Exisulind-induced Apoptosis in a Non-Small Cell Lung Cancer Orthotopic Lung Tumor Model Augments Docetaxel Treatment and Contributes to Increased Survival

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Abstract
We reported previously a significant increase in survival of nude rats harboring orthotopic A549 human non-small cell lung cancer tumors after treatment with a combination of exisulind (Sulindac Sulfone) and docetaxel (D. C. Chan, Clin. Cancer Res., 8: 904–912, 2002). The purpose of the current study was to determine the biochemical mechanisms responsible for the increased survival by an analysis of the effects of both drugs on A549 orthotopic lung tumors and A549 cells in culture. Orthotopic A549 rat lung tissue sections from drug-treated rats and A549 cell culture responses to exisulind and docetaxel were compared using multiple apoptosis and proliferation analyses [i.e., terminal deoxynucleotidyl transferase-mediated nick end labeling, active caspase 3, the caspase cleavage products cytokeratin 18 and p85 poly(ADP-ribose) polymerase, and Ki-67]. Immunohistochemistry was used to determine cyclic GMP (cGMP) phosphodiesterase (PDE) expression in cultured A549 cells. The cGMP PDE composition of cultured A549 cells was resolved by DEAE-Trisacryl M chromatography and the cGMP PDEs, and these results are consistent with a cGMP-regulated apoptosis pathway.

Introduction
Exisulind (Sulindac Sulfone, Aptosyn) is an antineoplastic drug being developed for the treatment of premalignant and malignant cancer. It is the lead drug of a family of compounds termed selective apoptotic antineoplastic drugs because they induce apoptosis in neoplastic but not normal cells. Exisulind has been shown to selectively induce apoptosis in cell lines derived from many cancers including colon, bladder, prostate, and breast (1–4). The drug also inhibits tumor growth in rodent models of mammary, bladder, prostate, and colon carcinogenesis (1, 3, 5, 6). In colon cancer cells exisulind induces apoptosis by a mechanism that involves the inhibition of cGMP2 phosphodiesterases, including PDE 2 and 5, activation of PKG and phosphorylation of selective substrates (7). Many cancers, including colon, pancreatic, lung, and bladder, have high levels of cGMP PDE compared with normal tissue (3, 8–10). Exisulind treatment of colon tumor cells in culture results in a sustained increase in intracellular cGMP levels that results in the activation of PKG (11, 12). Activated PKG directly phosphorylates mitogen-activated protein kinase kinase kinase 1, which leads to the activation of Jun kinase 1 and the initiation of apoptosis (13, 14). In addition, PKG phosphorylates β-catenin causing a reduction in its accumulation and anti-apoptotic effects (15).

Lung cancer is currently the leading cause of cancer death in the United States (16). Even with recent advances in treatment, response and remission rates remain unacceptably low. Whereas docetaxel treatment of NSCLC produces a modest improvement in survival (17–20), there remains a need for novel agents to improve the treatment and survival of NSCLC patients. A previous in vitro study reported that the combination of exisulind and paclitaxel produced a synergistic inhibition of growth in a variety of lung cancer cell lines (21). We reported recently the results of a study investigating the efficacy of exisulind and docetaxel treatment, alone or in combination, in a non-small cell lung cancer mouse model (22).

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2 The abbreviations used are: cGMP, cyclic G; PDE, phosphodiesterase; PKG, protein kinase G; NSCLC, non-small cell lung cancer; PARP, poly(ADP-ribose) polymerase; IHC, immunohistochemistry; TUNEL, terminal deoxynucleotidyl transferase-mediated nick end labeling; GST, glutathione-S-transferase; cAMP, cyclic AMP.
combination, in nude rats harboring orthotopic human A549 NSCLC tumors (22). We found that rats treated with the combination of exisulind (50 mg/kg/day) and docetaxel (5 mg/kg 6× or 5 mg/kg 3× reduced to 2.5 mg/kg 3×) had a 60% survival rate at the end of a 80-day study period. Conversely, untreated control rats or rats treated with each agent alone or in combination at a lower dose died at earlier time points (40–79 days). Lungs harvested from the above combination treatment groups at death or on sacrifice at the completion of the study displayed histological differences. Lungs harvested from rats treated with high dose combinations of exisulind/docetaxel had a reduction in both tumor mass and distal metastases compared with rats treated with either lower combination doses or each drug alone.

In our current study, these lung tissue specimens were additionally analyzed to determine the molecular mechanisms associated with the increase in survival observed previously. Tissue samples were labeled to determine the level of cGMP PDEs 5 and 1, apoptosis levels and proliferative rates within the in vivo orthotopic tumors. These data were supplemented by in vitro experiments on cultured human A549 NSCLC cells that determined cGMP PDE enzymatic activity and apoptosis levels. Comparing the effects of exisulind on A549 cells both in culture and within in vivo orthotopic tumors resulted in an additional understanding of its biological effects alone or in combination with docetaxel.

Materials and Methods

Cell Culture. The NSCLC line, A549, was obtained from the American Type Culture Collection (Rockville, MD) and maintained in F12K medium (Invitrogen, Carlsbad, CA) supplemented with 5% FCS, 1% glutamine, and 1% antibiotic/antimycotic solution. Cell cultures were grown in 5% CO2 with 100% humidity.

Exisulind and Docetaxel. Exisulind was provided by Cell Pathways, Inc. (Horsham, PA) and docetaxel by Aventis Pharmaceuticals (Bridgewater, NJ). Exisulind suspension was prepared for animal studies in 0.5% carboxymethylcellulose (low viscosity; Aldrich Chemical Company, Milwaukee, WI) and administered p.o. to the rats by single daily gastric gavage at doses of 25 or 50 mg/kg body weight. The stock solution of docetaxel was prepared in DMSO, and the working concentrations of 2.5 or 5.0 mg/kg were diluted in sterile water and injected i.p. once weekly for 5 or 6 consecutive weeks. For cell culture experiments, stock solutions of exisulind were prepared in 100% DMSO (500 mM) and diluted in growth medium with DMSO ≤0.1% or less.

Immunofluorescence. Subconfluent A549 cultures were treated, 24 h after seeding, with either DMSO alone (vehicle control) or exisulind as indicated. Floating cells were collected and pooled with adherent cells (detached by trypsin treatment). These total cell fractions were deposited on microscope slides with a Shandon cytospin (500 × g for 2 min). Cells to be labeled with antibodies to active caspase 3 and p85 PARP fragment (Promega, Madison, WI) were fixed in 3% paraformaldehyde for 10 min, and permeabilized with 0.5% Triton-100 for 2 min and washed for 2 min in PBS.

Cells to be labeled for caspase cleaved cytokeratin 18 using the M30 monoclonal antibody (Roche, Indianapolis, IN) were fixed for 30 min in −20°C methanol, air-dried, and rehydrated in PBS for 2 min. Primary antibodies were incubated as per the manufacturer’s protocols. Cells were washed twice for 5 min each in PBS between each antibody incubation. A species-specific FITC or Cy3-conjugated secondary antibody was used to visualize the distribution of the primary antibody within the cells. Slides were counterstained with 4′,6-diamidino-2-phenylindole (0.5 μg/ml in PBS), mounted in Vectashield antifade solution (Vector Laboratories, Inc., Burlingame, CA) and observed using an Olympus IX-70 microscope. Digital images were collected using a Spot camera (Diagnostic Instruments, Sterling Heights, MI), annotated, and arranged in Adobe Photoshop 7.

PDE Isozyme Fractionation and Assay. A549 cells were cultured, as described above, to confluence in 20 150-cm2 flasks. Approximately 400 million cells were manually homogenized in a buffer containing 20 mM Tris acetate, 5 mM magnesium acetate, 0.1 mM EDTA, 1.0% Triton X-100, and protease inhibitors (10 mM benzamidine, 10 μM tosyl-lysyl-chloromethylketone, 20 mM aprotinin, 2 μM leupeptin, and 2 μM pepstatin A) at pH 7.5 using a glass tissue grinder with a Teflon pestle. After ultracentrifugation at 100,000 × g at 4°C for 1 h, the supernatant was diluted 5-fold with a lower salt form of the homogenation buffer containing 5 mM Tris acetate and no Triton X-100. This sample was loaded at 1 ml/min onto an 18-ml DEAE-Trisacryl M column (BioSep, Rockland, MA) using a Pharmacia AKTA/fast protein liquid chromatography system. The column was washed with 8 ml Tris acetate, 5 mM magnesium acetate, and 0.1 mM EDTA (pH 7.5), and PDEs eluted with a gradient of 0–1 M sodium acetate in Tris acetate buffer at a flow rate of 1 ml/min into 1.5-ml fractions. [3H]cGMP substrate (0.25 μ Ci/mM; 300,000 dpm) were used to differentiate isozymes, as described previously (23). IC50s were determined using Graphpad Prism software. Reagents were from Sigma (St. Louis, MO), except calmodulin (bovine brain) was from Biomol (Plymouth Meeting, PA).

Orthotopic Rat Tumor Model. Animals were implanted with tumor cells and treated as described earlier (22). Briefly, 6–8-week old female athymic nude rats were obtained from the National Cancer Institute and maintained in pathogen-free conditions at the Animal Resources Center, University of Colorado Health Sciences Center. One day before the tumor implantation rats were treated with 400 cGy of total body irradiation (Co60) to increase the immunosuppression and the rate of tumor cell implantation. Human A549 adenocarcinoma cells (1 × 107 cells in 100 μl of saline) were carefully instilled into the left lung by intratracheal administration through a 3-inch 22-gauge catheter (Popper & Sons, Inc., New Hyde Park, NY) bypassing the trachea and left bronchus of each animal. Rats were anesthetized with Metofane inhalation and held upright at 60° on a slant platform with a rubber band through its incisor teeth. The animals usually recovered from the 2–3 min procedure in <5 min.

Seven days after tumor implantation, rats were randomly divided into groups of 8–15 animals and treated with vehicle solutions (controls) or experimental agents (exisulind and/or docetaxel). Rats were treated with exisulind (25 or 50 mg/kg) by single oral daily gavage with a feeding needle (18-gauge) until the day of sacrifice, alone or in combination with do-
cetaxel. Docetaxel alone or in combination (2.5 or 5 mg/kg) was injected i.p. once per week for 5–6 weeks. One combination group was treated with exisulind (50 mg/kg) and docetaxel 5 mg/kg for three injections followed by three 2.5 mg/kg injections. Body weights of animals were measured once a week, and the animals were monitored closely for clinical signs on a daily basis. On day 21 after tumor implantation, three rats from each group were sacrificed, their lungs were removed, fixed in 10% buffered formalin, and paraffin-embedded. The left lung lobe was longitudinally cut into 5-μm sections, mounted on microscope slides, and stored at room temperature until use.

**IHC and TUNEL Analysis.** Rat lung tissue slides were processed for either IHC or TUNEL analysis. IHC specimens were dewaxed in xylene and rehydrated through an ethanol series. Endogenous peroxidase activity was eliminated by preincubation in 0.3% H₂O₂. Primary antibodies used following the manufacturers recommended dilution and protocol included: Ki 67 (DAKO, Carpinteria, CA), M30 (Roche), active caspase 3 (Promega), and PDE1 (Fabgennix, Shreveport, LA). Antibodies to PDE5 (peptide sequence AQLYETSLLN-KRNQV amino acid residues 317–332 in PDE5A1) were raised in sheep, affinity-purified, and used at a dilution of 1:500. The slides were then incubated with an appropriate biotinylated secondary antibody, washed, and incubated with streptavidin-conjugated horseradish peroxidase. The slides were developed using ABC elite 3,3'-diaminobenzidine (Vector), counterstained with hematoxylin, dehydrated through an ethanol and xylene series, and mounted in Permount (Fisher Scientific, Fair Lawn, NJ). TUNEL analysis was performed using the terminal deoxynucleotidyl transferase-FragEL kit (Oncogene Research Products, La Jolla, CA) as per the manufacturer protocol. All of the IHC and TUNEL slides were scored using an Olympus BX-40 microscope. The numbers of positive cells per either 20× (TUNEL, M30 or active caspase 3) or 40× (Ki-67) field of views were scored. Statistics were determined using Graphpad Prism software (one-way ANOVA and Tukey’s multiple comparison test).

**Results**

**Exisulind Induces Apoptosis in A549 Cells.** The induction and progression of apoptosis can be cell type and condition dependent. Therefore, we investigated exisulind-induced apoptosis in cultured A549 cells using multiple apoptotic markers. Control and exisulind-treated cells were labeled with antibodies to the active form of caspase 3, caspase-cleaved cytokeratin 18, or the p85 fragment of PARP. This allowed us to determine apoptotic levels within treated A549 cells at multiple, early time points along the apoptotic pathway. Previous rat studies have determined that oral administration of exisulind results in serum drug concentrations ranging from 247 to 392 μM (5). Therefore, we treated A549 cells for 5 days with concentrations of exisulind that were under or within that range (100, 200, and 300 μM). To ensure that a representative distribution of the entire cell population was analyzed, both floating and adherent populations were collected and combined before processing. Apoptotic cells were found using all three of the markers (Fig. 1). Apoptotic rates within A549 cells were determined by scoring the number of cells that contained the apoptotic marker per 500 consecutive cells. The data from all three of the apoptotic markers were then averaged together. The A549 apoptosis rates increase in a dose-dependent fashion on treatment with either 100, 200, or 300 μM exisulind. Upon exisulind treatment the average levels of apoptosis increased from a background of 1.3% in the control cells to 2.9, 5.5, and 10.2% in the 100, 200, and 300 μM exisulind treated cultures, respectively (Fig. 1A). The increase in apoptosis observed in the 200 and 300 μM exisulind-treated cultures were both significant with Ps of <0.05 and <0.001, respectively. Representative images using all three of the apoptotic markers are shown in Fig. 1 (B–E). These data indicate that exisulind-induced apoptosis in A549 cells occurs at concentrations obtainable within in vivo rat serum and can be identified using multiple apoptotic markers.

**cGMP PDE Immunoreactivity within Orthotopic A549 Tumors.** The relative level of the cGMP PDEs 1 and 5, targets of exisulind, within the orthotopic A549 tumors was determined by IHC analysis. Lung tissue sections from the control group of nude rats containing orthotopic A549 tumors were harvested 21 days after implantation and processed for IHC. Affinity purified polyclonal antibodies specific for PDE1 (all subfamilies) or PDE5 A1 and A2 were used to label the specimens. Tumor cells displayed higher levels of PDE5 immunoreactivity compared with the surrounding normal bronchial epithelial or alveolar cells (Fig. 2A). The specificity of the immunoreactivity observed using our PDE5 antibodies was confirmed by comparing labeling intensities of serial lung tumor sections using the affinity-purified serum (Fig. 2B) or the serum preincubated with a PDE5 GST fusion protein corresponding to the bovine PDE5 (1) sequence Val¹⁵⁵-Asp³⁹³ (Fig. 2C). The vast majority of the labeling is lost after blocking indicating that the observed signal is because of increased levels of PDE5.

The immunolabeling of PDE1 within the lung tumors is also increased compared with normal alveolar cells, albeit not as intensely as PDE5 (Fig. 2D). This is not surprising given the high levels of endogenous PDE1 within normal lung tissue (24, 25). These data confirm that there is an increase in the levels of cGMP PDEs within the orthotopic A549 tumors.

**PDE Isozyme Expression in A549 Cells and Sensitivity to Exisulind.** We identified which PDE isoforms are expressed in the A549 cell line used for the formation of the orthotopic lung tumors. Cell lysates were fractionated by anion-exchange chromatography (Fig. 3A) and found to express three peaks of cGMP PDE activity. The first peak of activity (fraction 30) was activated by calcium and calmodulin, indicative of PDE1 isoform activity (Fig. 3B). The second peak (fraction 48) contained overlap from the first and third peaks, but also was partially inhibited with 100 nM of the PDE5-specific inhibitor E4021. The location of this second peak matched that seen previously for PDE5 from SW480 and HT29 colon cell lines, and the HT1376 bladder cell lines (3, 7, 13). Much of the cAMP PDE activity was inhibited by 10 μM rolipram, a PDE4-specific inhibitor. The cAMP activity...
that was not inhibited by rolipram coincided with the third peak of cGMP PDE activity (fraction 72). Addition of 5 μM cGMP showed inhibition of the cAMP activity consistent with the presence of the dual substrate isoform PDE3. In addition, the PDE3 selective inhibitors of 1 μM cilostamide and 1 nM trequinsin inhibited ~60% of the cGMP PDE activity (data not shown). High concentrations of exisulind (600 μM) inhibited most of the cGMP PDE activity from each of the three peaks even in the presence of calcium/calmodulin. PDE1 from fractions 24–33 when tested for exisulind inhibition on calcium/calmodulin activated cGMP PDE showed an IC_{50} of 70 μM (inset). Exisulind tested on the second peak with some overlap from the other peaks showed an IC_{50} of 110 μM (data not shown) similar to that seen previously for PDE5 from SW480, HT29, and HT1376 (3, 7, 13).

Drug-induced Apoptosis in Orthotopic A549 Tumors.

The apoptotic indexes in the orthotopic tumors from all of the treatment groups were determined in a manner similar to that performed on the A549 cells in vitro. Three independent apoptotic markers identified apoptotic cells: antibody markers to active caspase 3 (AC3), caspase cleaved cytokeratin 18 (M30), and p85 PARP fragment. The percentage of apoptosis for each dose of exisulind was obtained by averaging the data from the three apoptotic markers. Statistical significance is relative to control (0% exisulind) cultures. Representative images of apoptotic A549 cells identified by each marker are shown (B–E). Cells were treated with either 0.1% DMSO (B) or 600 μM exisulind for 24 h (C–E). Apoptotic cells were identified using antibodies to active caspase 3 (B and C), caspase cleaved cytokeratin 18 (D), and p85 PARP fragment (E). All of the cells were counterstained with 4',6-diamidino-2-phenylindole; bars, ±SD.
high-dose docetaxel, and the high-dose exisulind/docetaxel combination. Tissues labeled with antiactive caspase 3 antibodies or TUNEL gave similar images (data not shown). All three of the markers indicate that there was an increase in apoptosis, compared with controls, on treatment with exisulind and docetaxel. The apoptotic levels determined from each of the three markers for each treatment group is shown in Fig. 5, A–C. Treatment with exisulind and/or docetaxel resulted in a dose-dependent increase in the number of apoptotic cells as determined by each apoptotic marker. To quantify the cumulative apoptosis levels in all of the treatment groups using all three of the apoptosis markers, the data obtained from each individual marker was normalized relative to its own untreated control group and averaged between all three of the apoptosis markers. The resultant cumulative apoptotic index showed an exisulind dose-dependent increase in apoptosis (Fig. 5D). In addition, each treatment group showed increased apoptosis in the lung tumors when comparing higher versus lower exisulind doses. As single agents, exisulind-induced apoptosis was consistently higher than that observed for docetaxel. Additionally, exisulind increased the apoptotic rates at both doses with or without docetaxel treatment.

Only non-necrotic areas of the tumors were analyzed to ensure that only true apoptotic cells were scored. The percentage of each tumor that was necrotic within the control, high-dose exisulind, high-dose docetaxel, and the high-dose docetaxel, and the high-dose exisulind/docetaxel combination. Tissues labeled with antiactive caspase 3 antibodies or TUNEL gave similar images (data not shown). All three of the markers indicate that there was an increase in apoptosis, compared with controls, on treatment with exisulind and docetaxel. The apoptotic levels determined from each of the three markers for each treatment group is shown in Fig. 5, A–C. Treatment with exisulind and/or docetaxel resulted in a dose-dependent increase in the number of apoptotic cells as determined by each apoptotic marker. To quantify the cumulative apoptosis levels in all of the treatment groups using all three of the apoptosis markers, the data obtained from each individual marker was normalized relative to its own untreated control group and averaged between all three of the apoptosis markers. The resultant cumulative apoptotic index showed an exisulind dose-dependent increase in apoptosis (Fig. 5D). In addition, each treatment group showed increased apoptosis in the lung tumors when comparing higher versus lower exisulind doses. As single agents, exisulind-induced apoptosis was consistently higher than that observed for docetaxel. Additionally, exisulind increased the apoptotic rates at both doses with or without docetaxel treatment. Only non-necrotic areas of the tumors were analyzed to ensure that only true apoptotic cells were scored. The percentage of each tumor that was necrotic within the control, high-dose exisulind, high-dose docetaxel, and the high-dose exisulind/docetaxel combination. Tissues labeled with antiactive caspase 3 antibodies or TUNEL gave similar images (data not shown). All three of the markers indicate that there was an increase in apoptosis, compared with controls, on treatment with exisulind and docetaxel. The apoptotic levels determined from each of the three markers for each treatment group is shown in Fig. 5, A–C. Treatment with exisulind and/or docetaxel resulted in a dose-dependent increase in the number of apoptotic cells as determined by each apoptotic marker. To quantify the cumulative apoptosis levels in all of the treatment groups using all three of the apoptosis markers, the data obtained from each individual marker was normalized relative to its own untreated control group and averaged between all three of the apoptosis markers. The resultant cumulative apoptotic index showed an exisulind dose-dependent increase in apoptosis (Fig. 5D). In addition, each treatment group showed increased apoptosis in the lung tumors when comparing higher versus lower exisulind doses. As single agents, exisulind-induced apoptosis was consistently higher than that observed for docetaxel. Additionally, exisulind increased the apoptotic rates at both doses with or without docetaxel treatment. Only non-necrotic areas of the tumors were analyzed to ensure that only true apoptotic cells were scored. The percentage of each tumor that was necrotic within the control, high-dose exisulind, high-dose docetaxel, and the high-dose
exisulind/docetaxel combination groups were measured. Across the treatment groups necrosis ranged from 7.8 to 18.5% and was not statistically significant.

**Docetaxel but not Exisulind Decreases the Proliferation Rate within Orthotopic A549 Tumors.** The effects of exisulind and docetaxel on *in vivo* proliferation within the orthotopic A549 lung tumors were investigated by labeling tumor sections from all of the treatment groups with the proliferation marker Ki-67. Animals treated with 2.5 or 5.0 mg/kg docetaxel had a significant decrease in the number of proliferating cells. These observations were quantified by scoring the number of Ki-67-positive cells per ×40 field of view. The docetaxel effect was 21.5% inhibition with the lower dose (2.5 mg/kg) and 48.5% at the higher dose (5.0 mg/kg). Exisulind at either 25 or 50 mg/kg/day resulted in no significant decrease in the number of Ki-67-positive cells. Representative images from the control, exisulind 50 mg/kg, docetaxel 5 mg/kg, and combination exisulind 50 mg/kg docetaxel 5 mg/kg are shown in Fig. 6, A–D. Animals treated with a combination of both exisulind and docetaxel did not show an additional decrease in proliferation over that observed with the corresponding docetaxel treatment alone. Therefore, in this *in vivo* orthotopic model, exisulind does not lower the proliferative rate of the A549 tumor cells. The proliferation data for all of the treatment groups are summarized in Fig. 6E.

**Discussion**

These studies report that exisulind treatment of human NSCLC A549 cells grown in culture or administration to *in vivo* orthotopic tumors in a nude rat model increased apoptotic indexes. Determination of apoptosis was designed to minimize artifacts specific to a single apoptotic marker. The current standard of identifying apoptotic cells involves demonstration of specific oligonucleosomal DNA fragments either by double antibody ELISA or enzymatic TUNEL analysis. However, this method can give inaccurate results because of necrotic cells with fragmented DNA, early apoptotic cells lacking digested DNA, variation in sample preparation, and enzymatic activity (26–28). Recent reports also indicate potential shortcomings of relying on only the TUNEL apoptotic marker when quantifying apoptosis within tissue sections (29–31). To obviate these problems antibodies to specific apoptotic proteins present at various stages of the apoptotic pathway were used. Specifically, the active form of caspase 3 and caspase-cleaved cytokeratin 18 were studied. As expected, the specific number of apoptotic cells identified by each individual marker varied one to another; however, each marker indicated that the indexes for both *in vitro* and *in vivo* apoptosis increased in a dose-dependent fashion after exisulind exposure. These studies establish that apoptosis is induced by exisulind administration in A549 cells of *in vivo* tumors, as has been seen in numerous cancer cell lines.

Apoptosis, once initiated, is a relatively rapid event, and any resultant apoptotic bodies are engulfed by phagocytosis in a short period of time (32, 33). Therefore, within the orthotopic A549 tumors it is relatively difficult to identify the rare cells actively undergoing apoptosis. This would imply that the exisulind-induced increases in the *in vivo* apoptosis levels might be conservative indexes of its effects. Consequently, even moderate increases in apoptosis observed in a
single tissue section may only be a fraction of the total apoptotic effect occurring within the tumor as a whole. The variables inherent in this type of in vivo apoptotic analysis dictate that the results be interpreted with a degree of caution. The fact that the apoptotic rates of combination treated groups were not exactly additive of the single treatment dictates that the results be interpreted with a degree of caution. The fact that the apoptotic rates of combination treated groups increased their apoptosis levels quicker than the low-dose groups and, therefore, by day 21 the window of maximal apoptosis in the high-dose treatment groups may have passed. The lack of a statistical difference between the levels of apoptosis in the 5 mg/kg docetaxel plus 25 mg/kg or 50 mg/kg exisulind may reflect variable time courses.

The concentrations of exisulind used in our in vitro cell culture apoptosis analysis correlates with serum levels obtained in a previous rat cancer model (5). In the rat colon cancer model, exisulind serum levels of 346 µM were obtained without any observable weight decreasing toxicity. Our ability to induce apoptosis in A549 cells beginning with concentrations of 100 µM exisulind also correlates with the IC50 for PDE1 and 5 inhibition of 70 µM and 110 µM, respectively. The correlation between the increases in apoptosis within cultured A549 cells in vitro and the increases in apoptosis seen within the A549 orthotopic tumors in vivo is of importance, because it provides support for extrapolating the observations seen in culture to potential biological effects predicting in vivo drug activity. These data suggest that the inhibition of PDE1 and 5 by exisulind can induce apoptosis in both an in vitro and in vivo environment. In A549 cells exisulind also inhibits PDE1 expressed in this lung tumor along with PDE5 to prevent cGMP hydrolysis in this NSCLC model. This is supported by previous reports of exisulind inhibiting the cGMP hydrolysis by PDE family members including PDE2 and 5 in colon, pancreatic, and bladder cancer cell lines. In colon cancer cells the inhibition of cGMP PDE activity on exisulind treatment results in a sustained increase in cGMP, but not cAMP levels within the cells (7). This sustained cGMP elevation activates PKG, which plays a critical role in the mechanism by which exisulind induces apoptosis. To date, two direct effects of exisulind-dependent PKG activation have been described: first, the attenuation of β-catenin levels within cells treated with exisulind (7, 15). This facilitates a decrease in both nuclear and cytoplasmic levels of β-catenin through degradation by ubiquitin-dependent proteolysis. Second, PKG directly phosphorylates mitogen-activated protein kinase kinase kinase 1, which leads to the downstream activation of c-Jun NH2-terminal kinase 1 (13, 14, 34). Both of these molecular effects of exisulind inhibit cGMP hydrolytic activity. Therefore, the subsequent induction of apoptosis in A549 cells is likely because of a mechanism of action similar to that observed in colon cancer cells.
Docetaxel exhibits two major biological effects: the first being decreased proliferation because of direct binding to and stabilization of microtubules causing a G2/M block (35) and the second being apoptosis induction. The data obtained using the proliferation marker Ki-67 indicate that exisulind and docetaxel have disparate effects on in vivo tumor proliferation rates. Neither dose of exisulind used in this study lowered the proliferative rate within the tumors. In contrast, docetaxel lowered the proliferative rate within the tumor. Rats treated with the combination of exisulind and docetaxel displayed proliferative rates similar to rats treated with docetaxel alone. Therefore, exisulind does not affect the gross proliferative rate of A549 orthotopic tumors in vivo.

We reported previously an increase in survival of nude rats harboring orthotopic A549 tumors that were treated with a combination of exisulind (50 mg/kg/day) and docetaxel (5 mg/kg). In the same study rats treated with either exisulind at 25 or 50 mg/kg/day, or those treated with docetaxel at 2.5 or 5 mg/kg alone or in low-dose combinations did not survive significantly longer than control rats. As single agents the biological effects of docetaxel (decrease in proliferation) or exisulind (increase in apoptosis) were not sufficient to increase survival. However, these studies show that the treatment group with the highest survival (exisulind 50 mg/kg and docetaxel 5 mg/kg) had an exisulind-dependent increased apoptosis rate coupled with a docetaxel-dependent de-
increased proliferative rate. As commonly observed in the clinical setting, the growth rate of the A549 tumors and metastasis in this model could only be overcome through the combined effects of these compounds. These data support the use of combining exisulind and docetaxel as a rationale combination to increase apoptosis and decrease proliferation as a treatment for NSCLC.

References
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