Effectiveness and Anticancer Activity of a Novel Phenolic Compound from *Garcinia porrecta* Against the MCF-7 Breast Cancer Cell Line in vitro and in silico

Darwati Darwati
Ayu Nadila Safitri
Nurul Ambardhani
Tri Mayanti
Nurlelasari Nurlelasari
Dikdik Kurnia

Department of Chemistry, Faculty of Mathematics and Natural Sciences, Universitas Padjadjaran, Jatinangor, 45363, West Java, Indonesia

**Background:** Cancer is a leading cause of death worldwide, with breast cancer being the most common invasive cancer type in women. Several therapeutic strategies have been explored to reduce the mortality rates of breast cancer. Chemotherapy is the most commonly used systemic treatment, but associated with numerous side-effects. Development of anticancer agents with high efficacy and minimal negative effects is therefore an important focus of research. Natural materials provide an excellent source of bioactive compounds. For instance, *Garcinia porrecta* from the Clusiaceae family has multiple pharmacological activities, including antioxidant, anti-inflammatory, antibacterial, antiviral, anti-HIV, antidepressant, and anticancer properties.

**Purpose:** The main objective of this study was to investigate the potential anticancer effects of compounds extracted from the bark of *G. porrecta*.

**Materials and Methods:** Our experiments were divided into three steps: (1) chromatographic isolation of compounds using various separation techniques, such as extraction, separation and purification, (2) characterization via infrared (IR), nuclear magnetic resonance (NMR) and mass spectroscopy, and (3) evaluation of anticancer activity in vitro (MTT assay) and in silico (via analysis of molecular docking against caspase-9, tumor necrosis factor alpha (TNF-α), estrogen receptor alpha (ER-α), and human epidermal growth factor receptor 2 (HER-2)).

**Results:** Depsidone (1) and benzophenone (2) from the ethyl acetate extract of bark of *G. porrecta* were identified as bioactive components. Examination of the activities of these compounds against MCF-7 cells revealed an IC$_{50}$ value of 119.3 µg/mL for benzophenone, whereas IC$_{50}$ for depsidone could not be estimated. Benzophenone activity was lower than that of the positive control doxorubicin (6.9 µg/mL). Depsidone showed the highest binding affinity for HER-2 (−9.2 kcal.mol$^{-1}$) and benzophenone for ER-α (−8.0 kcal.mol$^{-1}$).

**Conclusion:** Benzophenone displays potency as an anticancer agent through blocking ER-α.

**Keywords:** *Garcinia porrecta*, Clusiaceae, phenolic, MCF-7 cells, in vitro, in silico

**Introduction**

Breast cancer is the most commonly diagnosed cancer type in Indonesia. Several therapeutic strategies have been explored over recent years in addition to surgery, radiotherapy and chemotherapy. Chemotherapy is the standard treatment regimen for multiple tumor types, including breast cancer. However, one major limitation is the bystander effect whereby normal cells are eliminated along with cancer cells. Moreover, a number of chemotherapeutic drugs induce negative side-effects and...
chronic toxicities that may be irreversible, particularly in the heart, lung, and kidney.\(^5\)

The MCF-7 cancer cell line has complex morphological and molecular characteristics. An important marker for prognosis is the presence of estrogen receptor (ER).\(^6\) Around 60% breast cancer cells overexpress estrogen receptor-α (ER-α),\(^7\) which plays an important role in transcription of nuclear DNA essential for mammary gland development\(^8\) and is necessary for the breast cancer signaling network.\(^9\) ER-α additionally regulates cell proliferation and differentiation through paracrine mechanisms.\(^10\) These findings support the utility of ER-α as a potential molecular target for inhibiting the progression of malignant cells. The development of molecular targeted drugs with anticancer efficacy is a predominant focus of current research efforts.

The search for novel bioactive compounds as therapeutic agents for deadly diseases, such as breast cancer, is an ongoing hot topic and numerous potential drug designs have been explored as pharmacological treatments.\(^11\) Active components of many herbal remedies are under extensive investigation in view of their affordability, with the aim of improving therapeutic efficacy and minimizing side-effects.\(^12\) In vitro and in vivo approaches and studies on molecular mechanisms are commonly employed to evaluate the active components of natural products. Additionally, computational methods have been widely used for the prediction and design of anticancer drugs.\(^13\)

Selection of drug candidates and isolation of novel bioactive compounds from plants are necessary preliminary steps for identification of effective anticancer agents. Structure–activity relationship (SAR) analyses are frequently utilized to determine candidates that are specific, effective, and selective for target receptors. To establish the molecular mechanisms of action of novel bioactive compounds, in silico clinical trials that effectively simulate and model the interactions between chemical molecules and binding targets are conducted,\(^14\) generating data that support in vitro and in vivo findings and accelerating the drug discovery process.

The Clusiaceae or Guttiferae family belongs to the Malpighiales order comprising 40 to 50 genera with 1000 species of herbs, shrubs, and trees that thrive throughout tropical and subtropical areas. One of the genera belonging to this family is *Garcinia*, distributed in India, Indonesia, West and Central Africa, and Brazil.\(^15\) Various genera of *Garcinia* have been used as folk medicine for treating cancer, oxidative stress, inflammation, and infectious disorders.\(^16\) Several chemical classes of *Garcinia* have been identified to date, including xanthones,\(^17\) benzophenones,\(^23\) anthocyanins and bioflavonoids.\(^24\)\(^,\)\(^25\) These phytochemical compounds have multiple pharmacological activities, including antioxidant, anti-inflammatory, antibacterial, antiviral, anti-HIV, anti-depressant, and anticancer properties.\(^16\)\(^,\)\(^26\)

*Garcinia porrecta* is a tropical-flowering tree that grows abundantly in Indonesia. Plant parts such as fruit, pericarp, leaf, bark, and stem, are traditionally used as medication for fever. Three xanthone (dulxanthone E-G) isolates from bark of *G. porrecta* with strong cytotoxic activity against murine leukemia L1210 cells have been identified.\(^26\) In the current study, the anticancer activity of phytochemical extracts of *G. porrecta* bark was investigated against breast cancer cells. Previously, we isolated a novel polyoxygenated dimer-type xanthone, 5,5’-oxybis (1,3,7-trihydroxy-9H-xanthen-9-one),\(^27\) from ethyl acetate extract of *G. porrecta*. Here, we present the isolation and structural determination of two compounds, depsidone (1) and benzophenone (2) derivatives, from *G. porrecta* (Figure 1). The cytotoxicity of these newly isolated compounds against the human breast cancer cell line MCF-7,
were examined with the MTT assay and molecular docking simulations conducted to validate our in vitro findings.

**Materials and Methods**

**Materials**

G. porrecta bark was collected from Bogor Botanical Garden (Bogor, Indonesia) in April 2019. The plant was identified and deposited in the Herbarium Bogoriense (No. IV.K.78a; Center of Biological Research and Development, National Institute of Science, Bogor, Indonesia). Distilled organic solvents used for extraction and purification included methanol, n-hexane, ethyl acetate, acetone, ethanol, and distilled water, while the chemicals for spectroscopic analysis were used pro-analyzed (p.a.) grades. Chromatographic separations were conducted on Silica G 60 (0.063–0.200 mm and 0.200–0.500 mm) (Merck, Darmstadt, Germany) and ODS RP-18 (0.040–0.063 mm) (Merck) columns. Silica G 60 F254 plates (0.25 mm, Merck) and ODS RP-18 F254S plates were used for thin-layer chromatography (TLC) and detection achieved by spraying with 5% AlCl3 in ethanol (v/v), followed by heating.

Human breast cancer cell line (MCF-7 ATCC-HBT-22TM) was used in this study. This cell line was a generous gift from Prof. Ahmad Faried, dr., PhD., SpBS(K), FICS (Faculty of Medicine, Universitas Padjadjaran) and the use of these cells was approved by the Research Ethics Committee Universitas Padjadjaran. MCF-7 cells were suspended in Roswell Park Memorial Institute (RPMI) 1640 medium supplemented with 10% fetal bovine serum solution (Gibco), 2% penicillin-streptomycin, and 0.5% Fungizone (Gibco) and added by RPMI 1640 media 100% and incubated for 2–3 days at 37°C in a 5% CO2 incubator.

The three-dimensional structure of caspase 9 used in the computational study was obtained from the Protein Data Bank (PDB) ID: 1NW9. TNF-α (PDB ID: 1TNF), ER-α (PDB ID: 1A52), and HER-2 (PDB ID: 3PP0) were used for screening of compounds with activity against breast cancer cells. 3D structures were retrieved from the RSCB Protein Data Bank (https://www.rcsb.org/) in PDB format. Doxorubicin (CID 31703) was used as a positive control ligand as for compounds 1 and 2 ligands retrieved from the PubChem compound database (https://pubchem.ncbi.nlm.nih.gov/) in SDF format and converted to PDB format with Open Babel 2.4.2 program.

**Isolation of Components of G. porrecta Bark**

Air-dried bark of G. porrecta (2 kg) was ground into powder and extracted with n-hexane (5×2 L), ethyl acetate (5×2 L) and methanol (5×2 L) at room temperature. Solvents were concentrated with a rotary evaporator under pressure to yield crude n-hexane (21 g), ethyl acetate (12.5 g), and methanol (25 g) extracts. The ethyl acetate extract (12.5 g) was fractionated via vacuum liquid chromatography on Silica G 60 using a gradient of n-hexane-ethyl acetate-methanol (0:100; v/v, each 1 L) to obtain eight fractions (A-H). Fraction F (2.07 g) was further fractionated via Silica G 60 F254 plates (0.25 mm, Merck) and ODS RP-18 F254S plates were used for thin-layer chromatography (TLC) and detection achieved by spraying with 5% AlCl3 in ethanol (v/v), followed by heating. The number of viable cells was counted using a hemocytometer.

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**Cytotoxicity Assay of Compounds 1 and 2**

The cytotoxicity of isolated compounds against human breast cancer MCF-7 cells was examined using the 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT) method. The stock culture was grown in flasks containing RPMI supplemented with 10% fetal bovine serum (FBS) and 1% penicillin-streptomycin and incubated at 37°C for 24 h. After the medium was changed, MCF-7 cells were detached and seeded in 96-well microtiter plates. After 24 h, the...
compounds were added to the wells. Cell viability was determined after 48 h by measuring the metabolic conversion of the yellow salt of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide to insoluble formazan, a purple colored product resulting from its reduction in viable cells. Insoluble formazan was diluted with DMSO.28

Data from the MTT assay were read using a microplate reader at 450 nm. Eight concentrations of all the compounds were evaluated (7.81, 15.63, 31.25, 62.50, 125.00, 250.00, 500.00, 1000.00 µg/mL) in 100% DMSO, with a final concentration of 2.5% DMSO in each well.

Cytotoxic activity was expressed as IC\textsubscript{50} and analyzed using a linear regression equation. The percentage of cell viability was calculated as follows:

\[
A = \frac{A_{\text{treated cell}} - A_{\text{medium}}}{A_{\text{control cell}} - A_{\text{medium}}} \times 100\%
\]

\(A = \text{Absorbance}\)

**Molecular Docking of Caspase 9/TNF-α/ER-α/HER-2 with Compounds 1 and 2**

Autodock Vina is open-source software in PyRx 0.8 used for ligand-protein docking and virtual screening for anticancer activity of compounds 1 and 2. Binding of compounds 1 and 2 and doxorubicin to caspase 9, TNF-α, ER-α, and HER-2 protein targets was examined and the ligand was free for bind docking. Conformations were selected based on binding energy, specifically, that with the lowest binding affinity score with a root-mean-square deviation value less than 1.0Å.

Docking results were visualized with PYMOL and analyzed using the Discovery Studio 2020 Client program. Ligand-residue interactions and docking poses in the 3-dimensional molecular images are shown in PYMOL program. Next, these interactions were observed in three-dimensional molecular images for optimal visualization using the Discovery Studio 2020 Client program. The docking pose of each protein-ligand complex was compared to the three-dimensional structures of caspase 9, TNF-α, ER-α, and HER-2 bound to ligands on fatty acid sites. The similarities with ligation pose of compounds bound to fatty acid sites were determined and the relation of the docking pose of the ligands and protein targets analyzed.

**Results**

**Structural Characterization of Compounds 1 and 2 from Bark of *G. porrecta***

Detailed structural characterization of compounds 1 and 2 are provided below based on the data interpretation spectroscopic results, including FTIR, HR-TOFMS \(^{1}\text{H}-\text{NMR}, \text{and} \quad ^{13}\text{C}-\text{NMR}.\)

Spectral data of compound 1 are as follows: IR: 3200, 2653, 1677, and 1600 cm\(^{-1}\) (Supplementary Material 1). HR-TOFMS (positive ion mode) (m/z): 273.0395. \(^{1}\text{H}-\text{NMR} \quad (500 \text{ MHz, acetone-d}_6): \delta_\text{H} \quad 6.8 \quad (1\text{H, d, } 8.0 \text{ Hz, H-1}), \quad 7.4 \quad (1\text{H, dd, } 8.0; 2.0 \text{ Hz, H-2}), \quad 7.5 \quad (1\text{H, d, } 2.0 \text{ Hz, H-4}), \quad 6.8 \quad (1\text{H, d, } 8.0 \text{ Hz, H-6}), \quad 7.4 \quad (1\text{H, dd, } 8.0; 2.0 \text{ Hz, H-7}), \quad 7.5 \quad \text{ppm (1H, d, } 2 \text{ Hz, H-9)} \quad \text{(Supplementary Material 3).} \quad ^{13}\text{C}-\text{NMR} \quad (125 \text{ MHz, acetone-d}_6): \delta_\text{C} \quad 115.6 \quad (\text{C-1}), \quad 123.6 \quad (\text{C-2}), \quad 150.6 \quad (\text{C-3}), \quad 117.4 \quad (\text{C-4}), \quad 145.5 \quad (\text{C-4a}), \quad 167.5 \quad (\text{C-5}), \quad 123.1 \quad (\text{C-5a}), \quad 115.6 \quad (\text{C-6}), \quad 123.6 \quad (\text{C-7}), \quad 150.6 \quad (\text{C-8}), \quad 117.4 \quad (\text{C-9}), \quad 145.5 \quad (\text{C-9a}), \quad 167.5 \quad (\text{C-10}), \quad 123.1 \quad \text{ppm (C-10a)} \quad \text{(Supplementary Material 4).}\)

Spectral data of compound 2 are as follows: IR: 3381, 2969, 1597, 1475 cm\(^{-1}\) (Supplementary Material 2). HR-TOFMS (positive ion mode) (m/z): 291.0859. \(^{1}\text{H}-\text{NMR} \quad (500 \text{ MHz, acetone-d}_6): \delta_\text{H} \quad 6.2 \quad (1\text{H, s, H-3}), \quad 6.2 \quad (1\text{H, s, H-5}), \quad 6.7 \quad (1\text{H, d, } 2.0 \text{ Hz, H-2}), \quad 6.5 \quad (1\text{H, t, } 2.0 \text{ Hz, H-4}), \quad 6.7 \quad (1\text{H, d, } 2.0 \text{ Hz, H-6}), \quad 3.6 \quad (\text{3H, s, 2-OMe}), \quad 3.6 \quad \text{ppm (3H, s, 6-OMe)} \quad \text{(Supplementary Material 5).} \quad ^{13}\text{C}-\text{NMR} \quad (125 \text{ MHz, acetone-d}_6): \delta_\text{C} \quad 111 \quad (\text{C-1}), \quad 161 \quad (\text{C-2}), \quad 92.6 \quad (\text{C-3}), \quad 159.3 \quad (\text{C-4}), \quad 92.6 \quad (\text{C-5}), \quad 161 \quad (\text{C-6}), \quad 195 \quad (\text{C-7}), \quad 141.7 \quad (\text{C-1}), \quad 108.3 \quad (\text{C-2}), \quad 159.4 \quad (\text{C-3}), \quad 108.3 \quad (\text{C-4}), \quad 159.4 \quad (\text{C-5}), \quad 108.3 \quad (\text{C-6}), \quad 55.9 \quad (\text{2-OMe}), \quad 55.9 \quad \text{ppm (6-OMe)} \quad \text{(Supplementary Material 6).}\)

Compound 1 was obtained as a pale yellow crystal and HR-TOFMS revealed an ion peak m/z 273.0395 [M+H]\(^{+}\) corresponding to the molecular formula C\textsubscript{11}H\textsubscript{8}O\textsubscript{6} with 11 degrees. The IR spectrum contained absorption bands at ν\textsubscript{max} 3200 (hydroxyl), 2653 (C-H stretching of aliphatic groups), 1677 (carbonyl), and 1600 cm\(^{-1}\) (aromatic C=C, respectively.

The \(^{1}\text{H}-\text{NMR} \) spectrum displayed signals corresponding to two doublet aromatic protons at δ\textsubscript{H} 6.8 (1H, d, J = 8.5 Hz, H-1,6) and 7.5 (1H, d, J = 2 Hz, H-2,7) as well as one doublet aromatic proton at δ\textsubscript{H} 7.4 (1H, dd, J = 8.5 and 2 Hz, H-4,9). The \(^{13}\text{C}-\text{NMR} \) spectrum contained 14 signals attributable to two
benzene rings and two carbonyl carbons. These signals were classified by their chemical shifts on distortionless enhancement by polarisation transfer (DEPT) and heteronuclear single-quantum correlation (HSQC) spectra as six sp² methine carbons, four sp² oxygenated carbons, two sp² quaternary carbons, and two carbonyl carbons at δ_C 167.5 ppm.

The heteronuclear multiple-bond correlation (HMBC) spectrum showed correlations from H-4,9 (δ_H 7.5) to C-4a,9a (δ_C 145.5) and C-3,8 (δ_C 150.6), confirming the location of a hydroxyl group at the 3,8-position (Figure 2). The carbonyl position was determined on the basis of correlations from H-4,9 (δ_H 7.5) to C-5,10 (δ_C 167.5). Another aromatic proton was identified at the 2,7-position based on correlations from H-2,7 (δ_H 7.4) to C-4,9 (δ_C 117.4), C-3,8 (δ_C 150.6) and at the 1,6-position based on correlations from H-1,6 (δ_H 6.8) to C-3,8 (δ_C 150.6), C-4a,9a (δ_C 145.5), C-2,7 (δ_C 123.6), and C-5a,10a (δ_C 123.1). 1H-1H COSY and further HMBC correlation data were in agreement with this finding, leading to the identification of the compound as a depsidone derivative (1), as shown in Figure 1.

![Figure 1](image1)

Compound 2 was obtained as a yellow-brown gum with a molecular weight (m/z) of 291.0859 [M+H]+ corresponding to a molecular formula of C_{15}H_{14}O_{6} with nine degrees. The IR spectrum exhibited absorption bands at V_{max} 3381 (hydroxyl), 2969 (C-H stretching of aliphatic bond), 1597 (carbonyl), and 1475 cm\(^{-1}\) (aromatic C=C), respectively.

The 1H-NMR spectrum contained signals corresponding to a singlet aromatic proton at δ_H 6.2 (1H, s, H-3,5), two aromatic methoxy groups at δ_H 3.6 (3H, s, OCH\(_3\)-2,6), two doublet aromatic protons at δ_H 6.7 (1H, d, J = 2 Hz, H-2',6') and a triplet aromatic proton at δ_H 6.5 (1H, t, J = 2 Hz, H-4'). The 13C-NMR spectrum exhibited 15 signals attributable to two benzene rings and one carbonyl carbon. These signals were classified by their chemical shifts on DEPT and HSQC spectra as five sp² methine carbons, five sp² oxygenated carbons, two sp³ oxygenated carbons (methoxy), and a carbonyl carbon at δ_C 195 ppm.

The HMBC spectrum showed correlations from OCH\(_3\)-2,6 (δ_H 3.6) to C-2,6 (δ_C 161), confirming the presence of a methoxy group at the 2,6-position (Figure 2). The carbonyl position was determined on the basis of the correlation from H-2',6' (δ_H 6.7) to C-7 (δ_C 195). Another aromatic proton was determined at the 3,5-position based on correlations from H-3,5 (δ_H 6.2) to C-1 (δ_C 111) and C-2,6 (δ_C 161). 1H-1H COSY and further HMBC correlations were in agreement with this finding, leading to the identification of the compound as a benzophenone derivative (2), as shown in Figure 1.

![Figure 2](image2)

Cytotoxic Bioassay of Compounds 1 and 2
The cytotoxic activities of compounds 1 and 2 were evaluated against the MCF-7 breast cancer cell line according to a previously published method. The isolated compounds were considered active inhibitors of cell growth at IC\(_{50}\) ≤ 20 μg/mL, moderately cytotoxic at 21–200 μg/mL, weakly

| Ligand | Caspase 9 | TNF-α | ER-α | HER-2 |
|--------|----------|-------|------|-------|
| Compound 1 | −6.3 | −6.0 | −6.7 | −9.2 |
| Compound 2 | −6.0 | −5.3 | −8.0 | −6.7 |
| Doxorubicin | −6.0 | −6.3 | NE | NE |
| Tamoxifen | NE | NE | −5.6 | NE |
| Herceptin | NE | NE | NE | −7.3 |

**Abbreviation:** NE, not estimable.
cytotoxic at 201–500 µg/mL, and not cytotoxic at IC$_{50}$ >501 µg/mL. 29,30 According to these thresholds, the cytotoxicity of depsidone (1) could not be estimated (NE) since cell viability was >80% while benzophenone (2) exerted a cytotoxic effect with an IC$_{50}$ value of 119.3 µg/mL. The anticancer activity of doxorubicin against MCF-7 was additionally compared as a positive control. The IC$_{50}$ value of doxorubicin was determined as ~6.9 µg/mL using the linear regression equation. The plot of the MTT assay of these compounds are shown in the Supplementary Materials 7–9.

Table 2 Hydrophilic Interactions of Ligand-Protein Complexes

| Ligand   | Caspase 9 | TNF-α | ER-α | HER-2 |
|----------|-----------|-------|------|-------|
| Compound 1 | Arg286, Tyr329, Tyr265, Arg268 | Asn46, Leu26, Ala134 | Glu330, Asp332, Arg335, Tyr331, Leu345 | Asp808, Gly729, Thr862, Val734, Cys805, Ala751, Lys753, Leu852 |
| Compound 2 | Thr308, Trp323, Tyr324, Asp309, Leu307, Trp310 | Leu26, Trp28, Gln47, Ser133 | Glu353, Gly521, His524, Phe404, Ala350, Leu387, Leu391, Leu525 | Cys805, Arg849, Leu726, Asp863, Val734, Leu852, Lys753, Ala751 |
| Doxorubicin | Tyr329 | Tyr119 | NE | NE |
| Tamoxifen | NE | NE | – | NE |
| Herceptin | NE | NE | NE | Asn850, Asp808, Arg849, Cys805 |

Abbreviation: NE, not estimable.

Figure 3 Binding sites on caspase 9 for 2 (A) and 1 (B).
Molecular Interactions of Compounds 1 and 2 with Caspase 9/TNF-α/ER-α/HER-2

Following the in vitro cytotoxicity assay of depsidone (1) and benzophenone (2), the molecular mechanisms of action were predicted via computational simulation. Estrogen receptors (ER) are a group of receptor proteins activated by estrogen hormones (17β-estradiol). The MCF-7 cell line is ER-α-positive and proliferates in response to estradiol. Here, we examined interactions of the novel isolated compounds with ligand binding domains (LBD) of caspase 9/TNF-α/ER-α/HER-2. The strength of interactions between each compound and ER-α was determined from the relative binding affinity (RBA).

Tables 1 and 2 present interpretations of the docking results (binding affinity and hydrophilic interactions) for compound 1, compound 2, doxorubicin, tamoxifen and herceptin against caspase-9, TNF-α, ER-α, and HER-2, respectively.

Binding affinity of the compound 1–HER-2 complex showed the greatest value (−9.2 Kcal/mol) relative to the other complexes while the highest value was obtained for the compound 2-ER-α (−8.0 Kcal/mol).

In terms of the ligand-caspase 9 complex, binding affinity of compound 1–caspase 9 was greater than doxorubicin-caspase 9 and compound 2–caspase 9 complexes, which had similar binding affinities. However, the binding pockets of compound 1–caspase 9 and doxorubicin–caspase 9 complexes were in the same position, as characterized by interactions with the same residue (Tyr329). Compound 1–Caspase 9 displayed more hydrophilic
interactions relative to the doxorubicin–caspase 9 complex (Figure 3).

The binding affinities of compounds 1 and 2 for TNF-α were lower than that of doxorubicin, but to a non-significant extent, with values of −6.0, −5.3, and −6.3 Kcal/mol, respectively. The ligand–TNF-α complexes utilized the same binding pocket, as shown in Figure 4.

For the ligand–ER-α complex, the binding affinity of compound 2–ER-α was higher compared to compound 1–ER-α and tamoxifen–ER-α complexes (−8.0, −6.7, and −5.6 Kcal/mol, respectively). Figure 5 shows the similarity in binding pockets between complexes of compounds 1 and 2 with ER-α although the residual bonds were different.

The binding affinities of compound 1, compound 2 and herceptin to HER-2 were −9.2, −6.7, and −7.3 Kcal/mol, respectively, with compound 1 displaying the highest affinity. We observed binding of Asp808, Arg849, and Cys805 residues to all the ligands tested (Figure 6).

**Discussion**

The cytotoxic activities of compounds 1 and 2 were evaluated via the MTT assay, a classic, simple, and economical colorimetric method for evaluating anticancer properties.

The viability of MCF-7 cell lines after exposure to a range of concentrations of each compound (7.81 to 1000 µg/mL) for 24 h was assessed using DMSO as a negative control. At a final concentration of 2.5% DMSO and lower (same percentage of DMSO at all compound concentrations), cell viability was not affected. Untreated cells represented the control group and were considered 100% viable.

According to the results, compound 1 induced no obvious inhibition of MCF-7 cell viability at all tested concentrations. The IC50 value could not be extrapolated and >80% cell viability was observed with all the experimental concentrations. Meanwhile, our novel compound 2 clearly exerted a cytotoxic effect, inducing inhibition of cell viability with an IC50 value of 119.3 µg/mL. However, the cytotoxic effect of
compound 2 was significantly lower than that of the available anticancer agent, doxorubicin (IC\textsubscript{50} of 6.99 µg/mL). The IC\textsubscript{50} value of doxorubicin against MCF-7 cell lines has been previously reported as 0.417 µg/mL.\textsuperscript{33}

Docking experiments showed that compound 2 was more effective in inhibiting ER-α activity while compound 1 was more effective against HER-2, as indicated by the greater binding affinity values compared to other compounds. Binding affinity depicts the strength of the ligand-protein bond, whereby greater negative affinity is correlated with a more stable and stronger bond. Hydrophilic interactions, such as hydrogen bonds, play an important role in determining binding affinity value and affect the stability of ligand-protein complexes owing to stronger interactions.\textsuperscript{34,35}

In experiments on ligand–ER-α complex formation, compound 2 had the highest binding affinity, suggesting efficient occupation of the active ER-α pocket. Compound 2 displayed more hydrophilic bonds (Glu353, Gly521, His524, Phe404, Ala350, Leu387, Leu391, Leu525) relative to compound 1 and tamoxifen. Glu353A, Ala350, His524, Phe404, Leu387, Leu391 and Leu525 residues are located at the active pocket of ER-α.\textsuperscript{36} No residual similarity was evident between the complexes of compound 1–ER-α and compound 2–ER-α. However, binding sides of the residues are sufficiently close to allow competition between the two.

Compound 1 showed the greatest binding affinity (–9.2 Kcal/mol) to HER-2, a growth factor receptor with

Figure 6 Binding sites on HER-2 for 2 (A) and 1 (B).
implications in cancer, compared to compound 2 and herceptin. Heterodimerization of cancer is mediated by HER-2. In our experiments, Asp808, Arg849, and Cys805 residues bound to all test ligands within similar binding pockets, which could potentially lead to competition, especially between compounds 1 and 2 displaying the same number of hydrophobic interactions.

Doxorubicin is often used in the treatment of breast cancer based on induction of higher activities of caspase 9 protein and TNF-α. The docking results of ligand–caspase 9 complexes showed that all ligands had similar binding affinity and activity against caspase 9. In particular, the Tyr329 residue was involved in hydrophobic interactions of compound 1–caspase 9 and doxorubicincaspase 9 complexes. In terms of ligand–TNF-α complexes, compound 1 was predicted to have equivalent activity as doxorubicin, but with different binding pocket positions, therefore leading to no competitive reactions.

**Conclusion**

Novel depsidone (1) and benzophenone (2) derivatives were isolated from stem bark of *G. porrecta* belonging to the Clusiaceae family for the first time. Compound 1 showed no inhibitory activity against the human breast cancer cell line MCF-7 while compound 2 induced inhibition with an IC₅₀ value of 119.3 µg/mL. Docking simulation and binding affinity experiments revealed that compound 1 was more effective in inhibiting HER-2 while compound 2 was more effective against ER-α. Our data support the utility of *G. porrecta* as a natural plant source of potentially therapeutic bioactive compounds. In view of the collective findings, further research on drug discovery from *Garcinia* genus, including in vitro, in vivo and clinical studies, is warranted.

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**Disclosure**

The authors have no conflicts of interest for this work to declare.

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