

Sequential treatment with celecoxib and bortezomib enhances the ER stress-mediated autophagy-associated cell death of colon cancer cells

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Abstract. Treatment with celecoxib and bortezomib as single chemotherapeutic agents reduces the viability and proliferation of colorectal cancer cells. The use of these agents in combination with other chemotherapeutic agents is usually associated with adverse effects. In the present study, a combination of celecoxib and bortezomib was investigated for potential synergistic effects in colon cancer cells. The sequential exposure to celecoxib with bortezomib synergistically induced apoptotic death in human colon cancer cells compared with groups treated with a single drug or other drug combinations. c-Jun N-terminal kinase/p38-mitogen-activated protein kinase-induced endoplasmic reticulum (ER) stress through serial exposure to celecoxib and bortezomib may have induced the intracellular Ca²⁺ release, leading to the generation of autophagosomes in p53-expressing HCT-116 cells. Targeted inhibition of p53 activity or ER stress or treatment with the Ca²⁺-chelating agent BAPTA-AM suppressed the ER stress-mediated Ca²⁺ release and apoptosis. Although p53^{-/-} HCT-116 cells were less sensitive to sequential treatment with celecoxib and bortezomib, co-localization of autophagosomes was detected in the absence of CCAAT-enhancer-binding protein homologous protein expression. Treatment of p53^{-/-} HCT-116 cells with BAPTA-AM did not inhibit apoptosis following serial treatment with celecoxib and bortezomib. These results suggest that the order of drug administration is important in treating cancer and that the sequential treatment with celecoxib and bortezomib enhances the ER stress-mediated

autophagy-associated cell death of colon cancer cells, regardless of p53 expression.

Introduction

Celecoxib, a cyclooxygenase-2 (COX-2)-specific inhibitor, reduces the risk of colorectal cancer, exhibiting anti-proliferative and chemoprophylactic effects on this cancer (1,2). Treatment with celecoxib induces the upregulation of endoplasmic reticulum (ER) chaperones, and promotes tumor cell death *in vitro* and *in vivo* through the ER stress response (3,4). This ER stress induces the nuclear phosphorylation and activation of p53, leading to ER stress-induced cell death in MCF-7 and HeLa cells (5). The co-treatment of p53-deficient colon cancer cells with zerumbone and celecoxib also induces ER stress and the transactivation of death receptor 5 (DR5) (6). The underlying molecular mechanisms by which celecoxib inhibits each cancer type have yet to be completely characterized. Therefore, it is necessary to investigate the downstream signaling pathways induced by treatment with celecoxib for clinical applications, and to examine whether it is more efficacious to treat cancer with a combination of drugs, rather than celecoxib alone.

The proteasome inhibitor bortezomib is a promising candidate for the treatment of hematological and solid cancer types (7). Bortezomib induces the unfolded protein response (UPR) to a limited extent, whereas the induction of binding immunoglobulin protein (BiP) and CCAAT/enhancer binding protein homologous protein (CHOP) by an ER stress-inducing agent is attenuated following exposure to this drug (8). Bortezomib activates downstream targets of p53, including p21, p53-upregulated modulator of apoptosis (PUMA) and Bcl-2-associated X (Bax); however, the induction of apoptosis by bortezomib is not affected by the deletion of p53 in colon cancer cells (9). Autophagy can protect cells from apoptotic stimuli, including growth factor deprivation and ER stress (10,11). Autophagy may also induce cell death, as the components of the autophagic and apoptotic machinery are interconnected and shared (12). The inhibition of cisplatin-induced autophagy by bortezomib has been shown to enhance the chemotherapeutic efficacy of cisplatin in ovarian

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cancer (13). The autophagy inhibitor 3-methyladenine (3-MA) enhances celecoxib-induced apoptosis in human colon cancer cells (14). On the basis of these reports, the effect and underlying mechanism of bortezomib or celecoxib on the induction of p53- and ER-stress-associated apoptosis in cancer cells remain controversial. Furthermore, the role of autophagy in cancer cells is complex and highly cell-type-dependent.

Despite the established connections between bortezomib or celecoxib treatment with ER stress or autophagy, it has yet to be determined whether combination treatment with celecoxib and bortezomib can improve the efficacy of treatment in colon cancer treatment by further promoting ER stress/autophagy-associated cell death. The present study focused on the development of novel chemotherapy combinations containing celecoxib and bortezomib for the treatment of colon cancer; it investigated whether the order of administration was critical for the induction of ER stress or stimulation of autophagy-associated cell death in colon cancer cells. In addition, the present study attempted to identify the role of p53 in the ER stress-mediated autophagy signaling pathway following the combination of celecoxib with bortezomib in HCT-116 and p53^{-/-} HCT-116 cells.

Materials and methods

Cell lines and reagents. The HCT-116, HCT-8 and HT-29 human colorectal cancer cell lines were purchased from the American Type Culture Collection (Manassas, VA, USA). p53^{-/-} HCT-116 cells were kindly provided by Professor Bert Vogelstein (Johns Hopkins University, Baltimore, MD, USA). All cells were maintained in RPMI-1640 medium (Corning Incorporated, Corning, NY, USA) supplemented with 10% FBS (HyClone; GE Healthcare, Chicago, IL, USA), streptomycin and glutamine at 37°C in 5% CO₂. Bortezomib was purchased from LC Laboratories (Woburn, MA, USA). Celecoxib was obtained from Selleck Chemicals (Houston, TX, USA). SP600125, a c-Jun N-terminal kinase (JNK) inhibitor, SB203580, a p38-mitogen-activated protein kinase (MAPK) inhibitor and salubrinal, an ER stress inhibitor, were purchased from Calbiochem (Merck KGaA, Darmstadt, Germany). BAPTA-AM and 3-MA were obtained from Sigma-Aldrich (Merck KGaA). Pifithrin (PFT)- α was purchased from Santa Cruz Biotechnology, Inc. (Dallas, TX, USA). To inhibit the activation of JNK or p38-mitogen-activated protein kinase, cells were pre-treated with SP600125 (20 μ M) or SB203580 (10 μ M) for 2 h at 37°C. To block endoplasmic reticulum stress, cells were pre-incubated with salubrinal (2 μ M) for 1 h at 37°C. To inhibit the autophagic signal, cells were pre-treated with 3-MA (10 mM) for 2 h at 37°C. To block the expression of p53, cells were pre-exposed to PTF- α (50 μ M) for 1 h at 37°C. To inhibit the effect of Ca²⁺, cells were pre-treated with BAPTA-AM (2 μ M) for 30 min at 37°C.

Cell viability assay with AlamarBlue and Cell Counting kit-8 (CCK-8). HCT-116, HCT-8 or HT-29 cells (2x10⁴ cells/well) were seeded in RPMI-1640 containing 10% FBS in 96-well plates. Cells were pre-treated with bortezomib (20 nM) for 6 h and then treated with celecoxib (20 μ M) for an additional 18 h. For comparison, cells were treated either with bortezomib (100 nM) or celecoxib (80 μ M) alone or co-treated

(20 nM bortezomib + 20 μ M celecoxib) for 24 h. Finally, cells pre-exposed to celecoxib (20 μ M) for 6 h were then treated with bortezomib (20 nM) for 18 h.

Cell viability was measured using an AlamarBlue assay (Serotec; Bio-Rad Laboratories, Inc., Hercules, CA, USA). AlamarBlue was added (10% by volume) to each well, and the relative fluorescence unit (RFU) values were determined 7 h later using a SpectraMax M2e Multi-Detection Microplate Reader (excitation, 530 nm; emission, 590 nm; Molecular Devices, LLC, Sunnyvale, CA, USA). Experiments were performed in triplicate, and RFU values were expressed as the mean $\pm$ standard deviation (SD) of three replicates.

Cell viability was also measured using a CCK-8 assay (Sigma-Aldrich; Merck KGaA) according to the manufacturer's protocol. Briefly, HCT-116, HCT-8 or HT-29 cells (2x10⁴ cells/well) in 96-well plates were pre-treated with celecoxib (20 μ M) for 6 h and then with bortezomib (20 nM) for an additional 18 h. For comparison, cells were treated with celecoxib (80 μ M). At 24 h, the cells were stained with 10 μ l CCK-8 dye in 90 μ l of culture medium for 2 h at 37°C. The absorbance was measured at 450 nm.

Analysis of apoptosis by flow cytometry. HCT-116 and p53^{-/-} HCT-116 cells (1x10⁵ cells/ml) were cultured in 6-well plates and pre-treated with 20 nM bortezomib for 6 h and then with 20 μ M celecoxib for an additional 18 h. For comparison, cells were treated either with 100 nM bortezomib or 80 μ M celecoxib alone or co-treated (20 nM bortezomib + 20 μ M celecoxib) for 24 h. Finally, cells were pre-exposed to 20 μ M celecoxib for 6 h and then treated with 20 nM bortezomib for 18 h.

The percentages of cells undergoing apoptosis were determined by flow cytometry using fluorescein isothiocyanate (FITC)-labeled annexin-V and 7-aminoactinomycin (7-AAD) (both BD Biosciences, San Diego, CA, USA). Cells were harvested, rinsed with PBS, and resuspended in 100 μ l of 1X annexin-V binding buffer (BD Biosciences). A total of 3 μ l annexin-V-FITC and 3 μ l 7-AAD were added, and cells were incubated at room temperature for 15 min in the dark, with gentle vortexing. The stained cells were analyzed using a FACSCalibur flow cytometer equipped with CellQuestpro software version 5.1 (BD Biosciences). Dot plot graphs were produced to quantify the percentage of viable cells (annexin-V⁻/7-AAD⁻), early-stage apoptotic cells (Annexin-V⁺/7-AAD⁻), late-stage apoptotic cells (annexin-V⁺/7-AAD⁺) and necrotic cells (Annexin-V⁻/7-AAD⁺).

Quantification of cytosolic and mitochondrial Ca²⁺ levels. Cytosolic Ca²⁺ levels were determined using the fluorescent dye Fluo3-AM (5 μ M; log mode in FITC setting; Molecular Probes; Thermo Fisher Scientific, Inc.). Mitochondrial Ca²⁺ levels were determined using the fluorescent dye Rhod2-AM (1 μ M; Molecular Probes; Thermo Fisher Scientific, Inc.). Cells were incubated with the fluorescent dyes for 20 min at 37°C, washed with calcium-free Dulbecco's PBS, and analyzed by flow cytometry. For certain experiments, cells were pretreated with BAPTA-AM (2 μ M) for 30 min.

Western blotting. Cells were washed in PBS and lysed in NP-40 buffer (Elpis Biotech, Inc., Daejeon, Korea) supplemented with a protease inhibitor cocktail (Sigma-Aldrich; Merck KGaA).

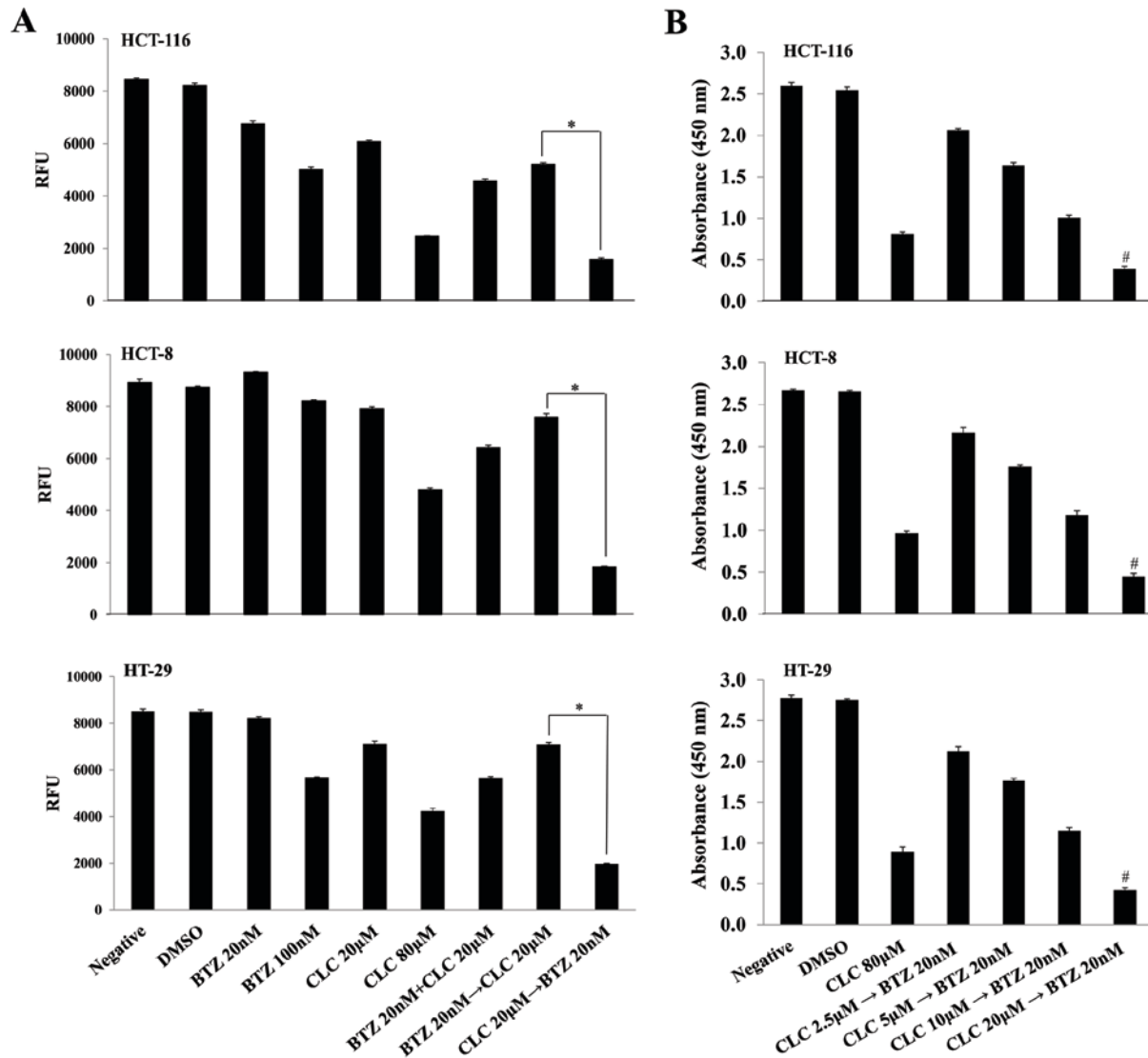


Figure 1. Effect of sequential treatment with CLC and BTZ on colorectal cancer cell survival. (A) Cells were treated with BTZ or CLC alone or in combination. Cell viability was measured using an AlamarBlue assay. The levels of fluorescence are presented. $n=3$. $^*P<0.01$. (B) Cells were sequentially treated with the indicated drug combinations for 24 h. For comparison, cells were treated with 80 μM CLC alone for 24 h. Cell viability was also measured using a Cell Counting kit-8 assay. The absorbance at 450 nm is presented. $n=3$. $^*P<0.01$ compared with CLC 2.5 μM followed by BTZ 20 nM. CLC, celecoxib; BTZ, bortezomib; negative, non-treated cell; DMSO, dimethyl sulfoxide control; RFU, relative fluorescence unit.

Protein phosphorylation states were preserved through the addition of phosphatase inhibitors (Cocktail II; Sigma-Aldrich; Merck KGaA) to the NP-40 buffer. Protein concentrations were determined using a BCA assay kit (Pierce; Thermo Fisher Scientific, Inc., Waltham, MA, USA). Proteins (10 $\mu\text{g}/\text{lane}$) were resolved through SDS-PAGE (12%) and transferred to a nitrocellulose membrane (EMD Millipore, Billerica, MA, USA). Membranes were blocked with 5% skimmed milk for 1 h at room temperature prior to western blot analysis. The following primary antibodies were used: Caspase-3 (cat. no. 9665; 1:1,000), caspase-9 (cat. no. 9502; 1:1,000), poly (ADP-ribose) polymerase (PARP; cat. no. 9542; 1:1,000), β -actin (cat. no. 4967; 1:1,000), Bcl-2 (cat. no. 2870; 1:1,000), Bax (cat. no. 2772; 1:1,000), Bcl-2 homologous antagonist/killer (Bak; cat. no. 6947; 1:1,000), Mcl-1 (cat. no. 4572; 1:1,000), Puma (cat. no. 4976; 1:1,000), survivin (cat. no. 2808; 1:1,000), p53 (cat. no. 2524; 1:1,000), phosphorylated (p)-extracellular-related

kinase (ERK)1/2 (Thr²⁰²/Tyr²⁰⁴; cat. no. 9101; 1:1,000), ERK1/2 (Tyr⁹²⁵; cat. no. 9102; 1:1,000), p-p38-MAPK (Thr¹⁸⁰/Tyr¹⁸²; cat. no. 9211; 1:1,000), p38-MAPK (cat. no. 9212; 1:1,000), p-JNK (Thr¹⁸³/Tyr¹⁸⁵; cat. no. 4671; 1:1,000), JNK (cat. no. 9258; 1:1,000), Beclin-1 (cat. no. 3495; 1:1,000), microtubule-associated protein 1A/1B-light chain 3 (LC3)-I/II (cat. no. 4108; 1:1,000). These antibodies were purchased from Cell Signaling Technology (Beverly, MA, USA), and CHOP (Santa Cruz Biotechnology, Inc.). Chemiluminescence was detected using an ECL kit (Advansta, Inc., Menlo Park, CA, USA) and a multiple Gel DOC system (Fujifilm, Tokyo, Japan). The membrane was probed with primary antibodies overnight at 4°C, followed by the application of the following secondary antibodies for 1 h at room temperature: Horseradish peroxidase (HRP)-conjugated goat anti-mouse IgG (cat. no. K0211589; 1:3,000) or HRP-conjugated goat anti-rabbit IgG (cat. no. K0211708; 1:3,000; both KOMABiotech, Seoul, Korea).

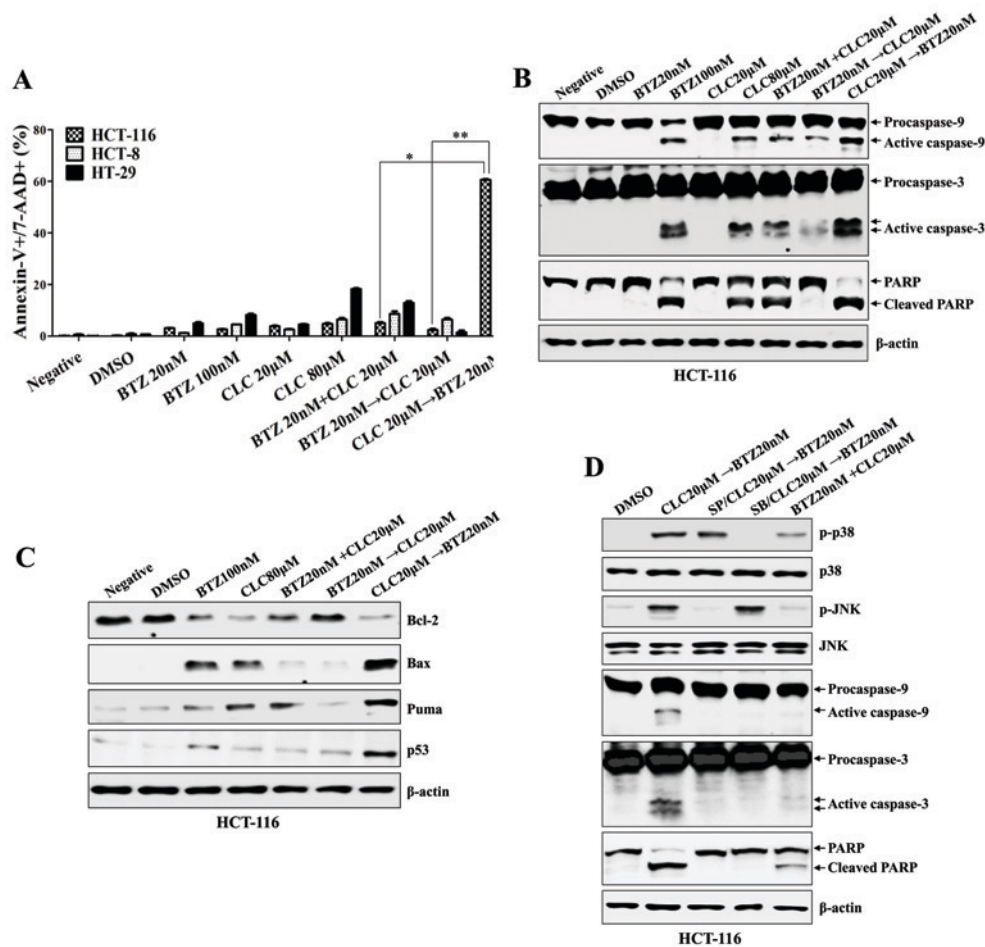


Figure 2. Sequential treatment with CLC and BTZ enhances apoptotic death of colon cancer cells. HCT-116 cells were treated with BTZ or CLC individually or in combination. (A) Annexin-V/7-AAD staining was used to estimate the rate of apoptosis. The number of late-stage apoptotic cells (annexin-V⁺/7-AAD⁺) was calculated by flow cytometry. * $P < 0.01$; ** $P < 0.005$. Cells from each indicated condition were harvested and blotted with the indicated antibodies, including (B) caspases (procaspase-9, active caspase-9, procaspase-3, active caspase-3 and PARP (full-length PARP, cleaved PARP), and (C) Bcl-2, Bax, PUMA and p53. β -actin served as an internal control. (D) To inhibit the activation of JNK or p38-mitogen-activated protein kinase, cells were pre-treated with SP600125 (20 μ M) or SB203580 (10 μ M) for 2 h. The expression of caspase-9, caspase-3 and cleaved PARP were detected by western blotting. The results are representative of three independent experiments. CLC, celecoxib; BTZ, bortezomib; 7-AAD, 7-aminoactinomycin D; PARP, poly (ADP-ribose) polymerase; negative, non-treated cell; DMSO, dimethyl sulfoxide control; p-, phosphorylated; JNK, c-Jun N-terminal kinase; PUMA, p53-upregulated modulator of apoptosis.

Confocal microscopy. Cells were permeabilized with permeabilization buffer (0.1% saponin in PBS), incubated with primary antibodies against LC3-I/II (cat. no. 4108; 1:200; Cell Signaling Technology) and lysosomal associated membrane protein 1 (LAMP1; cat. no. 4108; 1:100; Santa Cruz Biotechnology) for 30 min at 4°C, and incubated with FITC-conjugated goat anti-rabbit IgG (cat. no. F9887; 1:400) or PE-conjugated secondary goat anti-mouse IgG (cat. no. P9287; 1:400) (both Sigma-Aldrich; Merck KGaA) for 20 min at 4°C. Cells were mounted using Dako fluorescent mounting medium (Agilent Technologies, Inc., Santa Clara, CA, USA) and observed with a confocal laser scanning microscope at x400 magnification. Images were acquired using Confocal Microscopy Software version 3.0 (Carl Zeiss AG, Oberkochen, Germany).

Statistical analysis. Data are expressed as the mean $\pm$ SD. Statistical analysis was conducted using one-way analysis of variance (ANOVA) using SigmaPlot software (version 10.0; Systat Software, Inc., San Jose, CA, USA). Bonferroni post hoc analysis was performed following ANOVA for multiple

comparisons. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

Sequential treatment with celecoxib and bortezomib enhances the apoptotic signaling pathway through the up-regulation of p53, JNK, and p38-MAPK expression. Although sub-toxic doses of each drug in isolation (bortezomib 20 nM; celecoxib 20 μ M) slightly suppressed cell viability, the effect of the treatment with bortezomib (20 nM) and celecoxib (20 μ M) together on cell survival depended on the order in which the drugs were applied. When colon cancer cells were treated with celecoxib (CLC) then bortezomib (BTZ) (CLC 20 μ M then BTZ 20 nM), cell viability was significantly decreased compared with the group treated with bortezomib and then celecoxib (BTZ 20 nM then CLC 20 μ M) ($P < 0.01$, CLC 20 μ M then BTZ 20 nM vs. BTZ 20 nM then CLC 20 μ M) (Fig. 1A). The viability of colon cancer cells sequentially treated with celecoxib and bortezomib also decreased in a celecoxib dose-dependent manner. Notably, the effect of consecutive

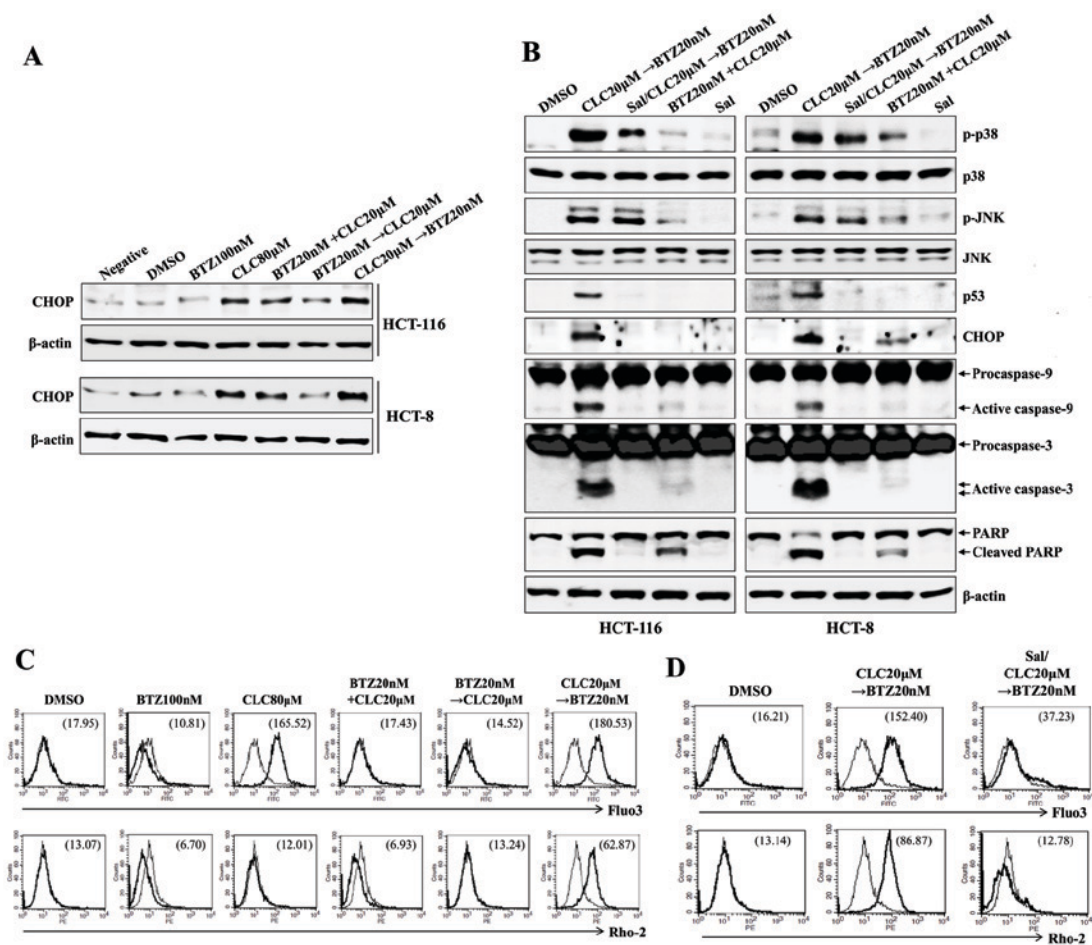


Figure 3. Sequential treatment with CLC and BTZ induces apoptosis in colon cancer cells through ER stress-mediated calcium translocation. HCT-116 cells were treated with BTZ or CLC alone or in combination. (A) Cells treated as indicated were blotted with antibodies against CHOP. (B) Cells treated as indicated were blotted with a range of antibodies. β -actin served as an internal control. To inhibit endoplasmic reticulum stress, cells were pre-incubated with Sal ($2 \mu\text{M}$) for 1 h. (C) Cytosolic and (D) mitochondrial Ca^{2+} levels in HCT-116 cells were determined using Fluo3-AM and Rhod2-AM, respectively. To block endoplasmic reticulum stress, cells were pre-incubated with Sal ($2 \mu\text{M}$) for 1 h. The results are representative of three independent experiments. CLC, celecoxib; BTZ, bortezomib; neg, non-treated cell; CHOP, CCAAT-enhancer-binding protein homologous protein; Sal, salubrinal; DMSO, dimethyl sulfoxide control; PARP, poly (ADP-ribose) polymerase.

treatment with celecoxib and bortezomib on cell viability differed significantly depending on the doses of celecoxib ($P < 0.01$, CLC $2.5 \mu\text{M}$ then BTZ 20 nM vs. CLC $20 \mu\text{M}$ then BTZ 20 nM) (Fig. 1B).

Sequential treatment with celecoxib followed by bortezomib synergistically increased the apoptotic cell death of colon cancer cells ($P < 0.01$, co-treatment with bortezomib and celecoxib vs. celecoxib $\rightarrow$ bortezomib; $P < 0.005$, bortezomib $\rightarrow$ celecoxib vs. celecoxib $\rightarrow$ bortezomib; Fig. 2A), which was associated with the activation and cleavage of caspase-9, caspase-3 and PARP (Fig. 2B). Furthermore, colon cancer cells treated with celecoxib followed by bortezomib exhibited decreased Bcl-2 expression, and increased Bax, p53 and PUMA expression, compared with any other single or combination treatment (Fig. 2C). The expression of p-JNK and p-p38-MAPK, which are required for the initiation of apoptosis, was induced in the HCT-116 cells sequentially treated with celecoxib followed by bortezomib; however, the expression of p-ERK was downregulated (Fig. 2D). With pre-exposure to SB203580 (a p38-MAPK inhibitor) or SP600125 (a JNK inhibitor), the activation of MAPKs and

caspases in the colon cancer cells by treatment with celecoxib followed by bortezomib was inhibited (Fig. 2D). These data suggest that sequential treatment with celecoxib and bortezomib enhances the synergistic induction of apoptotic death in human colon cancer cells in a manner that depends on the order in which the drugs are applied.

Sequential treatment with celecoxib followed by bortezomib induces apoptosis in colon cancer cells through ER stress-mediated calcium translocation. Whether treatment with celecoxib followed by bortezomib stimulated ER stress-mediated mitochondrial dysfunction, in which the translocation of calcium from the ER induces mitochondria-dependent apoptosis (15), was investigated next. The treatment of cells with celecoxib followed by bortezomib resulted in the increased induction of CHOP, compared with cells co-treated with alternative drug combinations, or with bortezomib followed by celecoxib (Fig. 3A). Pre-exposure to salubrinal (an ER stress inhibitor) of the colon cancer cells inhibited the expression of p53 and CHOP, and prevented the cleavage of caspase-9, caspase-3 and PARP subsequent to

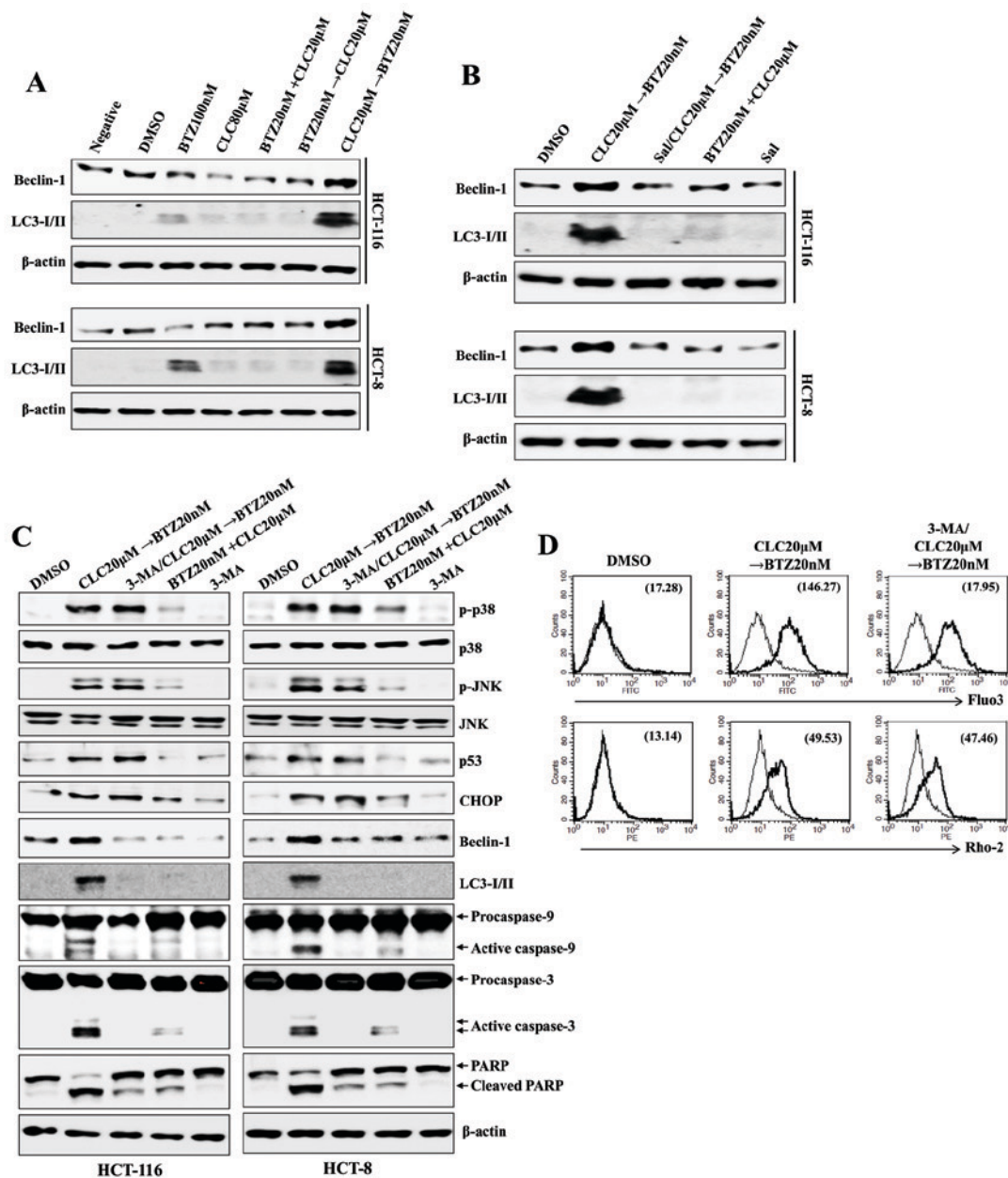


Figure 4. Sequential treatment with CLC and BTZ provokes ER stress-mediated autophagy-associated cell death in colon cancer cells. HCT-116 cells were treated with BTZ or CLC alone or in combination. (A-C) Total lysates of cells under each indicated condition were harvested and blotted with the indicated antibodies. β -actin served as an internal control. (B) To block ER stress, cells were pre-incubated with salubrinal ($2 \mu\text{M}$) for 1 h. (C) To inhibit the autophagic signal, cells were pre-treated with 3-MA (10 mM) for 2 h. (D) Cytosolic and mitochondrial Ca^{2+} levels in HCT-116 cells were determined using Fluo3-AM and Rhod2-AM, respectively. The results are representative of three independent experiments. CLC, celecoxib; BTZ, bortezomib; ER, endoplasmic reticulum; 3-MA, 3-methyladenine; negative, non-treated cell; LC3-I/II, microtubule-associated protein 1A/1B-light chain 3; DMSO, dimethyl sulfoxide control; PARP, poly (ADP-ribose) polymerase; p-, phosphorylated; JNK, c-Jun N-terminal kinase; CHOP, CCAAT-enhancer-binding protein homologous protein.

treatment with celecoxib followed by bortezomib (Fig. 3B). Treatment with celecoxib followed by bortezomib, and high doses of celecoxib alone, increased the cytosolic and mitochondrial Ca^{2+} levels in colon cancer cells; pre-treatment with salubrinal inhibited this effect (Fig. 3C and D). The data collectively suggest that ER stress-mediated apoptosis of colon cancer cells treated with celecoxib followed by bortezomib is associated, at least partially, with increases in cytosolic and mitochondrial Ca^{2+} .

Treatment with celecoxib followed by bortezomib induces ER stress-mediated autophagy-associated cell death in colon

cancer cells. Whether treatment with celecoxib followed by bortezomib was also connected to autophagy-associated cell death was examined. Treatment with celecoxib followed by bortezomib induced the expression of Beclin-1 and autophagosome-associated LC3-I/II proteins, which are associated with autophagy-associated cell death (Fig. 4A); however, pre-treatment with salubrinal decreased the expression of Beclin-1 and LC3-I/II by treatment with celecoxib followed by bortezomib (Fig. 4B). Although pre-exposure to the autophagy inhibitor 3-MA inhibited the autophagy-associated cell death of colon cancer cells subsequent to treatment with celecoxib followed by bortezomib, the expression levels of CHOP, p53 and p-MAPKs

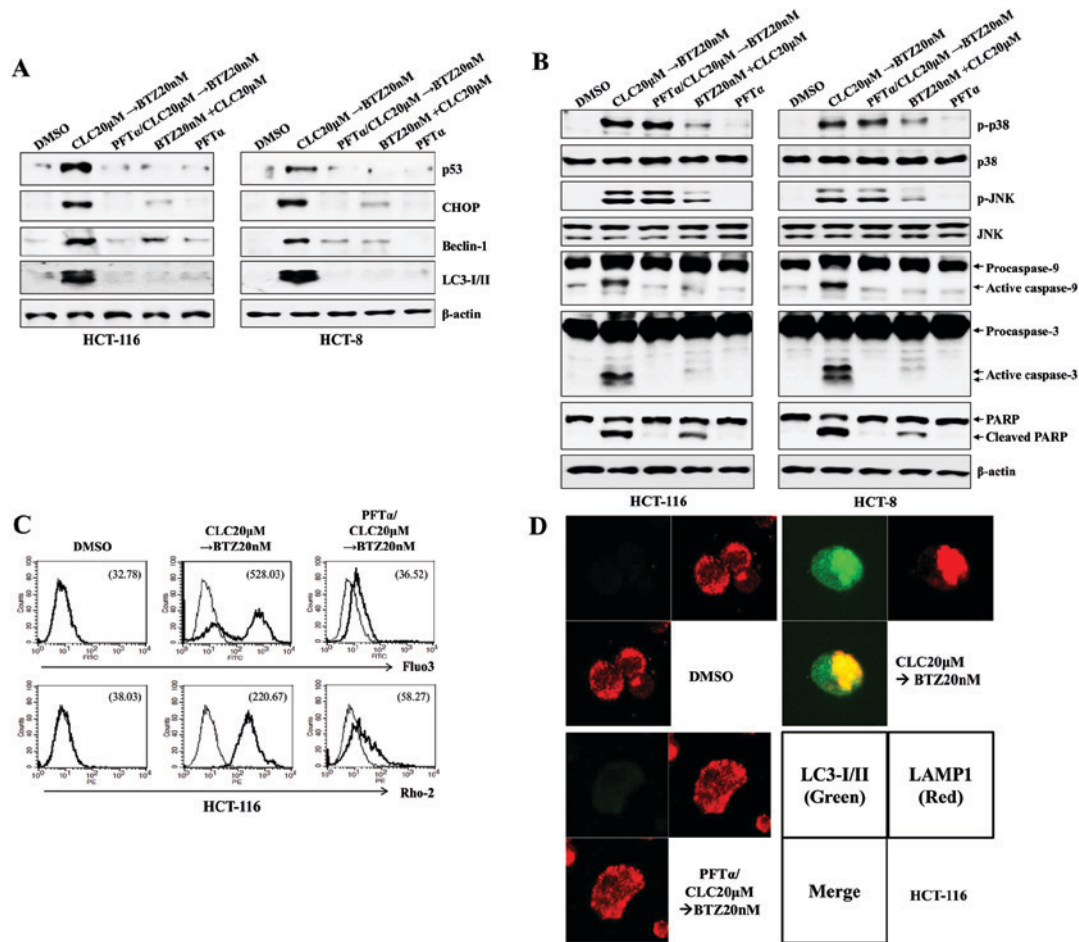


Figure 5. The expression of p53 is dispensable for the endoplasmic reticulum stress-mediated autophagy-associated cell death of colon cancer cells treated sequentially with CLC and BTZ. HCT-116 or HCT-8 cells were pre-incubated with PFT- α (50 μ M) for 1 h. (A and B) Cells in each indicated condition were harvested and blotted with the indicated antibodies. β -actin served as the internal control. (C) Cytosolic and mitochondrial Ca^{2+} levels in HCT-116 cells were determined using Fluo3-AM and Rhod2-AM, respectively. (D) Subcellular distribution of LC3-I/II and LAMP1 in HCT-116 cells. Cells were observed under a confocal microscope (magnification, $\times 400$). Green fluorescence indicates LC3-I/II, and red fluorescence indicates LAMP1. The results are representative of three independent experiments. CLC, celecoxib; BTZ, bortezomib; PFT- α , pifithrin- α ; LC3-I/II, microtubule-associated protein 1A/1B-light chain 3; LAMP1, lysosomal-associated membrane protein 1; DMSO, dimethyl sulfoxide control; CHOP, CCAAT-enhancer-binding protein homologous protein; p-, phosphorylated; JNK, c-Jun N-terminal kinase; PARP, poly (ADP-ribose) polymerase.

remained upregulated (Fig. 4C). Furthermore, treatment of the cells with 3-MA had no effect on the levels of cytosolic or mitochondrial Ca^{2+} subsequent to treatment with celecoxib followed by bortezomib (Fig. 4D). These data suggest that ER stress-associated apoptotic signaling in colon cancer cells leads to the induction of autophagy-associated cell death subsequent to treatment with celecoxib followed by bortezomib.

p53 expression regulates ER stress-mediated autophagy-associated cell death following treatment with celecoxib followed by bortezomib. ER stress is associated with the induction of p53 expression (16), and the level of cytoplasmic p53 serves a critical role in regulating ER stress-associated Ca^{2+} homeostasis (17). Treatment with celecoxib followed by bortezomib induced the expression of p53 and increased cytosolic and mitochondrial Ca^{2+} levels. Next, it was investigated whether the increased expression of p53 by treatment with celecoxib followed by bortezomib was responsible for the ER stress/autophagy-associated cell death due to the induction of apoptosis. Pre-treatment with PFT- α , a selective p53 inhibitor, not only prevented the increases

in CHOP, Beclin-1 and LC3-I/II expression (Fig. 5A), but also reduced the cleavage of caspase-9, caspase-3 and PARP in colon cancer cells treated with celecoxib followed by bortezomib (Fig. 5B). When PFT- α was applied in cells treated with celecoxib followed by bortezomib, intracellular and mitochondrial levels of Ca^{2+} were reduced (Fig. 5C), and the co-localization of LAMP1 and LC3-I/II was inhibited, compared with the cells treated with celecoxib followed by bortezomib (Fig. 5D). These data suggest that the expression of p53 in colon cancer cells serves a role in inducing ER stress/autophagy-associated cell death subsequent to treatment with celecoxib followed by bortezomib.

Treatment with celecoxib followed by bortezomib induces the autophagy-associated cell death of colon cancer cells even in the absence of p53 expression. Whether the treatment with celecoxib followed by bortezomib affects ER stress-mediated autophagy-associated cell death in p53 $^{-/-}$ colon cancer cells was then investigated. The treatment of p53 $^{-/-}$ HCT-116 cells with celecoxib followed by bortezomib induced the phosphorylation

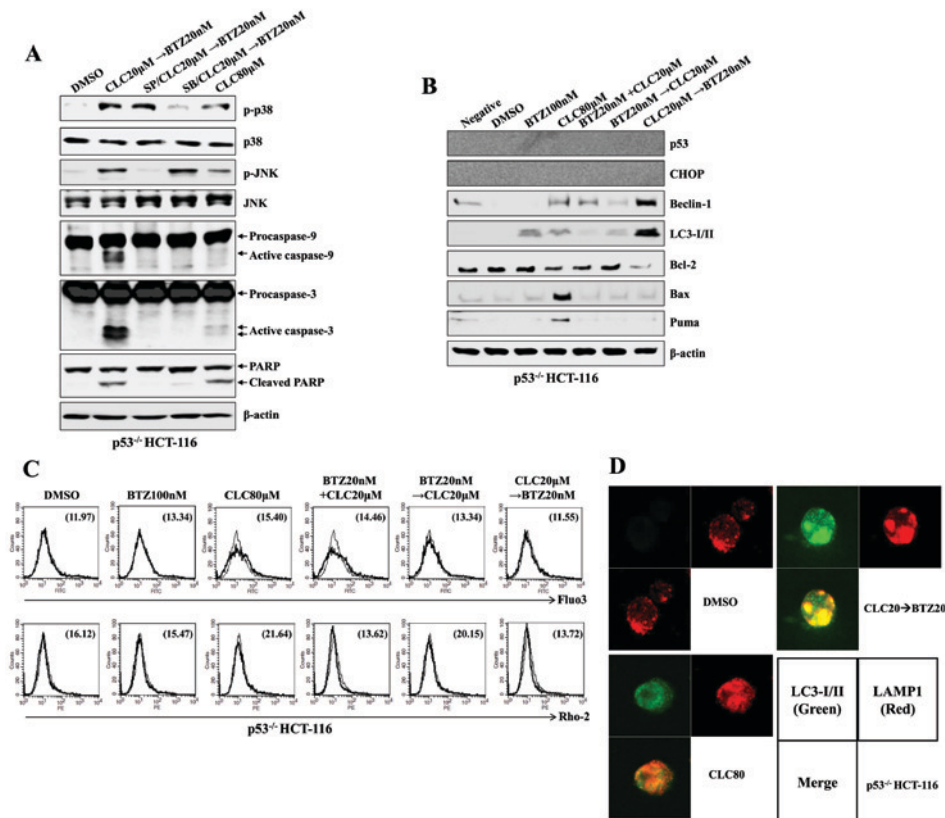


Figure 6. Sequential treatment with CLC and BTZ induces the autophagy-associated cell death of p53^{-/-} colon cancer cells. p53^{-/-} HCT-116 cells were treated with BTZ or CLC alone or in combination. (A and B) Total cell lysates of each condition were harvested and blotted with the indicated antibodies. β-actin served as an internal control. (C) Cytosolic and mitochondrial Ca²⁺ levels in HCT-116 cells were determined using Fluo3-AM and Rhod2-AM, respectively. (D) Subcellular distribution of LC3-I/II and LAMP1 in p53^{-/-} HCT-116 cells. Cells were observed under a confocal microscope (magnification, x400). Green fluorescence indicates LC3-I/II, and red fluorescence indicates LAMP1. The results are representative of three independent experiments. CLC, celecoxib; BTZ, bortezomib; LC3-I/II, microtubule-associated protein 1A/1B-light chain 3; LAMP1, lysosomal-associated membrane protein 1; DMSO, dimethyl sulfoxide control; p-, phosphorylated; JNK, c-Jun N-terminal kinase; PARP, poly (ADP-ribose) polymerase; negative, non-treated cells; CHOP, CCAAT-enhancer-binding protein homologous protein; Bax; Bcl-associated X; PUMA, p53-upregulated modulator of apoptosis.

of p38-MAPK and JNK and the expression of activated caspase-9 and caspase-3 (Fig. 6A). Furthermore, the induction of Beclin-1 and LC-3I/II expression was observed in p53^{-/-} HCT-116 cells treated with celecoxib followed by bortezomib, although the induction of CHOP was not detected (Fig. 6B). Notably, changes to the levels of Bcl-2, Bax and PUMA expression, which are induced by p53, were not observed (Fig. 6B). Although no Ca²⁺ increase was detected in the cytoplasm or mitochondria of p53^{-/-} HCT-116 cells (Fig. 6C), fusion of the autophagosomes and the lysosomes was detected subsequent to treatment with celecoxib followed by bortezomib (Fig. 6D). These data suggest that treatment with celecoxib followed by bortezomib may cause ER stress-mediated autophagy-associated cell death independent of p53 expression.

The expression of p53 regulates intracellular calcium release from the ER to enhance autophagy-associated cell death. As treatment with celecoxib followed by bortezomib increased the autophagy-associated cell death of colon cancer cells regardless of p53 expression, the role of, and association between, intracellular Ca²⁺ and p53 expression in the ER stress-induced autophagy-associated cell death of colon cancer cells was investigated. The p53-expressing HCT-116 colon cancer cells were more susceptible to apoptotic cell

death than p53^{-/-} HCT-116 colon cancer cells following treatment with celecoxib followed by bortezomib ($P < 0.05$, p53-expressing HCT-116 compared with p53-null HCT-116; Fig. 7A). In addition, treatment with BAPTA-AM, a calcium chelator, failed to inhibit apoptosis or the cleavage of caspase-3 in p53^{-/-} HCT-116 cells subsequent to treatment with celecoxib followed by bortezomib ($\#P < 0.05$, celecoxib→bortezomib in p53-expressing HCT-116 vs. pretreatment with BAPTA-AM, and then celecoxib→bortezomib in p53-expressing HCT-116; $\#\#P < 0.001$, pretreatment with BAPTA-AM, and then celecoxib→bortezomib in p53-expressing HCT-116 vs. pretreatment with BAPTA-AM, and then celecoxib→bortezomib in p53-null HCT-116; Fig. 7B). Exposure to BAPTA-AM not only reduced the expression of CHOP, Beclin-1 and LC3-I/II, but also prevented the cleavage of caspase-3 in p53-expressing HCT-116 colon cancer cells treated with celecoxib followed by bortezomib (Fig. 7C); however, treatment with BAPTA-AM failed to inhibit the activation of Beclin-1 and LC3-I/II and prevent the cleavage of caspase-3 in p53^{-/-} HCT-116 cells subsequent to treatment with celecoxib followed by bortezomib (Fig. 7C and D). These data suggest that p53 expression is critical for promoting autophagy-associated cell death through the regulation of intracellular Ca²⁺ released from the ER.

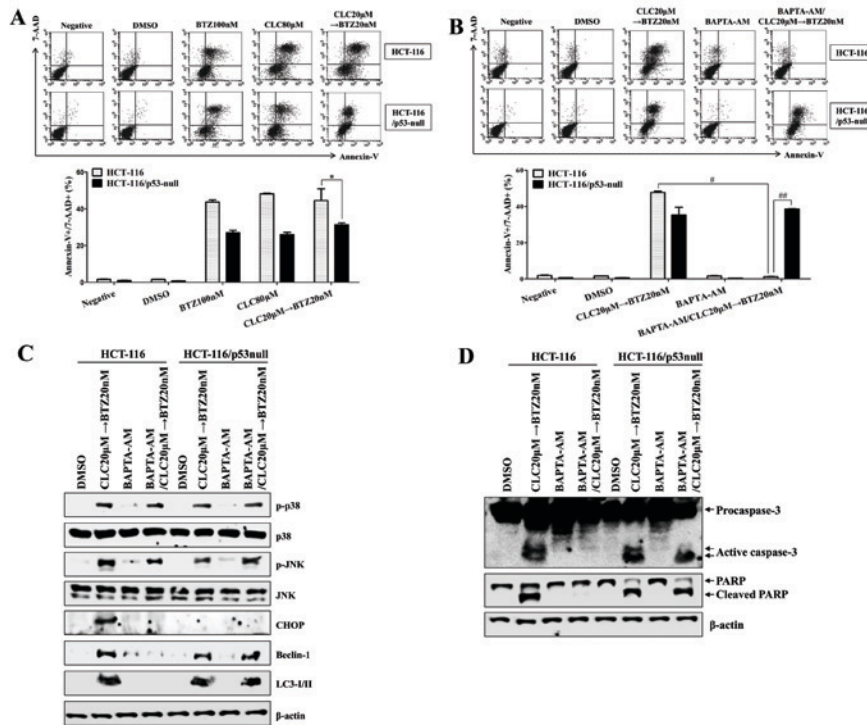


Figure 7. The expression of p53 is critical for endoplasmic reticulum stress-mediated autophagy-associated cell death through the regulation of intracellular calcium. HCT-116 or p53^{-/-} HCT-116 cells were treated with BTZ or CLC alone or in combination. (A and B) The percentage of apoptotic cells was estimated by annexin-V/7-AAD staining. Data are representative of three independent experiments. The number of late-stage apoptotic cells (annexin-V⁺/7-AAD⁺) was calculated by flow cytometry. *P<0.05; #P<0.05 and ##P<0.001. (B-D) To block the effect of Ca²⁺, HCT-116 or p53^{-/-} HCT-116 cells were pre-incubated with BAPTA-AM (2 μM) for 30 min. (C and D) Total cell lysates of each condition were harvested and immunoblotted with the indicated antibodies. β-actin served as an internal control. The results are representative of three independent experiments. BTZ, bortezomib; CLC, celecoxib; 7-AAD, 7-aminoactinomycin D; negative, non-treated cells; DMSO, dimethyl sulfoxide control; p-, phosphorylated; JNK, c-Jun N-terminal kinase; CHOP, CCAAT-enhancer-binding protein homologous protein; LC3-I/II, microtubule-associated protein 1A/1B-light chain 3; PARP, poly (ADP-ribose) polymerase.

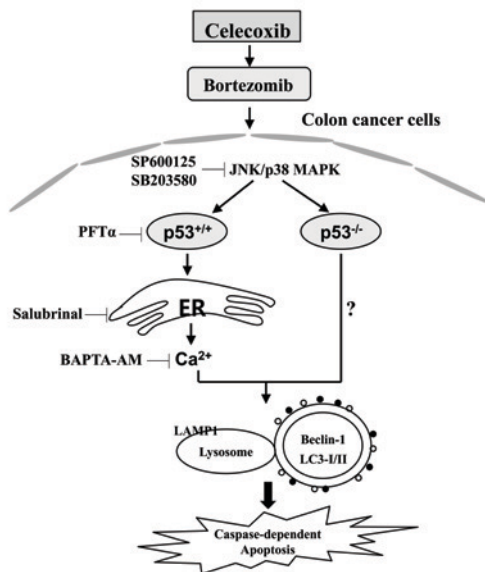


Figure 8. Schematic diagram of the intracellular signaling pathway following sequential treatment with celecoxib and bortezomib in human colorectal cancer cells. The expression of p53 is required for ER stress-mediated Ca²⁺ translocation to promote autophagy-associated cell death following the sequential treatment with celecoxib and bortezomib. The consecutive treatment of colon cancer cells with celecoxib and bortezomib also enhances autophagy-associated cell death in p53^{-/-} colon cancer cells. ER, endoplasmic reticulum; JNK, c-Jun N-terminal kinase; p38 MAPK, p38 mitogen-activated protein kinase; LAMP1, lysosomal-associated membrane protein 1; LC3-I/II, microtubule-associated protein 1A/1B-light chain 3.

Discussion

Bortezomib and celecoxib or dimethyl-celecoxib (DMC) as a novel drug combination enhance apoptotic cell death through the upregulation of ER stress in addition to the activation of JNK in glioblastoma (18). Pre-treatment with bortezomib enhances the rate of apoptosis induced by treatment with celecoxib through the induction of cell cycle arrest and ER stress (19). In the present study, the anticancer activity was enhanced by the treatment with celecoxib followed by bortezomib; this was associated with autophagy-associated cell death, regardless of p53 expression. Furthermore, the expression of p53 was required for ER stress-mediated Ca²⁺ translocation to promote autophagy-associated cell death following treatment with celecoxib followed by bortezomib (Fig. 8).

The anticancer activity of celecoxib has been demonstrated in various animal tumor models and may potentially be useful for the treatment of colorectal cancer and other tumor types (20,21). Although celecoxib toxicity in tumor cells occurs by toxic precipitation of increased doses of the drug (22), celecoxib also causes ER stress-induced apoptosis in a COX-2-independent manner (14,23). However, the long-term use of high-dose celecoxib is associated with severe gastrointestinal and cardiovascular adverse effects (24). Additionally, the combination treatment of colorectal cancer cells with celecoxib and other chemotherapeutic agents induces autophagy-mediated drug resistance (14). Although bortezomib exhibits a unique

mechanism for anticancer activity, it also blocks the anticancer effect of the ER stress-inducing agents thapsigargin and tunicamycin (8). The induction of autophagy and the ER stress response was demonstrated to protect MCF7 breast cancer cells from bortezomib-induced cell death (25). On the basis of these results, the limited effectiveness of celecoxib and bortezomib against cancer cells has led to a search for therapeutic combination regimens with other agents.

Persistent or intense ER stress induces different effects on normal ER function and induces cell death or apoptosis (26). The disturbance of autophagy also renders cells vulnerable to ER stress, and autophagy activation has been linked to cell death through the regulation of ER stress (11,27). In the present study, the combination of celecoxib followed by bortezomib was the most efficient regimen for the induction of ER stress-mediated autophagy-associated cell death, compared with the groups treated with single drugs alone or with other co-treatment regimens. Although the inhibition of autophagy by 3-MA suppressed the generation of autophagolysosomes, leading to autophagy-associated cell death, 3-MA had no effect on the inhibition of CHOP and p53 expression in colon cancer cells sequentially treated with celecoxib and bortezomib. Exposure to celecoxib followed by bortezomib induced the generation of autophagolysosomes in p53^{-/-} HCT-116 colon cancer cells. These results suggest that ER stress is not a critical regulation step for stimulating autophagy-associated cell death in colon cancer cells following the serial treatment with celecoxib and bortezomib.

Intracellular Ca²⁺ released due to ER stress as a result of various stimuli is absorbed by mitochondria. Once calcium is taken into the mitochondria, it causes the activation of the apoptotic pathway (28). Giorgi *et al* (17) demonstrated that wild-type cytoplasmic p53 at the ER binds to Ca²⁺ pumps to modulate Ca²⁺ homeostasis and mitochondrial function. In the present study, sequential treatment with celecoxib and bortezomib also promoted ER stress-mediated apoptosis through Ca²⁺ translocation and changes in the intracellular and mitochondrial Ca²⁺ concentration in p53-expressing HCT-116 cells. Calcium chelation with BAPTA-AM decreased the expression levels of CHOP and inhibited autophagy-associated cell death in wild-type HCT-116 colon cancer cells. Although p53-expressing HCT-116 cells were more sensitive to autophagy-associated cell death than p53^{-/-} HCT-116 cells, the treatment of p53^{-/-} HCT-116 cells with celecoxib followed by bortezomib also induced apoptosis and the expression of Beclin-1 and LC3-I/II, but not CHOP. These results suggest that the sequential drug combination can cause apoptosis in a p53-independent manner, although for full effectiveness, p53 expression is required to induce the release of Ca²⁺ from the ER.

The overexpression of Bcl-2 inhibits Ca²⁺ transfer from the ER to the mitochondria and prevents apoptosis (29). Bcl-2 proteins have been shown to inhibit autophagy by disrupting Bcl-2/Bcl-X_L-Beclin-1 complexes (30); however, it remains unclear how Bcl-2-family proteins are associated with autophagy-associated cell death in the absence of p53 expression. In the present study, only Bcl-2 was decreased in p53^{-/-} HCT-116 colon cells sequentially treated with celecoxib and bortezomib. On the basis of these results, further studies

are required to investigate how the combined treatment of celecoxib followed by bortezomib induces autophagy-mediated cell death in the absence of p53 expression.

Taken together, the results of the present study provide information about the optimal order for anticancer drug administration and the role of calcium and p53 on ER stress-mediated autophagy-associated cell death in colon cancer cells subsequent to treatment with celecoxib followed by bortezomib.

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