Synergistic antitumor effect between vorinostat and topotecan in small cell lung cancer cells is mediated by generation of reactive oxygen species and DNA damage-induced apoptosis

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Abstract
The topoisomerase-I (topo-I) inhibitor topotecan, derivative of camptothecin, is the only registered drug for relapsed small cell lung cancer (SCLC). The histone deacetylase inhibitor vorinostat has shown preclinical and clinical antitumor activities in hematologic malignancies and solid tumors, including SCLC, and has recently been approved for the treatment of cutaneous T-cell lymphomas. In this study, we analyzed the antitumor effect of vorinostat combined with topotecan or camptothecin in topo-I inhibitor-sensitive H209 and inhibitor-resistant H526 SCLC cells. Simultaneous or sequential exposure (24 h delay) to either agent resulted in strong synergistic cytotoxic effect in both cell lines, as shown by calculating combination index, and confirmed by growth in soft agar. Combination treatments increased S-phase cell cycle arrest paralleled by apoptosis as measured by hypodiploid peak formation, Annexin V binding, DNA fragmentation, and mitochondria destruction. The apoptotic process was triggered by a caspase-dependent mechanism and can be ascribed to the phosphorylation of H2AX, a reporter of DNA double-strand breaks. These effects were paralleled by an increase of topo-I/DNA covalent complexes induced by combination treatment and suggest a potentiation by vorinostat of topotecan-induced DNA damage. Finally, oxidative injury played a significant functional role in the observed enhanced lethality because coadministration of the antioxidant N-acetyl-L-cysteine blocked reactive oxygen species generation, apoptosis, and mitochondria destruction induced by the vorinostat/topotecan combination. To our knowledge, this is the first demonstration of a synergistic antitumor effect between topotecan and vorinostat in SCLC. Because no well-established treatment is available for recurrent SCLC patients, our results indicate that this drug combination should be explored clinically. [Mol Cancer Ther 2009;8(11):3075–87]

Introduction
Small cell lung cancer (SCLC) is an aggressive form of cancer, which accounts for ~12% to 16% of lung neoplasms (1). Although good response rates have been achieved using combination chemotherapy with cisplatin and etoposide as the most widely used regimen, only a small proportion of patients survive after 5 years (2). Most patients relapse within 1 year of starting first-line treatment due mainly to intrinsic or acquired resistance to chemotherapy. Recently, several new chemotherapeutics as well as targeted agents have been tested in SCLC; however, only minor positive results were obtained (2). Thus, alternative agents and new integrated therapeutics are needed.

Histone deacetylase inhibitors (HDACI) represent a new class of anticancer agents that, enhancing the acetylation of core histones, weaken histone-DNA interactions and consequently increase DNA accessibility and activate transcription (3, 4). These compounds act very selectively to alter the transcription of <2% to 5% of expressed genes (3, 5) and induce tumor cell growth arrest, differentiation, and apoptosis by mechanisms that cannot be solely attributed to the level of histone acetylation (6). In addition to histones, reversible acetylation of other proteins such as p53, α-tubulin, and heat shock protein 90, has also been described (7). Several HDACIs exhibit antitumor effects in preclinical animal models at amounts that have little or no toxicity (3), and some are in advanced clinical studies either as single agents or in combination with conventional chemotherapy or biologicals (3, 4, 8–10). One of the most promising HDACIs, vorinostat (suberoylanilide hydroxamic acid), has shown significant preclinical and clinical antitumor activities in both hematologic malignancies and solid tumors.
including SCLC and represents the first HDACI to be recently approved by the Food and Drug Administration for the treatment of cutaneous T-cell lymphoma with progressive, persistent, or recurrent disease during or following two systemic therapies (3, 9–11).

The nuclear topoisomerase-I (topo-I) and topo-II enzymes are responsible for regulating and maintaining DNA topology. In particular, by breaking and religating DNA strands, they relax DNA supercoiling generated by transcription, replication, and chromatin remodeling (12). Topoisomerase enzymes are targeted by several inhibitors currently used as antitumor agents. Several classes of these agents including topo-I inhibitors interfere with the reversible topo-DNA interaction by covalently trapping the enzyme on the DNA and inducing irreversible DNA damage during the replication process (13). It has been suggested that increased accessibility of the topoisomerase enzymes to chromatin DNA could result in increased sensitivity to the inhibitors (14, 15). In this regard, relaxation of chromatin structure induced by HDACIs may facilitate accessibility of the topoisomerase enzymes to DNA and increase the cytotoxicity of topoisomerase inhibitors.

In this study, we analyzed the antitumor effect of the HDACI vorinostat in combination with the topo-I inhibitors camptothecin and its water-soluble semisynthetic derivative topotecan, the only registered drug for second-line therapy in relapsed SCLC (16). Topotecan has shown antitumor activity in both chemosensitive and chemoresistant SCLC and has a nonoverlapping toxicity profile with other agents used in the treatment of SCLC (16).

We showed that vorinostat in combination with either camptothecin or topotecan exerted a strong synergistic cytotoxic effect in SCLC cells, including those resistant to topo-I inhibitors. This effect was not dependent on the schedule of treatment and was paralleled by a late S-phase cell cycle arrest and pronounced apoptosis triggered by a caspase-dependent mechanism. The induction of phosphorylated H2AX (γH2AX), which is a marker of DNA damage, the generation of reactive oxygen species (ROS), and the destruction of mitochondrial membrane were observed only in cells treated by the vorinostat/topotecan combination and were paralleled by an increase of topo-I/DNA covalent complexes, suggesting a potential mechanism for the observed synergistic antitumor interaction.

Materials and Methods

Materials
Clinical-grade vorinostat was provided by Merck & Co. Stock solutions were prepared in DMSO and diluted to appropriate concentrations in culture medium before addition to the cells. Topotecan (Hycammin) was supplied by GlaxoSmithKline and camptothecin was obtained from Alexis Biochemicals. Antibodies were purchased as follows: acetylated histone H3 and γH2AX from Upstate, poly(ADP-ribose) polymerase (PARP) from BD Biosciences, caspase-3 from Cell Signaling Technology, and topo-I, γ-tubulin, and p53 from Santa Cruz Biotechnology. MTT was from ICN Biomedicals. Enhanced chemiluminescence immunodetection reagents were from GE Healthcare. All media, serum, antibiotics, and glutamine were from Cambrex Bioscience. MitoTracker Red CMTMRos was by Invitrogen (Life Technologies). N-acetyl-l-cysteine (t-NAC) and hydroethidine were from Sigma-Aldrich and Annexin V-FITC was from Becton Dickinson.

Cell Culture and Cell Viability Assay
H126 mutant p53 and H209 wild-type p53 cell lines were obtained from the American Type Culture Collections and grown in suspension in RPMI supplemented with 10% heat-inactivated fetal bovine serum, penicillin (50 units/mL), streptomycin (500 μg/mL), and 4 mM/L glutamine in a humidified atmosphere of 95% air and 5% CO2 at 37°C. Cell viability was measured after treatment (72 or 96 h) in 96-well plates by a MTT assay as described previously (17).

Drug Combination Studies
Drug combination studies were based on concentration-effect curves generated as plot of the fraction of unaffected (surviving) cells versus drug concentration after 72 or 96 h (in the case of sequential experiments) of treatment. To explore the relative contribution of each agent to the synergism, serial dilutions of doses of the two agents (vorinostat/topotecan or camptothecin) in combination were tested with different molar ratios: equi-active doses (50:50 cytotoxic ratio) of the two agents (IC50), higher relative doses of topo-I inhibitors or camptothecin (25:75 cytotoxic ratio; IC25 of vorinostat/IC75 of topo-I or camptothecin), and higher relative doses of vorinostat (IC75 of vorinostat/IC25 of topotecan or camptothecin).

For sequential combinations, serial dilutions of equi-active doses were tested in three different sequences of treatment: simultaneous (both drugs were given together 24 h after seeding and cell growth assessment was done after 96 h) or when each one was applied 24 h before the other (vorinostat or topotecan or camptothecin was added 24 h after seeding followed by the indicated combined drug after 24 h, and cell growth assessment was done after 72 h), so that cells were exposed for 96 h to the first applied drug and for 72 h to the following combined agent.

Synergy for these combination effects was determined by the combination index (CI) calculated by the Chou-Talalay equation (Eq. 18) as described elsewhere (19) and by using the software CalcuSyn (Biosoft). Assuming 0.8 as the cutoff, CI < 0.8, CI = 0.8–1, and CI > 1 indicate synergistic, additive, and antagonistic effects, respectively. The dose reduction index (DRI) represents the measure of how much the dose of each drug in a synergistic combination may be reduced at a given effect level compared with the dose of each drug alone. The linear correlation coefficient (r) of the median-effect plot was always between 0.90 and 0.99, indicating good conformity of the data (18).

Analysis of Cell Cycle Kinetics
Analysis of cell cycle kinetics was done at the indicated times as described previously (20). Briefly, cells were harvested, fixed in 70% ethanol, and stored at -20°C until analysis. After nuclear DNA staining with propidium iodide, flow cytometry was done in duplicate by a FACSScan flow

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cytometer (Becton Dickinson). For each sample, 20,000 events were stored and cell cycle analysis was done by the ModFit LT software (Verity Software House). FL2 area versus FL2 width gating was done to exclude doubles from the G2-M region. The percentage of apoptotic cells was calculated in the subdiploid region of the DNA content, registered as FL2 signals in linear scale. To avoid cell debris contamination due to necrotic cell death, cells were selected by side scatter versus DNA signals (FL2) gating.

**Soft-Agar Clonogenic Assay**

The soft-agar clonogenic assay was done as reported previously (21). In brief, cells were plated at 2 \( \times 10^3 \) /mL in an upper layer of 0.3% bacto-agar in RPMI (1:1) and 20% fetal bovine serum over an underlayer of 0.8% agar in RPMI in 24-well plates, and after 3 h, the cells were treated with vorinostat at doses of 1/5 and 1/10 of the in vitro IC50 and/or topotecan at doses of 1/6 and 1/12 of the in vitro IC50. Cultures were maintained for 3 weeks in a dry atmosphere of 95% air and 5% CO2 at 37°C. Cellular colonies were then stained with 150 μL nitroblue tetrazolium and colonies >50 μm were scored with an Artek 880 colony counter (Artek). The data were analyzed by Image-Pro Plus software.

**Protein Extraction and Western Blotting**

Cells grown and treated as indicated were collected, lysed, and separated on SDS-PAGE, and proteins were transferred to nitrocellulose paper, immunoblotted with specific antibodies, and probed with the appropriate horse-radish peroxidase-linked IgG as described elsewhere (19). Immunoreactive bands were detected by enhanced chemiluminescence. For the immunodetection of topo-I, nuclear extracts were prepared as described previously (22).

**Apoptosis Assays**

Annexin V binding was identified by flow cytometry using Annexin V-FITC staining following the manufacturer’s instructions (Becton Dickinson). Internucleosomal DNA fragmentation (ladder) was measured after extraction of low molecular weight DNA, PAGE, and silver staining as described previously (23). Briefly, 1 \( \times 10^7 \) cells were resuspended in Tris-EDTA and lysed in 20% SDS and DNA was precipitated in 2 mol/L NaCl. The high molecular weight fraction was sedimented by high-speed centrifugation and fragmented DNA was extracted from the aqueous phase with phenol and chloroform and then precipitated with ethanol. After resuspension in loading buffer, DNA was electrophoresed using 10% polyacrylamide gel and visualized by silver staining.

**Measurement of ROS Generation**

ROS generation was analyzed by flow cytometry using the ROS-sensitive dye hydroethidine as described previously (24). Briefly, the cells, untreated or treated as indicated, were incubated with 20 ng/mL hydroethidine for 1 h at 37°C in the dark and then trypsinized from the culture dishes and resuspended in PBS. ROS generation was measured as significant increases in fluorescence intensity within 45 min by CellQuest data analysis software.

**Mitochondrial Potential Analysis**

Mitochondrial functions such as mitochondrial membrane potential can be studied in live cells using vital fluorescent lipophilic cationic dyes. We took advantage of the MitoTracker Red CMXROS fluorescent dye (Invitrogen), a specific mitochondria dye derived from X-rosamine that exhibits good photostability and is usually applied to analyze changes in mitochondrial membrane potential in fixed cells by confocal microscopy. We evaluated CMXROS staining of live cells by flow cytometry, which has been reported to be a more quantitative method to measure mitochondrial membrane potential (25). Cells were treated with the drugs at indicated concentrations and collected after 24, 48, and 72 h and then harvested, centrifuged, and washed with PBS. Cell suspensions were incubated in 1 mL PBS with 500 nmol/L CMXROS dye for 15 to 45 min in the dark at 37°C. After incubation, the cells were immediately analyzed. The transition of mitochondria from a polarized to depolarized status results in leakage of dye and a consequent decrease of red fluorescence as measured by CellQuest data analysis software as described previously (26, 27).

**DNA Relaxation Assays**

Nuclear extracts, prepared as described before (22), were incubated in 20 μL reaction volume containing 0.5 μg negatively supercoiled pBlueScript KSII (+) DNA and reaction buffer as described previously (28); to assess the effects of vorinostat on enzyme activity, increasing concentrations of nuclear proteins were added to the mixture. Reactions were stopped with a final concentration of 0.5% SDS after 1 h at 37°C. The samples were electrophoresed in a horizontal 1% agarose gel in 50 mmol/L Tris, 45 mmol/L boric acid, and 1 mmol/L EDTA. The gel was stained with ethidium bromide (5 μg/mL) and destained with water and the image was acquired under UV illumination by ChemiDoc XRS (Bio-Rad). Where indicated, purified topo-IB (28) and vorinostat on enzyme activity, increasing concentrations of nuclear proteins were added to the mixture. Reactions were stopped with a final concentration of 0.5% SDS after 1 h at 37°C. The samples were electrophoresed in a horizontal 1% agarose gel in 50 mmol/L Tris, 45 mmol/L boric acid, and 1 mmol/L EDTA. The gel was stained with ethidium bromide (5 μg/mL) and destained with water and the image was acquired under UV illumination by ChemiDoc XRS (Bio-Rad). Where indicated, purified topo-IB (28) and vorinostat, or camptothecin, where incubated at 37°C in the

**Detection of Covalent Topo-I/DNA Complexes**

Topo-I/DNA adducts were isolated using an Immuno Complex Enzyme bioassay as described previously (29). Briefly, 1 \( \times 10^7 \) treated and untreated cells were pelleted and immediately lysed with 1 mL of 1% sarkosyl. After syringe-drawing, cell lysates were gently layered on step gradients containing four different cesium chloride solutions (2 mL of each) of following densities: 1.82, 1.72, 1.50, and 1.45 (30). Tubes were centrifuged at 31,000 rpm for 21 h at 20°C. Half-milliliter fractions were collected from the bottom of the tubes and the DNA peak was localized by determining the absorbance at 260 nm. In all, 5 μg DNA was mixed with an equal volume of 25 mmol/L sodium phosphate (pH 6.5) and applied to nitrocellulose paper by using a slot-blot vacuum manifold (Bio-Rad). The membranes were equilibrated in TBST [10 mmol/L Tris-HCl (pH 7.5), 150 mmol/L NaCl, and 0.05% Tween 20] and then incubated overnight at 4°C with topo-I antibody diluted in TBST containing 5% nonfat dried milk.

**Statistical Analysis**

All determinations were done in quadruplicates or triplicates as indicated and each experiment was repeated at least three times. Representative results from flow cytometry analysis, Western blotting, DNA ladder analysis, DNA

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relaxation assays, and detection of covalent topo-I/DNA complexes from a single experiment are presented; additional experiments yielded similar results. Statistical analyses were done by paired \( t \) tests using Sigma Stat software (Systat Software). Differences were considered to be statistically significant at \( P < 0.05 \).

**Results**

**Vorinostat and Topo-I Inhibitors Interact Synergistically to Induce a Cytotoxic Effect in SCLC Cells**

In several *in vitro* cancer models, HDACIs in combination with drugs targeting DNA induce a synergistic inhibition of cell viability as well as apoptosis (4, 31). To evaluate the potential synergistic effect of vorinostat in combination with the topo-I inhibitors topotecan or camptothecin in SCLC cell lines, H209 (wild-type p53) and H526 (mutant p53) cells were exposed for 72 h to increasing concentrations of the drugs, alone or in combination (vorinostat/topotecan or vorinostat/camptothecin), and cell viability was evaluated by the MTT assay. Vorinostat reduced the viability of both H209 and H526 cells at micromolar concentrations (Fig. 1A) and induced a time-dependent histone H3 acetylation in both cell lines (Fig. 1A, inset). In contrast, the mutant p53 H526 cell line appeared strongly resistant to both topotecan and camptothecin, with respective IC\(_{50}\) values of \( \sim 1,000\)- and 160-fold higher than those observed in wild-type p53 H209 cells (Fig. 1B and C).

The interactions between vorinostat and either topotecan or camptothecin appeared to be strongly synergistic as shown by a median drug effect analysis calculating CIs in both cell lines (Table 1). In details, to explore the relative contribution of each agent to the synergism, different vorinostat/topotecan or vorinostat/camptothecin combinations were compared by testing equipotent doses of the two agents (50:50 cytotoxic ratio), higher relative doses of vorinostat (75:25 cytotoxic ratio), or higher relative doses of topotecan or camptothecin (25:75 cytotoxic ratio). A strong synergistic effect with low CIs (CIs < 0.8) was shown for both vorinostat/topotecan and vorinostat/camptothecin combinations when equipotent doses of the two drugs were used (50:50 cytotoxic ratio) in H209 cells. The same results were observed in the H526 cell line resistant to both topotecan and camptothecin (see Fig. 1B and C). Similar or even stronger effects were observed in both cell lines when lower doses of vorinostat were used (25:75 cytotoxic ratio). In contrast, when higher doses of vorinostat were combined with lower doses of either camptothecin or topotecan (75:25 cytotoxic ratio), a synergistic effect was observed only for the vorinostat/topotecan combination in H209 cells, whereas an additive or antagonistic effect was observed for the other combinations.

Furthermore, we explored different schedules of treatment combining equitoxic concentrations of vorinostat...
and topotecan in both H526 and H209 cell lines. As shown in Table 1, a sequential exposure with 24 h delay to the two agents in combination produced a strong synergistic cytotoxic effect similar to that observed for simultaneous treatment, indicating that the synergistic interaction between vorinostat and topotecan was not schedule-dependent. Interestingly, the evaluation of DRI values, which represent the order of magnitude (fold) of dose reduction obtained for IC₅₀ (DRI₅₀) in combination setting compared with each drug alone.

### Table 1. CI and DRI values according to the different cytotoxic ratios of vorinostat/topotecan and vorinostat/camptothecin or to the different schedules of exposure to vorinostat/topotecan combinations

<table>
<thead>
<tr>
<th>Cell lines</th>
<th>Vorinostat + topotecan</th>
<th>Vorinostat → topotecan</th>
<th>Topotecan → vorinostat</th>
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<tbody>
<tr>
<td>H209</td>
<td>CI₅₀ ± SD*</td>
<td>DRI₅₀ ± SD†</td>
<td>CI₅₀ ± SD*</td>
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<tr>
<td></td>
<td>vorinostat</td>
<td>topotecan</td>
<td>vorinostat</td>
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<td></td>
<td></td>
<td></td>
<td>topotecan</td>
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<tr>
<td>H209</td>
<td>0.38 (0.04)</td>
<td>4.5 (1.5)</td>
<td>6.94 (0.28)</td>
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<td></td>
<td>(0.15)</td>
<td>0.28 (0.015)</td>
<td>7.44 (0.9)</td>
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<td></td>
<td></td>
<td>0.41 (0.064)</td>
<td>9.4 (0.8)</td>
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<td>H209</td>
<td></td>
<td>0.39 (0.05)</td>
<td>17.19 (1)</td>
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<td></td>
<td></td>
<td></td>
<td>0.39 (0.056)</td>
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<td>H209</td>
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<td>3.6 (1.5)</td>
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<td>H209</td>
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<td></td>
<td>15.88 (1)</td>
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**NOTE:** 50:50, serial dilutions from vorinostat IC₀₅₀ [4 μmol/L (H209) and 3 μmol/L (H526)]/topotecan IC₀₅₀ [0.007 μmol/L (H209) and 4 μmol/L (H526)] or camptothecin IC₀₅₀ [0.008 μmol/L (H209) and 1.5 μmol/L (H526)]; 75:25, vorinostat IC₀₅₀ [10 μmol/L (H209) and 20 μmol/L (H526)]/topotecan IC₀₂₅ [0.001 μmol/L (H209) and 0.1 μmol/L (H526)]; and 25:75, vorinostat IC₀₂₅ [1.15 μmol/L (H209) and 0.8 μmol/L (H526)]/topotecan IC₀₇₅ [0.6 μmol/L (H209) and 64 μmol/L (H526)] or camptothecin IC₀₇₅ [10 μmol/L (H209) and 20 μmol/L (H526)]. Vorinostat + topotecan, simultaneous treatment; vorinostat → topotecan or topotecan → vorinostat, sequential treatment with 24 h delay to either drug.

**CI₅₀ (DRI₅₀)** values (mean ± SD from at least three separate experiments done in quadruplicates) were computed at 50% of cell kill (CI₅₀) by CalcuSyn software; CIs < 0.8 indicated synergism as described in Materials and Methods.

**DRI values (mean ± SD from at least three separate experiments done in quadruplicates)** represent the order of magnitude (fold) of dose reduction obtained for IC₅₀ (DRI₅₀) in combination setting compared with each drug alone.

**and topotecan in both H526 and H209 cell lines. As shown in Table 1, a sequential exposure with 24 h delay to the two agents in combination produced a strong synergistic cytotoxic effect similar to that observed for simultaneous treatment.**

**Interestingly, the evaluation of DRI values, which represent the order of magnitude (fold) of dose reduction obtained for IC₅₀ (DRI₅₀) in combination treatment compared with single-drug treatment, showed fold reductions of 4 to 9.4 for vorinostat and 3.6 to 0.8 for topotecan in all three sequences examined (Table 1).**

**Cell Cycle Effects Induced by Vorinostat in Combination with Topo-I Inhibitors**

To determine whether the mechanism underlying the observed synergistic interaction might involve cell cycle perturbations, we evaluated the effects of vorinostat and topotecan or camptothecin, either alone or in combination at IC₅₀ concentrations, on the cell cycle kinetics of both H209 and H526 cells (Fig. 2). Vorinostat alone after 48 h induced a slight increase in G₁ phase of the cell cycle in both cell lines in comparison with the controls. On the other hand, topotecan and camptothecin induced a late-middle S-phase cell cycle arrest in both cell lines, whereas an apoptotic effect, measured by the appearance of a hypodiploid population (sub-G₁/G₀ indicated by arrows), was observed only in H209 cells (Fig. 2A). Concomitant treatment with vorinostat did not change the cell cycle perturbations or the apoptotic effect induced by topotecan or camptothecin in H209 cells (Fig. 2A); however, a slight increase in the percentage of cells arrested in S phase and a clear apoptotic effect was observed in the H526 cells treated with a combination of vorinostat and topotecan (Fig. 2B). Similar results were obtained in H526 cells treated with the different sequential combination schedules described earlier (data not shown).

**Vorinostat in Combination with Topotecan Inhibits Soft-Agar Colony Formation and Induces Apoptosis and DNA Damage**

To confirm by a different method the synergistic effect observed thus far, a soft-agar clonogenic assay was used to determine the effect of vorinostat and topotecan, alone or in combination, in H526 cells. Vorinostat, when added alone at doses corresponding to 1/10 (250 nmol/L) or 1/5 (500 nmol/L) of the 72 h IC₅₀ values (IC₅₀ for the induction of apoptosis by the vorinostat/topotecan combination, to induce apoptosis in both H526 and H209 cells. Western blot analysis showed increased PARP and caspase-3 cleavage in H526 cells treated simultaneously for 48 h with vorinostat and topotecan at IC₅₀ doses compared with single-drug treatments (Fig. 3B). Notably, the induction of apoptosis by the vorinostat/topotecan combination in H526 cells was also confirmed by the following assays: Annexin V staining, which showed a 2.5-fold increase over the control compared with a slight increase.
induced by single drugs (Fig. 3C), DNA fragmentation as seen by ladder formation (Fig. 4B), and destruction of mitochondria (Fig. 4C). In the wild-type p53 H209 cell line, the increased cleavage of PARP was paralleled by an increased expression of p53 induced by combination of vorinostat and topotecan (Fig. 3B).

H2AX is phosphorylated on Ser\(^{139}\) by ATM kinase in response to DNA damage-induced DNA double-strand breaks and is
one of the earliest events in the DNA damage signaling and repair (32). An increase of γH2AX in S phase has been shown previously after treatment of cancer cells with topotecan or camptothecin (33). As shown in Fig. 3B, the induction of γH2AX in both H526 and H209 cell lines was observed, only in the combination setting after 48 h, indicating an evident potentiation by vorinostat of topo-I inhibitor-induced DNA damage.

These findings indicate that treatment of SCLC cells with vorinostat/topotecan combination potently induces DNA damage associated with caspase-dependent activation of the apoptotic program.

**Induction of Apoptosis Mediated by the Vorinostat/Topotecan Combination Is Correlated to Enhanced ROS Generation**

Previous studies have suggested that the lethal effect of both HDACI and topo-I inhibitors could be related to oxidative stress as measured by the increase of ROS levels (34–38). Studies were therefore performed to determine whether a similar mechanism might underlie the synergistic cytotoxic and apoptotic effects mediated by vorinostat/topotecan. As shown in Fig. 4A, vorinostat and topotecan (or camptothecin) combination induced a significant increase in ROS levels, compared with single-agent treatments, relative to control untreated H526 and H209 cells. This effect was evident with a time point peak statistically significant of 48 h in H526 cells and of 24 h in H209 cells, probably reflecting different sensitivity toward topo-I inhibitors between the two cell lines. H526 cells treated for 10 h with H2O2 were assumed as positive control (Fig. 4A).

Furthermore, coadministration of the antioxidant l-NAC substantially blocked H2O2-mediated as well as the vorinostat/topotecan (or camptothecin)–mediated increase of ROS levels.
**In vitro Synergism of Vorinostat plus Topotecan**

**A**

H526

<table>
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<tr>
<th>Time</th>
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<th>Topotecan</th>
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<tr>
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<td>+</td>
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H209

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<th>Topotecan</th>
<th>NAC</th>
<th>Vorinostat + Topotecan + NAC</th>
</tr>
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<tbody>
<tr>
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<tr>
<td>72 h</td>
<td>+</td>
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</table>

**B**

- Vorinostat
- Topotecan
- NAC

**C**

- H526 CTR 72h
- H526 NAC 72h
- H526 vor 72h
- H526 topo 72h
- H526 vor + topo 72h
- H526 vor + topo + NAC 72h

Flowerl.png
levels in both cell lines. Consistent with these findings, L-NAC also blocked the vorinostat/topotecan-mediated apoptosis in H526 cells as reflected by a reduction of DNA fragmentation shown after 48 h (Fig. 4C) as well as the destruction of mitochondria after 72 h (evaluated as a loss of membrane potential by MitoTracker Red staining and flow cytometric analysis; Fig. 4D), both observed only in combination-treated cells. The latter effect was not observed at early time points (data not shown), indicating that the loss of mitochondrial integrity is a consequence of the apoptotic pathway induced by the vorinostat/topotecan combination.

Collectively, these findings indicate that oxidative injury plays a significant functional role in the enhanced lethality induced by the vorinostat/topotecan combination in SCLC cancer cells.

**Vorinostat Enhanced Topo-I/DNA Cleavable Complex Formation Induced by Camptothecin but Do Not Affect Expression or Catalytic Activity of Topo-I**

To evaluate if vorinostat may affect either expression or activity of topo-I, we first performed Western blotting analysis, showing that treatment with vorinostat (48 h, IC$_{50}$ 2 h) of both H526 and H209 cells does not determine any significant change in topo-I protein level in comparison with control cells (Fig. 5A), in line with what was reported in other cell models (14). We next compared the catalytic activity and the camptothecin sensitivity of topo-I from nuclear extracts of untreated or vorinostat-treated H526 and H209 cells following the electrophoretic mobility of the DNA supercoiled plasmid topo-I substrate. As shown in Fig. 5B, control untreated or 24 h vorinostat-treated cells (IC$_{50}$ 22 h) showed comparable DNA relaxation activities. Similar results were obtained after 48 h of treatment (data not shown). Likewise, increasing concentrations and time titration of vorinostat up to 2 mmol/L do not affect the DNA relaxation monitored adding a purified topo-I protein (Supplementary Fig. S1). Moreover, the catalytic activities of both untreated and vorinostat-treated cells were similarly inhibited by the addition of camptothecin in the reaction mixture (Supplementary Fig. S2). These results indicate that vorinostat-treated cells have the same properties as control untreated cells, with respect to the topo-I catalytic activity and sensitivity to topo-I inhibitors.

To determine whether vorinostat may modulate the amount of topo-I/DNA cleavable complexes induced by topo-I inhibitors, such as camptothecin, we identified topo-I bound to cellular DNA by immunoblot after separating, by cesium chloride gradient centrifugation, cellular extracts of H526 and H209 cells incubated in different conditions (Fig. 5D). The amount of topo-I coincident with the DNA peak, as represented from control untreated H526 and H209 cells in Fig. 5C, is a measure of covalent topo-I/DNA complexes formed. The presence of topo-I/DNA cleavable complexes is detectable also in the control cells in the absence of camptothecin as shown in Fig. 5D (fractions 6-8, control lanes) in line with what was reported in literature (39). Vorinostat alone was unable to modify the amount of cleavable complexes compared with controls. In camptothecin-treated cells, a clear increase in the intensity of the spots corresponding to cleavable topo-I/DNA complexes is observed in H209 cells but not in H526 cells, confirming that they are camptothecin-sensitive and camptothecin-insensitive, respectively. Notably, an increase in topo-I/DNA covalent complexes is observed H526 cells, and slightly in H209 cells, when concomitantly treated with vorinostat and camptothecin relative to cells treated with camptothecin alone (Fig. 5D, fractions 6-8, two last lanes). Free topo-I (unbound) was found to be present in the low molecular weight fraction after cesium chloride centrifugation (Fig. 5D, fractions 15-17).

Collectively, these findings indicate that vorinostat does not affect directly topo-I activity nor changes the content of topo-I but rather potentiate the effect of topo-I inhibitors by increasing, at least in resistant H526 cells, the amount of topo-I/DNA cleavable complexes.

**Discussion**

In the present study, we have shown a synergistic cytotoxicity in SCLC cell lines, regardless of their p53 status, when the HDAC inhibitor vorinostat is combined with the topo-I inhibitor camptothecin or its derivative topotecan. Simultaneous exposures of SCLC cells to equitoxic doses of vorinostat and camptothecin or topotecan as well as low doses of vorinostat and higher doses of any topo-I inhibitors resulted in strong synergistic cytotoxicity. Moreover, the effect of combining vorinostat/topo-I inhibitors is not sequence-dependent, because an evident synergistic effect was achieved when both agents were applied simultaneously or when each was applied 24 h before the other. Notably, we have also shown a strong synergistic cytotoxicity in the H526 cell line that has a marked intrinsic resistance to topo-I inhibitors. The observation that vorinostat might enhance
topo-I inhibitors–induced cytotoxicity even at doses below the IC50 values suggests that the mechanism of the synergism might depend, at least in part, on the ability of vorinostat to modulate the sensitivity of the cells toward topo-I inhibitors rather than on the concurrent cell killing induced by both agents. Moreover, as shown by previous pharmacokinetic studies (10), the effective doses of vorinostat tested in our in vitro study can be easily reached in plasma concentrations.

Figure 5. Effects of vorinostat alone or in combination with camptothecin on topo-I expression, topo-I activity, and topo-I cleavage complexes in H526 and H209 cells. 

A, Western blot analysis of topo-I was done on H526 and H209 cells untreated or treated for 48 h with IC50 doses of vorinostat. Nuclear extracts (40 μg) were resolved by SDS-PAGE and immunoblotted. Ponceau staining (data not shown) and γ-tubulin immunoblotting ensured the equal loading of samples in each lane. B, DNA relaxation activity of topo-I in H209 and H526 cells untreated or treated for 24 h with IC50 doses of vorinostat. Substrate-negative supercoiled plasmid DNA, incubated with various amounts of nuclear extracts isolated from cells and resolved in a 1% agarose gel, was visualized with ethidium bromide. C, absorbance at 260 nm of the aliquots obtained in a cesium chloride gradient from untreated H526 and H209 cells. From the bottom of the gradient, the absorbance grows in two peaks; the first correspond to the purified DNA and the second one correspond to the free proteins. D, chromatin-associated topo-I cleavage complexes evaluated in H526 and H209 cells, untreated or treated, for 24 h with IC50 doses of vorinostat and/or camptothecin using Immuno Complex Enzyme bioassay as described in Materials and Methods. Fractions collected from cesium chloride gradients are indicated above the immunoblots.
in patients and with negligible toxicity. In addition, the lack of a schedule dependency in the synergistic interaction between vorinostat and topo-I inhibitors is clinically relevant because a less stringent condition of drug administration would make this combination easily adaptable for clinical applications.

Preclinical studies in multiple cancer cell lines have shown the potentiation by HDACIs of the effects of a wide range of DNA-damaging agents, including topoisomerase inhibitors (31). In vitro and in vivo preclinical synergistic effects were observed when HDACIs, including vorinostat, were combined with the topo-II inhibitors etoposide or epirubicin (14, 40, 41). A completed phase I clinical study of the HDACIs valproic acid and epirubicin has provided evidence for the safety of this combination (42). Scheduling appeared crucial in these studies and synergy was dependent on the sequence of drug administration (40). Our results are confirmed by the synergism observed in a panel of ovarian cancer cells or pancreatic cancer cells treated simultaneously with a HDACI (even in low doses; refs. 15, 43) and the topo-I inhibitors topotecan or CPT11 (irinotecan hydrochloride hydrate), respectively. On the other hand, in contrast to our data, Kim et al. reported in one glioblastoma cell line that the HDACI trichostatin A given before or simultaneously did not potentiate camptothecin-induced cell killing (41). Furthermore, Bevins and Zimmer showed that HDACIs, added before or simultaneously as camptothecin, limited the number of cells affected by camptothecin, whereas cells treated with a HDACI 24 h after camptothecin were efficiently killed with potentiation of apoptosis (44). The authors suggested that HDACI-induced apoptosis was most effective when HDACI was added to cells already arrested in the G2-M phase of the cell cycle by camptothecin. However, it is worth of mention that several reports clearly show that prolonged treatment of cancer cells with lower, therapeutically achievable camptothecin concentrations causes S-phase arrest, whereas only a brief exposure to high camptothecin concentrations produces a subsequent G2 arrest similar to that shown by Bevins and Zimmer (45–47). In agreement with these data, we showed a late-middle S-phase cell cycle arrest induced by topotecan or camptothecin in both H209 and H526 cells. Notably, this effect was either not changed by concomitant treatment with vorinostat, or a slight increase of the S-phase cell cycle arrest was observed in resistant H526 cells. Moreover, an apoptotic sub-G1 hypodiploid peak formation was observed in resistant H526 cells treated with a combination of vorinostat/topotecan or vorinostat/camptothecin compared with single drugs. Furthermore, we should point out that the majority of the studies mentioned above did not investigate increasing doses and different cytotoxic ratios of the two agents. Neither did those studies evaluate synergism by the median effect analysis calculating CIs, which is universally accepted as the most reliable and statistically robust approach to investigate the interaction between two or more drugs in a preclinical in vitro drug screening test (18). In addition, we showed synergistic interaction between vorinostat and either camptothecin or topotecan by several approaches, including agar colony forma-

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In summary, it is well known that acetylation of core nucleosomal histones on HDAC treatment induces a more relaxed winding of DNA around the histone core, increasing DNA accessibility to cellular enzymes that promote transcription, replication, and repair but eventually also increasing accessibility to DNA-damaging chemotherapeutics such as topo-I inhibitors (11, 31). It has been described that vorinostat as well as other HDACis are able to abrogate the DNA repair process and to inhibit survival mechanisms, thus increasing the cytotoxic effects of DNA-damaging agents and resulting in cell death (11). Together with these potential mechanisms, several lines of evidence indicate that enhanced oxidative damage plays an important functional role in the synergistic interactions observed between vorinostat and topo-I inhibitors in SCLC cells.

In conclusion, to our knowledge, this study is the first to show in a SCLC cell model, and with a novel potential mechanism, a synergistic cytotoxic effect between a HDACi such as vorinostat and a topo-I inhibitor such as topotecan, the latter being the only registered drug for relapsed SCLC. Our results also indicate that the vorinostat/topotecan combination is a promising chemotherapeutic strategy for killing SCLC cells regardless of whether the cells are sensitive or resistant to topo-I inhibitors. Based on our results and the fact that effective treatment options for SCLC patients remain inadequate, a clinical trial of vorinostat and topotecan combination, with a simultaneous schedule of administration, has been started in SCLC.

Disclosure of Potential Conflicts of Interest

The authors received a research grant from Merck and Co.

References


Synergistic antitumor effect between vorinostat and topotecan in small cell lung cancer cells is mediated by generation of reactive oxygen species and DNA damage-induced apoptosis

Francesca Bruzzone, Monia Rocco, Silvia Castelli, et al.