RPA Inhibition Increases Replication Stress and Suppresses Tumor Growth

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Abstract

The ATR/Chk1 pathway is a critical surveillance network that maintains genomic integrity during DNA replication by stabilizing the replication forks during normal replication to avoid replication stress. One of the many differences between normal cells and cancer cells is the amount of replication stress that occurs during replication. Cancer cells with activated oncogenes generate increased levels of replication stress. This creates an increased dependency on the ATR/Chk1 pathway in cancer cells and opens up an opportunity to preferentially kill cancer cells by inhibiting this pathway. In support of this idea, we have identified a small molecule termed HAMNO ((1Z)-1-[(2-hydroxyanilino)methyldiene]naphthalen-2-one), a novel protein interaction inhibitor of replication protein A (RPA), a protein involved in the ATR/Chk1 pathway. HAMNO selectively binds the N-terminal domain of RPA70, effectively inhibiting critical RPA protein interactions that rely on this domain. HAMNO inhibits both ATR autophosphorylation and phosphorylation of RPA32 Ser33 by ATR. By itself, HAMNO treatment creates DNA replication stress in cancer cells that are already experiencing replication stress, but not in normal cells, and it acts synergistically with etoposide to kill cancer cells in vitro and slow tumor growth in vivo. Thus, HAMNO illustrates how RPA inhibitors represent candidate therapeutics for cancer treatment, providing disease selectivity in cancer cells by targeting their differential response to replication stress. Cancer Res; 74(18); 5165–72. ©2014 AACR.

Introduction

One of the intrinsic features that distinguish cancer cells from surrounding normal cells is a deregulated cell cycle. Common cancer-associated mutations in genes such as p53, Myc, Rb, and EGFR in head and neck squamous cell carcinomas (HNSCC) promote a promiscuous S-phase entry, which leads to higher levels of replication stress (1–6). Replication stress occurs when the replication fork encounters aberrant DNA structures. Although these structures may differ greatly, they all lead to the formation of long stretches of ssDNA that are bound with replication protein A (RPA), the main eukaryote ssDNA binding protein (7, 8).

The major structural feature of heterotrimeric RPA is the presence of the oligonucleotide/oligosaccharide binding (OB) folds within each of its six DNA binding domains (DBD). This OB fold structure consists of β sheets, forming β-barrel structures that wrap around ssDNA, a common feature in ssDNA binding proteins (9, 10). These OB folds are responsible for RPA’s binding to long stretches of ssDNA, making RPA one of the first responders in the replication stress response. RPA has several functions other than stabilizing ssDNA. One of these functions is to recruit and interact with proteins needed for an effective replication stress response. RPA interacts with proteins needed for an effective replication stress response. This triggers a complex network of signaling pathways involved in cell-cycle checkpoints, DNA repair, and apoptosis that mediates the cellular response to replication stress. In the context of cancer therapy, the activation of these signaling pathways can limit the therapeutic efficacy of radiation and chemotherapeutics and enhance the survival of cancer cells (16).

The presence of higher replication stress in cancer cells compared with normal cells provides a therapeutic opportunity to target RPA. On the basis of observations of a DNA-damage response in early-stage cancerous lesions, two laboratories proposed a model referred to as the oncogene-induced DNA-damage model for cancer development, which posited that deregulation of growth regulation genes induces DNA replication stress and DNA damage (16–18). Their initial analyses provided evidence that loss of heterozygosity in
early-stage lesions was due to oncogene-induced replication stress preferentially targeting chromosomal fragile sites. More recent studies have supported these initial observations and provided more evidence of the presence of DNA replication stress in human cancers. These studies were able to examine a very large cancer specimen sample size using high-throughput genomic analyses to show that the most common deletions occur at fragile sites in large genes, and these deletions can be attributed to replication stress (19, 20). Combined, these results provide a strong foundation for developing a novel therapeutic approach that exploits the endogenous replication stress intrinsic to cancer cells (21).

Here, we present the identification and development of a small-molecule inhibitor that inhibits the protein interaction domain of RPA70 involved in the replication stress response. By targeting the cellular response to replication stress via RPA, we demonstrate that this approach is particularly toxic to cells that harbor cancer-associated mutations, which promote resistance to conventional chemotherapies.

Materials and Methods

Reagents

HAMNO, also known as NSC111847, was acquired from the Developmental Therapeutics Program of the NCI. The primary antibody for RPA32 was purchased from Santa Cruz Biotechnology. The primary antibodies for ATR and ATR-PT1989 were purchased from Genetex. Primary antibodies for RPA32-PS4S, RPA32-PS33, and GAPDH were purchased from Bethyl Laboratories. The Alexa Fluor 680 and Alexa Fluor 488 secondary antibodies were purchased from Invitrogen. The IRDye700-labeled polyT 30mer oligonucleotide and complementary polyA 30mer oligonucleotides were purchased from IDT.

Docking simulations

The Molegro Virtual Docker Program (CLC bio) was used to dock HAMNO to the DBD-F in silico. The HAMNO structure used for docking was created using Open Babel. The model of DBD-F chosen for docking was a modified version of the original crystal structure (PDB:2B29) optimized for ligand binding (22, 23). The entire protein structure was used for modeling under default conditions, resulting in five simulating dockings of high predictive affinity. Docking data and results are included in Supplementary Data.

Immunofluorescence and Western blot

For immunofluorescence studies, UMSCC38 cells were grown overnight before drug treatment, fixed and permeabilized with 100% ice-cold methanol for 10 minutes, washed and blocked 10% goat serum and 1% BSA in PBS for 30 minutes at room temperature and the primary antibodies to PS139-H2AX were applied in blocking solution for 1 hour at room temperature. An Alexa Fluor 488-conjugated secondary antibody was then incubated in blocking solution for 1 hour at room temperature. Cells were mounted in PermaFluor (Fisher) supplemented with 0.5 μg/mL 4',6-diamidino-2-phenylindole (DAPI; Roche). Images were captured digitally with a Zeiss Axiovert 200 M microscope, and scored using Mathematica software (Wolfram) as follows. DAPI staining was used to define the region of interest (ROI). H2AX signal within the ROI was determined and background intensity calculated from outside the ROI for each nuclei. A total of 200 cells were assayed per condition. Statistical analysis was then performed with Prism software (Graphpad). For Western blot, whole-cell lysates were separated by SDS–PAGE, blotted onto nitrocellulose membranes, and probed with primary antibodies to RPA32, followed by Alexa Fluor 680–conjugated anti-rabbit. Images were obtained with an Odyssey Imager (LI-COR).

Cells and clonogenic assays

The squamous cell carcinoma cell lines UMSCC38 and UMSCC11B (kindly obtained from Dr. Thomas G. Carey, University of Michigan, Ann Arbor, MI) were propagated in DMEM with 10% FBS. The immortalized primary oral keratinocyte cell line, OKF4 (obtained from Dr. James G. Rheinwald, Harvard Institutes of Medicine, Boston, MA), was propagated in fortified KBM-2 media (Lonza) with 10% FBS (Hyclone). For clonogenic assays, cells were trypsinized and diluted in media to 1,000 cells/mL, then dispersed into 60-mm dishes (3 mL) overnight. After addition of HAMNO, cells were grown for 9 days, then fixed in PBS containing 6% glacialdehyde for 30 minutes, and then dyed in 0.5% crystal violet for 30 minutes and rinsed. Colonies containing over 50 cells were counted. For studies requiring etoposide, HAMNO was added 1 hour before addition of 2 μmol/L etoposide. After 2 hours of etoposide exposure, media were removed and rinsed with PBS, before adding back media containing HAMNO. Data were analyzed using an unpaired 2-tailed Student t test to determine statistical significance.

Protein purification and electrophoretic mobility shift assays

RPA was purified using a published protocol as described (24). DBD-F fused to maltose binding protein was generated and purified as described (22). Quality of both proteins was assessed by SDS–PAGE, followed by coomassie staining (22). For ssDNA binding studies, 7 nmol/L RPA was added to 10 nmol/L labeled polyT 30mer in electrophoretic mobility shift assays (EMSA) buffer (10 mmol/L Tris, pH 7.5, 10 mmol/L KCl, 1% glycerol) for 10 minutes at 25°C. Samples were run on 1% agarose gels in 40 mmol/L Tris-Acetate buffer, pH 7.5, and then scanned on an infrared scanner. For DNA unwinding assays, 14 nmol/L RPA was added to 10 nmol/L PAGE purified annealed poly(A)-polyT 30mer oligonucleotides.

Flow cytometry

Cell-cycle assessment and γ-H2AX staining were monitored in UMSCC38 and OKF4 cells after 2-hour incubation with HAMNO and fixed in 70% ethanol overnight. Cells were washed with PBS and incubated overnight in PBS containing 1% BSA, 10% goat serum and PS139-H2AX antibodies (Millipore), washed and incubated in goat anti-mouse Alexa Fluor 647 antibody for 30 minutes at RT. Cells were incubated in 50 μg/mL propidium iodide and 100 μg/mL RNase A for 30 minutes, and 10,000 cells per sample were analyzed on a
BD FACSAarray (BD Biosciences) using 532- and 635-nm excitations and collecting fluorescent emissions with filters at 585/42 nm and 661/16 nm (yellow and red parameters, respectively). BD FACSAarray and WinList (Verity House) software were used for data collection and analysis, respectively.

Xenograft tumor model

Athymic nude mice were purchased from NIH and housed at the animal facility at the UNMC College of Dentistry. UMSCC38 and UMSCC11B cells were implanted into 6-week-old female mice by a single subcutaneous injection of tumor cells (2–6 x 10^5 cells in 100 mL of sterile PBS). The growth rates of tumors were determined by daily monitoring of tumor volume with vernier calipers [tumor volume = 1/2(length x width^2)]. Once the tumor size reached 50 mm^3, etoposide (10 mg/kg mouse) and HAMNO (2 mg/kg) were administered intraperitoneally every day for 3 days. Tumor size was monitored daily and the volume of the tumor was compared among all experimental groups. At least three mice were used per group. Data were analyzed using an unpaired 2-tailed Student t test to determine the statistical significance.

Results

HAMNO is selective for DBD-F

HAMNO (Fig. 1A) was first identified as an RPA DBD-F inhibitor in a high-throughput screen (HTS) that determined the ability of a small molecule to dissociate a RAD9–GST fusion protein from an RPA–ssDNA complex, an interaction that requires DBD-F (25). Binding of HAMNO to DBD-F was further investigated through in silico methods (Fig. 1B). These studies used a crystal structure of DBD-F (23) that was earlier optimized for binding to the in vitro DBD-F inhibitor, fumaropimaricar-micaric acid (FPA; ref. 22). The site of highest predicted affinity was to a position immediately adjacent to R43 on DBD-F (Fig. 1B, right), where the compound would predictively act to hinder protein–protein interaction, as this residue is essential for DBD-F protein binding (11).

To confirm an interaction of HAMNO with DBD-F in vitro, we took advantage of the ability of DBD-F to weakly bind a labeled oligonucleotide, and looked for changes in mobility and intensity of the complex in the presence of HAMNO using an EMSA (Fig. 1C; ref. 22). In the absence of DBD-F, HAMNO does not bind labeled oligonucleotide (Fig. 1C, left). In the presence of both DNA and DBD-F, the addition of HAMNO results in the formation of a band between the free ssDNA and protein-bound ssDNA bands (Fig. 1C, right; denoted by an asterisk). As HAMNO is uncharged at neutral pH, it is unlikely that binding of the compound to the protein–DNA complex alone would result in this newly formed band detected on the gel. Rather, this stimulation of DBD-F binding to DNA by HAMNO suggests a direct interaction of the small molecule with DBD-F, resulting in a conformational shift that alters DNA binding.

We then wanted to confirm that HAMNO is selective for inhibiting only DBD-F, and not DBDs A–E, which are important in RPA binding to ssDNA. This selectivity would ensure that HAMNO would affect protein recruitment more adversely than ssDNA binding. To determine this, we took advantage of a DNA unwinding activity by RPA that is dependent on the DBD-F, but does not require DBD-F for stable binding of the resultant ssDNA that is formed (22, 26). To do this, we used EMSAs with full-length RPA using ssDNA and dsDNA probes. HAMNO did not affect RPA binding to ssDNA at concentrations up to 200 μmol/L, but prevented dsDNA unwinding and subsequent ssDNA binding at 100 μmol/L (Fig. 1D). This inhibition of dsDNA unwinding by HAMNO is nearly equivalent to that by our previously identified in vitro inhibitor, FPA, whose subsequent dissociation constant for DBD-F was determined to be 9.0 μmol/L (22). Together, these data show a preference of HAMNO for selectively inhibiting DBD-F at micromolar levels, an ability that would predictively target the replication stress response in replication-stressed cancer cells over normal cells.

HAMNO induces γ-H2AX staining in a cell-cycle–specific manner

DBD-F binds ATRIP, RAD9, RAD17, and NBS1, thereby recruiting and stabilizing the proteins involved in ATR activation (11, 13, 15, 27, 28). Inhibition of DBD-F interactions with...
these proteins would likely short-circuit ATR signaling, leading to increased replication stress that can be monitored by pan-nuclear phosphorylation of H2AX in S-phase (29–31). This type of phosphorylation has been shown to be S-phase–specific as previously demonstrated with CHK1 inhibitors (32) and differs from the punctate foci that occur in response to double-strand breaks. We evaluated whether HAMNO induces replication stress as assessed through increases in pan-nuclear γ-H2AX staining (Fig. 2A). After UMSCC38 cells were exposed to HAMNO, increased pan-nuclear γ-H2AX staining occurred in a dose-dependent manner (Fig. 2A and B). When H2AX phosphorylation was assessed via Western blot, cancer-derived UMSCC38 cells, as well as another cancer cell line, UMSCC11B, had prominent γ-H2AX staining, particularly after incubation with 20 μmol/L HAMNO. In contrast, the telomerase-immortalized keratinocyte cell line, OKF4, did not show enhanced γ-H2AX staining at this concentration, suggesting that HAMNO is more effective in inducing H2AX phosphorylation in cancer cell lines that are potentiated for oncogene-induced stress. To further validate that pan-nuclear γ-H2AX represents replication stress and the difference in replication stress between HNSCC cells and OKF4 cells is S-phase–specific, we used flow cytometry. To measure cell-cycle–specific γ-H2AX staining, we assayed cells in the absence of HAMNO as a negative control, which determined the threshold between γ-H2AX–positive and γ-H2AX–negative cells for each cell-cycle phase (Fig. 2D). Both UMSCC38 and OKF4 cells presented increased γ-H2AX staining after addition of HAMNO, with the greatest increase in signal occurring in S-phase (Fig. 2D). In further comparison of the two cell lines, the ratio of γ-H2AX–positive to γ-H2AX–negative cells in S-phase was 6- to 8-fold greater in UMSCC38 cells than OKF4 cells at 20 μmol/L and 50 μmol/L, respectively (Fig. 2E). These data suggest that HAMNO selectively increased γ-H2AX staining in S-phase, indicative of increased replicative stress. These data also indicate that cells predicted to have high levels of oncogene-induced stress, such as mutant p53 squamous cell carcinoma cells, are selectively potentiated for increased replication stress by HAMNO.

**HAMNO affects RPA and ATR phosphorylation**

Inhibition of DBD-F by HAMNO is expected to inhibit RPA32 phosphorylation directly by inhibiting DBD-F interactions with replication stress response proteins involved in activating ATR (11, 12, 15, 27, 28, 33, 34). We tested this hypothesis under conditions that enhance RPA32 Ser33 phosphorylation (Fig. 3A). Ser33 of RPA32, an ATR substrate, is highly phosphorylated after 2 hours of treatment with 20 μmol/L of etoposide, which was reduced with the addition of 2 μmol/L HAMNO, and was nearly absent at higher concentrations, demonstrating an in vivo effect of HAMNO as an inhibitor of RPA32 phosphorylation by ATR.

Ser33 phosphorylation primes further phosphorylation of the N-terminus of RPA32 at other sites, including Ser4 and Ser8 by ATM and DNAPK, a requirement for competent RPA-dependent signaling (35–37). We tested whether inhibition of Ser33 phosphorylation would then inhibit phosphorylation of these downstream sites. As with Ser33, HAMNO also reduced phosphorylation of Ser4 and Ser8 after etoposide treatment (Fig. 3B).

The increase in γ-H2AX staining and decrease in RPA phosphorylation in HAMNO-treated cells suggest a deregulation of ATR signaling. To further describe this loss of competent DNA damage response (DDR) signaling, we assessed the autophosphorylation of ATR at T1989, a marker for ATR activity (Fig. 3C; refs. 38, 39). As expected, etoposide strongly induced ATR autophosphorylation, whereas treatment with HAMNO alone showed a slight increase in T1989 phosphorylation. Interestingly, the addition of etoposide and HAMNO...
combined with etoposide significantly inhibited colony formation to a greater degree than HAMNO alone (Fig. 4B). To further examine the potential of HAMNO as an anticancer agent, we tested HAMNO in a mouse xenograft model. In mice, HAMNO slowed the progression of UMSCC11B tumors (Fig. 4C). This inhibition was also seen in mice cotreated with sublethal etoposide as well as in similarly treated UMSCC38 cells (Fig. 4D and E). These results raise the exciting possibility that a DBD-F inhibitor alone can reduce tumors or can sensitize tumor cells to other chemotherapy agents.

**Discussion**

The inhibition of the ATR–CHK1 pathway has gained prominence as a potential therapeutic intervention for cancer. This strategy has the potential to strike at the disease in a multitude of ways. First, ATR inhibition alone increases replication stress, even in the absence of exogenous DNA damage, which is amplified in cells with active oncogenes or mutated tumor suppressors, resulting in cell death (30, 32). Second, ATR inhibition restrains replication and G1–M checkpoints, prematurely pushing cells along the cell cycle and increasing genomic instability (30). Third, ATR inhibition may act with synthetic lethality with cancers that have an incomplete or incompetent repertoire of DNA repair mechanisms (40). Therefore, the inhibition of ATR and CHK1 will have powerful consequences in the cell.

One major disadvantage of ATR–CHK1 inhibition is the off target effects that are not related to the replication stress response or the DDR, as both kinases phosphorylate hundreds of proteins, many of which are not DDR related (41, 42). CHK1 expression is essential for mouse embryonic cell survival, indicating that ATR and CHK1 inhibition would negatively affect normal cells (43). ATR and CHK1 null mice are embryonic lethal, further implicating that the inhibition of ATR–CHK1 directly may have unwanted consequences in the organism (44, 45). Therefore, a downstream ATR substrate that specifically modulates the replication stress response, such as RPA, may be a more appropriate target for cancer treatment. Inhibition of RPA phosphorylation by HAMNO allows the replication stress response–specific aspects of ATR signaling to be inhibited, allowing other aspects of ATR signaling to remain unimpeded.

Before the discovery of RPA protein inhibitors, inhibitors of RPA–ssDNA interaction have been characterized and have been shown to be cytotoxic and act with synergy with cisplatin, displaying the potential for RPA as a drug target (46, 47). The discovery of several other DBD-F binding proteins further increased the potential of such an inhibition (11, 14). Our pursuit of DBD-F inhibitors through HTS resulted in the first-identified in vitro inhibitor, FPA (25). FPA inhibited the DBD-F with high affinity and selectivity, but had a preponderance of high negative charge that restricted the compound from crossing cell membranes (22, 25). Souza-Fagundes and colleagues also used both HTS and a fragment-based nuclear magnetic resonance spectroscopy method to identify in vitro inhibitors of RPA with affinities similar to FPA (48, 49). HAMNO, with its neutral charge and hydrophobic ring structures, allows passage of the compound through the cell membrane, where it can bind to DBD-F, an improvement over FPA.

The ability of HAMNO to work effectively with etoposide attests to the effective strategy of inducing replication stress and reducing the replication stress response to increase cell death selectively in cancer cells that have constitutive DNA replication stress. This approach would be beneficial in the clinic, as the therapeutic efficacy would increase with the addition of an RPA inhibitor and reduce unwanted side effects. HAMNO also has the potential to be used as a stand-alone agent due to its ability to selectively increase cytotoxicity in cancer cells that already have oncogene-induced replicative stress.
This is the first in vivo study showing the potential of an RPA DBD-F inhibitor as a cancer chemotherapeutic agent. The ability of HAMNO to induce cytotoxicity alone and kill cells synergistically with etoposide suggests that HAMNO could be used alone or in combination with other chemotherapeutic agents. HAMNO is tolerated in mice at doses that affect tumor growth, attesting to the clinical potential of this compound. The structure of HAMNO allows for the addition and substitution of many possible moieties that could result in increased affinity to the DBD-F. We are currently testing several HAMNO derivatives for increased DBD-F inhibition.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

References


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