Cycloartobiloxanthone Induces Human Lung Cancer Cell Apoptosis via Mitochondria-dependent Apoptotic Pathway

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Abstract. Background: Lung cancer is one of the most malignant types of cancer and new anticancer agents are still required. Cycloartobiloxanthone, a flavonoid isolated from stem bark of Artocarpus gomezianus Wall. ex Tréc. (Moraceae), has potential for being developed for anticancer therapy. Materials and Methods: Cytotoxicity of cycloartobiloxanthone was evaluated by 3-(4,5-dimethylthiazol-2-yl)-2,5 diphenyltetrazolium bromide (MTT) assay against four human lung cancer cell lines (H23, H460, H292 and A549) and their half-maximal inhibitory concentrations (IC$_{50}$) were assessed. Apoptotic induction in H460 cells was investigated by Hoechst 33342/propidium iodide (PI) staining assay and protein hallmarks of mitochondria-dependent apoptotic pathway were examined by western blot analysis. Results: Cycloartobiloxanthone exhibited potent cytotoxic effect on both small and non-small cell lung cancer cells. Nuclear Hoechst/PI staining revealed that apoptotic cell death was the main mechanism of toxicity of cycloartobiloxanthone. The apoptosis-inducing potency of cycloartobiloxanthone was comparable to those of standard anticancer drugs cisplatin and etoposide at the same concentration. Protein analysis further showed that apoptosis was mediated via mitochondria-dependent pathway. p53 was activated in cells treated with cycloartobiloxanthone. Subsequently, pro-apoptotic protein B-cell lymphoma 2 (BCL2)-associated X protein (BAX) was found to be significantly increased, concomitantly with the decrease of anti-apoptotic proteins BCL2 and myeloid cell leukemia 1 (MCL1). Moreover, markers of the intrinsic apoptosis pathway, namely activated caspase-9, activated caspase-3, and cleaved poly(ADP-ribose)polymerase (PARP), dramatically increased in cycloartobiloxanthone-treated cells compared to the non-treated controls. Conclusion: Cycloartobiloxanthone has anticancer activity against human lung cancer cells by triggering mitochondrial apoptotic caspase-dependent mechanism. This compound might have promising effects for cancer therapy.

Lung cancer is recognized as the first cause of cancer-related death among all cancer types (1). Lung cancers are classified as small cell or non-small cell carcinomas according to whether they are non-epithelial or epithelial-derived, respectively. Currently, treatments for lung cancer are primarily surgical excision supplemented by radiation and chemotherapy. In terms of chemotherapy, cisplatin is a first-line chemotherapy used for both small cell and non-small cell lung cancer, with or without combination drugs such as etoposide or paclitaxel. Although these drugs have good efficacy against lung cancer, relapse and chemotherapy resistance occur (2) and remains a worldwide problem.

Discovery of novel anticancer drugs is needed and many interesting plant extracts have been studied for their anticancer effects (3, 4). Natural product-derived compounds have long been a major source for pharmacologically active compounds. Cycloartobiloxanthone (Figure 1), a flavonoid extracted from stem bark of Artocarpus gomezianus Wall. ex Tréc. (Moraceae), which is a tree widely distributed throughout Thailand locally known as “Hat-nun”, has been investigated for its possible pharmacological activities (5-7). However, to our knowledge, there are no reports about its anticancer activity against human lung cancer cells. The present study aimed to investigate the effects of this compound and its mechanism of actions against lung cancer cells.
Materials and Methods

**Cells and reagents.** Human-derived lung cancer cell lines H460, H292, A549, H23 and human keratinocyte cell line HaCaT were obtained from the American Type Culture Collection (Manassas, VA, USA). The cells were cultured in Roswell Park Memorial Institute 1640 (RPMI-1640) or Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum (FBS), 2 mM L-glutamine and 100 units/ml of each penicillin and streptomycin (Gibco, Waltham, MA, USA) at 37°C with 5% CO₂ in a humidified incubator. 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT), dimethyl sulfoxide (DMSO), Hoechst 33342, propidium iodide (PI), and bovine serum albumin (BSA) were purchased from Sigma Chemical, Inc. (St. Louis, MO, USA). Antibodies directed against caspase-3, caspase-9, poly (ADP-ribose) polymerase (PARP), B-cell lymphoma 2-associated X (BAX), myeloid cell leukemia 1 (MCL-1), B-cell lymphoma 2 (BCL2), β-actin, and their respective secondary antibodies were purchased from Cell Signaling (Danvers, MA, USA). Cycloartobiloxanthone extracted from the stem bark of *Artocarpus gomezianus* was obtained from Pharmacognosy and Pharmaceutical Botany department, the Faculty of Pharmaceutical Sciences, Chulalongkorn University, Thailand.

**Cell viability assay.** As described elsewhere, the MTT assay was performed to determine the cytotoxic effect of cycloartobiloxanthone on lung cancer cells. Lung cancer cells (H460, H292, A549 and H23) were seeded at a density of 1x10⁴ cells/well in 96-well plates and incubated overnight to allow the cells to attach to the plates. Next, the medium was removed and fresh medium with different concentrations of cycloartobiloxanthone (0-100 μM) was added to the cells. After 24 h of treatment, the media were aspirated and the cells were incubated with MTT (400 μg/ml in 1×PBS) for 4 h at 37°C, after which the supernatant solution was removed and 100 μl DMSO was added to dissolve the resulting formazan crystals. The formazan intensity was measured by spectrophotometry at 570 nm using a microplate reader (Perkin Elmer, Waltham, MA, USA). The percentage of cell viability was calculated as the absorbance of treated cells relative to that of untreated cells.

**Nuclear staining assay.** Cells were seeded in 96-well plates at a density of 1x10⁴ cells/well, incubated overnight, and treated with cycloartobiloxanthone (0-50 μM) for 24 h. Hoechst 33342 (10 μg/ml) and PI (5 μg/ml) were added for 30 min. Nuclear condensation and DNA fragmentation of apoptotic cells and PI-positive necrotic cells were visualized and scored using a phase-contrast and fluorescence microscope (Nikon eclipse Ts2; 40x) equipped with a Nikon’s DS-Fi3 digital camera system (Nikon, Tokyo, Japan).

**Western blot analysis.** H460 cells were treated with 0, 10, 20, 50 μM of cycloartobiloxanthone for 24 h. The cells were then incubated with lysis buffer containing 20 mM Tris-HCl (pH 7.5), 1% Triton X-100, 150 mM sodium chloride, 10% glycerol, 1 mM sodium orthovanadate, 50 mM sodium fluoride, 100 mM phenylmethylsulfonyl fluoride and protease inhibitor cocktail (Theera Trading, Bangkokkayai, BKK, Thailand) for 30 min on ice. The cellular lysates were collected by scraping and their protein content was determined using a BCA protein assay kit (Pierce Biotechnology, Rockford, IL, USA). The resultant lysates were added to Laemmli loading buffer and boiled at 95°C for 5 min. Equal amounts of protein from each sample were separated by sodium dodecyl sulfate–polyacrylamide gel electrophoresis and transferred to 0.45 μm nitrocellulose membranes (Bio-Rad, Pathumwan, BKK, Thailand). The resulting blots were blocked for 1 h with 5% non-fat dry milk in Tris-buffer saline with 0.1% Tween containing 25 mM Tris-HCl (pH 7.5), 125 mM NaCl and 0.1% Tween 20 (TBST) and incubated with the primary antibodies listed above at 4°C overnight. After three washes in TBST, the blots were incubated with horseradish peroxidase-conjugated secondary antibodies for 2 h at room temperature. Finally, protein bands were detected using an enhancement chemiluminescence substrate (Supersignal West Pico; Pierce) and quantified using ImageJ 1.51j8 software (National Institutes of Health, Bethesda, MD, USA).

**Statistical analysis.** Data from three or more independent experiments were presented as the mean±standard deviation (SD). Multiple comparisons for significant differences between multiple groups were performed using analysis of variance (ANOVA), followed by individual comparisons with Scheffe’s *post-hoc* test. Statistical significance was considered at *p*<0.05.

Results

**Cytotoxic effect of cycloartobiloxanthone on human lung cancer cells.** Anticancer activity of cycloartobiloxanthone (Figure 1) was first assessed in four types of human lung cancer cells. The human lung cancer H460, H292, A549 and H23 cells were seeded and treated with different concentrations (0-100 μM) of cycloartobiloxanthone for 24 h. The cell viability of treated and non-treated groups was evaluated by MTT assay. As shown in Figure 2, a significant decrease in cell viability of all cell types were observed in response to the treatment of cycloartobiloxanthone at 20 μM. The IC₅₀ of this compound against H460, H292, A549 and H23 cells was 59.19, 49.56, 110.93 and 91.19 μM, respectively (n=3).

**Apoptotic cell death induced by cycloartobiloxanthone.** Having shown the cytotoxic effect of cycloartobiloxanthone on human lung cancer cells, we next determined the mechanism of cell death induced by the compound and the mode of cell death, whether apoptosis or necrosis, was further...
investigated by Hoechst33342/PI nuclear co-staining. The non-treated H460 control cells exhibited normal intact cell morphology and adhered firmly to the culture surface with no cell debris under phase-contrast microscope. In contrast, cycloartobiloxanthone-treated cells exhibited rounded shape and detached from the plate with the formation of apoptotic bodies. Further examination of Hoechst/PI co-stained cells under fluorescent microscopy showed that the
Figure 3. A: Apoptotic and necrotic cell death after 24 h of cycloartobiloxanthone treatment were examined by Hoechst 33342/propidium iodide (PI) co-staining. B: Percentage of apoptotic and necrotic nuclei in cycloartobiloxanthone-treated cells. Data are presented as the mean±SD (n=3). *Significantly different at p<0.05 compared to the untreated control group.
Figure 4. A and B: Apoptotic and necrotic cell death after 24 h of 50 μM cisplatin, etoposide and cycloartobiloxanthone treatments were examined by Hoechst 33342/propidium iodide (PI) co-staining. C: Percentage of apoptotic and necrotic nuclei in each treatment group. Data are presented as the mean±SD (n=3). *Significantly different at p<0.05 compared to the untreated control group.
cycloartobiloxanthone-treated cells exhibited obvious characteristics of apoptosis (Figure 3). Apoptotic cells were indicated by bright blue dots of condensed/fragmented nuclei compared to other dimly blue-stained nuclei, which represented the intact cells. Necrotic cells were characterized by red spots of PI staining. PI cannot pass through intact cell membranes, unless there is any damage leading to loss of membrane permeability. However, late-apoptotic cells may be stained by PI, hence we confirmed the results by observing the cell morphology through merged images of Hoechst and PI staining (Figure 3A). The percentage of apoptotic and necrotic cells found under fluorescence microscopy were consistent with the MTT results and we concluded that cell death caused by cycloartobiloxanthone was mainly due to apoptosis. It is noteworthy that cycloartobiloxanthone rarely induced necrosis even at high concentrations (Figure 3B).

**Cytotoxic effect of cycloartobiloxanthone in comparison to standard anticancer drugs.** Cisplatin and etoposide were long used as anti-cancer agents for the treatment of lung cancer. Here we used cisplatin and etoposide as positive controls. Cell viability MTT assay and Hoechst 33342/PI nuclear co-staining were used to evaluate the potency of cycloartobiloxanthone. Cells were treated with the same dose (50 μM) of cisplatin, etoposide or cycloartobiloxanthone for 24 h and subjected to cell viability, apoptosis and necrosis detection. The results showed that cycloartobiloxanthone had comparable cytotoxic activity to cisplatin and etoposide.

**Figure 5. Western blot analysis of proteins involved in mitochondria-mediated apoptotic cell death.** H460 cells were treated with different doses of cycloartobiloxanthone for 24 h. B: Relative protein levels were quantified by densitometry. Data represent the mean±SD (n=3). *Significantly different at p<0.05 compared to the untreated control group.
Moreover, human keratinocytes, HaCaT cells, were used to evaluate the effect of cycloartobiloxanthone on non-cancerous cells. The results showed that 50 μM cycloartobiloxanthone significantly reduced HaCaT cell viability to a degree comparable to that of 50 μM cisplatin, and 50 μM etoposide (Figure 4A).

Cycloartobiloxanthone induces apoptosis via mitochondria-dependent pathway. To investigate the mechanism of cycloartobiloxanthone-induced apoptosis, apoptotic-related proteins were determined by western blot analysis. Human lung cancer H460 cells were treated with 0-50 μM for 24 h and p53, along with pro- and anti-apoptotic proteins related to mitochondria-mediated apoptosis were evaluated. In response to treatment with cycloartobiloxanthone, p53 was found to be activated and the cellular level of pro-apoptotic proteins was increased, while expression of anti-apoptotic proteins was suppressed. Pro-apoptotic protein BAX was up-regulated, while anti-apoptotic proteins MCL1 and BCL2 were down-regulated in cells treated with cycloartobiloxanthone (Figure 5). In addition, the mitochondria-dependent apoptosis pathway-specific caspase-9 was found to be activated, and apoptotic markers, caspase-3 and PARP were subsequently cleaved. The results indicated that cycloartobiloxanthone mediated apoptosis of lung cancer cells by increasing the ratio of pro- to anti-apoptotic proteins leading to death by mitochondria-dependent pathway.

Discussion

New drugs for cancer therapy are urgently needed, especially for lung cancer due to the limited efficacy obtained from the currently available therapy, and the 5-year survival rate of patients with lung cancer is low (8). Pharmacologically active compounds from natural products have great potential to be developed for anticancer therapies with an expectation of better efficacy and lower toxicity (9). We have demonstrated herein for the first time that cycloartobiloxanthone, a plant-derived compound, has a promising ability to kill lung cancer cells through apoptosis (Figures 2 and 3) at similar potency to the currently used standard drugs.

Apoptosis, or programmed cell death, can be caused by two major mechanisms: the extrinsic pathway via death receptors and the intrinsic pathway via mitochondrial pathways. The intrinsic apoptotic mechanism is very important for cancer inhibition and can be triggered by most anticancer drugs (10). Protein analysis revealed that cycloartobiloxanthone triggered mitochondria-mediated apoptosis via p53 induction, which in turn down-regulated anti-apoptotic BCL2 and MCL1 levels and induced that of pro-apoptotic BAX protein (Figure 6). In apoptosis, the reduction of BCL2 and MCL1, and the induction of BAX leads to the release of pro-apoptotic factors from mitochondria and activates caspase-9 as a downstream target. Activated caspase-9 activates the caspase-3 cascade, inhibits DNA repair, and accomplishes apoptosis (11, 12). Together with the fact that many cancer cells depend upon BCL2 and other anti-apoptotic proteins for their survival, the approach of molecular therapies that target BCL2 protein, such as cycloartobiloxanthone, could have promising activity in

Figure 6. Possible mechanism of action of cycloartobiloxanthone-induced apoptosis. Once cycloartobiloxanthone enters the cancer cell, it activates p53 and inhibits anti-apoptotic B-cell lymphoma 2 (BCL2) family proteins, BCL2 and myeloid cell leukemia 1 (MCL1), resulting in up-regulation of pro-apoptotic BCL2 family protein, BCL2-associated X protein (BAX). Alteration of the balance between these pro- and anti-apoptotic proteins triggers mitochondria-mediated apoptotic pathway through the caspase cascade. Cytochrome c released from mitochondria and apoptotic protease activating factor 1 protein forms the apoptosome with activated caspase-9 (cleavage form). Cleaved caspase-9 then cleaves caspase-3, which is a key caspase that triggers apoptotic processes, one of which is inhibition of DNA repair by poly (ADP-ribose) polymerase (PARP) activity.
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induced cell death in non-small cell lung cancer H460 cells overexpression of BCL2 has been showed to inhibit cisplatin-targeting MCL1 and BCL2 proteins, cycloartobiloxanthone (17). The findings of the current study indicate that, by of this compound itself should be further investigated. other underlying activity against cancer cells and toxicity test impairs intrinsic apoptotic signaling (16). Moreover, the malignancies, including small cell lung carcinoma, and protects cancer cells from various apoptotic stimuli (15). Apoptosis and anoikis (13, 14). In particular, non-small cell lung cancer cells taken from patients have been shown to have elevated MCL1 protein levels compared to the surrounding lung cells, and enhanced MCL1 expression protects cancer cells from various apoptotic stimuli (15).

Amplification and overexpression of BCL2 occurs in many malignancies, including small cell lung carcinoma, and impairs intrinsic apoptotic signaling (16). Moreover, the overexpression of BCL2 has been showed to inhibit cisplatin-induced cell death in non-small cell lung cancer H460 cells (17). The findings of the current study indicate that, by targeting MCL1 and BCL2 proteins, cycloartobiloxanthone may have utility in the treatment of lung cancer. However, other underlying activity against cancer cells and toxicity test of this compound itself should be further investigated.

Conclusion

This study revealed that cycloartobiloxanthone has anticancer activity against several types of lung cancer cells via induction of apoptosis by suppressing expression of anti-apoptotic BCL2 and MCL1 proteins and increasing that of pro-apoptotic BAX protein due to increased p53 protein expression. Understanding the mechanisms of action of this compound may benefit its further development for anticancer treatment.

Conflict of Interest

The Authors declare no conflict of interest in regard to this study.

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References


