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Keywords: prostate cancer; phospholipase D1; patient tissue; expression; inhibition; activity; cellular localisation

Phospholipase D inhibitors reduce human prostate cancer cell proliferation and colony formation

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Background: Phospholipases D1 and D2 (PLD1/2) hydrolyse cell membrane glycerophospholipids to generate phosphatidic acid, a signalling lipid, which regulates cell growth and cancer progression through effects on mTOR and PKB/Akt. PLD expression and/or activity is raised in breast, colorectal, gastric, kidney and thyroid carcinomas but its role in prostate cancer (PCa), the major cancer of men in the western world, is unclear.

Methods: PLD1 protein expression in cultured PNT2C2, PNT1A, P4E6, LNCaP, PC3, PC3M, VCaP, 22RV1 cell lines and patient-derived PCa cells was analysed by western blotting. PLD1 protein localisation in normal, benign prostatic hyperplasia (BPH), and castrate-resistant prostate cancer (CRPC) tissue sections and in a PCa tissue microarray (TMA) was examined by immunohistochemistry. PLD activity in PCa tissue was assayed using an Amplex Red method. The effect of PLD inhibitors on PCa cell viability was measured using MTS and colony forming assays.

Results: PLD1 protein expression was low in the luminal prostate cell lines (LNCaP, VCaP, 22RV1) compared with basal lines (PC3 and PC3M). PLD1 protein expression was elevated in BPH biopsy tissue relative to normal and PCa samples. In normal and BPH tissue, PLD1 was predominantly detected in basal cells as well in some stromal cells, rather than in luminal cells. In PCa tissue, luminal cells expressed PLD1. In a PCa TMA, the mean peroxidase intensity per DAB-stained Gleason 6 and 7 tissue section was significantly higher than in sections graded Gleason 9. In CRPC tissue, PLD1 was expressed prominently in the stromal compartment, in luminal cells in occasional glands and in an expanding population of cells that co-expressed chromogranin A and neurone-specific enolase. Levels of PLD activity in normal and PCa tissue samples were similar. A specific PLD1 inhibitor markedly reduced the survival of both prostate cell lines and patient-derived PCa cells compared with two dual PLD1/PLD2 inhibitors. Short-term exposure of PCa cells to the same specific PLD1 inhibitor significantly reduced colony formation.

Conclusions: A new specific inhibitor of PLD1, which is well tolerated in mice, reduces PCa cell survival and thus has potential as a novel therapeutic agent to reduce prostate cancer progression. Increased PLD1 expression may contribute to the hyperplasia characteristic of BPH and in the progression of castrate-resistant PCa, where an expanding population of neuroendocrine-like cells express PLD1.

Phospholipase D (PLD) hydrolyses major membrane glycerophospholipids to phosphatidic acid (PtdOH), an intermediate in complex lipid synthesis. PtdOH is also a key signalling lipid that activates proteins at membrane surfaces (Kooijman *et al*, 2003) and

regulates cell survival (Foster, 2009). Due to its inverted cone shape, PtdOH can also alter membrane curvature, facilitating the vesicle formation (Yang *et al*, 2008) needed to increase tumour cell proliferation.

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The PtdOH generated by PLD activity can regulate cell survival and motility through mTOR, a master regulator of cancer cell survival and growth, which controls protein synthesis, migration, invasion and metastasis (Foster, 2009). PtdOH formation also leads to inhibition of protein phosphatase 1 and upregulation of the ERK, PKB/Akt, NF κ B and Wnt signalling pathways, further promoting both cancer cell survival and metastasis (Foster, 2009; Winter *et al*, 2010; Kang *et al*, 2011; Yoon *et al*, 2011). Abnormal activation of Wnt/ β -catenin signalling and hyperactivation of transcriptional target genes has been linked to prostate cancer (PCa) and especially metastasis to bone in castration-resistant disease (Hall and Keller, 2006; Yokoyama *et al*, 2014).

PLD1 and PLD2 are the two main isoforms of PLD in human cells (Foster and Xu, 2003). PLD1 has a diffuse cytoplasmic localisation (Freyberg *et al*, 2001) and is activated by protein kinase C (PKC) (Kim *et al*, 2000), casein kinase-II (Ahn *et al*, 2006) and by the small GTPases ARF and RHO (Foster, 2009). In contrast, PLD2 is located at the plasma membrane and is activated by both PKC and by receptor (EGFR, PDGFR) and non-receptor (Src, JAK3) tyrosine kinases (Henkels *et al*, 2010). PLD2 has a higher basal activity than PLD1 and acts as both a phospholipase and as a guanine nucleotide exchange factor (Kang *et al*, 2014). Total PLD activity is increased in gastric cancer (Uchida *et al*, 1999) and breast cancer where levels correlate with disease grade (Uchida *et al*, 1997). PLD1 is overexpressed in human breast cancer tissue (Noh *et al*, 2000), while PLD2 expression is increased in renal (Zhao *et al*, 2000) and thyroid carcinomas (Kim *et al*, 2008). Expression of PLD2 is increased in colorectal cancer where it correlates with tumour size and prognosis (Saito *et al*, 2007). In a human breast carcinoma xenograft model, PLD2 drives cell invasion, tumour growth and metastasis (Henkels *et al*, 2013). Inhibition of either PLD1 or PLD2 also makes breast cancer cells more susceptible to radiotherapy (Cheol Son *et al*, 2013). New isoform-specific PLD inhibitors can reduce proliferation of breast cancer cells in mice and are well tolerated (Henkels *et al*, 2013). If similar overexpression of PLD is present, these new inhibitors have potential as alternative therapeutic agents to hormone-based and other current therapies used to treat PCa.

MATERIALS AND METHODS

Prostate epithelial cell lines. The prostate cell lines used with their growth media requirement, diagnosis and origin are listed in Table 1.

Patient-derived prostate epithelial and stromal cells. Primary prostate epithelial and stromal cells were cultured from human prostate tissues (Collins *et al*, 2001; Frame *et al*, 2016), which were obtained with patient consent and full ethical approval (LREC:07/H1304/121) from radical prostatectomies and transurethral resections of the prostate (TURPS). Histopathology information was provided for pre- and post-operative samples as a Gleason grade, or verifying morphology as normal. Epithelial cells were grown on collagen 1-coated 10 cm dishes in KSFM with supplements of 1% L-glutamine, bovine pituitary extract and epidermal growth factor at 37 °C with 5% CO₂. For the first passage, cells were co-cultured with irradiated (60 Gy) mouse embryonic fibroblast (STO) cells. Subsequent passages were free of STOs and all cultures were used at the lowest practical passage number after establishment in culture (p2–p5). Stromal cells were cultured in RPMI 1640 with 10% FCS and 1% glutamine.

Western blotting. Epithelial cell lysates were prepared using Cytobuster Protein Extraction Reagent (71009, EMD Millipore, Watford, UK) with protease inhibitors (cOmplete, EDTA-free Protease Inhibitor Cocktail Tablets, Roche, Burgess Hill, UK). Stromal cell lysates were prepared in a 1% TX-100 lysis buffer

containing protease inhibitors. SDS-PAGE and western blotting were as described previously (Rumsby *et al*, 2011). Primary antibodies were an anti-PLD1 rabbit polyclonal (sc25512, Santa Cruz, Insight Biotechnology, Wembley, UK, 1:750), and a rabbit anti-GAPDH polyclonal (Abcam, Cambridge, UK, ab9485, 1:20000). The secondary antibody was HRP-linked anti-rabbit IgG (7074S, Cell Signalling, New England Biolabs Ltd, UK, 1:6000). A kaleidoscope protein ladder (Bio-Rad, Watford, UK, Cat No 1610375) was used throughout.

PLD activity. PLD activity was measured with an Amplex Red PLD kit (A12219, Invitrogen, Fisher Scientific Ltd, Loughborough, UK) according to the manufacturer's instructions. Prostate tissue was chopped in the supplied buffer containing 1% peroxide-free Triton X-100 and protease inhibitors. The chopped tissue in buffer was put through three freeze/thaw cycles at –80 °C, homogenised and centrifuged at 1000 g and 4 °C for 5 min to remove cell debris. The supernatant was used for analysis. PLD activity was assayed with 50 μ g supernatant protein (measured by a bicinchoninic acid [BCA] method) in triplicate, where sufficient tissue was available. PLD from *Streptomyces chromofuscus* (P0065, Sigma Aldrich Company Ltd, Poole, UK) was used to produce a new standard curve for every set of measurements.

PLD inhibition and cell viability. The effects of PLD inhibition on the viability of prostate epithelial cell lines and patient-derived PCa cells was measured using an MTS ([3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium] assay (Promega, Southampton, UK). Wells of a 96-well plate were filled with 100 μ l of appropriate medium or medium containing DMSO vehicle or different PLD inhibitors over a 1–25 μ M concentration range. A total of 1×10^4 cells were then added. After 24 and 48 h treatment, cell viability was measured by adding 12 μ l MTS and incubating at 37 °C for 2 h prior to reading absorbance at 490 nm. The dual PLD1/PLD2 inhibitor FIPI was from Tocris, Bio-Techne Ltd, Abingdon, UK. Another dual PLD1/PLD2 inhibitor 5W0 (VU0155056) and the specific PLD1 inhibitor EVJ (VU0359595) were gifts from Prof Alex Brown, Vanderbilt University, USA (Lavieri *et al*, 2009; Lewis *et al*, 2009; Lavieri *et al*, 2010).

Colony recovery assays. Patient-derived PCa cells were seeded in collagen 1-coated 6 well plates at 2×10^5 cells per well in complete KSFM medium. The following day, cells were treated with either vehicle (DMSO) or a PLD inhibitor (17.5 μ M) for 4 h. Cultures were then rinsed, trypsinised, counted and seeded at 500 cells per well of collagen 1-coated 6 well plates with STO feeder cells (Frame *et al*, 2013). The complete KSFM medium was changed regularly and further STOs added when required. After 2–3 weeks, colonies of >32 cells (at least 5 population doublings) were scored after being visualised by staining with 1% crystal violet in 10% ethanol in PBS.

Immunohistochemistry. Sections of formalin-fixed paraffin-embedded normal, benign prostatic hyperplasia (BPH) and castrate-resistant prostate cancer (CRPC) tissue were warmed on a slide drier for 20 min at 45 °C, rehydrated through xylene and ethanol and then processed in a 2100 antigen-retriever (Aptum Biologics, Southampton, UK) for a two hour cycle in 10 mM sodium citrate buffer, pH6 containing 0.05% Tween-20. For immunoperoxidase staining, slides were then rinsed three times in phosphate-buffered saline (PBS), blocked with 10% foetal calf serum (FCS) in PBS for one hour at room temperature (RT) and incubated with primary antibodies in 10% FCS in PBS overnight at 4 °C or for 2 h at RT. Sections were rinsed three times in PBS, treated with 3% hydrogen peroxide in PBS for 30 min to remove endogenous peroxidases and rinsed well in PBS. Secondary biotinylated antibody in 10% FCS in PBS was then added for 30 min at room temperature followed by thorough rinsing in PBS.

Table 1. Details of cell lines, diagnosis, source and media requirements

Cell line	Diagnosis	Source/references	Culture medium
PNT2C2	Normal prostate epithelium	Obtained from ECACC (no longer available)	RPMI 1640 + 10% foetal calf serum (FCS) + 1% L-glutamine
PNT1A	Normal prostate epithelium	Kind gift to NJM's lab from P. Berthon. Currently available from Health Protection Agency Culture Collections.	RPMI 1640 + 10% FCS + 1% L-glutamine
P4E6	Prostate cancer	Derived in York. Currently available from Health Protection Agency Culture Collections	KSFM + 1% L-glutamine, bovine pituitary extract, epidermal growth factor and 2% FCS
LNCAp	Prostate cancer	ATCC	RPMI 1640 + 10% FCS + 1% L-glutamine
22RV1	Prostate cancer from a xenograft	RRIR: CVCL_1045	RPMI 1640 + 10% FCS + 1% L-glutamine
VCaP	Prostate cancer bone metastasis	ATCC	RPMI 1640 + 10% FCS + 1% L-glutamine
PC3	Prostate cancer bone metastasis	ATCC	Hams F12 + 7% FCS + 1% L-glutamine
PC3M	Prostate cancer bone metastasis	RRID:CVCL_9555	Hams F12 + 7% FCS + 1% L-glutamine

Finally, sections were treated with streptavidin-HRP in 10% FCS in PBS for 30 min at room temperature. After three rinses, sections were incubated with diaminobenzidine buffer (Vector kit or Sigma tablets) followed by extensive rinsing in PBS. Sections were counterstained with haematoxylin, rinsed with water and dehydrated through ethanol to xylene and mounted with DPX (Sigma Aldrich Company Ltd).

For immunofluorescence (IF), tissue sections were blocked with 10% FCS in PBS and incubated with primary antibody overnight at 4 °C in 10% FCS in PBS followed by further rinses. Secondary antibody in 10% FCS in PBS was added for 1 h followed by rinses. Slides were mounted using Vectashield with DAPI to define nuclei (Vector laboratories, Peterborough, UK) and examined using a Nikon Eclipse TE300 fluorescence microscope (Nikon, Surrey, UK). Antibodies used were a polyclonal anti-PLD1 (sc25512) at 1:100, an anti-p63 (Dako M7317, Agilent Technologies, Stockport, UK) at 1:100, an anti-AMACR at 1:100 (Dako M3616), an anti-chromogranin A at 1:400 (Invitrogen LK2H10 Fisher Scientific Ltd), a goat anti-rabbit immunoglobulin/biotinylated (Dako E0432) at 1:500 and a rabbit anti-mouse immunoglobulin/biotinylated (Dako E0354) at 1:200. Streptavidin-HRP (Dako, P0397) was used at 1:100.

Tissue microarray immunohistochemistry. A tissue microarray (TMA) of 41 kidney and 168 PCa tissue sections Gleason graded 6 ($n = 43$), 7 ($n = 97$), 8 ($n = 13$) or 9 ($n = 15$) from the Barts Cancer Institute was immunoperoxidase-stained for PLD1 as above. Stained sections were then scanned using a Zeiss AxioScan.Z1 slide scanner (ZEN 2012 software) with a Plan Apochromat 20 × / 0.8 objective. The composite.czi files were loaded into Tissue Gnostics, GmbH, HistoQuest software (V 3.5.3) for analysis. The intensity of DAB staining was quantified using the workable area (mean 0.3 mm²) of each tissue section. The results were analysed in a GraphPad Prism statistical package (Graphpad Software, CA, USA).

RESULTS

PLD1 protein expression in prostate cell lines. With equal protein loadings (30 µg), basal-like prostate epithelial cell lines PNT2C2, PNT1A, P4E6, and PC3 showed higher PLD1 protein expression than the luminal-like cell line LNCAp (Figure 1A). Two other luminal-like prostate epithelial cell lines VCaP and 22RV1 also expressed lower PLD1 protein compared with basal cell lines PNT2C2 and PC3 (Figure 1B). PLD1 protein expression in the

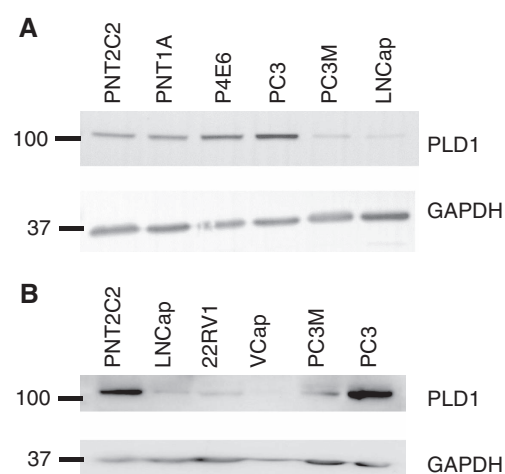


Figure 1. PLD1 protein expression in prostate epithelial cell lines. (A) PLD1 protein expression in normal and tumourigenic prostate epithelial cell lines. (B) PLD1 protein expression in luminal cell lines LNCAp, 22RV1 and VCaP. With all samples 30 µg protein were resolved by SDS-PAGE for western blotting with detection of GAPDH as a loading control. Markers are kDa. Blots shown are representative of several repeats. See Materials and Methods section for details.

metastatic PC3M cell line was much lower than in its parent cell line PC3.

PLD1 protein expression in patient-derived prostate epithelial and stromal cells. Cultured epithelial cells from Gleason 7 patient biopsies (Figure 2A) expressed PLD1 as did cells derived from prostate biopsies deemed to be normal (N) on pathological examination. PLD1 protein expression was usually higher in cells from Gleason 7 samples compared with cells from normal tissue but differences did not achieve statistical significance. Prostate epithelial cells derived from benign prostatic hyperplasia (BPH) tissue consistently had higher levels of PLD1 protein relative to cells from both normal and Gleason 7 prostate biopsies (Figure 2B). Stromal cells cultured from GL7 and GL9 tissue samples frequently expressed more PLD1 protein than the PC3 cell line or stromal cells purified from PCa tissue graded GL6 (Figure 2C).

PLD activity. Determination of PLD activity on tissue samples using the Amplex Red method gave results indicating that BPH,

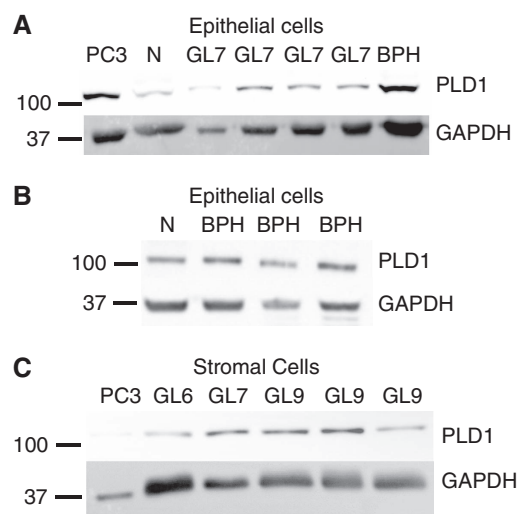


Figure 2. PLD1 protein expression in patient derived prostate cells. **(A)** PLD1 protein expression in epithelial cells derived from normal, BPH and Gleason 7 prostate tissue. **(B)** PLD1 protein expression in epithelial cells derived from normal and benign prostatic hyperplasia tissue. **(C)** PLD1 protein expression in stromal cells purified from prostate cancer tissue compared with PC3 cells. With all samples 30 µg protein were resolved by SDS-PAGE for western blotting with GAPDH as loading control. Markers are kDa. Blots shown are representative of several repeats. See Materials and Methods section for details. In Figure 2A and C, PC3 cells were included as a positive control for PLD1.

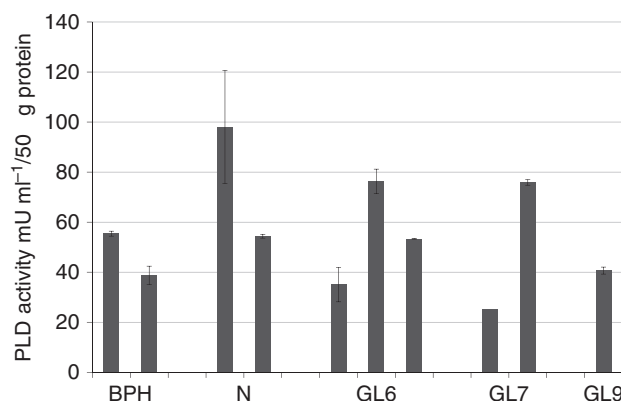


Figure 3. PLD activity in normal, BPH and Gleason-graded PCa tissue. Patient biopsy samples were extracted and analysed using a commercial Amplex Red PLD assay. 50 µg total protein were assayed per sample. Error bars on BPH and GL9 samples are standard deviation where triplicate measurements were made or on normal, GL6 and GL7 samples are a range where only duplicate readings were made per sample due to limited availability of tissue. Lack of an error bar for one GL7 sample is because there was only sufficient biopsy tissue for a single reading.

normal, and PCa tissue samples have similar levels of PLD activity (Figure 3). Error bars shown are the standard deviation ($n = 3$). The results without error bars are single results due to a limited amount of fresh biopsy tissue.

PLD inhibitors reduce cell viability in prostate epithelial cell lines. Changes in cell viability due to the actions of the dual PLD1/PLD2 inhibitors FIPI and 5W0 (VU0155056) and a PLD1-specific inhibitor EVJ (VU0359595) were evaluated after 24 and 48 h treatment (Figure 4A). All three inhibitors reduced the viability of prostate epithelial cell lines. The specific PLD1 inhibitor (EVJ) was the most effective since at 25 µM it reduced cell viability to almost zero after 48 h treatment. Cellular IC_{50} values calculated from the

MTS results using GraphPad prism (Table 2) confirmed that FIPI and 5W0 were not as effective at reducing cell viability as the specific PLD1 inhibitor EVJ.

PLD inhibitors reduce cell viability in patient-derived prostate epithelial cells. All the inhibitors reduced the viability of patient-derived epithelial cells over the 0–25 µM concentration range studied (Figure 4B). Again, the specific PLD1 inhibitor was more effective at reducing cell viability than FIPI and 5W0, especially with cells from patient 1 though in neither case was cell viability reduced to zero with 25 µM EVJ, some 20% of the cell population still remaining viable after 48 h treatment. Cellular IC_{50} values (Table 2) confirmed this finding.

PLD inhibitors reduce colony forming ability in patient-derived prostate cells. Exposure of cells from tissue from eight different PCa patients to 17.5 µM EVJ for 4 h significantly ($P = < 0.001$) reduced subsequent colony formation (> 32 cells) relative to DMSO vehicle controls (Figure 5) measured 2 weeks later. The median reduction relative to controls by the PLD1 inhibitor EVJ was about 30%. Of the eight patient samples treated with EVJ, five showed a uniform 25% reduction in colony-forming ability while three were more seriously affected.

PLD1 localisation in BPH tissue sections. Serial sections of BPH tissue were stained for PLD1, p63 as a positive control for basal cells and DAPI to reveal all cell nuclei. By IF, PLD1 protein was detected as perinuclear punctate staining predominantly in basal cells around prostate acini (Figure 6A, PLD1, arrows) compared with the nuclear p63 positive control (Figure 6A, p63, arrow). The merged images show that PLD1 is predominately expressed in basal cells. The enlarged IF image in Figure 6B (arrows) reveals the perinuclear punctate staining of PLD1 in the basal cells. This localisation of PLD1 to basal cells in BPH tissue was confirmed by immunoperoxidase staining. As shown in Figure 6C, PLD1-positive cells (arrows) matching the distribution of the basal cell marker p63 underlie a layer of light blue luminal cells lining the duct. Both immunohistochemical approaches revealed that some stromal cells in BPH tissue expressed PLD1.

PLD1 localisation in normal and tumourigenic prostate tissue samples. In tissue sections from a normal biopsy, PLD1 expression was detected by IHC in the nuclei and cytoplasm of rounded and elongated cells throughout the stroma, as well as prominently in basal cells round defined glands (Figure 7B, arrows). Lower PLD1 protein expression was also detected in luminal cell nuclei and cytoplasm (Figure 7B, arrow). In sections from a biopsy core graded Gleason 7, PLD1 protein was detected in elongated cells throughout the stroma as well as in the cytoplasm and nuclei of luminal cells (Figure 7C, arrows). In serial sections, these elongated PLD1-positive stromal cells also expressed smooth muscle actin (not shown). In CRPC tissue, PLD1 expression was detected in the cytoplasm and nuclei of many stromal cells (Figure 7D, arrows) and prominently in the cytoplasm of tightly packed cells (Figure 7D, *) that, in serial sections, stained positively for chromogranin A (Figure 7E *). A second antibody control of morphologically normal prostate tissue showed no peroxidase staining (Figure 7A).

PLD1 expression in a PCa tissue microarray. All PCa tissue sections graded Gleason 6–9 on a TMA showed PLD1 expression in the cytoplasm and nuclei of luminal cells in glands, some of which contained crystalloids (not shown). PLD1 expression was detected in scattered cells throughout the stroma in all sections examined. Some luminal cells showed intense PLD1 staining at their apical surfaces. Quantitative and statistical analysis using the GraphPad package revealed that the median peroxidase staining intensity of GL6 and GL7 sections was significantly ($*P < 0.05$) higher than that of GL9 sections (Figure 8). Control kidney tissue

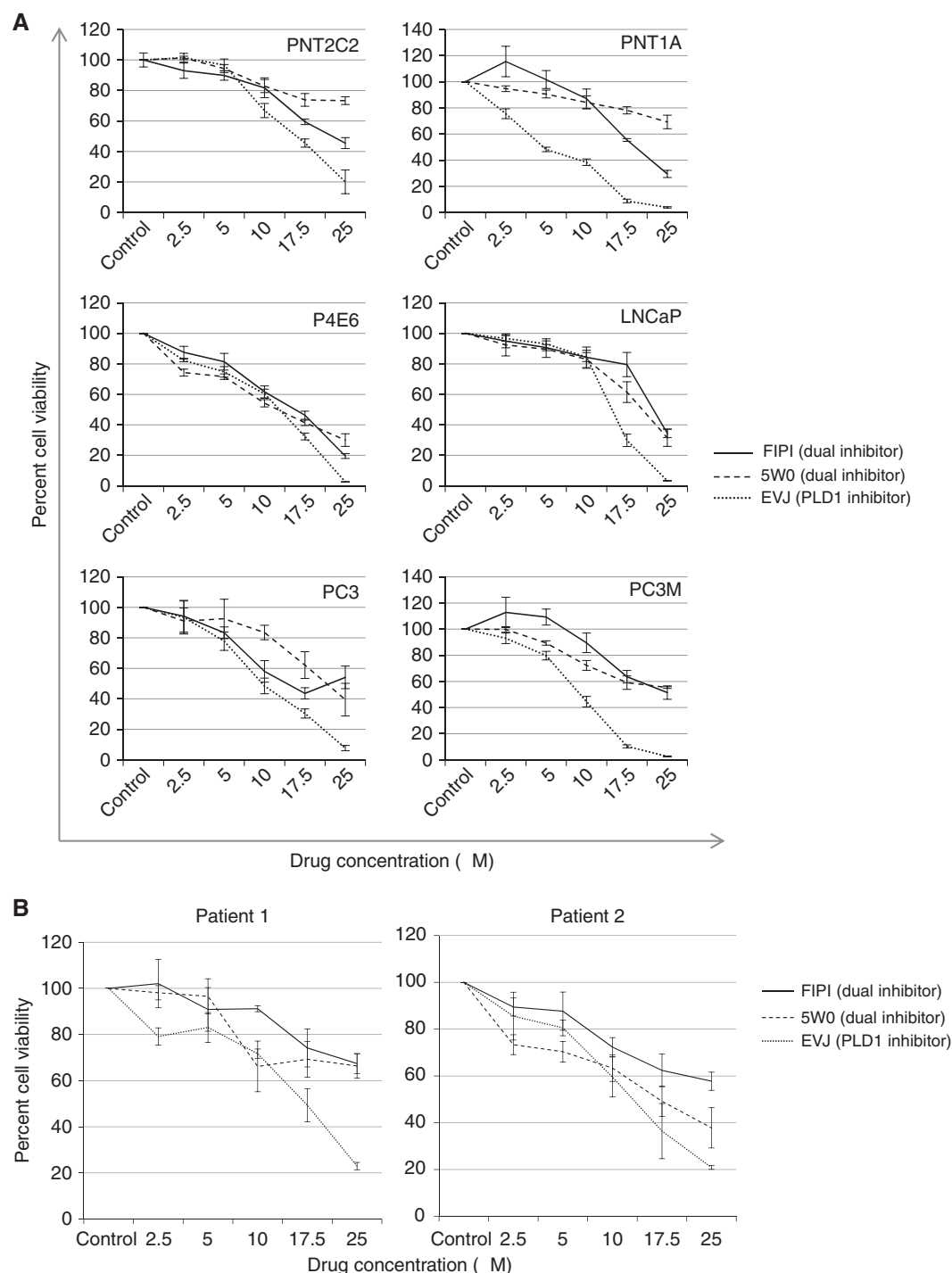


Figure 4. Effect of PLD inhibitors on prostate epithelial cell viability. **(A)** prostate cell lines, **(B)** patient-derived prostate epithelial cells. The cells were cultured with 0–25 μ M concentrations of the dual PLD1/PLD2 inhibitors FIPI and 5W0 and the PLD1-specific inhibitor EVJ. Inhibitors were dissolved in DMSO. Cell viability was measured by MTS assay after 48 h. Results are expressed as percentage cell viability relative to DMSO controls.

sections on the TMA showed positive PLD1 staining in epithelial layer cells lining most tubules.

DISCUSSION

The current consensus that PLD is involved in tumourigenesis (Jenkins and Frohman, 2005; Cazzolli *et al*, 2006; Wang *et al*, 2006; Foster, 2009; Winter *et al*, 2010; Kang *et al*, 2011; Henkels *et al*,

2013; Brown *et al*, 2017) has stimulated the development of new inhibitors specific for PLD1 and PLD2 that are well tolerated in mice (Scott *et al*, 2009). Although it is the major cancer of men in the Western world, surprisingly little is known about the expression and activity of PLD1 and PLD2 in PCa. In this first report on PLD isoforms in PCa, we have investigated PLD1 using a commercial anti-PLD1 antibody (sc-25512), which has been validated for western blotting (Disse *et al*, 2009; Scott *et al*, 2009; Chen *et al*, 2013) and which has also been used to detect PLD1 in cells by immunofluorescence (Disse

Table 2. Inhibitor IC ₅₀ values for FIPI, 5W0 and EVJ effects on cell viability as calculated from the MTS assay results using a Prism package			
Cell type	Inhibitor IC ₅₀ values (μm)		
	FIPI	5W0	EVJ
PNT2C2	28.2	60.4	17.3
PNT1A	27.1	56.8	4.7
P4E6	13.6	10.9	9.4
LNCaP	32	24.6	14.3
PC3	19.3	27.3	9.8
PC3M	42.9	29.8	8.0
Patient 1	57.6	39.8	14.8
Patient 2	30.1	13.9	12.0

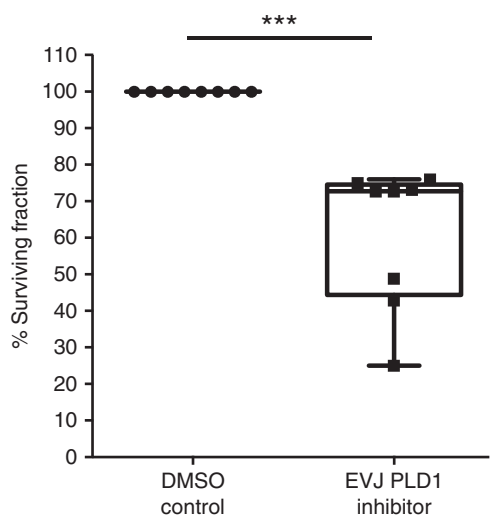


Figure 5. Effect of PLD1 inhibitor on colony recovery in patient-derived prostate epithelial cells. Patient-derived cells cultured from prostate cancer biopsies were treated with 17.5 μM EVJ (PLD1 inhibitor) for 4 h prior to re-plating cells at low density without inhibitor with STO feeder cells. Two weeks later, colonies of > 32 cells were counted and their number expressed relative to controls (DMSO vehicle only). A Mann–Whitney test gave a significance of *** $P < 0.001$. See Materials and Methods section for details.

et al, 2009; Han *et al*, 2011). We have not examined PLD2 expression in this work since, in agreement with Scott (Scott *et al*, 2009), we have found commercial anti-PLD2 antibodies to be unreliable.

PLD1 protein expression in prostate epithelial cell lines. We used the same panel of prostate epithelial cell lines as previously (Rumsby *et al*, 2011; Schmitt *et al*, 2014) to examine PLD1 protein expression in cells spanning the normal to metastatic extremes of PCa (Figure 1A). We added PC3M cells, a more metastatic cell line derived from PC3 cells. The finding that PLD1 protein expression was lower in PC3M cells compared with the parent PC3 cell line suggests that a PLD1-PtdOH-mTOR pathway may be less important for metastasis in prostate epithelial cells compared with other reported pathways such as an oncogene-tumour suppressor cascade (Min *et al*, 2010), DNA-PKcs-Mediated Transcriptional Regulation (Goodwin *et al*, 2015) and/or the tumour microenvironment (Mlecnik *et al*, 2016). Luminal LNCaP cells expressed the

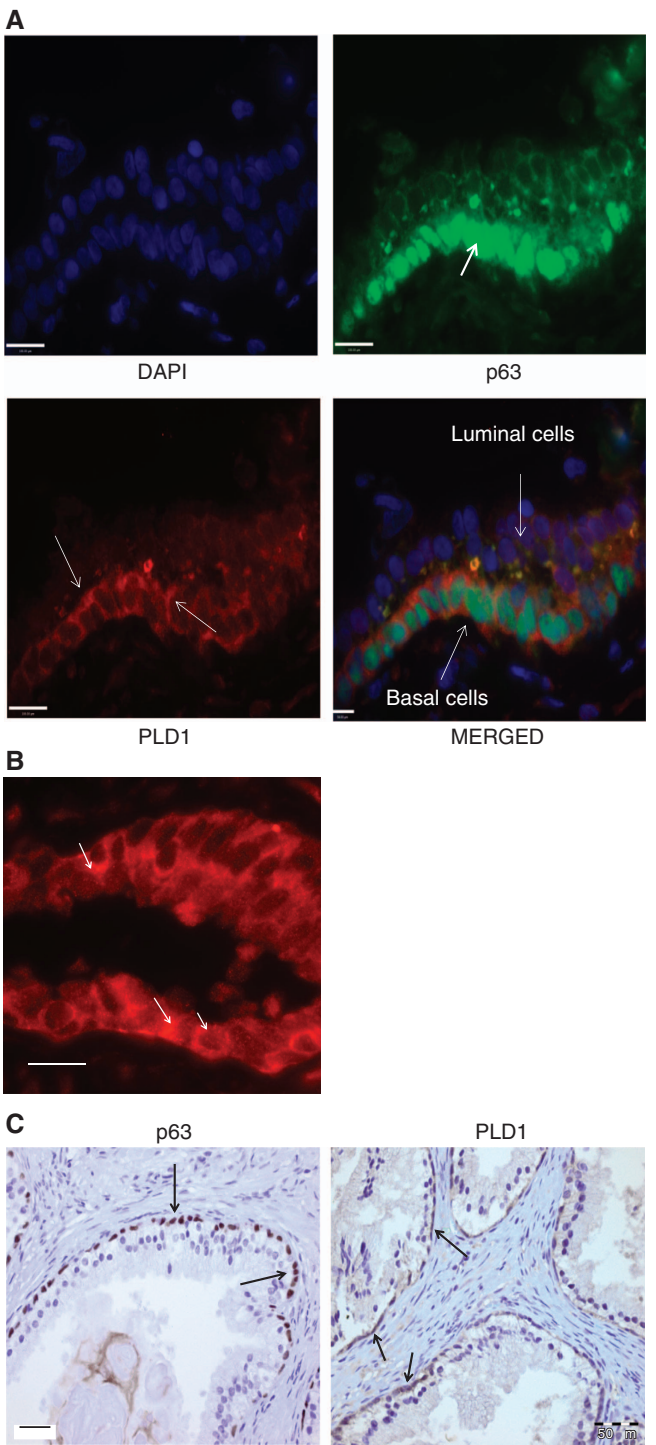


Figure 6. PLD1 is expressed in basal cells in BPH tissue. (A) Immunofluorescence: DAPI staining for nuclei, p63 marker for basal cells, PLD1 (arrows indicate perinuclear PLD1 staining), and merged images (scale bars 100 μm). (B) IF PLD1 in BPH tissue showing the perinuclear punctate pattern of expression (scale bar 90 μm) (C) Immunoperoxidase: BPH tissue stained for the basal cell marker p63 (arrows) and PLD1 (arrows). Scale bars: 50 μm. See Materials and Methods section for details of techniques.

lowest levels of PLD1 compared with the basal cell lines especially PC3 cells. This prompted us to check PLD1 protein expression in two other luminal prostate cell lines, 22RV1 derived from a tumour xenograft and VCaP derived from a PCa bone metastasis

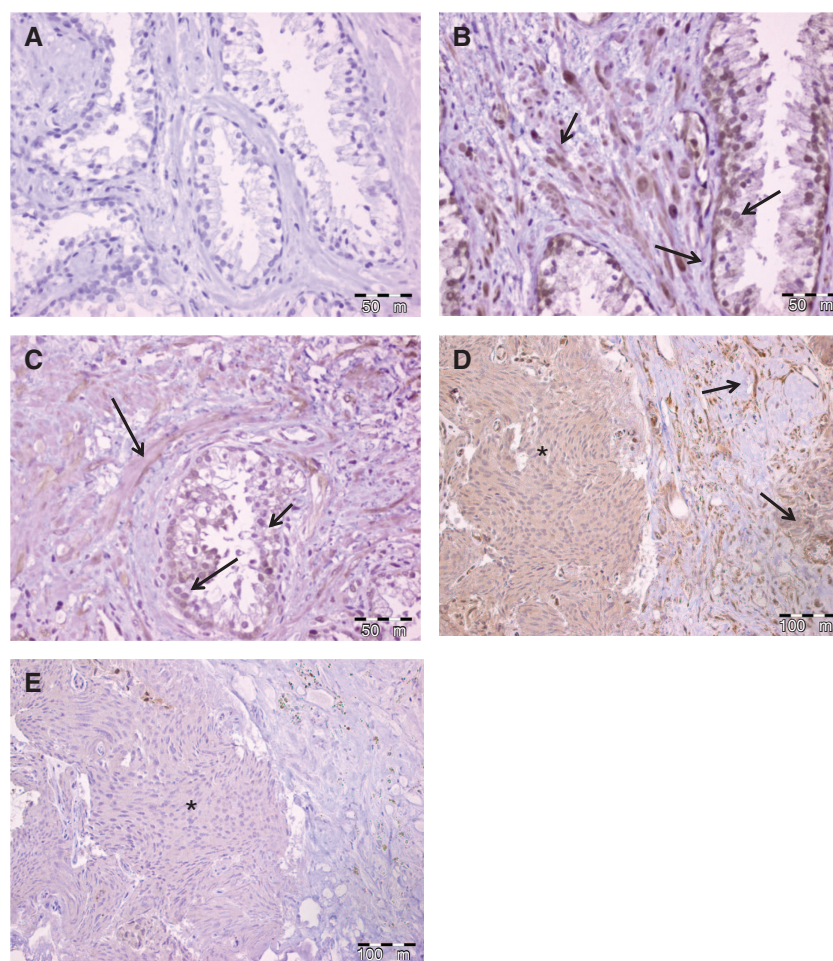


Figure 7. PLD1 protein expression in normal and tumourigenic prostate tissue. (A) Secondary antibody control, normal prostate tissue, (B) typical PLD1 expression in normal prostate tissue, (C) PLD1 expression in Gleason 7 PCa tissue, (D) PLD1 expression in castrate-resistant PCa tissue and (E) serial section of tissue in (D) showing cells staining positive for chromogranin (A) (*). Scale bar: (A–D) 50 μ m, (E) 100 μ m. See Materials and Methods section for details of techniques.

(Figure 1B). The finding that both had very low PLD1 protein expression similar to LNCaP cells reveals that basal and luminal prostate epithelial cell lines differ significantly in PLD1 protein expression.

This difference helps explain our previous results (Rumsby *et al*, 2011), where activation of protein kinase C (PKC) markedly stimulated choline headgroup release due to PLD1 activity in basal PC3 cells, while luminal LNCaP cells showed a very weak response. The difference would also explain our subsequent results (Schmitt *et al*, 2014) where stimulation of PKC in PC3 cells, but not in LNCaP cells, resulted in a rapid release of ethanolamine to the culture medium due to an increased turnover of ethanolamine phospholipids by PLD1. Our earlier results thus support this new finding that basal and luminal prostate epithelial cell lines differ significantly in their levels of PLD1 expression.

PLD1 protein expression in cells purified from prostate tissue.

Biopsy cores from different lobes of the prostate frequently show PCa in one lobe and no cancer in the other. We exploited such differences to isolate epithelial cells from biopsies of the same patient's prostate, which had been classified as malignant or non-malignant (normal). All cells whether cultured from non-malignant or malignant biopsy cores, or from TURPS (BPH or high Gleason samples) expressed PLD1 at levels that were usually lower than detected in the basal PC3 cell line. These biopsy-derived epithelial cell cultures are predominately basal in phenotype since

stromal and luminal cells are removed during preparation (Collins *et al*, 2001; Frame *et al*, 2016). PLD1 expression varied considerably (Figure 2A) between cell preparations cultured from PCa tissue of a similar Gleason grade. It was not routinely higher in cultures derived from malignant prostate biopsies as reported for some other cancers (Uchida *et al*, 1997; Uchida *et al*, 1999; Noh *et al*, 2000; Zhao *et al*, 2000; Saito *et al*, 2007). Such differences in PLD1 protein expression between cells isolated from PCa tissue of similar Gleason grade probably reflect inter-patient and biopsy variation. Significantly, however, epithelial cells cultured from BPH tissue consistently showed higher PLD1 protein expression compared with cells cultured from non-malignant or PCa tissue (Figure 2B). In BPH, basal, luminal and stromal cell numbers increase in the periurethral transition zone of the prostate (Dermer, 1978; Bonkhoff *et al*, 1994; Roehrborn, 2008; Rane *et al*, 2016) because of increased proliferation, or impaired programmed cell death, from the influence of androgens, oestrogens and reactive stromal growth factors (Roehrborn, 2008; Schauer and Rowley, 2011). The increased PLD1 protein expression we observe in these basal cell cultures could enhance the cellular proliferative changes observed in BPH tissue. This is because stromal growth factors such as EGF and IGF can stimulate PKC (Griner and Kazanietz, 2007; Lemmon and Schlessinger, 2010; Dowling and Kiely, 2015) thereby activating PLD1 and increasing PtdOH formation and mTOR signalling (Foster, 2009; Zhou and Huang, 2011).

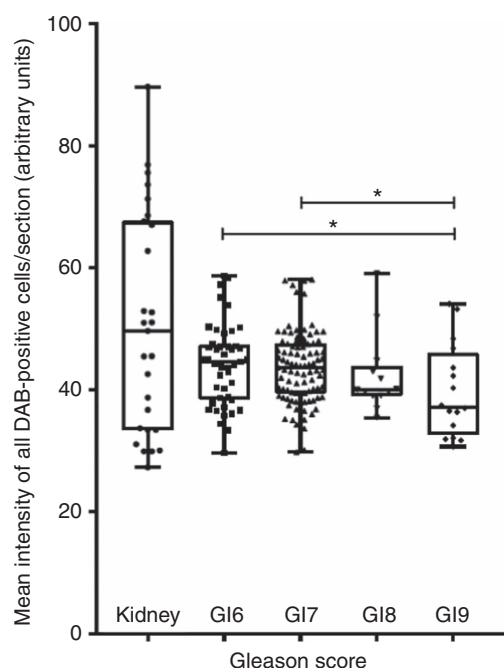


Figure 8. Quantification of DAB stain intensity of PLD1 protein in relation to Gleason grade in a tissue microarray. Over 150 PCa tissue sections were analysed for PLD1 expression in a tissue microarray as described in Materials and Methods section. Tissue sections graded GL6 and 7 have a significantly higher mean peroxidase stain intensity compared with GL9 sections (* $P < 0.05$).

PLD1 protein expression in non-malignant, malignant and BPH prostate tissue. Our IF and IHC results show that PLD1 is preferentially expressed in basal rather than luminal epithelial cells *in situ* in non-malignant and BPH tissue (Figures 6 and 7) in agreement with western blot results on cells. Basal cells expressing PLD1 are not observed in malignant tissue (Figure 7C) where proliferative luminal cells predominate (Jonathan and Epstein, 2008). The increased PLD1 expression observed in the expanding luminal compartment detected in PCa tissue (Figure 7C) may be regulating part of this proliferation process. If so, the TMA results suggest that PLD1 expression may play a more significant role in prostate tumours graded Gleason 6 or 7 compared with the more severe Gleason 9 stage. This agrees with our finding that more metastatic PC3M cells had lower levels of PLD1 expression than the less metastatic PC3 parental cell line.

The perinuclear punctate distribution of PLD1 in the cytosol of prostate basal cells as revealed by IF (Figure 6B) is in keeping with results by others using IF and overexpression methods (Brown *et al*, 1998; Cockcroft, 2001; Freyberg *et al*, 2001; Lucocq *et al*, 2001; Du *et al*, 2003; Disse *et al*, 2009; Han *et al*, 2011) or other anti-PLD1 antibodies (Freyberg *et al*, 2001; Vitale *et al*, 2001). These results suggest a role for PLD1 in regulated exocytosis involving lysosomes and endosomes and the generation of PtdOH for second messenger effects at the plasma membrane notably on mTOR and Raf (Foster *et al*, 2014). Our IHC detection of PLD1 in the nucleus of stromal and luminal cells in PCa is significant because nuclear PLD1 can activate a local nuclear PKC α and ERK signalling pathway to stimulate cell proliferation (Jang and Min, 2012). This can be regulated by several cell surface signalling pathways (Baldassare *et al*, 1997; Gayral *et al*, 2006; Ushio-Fukai, 2006; Jang and Min, 2012). Nuclear PLD1 is also linked to diacylglycerol formation, which regulates entry of PKC isoforms into the nucleus with effects on cell proliferation, gene expression and malignancy (Martelli *et al*, 2004; Tu-Sekine and Raben, 2004; Martelli *et al*, 2006). Stroma in non-malignant prostate tissue

contains mainly smooth muscle cells (Hagglof and Bergh, 2012) and our finding that such cells co-express PLD1 is not surprising since vascular smooth muscle cells also express PLD1 (Gayral *et al*, 2006). However, detection of PLD1 in CRPC, an incurable and aggressive terminal stage of PCa (Katsogiannou *et al*, 2015), is intriguing. Stromal cells that express PLD1 (Figures 2C and 7) also express smooth muscle actin so are likely to be myofibroblasts (Figure 7D) as smooth muscle cells decline in CRPC (Hagglof and Bergh, 2012). Such cells secrete a variety of growth factors (Hagglof and Bergh, 2012), which will activate PLD1 stimulating local cell proliferation. In CRPC tissue, these growth factors may also influence proliferation of the densely packed cells, which express PLD1 in their cytoplasm (Figure 7D and E*). We showed that these cells co-express chromogranin A (CgA), a marker for neuroendocrine cells (Deftos *et al*, 1996; Sun *et al*, 2009) and thus show focal neuroendocrine differentiation (Terry and Beltran, 2014). We find that similar PLD1-expressing cells in CRPC tissue also co-express neurone-specific enolase (NSE), another neuroendocrine cell marker (Deftos *et al*, 1996; Sun *et al*, 2009). These CgA- and NSE-expressing neuroendocrine-like cells are indicative of a poor prognosis (Terry and Beltran, 2014) and may arise from adenocarcinoma cells by neuroendocrine transdifferentiation involving IL-6 among other factors (Terry and Beltran, 2014; Grigore *et al*, 2015). PLD1 in the cytosol of these neuroendocrine-like cells may play a role in exocytosis important for cell proliferation (Vitale *et al*, 2001) and may also regulate cell growth through the generation of PtdOH and its links to Wnt signalling pathways (Kang *et al*, 2011).

PLD activity. The anomaly that PLD1 protein expression in BPH tissue samples is higher than in normal or PCa tissue, while PLD activity in the two BPH tissue samples measured is not raised above values for normal and PCa tissue may arise for two reasons. Firstly, PLD1 protein expression was measured in cultured cells from BPH tissue that are largely basal in phenotype, while PLD activity was assayed in whole BPH tissue samples which contain stromal and luminal cells as well as basal cells (Schauer and Rowley, 2011). Secondly, any nuclear PLD1 detected in BPH tissue by IHC would not have been assayed since these organelles would be removed during centrifugation to pellet cell debris. With these caveats, our activity results suggest that, unlike in breast adenocarcinomas and other cancers (see Introduction), PLD activity in PCa is not raised compared to normal tissue.

PLD inhibition. The potent effects of the new generation of PLD1 and PLD2 inhibitors (Monovich *et al*, 2007; Lavieri *et al*, 2009; Lewis *et al*, 2009; Scott *et al*, 2009; Lavieri *et al*, 2010) on cell survival show the importance of PLD isoforms in controlling prostate epithelial cell proliferation (Foster, 2009; Winter *et al*, 2010; Kang *et al*, 2011; Yoon *et al*, 2011). It should be noted that our IC₅₀ values (Table 2) are higher than those reported for the specific inhibition of PLD1 (Scott *et al*, 2009; Lavieri *et al*, 2010). This is because our cells were cultured in serum-containing medium under which conditions greater than 95% of these lipophilic inhibitors can become protein bound (Lavieri *et al*, 2010; Mathews *et al*, 2015). Our inhibitor concentrations in the μ M range are in agreement with other reports for the effects of these PLD inhibitors on cells growing in serum-supplemented medium (Lavieri *et al*, 2010; Liu *et al*, 2013; Mathews *et al*, 2015). Mathews *et al* (2015) report that at 5–10 μ M sufficient inhibitor is still available for subtype-selective inhibition of PLD1 in cells growing in serum-supplemented medium. The IC₅₀ values in Table 2 indicate that basal PC3 cells are more sensitive to PLD inhibitors than luminal LNCaP cells presumably because they express more PLD protein. The cells from two Gleason 7 patient tissues growing in serum free medium plus additives are also sensitive to the specific PLD1 inhibitor. Their IC₅₀ values are similar to those for metastatic PC3 and PC3M cells because such patient derived

prostate cells are often more resistant to treatments than cell lines (Ulukaya *et al*, 2013; Butler *et al*, 2017). We are now investigating the effects of the specific PLD2 inhibitor JWJ (VU0364739) on prostate cancer cell survival since this isoform has both enzymatic and GEF functions (Kang *et al*, 2014), which may give it a more direct role in proliferation, cell adhesion, migration and invasion (Chae *et al*, 2010).

The fact that a brief four hour exposure of cells to EVJ had a damaging effect on colony formation 2 weeks later indicates that short term inhibition of PLD1 has significant long term effects on PCa cell viability. That subsequent colony formation by PCa cells from three patients was more seriously affected by initial EVJ treatment than the five others must reflect differences such as stage of disease in the PCa biopsies from which the cells were purified. Our observations on EVJ effects on colony formation and cell viability suggest that this new specific PLD1 inhibitor has similar potential for treating PCa as already shown for breast cancer models in mice (Scott *et al*, 2009). In this connection, it is relevant that the antipsychotic agent haloperidol, on which these new inhibitors are based, is well tolerated in humans and has been tested in clinical trials (Loonen and Soudijn, 1985; Lindsley and Brown, 2012). EVJ could be used as a combination therapy alongside hormone/anti-androgen treatments that target luminal cells in PCa. PLD inhibition has also been shown to sensitise cancer cells to radiotherapy (Cheol Son *et al*, 2013) opening up further possible therapeutic applications.

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

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