Inhibition of Snail1-DNA-PKcs Protein-Protein Interface Sensitizes Cancer Cells and Inhibits Tumor Metastasis §

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Background: Previously we identified the interaction between DNA-PKcs and Snail1.

Results: Inhibition of the interaction between two molecules using Snail1 peptide (7 amino acid construct) sensitized cancer cells and inhibited tumor migration.

Conclusion: Interfering with the protein-protein interaction between DNA-PKcs and Snail1 may be an effective strategy for cancer treatment.

Significance: Interfering with the interaction between DNA-PKcs and Snail1 may be a novel target for cancer treatment.

SUMMARY

Our previous study suggested that DNA-PKcs interacts with Snail1, which affects genomic instability, sensitivity to DNA-damaging agents, and migration of tumor cells by reciprocal regulation between DNA-PKcs and Snail1. Here, we further investigate that peptide, which contains 7 amino acid sequences 15-21 of Snail1 (KPNYSEL, SP), inhibits the endogenous interaction between DNA-PKcs and Snail1 through primary interaction with DNA-PKcs. SP restored the inhibited DNA-PKcs repair activity and downstream pathways. On the other hand, DNA-PKcs-mediated phosphorylation of Snail1 was inhibited by SP, which resulted in decreased Snail1 stability and Snail1 functions. However, these phenomena were only shown in p53-wild type cells, not in p53-defective cells. From the results, it is suggested that interfering with the protein interaction between DNA-PKcs and Snail1 might be an effective strategy for sensitizing cancer cells and inhibiting tumor migration, especially in both Snail1- and DNA-PKcs-overexpressing cancer cells with functional p53.

The epithelial-mesenchymal transition (EMT), a major mechanism of cancer metastasis, is initiated by repression of the epithelial adhesion molecule, E-cadherin. This repression is mediated by several transcription factors including Snail (also known as Snail1) and Slug (also known as Snail2) (1-5). In most human cancers, metastatic tumors are resistant to radiation or chemotherapy, and emerging evidence suggests a correlation between EMT and resistance of cancer cells to chemo and radiotherapy (6-11).

The DNA-dependent protein kinase catalytic
subunit (DNA-PKcs) is a DNA-activated serine/threonine protein kinase, and is abundantly expressed in almost all mammalian cells. DNA-PKcs is most commonly known as a critical component in the DNA-damage repair pathway, including nonhomologous end-joining (NHEJ) repair and homologous recombinant (HR) repair. DNA-PKcs has been shown to phosphorylate several cellular proteins, including itself, the variant histone H2AX and p53 (12-15). Homozygous knockout mice of DNA-PKcs catalytic subunit (DNA-PKcs−/−) are hypersensitive to ionizing radiation (IR) and chemical treatment, and have defects in DNA repair activity (16-18). From this observation, one might hypothesize that DNA repair inhibitor would sensitize the tumor cells, and in some cases, reverse the resistance phenotype, when administered in combination with drugs and/or irradiation. However, a DNA repair inhibitor generally exhibits no specific activity in killing the tumor cells, and the screening procedure should include a DNA-damaging treatment (19, 20). Moreover, a number of studies have reported a lack of correlation between cell sensitivity/resistance to IR and down/up regulation of DNA-PKcs expression (18, 21).

DNA repair is also involved in the maintenance of genome stability and deficient activity may contribute to neoplastic transformation, as it has been repeatedly documented in relation to mismatch repair and nucleotide excision repair. DNA-PKcs is found at the end of chromosomes, suggesting a role in telomere maintenance and prevention of chromosomal end-to-end fusion (22-26). Interestingly, although DNA-PKcs is highly expressed in human cells, the other NHEJ factors are not as abundant, and the high levels of DNA-PKcs in human cells do not impart any increased ability to repair DNA damage. If the amount of expressed DNA-PKcs essentially exceeds the demand for DNA-damage repair, it is not yet understood why human cells universally express such high levels of this complex. Indeed, DNA-PKcs is also reported to be involved in metabolic gene regulation in response to insulin, independently from the NHEJ repair system (27-29).

In our previous study, we identified a novel function of DNA-PKcs as a regulator of Snail1. The direct interaction between DNA-PKcs and Snail1 induced Snail1 phosphorylation at Ser100, which stabilized the Snail1 protein and potentiated Snail1 functions, such as E-cadherin promoter repression and metastasis properties. Snail1 phosphorylation exhibited reciprocal inhibition of DNA-PKcs kinase activity, resulting in impaired DNA-damage repair and induction of chromosomal or genomic instability (30). In this study, we further utilized a novel, seven amino acid-containing Snail-peptide (SP), which is included in the binding site for interaction with DNA-PKcs. In addition, SP can abrogate the DNA-PKcs-mediated Snail1 phosphorylation at Ser100 that resulted in the blocking of Snail1-mediated radioresistance and metastasis. SP also restored DNA-PKcs-mediated repair activity, which was due to direct interaction with endogenous Snail1. These results suggest a novel strategy for anticancer therapy through inhibition of binding between DNA-PKcs and Snail1 in cancer cells with overexpression of both DNA-PKcs and Snail1. This approach highlights how protein-protein interfaces can be exploited to develop novel targets, which can be used to dissect signal transduction mechanisms and provide leads for novel therapeutics to target specific Snail1-DNA-PKcs protein-protein interactions in cancer cells.

**EXPERIMENTAL PROCEDURES**

**Plasmids** - DNA-PKcs deletion constructs were generated using PCR, by isolating internal regions from the full-length cDNA for DNA-PKcs, kindly given to us by Dr. Katheryn Meek (Michigan State University, East Lansing, MI). PCR fragments were inserted into the pCMV6 (31) expression vector using NotI and KpnI restriction sites; products were sequenced for verification (Cosmogenetech co, Ltd., Seoul, Korea). Expression vectors pCR3.1–Snail1–Flag and Snail1 mutant proteins with Flag epitope tails, including the S100A mutant, were prepared as described previously (32, 33).

**Cell culture and transfection** - DLD-1 human colorectal adenocarcinoma cells, M059J/M059K human glioblastoma cells, and CT26 murine colon carcinoma cells were cultured in Dulbecco’s minimal essential medium (DMEM; GIBCO, Gaithersburg, MD, USA), supplemented with heat-inactivated 10% fetal bovine serum (FBS; GIBCO) and antibiotics. The expression of peptides (4 μM) were introduced into cells using Lipofectamine® 2000 transfection reagent (Invitrogen, CA, USA).
Antibodies and Reagents - Cycloheximide (CHX), propidium iodide (PI), and a monoclonal antibody for Flag were purchased from Sigma (St. Louis, MO, USA). Monoclonal antibodies for DNA-PKcs, Ku86, p53, pre-designed siRNAs for human DNA-PKcs and a negative control Si-RNA were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Anti-Snail1 and p-p53 were purchased from Cell Signaling Technology (diluted 1:1000, Danvers, MA, USA). Anti-p-DNA-PKcs (Ser2056) was purchased from Abcam (diluted 1:1000, Cambridge, MA, USA) and γ-H2AX was purchased from Millipore (diluted 1:1000, Young In Frontier, Seoul, Korea). Control peptide and Snai1 peptide were purchased from AnyGen (Jangseong-gun, Korea). TGF-β was purchased from BD Biosciences (San Jose, CA, USA).

Immunoblotting and immunoprecipitation – For immunoblotting, cells were lysed with RIPA buffer (50mM Tris-HCl [pH 7.5], 150mM NaCl, 1% NP-40, 0.1% sodium dodecyl sulfate (SDS), and 1% sodium deoxycholate) supplemented 1mM Na3VO4, 1mM dithiothreitol (DTT), 1mM NaF, and protease inhibitor cocktail (Roche, IN, USA). The samples were boiled for 5 min, and an equal amount of protein was analyzed on SDS-PAGE. For immunoprecipitation, cells (1x10⁷) were lysed in immunoprecipitation buffer (50 mM HEPES pH 7.6, 150mM NaCl, 5mM EDTA, 0.1% NP-40). After centrifugation (10 min at 15,000 g) to remove particulate material, supernatants were incubated with antibodies (1:100) against DNA-PKcs, biotin, and Flag with constant agitation at 4°C. Immunocomplexes were precipitated with protein A–Sepharose (Sigma) and analyzed by SDS-polyacrylamide gel electrophoresis using enhanced chemiluminescence detection.

Irradiation – Cells or mice were exposed to γ-ray with a 137Cs γ-ray source (Atomic Energy of Canada, Mississauga, Ontario, Canada) with a dose rate of 3.18Gy/min.

DNA-PKcs kinase assay - DNA-PKcs kinase activity was analyzed, using the signaTECT DNA-PKcs assay system (Promega, Madison, WI, USA). Nuclear extracts were isolated for DNA-PKcs kinase activity analysis. The phosphorylation reactions were carried out in the presence of γATP-3²P (NEN Life Science, Boston, MA, USA) for 5 min at 30°C with 25 μg of biotinylated-p53 peptide (biotin-EPPLSQEAFADLWKK; Promega) as the substrate. The efficiency of ³²P labeling of the substrate was measured using a phosphor-imager.

Ubiquitination assay - For cell-based ubiquitination assays, DLD-1 cells were transfected with HA-ubiquitin constructs. After transfection, the cells were lysed with immune precipitation assay buffer, and samples were briefly sonicated. After centrifugation, clear lysates were immunoprecipitated with anti-Ub (Santa Cruz) antibodies. Immunocomplexes were analyzed by Western blot, using anti-Ub or anti-Snail1 antibodies.

E-cadherin promoter assays - Cells growing at ~70% confluence in 60 mm dishes were transfected using Lipofectamine® 2000 transfection reagent (Invitrogen) according to the manufacturer’s protocol. For experiments assessing activation of 1 μg pGL3-control vector (Promega, Madison, WI, USA) or E-cadherin reporter gene construct by endogenous factors, 1 μg E-cadherin reporter gene construct and 200 ng of β-galactosidase expression vector were transfected into cells. The cells were harvested 48h after transfection using reporter lysis buffer (Promega) followed by determination of luciferase and β-galactosidase activities. All results represent the mean of three independent experiments.

Migration assay - Cell migration was assayed using a Boyden chamber assay, as described previously (34). Cells were harvested with trypsin (0.1 mg/mL), centrifuged, counted, and resuspended in serum-free media. The cells were plated on the upper side of a collagen-treated, polycarbonate membrane, separating the two chambers of 6.5 mm Transwell culture plates (Costar, Corning, NY, USA). After 16 h, the cells on the upper face of the membrane were scraped using a cotton swab and the cells that migrated to the lower face of the membrane were stained with DiffQuick Wright-Giemsa solution (Baxter, Deerfield, IL, USA).
**Linear dsDNA-associated protein pull-down assay** - Nuclear extracts were isolated for a dsDNA pull-down assay. A biotinylated oligonucleotide (1 kb), which was generated by PCR amplification of pcDNA3 (5.4 kb) with the biotinylated forward primer 5’-GACTCTCAGTACAATCTGCTCTG-3’ and reverse primer 5’-AGCTCTAGCATTTAGGTGACACT-3’, was immobilized on streptavidin beads (Sigma). Immobilized DNA was mixed with nuclear extracts and incubated for 3 h at 4°C. The beads were washed with buffer D (10mM Tris, pH 7.6, and 100 mM NaCl) and boiled for 5 min in 2x SDS sample buffer. Double-stranded DNA-associated proteins were analyzed by SDS-PAGE using enhanced chemiluminescence detection (ECL, Amersham).

**Immunofluorescence analysis** - For immunofluorescence analysis, cells were fixed with 4% paraformaldehyde, permeabilized with 0.1% Triton X-100 in PBS, and then washed three times with PBS. The cells were then incubated with anti-DNA-PKcs or anti-biotin diluted to 1:200 in PBS with 5% FBS for 1 h at room temperature in a humidified chamber. Excess antibody was removed by washing the coverslips three times with PBS. The cells were then incubated with Alexa Fluor-488 (green) and Alexa Fluor-568 (red) secondary antibodies (Invitrogen, CA, USA) at a 1:200 dilution in PBS with 5% FBS for 2 h. After washing three times with PBS, coverslips were mounted onto microscope slides using ProLong anti-fade mounting reagent (Molecular Probes, Eugene, OR, USA). The slides were analyzed using a confocal laser-scanning microscope (Leica Microsystems, Wetzlar, Germany).

**Cell cycle analysis** - For cell cycle analysis, the cells were fixed in 70% ethanol at -20°C for at least 18 h. The fixed cells were washed once with PBS-EDTA and resuspended in 1 mL of PBS. After the addition of 10 µL each of PI (5 mg/mL) and RNase (10 mg/mL), the samples were incubated for 30 min at 37°C and analyzed using a FACSScan flow cytometer (Becton Dickinson).

**Comet assay** - The cells were exposed to IR (5 Gy) and subjected to a comet assay to detect DNA damage and repair at the single-cell level, using a commercially available assay system (Trevisgen, Gaithersburg, MD, USA). Briefly, after treatment with IR, cells were harvested and mixed with low melting temperature agarose. After lysis, electrophoresis was performed at 1 V cm⁻¹ and 15 mA for 40 min. Slides were stained with SYBG green dye for 10 min. Twenty randomly selected cells per sample were captured under a Zeiss fluorescent microscope and digital fluorescent images were obtained using Comet software (Andor, Belfast, UK). The comet distance moment was proportional to the amount of DNA damage present in the individual nuclei and was measured by the Olive tail moment using the Comet software.

**Lung colonization assay** - All protocols involving mice were approved by the Institutional Animal Care and Use Committee of Ewha Womans University. To generate lung metastases, 2×10⁶ CT26 cells were injected into the tail vein of C57BL6 mice (Central Lab. Animal Inc, Seoul, Korea). Two days later, whole body 3 Gy irradiation was performed. Intraperitoneal injection of CP or SP at 3 mg/kg in PBS was performed once before IR and 4 times after IR. Lungs were harvested 2 weeks after cell injection, weighed, and fixed in formalin. Colony number and colony area in lung were counted.

**Statistical analyses** - A student’s t-test and an analysis of variance (ANOVA) were used to determine significant differences between the experimental groups. Statistical analyses were performed using GraphPad Prism 5.0 (GraphPad Software, Inc.).

**RESULTS**

**DNA-PKcs interacts with both Snail1 and Slug** - Our previous study suggested that Snail1 interacts with DNA-PKcs (30), and Snail1 and Slug belong to the same Snail super family of zinc finger transcription factors (35-38). When DLD-1 cells overexpressed Flag-Snail1 or Flag-Slug, immunoprecipitated DNA-PKcs bound to both Snail1 and Slug (Fig. 1A). Sequence analysis of Snail1 and Slug revealed that only the amino acid sequences 15-21 of Snail1 and Slug showed sequence homology. Because these sequences have only seven overlapping amino acids (KPNYSEL), we were able to synthesize these Snail peptides (SP). A control peptide (CP), made up of an amino acid sequence that was not
homologous to either Snail1 or Slug, was also prepared (NPEFTFQ). Biotin-labeled SP interacted with DNA-PKcs; however, biotin-labeled CP did not (Fig. 1B, left). Interaction between Snail1 and DNA-PKcs was inhibited by an increased addition of SP to Snail1-overexpressing cells (Fig. 1B middle and right). Immunofluorescence data also revealed that biotin labeled SP co-localized with DNA-PKcs in nucleus of the cells, but biotin labeled CP did not (Fig. 1C). When we added Site 7, amino acids 3534-4129 of the kinase domain of DNA-PKcs, which is the binding site of DNA-PKcs for interaction with Snail1, binding of biotin-labeled SP with Site-7 occurred. This suggests that SP interacted with DNA-PKcs at the Snail binding site (Fig. 1D). To examine the physiological relevance, TGF-β, transcriptional activator of Snail1 (39-41), was treated and binding activity was detected. Increased binding activity between DNA-PKcs and Snail1 by TGF-β was inhibited by additional SP transfection (Fig. 1E). The peptide sequences selected from Snail1 and Slug, hypothesized to interact with DNA-PKcs, included seven amino acids and treatment of this peptide inhibited endogenous binding activity between Snail1 (or Slug) and DNA-PKcs.

**SP activates DNA repair activity and inhibits DNA-PKcs-mediated Snail1 phosphorylation at Ser100 after IR** - We hypothesized that SP might interfere with direct interaction between DNA-PKcs and Snail1. To confirm this, we examined the downstream pathway of DNA-PKcs after IR. SP treatment potentiated IR-induced phosphorylation of DNA-PKcs at Ser2056, p53 at Ser15 and γ-H2AX, when compared to those of CP treated cells. However, in the case of DNA-PKcs knockdown cells, these differences were not observed (Fig. 2A). IR-induced DNA-PKcs kinase activity, using p53 peptide as a substrate, was inhibited by Snail1 overexpression; however, SP treatment restored the DNA-PKcs kinase activity, when compared to the CP treated cells. Moreover, adding Site 7 of DNA-PKcs, which can interact with SP, blocked the restoring DNA-PKcs kinase activity, which was possibly due to an endogenous interaction between Snail1 and DNA-PKcs (Fig. 2B). We also observed recruitment of DNA-PKcs to double-stranded breaks (DSB) in DNA after transfection of DNA with double-stranded (ds) oligonucleotides (DSB pull down assay). SP increased recruitment of DNA-PKcs or Ku86 to DSB sites, while CP did not. In the case of cells in which DNA-PKcs was knocked-down through treatment with Si-DNA-PKcs, no recruitments were observed (Fig. 2C).

To examine its physiological relevance, we examined DSB repair activity. We observed that SP treatment significantly inhibited IR-mediated comet tail production, and reduced IR-mediated aneuploidy (>4N), when compared to cells treated with CP; however these phenomena were not detected in DNA-PKcs-knockdown cells (Fig. 2D).

**Phosphorylation of Snail1 at Ser100 is blocked by SP, which results in Snail1 protein degradation by the ubiquitination pathway** - Our previous data suggested that DNA-PKcs phosphorylated Snail1 at Ser100, thereby increasing its protein stability (30). Since SP blocked interaction between DNA-PKcs and Snail1, we examined whether SP would affect DNA-PKcs-mediated Snail1 phosphorylation at Ser100. The phosphorylation of Snail1 at Ser100 was increased by IR in CP treated cells; however, in the case of SP treated cells, IR did not induce an increase in phosphorylation at Ser100. When DNA-PKcs was knocked down, the potential of Snail1 phosphorylation at Ser100 by IR was attenuated and treatments of CP and SP showed no differences in the patterns of Snail1 phosphorylation (Fig. 3A).

Since Snail1 phosphorylation at Ser100 increased the stability of Snail1 protein, we compared the half-life of Snail1 after treatment with SP by using cycloheximide (CHX) to block protein synthesis. SP treatment appeared to result in a shorter half-life of Snail1 when compared to CP treated cells. However, these differences were not shown in DNA-PKcs-knockdown cells (Fig. 3B). Indeed, ubiquitination patterns also indicated that ubiquitinated Snail1 was increased by SP treatment, when compared to the CP treated cells (Fig. 3C).

**SP increases E-cadherin promoter activity and inhibits tumor migration** - To elucidate whether the inhibition of Snail1 phosphorylation at Ser100 by SP affected Snail1 function, we examined E-cadherin promoter activity, which is reported to be inhibited by Snail1. E-cadherin promoter activity was higher in SP treated cells...
than in CP treated cells. The knock down of DNA-PKcs increased E-cadherin promoter activity; however, SP treatment did not increase E-cadherin promoter activity. These phenomena were also observed when wild-type (M059K) and DNA-PKcs deleted (M059J) cells were used (Fig. 4A). We also examined the protein expression of E-Cadherin and similar effects were observed (Fig. 4B). Also, migration activity was inhibited by SP treatment; however, migration activity of DNA-PKcs-knockdown cells was not affected by SP treatment (Fig. 4C). As Snail1 function is necessary for tumor metastases and expansion, we examined whether SP affected the Snail1-mediated growth of lung metastasis in animal models. Stable transfection of murine CT26 colon carcinoma cells with Snail1-WT, or Snail1-S100A (phospho-defective mutant) was performed and controlled. Transfected cells were injected into mice by tail vein to generate lung colonies. When compared to irradiated mice after injection with control CT26 cells (3Gy IR alone), Irradiated mice after injection with Snail1-WT overexpressed cells did not show any significant differences in lung colonies, which was observed in lung weights, lung tumor nodule counts, and lung tumor area in lung tissues. Interestingly, SP treatment in combination with IR significantly inhibited lung colonies; these data were similar to those of S100A transfected cells. However, in the case of CP treatment, no inhibition was observed (Fig. 4D). In summary, SP inhibited tumor metastasis by inhibition of Snail1 function.

SP sensitizes IR-mediated cell death - Because SP treatment restored DNA-PKcs kinase activity and repair activity, and p53 is one of the DNA-PKcs substrates, we decided to examine whether the effects of SP were affected by p53 status. SP potentiated phosphorylation of DNA-PKcs and decreased protein stability of Snail1 in both p53+/+ and p53/-/- cells. Phosphorylation of p53 and γ-H2AX was also potentiated in p53+/+ cells. However, in the case of p53/-/- cells, no increase of phosphorylation of γ-H2AX was observed (Fig. 6A). A colony forming assay, cell death detection by FACS and caspase cleavage data also suggest that the SP sensitization of IR-mediated cell death is p53 dependent (Figs. 6B, 6C and 6D).

DISCUSSION

The Snail1 and Slug proteins play an essential role in cancer associated EMT (1-5). Even though many functions about the Snail family and EMT are well recognized, there are still unknown mechanisms of Snail. In this study, we identified that the seven amino acid (KPNYSEL, SP) peptide sequence, contained by both Snail1 and Slug, can interact with DNA-PKcs. Treatment with SP blocked the endogenous binding activity between Snail1 and DNA-PKcs, resulting in abolishment of DNA-PKcs-mediated Snail1 phosphorylation at Ser100 and Snail protein stability. Additionally, SP treatment provided effective sensitization of cancer cells and inhibited tumor metastasis in both Snail- and DNA-PKcs-overexpressed cancer cells.

In our previous study, we identified a novel function of DNA-PKcs as a regulator of Snail1. The direct interaction between DNA-PKcs and Snail1 induced Snail1 phosphorylation at Ser100, which stabilized the Snail1 protein and potentiated Snail1 functions, such as E-cadherin promoter repression and metastasis properties. Snail1 phosphorylation exhibited reciprocal inhibition of DNA-PKcs kinase activity, resulting in impairment of DNA damage-repair and induction of chromosomal and genomic instability (30).

In this study, we aimed to determine whether disruption of the protein-protein interaction between DNA-PKcs and Snail1 would give rise to similar and/or novel biological insights on the sensitization of cancer cells and inhibition of tumor metastasis. Because DNA-PKcs kinase activity is very important to
DNA damage control, kinase inhibition by Snail1-phosphorylation, which was induced by TGFβ treatment, or IR, may lead to more serious genomic instability of the cancer cells. Because DNA-PKcs interacted with both Snail1 and Slug, we searched sequence homology between two proteins and found that only seven amino acids (SP) showed sequence homology. Not only did SP interact with DNA-PKcs, it inhibited the endogenous interaction between DNA-PKcs and Snail1. SP attenuated DNA-PKcs-mediated phosphorylation of Snail1 at Ser100, which resulted in decreased protein stability of Snail1.

In terms of physiological relevance, defective DNA-damage repair and aneuploidy production can result from defective DNA-PKcs kinase activity through the interaction with Snail1. In addition, tumor migration is increased by DNA-PKcs-induced phosphorylation of Snail1 at Ser100; however, SP treatment restored DNA-PKcs kinase activity and blocked Snail1 phosphorylation. SP treatment actually sensitized cancer cells, and in combination with IR, inhibited tumor migration, or metastasis, in both in vitro and in vivo systems.

One of the targets of DNA-PKcs is p53; function of this protein is essential in DNA repair and cancer cell sensitization (42). Indeed, SP effects were only observed in p53-functional cells, not in p53-defective cells, suggesting that intact p53 function is necessary for SP-mediated inhibition of migration and sensitization of tumor cells. Of course, our finding has limitations because SP treatment alone only exhibited minor effects on the sensitization of cancer cells and inhibition of tumor metastasis. These results suggest that DNA-PKcs may be involved in only a small portion of Snail functions, especially after a DNA-damaging incidence such as IR. Another limitation of our findings is that cancer cells showed p53 deletion or mutation, however, SP showed sensitization of cancer cells and inhibition of tumor metastasis only with wild type p53. Nonetheless, since Snail1 acts as a critical role in cancer metastasis (1-5) and DNA-PKcs, which we do not know much about in cancer, is an abundant protein in cancer tissue (28), inhibition of these two proteins might be an effective strategy for cancer treatment, focusing on our identification of a peptide motif as a key binding site for DNA-PKcs involved in protein-protein interactions with Snail1. This peptide found in Snail1, that competes with endogenous Snail1 for interaction with DNA-PKcs, can induce sensitization of cancer cells and inhibition of tumor metastasis. Therefore, interfering with the protein-protein interaction between DNA-PKcs and Snail1 may be an effective strategy for sensitizing cancer cells to treatment and inhibiting tumor metastasis. This concept is relatively important to address due to a growing interest in pharmaceutical development to target protein-protein interactions as a therapeutic strategy against diseases such as cancer (43).
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**FOOTNOTES**

DNA-PKcs, DNA-dependent protein kinase catalytic subunits; NHEJ, non-homologous end joining; DSBs, DNA double-stranded breaks; HR, Homologous recombination; EMT, epithelial-mesenchymal transition; CHX, cycloheximide

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**FIGURE LEGENDS**

**Figure 1. DNA-PKcs interacts with both Snail1 and Slug.** (A) Flag-tagged Snail1 or Flag-tagged Slug was transfected into DLD-1 cells. Western blotting or immunoblotting was conducted, following immunoprecipitation using cell extracts. (B) Control-peptide (CP, 4 μM) and Snail1-peptide (SP, 4 μM) were transfected into DLD-1 cells (left) or SP (0, 1.5, 3 and 6 μM) was transfected into a vector control or Flag-tagged Snail1-overexpressing cells. After 24 h, the cells were harvested and lysates were generated for immunoprecipitation and immunoblotting (middle). The relative binding activity between Snail1 and DNA-PKcs was calculated by comparing densitometric scans of the sample immunoblots after immunoprecipitation with the values of control samples set at 1 (right). (C) DLD-1 cells were transfected with biotin-labeled CP or SP (each 4 μM). The cells were immunostained for detection of peptide (green) and DNA-PKcs (red). DNA was stained with DAPI. Localization was analyzed by confocal microscopy. (D) c-Myc-tagged, truncated DNA-PKcs constructs (amino acids 3534-4129 of the kinase domain, Site 7) were transfected into CP and SP transfected cells. Western blotting or immunoblotting was conducted, following immunoprecipitation using cell extracts. (E) After transfection of CP or SP to DLD-1 cells, TGF-β (10 ng/ml) were incubated for 24 h. Western blotting and immunoprecipitation were performed.

**Figure 2. SP activates DNA repair activity and inhibits DNA-PKcs-mediated Snail phosphorylation at Ser100 after IR.** (A) After transfection of Control peptide (CP, 4 μM) or Snail1 peptide (SP, 4 μM) into control (Si-CON) or SiRNA of DNA-PKcs (Si-DNA-PKcs)-transfected DLD-1 cells, immunoblotting was performed at the indicated time points after exposure of the cells to 5 Gy of radiation (IR) (upper). The relative protein band intensity was calculated from densitometric scans of immunoblots with the control values set to 1. (n=3, data are presented as mean ± s.d.) (lower). *p<0.05 vs CP treated cells. (B) CP or SP was transfected into c-Myc-tagged, truncated DNA-PKcs
construct (amino acids 3534-4129 of the kinase domain, Site 7) transfected cells. An in vitro DNA-PKcs kinase assay was performed. The SAM membrane or 3 mm filter disc was analyzed using a phosphoimager system and values of Si-Con using p53 peptide as a substrate were set to 1. (C) The protein levels of DNA-PKcs and Ku86 in the dsDNA pull-down lysates, as well as in complete whole nuclear extracts (WNE), were analyzed. (D) Sh-control (Sh-CON) or Sh-DNA-PKcs (Sh-DNA-PKcs) was stably transfected into DLD-1 cells. After 12 h of 5 Gy IR, immunoblotting (left), a comet assay (middle) and flow cytometry analysis, after PI staining for detection of >4N cells (right), were performed. Experiments were performed more than 3 times and data are presented as mean ± s.d. *p<0.05

Figure 3. Phosphorylation of Snail at Ser100 is blocked by SP and results in Snail1 degradation by ubiquitination pathway. (A) At the indicated time points of 5 Gy of radiation (IR) after transfection of Control-peptide (CP, 4 μM) or Snail1-peptide (SP, 4 μM), western blotting was performed using anti p-Snail1 (upper). Relative protein band intensity was calculated by comparing densitometric scans of the sample immunoblots with the values of control samples set at 1. Result is the representative of 3 independent experiments (lower). (B) Protein extracts were prepared at the indicated time points, following cycloheximide (CHX) treatment (20 μg/mL) to control (Si-CON)- or SiRNA of DNA-PKcs (Si-DNA-PKcs)-transfected DLD-1 cells, also transfected with either CP or SP. Western blot analysis was performed. The relative protein band intensity was calculated from densitometric scans of immunoblots with the control values set at 1 (n=3, data are presented as means ± s.d. *p<0.05 vs CP treated cells). (C) For ubiquitination assays, CP or SP was transfected into control (Si-CON) or SiRNA of DNA-PKcs (Si-DNA-PKcs) transfected DLD-1 cells. Cell lysates were immunoprecipitated and immunoblotted.

Figure 4. SP increases E-cadherin promoter activity and inhibits tumor migration. (A) Control (Si-Con) or SiRNA of DNA-PKcs (Si-DNA-PKcs)-treated DLD-1 cells (left) or M059K and MO59J cells (right) with transfection of Control-peptide (CP, 4 μM) or Snail1-peptide (SP, 4 μM) were transiently transfected with E-cadherin promoter construct fused to a luciferase reporter-gene. A luciferase assay was then performed. The E-cadherin promoter activity of CP treated cells was set at 1. Data are presented as mean ± s.d. *p<0.05. (B) After transfection of CP or SP to Si-Con or Si-DNA-PKcs-transfected DLD-1 cells, western blotting was performed. (C) After transfection of CP or SP to Si-Con- or Si-DNA-PKcs-transfected DLD-1 cells, cellular invasiveness was compared. Data are presented as mean ± s.d. *p<0.05 (D) After stable transfection of Snail1-WT or mutant Snail1 S100A plasmids to CT26 colon carcinoma cells, western blot analysis was performed (upper). CT26 cells were intravenously injected through tail vein of the mice and 3 Gy radiation (IR) was administered. Intraperitoneal injection of CP or SP at 3 mg/kg in PBS was performed once before IR and 4 times after IR. The metastasis of CT26 cells was assessed by lung histology, lung weight, the number of lung metastatic nodules, and lung tumor area present 14 days after cell injection. Quantitative analysis of the number of lung weights (upper), metastatic nodules (middle) and tumor area (lower) (Normal, no tumor cell injection; Con, CT26 cell injection; WT, Snail1-WT; S100A, Snail1-S100A). Data are presented as mean ± s.d. of 5 mice from each group. *p<0.05

Figure 5. SP sensitizes IR-mediated cell death. (A) After 48 h of 5 Gy radiation (IR) was applied to DLD-1 cells transfected with Control-peptide (CP, 4 μM) or Snail-peptide (SP, 4 μM), cell death was analyzed by FACS. Data are presented as mean ± s.d. *p<0.05 (B) Western blot analysis was performed using anti-cleaved-caspase 3 and PARP antibodies. (C) CP or SP were transfected into DLD-1 cells with or without SiRNA of DNA-PKcs (Si-DNA-PKcs). A colony forming assay was then conducted after the cells were treated with various doses of radiation. Data are presented as mean ± s.d. *p<0.05

Figure 6. SP shows sensitization of cancer cells in p53 dependent manner. (A) p53+/+ and p53-/- HCT116 cells or SiRNA p53 (Si-p53)-transfected p53+/+ HCT cells were transiently transfected with Control peptide (CP, 4 μM) or Snail peptide (SP, 4 μM) at the indicated time points after exposure of the cells to 5 Gy of radiation (IR). Cell lysates were obtained and immunoblotting was performed.
(upper). The relative protein band intensity was calculated from densitometric scans of immunoblots with the control values set at 1. (n=3, data are presented as mean ± s.d., *p<0.05 vs corresponding CP treated cells). (B) A colony forming assay was then conducted after the cells were treated with various dose of IR. (C) After 48h of 5Gy IR, cell death was analyzed by FACS. Data are presented as mean ± s.d., *p<0.05 (D) Western blot analysis was performed.

Figure 7. A model illustrating the inhibition of Snail1-DNA-PKcs protein-protein interface decreases aneuploidy, sensitizes to IR and inhibits tumor metastasis The peptide sequences with 7 amino acids (KPNYSEL, SP), contained in both Snail1 and Slug, can interact with DNA-PKcs. Treatment with SP blocks the endogenous binding activity between Snail1 and DNA-PKcs, which results in abolishment of DNA-PKcs-mediated Snail1 phosphorylation at Ser100 and Snail protein stability, and finally provides effective sensitization of cancer cells and inhibition of tumor metastasis in both Snail- and DNA-PKcs-overexpressed cancer cells with functional p53.
Figure 2.

A

|         | SI:CON       | SI:DNA-PKcs |
|---------|--------------|-------------|
|         | CP           | SP          | CP           | SP          |
| IR (min)| 0 30 60 90   | 0 30 60 90  | 0 30 60 90   | 0 30 60 90  |
| p-DNA-PKcs (Ser2056) | ![Image](image1) | ![Image](image2) | ![Image](image3) | ![Image](image4) |
| DNA-PKcs | ![Image](image5) | ![Image](image6) | ![Image](image7) | ![Image](image8) |
| p-p53 (Ser15) | ![Image](image9) | ![Image](image10) | ![Image](image11) | ![Image](image12) |
| γ-H2AX | ![Image](image13) | ![Image](image14) | ![Image](image15) | ![Image](image16) |
| Snail1 | ![Image](image17) | ![Image](image18) | ![Image](image19) | ![Image](image20) |
| β-Actin | ![Image](image21) | ![Image](image22) | ![Image](image23) | ![Image](image24) |

B

![Image](image25)

C

WNE

DSB pull down

DNA-PKcs

Ku66

D

|         | Sh:CON       | Sh:DNA-PKcs |
|---------|--------------|-------------|
|         | CP           | SP          | CP           | SP          |
| IR      | + + + +      | + + + +     | + + + +      | + + + +     |
| p-DNA-PKcs | ![Image](image26) | ![Image](image27) | ![Image](image28) | ![Image](image29) |
| Cyclin B1 | ![Image](image30) | ![Image](image31) | ![Image](image32) | ![Image](image33) |
| Snail1  | ![Image](image34) | ![Image](image35) | ![Image](image36) | ![Image](image37) |
| β-Actin | ![Image](image38) | ![Image](image39) | ![Image](image40) | ![Image](image41) |

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Figure 3.

(A) Western blot analysis of p-Snail1 (Ser100) and Snail1 in cells treated with or without siRNA targeting DNA-PKcs. The blots were probed with antibodies against p-Snail1 (Ser100), Snail1, and β-Actin. The graph shows the relative band intensity of p-Snail1 (fold induction) over time (CP and SP conditions).

(B) Western blot analysis of Snail1, β-Actin, and DNA-PKcs in cells treated with or without cycloheximide (CHX). The blots were probed with antibodies against Snail1, DNA-PKcs, and β-Actin. The graph shows the relative band intensity of Snail1 (fold induction) over time (CP and SP conditions).

(C) Western blot analysis of ubiquitin (Ub) and DNA-PKcs in cells treated with or without siRNA targeting DNA-PKcs. The blots were probed with antibodies against Ub and DNA-PKcs. The graph shows the relative band intensity of DNA-PKcs and Snail1 (fold induction) over time (CP and SP conditions).
Figure 4.

A

B

C

D

E-cadherin promoter activity (Fold increase)

E-cadherin (Fold increase)

CT26-Snail1

Con

WT

S100A

Snail1

β-Actin

CPS

PD-1

CP

SP

CT26

Con

WT

S100A

IR (3Gy)

The number of lung tumor nodules

Con

CP

SP

CT26

CT26

Con

CP

SP

WT

S100A

IR (3Gy)

Tumor area

Con

CP

SP

CT26

CT26

Con

CP

SP

WT

S100A

IR (3Gy)
Figure 5.

A

B

C

[Graphs and images showing experimental results related to cell death, protein expression, and survival fraction after different treatments and irradiation levels.]
Figure 7.

DNA-PKcs Positive/\textit{Snail1} Positive cancer cells

IR \rightarrow DNA-PKcs \rightarrow \textit{Snail1} \rightarrow S100 \rightarrow Aneuploidy Resistance to IR \rightarrow Invasion

IR \rightarrow DNA-PKcs \rightarrow \textit{Snail1} \rightarrow S100 \rightarrow E-cadherin \rightarrow Invasion

DNA-PKcs Negative/\textit{Snail1} Positive cancer cells

IR \rightarrow DNA-PKcs \rightarrow \textit{Snail1} \rightarrow E-cadherin \rightarrow Invasion

IR \rightarrow DNA-PKcs \rightarrow \textit{Snail1} \rightarrow E-cadherin \rightarrow Invasion
Inhibition of Snail1-DNA-PKcs Protein-Protein Interface Sensitizes Cancer Cells and Inhibits Tumor Metastasis
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