Research Article

In Vitro Antiplasmodial, Heme Polymerization, and Cytotoxicity of Hydroxyxanthone Derivatives

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1.Introduction

Malaria is an infectious disease caused by a Plasmodium parasite transmitted through female Anopheles mosquito bites [1]. There are five types of Plasmodium species that cause malaria in humans: Plasmodium falciparum, Plasmodium vivax, Plasmodium ovale, Plasmodium malariae, and Plasmodium knowlesi. Plasmodium falciparum (P. falciparum) is a cause of malaria with severe symptoms which can lead to death [2].
hemozoin to protect the *Plasmodium* life. Hemozoin formation occurs only in the *Plasmodium* infection of the erythrocytic cycle. Hemozoin has a similar structure to β-hematin, while heme is similar to hematin. Therefore, by assessing this process, a compound can be developed into an antimalarial drug using the mechanism of action of heme polymerization. *In vitro* hematin can be polymerized into β-hematin in the acidic conditions, which has the same properties as existing hemozoin in the *Plasmodium* [6].

*Plasmodium falciparum* resistance to antimalarials is one of the factors of malaria treatment failure [7–9]. More advanced research is required to find new antimalarials. One of the strategies is to synthesize new compounds from the guiding compounds that have been known to have antiplasmodial activity. The determination of a compound to be used as a guide compound can be based on the resemblance of its chemical structure with other compounds that have been known to have high antiplasmodial activity [10].

One of the compounds that are promising to be developed as an antimalarial alternative is the xanthone derivative compound. Xanthone is a natural phenolic compound known to have antiplasmodial activity. Various studies have shown that the xanthone compounds of natural materials are proven to have the inhibition of *Plasmodium* growth [11, 12]. While encouraging as candidates, these potential antimalarials, derived from natural compounds, have a limited amount, so that their availability cannot be guaranteed. Therefore, to ensure the availability and sustainability of its production, it is necessary to develop a synthetic compound that can be remanufactured in large quantities. One study conducted by Amanatie et al. reported that the xanthone derived 2-hydroxyxanthone compound had an antiplasmodial activity with inhibitory concentration of 50% (IC50) of 4.385 μg/mL. The study stated that the hydroxyl group influenced the high antiplasmodial activity of hydroxyxanthone in the xanthone framework. The addition of a hydroxyl group to the xanthone derivatives causes the antiplasmodial activity of the compound to be higher than the xanthone compound before it is tied to the hydroxyl group [13].

Some hydroxyxanthone derivative compounds have been synthesized by Fatmasari [14], however, their antiplasmodial activity is not known. Previous studies showed that an interaction occurs between the phenol compounds with the hematin electronic system. The phenol compound with a hydroxyl group can bind the heme iron [6]. The hydroxyl clusters that bind to hematin can form complexes that will inhibit the formation of β-hematin *in vitro*. The inhibitory test of the polymerization of heme can be used to identify the mechanisms of action of antimalarials. Testing cytotoxicity on Vero cells can be conducted to evaluate the safety of hydroxyxanthone derivative compounds. To develop the hydroxyxanthone derivative compounds as malaria treatment, it is necessary to test their antiplasmodial activity, heme polymerization inhibitory activity, and cytotoxic effect on the Vero cells.

### 2. Materials and Methods

#### 2.1. Testing Compounds and Plasmodium

Five hydroxyxanthone derivatives have been synthesized by Fatmasari [14], i.e., 1,6,8-trihydroxyxanthone (HX1); 1,6-dihydroxyxanthone (HX2); 1,5,6-trihydroxyxanthone (HX3); 1-hydroxy-5-chloro xanthone (HX4); and 1,6-dihydroxy-5-methylxanthone (HX5). The *Plasmodium falciparum* 3D-7 (chloroquine-sensitive strain) and FCR-3 (chloroquine-resistant strain) were obtained from laboratory collection of Department of Pharmacology and Therapy, Faculty of Medicine Public Health and Nursing, Universitas Gadjah Mada, Yogyakarta, Indonesia.

#### 2.2. Plasmodium Culture

The test of antiplasmodial activity began with the culture of *P. falciparum* 3D-7 and FCR-3 using the modified Trager and Jensen method [15]. The plasmodium was cultured in human O red blood cells diluted to 3% haematocrit in RPMI 1640 medium complemented with 10% human O serum. The medium was made by adding 10.43 g of powder RPMI 1640, 6 g HEPES, 2 g NaHCO3, 25 mg gentamycin, and sterile distilled water up to 1 L. The medium pH was adjusted so that it reached ±7.2. It was sterilized using a 0.22 μm filter and stored at 4°C. The complete plasmodium culture medium was made by adding human serum with a concentration of 10% in the medium. The plasmodium culture was incubated in candle jar in an incubator with a temperature of 37°C and was observed every 24 hours.

#### 2.3. In Vitro Antiplasmodial Activity Assay

The plasmodium was synchronized to obtain the ring stage by adding 5% of D-sorbitol. The plasmodium was transferred from culture flask to a conical tube, and then it was centrifuged with a speed of 1000 rpm for 10 minutes. After the supernatant was disposed, the sterile 5% sorbitol was added and incubated for 10 minutes at a temperature of 37°C. The plasmodium suspension was centrifuged again; the supernatant was disposed and the plasmodium was washed by adding culture medium. Then, the plasmodium suspension was centrifuged again, and the supernatant was discarded, resulting in a plasmodium in ring stage only. The parasitemia was calculated from a thin blood preparation. The test used 1% parasitemia at 2% haematocrit in RPMI medium which complemented with 10% human O serum.

Each testing compound was dissolved in RPMI medium. The testing compound in various concentrations with volume of 100 μL was incorporated into the 96-well microplate, and then 100 μL of *Plasmodium* suspension was added. Each series of concentrations was replicated three times. The microplate was incubated at a temperature of 37°C for 72 hours. At the end of the incubation, a thin blood smear was made using 10% Giemsa stain and observed under a light microscope at 1000x magnification. The percentage of parasitemia (counting a minimum of 1,000 erythrocytes) was calculated from the preparation of thin blood and then used to calculate the percentage of plasmodium growth.
inhibition. As a control, plasmodium culture without any testing compounds was considered to have a growth of 100%. Antiplasmodial activity is expressed as IC₅₀, which is the concentration of a compound that is required for 50% inhibition of plasmodium growth. The IC₅₀ value was calculated by probit analysis using SPSS software (IBM Corp., Chicago). The lower the IC₅₀ value obtained, the greater the antiplasmodial activity. The antiplasmodial activity was classified into 5 categories, i.e., excellent (IC₅₀ < 1 μM), good (IC₅₀ 1–20 μM), moderate (IC₅₀ 20–100 μM), low (IC₅₀ 100–200 μM), and inactive (IC₅₀ > 200 μM) [16, 17].

2.4. Flow Cytometry Method. The Plasmodium falciparum strains of 3D-7 were used in this method. The plasmodium synchronized to obtain the ring stage by adding 5% of D-sorbitol. Plasmodium with 1% parasitemia was cultured in a 96-well microplate. The testing compounds were added in duplicate and incubated at 37°C for 72 hours. The samples were centrifuged with a speed of 1000 rpm for 10 minutes and washed twice in 100 μL of phosphate-buffered saline (PBS). Samples were incubated with 50 μL of 1:1000 SYBR green I and 20 μL of CD235A-PE for 15 minutes at room temperature. Cells were washed and resuspended in PBS. Data were obtained using a FACSCalibur with the acquisition of 1,00,000 events per sample. Initial gating was done with uninfected and unstained erythrocytes to account for erythrocyte autofluorescence. The control plasmodium-infected erythrocyte was referred as 100% growth to calculate the percentage of growth inhibition after treated with testing compound.

2.5. In Vitro Heme Polymerization Inhibitory Activity Assay. The heme polymerization inhibitory activity (HPIA) was conducted according to Basilico et al.’s modified method [6]. The 100 μL of 1 mM hematin in 0.2 M NaOH was added into a microtube. Then, 50 μL of testing compound at various concentrations (20.475; 10.238; 5.119; 2.580; 1.269 mM) was added in triplicates. Distilled water was used as a negative control. The 50 μL solution of glacial acetic acid (pH 2.6) was added into the microtube to initiate the polymerization reaction and incubated at a temperature of 37°C for 24 hours. Microtubes were centrifuged at 8000 rpm for 10 minutes and the supernatant was discarded and then washed three times using the 200 μL dimethyl sulfoxide (DMSO). Then, the deposition of the hematin crystal was dissolved with 200 μL of 0.1 M NaOH and 100 μL of solution was added into the 96-well microplate.

Absorbance was read using ELISA reader at λ 405 nm. A standard curve was made to illustrate the relationship between the concentration of hematin and its absorbance. The various concentrations of hematin (250; 125; 62.5; 31.25; 15.6; 7.8; and 3.9 mM) were used to make the standard curve. The heme polymerization inhibition was expressed as IC₅₀, which is the concentration of testing compounds that can inhibit 50% of the formation of β-hematin. The IC₅₀ value was calculated by probit analysis using SPSS software (IBM Corp., Chicago). A compound shows to have heme polymerization inhibition activity if it has an IC₅₀ value lower than the IC₅₀ value of chloroquine as in reference (37.5 mM) [17].

2.6. In vitro Cytotoxicity Test on Vero Cell. The in vitro cytotoxicity test was conducted using the 3-(4,5-dimethyl-thiazol-2-yl)2,5-diphenyltetrazolium bromide (MTT) assay method on Vero cell line culture using the Anderson et al.’s modified method [18].

2.7. Vero Cell Line Culture. The cell was removed from the cryo medium. Then, the cell was thawed at 37°C. The liquid cell was subsequently transferred to the tube and a culture medium was added to a volume of 10 mL. Then, the suspension was centrifuged for 15 minutes so that it would form cell pellets. The supernatant was removed and 1 mL of culture medium was added to the cell pellets. Finally, the suspension was transferred to the Petri dish and incubated in the incubator with 5% of CO₂ at 37°C.

2.8. Cell Harvesting. The medium of Vero cell culture was removed, and then the cell was washed using a PBS solution with a volume of PBS ± 1/2 of the initial medium volume. This step was done twice and 1 mL of trypsin 0.25% was added. Subsequently, the cell was incubated in the CO₂ incubator at 37°C for 3 minutes. Then, the cell was resuspended using 2 mL of culture medium and PBS solution was added up to 10 mL. The suspension was centrifuged for 15 minutes so that it formed pellets. Pellets were added with 1 mL of culture medium and resuspended. Take 10 μL of the cell suspension and density of cells was calculated. Then, the cell was added with culture medium to 10 mL. A total of 100 μL of such suspensions was transferred into each well of the 96-well microplate, but three of the wells were emptied as medium controls. Subsequently, the microplate was incubated in a 5% CO₂ incubator at 37°C.

2.9. Preparation of Testing Compounds. Each hydroxyanthone derivative compound was dissolved in DMSO, and a variety of concentrations were prepared in the medium. Each concentration of the testing compound was added into the 96-well microplate triplicately. Cell control and medium control were not given a testing compound solution. The medium control remained blank, while the cell control contained culture medium only. Chloroquine was used as a positive control. Then, the cell was incubated in a 5% CO₂ incubator at 37°C for 24 hours.

2.10. MTT and SDS Addition. After the cell was incubated for 24 hours, the medium was discarded and washed with PBS. Then, each well was added with 100 μL of 5 mg/mL of MTT solution. The 96-well microplate was incubated again in the incubator of CO₂ at 37°C for 4 hours. Then, 100 μL of 10% SDS in 0.01 M HCI was added. The cells were placed in a dark room at room temperature for 24 hours.
2.11. Measurement of the Absorbance and IC\textsubscript{50} Value. The absorbance was measured using the enzyme-linked immunosorbent assay (ELISA) reader at 595 nm wavelength. The cytotoxicity test on the Vero cells was done by calculating the percentage of Vero cell inhibition of the absorption value of the test compound, cell control, and medium control. The cytotoxic effect was expressed as IC\textsubscript{50}. The IC\textsubscript{50} value was obtained from the probit analysis by using SPSS software (IBM Corp., Chicago). The degree of selectivity was expressed as a selectivity index. The selectivity index was obtained from the ratio between the IC\textsubscript{50} cytotoxic effect on Vero cells and IC\textsubscript{50} antiparasomal activity on \textit{Plasmodium} [19]. A compound is said to be safe if the value of the selectivity index is $>10$ [20].

3. Results

3.1. In Vitro Antiplasmodial Activity Assay. The increasing concentrations of testing compounds showed increasing the percentage of \textit{Plasmodium} growth inhibition. The percentages of \textit{Plasmodium} growth inhibition of hydroxyxanthone derivatives and chloroquine on \textit{P. falciparum} 3D-7 and FCR-3 are presented in Table 1. The IC\textsubscript{50} values of hydroxyxanthone derivatives on \textit{P. falciparum} 3D-7 and FCR-3 are presented in Table 2. The lowest IC\textsubscript{50} value of the hydroxyxanthone derivative compound on \textit{P. falciparum} 3D-7 $6.10 \pm 2.01 \mu M$ was found in HX1. In contrast, the highest IC\textsubscript{50} value of hydroxyxanthone derivative compounds on \textit{P. falciparum} strain 3D7 $85.30 \pm 4.87 \mu M$ was found in HX4. The lowest IC\textsubscript{50} value of the hydroxyxanthone derivative compound with the microscopic method on \textit{P. falciparum} FCR-3 was also found in HX1 ($6.76 \pm 2.38 \mu M$) while the highest IC\textsubscript{50} value of hydroxyxanthone derivatives on \textit{P. falciparum} FCR-3 was $89.85 \pm 17.69 \mu M$ found in HX4. These results showed that the HX1 showed good antimalarial activity, whereas HX2, HX3, HX4, and HX5 had moderate antiparasomal activity on \textit{P. falciparum} strain 3D-7 and FCR-3. Chloroquine as positive control had IC\textsubscript{50} value $0.01 \pm 0.001$ on \textit{P. falciparum} strain 3D-7 and $0.11 \pm 0.052$ on \textit{P. falciparum} strain FCR-3.

3.2. Flow Cytometry Method. In flow cytometry assay, the compound HX1 with an optimal concentration of $102.38 \mu M$ showed inhibition activity of \textit{P. falciparum} 3D-7 growth. The higher concentration of compound HX1 with a concentration of $204.75 \mu M$ showed the higher inhibition activity by more than 90%. The flow cytometric analysis diagram of compound HX1 on \textit{P. falciparum} 3D-7 is presented in Figure 1.

3.3. In Vitro Heme Polymerization Inhibitory Activity Assay. The results showed that the higher the concentration of the HX1, the lower the amount of $\beta$-hematin formed, and the higher the inhibitory percentage of $\beta$-hematin. The HX1 at concentration of $20.475 \mu M$ were able to inhibit the polymerization of heme with IC\textsubscript{50} value which is $2.854 \mu M$. In comparison, the IC\textsubscript{50} value of chloroquine as the positive control is $1.478 \mu M$. The IC\textsubscript{50} values of 1,6,8-trihydroxyxanthone (HX1) compound and chloroquine in inhibiting the formation of the $\beta$-hematin are presented in Table 3.

3.4. In Vitro Cytotoxicity Test on Vero Cell. Among 5-hydroxyxanthone compounds tested, the compound HX1 showed the highest IC\textsubscript{50} value in \textit{in vitro} cytotoxicity test on Vero cells ($3394.90 \pm 435.44 \mu M$). Compound HX1 showed the highest value of the selectivity index (cytotoxicity/antiplasmodium ratio on \textit{P. falciparum} strain 3D-7 and FCR-3) with range of 502.20–556.54. The IC\textsubscript{50} values of hydroxyxanthone compounds on Vero cells are presented in Table 4. The selectivity index of hydroxyxanthone compounds and chloroquine is presented in Table 5.

4. Discussion

4.1. In Vitro Antiplasmodial Activity of Hydroxyxanthone Derivative Compounds. The study showed that the hydroxyxanthone derivative compounds tested showed different levels of antimalarial activity. Compound 1,6,8-trihydroxyxanthone (HX1) showed the best antimalarial activity among five hydroxyxanthone compounds tested using microscopic method. The category of antimalarial activity of the hydroxyxanthone compound was determined according to Batista et al.’s criteria [17] that divide a compound into five categories, i.e., very good (IC\textsubscript{50} < 1 \mu M); good (IC\textsubscript{50} 1–20 \mu M); moderate (IC\textsubscript{50} 20–100 \mu M); low (IC\textsubscript{50} 100–200 \mu M); and inactive (IC\textsubscript{50} > 200 \mu M). In \textit{in vitro} antimalarial activity assay conducted on \textit{Plasmodium} strains of 3D-7 and FCR-3 showed an IC\textsubscript{50} value of $6.10 \pm 2.01 \mu M$ and $6.76 \pm 2.38 \mu M$ for the 1,6,8-trihydroxyxanthone(HX1), which showed that this compound has a good antimalarial activity. The flow cytometry analysis with SYBR green I and CD235A demonstrated a clear separation of infected erythrocytes with \textit{P. falciparum} and uninfected erythrocytes (Figure 1). This method identifies \textit{P. falciparum} based on nucleic acid contents [21].

Based on its structure, HX1 consists of a hydroxyl group, C-H, aromatic (C-C), cluster that characterizes the xanthone group, which is the top of the carbonyl group (C=O) (carbonyl number 9) and the other group (C-O-C) (Carbon numbers 4a and 4b) [14]. The hydroxyl of the xanthone compound can bind to the carbohydrate heme group, an aromatic role in the stability of the hydroxyxanthone-heme complex, and the carbonyl group of the xanthone compound can bind with the heme iron so that the hydroxyxanthone-heme complex is formed [22]. The interaction with the free heme is most likely to contribute to the mechanism of detoxification of the heme. It is one of the known mechanisms of antimalarial action.

Amanat reported the activity of the hydroxyxanthone compound, which had a distinct number of hydroxyl clusters in the \textit{P. falciparum}, and identified that the 2-hydroxyxanthone compound had an antimalarial activity on \textit{P. falciparum} strain 3D-7 with IC\textsubscript{50} 20.69 \mu M, whereas IC\textsubscript{50} value of 2,7-dihydroxyxanthone was 1.342 \mu M [13]. The more the hydroxyl clusters on the xanthone framework, the...
Table 1: The percentage of Plasmodium growth inhibition of hydroxyxanthone derivatives and chloroquine on *P. falciparum* 3D-7 and FCR-3.

| Compound                        | Concentration (μM) | Growth inhibition of *P. falciparum* (%) |
|---------------------------------|--------------------|----------------------------------------|
|                                 | Strain 3D-7        | Strain FCR-3                            |
| 1,6,8-Trihydroxyxanthone (HX1) |                    |                                        |
|                                 | 102.38             | 75.864 ± 0.471                         |
|                                 | 51.19              | 69.033 ± 0.888                         |
|                                 | 25.59              | 64.726 ± 0.861                         |
|                                 | 12.8               | 61.479 ± 0.710                         |
|                                 | 6.39               | 47.405 ± 5.857                         |
| 1,6-Dihydroxyxanthone (HX2)    | 876.42             | 90.560 ± 0.665                         |
|                                 | 438.21             | 73.618 ± 0.713                         |
|                                 | 219.11             | 67.164 ± 0.601                         |
|                                 | 109.55             | 54.805 ± 2.433                         |
|                                 | 54.78              | 49.172 ± 1.606                         |
| 1,5,6-Trihydroxyxanthone (HX3) | 409.5              | 87.438 ± 1.154                         |
|                                 | 204.75             | 80.923 ± 0.679                         |
|                                 | 102.38             | 68.886 ± 0.151                         |
|                                 | 51.19              | 63.400 ± 0.421                         |
|                                 | 25.59              | 47.295 ± 0.611                         |
| 1-Hydroxy-5-chloroxanthone (HX4)| 810.86             | 74.002 ± 0.923                         |
|                                 | 405.43             | 64.513 ± 0.418                         |
|                                 | 202.71             | 57.136 ± 1.574                         |
|                                 | 101.36             | 50.963 ± 0.361                         |
|                                 | 50.68              | 46.473 ± 0.800                         |
| 1,6-Dihydroxy-5-methylxanthone (HX5)| 330.25              | 93.288 ± 0.956                         | 84.664 ± 0.902 |
|                                 | 165.13             | 75.364 ± 1.287                         | 73.462 ± 1.396 |
|                                 | 82.56              | 54.180 ± 1.574                         | 46.519 ± 0.816 |
|                                 | 41.28              | 45.041 ± 0.563                         | 41.140 ± 0.396 |
|                                 | 20.64              | 37.886 ± 0.441                         | 33.632 ± 0.327 |
| Chloroquine                     |                    |                                        |
|                                 | 0.037              | 97.766 ± 0.690                         | 84.189 ± 2.449 |
|                                 | 0.029              | 97.490 ± 0.517                         | 82.599 ± 2.945 |
|                                 | 0.023              | 96.581 ± 1.330                         | 79.923 ± 3.863 |
|                                 | 0.016              | 96.354 ± 1.131                         | 78.570 ± 4.507 |
|                                 | 0.008              | 54.415 ± 12.42                         | 18.748 ± 9.233 |

Table 2: The IC<sub>50</sub> values of hydroxyxanthone derivatives on *Plasmodium falciparum* 3D-7 and FCR-3 after 72 hours incubation period using microscopic method.

| Compound                        | Structure | IC<sub>50</sub> (μM) on *P. falciparum* |
|---------------------------------|-----------|----------------------------------------|
|                                 |           | 3D-7                                   | FCR-3                                  |
| 1,6,8-Trihydroxyxanthone (HX1)  | ![Structure](image1) | 6.10 ± 2.01                           | 6.76 ± 2.38                           |
| 1,6-Dihydroxyxanthone (HX2)     | ![Structure](image2) | 71.78 ± 0.31                           | 81.77 ± 5.78                           |
| 1,5,6-Trihydroxyxanthone (HX3)  | ![Structure](image3) | 27.64 ± 0.19                           | 64.09 ± 5.08                           |
better the antiplasmodial activity [23]. These results are in line with the results in this study that found HX1 and HX3 have lower IC50 values which showed that these compounds are more active than HX2, HX5, and HX4 which have fewer number of hydroxyl clusters. In addition to the number of hydroxyl groups, differences in antiplasmodial activity can be due to differences in substituent and position of the hydroxyl group in the hydroxynanthone derivatives. The HX5 compound has a methyl group, which is an electron donor. Still, the cluster’s electron donor strength was weaker than the hydroxyl group, and the HX4 compound was an electron puller, chlorosubstituent [14]. The study showed that the addition of methyl and chlorogroups to the carbon position of 5-hydroxyxanthone compounds had not been

| Compound                                | Structure | IC50 (μM) on P. falciparum 3D-7 | IC50 (μM) on P. falciparum FCR-3 |
|-----------------------------------------|-----------|---------------------------------|----------------------------------|
| 1-Hydroxy-5-chloroxanthone (HX4)        | ![Structure](image1) | 85.30 ± 0.87                    | 89.85 ± 7.69                     |
| 1,6-Dihydroxy-5-methylxanthone (HX5)   | ![Structure](image2) | 46.69 ± 0.29                    | 59.73 ± 0.78                     |
| Chloroquine                             | ![Structure](image3) | 0.01 ± 0.001                    | 0.11 ± 0.052                     |

**Figure 1**: The flow cytometric analysis diagram of compound HX1 on *Plasmodium falciparum* 3D-7 with 72 hours incubation. (a) Normal unstained erythrocytes. (b) Normal stained red erythrocytes (uninfected). (c) *P. falciparum*-infected erythrocytes control. (d) *P. falciparum*-infected erythrocytes with HX1 (concentration of 102.38 μM). (e) *P. falciparum*-infected erythrocytes with HX1 (concentration of 204.75 μM). The diagram was divided into four regions: Q1 (FITC/PE. ±); Q2 (FITC/PE. +/+); Q3 (FITC/PE. -/-); Q4 (FITC/PE. ±).
able to provide better activity shown in HX5 and HX4 compounds, which had a higher IC₅₀ value than HX2.

Previous study on the hydroxyxanthone compounds reported that xanthone compounds substituted hydroxyl clusters at carbon positions 2, 3, 4, 5, and 6 had in vitro potential of antiplasmodial [13]. The same study was reported by Ignatuschenko et al. [11], which found xanthone, with the addition of 1-hydroxyl group at position 4 or 5, and had moderate antiplasmodial activity. In contrast, xanthone with the addition of hydroxyl groups in both locations 4 and 5 have more active antiplasmodial activity. The data were in

### Table 3: The IC₅₀ value of 1,6,8-trihydroxyxanthone (HX1) compound and chloroquine on the formation of the β-hematin using method of HPIA.

| Compounds | Concentration (mM) | β-Hematin (mM) | Inhibition of β-hematin (%) | IC₅₀ (mM) |
|-----------|--------------------|----------------|-----------------------------|----------|
| HX1       | 20.475             | 9.684          | 91.171                      |          |
|           | 10.238             | 14.376         | 86.893                      |          |
|           | 5.119              | 54.448         | 50.359                      |          |
|           | 2.580              | 55.776         | 49.149                      |          |
|           | 1.269              | 72.500         | 33.901                      |          |
| Chloroquine | 9.692             | 12.500         | 88.604                      | 1.478    |
|           | 4.846              | 22.040         | 79.906                      |          |
|           | 2.423              | 47.845         | 56.379                      |          |
|           | 1.221              | 57.787         | 47.315                      |          |
|           | 0.601              | 77.787         | 29.080                      |          |

### Table 4: The IC₅₀ values of hydroxyxanthone compounds on Vero cell after 24 hours incubation using the MTT assay method.

| Compounds | Concentration (µM) | Vero cell growth inhibition (%) after 24 hours incubation | IC₅₀ (µM) |
|-----------|--------------------|----------------------------------------------------------|----------|
| HX1       | 2047.50            | 89.43 ± 1.90                                             | 3394.90 ± 435.44 |
|           | 1023.75            | 45.86 ± 5.51                                             |          |
|           | 511.88             | 27.96 ± 5.69                                             |          |
|           | 255.94             | 4.13 ± 8.21                                              |          |
| Chloroquine | 127.76             | 38.93                                                   | 308.32   |
|           | 63.88              | 31.75                                                   |          |
|           | 34.74              | 15.63                                                   |          |
|           | 20.475             | 67.03                                                   |          |
|           | 1023.75            | 59.75                                                   |          |
|           | 511.88             | 38.93                                                   |          |
|           | 255.94             | 31.75                                                   |          |
|           | 127.76             | 15.63                                                   |          |
|           | 63.88              | 20.70                                                   |          |
| HX3       | 2047.50            | 88.86                                                   | 224.45   |
|           | 1023.75            | 79.92                                                   |          |
|           | 511.88             | 68.04                                                   |          |
|           | 255.94             | 55.70                                                   |          |
|           | 127.76             | 39.58                                                   |          |
|           | 63.88              | 20.70                                                   |          |
| HX4       | 2047.50            | 89.96                                                   | 330.68   |
|           | 1023.75            | 79.92                                                   |          |
|           | 511.88             | 68.04                                                   |          |
|           | 255.94             | 55.70                                                   |          |
|           | 127.76             | 39.58                                                   |          |
|           | 63.88              | 20.70                                                   |          |
| HX5       | 2047.50            | 89.96                                                   | 417.57   |
|           | 1023.75            | 79.92                                                   |          |
|           | 511.88             | 68.04                                                   |          |
|           | 255.94             | 55.70                                                   |          |
|           | 127.76             | 39.58                                                   |          |
|           | 63.88              | 20.70                                                   |          |

### Table 5: The selectivity index (ratio IC₅₀ on Vero cell/IC₅₀ on plasmodium) of hydroxyxanthone compounds and chloroquine.

| Compounds | IC₅₀ on Vero (µM) | Selectivity index | IC₅₀ on plasmodium (µM) | Selectivity index |
|-----------|-------------------|-------------------|-------------------------|-------------------|
| HX1       | 3394.90           | 556.54            | 502.20                  | 556.54            |
| HX2       | 308.32            | 4.30              | 3.77                    | 4.30              |
| HX3       | 224.45            | 8.12              | 3.50                    | 8.12              |
| HX4       | 330.68            | 3.88              | 3.68                    | 3.88              |
| HX5       | 417.57            | 8.94              | 6.99                    | 8.94              |
| Chloroquine | 2216.92           | 221692            | 20153.82                | 221692            |
line with this study, i.e., the addition of a hydroxyl group in carbon position 6 in compounds HX1, HX3, HX2, and HX5 which had lower IC\textsubscript{50} value than the HX4 that has a hydroxyl group in the position of carbon 6.

4.2. Polymerization Inhibitory Activity of Hydroxyxanthone Derivative Compounds. When Plasmodium infects human erythrocytes, it brings hemoglobin from erythrocytes to the Plasmodium’s digestive vacuoles. Plasmodium has acid pH through the process of pinocytosis [24]. The hemoglobin is oxidized to methemoglobin, which is hydrolyzed into free heme and globin by the aspartic proteases (plasmepsin I, II, IV and histo-aspartic protease). Globin is hydrolyzed to peptides by cysteine proteases (falcipain) and zinc-containing metallopeptidase. Then, peptides are hydrolyzed to amino acids by cytosol exopeptidase, which will be used by Plasmodium for protein synthesis from the vacuole membranes to the Plasmodium cytoplasm [25, 26].

The result of the degradation of hemoglobin is that the free heme is toxic to the Plasmodium because it produces reactive oxygen species (ROS) and induces oxidative stress, which leads to cell lysis and the parasite’s death [27]. Therefore, Plasmodium resolves the toxicity of free heme with the heme detoxification system [28]. The primary heme detoxification system is the formation of hemozoin, which occurs in the Plasmodium’s digestive vacuole. In contrast, the secondary heme detoxification system is the formation of hematin, which is stable and nontoxic. The heme polymer is interconnected with the iron-carboxylase bond that connects the Ferrihematin to the other free heme passes through the Plasmodium’s digestive vacuole, then neutralized by GSH, H\textsubscript{2}O\textsubscript{2}, and heme-protein bond [29].

The free heme that is the heme monomer will be converted into the form of a polymer heme (hemozoin) by the enzyme of heme polymerase. Hemozoin (malaria pigment) is stable and nontoxic. The heme polymer is interconnected with the iron-carboxylase bond that connects the Ferri center of one heme with the side chain propionate of the other heme. Hemozoin formation is antimalarial’s target, such as chloroquine. The target activity is to inhibit the polymerization of heme [10, 30, 31].

The HX1 compound, which had the best antiplasmodial activity in this research, was tested in vitro for its heme polymerization inhibitory activity. The test involves a spontaneous hemozoin formation mechanism (β-hematin). Hematin was used in this assay because it has a similar structure to the free heme, whereas to mimic the model of the Plasmodium’s digestive vacuoles, which has acidic pH and initiates the reaction of the formation of β-hematin; glacial acetic acid was administered. In acidic environments, the hematin will polymerize into the crystal β-hematin [24].

The lesser the amount of β-hematin formed, the more inhibition of the polymerization of heme. This test suggested that the administration of HX1 can inhibit the formation of β-hematin. The higher the concentration of compounds, the greater the inhibitory percentage of β-hematin will be. The HX1 compound can inhibit the formation of β-hematin with an IC\textsubscript{50} value of 2.854 mM. Xanthone’s molecular characteristics make it insoluble in water, as well as having a high solubility to organic solvents. Nonpolluting characters cause most xanthone compounds to have some biological effects that can affect the membrane [32].

Chloroquine will diffuse the Plasmodium’s digestive vacuole membrane lipid bilayer. Inside the Plasmodium vacuole, chloroquine is protonated and cannot be diffused out so that chloroquine accumulates in the Plasmodium’s digestive vacuole. Chloroquine then binds to the heme forming a stable chloroquine-heme complex that will inhibit the polymerization of the heme resulting in disruption of the structure and damage of the digestive vacuole as well as the Plasmodium death. Consistent results are demonstrated from the activity structure profile of the 4,5-dihydroxynanthone (45X2) compound that has in vitro antiplasmodial activity [11, 24].

Spectroscopic nuclear magnetic resonance (NMR) research reported that the structure of hydroxyxanthone (45X2)-heme complex was formed from an interaction between the iron heme and the carbonyl group in the xanthone compound. Two aromatic systems contribute to the stability of the heme-45X2 complex along with the hydrogen bond between the hydroxyl clusters on the xanthone compound with the carbonyl groups in the heme. The bond can depict a 45X2 model with μ-oxo dimer hematin. The study expressed a 45X2 bond affinity with heme almost identical to the heme-antimalaria complex, such as chloroquine and quinine. Interaction stoichiometry 45X2-heme showed two units of heme bonded with one 45X2 molecule [22]. The data gave an overview of how the hydroxyxanthone compound can bind to the heme.

Earlier research conducted with H-NMR spectroscopy also showed the interaction of chloroquine with hematin. The study reported the interaction of the chloroquine molecule that binds four hematin molecules. Chloroquine binds mainly with the form of μ-oxodimerhematin. The structure and stability of the μ-oxodimerhematin obstruct the formation of hemozoin. Its quinoline derivatives that bond to hematin can stabilize hematin in the form of the μ-oxo dimer, thereby reducing the amount of hematin monomer to be combined into hemozoin and inhibit the polymerization of hematin [33]. However, there are aminoquinoline derivatives that interact with hematin but do not impede hematin polymerization. Thus, binding of compounds with hematin does not necessarily cause inhibition of hematin polymerization. Inhibitory polymerization of heme is not only determined from the ability of the compound in binding heme but its expertise in inhibiting heme polymer bonding with other heme [31]. The displacement of hematin can be caused by bonding with unstable hematin, such as a particular cluster that is at position 7 of the quinoline derivatives ring which has a firm bond with hematin. Still, when the group is moved at location six on a quinoline derivatives ring, the bond with hematin becomes unstable and cannot inhibit the polymerization of hematin [24].

Heme polymerization inhibitory activity (HPIA) assay showed that HX1 had low IC\textsubscript{50} (2.854 mM), as did
chloroquine (1.478 mM). Chloroquine had a potential antiplasmodial activity with low IC_{50} value on *Plasmodium* 3D-7 and FCR-3 (0.001 and 0.11 μM, respectively). The HX1 had less active antimalarial activity than chloroquine with IC_{50} value on *Plasmodium* 3D-7 and FCR-3 (6.10 and 6.76 μM, respectively). The data indicated there was a correlation between inhibitory polymerization of heme with antimalarial activity. However, these results are not necessarily linear. Ignatuschenko et al. stated that there were differences in the antimalarial activity and heme polymerization inhibitory activity obtained from the derivatives of xanthone, namely, pentametoxyxanthone and pentaacetxyxanthone. They were inactive in the heme polymerization inhibitory but had a potential or excellent antimalarial ability [11]. These data indicated that there was no direct correlation between heme polymerization inhibitory activity and antimalarial activity. Compounds that can inhibit the heme polymerization do not necessarily have good antiplasmodial activity. Compounds that have good antiplasmodial activity not necessarily inhibit the heme polymerization.

Some theories describe other mechanisms of action of antimalarial through the pathway of glycolysis. It inhibits the lactate dehydrogenase (LDH) enzyme and reduces adenosine triphosphate (ATP) production, which can cause a major death. Besides, the different mechanisms of action can be through DNA replication pathways by inhibiting the topoisomerase II enzyme, which will eventually lead to *Plasmodium* death [34, 35].

4.3. Cytotoxicity of Hydroxyxanthone Derivatives on Vero Cells. This cytotoxicity test was performed to determine the safety of hydroxyxanthone compounds on normal cells (Vero cells). The Vero cell culture which given the hydroxyxanthone compound indicates that the higher the concentration given, the higher the percentage of growth inhibition of Vero cells and becomes more toxic to the cells depending on the concentration.

Cytotoxicity of compounds HX1, HX2, HX3, HX4, and HX5 on the Vero cell was shown from the IC_{50} values (3283.72; 389.66; 230.28; 334.90; 458.94 μM or 801.88; 88.92; 56.23; 82.60; 111.17 μg/mL, respectively). The five hydroxyxanthone compounds have weak cytotoxic effects (IC_{50} ≥ 30 μg/mL) on the Vero cell [36]. The same results were presented in the research conducted by Fatmasari [14], showing weak cytotoxicity of compounds HX1, HX2, HX3, HX4, and HX5 on MCF-7, WIDR, and Hela cell lines.

*Insilico* interactions between the HX1 compound with the topoisomerase enzyme II have been conducted through the molecular docking process. The process of molecular docking involves the prediction of conformation and orientation of the ligands with binding sites (binding site) of the target protein. Compound or ligand serves as a complementary pair of proteins, while protein acts as a receptor. Mitoxantrone (MIX) is a ligand that has a basic frame similar to xanthone found in the enzyme topoisomerase II. Also, both have a match of shape and volume on the active side then interaction ligand and protein occur. The lower the energy value, the more stable the ligand so that the interaction produced with protein is stronger. Thus, the presence of a variety of strong interactions can cause the performance of the topoisomerase II enzyme. The enzyme is found in viable cells and serves to improve the structure of damaged DNA. When the enzyme function is interrupted, the process of replication and transcription of DNA into RNA cannot take place so that the cell will experience apoptosis. The process is also one of the mechanisms of action from antimalarials, such as chloroquine.

The result of the molecular docking indicates the interaction of the DNA chains, namely, hydrogen interactions (DG C:13 and pi-pistacked (DA f:12 and DC c:8), with the binding affinity value −8.0 kcal/mol. However, the *in vitro* cytotoxicity test showing HX1 had a weak cytotoxic activity on MCF-7, WIDR, and Hela cell lines with IC_{50} 45; 61.9; 67.7 μg/mL. These results demonstrated that the mechanism of action of the HX1 compound might also work in inhibition of the topoisomerase enzyme II. To determine the presence of the bond with DNA *in vitro* can be tested with DNA-methyl green assay with spectrophotometry [37].

The selectivity of a compound is crucial in the development of new drugs. The compound is said to be selective if the selectivity index is more than 10 and has a good antimalarial effect, but is not toxic to the host. The selectivity index was obtained from the ratio between (IC_{50} of cytotoxic effect on *Vero* cells) and (IC_{50} of antimalarial activity on *P. falciparum* 3D-7 and FCR-3). The HX1 has an index of selectivity >10, which means it is safe for the host so that it is potentially to be developed as an antimalarial.

5. Conclusions
Among 5-hydroxyxanthone derivatives tested, the 1,6,8-trihydroxyxantone showed the best antimalarial activity and has heme polymerization inhibition activity and high selectivity.

Data Availability
The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest
The authors declare that there are no conflicts of interest regarding the publication.

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