The effect of $\alpha$-terpineol on cell cycle, apoptosis and Bcl-2 family protein expression of breast cancer cell line MCF-7

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ABSTRACT

The cytotoxic activity of $\alpha$-terpineol on T47D and HeLa cancer cell lines have been reported. This study was conducted to evaluate the effect of $\alpha$-terpineol on cell cycle, apoptosis and Bcl-2 as well as Bax expression on MCF-7 cell line. The cytotoxic activity of $\alpha$-terpineol was determined using MTT cell assay. Cell cycle and apoptosis were analysed using flowcytometry, whereas Bcl-2 and Bax expression were evaluated using immunohistochemistry. The results showed that $\alpha$-terpineol had cytotoxic effect on the MCF-7 cell lines with an IC$_{50}$ value of 33.0 ± 5.4 $\mu$g/mL. $\alpha$-Terpineol induced cell accumulation in Sub-G1 lead to apoptosis of the MCF-7 cell. Moreover, $\alpha$-terpineol inhibited Bcl-2 and induced Bax expressions. In conclusion, $\alpha$-terpineol has potential anticancer activity against MCF-7 cancer cell line trough through cells cycle inhibition and apoptosis stimulation.

Keywords: $\alpha$-terpineol - MCF-7 cell line – cytotoxicity – apoptosis – Bcl-2 - Bax

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INTRODUCTION

Breast cancer is the most frequently diagnosed cancer and the leading cause of cancer deaths in females worldwide, accounting for 23% (1.38 million) of the total new cancer cases and 14% (458,400) of the total cancer deaths in 2008. In Indonesia in 2008, the incidence rate of breast cancer per 100,000 is 36.2, while the mortality rate per 100,000 is 18.6. In the 2007 data obtained from Dharmais Hospital, National Cancer Center, Jakarta, it was reported that 437 breast cancer patients had been hospitalized among a total of 1,264 out patients.

Chemotherapy is a treatment option for most types breast cancer. It is normally performed in conjunction with surgery and radiotherapy. However, chemotherapy owns disadvantages ranging from adverse effects of drugs to patients’ death. In addition, resistance to anticancer remains a major problem in chemotherapy. Face the problems in this chemotherapy, efforts to find a new anticancer that more sensitive and specific is urgently needed. Medicinal plant has been a source of new anticancer agents during last few decades. Many anticancers used in clinic have been developed from medicinal plants like vincristine, camptothecin, and docetaxel.

α-Terpineol is, a monoterpenooid alcohol, one of natural agents that has potential anticancer activity. α-Terpineol is the major components of terpineol that has been isolated from a variety of plants such as Eucalyptus globulus (Eucalyptus) and Pinus merkusii (Pinus). α-Terpineol is usually present in a mixture of three isomers namely β-, γ- and terpinen-4-ol. The potency of α-terpineol as anticancer candidate has been reported by some authors. α-Terpineol has been proven to be able to induce cell cycle and apoptosis of colon cancer HCT-116 cells in vitro through caspase activation, cytochrome C release and PARP cleavage. Whereas, Hassan et al. proved that α-terpineol is able to prevent the MCF-7 and HeLa cells growth by supressing NF-kB signalling pathway.

In order to develop α-terpineol as a potential anticancer, Budiman et al. synthesized α-terpineol from α-pinene isolated from Indonesian crude turpentine. Furthermore, this α-terpineol synthesized has been proven its cytotoxicity on T47D and HeLa cancer cell lines by induce the cell apoptosis and inhibit cell cycle. In this study we continued to investigate the cytotoxicity of α-terpineol on MCF-7 and its effect on cell cycle, apoptosis and Bcl-2 as well as Bax proteins expression.

MATERIALS AND METHODS

Materials

α-Terpineol was synthesized by Prof Arief Budiman from Department of Chemical Engineering, Faculty of Engineering, Universitas Gadjah Mada, Yogyakarta. MCF-7 cell line was kindly provided by Stem Cell and Cancer Institute Kalbe Farma, Jakarta.

Cell cultures

MCF-7 cells line were culture in tissue culture flask 25 cm² containing DMEM high glucose, insulin, 10% FBS, 2% penicillin-streptomisin, and 0.5% fungizone. Cultures were maintained in 5% CO₂ incubator at 37°C. After 24 hours, medium was replaced and cells were grown until 70% - 80% confluency for further experiments.

Cytotoxicity assay

Cytotoxicity of α-terpineol was evaluated on MCF-7 cells using the MTT assay as reported by Mosmann after modification. Cells were distributed in 96-wells microplates at 2 x 10⁴ cells per well in 100 mL and 100
mL of complete DMEM medium were added. The cell cultures were then incubated in 5% CO₂ incubator at 37°C for 24 hours. After incubation, the medium was removed and replaced with new complete DMEM medium with various concentrations of α-terpineol. Cells culture and α-terpineol were incubated again in 5% CO₂ incubator at 37°C for 24 hours. After the incubation, the medium was removed and the cells were resuspended in DMEM medium, 10 µL of 5 mg/mL MTT [3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide] and then further incubated for 4 hours. The reaction was stopped by adding 100 µL of 10% sodium dodecyl sulfate (SDS) in 0.01N HCl. Microplate was then shaken gently for 5 minutes, covered with aluminium foil and incubated at room temperature overnight. Absorbance of the microplate was measured in an ELISA plate reader at λ₅₉₅ nm. The absorbance values were directly proportional to the number of live cells. The absorbance values in the presence of α-terpineol were compared with that of control cultures without α-terpeniol to obtain cells growth inhibition. For this MTT method, IC₅₀ (inhibitory concentration of 50% cell growth) values were determined by probit analysis based on the relationship between log concentrations versus the percentage of cells growth inhibition.

**Cell cycle analysis**

Cells cycle analysis was performed by flowcytometry. MCF-7 cell line were seeded in 6-well plates at 5x10⁵ cells/well and incubated 24 hours, 37°C, 5% CO₂. After 24 hours, cells were treated with α-terpineol in triplicate at 2 concentrations: ½ IC₅₀ and IC₅₀ for 24 hour. At the end of the incubation period, cells were collected and harvested. After centrifugation, cell pellets were washed twice with 500 µL of cold PBS and then added 1 mL of 70% ice-cold ethanol and stored at -20°C for 30 minutes. The fixed cell centrifugated, the pellet was washed once with PBS and then incubated with Propidium Iodide reagen for 10 minutes in 37°C and transferred to flowcytoube. The cells were immediately analyzed by FACS Calibur flowcytometer to evaluate cell cycle profile. Flowcytometric data were analysed using Cell Quest to evaluate the cells distribution at each phase of the cell cycle namely the sub G1 (apoptosis), S, G2/M, and the cells undergoing polyploidy. The cell cycle inhibition was observed by comparing the cells distribution at G0/G1 and G2/M phases of treated and untreated cells. Ethical approval of the study was obtained the Medical and Health Research Ethics Committee, Faculty of Medicine, Universitas Gadjah Mada, Yogyakarta.

**Apoptosis**

Cells cycle analysis was performed by flowcytometry. MCF-7 cell line were suspended at a final concentration of 5 x 10⁵ cells/well in complete DMEM medium and distributed in 24 wells. The cells were then incubated for 24 hours, at 37°C in 5% CO₂. After 24 hours of incubation, cells were treated with α-terpineol in triplicate at 2 concentrations: ½ IC₅₀ and IC₅₀ for 24 hour. At the end of the incubation period, cells were collected, washed twice with 500 µL PBS, and then incubated with Annexin V Fluos-Propidium Iodide reagent for 10 minutes. The result of the staining was then detected using FACS Calibur.

**Immunohistochemical analysis of Bcl-2 and Bax proteins expression**

Bcl-2 and Bax proteins expression was performed by immunohistochemistry. MCF-7 cell line were distributed in 24-well plates at 75 x 10³ cells/well and incubated at 37°C
in 5% CO₂ for 24 hours. Followed after incubation, cells were treated with α-terpineol in triplicate at 2 concentrations: ½ IC₅₀ and IC₅₀ for 24 hour. At the end of incubation period, cells were washed twice with PBS. The cells were incubated at 4°C for one hour with dilution of primary antibodies against Bcl-2 (dilution 1:400) or Bax (dilution 1:400) and then were washed with PBS three times. The cells were subsequently incubated with biotinylated secondary antibodies for five minutes, washed with PBS and incubated with HRP-conjugated streptavidin. The cells were washed with PBS and then visualized using 3,3’-diaminobenzidine (DAB) chromogen and counterstained with Harry’s haematoxylin. The cells were washed with PBS, dried, mounted in Canada balsam, and observed at 400 x magnification using an image analysis system.

**Statistical analysis**

Data were expressed as the mean ± standard deviation (SD). Statistical comparisons were performed using Student’s t-test. A p value less than 0.05 was considered to indicate statistically significant.

**RESULTS**

Cytotoxicity of α-terpineol on MCF-7 cell line

The inhibition of MCF-7 cells growth after incubation with α-terpineol in various different concentrations for 24 hours is presented in TABLE 1. A dose-dependent manner in MCF-7 cells growth after incubation with α-terpineol was observed. The maximum growth inhibition (100 %) was observed after incubation with α-terpineol in concentration of 200 µg/mL. Probit analysis showed that the IC₅₀ value of α-terpineol on MCF-7 cell line was 33.0 ± 5.4 µg/mL.

![FIGURE 1. Growth inhibition of MCF-7 cells (% ± SD) after 24 hours incubation with the α-terpineol](image-url)
The effect of α-terpineol on MCF-7 cell cycle

The effect of α-terpineol on the MCF-7 cell cycle changes was analyzed by flowcytometry. FIGURE 2 shows the MCF-7 cell cycle profile after incubation with α-terpineol at 2 different concentrations (16.5 and 33.0 µg/mL). α-Terpineol at concentration of 33.0 µg/mL (IC₅₀) induced MCF-7 cell accumulation until 18.59% at Sub-G1 phase. This accumulation was accompanied with the reduction of cycling cells in G0/G1 phase (51.24%), S phase (15.4%) and G2M phase (9.69%) as compared to control. At concentration of 16.5 µg/mL (½ IC₅₀), α-terpineol induced MCF-7 cell accumulation until 24.78% at S phase, as compared to 21.15% in the control cells. The cell accumulation in S phase caused reduction of cycling cells in G2M phase.

The effect of α-terpineol on apoptosis

The effect of α-terpineol on the MCF-7 cell apoptosis was analyzed by flowcytometry using Annexin V Fluos-Propidium Iodide staining. TABLE 1 shows MCF-7 cell distribution in four quadrant, whereas its percentage of early apoptotic cell after 24 hours α-terpineol incubation compared to control is presented in FIGURE 2. The percentage of apoptotic cell after 24 hours incubation with α-terpineol at concentration of 33.0 µg/mL (5.92 ± 0.13%) was significantly higher than that at concentration 16.5 µg/mL (2.57 ± 0.12%) and that control (1.03 ± 0.06%) (p < 0.05).

TABLE 1. MCF-7 cell distribution in four quadrant (lower left/live, lower right/early apoptotic cell, upper right/necrotic cell, upper left/necrotic cell)

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Concentration (µg/mL)</th>
<th>Cell percentage (% ± SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Live</td>
</tr>
<tr>
<td>Control</td>
<td>0</td>
<td>96.42±0.43</td>
</tr>
<tr>
<td>Alpha terpineol</td>
<td>16.5</td>
<td>87.48±0.05</td>
</tr>
<tr>
<td></td>
<td>33.0</td>
<td>86.45±0.37</td>
</tr>
</tbody>
</table>
The effect of α-terpineol on Bcl-2 and Bax expressions

The effect of α-terpineol on Bcl-2 and Bax expressions of the MCF-7 cell were analysed by immunohistochemistry. The expression of Bcl-2 and Bax protein on MCF-7 cell line after 24 hours α-terpineol incubation are presented in FIGURE 4 and 5, respectively. The percentage of Bcl-2 expression after 24 hours incubation with α-terpineol at concentration of 16.5 and 33.0 µg/mL (59.8 and 56.2%) were lower than that control (95.8%). Inversely, the percentage of Bax expression after 24 hours incubation with α-terpineol at concentration of 33.0 µg/mL (100 %) was significantly higher than that at concentration 16.5 µg/mL (68.0 %) and that control (53.2%).
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**FIGURE 5.** The effect of 24 hours a-terpineol incubation on Bax expression of MCF-7 cell line

**DISCUSSION**

This study showed that a-terpineol inhibited the MCF-7 cell growth in a dose-dependent manner with an IC$_{50}$ value of 33.0 µg/mL. The cytotoxicity of a-terpineol on different cancer cell lines has been reported in the previous studies by some authors. Hasan et al. reported that a-terpineol was most active on NCI-H69 lung cancer cell with IC$_{50}$ of 39.4 µg/mL among 14 human cancer cell lines tested. a-Terpineol was also reported active against T47D breast cancer cell line (IC$_{50}$: 20.5 µg/mL) and HeLa cervical cancer cell line (IC$_{50}$: 12.5 µg/mL). Based on a criteria proposed by America National Cancer Institute, a-terpineol can be classified as a potential anticancer agent (IC$_{50}$ about 30 µg/mL).

The effect a-terpineol on MCF-7 cell cycle in this study showed that a-terpineol induced cell cycle arrest in the cell line tested in a dose-dependent manner. This result is consistent with the previous reports which showed that a-terpineol is active in inducing cell cycle arrest. Itani et al. reported that *Salvia libanotica* essential oil containing three bioactive compounds i.e. linalyl acetate, terpeniol and camphor caused significant growth suppression of colorectal cancer cell lines HCT116 (p53$^{+/+}$) in pre G1 phase and in p53$^{-/-}$ cells caused cell accumulation in pre G1 and G2/M phases. Moreover, Hassan et al. also reported that a-terpineol inhibited the proliferation of lymphoma U937-GTB cancer cells in G0/G1 phase lead to reduction in the number of cells in the later stages of cell cycle (S, G2 and M) of the cells.

This study also showed that a-terpineol induced apoptosis of the cell line tested in a dose-dependent manner. The apoptosis induced by a-terpineol may be through Bcl-2 protein family as demonstrated by the decrease of Bcl-2 ad the increase of Bax expressions in this study. The effect of a-terpineol on cancer cell apoptosis has been demonstrated previously. Itani et al. reported that a-terpineol, linalyl acetate and camphor synergised to induce cell cycle arrest and apoptosis, mainly via mitochondrial damage (cytochrome c release), caspase activation, and PARP cleavage, in human colorectal cancer cells. Furthermore, Hassan et al. also reported that a-terpineol exhibited a potential anticancer which acts by suppressing NF-κB which signals various cancer cells line. NF-κB protein is one of transcription factors that
are involved in the control of inflammatory responses, developmental processes, cellular growth and apoptosis.\textsuperscript{18} $\alpha$-Terpineol inhibits NF-$\kappa$B translocation and activity and down-regulates the expression of several NF-$\kappa$B-related genes such as IL-1$\beta$ and IL1R1 resulting in the inhibition of cancer cells growth.\textsuperscript{14}

**CONCLUSION**

In conclusion, $\alpha$-terpineol has potential anticancer activity against MCF-7 cancer cell line trough through cells cycle inhibition and apoptosis stimulation. The cells cycle inhibition is indicated by the induction of cell accumulation in Sub-G1, whereas cells apoptosis induction may be through Bcl-2 protein family as demonstrated by the decrease of Bcl-2 ad the increase of Bax expressions.

**ACKNOWLEDGMENTS**

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**REFERENCES**


