Curcumin Enhances Docetaxel-Induced Apoptosis of 8505C Anaplastic Thyroid Carcinoma Cells

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Background: Anaplastic thyroid cancer (ATC) is one of the most aggressive malignancies in humans, and its progression is poorly controlled by existing therapeutic methods. Curcumin has been shown to suppress inflammation and angiogenesis. In this study, we evaluated whether curcumin could augment docetaxel-induced apoptosis of ATC cells. We also analyzed changes in nuclear factor κB (NF-κB) and cyclooxygenase-2 (COX-2) expression levels to delineate possible mechanisms of their combined action.

Methods: ATC cells were cultured and treated with curcumin and docetaxel alone or in combination. The effects on cell viability were determined by MTS assay. Apoptosis was assessed by annexin V staining and confirmed by flow cytometric analysis. Caspase, COX-2, NF-κB levels were assayed by Western blotting.

Results: Curcumin combined with docetaxel led to lower cell viability than treatment with docetaxel or curcumin alone. Annexin V staining followed by flow cytometric analysis demonstrated that curcumin treatment enhanced the docetaxel-induced apoptosis of ATC cells. Additionally, curcumin inhibited docetaxel-induced p65 activation and COX-2 expression.

Conclusion: We conclude that curcumin may enhance docetaxel’s antitumor activity in ATC cells by interfering with NF-κB and COX-2. Our results suggest that curcumin may emerge as an attractive therapeutic candidate to enhance the antitumor activity of taxanes in ATC treatment.

Keywords: Thyroid cancer, anaplastic; Apoptosis; Curcumin; Cyclooxygenase 2; Docetaxel; NF-kappa B

INTRODUCTION

Anaplastic thyroid cancer (ATC) is one of the most aggressive malignancies in humans. Multidisciplinary treatments, including radiotherapy and chemotherapy, poorly control the progression of this disease, and the mean survival is less than 1 year after initial diagnosis [1]. Therefore, discovery of new and improved therapeutic strategies are critical for improving the quality of life and survival rates of patients with ATC.

Curcumin is a polyphenolic natural product found in turmeric, a commonly used food coloring and flavoring agent. The active component of turmeric, curcumin, has been shown to have antiviral, antibacterial, antioxidant, anti-inflammatory, antiproliferative, and antiangiogenic activities [2,3]. Studies
conducted in both animals and humans have suggested that curcumin is a promising candidate for the treatment of inflammation and cancer [4–7].

Docetaxel is an anticancer microtubule-stabilizing agent that induces apoptosis by suppressing the microtubule dynamics of the mitotic apparatus [8,9]. Docetaxel arrests cells in the G2/M phase, leading to increased susceptibility of cancer cells to radiotherapies and chemotherapies [10–12]. It is severely dose-limited because of its adverse side effects, including neutropenia, diarrhea, nausea, and fatigue [13]. Combination therapies to reduce its toxicity at higher doses and/or to increase its efficacy at lower doses are highly desirable, and could have a significant impact on therapeutic outcome. Studies have shown that docetaxel alone was only modestly effective against ATC in clinical trials [14]. However, available evidence suggests that combined therapy with radiation and/or other pharmacological agents may be more effective [15–17].

In this study, we attempted to increase the sensitivity of ATC cells to docetaxel by simultaneous treatment with curcumin and analyzed the apoptotic signaling pathways in treated cells. Here, we report for the first time that curcumin augments the growth-inhibitory and proapoptotic effects of docetaxel, possibly by interfering with nuclear factor κ light-chain enhancer of activated B cells (NF-κB) and cyclooxygenase-2 (COX-2) signaling pathways. Our results suggest that a curcumin-docetaxel combination could be a useful therapeutic agent for the treatment of ATC.

METHODS

Materials

Human anaplastic thyroid carcinoma cell lines 8505C and CAL62 were a gift from Dr. Shong (Asan Medical Center, University of Ulsan College of Medicine, Seoul, Korea). The ATC cell line FRO (wild type) was provided by J.A. Fagin (University of Cincinnati College of Medicine, Cincinnati, OH, USA), and the cell line KTC-2 was obtained from Kawasaki Medical School (Okayama, Japan) [18].

Docetaxel and curcumin were purchased from Sigma (St. Louis, MO, USA). They were dissolved in dimethyl sulfoxide (DMSO, Sigma) at appropriate concentrations prior to dilution in culture medium such that the final concentration of DMSO would not exceed 0.1%.

Antiactin, antirabbit immunoglobulin G (IgG), and antimouse IgG horseradish peroxidase-conjugated secondary antibody were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Antibodies against NF-κB, cleaved caspase-3 and caspase-9 were obtained from Cell Signaling Technology (Beverly, MA, USA). A polyclonal antibody against COX-2 was procured from Abcam (Cambridge, MA, USA).

Cell culture

8505C, FRO, and KTC2 cells were grown in RPMI 1640 medium, and CAL62 cells were grown in Dulbecco’s modified Eagle’s medium and both media types were supplemented with 10% fetal bovine serum and 1% penicillin/streptomycin (all reagents from Invitrogen Life Technologies, Paisley, UK). Cells were grown in a 5% CO₂ humidified incubator maintained at 37°C.

Cell proliferation assay

Cell viability was measured by a CellTiter 96 AQueous One Solution Cell Proliferation Assay according to the manufacturer’s instructions (Promega, Madison, WI, USA). Aliquots of 4 × 10⁴ cells were plated onto each well of a 96-well microplate (Falcon, Franklin Lakes, NJ, USA) in 100 µL complete growth medium and incubated with 1, 2, 5, 10, or 20 nM of docetaxel in the presence or absence of 1, 5, 10, 25, or 50 µM curcumin or 0.1% DMSO (control) for 24 to 48 hours. Following this, 20 µL of 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium inner salt was added to each well and incubated for an additional 3 hours. The relative cell proliferation rates were determined indirectly by quantifying the color developed at 490 nm using a 96-well plate reader. Each experiment was performed in triplicate, and repeated a minimum of three times with three independent cultures.

Annexin V staining

Annexin V staining was performed using the Annexin-V-FITC apoptosis detection kit (BD Pharmingen, San Diego, CA, USA) according to the manufacturer’s protocols. Briefly, cells were harvested, resuspended in 100 µL binding buffer, and stained with 5 µL Annexin-V-FITC staining solution in the dark at room temperature (RT) for 15 minutes. The samples were analyzed by flow cytometry (Becton Dickenson, Franklin Lakes, NJ, USA) using CellQuest software.

Preparation of cell extracts

Cells were grown in 150-mm dishes in medium containing 10% FBS at 37°C. After 24 hours, cells were treated with docetaxel (5 nM) and/or curcumin (10 µM) or vehicle for 24 hours. Follow-
ing this, cells were washed twice with ice-cold phosphate buffered saline (PBS), collected in 1 mL PBS and centrifuged at 4°C for 5 minutes at 1,200 rpm. The cell pellet was then resuspended in 50 µL of M-PER mammalian protein extraction reagent (Pierce, Rockford, IL, USA) containing a cocktail of protease inhibitors (Pierce). After incubating for 20 minutes on ice, the lysate was centrifuged for 10 minutes at 14,000 rpm. The supernatant was stored at -80°C until use. Protein concentrations were determined using a BCA protein assay kit (Pierce) according to manufacturer’s protocol. Nuclear extracts was prepared from cells using the NE-PER nuclear and cytoplasmic extraction kit (Pierce) following the manufacturer’s instructions.

Western blotting
Forty micrograms of protein was loaded onto 10% to 15% polyacrylamide gels and separated by sodium dodecylsulfate-polyacrylamide gel electrophoresis (SDS-PAGE). Separated proteins were transferred onto a nitrocellulose membrane (Millipore Corp., Bedford, MA, USA) and blocked with Tris-buffered saline/0.1% Tween 20 (TBST) containing 5% nonfat dry milk for 1 hour at RT. After washing four times with TBST, membranes were incubated with desired concentrations of primary antibodies in blocking solution at 4°C overnight. After washing four times with TBST, the blots were incubated with horseradish peroxidase-conjugated species-specific secondary antibodies (Santa Cruz Biotechnology) in TBST for 1 hour at RT. The blots were washed again four times with TBST, and the complexes were visualized using an Immun-Star Western Chemiluminescent kit (Bio-Rad, Hercules, CA, USA) in a Bio-Rad Molecular Imager ChemiDoc XRS imaging system (Bio-Rad). Western blotting was performed using lysates from at least three independent experiments. Intensities of the bands developed after the application of HRP substrate were quantified using the Bio-Rad Molecular Imager ChemiDoc XRS imaging system (Bio-Rad). Relative expression levels of various proteins were determined using a BCA protein assay kit (Pierce) according to manufacturer’s protocol. Nuclear extracts was prepared from cells using the NE-PER nuclear and cytoplasmic extraction kit (Pierce) following the manufacturer’s instructions.

**Fig. 1.** Effects of docetaxel and/or curcumin treatment on the proliferation of 8505C, CAL62, FRO, and KTC2 cells. (A) 8505C cells were grown on 96-well plates and treated with varying doses of docetaxel (DOC) or curcumin (CUR) for 24 hours. (B) 8505C cells were grown on 96-well plates and treated with a combination of docetaxel (5 nM) and curcumin (10 µM) for 24 hours. (C) CAL62, FRO, and KTC2 cells were treated with docetaxel (5 nM) and curcumin (10 µM). Bars represent mean ± SD. CTL, cell only; NS, no significant. *P < 0.001; †P < 0.05.
proteins were estimated by densitometric analysis using actin expression levels as a standard.

**Statistical analysis**
All data are expressed as the mean±SD. Differences between groups were examined for statistical significance by performing analysis of variance (GraphPad Prism program) and/or Student t tests. $P<0.05$ indicated a statistically significant difference.

**RESULTS**

**Docetaxel in combination with curcumin induces enhanced cell death in 8505C anaplastic thyroid carcinoma cells**
Both docetaxel (1, 2, 5, 10, and 20 nM) and curcumin (1, 5, 10, 25, and 50 µM) used independently showed inhibitory effects on ATC cell proliferation. Compared to controls, cotreatment with docetaxel and curcumin significantly reduced the number of cells (Fig. 1A, B). Treatment of 8505C cells with 25 µM curcumin resulted in 50% growth inhibition. Importantly, the addition of treatment with 5 nM docetaxel resulted in a 2.5-fold decrease in the curcumin concentration that was needed to achieve 50% growth-inhibition. Therefore, we conducted experiments using 10 µM curcumin and 5 nM docetaxel. Although we attempted to extend our findings to other well-established cellular models of human ATCs (CAL62, FRO, and KTC2), we did not observe the synergistic effects of docetaxel and curcumin as observed in 8505C cells (Fig. 1C).

**Docetaxel in combination with curcumin enhances apoptosis in 8505C thyroid carcinoma cells**
To confirm the effects of cotreatment with docetaxel and curcumin on 8505C cells, the cells were treated with the drugs for 24 hours. After Annexin V staining, cells were analyzed by flow cytometry. Cells treated with a combination of docetaxel and curcumin showed increased Annexin V staining as compared with cells treated with either docetaxel or curcumin alone (Fig. 2).

**Increased expression of proapoptotic proteins in 8505C cells treated with docetaxel and curcumin**
To identify the mechanism underlying the enhanced apoptotic response of 8505C cells to co-treatment with docetaxel and curcumin, we assessed the expression of proapoptotic proteins by SDS-PAGE followed by Western blotting. Cells were treated with 5 nM docetaxel and 10 µM curcumin for 24 hours, and whole-cell lysates were subjected to Western blotting. Cotreatment with curcumin and docetaxel increased the expression of

![Fig. 2. Apoptosis-promoting effects of docetaxel (DOC) and curcumin (CUR) treatment. 8505C cells were cultured in 100-mm petri dishes and treated with docetaxel (5 nM) and/or curcumin (10 µM) for 24 hours. Apoptosis was assessed by Annexin V staining followed by flow cytometry analysis. Bars represent mean±SD. CTL, control.](image-url)
proapoptotic proteins. Compared to controls, an active form of caspase-3 (cleaved caspase-3, 17- and 19-kDa bands), and caspase-9 (including cleaved caspase-9, 37 kDa) were significantly increased in cells treated with docetaxel and curcumin (Fig. 3).

**Effects of docetaxel and curcumin on NF-κB activation and COX-2 expression**

To examine the effects of docetaxel on NF-κB signaling, cells were treated with docetaxel and/or curcumin for 24 hours, and nuclear extracts from drug-treated cells were subjected to Western blotting. Compared to controls, the presence of p65 in the nuclear fraction was markedly decreased in cells treated with curcumin alone or with curcumin and docetaxel (Fig. 4). These results confirmed not only the phenomenon of docetaxel-induced NF-κB activation but also that curcumin inhibited this process (Fig. 4). Further, we found that curcumin treatment resulted in the down-regulation of COX-2 protein expression (Fig. 5).

![Fig. 3. Effects of docetaxel (DOC) and curcumin (CUR) treatment on the expression of proapoptotic proteins. 8505C cells were treated with docetaxel (5 nM) and/or curcumin (10 µM) for 24 hours. Expression of cleaved caspase-3 and cleaved caspase-9 were detected by Western blot analysis. Actin was used as an internal control. Relative expression values are presented in the bar graph. Bars represent mean±SD. CTL, control.](image)

![Fig. 4. Effect of docetaxel (DOC) and curcumin (CUR) treatment on the nuclear factor κB (NF-κB) pathway. 8505C cells were treated with docetaxel (5 nM) and/or curcumin (10 µM) for 24 hours. Nuclear NF-κB was detected by Western blot analysis. Actin was used as control. Relative expression values are presented in the bar graph. Bars represent mean±SD. CTL, control.](image)

![Fig. 5. Effect of docetaxel (DOC) and curcumin (CUR) treatment on the expression of cyclooxygenase-2 (COX-2) protein. 8505C cells were treated with docetaxel (5 nM) and/or curcumin (10 µM) for 24 hours. Expression of COX-2 was detected by Western blot analysis. Actin was used as an internal control. Relative expression values are presented in the bar graph. Bars represent mean±SD. CTL, control.](image)
DISCUSSION

ATC is highly resistant to standard therapeutic interventions, including surgical treatment, radiation therapy, and chemotherapy. Recently, it has been found that targeted therapeutic agents enhance the susceptibility of tumors to ionizing radiation by interfering with intracellular signal transduction pathways and DNA repair, thereby leading to senescence-like terminal growth arrest [19-21].

Previous studies have shown that paclitaxel is a promising candidate for therapeutic intervention in anaplastic thyroid carcinoma cases [21,22]. Docetaxel, similar to paclitaxel, has been shown to be clinically effective against solid tumors. However, chemotherapeutic agents such as taxanes and gemcitabine induce NF-κB activation, resulting in drug resistance and therapeutic failure.

The transcription factor NF-κB regulates genes involved in cell survival and proliferation [23]. Thus, NF-κB suppresses apoptosis by up-regulation of antiapoptotic proteins [24]. In addition, elevated basal NF-κB activity, as found in various types of human cancers, causes resistance to chemotherapy [25-27]. Therefore, the NF-κB signaling pathway has been an attractive target for the development of chemotherapeutic drugs. Combination therapy has the potential of producing synergistic effects on tumor suppression without adverse events. Meng et al. [15] reported that in nude mice xenograft models, a combination of NF-κB inhibitor (DHMEQ) and docetaxel was more effective in inhibiting ATC tumor growth than treatment with either of the drugs alone. However, the NF-κB inhibitor caused only a transient inhibition of NF-κB activity, resulting in an early restoration of p65 DNA binding [28].

Curcumin has been reported to inhibit a number of targets, including NF-κB, COX-2, kinases associated with survival signaling (IKK, NIK, and AKT), ERK, and other proteins involved in cell cycle regulation. Thus, curcumin shows a broad range of activities owing to its ability to target multiple pathways involved in regulating diverse cellular processes [29]. In a previous study, NF-κB was bound constitutively in melanoma cell lines. NF-κB binding activity was decreased when treated with curcumin compared with controls [30]. The mechanisms responsible for the inhibition of tumor growth by curcumin are diverse and appear to involve a combination of anti-inflammatory, antioxidant, immunomodulatory, proapoptotic, and antiangiogenic properties via pleiotropic effects on genes and cell signaling pathways at multiple levels [31]. At the molecular level, curcumin disrupts functional and physical interactions between α6β4 integrins and human epidermal growth factor receptor (EGFR) and blocks integrin α6β4/EGFR-dependent pathways in carcinoma cells [32], in addition to inhibiting MMP-2, MMP-9, and vascular endothelial growth factor [33].

The potential therapeutic role of curcumin in conjunction with other therapeutic agents has previously been explored [34]. In bladder cancer cell lines, curcumin was shown to act as a counterbalance to reverse paclitaxel-induced NF-κB activation [35]. Here we demonstrated, for the first time, the synergistic antiproliferative and proapoptotic effects of a combination of docetaxel and curcumin treatment on 8505C anaplastic thyroid carcinoma cells. In the present study, we found that treatment with docetaxel in combination with curcumin enhanced the inhibition of cell proliferation and increased induction of apoptosis in ATC cells as compared to treatment with docetaxel alone. We also showed that curcumin counteracts the effects of docetaxel on NF-κB activation and COX-2 expression in 8505C cells. Thus, docetaxel activated NF-κB and increased COX-2 expression, and treatment with curcumin suppressed both docetaxel-induced NF-κB activation and COX-2 expression in 8505C ATC cells.

Induction of COX-2 in tumor cells by docetaxel may be a protective mechanism, counterproductive to the overall antiangiogenic and antitumor effects of docetaxel [36]. COX-2 plays an important role in tumor progression by promoting angiogenesis, stimulating cell growth, and inhibiting apoptotic cell death [37-39]. Cornetta et al. [40] reported that the expression of COX-2 was upregulated in well-differentiated thyroid cancer, and this was not observed in the normal thyroid. Overexpression of COX-2 may inhibit apoptosis, enhance angiogenesis, and promote cell invasion. Our results suggest that curcumin may counteract the adverse clinical outcomes arising from increased expression of COX-2 in ATC cells.

In conclusion, our study results suggest that curcumin augments the therapeutic efficacy of docetaxel by lowering its threshold concentration required for inhibiting cell proliferation and inducing cell death. Additionally, inhibition of docetaxel-induced NF-κB activation and COX-2 expression by curcumin in 8505C cells revealed a potential anti-angiogenic effect of the combination therapy. Taken together, our findings will lead to improved clinical outcomes in ATC therapy.

CONFLICTS OF INTEREST

No potential conflict of interest relevant to this article was reported.
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