Research Article

Cytotoxic and Antimigration Activity of *Etlingera alba* (A.D.) Poulsen Rhizome

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*Etlingera alba* is one of the *Etlingera* plants that might have anticancer activity. This study aims to investigate the cytotoxic and antimetastatic activity of *E. alba* rhizome fractions and migration cell assay against MDA-MB-231 cell lines, which are used for triple-negative breast cancer (TNBC) treatment assay. The cytotoxic activity was assayed using CCK-8 assay, while the antimetastatic was assayed using migration cell assay for the fractions A–F. They were followed by LCMS/MS profiling to determine the chemical contents in the most active fraction. According to results obtained, fraction B was the most active fraction for cytotoxic activity with an IC50 value of 65.43 μg/mL, while fraction E was the most active fraction for antimetastasis activity against migration rates of 50, 100, and 200 ppm which were 6.80, 3.66, and 3.00%, respectively. Several compounds in fraction B, such as rengyolone, licochalcone A, sugiol, and spinasterol, might have been known to have activity against cancer cells, as well as aschantin and lirioresinol B dimethyl ether from fraction E. In conclusion, the chemical components from *E. alba* rhizome fractions provided potency for discovering new agents for cancer treatment, specifically for TNBC.

1. Introduction

Cancer is the first or second leading cause of death, mainly worldwide, according to WHO (World Health Organization). The incidence and mortality gradually increase each year, including breast cancer. In 2020, the new cases and deaths reported were 2,261,419 cases (11.7%) and 684,999 deaths (6.9%) because of breast cancer [1]. Breast cancer is commonly metastatic cancer, thus it can transfer to other organs and cause death [2].

Triple-negative breast cancer (TNBC) is a kind of breast cancer diagnosed in 15–20% of breast cancer cases. It is categorized with lack or absence of the presence of estrogen (ER), progesterone (PR), and HER-2 (human epidermal growth receptor-2) receptor expression. The TNBC contributes to 170,000 cases with more aggressive behavior, high risk of recurrence, and high mortality than the non-TNBC [3–6].

In treating breast cancer, surgery, radiation therapy, and chemotherapy are the options. The chemotherapy is administered to shrink the tumor before the surgery. However, they can cause many side effects, such as nausea and vomiting, neutropenia, rash and redness, nephrotoxicity, and cardiotoxicity[7]. Therefore, discovering a new agent with the lowest side effect and high efficacy is required because the TNBC has poor prognosis than other breast cancer types. Many compounds from terrestrial plants are the potential for providing anticancer activity. For example, chemotherapy agents such as taxol, vincristine, and vinblastine are from terrestrial plants [6, 8, 9].

*Etlingera alba* (A.D.) Poulsen is one of the newest Etlingera species found in Southeast Sulawesi, and not many studies are conducted on it. The *E. alba* is well known medicinal herb similar to *E. elatior*, mainly used to treat wounds, as deodorant, and as earache medicine. A previous
study showed that *E. alba* rhizome extract has antibacterial and anti-inflammatory properties [10, 11]. *E. alba* becomes a promising candidate as a source of therapeutic agents. Many studies showed that species that belong to similar genus provide similar activities. For example, *E. elatior* is a species that belongs to a similar genus with *E. alba*. It provides antibacterial, antioxidant, anti-inflammatory, and antipyretic properties. Secondary metabolites, including phenolics, alkaloids, flavonoids, steroids, saponins, and volatile oil, are responsible for providing these properties [11].

*E. elatior* flower provides cytotoxic activity against MDA-MB-231 cell line with IC 

50 196.2 μg/mL [12], *E. belalongensis* rhizome and stem with IC 

50 51.00 ± 4.24 and 74.00 ± 2.83 μg/mL, respectively [13], and *E. velutina* rhizome and stem with IC 

50 4.24 and 7.40 ± 4.85 μg/mL, respectively [13]. In addition, bioactive compound trans-4-methoxycinnamaldehyde (4-MCA) identified in *E. pavieana* rhizome extracted IC50 values 39.33 ± 1.53 μg/mL against MDA-MB-231 cell lines [14]. MDA-MB-231 cell is an invasive ductal/breast carcinoma cell which is one of the tools for researching TNBC because it lacks amplifying estrogen receptor (ER) and progesterone receptor (PR) expression, as well as HER2 (human epidermal growth factor receptor 2). It is highly metastatic and aggressive [15, 16]. Thus, we aim to investigate the cytotoxic activity of fractions from extract *E. alba* rhizomes by using CCK-8 assay against MDA-MB-231 cell lines and their metastatic activity by using migration cell assay.

2. Materials and Methods

2.1. Plant Material. *Etlingera alba* rhizomes were obtained from Punggaluku of South Konawe Regency in Southeast Sulawesi (40°19’26″S 122°02’58″E, 325 m). The rhizomes were dried at 40°C and direct sunlight was avoided. The rhizomes were powdered and stored in a sealed jar for further analysis.

2.1.1. Cell Lines Preparation and Culture. MDA-MB-231 cell line was grown in 100 mm plate in CO2 incubator and observed under an inverted microscope. The cell line was seeded in 6-well plates with 2 mL of media per well. A total of 2 × 104 cells/well were needed for cytotoxicity assay and migration assay using scratch wound-healing assay. Cell distribution was observed under an inverted microscope to obtain the even distribution until attaining ~70–80% of confluence.

2.2. Chemical Reagents. Ethanol 95% (technical grade), silica gel 60 (0.063–0.200 mm) Merck 107734, silica gel 60 GF254 Merck 1.07730, TLC silica gel 60 F254 Merck 1.05554.0001, n-hexan (technical grade), ethyl acetate (technical grade), methanol (technical grade), formic acid, water, acetonitrile, Dulbecco’s modified Eagle medium (DMEM)-high glucose (Sigma); fetal bovine serum (Sigma); trypsin TrypLE (Gibco); penicillin-streptomycin (Sigma); phosphate buffer saline 10X (Lonza); cell counting kit-8 (product code: CK04).

2.3. Apparatus. Vacuum rotary evaporator (Buchi®), vacuum liquid chromatography (VLC), LC/MS/MS, Xevo G2-XS QTOF (Waters Corporation, Milford, USA), microplate reader, 96-well microplate; 6-well plate (Nest); plate 100 mm (Nest); serological pipettes 5 mL, 10 mL, and 25 mL (Nest); micropipette (Eppendorf) in various measurements, pipette tips in 1000 μL (GenFollower), 200 μL (GenFollower), and 10 μL (Biogel); CO2 incubator (Thermo Scientific); centrifuge tubes 15 mL and 50 mL (Nest); haemocytometer (Assistant-Germany); light microscope Axio Vert.A1 (Zeiss) camera build-in, and Zen microscope software (Zeiss); biological safety cabinet-level II (ESCO); centrifuge PLC series; Multi-Vortex V-32 (BioSan).

2.4. Extraction and Fractionation. A total of 5.5 kg of powdered *E. alba* rhizome was macerated with 96% ethanol for 3 × 24 hrs at room temperature. The filtrate obtained was concentrated under a low-pressure vacuum with a rotary evaporator at 40°C in 160 g of concentrated extract (2.9%). The extract was impregnated into silica gel and fractionated using vacuum liquid chromatography (VLC, 10 × 5 cm) and the mixture of n-hexane: ethyl acetate with increased polarity (9:1 to 0:10 v/v) and pure MeOH till obtained 6 fractions. The fractions were fraction A (3,239 g); fraction B (4,345 g); fraction C (5,362 g); fraction D (1,563 mg); fraction E (4,432 mg); and fraction F (74,224 mg).

2.5. Cytotoxic Assay. The triple-negative breast cancer cell line (MDA-MB-231) was obtained from the collection of the Department of Pharmacology and Clinical Pharmacy, Universitas Padjadjaran. The cells were grown in Dulbecco’s modified Eagle medium (Sigma-Aldrich, St. Louis, MO) containing 10% fetal bovine serum (Sigma-Aldrich, St. Louis, MO) and 1% penicillin-streptomycin (Sigma-Aldrich, St. Louis, MO). The cell line was seeded in a 96-well plate and cultured for 24 h at 37°C in a humidified atmosphere of 5% CO2. The cells were refreshed with new media and treated with various concentrations of samples, cisplatin was used as control, and DMSO was used as blank. After 24 h, cell counting kit-8 (CCK) reagent (Dojindo, Rockville, MD, USA) was added, and the mixtures were incubated for 4 h. The absorbances were measured using a Tecan Infinite spectrophotometer (λ 450 nm).

2.6. Migration Cell Assay. The migration rates of MDA-MB-231 cells were assessed by the scratch assay method. The cells were seeded in 6-well plates, and after 70–80% confluence, the media was replaced with a low FBS growth medium (concentration 0.5%) and cultured at 37°C in a humidified atmosphere of 5% CO2. After 24 h, the monolayer cells were scratched with sterile yellow micropipette tips, and the cells’ debris was washed with PBS. The cells were treated with various concentrations of a sample by diluting with serum-free DMEM. The cells treated with cisplatin were used as a positive control. The scratch induced that represented wound was photographed at 0 h using phase-contrast microscopy at ×40 magnification at 0 h. After 24 h of...
 incubation, the second set of images was photographed, and the percentage of the closed area was measured and compared with the value obtained at 0 h. An increase in the percentage of the closed area indicated the migration of cells.

3. Results and Discussion

The inhibition of each fraction against MDA-MB-231 cells exhibited various activities. The chemical compounds in each fraction influence these results. Each chemical compound will have a molecular structure with specific physical and chemical properties. The different properties of each of these molecular structures, of course, provide different pharmacological effects. According to Figure 1, the extract and fractions exhibited concentration-dependent inhibition with the higher concentration which gave higher inhibition activity. Fraction B provides better inhibitory activity similar to cisplatin than another fraction against MDA-MB-231 cells. Fraction B provides similar activity to control, which is cisplatin observed 12 hr after treatment. The mechanism itself is similar to cisplatin by blocking the filopodia formation in cell movement. Often in cell culture, the filopodia formation is associated with dynamic cytoskeleton rearrangement induced by TGF-β, which acts as a tumor promoter that leads to metastasis. On MDA-MB-231 cells, cisplatin blocks the cell migration induced by transforming growth factor-beta (TGF-β), which is shown in Figure 3, in which cells migrate towards the center of the wound. The study results demonstrated that fraction E showed the lowest migration rate compared to other fractions and the extract itself by the concentrations 50, 100, and 200 ppm in a dose-dependent manner, which were 6.80, 3.66, and 3.00%. They were not significant to cisplatin as control, which was 5.66, 4.84, and 1.61%, respectively. It suggests that fraction E can prevent the metastasis that produces secondary tumors in an organism [19].

A previous study showed that cisplatin could prevent metastasis by blocking the early step of EMT (epithelial-mesenchymal transition). Cisplatin blocks the cell migration induced by transforming growth factor-beta (TGF-β), which is shown in Figure 3, in which cells migrate towards the center of the wound. The study results demonstrated that fraction E showed the lowest migration rate compared to other fractions and the extract itself by the concentrations 50, 100, and 200 ppm in a dose-dependent manner, which were 6.80, 3.66, and 3.00%. They were not significant to cisplatin as control, which was 5.66, 4.84, and 1.61%, respectively. It suggests that fraction E can prevent the metastasis that produces secondary tumors in an organism [19].

Experiments were performed in a triplicate manner, and the data were recorded and analyzed statistically using SPSS.

To determine the migration rate, the migration percentage (%) was calculated as

\[
\text{%migration} = \frac{(\text{diameter at 0 - hrs}) - (\text{diameter at 24 - hrs})}{(\text{diameter at 0 - hrs})} \times 100\%.
\]
Figure 1: Inhibitory activity of extract and fractions on MDA-MB-231 cell line growth in various concentrations.

Figure 2: IC_{50} value of extract and fraction on MDA-MB-231 cell growth.

Figure 3: The migration rate in percentage for MDA-MB-231 cell line.
Figure 4: *In vitro* scratch assay (×40 magnification) of MDA-MB-231 cells towards extract, fractions, and cisplatin in various concentrations (50 ppm; 100 ppm; and 200 ppm).
Figure 5: LCMS chromatogram of the chemical constituents of fraction B (a) and fraction E (b).

### Table 1: Chemical composition of fraction B.

| No. | Observed Rt (min) | Observed (m/z) [(+)-ESI] | Neutral mass (Da) | Formula | Compound name |
|-----|-------------------|--------------------------|-------------------|---------|---------------|
| 1   | 5.99              | 153.0534 [M+Na]^+       | 130.06299         | C_6H_{10}O_3 | 3-Hydroxy-2,5-hexadione |
| 2   | 6.16              | 183.0640 [M+H]^+        | 182.05791         | C_9H_{10}O_4 | Methyl-3-hydroxy-4-methoxybenzoate |
| 3   | 7.15              | 155.0690 [M+H]^+        | 154.06299         | C_8H_{10}O_3 | Rengyolone |
| 4   | 8.08              | 216.1736 [M+H]^+        | 215.16740         | C_{12}H_{22}N | Candidate mass C_{12}H_{22}N |
| 5   | 8.61              | 179.0691 [M+H]^+        | 178.06299         | C_{10}H_{10}O_3 | tran-Ferulaldehyde |
| 6   | 9.66              | 195.1004 [M+H]^+        | 194.09429         | C_{11}H_{14}O_3 | Ferrozin |
| 7   | 9.74              | 235.0956 [M+H]^+        | 234.08921         | C_{12}H_{16}O_2 | 2-Methoxyvanillic acid |
| 8   | 9.82              | 339.1582 [M+H]^+        | 338.15181         | C_{13}H_{22}O | Licochalcone A |
| 9   | 10.12             | 233.1889 [M+H]^+        | 232.18272         | C_{16}H_{22}O | Candidate mass C_{16}H_{22}O |
| 10  | 10.18             | 301.2154 [M+H]^+        | 300.20893         | C_{20}H_{22}O_3 | Sugiol |
| 11  | 10.34             | 477.3655 [M+Na]^+       | 454.37706         | C_{26}H_{50}N_2O_4 | Candidate mass C_{26}H_{50}N_2O_4 |
| 12  | 10.48             | 505.3973 [M+Na]^+       | 482.40836         | C_{26}H_{50}N_2O_4 | Candidate mass C_{26}H_{50}N_2O_4 |
| 13  | 10.82             | 573.3923 [M+Na]^+       | 550.40221         | C_{29}H_{54}O_2 | Adhyperforin |
| 14  | 11.32             | 429.3721 [M+H]^+        | 428.36543         | C_{29}H_{54}O_2 | Stigmastan-3,6-dione |
| 15  | 11.43             | 413.3771 [M+H]^+        | 412.37052         | C_{29}H_{54}O | Spinasterol |
| 16  | 11.50             | 639.4974 [M+Na]^+       | 616.50668         | C_{39}H_{68}O_2 | Candidate mass C_{39}H_{68}O_2 |
| 17  | 11.69             | 615.4965 [M+H]^+        | 614.49103         | C_{39}H_{68}O_2 | Candidate mass C_{39}H_{68}O_2 |
compounds in fraction B, such as rengyolone [22], licochalcone A [23], sugiol [24, 25], and spinasterol [26], have been known to have activity against cancer cells. For fraction E, thymine [27], aschantin [28], lirioresinol B dimethyl ether [29], and others have potency as anticancer, as well.

For the compound licochalcone A, the antiproliferative and apoptotic effects of MDA-MB-231 cancer cells have been investigated. Licochalcone A provides an antiproliferative activity by inducing cell cycle arrest at G0/G1 in MDA-MB-231 cells [23, 30]. In addition, licochalcone A exhibits apoptotic effects in MDA-MB-231 cells by inducing cleaved caspase-3 and caspase-8 and promoting caspase-3 activity in MDA-MB-231 cells through partial extrinsic signal pathway activation [23, 30]. It also provides an antitumor effect, although our finding resulted in the antitumor effect by wound-healing methods which was not better than fraction E. Licochalcone A decreased phosphorylation of AKT, JNK, p38, and NF-κB. It also enhances the expressions of E-cadherin and reduces vimentin levels in MDA-MB-231 cells. This activity was in a concentration-dependent manner [23]. Thus, licochalcone A is a chemical component in fraction B, influencing the cytotoxic effect on MDA-MB-231 cells.

Fraction E demonstrated the antimetastasis effect through antimigration activity compared to other fractions because of the lirioresinol B dimethyl ether (LBDE) contained in fraction E. LBDE provides antimetastasis and bone microenvironment by inhibiting parathyroid hormone-related protein (PTHrP), matrix metallopeptidase 9 (MMP-9), and cathepsin K activities in breast cancer cells and resorption of osteoclastic bone [29]. LBDE belongs to tetrahydrofurofuran lignans that prevent metastasis by suppressing TGF-β that induced cancer cell viability, migration, invasion, and PTHrP expression. PTHrP has a role in cancer progression and metastasis through a vicious cycle [29]. Fraction E also provides cytotoxic activity, although it was not efficacious than fraction B. This activity might be affected by NF-κB signaling pathway inhibition [31].

Figure 6: Structure molecule of the chemical constituents of fraction B.
Figure 7: Structure molecule of the chemical constituents of fraction E.

| No. | Observed Rt (min) | Observed (m/z) [(+)-ESI] | Neutral mass (Da) | Formula | Compound name |
|-----|-------------------|--------------------------|-------------------|---------|---------------|
| 1   | 1.38              | 113.0332 [M+H]^+         | 112.02728         | C_4H_4N_2O_2 | Candidate mass C_4H_4N_2O_2 |
| 2   | 1.86              | 127.0489 [M+H]^+         | 126.04293         | C_6H_6N_2O_2 | Thymine       |
| 3   | 4.16              | 237.1113 [M+H]^+         | 236.10486         | C_{13}H_{16}O_4 | Candidate mass C_{13}H_{16}O_4 |
| 4   | 4.22              | 167.0692 [M+H]^+         | 166.06299         | C_{9}H_{10}O_3 | 2-Hydroxy-5-methoxy acetophenone |
| 5   | 4.37              | 227.0905 [M+H]^+         | 226.08412         | C_{11}H_{14}O_3 | Dehydrormorroniaglycone |
| 6   | 4.47              | 219.1005 [M+Na]^+        | 196.10994         | C_{11}H_{16}O_3 | Digiprolactone |
| 7   | 4.56              | 197.1273 [M+H]^+         | 196.12118         | C_{10}H_{16}N_2O_2 | Cyclo(Pro-Val) |
| 8   | 4.75              | 203.1055 [M+Na]^+        | 180.11503         | C_{11}H_{16}O_3 | 3-Tert-butyl-4-methoxyphenol |
| 9   | 4.92              | 185.1273 [M+H]^+         | 184.12118         | C_{9}H_{16}N_2O_2 | Cyclo(Ala-Ile) |
| 10  | 5.05              | 209.1525 [M+Na]^+        | 186.16198         | C_{11}H_{18}O_3 | Undecanoic acid |
| 11  | 5.17              | 353.1375 [M+Na]^+        | 330.14672         | C_{19}H_{22}O_3 | Imperanene |
| 12  | 5.65              | 371.1480 [M+H]^+         | 370.14164         | C_{21}H_{24}O_8 | Fragransol D |
Table 2: Continued.

| No. | Observed Rt (min) | Observed (m/z) [(+)-ESI] | Neutral mass (Da) | Formula | Compound name |
|-----|------------------|--------------------------|-------------------|---------|---------------|
| 13  | 5.78             | 401.1584 [M+H]^+         | 400.1522          | C_{23}H_{32}O_{12} | Aschantin    |
| 14  | 6.19             | 265.1426 [M+H]^+         | 264.1361          | C_{13}H_{20}O_{4} | 1,10,4,5-diepoxym-germacren-12,8-olide |
| 15  | 6.29             | 321.2415 [M+H]^+         | 320.2354          | C_{20}H_{12}O_{2} | ent-16a,17-Hydroxy-19-kauranoic acid |
| 16  | 6.49             | 323.2573 [M+H]^+         | 322.2509          | C_{20}H_{14}O_{4} | Darutigenol  |
| 17  | 6.81             | 597.1969 [M+H]^+         | 596.1893          | C_{31}H_{32}O_{12} | Candidate mass C_{31}H_{32}O_{12} |
| 18  | 7.00             | 211.0955 [M+H]^+         | 210.0892          | C_{11}H_{14}O_{4} | Ethyl-5-ethoxy-2-hydroxy benzooate |
| 19  | 7.30             | 443.1072 [M+H]^+         | 442.0938          | C_{13}H_{14}O_{4} | Candidate mass C_{31}H_{32}O_{12} |
| 20  | 7.37             | 425.1582 [M+Na]^+        | 420.1678          | C_{22}H_{26}O_{6} | Neoecuviaphenol |
| 21  | 7.97             | 621.2307 [M+Na]^+        | 598.2414          | C_{33}H_{32}O_{12} | Candidate mass C_{31}H_{32}O_{12} |
| 22  | 8.30             | 303.2311 [M+H]^+         | 302.2248          | C_{20}H_{10}O_{2} | (E)-Labda-(817,15,16-dial |
| 23  | 8.69             | 209.1161 [M+H]^+         | 208.1099          | C_{12}H_{14}O_{4} | Elemicin |
| 24  | 9.04             | 537.2112 [M+H]^+         | 536.2046          | C_{30}H_{32}O_{13} | Isolappao A |
| 25  | 9.16             | 447.2015 [M+H]^+         | 446.1940          | C_{24}H_{18}O_{8} | Lirioresinol B dimethyl ether |
| 26  | 9.28             | 457.1844 [M+H]^+         | 456.1784          | C_{25}H_{26}O_{12} | Kushenol G |

4. Conclusions

Our finding suggests that fraction B of *E. alba* rhizome extract has an antiproliferative effect on the MDA-MB-231 cell lines. On the other hand, fraction E displayed better activity, but not as efficacious as cisplatin in blocking tumor growth with licocalchone A might provide this activity, but not as efficacious as cisplatin in blocking tumor growth. On the contrary, fraction E is still not more efficacious than cisplatin in suppressing tumor growth but has similar activity in blocking metastasis. The lirioresinol B dimethyl ether (LBDE) contained in this fraction E might provide this antimetastasis activity.

Data Availability

The raw data used to support the findings of this study are accessible from the corresponding author.

Conflicts of Interest

The authors declare no conflicts of interest.

Acknowledgments

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