Investigation of Chlorophyl-a Derived Compounds as Photosensitizer for Photodynamic Inactivation

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Abstract

Chlorophyll has unique physicochemical properties which makes them good as photosensitizer of Photodynamic Inactivation (PDI). The physicochemical properties of chlorophyll as photosensitizer can be optimized through several routes. One of the possible routes is by replacing the metal ion center of chlorophyll with other ions. In this research, the effect of coordinated metal ion in the natural chlorophyll-a was studied for bacterial growth (S. aureus) inhibition. The replacement of metal in the center of chlorophyll hopefully can improve the intensity of Intersystem Crossing Mechanism (ISC) lead to the formation of singlet oxygen species. The chlorophyll a and b were isolated from spinach via precipitation technique using 1,4 dioxane and water. The chlorophyll a and b were separated using sucrose column chromatography. The thin layer chromatography result showed that chlorophyll a (Rf: 0.57) had been well separated with chlorophyll b (Rf: 0.408). The absorption spectra of chlorophyll a and b showed that the Soret band was observed at 411 and 425 nm, while the Q band appeared at 663 and 659 nm. Replacement of metal ion center shifted the Soret band of chlorophyll a derivatives to lower energy region, while Q-band was slightly shifted to the higher energy region. The absorption and the fluorescence intensity were also observed decreasing after ion replacement. The Inhibition activity investigation over S. aureus showed the highest inhibition activity was exhibited by Zn-pheophytin-a (66.8%) followed by chlorophyll a (30.1 %) and Cu-pheophytin-a (0%). The inhibition activity is correlated with decreasing fluorescence intensity. The formation of singlet oxygen by ISC mechanism is hypothesized to deactivate the excitation state of Cu-pheophytin-a.

Keywords: Chlorophyll a; Pheophytin; Photosensitizer; Photodynamic inactivation; Sucrose

1. Introduction

Antibiotics are used to control, treat, or prevent bacterial infection, however, in the recent year, anti-bacterial resistance phenomenon is becoming more of a problem in combating the infectious disease caused by bacteria [1]. To restraint this emerging public health problem, it is important to develop a non-toxic treatment that acts more effectively than current antibiotic treatment [2–4]. One of these effective methods is Photodynamic therapy for bacterial inactivation or widely known as Photodynamic Inactivation (PDI) [5,6]. In addition, PDI was reported to exhibit a fast and effective approach to inactivate multi-resistant bacteria. The in vitro studies showed the possibility of bacterial
reduction up to $6-\log_{10^6}$ CFU within seconds during the incubation and irradiation [7].

The bacterial treatment using PDI is a treatment that use the light-sensitive material, photosensitizer (PS) along with visible or ultraviolet light to produce singlet oxygen ($^1\text{O}_2$) and radical species that induces phototoxic damage to the bacterial cell immediately during the irradiation [8,9]. The toxic singlet oxygen can be produced when the PS is irradiated with the appropriate light with a particular wavelength (visible or UV light) that led to the excitation of PS ground state. The excited single state is unstable and lives for less than 1 µs. It can return back to the ground state via emission of a secondary photon in the form of fluorescence or it can undergo intersystem crossing into the excited triple-state to produce singlet oxygen species [10]. The PS triple-state can produce the chemical changes in the bacterial cell via two competitive pathways (type 1 and type II) [2]. Type I reaction produces radical ions that can pass quickly through cell membranes causing the detrimental damage to the bacteria cell [11]. Type II reaction is characterized by the presence of appropriate oxygen concentration. The triple state of PS can transmit its energy directly to the triplet ground state of molecular Oxygen found mostly in the cells [12]. The transmitted energy gained by triple state oxygen molecule ($^3\text{O}_2$) excites the ground state of $^3\text{O}_2$ to produce singlet oxygen ($^1\text{O}_2$) [13,14]. PDI relies on the production of $^1\text{O}_2$ by type II reaction as the predominant cytotoxic ROS (Reactive Oxygen Species) that has significant lethal effect [15].

The PDI technique exhibits some advantages for the treatment of microbial infections such as a broad spectrum of actions compared to the antibiotics, the efficient inactivation of antibiotic-resistant strains, the low mutagenic potential, and the absence of selection of the photo-resistant microbial cells [9,16]. Kasf er and Hamblin [17] mentioned that the development of resistance to PDI treatment is unlikely due to the short duration of treatment and the photo-generated ROS could attack multiple cellular targets. The tolerance enhancement of bacteria to PDI is considered caused by the induction of genes responsible for defense against oxidative stress [8]. Furthermore, the absence of specific defense systems against singlet oxygen in the organism is also explained for the reason why PDI is less possible to develop resistance strain [18]. Another study [19] proposed that repeated exposure of E. coli and S. aureus to ZnTnHex-2-Pyp (photosensitizer), did not change the tolerance of the microbes for the PS.

The PDI efficacy depends on the photosensitizer (PS) material. The most appropriate photosensitizer is characterized as positively-charged, water-soluble and photo-stable [7]. Many PS have been introduced for the treatment ranging from synthetic PS to naturally available PS. One class of PS attracting many researchers is naturally occurring porphyrin-type compound. The porphyrin nucleus contains 22 π electrons, with 18 π electrons delocalized the macrocycle. Since their aromatic properties, porphyrins commonly participate in electrophilic substitution reaction at the meso positions, which possess the most electron density, as such, are also the most reactive [2]. As the compliance of the delocalized electrons, porphyrins show have very intense absorption bands in the visible region which is favorable properties as PS for PDI. Porphyrins possess a good amphiphilicity and ability for numerous chemical modifications [20]. Porphyrin molecules were reported to produce high quantum yield of $^1\text{O}_2$ production and high one-photon absorption coefficient ($\approx 500,000$ M$^{-1}$cm$^{-1}$) [21]. Moreover, porphyrins were reported to possess a high binding affinity to cellular components, membranes, proteins, and DNA [22,23].

One of the prominent natural porphyrin-type molecules is chlorophyll. The most abundant and widely distributed green pigment in green plants, chlorophyll, has been studied in decades for its unique physicochemical properties [24,25]. Moreover, The wide range of absorption spectrum of chlorophyll and chlorophyll derivative (CD) compounds, high absorption spectra at the range of 400–800 nm, high rate of ROS production and easy chemical modification make chlorophyll appropriate to be investigated for the next potential PS for PDI [26,27]. Chemical modifications either by ion center replacement or hydrophobic group (phyt yl group) substitution were reported to generate the chlorophyll derivatives that have more beneficial properties for PDI photosensitizer [25].

In this study, the potency of natural chlorophyll-a and its derivatives will be evaluated as PS for photodynamic Inactivation (PDI). The Mg$^{2+}$ ion in the center of the chlorophyll (Figure 1) will be replaced with Zn$^{2+}$ and Cu$^{2+}$ and the antimicrobial effect of those CDs will be evaluated. Ion Cu and Zn were chosen in this experiment since the ion size of both ions is similar enough with Mg$^{2+}$. In addition, the chemical stability of the porphyrin-metal ion.
complex is also considered for ion replacement [25]. Moreover, the previous study reported Zn-porphyrin complex generated more \(^1\)O\(_2\) compare to its parent compound [28,29]. The unusual chlorophyll-a isolation was also introduced here, we applied sucrose column chromatograph instead of silica for the stationary phase [30]. This study hopefully can be a preliminary study for exploring the potency of natural chlorophyll for photosensitizer of Photodynamic Inactivation.

2. Materials and Methods

2.1 Extraction and Isolation of Chlorophyll a

The commercially available green spinach (Amarantus tricolor L.) was immersed in cold acetone (C\(_6\)H\(_4\)O) (Merck) and was mashed up by the mechanic blender creating the green slurry suspension. The crude chlorophyll was obtained by precipitation of spinach residue with distilled water (1/4 from the total volume) and 1,4-dioxiane (1/8 from the total volume) (Merck) and let to settle for 7 days at \(-20^\circ\)C. This precipitation process was repeated 2 times. The precipitated material then was separated and diluted with acetone. The crude extract was obtained by removing the solvent with reduced pressure evaporation.

The chlorophyll a was isolated by column chromatography using manually powdered sucrose C\(_{12}\)H\(_{22}\)O\(_{11}\) (commercial sucrose) as a stationary phase. The mobile phase system was used petroleum ether C\(_{6}\)H\(_{14}\) (Merck) followed by 10% diethyl ether (C\(_4\)H\(_{10}\)O) (Merck) in petroleum ether. The collected fraction was separated from the solvent by reduce pressure evaporation.

![Figure 1](image)

Figure 1. The chemical structure of chlorophyll a and b which are different by methyl and aldehyde. Pheophytin-a is chlorophyll structure without Mg\(^{2+}\) in the center and just replaced by 2 hydrogen. (Adopted from [31]).

2.2 Zn-pheophytin-a (Zn-Pa) Synthesis

Isolated chlorophyll-a (Cla) (200 mg) was diluted with acetone (150 mL) and mixed gently with 10 mL of Hydrochloric acid (HCl 37%, Merck) 0.1 M. The mixture was mixed for 30 minutes at 40 °C to form Mg\(^{2+}\)-free chlorophyll-a, pheophytin-a. Hexane (Merck) was used to extract pheophytin-a from the mixture. Collect-
ed pheophytin-a (20 mg) diluted in chloroform (CHCl\(_3\), Merck) was mixed with Zinc acetate dihydrate ((ZnCH\(_3\)CO\(_2\))\(_2\)2H\(_2\)O, Merck) in methanol. The mixture was stirred at 40 °C under nitrogen atmosphere for 1 h. After removing chloroform-methanol solvent, the residue was extracted with hexane (C\(_6\)H\(_{14}\), Merck) and the extract was washed 2 times with distilled water. The hexane was removed by rotary evaporator at 40 °C to obtain Zn-Pa [32]. The purification of Zn-Pa was used TLC preparative method using silica plates (TLC Silica Gel 60 F\(_{254}\), Merck) and hexane:acetone 7:3 as a mobile phase.

2.3 Cu-Pheophytin-a (Cu-Pa) Synthesis

The Cu-pheophytin-a was prepared by inserting the Cu (II) to pheophytin-a. The preparation of pheophytin-a was similar as described above. Pheophytin-a was added with copper chloride dihydrate (CuCl\(_2\)2H\(_2\)O, Merck) and acetate buffer pH 5. The reaction was conducted by reflux for 3 h at the room temperature under nitrogen atmosphere. The reaction generated two separated phase. The bottom layer was collected and the solvent was removed by rotary evaporatory. The purifica-
tion was also conducted by preparative TLC as described above.

2.4 Photo-physical Properties Characterization

The absorption spectra of CDs was measured by single beam visible and UV/Visible Spectrophotometer Jenway 6305 and sample was diluted in chloroform. The absorption measurement was conducted from 350–700 nm. The fluorescence spectra was investigated at the wavelength (\(\lambda\)) 430–800 nm using Spectrofluorophotometer RF-5301 (Shimadzu Eu-

2.5 Bioactivity Investigation of CDs

Bioactivity investigation was carried out based on the method described [33] using S. aureus culture in Nutrient Broth (NB) medium.
containing peptone (NB for microbiology, Merck), yeast extract, sodium chloride (NaCl ≥99%, Merck). In brief, from a calibrated inoculum at 10^8 CFU/mL, a volume of 50 µL was transferred to a test tube containing 950 µL of CDs at concentration 40 µg/L diluted with aqueous Tween® 80 (Polisorbat, Merck) 1% (wt/vol). Aliquots of 500 µL of each suspension were transferred to the test tube for irradiation, and the remaining 500 µL sample was used as non-irradiated sample. The irradiation process was conducted using Natrium lamp (λ: 650 nm) using intensity 2000 lux/cm^2 for 30 minutes. During irradiation, oxygen was added to the samples. After irradiation, each sample was adjusted with NB to create 4×10^4 CFU/mL of bacteria. The diluted inoculum (50 µL) was plated in Nutrient Agar (NA) for 24 h for incubation at 37 °C, and the CFU number was counted. The same experiment was carried out for control groups using the same amount of inoculum and 1.0% aqueous Tween 80 solution without CDs. The control group without any treatment was considered as 0% of death.

3. Results and Discussions

The green spinach was chosen due to its abundant, easy to get and relatively low price natural sources. The cold precipitation technique by addition of 1,4 dioxane and water is to optimize the chlorophyll extraction by formation of chlorophyll-dioxane complex and to eliminate the unnecessary pigments and plant materials. Chlorophyll a, b and β-carotene are known as major pigments from green spinach leaves [34]. The chlorophyll a (Cla) was chosen for future treatment since its higher availability in the natural sources compared to chlorophyll b (Clb) and also Cla is reported to be more photostable. The yield of the crude extract from green spinach was 1.4%.

The supersensitive column chromatography with sucrose as stationary phase was performed. It was conducted since slight polarity difference between Cla and Clb. Sucrose column chromatography was reported successful to separate chlorophyll a from other pigments.

![Figure 2: The absorption spectra of Cla (chlorophyll-a) and Clb (chlorophyll-b) in chloroform.](image)

![Figure 3. Synthesis route of Zn-Pa and Cu-Pa.](image)
from spinach extract [35]. The elution order was applied according to the polarity of the target compounds. β-carotene is the most non-polar compound, β-carotene in yellow color came out first from the column followed by Cla (dark green) and the last pigment was Clb (light-green). The appearance of one spot after TLC observation indicated the purity of the isolated compounds. The retention factor (R) is 0.571 and 0.408 for Cla and Clb, respectively. Figure 2 described the absorption spectra of isolated Cla and Clb.

Cla exhibited the maxima peak at 411 nm (Soret band) and 663 nm (Q band), while for Clb, the maxima peak appeared at 423 nm (Soret band) and 659 nm (Q band). The difference maxima peak showed that Cla and Clb were separated. Cla and Clb showed good absorption ability in the 400–800 nm. This physicochemical property is an essential characteristic for PS. At the same concentration, Cla exhibited higher absorption intensity that Clb, as one of the ideal PS requirements is the high absorption ability in the visible wavelength. According to the absorption analysis, Cla was showed more beneficial properties compared to Clb, thus in this study, the modified chlorophyll compounds are derived from Cla.

The replacement of metal ion center of Cla was purposed to improve the physicochemical properties and chemical stability of Cla. Hydrochloric acid addition to Cla releases Mg²⁺ from the center and turned to pheophytin-a as pictured in Figure 3. Metal insertion finally was proceed under inert reaction (N₂ atmosphere) by addition of Zu and Cu ion in their salt form.

The physicochemical properties of CDs were evaluated by absorption and fluorescence spectra. The electronic absorption spectra pattern of CDs as depicted in Figure 4 were almost similar to the chlorophyll a. That indicates the presence of the same chromophore on CDs. The ion replacement gives an impact to their absorption intensity. As shown in the Figure 3, the absorption intensity of Cla was the highest among other CDs and then followed by Cu-Pa, Zn-Pa, Clb and pheophytin-a (pheo-a). The decrement of absorption intensity might be related with the photobleaching of the CDs [17,30].

The replacement of Mg²⁺ from Cla obviously shifted the Soret peak to the lower energy area, while Q maxima peak was observed to shift up to the higher energy as written in the Table 1. The absence of the ion center in the chlorophyll (pheo-a) shifted the Soret peak to 421 nm. Cu-Pa and Zn-Pa exhibited the Soret peak at 414 nm and 420 nm, respectively. The paramagnetic properties of Cu might change the electron delocalization on the porphyrin, so thus the Soret peak of Cu-Pa comes to a higher energy than Zn-Pa. The Q peak of Pheo-a, Zn-Pa, and Cu-Pa were observed at 664 nm, 653 nm and 649 nm, respectively.

\[ \text{Table 1. Absorption and fluorescence peak data for Cla, Clb and Cla Derivatives in chloroform.} \]

|       | Absorption       | Fluorescence peak (nm) |
|-------|------------------|------------------------|
|       | Soret peak (nm)  | Q peak (nm)            |
| Cla   | 411              | 663                    | 672                     |
| Clb   | 423              | 659                    | 669                     |
| Pheo-a| 421              | 664                    | 673                     |
| Cu-Pa | 414              | 649                    | 667                     |
| Zn-Pa | 420              | 653                    | 662                     |

Figure 4. Absorption spectra of Cla, Clb, and all CDs in chloroform.

Figure 5. Absorption spectra of Cla, Clb, and all CDs in methanol.
The fluorescence peaks of Cla, Clb, and all CDs exhibited in the range of 660–673 nm (Table 1). It was observed that the position of fluorescence peaks is slightly shifted. According to Figure 5, the Cla and Clb exhibited a higher fluorescence intensity than other CDs. The low fluorescence intensity of other CDs might be due to the Intersystem Crossing reaction (ISC) from their excited state. The ISC reaction that happened in the CDs hopefully can induce the formation of \( 1^O_2 \).

The synthesized CDs were treated to the bacterial suspension to understand the effect of the sample for bacterial inhibition. In this study, the non-pathogenic Gram-positive bacteria, \( S. aureus \) was used as the testing bacteria. The tested compounds+bacteria colony were irradiated with short wavelength energy (sodium lamp) at 650 nm since it has low level of harm to the human tissue and also considerably low in price. During the treatment, oxygen was supplied since the type II ISC requires enough \( O_2 \) to optimize the \( 1^O_2 \) formation. The dark toxicity was measured to understand the potency of the tested samples to inhibit the bacterial growth without irradiation. Clb was not included in bioactivity test due to lack of amount and photostability issue. Tween 80 1% was used as a surfactant to help immobilization of CDs to the bacterial.

The effect of CDs on bacterial growth was described in the Figure 6. All of CDs showed a dark toxicity potency as shown the bacterial inhibition even without irradiation. The number of bacteria for Cla and Zn-Pa without irradiation was observed at \( 1.02 \times 10^7 \) and \( 1 \times 10^7 \) (CFU/mL), respectively. After irradiation, the number of bacteria decreased to \( 9 \times 10^6 \) and the Zn-Pa presented the promising effect by inhibiting the bacterial growth as the number of bacteria becomes \( 4.3 \times 10^6 \) after irradiation. That result indicates the potency of Zn-Pa as a photosensitizer for PDI.

There is no significant effect showed by Cla for irradiated and non-irradiated photosensitizer as shown in the Figure 7. Inhibition activity against of \( S. aureus \) for irradiated Cla was 30%, just slightly higher from non-irradiated Cla which exhibited 27% of inhibition activity. Effect of irradiation on inhibition of \( S. aureus \) was shown by Zn-Pa. Irradiated Zn-Pa was able to suppress 66.8% of bacterial growth while non-irradiated Zn-Pa just showed 28% of growth inhibition. The ability of Cla and Zn-Pa to inhibit \( S. aureus \) growth even in dark conditions showed their dark toxicity potency of both compounds. Cu-Pa didn’t show any inhibition at the radiated and non-radiated condition.

Figure 7 described the inhibition rate of \( S. aureus \) after treated with CDs with irradiation and without irradiation. The unique behavior was observed in Cu-Pa, as it does not show any toxicity even and after radiation. The low fluorescence intensity of Cu-Pa was contradictory with its bioactivity. Non-toxic properties of Cu-Pa might be because of copper due to its paramagnetism that favors the deactivation of the excited states, decreasing the production \( 1^O_2 \) [25]. Moreover, the absence of dark toxicity of Cu-Pa might be due to low microbial uptake. Otherwise, these hypotheses need to be confirmed by further analysis such as the quantum yield of \( 1^O_2 \) and cellular uptake measurement.

The finding of this study indicated that the modification of the metal ion center of porphyrin-based compounds influences the physicochemical properties of CDs, while maintaining or even improve their biological activity. A study by Rahimi et al. [36] stated zinc porphyrin complex (ZnTNPP) showed the improvement of photostability properties while main-
taining the inhibition activity of \( P. \ aeruginosa \). Zoltan et al. [37] described improvement of the biological activity of metal-based porphyrin compared to the metal-free porphyrin because the role of metal ion to the primordial factor to interfere the spin-orbital coupling as the consequence of the multiple polarisable electron shells and a high nuclear charge density. The change in the sin-orbital coupling thus affects the many physicochemical properties such as, absorption, and fluorescence intensity, energy transfer at the triplet state, and production of singlet oxygen species. That study also showed an efficient inactivation of \( E. \ coli \) when it was treated by meso-tetra(pyren-1-yl)porphyrin complexes of Ni(II), Cu(II), and Zn [37].

Even though this study showed a promising photosensitizer candidate among the tested CDs, the further advanced studies are still necessary to be conducted in the future. This will cover several advanced measurements such as are \( O_2 \) quantification, photobleaching analysis, and microbial uptake that provide a comprehensive understanding of PDI mechanism in inhibiting the bacterial growth. Moreover, the targeted delivery of the photosensitizer is also needed to be concerned for the proper accumulation of the photosensitizer complexes in the target infected cells. Proper accumulation in the infected cells is important in order to gain effective treatment without causing any harm to the healthy cells. The targeted delivery of photosensitizer could be accomplished by conjugation with antibodies, synthesis of molecules with a specific structure or attached with the magnetic nanoparticles [2].

4. Conclusions

Chlorophyll-a was isolated successfully isolated from raw spinach leaves by the formation of complex precipitate between chlorophyll and 1,4 dioxane then followed by the sucrose gravity column. Natural chlorophyll-a derivatives (CDs) were obtained by introducing the \( Zn^{2+} \) and \( Cu^{2+} \) ions into the phaeophytin-a porphyrin ring. Absorption and fluorescence spectra of CDs exhibited favorable physicochemical properties as photosensitizer. Zn-Pa displayed a promising candidate as a photosensitizer for PDI as it reduced the number of bacteria by 66.8%. Cla showed 30% bacterial inhibition, while Cu-Pa did not show any bacterial inhibition activity. Inactivity of Cu-Pa to inhibit the bacterial growth was might be due to the deactivation of the excited states. This study hopefully can bring new insight for developing new and effective PS from naturally occurring compounds and their derivatives. Although, some results in this study showed promising result, more advanced studies are necessary to have a better understanding of PDI mechanism in inhibiting bacterial growth. The targeted delivery scheme also is important to be noted for gaining effective and safe PDI treatment.

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