Removal of uracil by uracil DNA glycosylase limits pemetrexed cytotoxicity: overriding the limit with methoxyamine to inhibit base excision repair

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Uracil DNA glycosylase (UDG) specifically removes uracil bases from DNA, and its repair activity determines the sensitivity of the cell to anticancer agents that are capable of introducing uracil into DNA. In the present study, the participation of UDG in the response to pemetrexed-induced incorporation of uracil into DNA was studied using isogenic human tumor cell lines with or without UDG (UDG+/+/UDG−/−). UDG−/− cells were very sensitive to pemetrexed. Cell killing by pemetrexed was associated with genomic uracil accumulation, stalled DNA replication, and catastrophic DNA strand breaks. By contrast, UDG+/+/ cells were >10 times more resistant to pemetrexed due to the rapid removal of uracil from DNA by UDG and subsequent repair of the resultant AP sites (abasic sites) via the base excision repair (BER). The resistance to pemetrexed in UDG−/− cells could be reversed by the addition of methoxyamine (MX), which binds to AP sites and interrupts BER pathway. Furthermore, MX-bound AP sites induced cell death was related to their cytotoxic effect of dual inactivation of UDG and topoisomerase IIX, two genes that are highly expressed in lung cancer cells in comparison with normal cells. Thus, targeting BER-based therapy exhibits more selective cytotoxicity on cancer cells through a synthetic lethal mechanism.

Keywords: UDG, uracil DNA glycosylase; BER, base excision repair; MX, methoxyamine; Topo II α, topoisomerase II α; PCNA, proliferating cell nuclear antigen; RPA, replication protein A

Subject Category: Cancer

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UDG is a conserved DNA repair protein expressed in all types of human cells. It specifically removes uracil from DNA and protect cells from cytotoxicity and mutagenicity. Human UDG is encoded by the UNG gene. Alternative promoter usage and splicing of this gene produces two different isoforms: the mitochondrial UNG1 and the nuclear UNG2. Nuclear UDG (UNG2) is the predominant form in cells and represents >90% of the total enzyme activity. Therefore, UDG used in this article refers to UNG2. Nuclear UDG activity is subject to cell cycle-dependent regulation and shows a marked increase during the S-phase. During the S-phase, UDG is localized in replication fork and interacts with PCNA (proliferating cell nuclear antigen) and RPA (replication protein A), two proteins that are required to form functional replication forks. This suggests that the UDG removal of incorporated uracil may directly link to the progression of the replication fork. In addition, UDG has recently been shown to promote the assembly of human centromere protein A (CENP-A). As CENP-A is an essential protein required for chromosome segregation during mitosis, the association between UDG and CENP-A implies that UDG may be involved in cell proliferation.

The base excision repair (BER) pathway is initiated following the removal of a base lesion by a DNA glycosylase. Glycosylase excision of the damaged base proceeds via hydrolytic cleavage of the N-glycosidic bond, leaving the sugar–phosphate backbone intact and producing an abasic site (AP site). The resultant AP site is processed by AP endonuclease 1 (APE1), which generates a single-strand break containing a 5′-dRP residue and a 3′-hydroxyl group. The completion of BER occurs by short-patch (single nucleotide) or long-patch (2–13 nucleotides) repair pathways involving DNA polymerase gap filling and DNA ligation. BER is the most efficient mechanism for repairing a variety of base lesions, including those induced by chemotherapeutic agents, thereby rendering tumor cells resistant to DNA-damaging chemotherapeutic agents. To overcome BER-confferred drug resistance, methoxyamine (MX) has been developed as an active inhibitor of BER. MX reacts specifically with the aldehyde group in the sugar moiety of the AP site, forming an MX-bound AP-site. This structurally modified AP site is refractory to the repair activity of APE1 leading to the interruption of the BER pathway. MX has been shown to potentially enhance the therapeutic efficacy of various
DNA-damaging chemotherapeutic agents.\textsuperscript{22–25} It is currently being evaluated in multiple clinical trials (as TRC102, TRACON Pharmaceuticals, Inc., San Diego, CA, USA) and has completed phase-I testing in combination with pemetrexed.

In the present study, we investigated the impact of UDG and BER on cell sensitivity to pemetrexed using isogenic UDG-proficient and -deficient human cancer cells. Although pemetrexed has multiple targets, the different responses to pemetrexed UDG has multiple targets, the different responses to pemetrexed UDG and has completed phase-I testing in combination with pemetrexed TRACON Pharmaceuticals, Inc., San Diego, CA, USA) being evaluated in multiple clinical trials (as TRC102, UDG activity determines the level of uracil retained in DNA. To correlate UDG activity with the cytotoxicity of pemetrexed, comparative studies were performed in UDG\textsuperscript{+/+} and UDG\textsuperscript{−/−} cells. We first confirmed the UDG activity in these cells using an \textit{in vitro} glycosylase cleavage assay, in which oligonucleotide substrates containing uridine residues were incubated with either purified UDG/APE1 enzymes or cell extracts. As shown in Figure 1a, after the reaction with fluorescent probe-labelled oligonucleotide substrates (40-mer) containing U:G mispairs, both purified UDG/APE1 enzymes and cell extracts from UDG\textsuperscript{+/+} cells produced cleaved DNA fragments as an 18-mer band, which resulted from the removal of uracil bases by UDG and subsequent incision of the resultant AP sites by APE1. By contrast, no cleaved fragments were observed in UDG\textsuperscript{−/−} cell extracts after incubation with an even higher concentration of cell extracts. D\textsuperscript{flag} cells were capable of removing uracil bases, which were derived from UDG\textsuperscript{−/−} cells by restoring UDG activity.

We next determined the levels of uracil in the DNA of UDG\textsuperscript{+/+} and UDG\textsuperscript{−/−} cells following pemetrexed exposure using HPLC/MS/MS. There was an inverse relationship between UDG activity and the level of uracil bases in the DNA (Figure 1b). A significant amount of uracil was detected in UDG\textsuperscript{−/−} cells, which correlated with the duration of cell extracts (5–10 \(\mu\)g) from UDG\textsuperscript{+/+}, D\textsuperscript{flag}, and UDG\textsuperscript{−/−} cells at 37 °C for 1 h. Reaction products were resolved by electrophoresis through denaturing 20% polyacrylamide gels. (b) Incorporated uracil detected in UDG\textsuperscript{+/+} and UDG\textsuperscript{−/−} cells by HPLC/MS/MS analysis. Cells were treated with pemetrexed (10 \(\mu\)M) for 6, 24, 48, and 72 h. Cells were harvested and 40 \(\mu\)g of extracted DNA were incubated with either purified UDG/APE1 enzymes or cell extracts after reacted with purified UDG. Results are representative of three independent experiments.

**Results**

UDG activity determines the level of uracil retained in DNA. To correlate UDG activity with the cytotoxicity of pemetrexed, comparative studies were performed in UDG\textsuperscript{+/+} and UDG\textsuperscript{−/−} cells. We first confirmed the UDG activity in these cells using an \textit{in vitro} glycosylase cleavage assay, in which oligonucleotide substrates containing uridine residues were incubated with either purified UDG/APE1 enzymes or cell extracts. As shown in Figure 1a, after the reaction with fluorescent probe-labelled oligonucleotide substrates (40-mer) containing U:G mispairs, both purified UDG/APE1 enzymes and cell extracts from UDG\textsuperscript{+/+} cells produced cleaved DNA fragments as an 18-mer band, which resulted from the removal of uracil bases by UDG and subsequent incision of the resultant AP sites by APE1. By contrast, no cleaved fragments were observed in UDG\textsuperscript{−/−} cell extracts after incubation with an even higher concentration of cell extracts. D\textsuperscript{flag} cells were capable of removing uracil bases, which were derived from UDG\textsuperscript{−/−} cells by restoring UDG activity.

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**Figure 1** UDG activity determines the levels of uracil and AP sites in DNA. (a) UDG activity assay in vitro. Oligonucleotide duplexes containing U:G were incubated with cell extracts (5–10 \(\mu\)g) from UDG\textsuperscript{+/+}, D\textsuperscript{flag}, and UDG\textsuperscript{−/−} cells at 37 °C for 1 h. Reaction products were resolved by electrophoresis through denaturing 20% polyacrylamide gels. (b) Incorporated uracil detected in UDG\textsuperscript{+/+} and UDG\textsuperscript{−/−} cells by HPLC/MS/MS analysis. Cells were treated with pemetrexed (10 \(\mu\)M) for 6, 24, 48, and 72 h. Cells were harvested and 40 \(\mu\)g of extracted DNA were incubated with either purified UDG/APE1 enzymes or cell extracts after reacted with purified UDG. Results are representative of three independent experiments.
pemetrexed exposure. By contrast, the detectable uracil in the DNA was very low in UDG+/+ cells, suggesting rapid and efficient removal of the incorporated uracil. Similarly, a greater retention of uracil in UDG−/− cells than in UDG+/+ cells (Figure 1c) was detected following exposure to 5-fluorouracil (5-FU), a well-known thymidylate synthase inhibitor capable of introducing uracil into DNA through imbalanced nucleotide pools.

The AP sites formed by pemetrexed were measured in cells, which are a surrogate marker for UDG activity in the cells. As shown in Figure 1d, a dose-dependent formation of AP sites in DNA was detected in UDG+/+ but not in UDG−/− cells. The lack of detectable AP sites in UDG−/− cells is presumably due to the absence of UDG activity to remove uracil from the DNA, resulting in the accumulation of uracil in DNA. To confirm this, pemetrexed-induced AP sites in UDG−/− DNA were analyzed after incubation with purified UDG enzymes in vitro. As shown in Figure 1e, increased AP sites were detected when UDG activity was restored in D flag cells. Similarly, increased AP sites were also detected in UDG−/− cells treated with 5-fluorouracil (5-FU), a well-known thymidylate synthase inhibitor capable of introducing uracil into DNA through imbalanced nucleotide pools.

UDG activity determines cell sensitivity to pemetrexed. To determine the correlation between the expression of UDG and the sensitivity to pemetrexed, the cytotoxic effect of pemetrexed was examined in UDG+/+ and UDG−/− cells with a clonogenic assay. UDG−/− cells were 10 times more sensitive to pemetrexed than UDG+/+ cells (Figure 2a). The IC₅₀ value for pemetrexed was 20 nM in UDG−/− cells, compared with 210 nM in UDG+/+ cells. The killing effect of pemetrexed in UDG−/− cells was reversed when UDG activity was restored in D flag cells. Similarly, UDG+/+ cells were more sensitive to 5-FU than either UDG+/+ or D flag cells (Figure 2b). By contrast, there was no significant difference in sensitivity to temozolomide, an alkylating agent, or cisplatin, a crosslinking agent (Figures 2c and d), between UDG+/+ and UDG−/− cells. Thus, results suggest that UDG activity specifically impacts the cytotoxicity of anticancer agents that are capable of inducing the incorporation of uracil bases in DNA.

The accumulation of incorporated uracil in DNA stalls DNA replication. To elucidate the underlying mechanisms responsible for pemetrexed’s cytotoxicity, we first examined cell cycle progression in response to pemetrexed. As shown in Figure 3a, pemetrexed (25 nM) caused the arrest of ~25% and 36% of UDG−/− cells in the S-phase at 6 and 24 h, respectively (Figure 3a). The S-phase arrest lasted more than 72 h. At this time point, 35% of the cells were still in the S-phase and ~22% of the cells had undergone apoptosis (subG1). By contrast, progression of the cell cycle in UDG+/+ cells was only slightly affected by this dose of pemetrexed (Figure 3b). The effect of pemetrexed on cell killing was further analyzed by annexin V staining at 24 h after cells were treated with 50 nM of pemetrexed. Results revealed that pemetrexed at the same concentration induced 30% of cell death in UDG−/− cells compared with 5% in UDG+/+ cells (Figures 4a and b).

We next examined a network of proteins responsible for DNA damage checkpoints following exposure to pemetrexed. We found that phospho-Chk1 (Ser345) was significantly increased in UDG−/− cells (Figure 3c) but not in UDG+/+ cells (Figure 3d). As a DNA damage checkpoint kinase, the induction of phospho-Chk1 suggests that pemetrexed stalls replication forks in UDG−/− cells. We also noted that in both cell lines, phospho-histone H3 (Ser 10), a mitotic marker, increased at 6 h and decreased at 24 h following exposure to pemetrexed. Interestingly, the periodic fluctuation of phosphorylation of histone H3 was not directly associated with mitotic cell cycle progression. In addition, the induction of phospho-histone H3 was concomitant with the increase in G1–S-phase–specific cyclin D1. Subsequently, the S-phase–specific cyclin E was upregulated following the exposure to pemetrexed (Figures 3c and d). It has been reported that inhibition of cell proliferation can regulate histone H3 phosphorylation and that this modification enables the transcription of genes that are activated as a consequence of a variety of cell-signaling events. Therefore, our data may suggest that phosphorylation of histone H3 is an important regulatory mechanism of gene transcription. However, whether the phosphorylation of histone H3 is linked to transcriptional activation of cyclin proteins in response to DNA damage induced by pemetrexed needs to be further investigated. Although the role of cyclin D1 and E remains to be fully elucidated, they function as the regulators of the G1–S phase entry in response to inhibition of replication-stalled S-phase. Furthermore, the upregulation of phospho-cdc2 and cyclin B1 also indicates that pemetrexed-induced DNA damage activates the S- and M-phase checkpoints.

In addition, pemetrexed induced an increase in the expression of topoisomerase IIα (topo IIα) that was much greater in UDG−/− cells than in UDG+/+ cells.
The induction of topo IIα may be associated with either a global signal of DNA damage or a more specific response to the S-phase arrest. As expected, γH2AX was markedly increased in UDG−/−/− cells, which is a well-known marker of DNA double-strand breaks (DSBs) and an indicator of stalled replication forks and replication fork collapse. In company with the induction of γH2AX, a significant cleaved PARP (a hallmark of apoptotic cell death) was detected in UDG−/−/− cells. The activation of caspase 3, 8 and 9 (Figures 4c and d) was then measured. Those are considered to be the key effect of proteases on apoptosis through the degradation of numerous cellular substrates including the cleavage of PARP. Surprisingly, the pemetrexed-induced PARP cleavage appeared to be independent of the caspase pathway. Moreover, survivin was noted to be upregulated in response to pemetrexed. As a member of the inhibitor of apoptosis protein family, survivin has been reported to bind to and inhibit effector caspase-3 and -7. Although the direct effect on caspases still remain to be further determined, the induction of survivin may link to the inhibition of caspase 3. Taken together, our results suggest that pemetrexed-induced apoptotic cell death may not be mediated through the caspase pathway or that other cell death regulators are more important in pemetrexed-treated cells.

We further investigated whether uracil-DNA lesions could inhibit DNA replication. Cells were sequentially labeled with the halogenated nucleotides chlorodeoxyuridine (CldU) and iododeoxyuridine (IdU) to stain replication foci. Representative cells are depicted in Figures 5b and c. Compared with replication foci stained with CldU and IdU in untreated cells, a significant reduction in IdU incorporation was observed in UDG−/−/− cells following a 6-h exposure to pemetrexed and a further 18-h after incubation, suggesting that pemetrexed-induced accumulation of uracil has the potential to inhibit DNA replication. By contrast, there was no significant change in CldU and IdU incorporation in the replication foci of UDG+/+ cells before and after pemetrexed treatment (25 nM). The fluorescence density of IdU or CldU was quantified in UDG−/−/− and UDG+/+ cells by using the NIH Image J software (NIH, Bethesda, MD, USA; Figure 5d). The values of the ratio of IdU to CldU indicate that accumulated uracil bases in DNA block DNA replication in UDG−/−/− cells.

Blocking BER enhances pemetrexed cytotoxicity. We have shown above that the lack of UDG activity sensitizes tumor cells to antimetabolites. However, UDG deficiency appears to be surprisingly rare in human tumors. In fact, human lung cancer express higher UDG than its corresponding normal tissues. Thus, the removal of
uracil by UDG and subsequent BER activity limits the efficacy of pemetrexed in lung cancer treatment. We next studied a therapeutic strategy to override the UDG-conferred resistance to pemetrexed by interrupting the BER pathway using MX. Cytotoxicity was measured in UDG+/+ and UDG−/− cells after treatment with pemetrexed alone or in combination with MX. MX greatly sensitized UDG+/+ cells to pemetrexed and reduced the pemetrexed IC50 value from 220 to 80 nM (Figure 6a). By contrast, no differential sensitivity between pemetrexed alone and in combination with MX was observed in UDG−/− cells (Figure 6b). The failure of MX to potentiate pemetrexed toxicity in UDG−/− cells can be explained by the fact that, in UDG−/− cells, there are no DNA-binding sites available for MX due to the absence of AP sites in the DNA (Figure 1d). Similar experiments were performed in the human nonsmall cell lung cancer cell lines H460 and A549 (Figures 6c–e). These two cell lines retain wild-type p53 and harbor a mutation in K-ras but express different levels of UDG. Western blotting revealed that UDG protein levels in A549 were approximately 9- and 17-fold higher than in H460 cells and normal lung epithelial cells, respectively (Figure 6d). A549 cells were obviously more resistant to pemetrexed than H460 cells. IC50 values for pemetrexed were 1200 nM in A549, compared with 110 nM in H460 cells. MX was capable of enhancing pemetrexed cytotoxicity in both cell lines four- to fivefold (Figure 6e). Thus, although multiple mechanisms may confer resistance to pemetrexed, our results indicated that UDG activity in lung cancer cells is an important factor in pemetrexed-resistance. This was further confirmed by the studies, in which UDG, in H460 cells, was knocked down by siRNA, resulting in the increase in cytotoxicity by threefold (data not shown). Importantly, MX reverses this resistance.

AP sites were detected in H460 cells following treatment with pemetrexed. As shown in Figure 7a, the formation of AP sites increased as the concentration of pemetrexed increased. Co-treatment with MX formed MX-bound AP sites, resulting in the reduction of ARW (aldehyde-reactive probe)-detected AP sites. This is because ARW and MX react competitively with the aldehyde group in AP sites and binding of MX to the AP sites makes them unavailable for ARW binding (Figure 7a). Furthermore, the levels of UDG protein were significantly induced in cells treated with the combination of pemetrexed and MX (Figure 7b). Immunofluorescent staining revealed that the UDG protein was increased in both the cytosol and the nucleus but predominantly accumulated in the nucleus at 24 h after treatment with the combination (Figure 7c), suggesting that MX-bound AP sites were able to

Figure 4 Pemetrexed-induced apoptotic death is independent of caspase pathway. The percentage of cell death induced by pemetrexed using Annexin V staining in UDG−/− (a) and UDG+/+ cells (b). Protein alterations involved in regulation of apoptotic cell death were detected in (c) UDG−/− cells and (d) UDG+/+ cells treated with pemetrexed. Cells were collected at 6 and 24 h after pemetrexed treatment. Results are representative of two independent experiments
trap or stabilize UDG in DNA. In addition, concomitant inductions of topo IIα, γH2AX, and cleaved PARP were seen in cells treated with the combination of pemetrexed and MX (Figures 7d and e). These results could be explained by our previous findings that MX-bound AP sites were capable of poisoning topo IIα and inducing topo IIα-mediated DNA DSBs,33 triggering apoptosis.

MX potentiates the anti-tumour effects of pemetrexed in vivo. The MX-potentiated antitumour effect of pemetrexed was further tested in vivo using human lung cancer xenografts. As shown in Figure 8, both H460 and A549 tumors were moderately sensitive to pemetrexed alone. However, the antitumour activity of pemetrexed was significantly enhanced by the combination of pemetrexed (150 mg/kg) and MX (4 mg/kg) in these two xenograft tumors. At 15 days, H460 xenografts treated with PBS (control group) had a mean tumor volume of 2100 ± 106 mm³, compared with a mean tumor volume of 1726 ± 176 mm³ or 543 ± 82 mm³ (*P-value < 0.02) in mice treated with either pemetrexed alone or in combination with MX, respectively. Similarly, MX-potentiated antitumor effect of pemetrexed was observed in A549 xenografts, which was consistent with the in vitro data. In addition, at these doses, mice did not present evidence of systemic toxicity as evaluated by body weight measurements and complete blood count tests (data not shown).

Discussion

The current study provides evidence that UDG has a profound impact on the cytotoxicity of uracil incorporation into DNA induced by pemetrexed. Because the expression of UDG in lung cancer tissue is high but varies between individuals, UDG activity may be exploitable as a biomarker for predicting susceptibility and response to pemetrexed-related treatment. Moreover, our results demonstrated that the loss of UDG results in the accumulation of uracil in DNA, which stalls replication forks and subsequently induces replication

![Figure 5](https://example.com/figure5.png)
collapse, leading to remarkable increases in the killing effect of pemetrexed. These studies lead us to propose that, as a therapeutic target, UDG may offer new avenues for the development of novel anticancer agents.

UDG removal of incorporated uracil links the DNA replication process. The progression of the DNA replication fork can be stopped by various factors, including DNA damage, protein–DNA complexes, and depletion of...
UDG activity determines the levels of AP sites, key targets of MX. When uracil bases are present in DNA, UDG makes almost all of its contacts with the uracil-containing DNA strand. The binding of UDG transiently forces DNA to undergo local distortions to flip out the uracil base. By contrast, normal bases resist the distortion, so that the UDG active site only allows uracil to bind in extra helical DNA, facilitating the UDG enzymatic excision to produce AP sites. Interestingly, UDG remains bound to the AP site after release of the uracil base, making them unrecognizable by APE1, blocking the crosstalk of APE1 with other BER proteins and interrupting BER activity. Importantly, the failure of UDG-mediated repair may directly and indirectly affect the progression of DNA replication, leading to cell death. The present study demonstrates that targeting UDG activity on pemetrexed cytotoxicity.

Targeting UDG/Ber and Inducing Apoptosis in Cancer Cells. UDG activity is critical for MX-potentiation of pemetrexed cytotoxicity. The combination of MX and UDG enhances pemetrexed cytotoxicity by the dual mechanism of blocking DNA replication and inducing apoptosis. MX inhibits DNA replication by producing AP sites, which are recognized by UDG, leading to the formation of protein–DNA complexes and the inactivation of UDG activity. This results in the accumulation of AP sites, which are then recognized by APE1, leading to the formation of protein–DNA complexes and the inactivation of APE1 activity. This dual mechanism is mediated by the binding of UDG to MX, which stabilizes the AP site and blocks its recognition by APE1. Additionally, MX inhibits DNA repair by targetting UDG, which is essential for MX-potentiation of pemetrexed cytotoxicity.

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inactivation of UDG and topo IIα, through trapping in MX-AP sites. As the expression of UDG and topo IIα is much higher in tumor cells than in normal bone marrow cells,33,36 the MX-potenti…

Materials and Methods

Cells and reagents. Stable and complete knockdown of UDG expression in DLD1 colon cancer cells (DLD1/UDG−/−) cells was achieved by homologous recombination, as described by Zhang et al.41 Briefly, homologous recombination resulted in insertion of a DNA construct containing a series of stop codons within Exon 1 of the UNG gene to interrupt transcription of both the mitochondrial and nuclear UNG isoforms. Recombinant virus was grown in 293T cells and subsequently used to infect DLD1 cells. Selection of positively transfected clones was achieved by the addition of G418 to the culture medium. A second infection of transfected clones with Cre-recombinase adenovirus caused excision of the neomycin-resistant cassette. D1/H3 cell line was produced by transfection of UDG expression vector in UDG−/− cells to restore UDG activity.

Pemtrexed was obtained from LC Laboratories (Woburn, MA, USA), 5-Fluorouracil and cisplatin were purchased from Sigma-Aldrich (St. Louis, MO, USA) and temozolomide was purchased from Ochem Inc. (Des Plaines, IL, USA). Uracil DNA glycosylase was purchased from New England Biolabs (Ipswich, MA, USA) and APE1 was purchased from Trevigen (Gaithersburg, MD, USA). MX, uracil, and uracil-1,3,14-N2 were purchased from Sigma-Aldrich. MX was dissolved in sterilized water (pH 7.0) at a stock solution of 2.5 M and stored at −20°C. Working solutions were generated before experimental use. Fluorescent dye-labeled 40-mer oligonucleotides containing U:G mispairs were purchased from Operon Biotechnologies (Huntsville, AL, USA).

Colonial survival assay. Tumor cells (500–2000/dish) were plated and treated with pemtrexed (0–400 nM), 5-FU (0–20 μM), temozolomide (0–1500 μM) or cisplatin (0–40 μM). After 7 days, surviving colonies were stained with methylene blue for 30 min at room temperature and the colonies containing more than 50 cells were counted to generate survival curves.

Glycosylase activity assay. UDG activity (purified protein or whole cell extract) was measured using an oligodeoxynucleotide containing a single uracil. S′-HEX[GTAAACGAGCGCCAGTGATTCGGAGCTCGTGATACCGG/GG C′-CATTTTGTGCCCGTGACACGGAGCTCGATGGGCCC(CCT)[Cy5]. The fluorescent dye-labeled duplex oligonucleotides were incubated with purified UDG at 37°C for 30 min, followed by 30 min incubation with APE1 or with whole-cell extract. The reaction was stopped by incubation at 95°C for 5 min. Reaction products, 18-mer fragments, were resolved by electrophoresis on denaturating 20% polyacrylamide gels (7 M urea, 1 × Tris-borate-EDTA). Visualization and quantitation were achieved using a Typhoon 9200 fluorescence imager (Amersham Bioscience, Piscataway, NJ, USA). UDG activity was determined based on fluorescence density quantified using the ImageQuant software (Amersham BioScience).

AP site assay. The number of AP sites was measured using ARP reagent. The assay was performed as previously described.42 Briefly, cells (2 × 106) were collected after drug treatment and DNA was extracted. DNA (10 μg) was incubated with 15 μl of 1 mM ARP (Dojindo Laboratories, Kumamoto, Japan), and then precipitated and washed with ice-cold ethanol. The ARP-labeled DNA was then heat-denatured at 100°C for 5 min, quickly chilled on ice and mixed with an equal volume of 2 M ammonium acetate. The DNA was then immobilized on a BA-S 85 nitrocellulose membrane (Schleicher and Schuell, Dassel, Germany) using a minifold II vacuum filter device (Schleicher and Schuell). The membrane was incubated with 0.25% BSA-PBS containing streptavidin-conjugated horseradish peroxidase (BioGenex, San Ramon, CA, USA) at room temperature for 40 min with gentle shaking. ARP-labeled AP sites were visualized by chemiluminescence (Amersham) followed by quantitative densitometry using the NIH ImageJ software.

Detection and quantification of uracil using HPLC/MS/MS analysis. UDG−/− and UDG+/− cells were exposed to pemtrexed (10 μM) or 5-FU (10 μM) for 6, 24, 48 and 72 h. At indicated time points, cells were harvested and genomic DNA was extracted by phenol/chloroform. DNA (40 μg) was incubated with 10 U of purified UDG (New England Biolabs) in 60 μl of reaction buffer at 37°C for 2 h. The reaction products were dried at 35°C in a Turboblow under a stream of nitrogen and reconstituted in 150 μl 90% acetonitrile. The analysis was retained by an Atlantis HILIC Silica analytical column (Waters Corporation, Milford, MA USA) (2.1 × 100 mm, 3.5 μM) and eluted isocratically by a mixture of 90% acetonitrile and 10% 0.2 mM ammonia formate at a flow rate of 0.2 ml/min. The detection was carried out by an API 3200 MS/MS mass spectrometer (AB SCIEX, Foster City, CA, USA).

Immunofluorescence microscopy. Cells were grown on coverslips and were treated with drugs for 6–24 h. Then cells were fixed in 2% paraformaldehyde and permeabilized with 0.2% Triton X-100. Cells were incubated with primary antibody (Abcam, Cambridge, MA, USA) or H2AX (Bethyl, Montgomery, TX, USA) antibody for 1 h at room temperature, followed by incubation with a secondary antibody conjugated with Alexa 488 (green) (Molecular Probes, Carlsbad, CA, USA). The nucleus was stained using Hoechst 33258 for 15 min at room temperature. Images were digitally captured using an Olympus microscope (Olympus, Westmont, IL, USA) equipped with a digital camera.

Cell survival curve analysis. For cell cycle analysis, 106 cells (DLD1/UDG−/− or DLD1/UDG+/+) were plated in 100-mm tissue culture dishes and exposed to pemtrexed (25 nM). After 6, 24, and 72 h of culture, cells were fixed with 80% methanol and washed with ice-cold 1% BSA/PBS. DNA was stained with 20 μg/ml propidium iodide (Sigma-Aldrich) and 2.5 μg/ml of DNase-free RNaseA (Roche, Branford, CT, USA). The DNA fluorescence of propidium iodide–stained cells was measured with an Elite ESP flow cytometer/cell sorter (Coulter, Miami, FL, USA).

Immunofluorescence microscopy of replication foci stained with CiDu and IdU. DNA replication sites were visualized by incorporation of CiDu and IdU into DNA. UDG−/− and UDG+/− cells were labeled with 100 μM CiDu (ICN, Irvine, CA, USA) or IdU (Sigma-Aldrich) for 45 min at different time intervals. Cells were washed with PBS, fixed with cold 70% ethanol and stored at 4°C. For antibody staining, the ethanol was removed, and 100% methanol was added for 5 min. Cells were washed twice with PBS and incubated with 1.5 μM HCl for 30 min to denature the DNA. Cells were washed with PBS, permeabilized with 0.5% Tween-20 in PBS for 5 min, and then incubated in 5% BSA (Sigma-Aldrich) and in PBS for 20 min to reduce nonspecific binding. Primary antibodies CiDu (rat anti-BrdU; Accurate Chemical and Science Co., Westbury, NY, USA) and IdU (mouse anti-BrdU; BD Biosciences) were diluted in 1% BSA PBS buffer, added to the slides, and incubated in a humid environment for 2 h. Slides were washed with PBS-Tween20 and then in a high-salt buffer (200 mM NaCl, 0.2% Tween-20, and 0.2% NP-40 in PBS) for 15 min. The samples were incubated with secondary antibodies for 1 h. Finally, slides were washed with PBS-Tween 20, mounted with Vectashield antifade mounting medium (Vector Laboratories, Inc., Burlingame, CA, USA), and stored at 4°C. Images were visualized on a Nikon Eclipse TE-2000 confocal microscope (Jenoptik Laser Technologies Corp., Brighton, MI, USA). Fluorescence density of CiDu or IdU was quantified by the NIH Image J software.

Xenograft tumors in nude mice. H460 or A549 tumor cells (5 × 106) were injected into bilateral flanks of female athymic nCr (nu/nu) mice (6 weeks old). When the tumor volumes reached 100–150 mm3, mice were divided into control and treatment groups (6–9 mice/group). Nude mice carrying tumors were treated with pemtrexed (150 mg/kg) alone, MX (4 mg/kg) alone, or the combination of the two agents, by daily intraperitoneal injection (i.p.) for 5 consecutive days. Tumor measurements were taken every 2 days. Tumor responses were quantified by tumor volume.
Statistical analysis. Results are presented as the mean ± s.e.m. with significance calculated by Student’s t-test with standard software (GraphPad Prism, San Diego, CA, USA). Significance was assigned for a P-value < 0.05.

Conflict of Interest

Stanton Gerson and Lili Liu own a patent on the methoxyamine. The intellectual property of methoxyamine is licensed by Tracon Pharmaceutical Inc.

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