Interaction between 3-Bromopyruvate and SC-514 in prostate cancer treatment

Abstract

Background: Prostate cancer (PCA) is the second most diagnosed cancer in men. The high incidence of prostate cancer has been attributed to failures in single treatment of chemotherapy. Failure of mono treatment is mediated by heterogeneity and plasticity of prostate cancer cells. 3-Bromopyruvate is one of the most common chemotherapeutic drugs used for the treatment of prostate cancer. However, its clinical therapeutic efficiency has been limited due to numerous side effects and drug resistance. SC-514 is a relatively new drug. Very little information exists on the anti-cancer effects of SC-514. Nevertheless, SC-514 might be able to overcome side effects of conventional chemotherapy. 3-BPA is strong potentiators of chemotherapeutic drug. 3-BPA has the potential to potentiate the anti-cancer activity of SC-514. The combination of 3-BPA and SC-514 might be able to inhibit prostate cancer carcinogenesis despite existence of heterogeneity and plasticity of prostate cancer cells.

Objective of study: This study aims to investigate the potential interaction between 3-BPA and SC-514 during treatment of prostate cancer.

Method: The bioassays used in this study include: trypan blue exclusion, MTT tetrazolium, NBT, LDH cytotoxicity, and poly caspase assay. Combination Index (CI) calculation was used to investigate the antagonistic, synergistic or additive interaction between 3-BPA and SC-514. One-way ANOVA was utilized to compare the cytotoxic effects of 3-BPA, SC-514 and the combination of 3-BPA and SC-514 on DU-145 cells and PC-3 cells prostate cancer cells.

Results: Results suggested a weak negative (r=-0.29) to moderate negative (r=-0.42) correlation between ROS released and cell death. In addition, there was a weak correlation (r=0.19) between percentage ROS induced and percentage apoptotic death. There was a positive correlation between the concentration of drug and cell death in DU-145 and PC-3 prostate cancer cells.

Conclusion: The overlap in mechanisms of action of 3-BPA and SC-514 increased the impact of SC-514 on prostate cancer cells. Hence, the combination of 3-BPA and SC-514 was more therapeutically effective (synergistic effect) than the single treatments of either 3-BPA or SC-514. The synergistic effect between 3-BPA and SC-514 did not occur by ROS induction. The apoptotic induction in DU-145 and PC-3 prostate cancer cells appears to occur via a mechanism other than reactive species (ROS) induction. This study suggests that the combination of 3-BPA and SC-514 as a therapeutic regimen can inhibit prostate cancer carcinogenesis effectively.

Introduction

Prostate cancer

Prostate cancer (PCAs) still remains a medical challenge to the world and it is becoming a common disease in the world. PCa account for 15% of all cancers in men worldwide. Prostate cancer is the most prevalent among the cancer related diseases and the second leading cause of cancer related death in USA. Prostate cancer poses a major economic limitation in humans. The economic limitation on patients makes prostate cancer one of the deadliest diseases. Patients with advanced stage of the disease have a low quality of life; they constantly visit the hospital for extremely expensive treatments. Regardless of the costly treatment of prostate cancer, there is life threatening transformations in the patient’s body system.

The life-threatening transformation in the patient’s system is because of uncontrollable division of the prostate cancer cells even after chemotherapy. Chemotherapy may be ineffective as a result of spontaneous accretion of tissues with polymorphous phenotype during treatment. This is illustrated in Figure 1A and 1B. The blue arrow represents epithelial cells and the red arrow represents fibroblastic cells. Biologically, prostate cancer cells possess six defining characteristic features that favor continuous proliferation. These features include invasion and metastasis, proliferative signaling, evasion of growth suppression, resistance to cell death, replicative immortality, and angiogenesis. These features differentiate them from healthy cells. Cancer researchers have made significant efforts to reduce deaths caused by prostate cancer. Conventional treatment modalities comprising radiotherapy, chemotherapy, gene therapy, immunotherapy, surgery and prostate specific membrane antigen (PSMA) targeted therapy have been developed to treat prostate and other human cancers. Unfortunately, the standard treatment
regimens frequently destroy healthy cells and thus causes considerable harmful side effects. Specific challenges faced by chemotherapeutic agents in cancer treatment include poor solubility, rapid deactivation, restricted bio-distribution, low therapeutic index, severe side effects, poor pharmacokinetic and poor pharmacodynamics performance.\textsuperscript{2,18--21} Studies have shown that androgen dependent prostate cancer can be treated with chemotherapeutic agents. However, androgen independent and metastatic prostate cancers are not responsive.\textsuperscript{22--24} In this perspective, the materialization of combination therapy provide a new opportunity for researchers to solve these shortcomings in conventional chemotherapy.\textsuperscript{25}

\textbf{Figure 2} A clear knowledge of the mechanism of ATP production in prostate cancer cells and normal prostate cells is important in achieving better treatment outcome in combination therapy.\textsuperscript{29 -- 40, 42 -- 58}
The mechanism of anticancer effects of 3-BPA

A potent anticancer agent known as 3-Bromopyruvate (3-BPA) has been reported to inhibit ATP production in prostate cancer (PCa) cells by targeting glycolysis, promoting mitochondrial destruction, and consequently increase oxidative stress. 3-BPA is an antineoplastic compound that targets both the "Warburg effect", (elevated glycolysis even in the presence of oxygen), as well as mitochondrial oxidative phosphorylation in cancer cells. 3-BPA is a lactic acid analog of private (the simplest of the alpha keto acids and intermediate in several metabolic pathways), that is transported through the same mono carboxylate transporters (MCT) as pyruvate. 3-BPA mimics lactic acid. Lactic acid is taken up by the cells’ lactate transporters and inhibits hexokinase. MCTs are involved in the efflux of lactic acid out of the cells. A study suggested that 3-BPA uptake is particularly effective because of the overexpression of MCTs in PCa cells. The entry of 3-BPA is successfully achieved because lactic acid and 3-BPA differ in only a single atom (Br), making it impossible for the cancer cells’ MCTs to distinguish between Br and O. Once inside the PCa cells, 3-BPA inhibits glycolysis (2 ATP production machinery). Normal cells are not inhibited by 3-BPA, as they have a deficiency of MCTs. This deficiency does not allow 3-BPA to enter the normal cells easily.

Furthermore, 3-BPA was shown to inhibit mitochondrial bound hexokinase II (HKII). The mitochondrial phosphate transporter, that is essential for ATP synthesis is also inhibited by compounds reacting with sulphydryl groups (-SH) such as 3-BPA. The second most important cellular target for 3-BPA is the Glyceraldehyde 3-phosphate dehydrogenase (GAPDH) enzyme. GAPDH like HK2 is also important for glycolysis. 3-BPA can compete with glucose in the first step of glucose conversion into glucose-6-phosphate via the enzyme hexokinase II. 3-BPA inactivates HK II such that glucose cannot be metabolized any further in the glycolytic pathway, causing inhibition of glycolysis. Detailed examination of 3-BPA treated cells showed release of cytochrome c which is an apoptotic marker indicating that there has been a disruption of the mitochondrial membrane. 3-BPA is a potent anti-glycolytic drug, able to induce severe ATP reduction and viability loss in many PCa cell lines; and it is less toxic to normal hepatocytes while depleting ATP in hepatocellular carcinoma (HCC) cells.

Over the past 16years 3-BPA have been frequently studied as a promising antitumor agent. 3-BPA regressed advance abdominal tumor. It eradicated xenograft tumors of HCC in all tested animals. There were no reported corrosive effects of 3-BPA. Furthermore, 3-BPA inhibits angiogenesis. 3-BPA induced the reversal of cancer cell chemo-resistance, where 3-BPA was reported to inhibit the efflux of chemotherapy through the ATP-binding cassette transporters; and antagonized the P-glycoprotein-mediated efflux in cancer cells. Multidrug resistance reversal, using 3-BPA might take place through decreasing ATP content in cancer cells, decreasing HK II activity, inhibiting ATPase activity, and reducing the expression of P-glycoprotein in chemo-resistant prostate cancer cells and thus results in a chemo-sensitization effect. Furthermore, 3-BPA has also been shown to inhibit cell viability by increasing reactive oxygen species (ROS) and inhibiting translation. The response to 3-BPA at 50mM and 100 mM was dose-dependent, leading to loss of ATP at 65% and 90% loss, respectively. It was reported in the study that 3-BPA treated cells revealed an increase in cellular ROS levels. The properties of 3-BPA that promote mitochondrial destruction and increase in ROS, highlight its potential benefits in treatment of carcinoma. 3-BPA impacts rapid depletion of glutathione (GSH) pool, and consequent increase in cell oxidative stress. Clinically, 3-BPA kills prostate cancer cells, prevents cancer recurrence, and reduces chemo-resistance and radio-resistance commonly encountered in clinical oncology. 3-BPA dramatically improve the therapeutic outcome of a patient having fibro lamellar hepatic carcinoma and/or metastatic melanoma. Significant improvements in late stage cancer patients suggest the potential efficacy of 3-BPA to differentiate between tumor and healthy tissues.

The mechanism of anticancer effects of SC-514

Not much is known about SC-514. SC-514 is an orally active, ATP-competitive inhibitor of nuclear factor kappa-B kinase subunit beta inhibitor (IKK-2 or IKKβ). SC-514 blocks nuclear factor kappa-light-enhancer of activated B cells (NF-κB)-dependent gene expression with IC-50 of 3-12nM. Previous studies have also reported the inhibitory effects of SC-514 on IKKβ in the treatment of tumors and inflammation. However, the effect of SC-514 on proliferation of PCa cell lines, multidrug resistance and RANKL-induced NF-κB signaling pathways is hitherto unknown. RANKL-inducing IKKβ inhibitor SC-514 enhanced nitrosourea-induced cell death in melanoma cells. SC-514 has been reported to be a selective IKKβ inhibitor and displayed >10-fold selectivity against
28 other kinases, including both tyrosine kinases and other serine-threonine kinases.\textsuperscript{71} Inhibition of NF-κB pathway may influence cell survival.\textsuperscript{72} Although kinases have a number of similarities, IKKβ has a 20- to 50-fold higher level of kinase activity for IkB than IKKa.\textsuperscript{73,74} Hence, this study investigates the impact of inhibitor of IKKβ on prostate cancer carcinogenesis. SC-514 dose-dependently inhibits RANKL-induced osteoclastogenesis with an IC-50 <5µM.\textsuperscript{20} SC-514 inhibits transcription of NF-kappa B-dependent genes in IL-1 beta-induced rheumatoid arthritis-derived synovial fibroblasts in a dose-dependent manner.\textsuperscript{89} High concentrations, SC-514(12.5 mM) induced apoptosis and caspase 3 activation in RAW 264.7 cells. Moreover, SC-514 significantly suppressed NF-kB activity owing to delayed RANKL-induced degradation of IkBα and inhibition of p65 nuclear translocation.\textsuperscript{90} Studies indicate that SC-514 impairs RANKL-induced osteoclastogenesis and NF-kB activation. Consistent with this observation, blocking IKKβ kinase activity by SC-514 decreases the phosphorylation of p65/RelA at Ser-536 in human T-cell lymphotrophic virus type I-transformed cells.\textsuperscript{44} Co-treatment of mouse skin with the IKKβ-specific inhibitor SC-514 (1µM) attenuated TPA-induced activation of Akt and NF-kappaB, and also the expression of COX-2 in hairless mouse skin.\textsuperscript{45} SC-514 is a reversible and highly selective inhibitor of IKK-2.\textsuperscript{82} SC-514 inhibited proliferation of DU-145 prostate cancer cells.\textsuperscript{83} SC-514 potentiated the effect of tyrosine receptor kinase (Axl) knockdown on proliferation (up to 10 fold) and apoptosis.\textsuperscript{82}

**Methods**

**Experiment 1:** MTT Tetrazolium assay was performed to assess the cell viability of the prostate cancer cells after treatment with 3-BPA and/or SC-514. Results from this experiment will answer the question “will the combination of 3-BPA and SC-514 synergistically reduced cell viability compared to the single treatment of 3-BPA or SC-514?”

Briefly, prostate cancer cells (DU-145 and PC-3) were seeded at a density of 2500 cells/well in 96-well plate. These cells were incubated at 37 °C and 5% CO2 for 48hrs (until they reach their log phase and 80-90% of confluence). These cells were treated with 3-BPA and/or SC-514 at their log phase for 48hrs. 20 µl of MTT (5 µg/ml) were added to the wells in the 96 well plate after the media was removed from the wells. The cells and MTT solution in the wells was incubated for 4hrs at 37°C. The yellow tetrazolium MTT (3-(4, 5-dimethylthiazol-2)-2, 5-diphenyltetrazolium bromide) (Invitrogen) was reduced by metabolically active prostate cancer cells after drug treatment (3-BPA, SC-514 and 3-BPA + SC-514), by the action of dehydrogenase enzymes, to generate reducing equivalents such as NADH and NADPH. The resulting intracellular purple formazan was solubilized by adding 50µl DMSO to each well and quantified by spectrophotometric means using ELISA plate reader (Biotek ELx800). The absorbance was measured at 570 nm. The absorbance values recorded was a measure of live cells in each well after drug treatment.

**Computation of the combination index for quantitative determination of drug interactions**

The percentage cell viability values from MTT Tetrazolium assay was used to determine drug interaction between 3-BPA and SC-514. The combination index values were used to quantify drug interactions between 3-BPA and SC-514. The classification of the interactions into categories of synergy, additivity, or antagonism was based on applying the formula in equation (1) and equation (2). Combination index (CI) analyses are widely used methods for evaluating drug interactions in combination cancer chemotherapy.\textsuperscript{83} The Loewe additivity model has been largely used as a reference model when the combined effect of two drugs is additive.

The model can be written as in Equation (1)

\[
CD_1/(Dx_1) + CD_2/(Dx_2) = 1
\]

where \(CD_1\) and \(CD_2\) are the respective combination doses of drug 1 (3-BPA) and drug 2 (SC-514) that yielded an effect of 50% growth inhibition, with \(Dx_1\) and \(Dx_2\), being the corresponding single doses for drug 1 (3-BPA) and drug 2 (SC-514) that resulted in the same effect, which is by definition the concentrations of drug 1 (3-BPA) and drug 2 (SC-514) that will impact 60% reduction in cell viability. When Equation 1 holds, it can be concluded that the combined effect of the two drugs is additive. Based on Equation 1, the combination index, defined in Equation 2, can be used to classify drug interactions as synergistic, additive, or antagonistic.

\[
CI = CD_1/(Dx_1) + CD_2/(Dx_2)
\]

A CI of less than, equal to, and more than 1 indicates synergy, additivity, and antagonism, respectively.\textsuperscript{83}

**Experiment 2:** To further evaluate the cytotoxic effects of 3-BPA and SC-514, we used Lactate Dehydrogenase (LDH) assay to quantify the LDH activity in the drug treated prostate cancer cells. This is a more sensitive assay to investigate the cell viability of the drug treated prostate cancer cells.

Lactate dehydrogenase (LDH) is a cytosolic enzyme present in prostate cancer cells. Plasma membrane damage releases LDH into the prostate cell culture media after treatment with 3-BPA and/or SC-514. Extracellular LDH in the media was quantified by a coupled enzymatic reaction in which LDH catalyzes the conversion of lactate to pyruvate via NAD+ reduction to NADH. Diaphorase then uses NADH to reduce a tetrazolium salt (INT) to a red formazan product that can be measured at 490nm. The level of formazan formation is directly proportional to the amount of LDH released into the medium, which is indicative of extent of cytotoxicity after 3-BPA and/or SC-514 treatment. Briefly, cultured prostate cancer cells (DU-145 and PC-3) were incubated with 3-BPA and/or SC-514 to induce cytotoxicity and subsequently release LDH. The LDH released into the medium was transferred to a new microtiter plate and mixed with reaction mixture. After a 30-minute room temperature incubation, reactions were stopped by adding stop solution. Absorbance at 490nm and 630nm were measured using a plate-reading spectrophotometer (Biotek ELx800) to determine LDH activity.

**Experiment 3:** After determining the cytotoxic effects of 3-BPA and/or SC-514. We initiated studies on the mechanism of action of 3-BPA and SC-514. This study indicated impairment in the biological activities of prostate cancer cells after drug treatment. Extremely low or high ROS levels in prostate cancer cells may be correlated to impaired cell functions. NBT assay was performed to measure the ROS level in the treated prostate cancer cells after treatment with 3-BPA and/or SC-514. This assay answers the question “will the combination of 3-BPA and SC-514 modulate ROS levels significantly compared to the single treatment of 3-BPA or SC-514?”

The bioactivity of 3-BPA and/or SC-514 generated Superoxide ions \(O_2^ {-}\). Superoxide ions \(O_2^ {-}\) converted NBT to NBT dimazon.
On the other hand, SOD reduces the O$_2^-$ concentration and thereby lowers the rate of NBT-diformazan formation. The extent of reduction in the appearance of NBT diformazan is a measure of SOD activity present in the drug treated prostate cancer cells (DU-145 and PC-3). Increase in absorbance reading from absorbance reader (Biotek ELx800) reflects an increased level of intracellular ROS in the drug treated prostate cancer cells.

**Experiment 4:** We further investigated the mode of cell death in 3-BPA and/or SC-514 treated prostate cancer cells by detecting caspase activity in the treated prostate cancer cells. Caspases activities are one of the indicators of apoptosis. Live red image-poly caspase detection assay was used to detect apoptosis in prostate cancer cells after treatment with 3-BPA and/or SC-514.

This assay is based on a fluorescent inhibitor of caspases (FLICA™) methodology, essentially an affinity label. The reagent associates a fluoromethyl ketone (FMK) moiety, which can react covalently with a cysteine, with a caspase-specific amine acid sequence. For poly caspases, this recognition sequence is valine-alanine-aspartic acid (VAD). A sulforhodamine group (SR) is attached as a reporter. The FLICA reagent is thought to interact with the enzymatic reactive center of an activated caspase via the recognition sequence, and then to attach covalently through the FMK moiety. The FLICA inhibitor is cell permeant and noncytotoxic. Unbound FLICA molecules diffuse out of the cell and are washed away; the remaining red-fluorescent signal is a direct measure of the amount of active caspase that was present at the time the inhibitor was added. The amount of caspase is an indication of the extent of apoptosis in the treated prostate cancer cells.

Briefly, 2500 cells/well were seeded in the 96 well microtiter plate and drug treatments were performed as described earlier. To estimate the number of cells with caspase activities, 100 cells were counted in five different fields under the fluorescence microscope. Percentage of cells showing caspase activities was calculated. The percentage of cells indicating caspase activity was plotted against the concentration of the drugs.

**Results and discussion**

**MTT Tetrazolium assay**

To evaluate the impact of 3-BPA and SC-514 on percentage cell viability of prostate cancer cells in vitro, PCa cells were treated with 3-BPA and/or SC-514. Results are shown in Figures 4A & 4B.

**Calculation of combination index in DU-145 Prostate cancer cells**

Equation (2)

$$CI = \frac{D_1}{Dx_1} + \frac{D_2}{Dx_2}$$

Substituting values extrapolated from Figure 5 for equation (2):

$$D_1 = 150 \mu M, (Dx)_1 = 250 \mu M, D_2 = 0.4 \mu M, (Dx)_2 = 1000 \mu M.$$ 

$$CI = \frac{150}{250} + \frac{0.4}{1000} = 0.6 + 0.0004 = 0.6004$$

Based on the standard that CI <=1 synergy; CI=1 additivity; CI > antagonism. Combination treatment using SC-514 and 3-BPA was synergistic because 0.6004<1.

**Calculation of combination index in PC-3 Prostate cancer cells**

Equation (2)

$$CI = \frac{D_1}{Dx_1} + \frac{D_2}{Dx_2}$$

Substituting values extrapolated from Figure 5 for equation (2):

$$D_1 = 110 \mu M, (Dx)_1 = 150 \mu M, D_2 = 31.25 \mu M, (Dx)_2 = 500 \mu M.$$ 

$$110/150 + 31.25/500 = 0.73 + 0.0625 = 0.8025$$

Based on the standard that CI <=1 synergy; CI=1 additivity; CI > antagonism. Combination treatment using SC-514 and 3-BPA was synergistic because 0.8025<1.

Interaction between 3-Bromopyruvate and SC-514 in prostate cancer treatment

NBT assay results showing treatment-induced inhibition of SOD/ROS production in DU-145 prostate cancer cells. Cells were treated as described earlier and subjected to the NBT assay for ROS determination. The results indicated no statistical differences (P > 0.05) between the different treatment regimens at all concentration points. The results/data points were the means of six independent experiments performed in triplicates.

Figure 5 (A) Comparison of LDH cytotoxicity and MTT cell viability of 3-BPA treated DU-145 prostate cancer cells. DU-145 prostate cancer cells (2500 cells per well) were plated in a 96-well plate in DMEM medium supplemented with 10% fetal bovine serum and incubated at 37°C and 5% CO₂. After 48hrs, varying concentrations of 3-BPA (0.488µM, 0.976µM, 1.953µM, 3.906µM, 7.812µM, 15.625µM, 31.25µM, 62.5µM, 125µM, 250µM, 500µM, 1000µM) were added to the 96 well plates and incubated for 48hrs at 37°C, and 5% CO₂. LDH Cytotoxicity was measured using the Pierce LDH cytotoxicity assay and cell viability measured by MTT cell proliferation assay. (B) Comparison of LDH cytotoxicity and MTT cell viability of SC-514 treated DU-145 prostate cancer cells. DU-145 prostate cancer cells (2500 cells per well) were plated in a 96-well plate in DMEM medium supplemented with 10% fetal bovine serum and incubated at 37°C and 5% CO₂. After 48hrs, varying concentrations of SC-514 (0.488µM, 0.976µM, 1.953µM, 3.906µM, 7.812µM, 15.625µM, 31.25µM, 62.5µM, 125µM, 250µM) were added to the 96 well plates and incubated for 48hrs at 37°C and 5% CO₂. LDH Cytotoxicity was measured using the Pierce LDH cytotoxicity assay and cell viability measured by MTT cell proliferation assay. (C) Comparison of LDH cytotoxicity and MTT cell viability of 3-BPA + SC-514 treated DU-145 prostate cancer cells. DU-145 prostate cancer cells (2500 cells per well) were plated in a 96-well plate in DMEM medium supplemented with 10% fetal bovine serum and incubated at 37°C and 5% CO₂. After 48hrs, varying concentrations of SC-514 (0.488µM, 0.976µM, 1.953µM, 3.906µM, 7.812µM, 15.625µM, 31.25µM, 62.5µM, 125µM, 250µM, 500µM, 1000µM) + 110µM 3-BPA (IC-50) were added to the 96 well plates and incubated for 48hrs at 37°C and 5% CO₂. LDH Cytotoxicity was measured using the Pierce LDH cytotoxicity assay and cell viability measured by MTT cell proliferation assay. (D) Comparison of LDH cytotoxicity and MTT cell viability of 3-BPA treated PC-3 prostate cancer cells. PC-3 prostate cancer cells (2500 cells per well) were plated in a 96-well plate in DMEM medium supplemented with 10% fetal bovine serum and incubated at 37°C and 5% CO₂. After 48hrs, varying concentrations of 3-BPA (0.488µM, 0.976µM, 1.953µM, 3.906µM, 7.812µM, 15.625µM, 31.25µM, 62.5µM, 125µM, 250µM, 500µM, 1000µM) were added to the 96 well plates and incubated for 48hrs at 37°C, and 5% CO₂. LDH Cytotoxicity was measured using the Pierce LDH cytotoxicity assay and cell viability measured by MTT cell proliferation assay. (E) Comparison of LDH cytotoxicity and MTT cell viability of SC-514 treated PC-3 prostate cancer cells. PC-3 prostate cancer cells (2500 cells per well) were plated in a 96-well plate in DMEM medium supplemented with 10% fetal bovine serum and incubated at 37°C and 5% CO₂. After 48hrs, varying concentrations of SC-514 (0.488µM, 0.976µM, 1.953µM, 3.906µM, 7.812µM, 15.625µM, 31.25µM, 62.5µM, 125µM, 250µM, 500µM, 1000µM) + 110µM 3-BPA (IC-50) were added to the 96 well plates and incubated for 48hrs at 37°C, and 5% CO₂. LDH Cytotoxicity was measured using the Pierce LDH cytotoxicity assay and cell viability measured by MTT cell proliferation assay. (F) Comparison of LDH cytotoxicity and MTT cell viability of SC-514 treated PC-3 prostate cancer cells. PC-3 prostate cancer cells (2500 cells per well) were plated in a 96-well plate in DMEM medium supplemented with 10% fetal bovine serum and incubated at 37°C and 5% CO₂. After 48hrs, varying concentrations of SC-514 (0.488µM, 0.976µM, 1.953µM, 3.906µM, 7.812µM, 15.625µM, 31.25µM, 62.5µM, 125µM, 250µM, 500µM, 1000µM) + 150µM 3-BPA (IC-50) were added to the 96 well plates and incubated for 48hrs at 37°C, and 5% CO₂. LDH Cytotoxicity was measured using the Pierce LDH cytotoxicity assay and cell viability measured by MTT cell proliferation assay.

NBT assay measuring the ROS level

Figure 6 (A) NBT assay results showing treatment-induced inhibition of SOD/ROS production in DU-145 prostate cancer cells. Cells were treated as described earlier and subjected to the NBT assay for ROS determination. The results indicated no statistical differences (P > 0.05) between the different treatment regimens at all concentration points. The results/data points were the means of six independent experiments performed in triplicates. Figure 6 (B) NBT assay results showing treatment-induced inhibition of SOD/ROS production in PC-3 prostate cancer cells. Cells were treated as described earlier and subjected to the NBT assay for ROS determination. The results indicated statistical differences (P < 0.05) between the different treatment regimens at all concentration points. The results/data points were the means of six independent experiments performed in triplicates.


Lactate dehydrogenase (LDH) assay
Live red image-poly caspase detection assay in PC-3 cells

Image-iT™ LIVE Red Poly Caspases Detection in DU-145 cells (Figure 7C)

This study suggests that 3-BPA and/or SC-514 treatment can inhibit prostate cancer carcinogenesis. The combination treatment of 3-BPA and SC-514 was more effective in inhibiting prostate cancer cell proliferation compared to the monotreatment of either 3-BPA or SC-514 as shown in Figure 4A and Figure 4B. Figure 4A shows a dose-dependent effect for the combined drug treatment, while Figure 4B shows no clear dose-dependence. There is a possible explanation for this observation. The cell line used in figure 4B is DU-145 prostate cancer cell line while the cell line used in figure 4A is PC-3 prostate cancer cell line. DU-145 prostate cancer cell has an active nuclear factor kappa light chain enhancer of activated B cells (NF-KB) receptor, which is not as active in PC-3 prostate cancer cells. The NF-KB receptor in DU-145 prostate cancer cells could be interfering with the activity of the drugs (3-BPA and SC-514) in the combination treatment. This is highly possible since SC-514 works by inhibiting NF-KB. Overexpression of NF-KB receptor in DU-145 prostate cancer cells potentially hinders/impacts the mechanism of action of SC-514 drug in DU-145 prostate cancer cells.

The interaction between 3-BPA and SC-514 was synergistic in DU-145 and PC-3 prostate cancer cells as calculated from Loewe additivity model and combination index. Comparison of LDH cytotoxicity and MTT cell viability results in figures 5A-5F further confirm the cytotoxicity effect of 3-BPA and/or SC-514 on prostate cancer cells. The result show that as cell proliferation decreased the quantity of lactate dehydrogenase (LDH) released into the culture media from damaged prostate cancer cells increased at most treatment.

concentrations. The LDH released serve as a biomarker for cellular cytotoxicity and cytolyis of the drug treated DU-145 and PC-3 prostate cancer cells. \cite{38,39} The combination of these two assays provide a safe alternative to radioactive cytotoxicity assays.

LDH cytotoxicity level in DU-145 cells treated with combination of 3-BPA and SC-514 (Figure 5C) did not impact the highest level of cytotoxicity at all concentrations when compared to the single treatment of 3-BPA (Figure 5A) or SC-514 (Figure 5B). Similarly, LDH cytotoxicity level in PC-3 cells treated with combination of 3-BPA and SC-514 (Figure 5F) did not impacted highest level of cytotoxicity at all concentrations when compared to the single treatment of 3-BPA (Figure 5D) or SC-514 (Figure 5E). This observation suggests that there are other mechanisms of actions used by these 3-BPA and/or SC-514. These undiscovered mechanisms may not involve destruction of cell membrane and leakage of lactate dehydrogenase enzymes to the extra-cellular spaces. For example, shrinking of the prostate cancer cells in manner that allow other cell contents to be released more than the release of lactate dehydrogenase enzymes. The synergistic effect observed earlier could have occurred as result multiple mechanisms that are not fully understood. Future studies will use other cell viability assays such as alamar Blue and cell titer glo to confirm this suggestion.

MTT cell viability in DU-145 prostate cancer cells indicated the lowest level at combination treatment of 3-BPA and SC-514 (Figure 5C) when compared to the single treatment of 3-BPA (Figure 5A) or SC-514 (Figure 5B). Similarly, MTT cell viability in PC-3 prostate cancer cells indicated the lowest level at combination treatment of 150µM 3-BPA and 1000µM SC-514 (Figure 5F) when compared to the single treatment of 1000µM 3-BPA (Figure 5D) or 1000µM SC-514 (Figure 5E). However, the over all trend of cell viability in PC-3 prostate cancer cells do not show that the combination treatment of 3-BPA and SC-514 impacted the lowest level of cell viability at all concentrations. For example, the combination treatment of 150µM 3-BPA and 125µM SC-514 in PC-3 recorded higher cell viability when compared to the single treatment of 125µM 3-BPA or 125µM SC-514. This may be as result of resistance to the combination treatment at 125 µM concentration.

Results from the NBT assay suggests that ROS induction only cannot account for the mechanism of action of 3-BPA and/or SC-514. ROS modulation in DU-145 and PC-3 prostate cancer cells indicated a weak correlation between ROS released and concentration of 3-BPA and/or SC-514 (Figure 6A & Figure 6B). However, SC-514 monotreatment induced elevated level of ROS at 125µM concentration and higher concentrations. Oxidative stress in cells could mean that the ROS is too low or too high. In this study, 3-BPA at low concentrations (0.48µM–62.5µM) appears to work by increasing the ROS levels,\cite{40,41} but 3-BPA at higher concentrations (62.5µM-1000µM) appears to work by decreasing the ROS levels.\cite{42} On the other hand, higher concentrations of SC-514 (125µM) appear to upregulate ROS levels. This conflicting effect between 3-BPA and SC-514 explains why the combination treatment (SC-514 + 3-BPA) gave a lower ROS level compared to the single treatment with either 3-BPA or SC-514. The ROS modulation result shows that the interaction between the 3-BPA and SC-514 is not through ROS only. The synergistic effect between 3-BPA and SC-514 could be through a mechanism that is not directly linked to ROS induction. Future studies will elucidate the mechanism(s) of action that may be involved. Survival pathways such as NF-KB pathway, IL-6 pathway and AKT-p13 pathway can be investigated to further understand the mechanism of action of 3-BPA and/or SC-514.

It appears that combination of 3-BPA and SC-514 was more effective than SC-514 monotreatment in inducing apoptosis in DU-145 and PC-3 prostate cancer cells (Figure 7A–7C). But the combination treatment of 3-BPA and SC-514 did not induce apoptosis more than 3-BPA monotreatment at most concentration in DU-145 and PC-3 prostate cancer cells (Figure 7A–7C). 3-BPA appears to induce more apoptosis than SC-514 in DU-145 and PC-3 cells at most concentrations as indicated by caspase activation assay (Figure 7A–7C). However, other modes of cell death may exist with SC-514 drug treatment that may make SC-514 therapeutically more effective than 3-BPA. Future studies will investigate modes of cells deaths such as necrosis, autophagy, necroptosis, and ferroptosis in prostate cancer cells treated with 3-BPA and/or SC-514.\cite{43}

The existence of multidrug resistance (MDR) may influence the mode of cell deaths or survival pathways in prostate cancer cells as mentioned above. Metabolic activities of prostate cancer cells favor the sustained production of ATP energy required for MDR (Figure 2). Specifically, catabolic activity in prostate cancer cells produces ATP from break down of fats proteins and carbohydrate (Figure 3). MDR occurs when ATP Binding Cassette (ABC) proteins use the ATP energy produced to pump drug from inside of the prostate cancer cells to the extracellular surroundings of the prostate cancer cells. Future studies will investigate the role of BCL2 family on ABC proteins-mediated MDR during the treatment of prostate cancer with 3-BPA and/or SC-514.

Furthermore, drug delivery of 3-BPA and/or SC-514 will be explored for better therapeutic efficiency, reduction of MDR and better treatment outcome because nanoparticle drug delivery systems (NDDS) are effective in delivering anti-cancer drugs such as 3-BPA and/or SC-514 to prostate cancer.\cite{44,45}

Conclusion

In conclusion, combination treatment remains a viable option when conventional monotherapy fails. Results from combination of 3-BPA and SC-514 in prostate cancer therapy will potentially encourage further combination possibilities between other chemotherapeutic drugs.

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Conflict of interest

Author declares that there is no conflict of interest.

References

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