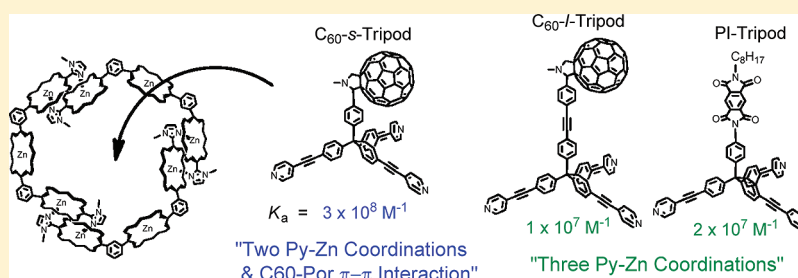


## Fullerene- and Pyromellitdiimide-Appended Tripodal Ligands Embedded in Light-Harvesting Porphyrin Macrorings

Yusuke Kuramochi,<sup>†</sup> Akiharu Satake,<sup>†</sup> Atula S. D. Sandanayaka,<sup>‡</sup> Yasuyuki Araki,<sup>‡</sup> Osamu Ito,<sup>‡</sup> and Yoshiaki Kobuke<sup>\*,†,§</sup><sup>†</sup>Graduate School of Materials Science, Nara Institute of Science and Technology, Takayama 8916-5, Ikoma, Nara 630-0101, Japan<sup>‡</sup>Institute of Multidisciplinary Research for Advanced Materials, Tohoku University, Katahira, Sendai, 980-8577, Japan

§ Supporting Information

## ABSTRACT:



Three new tripyridyl tripodal ligands appended with either fullerene or pyromellitdiimide moieties, named  $C_{60}$ -s-Tripod,  $C_{60}$ -l-Tripod, and PI-Tripod, were synthesized and introduced into a porphyrin macroring  $N-(1-Zn)_3$  (where  $1-Zn$  = trisporphyrinatozinc(II)). From UV-vis absorption and fluorescence titration data, the binding constants of  $C_{60}$ -s-Tripod,  $C_{60}$ -l-Tripod, and PI-Tripod with  $N-(1-Zn)_3$  in benzonitrile were estimated to be  $3 \times 10^8$ ,  $1 \times 10^7$ , and  $2 \times 10^7$   $M^{-1}$ , respectively. These large binding constants denote multiple interactions of the ligands to  $N-(1-Zn)_3$ . The binding constants of the longer ligand ( $C_{60}$ -l-Tripod) and the pyromellitdiimide ligand (PI-Tripod) are almost the same as those without the fullerene or pyromellitdiimide groups, indicating that they interact via three pyridyl groups to the porphyrinatozinc(II) coordination. In contrast, the larger binding constants and the almost complete fluorescence quenching in the case of the shorter ligand ( $C_{60}$ -s-Tripod) indicate that the interaction with  $N-(1-Zn)_3$  is via two pyridyl groups to the porphyrinatozinc(II) coordination and a  $\pi$ - $\pi$  interaction of the fullerene to the porphyrin(s). The fluorescence of  $N-(1-Zn)_3$  was quenched by up to 80% by the interaction of  $C_{60}$ -l-Tripod. The nanosecond transient absorption spectra showed only the excited triplet peak of the fullerene on selective excitation of the macrocyclic porphyrins, indicating that energy transfer from the excited  $N-(1-Zn)_3$  group to the fullerenyl moiety occurs in the  $C_{60}$ -l-Tripod/ $N-(1-Zn)_3$  composite. In the case of PI-Tripod, the fluorescence of  $N-(1-Zn)_3$  was quenched by 45%. It seems that the fluorescence quenching probably originates from electron transfer from the excited  $N-(1-Zn)_3$  group to the pyromellitdiimide moiety.

## INTRODUCTION

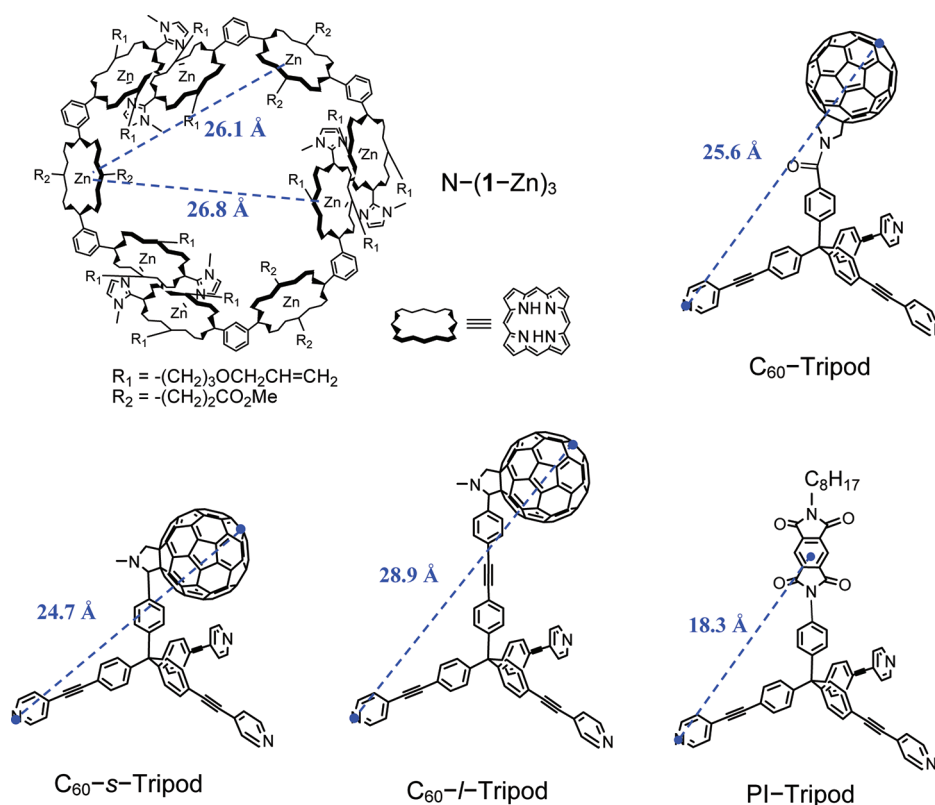
In photosynthesis, sunlight is converted into biochemical energy with a high efficiency. These light reactions begin with the capture of dilute photons from the sun by a network of chromophores, known as light-harvesting (LH) complexes. Photosynthetic purple bacteria have well-defined cyclic bacteriochlorophyll as architectures, such as LH1 and LH2,<sup>1,2</sup> and carry out isotropic energy migration among the LHs in the photosynthetic membrane.<sup>3</sup> The excited energy migrates along the LHs until it finally reaches a special pair of the reaction center (RC) that can produce a stable transmembrane-spanned charge-separated state through a multistep photoinduced electron transfer process.<sup>4</sup> The efficient collection of sunlight and conversion of energy into electrochemical potential enable the photosynthetic living organism to use solar energy for metabolic reactions with a quantum yield of almost 100%. The construction of artificial photosynthetic systems has attracted considerable

attention, not only from a pure scientific interest but also from potential applications of photosynthesis in solar energy conversion systems, which could help resolve the energy and environmental problems of the world. Porphyrin is a good chromophore for use in artificial photosynthesis because of its structural and functional similarities to natural chlorophyll. Thus, a number of researchers have studied various types of multiporphyrins and porphyrin-electron acceptor conjugated systems as models for LH and RC systems.<sup>5-7</sup>

We have reported on LH porphyrin macrorings<sup>8</sup> using a supramolecular methodology based on the complementary coordination of spacer-linked bisimidazolylporphyrinatozinc(II),<sup>9</sup> and have developed further the host macroring  $N-(1-Zn)_3$  having three noncoordinated porphyrinatozinc(II) units as

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**Figure 1.** Molecular structures of the macrocyclic  $N-(1-Zn)_3$  and tripyridyl tripodal ligands. The distances were estimated from molecular models using the Material Studio software package.<sup>13</sup>

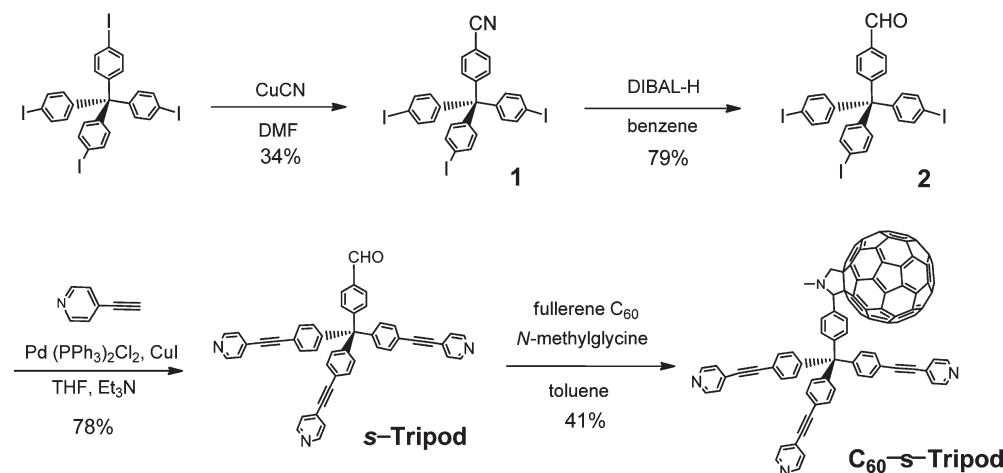
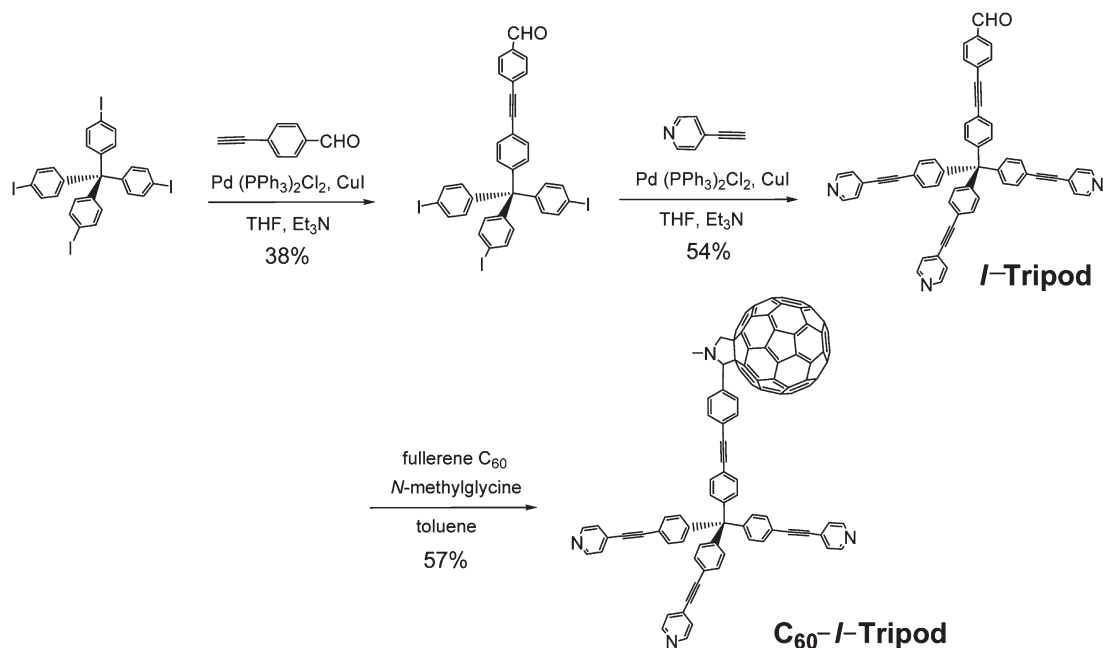
scaffolds that can accommodate the tripyridyl tripodal ligands.<sup>10,11</sup> We employed a tripodal pyridyl ligand with an energy acceptor porphyrin appended to a fullerene ( $C_{60}$ -ZnP-Tripod) as the specific guest of  $N-(1-Zn)_3$ .<sup>11c</sup> The composite exhibited a smooth singlet-singlet energy transfer process from the excited  $N-(1-Zn)_3$  group to the acceptor porphyrin, followed by the production of a stable charge-separated state between the acceptor porphyrin and the appended fullerene in  $C_{60}$ -ZnP-Tripod. We also synthesized a tripodal ligand with a fullerene linked via an amide ( $C_{60}$ -Tripod), and incorporated it into  $N-(1-Zn)_3$ , aiming to induce the direct electron transfer from the excited  $N-(1-Zn)_3$  unit to the fullerene moiety.<sup>11a</sup> Although  $C_{60}$ -Tripod was smoothly accommodated into  $N-(1-Zn)_3$  with a binding constant up to  $3 \times 10^8 \text{ M}^{-1}$ , even in a polar benzonitrile environment, and the fluorescence of  $N-(1-Zn)_3$  was almost completely quenched, no peaks from the radical ion were detected in either the nanosecond or the picosecond transient absorption spectral measurements. These results suggest that the fullerene moiety of  $C_{60}$ -Tripod is in direct contact with the porphyrin plane(s) of  $N-(1-Zn)_3$  through a fullerene-porphyrin  $\pi-\pi$  interaction along with coordination with two pyridyl groups, resulting in fast back electron transfer.<sup>12</sup>

Here, we report on the synthesis of three novel tripodal acceptor ligands:  $C_{60}$ -*l*-Tripod,  $C_{60}$ -*s*-Tripod, and PI-Tripod (Figure 1).  $C_{60}$ -*l*-Tripod has a long phenylene-ethynylene-phenylene linker in between a fullerene-pyrrolidine and a tris-[4-(2-(4-pyridyl)ethynyl)phenyl]methane part. In  $C_{60}$ -*l*-Tripod, the distance from the pyridyl nitrogen atom to the fullerene edge is large enough to prevent the fullerene moiety from being enclosed by the  $N-(1-Zn)_3$  groups, and also to prevent a direct  $\pi-\pi$

interaction between the fullerene moiety and the porphyrin groups of  $N-(1-Zn)_3$ , which causes a fast back electron transfer.  $C_{60}$ -*s*-Tripod was the reference compound, as it has a short phenylene linker, where the distance from the pyridyl nitrogen atom to the fullerene edge is small enough for the fullerene moiety to be enclosed by the  $N-(1-Zn)_3$  group. PI-Tripod has a pyromellitimide as an electron acceptor. In a PI-Tripod/ $N-(1-Zn)_3$  composite, a three-point coordination from the pyridyl groups to the porphyrinatozinc(II) groups is expected to be an exclusive binding mode, because the  $\pi-\pi$  interaction between the pyromellitimide and the porphyrin would be weak, and furthermore, a  $\pi-\pi$  stacking between PI-Tripod and  $N-(1-Zn)_3$  is structurally not possible. By comparing the properties of these novel tripodal ligands, we attempted to construct supramolecular composites that would perform an optimal photoinduced electron transfer from LH macrocyclics to a central acceptor.

## RESULTS AND DISCUSSION

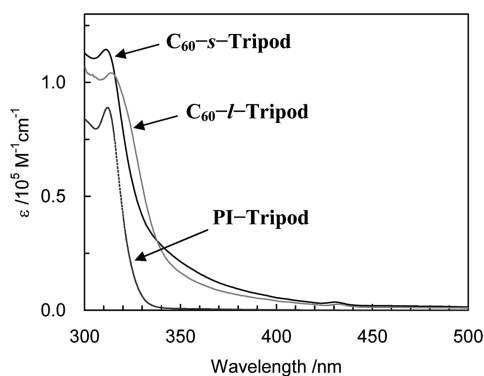
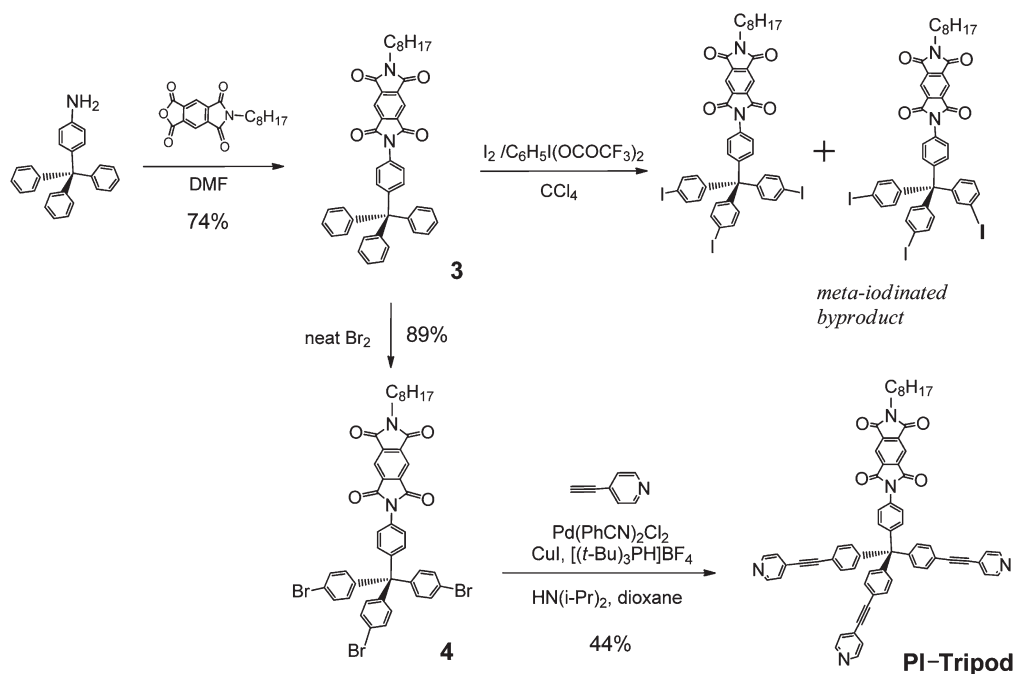
**Synthesis of  $C_{60}$ -*s*-Tripod,  $C_{60}$ -*l*-Tripod, and PI-Tripod.** The  $C_{60}$ -*s*-Tripod,  $C_{60}$ -*l*-Tripod, and PI-Tripod were prepared according to Schemes 1, 2, and 3, respectively.  $C_{60}$ -*s*-Tripod was synthesized from tetrakis(4-iodophenyl)methane<sup>14</sup> in four steps. The cyanation of tetrakis(4-iodophenyl)methane was carried using 1 equiv of CuCN in dimethylformamide (DMF) to give a mixture of **1** (35%), unreacted starting material (50%), and a dicyanated byproduct (15%). The crude mixture was purified using a silica gel column to afford **1** in a yield of 34%. Cyanide **1** was reduced by DIBAL-H to the corresponding aldehyde **2** in a yield of 79%. The cross-coupling reaction of **2** with 4-ethynylpyridine<sup>15</sup> was carried out in the presence of a

Scheme 1. Synthetic Route of C<sub>60</sub>-*s*-TripodScheme 2. Synthetic Route of C<sub>60</sub>-*l*-Tripod

Pd(PPh<sub>3</sub>)<sub>2</sub>Cl<sub>2</sub>/CuI catalyst<sup>16</sup> to afford *s*-Tripod in a yield of 78%. The succeeding 1,3-dipolar cycloaddition with C<sub>60</sub> and *N*-methylglycine<sup>17</sup> afforded C<sub>60</sub>-*s*-Tripod in a yield of 41%. C<sub>60</sub>-*l*-Tripod was synthesized from tetrakis(4-iodophenyl)methane in three steps. *l*-Tripod was synthesized from tetrakis(4-iodophenyl)methane in a total yield of 21%, as has been described elsewhere.<sup>11c</sup> C<sub>60</sub>-*l*-Tripod was synthesized using a 1,3-dipolar cycloaddition employing *l*-Tripod, C<sub>60</sub> and *N*-methylglycine in a yield of 57%. PI-Tripod was synthesized according to Scheme 3. Because the starting material 4-(triphenylmethyl)aniline has an NH<sub>2</sub> group that strongly activates electrophilic substitution at the position *ortho* to the NH<sub>2</sub> group in the aniline moiety, pyromellitic monoanhydride was first introduced at the *para*-positions of the three other phenyl groups before halogenation.<sup>18</sup> The condensation of 4-(triphenylmethyl)aniline<sup>14</sup>

and pyromellitic monoanhydride<sup>19</sup> in DMF afforded 3 in a yield of 74%. In the final cross-coupling reaction for PI-Tripod, the reactivity of aryl iodide was expected to be much higher than that of aryl bromide.<sup>20</sup> Thus, we first attempted to synthesize the iodinated precursor for PI-Tripod. Iodination at the *para*-positions of the three phenyl groups of 3 was first attempted using bis(trifluoroacetoxy)iodobenzene (3–5 equiv) and iodine (3–5 equiv) in CCl<sub>4</sub>.<sup>14</sup> However, the iodination reaction gave a mixture of *para*- and *meta*-iodinated products (Scheme 3). The *meta* byproduct could not be removed using column chromatography, even after the successful coupling reaction with 4-ethynylpyridine. On the other hand, the bromination reaction only proceeded at the targeted *para*-positions of the three phenyl groups after treatment with neat bromine at room temperature.<sup>21</sup> Purification using a silica gel column gave 4 in a yield of 89%. The

Scheme 3. Synthetic Route of PI-Tripod



**Figure 2.** UV-vis absorption spectra of  $C_{60}$ -*s*-Tripod ( $CHCl_3$ ),  $C_{60}$ -*l*-Tripod (benzonitrile), and PI-Tripod ( $CHCl_3$ ) at 25 °C.

cross-coupling reaction of **4** with 4-ethynylpyridine hardly proceeded when  $Pd(PPh_3)_2Cl_2$  and CuI were used as catalysts. To overcome this problem, the bulky and strong electron-donating phosphine ligand  $(P(t-Bu)_3)_3$  was used instead of  $PPh_3$ .<sup>20</sup> The coupling reaction proceeded at room temperature in the presence of  $Pd(PhCN)_2Cl_2$ ,  $[(t-Bu)_3PH]BF_4$  and CuI, and afforded PI-Tripod in a yield of 44%.  $C_{60}$ -*s*-Tripod,  $C_{60}$ -*l*-Tripod, and PI-Tripod were characterized using  $^1H$  and  $^{13}C$  NMR, Matrix-Assisted Laser Desorption/Ionization-Time-of-Flight (MALDI-TOF) mass spectrometry, high-resolution mass spectrometry, and UV-vis absorption spectroscopy.

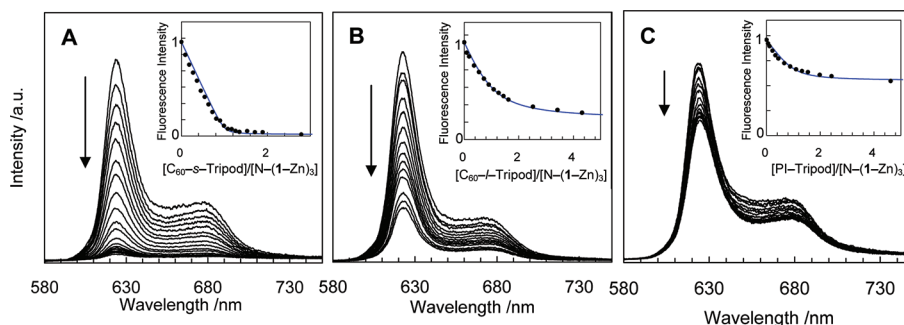
The UV-vis absorption spectra of  $C_{60}$ -*s*-Tripod,  $C_{60}$ -*l*-Tripod, and PI-Tripod are shown in Figure 2. The spectra of  $C_{60}$ -*s*-Tripod and  $C_{60}$ -*l*-Tripod show a strong absorption in the UV region accompanied by a weak absorption that extended continuously to the visible region, which originated from the absorption of the fullereryl moiety. In the case of

PI-Tripod, there was no absorption extending to the visible region ( $>450$  nm).

**Binding Modes of  $C_{60}$ -*s*-Tripod,  $C_{60}$ -*l*-Tripod, and PI-Tripod to  $N-(1-Zn)_3$ .** Incorporation of the acceptor tripodal ligands into the  $N-(1-Zn)_3$  group was determined using UV-vis absorption and fluorescence titration. The experiments were performed in benzonitrile as a suitable solvent to stabilize the charge-separated species. The binding mode of the tripodal ligands to  $N-(1-Zn)_3$  was estimated by comparing their binding constants with the corresponding tripodal ligand without their acceptors. If almost the same values were observed between the two tripodal ligands, with and without the acceptor, the binding mode was concluded to maintain the three-point coordination of the pyridyl group to the zinc atom. The changes in the UV-vis absorption spectra are shown in Supporting Information, Figures S15–S17. In all three spectra, a small decrease and a red-shift of the Sorbet band region was observed. No change in the Q-band was observed, because benzonitrile was already coordinated to the uncoordinated porphyrinatozinc(II).<sup>11a</sup> The change in the fluorescence spectra of  $N-(1-Zn)_3$  on addition of the acceptor ligands is shown in Figure 3. The fluorescence of  $N-(1-Zn)_3$  ( $4.0 \times 10^{-7}$  M) is efficiently quenched by 98% with 2.8 equiv of  $C_{60}$ -*s*-Tripod (Figure 3A, inset). The binding constant estimated from the UV-vis absorption and fluorescence titration data was almost same:  $3 \times 10^8$  and  $4 \times 10^8$   $M^{-1}$ , respectively.<sup>22</sup> These values are 10 times the value of  $2.9 \times 10^7$   $M^{-1}$  of the precursor *s*-Tripod, suggesting a switch of the binding mode to a two-point coordination of the pyridyl group to the zinc atom and a  $\pi$ - $\pi$  interaction between the fullereryl moiety and the porphyrin(s) in  $N-(1-Zn)_3$ , as was observed previously.<sup>11a</sup> The almost complete fluorescence quenching is similar to the previous case, consistent with this binding mode.

On the titration of  $C_{60}$ -*l*-Tripod, the binding constant was estimated from the UV-vis absorption and fluorescence titration





**Figure 3.** Steady-state fluorescence quenching of  $N-(1-Zn)_3$  from the addition of: (A)  $C_{60}$ - $s$ -Tripod, (B)  $C_{60}$ - $l$ -Tripod, and (C) PI-Tripod in benzonitrile at 25 °C. When excited at 567 nm: (A)  $[N-(1-Zn)_3] = 4.0 \times 10^{-7}$  M,  $[C_{60}$ - $s$ -Tripod] = 0–2.8 equiv, (B)  $[N-(1-Zn)_3] = 2.5 \times 10^{-7}$  M,  $[C_{60}$ - $l$ -Tripod] = 0–4.3 equiv, (C)  $[N-(1-Zn)_3] = 3.9 \times 10^{-7}$  M and  $[PI-Tripod] = 0$ –4.6 equiv. The insets show the change in fluorescence intensity with the signal integration (●) and the theoretical curves (blue lines) for: (A)  $K_a = 4 \times 10^8$  M $^{-1}$ , (B)  $K_a = 1.3 \times 10^7$  M $^{-1}$ , and (C)  $K_a = 2.0 \times 10^7$  M $^{-1}$ .

**Table 1.** Binding Constants,  $K_a$ , and Fluorescence Quenching Efficiencies of  $N-(1-Zn)_3$  of Tripodal Ligands in Benzonitrile at 25 °C, and Distances from the Pyridyl Nitrogen Atom and the Fullerene Edge of the Tripodal Ligands

		$s$ -Tripod	$C_{60}$ - $s$ -Tripod	$C_{60}$ -Tripod <sup>a</sup>	$l$ -Tripod	$C_{60}$ - $l$ -Tripod	PI-Tripod
$K_a$ /M $^{-1}$	fluorescence <sup>b</sup>		$4 \times 10^{8c}$	$3.4 \times 10^8$		$1.3 \times 10^7$	$2.0 \times 10^7$
	UV-vis	$2.9 \times 10^7$	$3 \times 10^{8c}$	$3.1 \times 10^8$	$1.1 \times 10^7$	$1.0 \times 10^7$	$2.3 \times 10^7$
quenching efficiency <sup>b,c</sup>		$f$	98%	97%	$f$	80%	45%
distance <sup>d</sup>			24.7 Å	25.6 Å		28.9 Å	

<sup>a</sup> Ref 11a. <sup>b</sup> The data were obtained using an excitation wavelength at 567 nm. <sup>c</sup> These values were calculated assuming a complexation rate of 100%. <sup>d</sup> See Figure 1. <sup>e</sup> Ref 22. <sup>f</sup> No fluorescence quenching was observed.

data to be  $1.0 \times 10^7$  and  $1.3 \times 10^7$  M $^{-1}$ , respectively. The fluorescence of  $N-(1-Zn)_3$  was quenched by 80% (Figure 3B, inset). The binding constant of  $C_{60}$ - $l$ -Tripod was similar to that of the precursor,  $l$ -Tripod ( $1.1 \times 10^7$  M $^{-1}$ ), indicating that it maintained the three-point coordination of the pyridyl group to the zinc atom.

Fluorescence titration using PI-Tripod in benzonitrile is shown in Figure 3C. From the UV-vis absorption and the fluorescence titration curves, the binding constants were estimated to be  $2.3 \times 10^7$  and  $2.0 \times 10^7$  M $^{-1}$ , respectively. The fluorescence of  $N-(1-Zn)_3$  was quenched by 45% (Figure 3C, inset). The binding constant of PI-Tripod was almost the same as that of the tripodal ligands,<sup>10,23</sup> indicating a three-point coordination of the pyridyl group to the zinc atom.

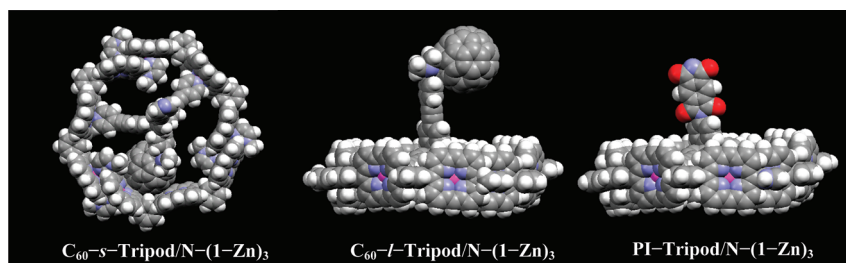
All the binding constants and the fluorescence quenching efficiencies in benzonitrile are summarized in Table 1. Judging from these data, it was estimated that  $C_{60}$ - $s$ -Tripod interacted with the  $N-(1-Zn)_3$  group as a two-point pyridyl coordination along with a fullerene-porphyrin contact, whereas the other two ligands interacted as a three-point coordination.

According to the molecular model created using the Material Studio software package,<sup>13</sup> the distance in  $C_{60}$ - $s$ -Tripod from the pyridyl nitrogen atom to the fullerene edge was estimated to be 24.7 Å, which is smaller than the cavity of  $N-(1-Zn)_3$  (Figure 1) based on the coordination bond length (approximately 2.1 Å)<sup>24</sup> between the pyridyl nitrogen atom and the porphyrinatozinc atom. Thus, the fullerene moiety is incorporated inside the  $N-(1-Zn)_3$  ring and so gains a high degree of entropy from the release of constricted solvent molecules located inside the cavity.<sup>23</sup> In the case of  $C_{60}$ - $l$ -Tripod, the distance from the pyridyl nitrogen atom to the fullerene edge was estimated to be 28.9 Å, which is too large for the fullerene moiety to be enclosed

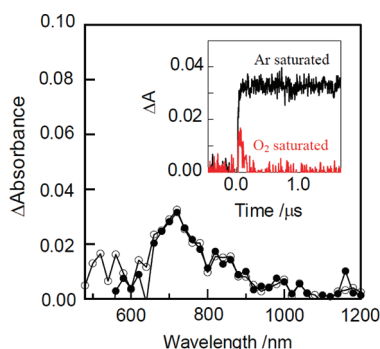
by the ring. The pyromellitimide moiety of PI-Tripod cannot be located inside the cavity, because weak  $\pi$ - $\pi$  interaction between the pyromellitimide and the porphyrin ring cannot replace the three cooperative pyridyl-porphyrinatozinc coordinations. The prospective structures of their composites are shown in Figure 4.

**Photodynamic Properties of Macroring/Tripodal Ligand Composites.** The results of the binding constant and the fluorescence quenching experiments suggest that the binding mode of  $C_{60}$ - $s$ -Tripod is the same as that of  $C_{60}$ -Tripod. Thus, a stable charge-separated state is not expected, as observed previously.<sup>11a</sup> On the other hand, the fullerene moiety in the  $C_{60}$ - $l$ -Tripod/ $N-(1-Zn)_3$  composite must locate not inside, but over the cavity. Because the nearest edge-to-edge distance between the porphyrin and the fullerene moiety in this composite is estimated to be 13 Å, the formation of a stable charge-separated state is expected if the photoinduced electron transfer from the excited  $N-(1-Zn)_3$  group to the fullerene moiety occurs.<sup>6</sup>

The time-resolved fluorescence decay and nanosecond transient absorption were measured to investigate the photophysical dynamics of the  $C_{60}$ - $l$ -Tripod/ $N-(1-Zn)_3$  composite. Under these conditions, the binding constants ensured that more than 95% of the macroring existed as a complex with  $C_{60}$ - $l$ -Tripod. The fluorescence time profile of the  $C_{60}$ - $l$ -Tripod/ $N-(1-Zn)_3$  composite decayed biexponentially with a time constant of 184 ps (70%) and 1220 ps (30%). The minor longer-lived component may arise from impurities or decomposed products. The quenching rate ( $k_q$ ) and the quantum yield ( $\Phi_q$ ) were evaluated to be  $5.0 \times 10^9$  s $^{-1}$  and 0.9 from the shorter-lifetime component using the lifetime of 2100 ps of free  $N-(1-Zn)_3$ .<sup>11a</sup> The nanosecond transient absorption spectra were measured to elucidate the quenching mechanism of the



**Figure 4.** Prospective structures of  $C_{60}$ -*s*-Tripod,  $C_{60}$ -*l*-Tripod, and PI-Tripod accommodated by  $N-(1-Zn)_3$ . (The left image is depicted from the top. The others are side views).



**Figure 5.** Nanosecond transient absorption spectra of the composite formed by mixing  $N-(1-Zn)_3$  ( $1.2 \times 10^{-6}$  M) and  $C_{60}$ -*l*-Tripod ( $2.6 \times 10^{-6}$  M) in benzonitrile at 0.1  $\mu$ s (●) and 1.0  $\mu$ s (○) after laser irradiation at 532 nm. Inset = absorption-time profiles at 720 nm in Ar- and  $O_2$ -saturated benzonitrile.

excited singlet state of  $N-(1-Zn)_3$ . The spectra were obtained using laser irradiation at 532 nm so that the  $N-(1-Zn)_3$  group was excited selectively. The transient absorption spectra showed a peak occurring at 720 nm without a signal occurring around 1000 nm corresponding to the fullereryl radical anion (Figure 5), whose spectrum is similar to that observed in only  $C_{60}$ -*l*-Tripod (Supporting Information, Figure S18). The decay profile at 720 nm was reduced markedly on exposure to oxygen (Figure 5 inset). These results indicate that the species resulting from photoexcitation of the macroring porphyrins is the excited triplet state of fullerene formed by intersystem crossing (ISC) from  $^1C_{60}^*$ , which is produced after the singlet–singlet energy transfer from  $N-(1-Zn)_3$  to fullerene. Such energy transfer and ISC processes may occur competitively with the photoinduced electron transfer from  $C_{60}/^1N-(1-Zn)_3^*$  and  $^1C_{60}^*/N-(1-Zn)_3^*$  to yield  $C_{60}^{\bullet-}/N-(1-Zn)_3^{\bullet+}$ , which is thermodynamically possible. Absence of a clear absorption peak from  $C_{60}^{\bullet-}$  at 1000 nm, even in the spectrum at 0.1  $\mu$ s in Figure 5, suggests that  $C_{60}^{\bullet-}/N-(1-Zn)_3^{\bullet+}$  has a considerably shorter lifetime, and mostly returns to the neutral ground state within a period of 0.1  $\mu$ s, leaving only  $^3C_{60}^*$ .

The energy transfer process is also supported by the result that the fluorescence quenching efficiency of  $N-(1-Zn)_3$  by the addition of  $C_{60}$ -*l*-Tripod in less polar benzene is almost same (83%) as that in benzonitrile (80%). Since fullerene has a weak, but steady absorption band in the visible region (Supporting Information, Figure S19), the through-space singlet–singlet energy transfer rate,  $k_{TS}$ , from  $N-(1-Zn)_3$  to the fullereryl moiety of  $C_{60}$ -*l*-Tripod can be calculated using the following

Förster equation:

$$k_{TS} = \frac{8.8 \times 10^{-25} \kappa^2 \Phi}{n^4 R^6 \tau} \quad (1)$$

where  $\kappa^2$ ,  $\Phi$ ,  $\tau$ ,  $R$ , and  $J$  are the orientation factor of the donor and acceptor, the fluorescence quantum yield, the fluorescence lifetime of the donor, the distance between the donor and the acceptor, and the spectral integral of the fluorescence of the donor and the absorption of the acceptor, respectively. The orientation factor of  $\kappa^2 = 2/3$  was used for randomly orientated dipoles<sup>25</sup> to estimate the energy transfer rate. The fluorescence quantum yield of  $\Phi = 0.039$  and the fluorescence lifetime of  $\tau = 2.1$  ns were obtained from the experiments.<sup>11a</sup> The distance between the center of the acceptor and the center of the nearest donor,  $R = 16$  Å, was estimated from molecular modeling using the Material Studio software package, and the spectral integral was calculated to be  $J \approx 5 \times 10^{-15}$  cm<sup>6</sup> mmol<sup>-1</sup> from the experimental data. The energy transfer rate,  $k_{TS}$ , from  $N-(1-Zn)_3$  to the fullereryl moiety was calculated to be approximately  $6 \times 10^8$  s<sup>-1</sup> using eq 1. The calculated value is comparable to the experimental value ( $5.0 \times 10^9$  s<sup>-1</sup>). The difference between the two values may come from contributions of Dexter-type through-bond energy transfers or higher-order multipole–multipole interactions, which is neglected in the Förster approximation,<sup>26</sup> otherwise, of electron transfer from  $C_{60}/^1N-(1-Zn)_3^*$  to yield  $C_{60}^{\bullet-}/N-(1-Zn)_3^{\bullet+}$ . The three-point coordination from the pyridyl groups to the porphyrins makes fullereryl moiety of  $C_{60}$ -*l*-Tripod to be fixed tightly over the center of the cavity of the  $N-(1-Zn)_3$ . Since the through-bond pathway from the porphyrins to the fullerene moiety is very long, the electron transfer is expected to occur via the through-space pathway. However, the superexchange interaction involving only solvent molecules would be too small for the electron transfer to compete the energy transfer, resulting in the singlet–singlet energy transfer mostly.

In the PI-Tripod/ $N-(1-Zn)_3$  composite, the possibility of a through-space singlet–singlet energy transfer from an excited  $N-(1-Zn)_3$  group to a pyromellitimide moiety can be excluded owing to both the zero absorption of PI-Tripod in the visible region where the fluorescence of  $N-(1-Zn)_3$  occurs (Figure 2) and the higher energy of  $^1PI^*/N-(1-Zn)_3$  (approximately 3.7 eV) than  $PI/^1N-(1-Zn)_3^*$  (2.0 eV). The energy level of the ion-pair state,  $PI^{\bullet-}/N-(1-Zn)_3^{\bullet+}$ , was estimated to be 1.34 eV above the ground state.<sup>28</sup> Therefore, the photoinduced electron transfer from  $PI/^1N-(1-Zn)_3^*$  to yield  $PI^{\bullet-}/N-(1-Zn)_3^{\bullet+}$  should have sufficient driving force. Table 2 shows the solvent dependency of the binding constant of PI-Tripod and the fluorescence quenching. The fluorescence

Table 2. Solvent Dependency of the Steady-State Fluorescence Quenching Rate of PI-Tripod/N-(1-Zn)<sub>3</sub> Composite<sup>a</sup>

solvent	$\epsilon^b$	binding constant <sup>c</sup> /M <sup>-1</sup>	N-(1-Zn) <sub>3</sub>			
			$\Phi_f^d$	$\tau_f^e$ /ns	quenching efficiency <sup>f</sup>	quenching rate <sup>g</sup> /s <sup>-1</sup>
toluene	2.38	>10 <sup>8</sup> (>4 × 10 <sup>8</sup> )	5.1%	2.2	31%	2.0 × 10 <sup>8</sup>
chloroform <sup>h</sup>	4.81	4.5 × 10 <sup>7</sup> (3.0 × 10 <sup>7</sup> )	3.4%	2.0	38%	3.1 × 10 <sup>8</sup>
benzonitrile	25.2	2.0 × 10 <sup>7</sup> (2.3 × 10 <sup>7</sup> )	3.9%	2.1	45%	3.9 × 10 <sup>8</sup>

<sup>a</sup> Excited at 567 nm. <sup>b</sup> Dielectric constant. <sup>c</sup> Estimated from fluorescence titration. The values in parentheses were estimated from UV-vis titration data.

<sup>d</sup> The fluorescence quantum yield of free N-(1-Zn)<sub>3</sub> was determined using ZnTPP of = 3.3% as the standard value.<sup>27</sup> <sup>e</sup> Fluorescence lifetime of free N-(1-Zn)<sub>3</sub>. <sup>f</sup> These values were calculated assuming a complexation rate of 100%. <sup>g</sup> The quenching rate constant was calculated using the following equation,  $k_q = 1/\tau_f \times \Phi_q/(1 - \Phi_q)$ , where  $\tau_f$  is the fluorescence lifetime of free N-(1-Zn)<sub>3</sub> and  $\Phi_q$  is the steady-state fluorescence quenching efficiency. <sup>h</sup> Chloroform containing 0.5% EtOH as a stabilizer.

quenching was more efficient in a polar solvent, such as benzonitrile. This observation suggests that photoinduced electron transfer occurs from N-(1-Zn)<sub>3</sub> to the pyromellitimide moiety. Since the center-to-center distance between the nearest porphyrin ring and the pyromellitimide moiety in the composite was estimated to be 16 Å, photoinduced electron transfer in the PI-Tripod/N-(1-Zn)<sub>3</sub> composite is considered most likely to be the result of a superexchange coupling via the linker moieties of PI-Tripod. The transient absorption spectrum at 0.1 μs after laser irradiation showed neither a definitive charge-separated species nor a triplet excited state of N-(1-Zn)<sub>3</sub> formed by ISC from <sup>1</sup>N-(1-Zn)<sub>3</sub>\*. This strongly indicates that a fast charge recombination to the ground state occurred after the initial charge separation.

## CONCLUSIONS

We have synthesized three new acceptor tripodal ligands having fullerene or pyromellitimide moieties. The tripodal ligands having fullerene through a long linker (C<sub>60</sub>-l-Tripod) and the tripodal ligand containing pyromellitimide (PI-Tripod) were accommodated via a three-point coordination from a pyridyl group to a porphyrinatozinc(II) group. In a C<sub>60</sub>-l-Tripod/N-(1-Zn)<sub>3</sub> composite, the singlet-singlet energy transfer from the excited porphyrin in N-(1-Zn)<sub>3</sub> to the fullerene occurs in preference to electron transfer, even in polar benzonitrile. In this case, the fullerene works as a good energy acceptor because the fullereryl moiety of C<sub>60</sub>-l-Tripod is tightly fixed over the center of the cavity of the N-(1-Zn)<sub>3</sub> group with a lower electronic orbital coupling with the donor N-(1-Zn)<sub>3</sub> group, and the fullereryl moiety has an absorption peak in the region where the fluorescence of the N-(1-Zn)<sub>3</sub> group appears. In the case of the PI-Tripod/N-(1-Zn)<sub>3</sub> composite, where the pyromellitimide moiety has no absorption in the visible region, a solvent dependence of the quenching rate was observed, indicating that electron transfer from the excited N-(1-Zn)<sub>3</sub> group to the pyromellitimide moiety occurred.

## EXPERIMENTAL SECTION

**General Procedure.** All chemicals and solvents were of commercial reagent grade, and used without further purification unless otherwise stated. Dry tetrahydrofuran (THF), 1,4-dioxane, benzene, and toluene were prepared by distillation over benzophenone-Na. Dry dimethylformamide (DMF) and triethylamine (Et<sub>3</sub>N) were prepared by distillation over CaH<sub>2</sub>. <sup>1</sup>H NMR spectra (600 MHz) were recorded on JEOL ECP-600, and chemical shifts were recorded in parts per million (ppm) relative to tetramethylsilane. UV-vis absorption spectra were recorded on a Shimadzu UV-3100PC spectrometer. Steady-state fluorescence

emission spectra were recorded on a Hitachi F-4500 spectrometer. The fluorescence intensities were normalized at the absorption of their excitation wavelengths. UV-vis  $\lambda_{\max}$  (log  $\epsilon$ ) values are reported in nanometers (nm). Quantum yields were determined by corrected integrated ratios of steady-state fluorescence spectra relative to that of ZnTPP ( $\Phi_f$  = 3.3% in benzene; tetraphenylporphyrinatozinc(II)).<sup>27</sup> MALDI-TOF mass spectra were obtained on PerSeptive Biosystems Voyager DE-STR with dithranol (Aldrich) as the matrix. Reactions were monitored on silica gel 60 F<sub>254</sub> TLC plates (Merck). The silica gel utilized for column chromatography was purchased from Kanto Chemical Co. Inc.: Silica Gel 60N (Spherical, Neutral) 60–210 μm and 40–210 μm (Flash). The nanosecond transient absorption spectra in the NIR region were measured by means of laser-flash photolysis; 565 nm light from Nd:YAG laser was used as the exciting source, and a Ge-avalanche-photodiode module was used for detecting the monitoring light from a pulsed Xe-lamp. The details of the experimental setup are described elsewhere.<sup>29</sup>

**4-Cyanophenyl-tris(4-iodophenyl)methane (1).** In a 300 mL three necked flask equipped with a dropping funnel, a reflux condenser, and a N<sub>2</sub> balloon were placed tetrakis(4-iodophenyl)methane (1 g, 1.2 × 10<sup>-3</sup> mol)<sup>14</sup> and dry DMF (100 mL) under N<sub>2</sub> atmosphere. The mixture was heated at 140 °C to become a homogeneous solution. CuCN (78 mg, 1.2 × 10<sup>-3</sup> mol) dissolved in dry DMF (40 mL) was then added dropwise. The reaction mixture was stirred for 24 h at 140 °C. After the solvent was evaporated, toluene (approximately 100 mL) and aqueous ammonia solution (approximately 100 mL) were added and stirred overnight. The organic layer was collected, washed with aqueous ammonia solution and water, and dried over anhydrous Na<sub>2</sub>SO<sub>4</sub>. The residue obtained by evaporation of the solvent was purified with a short silica gel column chromatography (CHCl<sub>3</sub>/hexane 1:1 to CHCl<sub>3</sub>). The fraction eluted with CHCl<sub>3</sub> was collected and evaporated to afford 295 mg (34%) of the titled compound: TLC (silica gel, CHCl<sub>3</sub>/hexane 1:2)  $R_f$  = 0.2, (silica gel, CHCl<sub>3</sub>)  $R_f$  = 0.7; <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>, 25 °C, TMS)  $\delta$  7.60 (d,  $J$  = 8.4 Hz, 6H, Ph), 7.56 (d,  $J$  = 8.4 Hz, 2H, Ph-CN), 7.29 (d,  $J$  = 8.4 Hz, 2H, Ph-CN), 6.86 (d,  $J$  = 8.4 Hz, 6H, Ph).

**4-Formylphenyl-tris(4-iodophenyl)methane (2).** In a 50 mL flask was placed **1** (295 mg, 4.1 × 10<sup>-4</sup> mol), and the flask was evacuated and replaced with Ar gas. Dry benzene (5 mL) was added to it, and the solution was cooled to 0 °C, and then 0.95 M hexane solution of diisobutylaluminum hydride (DIBAL-H, 1.2 mL, 1.1 × 10<sup>-3</sup> mol) was added at 0 °C. The reaction mixture was warmed to room temperature and stirred for 1.5 h. The reaction was quenched by addition of 5% H<sub>2</sub>SO<sub>4</sub> aqueous solution. The organic layer was collected, and the aqueous layer was extracted with benzene several times (Benzene is a cancer suspect agent and flammable liquid. Toluene should be substituted for benzene.). The combined organic layer was washed with water and dried over anhydrous Na<sub>2</sub>SO<sub>4</sub>, and then the solvent was evaporated. The residue was purified with a silica gel column chromatography (CHCl<sub>3</sub>), giving 235 mg (79%) of the titled compound: TLC (silica gel, CHCl<sub>3</sub>)



$R_f = 0.6$ ;  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ , 25 °C, TMS)  $\delta$  9.99 (s, 1H, CHO), 7.78 (d,  $J = 7.0$  Hz, 2H, Ph-CHO), 7.60 (d,  $J = 7.3$  Hz, 6H, Ph), 7.35 (d,  $J = 7.0$  Hz, 2H, Ph-CHO), 6.91 (d,  $J = 7.3$  Hz, 6H, Ph).

**4-Formylphenyl-tris[4-(2-(4-pyridyl)ethynyl)phenyl]methane (s-Tripod).** In a Schlenk flask were placed 2 (116 mg,  $1.6 \times 10^{-4}$  mol), 4-ethynylpyridine (82 mg,  $8.0 \times 10^{-4}$  mol),<sup>15</sup>  $\text{Pd}(\text{PPh}_3)_2\text{Cl}_2$  (13 mg,  $1.9 \times 10^{-5}$  mol), and  $\text{CuI}$  (3.7 mg,  $1.9 \times 10^{-5}$  mol) under Ar atmosphere. Dry  $\text{Et}_3\text{N}$  (0.5 mL) and dry THF (0.5 mL) were added to it, and the mixture was degassed by freeze–thaw cycles. The reaction mixture was stirred at room temperature under Ar atmosphere. After stirring for 1 h, further dry THF (1 mL) was added to the reaction mixture to dissolve the insoluble solid. After stirring for 20 h, the reaction mixture was diluted with chloroform, washed with water and dried over anhydrous  $\text{Na}_2\text{SO}_4$ . The residue obtained by evaporation of the solvent was purified with a silica gel column chromatography ( $\text{CHCl}_3/\text{MeOH}$  9:1). The fractions showing the signals of the target compound on MALDI-TOF mass spectra were concentrated and reprecipitated with hexane to remove a byproduct of 1,4-bis(4-pyridyl)butadiyne. Further purification was performed with a silica gel column chromatography ( $\text{CHCl}_3/\text{MeOH}$  9:1), giving 80.9 mg (78%) of the titled compound: TLC (silica gel,  $\text{CHCl}_3/\text{MeOH}$  9:1)  $R_f = 0.3$ ; MALDI-TOF MS found  $m/z$  652.6 ( $\text{MH}^+$ ); HRMS (FAB) calcd for  $\text{C}_{47}\text{H}_{30}\text{N}_3\text{O}$  ( $\text{MH}^+$ ) 652.2389, found 652.2383;  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ , 25 °C, TMS)  $\delta$  10.02 (s, 1H, CHO), 8.60 (d,  $J = 6.0$  Hz, 6H, Py), 7.83 (d,  $J = 8.2$  Hz, 2H, Ph-CHO), 7.50 (d,  $J = 8.5$  Hz, 6H, Ph), 7.43 (d,  $J = 8.2$  Hz, 2H, Ph-CHO), 7.37 (d,  $J = 6.0$  Hz, 6H, Py), 7.25 (d,  $J = 8.5$  Hz, 6H, Ph);  $^{13}\text{C}$  NMR (150 MHz,  $\text{CDCl}_3$ , 25 °C)  $\delta$  191.5, 152.0, 149.8, 146.1, 134.6, 131.6, 131.3, 131.1, 130.8, 129.3, 125.5, 120.6, 93.16, 87.31, 65.36.

**$\text{C}_{60}$ –s–Tripod.** In a Schlenk flask were placed s-Tripod (15 mg,  $2.3 \times 10^{-5}$  mol),  $\text{C}_{60}$  (83 mg,  $1.2 \times 10^{-4}$  mol), and *N*-methylglycine (107 mg,  $1.2 \times 10^{-3}$  mol). The flask was evacuated and replaced with Ar gas. Toluene (10 mL) was added to it, and the mixture was heated at 100 °C and stirred for 21 h. The solvent was evaporated to give a crude residue, which was passed through a short silica gel column by using toluene, and then,  $\text{CHCl}_3/\text{MeOH}$  9:1 as eluents to remove excess  $\text{C}_{60}$ . The fractions eluted with  $\text{CHCl}_3/\text{MeOH}$  9:1 was collected and evaporated. Further purification was performed with a silica gel column chromatography ( $\text{CHCl}_3/\text{MeOH}$  99:1 to 98:2), giving 13 mg (41%) of the titled compound: TLC (silica gel,  $\text{CHCl}_3/\text{MeOH}$  9:1)  $R_f = 0.6$ ; MALDI-TOF MS found  $m/z$  1400.6 ( $\text{MH}^+$ ); HRMS (ESI) calcd for  $\text{C}_{109}\text{H}_{35}\text{N}_4$  ( $\text{MH}^+$ ) 1399.2862, found 1399.2861;  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ , 25 °C, TMS)  $\delta$  8.60 (brs, 6H, Py), 7.52–7.77 (brs, 2H, Ph-pyrrolidine), 7.41 (d,  $J = 8.2$  Hz, 6H, Ph), 7.34 (brd,  $J = 4.8$  Hz, 6H, Py), 7.25 (brs, Ph-pyrrolidine), 7.19 (d,  $J = 8.2$  Hz, 6H, Ph), 4.98 (d,  $J = 9.6$  Hz, 1H, pyrrolidine  $\text{CH}_2$ ), 4.96 (s, 1H, pyrrolidine CH), 4.28 (d,  $J = 9.6$  Hz, 1H, pyrrolidine  $\text{CH}_2$ ), 2.90 (s, 3H,  $\text{N}-\text{CH}_3$ );  $^{13}\text{C}$  NMR (150 MHz,  $\text{CDCl}_3$ , 25 °C)  $\delta$  156.19, 153.85, 153.29, 152.96, 149.76, 147.32, 147.28, 146.89, 146.65, 146.43, 146.27, 146.20, 146.11, 146.09, 145.95, 145.88, 145.72, 145.56, 145.52, 145.41, 145.36, 145.29, 145.21, 145.14, 144.98, 144.70, 144.68, 144.40, 144.30, 143.16, 143.01, 142.69, 142.58, 142.44, 142.21, 142.15, 142.07, 142.03, 141.92, 141.72, 141.67, 141.46, 140.21, 140.12, 139.71, 139.06, 136.50, 136.41, 135.97, 135.78, 135.72, 131.42, 131.26, 130.82, 128.79, 125.54, 120.20, 93.54, 87.18, 82.78, 77.37, 70.04, 68.93, 65.11, 40.05; UV–vis ( $\text{CHCl}_3$ ) 257 (5.17), 292 (5.08), 311 (5.06), 430 (3.56), 703.5 (2.60).

**$\text{C}_{60}$ –l–Tripod.** In a test tube with screw cap were placed l-Tripod (4.2 mg,  $5.6 \times 10^{-6}$  mol),  $^{11}\text{C}$   $\text{C}_{60}$  (20 mg,  $2.8 \times 10^{-5}$  mol), *N*-methylglycine (25 mg,  $2.8 \times 10^{-4}$  mol), and toluene (1 mL), and the tube was purged with  $\text{N}_2$  gas. The reaction mixture was heated to 100 °C and stirred for 32 h. The resulting solution was passed through a short silica gel column (toluene to  $\text{CHCl}_3/\text{MeOH}$  9:1) to remove excess  $\text{C}_{60}$ . The fraction eluted with  $\text{CHCl}_3/\text{MeOH}$  9:1 was collected and evaporated. Further purification was performed with a silica gel column chromatography ( $\text{CHCl}_3/\text{MeOH}$  99:1 to 98:2), giving 4.8 mg (57%) of the titled

compound: TLC (silica gel,  $\text{CHCl}_3/\text{MeOH}$  9:1)  $R_f = 0.8$ ; MALDI-TOF MS found  $m/z$  1500.0 ( $\text{MH}^+$ ); HRMS (ESI) calcd for  $\text{C}_{117}\text{H}_{39}\text{N}_4$  1499.3175 ( $\text{MH}^+$ ), found 1499.3170;  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ , 25 °C, TMS)  $\delta$  8.60 (d,  $J = 5.4$  Hz, 6H, Py), 7.81 (brs, 2H, Ph-pyrrolidine), 7.59 (d,  $J = 7.8$  Hz, 2H, Ph-pyrrolidine), 7.47 (d,  $J = 9.0$  Hz, 6H, Ph), 7.44 (d,  $J = 9.0$  Hz, 2H,  $\text{Ph}'$ ), 7.36 (d,  $J = 5.4$  Hz, 6H, Py), 7.23 (d,  $J = 9.0$  Hz, 6H, Ph), 7.18 (d,  $J = 9.0$  Hz, 2H,  $\text{Ph}'$ ), 4.99 (d,  $J = 9.6$  Hz, 1H, pyrrolidine  $\text{CH}_2$ ), 4.95 (s, 1H, pyrrolidine CH), 4.27 ( $J = 9.6$  Hz, 1H, pyrrolidine  $\text{CH}_2$ ), 2.81 (s, 3H,  $\text{N}-\text{CH}_3$ );  $^{13}\text{C}$  NMR (150 MHz,  $\text{CDCl}_3$ , 25 °C)  $\delta$  156.1, 153.9, 153.1, 153.0, 149.8, 147.33, 147.31, 146.6, 146.4, 146.32, 146.30, 146.23, 146.20, 146.15, 146.01, 145.95, 145.7, 145.6, 145.52, 145.48, 145.42, 145.37, 145.29, 145.27, 145.24, 145.18, 144.7, 144.6, 144.42, 144.36, 143.2, 143.0, 142.7, 142.60, 142.56, 142.25, 142.24, 142.18, 142.14, 142.10, 142.05, 141.99, 141.9, 141.8, 141.7, 141.6, 140.21, 140.19, 139.9, 139.5, 140.22, 140.19, 139.9, 139.5, 137.6, 136.9, 136.4, 135.9, 135.7, 131.9, 131.5, 131.3, 131.2, 130.9, 130.8, 125.5, 123.1, 121.5, 120.4, 93.37, 89.83, 89.45, 87.19, 83.27, 70.00, 69.06, 65.03, 40.00; UV–vis (benzonitrile) 314 (5.02), 432 (3.42).

***N*-(4-(Triphenylmethyl)phenyl)-*N'*-(*n*-octyl)pyromellitimide (3).** In a 20 mL flask were placed *N*-(*n*-octyl)pyromellitic monoanhydride (100 mg,  $3.0 \times 10^{-4}$  mol),<sup>19</sup> 4-(triphenylmethyl)aniline (102 mg,  $3.0 \times 10^{-4}$  mol), and dry DMF (2 mL). The mixture was purged with  $\text{N}_2$  gas and then heated at 130 °C for 6 h. A white solid generated during cooling was collected by filtration and washed with hexane, affording 117.6 mg (60%) of the titled compound. The filtrate of the reaction mixture was evaporated and the residue was recrystallized in acetic anhydride. The crystal was collected by filtration and dried in vacuo, giving 26.7 mg (14%) of the titled compound. Total yield was 144.3 mg (74%): TLC (silica gel,  $\text{CHCl}_3$ )  $R_f = 0.5$ ;  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ , 25 °C, TMS)  $\delta$  8.57 (s, 2H, Ph-PI), 7.39 (d,  $J = 9.0$  Hz, 2H,  $\text{Ph}'$ ), 7.36 (d,  $J = 9.0$  Hz, 2H,  $\text{Ph}'$ ), 7.20–7.29 (m, 15H, Ph), 3.75 (t,  $J = 7.5$  Hz, 2H,  $\text{C}_\alpha\text{H}_2$ ), 1.71 (quintet,  $J = 6.9$  Hz, 2H,  $\text{C}_\beta\text{H}_2$ ), 1.24–1.34 (m, 10H,  $\text{CH}_2$ ), 0.87 (t,  $J = 6.9$  Hz, 3H,  $\text{CH}_3$ );  $^{13}\text{C}$  NMR (150 MHz,  $\text{CDCl}_3$ , 25 °C)  $\delta$  166.2, 165.3, 147.3, 146.3, 137.6, 136.7, 131.9, 131.1, 128.8, 127.6, 126.1, 125.0, 118.7, 64.86, 38.82, 31.72, 29.10, 29.06, 28.41, 26.83, 22.58, 14.04.

***N*-(4-(Tris(4-bromophenyl)methyl)phenyl)-*N'*-(*n*-octyl)pyromellitimide (4).** In a 20 mL flask were placed 3 (165 mg,  $2.6 \times 10^{-4}$  mol) and neat bromine (1 mL), and then the reaction mixture was stirred for 30 min at room temperature. The reaction mixture was diluted with  $\text{CHCl}_3$  (20 mL) and washed successively with an aqueous sodium bisulfite solution and water. The organic layer was dried over anhydrous  $\text{Na}_2\text{SO}_4$ , and the solvent was evaporated. The residue was purified with silica gel column chromatography ( $\text{CHCl}_3/\text{hexane}$  1:2 to  $\text{CHCl}_3/\text{MeOH}$  9:1) to afford 201 mg (89%) of the titled compound: TLC (silica gel,  $\text{CHCl}_3$ )  $R_f = 0.8$ ;  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ , 25 °C, TMS)  $\delta$  8.37 (s, 2H, Ph-PI), 7.41 (d,  $J = 8.2$  Hz, 8H, Ph and  $\text{Ph}'$ ), 7.32 (d,  $J = 8.5$  Hz, 2H,  $\text{Ph}'$ ), 7.08 (d,  $J = 8.2$  Hz, 6H, Ph), 3.75 (t,  $J = 7.1$  Hz, 2H,  $\text{C}_\alpha\text{H}_2$ ), 1.71 (brquintet, 2H,  $\text{C}_\beta\text{H}_2$ ), 1.26–1.34 (m, 10H,  $\text{CH}_2$ ), 0.87 (t,  $J = 6.6$  Hz, 3H,  $\text{CH}_3$ );  $^{13}\text{C}$  NMR (150 MHz,  $\text{CDCl}_3$ , 25 °C)  $\delta$  166.0, 165.1, 145.6, 144.4, 137.7, 136.6, 132.4, 131.4, 131.1, 129.4, 125.4, 120.8, 118.7, 63.80, 38.80, 31.69, 29.06, 29.02, 28.37, 26.78, 22.55, 14.02.

**PI–Tripod.**  $\text{Pd}(\text{PhCN})_2\text{Cl}_2$  (2.6 mg,  $6.8 \times 10^{-6}$  mol),  $[(t\text{-Bu})_3\text{P}]\text{BF}_4$  (3.9 mg,  $1.4 \times 10^{-5}$  mol), and  $\text{CuI}$  (1.3 mg,  $6.8 \times 10^{-6}$  mol) were placed in a Schlenk flask and dissolved in dry dioxane (3 mL) and dry (*i*-Pr) $_2\text{NH}$  (100  $\mu\text{L}$ ,  $6.9 \times 10^{-4}$  mol) under Ar atmosphere. The solution was degassed by freeze–thaw cycles, and then 4 (120 mg,  $1.4 \times 10^{-4}$  mol) and 4-ethynylpyridine (70 mg,  $6.8 \times 10^{-4}$  mol)<sup>15</sup> were added to it. The reaction mixture was stirred for 11 h at room temperature. The solvent was evaporated in vacuo, and the residue was extracted with  $\text{CHCl}_3$  and washed with water. The solvent was evaporated, and the residue was purified with flash silica gel column chromatography ( $\text{CHCl}_3/\text{MeOH}$  95:5) to give a mixture including the target compound.



Further purification of the mixture by reprecipitation with hexane afforded 56.7 mg (44%) of the titled compound: MALDI-TOF MS found  $m/z$  951.2 ( $MH^+$ ); HRMS (FAB) calcd for  $C_{64}H_{48}N_5O_4$  950.3706, found 950.3713;  $^1H$  NMR (600 MHz,  $CDCl_3$ , 25 °C, TMS)  $\delta$  8.64 (brs, 6H, Py), 8.39 (s, 2H, Ph-PI), 7.50 (d,  $J$  = 8.5 Hz, 6H, Ph), 7.45 (d,  $J$  = 8.8 Hz, 2H, Ph'), 7.38–7.40 (m, 8H, Py and Ph'), 7.28 (d,  $J$  = 8.2 Hz, 6H, Ph), 3.76 (t,  $J$  = 7.4 Hz, 2H,  $C_{\alpha}H_2$ ), 1.72 (quintet,  $J$  = 7.0 Hz, 2H,  $C_{\beta}H_2$ ), 1.25–1.34 (m, 10H,  $CH_2$ ), 0.87 (t,  $J$  = 7.0 Hz, 3H,  $CH_3$ );  $^{13}C$  NMR (150 MHz,  $CDCl_3$ , 25 °C)  $\delta$  166.0, 165.1, 149.7, 146.5, 145.5, 137.7, 136.6, 131.54, 131.49, 131.2, 130.9, 129.5, 125.5, 120.5, 118.7, 93.32, 87.21, 64.90, 38.82, 31.68, 29.06, 29.02, 28.38, 26.79, 22.55, 14.01; UV–vis. ( $CHCl_3$ ) 292.0 (4.97), 311.8 (4.94).

## ■ ASSOCIATED CONTENT

**S Supporting Information.** Listings of  $^1H$ ,  $^{13}C$  NMR of 1–4,  $s$ -Tripod,  $C_{60}$ - $s$ -Tripod,  $C_{60}$ - $l$ -Tripod, and PI-Tripod, UV–vis spectral changes of  $N-(1-Zn)_3$  by bindings of  $C_{60}$ - $s$ -Tripod,  $C_{60}$ - $l$ -Tripod, and PI-Tripod, nanosecond transient absorption spectra of  $C_{60}$ - $l$ -Tripod, spectral overlap of the absorption of  $C_{60}$ - $l$ -Tripod, and the fluorescence of  $N-(1-Zn)_3$ . This material is available free of charge via the Internet at <http://pubs.acs.org>.

## ■ AUTHOR INFORMATION

### Corresponding Author

\*Fax: (+81)774-38-3508. E-mail: [kobuke@iae.kyoto-u.ac.jp](mailto:kobuke@iae.kyoto-u.ac.jp).

### Present Addresses

<sup>§</sup>Institute of Advanced Energy, Kyoto University, Gokasho, Uji, Kyoto 611–0011, Japan.

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