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HDAC inhibitor confers radiosensitivity to prostate stem-like cells

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Background: Radiotherapy can be an effective treatment for prostate cancer, but radiorecurrent tumours do develop. Considering prostate cancer heterogeneity, we hypothesised that primitive stem-like cells may constitute the radiation-resistant fraction.

Methods: Primary cultures were derived from patients undergoing resection for prostate cancer or benign prostatic hyperplasia. After short-term culture, three populations of cells were sorted, reflecting the prostate epithelial hierarchy, namely stem-like cells (SCs, $\alpha_2\beta_1$ integrin^{hi}/CD133⁺), transit-amplifying (TA, $\alpha_2\beta_1$ integrin^{hi}/CD133[−]) and committed basal (CB, $\alpha_2\beta_1$ integrin^{lo}) cells. Radiosensitivity was measured by colony-forming efficiency (CFE) and DNA damage by comet assay and DNA damage foci quantification. Immunofluorescence and flow cytometry were used to measure heterochromatin. The HDAC (histone deacetylase) inhibitor Trichostatin A was used as a radiosensitiser.

Results: Stem-like cells had increased CFE post irradiation compared with the more differentiated cells (TA and CB). The SC population sustained fewer lethal double-strand breaks than either TA or CB cells, which correlated with SCs being less proliferative and having increased levels of heterochromatin. Finally, treatment with an HDAC inhibitor sensitised the SCs to radiation.

Interpretation: Prostate SCs are more radioresistant than more differentiated cell populations. We suggest that the primitive cells survive radiation therapy and that pre-treatment with HDAC inhibitors may sensitise this resistant fraction.

Precision radiotherapy (external beam or brachytherapy) is an important treatment for localised prostate cancer. However, despite improvements in targeting and dose, one-third of patients still fail because of local or systemic resistance (Brawer, 2002; Catton *et al*, 2003; Ishkanian *et al*, 2010). Metastases following radiotherapy are a significant clinical problem that requires an improved understanding of the biology of treatment-resistant phenotypes (Zafarana and Bristow, 2010; Jones, 2011). The use of *in vitro* cell line models of prostate cancer has demonstrated that

clones of radioresistant cells emerge post irradiation (Bromfield *et al*, 2003; van Oorschot *et al*, 2013). There have also been radiation response studies using normal prostate tissue and primary cells from normal biopsies (Kiviharju-af Hallstrom *et al*, 2007; Jaamaa *et al*, 2010; Zhang *et al*, 2011). However, none of these studies addressed the phenotype of the resistant clones. We, and others, have shown that a cellular hierarchy exists in several cancer types and in both prostate cancer epithelium and normal prostate epithelium (Collins *et al*, 2001; Hudson *et al*, 2001;

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Miller *et al*, 2005; Visvader and Lindeman, 2008; Maitland *et al*, 2010; Clevers, 2011). These studies demonstrated that the more stem-like cells (SCs), at the beginning of the hierarchy, have more clonogenic and tumourigenic potential than the more differentiated cells. Moreover, in glioblastoma, it was demonstrated that the primitive cells were more resistant to radiotherapy than the majority of cells within the tumour (Bao *et al*, 2006). This finding has been backed up by other studies suggesting that the SCs may be directly responsible for tumour recurrence (Chiou *et al*, 2008; Diehn *et al*, 2009b; Conley *et al*, 2012; Chen *et al*, 2013). In light of these findings, we hypothesised that the SCs in prostate cancer would be more resistant to irradiation than the more differentiated populations. Using the same markers we had previously used to isolate the normal and malignant prostate hierarchy (Collins *et al*, 2001; Richardson *et al*, 2004; Collins *et al*, 2005), we show here that the most undifferentiated cells in both benign and malignant primary cultures are more resistant to irradiation. This resistance is

conferred by heterochromatin, which protects the cells from the DNA-damaging effects of radiation.

MATERIALS AND METHODS

Tissue collection, isolation and culture of tumour cells. Human prostate tissue was obtained with patient's consent and full ethical approval from patients undergoing radical prostatectomy and channel transurethral resection (TURP) for prostate cancer and from patients undergoing transurethral resection for benign prostatic hyperplasia (BPH) (Table 1). Grade and stage of tumour were confirmed by histologic examination of representative fragments by a uropathologist. Epithelial cultures were prepared and characterised as described previously (Collins *et al*, 2001). Cell cultures were maintained in stem cell media (SCM) consisting of

Table 1. Patient samples

Sample	Passage	Operation	Patient age	Diagnosis	Hormone status
Benign samples					
B1	8 + 5	C	—	Benign	—
B2	5	T	58	Benign	—
B3	5	T	77	Benign	—
B4	4	T	88	Benign	—
B5	2 + 6	C	72	Benign	—
B6	5	T	83	Benign	—
B7	3	T	82	Benign	—
B8	4	T	65	Benign	—
B9	3	T	83	Benign	—
B10	6	T	77	Benign	—
B11	5	T	67	Benign	—
B12	6	T	74	Benign	—
B13	1	T	—	Benign	—
B14	2	T	72	Benign	—
B15	3	T	71	Benign	—
B16	3	T	62	Benign	—
B17	3	T	84	Benign	—
B18	4	R	61	Benign	—
Cancer samples					
C1	3 + 4	T	71	Cancer G16 (3 + 3)	Naive
C2	3	R	68	Cancer G16 (3 + 3)	Naive
C3	3	R	68	Cancer G16 (3 + 3)	Naive
C4	5 + 6	R	62	Cancer G17 (3 + 4)	Naive
C5	10	R	53	Cancer G17 (3 + 4)	Naive
C6	3	R	65	Cancer G17 (3 + 4)	Naive
C7	2	R	70	Cancer G17 (3 + 4)	Naive
C8	4	R	64	Cancer G17 (3 + 4)	Naive
C9	4	R	64	Cancer G17 (3 + 4)	Naive
C10	6	R	—	Cancer G17 (4 + 3)	Naive
C11	3	R	58	Cancer G17 (3 + 4)	Naive
C12	6	R	47	Cancer G17 (4 + 3)	Naive
C13	8	R	—	Cancer G17 (3 + 4)	Naive
C14	9	R	66	Cancer G17 (3 + 4)	Naive
C15	4	R	70	Cancer G17 (3 + 4)	Naive
C16	4	R	70	Cancer G17 (3 + 4)	Naive
C17	3 + 6	T	59	Cancer G17 (3 + 4)	Sensitive
C18	4	T	73	Cancer G18 (3 + 5)	Sensitive
C19	3	R	—	Cancer G18 (3 + 5)	Naive
C20	3	R	—	Cancer G18 (3 + 5)	Naive
C21	7	T	85	Cancer G19 (4 + 5)	Sensitive
C22	4 + 5	T	80	Cancer G10 (5 + 5)	Sensitive

Abbreviations: C = cystectomy; R = radical prostatectomy; T = transurethral resection of the prostate.

keratinocyte growth medium supplemented with EGF, bovine pituitary extract (Life Technologies Ltd, Paisley, UK), 2 ng ml⁻¹ stem cell factor (SCF) (First Link UK Ltd, Wolverhampton, UK), 100 ng ml⁻¹ cholera toxin (Sigma-Aldrich Company Ltd, Gillingham, UK) and 1 ng ml⁻¹ granulocyte macrophage colony-stimulating factor (GM-CSF) (First Link UK Ltd). Cells were cultured in the presence of irradiated (60 Gy) STO (mouse embryonic fibroblast) cells. After expansion, CD133⁺/α₂β₁integrin^{hi} (stem-like (SC)), CD133⁻/α₂β₁integrin^{hi} (transit-amplifying (TA)) and α₂β₁integrin^{lo} (committed basal (CB)) cells were isolated by magnetic-activated cell sorting (MACS; Miltenyi Biotec, Surrey, UK) as described previously (Richardson *et al*, 2004; Collins *et al*, 2005). SC cells are the most primitive cells, with TA cells being a progenitor population and CB cells being further along the differentiation hierarchy.

Irradiation of cells. To irradiate cells, an RS2000 X-Ray Biological Irradiator was used that contains a Comet MXR-165 X-Ray Source (Rad-Source Technologies Inc., Suwanee, GA, USA). A dose of 2 or 10 Gy was administered with a dose rate of 0.02 or 0.08 Gy s⁻¹. To determine colony-forming ability post irradiation, primary cultures were irradiated as a whole population and subsequently sorted. To assay DNA damage, in response to radiation, primary cells were sorted into their respective populations before irradiation because of the rapid nature of DNA damage formation.

Clonogenic recovery. Primary prostate cultures were irradiated (2 Gy) and immediately sorted into subpopulations (SC, TA and CB), counted and plated on to 35-mm collagen-coated plates (BD Biocoat, BD Biosciences, Oxford, UK) at a density of 100 cells per well in the presence of irradiated STO feeder cells. For treatment with HDAC inhibitor, cells were treated with 0.6 μM of Trichostatin A (TSA; Sigma-Aldrich Company Ltd, T1952) for 1 h and 30 min and then irradiated (2 Gy) and treated as above. Colonies were subsequently scored if they contained >32 cells (at least 5 population doublings, which are considered as self-sustaining colonies with proliferation potential (Puck and Marcus, 1956; Francipane *et al*, 2008), usually ~14 days after treatment. Colonies were visualised by staining with 1% crystal violet/10% ethanol/PBS.

Alkaline and neutral comet assays. Comet assays were modified from Sturmey *et al* (2009). Primary prostate epithelial cells were separated into SC, TA and CB cells, resuspended in 25 μl PBS and irradiated. At 30 min post irradiation, 225 μl of low-melting-point (LMP) agarose was added to the cells, mixed and pipetted onto an agarose-coated slide (1% agarose in PBS). A clean coverslip was placed on top and slides placed at 4 °C for 15 min. Once set, coverslips were removed and slides placed in lysis buffer (2.5 M NaCl, 1 mM EDTA, 10 mM Tris, 10% DMSO, 1% Triton X-100, pH 10) overnight at 4 °C. For the alkaline comet assays, slides were incubated in alkaline solution (0.3 M NaOH, 1 mM EDTA, pH 13) for 40 min at 4 °C, and then electrophoresed at 23 V, 300 mA in the same alkaline solution (for 40 min on ice). Subsequently, they were incubated for 2 × 10 min in neutralising buffer (0.4 M Tris, pH 7.5). For the neutral comet assays, slides were incubated in TBE electrophoresis buffer at 4 °C for 40 min and electrophoresed at 25 V for 20 min. For both assays SYBRgold was applied (1:10 000 in TE buffer, 10 mM Tris-Cl, pH 7.5, 1 mM EDTA) to stain the DNA. Comets were measured using CometScore freeware (TriTek Corp., Sumerduck, VA, USA) following image capture using a Nikon Eclipse TE300 fluorescent microscope (Nikon, Surrey, UK) with a ×10 lens.

Immunofluorescence. Immunocytochemistry was conducted on sorted α₂β₁integrin^{hi}/CD133⁺ (SC), α₂β₁integrin^{hi}/CD133⁻ (TA) and α₂β₁integrin^{lo} (CB) primary cells. The cell isolates were allowed to adhere for 2 h onto collagen-coated slides, and in some experiments were treated with 0.6 μM of TSA (Sigma-Aldrich Company Ltd T1952) for 1 h and 30 min and irradiated (2 Gy) before fixing in 2% paraformaldehyde/PBS/0.2% Triton X-100, pH

8.2 (20 min), then permeabilised with 0.5% NP40/PBS (20 min). Cells were then blocked (2% BSA/1% normal goat serum/PBS) and incubated in primary antibody in 3% BSA/PBS at 4 °C overnight followed by washes in 0.5% BSA/0.175% Tween-20/PBS. Secondary antibody was added for 45 min in 3% BSA/PBS followed by washes. Slides were mounted with Vectashield with DAPI (Vector Laboratories, Peterborough, UK). Primary antibodies used were as follows: anti-phospho-Histone H2A.X (Ser139), clone JBW301 (Millipore, Watford, UK); 53BP1 (Alexis Biochemicals, Exeter, UK, ALK-210-419 or Abcam, Cambridge, UK, ab36823); phospho-Chk2 (Thr68; Cell Signaling Technology, Hitchin, UK, 2661); phospho-(Ser/Thr) ATM/ATR substrate antibody (Cell Signaling Technology 2851); H3K9me3 (histone H3(trimethylated at lysine 9); Diagenode, Denville, NJ, USA, pAb-056-050); Ki67 (Abcam ab15580); H3K27me3 (histone H3(trimethylated at lysine 27); Millipore 17-622). Secondary antibodies were goat anti-mouse AlexaFluor-488 (Life Technologies Ltd A11029) and goat anti-rabbit AlexaFluor-568 (Life Technologies Ltd A11036).

Confocal images were acquired on Zeiss LSM 510 and Zeiss LSM Meta Laser confocal microscopes (Zeiss, Cambridge, UK). Colocalisation analysis, Pearson's correlation, line scan analysis and fluorescence intensity measurements were carried out using Velocity software (Improvision, Perkin Elmer, Waltham, MA, USA). For foci counting, images were captured using a Nikon Eclipse TE300 fluorescent microscope (Nikon, Surrey, UK) with a ×63 oil immersion lens and foci were counted using the multi-point tool in ImageJ software (Town of Mount Royal, QC, Canada).

Flow cytometry. Primary cultures were trypsinised, resuspended in MACs buffer and incubated with antibodies to the α₂ subunit of integrin α₂β₁ (CD49b-RPE, MCA743PET, AbD Serotec, Kidlington, UK) and CD133/2 (293C)-APC (130-090-854, Miltenyi Biotec) for 10 min at 4 °C.

For the detection of heterochromatin, the cells were then permeabilised in MACs buffer with 0.5% W/V saponin and 20% normal goat serum (NGS) and then incubated with the H3K9me3 antibody (Diagenode) or H3K27me3 antibody (Millipore 17-622) in MACs buffer with 20% NGS. Cells were then washed and incubated with goat anti-rabbit AlexaFluor-488 secondary antibody (Life Technologies Ltd) in MACs buffer with 20% NGS. Finally, cells were analysed on a CyAn-ADP flow cytometer (Beckman Coulter, High Wycombe, UK) and data processed using Summit v4.3 software (Beckman Coulter).

Statistical analysis. Median values of each analysis were calculated. Significance was assessed using the Wilcoxon rank-sum test or *t*-test for smaller sample numbers where stated (GraphPad Prism, La Jolla, CA, USA). The *P*-values indicating statistical significance are displayed.

RESULTS

SCs are more radioresistant than progenitor cells. To establish whether prostate SCs are more resistant to irradiation than progenitors, we determined the potential of each population to initiate colonies (at clonal density) following irradiation. Our rationale for using 2 Gy was to mimic the clinical daily dose of 2 Gy that has been routinely administered to patients (Bromfield *et al*, 2003). Primary cells, derived from patients with BPH, and prostate cancer, including those who had undergone androgen-ablation therapy, were irradiated, sorted into SCs and progenitors (TA and CB) and plated to determine colony-forming efficiency (CFE) that is represented as surviving fraction (SF) after 2 Gy (SF2Gy; Figure 1A and B). The highest SF2Gy (post irradiation) was observed in the SCs, independent of disease status (Figure 1A and B). To investigate the consequences of irradiation on each cell type, DNA damage was measured following radiation in each

Figure 1. Stem-like cells (SCs) from benign and cancer cultures are less radiosensitive and have a differential DNA damage response compared to progenitor TA cells and more differentiated CB cells. (**A** and **B**) Benign and malignant primary prostate epithelial cultures were treated with 2 Gy radiation and cell populations sorted and plated for clonogenic assays. Surviving fraction following 2 Gy (SF2Gy) was calculated and plotted for each cell subpopulation. Each symbol represents a different patient sample (red symbols indicate prostate cancer and blue symbols indicate BPH) and the black bar represents the median. (**C** and **D**) Benign and malignant primary prostate epithelial cultures were sorted into cell subpopulations and treated with 2 Gy radiation, and then processed in alkaline (**C**) and neutral (**D**) comet assays. Each symbol represents a different patient sample (red symbols indicate prostate cancer and blue symbols indicate BPH) and the black bar represents the median. Statistical significance values were measured using the Wilcoxon rank-sum test (* $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$).

population (Figure 1C and D). To do this we used comet assays to measure DNA damage directly, following radiation doses of 2 and 10 Gy. Alkaline comet assays were used to measure single-strand breaks (SSBs), double-strand breaks (DSBs), alkali labile sites, DNA crosslinks and incomplete excision repair (Singh *et al*, 1988; Tice *et al*, 2000; Nandhakumar *et al*, 2011; Swain and Subba Rao, 2011). We observed comparable damage in each population, with a clear dose-dependent increase (up to 10 Gy; Figure 1C). In contrast, using the neutral comet assay, we observed that the SCs incurred significantly less damage than either the TA or CB populations (Figure 1D). Moreover, there was no significant difference in DNA damage with increasing dose. The neutral comet assay has a bias for

measuring lethal DSBs (Ostling and Johanson, 1984; Singh *et al*, 1988; Lemay and Wood, 1999; Wojewodzka *et al*, 2002; Van Kooij *et al*, 2004; Swain and Subba Rao, 2011). This suggests that the SCs are protected from this specific type of damage compared with the progenitor (TA) and more differentiated (CB) cells. As observed with the colony-forming assay, this effect on DNA damage was independent of disease status.

SCs sustain less DSBs than progenitors. To confirm this finding, we investigated the DSB DNA damage response further. Primary cells were sorted, irradiated and fixed at 30 min post irradiation. Cells were then stained for γ H2AX and 53BP1 DNA damage nuclear foci,

Figure 2. Fewer SCs from benign and malignant primary epithelial cultures contain DNA damage foci relative to TA and CB cells post irradiation. **(A)** Cells from benign and malignant primary prostate epithelial cultures were sorted then fixed and stained for γ H2AX and 53BP1 at 30 min post irradiation. Representative images are shown of nuclei showing (i) negative (ii) and positive staining for DNA damage foci (γ H2AX (green), 53BP1 (red) and DAPI nuclear staining (blue)). Image squares containing a single nucleus measured $27.35 \mu\text{m}$ by $27.35 \mu\text{m}$. **(B–E)** Cell nuclei were scored for foci and ~ 100 nuclei per sample were counted. The graphs indicate the percentage of cell nuclei containing foci. Each symbol represents a different patient sample (red symbols indicate prostate cancer and blue symbols indicate BPH). Boxplots show minimum, 25%, median, 75% and maximum. Statistical significance values were measured using the Wilcoxon rank-sum test (** $P < 0.01$, *** $P < 0.001$).

indicative of DSBs (Paull *et al*, 2000; Belyaev, 2010; Mah *et al*, 2010) (Figure 2A). Approximately 90% of TA and CB cell nuclei showed evidence of DSBs, whereas significantly fewer (30–50%) SCs sustained DSBs (Figure 2B–E). These results are in agreement with our findings using the neutral comet assay, showing evidence of DSBs in the majority of TA and CB cells, but only in a minority of SCs.

We also investigated the downstream events following induction of DNA damage by staining the cell populations for members of the ATM-Chk2 DSB DNA damage pathway (Figure 3). Again, we observed that the percentage of SCs containing ATM/ATR phosphorylated substrates and phospho-Chk2^{Thr68} was significantly less than either of the progenitor populations, again reinforcing our findings that SCs sustain less DSBs.

Although the majority of SCs appear to sustain no damage, we wanted to know if the minority subpopulation that did sustain damage could undertake DNA repair. To do this, the number of

foci *per cell nucleus* was quantified. It was clear that in all cell types (SC, TA, CB), and in all patients, DNA repair was taking place, as evidenced by a significant reduction in the number of foci *per cell nucleus* at 24 h compared with 30 min (Figure 4).

SCs are less proliferative and have increased heterochromatin.

As SCs appeared to sustain less DNA damage following irradiation, we hypothesised that they may have an inherent protection mechanism. We had previously demonstrated that the SCs are less proliferative (Lang *et al*, 2010), and hence we quantified the number of cells expressing the proliferation marker Ki67 (Figure 5A). Primary cells from six patients were sorted as previously described and stained with an antibody against Ki67. We found that the percentage of SCs expressing Ki67 was significantly less than either the TA or CB cells, suggesting that the progenitor (TA) and more differentiated (CB) cells are more proliferative than the SCs (Figure 5A).

Figure 3. Fewer prostate SCs from benign and malignant primary epithelial cultures have an active ATM/Chk2 DNA damage response post irradiation. Nuclear DNA damage foci were scored as in Figure 2, after staining with ATM/ATR substrate antibody (A) and pChk2^{Thr68} antibody (B). Percentage of cell nuclei that contain foci is represented. Each symbol represents a different patient sample (red symbols indicate prostate cancer and blue symbols indicate BPH). Representative images of nuclei showing co-staining of γ H2AX antibody (green), (Ai) ATM/ATR substrate antibody or (Bi) pChk2^{Thr68} antibodies (red) and DAPI nuclear stain (blue) are shown. Boxplots show minimum, 25%, median, 75% and maximum. Statistical significance values were measured using the Wilcoxon rank-sum test (* $P < 0.05$, ** $P < 0.01$).

Figure 4. All cell types are equally capable of undergoing DNA repair post irradiation. Number of foci per cell nucleus were counted at 30 min and 24 h post irradiation in SCs (**A**), TA (**B**) and CB (**C**) cells. Each symbol represents a single cell (red symbols indicate prostate cancer and blue symbols indicate BPH). Boxplots show minimum, 25%, median, 75% and maximum. Statistical significance values were measured using the Wilcoxon rank-sum test (* $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$).

One other mechanism by which a cell could sustain less damage is by packing more DNA into heterochromatin. Storch *et al* (2010) found that growing cells in a 3D formation led to increased chromatin condensation, which subsequently resulted in radioresistance. Furthermore, when measuring the number of DSBs in euchromatin vs heterochromatin in monolayer cells, the authors found a ratio of 2:1, that is, fewer breaks in heterochromatin. When we examined heterochromatin marks (H3K27me3 and H3K9me3) in each cell population, we observed that a proportion of SCs appeared to have increased heterochromatin content compared with the other cell types (TA and CB) (Figure 5B). We quantified heterochromatin by measuring the fluorescence intensity of sorted cells (Figure 5C) as well as flow cytometry where cells were co-stained with CD49b ($\alpha_2\beta_1$ integrin) and CD133 to

distinguish the three populations (Figure 5D). This showed that SCs had significantly increased heterochromatin compared with the progenitor (TA) and more differentiated (CB) cells.

To confirm that heterochromatin conferred a protective effect to the cells, we carried out dual staining of DNA damage foci and heterochromatin that showed preferential formation of foci located almost exclusively at nuclear sites devoid of heterochromatin staining (Figure 5Ei). In addition, several SCs were observed to have high levels of heterochromatin throughout the nucleus that corresponded with an absence of foci (Figure 5Eii).

Combination treatment of HDAC inhibitor and radiation results in increased DNA damage and reduced clonogenic survival in the SCs. Based on the heterochromatic patterns, we

Figure 5. Stem-like cells (SCs) from benign and malignant primary epithelial cultures are less proliferative and have more heterochromatin than more differentiated TA and CB cells. **(A)** Cells were stained for Ki67 and scored for positive staining and presented as a percentage of the total cell number. Each symbol represents a different patient sample (red symbols indicate prostate cancer and blue symbols indicate BPH). Boxplots show minimum, 25%, median, 75% and maximum. **(B)** Staining for H3K9me3 (heterochromatin). **(C)** Cells were analysed for amount of heterochromatin using intensity of immunofluorescence staining (analysis using Velocity software) (i) H3K9me3 (ii) H3K27me3. **(D)** Cells were analysed for heterochromatin content using flow cytometry analysis (i) H3K9me3 (ii) H3K27me3. Each symbol represents a different patient sample (red symbols indicate prostate cancer and blue symbols indicate BPH) and the black bar represents the mean of the total fluorescence of each cell population divided by the median of the CB value. **(E)** Co-staining of γ H2AX and H3K9me3 (heterochromatin). (i) The γ H2AX-positive SC, TA and CB cells were analysed for colocalisation with heterochromatin (analysis using Velocity software). (ii) Representative images of SCs negative for γ H2AX nuclear foci. Statistical significance values were measured using the Wilcoxon rank-sum test (* $P < 0.05$, ** $P < 0.01$).

hypothesised that manipulating chromatin status in the SCs would render them more sensitive to radiation treatment. To achieve this, cells were sorted and treated with a low dose of HDAC inhibitor ($0.6 \mu\text{M}$ TSA) for 1 h and 30 min followed by 2 Gy irradiation. The treated cells were then fixed 30 min post irradiation and subsequently stained for DNA damage foci (Figure 6A). The effect

of TSA was confirmed by western blot using lysates from unsorted cells for acetylated histones where we observed an increase in histone acetylation following treatment (Figure 6B). This is known to correlate with chromatin decondensation (Toth *et al.*, 2004). Following the combination treatment, the number of SCs sustaining DSBs, as evidenced by increased number of cells containing foci, significantly increased (Figure 6Ai). There was no significant increase in the percentage of TA or CB cells containing foci as the majority of these cells contain foci following radiation alone, and hence the effect is already saturated. On quantifying foci number in the SCs we found no increase in number of foci *per cell nucleus* following TSA treatment (Figure 6Aii). We then assessed the effect of combination treatment on clonogenic recovery and found that the SCs formed significantly fewer colonies following combination treatment compared with irradiation alone (Figure 6C). We concluded from this experiment that the HDAC inhibitor had sensitised the SCs to radiation treatment.

DISCUSSION

In this study, we present evidence demonstrating that primitive prostate SCs, from freshly cultured patient tissues, are more radioresistant than more differentiated cells, which we show to be independent of patient disease status. We propose that the SCs, by having higher levels of heterochromatin, sustain fewer lethal DSBs, which contributes to increased survival. Significantly, by treating this population with an HDAC inhibitor, DNA damage was increased, resulting in sensitisation of the cells to radiation, thus reducing survival.

Using colony-forming assays, we were able to demonstrate that SCs were less affected by irradiation as compared with the progenitor cells. Although the overall induction of DNA damage was similar between populations (alkaline comet assays), we observed a significant reduction in the percentage of SCs sustaining lethal DSBs (neutral comet assays and DNA damage foci). Lack of activation of the ATM/ATR DNA damage signaling pathways correlated with this population. By quantifying the number of foci in the minority of SCs positive for foci, as well as TA and CB cells, we confirmed that DNA repair was undertaken in all cell types.

We present evidence here that SCs sustain less DNA damage because of increased heterochromatin content. Our results pointed towards the use of HDAC inhibitors, in combination with radiation, as a therapy for prostate cancer. HDAC inhibitors have been heavily investigated for their clinical use and are also in

Figure 6. Treatment with an HDAC inhibitor radiosensitises stem cells from benign and malignant primary epithelial cultures (**Ai**) Stem-like cells (SCs), TA and CB cells were treated with $0.6 \mu\text{M}$ TSA for 90 min, irradiated (2 Gy) and then fixed 30 min post IR and stained for γ H2AX. Each symbol represents a different patient sample (red symbols indicate prostate cancer and blue symbols indicate BPH) and the black bar represents the median. (**Aii**) Number of foci per cell nucleus of SCs, following irradiation or TSA plus irradiation. (**B**) Western blot for acetyl-H3 in unsorted cell lysates following treatment with $0.6 \mu\text{M}$ TSA for 2 h. (**C**) Whole populations of primary epithelial cell cultures were treated with $0.6 \mu\text{M}$ TSA for 90 min, irradiated (2 Gy) and sorted into subpopulations 30 min post IR. Surviving fraction following 2 Gy (SF2Gy) and surviving fraction following TSA treatment and 2 Gy (SF-TSA-2Gy) were calculated and plotted for SCs following clonogenic assays. Each symbol represents a different patient sample (red symbols indicate prostate cancer and blue symbols indicate BPH), Boxplots show minimum, 25%, median, 75% and maximum. Statistical significance values were measured using the Wilcoxon rank-sum test (* $P < 0.05$).

clinical trials for prostate cancer (Marchion and Munster, 2007; Frew *et al*, 2009; Atadja, 2011). There are encouraging results with HDAC inhibitors targeting proliferating and nonproliferating cells (Burgess *et al*, 2004). However, they are often being employed as toxins, rather than response modifiers, the latter being what we propose based on the data in Figure 6. Our work shows that a short exposure to sublethal doses of HDAC inhibitors may be enough to sensitise prostate SCs.

It is also possible that less damage is sustained because of the less proliferative nature of SCs. It has previously been shown that adult SCs are predominantly quiescent and that this can reduce accumulation of mutations through DNA replication and contribute to therapy resistance (Mohrin *et al*, 2010; Blanpain *et al*, 2011; Li and Bhatia, 2011). Using the vital dye PKH26, we were able to show that prostate stem cells undergo up to four population doublings (in monolayer culture) before exiting the cell cycle (Lang *et al*, 2010). We also observed here that significantly more progenitor and more differentiated cells (TA and CB cells) are in cycle compared with the SCs.

In summary, radiorecurrent prostate cancer is a significant problem for both physicians and patients because of local and systemic resistance (Ishkanian *et al*, 2010). The heterogeneity of prostate cancers has been well documented, yet we are the first to report the radiation-resistant phenotype of the primitive SCs and the mechanism behind this resistance. Other radiation resistance mechanisms have been documented, such as increased scavenging of reactive oxygen species (ROS) in breast cancer stem cells and increased efficiency of DNA repair and DNA damage response in glioblastoma cancer stem cells (Bao *et al*, 2006; Woodward and Bristow, 2009; Diehn *et al*, 2009a). It remains to be seen whether those play an additional role in radiation resistance of prostate tumours.

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