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<td>Author(s)</td>
<td>Ikeda, Atsushi; Satake, Shuhei; Mae, Tomoya; Ueda, Masafumi; Sugikawa, Kouta; Shigeto, Hajime; Funabashi, Hisakage; Kuroda, Akio</td>
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<tr>
<td>Citation</td>
<td>ACS Medicinal Chemistry Letters, 8 (5) : 555 - 559</td>
</tr>
<tr>
<td>Issue Date</td>
<td>2017-05-11</td>
</tr>
<tr>
<td>DOI</td>
<td>10.1021/acsmedchemlett.7b00098</td>
</tr>
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<td>Self DOI</td>
<td></td>
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<tr>
<td>URL</td>
<td><a href="http://ir.lib.hiroshima-u.ac.jp/00046209">http://ir.lib.hiroshima-u.ac.jp/00046209</a></td>
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| Relation | |
Photodynamic Activities of Porphyrin Derivative–Cyclodextrin Complexes by Photoirradiation

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KEYWORDS: porphyrin, cyclodextrin, photodynamic therapy, photosensitizer

ABSTRACT: Water-soluble cyclodextrin (CyD)-complexed with porphyrin derivatives with different substituents in the meso-positions showed different photodynamic activities toward cancer cells under illumination at wavelengths over 600 nm, the most suitable wavelengths for photodynamic therapy (PDT). In particular, aniline- and phenol-substituted derivatives had high photodynamic activity because of the efficient intracellular uptake of the complexes by tumor cells. These complexes showed greater photodynamic activity than photos, currently the main drug in clinical use as a photosensitizer. These results represent a significant step towards the optimization of porphyrin derivatives as photosensitizers.

Porphyrins have been widely investigated as potential photosensitizers (PSs).1–3 During photodynamic therapy (PDT), an intravenously administered PS is allowed to accumulate in a tumor. Selective light-irradiation is then used to produce reactive oxygen species (ROS) by photochemical reactions between photoexcited PS and molecular oxygen.4 Therefore, water-solubilization of PSs is very important for biological applications such as PDT. Water-soluble porphyrins and their derivatives have been synthesized by introduction of hydrophilic moieties.1–3 However, the introduction of the hydrophilic moieties frequently results in insufficient water-soluble porphyrin derivatives due to self-aggregation of the compounds. Therefore, these porphyrin derivatives are commonly injected into cells or the body as an aqueous solution with dimethyl sulfoxide or ethanol solutions, both of which are highly cytotoxic.5–7 An alternative approach is to use solubilizing agents such as water-soluble polymers,8 cyclodextrins,9–11 or liposomes.12 The complexes of the porphyrin derivatives with cyclodextrins are especially easy to prepare with a mechanochemical high-speed vibration milling apparatus.12–14 In this paper, we compare the photodynamic activities of porphyrin derivatives with different substituents in meso-positions, complexed with trimethyl-β-cyclodextrin (TMe-β-CyD; Figure 1). Photodynamic activities were evaluated with HeLa cells through irradiation with visible-light at long wavelengths (610–740 nm), the optimal wavelengths for PDT.

Complexes of porphyrin derivatives (1–6) with TMe-β-CyD (Figure 1) were formed as previously described.12–14 The formation of these materials was confirmed by UV-vis absorption and 1H NMR spectroscopy (Figures 2, S1, and S2). These spectra indicated that compounds 1–6 all formed complexes with two TMe-β-CyDs. They were all soluble in water and stable at room temperature (Figure 1).

Figure 1. Compound structures and schematic illustration of the TMe-β-CyD-complexed with porphyrin derivatives.

We first investigated the levels and kinds of generated cytotoxic ROS and the stability of the TMe-β-CyD-complexed with porphyrin derivatives under visible-light irradiation at a wavelength greater than 620 nm. Photoexcited porphyrins transfer an electron to molecular oxygen (O2) to give superoxide anions (O2−) (electron-transfer pathway Type I), and transfer energy to O2 to give singlet oxygen...
molecules (\( {\text{O}_2} \)) (energy-transfer pathway Type II). To identify the reactive oxygen species, the presence of \( {\text{O}_2} \) and \( {\text{O}_2}^{•–} \) was analyzed with 9,10-anthracenediyldis(methylene)dimalonic acid (ABDA) and nitroblue tetrazolium (NBT), respectively. Conversion to an endoperoxide from ABDA by reaction with \( {\text{O}_2} \) leads to a decrease in absorbance at 380 nm (Figures S3 and S4). Generation of formazan, reduced NBT by \( {\text{O}_2}^{•–} \), is observed as an increase in absorbance at 560 nm (Figure S5). The decrease of ABDA was plotted as a function of photoproduction time of the complexes in aqueous solutions. The TMe-β-CyD-complexed with 1, 5, and 6 precipitated out not only after vigorous bubbling of oxygen but also after photolysis, suggesting that these complexes were destabilized by photoactivation (Figures S3a, S4b, and S4c). As shown by their precipitation with oxygen gas bubbling and photolysis, these complexes are unstable due to the effect of the substituents in the meso-positions. Therefore, compound 2 was employed as a neutral guest molecule in the place of 1. The TMe-β-CyD-complexed with 2 was stable in water after photolysis. Cationic 3 and anionic 4 were also stable as complexes with TMe-β-CyD after photolysis. The absorption level of ABDA at 380 nm (the absorption maximum of ABDA) was plotted as a function of photolysis time of the TMe-β-CyD-complexed with 2-4 (15 \( \mu \text{M} \)) in \( {\text{O}_2} \)-saturated aqueous solutions (Figure 3a). The results show that these TMe-β-CyD complexes generated \( {\text{O}_2} \) in the order of \( \text{3} \approx \text{4} \approx \text{2} \); however, the differences were very small. The difference between 2 and 3 in the photoproduction abilities of \( {\text{O}_2} \) simply reflects the amount of light absorption over 620 nm (Figure 2 inset, black and red lines). However, 4 displayed relatively high \( {\text{O}_2} \) photoproduction, despite the low absorption over 620 nm (Figure 2 inset, blue line).

**Figure 2.** UV-vis absorption spectra of the TMe-β-CyD-complexed with 2 (black line), the TMe-β-CyD-complexed with 3 (red line), and TMe-β-CyD-complexed with 4 (blue line; [TMe-β-CyD-complexed with 2-4] = 30 \( \mu \text{M} \)). All absorption spectra were measured in H₂O at 25 °C (1 mm cell).

In contrast, reduction of NBT by \( {\text{O}_2}^{•–} \) could not be detected in the photolysed TMe-β-CyD-complexed with 2-4, even though formazan was readily detected in the positive control sample TMe-β-CyD-complexed with 2 in the presence of NADH (Figure 3b). These results suggest that ROS produced by the TMe-β-CyD-complexed with porphyrin derivatives of 2-4 are mostly \( {\text{O}_2} \) generated by a Type II reaction.

Next the photodynamic activities of the TMe-β-CyD-complexed with porphyrin derivatives were evaluated in vitro on HeLa (human cervical carcinoma) cells. The cells were incubated with 0.02–0.4 \( \mu \text{M} \) of the TMe-β-CyD-complexed with porphyrin derivatives for 24 h before photolysis with wavelengths greater than 610 nm (610–740 nm) for 30 min. Photolysis was carried out under 9 mW cm⁻² light power at the cell level. The cell survival rate was determined using a WST-8 assay with a Cell Counting Kit-8. The
results are summarized in Figure 4. Under dark conditions, the TMe-β-CyD-complexed with porphyrin derivatives of 2–4 show no cytotoxicity to the HeLa cells (Figure 4a). The TMe-β-CyD that remained in the HeLa cells did not influence the results because the toxicities of β-CyD and their derivatives followed the general trend DMe-β-CyD >> β-CyD > TMe-β-CyD. Furthermore, the TMe-β-CyD-complexed with 4 showed no photodynamic activity. In contrast, the TMe-β-CyD-complexed with 2 and 3 showed photodynamic activities in a dose-dependent manner, with half maximal inhibitory concentration (IC50) values of 0.15 and 0.08 μM, respectively (Figure 4b). The TMe-β-CyD-complexed with 4 had a very low photodynamic activity in spite of a similar structure to TMe-β-CyD-complexed with 2. Because the TMe-β-CyD-complexed with 2 and 4 generate a similar amount of 1O2, we suggest the phenol moieties of 2 play a key role in intracellular uptake of the complex by HeLa cells. Detail of this intracellular uptake is described below. In these TMe-β-CyD complexes, the order of the photodynamic activity was 3 >> 2 >> 4. The IC50 values of the TMe-β-CyD-complexed with 2 and 3 are lower than that of photofrin, presently used as the main drug in clinical photosensitizers. Here, the molecular numbers of photofrin were converted to moles of porphyrin units because photofrin consists of hematoporphyrin esters and others containing monomers, dimers and oligomers. Figure 4b shows IC50 value of 2.1 μM for photofrin under the same conditions as described above. The TMe-β-CyD-complexed with 2 and 3 had approximately 14 and 26 times higher photodynamic activities than photofrin, respectively.

The ROS in HeLa cells were identified through analyses of the inhibitory effect of L-histidine as a 1O2 scavenger and D-mannitol as a hydroxyl radical scavenger, respectively. Figure 5 shows that the photocytotoxicity of the TMe-β-CyD-complexed with 2 and 3 was effectively inhibited by L-histidine but was only weakly inhibited by D-mannitol. These results indicate that 1O2 played a major role in the photodynamic activity of the TMe-β-CyD-complexed with 2 and 3 against HeLa cells.
Figure 5. Inhibitory effect in the presence of ROS scavengers on the photodynamic activities of the TMe-β-CyD-complexed with 2 and 3. The photodynamic activities of the TMe-β-CyD-complexed with 2 and 3 (0.8 μM) were measured in the absence (gray bar) and presence of 50 mM L-histidine (red bar) or 50 mM D-mannitol (blue bar). Error bars represent the mean ± standard deviation (SD) for n = 3.

The intracellular uptake by HeLa cells of the TMe-β-CyD complexed with 2–4 was observed with fluorescence microscopy at the emission wavelength of the porphyrin derivatives. As shown in Figure S6, all the TMe-β-CyD complexed with 2–4 possessed fluorescence peaks around 650 nm when they were excited at the wavelength (λex = 540 nm) used in the fluorescence microscopy. The fluorescence quantum yields were estimated to be 11.0% for 2, 5.5% for 3, and 9.2% for 4. Although these values have some scatter, we judged that this would not affect the fluorescence microscopy results. The cells were incubated with the TMe-β-CyD-complexed with 2–4 at a concentration of 0.8 μM on a glass dish for 24 h in air with 5% CO2 at 37 °C (Figure 6). Because fluorescences were observed by treatment with the TMe-β-CyD-complexed with 2 and 3, neutral 2 and cationic 3 were taken up into HeLa cells (Figure 6b and 6d). Because the polar and hydrophilic groups of the compounds prohibit the transfer from the CyD cavities into the lipid membranes via an exchange reaction,12 it is thought that compounds 2 and 3 released from the TMe-β-CyD cavities may not be directly incorporated into the plasma membranes of the HeLa cells. In contrast, anionic 4 was not taken up into HeLa cells (Figure 6f). To understand the mechanism of uptake of 2 and 3 by the cells, we investigated temperature dependence of uptake, as it is well-known that endocytosis can be inhibited at low temperatures.

After TMe-β-CyD-complexed 2 and 3 was added, HeLa cells were incubated for 30 min at 4 °C (Figure S7). The results clarified that the fluorescence of 2 and 3 (Figure S7d and S7h) was not observed, in contrast with that of the uptake at 37 °C (Figure S7b and S7f). This result indicated that 2 and 3 were taken up into the HeLa cells by endocytotic processes. Consequently, the result suggests that TMe-β-CyD-complexed with 2 and 3 did not decompose after uptake by HeLa cells. The results indicate that (i) intracellular uptake of the neutral TMe-β-CyD-complexed with 2 increased due to interactions between the phenol moieties and the surface of the HeLa cells, (ii) uptake of the cationic TMe-β-CyD-complexed with 3 increased because of electrostatic interactions with the anionic cellular surface, while (iii) that of the neutral TMe-β-CyD-complexed with 2 decreased with the absence of phenolic hydroxy groups, and (iv) uptake of the anionic TMe-β-CyD-complexed with 4 decreased because of electrostatic repulsion. These results are consistent with those described in previous reports.23,24 However, why was the neutral TMe-β-CyD-complexed with 2 taken up into HeLa cells? To investigate the effect of the phenol moiety, we used a TMe-β-CyD-complexed with 5, with anisole moieties. Although the neutral TMe-β-CyD-complexed with 5 was also taken up into HeLa cells (Figure 6h), the fluorescence intensity was considerably weaker than that of the TMe-β-CyD-complexed with 2. This result indicates that a hydroxy group in the phenol moieties of 2 is important for interaction with the surface of HeLa cells. Although the details are not yet clear, we suggest the existence of phenol receptors on HeLa cells. To show the existence of phenol receptors, we tried to evaluate the inhibitory effect using phenol. However, this led to cell death due to the high cytotoxicity of phenol.
Figure 6. Phase contrast (a, c, e, and g) and fluorescence (b, d, f, and h) images of HeLa cells after being treated with (a and b) the TMe-β-CyD-complexed with \(2\), (c and d) the TMe-β-CyD-complexed with \(3\), (e and f) the TMe-β-CyD-complexed with \(4\), and (g and h) the TMe-β-CyD-complexed with \(5\) for 24 h at 37 °C. The scale bar represents 50 μm.

In summary, we have demonstrated that the photodynamic activity of TMe-β-CyD-complexed with \(2\) and \(3\) was remarkably greater than that of TMe-β-CyD-complexed with \(4\) toward HeLa cells under photoradiation with wavelengths greater than 610 nm. The TMe-β-CyD-complexed with \(2\) and \(3\) were approximately 14 and 26 times higher, respectively, in photodynamic activities than photofrin. The reasons for the high photodynamic activities of the TMe-β-CyD-complexed with \(2\) and \(3\) can be attributed to high intracellular uptake into HeLa cells, rather than high \(1\)O\(_2\) photoproduction abilities of the compounds. The fact that only a small amount of neutral TMe-β-CyD-complexed with \(5\) were taken up into the HeLa cells suggests that the phenol moieties in \(2\) are important for interaction with the surface of HeLa cells. Our findings therefore afford useful information to optimize the molecular design of TMe-β-CyD-complexed with porphyrin derivatives as photosensitizers. Future research will include the development of TMe-β-CyD-complexed with porphyrin derivatives with other substituents in meso-positions, for example hydroxy- and methoxy-substituted benzenes such as catechol, guaiacol, or 1,2-dimethoxybenzene.

ASSOCIATED CONTENT
Supporting Information
Supporting Information is available free of charge on the ACS Publications website.
Experimental procedures and UV-vis absorption and NMR spectra (PDF).

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Notes
The authors declare no competing financial interests.

ACKNOWLEDGMENTS
This work was supported by JSPS KAKENHI Grant-in-Aid for Scientific Research (B) (Grant No. JP16H04133) and Grant-in-Aid for Challenging Exploratory Research (Grant No. JP16K13982) and the Electric Technology Research Foundation of Chugoku. We would like to thank Prof. J. Ohshita and Dr. Y. Adachi of Hiroshima University for the fluorescence measurements.

ABBREVIATIONS
PS, photosensitizer; PDT, photodynamic therapy; ROS, reactive oxygen species; CyD, cyclodextrin; ABDA, 9,10-anthracenediyl-bis(methylene)dimalonic acid; NBT, nitroblue tetrazolium; IC\(_{50}\), half maximal inhibitory concentration.

REFERENCES
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