

Lysosome-specific one-photon fluorescence staining and two-photon singlet oxygen generation by molecular dyad†

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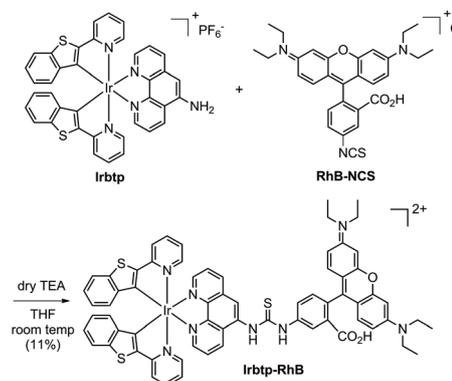
A dual functional molecular dyad, consisting of a cyclometalated Ir(III) complex and rhodamine B, has been synthesized and evaluated for its ability for independent operations of fluorescence staining and photodynamic therapy.

Image-guided photodynamic therapy holds considerable promise as a reliable strategy to treat cancer because the technique can avoid undesired photodamage.^{1–4} Dual functional molecules that produce photoluminescence and reactive oxygen species under photoirradiation are a prerequisite for the operation of the image-guided photodynamic therapy. Although phosphorescent transition metal complexes, such as those containing Ru(II),^{5–8} Ir(III),^{9,10} and Pt(II),¹¹ have been demonstrated for the dual function, their practicability is limited because singlet oxygen (¹O₂) produced during phosphorescence imaging provokes deleterious effects. Thus, the development of molecules with controllable phototoxicity remains a significant challenge, requiring the judicious molecular design that allows for independent operations of the imaging and photosensitization of ¹O₂. Specifically, utility of modulating function by selecting the photoirradiation wavelength is urgently required.

To address this challenge, we designed and synthesized a molecular dyad comprising a biscyclometalated Ir(III) complex and rhodamine B (RhB). In this dyad structure, the former sensitizes ¹O₂ under two-photon irradiation at 800 nm, while the latter produces strong red fluorescence under one-photon irradiation at 550 nm. Notably, the two photofunctions can be

operated in a fully independent manner because the electronic spin selection rule effectively prevents the bidirectional energy transfer between the molecular entities. The unique spin-gated energy transfer has been proved by steady-state and transient photoluminescence studies. Finally, photobiological utility of the dyad has been fully evaluated *in vitro*. It should be noted that this is the first report on the single molecule containing an Ir(III) complex capable of independent operation of fluorescence imaging and ¹O₂ sensitization for potential applications in image-guided photodynamic therapy.

The dyad (Irbtp–RhB) was prepared by forming a thiourea linkage between an amine-functionalized Ir(III) complex (Irbtp)¹² and rhodamine B isothiocyanate (RhB–NCS) in the presence of triethylamine (Scheme 1). Structures and purity of Irbtp–RhB and its precursors were examined by standard characterization methods, and the spectral identification data fully agreed with the proposed structure (ESI†). UV-vis absorption spectrum of a deaerated acetonitrile solution of 10 μM Irbtp–RhB is a simple superposition of the absorption spectra of the individual components of Irbtp and RhB (Fig. 1), excluding any ground-state interaction between two units. The dyad exhibits strong red fluorescence (λ_{em} = 570 nm,



Scheme 1 Synthesis of Irbtp–RhB.

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† Electronic supplementary information (ESI) available: Experimental details and Fig. S1–S10, plotting photoluminescence decay traces after nanosecond and picosecond laser excitation, determination of Φ(¹O₂) values, cyclic voltammograms, photostability of Φ_{PL} and Φ(¹O₂), cellular uptake, subcellular localization of Irbtp–RhB, and copies of ¹H and ¹³C NMR spectra. See DOI: 10.1039/c4ra02354d

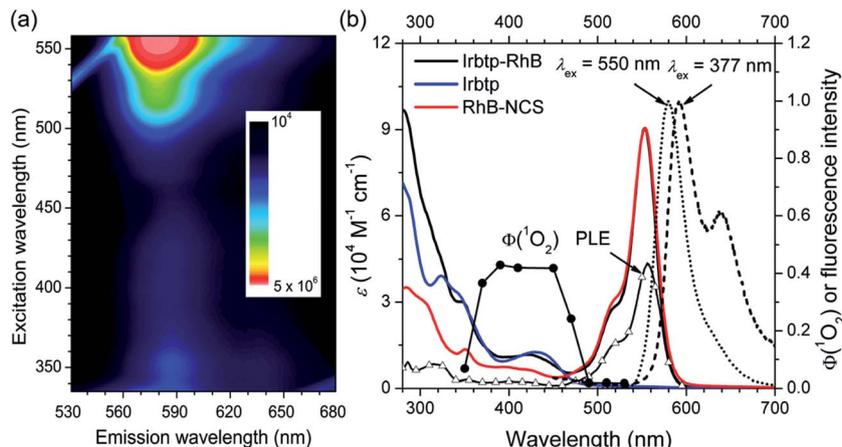


Fig. 1 Spectral properties of 10 μM Irbtp-RhB in deaerated acetonitrile. (a) Contour plot of photoluminescence intensities obtained with varying the excitation (330–560 nm) and emission (530–680 nm) wavelengths. (b) UV-vis absorption spectra (solid lines) of acetonitrile solutions of 10 μM Irbtp-RhB (black), 10 μM Irbtp (blue), and 10 μM RhB-NCS (red). Empty triangles, filled circles, and broken lines are the PLE spectrum ($\lambda_{\text{obs}} = 570 \text{ nm}$), the $\Phi(^1\text{O}_2)$ values at photoirradiation of $\lambda_{\text{ex}} = 350\text{--}550 \text{ nm}$, and photoluminescence spectra ($\lambda_{\text{ex}} = 377$ and 550 nm) of 10 μM Irbtp-RhB, respectively.

photoluminescence lifetime ($\tau_{\text{obs}} = 1.84 \text{ ns}$, photoluminescence quantum yield ($\Phi_{\text{PL}} = 2.3 \pm 0.2\%$) upon photoexcitation of the RhB moiety ($\lambda_{\text{ex}} = 490\text{--}550 \text{ nm}$), whereas weaker phosphorescence from Irbtp ($\lambda_{\text{em}} = 592 \text{ nm}$, $\tau_{\text{obs}} = 1.68 \mu\text{s}$, $\Phi_{\text{PL}} = 1.2 \pm 0.1\%$) with a characteristic vibronic progression of $\Delta\nu = 1340 \text{ cm}^{-1}$ is observed when the photoexcitation beam at $\lambda_{\text{ex}} = 330\text{--}450 \text{ nm}$ is provided (Fig. 1).^{12,13} The photoluminescence excitation spectrum (PLE) for the 570 nm emission overlaps exclusively with the absorption band of the RhB unit (Fig. 1b), supporting that the fluorescence emission is due to selective photoexcitation of RhB. The minimal contribution of the Irbtp unit to the PLE spectrum precludes occurrence of singlet-singlet energy transfer from Irbtp to RhB. In addition, spin-restricted energy transfer from the singlet excited state of the RhB unit to the triplet state of the Irbtp unit is also ruled

out, because the fluorescence lifetimes for the RhB fluorescence of Irbtp-RhB and RhB-NCS are identical (ESI, Fig. S1†).

Ir btp-RhB shows high ability for photosensitization of $^1\text{O}_2$, with a quantum yield ($\Phi(^1\text{O}_2)$) of 43% ($\lambda_{\text{ex}} = 365 \text{ nm}$; $\text{abs}(365 \text{ nm}) = 0.17$; ESI, Fig. S2†). To identify the $^1\text{O}_2$ -photosensitizing moiety, $\Phi(^1\text{O}_2)$ values were determined with increasing the photoexcitation wavelength (λ_{ex}) from 350 nm to 550 nm, using

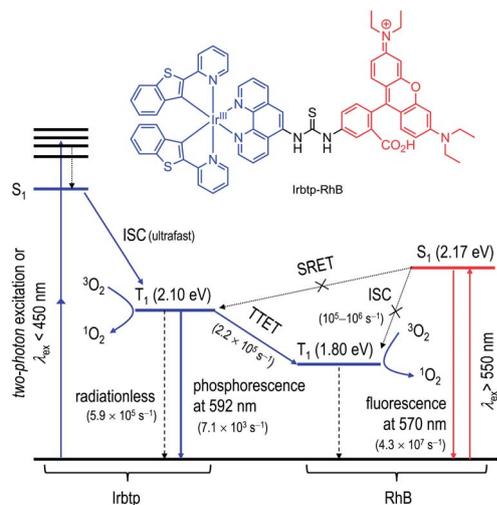


Fig. 2 Proposed mechanism for the photophysical processes of Irbtp-RhB: ISC, intersystem crossing; SRET, spin-restricted energy transfer; TTET, triplet-triplet energy transfer.

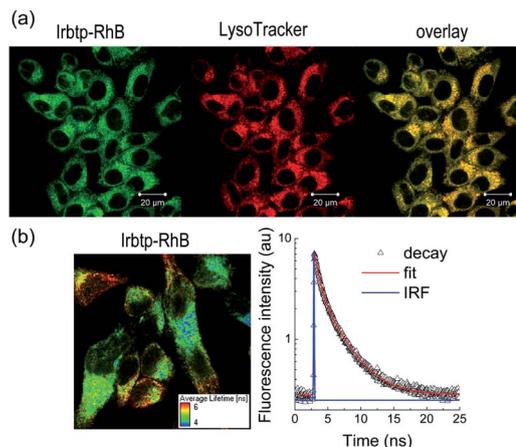


Fig. 3 Lysosomal staining of HeLa cells with Irbtp-RhB. (a) Confocal laser-scanning micrographs of HeLa cells co-stained with 10 μM Irbtp-RhB (10 min incubation at 37°C) and 500 nM LysoTracker Red (2 h incubation at 37°C): left, fluorescence image from Irbtp-RhB ($\lambda_{\text{ex}} = 488 \text{ nm}$; $\lambda_{\text{em}} = 551\text{--}591 \text{ nm}$); middle, fluorescence image from LysoTracker Red ($\lambda_{\text{ex}} = 577 \text{ nm}$; $\lambda_{\text{em}} = 590\text{--}747 \text{ nm}$); right, overlay image. Scale bar = 20 μm . (b) Fluorescence lifetime-imaging micrograph of HeLa cells treated with 10 μM Irbtp-RhB (10 min incubation at 37°C): left, lifetime image (250 pixel \times 250 pixel and 80 μm \times 80 μm) of the cells acquired after picosecond laser excitation at 470 nm (repetition rate = 40 MHz, pulse power = 1 μW , and acquisition window = 5 ms per pixel); right, fluorescence decay traces (black symbols) and their fit (red curve) to a biexponential decay model ($\tau_{\text{avg}} = 3.2 \text{ ns}$). Blue curve is the instrumental response function.

1,3-diphenylisobenzofuran^{14,15} and methylene blue¹⁶ as a ¹O₂ trap and a reference material ($\Phi(^1\text{O}_2) = 52\%$), respectively. As shown in Fig. 1b, the action profile (*i.e.*, $\Phi(^1\text{O}_2)$ vs. λ_{ex}) matches perfectly with the singlet metal-to-ligand charge-transfer (¹MLCT) absorption band of Irbtp.¹² On the contrary, direct photoexcitation of the RhB moiety produces smaller quantum yields ($\Phi(^1\text{O}_2) < 9\%$), because the triplet state formation in RhB is possible only by triplet-triplet energy transfer from the Irbtp unit at a rate constant, $k_{\text{TET}} = 2.2 \times 10^5 \text{ s}^{-1}$ (ESI, Fig. S1†).^{17,18} These results demonstrate that the photoactions of the Irbtp and RhB entities can be performed independently, while intramolecular photoinduced electron transfer from Irbtp to RhB may occur with a positive driving force of 0.56 eV to produce radical ion pair of Irbtp–RhB (*i.e.*, Irbtp^{•+}–RhB^{•-}; ESI, Fig. S3†). This decoupling nature also affords significant photostability, as demonstrated by the Φ_{PL} and $\Phi(^1\text{O}_2)$ values being unaffected by the continuous exposure to a 550 nm beam for 10 min (ESI, Fig. S4†). Taking the results into consideration, a photophysical scheme of Irbtp–RhB is summarized as Fig. 2. It is noteworthy that the unidirectionality in the energy transfer is the key feature that enables the independent actions of fluorescence and ¹O₂ photosensitization.

Having understood the photophysical mechanism, photo-biological utility of Irbtp–RhB was examined. Irbtp–RhB is cell-permeable when a 10 μM stock solution in DMSO is added to

the culture medium of live HeLa cells at 37 °C. The intracellular uptake of Irbtp–RhB may mainly involve passive diffusion, because incubation at 4 °C and treatments of the cells with endocytic inhibitors (50 μM chloroquine or 50 mM NH₄Cl)¹⁹ and metabolic inhibitors (50 mM 2-deoxy-D-glucose + 5 μM oligomycin)²⁰ do not alter the cellular entry (ESI, Fig. S5†). Bright punctate patterns are localized within the lysosomes of the treated HeLa cells, as seen from the fluorescence images obtained through the excitation ($\lambda_{\text{ex}} = 488 \text{ nm}$) and emission ($\lambda_{\text{em}} = 551\text{--}591 \text{ nm}$) channels adjusted for the RhB fluorescence (Fig. 3a; see ESI, Fig. S6† for more images). Co-staining experiments with 500 nM LysoTracker Red (Molecular Probe) reveal the lysosomal staining capability of Irbtp–RhB (overlap coefficient = 0.99). In addition to the confocal laser-scanning microscopy, Irbtp–RhB is compatible with the standard set-up for fluorescence lifetime-imaging microscopy (Fig. 3b).²¹

Finally, photoinduced cytotoxicity by intracellular generation of ¹O₂ has been evaluated. HeLa cells pretreated with different concentrations (0–20 μM , 1 h at 37 °C) of Irbtp–RhB were photoirradiated using a two-photon beam at 800 nm (32 nJ, 80 MHz). The excitation wavelength corresponded to the two-photon absorption of Irbtp, although quantitative evaluations require higher precision of the measurements. Cells that were kept under dark served as controls to estimate dark cytotoxicity of Irbtp–RhB. As shown in Fig. 4a, the 800 nm photoirradiation resulted in vacuolization of the Irbtp–RhB-treated cells. In sharp contrast, neither 561 nm photoirradiation (one-photon) of the treated cells nor 800 nm excitation (two-photon) of untreated cells led to such changes. MTS assays were employed to quantitate the photoinduced cytotoxicity. As shown in Fig. 4b, the 800 nm two-photon photoexcitation significantly attenuates the cell viability value < 30% (10 μM Irbtp–RhB, 15 min photoirradiation). The efficacy of phototoxicity, estimated by the cell viability ratio of the control cells over the photoirradiated cells, reaches as high as 2.72 after 15 min photoirradiation.

To summarize, we synthesized a molecular dyad comprising a biscyclometalated Ir(III) complex and rhodamine B, and characterized its dual functionality for fluorescence imaging and photodynamic therapy. The unidirectional energy transfer and the ability for selective photoexcitation enable independent operation of the lysosomal staining and the photosensitization of ¹O₂.

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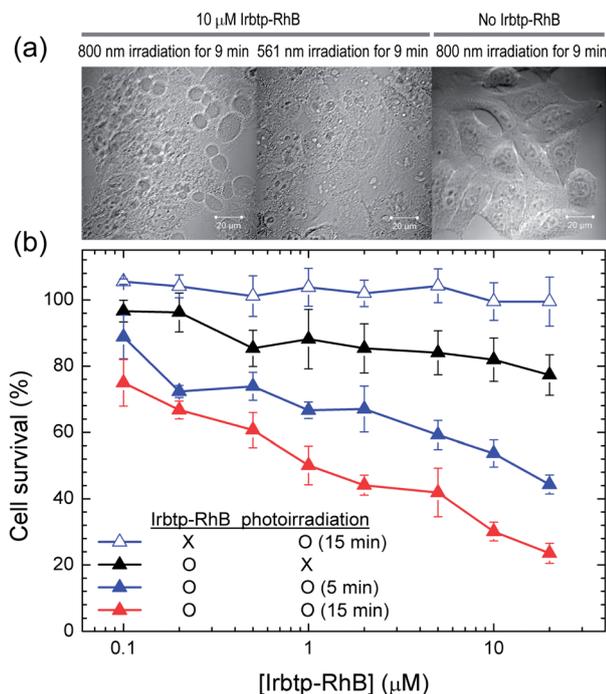


Fig. 4 Photoinduced cell death by Irbtp–RhB. (a) Changes in the cell morphology of HeLa cells under two-photon photoirradiation (80 MHz, 32 nJ) at 800 nm (left and right panels) and one-photon photoirradiation at 561 nm (middle panel). The cells in the left and middle panels were preincubated with 10 μM Irbtp–RhB. See ESI† for the movie clips showing temporal changes of the HeLa cells upon photoirradiation. (b) MTS cell viability data for HeLa cells incubated in the presence and absence of 0.1–20 μM Irbtp–RhB (1 h at 37 °C). The cells were photoirradiated for 5 and 15 min prior to the MTS assay.

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