 7.5, 10% glycerol, 2 mM dithiothreitol) containing 50 μM of peptide substrates for caspase-3 (DEVD-AFC), caspase-8 (IETD-AFC), or caspase-9 (LEHD-AFC) (all from Biovision Inc.) were added to each well. In total, 50 μL of cell lysate were added to initiate the reactions. Backgrounds were measured in wells that contained assay buffer, substrate, and lysis buffer. Fluorescence was measured on a CytoFluor 4000 fluorescence plate reader (Applied Biosystems) set at 400 nm excitation and 508 nm emission. Backgrounds were measured in wells that contained assay buffer, substrate, and lysis buffer. Fluorescence was measured on a CytoFluor 4000 fluorescence plate reader (Applied Biosystems) set at 400 nm excitation and 508 nm emission. Caspase activities were calculated as fold increases relative to control wells.

2.10 | Software, data manipulation and statistical analyses

At least three experiments of each type were performed with triplicate replicates. Results are expressed as means ± SD (± SD). Statistical significance between groups of data was analyzed by unpaired t-test or 2-way ANOVA as appropriate using GraphPad Prism (GraphPad Software, La Jolla, CA) or SigmaStat (Systat Software, San Jose, CA). Statistical significance is represented as follows: ns (not significant), * P ≤ .05; ** P < .01; *** P < .001; **** P ≤ .0001. Dose-response curves for compounds were plotted using the kinetics module of SigmaPlot (Systat Software, San Jose, CA). Marvin was used for drawing and displaying chemical structures of ATRA and synthetic retinoids (Marvin 19.12, 2019, ChemAxon).

3 | RESULTS

3.1.1. Low concentrations of all-trans retinoic acid stimulate proliferation and colony forming ability of prostate cancer cells

In light of our hypothesis that PC cells may be adapted to survive and proliferate in a low ATRA concentration environment, we initially sought to titrate ATRA against three model PC cell lines derived from metastatic prostate carcinoma: LNCaP, PC-3, and DU145. We found that in all three PC cell lines, ATRA treatment for 5 days within the range of 10⁻¹¹⁻¹⁻⁰ M resulted in an increase in cell numbers of up to 1.5-fold (Figure 2A). By contrast, treatment ATRA at concentrations of 10⁻⁹ M and above progressively diminished cell growth. In order to assess the long-term effects of low vs high concentrations of ATRA on cell growth and survival, we performed single-cell clonogenic assays on LNCaP, PC-3 and DU145 cells and RWPE-1 cells, which are derived from HPV-18-transformed epithelial cells from the peripheral zone of normal prostate. Consistent with our 5-day ATRA treatment results, we found that for all three PC cell lines treatment with 10⁻¹⁰ M ATRA increased the colony forming efficiency (CFE) while treatment with 10⁻⁶ M ATRA decreased CFE (Figure 2B). By contrast, non-tumorigenic RWPE-1 cells, which untreated had a low CFE of 0.43, failed to respond to either high or low concentrations of ATRA. These results indicated that treatment of malignant PC cell lines with low nanomolar concentrations of ATRA favors survival/proliferation of the clonogenic fraction, whereas higher pharmacological concentrations promote growth arrest and/or differentiation.

When subjected to single cell clonogenic assays, malignant PC cells produce colonies of three different types: holoclone (comprising large cells with homogeneous morphology), meroclone (comprising cells with intermediate size and heterogeneous morphology), and paracline (comprising smaller cells with homogeneous morphology). These morphologically distinct types of colonies are considered to contain cells that possess stem cell-like (holoclone), early progenitor (meroclone), or late progenitor/differentiation-committed (paracline) characteristics. In the absence of ATRA, around 70% of colonies formed by serum-free adapted LNCaP displayed typical large holoclone/meroclone morphologies with the remaining 30% containing cells with the smaller, differentiated paracline morphology (Figure 2C). Treatment with 10⁻¹⁰ M ATRA increased the proportion of colonies displaying morphologies consistent with stem cell-like or early progenitor status to around 85%. Treatment with 10⁻⁶ M ATRA, on the other hand, increased the proportion of colonies displaying late progenitor/differentiation-committed (paracline) characteristics to more than 60%. Taken together, these results suggested that the low physiological concentrations of ATRA found in PC serve to either increase the clonogenic (stem cell) population by providing a stem-cell specific survival stimulus or block entry of this cell population into a spontaneous differentiation/apoptosis programme.
3.1.2. | Agonism of RARγ promotes growth and inhibits adipogenic differentiation in prostate cancer cells

ATRA is the natural endogenous ligand for all three RAR isotypes and they display similar binding affinities for this retinoid (RARα, $0.37 \times 10^{-9}$ M; RARβ, $0.37 \times 10^{-9}$ M; RARγ, $0.22 \times 10^{-9}$ M). However, we and others have previously shown that while ATRA stimulates transactivation in CV-1 cells via RARβ and RARγ at sub-nanomolar concentrations, much higher concentrations are required for RARα-mediated transactivation. We also observed this phenomenon in LNCaP prostate cancer cells (that do not express RARβ), where RARγ displayed a half-maximal response (EC50) value of $0.24 \times 10^{-9}$ M while RARα displayed an EC50 value of $19 \times 10^{-9}$ M, which was 79-fold higher. This prompted us to investigate whether individual RAR isotypes played a role in the differential response of PC cells to low vs high concentrations of ATRA. Consistent with our prior data, we found that RARβ and RARγ displayed EC50 values of $0.39 \times 10^{-9}$ M and $0.36 \times 10^{-9}$ M, respectively, while ATRA had a minimal effect on transactivation via RARα at concentrations below $10^{-9}$ M (EC50 = $12.4 \times 10^{-9}$ M) (Figure 3A-C). This represented a 30-fold difference and maximal transactivation by RARα did not occur until $10^{-6}$ M. We then examined the growth response of LNCaP cells of RAR isotype-selective agonists in comparison to the pan-RAR agonist ATRA (Figure 3D). In line with results suggesting opposing roles for RARα and RARγ in prostate growth and differentiation (and potentially in PC), the RARγ-selective agonist AGN205327 elicited an effect on LNCaP cells that mirrored ATRA while the RARα-selective agonists AM580 and AGN195183 failed to promote cell growth at low concentrations. It is also important to note that the modest inhibition in the growth of LNCaP cells and concentrations greater than $10^{-6}$ M can be explained by a loss of receptor specificity and subsequent agonism of RARα at high concentrations (Table 1). Further supporting the notion that, in contrast to RARγ, RARα has anti-growth and pro-differentiative role PC, co-treatment of AGN195183 with an RARα-selective antagonist (AGN196996) abolished AGN195183-mediated growth inhibition in LNCaP cells (Figure 3E). Moreover, co-treatment of ATRA with AGN196996 promoted proliferation of LNCaP cells, likely due to activity of ATRA on RARγ.

PC cell lines contain a sub-population of cells that possess characteristics similar to mesenchymal stem cells that, upon appropriate stimulation, can undergo neuroendocrine, adipogenic, or osteoblastic differentiation. When grown in a pro-differentiative culture medium containing dexamethasone, indomethacin, and the phosphodiesterase inhibitors...
inhibitor IBMX, lipid droplets began to form within cells and 10-12 days post-induction of differentiation, lipid accumulation may be quantitatively analyzed. In agreement with our results suggesting that, via either RARα or RARγ, ATRA can elicit either pro- or anti-growth and differentiation responses in PC, it has also been reported that ATRA can block the differentiation of pre-adipocyte-like cells into lipid-loaded adipocytes in several model systems. We therefore examined whether RARγ agonism could inhibit adipogenic differentiation in PC cell lines. Oil red O staining showed that treatment with AGN205327 (10^{-7} M) was able to inhibit the accumulation of lipid droplets by around 50% (Figure 3F). These results were of a similar order to those obtained with the PPARγ antagonist GW9662 (10^{-5} M). We also examined the ability of AGN205327 to diminish adipogenic differentiation arising from PPARγ agonism (Figure 3G). We found that induction of adipogenic differentiation in LNCaP cells by the PPARγ agonist ciglitazone (10^{-7} M) was inhibited by up 70% when AGN205327 (10^{-7} M) was given at time points ranging between 48 hours pre- and post-addition of ciglitazone. Addition of
AGN205327 later than 48 hours post-induction of adipogenesis failed to have any effect on cigitazone-induced adipogenesis, which is consistent with studies showing that retinoids can only block adipogenesis when added during the early commitment phase.\textsuperscript{47,50,51}

### 3.1.3. Antagonism of RARγ inhibits growth and colony formation in prostate cancer cells

Our results showing that RARγ agonism could promote growth and clonogenicity, and inhibit differentiation strongly suggested that RARγ-selective antagonism could have the opposite effects. This is in line with our previous data showing that treatment of LNCaP cells with pan-RAR antagonist AGN194310 (Figure 1, S1A, S1D and Table 1) inhibited colony formation.\textsuperscript{25} Consistent with these observations, the RARγ-selective antagonist AGN205728 (Figure 1, S1B, S1E and Table 1) diminished the CFE of all three PC cell lines in a fashion similar to AGN194310 (Figure 4A). Treatment with neither ATRA, nor the RARα-selective antagonist AGN196996 (Figure 1, S1C, S1F and Table 1) proved as effective AGN205728 or AGN194310. These data indicate that antagonism of RARγ alone is sufficient to reduce colony formation in PC cell lines. The growth inhibitory effect of treatment AGN205728 in PC cell lines was observed also in short-term assays, with concentrations of 10^{-6} M and above abolishing metabolic activity (Figure 4B). By contrast, non-tumorigenic RWPE-1 cells showed significantly diminished response to AGN205728. Consistent with this, AGN194310 and AGN205728 but not AGN196996 were able to diminish RAR-mediated transactivation stimulated by ATRA (Figure 4C).

While the EC_{50} data indicated a high degree of selectivity AGN205728 towards RARγ (Table 1), strongly supporting a role for this RAR isotype in the pathology of PC, we nevertheless wanted to demonstrate the role of RARγ by other means. To this end, we took a completely different approach to abrogating activation of RARγ and used an anti-RAR antibody to interfere with its function (Figure 4D). Transfection with this “neutralizing” antibody phenocopied treatment with AGN205728 and diminished CFE, both with and without low dose (10^{-11} M) ATRA. This result validated RARγ as a bona fide therapeutic target, at least as far as in vitro studies permit. By contrast, and consistent with results from RARα-selective antagonism, treatment with a neutralizing antibody raised against RARα failed to block increases in CFE arising from low dose ATRA treatment. These data reveal that antagonism of RARγ is sufficient to inhibit the growth PC cell lines in vitro and non-malignant prostate cells display reduced sensitivity to this treatment.

### 3.1.4. Antagonism of RARγ in LNCaP cells induces growth arrest followed by caspase-independent apoptosis

In order to understand the mechanisms underlying the inhibition of growth and clonogenic potential we observed upon abrogation of proper RARγ function, we first analyzed the effects of treatment with AGN194310 (10^{-8} M) or AGN205728 (10^{-8} M) on the cell cycle (Figure 5A). The proportion of cells in G1 phase of the cell cycle gradually increased following treatment with AGN194310 or AGN205728, which was paralleled by a fall in the proportion of cells found in S-phase. After 12 hours the proportion of cells identified as sub-G0/G1 (indicative of apoptosis) started to rise until at the 48-hour timepoint almost 60% of cells were sub-G0/G1 following AGN194310 treatment and almost 40% of cells were sub-G0/G1 following AGN205728 treatment. The accumulation propidium iodide-stained DNA in sub-G0/G1 compartment indicated that cellular DNA had begun to fragment, a biomarker of apoptotic or necrotic cell death. Time-course agarose gel electrophoresis analysis of DNA from AGN205728-treated LNCaP cells revealed a discrete laddering pattern that is typical of cells undergoing apoptosis rather than the more generalized smeared pattern that is associated with necrosis (Figure 5B). A quantitative readout of the percentage of cells undergoing apoptosis following treatment with AGN194310 (10^{-6} M) or AGN205728 (10^{-6} M) was obtained using the terminal deoxynucleotide transferase...
end-labeling (TUNEL) assay (Figure 5C). After 24 hours, the percentage of cells undergoing apoptosis gradually increased until 60 hours post-treatment the proportion of apoptotic cells reached 35% and 40% for AGN194310 and AGN205728, respectively. Again, among the untreated cells and the AGN196996-treated sample there was little evidence of induction of apoptosis. We then assessed the contribution of caspases to AGN205728-associated induction of apoptosis. Here, we found that pre-treatment with the pan-caspase inhibitor Z-VAD-FMK (5 $\times$ 10$^{-5}$ M) failed to diminish induction of apoptosis by AGN205728 (Figure 5D) and, furthermore, AGN205728 failed to activate caspases-3, -8 or -9 (Figure 5E).

Chemotherapy has been one of the mainstays in the treatment of castration-resistant PC, including the taxanes, carbazitaxel, and docetaxel.52 These agents induce caspase-dependent apoptosis (similar to RAR$\alpha$ agonism) in numerous PC cell lines, which prompted us to test docetaxel in combination with low-dose AGN205728 (10$^{-7}$ M). We found that the combination of AGN205728 and docetaxel led to a significant reduction in cell viability compared to their use as single agents (Figure 5F). Similar results were obtained with the pyrimidine analog 5-fluorouracil (5-FU) and the topoisomerase 2 inhibitor etoposide.

### DISCUSSION

Retinoic acid signaling in prostate carcinoma cells is complex and our data strongly suggest that is at least in part because RAR$\alpha$s and RAR$\gamma$s play opposing roles in the promotion of differentiation and proliferation/survival, respectively. This notion is in agreement with studies in breast carcinoma cell lines showing that RAR isoforms have opposing and, perhaps, mutually antagonistic roles in controlling proliferation.14,53-56 While RAR$\gamma$s supports the proliferation and survival of malignant breast epithelial cells RAR$\alpha$s, on the other hand, inhibits proliferation and promotes differentiation. Our results suggest that this
complexity is due, at least in part, to concentration-dependent differential responses of RAR isotypes to ATRA. At low nanomolar levels, RAR\(_\gamma\) is activated preferentially, while higher concentrations of ATRA (including pharmacological levels) activate both. These findings are in keeping with studies that have reported that ATRA exerts concentration-related biphasic effects on the proliferation of PC cell lines when these cells are grown under serum free conditions.\(^{11,35,57-63}\) Furthermore, we have shown that low levels of ATRA, ...
probably acting via RARγ, increase colony forming efficiency, promote proliferation, and survival and oppose differentiation, suggesting that stem-cells or early progenitors may be the target cell populations.

It is well-established that PC cell populations contain fractions possessing mesenchymal stem cell-like properties and can be differentiated into adipocytes, and we demonstrate that this can be blocked by activation of RARγ. The differentiation cocktail used in this study for the induction of adipogenesis in LNCaP cells included the glucocorticoid agonist dexamethasone and the phosphodiesterase inhibitor IBMX, the combination of which down-regulates expression of RARγ in pre-adipocyte models. The down-regulation of RARγ expression prior the onset of adipogenesis is an essential molecular signature of several in vitro pre-adipocyte differentiation models, while in vivo, an inverse correlation between RARγ expression and adiposity has been demonstrated. Moreover, RARγ agonism inhibits the expression of the pro-adipogenic nuclear receptor PPARγ and thereby blocks the adipogenic signals emanating from this receptor. Our observation that the addition of an RARγ agonist prior to adipogenesis induction blocks PPARγ-stimulated adipogenesis of LNCaP cells is consistent with these findings.

The deregulation of retinoic acid signaling in malignant cells also has implications for the use of retinoids as chemotherapeutic agents. It has long been known that pharmacological concentrations of ATRA (10^{-7} M), acting via RARα, can drive malignant prostate cells into growth arrest and apoptosis both in vitro and in vivo. However, high dose retinoid treatment for PC, as well as most other solid tumors, has had limited success in the clinic. Several issues contribute to this failure, including acquisition of retinoid-resistance due to promoter hypermethylation-mediated loss of RARγ expression in patients with advanced PC. RARγ is known to play a role in driving growth arrest in malignant prostate cells at physiological ATRA concentrations, so its absence may provide a growth/survival advantage to the malignant cells over their normal counterparts. Second, the altered biology of cancer cells may negatively impact uptake, metabolism, storage of retinoids, thus impeding the ability to achieve a high enough local concentrations of ATRA to activate RARα. Indeed, pharmacodynamic studies have shown that the plasma concentrations of ATRA (in the region of 2.2-0.4 μM) that have to be achieved before marginal clinical benefits are observed are often associated with therapy-limiting toxicities.

The pan-RAR agonistic activities of ATRA may well preferentially activate RARγ over RARα in the tumor microenvironment and our results indicate that antagonism RARγ may represent a better therapeutic strategy. This notion is in agreement with our previous findings that pan-RAR antagonists can induce caspase-independent apoptosis more effectively that ATRA in PC cell lines. Our results also suggest that RARγ antagonists may be effective in combination with cytotoxic chemotherapy. We showed that a sub-maximally effective concentration of AGN205728 (10^{-7} M) significantly increased the degree of cell death induced in serum free adapted-LNCaP cells produced by sub-maximal concentrations of the taxane docetaxel, as well as sub-optimal concentrations of the commonly used chemotherapeutic agents 5-fluorouracil and etoposide. Docetaxel is used for the treatment of castration-resistant PC but this is limited due to high levels of toxicity (particularly neutropenia) and either innate or acquired resistance in a significant percentage of patients. If the sensitivity of PC towards taxanes and other therapeutic agents could be increased, both the incidence of toxicity and the acquisition of resistance may be reduced. Finally, while we performed experiments in defined media in order to prevent ATRA and, potentially, androgens or other factors found in fetal bovine serum from interfering with our data, the androgen-deprived environment created mimics the clinical scenario. Recent research indicates that androgen deprivation induces a neuroendocrine phenotype in LNCaP cells that is followed by the acquisition of a "stem cell-like" state characterized by expression of markers including markers CD133, ALDH1A1, and ABCB1A. Many PC patients who are not cured with surgery or radiation and subsequently undergo androgen deprivation therapy (ADT) develop resistance to ADT and relapse with castration resistant prostate cancer (CRPC) that is associated with very poor outcome. Our data indicate RARγ antagonism to be an effective approach in this context and thus merits further investigation.

In conclusion, we have demonstrated that lower than normal concentrations of ATRA oppose differentiation, promote the survival and proliferation of several PC cell lines by activating RARγ. We also showed that RARγ antagonists alone can block proliferation of both androgen-dependent and independent PC cell lines and eventual induce caspase-independent apoptosis. Hence, RARγ antagonism alone, or in combination with cytotoxic chemotherapy, may represent a novel approach to the treatment of advanced prostate that targets both androgen-dependent and androgen-independent PC cells.

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CONFLICT OF INTEREST
The authors have stated explicitly that there are no conflicts of interest in connection with this article.

AUTHOR CONTRIBUTIONS
Kevin Petrie: Formal analysis; investigation; methodology; supervision; writing-original draft; writing-review and editing. Zuzanna Urban-Woźniuk: Formal analysis; investigation; methodology; writing-original draft; writing-review and editing. Yordan Sbirkov: Formal analysis; investigation; methodology; writing-original draft; writing-review and editing. Amy Graham: Formal analysis; investigation; methodology; writing-original draft; writing-review and editing. Annika Hamann: Formal analysis; investigation; writing-original draft; writing-review and editing. Geoffrey Brown: Conceptualization; formal analysis; investigation; methodology; project administration; supervision; writing-original draft; writing-review and editing.

ETHICAL STATEMENT
Not Applicable.
DATA AVAILABILITY STATEMENT
The data that support the findings of this study are available from the corresponding author upon reasonable request.

ORCID
Kevin Petrie https://orcid.org/0000-0002-9805-9152

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SUPPORTING INFORMATION
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