

Ataxia Telangiectasia Mutated and Rad3 Related (ATR) Protein Kinase Inhibition Is Synthetically Lethal in XRCC1 Deficient Ovarian Cancer Cells

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Abstract

Introduction: Ataxia telangiectasia mutated and Rad3 Related (ATR) protein kinase is a key sensor of single-stranded DNA associated with stalled replication forks and repair intermediates generated during DNA repair. XRCC1 is a critical enzyme in single strand break repair and base excision repair. XRCC1-LIG3 complex is also an important contributor to the ligation step of the nucleotide excision repair response.

Methods: In the current study, we investigated synthetic lethality in XRCC1 deficient and XRCC1 proficient Chinese Hamster ovary (CHO) and human ovarian cancer cells using ATR inhibitors (NU6027). In addition, we also investigated the ability of ATR inhibitors to potentiate cisplatin cytotoxicity in XRCC1 deficient and XRCC1 proficient CHO and human cancer cells. Clonogenic assays, alkaline COMET assays, γ H2AX immunocytochemistry, FACS for cell cycle as well as FITC-annexin V flow cytometric analysis were performed.

Results: ATR inhibition is synthetically lethal in XRCC1 deficient cells as evidenced by increased cytotoxicity, accumulation of double strand DNA breaks, G2/M cell cycle arrest and increased apoptosis. Compared to cisplatin alone, combination of cisplatin and ATR inhibitor results in enhanced cytotoxicity in XRCC1 deficient cells compared to XRCC1 proficient cells.

Conclusions: Our data provides evidence that ATR inhibition is suitable for synthetic lethality application and cisplatin chemopotential in XRCC1 deficient ovarian cancer cells.

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Introduction

Targeting DNA repair for synthetic lethality is an exciting new strategy for personalized therapy in ovarian cancer. DNA repair is essential for processing DNA damage induced by chemotherapy such as platinating agents (carboplatin, cisplatin) [1]. Intra-strand crosslink DNA adducts induced by platinating agents, if unrepaired, ultimately result in cell death [2,3]. DNA intra-strand crosslinks are repaired predominantly by nucleotide excision repair (NER) in cells [4,5]. Platinating agents can also generate oxygen free radicals that induce oxidative base damages that are processed by the DNA base excision repair (BER) pathway in cells [6,7].

The XRCC1 (X-ray repair cross-complementing gene 1) protein is a critical factor in BER and single strand break repair pathway (SSBR). XRCC1-LIG3 complex is also an important contributor to the ligation step of the nucleotide excision repair (NER) response. XRCC1, a 70-kDa protein, has no known enzymatic activity (reviewed in [8,9,10]). XRCC1 functions as a

molecular scaffold protein and coordinates DNA repair by interacting with several components of BER/SSBR such as PARP-1 [Poly(ADP-ribose)polymerases 1], DNA glycosylases, AP endonuclease (APE1) and others (reviewed in [8,9,10]). XRCC1 deficiency in cells lead to accumulation of DNA single strand breaks (SSBs), induce mutations and result in elevated levels of sister chromatid exchanges. XRCC1 deficiency in cell lines result in hypersensitivity to ionizing radiation and chemotherapy [9]. In human association studies, germline polymorphisms in XRCC1 may influence cancer risk [11,12] and influence response to platinum based chemotherapy [13,14,15,16]. In human ovarian cancer we have recently demonstrated that tumours frequently over-express XRCC1 (48%) and significantly associated with higher stage ($p = 0.006$), serous type tumours ($p = 0.008$), sub-optimal de-bulking ($p = 0.004$), a two fold increase of risk of death ($p = 0.007$) and progression ($p < 0.0001$) [17]. In the multivariate analysis, XRCC1 expression was independently associated with survival in ovarian cancer patients [HR 2.3, $p = 0.002$]. XRCC1 negative tumours were associated with platinum sensitivity

($p < 0.0001$). Pre-clinically we also confirmed that XRCC1 negative cells are hypersensitive to cisplatin compared to XRCC1 positive cells [17]. Hypersensitivity to cisplatin in XRCC1 negative cells was associated with accumulation of DNA strand breaks and G2/M cell cycle arrest [17]. Our data therefore suggests that XRCC1 is a promising biomarker in ovarian cancer.

Ataxia telangiectasia mutated and Rad3 Related (ATR) protein kinase is a key sensor of single-stranded DNA associated with stalled replication forks as well as generated during BER and double strand break repair as DNA repair intermediates. Activated ATR in turn phosphorylates a number of substrates involved in cell cycle regulation, DNA replication, DNA repair and apoptosis (reviewed in [18,19,20,21,22]). In preclinical studies, ATR inhibition may result in cytotoxic therapy sensitization [22,23,24]. Small molecule inhibitors of ATR are currently under development for therapeutic application in cancer [20,21,22].

The ability of PARP inhibitors to induce synthetic lethality in BRCA deficient ovarian cancers [25,26,27] suggests that additional factors within BER/SSBR may be suitable for such personalized approaches. XRCC1 is a critical factor in BER, SSBR and NER. ATR is a key sensor of SSBs. In the current study we have investigated and confirmed synthetic lethality in XRCC1 deficient cells treated with ATR inhibitors. Moreover, compared to cisplatin alone, combination of cisplatin and ATR inhibitor treatment results in enhanced cytotoxicity in XRCC1 deficient cells compared to XRCC1 proficient cells.

Materials and Methods

Compounds and Reagents

Small molecule ATR inhibitors NU6027 and VE-821 were purchased from Tocris Bioscience, UK and Tinib-Tools, Czech Republic respectively. The compounds were dissolved in 100% DMSO and stored at -20°C . Cisplatin (1 mg/ml) was obtained from the Department of Pharmacy, Nottingham University Hospitals, UK.

Cell Lines and Culture

Previously well characterized Chinese hamster ovary (CHO) cells; CHO9 (Wild type), EM-C11 (XRCC1-mutant: C389Y substitution leading to XRCC1 protein instability), EM-C12 (XRCC1-mutant: E98K substitution resulting in reduced XRCC1 protein integrity) [28] were provided by Professor Małgorzata Z. Zdżienicka, Department of Molecular Cell Genetics, Nicolaus-Copernicus University in Torun, Bydgoszcz 85-094, Poland. Cells were grown in Ham's F-10 media (PAA, UK) [supplemented with 10% fetal bovine serum (FBS) (PAA,UK) and 1% penicillin/streptomycin]. EM9-V (XRCC1 mutant) and EM9 cells stably transfected with a human XRCC1 expression vector (EM9-XH cells) [29] were provided by Professor Keith Caldicott, Genome Damage and Stability Centre, University of Sussex, UK. Cells were grown in DMEM media (PAA, UK) [supplemented with 10% fetal bovine serum (FBS) (PAA,UK) and 1% penicillin/streptomycin]. Ovarian cancer cells OVCAR-3 and OVCAR-4 were grown in RPMI media (PAA, UK) [supplemented with 10% fetal bovine serum (FBS) (PAA,UK) and 1% penicillin/streptomycin].

XRCC1 Knockdown Using siRNAs

Three XRCC1 siRNA constructs (sequences listed in Table 1) and one negative scrambled control and siRNA for Glyceraldehyde 3-phosphate dehydrogenase (GAPDH) (positive control) were used in these studies. The siRNA constructs were purchased from Ambion life technologies, UK. The transfection protocol was as

described previously by Fan et.al [30]. Cells were plated in 6-well plates (2 ml medium/well without antibiotics). At 50% confluence, transfection was achieved using Lipofectamine TM 2000 (Invitrogen) according to the manufacturer's protocol. Briefly, siRNA (100 pmol) and Lipofectamine (5 μl) were each separately mixed with 250 μl Opti-MEM1 (GIBCO/Invitrogen) without FBS. After 5 minute incubation at room temperature, the siRNA and Lipofectamine solutions were combined and incubated for another 20 min at room temperature. This mixture was then added to plated cells, cultured at 37°C overnight and the medium was later replaced with fresh medium plus penicillin/streptomycin (1%). When the cells attained 100% confluence, they were trypsinized and subsequently transferred into 75 cm^2 flasks for continued growth and/or treatment. XRCC1 Knockdown was evaluated by western blotting at various time points after transfection (days 3, 5 and 7).

Western Blot Analysis

Protein samples were prepared by lysing cells in RIPA buffer (20 mM Tris, 150 mM NaCl, 1% Nonidet p-40, 0.5% sodium deoxycholate, 1 mM EDTA, 0.1% SDS) containing protease inhibitor (Sigma) and phosphatase inhibitor cocktail 2 and 3 (Sigma) and then taken to western blot analyses as described previously [29]. Primary antibody were a mouse anti -XRCC1 (Thermo Fisher Scientific, Waltham, MA, USA) and rabbit anti-ATR (Novus Biologicals, USA). HRP conjugate secondary antibody were a rabbit anti-mouse and goat anti-rabbit respectively (Dako, Glostrup, Denmark).

Clonogenic Survival Assay

Two hundred cells per well were seeded in six-well plates. Cells were allowed to adhere for 4 hours. NU6027 or VE-821 were added at the indicated concentrations and the plates were left in the incubator for 10 days for CHO cells. For siRNA transfected OVCAR-3 and OVCAR-4 cells, three days after transfection, NU6027 or VE-821 was added at indicated concentrations and the plates were left in the incubator for 14 days. For cisplatin and ATR inhibitor combination studies, cells were initially treated with cisplatin for 16 hours and then gently washed twice with 1X phosphate buffered saline and incubated in fresh media with or without NU6027 (4 μM for CHO cells and 6 μM for OVCAR-3 cells) for 10 days (CHO cells) or 14 days (human cancer cells). After incubation, the media was discarded, fixed (with methanol and acetic acid mixture) stained with crystal violet and counted. Surviving Fraction = [No. of colonies formed/(No. of cells seeded \times Plating efficiency)]. All clonogenic assays were done in triplicate.

Alkaline COMET Assay

This assay was performed as described previously [31]. Briefly, sub-confluent cells were exposed to NU6027 (4 μM). At 24 hours, cells were extracted and alkaline comet assays were performed. Alkali electrophoresis buffer consisted of 200 mM NaOH, 1 mM EDTA and pH 13. The slides were then stained with SYBR[®] green (1:10,000 dilution) (Molecular Probes) for 10 minutes and images were visualized under a rhodamine filter with an Olympus BX40 microscope. The comets were analysed using Comet Assay III image analysis software (Perceptive Instruments, Suffolk, UK). A total of 200 comet images were evaluated for olive tail moment.

γH2AX Immunocytochemistry

Cells were treated for 48 hours with NU6027 (4 μM for CHO cells and 6 μM for OVCAR-3 cells) and the assay was performed as described previously [31]. For cisplatin and NU6027 combi-

Table 1. XRCC1 siRNA constructs.

siRNA	Sequence
XRCC1_1 siRNA (ID s14940)	5'GGCAGACACUUACCGAAAAtt 3'-sense sequence 3'ttCCGUCUGUGAAUGGCUUUU5'-antisense sequence
XRCC1_2 siRNA (ID s14941)	5'GGCAAGCACUUCUUUUUtt 3'-sense sequence 3'tcCCGUUCGUGAAGAAAAGAA5'-antisense sequence
XRCC1_3 siRNA (ID s14942)	5'GCUUGAGUUUUGUACGGUtt 3'-sense sequence 3'acCGAACUCAAAAACAUGCCAA5' -antisense sequence

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nation studies, initially the cells were treated for 16 hours with cisplatin (1.5 μ M for CHO cells and 1 μ M for OVCAR-3 or OVCAR-4 cells) and then gently washed twice with 1X phosphate buffered saline and incubated in fresh media with or without NU6027 (4 μ M for CHO cells and 6 μ M for OVCAR-3 or OVCAR-4 cells) and the assay was performed as above. The frequencies of cells containing γ H2AX foci were determined in 100 cells per slide in three separate experiments. Nuclei containing more than six γ H2AX foci were considered positive.

Flow Cytometric Analyses (FACS) for Cell Cycle Progression

Cells were treated for 24 hour with NU6027 (4 μ M for CHO cells and 6 μ M for OVCAR-3 or OVCAR-4 cells). Cells were later collected by trypsinization and centrifugation (1000 rpm for 5 minutes) and FACS analyses were performed as described previously [31].

Apoptosis Detection by FITC-annexin V Flow Cytometric Analysis

Cells were treated for 48 hours with NU6027 (4 μ M for CHO cells and 6 μ M for OVCAR-3 or OVCAR-4 cells). Cells were later collected by trypsinization and centrifugation (1000 rpm for 5 minutes) and were washed twice with cold PBS and then resuspended cells in 1X Binding buffer at a concentration of 1×10^6 cells/ml. Then 100 μ l of the solution (1×10^5 cells) was transferred to a 5 ml culture tube and 5 μ l of FITC Annexin V and 5 μ l PI were added. The cells were then gently vortex and incubated for 15 minutes at room temperature (25°C) in the dark. After the incubation, 400 μ l of binding buffer was added to each tube and was analyzed by flow cytometry within 1 hour. For cisplatin and NU6027 combination studies, the cells were initially treated for 16 hours with cisplatin (1.5 μ M for CHO cells and 1 μ M for OVCAR-3 or OVCAR-4 cells) and then gently washed twice with 1X phosphate buffered saline and incubated in fresh media with or without NU6027 (4 μ M for CHO cells and 6 μ M for OVCAR-3 or OVCAR-4 cells) and the assay was performed as described previously. Data was analysed using FlowJo7.6.1 software.

Evaluation of Drug Interaction (Combination Index)

To investigate synergistic and additive activity, combination index was calculated as described previously [30]. Dose-response curves for cisplatin or NU6027 alone were first generated. The effect of the combined treatment was then analysed for the combination of drug A (cisplatin) and B (NU6027), by applying the following equation: $Ac/Ae+Bc/Be = D$, where Ac and Bc correspond to the concentrations of drugs used in the combination treatment, and Ae and Be corresponds to the concentrations of drugs able to, by themselves, produce the same magnitude of effect. If D (combination index) is < 1 the effect of the combination

is synergistic, whereas if $D = 1$ or $D > 1$ the effect is additive or antagonistic respectively [32].

Results

Synthetic Lethality

To evaluate synthetic lethality pre-clinically, a panel of XRCC1 deficient and XRCC1 proficient Chinese Hamster Ovary and human ovarian cancer cell lines were treated with small molecule inhibitors of ATR (NU6027 and VE-821).

Chinese Hamster Ovary (CHO) cells. CHO9 (Wild type), EM-C11 (XRCC1 deficient) and EM-C12 (XRCC1 deficient) were investigated in clonogenic survival assays. We initially evaluated XRCC1 and ATR expression status in CHO9, EM-C11 and EM-C12 cells. Western blot analysis in Figure 1A demonstrates that EM-C11 and EM-C12 have no measurable XRCC1 protein expression compared to CHO9. EM-C11, EM-C12 and CHO9 are proficient in ATR expression. Figure 1B shows that EM-C11 and EM-C12 cells are sensitive to NU6027 treatment compared to CHO9 cells. Similarly, EM-C11 and EM-C12 cells are also sensitive to VE-821 compared to CHO9 cells (Figure 1C). To investigate if sensitivity of XRCC1 deficient cells to the ATR inhibitors can be corrected by expression of wild-type XRCC1 protein in XRCC1 deficient CHO cells, we performed clonogenic assays in EM9-V (XRCC1 mutant) and EM9-XH (cells stably transfected with a human XRCC1 expression vector). Figure 1D demonstrates that EM9-V cells are sensitive to NU6027 compared to EM9-XH.

We then conducted functional analysis in cells. ATR inhibition leads to DNA single strand break (SSB) accumulation. Therefore Alkaline COMET assay was performed. Figure 1E summarizes the results for CHO9, EM-C11 and EM-C12 cells treated with 4 μ M of NU6027. Compared to pre-treatment samples, after 24 hours of exposure to ATR inhibitor, EM-C11 and EM-C12 cells demonstrate a significantly higher mean tail moment compared to CHO9 ($p < 0.01$). The data confirms SSB accumulation following ATR inhibition in XRCC1 deficient cells.

DNA double strand breaks (DSBs) induce phosphorylation of H2AX at serine 139 (γ H2AX). Accumulation of γ H2AX foci in the nucleus is a marker of DSBs. Therefore, γ H2AX immunocytochemistry was performed in EM-C11, EM-C12 and CHO9 cells treated with 4 μ M of NU6027. Nuclei containing more than six γ H2AX foci were considered positive. γ H2AX immunocytochemistry confirmed that XRCC1 deficient EM-C11 and EM-C12 cells accumulated more γ H2AX foci at 48 hours ($p = 0.02$ and $p = 0.05$) compared to CHO9 cells (Figures 1D).

Accumulation of DSBs may delay cell cycle progression. FACS analyses were therefore performed in EM-C11, EM-C12 and CHO9 cells treated with NU6027 (4 μ M). Cell cycle progression was evaluated and compared to control samples. At 24 hours, EM-C11 and EM-C12 were shown to be arrested in G2/M phase of

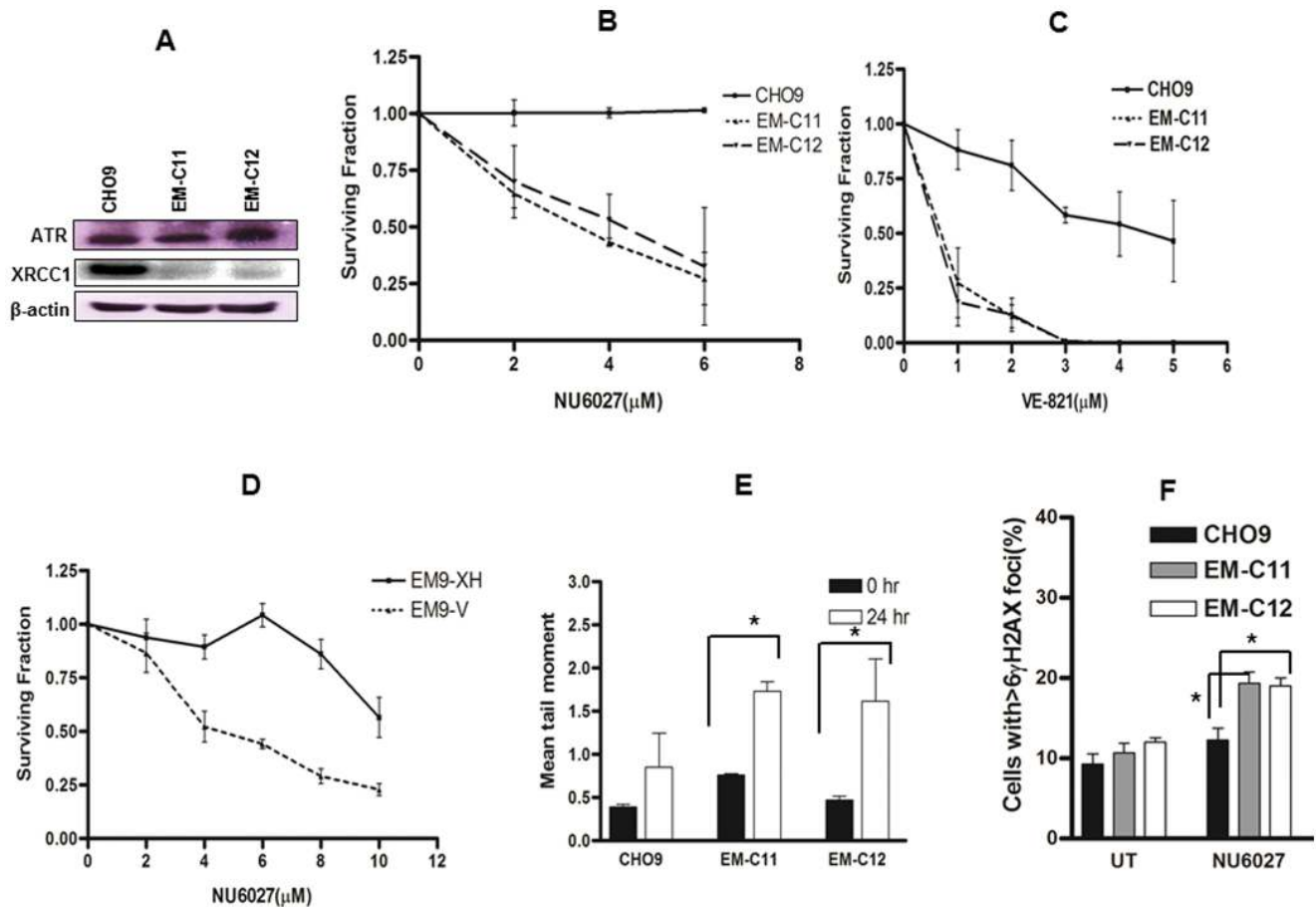


Figure 1. Western blot analysis in chinese hamster (CH) cells (CHO9, EM-C11, EM-C12) (A). Clonogenic survival assays for CH cells treated with NU6027 (B) and VE-821 (C) at indicated concentrations (see methods for details). D. Clonogenic survival assays for EM9-V and EM9-XH cells treated with NU6027. E. Alkaline COMET assay in CH cells treated with NU6027. EM-C11 and EM-C12 demonstrated a higher mean tail moment compared to CHO9 cells. F. EM-C11 and EM-C12 cells accumulate significantly higher γ H2AX foci compared to CHO9 cells upon NU6027 treatment. Data represent mean values \pm SEM (n=6). Results were analysed using Students t-test. * $p < 0.05$. doi:10.1371/journal.pone.0057098.g001

the cell cycle ($p = 0.01$ and $p = 0.03$) compared to CHO9 cells (Figure 2A and 2B).

Accumulation of DSBs, if unrepaired, induces apoptosis in cells. Therefore, FITC-annexin V flow cytometric analysis was conducted to quantify apoptosis in cells treated with $4 \mu\text{M}$ of NU6027 and apoptotic cells quantified at 48 hours. In EM-C11 cells the proportion of cell in early apoptosis increased to 7.5% after 48 hours treatment with NU6027 compared to 1.73% in untreated cells. Similarly in EM-C12 cells, percentage of early apoptotic cells increased from 2.62% to 9.88%. On the other hand in CHO9 cells, there was no change in the percentage of early apoptotic cells (3.22% in untreated and 3.38% after 48 hours of NU6027 treatment (Figure 2C).

The data presented in Chinese Hamster cells suggests that XRCC1 deficient cells are sensitive to ATR inhibitors. ATR inhibition leads to increased DSB accumulation, G2/M cell cycle arrest and apoptosis. This suggests a synthetic lethality relationship between XRCC1 and ATR. To confirm this data further we investigated in human ovarian cancer cell lines.

Human ovarian cancer cells. We generated XRCC1 knockdown human ovarian cancer cell lines using three siRNA constructs (Table 1). After transfection, cell lysates were sampled on days 3, 5 and 7 for XRCC1 knock down by western blot analysis. Figure 3A confirms that all three constructs (siRNA-1,

siRNA-2, siRNA-3) induce efficient knockdown (more than 80%) of XRCC1 in OVCAR-3 cells on day 3 compared to scrambled negative control and GAPDH positive control. In clonogenic survival assays, NU6027 treatment reduced survival in XRCC1 deficient cells compared to proficient cells (Figure 3B). Similar results were also seen in OVCAR-4 cells (Figure S1 A). γ H2AX immunocytochemistry confirmed that XRCC1 deficient cells accumulated more γ H2AX foci at 48 hours ($p = 0.05$, $p = 0.01$ and $p = 0.007$ respectively) compared to scrambled control (Figure 3C). Moreover, at 24 hours, XRCC1 deficient cells were shown to be arrested in G2/M phase of the cell cycle ($p = 0.05$, $p = 0.04$ and $p = 0.05$) compared to CHO9 cells (Figure 3D). In XRCC1 deficient cells the proportion of cells in apoptosis increased substantially compared to scrambled controls upon NU6027 treatment (Figure 3E).

Taken together, the data from CHO cells and human cell lines provide convincing evidence that ATR inhibitors induce synthetic lethality in XRCC1 deficient cells.

Cisplatin Chemopotentiation

We have previously demonstrated that XRCC1 deficient cells are sensitive to cisplatin [17]. In the current study, we first confirmed this observation in XRCC1 deficient EM-C11 and EM-C12 cells compared to CHO9 wild type cells (Figure 4A). We then

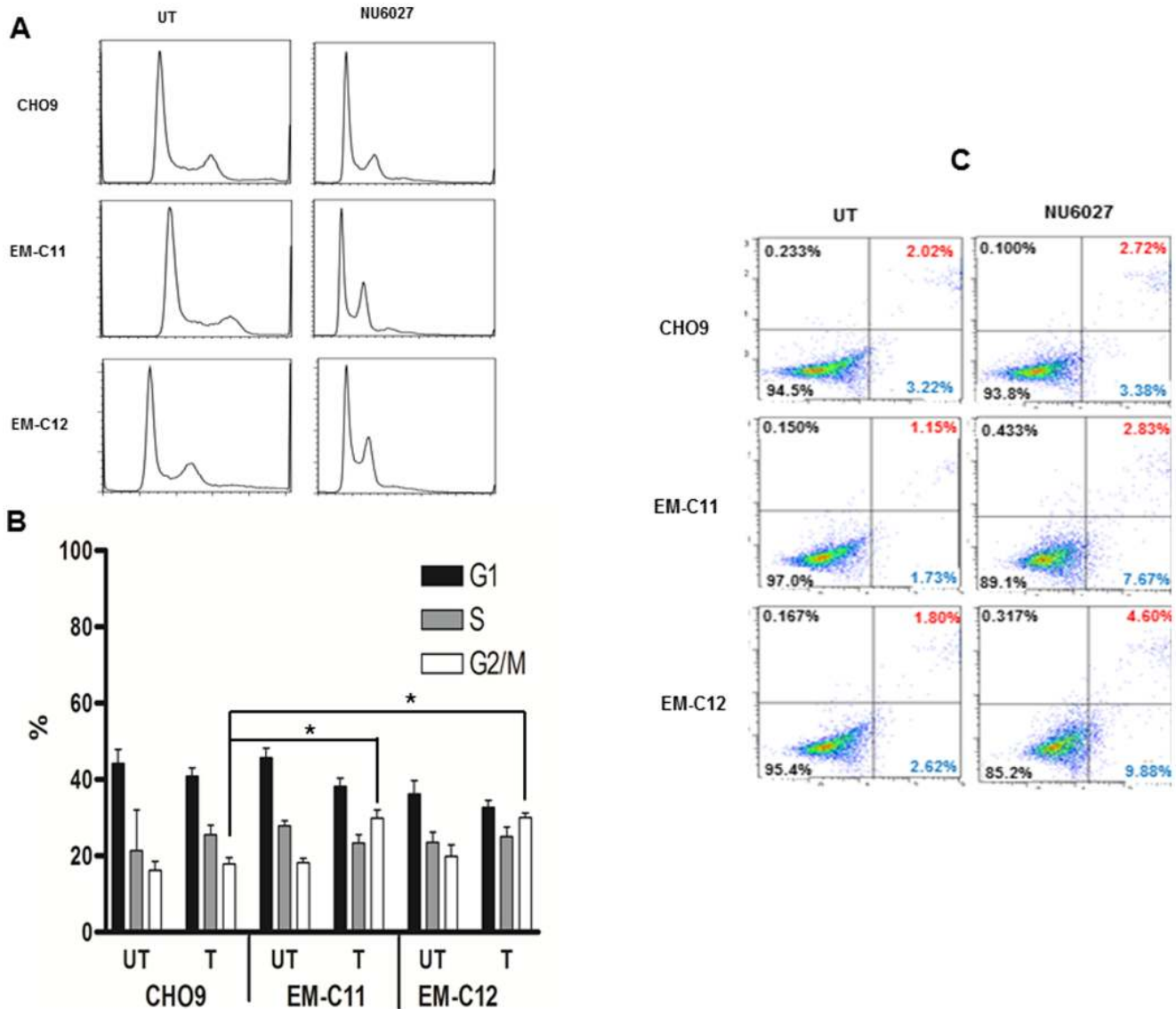


Figure 2. FACS read out in CH cells treated with 24 hours of NU6027 is shown here (A). B. Quantification of various phases of the cell cycle is shown for CH cell treated with NU6027. Data represent mean values \pm SEM (n=6). Results were analysed using Students t-test. * p<0.05. **C.** FITC-Annexin V apoptosis assay is shown here. The proportion of cells in early phase apoptosis is higher in XRCC1 deficient cells treated with NU6027 compared to wild type cells. doi:10.1371/journal.pone.0057098.g002

evaluated combination strategies. The cytotoxicity of cisplatin was enhanced by NU6027 in XRCC1 deficient CH cells compared to XRCC1 proficient cells. In order to evaluate the interaction between NU6027 and cisplatin, combination index was calculated as described previously [32]. Cells were cultured in the presence of increasing doses of cisplatin (range 0.5–3 μ M) in combination with a concentration of NU6027 able to induce a 50% growth inhibition. NU6027 potentiated the cytotoxic effect of cisplatin on XRCC1 deficient cells (Table 2). If combination index (D) is <1 the effect of the combination is synergistic, whereas if D = 1 or D is >1 the effect is additive or antagonistic respectively [32]. In the current study, the combination index was one in EM-C11 and EM-C12 cells. We concluded that enhancement of cisplatin cytotoxicity by NU6027 in CHO cells was additive rather than synergistic. We then proceeded to conduct functional analyses in cells. Cisplatin alone treatment increased DSB accumulation in

XRCC1 deficient cells which was further increased by NU6027 (p = 0.01 and p = 0.02) (Figure 4B). The DSB accumulation seen in XRCC1 deficient cells was also associated with accumulation of apoptotic cells as shown in Figure 4C.

We then conducted similar studies in siRNA transfected human OVCAR-3 or OVCAR-4 cells. Similar to the results seen in CH cells, XRCC1 deficient OVCAR-3 or OVCAR-4 cells were sensitive to cisplatin. NU6027 enhanced cytotoxicity of cisplatin in XRCC1 deficient OVCAR-3 cells compared to XRCC1 proficient cells (Figure 5A). Similar results were also seen in OVCAR-4 cells (Figure S1 B). Combination index studies (Table 2) demonstrated that in most cells was one, except for OVCAR-3 cells treated with Si RNA_3 (D = 0.93) and OVCAR-4 cells SiRNA_1 (D = 0.99). Taken together, we concluded that human ovarian cancer cells the potentiating effect is likely to be additive. Cisplatin alone treatment increased DSB accumulation in cells which was

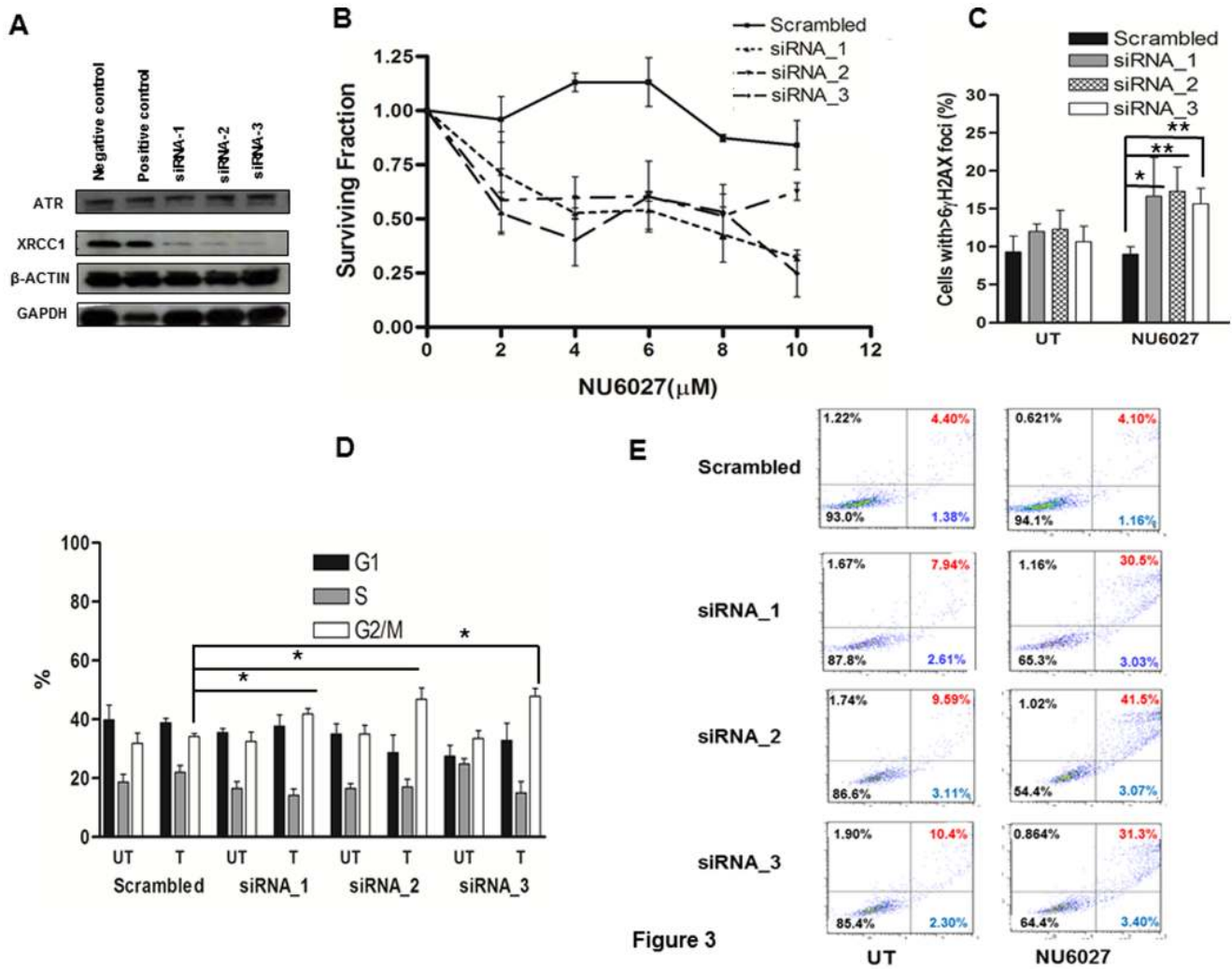


Figure 3

Figure 3. Western blot analysis in siRNA transfected OVCAR-3 cells is shown here (A). B. Clonogenic survival assays for siRNA transfected OVCAR-3 cells treated with NU6027 at indicated concentrations is shown here (see methods for details). **C.** XRCC1 deficient cells accumulate significantly higher γ H2AX foci compared to scrambled control cells upon NU6027 treatment. Data represent mean values \pm SEM (n = 6). Results were analysed using Students t-test. * p < 0.05, ** p < 0.01. **D.** Quantification of various phases of the cell cycle is shown for siRNA transfected OVCAR-3 cell treated with NU6027 is shown here. Data represent mean values \pm SEM (n = 3). Results were analysed using Students t-test. * p < 0.05. **E.** FITC-Annexin V apoptosis assay for siRNA transfected OVCAR-3 cells is shown here. The proportion of cells in late phase apoptosis is higher in XRCC1 deficient cells treated with NU6027 compared to scrambled control cells. doi:10.1371/journal.pone.0057098.g003

Table 2. Effect of NU6027 and Cisplatin in Chinese hamster ovary and human ovarian cancer cells.

Cell lines	NU6027 (μ M;Ac)	Cisplatin (μ M;Bc)	NU6027 (μ M;Ae)	Cisplatin (μ M;Be)	D	
CHO	EM-C11	4	1.5	8	3	1
	EM-C12	4	1.5	10	2.5	1
OVCAR-3	siRNA_1	6	1	10	2.5	1
	siRNA_2	6	1	12	2	1
	siRNA_3	6	1	10	3	0.93
OVCAR-4	siRNA_1	6	1	9	3	0.99
	siRNA_2	6	1	10	2.5	1
	siRNA_3	6	1	10	2.5	1

Ac and Bc, concentrations of drugs used in the combination treatment; Ae and Be concentrations of NU6027 alone and cisplatin alone respectively that produce a similar magnitude of effect; D (combination index).

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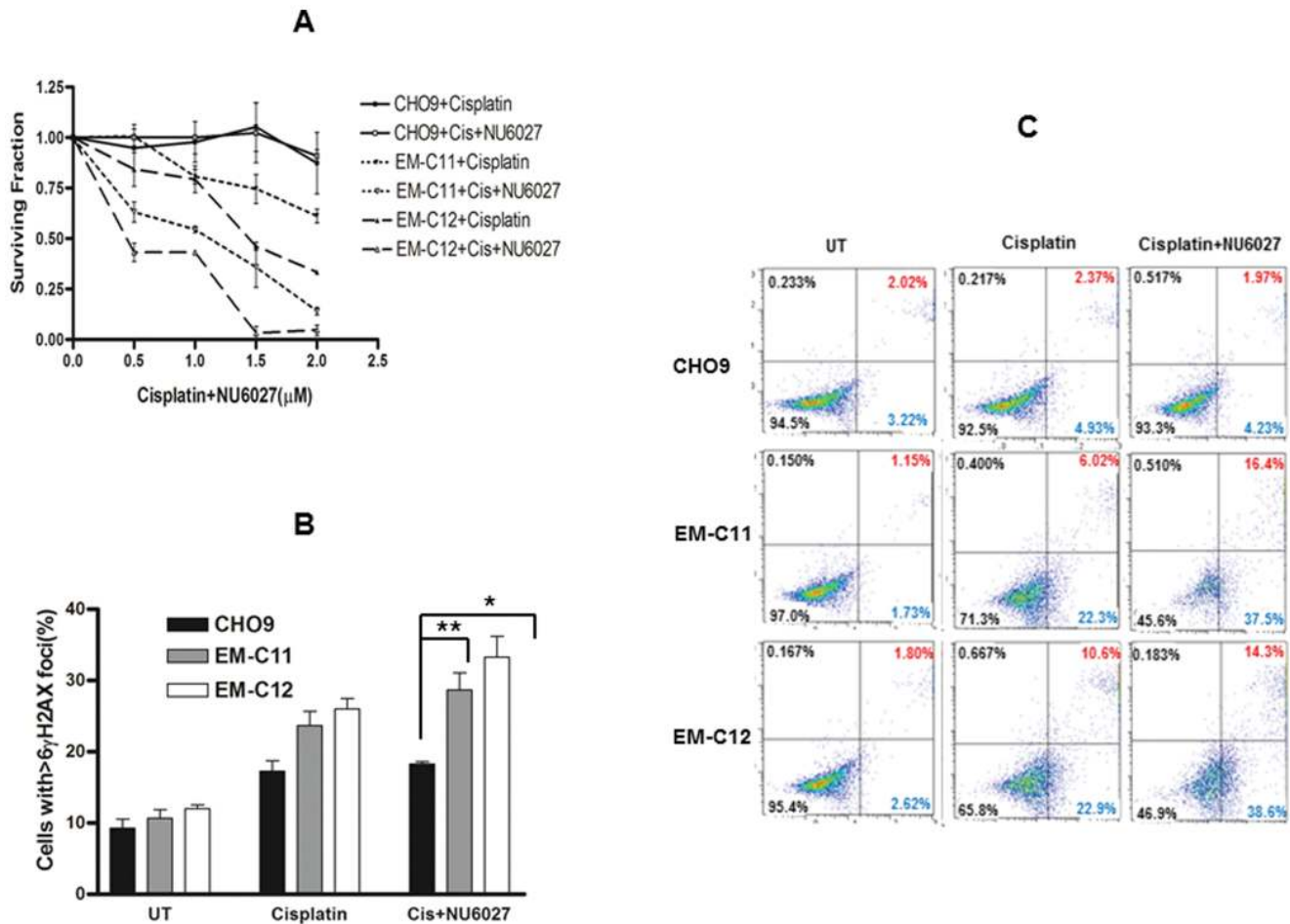


Figure 4. Clonogenic survival assays for CH cells treated with cisplatin alone or in combination with NU6027 is shown here (A). X-axis designates increasing concentration of cisplatin only. NU6027 was fixed at 4 μM. **B.** XRCC1 deficient CH cells accumulate significantly higher γH2AX foci compared to XRCC1 proficient CH cells upon cisplatin treatment alone or a combination of cisplatin and NU6027. Data represent mean values ± SEM (n=6). Results were analysed using Students t-test. * p<0.05, ** p<0.01. **C.** FITC-Annexin V apoptosis assay is shown here. The proportion of cells in early phase as well as late phase apoptosis is higher in XRCC1 deficient cells treated with cisplatin alone or a combination of cisplatin and NU6027 compared to wild type cells. doi:10.1371/journal.pone.0057098.g004

further increased by NU6027 ($p=0.02$, $p=0.007$ and $p=0.004$) (Figure 5B). The DSB accumulation seen in XRCC1 deficient cells was associated with accumulation of substantial apoptotic cells as shown in Figure 5C.

The data presented here not only provides further evidence that XRCC1 deficient cells are sensitive to cisplatin chemotherapy but also suggests that ATR inhibition additively enhances cisplatin toxicity in XRCC1 deficient cells compared to XRCC1 proficient cells.

Conclusions

ATR protein kinase is a key sensor of single-stranded DNA associated with stalled replication forks and repair intermediates generated during BER and DSB repair. ATR activation regulates several cellular processes including cell cycle regulation, DNA replication, DNA repair and apoptosis. XRCC1 is essential for BER and SSB repair and contributes to the ligation step of the NER response. We hypothesised that ATR inhibition could be synthetically lethal in XRCC1 deficient cells.

In the current study we have confirmed that ATR inhibitors are synthetically lethal in XRCC1 deficient cells. We have concluded

synthetic lethality for the following reasons. First, CHO cells as well as human cancer cells deficient in XRCC1 were highly sensitive to ATR inhibitors. Second, functional analyses demonstrated that ATR inhibition in XRCC1 deficient cells led to an accumulation of DNA DSBs, G2/M cell cycle arrest and increased apoptosis. This data is consistent with a study by Peasland et al [22] who demonstrated that NU6027 is synthetically lethal in cells treated with a PARP inhibitor that blocks BER. Moreover, the authors also demonstrated that EM9 Chinese hamster cells lacking XRCC1 are also sensitive to NU6027 [22]. The data, including ours, therefore provides compelling evidence that ATR inhibition is synthetically lethal in BER deficient cells. We present a working model for ATR inhibition as a synthetic lethality strategy in XRCC1 deficient cells. In brief, ATR inhibition leads to SSB accumulation. Cells deficient in XRCC1 are unable to process SSBs which are eventually converted to toxic DSBs at replication forks. Overwhelming DSBs may not only saturate DSB repair, but ATR inhibition is also known to modulate DSB repair directly [33,34] contributing to synthetic lethality observed in cells.

We also found that XRCC1 deficient cells are sensitive to cisplatin. The cisplatin sensitivity in XRCC1 deficient cells observed in our study is consistent with a recent study in HepG2

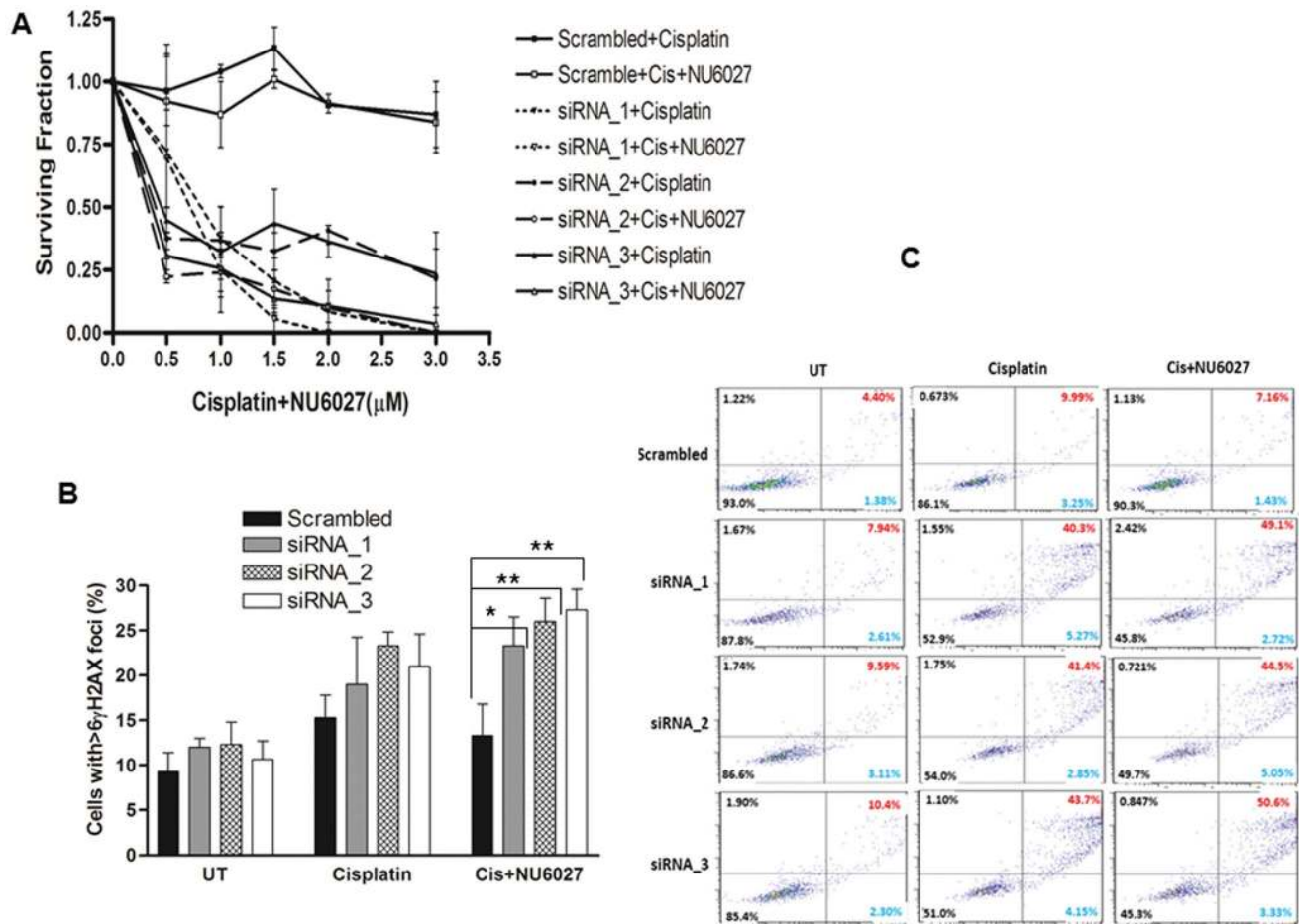


Figure 5. Clonogenic survival assays for siRNA transfected OVCAR-3 cells treated with cisplatin alone or in combination with NU6027 is shown here (A). B. XRCC1 deficient cells accumulate significantly higher γ H2AX foci compared to XRCC1 proficient cells upon cisplatin treatment alone or a combination of cisplatin and NU6027. Data represent mean values \pm SEM (n=6). Results were analysed using Students t-test. * $p < 0.05$, ** $p < 0.01$. C. FITC-Annexin V apoptosis assay is shown here. The proportion of cells in late phase apoptosis is higher in XRCC1 deficient cells treated with cisplatin alone or a combination of cisplatin and NU6027 compared to wild type cells.
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cells where cisplatin sensitivity was demonstrated following XRCC1 depletion [35]. We did not observe any potentiation of cisplatin cytotoxicity by ATR inhibitor in XRCC1 wild type cells. This is in contrast to previous preclinical studies where ATR inactivation (genetically or with inhibitors) has demonstrated increased platinum sensitivity in a panel of cell lines [22]. However, a limitation of our study is that our investigation was restricted to a few cell types only. Nevertheless, our data suggests that genetic background (such as XRCC1 status) may influence platinum sensitivity. In conclusion, we have demonstrated a

synthetic lethality application for ATR inhibitors in XRCC1 deficient cells. ATR inhibition may also influence platinum sensitivity in XRCC1 deficient cells.

Author Contributions

Reviewed and approved the final version of the manuscript: RS TAF CP PM NA VM CS SC SM. Conceived and designed the experiments: SM TAF. Performed the experiments: RS CP VM PM. Analyzed the data: RS CP VM SM SC NA. Contributed reagents/materials/analysis tools: CS. Wrote the paper: RS SM CP TAF CS.

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