

2.5. Electrostatic interactivity

The electrostatic interactivity of the anionic vesicles was evaluated using the change in the zeta potential in presence of Ca^{2+} , pentyllysine, and poly-L-lysine (Mw. 15–30 kDa) as an index. A 10 μL aliquot of vesicles (lipid concentration: 2 g/dL) was diluted in 2 mL of 10 mM 2-[4-(2-Hydroxyethyl)-1-piperazinyl] ethanesulfonic acid (HEPES) buffer (pH 7.4, 37 °C) containing 0–3 mM CaCl_2 and 17–20 mM NaCl (total 20 mM), or containing varying amounts of pentyllysine or poly-L-lysine (Mw. 15–30 kDa) with 20 mM NaCl. The dispersions were incubated for 1 h at 37 °C before mobility measurement of the vesicles was performed by electrophoresis as described in Section 2.4.

2.6. Animal experiments

Animal experiments were conducted under the guidelines recommended by the National Institutes of Health, Animal Use and Care and the protocol was approved by the Steering Committee for Animal Experimentation at Waseda University. Male Wistar rats (250±20 g) were anesthetized with ether. The vesicular dispersion (5 g/dL) was introduced into rats through the tail veins at 1 mL/min ($n=5$ for each sample). Each rat received 5.6 mL/kg of body weight of vesicle dispersion (lipids: 280 mg/kg of body weight). At 1 or 24 h after injection, the blood was collected and centrifuged to separate the serum (1×10^5 g, 10 min). The collected serum was further ultracentrifuged to remove the vesicles (3×10^5 g, 30 min). The 50% hemolytic unit of complement serum (CH50) was determined in accordance with general procedures for clinical laboratory tests by a commercial company (BML, Japan).

2.7. Statistical methods

The data from the animal experiments are reported as means±standard error of the mean. Statistical analysis was performed using Microsoft Excel for Windows and CH50 values were compared using Student's unpaired *t* test.

3. Results and discussion

3.1. Samples

Two characteristic acidic lipids used in this study are shown in Fig. 1. SA and PG each have carboxylic acid and phosphoric acid as ionized groups, respectively. Both lipids have a strongly hydrophobic dialkyl structure to fix the ionized groups on vesicle surface. The molecular length indicated by the CPK model showed that the carboxylate anion of SA and the phosphate anion of PG would be located at the surface of PC-vesicles (Fig. 1B). Various amounts of SA and PG were incorporated into the PC/CH membrane (1:1, molar ratio) of the anionic vesicles. The size of vesicles was controlled by extrusion methods (final pore size: 0.2 μm), with final mean diameters of approximately 200 nm (Table 1). The vesicles prepared without acidic lipids

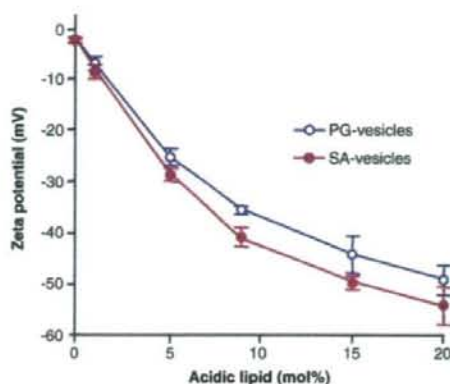


Fig. 2. Zeta potential of vesicles modified with SA (SA-vesicles) and PG (PG-vesicles) as a function of acidic lipid content. Zeta potentials were measured in 10 mM phosphate buffer (pH 7.4, NaCl; 20 mM) at 37 °C.

or with 1 mol% of acidic lipids tended to be slightly larger (ca 30 nm) than vesicles containing more acidic lipids. This effect of acidic lipids on the size of vesicles could be due to the improved dispersion stability of vesicles and electrostatic repulsion of the anionic surfaces. The vesicles without acidic lipids were observed to precipitate in a day, indicating poor dispersion stability. Vesicles without acidic lipids containing small amount of PEG-DSPE to prevent aggregation had diameters of 202±49 nm. Therefore, the slightly large diameter of vesicles without acidic lipids or with 1 mol% of acidic lipids may be the cause for their increased aggregability. The low aggregability of vesicles having large zeta potential is due to their electrostatic repulsive interaction causing anionic vesicles to exist as stable dispersions.

3.2. Zeta potential of vesicles

The zeta potential is the electrostatic potential at the hydrodynamic slip plane, and is characterized as having an electrical double-layer consisting of the Stern layer and the diffuse layer. Fig. 2 shows the zeta potential of prepared vesicles as a function of acidic lipid content at pH 7.4. Vesicles containing PC/CH (1:1, molar ratio) have an almost neutral surface (zeta potential: -2.22 ± 0.62 mV), indicating that the surface is inactive for electrostatic events. The magnitude of the negative charge on the surface increased with the incorporation of SA or PG, indicating that the ionized groups of SA and PG act to characterize the vesicle surface depending on their content. The zeta potentials of SA-vesicles and PG-vesicles reached -54.2 ± 3.68 mV and -49.0 ± 2.89 mV for acidic lipids of 20 mol%, respectively, with the negative zeta potential of SA-vesicles being relatively higher compared to that of PG-vesicles at any concentration. In theory, the electrostatic potential is dependent upon distance from the membrane surface as well as surface charge density [22]. The slightly extended negative charge of SA from the surface shown in Fig. 1B, would reduce the distance between the change to slip plane, resulting in the higher negative zeta potential of SA-vesicles.

Table 1
Diameter of prepared vesicles containing various amounts of acidic lipid

Acidic lipid (mol%)	Mean diameter±SD (nm)	
	SA-vesicles	PG-vesicles
0 (PC-vesicles)	232±60	232±60
1	224±58	230±60
5	193±50	196±58
9	205±40	204±49
15	194±45	199±55
20	198±52	194±64

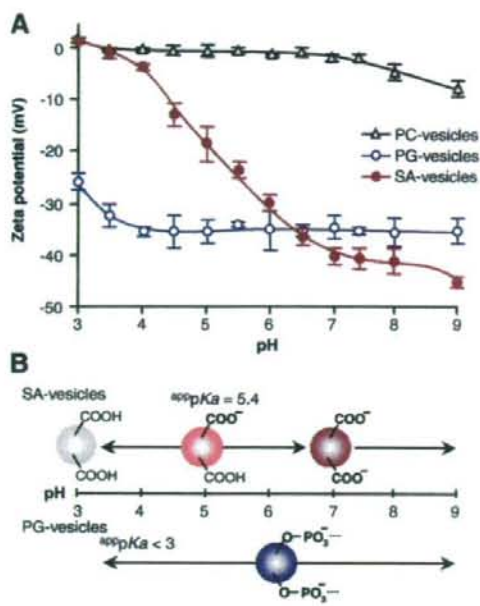


Fig. 3. Ionization state of acidic groups as a function of pH. (A) Zeta potential of vesicles (PC/CH, 1:1, molar ratio) (PC-vesicles), and vesicles containing 9 mol% of SA (SA-vesicles) or PG (PG-vesicles) at various pH. Zeta potentials were measured in 10 mM phosphate buffer (NaCl; 20 mM) at various pH (37 °C). (B) Schematic representation of the ionization state of acidic groups with pH. The apparent pK_a s of SA-vesicles and PG-vesicles were calculated to be 5.4 and <3.

This result demonstrated that the capacity of SA as an anionic component of vesicles is equal to acidic phospholipids at pH 7.4.

3.3. Ionization properties of acidic lipids

Stability of the ionization state conferred by acidity is an important characteristic of acidic compounds. We examined the ionization properties of acidic lipids as a function of pH using SA-vesicles and PG-vesicles containing acidic lipids of 9 mol% with zeta potentials of -40.7 ± 2.09 mV and -35.4 ± 0.61 mV (at pH 7.4), respectively. As shown in Fig. 3A, the zeta potential of SA-vesicles varied markedly depending on the environmental pH (pH 3–7). The change in the zeta potential of SA-vesicles is thought to reflect the ionization state of SA, as control vesicles without SA, namely PC-vesicles, maintained almost neutral surfaces irrespective of pH. The relationship between pH and the pK_a of the acid is expressed using the well-known Henderson–Hasselbalch equation as follows:

$$\text{pH} = \text{p}K_a + \log \frac{[-\text{COO}^-]}{[-\text{COOH}]} \quad (2)$$

When we analyzed the data shown in Fig. 3A using Eq. (2) and the assumption that the zeta potential was linearly correlated with the ionization acid, the pK_a of the carboxyl group of SA was estimated as 5.4 (Fig. 3B). Above pH 7, the zeta potential of SA-vesicles was almost constant, indicating that the carboxyl

group of SA would mostly be ionized above pH 7. The zeta potentials of PG-vesicles were almost constant in the range pH 4–9, indicating that the ionized form of the phosphoric acid moiety is stable in this range. The change in the zeta potential observed at a pH lower than pH 4, and its pK_a would be lower than pH 3 [23,24]. Thus, we confirmed that the surface of SA-vesicles and PG-vesicles exhibited the characteristics of a weak acid with SA and a strong acid with PG, respectively, indicating that the individual characteristics of acidic groups are expressed on the surface of vesicles. We also observed that the magnitude of the negative electrostatic charges in SA-vesicles was equal to that observed in PG-vesicles at approximately neutral pH.

3.4. Electrostatic interactivity

Ca^{2+} is found in biological fluids (normally 2–3 mM in plasma) and is known to mediate biological processes by binding to the anionic domains such as those involved in the specific binding of proteins to membranes [25,26]. Acidic phospholipids, such as PG and PS, are also known to bind Ca^{2+} [27–29]. As shown in Fig. 4, the negative charge on anionic vesicles was suppressed by increasing the concentration of Ca^{2+} . When the concentration of Ca^{2+} was increased to 3 mM, the zeta potentials of SA-vesicles and PG-vesicles were -16.4 ± 1.9 mV and -13.8 ± 1.4 mV, respectively. Recently, Hautala et al. reported that vesicles containing phosphatidic acid possess a specifically strong affinity for Ca^{2+} , and that the zeta potential of these vesicles changed from being strongly negative to positive after binding Ca^{2+} [29]. These authors also showed that other acidic phospholipids, including PG, do not exhibit a strong affinity towards becoming cationic. Consequently, one aim of this experiment was to determine whether the binding of Ca^{2+} is a specificity factor between SA-vesicles and PG-vesicles, and also whether the surface of SA-vesicles remained negative in the presence of Ca^{2+} . Our experiment showed that the surface of the SA-vesicles remained

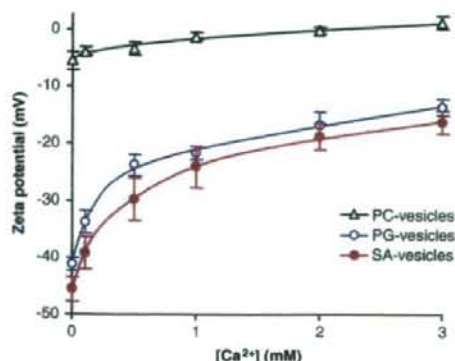


Fig. 4. Change in zeta potential of vesicles as a function of Ca^{2+} concentration. Vesicles were dispersed at 100 $\mu\text{g}/\text{mL}$ in 10 mM HEPES buffer (pH 7.4, at 37 °C) containing NaCl and CaCl_2 (total: 20 mM). PC-vesicles: PC/CH (1:1, molar ratio), SA-vesicles: PC/CH/SA (1:1:0.2, molar ratio), and PG-vesicles: PC/CH/PG (1:1:0.2, molar ratio).

negative in the presence of Ca^{2+} . In addition, comparisons of SA-vesicles and PG-vesicles also showed that the specificity of binding Ca^{2+} was not observed.

Additional model-based studies of electrostatic interactivity, pentyllysine and poly-L-lysine (Mw. 15–30 kDa with a repeating primary lysine amine) were also conducted. Oligomers or polymers of lysine are often used to model basic peptides or macromolecules and their electrostatic interactions on membranes [30,31]. As shown in Fig. 5, the zeta potential is a linear function of the concentration of pentyllysine. The lines, which represent the least-squares best fit, have slopes of 9.3 and 7.4 mV per decade for the pentyllysine concentrations in SA-vesicles and PG-vesicles. The slope for PC-vesicles was as little as 0.26 mV per decade of pentyllysine concentration (data not shown), indicating that the negative charge of acidic lipids mediate the interaction with basic pentyllysine. A change in the zeta potential is due to binding of basic peptide [31] and a similar decay slope of the zeta potential would indicate that the binding constant of a basic peptide to SA-vesicles and PG-vesicles was similar. As shown in Fig. 6, the change in the zeta potential of these vesicles due to the interaction with poly-L-lysine increased drastically, changing from a negative to a positive in presence of 1.5–2 $\mu\text{g}/\text{mL}$ poly-L-lysine. Conversely, the change in the surface potential of PC-vesicles was negligible, indicating that the acidic lipids mediate the interaction with basic macromolecules. This experiment also demonstrates that the interactivity of SA-vesicles and PG-vesicles to basic macromolecules is equal at pH 7.4, and that within an electrostatic context, SA-vesicles and PG-vesicles interact similarly with basic compounds at physiological pH. It has been shown that electrostatic interactions are involved in the binding of C1q to the surface of anionic vesicles containing acidic phospholipid [6]. In the event that the negative

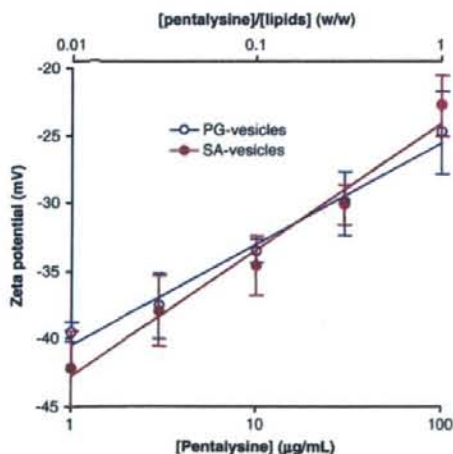


Fig. 5. Change in zeta potential of vesicles as a function of the concentration of basic oligomer (pentyllysine). Vesicles were dispersed at 100 $\mu\text{g}/\text{mL}$ in 10 mM HEPES buffer (pH 7.4, at 37 °C, NaCl, 20 mM) containing various amount of pentyllysine. The lines have slopes of 9.3 and 7.4 mV per decade for the pentyllysine concentrations in SA-vesicles and PG-vesicles. SA-vesicles: PC/CH/SA (1:1:0.2, molar ratio) and PG-vesicles: PC/CH/PG (1:1:0.2, molar ratio).

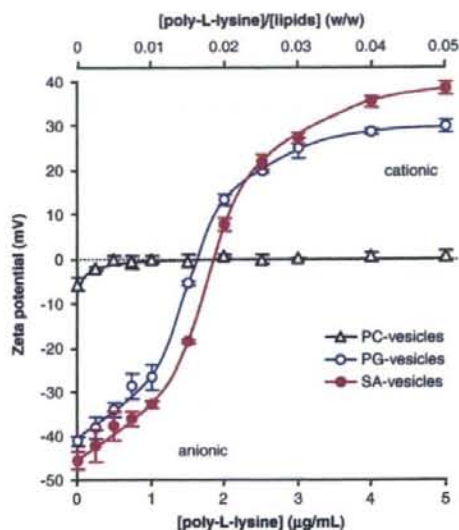


Fig. 6. Change in zeta potential of vesicles as a function of the concentration of basic macromolecule (poly-L-lysine, Mw. 15–30 kDa). Vesicles were dispersed at 100 $\mu\text{g}/\text{mL}$ in 10 mM HEPES buffer (pH 7.4, at 37 °C, NaCl, 20 mM) containing various amount of poly-L-lysine. PC-vesicles: PC/CH (1:1, molar ratio), SA-vesicles: PC/CH/SA (1:1:0.2, molar ratio), and PG-vesicles: PC/CH/PG (1:1:0.2, molar ratio).

charge on the surface of vesicles is critical for complement activation, both the SA-vesicles and PG-vesicles should be capable of activating the complement system to similar degree. We therefore conducted animal experiments to clarify the issue of complement activation by the anionic electrostatic charge of vesicles.

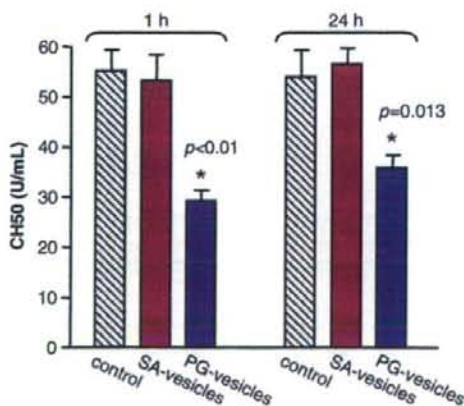


Fig. 7. Comparison of the 50% hemolytic unit of complement (CH50) in rat serum at 1 or 24 h after infusion of saline (control), SA-vesicles, or PG-vesicles. CH50 value for PG-vesicles was significantly lower than that of the control ($p < 0.01$ at 1 h, $p = 0.013$ at 24 h), indicating the complement consumption in serum after activation. Equal CH50 value for SA-vesicles with the control indicates that SA-vesicles failed to activate complement.

3.5. Complement activation

Serum was collected at 1 and 24 h after the infusion of SA-vesicles or PG-vesicles in experimental rats to determine CH50 levels. The control group received saline as a vehicle. The values of CH50 in control rats, and SA-vesicles- and PG-vesicles-administered rats, were 55.1 ± 4.1 U/mL, 53.2 ± 4.8 U/mL, and 29.3 ± 1.6 U/mL at 1 h after infusion of sample, respectively. At 24 h, the values of CH50 in control rats, and SA-vesicles- and PG-vesicles-administered rats, were 54.0 ± 4.7 U/mL, 56.5 ± 3.3 U/mL, and 35.8 ± 2.3 U/mL, respectively (Fig. 7). The lower CH50 levels observed in the PG-vesicles-administered group in comparison with the control group indicate that complement consumption occurred after activation. These findings imply that significant complement activation is induced in rats receiving PG-vesicles compared to the control group ($p < 0.01$ at 1 h, $p = 0.013$ at 24 h). Complement consumption was not observed in rats administered SA-vesicles.

Since the negative charge and electrostatic interactivity of SA-vesicles were the same as in PG-vesicles (Figs. 2–6), the data obtained from the animal experiments indicates that the negative charge on the anionic vesicle is not a critical factor underlying the activation of complement. The first step in the activation of the classical complement pathway involves the binding of an activator to C1q, resulting in the activation of serine proteases C1r and C1s. It has been suggested that the negative charge of an activator such as PG-vesicles is involved in some way with the binding of the activator to C1q [1–8]. Assuming that the electrostatic interaction is non-specific, SA-vesicles should interact with C1q electrostatically. Since the action of complement proteases, which follows the binding of the activator to the C1q, is known to be highly specific [32,33], it seems likely that complement activation on anionic surface is limited to an activation step rather than a binding step. Such specific activation of complement by the anionic vesicles in the present study may be involved in the physiological regulation of complement activation on anionic biomembranes.

4. Conclusions

The carboxylic acid of SA and phosphoric acid of PG have equal capacity as anionic components of vesicles at neutral pH. The results presented in this investigation demonstrated that the negative electrostatic charge of anionic vesicles is not a critical factor in the activation of complement. Rather, the induction of complement activation by anionic vesicles is dependent on the structure of acidic lipids. This finding may facilitate development and various biological applications of anionic vesicles.

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Structure, Photophysical Property, and Cytotoxicity of Human Serum Albumin Complexed with Tris(dicarboxymethylene)[60]fullerene

Xue Qu,[†] Teruyuki Komatsu,^{*†‡} Takaaki Sato,[†] Otto Glatter,[§] Hirohisa Horinouchi,^{||} Koichi Kobayashi,^{||} and Eishun Tsuchida^{*†}

Research Institute for Science and Engineering, Waseda University, 3-4-1 Okubo, Shinjuku-ku, Tokyo 169-8555, Japan, PRESTO, Japan Science and Technology Agency (JST), Institute of Chemistry, University of Graz, Heinrichstrasse 28, A-8010 Graz, Austria, and Department of Thoracic Surgery, School of Medicine, Keio University, 35 Shinanomachi, Shinjuku-ku, Tokyo 160-8582. Received February 6, 2008; Revised Manuscript Received June 18, 2008

We present structure, photoexcited triplet-state property, and singlet oxygen ($^1\text{O}_2$) generation capability of human serum albumin complexed with a tris(dicarboxymethylene)[60]fullerene C₃-isomer (HSA-CF). Small angle X-ray scattering measurements showed that a globular size-dimension and surface charge distribution of HSA were unaltered by monomolecular complexation of CF into the hydrophobic cavity. Laser flash photolysis to the HSA-CF solution yielded a photoexcited triplet state of the CF chromophore ($^3\text{CF}^*$) with lifetime of 46 μs . In the presence of O₂, energy transfer occurred from HSA- $^3\text{CF}^*$ to O₂ to generate $^1\text{O}_2$; the quenching rate constant [$k_q(\text{O}_2)$] was determined to be $2.2 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$. The HSA-CF showed strong photoresistance relative to HSA complexed with a protoporphyrin IX. The quantum yield of $^1\text{O}_2$ production for this artificial protein was compared to those of other photosensitizing agent. The HSA-CF did not show a dark cytotoxicity, but induced cell death efficiently under visible light irradiation.

INTRODUCTION

Photodynamic therapy (PDT) is a nonsurgical cancer treatment that combines a photosensitizing medication with exposure to visible light (1, 2). Highly cytotoxic singlet oxygen ($^1\text{O}_2$) formed by energy transfer from photoexcited sensitizer to molecular O₂ has been implicated as an intermediary species leading to cell death in tumors. To accelerate this $^1\text{O}_2$ generation, a range of organic dyes, especially porphyrin compounds, have been synthesized as photosensitizers (3). The most widely used reagent in clinical PDT is Photofrin, which is a heterogeneous mixture of hematoporphyrin oligomers (1, 2, 4). Several porphyrin or chlorin derivatives are also being tested as second generation (1, 2, 5, 6). Another well-known agent approved by the FDA is Levulan; 5-aminolevulinic acid (ALA) (7–9). ALA-induced photosensitization is unique approach that utilizes the heme biosynthetic pathway to produce endogenous protoporphyrin IX (PP). When ALA enters cancer cells, it triggers the synthesis of PP, which acts as an active sensitizer. Wavelengths typically used in PDT are between 600 and 800 nm, because light below 600 nm is absorbed by tissues and hemoglobin (Hb). Photofrin is normally excited by a red light (630 nm) using an Excimer dye-laser (1, 2). The ALA-PDT also uses the 630 nm laser, while the irradiation using a metal halide lamp or xenon arc lamp is still popular, because this is reasonable in comparison to the expensive laser pulse system and suitable for irradiation of the wide area of the diseased part (9). In fact, the light source approved in the U.S. for ALA-PDT was halogen lamp ($\lambda_{\text{exc}} = 410 \text{ nm}$). Buckminster[60]fullerene (C₆₀) produces $^1\text{O}_2$ by energy transfer with extremely high quantum yield (Φ_{Δ} : 0.96,

$\lambda_{\text{exc}} = 532 \text{ nm}$) (10, 11) and generally shows strong photoresistance compared to the porphyrins. Consequently, various water-soluble fullerenes and fullerene-polymer hybrids have been prepared as a new class of photosensitizer for $^1\text{O}_2$ production (11–16). However, we would like to emphasize herein that most exogenous compounds administered into our bloodstream are immediately captured by human serum albumin (HSA, Mw: 66.5 kDa), which is the major protein component in our blood plasma (17, 18). To evaluate the biological performance of the fullerene, we must investigate the structure and function of the HSA-fullerene complex. Despite that necessity, photochemistry of the HSA-fullerene complex has never been studied to date. In this paper, we report for the first time the structure and photoexcited triplet-state properties of HSA complexed with a tris(dicarboxymethylene)[60]fullerene C₃-isomer (HSA-CF) and highlight its photoinduced energy transfer to O₂ to produce $^1\text{O}_2$. Furthermore, cytotoxicity of HSA-CF in vitro has also been evaluated. It is known that CF acts as a neuroprotective agent (19) and binds to HSA (20), but its photophysical behavior and cytotoxicity under visible light irradiation have not been elucidated.

EXPERIMENTAL PROCEDURES

Materials and Apparatus. All reagents were purchased from commercial sources as special grades and used without further purification. Recombinant HSA expressed in *Pichia pastoris* was purchased from Sigma-Aldrich Corp. Tris(dicarboxymethylene)-[60]fullerene C₃-isomer (CF) was purchased from Axxora LLC and 2-[4-(2-hydroxyethoxy)phenyl]-1,3-dioxolo-4a,5a-[60]fullerene (HF) was purchased from FLOX Corp. Protoporphyrin IX (PP), protoporphyrin IX dimethyl ester (PPDME), and [60]fullerene (C₆₀) were purchased from Sigma-Aldrich Corp. The water was deionized using a Millipore Elix and Simpli Laboratory-UV. The UV-vis absorption spectra for the potassium phosphate buffered (PB) solution (pH 7.0, 50 mM) of the HSA-fullerene complex (normally 20 μM) were recorded using an Agilent 8453 UV-visible spectrophotometer with an Agilent

* Corresponding author. (E.T.) Tel: +81-3-5286-3120, Fax: +81-3-3205-4740, E-mail: eishun@waseda.jp. (T.K.) E-mail: teruyuki@waseda.jp.

[†] Waseda University.

[‡] PRESTO, JST.

[§] University of Graz.

^{||} Keio University.

89090A temperature control unit. The CD spectra for the PB solution of the HSA–fullerene complex were measured using a JASCO J-820 circular dichrometer. The concentration of the HSA sample was 2 μ M and a quartz cuvette with a 2.0 mm thickness was used for measurements in the range 195–250 nm. Mass spectroscopic analyses were performed using a JEOL JMS-T100CS ESI-TOF mass spectrometer.

Preparations of HSA–Fullerene Complexes. Typically, 3.92 mL of a PB solution (pH 7.0, 50 mM) of HSA (0.25 mM) was mixed with 1.0 mL of 0.98 mM CF in PB (CF/HSA molar ratio of 1:1) and incubated for 12 h with rotation in the dark at room temperature. The complex was then diluted with 50 mM PB solution and concentrated to the initial volume using a Vivaspin 20 centrifuge filter (5 kDa MW cutoff) at 4000 \times *g* using a Beckman Coulter Allegra X-15R centrifuge. In the case of HSA–HF, 4.29 mL of PB solution (pH 7.0, 50 mM) of HSA (42 μ M) was mixed with 0.67 mL of 0.267 mM HF in DMSO (HF/HSA molar ratio of 1:1) and incubated for 12 h with rotation in the dark at room temperature ([DMSO] = 13.5 vol %). The complex was then diluted using 50 mM PB solution and concentrated to the initial volume using a Vivaspin 20 centrifuge filter at 4000 \times *g*. These dilution/concentration cycles were repeated to reduce the DMSO concentration to <0.1 vol %. The obtained protein solution was loaded on a Superdex 75 g gel-permeation column by elution of 50 mM PB using an ÄKTA Prime Plus FPLC system; only one band was observed during the chromatography. The PB solutions (pH 7.0, 50 mM) of HSA–CF and HSA–HF ([HSA]: 20 μ M) in a 10-mm-path length optical quartz cuvette sealed with a rubber septum was degassed and purged using N₂ prior to use. The HSA complexed with PP (HSA–PP) was also prepared by the same procedure as HSA–HF.

HPLC. HPLC measurement of the HSA–fullerene complex was carried out using an HPLC system consisted of a Shimadzu LC-8A pump and a Shimadzu SPD-10A UV detector. A Shodex Protein KW-803 column was used with PB (pH 7.0, 10 mM) as the mobile phase at 22 °C (1.0 mL min⁻¹).

Isoelectric Focusing (IEF). The IEF was performed using an Amersham Biosciences Electrophoresis Power Supply EPS 301 with an Invitrogen Novex pH 3–7 IEG Gel. The protein marker used was a GE Healthcare IEF calibration kit Low Range pI (pH 2.5–6.5). The voltage was gradually raised to 300 V for 4 h. The gels were stained by a Daiichi Pure Chemicals PAGE Blue 83 and destained by washing with aqueous ethanol/acetic acid and 10% acetic acid.

Small-Angle X-ray Scattering (SAXS). SAXS experiments were carried out using an Anton Paar SAXSess camera in the *q*-range 0.10–9.0 nm⁻¹. Deionization of the protein sample was performed by dilution/concentration cycles with pure water using Vivaspin 20 centrifuge filter (5 kDa MW cutoff) at 4000 \times *g*, giving an aqueous solution of the HSA–CF complex. Washing with phosphate buffered saline (PBS) solution (pH 7.4, 150 mM) gave the PBS solution of the protein. The scattered intensity $I(q)$ was generally given as $I(q) = nP(q)S(q)$ for globular particle system, where *n* is the particle number density, $P(q)$ is the form factor, and $S(q)$ is static structure factor. The measured intensities were normalized to give the uniform primary intensity at $q = 0$ and converted into absolute scale using water as a secondary standard. A model-independent collimation-correction procedure was made via an indirect Fourier transformation (IFT) technique. The pair-distance distribution functions $p(r)$ of HSA and HSA–CF in solution, which are given as inverse Fourier transformations of the form factor $P(q)$ into real space, were obtained using IFT technique for a highly diluted sample ($S(q) \sim 1$) and its generalized version (GIFT) for $c > 1$ g dL⁻¹. The GIFT technique enables splitting of $I(q)$ into $P(q)$ and $S(q)$, letting $P(q)$ and the resulting $p(r)$ virtually model free, where

we assumed a screened Coulomb (Yukawa) potential and Rogers-Young (RY) closure relation. We made a complementary analysis to deduce the experimental structure factor, $S(q)^{exp}$, by dividing the normalized intensity $I(q)/c$ by the experimental form factor $P(q)^{exp}$ determined at very low concentration, which confirms the validity of the GIFT results.

Laser Flash Photolysis. The transient absorption spectra and triplet lifetime measurements were carried out using a Unisoku TSP-1000WK time-resolved spectrophotometer with a Spectron Laser Systems SL803G-10 Q-switched Nd:YAG laser, which generated a second-harmonic (532 nm) pulse of 6 ns duration (10 Hz) (21). The samples (20 μ M) of the PB solutions (50 mM, pH 7.0) of HSA–CF, HSA–HF, HSA–PP, THF solution of PPDME, and toluene solution of C₆₀ were held in a quartz cuvette (optical path length: 10 mm); experiments were carried out at 22 °C. The gas mixture with the desired partial pressure of O₂/N₂ prepared by a KOFLOC Gasbender GB-2C was flowed into the sample cuvette for 20 min for equilibration.

Near-Infrared Photo-Counting Measurement. The emission spectrum of ¹O₂ (¹Δ_g→³Σ_g⁻) was measured using a Hamamatsu Photonics C8282W Near-Infrared Photo-Counting System equipped with a Hamamatsu Photonics R5509 photomultiplier and a Q-switched Nd:YAG laser, which generated a second-harmonic (532 nm) pulse (20 Hz, 60 μ J/pulse at the sample position). The gate time width was 2–100 μ s. The samples (40 μ M) were held in a quartz cuvette (optical path length 10 mm). The experiments were carried out in air at 22 °C. Quantum yield of ¹O₂ (Φ_{Δ}) production by HSA–CF was determined using rose bengal as the standard compound.

In Vitro Cytotoxicity Evaluation. The LY80 tumor cell, a variant of the Yoshida sarcoma, was kindly gifted from Dr. Katsuyoshi Hori, Tohoku University. The cells were grown at 37 °C in DMEM supplemented with 10% fetal bovine serum (FBS) and penicillin (100 IU/cm³) in 95% air–5% CO₂ humidified incubator. The cell solution (2.5 \times 10⁵ cells/mL, 0.16 mL) was mixed with PBS (0.04 mL) or PBS solution of HSA–CF (100 μ M, 0.04 mL) in a Corning Costar 3596 cell culture plate (96 well) and incubated in the dark. Then the plate was irradiated by visible light using a 500 W halogen lamp in conjunction with a HOYA HA30 heat absorption filter (330–700 nm). The power at the cell level was monitored using a Gentec Power Meter and adjusted to 20 mW cm⁻². The temperature was kept at 36 \pm 1 °C during the irradiation. As a negative control, we examined another PBS group in the same culture plate without light irradiation. After 2 h, the cell numbers were counted using microscopy under tripan blue staining.

RESULTS AND DISCUSSION

Structure of HSA–Fullerene Complexes. The HSA–fullerene complexes were prepared essentially as described previously for HSA–hemin complexes (21). A phosphate buffered (PB) solution (pH 7.0, 50 mM) of HSA was mixed with PB solution of CF (CF/HSA molar ratio of 1:1) and incubated for 12 h. The complex was then concentrated to the initial volume using a centrifuge filter (Figure 1 inset photograph). The filtrate was colorless during the centrifugation. The gel permeation chromatogram (Superdex 75 g) of the orange-colored protein exhibited only a single elution peak. These results indicate that CF is efficiently incorporated into HSA. For comparison, hydrophobic fullerene, 2-[4-(2-hydroxyethoxy)-phenyl]-1,3-dioxolo-4a,5a-[60]fullerene (HF), was also complexed with HSA to give HSA–HF. The obtained HSA–fullerene solutions can be concentrated up to 3.75 mM (25 g dL⁻¹); they were stable for a year without precipitation.

The UV–vis absorption spectroscopic features of the HSA–CF solution are the sum of those from the individual HSA and CF (Figure 1), which indicates that CF is monomolecularly

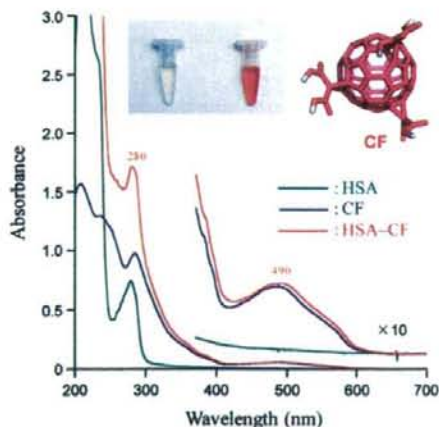


Figure 1. UV-vis absorption spectrum of HSA-CF complex, CF and HSA (20 μM) in PB solution (pH 7.0, 50 mM) at 22 $^{\circ}\text{C}$. Inset photographs are 5 g dL^{-1} HSA (left) and HSA-CF (right) solutions.

incorporated into HSA and no specific interaction between the two molecules at the ground state. In contrast, the absorption spectrum of the brown-colored HSA-HF solution did not coincide with the superposition of the HSA and HF spectra (Figure S1). The increased absorbance at 380–550 nm suggests the formation of HF clusters in the protein (11, 16).

The HSA-CF complex was sufficiently stable to apply the HPLC measurement. In the elution profile, a single peak appeared at a retention time of 7.64 min, which was slightly earlier than that of HSA itself (7.75 min) (Figure S2). The ratio of the peak intensity monitored at 280 nm (based on HSA) and at 490 nm (based on CF) ($I_{280}/I_{490} = 30$) was exactly the same as the absorbance ratio at 280 and 490 nm in the UV-vis absorption spectrum (A_{280}/A_{490}) of HSA-CF, which implies that all CF molecules were eluted within the HSA fraction.

The ESI-TOF mass spectrometry of HSA-CF showed a distinct ion peak at 67 587 Da, which corresponds to the mass of the equivalent complex of HSA-CF. On the other hand, two separated peaks were observed in the HSA-HF spectrum at 66 585 (free HSA) and 68 296 Da, that reveals 1:2 stoichiometry for HSA/HF and the formation of HF clusters in the protein. The CD spectral patterns and intensities of HSA-CF and HSA-HF were identical to those of HSA (Figure S3). We inferred that fullerene binding did not change the highly ordered structure of the host HSA.

The incorporation of negatively charged CF having hexa-carboxyl groups might influence the surface charge distribution of albumin. However, isoelectric focusing of HSA-CF indicated the same isoelectric point ($p\text{I}$) with HSA. This contrasts sharply to the fact that fatty acid binding induced the reduction of the $p\text{I}$ value because of partial neutralization of the molecular charge (22). We then used small-angle X-ray scattering (SAXS) to evaluate the globular particle structure and protein-protein interactions of HSA-CF (Figure 2) (23). The pair-distance distribution functions [$p(r)$] of HSA-CF and HSA were almost identical to the curve calculated from crystallographic data of HSA (Iuor) (Figure 2 upper inset). It demonstrates that the maximum diameter (D_{max} : ca. 8 nm) and three-dimensional particle shape of HSA were unaltered by complexation of CF. The normalized SAXS intensities [$I(q)/c$] of HSA-CF at the low q range [$I(q \rightarrow 0)/c$] monotonously decreases with increasing protein concentration. In the PBS solution, the long-range electrostatic repulsion between HSA-CFs is efficiently screened, so that the suppressed forward intensity could be mainly due

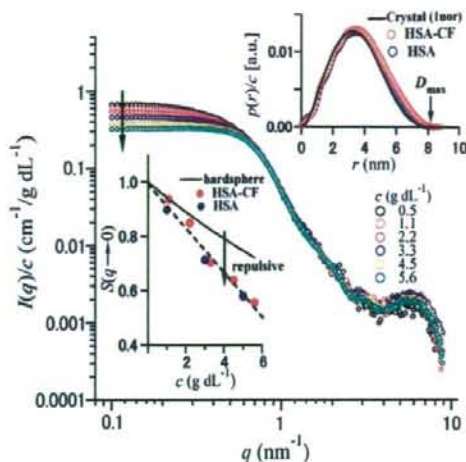


Figure 2. Normalized SAXS intensities [$I(q)/c$] of HSA-CF complex in 150 mM PBS solutions ($c = 0.5\text{--}5.6 \text{ g dL}^{-1}$). The upper and lower insets show pair-distance distribution functions [$p(r)$] and extrapolated structure factors [$S(q \rightarrow 0)$] of HSA-CF and HSA, respectively.

to excluded volume effect. However, the extrapolated structure factors [$S(q \rightarrow 0)$] of HSA-CF and HSA, which reflect the net repulsive forces between the protein molecules, were systematically lower than that predicted for hardsphere with an identical volume fraction (Figure 2 lower inset). The negative deviations are attributed to the electrostatic repulsion between the proteins. It is remarkable that the plots of $S(q \rightarrow 0)$ for HSA-CF and HSA perfectly lie on the same line, suggesting that HSA preserves its surface net charges upon the CF binding. On the basis of these results, we conclude that (i) CF is accommodated into the deep hydrophobic cavity of HSA with internal charge neutralization, and (ii) it does not induce marked changes in the globular particle size and surface charge distribution of HSA.

Photoexcited Triplet State of HSA-CF Complex and $^1\text{O}_2$ Formation. Photoexcitation of the fullerenes generates the singlet state, which undergoes intersystem crossing to the triplet state in high yield (10, 11). Laser flash photolysis (SHG of Nd:YAG, 532 nm, 6 ns) of the HSA-CF solution under an N_2 atmosphere gave a triplet-triplet (T-T) absorption spectrum of the $^3\text{CF}^*$ chromophore ($\lambda_{\text{max}} = 740 \text{ nm}$) (Figure 3) (11, 13, 24). The time course of the absorbance decay was composed of a single exponential kinetics with a lifetime (τ_T) of 46 μs (Figure 3 inset). The peaks of maximum absorption and τ_T value resembled those of the aqueous CF solution (τ_T : 43 μs). These observations again support that CF is held monomolecularly within the HSA interior.

Although HF in DMSO exhibited clear triplet absorption (τ_T : 31 μs), HSA-HF showed no signals in this wavelength range (Figure S4). The extremely short triplet lifetime of HSA-HF is probably attributed to T-T annihilation of the HF aggregates in the protein (11). If this is the case, the lifetime should become longer when the laser pulse energy is decreased. However, the decay was too fast to analyze precisely even under the weak laser flash photolysis.

In the presence of O_2 , the triplet lifetime of HSA-CF significantly decreased. Energy transfer occurred from HSA- $^3\text{CF}^*$ to the O_2 molecule to generate active $^1\text{O}_2$. The Stern-Volmer plot clearly depicts a linear correlation for O_2 concentrations of 0–1.0 mM, giving the Stern-Volmer constant (K_{SV}) and quenching rate constant [$k_q(\text{O}_2)$] (Figure 4) [Note: $\tau_T^0/\tau_T = K_{\text{SV}}[\text{O}_2]$ (τ_T^0 and τ_T represent triplet lifetimes in the absence and presence of O_2)]. The $k_q(\text{O}_2)$ of HSA-CF was $2.2 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$, which is one-seventh that of the monomeric

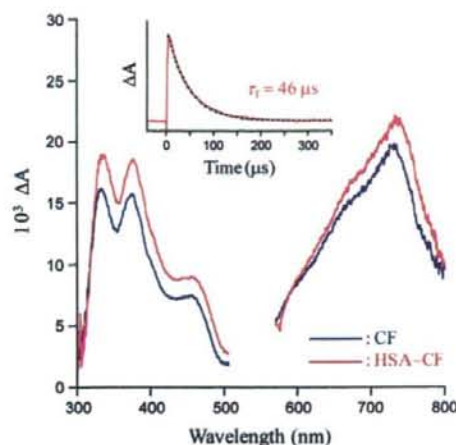


Figure 3. Triplet-triplet absorption spectra of HSA-CF complex and CF in PB solution (pH 7.0, 50 mM) under N_2 atmosphere at 22 °C; Ex. at 532 nm. The inset shows absorption decay of HSA-CF complex at 734 nm, which was fitted by single exponential relaxation curve (black dotted line).

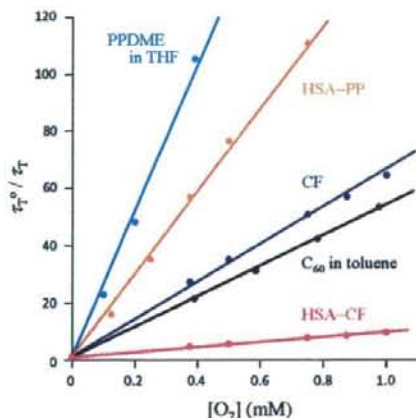


Figure 4. Stern-Volmer plots of triplet state quenching of HSA-CF complex and other compounds by O_2 at 22 °C. The HSA-CF, HSA-PP, and CF were in PB solution (pH 7.0, 50 mM).

Table 1. Quenching Rate Constants of the Photoinduced Excited Triplet States of HSA-CF Complex and Other Compounds by O_2 at 22 °C

compound	solvent	$k_q(O_2)$ ($M^{-1} s^{-1}$)
HSA-CF	PB (pH 7.0, 50 mM)	2.2×10^8
CF	PB (pH 7.0, 50 mM)	1.6×10^9
HSA-PP	PB (pH 7.0, 50 mM)	1.3×10^8
PPDME	THF	1.6×10^9
C_{60}	toluene	1.6×10^9
		1.9×10^{10}

^a ref 10.

CF solution ($1.6 \times 10^9 M^{-1} s^{-1}$) (Table 1). The HSA-PP complex in PB also showed smaller $k_q(O_2)$ ($1.3 \times 10^8 M^{-1} s^{-1}$) compared to that of the protoporphyrin IX dimethylester (PPDME) in THF (25). The slow $k_q(O_2)$ values of the protein systems suggest that the quenching reaction is almost diffusion controlled. The intensity of the visible band of HSA-CF ($\lambda_{max} = 490$ nm) did not change after 10^3 laser flash photolysis ($\lambda_{ex} = 532$ nm, 0.5 W) in air, indicating no photodegradation of the CF chromophore. In contrast, the Soret band of HSA-PP was

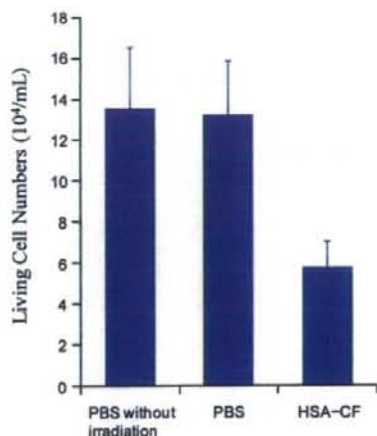


Figure 5. Living cell numbers of LY80 with HSA-CF and PBS after visible light irradiation ($20 mW cm^{-2}$, 2 h, 36 ± 1 °C). Each value represents the mean \pm SD ($n = 4$).

$\approx 6\%$ photobleached after the identical flash irradiations. It can be concluded that the light resistance of CF is significantly higher than that of PP, which is an active photosensitizer in ALA-PDT.

We observed the near-infrared luminescence of the produced 1O_2 (λ_{max} 1270 nm) and evaluated the quantum yield of 1O_2 production (Φ_Δ) for HSA-CF from the value of emission intensity at 1270 nm (I) using rose bengal as a standard (26). The Φ_Δ value of HSA-CF (0.46) was in the same range as those of monomeric CF (0.48), methylene blue (0.52), PPDME in benzene (0.59) (26), and substituted fullerenes in organic solvent (27, 28), but somewhat lower than those of rose bengal (0.75), hematoporphyrin in methanol (0.74) (29), and C_{60} (0.96) in benzene (10). It is known that the substitution of the pristine C_{60} causes a perturbation of the electronic structure of the fullerene core, and thereby results in a decrease in the quantum yield for the 1O_2 formation (27, 28).

Cytotoxicity of HSA-CF Complex. Finally, we measured the cytotoxicity and photodynamic activity of the HSA-CF complex to the LY80 tumor cells in vitro. The cell cultures were first incubated for 24 h in the dark with HSA-CF ($20 \mu M$) at 37 °C under 5% CO_2 . The cell numbers after the incubation were identical to that of control group with the PBS solution, indicating that the HSA-CF complex does not have dark cytotoxicity. The cell culture plates were then exposed to visible light of 350–600 nm ($20 mW cm^{-2}$) for 2 h at 36 ± 1 °C. Some of the cells mixed with HSA-CF showed abnormal shape after the light irradiation, whereas the PBS group did not show any morphological changes of the cells. The living cell numbers of the PBS groups with and without light irradiation were almost the same (Figure 5), which means that the light exposure did not affect the LY80 tumor cells in this experimental condition. On the other hand, the living cell numbers of the HSA-CF group were markedly lower than that of the PBS group; 57% cell death occurred by visible light irradiation. This result clearly implies that the HSA-CF complex acts as a photosensitizer for PDT.

CONCLUSIONS

The HSA-CF complex having identical molecular shape and surface charge distribution with HSA is easily excited by visible light and shows a high charge transfer rate constant for molecular O_2 . The efficiency of the 1O_2 production of this artificial protein is in the same range as well-known dye

sensitizers, such as methylene blue and PPDME. In general, water-soluble drugs administered into our bloodstream are quickly eliminated from the body. On the contrary, the HSA hybrid can remain in the circulatory system and/or tissues with a long persistence; therefore, the $^1\text{O}_2$ generation by albumin-based photosensitizer might be of clinical importance for photodynamic cancer therapy. The HSA-CF complex does not show dark cytotoxicity, but engenders cell death efficiently under visible light irradiation. Recombinant HSA is currently manufactured on an industrial scale using yeast species *Pichia pastoris* (30), which allows the use of the HSA-fullerene complex in practical applications.

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Supporting Information Available: Figures S1–S4. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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COMMUNICATIONS

O₂ Binding to Human Serum Albumin Incorporating Iron Porphyrin with a Covalently Linked Methyl-L-Histidine Isomer

Akito Nakagawa,[†] Teruyuki Komatsu,^{*,†,‡} Makoto Iizuka,[†] and Eishun Tsuchida^{*,†}

Research Institute for Science and Engineering, Waseda University, 3-4-1 Okubo, Shinjuku-ku, Tokyo 169-8555, Japan, and PRESTO, Japan Science and Technology Agency (JST), 4-1-8 Honcho, Kawaguchi-shi, Saitama 332-0012, Japan. Received October 30, 2007; Revised Manuscript Received December 1, 2007

We describe the significant difference in the O₂ binding affinities of human serum albumin (HSA) incorporating 5,10,15,20-tetrakis($\alpha,\alpha,\alpha,\alpha$ -*o*-(1'-methylcyclohexanamido)phenyl)porphinatoiron(II) with a covalently linked 1-methyl-L-histidine or 3-methyl-L-histidine [HSA-FeP(1-MHis), HSA-FeP(3-MHis)]. The HSA-FeP(3-MHis) showed an extraordinarily high O₂ binding affinity ($P_{1/2} = 0.2$ Torr, 25 °C, pH 7.4), which is close to those of relaxed-state hemoglobin and myoglobin. However, replacement of the 3-methyl-L-histidine moiety in FeP(3-MHis) by 1-methyl-L-histidine caused a 35-fold reduction in O₂ affinity; the $P_{1/2}$ value of HSA-FeP(1-MHis) (22 Torr, 37 °C, pH 7.4) is almost identical to that of human red blood cells. Results of kinetic studies indicate that the low O₂ binding affinity of FeP(1-MHis) is predominantly manifested in the high O₂ dissociation rate constant. In a toluene solution, an identical relationship in the O₂ binding property was similarly observed for FeP(1-MHis) and FeP(3-MHis). The axial Fe-N(1-MHis) coordination might be restrained by steric interaction between the 4-methylene group of the histidine and the porphyrin plane.

Numerous synthetic model hemes have been prepared to mimic the O₂ binding capability of hemoglobin (Hb) and myoglobin (Mb) over the past few decades (1–3). In these highly modified iron(II) porphyrins, axial coordination of the nitrogenous base plays a crucial role in regulating O₂ binding affinity. Collman and co-workers demonstrated that picket-fence porphyrin¹ ligating a 1,2-dimethylimidazole (1,2-DMIm) showed a 70-fold lower O₂ binding affinity [$P_{1/2}$ (O₂ partial pressure at

which 50% of porphyrin was dioxygenated) = 38 Torr, 25 °C] than that of the complex with 1-methylimidazole (1-MIm) (4, 5). This reduction was ascribed to the steric interaction between the 2-methyl group of the imidazole and the porphyrin plane. Furthermore, they showed that the replacement of imidazole by pyridine in the tailed picket-fence porphyrin lowered the O₂-binding affinity by a factor of 40 (6). The weaker π -basicity of pyridine and the increased steric repulsion of the six-atom pyridine ring compared to that of the five-atom imidazole reduced the O₂ binding of the iron(II) porphyrins (7–10). We previously reported that 2-[[8-*N*-(methylimidazolyl)octanoyloxy]methyl]-5,10,15,20-tetrakis($\alpha,\alpha,\alpha,\alpha$ -*o*-(1'-methylcyclohexanamido)phenyl)porphinatoiron(II) [FeP(2-MIm)] formed a stable O₂ adduct in toluene solution, and human serum albumin (HSA) incorporating FeP(2-MIm) [HSA-FeP(2-MIm)]

* Corresponding author. Tel: +81-3-5286-3120. Fax: +81-3-3205-4740. E-mail: eishun@waseda.jp (E.T.). E-mail: teruyuki@waseda.jp (T.K.).

[†] Waseda University.

[‡] Japan Science and Technology Agency (JST).

¹ Picket-fence porphyrin: 5,10,15,20-tetrakis($\alpha,\alpha,\alpha,\alpha$ -*o*-pivalamido-phenyl)porphinatoiron.

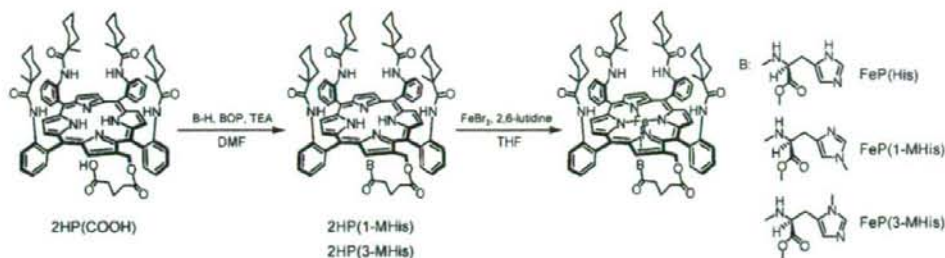


Figure 1. Synthetic scheme of iron(II) porphyrin bearing a covalently linked histidine derivative as an axial base at the β -pyrrolic position.

can bind O_2 not only in aqueous media (pH 7.4) but also in the blood stream as an artificial red blood cell (RBC) substitute (11, 12). For practical biomedical applications of the synthetic iron(II) porphyrin as an O_2 carrier, two physiological requirements exist. First, O_2 binding affinity should be close to that of human RBC ($P_{1/2} = 27$ Torr, 37 °C). Second, it is favorable to use natural histidine as an axial base because some imidazole and pyridine derivatives show toxicity to respiratory organs and the central nervous system (13). Nevertheless, few studies have addressed O_2 binding of the synthetic model heme having a histidyl group (14–16). We have demonstrated the O_2 binding equilibrium and kinetics of 5,10,15,20-tetrakis($\alpha,\alpha,\alpha,\alpha$ -o-(1'-methylcyclohexanamido)phenyl)porphyrinatoiron(II) with an ω -histidyl-alkyl chain [FeP(His)²] (17). A new class of biocompatible O_2 carriers would be realized if we can regulate O_2 binding affinity by changing the steric and electronic natures of the attached histidine. We now report a significant difference in the O_2 binding properties of HSA hybrid with synthetic iron(II) porphyrin bearing a covalently linked 1-methyl-L-histidine or 3-methyl-L-histidine [HSA-FeP(1-MHis) and HSA-FeP(3-MHis)] (Figure 1) under physiological conditions (pH 7.4, 37 °C).

The 1-methyl-L-histidine methylester or 3-methyl-L-histidine methylester was introduced into the carboxyl group of the parent free-base porphyrin, 2-((4-carboxybutanoyloxy)methyl)-5,10,15,20-tetrakis($\alpha,\alpha,\alpha,\alpha$ -o-(1'-methylcyclohexanamido)phenyl)porphyrin [2HP(COOH)] (11), via an amide linkage, yielding histidine-terminated porphyrins (>60% yield) (Figure 1). The cyclohexanoyl substituents stabilized the O_2 adduct complex rather than the pivaloy group (11). The central iron was inserted as iron(II) using $FeBr_2$ and 2,6-lutidine in anhydrous THF. The analytical data of all compounds were satisfactorily obtained (see Supporting Information).

Both FeP(1-MHis) and FeP(3-MHis) are readily incorporated into HSA (Mw: 66.5 kDa), producing stable artificial hemoproteins: HSA-FeP(1-MHis) and HSA-FeP(3-MHis) [in phosphate-buffered saline (PBS) solution at pH 7.4, [porphyrin]/[HSA] = 4 (mol/mol)]. The UV–vis absorption spectra of the aqueous HSA-FeP(1-MHis) and HSA-FeP(3-MHis) solutions under an N_2 atmosphere showed a similar feature of HSA-FeP(2-MIm) and HSA-FeP(His) (Figure 2) (Table S1) (11, 17). The spectral shapes resembled those of the five-N-coordinate high-spin ferrous complex of picket-fence porphyrin and other tetraphenylporphyrin derivatives (5). These results indicated that FeP(1-MHis) and FeP(3-MHis) formed a five-N-coordinate high-spin complex with an intramolecularly coordinated methyl-

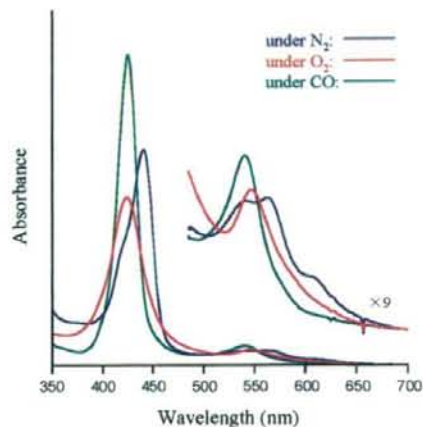
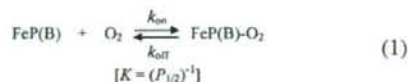


Figure 2. UV–vis absorption spectral changes of HSA-FeP(1-MHis) in PBS solution at 25 °C.

L-histidine moiety in the protein matrix. Upon bubbling of O_2 gas into the solution, the absorption pattern changed immediately to that of the O_2 adduct complex. The dioxygenations of these artificial hemoproteins have been observed reversibly under physiological conditions (pH 7.4, 37 °C). After passing the CO gas, the absorption maxima shifted to that of the carbonyl complex.

The O_2 binding affinities [$K = (P_{1/2})^{-1}$] of HSA-FeP(1-MHis) and HSA-FeP(3-MHis) were determined by the absorption spectral changes at various O_2 concentrations (eq 1).



The HSA-FeP(3-MHis) showed an extraordinarily high O_2 binding affinity ($P_{1/2} = 0.2$ Torr, 25 °C), which is 5-fold greater than that of HSA-FeP(His), which was close to the values of the relaxed-state Hb(α) and Mb (Table 1) (18–20). However, replacement of the 3-methyl-L-histidine moiety in FeP(3-MHis) by the 1-methyl-L-histidine isomer [FeP(1-MHis)] caused a 35-fold reduction of O_2 binding affinity ($P_{1/2} = 7$ Torr, 25 °C); the $P_{1/2}$ value of 22 Torr at 37 °C was almost identical to that of human RBC (27 Torr) (21).

Laser flash photolysis experiments gave the association and dissociation rate constants for O_2 (k_{on} , k_{off}) (5, 11). The time course of the absorption change accompanying the O_2 recombinations to HSA-FeP(1-MHis) and HSA-FeP(3-MHis) after the laser pulse irradiation comprised two phases of first-order kinetics (Figure S1a). The decay was fitted by a double exponential profile, which provided the fast and slow association rate constants of O_2 (k_{on} and k'_{on}) (Table 1). This behavior was similarly observed in HSA-

² Abbreviations: FeP(His), 2-((4-methoxycarbonyl-histidinamidobutanoyloxy)methyl)-5,10,15,20-tetrakis($\alpha,\alpha,\alpha,\alpha$ -o-(1'-methylcyclohexanamido)phenyl)porphyrinatoiron; FeP(1-MHis), 2-[[4-methoxycarbonyl(1-methyl)histidinamidobutanoyloxy]methyl]-5,10,15,20-tetrakis($\alpha,\alpha,\alpha,\alpha$ -o-(1'-methylcyclohexanamido)phenyl)porphyrinatoiron; FeP(3-MHis), 2-[[4-methoxycarbonyl(3-methyl)histidinamidobutanoyloxy]methyl]-5,10,15,20-tetrakis($\alpha,\alpha,\alpha,\alpha$ -o-(1'-methylcyclohexanamido)phenyl)porphyrinatoiron.

Table 1. O₂ Binding Parameters of Human Serum Albumin Hybrid with Iron(II) Porphyrin Bearing a Methyl-L-Histidine Isomer in PBS Solution (pH 7.4) at 25 °C

HSA-iron(II) porphyrin	k_{on} (M ⁻¹ s ⁻¹)	k'_{on} (M ⁻¹ s ⁻¹)	k_{off} (s ⁻¹)	k'_{off} (s ⁻¹)	$P_{1/2}^a$ (Torr)
HSA-FeP(1-MHis)	5.4×10^7	8.1×10^6	620	93	7 (22)
HSA-FeP(3-MHis)	5.4×10^7	6.8×10^6	20	2.4	0.2 (1)
HSA-FeP(His) ^b	5.4×10^7	8.8×10^6	89	14	1 (3)
Hb(α) (R-state) ^c	3.3×10^7 ^d		13 ^e		0.24
Mb ^{f,g}	1.4×10^7		12		0.51
RBC ^h					8 (27)

^a At 37 °C in parenthesis. ^b Ref 11. ^c Human Hb α-subunit. ^d In 0.1 M phosphate buffer (pH 7.0, 21.5 °C); ref 18. ^e In 10 mM phosphate buffer (pH 7.0, 20 °C); ref 19. ^f Sperm whale Mb. ^g In 0.1 M phosphate buffer (pH 7.0, 20 °C); ref 20. ^h Human red blood cell suspension. In isotonic buffer (pH 7.4); ref 21.

Table 2. O₂ Binding Parameters of Iron(II) Porphyrin Bearing a Methyl-L-Histidine in Toluene Solution at 25 °C

iron(II) porphyrin	k_{on} (M ⁻¹ s ⁻¹)	k_{off} (s ⁻¹)	$P_{1/2}$ (Torr)
FeP(1-MHis)	9.8×10^7	3.5×10^4	27
FeP(3-MHis)	1.6×10^8	5.2×10^2	0.3
FeP(His) ^a	2.0×10^8	4.3×10^3	1.7

^a In benzene solution, ref 17.

FeP(His) and HSA-FeP(2-MIm) (11). It has been interpreted that the O₂ associations to the iron(II) porphyrins in HSA were affected by microenvironments around the accommodation site (steric hindrance of the amino acid residue and difference in polarity) (22). The kinetic data indicated that the low O₂ binding affinity of HSA-FeP(1-MHis) is predominantly manifested in the high O₂ dissociation rate constant (Table 1).

To evaluate the significant differences in FeP(1-MHis) and FeP(3-MHis), their O₂ binding parameters in toluene solution were determined. The UV-vis absorption spectra of ferrous FeP(1-MHis) (λ_{max} , 440, 542, 563 nm) and FeP(3-MHis) (λ_{max} , 440, 540, 565 nm) under an N₂ atmosphere showed the formation of a five-N-coordinate high spin complex. Dioxygenation was sufficiently stable and reversible at 25 °C depending on O₂ partial pressure. The recombination process of O₂ to the iron(II) porphyrins in toluene solution after laser flash photolysis was fitted by a single exponential (Figure S1b). FeP(3-MHis) exhibited a 6-fold higher O₂ binding affinity ($P_{1/2}$ = 0.3 Torr, 25 °C) compared to that of FeP(His) (Table 2) (11). Kinetically, this high O₂ binding affinity is attributable to the low O₂ dissociation rate. The high σ -basicity of 3-methyl-L-histidine (pK_a, 6.48) in comparison to that of histidine (pK_a, 6.00) (23) can serve to increase the electron density on the central metal, which would enhance O₂ uptake. FeP(1-MHis) showed a 90-fold lower O₂ affinity ($P_{1/2}$ = 27 Torr, 25 °C) relative to FeP(3-MHis), which is mainly caused by the high k_{off} value, as observed in aqueous media. In general, the steric hindrance to the axial base coordination deforms the six-coordinate structure of the iron(II) porphyrin and increases the O₂ dissociation rate constant (1, 2). For example, the 1,2-DMIm complexes showed 11–58-fold higher k_{off} values than the 1-MIm ligated complexes (3, 5, 24).

We simulated the geometry of histidine coordination in dioxygenated porphyrin.³ The bond angle of N(por)–Fe–N(His) was 90° for FeP(His) and FeP(3-MHis), but it was markedly tilted in FeP(1-MHis) (a maximum of 98°) (Figure 3). On the basis of these results, it can be concluded that the axial coordination of 1-methyl-L-histidine to the central iron is

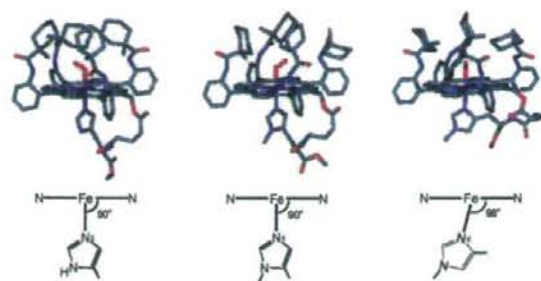


Figure 3. Simulated structures of the O₂ adduct complexes of FeP(His), FeP(1-MHis), and FeP(3-MHis). In FeP(His) and FeP(3-MHis), the bond angles of N(por)–Fe–N(His) were 90°, but 98° in FeP(1-MHis) because of the steric interaction between the 4-methylene group of histidine and the porphyrin plane. Molecular dynamics and minimization (force field: esff) were performed using an Insight II system. Hydrogens were omitted for clarification.

restrained by the steric interaction between the 4-methylene group of the histidine and the porphyrin plane, which significantly increased the dissociation rate of O₂.

In conclusion, a variation in the methyl-L-histidine isomer covalently linked at the porphyrin periphery changed the O₂ binding affinity of the iron(II) porphyrin by a factor of 90 in toluene. This difference is greater than the previous finding for a picket-fence porphyrin (2-MIm) complex-bound O₂ with a 70-fold lower affinity than that of the 1-MIm complex (4). The increased k_{off} value of FeP(1-MHis) is caused by steric repulsion between the 4-methylene group of histidine and the porphyrin plane. In aqueous media, the O₂ binding ability of HSA incorporating the iron(II) porphyrin can also be modulated by structural isomerization of the axial methyl-L-histidine. HSA-FeP(1-MHis) with a $P_{1/2}$ value similar to that of human RBC under physiological conditions can become a promising O₂ carrier, which satisfies both the biocompatibility and clinical requirements for efficient O₂ delivery to the tissue cells.

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Supporting Information Available: Experimental details, absorption maximum wavelengths of the iron(II) porphyrins, and absorption change accompanying O₂ rebinding to FeP(3-MHis) and HSA-FeP(3-MHis) after laser flash photolysis. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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³ The esff forcefield simulation was performed using an Insight II system (Molecular Simulations Inc.). The structure was generated by alternative minimizations and annealing dynamic calculations from 1,000 to 100 K.

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Heme Pocket Architecture in Human Serum Albumin: Regulation of O₂ Binding Affinity of a Prosthetic Heme Group by Site-Directed Mutagenesis

Teruyuki Komatsu,^{*1,2} Akito Nakagawa,¹ Eishun Tsuchida^{*1}

Summary: We present the O₂ binding properties of recombinant human serum albumin (rHSA) mutants complexed with an iron(II) protoporphyrin IX as a prosthetic heme group. Iron(III) protoporphyrin IX (hemin) is bound within subdomain IB of HSA with weak axial coordination by Tyr-161. In order to confer O₂ binding capability to this naturally occurring hemoprotein: (i) a proximal histidine was introduced into position Ile-142; and (ii) the coordinated Tyr-161 was replaced with hydrophobic Leu using site-directed mutagenesis. It provided a recombinant HSA double-mutant [rHSA(I142H/Y161L) = rHSA(HL)]. The rHSA(HL)-heme formed a ferrous five-coordinate high-spin complex with axial ligation of His-142 under an Ar atmosphere. This artificial hemoprotein binds O₂ at room temperature. Laser flash photolysis experiments demonstrated that O₂ rebinding to rHSA(HL)-heme displays monophasic kinetics, whereas the CO recombination process obeyed a double-exponential pattern. This might be attributable to the two different geometries of the axial imidazole coordination arising from the two orientations of the porphyrin plane in the heme pocket. The O₂ binding affinity of rHSA(HL)-heme was considerably lower than those of R-state hemoglobin (Hb) and myoglobin (Mb), principally because of the high O₂ dissociation rate constant. The third mutations have been introduced into the distal side of the heme (at position Leu-185 or Arg-186) to increase the O₂ binding affinity. The rHSA(HL/L185N)-heme showed high O₂ binding affinity ($P_{1/2}^{O_2}$: 1 Torr), which is 18-fold greater than that of the original double mutant rHSA(HL)-heme and which is rather close to those of Hb (R-state) and Mb. Furthermore, replacement of polar Arg-186 with Leu or Phe adjusted the O₂ binding affinity ($P_{1/2}^{O_2}$) to 10 Torr, which is almost equivalent to value for human red blood cells.

Keywords: biomimetics; heme; human serum albumin; O₂ binding; proteins

Introduction

Human serum albumin (HSA), the most abundant plasma protein (4–5 g/dl) in our circulatory system, is characterized by its remarkable ability to bind widely various endogenous and exogenous compounds^[1]

such as fatty acids, bilirubin, bile acids, thyroxine,^[2,3] and a wide range of drugs.^[4] Hemin [iron(III) protoporphyrin IX] released from methemoglobin is also captured by HSA with a high binding constant ($K \approx 10^8 \text{ M}^{-1}$).^[5] This strong affinity of HSA for hemin has stimulated efforts to develop albumin as an artificial hemoprotein which can mimic the O₂ binding ability of hemoglobin (Hb) and myoglobin (Mb).^[6,7] HSA consists of a helical monomer of 66.5 kDa containing three homologous domains (I–III), each of which comprises of A and B subdomains.^[8] Crystallographic

¹ Research Institute for Science and Engineering, Waseda University, 3-4-1 Okubo, Shinjuku-ku, Tokyo 169-8555, Japan

Fax: (+81) 3-3205-4740;

E-mail: teruyuki@waseda.jp

² PRESTO, Japan Science and Technology Agency (JST)

studies have revealed that heme is bound within a narrow D-shaped hydrophobic cavity in subdomain IB with axial coordination of Tyr-161 to the central ferric ion and electrostatic interactions between the porphyrin propionates and a triad of basic amino acid residues (Arg-114, His-146 and Lys-190) (Figure 1).^[9,10] In terms of the general hydrophobicity of this α -helical heme pocket, the subdomain IB of HSA potentially has similar features to the heme binding site of Hb or Mb. However, if one reduces the HSA–hemin to obtain the ferrous complex, it is rapidly oxidized by O₂, even at low temperature, because HSA lacks the proximal histidine that enables the prosthetic heme group to bind O₂ and serves to regulate the O₂ binding affinity (Figure 1). In order to confer the O₂ binding capability to this naturally occurring hemoprotein, we have introduced a proximal histidine into the heme binding site of HSA by site-directed mutagenesis; it would provide axial coordination to the central ferrous ion of the heme and thereby promote O₂ binding.^[11] Moreover, to modulate its O₂ binding affinity, we have added further modification to the distal side of the heme. The O₂ binding properties of several rHSA(mutant)–heme complexes have been characterized kinetically and compared to those of the natural Hb, Mb, and red blood cells (RBC). We have shown that our mutagenesis approach can create a new

class of albumin-based artificial hemoprotein which would serve as an O₂ carrier.

Results and Discussion

Double-Mutations to Confer the O₂ Binding Capability

The detailed structure of the heme binding site in HSA revealed by crystallographic studies allows the design of mutagenesis experiments to construct a tailor-made heme pocket for O₂ binding.^[9,10] In fact, Tyr-161 was the first candidate to introduce a proximal histidine (Figure 1). However, the Y161H mutation was not done because our simulation indicated that the distance from His-161 to the central Fe would be too great (4.0 Å). Instead, modeling experiments suggested that the favorable positions for the axial imidazole insertion would be Ile-142 (Figure 2a). The N(histidine)–Fe distance was estimated as 2.31 Å for H142 (compared to 2.18 Å for Mb). We therefore designed a recombinant HSA (rHSA) double-mutant I142H/Y161L [=rHSA(HL)].

The specific mutations were introduced into the HSA coding region in the plasmid vector (pHIL-D2 HSA) using the Quik-Change mutagenesis kit (Stratagene), and clones were expressed in the yeast *Pichia pastoris* (Invitrogen Corp.). The rHSA–hemin complexes were prepared funda-

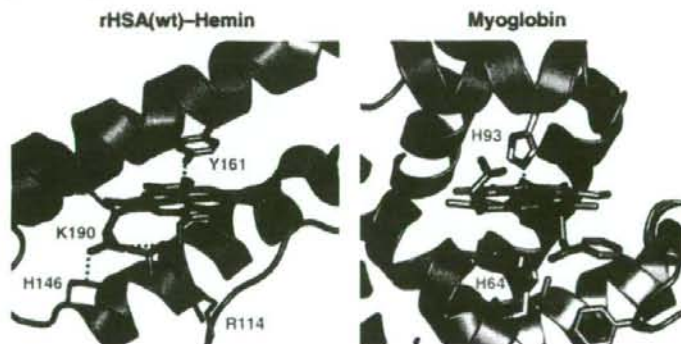


Figure 1. Heme pocket structure in subdomain IB of HSA (left; 1O9X from ref. [9]) and heme pocket structure of Mb (1MBO).

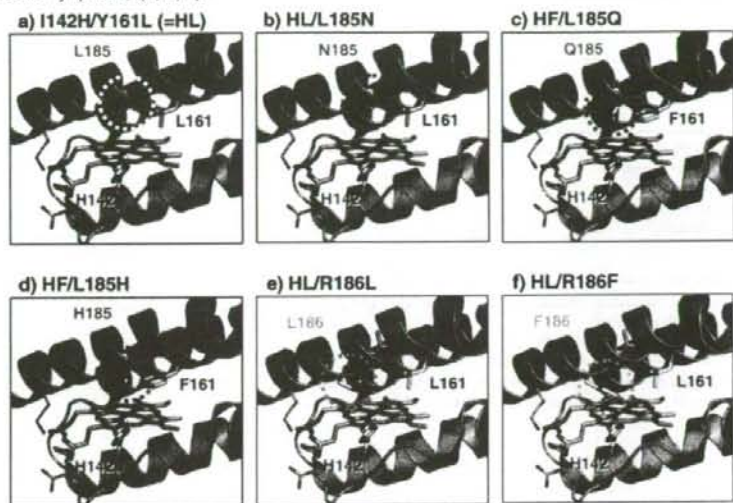


Figure 2. Structural models of the heme pocket in rHSA(mutant)-heme complexes.

mentally according to our previously reported procedures.^[9,11]

The MCD spectra of the rHSA(HL)-hemin showed S-shaped patterns in the Soret band region, which resembled that of ferric Mb.^[12] One water molecule is known to coordinate axially to the sixth position of the central ferric ion of the heme in metMb. Our MCD result suggests that the rHSA(HL)-hemin is also in a ferric high-spin complex, having a water molecule as the sixth ligand.

O₂ Binding Affinity of rHSA(HL)-Heme

The rHSA(HL)-hemin was reduced to a ferrous complex by adding a small molar excess of aqueous sodium dithionite under an Ar atmosphere. A single broad absorption band (λ_{max} : 559 nm) in the visible region of rHSA(HL)-heme resembled that observed for deoxy Mb^[13] or the chelated protoheme in DMF,^[14] indicating the formation of a five-coordinate high-spin complex. The heme therefore appears to be accommodated in the mutated heme pocket with an axial coordination involving His-142. Upon exposure of the rHSA(HL)-

heme solution to O₂, the UV-vis absorption immediately changed to that of the O₂ adduct complex at 0–25 °C.^[13,14] After introduction of CO gas, the hemoprotein produced a stable carbonyl complex.

Laser flash photolysis experiments were carried out to evaluate the kinetics of the O₂ binding to the rHSA(HL)-heme.^[11,15,16] The absorbance decay accompanying the O₂ recombination to rHSA(HL)-heme was composed of single-exponential. On the other hand, the rebinding of CO followed biphasic decay, which is normally not observed in Mb. Results of numerous investigations of synthetic model hemes have shown that a bending strain in the proximal base coordination to the central ferrous ion, the “proximal-side steric effect”, can decrease the association rate for CO without greatly altering the kinetics of O₂ association.^[15,16] Therefore, a possible explanation is that there are two different geometries of the axial histidine (His-142) coordination to the central ferrous ion of the heme in rHSA(HL), each one accounting for the biphasic kinetics of CO rebinding.

Table 1.
O₂ binding parameters of rHSA(mutant)-heme complexes.^a

Hemoproteins	10 ⁻⁶ k _{on} ^{O₂} (M ⁻¹ s ⁻¹)	10 ⁻³ k _{off} ^{O₂} (s ⁻¹)		p _{1/2} ^{O₂} (Torr)	
		I	II	I	II
rHSA(HL)-Heme	7.5	0.22	1.70	18	134
rHSA(HL/L185N)-Heme	14	0.02	0.29	1	14
rHSA(HL/R186L)-Heme	25	0.41	8.59	10	209
rHSA(HL/R186F)-Heme	21	0.29	7.01	9	203
Hb(α) (R-state) ^b	33 ^c	0.013 ^d		0.24	
Mb ^e	14	0.012		0.51	
RBC ^f				8	

^a In 50 mM potassium phosphate buffered Solution (pH 7.0) at 22 °C. I or II indicates species I or II.

^b Human Hb α-subunit.

^c In 0.1 M phosphate buffer (pH 7.0, 21.5 °C); ref. [17].

^d In 10 mM phosphate buffer (pH 7.0, 20 °C); ref. [18].

^e Sperm whale Mb. In 0.1 M potassium phosphate buffer (pH 7.0, 20 °C); ref. [19].

^f Human red cell suspension. In isotonic buffer (pH 7.4, 20 °C); ref. [23].

By analyzing the CO/O₂ competitive binding following laser flash photolysis,^[15,16] we obtained the association rate constants for O₂ (k_{on}^{O₂}) and the O₂ binding affinities [P_{1/2}^{O₂} = (K^{O₂})⁻¹] for rHSA(HL)-heme (Table 1).^[11] The faster phase, defined as species I, and the slower phase, defined as species II, yielded two different O₂ binding affinities. In species I, the proximal His might coordinate to the central ferrous ion without strain, whereas in species II, the ligation might involve some distortion, resulting in weaker O₂ binding (Figure 3).

The p_{1/2}^{O₂} value of rHSA(HL)-heme was determined to be 18 Torr (species I), which was 35–75-fold higher (O₂ binding affinity is lower) than those of Hbα (R-state) and Mb.^[17–19] This low affinity for O₂ was kinetically attributable to a 17–18-fold

increases in the O₂ dissociation rate constants. The O₂ binding affinity should be adjusted to similar values for Hb and human RBC to develop this artificial hemoprotein as a blood substitute.

Introduction of Distal Base into the 185 Position

The His-64 in Hb and Mb on the distal side of the heme plays an important role for tuning their ligand affinities. A neutron diffraction study of MbO₂ showed that the N–H bond of the distal His-64 is restrained from optimal alignment for strong H-bonding with the coordinated O₂.^[20] Olson and co-workers reported that the substitution of Gly for His-64 in Mb and Hb(α) caused a marked decrease in the O₂ binding affinity.^[21] In view of these investigations, we reasoned that systematic variation of the

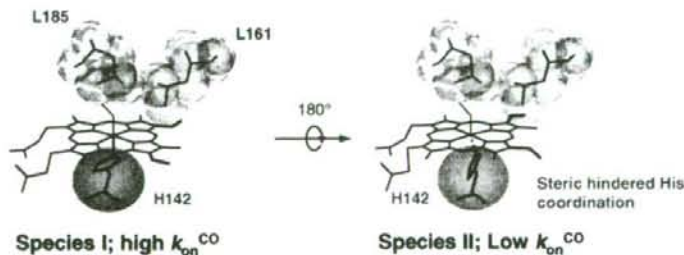


Figure 3.
Structural models of heme-O₂ site of rHSA(HL)-heme complex.

steric hindrance and local polarity of the heme pocket in subdomain IB of HSA would allow modulation of the O₂ binding affinity. One approach to enhancing the O₂ binding affinity of rHSA–heme would be to introduce a basic amino acid into an appropriate position on the distal side of the heme. Our modeling results showed that the favorable position for the distal base insertion was Leu-185. Consequently, we replaced Leu-185 in rHSA(HL) [or rHSA(I142H/Y161F) [=rHSA(HF)]] with Asn, Gln, and His using site-directed mutagenesis (Figure 2b–2d).^[22]

The rHSA(HL/L185N)–heme under Ar atmosphere showed a visible absorption band at 559 nm with a small shoulder at 530 nm, which was similar to the spectrum observed for rHSA(HL)–heme,^[11] deoxy Mb,^[13] and chelated protoheme.^[14] The spectral pattern clearly indicated the formation of a five-coordinate high-spin complex. On the contrary, in the spectra of rHSA(HF/L185Q)–heme and rHSA(HF/L185H)–heme, the β band at 528 nm appeared relatively sharp, suggesting partial formation of a six-coordinate heme complex. Upon exposure of the rHSA(HL/L185N)–heme solution to O₂, the UV-vis absorption changed immediately to that of the O₂ adduct complex at 22 °C. In contrast, rHSA(HF/L185Q)–heme and rHSA(HF/L185H)–heme were oxidized by O₂, even at low temperature (5 °C). After introducing CO gas, all the hemoproteins produced stable carbonyl complexes with identical absorption spectral patterns.

Marked differences are apparent in the comparison of the O₂ binding parameters for rHSA(HL)–heme and rHSA(HL/L185N)–heme. The presence of Asn rather than Leu at position 185 resulted in 18-fold and 10-fold increases in the O₂ binding affinity, respectively, for species I and II (Table 1). These increases were predominantly attributable to the 6–11-fold diminution of the $k_{\text{off}}^{\text{O}_2}$ values. The high O₂ binding affinity ($P_{1/2}^{\text{O}_2}$: 1 Torr) for rHSA(HL/L185N)–heme is now close to that of Hb (R-state) (0.24 Torr) and Mb (0.5 Torr) (Table 1).

Introduction of Leu or Phe into the 186 Position

For rHSA–heme to provide effective O₂ transport *in vivo*, the affinity should be more similar to that of human RBC ($P_{1/2}^{\text{O}_2}$: 8 Torr).^[23] This requires an O₂ binding affinity that is intermediate between the values of rHSA(HL)–heme and rHSA(HL/L185N)–heme. An effective means to control the O₂ binding affinity of the heme is introduction of a different polar amino acid around the O₂ binding site. A polar Arg-186 exists at the entrance of the heme pocket; we expected that insertion of a nonpolar residue at this position would change the O₂ binding affinity of rHSA–heme. Consequently, we designed new triple mutants, rHSA(HL/R186L) and rHSA(HL/R186F) (Figure 2e, 2f).

The MCD in the Soret band region of the ferric rHSA(HL/R186L)–hemin and rHSA(HL/R186F)–hemin both showed low intensity, which is fundamentally equivalent to that observed for rHSA(HL)–hemin. The reduced ferrous form demonstrated the characteristic UV-vis absorption and MCD spectra of the five-coordinate high-spin complex under an Ar atmosphere. Upon bubbling O₂ gas through the solutions, the spectral patterns were shifted to that of the O₂ adduct complex. The distinct features of all the spectra were quite similar to those of the rHSA(HL)–heme. Fortunately, the O₂ binding affinities of rHSA(HL/R186L)–heme and rHSA(HL/R186F)–heme were more similar to that of human RBC ($P_{1/2}^{\text{O}_2}$: 8 Torr) (Table 1). We can conclude that the Arg-186 is an important key amino acid to control the O₂ binding property of the heme and the obtained triple mutants could become RBC substitutes.

Conclusion

We have shown clearly that rHSA–heme can be engineered to bind O₂ reversibly. However, the complex did not display optimal O₂ binding affinity. By emphasizing modification on the distal side of the heme pocket in rHSA, we have prepared distinct

rHSA(triple mutant)-heme complexes with a broad range of O₂ binding affinities. The highest affinity mutant rHSA(HL/L185N) contains Asn-185, which has a short amide side-chain that enhances the O₂ binding affinity. On the other hand, introduction of the larger Gln and His side-chains at position 185 partly provided a six-coordinate heme character, and did not stabilize O₂ binding. In a different approach, substitution of Arg-186 at the entrance of the heme pocket with Leu or Phe provided a useful reduction in the O₂ binding affinity, yielding $P_{1/2}^{O_2}$ values that are closely resemble that of the human RBC.

The transport of O₂ by rHSA-heme could be of great clinical importance, not only as a blood substitute, but also as an O₂-providing therapeutic reagent. If the HSA-based O₂ carrier is realized, it has the potential of acting not only as a RBC substitute, but also as an O₂ providing therapeutic reagent.

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