

Protein Photoconductors and Photodiodes**

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Long-range electron transfer (ET) through proteins plays important roles in living systems. ET proteins can be regarded as wide-band-gap molecular semiconductors with “dopants” (redox or photoactive species) that provide an electron- or hole-localizing site, though the protein scaffold has insulating character.^[1–10] Intramolecular ET in proteins has been investigated in detail by many groups to elucidate the factors that control the rates of these nonadiabatic reactions.^[1–9,11–13] Dutton and co-workers proposed a square barrier ET rate model,^[11,12] while Gray and colleagues showed that ET depends on the structure of the medium between an electron donor and acceptor.^[5–9]

We have investigated photoinduced ET in two proteins, zinc-substituted cytochrome *b*₅₆₂ (Zn-cyt *b*₅₆₂) and zinc-substituted cytochrome *c* (Zn-cyt *c*), the semiconductor properties of which depend on the charge distribution on their molecular surfaces: Zn-cyt *b*₅₆₂ has n-type semiconductor character while Zn-cyt *c* is p-type, although the active center of the two systems is virtually identical. This finding may open up the world of protein-based electronics, because the semiconductor character of proteins could be controlled by variations in surface charge.

Zn-cyt *b*₅₆₂ on the H₂N-SAM/Au electrode (SAM = self-assembled monolayer) was prepared according to the literature (see the Supporting Information). The orientation of Zn-cyt *b*₅₆₂ on the H₂N-SAM/Au surface (Figure 1) is optimal for favorable heme-region (acidic patch) electrostatic interactions with the SAM positive charge.^[14]

Figure 2a shows cathodic photocurrents in response to switching on and off of 420 nm illumination for bias voltages of +300, 0, and –300 mV (vs. Ag|AgCl). The action spectrum (Figure 2b) is similar to the absorption spectrum of Zn-cyt *b*₅₆₂ (Figure 2b, inset), which suggests that the photocurrent originates in the protein. Incidentally, there is no photocurrent response in the case that protein changes Zn-cyt *b*₅₆₂ into Fe-cyt *b*₅₆₂. Notably, the Q-band (550 nm) photocurrent response is more prominent in comparison with the

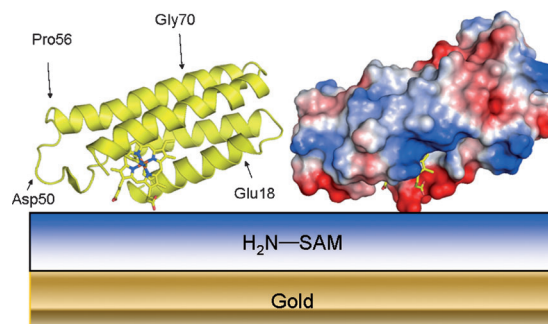


Figure 1. Schematic view of Zn-cyt *b*₅₆₂ immobilized on H₂N-SAM fabricated on a gold electrode. Left: ribbon model with the porphyrin moiety in rod model; right: charge distribution of Zn-cyt *b*₅₆₂. Red and blue indicate negative and positive charges, respectively.

absorption spectrum, that is, the Soret/Q ratio in absorbance (I_{420}/I_{550}) of the action spectrum is 3.7 whereas that of the UV/Vis spectrum is 11.1, thus implying that the photocurrent is derived from migration of the photoexcited electron.^[15,16] These findings are consistent with this mechanism: 1) the photocurrent was observed only in the cathodic direction under the range of bias potential employed here (Figure 2c), and 2) addition of methyl viologen ($E_0 = -0.62$ V vs. Ag|AgCl) to the solution as a redox reagent enhanced the cathodic photocurrent, while addition of potassium ferri/ferrocyanide ($E_0 = +0.17$ V vs. Ag|AgCl) did not show such an effect.^[17,18]

This is quite in contrast to the Zn-cyt *c*/HOOC-SAM/Au system, where the photocurrent action spectrum can be superimposed on the UV/Vis spectrum and the addition of the ferri/ferrocyanide effectively increases the photocurrent that derives from the hole in the occupied molecular orbitals (MOs), or the valence band.^[13] Thus, although Zn-cyt *b*₅₆₂ and Zn-cyt *c* are derivatives from similar ET proteins, the systems have a different electronic nature: a photodiode (n-type) and a photoconductor (p-type) character, respectively (Figure 2c).

To elucidate the electronic structure and the factors that control the semiconductor properties of the two Zn-cytochrome proteins, we carried out all-electron DFT calculations for Zn-cyt *b*₅₆₂. We estimated the photoinduced ET rate constants for Zn-cyt *c* [Eq. (1)]:

$$k_{i \rightarrow f} = 2\pi/\hbar \times |H_{fi}|^2 \delta(E_{fi}) \quad (1)$$

where H_{fi} is the electronic coupling matrix between effective initial and final MOs, and E_{fi} is the corresponding orbital energy difference.^[13,19]

It is noteworthy that the energy levels of MOs undergo aggregation with a band gap (ca. 2 eV) like a typical

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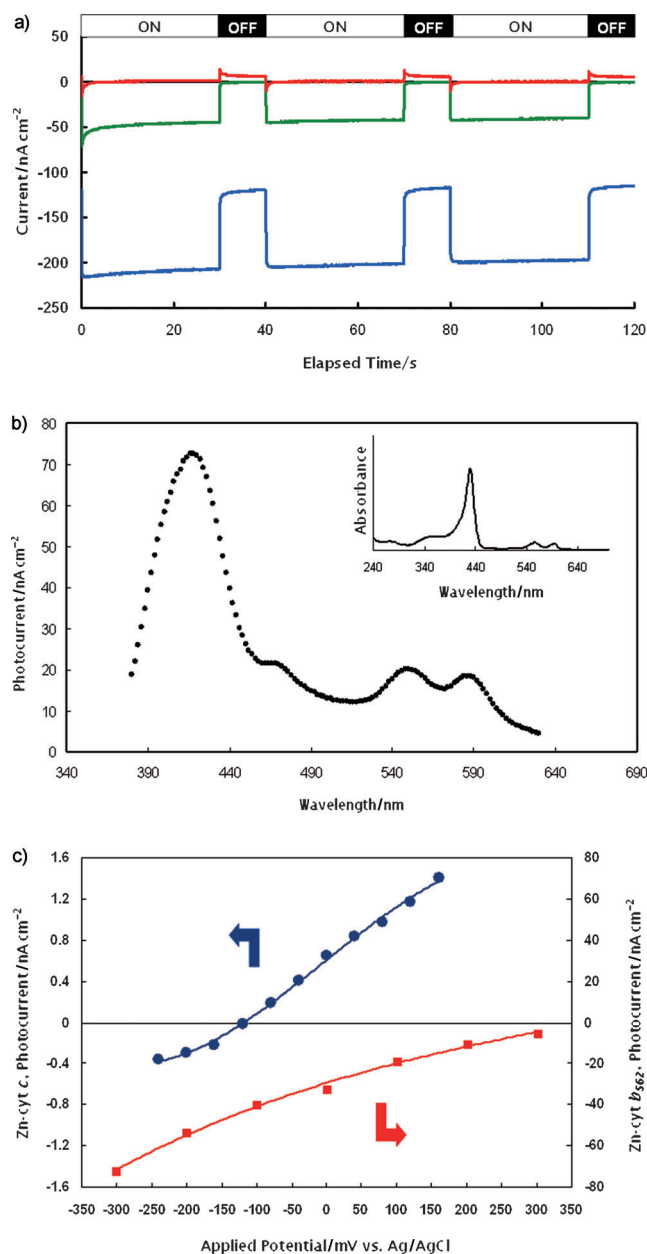


Figure 2. a) Photocurrents of the Zn-cyt *b*₅₆₂/H₂N-SAM/Au electrode upon illumination at 420 nm in N₂-saturated 10 mM sodium phosphate buffer (pH 7.0). Applied bias potentials (vs. Ag|AgCl) are +300 (red), 0 (green), and -300 mV (blue). b) Plots of the photocurrents of the Zn-cyt *b*₅₆₂/H₂N-SAM/Au electrode in N₂-saturated 10 mM sodium phosphate buffer (pH 7.0). The applied potential was held at 300 mV (vs. Ag|AgCl). Inset: UV/Vis absorption spectrum of Zn-cyt *b*₅₆₂ in the buffer solution. c) Plots of photocurrent against applied bias potential for the Zn-cyt *b*₅₆₂/H₂N-SAM/Au (red) and Zn-cyt *c*/HOOC-SAM/Au (blue) electrodes in N₂-saturated 10 mM sodium phosphate buffer (pH 7.0).

semiconductor's band structure, which is also seen in previously studied cyt *c* and Zn-cyt *c* systems.^[13,20] The calculated band gap (ca. 2 eV) is somewhat smaller than the estimated value from polypeptide (ca. 4 eV).^[8] This might be attributed to the difference between the target sizes for estimation, polypeptide (small) and protein (large), and the effect of a DFT method that is known to underestimate the band

gap.^[20,21] Among the 8758 MOs, those responsible for photoexcitation of the porphyrin moiety (Gouterman's four-orbital model)^[22] in Zn-cyt *b*₅₆₂ can be assigned to occupied MOs 3302 and 3304 having porphyrin π nature with zinc-sulfur π -bonding and unoccupied MOs 3326 and 3329 having porphyrin π^* nature (Figure S2 and Table S2 in the Supporting Information), on the basis of the time-dependent DFT calculation previously obtained (Figure S3, Table S2).^[13]

The intramolecular ET rate constants of photoexcited Zn-cyt *b*₅₆₂ were estimated from Equation (1) and related to Gouterman's 4-orbitals.^[13,19,21,23] As is seen in the case of Zn-cyt *c* (Figure S5 in the Supporting Information), distinct ET rates for specific couples of MOs in Zn-cyt *b*₅₆₂ are observed (Figure S4). Dominant intramolecular ET rate constants for each of Gouterman's 4-orbitals of Zn-cyt *b*₅₆₂ and Zn-cyt *c* are summarized in Table 1. In Zn-cyt *b*₅₆₂, the rate constants

Table 1: Calculated ET parameters for Zn-cyt *b*₅₆₂ and Zn-cyt *c*.

Proteins	MO pairs	H_{fi} [eV]	E_{fi} [eV]	$k_{i \rightarrow f}$ [s ⁻¹]
Zn-cyt <i>b</i> ₅₆₂	3302_3300	-8.52×10^{-1}	-4.41×10^{-2}	1.57×10^{17}
	3304_3307	-6.94×10^{-1}	$+1.11 \times 10^{-1}$	4.15×10^{16}
	3326_3325	-5.71×10^{-1}	-5.50×10^{-2}	5.66×10^{16}
	3329_3331	$+1.15 \times 10^0$	$+9.11 \times 10^{-3}$	1.39×10^{18}
	3329_3332	-7.33×10^{-1}	$+1.48 \times 10^{-2}$	3.46×10^{17}
Zn-cyt <i>c</i> ^[a]	3268_3270	-3.51×10^{-1}	$+2.27 \times 10^{-2}$	5.17×10^{16}
	3272_3271	-6.15×10^{-2}	-9.24×10^{-5}	3.91×10^{17}
	3297_3296	-4.91×10^{-2}	-1.59×10^{-2}	1.45×10^{15}
	3299_3298	$+9.40 \times 10^{-2}$	-1.14×10^{-1}	7.42×10^{14}

[a] Reference [13].

between unoccupied MOs are one order of magnitude larger than those between the occupied MOs. The two unoccupied MOs are localized on the outermost amino acid residues around Gly70 (MO3331) and Pro56 (MO3332), respectively, which are located on the opposite side of the protein/SAM interface and exposed to the solution (Figures 1 and 3a; Figure S2 and Table S2 in the Supporting Information). On the other hand, the two occupied MOs are localized on the Glu18 (MO3300) and Asp50 (MO3307), respectively, and both reside on the electrode-contact side (Figure 1, and Figure S2 and Table S2 in the Supporting Information). Hence the major mechanism of ET in the photoexcited Zn-cyt *b*₅₆₂ is "ET in the conduction band" through two-state coupling between unoccupied MOs, which shows that the photoexcited Zn-cyt *b*₅₆₂ is an n-type semiconductor.^[24] This is quite different from Zn-cyt *c*, where "hole transfer (HT) in the valence band" between occupied MOs dominates. This protein is a p-type semiconductor (Figure 2c, Table 1), even though the "dopant" is the same as in Zn-cyt *b*₅₆₂.

Why is one protein n-type and the other a p-type semiconductor? We analyzed the ground-state electronic structures of both proteins (Figure 3b). MO energies of Zn-cyt *b*₅₆₂ are generally higher than those of Zn-cyt *c*, which reflects a difference in charge of the two proteins: Zn-cyt *b*₅₆₂ and Zn-cyt *c* have negative (isoelectric point, pI = 5.5) and positive (pI = 9.6) charges at pH 7.0, respectively. Importantly, the effect of the negative charge of Zn-cyt *b*₅₆₂ on its 4-orbital energies is more prominent than that of the other porphyrin MOs assigned to porphyrin σ orbitals and the MOs localized on amino acid residues. The differences in MO

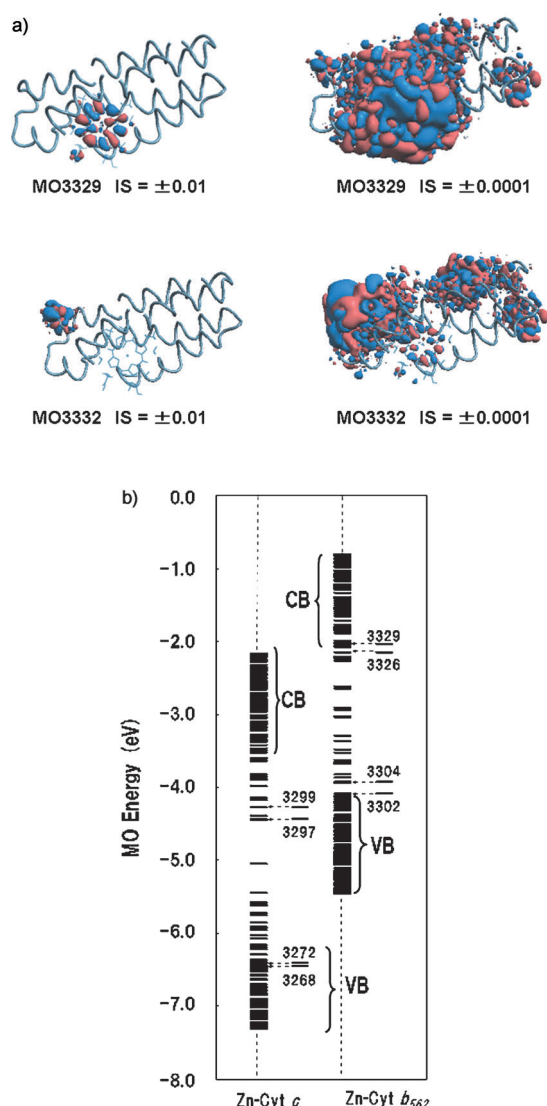


Figure 3. a) Main MOs responsible for photoexcited ET in Zn-cyt b_{562} . MO3329 (top) has the characteristics of one of Gouterman's 4-orbitals and MO3332 (bottom) originates in the protein scaffold for which the energy level is involved in the "conduction band" of the protein and is quite close to MO3329. IS: isosurface value. b) Kohn-Sham orbital distributions around the band gap of Zn-cyt b_{562} and Zn-cytc. Numbers represent Gouterman's 4-orbitals of the proteins. CB: conduction band; VB: valence band. The MO3332 shown in (a) is in the CB and its energy level (-2.02 eV) is very close to that of MO3329 (-2.04 eV).

energy levels of each corresponding 4-orbital between the two systems are almost the same, +2.34 eV, although the average energy shift from Zn-cytc to Zn-cyt b_{562} is +1.38 eV. Hence the relative energy level of the 4-orbitals against protein bands consisting of MOs localized on amino acid residues is very different for the two proteins. In Zn-cyt b_{562} , the unoccupied 4-orbitals (MOs 3326 and 3329) locate in the lower region of the protein's conduction band, whereas such an overlap is not found in Zn-cytc (Figure 3b). On the other hand, the occupied 4-orbitals (MOs 3302 and 3304) are placed at a shallow position in the higher valence band of Zn-cyt b_{562} . In comparison to Zn-cytc, the MOs with holes and the valence band with amino acid character are more weakly

coupled in Zn-cyt b_{562} (Figure 3b). These electronic structure differences between the two proteins dictate the molecular-semiconductor nature of the proteins.

Why are the 4-orbitals so much more sensitive to charge than the other MOs? We compared the diffusibilities of Gouterman's 4-orbitals and those of MOs with amino acid character (Figure 3a). Gouterman's 4-orbitals are more expansive and delocalized than the other MOs, thereby implying that the protein charge can more easily and homogeneously affect their orbital energies, even though the charged amino acid residues have an outermost distribution in the protein shell. In contrast, since the other MOs have compact and localized characters, the effect of the charges of the protein surface could be relatively small. Furthermore, due to the anionic character of the porphyrin ring,^[25] in which the gross net charge is -1.59, the 4-orbitals would be more sensitive to the charge of the protein shell.

This expandable property of the 4-orbitals can be explained by the highly ordered polarizability of the porphyrin π or π^* nature. Since the unoccupied 4-orbitals correspond to a porphyrin π^* orbital derived from the e_g orbital of metalloporphyrins with D_{4h} symmetry having quadratic function (xz , yz),^[22] they can be easily mixed with MOs of the protein shell and thus its effect can be homogeneously expanded into the whole molecule (Figure 4). In the occupied 4-orbitals, a similar expansion effect could be realized: by mixing the porphyrin π orbital derived from a_{1u} and a_{2u} orbitals not having quadratic functions^[22] with zinc-sulfur (π) hybridized by sulfur p_x or p_y and zinc d_{xz} or d_{yz} , they can acquire xz , yz characters (Figure 4).

Because the band-like structure would be a common feature among various kinds of proteins, the type of semiconductor-like behavior can be mostly described by the energy level relationship between the redox center and the protein scaffold. On the other hand, Beratan and co-workers used a donor-acceptor (D-A) coupling indicator, $|H_{fi}|^2$, to

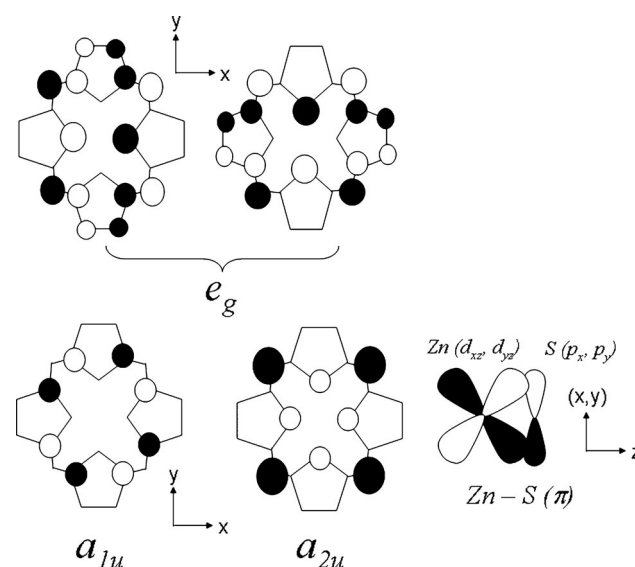


Figure 4. The π -electron densities of Gouterman's 4-orbitals derived from D_{4h} symmetry and Zn-S π -bonding nature. White and black circles are positive and negative aspects, respectively, and the areas are proportional to the π -electron densities.

analyze ET in nine ruthenized cyt b_{562} , in which they found a multiple edge-coupling regime and a dominant pathway regime.^[26,27] What we have found here is that ET pathways can be more precisely described by the combination of the energy-level relationship and D–A coupling between the responsible MOs [Eq. (1)], that is, analyses of ET rates on the basis of Fermi's Golden Rule by using all-electron wave functions of the protein. Besides the MO coupling term (H_{fi}), the proximity in the energy levels [$\delta(E_{fi})$] of donor and acceptor should be tuned for effective electron pathway(s) in the ET protein.

In summary, we have shown that two zinc-substituted cytochromes employ different photoinduced ET mechanisms. The effect of charge on the protein scaffold on the energy of Gouterman's 4-orbitals determines the semiconductor property (n- or p-type) of Zn-cyt b_{562} and Zn-cyt c . The energy shift of the 4-orbitals is highly sensitive to changes in the pI because of the expandable and delocalized nature of the MO electron density, most likely because of the highly ordered polarizability of the porphyrin π or π^* orbitals. We suggest that a protein semiconductor with a p–n junction could be constructed by fusing two suitably engineered cytochromes.

Experimental Section

Photocurrent measurements of Zn-cyt b_{562} : Photocurrent measurements were carried out using the same system as for previous Zn-cyt c gold electrode measurements.^[13] A xenon lamp (Ushio SX-UI150XQ) was used as light source, and monochromated incident light with full width at half maximum of 10 nm was generated by a monochromator (JASCO CT-151T). The incident light was blocked intermittently by a mechanical shutter and focused on the Zn-cyt b_{562} gold electrode in a quartz electrochemical cell (10 mm phosphate buffer solution; counter electrode: Pt mesh; reference electrode: Ag|AgCl). A potentiostat (Hokuto Denko HA150G) was used to control the bias voltage applied to the Zn-cyt b_{562} gold electrode against the reference electrode, and the time course of the current was recorded. All measurements were carried out under nitrogen after 15 min nitrogen bubbling of 10 mm sodium phosphate buffer solution.

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