PHYTOCHEMICALS CONSTITUENT ANALYSIS AND CELL CYCLE INHIBITION EFFECT OF ETHANOL EXTRACT OF Litsea cubeba Lour. HEARTWOOD TOWARDS MCF-7/HER-2 CELL LINE

A. Dalimunthe1, P.Sitorus2 and D. Satria2

1Department of Pharmacology, Faculty of Pharmacy, Universitas Sumatera Utara, Medan 20155, Indonesia
2Department of Pharmaceutical Biology, Faculty of Pharmacy, Universitas Sumatera Utara, Medan 20155, Indonesia
3Department of Pharmaceutical Chemistry, Faculty of Pharmacy, Universitas Sumatera Utara, Medan 20155, Indonesia

Corresponding Author: aminah@usu.ac.id

ABSTRACT
Attarasa (Litsea cubeba Lour.) heartwood is potential anticancer especially for breast cancer which is overexpressed in HER-2. This study evaluated phytochemicals analysis of ethanol extract, cytotoxicity, cell cycle inhibition and p53 expression of ethanol extract (EE) of Litsea cubeba Lour. heartwood. EE was analyzed for phytochemicals content with LC-HRMS, cytotoxicity activity was determined by MTT method towards breast cancer cell line MCF-7/Her-2. Cell cycle inhibition and p53 expression were analyzed with the flow cytometry method. EE was found to contain 24 compounds. EE was found to have IC50 191.63 ± 1.56 µg/mL, at concentration 40 µg/mL caused accumulation in G2-M phase (17.71% to 23.34%) and increased p53 expression (0.99% to 17.63%). The results utter that EE of Litsea cubeba Lour. heartwood has a cytotoxic effect through inhibition of the cell cycle. Our further study is to assess the molecular mechanism that responsible for anticancer activity.

Keywords: Breast Cancer, MCF-7/Her-2, Litsea cubeba Lour., Heartwood, Ethanol Extract.

INTRODUCTION
Breast cancer is one of the most incidence rates of cancer in women after cervical cancer. Alteration in lifestyle and daily diet is the most cause that affecting the number of breast cancer patients. The high number of incidence and the high charge treatment for cancer patients, therefore, is a serious problem for looking for other sources of medicine notably through traditional medicine.1-3 HER2 signaling becomes an interesting field to develop a molecular targeted therapy for cancer. Several chemotherapeutic agents targeted on HER2 receptor are developed. Trastuzumab is found to be the first antibody targeting HER2. However, 70% of patients who prolonged use of trastuzumab shows a resistance phenomenon leading to a progression of metastatic cancer due to the increasing of EGFR expression.4,5

Litsea cubeba (Lour.) is a Lauraceae family plant which has contains volatile oils which used as antimicrobial, anticancer on breast cancer, pesticide, antidepressants, antiinflammation, antioxidant, and neuropharmacology. It was showed to be active on HeLa cellse which causes apoptosis through the initiation of activation of caspase 3/7.6,7 Isoquinoline alkaloids in Litsea genus are active as antibacterial.8 The heartwoods contained a high concentration of flavonoid and phenolic and active as an antioxidant and inhibit breast cancer progression through cell cycle arrest. Alkaloid fractions of heartwoods and fruits can decrease PI3KCA, Akt-1 and Akt-2 gene expression. Alkaloid compounds from heartwood have antioxidant effects with ABTS and DPPH methods.9-12 The purpose of our study was to analyze phytochemicals constituent and assess the anti-breast cancer activity of ethanol extract of Litsea cubeba Lour. heartwood on MCF-7/Her-2 cells.
EXPERIMENTAL

Preparation of Extract
The air-dried and powdered heartwoods of *Litsea cubeba* (Lour.) (1000 g) were macerated with ethanol 96% (3x3 d, 7.5 L), The filtrate was evaporated to give a viscous extract.  

Phytochemicals Constituent Analysis with LC-HRMS
Analysis of phytochemicals from EE was analyzed with TSQ executive (Thermo) (LSIH, Brawijaya University) with mobile phase A (0.1% formic acid in water and B (0.1% formic acid in acetonitrile) with gradient method and flow rate 40 µL/minute for the column using Hypersil GOLD aQ 50 x 1 mm x 1.9 µm and the time for analysis was 70 minute. The results were analyzed compound discover with mz cloud software. 

Cytotoxicity Activity
MCF-7/Her-2 cells (1x10^4 cells) were grown in DMEM complete medium. After 24 h incubated, the medium was discharged and treated by EE. The further procedure was followed as previously described. 

Cell Cycle Inhibition Analysis
MCF-7/Her-2 cell line (5x10^5 cells) were seeded and incubated for 24 hours in incubator CO₂ 5%. After treatment and incubation, cells were harvested and analyzed with a flow cytometer were followed the procedure from the previous study.

p53 Expression
MCF-7/Her-2 cell line (5x10^5 cells) were seeded and incubated for 24 hours in incubator CO₂ 5%. After treatment and incubation, cells were collected in a conical tube and washed thrice with cold PBS. The sediment of cells was collected in a microtube. The cells were fixated and p53 FITC antibody was added and incubated at 37°C for 10 min and analyzed using FACScan flow cytometer.

RESULTS AND DISCUSSION

Phytochemicals Constituent Analysis Result of Ethanol Extract
Phytochemicals constituent analysis from EE were determined to obtain the information of compounds contain in Ee with LC-HRMS. The results were given in (Table-1).

| No | Name            | Formula     | Molecular Weight | Retention Time (Min) |
|----|-----------------|-------------|------------------|----------------------|
| 1  | N-methylhernagine | C20H23NO4   | 341.1623         | 0.835                |
| 2  | Salsolinol       | C9H13NO2    | 179.0945         | 1.198                |
| 3  | Pipelic acid     | C6H11NO2    | 129.0789         | 1.228                |
| 4  | 2-methoxresorcinol | C7H6O3  | 108.0210         | 1.786                |
| 5  | Citral           | C10H12O     | 152.1201         | 2.607                |
| 6  | Ethyl cinnamate  | C11H12O2    | 176.0840         | 3.925                |
| 7  | Jasmonic acid    | C12H10O1    | 210.1229         | 4.19                 |
| 8  | Senkyunolide H   | C12H10O4    | 224.1024         | 4.293                |
| 9  | Sinomenine       | C9H12NO4    | 329.1627         | 4.397                |
| 10 | Methyl jasmonate | C13H2O7     | 224.1410         | 4.443                |
| 11 | Quercetin        | C15H10O5    | 302.042          | 4.492                |
| 12 | Merpedine        | C13H2NO2    | 247.1571         | 5.234                |
| 13 | Kaemferol-7-O-glucoside | C17H15O1 | 448.1004         | 5.36                 |
| 14 | p-cymene         | C10H14      | 134.1100         | 5.735                |
| 15 | Alphaprodine     | C10H2NO2    | 261.1727         | 5.803                |
| 16 | Kaemferol        | C15H10O6    | 286.0476         | 5.963                |
| 17 | Carbone          | C16H10O4    | 168.1150         | 6.002                |
| 18 | (-)-caryophyllene oxide | C15H2O    | 220.1827         | 7.136                |
| 19 | Drofenine        | C20H21NO2   | 317.2353         | 7.456                |
| 20 | Samandarone      | C19H29NO2   | 303.2200         | 7.647                |
The results were showed that ethanol extract of *Litsea cubeba* Lour. heartwood contains alkaloids, phenolics, flavonoids, volatile oils and coumarins.

**Cytotoxic Activity**
In treatment, EE was shown to exhibit cell growth. The IC\(_{50}\) value of EE was 191.63 ± 1.56 µg/mL. The cytotoxicity estimate of a natural product is related to the content of active compounds in these plants. Alkaloids, phenolics, flavonoids, volatile oils and coumarins are estimated as active compounds.\(^{19,20}\)

**Analysis of Cell Cycle**
The effect of EE is given in Fig.-1. Treatment with EE in 40 µg/mL caused cell accumulation at G\(_2\)-M phase (23.34%) and for the control cell (17.71%). This fact was to indicate that EE can inhibit cell growth at G\(_2\)-M phase. Flavonoids induce apoptosis and inhibition cell cycle through inhibition of survival signaling proteins such as protein kinase c (PKC-α) and the activation of death signals (PKC-δ).\(^{21}\) Evaluation of cell cycle inhibition was analyzed with flow cytometry using propidium iodide as shown in Fig.-1.

**p53 Expression**
The analysis of the impact of EE on p53 expression was conducted by the flow cytometry method, and the results is given in Fig.-2. Treatment with EE 40 µg/mL caused cell accumulation in M1 area (17.63%) and for the control cell (0.99%). Quercetin is one of the flavonoids compounds that have an anticancer activity which induced expression of p53 and modulation of ROS expression and the cells will be death cause of apoptosis process.\(^{21}\)

**CONCLUSION**
The results reveal that ethanol extract of *Litsea cubeba* Lour. heartwood contains various active compounds and active as an anticancer against MCF-7/Her-2 cell lines by cell cycle inhibition and increase of p53 expression.
ETHANOL EXTRACT OF *Litsea cubeba* Lour.  

A. Dalimunthe et al.