Effect of Proton Pump Inhibitor Pretreatment on Resistance of Solid Tumors to Cytotoxic Drugs

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Background: Resistance to antitumor agents is a major cause of treatment failure in patients with cancer. Some mechanisms of tumor resistance to cytotoxic drugs may involve increased acidification of extracellular compartments. We investigated whether proton pump inhibitors (PPIs), currently used in the anti-acid treatment of peptic disease, could inhibit the acidification of the tumor microenvironment and increase the sensitivity of tumor cells to cytotoxic agents.

Methods: We pretreated cell lines derived from human melanomas, adenocarcinomas, and lymphomas with the PPIs omeprazole, esomeprazole, or pantoprazole and tested their response to cytotoxic drugs in cell death assays. We also evaluated extracellular and intracellular pH and vacuolar-H^+-ATPase (V-H^+-ATPase) expression, distribution, and activity in PPI-pretreated cells by using western blot analyses, immunocytochemistry, laser scanning confocal analysis, and bioluminescence assays. Finally, we evaluated human melanoma growth and cisplatin sensitivity with or without omeprazole pretreatment in xenografted SCID/SCID mice.

Results: PPI pretreatment sensitized tumor cell lines to the effects of cisplatin, 5-fluorouracil, and vinblastine, with an IC_{50} value reduction up to 2 logs. PPI pretreatment was associated with the inhibition of V-H^+-ATPase activity and increases in both extracellular pH and the pH of lysosomal organelles. PPI pretreatment induced a marked increase in the cytoplasmic retention of the cytotoxic drugs, with clear targeting to the nucleus in the case of doxorubicin. In in vivo experiments, oral pretreatment with omeprazole was able to induce sensitivity of human solid tumors to cisplatin.

Conclusion: Our results open new possibilities for the treatment of drug-resistant tumors through combination strategies based on the use of well-tolerated pH modulators such as PPIs.

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agents may reverse anthracycline resistance in multidrug-resistant cells with an expanded acidic lysosomal compartment (18). Another approach may be to inhibit the function of the pumps that establish the pH gradient. Vacuolar-H⁺-ATPases (V-H⁺-ATPases) represent a major mechanism in the regulation of cellular pH (19). V-H⁺-ATPases pump protons across the plasma membrane and across the membranes of a wide array of intracellular compartments (19). Some human tumor cells, particularly those selected for multidrug resistance, exhibit enhanced V-H⁺-ATPase activity (20–25). These data suggest that the enhanced V-H⁺-ATPase activity increases the acidity of intracellular vesicles, allowing drug sequestration and consequently the development of multidrug resistance. Some molecules that inhibit V-H⁺-ATPases and may reverse tumor resistance to cytotoxic drugs have been identified (5,6,18,26). However, their toxicity and poor results in preclinical testing have limited their development as therapeutic agents.

A class of H⁺-ATPase inhibitors called proton pump inhibitors (PPIs) has emerged as the drug class of choice for treating patients with peptic disease, including gastroesophageal reflux disease and duodenal or gastric ulcers. These anti-acid drugs inhibit gastric acid secretion by targeting the gastric acid pump (26–30). Their effects at the cellular level are mediated by direct inhibition of V-H⁺-ATPase (31–33). PPIs, which include omeprazole, esomeprazole, lansoprazole, pantoprazole, and rabeprazole, are substituted 2-pyridyl-methylsulfinyl benzimidazoles that share a similar core structure (30). These agents are protonatable weak bases with pKᵦ (negative logarithm of the acid ionization constant) values of approximately 4, with the exception of rabeprazole, which has a pKᵦ of 5. Therefore, PPIs accumulate selectively in acidic spaces with a pH of less than 4. In such acidic environments, protonation of the pyridine and benzimidazole nitrogens results in formation of a tetracyclic sulfenamide, which is the active form of the drug (30).

The aim of our work was to determine whether PPIs can restore drug sensitivity to drug-resistant cells by inhibiting the increased acidification of both the intracellular compartments and the extracellular spaces in the tumor, possibly through a V-H⁺-ATPase–mediated mechanism. We examined this hypothesis both in vitro, using human tumor cell lines displaying intrinsic or acquired resistance to antitumor drugs, and in vivo, in an animal model represented by CB.17 SCID/SCID mice engrafted with human tumor cells.

**Materials and Methods**

**Drugs**

Omeprazole and esomeprazole (AstraZeneca, Mölndal, Sweden) and pantoprazole (Byk Gulden, Konstanz, Germany) sodium salts were resuspended in normal saline (0.85%) at a concentration of 1 mg/mL immediately before use. Cisplatin (Aventis, Schiltigheim, Germany) was resuspended in phosphate-buffered saline (PBS) at a stock concentration of 1 mg/mL and stored at −20 °C. This stock solution was thawed immediately before use and not refrozen. 5-Fluorouracil (5-FU; Teva Pharma, Haarlem, The Netherlands) was supplied in solution at a concentration of 50 mg/mL and was stored at room temperature, as indicated by the supplier. Vinblastine sulfate (Eli Lilly, Paris, France) was suspended in ethanol:distilled water (1:1000) at a concentration of 0.1 mg/mL; this stock solution was stored at 4 °C and used within 3 days. Vinblastine-bodipy was obtained from Molecular Probes (Eugene, OR) and stored at −20 °C as stock solution at 0.1 mg/mL in dimethyl sulfoxide (DMSO).

**Tumor Cells**

Human drug-resistant tumor cell lines (from 22 melanomas, two colon adenocarcinomas, two breast cancers, and two ovarian carcinomas) were supplied by Istituto Nazionale per lo Studio e la Cura dei Tumori, Milan, Italy, and were obtained from primary lesions. HeLa and CEM-CCRF T-lymphoblastoid cells (referred to as CEM cells) were obtained from American Type Culture Collection (Manassas, VA). The CEM-VBL100 cell line, a multidrug-resistant P-glycoprotein–overexpressing variant of CEM cells, was selected in our laboratory. These cells were produced by exposing parental CEM cells to increasing sublethal concentrations of vinblastine up to 100 ng/mL (34). All cells were cultured in RPMI-1640 medium supplemented with 10% fetal calf serum (FCS) and antibiotics, in humidified 5% CO₂.

**Dose–Response Curves**

Tumor cells growing in suspension (CEM and CEM-VBL100 cells) were plated at 1.5 × 10⁵ cells per milliliter in 24-well plates at one milliliter per well. Tumor cells growing in adherence (melanoma and colon, breast, and ovarian cancer cell lines) were plated at 3 × 10⁵ cells per well in 24-well plates. After 24 hours, each cell line was treated with three to five logarithmic dilutions of each of the drugs. Each drug was tested on each cell type in triplicate. In combined treatment experiments, cells were pretreated for 24 hours with the PPI and then treated for an additional 24 hours with the antitumor drug.

**Cytotoxicity Assays**

Trypan blue exclusion method. After treatment, cells growing in suspension were collected, centrifuged 5 minutes at 500g, and resuspended in 30 µL of PBS. Cells growing in adherence were collected by pooling cells from the medium (i.e., dead cells) and adherent (live) cells obtained by trypsinization. Cells were then centrifuged (10 minutes at 500g) and resuspended in PBS (50–100 µL). An aliquot of each cell line suspension was diluted 1:1 (vol/vol) with 0.4% trypan blue. After 5 minutes, cells were loaded on a hemocytometer, and both live (unstained) and dead (blue-stained) cells were counted under a light microscope. The percentage of dead cells was then determined. Each treatment condition was tested at least in triplicate, and the mean value (% dead cells) was determined.

**LIVE/DEAD Viability/Cytotoxicity assay.** The LIVE/DEAD Viability/Cytotoxicity Assay (Molecular Probes, Eugene, OR) is a two-color fluorescence cell viability assay that is based on the simultaneous determination of live and dead cells with two probes (i.e., calcine AM and ethidium homodimer 1, respectively) that measure two recognized parameters of cell viability (i.e., intracellular esterase activity and plasma membrane integrity, respectively). The optimal dye concentrations for the cell types used in this study were determined according to manufacturer’s instructions. After treatment, cells were collected, centrifuged, and resuspended in PBS as described above, treated with calcine AM and ethidium homodimer 1 at the final concentrations of 0.1 µΜ and 1 µΜ, respectively, and left at room temperature for 30 minutes. The cells were then washed in PBS,
and the samples were analyzed with a FACScan cytometer (Becton Dickinson, Franklin Lakes, NJ). At least 20,000 events were acquired with at least three replicates for each treatment condition.

**Determination of Cell Cycle**

Cell cycle distribution was analyzed as described (35). Briefly, 0.2 × 10^6 cells were washed in PBS and incubated overnight in 400 μL of ice-cold 70% ethanol. After two washes in PBS, the cells were incubated for 30 minutes at 37 °C in PBS containing 100 μg/mL DNase-free RNAse and 40 μg/mL propidium iodide. The samples were then acquired with a FACScan cytofluorimeter that collected the fluorescence signal in FL2 channel on a linear scale and analyzed using CellQuest and ModFIT software (Becton Dickinson).

**Laser Scanning Confocal Microscopy Analysis**

For laser scanning confocal microscopy (LSCM) analysis of drug uptake and efflux, cells (3 × 10^4 cells per well) were plated on glass coverslips in 24-well plates. After the cells had adhered, the medium was replaced with fresh medium (1 mL) supplemented with 0.1 μg/mL vinblastine-bodipy or 5 μM doxorubicin was added to cells. After 6 hours (uptake phase), drug-containing medium was removed and the cells were incubated in drug-free medium for up to 36 hours (efflux phase). Cells were observed with a Leica TCS SP2 spectral confocal microscope, as previously described (36). Doxorubicin fluorescence was excited with a 488-nm argon laser, and emission lines were collected after passage through a DD 488/543 filter in a spectral window ranging from 515 to 600 nm. Vinblastine-bodipy fluorescence was excited at the 488-nm argon laser line, and emission lines were collected after passage through a DD 488/543 filter in a spectral window ranging from 515 to 565 nm. Pixel intensity was analyzed with the Quantify Leica TCS SP2 program. Mean pixel intensities, evaluated on 255 gray levels, were calculated by analyzing a total cell area of 25,000 μm^2 for each sample. The calculation was performed on images representing orthogonal maximum projections of 20 optical sections (0.5 μm thick) acquired with the following acquisition parameters: 63.0/1.4 NA objective; image size: 1024 × 1024 pixels; pinhole size: 1 Airy.

**Sodium Dodecyl Sulfate–Polyacrylamide Gel Electrophoresis and Western Blotting**

Sodium dodecyl sulfate–polyacrylamide gel electrophoresis and western blot analysis of V-H^+-ATPase was performed as previously described (35). Briefly, melanoma cells were lysed in Akt buffer (150 mM NaCl, 20 mM Tris–HCl [pH 7.4], 1% NP40, 10% glycerol) supplemented with protease and phosphatase inhibitors (aprotinin, leupeptin, phenylmethylsulfonyl fluoride, sodium orthovanadate) incubated for 15 minutes on ice, and centrifuged at 13,000g for 15 minutes to remove nuclei and cell debris. The protein concentration of the extracts was determined by the Bio-Rad Protein Assay (Bio-Rad, Hercules, CA), following the manufacturer’s instructions. Fifty micrograms of each cell extract was separated on 10% gels and electropholated to a nitrocellulose membrane. Nonspecific binding was blocked by incubating the membrane in 1× PBS with 5% nonfat dry milk. Blots were then incubated with a polyclonal goat anti-human antibody to V-H^+-ATPase subunit C (Santa Cruz Biotechnology, Santa Cruz, CA) and a monoclonal mouse anti-human antibody to actin (Chemicon, Temecula, CA) as a control for protein loading. Antibody binding was detected by incubating the blot with a horseradish peroxidase–conjugated rabbit anti-goat antibody (Jackson Immunoresearch, West Grove, PA) and then with a horseradish peroxidase–conjugated sheep antimouse antibody (Amersham, Piscataway, NJ), respectively. Antibody staining was visualized by enhanced chemiluminescence (Pierce, Rockford, IL).

**Determination of Intracellular ATP Concentration**

We measured the amount of available intracellular ATP in melanoma cell lines as an indirect parameter (37) of the activity of V-H^+-ATPases. We used a commercially available ATP Determination Kit (Molecular Probes) that is based on luciferase activity. Cells were cultured for 24 hours in 24-well plates at a density of 0.05 × 10^5 cells per well in the presence of a PPI (1 μg/mL). ATP determination was performed by using the ATP Determination Kit, as reported elsewhere (37). Briefly, 10,000 cells were collected, washed in PBS, resuspended in 100 μL of distilled water, and boiled for 5 minutes. Ten μL of each sample (1000 cells) or 10 μL of each dilution point of ATP standard solution (0–1000 nM) was added to 90 μL of reaction solution in the wells of 96-well plates. After 5 minutes, plates were analyzed with a luminometer (Wallac 1420 VICTOR; Wallac, Boston, MA). The amount of intracellular ATP was determined by plotting the light output values of each sample against those of the ATP standard solutions. All experiments were run in duplicate, and mean values were calculated.

**Determination of Extracellular pH**

MelM1, MelM6, MelM9, and MelP6 cells were incubated in a medium (pHmed) made up of normal saline (80%), RPMI-1640 (10%), and FCS (10%). This composition minimized the buffering activity of phosphate and bicarbonate in the medium but still contained sufficient nutrients and growth factors to support cell growth (data not shown). Untreated cells were harvested, washed twice in pHmed, and then incubated at 2 × 10^6 cells per milliliter in pHmed for 3 hours at 37 °C. The cells were then collected by centrifugation (10 minutes at 500g), and the supernatant was harvested for pH measurements. pH was determined by reading each sample at 32 °C, in triplicate, using a Titroprocessor 726 pH meter (Metrohm, Herisau, Switzerland) equipped with a glass microelectrode (LongLife; Metrohm). Data are presented as means with 95% confidence intervals (CIs).

**Staining of Acidic Vesicles With a pH Indicator**

The LysoSensor Green DND-189 probe (Molecular Probes), which accumulates in acidic vesicles and exhibits a pH-dependent increase in fluorescence intensity on acidification, was used according to the manufacturer’s indications to measure the effects of omeprazole treatment on acidic vesicles. Briefly, 5 × 10^5 MelM6 cells were collected after 24 hours of omeprazole treatment (1 μg/mL) and washed twice in PBS. Cells were then incubated for 5 minutes at 37 °C with 500 μL of prewarmed PBS containing 1 μM LysoSensor probe and analyzed by flow cytometry collecting FL1 fluorescence. Untreated and unstained cells were used to set the background fluorescence. The exper-
iment was repeated twice. For analysis of the localization of LysoSensor-positive vesicles, cells were incubated for 1 hour at 37 °C with prewarmed PBS containing the LysoSensor probe and analyzed by LSCM, as described above.

Histology and Immunocytochemistry

Human tumors were fixed in 10% formalin and embedded in paraffin. Four-micrometer sections were cut, stained with hematoxylin and eosin, and examined under the microscope. MelM6 cells (3 × 10^4 cells per chamber) were attached to sterile glass chamber slides (LabTek, Naperville, IL) by overnight incubation in 150 μL of RPMI-1640 per well in a humidified 5% CO₂ atmosphere. Cells were then fixed in cold 70% methanol (10 minutes at 4 °C) and stained with polyclonal goat anti-human V-H⁺-ATPase antibodies (Santa Cruz, CA) using the alkaline phosphatase anti–alkaline phosphatase (APAAP) method (Dako, Glostrup, Denmark), as described (38).

In Vivo Tumor Growth Analyses

Female CB.17 SCID/SCID mice aged 4–5 weeks (Harlan; Correzzana, Milan, Italy) were kept under specific pathogen-free conditions and fed ad libitum. The mice (39) were housed in micro-isolator cages, and all food, water, and bedding were autoclaved prior to use. Each mouse was injected subcutaneously in the right flank with 3 × 10⁶ human melanoma or colon adenocarcinoma cells derived from metastatic lesions that had been resuspended in 0.2 mL of RPMI-1640 containing 10% FCS. Once tumors became evident (at least 0.10 cm, approximately 10 days after the tumor cell injection), PPI (omeprazole, esomeprazole, or pantoprazole sodium salts resuspended in normal saline at a concentration of 15 mg/mL immediately before use) was orally administered by gavage (40) at a dose of 75 mg/kg. Cisplatin was administered by intraperitoneal injection at a dose of 5 mg/kg (41) simultaneously with PPI oral treatment, 24 hours after PPI treatment, or in mice that did not receive any PPI treatment. Tumor dimensions were measured three times per week with calipers. Tumor weight was estimated according to Geran et al. (42) using the following formula: tumor weight (mg) = length (mm) × width² (mm)/2.

At least 10 mice were used for each treatment group. Data are expressed as the mean value of tumor weight with 95% confidence intervals. Mice were monitored for the duration of the in vivo experiments for body weight, hair ruffling, and the presence of diarrhea. All mice were killed at the end of the experiments, within months after the injection of the human tumor cells (following the guidelines of the Istituto Superiore di Sanità).

Statistical Analysis

Data from the LIVE/DEAD Viability/Toxicity assay were recorded and statistically analyzed on a Macintosh computer with CellQuest software. Fluorescence intensity (expressed as median values) was calculated after logarithmically amplified signals were converted to a linear scale. Statistical significance of the difference in mean fluorescence intensity between PPI pretreated and untreated cells was calculated with the parametric Kolmogorov–Smirnov test. Statistical analysis of data from the trypan blue exclusion test was performed with the Student’s t test. Only P values of less than .01 were considered statistically significant. One-way analysis of variance with a pairwise multiple comparison procedure (Tukey test) was used to analyze the statistical significance of tumor weight differences between the treatment groups in the in vivo experiments carried out in tumorgenrafted SCID mice.

RESULTS

Effect of PPI Pretreatment on Drug-Resistant Human Tumor Cells

We first examined whether the PPIs omeprazole, esomeprazole, or pantoprazole could reverse the intrinsic resistance of human tumor cells to cytotoxic drugs. In designing the experiments, we considered that cisplatin can enhance the activity of proton pumps (25), that PPIs must be protonated in an acidic environment to function as PPIs (30), and that PPIs and antitumor drugs, such as cisplatin, 5-FU, and vinca alkaloids, are all weakly basic molecules that would thus compete for sequestration by protonation in acidic microenvironments. However, an important difference between PPIs and the anticancer drugs is that protonation activates PPIs but neutralizes (i.e., inactivates) the weakly basic antitumor drugs. Moreover, tumor cells were cultured in buffered media, which maintains the pH of tumor cell cultures approximately at neutrality. Therefore, under culture conditions, the only possible place where the acidity of the tumor cell microenvironment could reach levels more suitable for PPI protonation (i.e., a pH of <4) is in close proximity to the plasma membrane of the tumor cells. Thus, we hypothesized that the best approach to test the effect of PPIs in inhibiting intrinsic resistance to antitumor drugs would be to avoid possible competition between PPIs and the tumor drugs at the tumor cell level.

We therefore performed a first set of experiments to compare the effect of pretreating tumor cells with a PPI and then treating them with a cytotoxic drug with the effect of treating tumor cells with a PPI and an antitumor drug simultaneously. In these experiments, human tumor cell lines of different histologies (22 melanoma, two colon adenocarcinoma, two breast cancer, and two ovarian carcinoma), all of which had been determined by a trypan blue exclusion assay to be resistant to the cytotoxic effects of cisplatin (Table 1), were treated with cisplatin after a 24-hour pretreatment with omeprazole. The 24-hour PPI pretreatment was chosen on the basis of preliminary experiments showing more reproducible results than those obtained with shorter (6 or 12 hours) or longer (36 hours) PPI pretreatments (data not shown). We used two different approaches to evaluate the cytotoxic effects of cisplatin (% of dead cells after 24 hours of treatment with the drug) that provided fully comparable and reproducible results. Dose–response curves for cisplatin were obtained by pretreating cells for 24 hours with omeprazole or with saline and then treating them with one of three logarithmic dilutions of cisplatin. The results of repeated experiments with three melanoma lines (Fig. 1, A–C) indicated that pretreatment with omeprazole induced susceptibility of melanoma cells to the cytotoxic effect of cisplatin. Similar results were obtained with other tumor cell lines and other PPIs (Table 1). Figure 1, D, shows the results of repeated experiments in which a melanoma cell line was pretreated with esomeprazole and then with cisplatin, providing data similar to those obtained with omeprazole.
Consistent with our hypothesis that omeprazole and antitumor drugs (e.g., cisplatin) would compete for cellular uptake and localization, thus weakening or inhibiting the effects of PPIs, omeprazole did not induce any change in the responsiveness of the same melanoma cells to cisplatin when administered simultaneously with the anticancer drug (Fig. 1, E).

To verify that PPIs could induce effectiveness of different classes of antitumor drugs, we tested the efficacy of omeprazole on tumor cell resistance to 5-FU, using cell lines resistant to this drug (Table 1). Dose–response curves were obtained from tumor cell lines of different histologies that had not been pretreated with omeprazole or that had been pretreated with this PPI for 24 hours and were then treated with five logarithmic dilutions of 5-FU (Fig. 2). Omeprazole pretreatment induced susceptibility to the cytotoxic effect of 5-FU in all the tumor cell lines tested, including two melanoma lines (Fig. 2, A and B) and one colon carcinoma line (Fig. 2, C). Again, simultaneous treatment with the PPI and 5-FU was ineffective (data not shown).

Using the same experimental protocol, we obtained dose–response curves of the effects of five logarithmic dilutions of vinblastine on tumor cell lines pretreated with omeprazole. Again, omeprazole pretreatment of melanoma cell lines intrinsically resistant to vinblastine (Table 1) resulted in their becoming sensitive to the cytotoxic effects of this drug (Fig. 3, A). Similar results were obtained with other PPIs (i.e., esomeprazole and pantoprazole) and other tumor cell lines (Table 1).

We also tested the effect of PPI pretreatment in cells that had been selected in vitro for a multidrug-resistant phenotype. For this analysis, we used CEM-VBL100 cells, which were obtained by selection of the parental human T-lymphoblastoid cell line CEM in a medium containing increasing concentrations of vinblastine (34). The results clearly showed that vinblastine sensitivity was restored in CEM-VBL100 cells after pretreating them with omeprazole (Fig. 3, B). Similar results were obtained with esomeprazole and pantoprazole (not shown). Again, omeprazole was ineffective when administered at the same time as vinblastine (data not shown). Interestingly, omeprazole was able to lower the minimal cytotoxic dose of vinblastine on the CEM drug-sensitive parental line (Fig. 3, C). Thus, PPI pretreatment not only induced susceptibility to anticancer drugs in tumor cells intrinsically resistant to such drugs but also reversed acquired multidrug resistance and increased cytotoxicity of antitumor treatments in drug-sensitive human tumor cells.

### Effects of Omeprazole Treatment on Human Tumor Cell Lines

We next investigated how PPIs interfered with cellular functions that otherwise prevent cytotoxic drugs from exerting their cytotoxic effects. Thus, we analyzed the effects of 24 hours of PPI (omeprazole or esomeprazole) treatment on two human melanoma cell lines (MelP6 and MelM6). PPI treatment had no effect on cell viability or cell cycle progression (Fig. 4, A). We next compared the pH of the medium of PPI-treated and untreated human tumor cells. The results demonstrated that omeprazole impaired the ability of tumor cells to acidify the extracellular medium.
cellular medium as early as 3 hours after treatment (Fig. 4, B). Omeprazole treatment also induced an increase in lysosomal pH (Fig. 4, C). Moreover, LSCM analysis showed that the LysoSensor-positive acidic vesicles lost their secretory behavior and accumulated within the cell cytoplasm (Fig. 4, D). Given the inhibitory effect of PPIs on V-H\textsuperscript{+}\textsubscript{-}ATPase activity in other cellular systems (31–33), we also analyzed both the expression and activity of V-H\textsuperscript{+}\textsubscript{-}ATPases in four human tumor cell lines (MelM1, MelM6, MelM9, MelP6) treated with omeprazole. Treatment with different doses of omeprazole did not induce a change in V-H\textsuperscript{+}\textsubscript{-}ATPase protein levels (Fig. 5, A). However, the subcellular localization changed, with V-H\textsuperscript{+}\textsubscript{-}ATPase-expressing vesicle-like structures accumulating in perinuclear regions of the omeprazole-treated cells (Fig. 5, B, lower panel). In fact, in untreated cells, V-H\textsuperscript{+}\textsubscript{-}ATPase staining appeared widely diffuse in the cytoplasm and beneath the cell membrane (Fig. 5, B, upper panel). This change in localization was accompanied by a change in levels of intracellular ATP. In fact, cells treated with omeprazole or esomeprazole had higher levels of intracellular ATP than untreated cells, suggesting potent inhibition of the V-H\textsuperscript{+}\textsubscript{-}ATPase activity (Fig. 5, C).

**PPI Effect on Drug Efflux**

We next investigated possible mechanisms responsible for the effect of PPI on tumor cell resistance to antitumor drugs. We used LSCM to monitor the intracellular distribution of fluorescently labeled (vinblastine-bodipy) or spontaneously fluorescent (doxorubicin) antitumor drugs in untreated or omeprazole-pretreated tumor cells in both the uptake and efflux phases. We first evaluated the effect of omeprazole on vinblastine-bodipy accumulation. Cells pretreated with omeprazole displayed a marked vinblastine accumulation in cytoplasmic vesicles 6 hours after the vinblastine-bodipy was added (Fig. 6, B), whereas cells that had not been pretreated displayed fluorescence diffused throughout the cytoplasm and in vesicle-like formations (Fig. 6, A). When cells were transferred to drug-free medium to allow drug efflux, cells that had been pretreated with omeprazole retained vinblastine-bodipy in vesicle-like structures (Fig. 6, D), whereas most vinblastine-bodipy was lost from cells that had not been pretreated (Fig. 6, C). The amount of retained antitumor drug was evaluated with LSCM. This analysis showed that, at the end of the uptake phase, the difference in mean pixel inten-
sities between cells that were pretreated with omeprazole (45.78) and cells that were not pretreated (48.84) was not statistically significant (3.06, 95% CI 3.46 to 9.58; P = .55). However, at the end of the efflux phase, the difference in mean pixel intensities of the retained fluorescent drug between tumor cells pretreated with omeprazole (52.71) and cells not treated with omeprazole (20.09) was statistically significant (32.62, 95% CI 23.08 to 42.16; P<.001). These results suggest that omeprazole pretreatment does not influence the uptake phase of the antitumor drug but strongly inhibits the elimination of antitumor drugs through the secretory pathway.

We next investigated the localization of a nuclear-targeted antitumor drug (i.e., doxorubicin) (Fig. 6, E–L). Whereas omeprazole pretreatment only slightly increased doxorubicin uptake in tumor cells as compared with untreated cells (Fig. 6, E and F), doxorubicin was completely retained in omeprazole-pretreated cells within vesicle-like structures, even after 24 hours in drug-free medium (Fig. 6, G and H). By 36 hours, doxorubicin was detectable in the nuclei of PPI-treated cells (Fig. 6, I and L). Thus, PPI treatment induced a massive retention of doxorubicin within vesicle-like structures, allowing this drug to get to the nucleus.

**Effects of Omeprazole on Sensitivity of Human Tumors to Antitumor Agents in SCID Mice Engrafted With Human Tumor Cells**

To assess the potential clinical relevance of the *in vitro* results, we performed *in vivo* experiments in a human/mouse...
model system represented by CB.17 SCID/SCID mice injected subcutaneously with human melanoma cells (MelM6). These mice have proved useful in assessing the in vivo efficacy of local and systemic antitumor treatments (39,43–45). Mice engrafted with human tumor cells were pretreated in groups of 10 with omeprazole administered orally (by gavage); 24 hours later, they were injected intraperitoneally with a single dose of cisplatin. Tumor growth was then followed three times per week. Figure 7, A, shows that tumors in mice pretreated with omeprazole and then treated with cisplatin grew more slowly (mean tumor weight 16 days after treatment /H11005/2788 mg) than tumors in mice that were treated with cisplatin but had not been pretreated with omeprazole (mean tumor weight 16 days after treatment /H11005/7373 mg) for a difference at 16 days after treatment of 4585 mg (95% CI /H11005/3711 to 5459) (Fig. 7, A). Again, consistent with the in vitro results, simultaneous treatment with cisplatin and omeprazole of melanoma-bearing SCID mice in vivo did not have any statistically significant effect on tumor growth (mean tumor weight 16 days after treatment /H11005/7749 mg; difference versus non-pretreated mice /H11005/376 mg, 95% CI /H11005/H11002/1062 to 1814) (Fig. 7, A) Similar results were obtained with esomeprazole (data not shown). Histologic examination of the human tumors after the animals were killed showed that, in the omeprazole–cisplatin-treated mice, the tumor mass was occupied by large necrotic areas that accounted for most of the tumor size (Fig. 7, B), suggesting that the cytotoxic effect was greater than that quantified by the in vivo tumor size measurements. Mice were followed for 1 month, and the tumors of mice pretreated with omeprazole slowly regained their growth but never reached the size of the tumors in non-pretreated mice or in mice treated with ineffective therapeutic regimens (i.e., normal saline; data not shown). Melanoma-bearing SCID mice treated with cisplatin after omeprazole pretreatment did not show any signs of systemic toxicity such as weight loss, diarrhea, or hair ruffling (data not shown).

**DISCUSSION**

The major hypothesis of this study was that the intrinsic resistance of tumors to cytotoxic drugs could be inhibited by
agents that affect the function of proton pumps that regulate cellular pH gradients. This hypothesis was based on previous findings that a marked alteration of the intra- to extracellular pH gradient occurs in malignant tumors, such that the extracellular compartments become highly acidic (4,5,46), and the acidic microenvironment and acidic vesicles (2,3,6,7,9) exert a major role in tumor resistance to cytotoxic drugs. We studied PPIs because these drugs inhibit the activity of V-H⁺/H₁₁₀⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻˓˓˓(182,530),(597,622)(182,638),(597,730)(182,742),(597,836)(182,847),(597,941)(182,956),(597,1050)(182,1062),(597,1156)(182,1169),(597,1263)(182,1275),(597,1369)
degranulation of microvesicles that are able to kill lymphocytes through apoptotic pathways (49). It is therefore conceivable that the traffic of acidic vacuoles and microvesicles may have a prominent role in tumor homeostasis.

Our findings add to the understanding of the resistance of human tumors to cytotoxic drugs in an additional way. We have shown here that human melanoma and adenocarcinoma cells express substantial levels of V-H+\textsuperscript{+}-ATPases and that PPI treatment of tumor cells, while not affecting the level of V-H+\textsuperscript{+}-ATPase protein, inhibits its activity and changes its subcellular localization. Thus, inhibition of V-H+\textsuperscript{+}-ATPase activity may represent an important mechanism of action of PPIs in their effect on tumor drug resistance. In fact, V-H+\textsuperscript{+}-ATPases carry out ATP-dependent proton transport from the cytoplasmic compartment to the opposite side of the membrane (the lumen of an intracellular vesicle or the extracellular space) (19), thus contributing to the creation and maintenance of the acidic microenvironment of tumors. Recent data suggest that V-H+\textsuperscript{+}-ATPase may be anchored, via its subunit C or ezrin adaptor protein, to the actin cytoskeleton (50,51). We have recently shown that ezrin, radixin, and moesin may exert an important role in multidrug resistance by mediating the P-glycoprotein linkage to actin (36); indeed, the function of a growing number of efflux pumps seems to depend on their connection to the cytoskeleton (52). Moreover, the ezrin and actin connection seems to have an important role in the trafficking of
acidic vesicles in human malignant tumors (38). It is therefore conceivable that connection to the cytoskeleton may have a key role for V-H+-ATPase activity as well. This, in turn, suggests that combined strategies aimed at inhibiting tumor acidity or at inhibiting connections between actin and the ion pumps may be extremely effective in depriving tumors of drug resistance strategies.

An important finding from our study is that PPI pretreatment predisposes melanoma and adenocarcinoma cells to the effects of different classes of antitumor agents, including cisplatin, whose mechanism of resistance is poorly known; 5-FU, whose of different classes of antitumor agents, including cisplatin, drug resistance strategies.

The multidrug-resistant human (14)

REFERENCES


NOTES

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