Mild antagonistic effect of Valproic acid in combination with AZD2461 in MCF-7 breast cancer cells

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Abstract

Background: Breast cancer (BC) is a complex disease, but current treatments are not efficient enough considering increased relapse and decreased survival rate among patients. Poly (ADP-ribose) polymerase inhibitors are recently developed anticancer agents which target cells with defects in homologous recombination (HR) pathway. This study wishes to assess whether the combination of AZD2461 as a newly developed PARP1 inhibitor and valproic acid (VPA), a histone deacetylase inhibitor could effectively reduce the growth of MCF-7 cells with no fundamental DNA repair defect.

Methods: Both trypan blue dye exclusion assay and MTT viability test were used to evaluate cell death. γ-H2AX levels, as a marker of DNA repair, were measured using in cell ELISA method. The Student’s t-test and non-parametric analysis of variance (ANOVA) were applied for our data analyses where p-value <0.05 was considered statistically significant.

Results: As calculated by CompuSyn software, IC50 values for VPA and AZD2461 were 4.89 mM and 42.83 µM respectively following 48 hours treatment. Also, the trypan blue exclusion assay results showed a concentration- and time-dependent decrease when MCF-7 cells were treated with both agents (p<0.05). Combination analysis showed a mild antagonism (CI>1.1) while γ-H2AX levels found not to be significantly increased in MCF-7 cells co-treated with VPA+AZD2461 compared to each agent alone (p=0.29).

Conclusion: Our findings revealed that the combination of VPA and AZD2461 could decrease cell viability of MCF-7 cells, but it was not able to significantly increase unrepaired DNA damage sites. The mechanism responsible for drugs combination was not of synergism or addition. Determining the type of involved cell death mechanisms might be followed in further studies.

Keywords: Breast cancer, Valproic acid, Combination therapy, AZD2461

Introduction

Breast cancer (BC) is the most frequent female tumor worldwide (1). The range of mortality rates due to this
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Type of cancer is approximated up to 6-19 per 100,000 (2). Recent studies have shown that BC associated mortalities in industrialized countries (i.e., the United States) has been decreased but in developing countries, the disease is still on the rise, especially in Iran. It is also estimated that BC incidence among Iranian females is almost 10 years earlier than their western counterparts (3). Depending on subtype and stage of BC, there are various strategies to treat it, including Surgery, hormone therapy, radiation therapy, chemotherapy and finally targeted therapy and many patients get more than one type of treatment for the enhancement of their survival (4). There is an increasing number of genes that increase the risk of developing BC when they carry a mutation, such as *CTLA4, CYP19A1, FGFR2, ATM, BARD1, BRIP1, CASP8*, and *TERT*(5). Accordingly, one of the crucial reasons for the failure of cancer treatment methods is the presence of one or more of these mutations (6). About 5% to 10% of breast cancer patients have a mutation in such important genes (7). For example, *BRCA1* (breast cancer responsibility gene 1) and *BRCA2* (breast cancer responsibility gene 2) are two key genes which their product is involved in repair cell damage response (8) but many tumor cells derived BC cells do not carry such (i.e., MCF-7 cell line) mutations. Advanced Breast cancers are often attributed to defects in DNA damage response (9), and unrepaired DNA lesions can lead to cancer development and progression (10). Poly (ADP-ribose) polymerases (PARPs) is an enzymatic protein family which is responsible for poly (ADP-ribose) polymerization and transfer of ADP-ribose to target proteins at the sites of DNA damage (11). Their roles in DNA repair, programmed cell death (PCD), and genomic stability has been well established (12). Contributing to repair of single- or double strand DNA breaks (SSBs and DSBs respectively) is considered to be the most critical functions of this enzyme family consisting of 17 members (13). Recent studies on the field of drug development are focused on using PARP inhibitors (PARPis) in order to perturbate the DNA damage response. Currently, clinical trials suggest that PARP inhibitors may be useful in the treatment of BRCA-related and triple-negative breast cancer (14). Rucaparib (PF-01367338), Niraparib (MK-4827) and Olaparib (AZD-2281) are three classical PARPi which approved by FDA while others are still in early or mid-phases of clinical trials (15). *AZD2461* is a novel PARP1, PARP2 and PARP3 inhibitor which initially developed by AstraZeneca but its phase I clinical studies for treating solid malignancies was terminated in 2011 due to efficacy problems (16). As a novel and well tolerated structural analogue for olaparib, *AZD2461* displayed high antitumor activity in BC patients (17).

On the other hand, the co-treatment of cancer cells with histone deacetylases (HDACs) inhibitors showed promising advances in the field of cancer therapies (18). Histone deacetylases (which consist of HDAC1 to HDAC11 and sirtuins) are a family of enzymes that remove acetyl groups from histones, allowing them to wrap the DNA tightly (19). HDACs inhibitors (HDACis) selectively inhibit HDACs, lead to reduce chronic inflammation and induction of apoptosis in the intestine and colorectal cancer (20). Valproic acid (VPA) is a member of HDAC inhibitors that proved efficacious in inhibiting the growth of breast cancer cells alone or in combination with ionizing radiation (IR) in order to sensitize the tumor cells to cell death-inducing agents (21). Recent experiments unraveled the synergism relationship between HDACis and Poly (ADP-ribose) polymerase (PARP) by suppressing cell growth *in vitro*. For example, triple-negative breast cancers (TNBCs) are conceived to be resistant to PARP inhibitors, but HDAC inhibitors could sensitize TNBC cells to olaparib by down-regulating proteins involved in DNA repair pathways (22). The phosphorylation of histone H2AX is the first event in response to DNA damage by HDACs and PARPs in cancer cells (23). Hence, the detection of phosphorylated H2AX can potentially assist the transformation of non-tumor cells to malignant cells.

We hypothesized that VPA and AZD2461 together could inhibit the growth of MCF-7 breast cancer cells with no hampered DNA-repair capacity. The current study is conducted to investigate whether the combined use of these inhibitors could improve their antiproliferative and DNA repair efficacy in MCF-7 cell, not carrying such mutations in their profile of DNA repair genes.

**Methods**

**Chemicals, Cell line and Culture Methods**

MCF-7, human breast (adenocarcinoma) cell line was sourced from Pasteur Institute of Iran and were grown in RPMI 1640 medium obtained from INOCLON (G. Innovative Biotech Co (INOCLON), Iran) supplemented with 10% FBS (Gibco, Rockville, MD, USA), antibiotic-antimycotic solution (containing 100 mg/ml of penicillin, 2.5 mg/L of amphotericin B, and 100 U/ml of streptomycin. 0.25% Trypsin-EDTA solution (G. Innovative Biotech Co (INOCLON), Iran) was used to detach cells from the surface. AZD2461, VPA, TWEEN 20, Trypan blue and Triton X-100 was procured from Sigma-Aldrich (St. Louis, MO, USA). Cell culture flasks and microtiter plates were supplied by Biofill (Jet Biofill, China). All other chemicals were of certified reagent analytical grade.

**MTT Assay**

For evaluation of the viability of MCF-7 cells in response to both drugs alone or combined, MTT assay was performed (25). At first, 6000 cells/well were seeded. Af-
after one day, cells were exposed to AZD2461 and VPA at the concentration ranging from 0.5mM to 16mM for VPA and 5μM to 160μM for AZD2461. After 24, 48 and 72 hours incubation, 20μL of 5 mg/ml tetrazolium dye was added to each micro-well and cells were incubated for 3 minutes. Then, the culture was replaced by 200μL of DMSO and kept for 20 minutes in the dark. Using a STAT FAX 2100 microplate reader, the absorbance at 570 nm wavelength was calculated. The viability percentage was calculated as below mentioned.

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\text{Mean OD treatment/Mean OD control } \times 100 = \% 
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**Trypan Blue Dye Exclusion Assay**

The trypan blue dye exclusion assay is used to find the number of viable cells in a cell suspension (26). Firstly, 50μL of cell suspension was added into a cryo-vial. Then, an equal volume of 0.4% trypan blue dye was placed to the cell suspension and mixed. Mixture Incubated for at least 3 minutes at room temperature. With the coverslip, one side of a hemacytometer counter filled with the cell suspension and using a conventional light microscope, coloured (death) cells were detected, and therefore the percentage of viable cells was calculated.

**H2AX In Cell Elisa Assay**

We used the DuoSet IC ELISA kit (R&D Systems, Minneapolis, USA) according to the kit manual to measure phosphorylated H2AX in cell lysates (27). Briefly, the captured antibody was diluted in PBS and immediately coated to a 96 well microplate (100 μL/well) and incubated 24 hours at room temperature. Following 3 times washing, 300 μL of block buffer was added and incubated at room temperature for 90 minutes. Then 100 μL of sample or standard in IC Diluent added and incubated 2 hours at the same temperature. Then, washing steps were repeated and by placing 100 μL of the diluted detection antibody into each well, the plate was again incubated 2 hours at room temperature. Next, 100 μL of the diluted Streptavidin-HRP placed into each well before incubating for 20 minutes at the mentioned temperature. In next step after washing, 100 μL of substrate solution was added, and after 20 minutes, 50 μL of stop solution added to each micro-well, and the optical density (OD) of each well was directly measured at 450nm. The fold changes were calculated by dividing absorbance values (OD) of treated cells by the OD values of adjusted untreated cells in a time-dependent manner.

**Data Analysis and Statistical methods**

Using SPSS16 software for Windows (release 16, SPSS Inc., Chicago, Illinois), the Student’s t-test and non-parametric analysis of variance (ANOVA) were performed for our data analyses when appropriate. In all assays, p-value <0.05 was considered significant.

**Analysis of Drug Combination**

MCF-7 cells were exposed to both agents (VPA: 0.15mM - 20mM and AZD2461: 1.5μM - 200μM) with the combined ratio of 1:115, each diluted 1:2. The drug interactions between constant ratios of both inhibitors were calculated using CompuSyn software (Version 1.0, Combo-Syn In., US) based on Chou-Talalay principles for drugs combination (28) where CI (combination index) values indicate the mechanism of drugs interaction. Briefly, CI > 1.1, CI = 0.9–1.1, CI < 0.9 represent antagonism, an additive effect, and synergism respectively. The concentration-effect relationships for this combination regimen were plotted using the Median-Effect principles (29).

**Results**

**Anti-proliferative Effects of VPA, AZD2461 and Their Combination on MCF-7 Cells**

As shown in (Figs. 1A and 1B), both agents diminished cell viability of MCF-7 cells in concentration and time-dependent manners which was in agreement with the results of trypan blue dye exclusion assay (Figs. 2A and 2B). VPA significantly decreased the number of viable cells following 24 hours (p=0.005), 48 hours (p=0.002), 72 hours (p=0.003).
and 72 hours (p=0.001) of treatment while these p-values were p=0.008, p=0.006, and p=0.003 in cells exposed to AZD2461 after the same periods, respectively.

Analyses of drug combination using combination index and median-effect plots indicate that combination of these two agents would instead exert mild antagonism and not benefits from additive or synergistic interactions (CI>1.1) (Figs. 2D, E). Also, various effective doses (EDs) of VPA and AZD2461 showed CI values higher than 1.1 (Fig. 2F) which indicates the type of drug relationship based on different mechanisms of action. Therefore, the co-treatment of MCF-7 cells with both agents was not sufficiently able to suppress cell proliferation. Table 1 shows the IC50 values calculated by CompuSyn software for both agents alone or in combination.

Effects of Combination with VPA and AZD2461 on γ-H2AX Levels of MCF-7 Cells

As shown in Figure 3, following 24 h treatment, the levels of γ-H2AX were significantly increased up to 1.9, 2.8 and 1.7 fold compared to adjusted untreated cells when treating MCF-7 cells with VPA, AZD2461, and their combination, respectively; but co-treatment of two agents did not significantly increase phosphorylated H2AX levels compared to each drug alone (p=0.290).

Discussion

Drug combination has proved to be one potential strategy towards increasing therapeutic index and overcome drug resistance against tumor cells (30). Although it is well established that BC cells, could be more responsive to a combination of HDACi and PARPi, compared to each agent alone (31), our information concerning this procedure and possible drug interaction effects still remained incomplete. So far, several studies examined the combination of VPA with another agent in cancer therapy (32-34). Otherwise, some studies have concluded that VPA is capable of inhibiting HDACs (35). Kuendgen indicated VPA combined with all-trans retinoic acid synergistically suppressed tumor cells growing in patients with acute myeloid leukemia (36). Rottenberg found that olaparib has efficacy against BRCA1-deficient breast cancer in combination with platinum drugs or when used as a mono-
therapeutic agent (37). O’Connor and colleagues investigate the efficacy of AZD2461 against olaparib-resistant tumors that overexpress P-glycoprotein, suggesting that use of this specific PARPi-3 inhibitor may have benefits in terms of causing lower drug resistance than olaparib in vitro (38). Konstantinopoulos and colleagues investigated the effects of suberoylanilide hydroxamic acid (SAHA) as an HDACi and olaparib in ovarian cancer cells harboring mutations in BRCA. Based on their findings, SAHA combined with olaparib induced higher apoptosis rates and H2AX foci formation than each other drug alone (39). Phosphorylated H2AX (γ-H2AX) is a marker of unrepaired DNA damage that found to be decreased in cells treated with PARPi (40). Also, VPA and PARPi lead to a decrease in mRNA levels of some important DNA repair genes (41).

In the current study, we investigated the combined effects of VPA and AZD2461 in MCF-7 BC cell line besides measuring phosphorylated H2AX in cell lysates as a marker of DNA damage response perturbation. While many prior studies discovered a synergistic relationship in case of combining other HDACi and inhibitors of PARP family members (42, 43), our results did not indicate the probable interaction between VPA and AZD2461. Our data regarding the quantification of phosphorylated H2AX in MCF-7 cells was also indicative of such mentioned drug interaction. In an experiment conducted by Ha et al. (2014), the synergistic lethal effect of combining ABT-888, a PARPi, in combination with pan-histone deacetylase inhibitor (HDi) and cisplatin was evaluated in human triple negative breast cancer (TNBC) cells (24). The outcomes of this study were in agreement with our findings, suggesting that such co-treatment strategies could prove efficacious in MCF-7 cells irrespective of BRCA1 expression status. This matter is important since a majority of TNBC cells lack germ-line mutations of HR-related factors. To best of our knowledge, the combination of these two drugs in this BC cell line with this genetic profile did not study before since AZD2461 is known as a recently designed analogue for olaparib. Considering the novelty of combining AZD2461 with other HDACis or classical chemotherapeutic agents, different combinations of strategies may be followed. Our findings support the rationale to use this novel combination against human BC cells and other malignancies.

**Conclusion**

We concluded from our data that although combination regimen of VPA and AZD2461 could decrease cell viability of MCF-7 cells, it was not able to significantly increase unrepaired DNA damage sites and the mechanism responsible for drugs combination was not of synergism or addition. Assessment of other cell death markers, flow-cytometric analyses of cell death and evaluating the relative expression of main DNA repair genes and proteins might be followed in further experiments.

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**Conflict of Interests**

The authors declare that they have no competing interests.

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