Functionalizing Collagen with Vessel-Penetrating Two-Photon Phosphorescence Probes: A New In Vivo Strategy to Map Oxygen Concentration in Tumor Microenvironment and Tissue Ischemia

Cheng-Ham Wu, Kristina S. Kisel, Muthu Kumar Thangavel, Yi-Ting Chen, Kai-Hsin Chang, Ming-Rung Tsai, Chia-Yu Chu, Yu-Fang Shen, Pei-Chun Wu, Zhiming Zhang, Tzu-Ming Liu,* Janne Jänis, Elena V. Grachova, Julia R. Shakirova, Sergey P. Tunik,* Igor O. Koshevoy,* and Pi-Tai Chou*

1. Introduction

Sufficient oxygen (O$_2$) supply is critical to maintaining an organism’s homeostasis. Chronically hypoxic niche of poorly perfused tissues may transform tumor biology to develop drug resistance, promote cancer cell stemness, invasion, metastasis, dormany, and enhance the clonal heterogeneity.$^{[1]}$ Satellite tumors around in this transitional microenvironment may be neglected in the surgical removal of major sites. Measuring the O$_2$ concentration in these biomedical contexts could help to understand the dynamics of pathophysiology in situ and identify the problematic niches for the relapse of cancers. Conventionally, the measurement of tissue oxygen levels relies on O$_2$ microelectrodes.$^{[2]}$ Despite the accuracy and immediacy, invasive insertion...
of an electrode stimulates the studied organ and causes inevitably metabolic disorders (i.e., bleeding induced microinflammations). That artifact may change the physiological status and the actual read-out of the O₂ level. Besides, the lack of spatial information limits the use to navigate O₂ distribution in vivo.

To achieve the least invasive mapping of O₂ concentration in biomedical microenvironments, chemists have developed many transition metal complexes. Their long T₁, S₀ phosphorescence lifetime allows the collisional quenching by dissolved oxygen molecules. The quenching rate is proportional to O₂ concentration, and the corresponding partial oxygen pressure (pO₂) could be optically read-out by phosphorescence lifetime measurement. Combined with laser-scanned microscopes, the phosphorescence lifetime imaging microscopy (PLIM) can further map the microscopic distribution of oxygen in vivo. For increased measurement depth, least photodamage, and high spatial resolution, near-infrared (NIR) excited two-photon (TP) and three-photon phosphorescence probes were developed for in vivo oxygen sensing. Strongly emissive complexes of Pt, Ru, and heterometallic Ir–Eu dyads commonly show rather poor (≤100 GM) two-photon absorption (TPA) cross-sections (σ₂). The TPA could be significantly enlarged by coupling the metal ions with extended two-photon absorbing organic ligands in a metal-porphyrin (D–π)n–A architectures. Recently, large TPA and high quantum yields (Q.Y.) are simultaneously achieved in metalloporphyrin complexes.

However, probes’ phosphorescence properties could be easily affected by molecules’ structural rigidity, solvent viscosity, environment temperatures, and pH values of solutions. The binding of probes on surrounding proteins could easily change the rate of oxygen quenching. Besides, some of the probes are hydrophobic and not biocompatible if directly applied. Under these considerations, the biomedical phosphorescence probes usually need an encapsulation or surface modification to stabilize the permeation and reduce the toxicity. This protection layer unavoidably enlarges the hydrodynamic sizes up to 5–100 nm, which is >5-nm intercellular cleft of non-fenestrated capillary and 1-nm tight junctions in the brain and spinal cord. As a result, many of them need to be injected at or near the studied site with minimally invasive surgery instead of systemically intravenous administration. Least-invasive delivery can only be achieved either through ligand installation, enhanced vessel permeation in the tumor microenvironment or fenestrated capillaries of bone marrow cavities. This encapsulation or surface modification reduces the trans-vascular permeability and restricts their broad applications in biomedicine.

To overcome this last-mile hurdle of non-invasive probe delivery, in this work, we design a new class of hydrophilic Re₁-diamine luminescent complexes employing water-soluble phosphine ligands and collagen-targeting sulfonic groups. Instead of pre-encapsulation or dendrimer surface modification, we plan to let the non-toxic probes penetrate the vessels first and then stabilized with extracellular matrices. Despite a simple configuration of the target compounds, accessible modulation of their coordination sphere allowed to simultaneously achieve large two-photon absorption (σ₂ = 300 GM) and intense triplet luminescence (Φem = 0.34) in deoxygenated water at room temperature.

The deoxygenated phosphorescence lifetime at the 37°C body temperature is insensitive to the pH value. It keeps around 4 μs in water, which is long enough to sense the change of pO₂ in the microenvironment. The in vitro cell assay and in vivo animal studies showed good biocompatibility and dispersed circulation in angiography. Most importantly, without the protective carriers and encapsulation, the Re₁-diamine luminescent probes can easily penetrate typical vessel walls and diffuse into the dermis space of tissues. Owning to the high-affinity of sulfonic groups to collagens, a mechanism typically used in histopathological labeling, the probe is stabilized on tissue collagens after extravasation. Considerable enhancement in Q.Y. and a prolonged lifetime (7.5 μs) in deoxygenated conditions are achieved. No other TP phosphorescence probes ever achieved such kind of biomaterial enabled stabilization after extravasation. With high two-photon phosphorescence yields, properly long phosphorescence lifetime, high Stern-Volmer quenching constant for oxygen, good biocompatibility, and high vessel permeability, we believe the holistically advanced Re₁-diamine luminescent probes have great advantages to be applied in the in vivo studies of cell metabolism, stem cell niche, tumor hypoxic niche in poorly-perfused tissues, blood perfusion or oxygen transportation.

2. Results and Discussion

2.1. Synthesis and Characterization

Among the family of Re₁-diamine carbonyl complexes, those with phenanthroline ligand are capable of showing higher phosphorescence quantum yields in comparison to their bipyridine congeners. In particular, the dicarbonyl derivatives of [Re(phen)₂(CO)₃(PR₃)₂]⁺ reach the values of Φem above 0.4 in fluid medium at room temperature. The photo-physical properties of these species in solution were shown to be poorly sensitive to the electronic and stereochemical features of structurally similar ancillary phosphine ligands. Thus, in order to introduce hydrophilic groups with minimum interference into the optical characteristics of the [Re(phen)] chromophore, we have chosen two water-soluble phosphines, 1,3,5-triaza-7-phenyphosphetamantane (PTA) and Na₃-tris-(3-sulfophenyl)phosphine (TPPTS), which have been earlier successfully applied for the aqua-solubilization of the photoactive organometallic compounds.

The TPPTS relatives, [Re(phen)(CO)₃(TPPTS)]Na₂ (1) and [Re(phen)(CO)₃(TPPTS)]Na₅ (2) were prepared in an autoclave in water/ethanol mixtures at 130 and 220 °C, respectively and were isolated as polyamionic sodium salts (Scheme 1). The mass spectra of 1 and 2 were recorded in a negative mode and revealed the peaks at m/z 972.87 (1) and 1512.75 (2), which correspond to the monoanionic [Re(phen)(CO)₃(TPPTS)]Na₂⁻ and [Re(phen)(CO)₃(TPPTS)]Na₅⁻ (Figure S1). The NMR data obtained for 1 and 2 are completely compatible with the structural patterns suggested in Scheme 1. Each of the ⁵¹P NMR spectra displays a single resonance indicating that mono-(1) and disubstituted phosphine compounds (2) correspond to C₁ (1) and C₂ (2) symmetry point groups. The ¹H NMR spectroscopic patterns, see the Experimental section, also fit well these structural hypotheses.
Scheme 1. Synthesis of complexes 1–3 (a: TPPTS, water/ethanol 1:1 v/v, autoclave, 130 °C, 12 h, N₂; b: TPPTS, water/ethanol 1:1 v/v, autoclave, 220 °C, 12 h, N₂; c: PTA, toluene, reflux, 12 h, N₂; d: PTA, ONMe₃, 2H₂O, acetone, reflux, 12 h, N₂).

Table 1. Photophysical properties of 1–3 in H₂O (pH = 7.4) and in solid-state at 310 K.

<table>
<thead>
<tr>
<th>H₂O Solution</th>
<th>λₘₐₓ [nm] [ε×10⁻³ M⁻¹ cm⁻¹]</th>
<th>λₘᵦ [nm]</th>
<th>Φₑᵦ [a]</th>
<th>Φₑᵦ [b]</th>
<th>r [μs]</th>
<th>λₑᵦ [nm]</th>
<th>Φₑᵦ</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>366 (2.6); 319 (6); 275 (22)</td>
<td>515</td>
<td>0.05</td>
<td>0.10</td>
<td>2.91</td>
<td>540</td>
<td>1.21</td>
</tr>
<tr>
<td>2</td>
<td>379 (3.8); 296 (20); 271 (25)</td>
<td>583</td>
<td>0.15</td>
<td>0.34</td>
<td>4.05</td>
<td>537</td>
<td>9.30</td>
</tr>
<tr>
<td>3</td>
<td>404 (2.6); 293 (8); 263 (22); 221 (28)</td>
<td>634</td>
<td>0.02</td>
<td>0.03</td>
<td>0.33</td>
<td>645</td>
<td>2.19</td>
</tr>
</tbody>
</table>

[a] Aerated; [b] degassed solution; [c] excitation wavelengths λₑᵦ = 375 nm.

The above synthetic approach works well for the anion type of Re(I) complexes. For reference, we also synthesized the cationic type compound 3 (see Scheme 1), and its monophosphine predecessor 3i, which has been described previously. Both ¹H and ³¹P NMR data fit well the composition and structure of 3.

2.2. Photophysical Properties

The optical behavior of the title complexes was studied at room temperature in water solutions and in the solid-state; the pertinent data are summarized in Table 1. Absorption spectra of 1 and 2 (Figure 1a) display intense bands in the region 260–280 nm, which can be assigned to the π→π* transitions localized at the aromatic system of the phen ligand. The lower energy broad absorptions around 365–404 nm are assigned to Re→phen charge transfer (1MLCT), which shows a bathochromic shift upon the increase of the number of Re-bound phosphines. This redshift is further enhanced by the aliphatic PTA versus a triaryl TPPTS, a similar effect was noticed for congener compounds with PEt₃Ph₃ ligands.

The emission energies of the studied complexes in solution are in line with those observed in absorption spectra and with the trends typical for the Re diimine luminophores, where an increase in the number of phosphine ligands in the coordination sphere or in their donor ability (aliphatic versus aromatic phosphines) leads to a significant red shift of emission maxima. Expectedly, the dicarbonyl compounds 2 and 3 exhibit visibly red-shifted luminescence than their tricarbonyl monophosphine relatives 1 and 3i in both solution and solid phase. The luminescence intensity for these species in solid is approximately as strong as that in the degassed solution. The microcrystalline complex 2 shows high quantum efficiency (Φₑᵦ = 0.3), nearly an order of magnitude larger than that of 1. In solution, the lifetimes for 1 and 2 span the range from 2.91 to 4.14 μs (Figure 1c–f) at room temperature that points to a triplet character of the observed emission, assigned predominantly to the metal-to-ligand charge transfer (1MLCT[dₓₓ(Re)→π→π*(phen)]). Upon aeration, the quantum yield of 2 in water solution drops from 0.34 to 0.15, accompanied by a visible shortening of the lifetime (from 4.05 to 1.43 μs, Figure 1d,f). This phosphorescence property indicates collisional quenching by molecular oxygen via the triplet-triplet annihilation pathway.

As for the reference compound 3, the emission peaks are at 634 nm in solution and 645 nm in the solid phase. Because 3 turns out to be non-selectively spread in tissue fluid (vide infra), it is used as a reference compound to support the mechanism of bioconjugation suggested for 1 and 2. Therefore the photophysical and nonlinear properties of 3 are not elaborated in detail.
Figure 1. a) Normalized absorption (dashed line) and emission (solid line) spectra of 1–3 in water. \( \lambda_{ex} = 375 \text{ nm} \), b) The z-scan experimental data of 1 (green circles) and 2 (orange circles) in neutral water. Samples were held in a cuvette cell with 1-mm thickness. Solid red lines are the corresponding fit of the data points (pulse energy = 1.80 \( \mu \text{J} \), pulse width: 180 fs). \( \lambda_{ex} = 800 \text{ nm} \). c–f) Dependence of phosphorescence lifetime on oxygen concentration for complex 1 and 2 (PBS buffer, 37 °C), respectively. The \( \tau_0 \) represents the phosphorescence lifetime at 0% oxygen concentration, \( K_{SV} \) – Stern-Volmer quenching constants. The phosphorescence of 1 and 2 were single-photon excited at 375 nm and detected at around 500 nm.

2.3. Biomedical Oxygen Sensing and Imaging

High photostability of 2, intense triplet emission in water, and excellent water solubility prompted us to investigate its biomedical application in the mapping of oxygen concentration in vitro and in vivo. First of all, we established the relationship between the phosphorescence lifetime and \( O_2 \) concentration by the Stern–Volmer equation:

\[
\frac{\tau_0}{\tau} = 1 + K_{SV}[O_2]
\]

where \( \tau_0 \) represents the phosphorescence lifetime under \( O_2 \)-free conditions, and \( K_{SV} \) is the quenching constant. In the physiological medium (PBS buffer, pH = 7.4 at 37 °C), the Stern–Volmer plots of 1 and 2 are approximately linear in the range of 0–30% oxygen mole fraction in 1 atm gas, and the \( K_{SV} \) values are 6.1 and 9.2 atm\(^{-1}\), respectively (Figure 1c–f), which indicate a moderate quenching sensitivity. We also investigated the temperature and pH dependence of phosphorescence lifetime (Figure S3, Supporting Information). The data obtained indicate that lifetime decreases as temperature increases. In contrast, the lifetime of 2 is insensitive to the pH variations in the range from 5.1 to 8.5. (Figure S3c, Supporting Information), that prevent the lifetime signal crosstalk and is evidently beneficial to the oxygen sensing.

For deep-tissue in vivo bio-imaging and sensing, the probe needs to have strong two-photon absorption and efficient two-photon phosphorescence. We investigated the two-photon absorption properties of 1 and 2 in pure water by the open-aperture Z-scan method.[57] Upon fitting the data (Figure 1b), the two-photon absorption cross-section 2 (was then calculated to be 173 ± 7 GM and 300 ± 5 GM for 1 and 2, respectively. These
values are comparable to that of Coumarin 480 of 148 GM in ethanol upon excitation at 800 nm.\textsuperscript{[57]}

To verify their biocompatibility, we carried out the MTT test to assess the viability of human liver carcinoma (HepG2) cells. Incubating cells with the probes for 15 hours, we observed no obvious cytotoxicity even at a high concentration up to 500 μM (Figure S4, Supporting Information). We also confirmed the uptake of probes by HepG2 cells via bright-field, two-photon phosphorescence images and the corresponding two-photon luminescent spectra, which are shown in Figure S5, Supporting Information. The intracellular luminescence spectra fit well those obtained for 1 and 2 in an aqueous solution. These observations clearly indicate that probes 1 and 2 are biocompatible and could be successfully internalized by cells.

To verify the collagen-targeting specificity, as a first step, we measured the phosphorescence lifetime of 2 in various biological media in vitro. Results showed that phosphore buffer saline (PBS), albumin solution and fetal bovine serum (FBS) did not change the lifetime properties of a lot (Table S1, Supporting Information) either at 0% O\(_2\) or at 20% O\(_2\) condition. The pure Matrigel, containing extracellular matrix proteins, slightly decreases the lifetime at 0% O\(_2\), which may be due to the change of solution viscosity. We then mixed the Matrigel with lipopolysaccharide (LPS;10 mg mL\(^{-1}\)) interferon-Gama (IFN-γ; 20 ng mL\(^{-1}\)) and 30% L929 fibroblast conditioned medium and plugged it into the abdomen space of a C57BL/6-C2J mouse. On the 7th day post-implantation, we took the Matrigel out for imaging. As we previously demonstrated, the Matrigel recruited M1 macrophages and fluorescence lifetime was found at 7.5 μs at 0% O\(_2\), and dropped to 5 μs at 21.5% O\(_2\). Following the \textit{ex vivo} calibration curve, the 6.5 μs lifetime around the vessel corresponds to a 6.46% O\(_2\) (pO\(_2\) = 49.1 mm Hg) under tissue normoxia conditions. It is well-known that the tissue oxygenation level highly depends on the blood flow and cellular consumption rate to give O\(_2\) concentration in normal oxygenation from 1% to 9.5%.\textsuperscript{[60]} Thus, the \textit{in vivo} obtained lifetime value (6.5 μs) corresponds to oxygen concentration, which falls in the range typical for normoxia conditions.

To evaluate whether the probes can detect the variation of tissue oxygen levels in vivo, we induced partial hypoxia in the tissues by pressing the vessels at the root of the mice ear. Under these conditions, the tissues consume more oxygen than supply, and the phosphorescence lifetime histogram in PLIM finally showed an increased peak around 7 μs (Figure 3e). This result corresponds to a 22.8 mm Hg pO\(_2\) (3% O\(_2\)). Since the circulation might not be fully blocked, the complete hypoxic conditions were not reached. The oxygen pressure returned to a normal level of 53.2 mm Hg (PLIM histogram peaks at 6 μs) 10 min after reperfusion (Figure 3e). Accordingly, long-term inhalation of \(\text{O}_2\) 100% at 760 mm Hg gave peak of lifetime histogram downshifted to 5.25 μs (pO\(_2\) = 137 mm Hg; 18% O\(_2\)). We then investigated the gradient of oxygen concentration near large vessels (green color in Figure 4) was 10% (pO\(_2\) = 76 mm Hg), which is obviously higher than those in the regions far away from the vessel (blue color in Figure 4).\textsuperscript{[61]} At a depth of 29 μm away from the vessel, the average oxygen concentration can be decreased down to 5% (pO\(_2\) = 76 mm Hg) level (Figure 4 right).\textsuperscript{[62]}

Finally, the hypoxia conditions in the melanoma microenvironment were carefully investigated (Figure 5). We implanted mCherry labeled melanoma cells into the C57BL/6-c2J (c2J) mice...
ear, and the tumor nodule appeared after 2–4 weeks. After locating the tumor position in vivo through the mCherry fluorescence, we tail-vein injected the probes for the PLIM. We found pathological hypoxia (pO$_2$ = 7.6 mm Hg; 1% O$_2$) at the center of the tumor, where PLIM display the average lifetime of 7.32 μs. At the edge of the tumor, the oxygen tension increased to 22.8 mm Hg (7 μs; 3% O$_2$) and up to 38.1 mm Hg at the region 5-mm away from the edge (6.7 μs; 5% O$_2$). These results agree well with the change in oxygen tension during tumor development.[63]

Evidently, it is of particular importance to double-check the reasons for the high binding affinity of 2 to collagens. For the sake of comparison, we performed angiography experiments with complexes 1, 2, and 3 and found they all penetrated the vessel walls. Two-photon phosphorescence signals of 1 and 2 (Figure S10, Supporting Information, green and yellow colors, respectively) are co-localized with the second harmonic generation signals of collagen fibers (blue colors in Figure S10, Supporting Information). In sharp contrast, complex 3 tends to spread evenly over the extravascular space without any clusterization. Because (1, 2) and (3) contain the same 1,10-phenanthroline and –CO ligands in their coordination sphere, the difference in collagen affinity cannot be attributed to these groups. A similar chemical design also excludes the intercalation of 1,10-phenanthroline into the triple helix region of collagens. Therefore, most plausibly, the outer shell ionic properties account for this discrepancy. The most reasonable hypothesis might be that in the case of 1 and 2 the peripheral sulfonic anions act as tripods, which can anchor the complexes to collagen by either electrostatic interaction or hydrogen bonding with free amino or ammonium residues.[64] This viewpoint can be envisaged by a cartoon-like schematic diagram shown in Scheme 2. On the other hand, the positive charge in the cationic complex 3 is located at the central Re ion, the peripheral ligand of which may only be the subject of a weak van der Waals interaction. As a result, 3 spreads evenly in the entire tissue fluid without specific affinity to collagen-binding. Accordingly, we report an intriguing case that collagen can be functionalized by vessel-penetrating probe 2 (and 1), and then successfully applied in TP-PLIM to visualize tumor hypoxia and tissue ischemia in vivo via oxygen mapping.

### 3. Conclusions

In this study, we developed a series of water-soluble and biocompatible Re$^{II}$-diimine carbonyl complexes that can easily penetrate...
the vessel walls and detect oxygen tension in vitro and in vivo for a long period. Among these probes, complex 2 exhibits excellent two-photon absorption cross-sections (300 GM) under the excitation with an 800-nm femtosecond laser. High two-photon emission intensity allows visualizing the distribution of oxygen in the deep tissue regions. Moreover, a wide range of phosphorescence lifetime ($\Delta \tau \approx 3 \mu s$) between oxygenated and deoxygenated conditions makes it sensitive enough to quantify the oxygenation in biological tissues, which is typically lower than 10%. Most importantly, complex 2 can easily permeate into tissues and form a stabilized bioconjugate with collagens for at least 6 h. This “2-collagen” bioconjugate outperforms the starting complex with enhanced quantum yields and lengthened phosphorescence lifetime. Since complex 2 was stuck on collagen and did not diffuse away quickly, it could serve as stationary oxygen sensors at local tissues. Then in the in vivo studies, we found that the phosphorescence lifetime of 2 increased from 5.25 $\mu s$ nearby vessels to 7 $\mu s$ in the avascular area. Based on the ex vivo calibration, the corresponding oxygen tension decreased from 10.17% to 5.32%, which agrees well with the oxygen level in skin tissue under physiological conditions.$^{(60)}$ A lower level (<5%) of oxygen was found in pathological hypoxia. Lifetimes around 7.3 $\mu s$ and 6.9 $\mu s$ were found at the center and at the edge of the melanoma, corresponding to the oxygen level of 1% and 4%, respectively. These results suggest that the water-soluble and biocompatible ReI-complex 2 is, with the help of deep-tissue TP-PLIM, suitable for monitoring tissue metabolism, ischemia, and tumor hypoxia. This post-extravasation stabilization and lifetime lengthening by collagen render a new strategy for designing phosphorescence probes in biological tissues.

4. Experimental Section

**Synthesis General Comments:** 1,3,5-Triaza-7-phosphaadamantane (PTA),$^{(63)}$ [Re(phen)(CO)$_3$(NCMe)](CF$_3$SO$_2$)$_2$ were prepared according to published procedures. Toluene was distilled from Na-benzophenone ketyl under nitrogen prior to use. Other reagents and solvents were used as received. The solution $^1$H and $^{31}$P($^1$H) spectra were recorded on a Bruker Avance 400 spectrometer. Mass spectra were measured on a Bruker APEX-Qe Qb-FT-ICR instrument in ES$^+$ (1, 2) modes. Microanalyses were carried out at the analytical laboratory of the University of Eastern Finland.
Figure 4. Two-photon PLIM tomography of normal skin at various depths. The emission intensity of the complex 2 in tissue was shown in grayscale images. The vertical distance (in μm) from the giant vessel was recorded by the objective stage, and the corresponding oxygen concentration (estimated from the peak value in the lifetime histogram) was presented. Oxygen decreases rapidly as the distance increase. Field of view: 240 × 240 μm.

Figure 5. a) TP-PLIM results of complex 2 at the center of tumor and 5-mm away from tumor in a mouse’s ear. b) Corresponding lifetime histograms measured around the region indicated by yellow arrow. Field of view: 100 × 100 μm.

Scheme 2. The proposed structure of 2 complex anchoring on collagen via electrostatic and hydrogen bonds. His: Histidine; Arg: Arginine.

[Re(phen)(CO)3(TPPTS)2]Na5 (2): [Re(phen)(CO)3(CH3CN)](CF3SO3) (118 mg, 0.184 mmol), TPPTS (262 mg, 0.461 mmol) and the mixture of water with ethanol (1:1 v/v, 15 mL) were placed in a 60 mL autoclave, pressurized with nitrogen to 25 atm and heated at 220 °C overnight. The resulting yellow solution was evaporated and the residue was washed with cold methanol (2 mL). Recrystallization by a gas phase diffusion of acetone into water solution of 2 at room temperature to afforded bright yellow fine crystalline material (125 mg, 44%). ESI-MS (m/z): [M+Na]− 1512.75 (calcd 1512.85), [M–TPPTS+Na]− 944.88 (calcd 944.94). IR (CH3CN, ν(CO), cm−1). 1H NMR (D2O, 298 K; δ): 8.43 (d, JHH 5.0 Hz, 2H, 2,9-H phen), 8.23 (d, JHH 8.1 Hz, 2H, 3,8-H phen), 7.60 (m, 6H, TPPTS), 7.41 (m, 6H, TPPTS), 7.30 (m, 12H, TPPTS), 7.17 (dd, JHH 8.1 and 5.0 Hz, 2H, 5,6-H phen). 31P{1H} NMR (D2O, 298 K; δ): 24.1 (s). Anal. Calcd for C50H32N2Na5O20P2ReS6.4H2O: C 37.33; H 2.50; N 1.74; S 11.96. Found: C 37.22; H 2.52; N 1.69; S 12.08.

[Re(phen)(CO)3(TPA)](CF3SO3) (3i): [Re(phen)(CO)3(CH3CN)](CF3SO3) (500 mg, 0.78 mmol) and PTA (126 mg, 0.803 mmol) were dissolved in toluene (60 mL). The mixture was refluxed overnight under a nitrogen atmosphere. The solvent was removed and the residue was recrystallized by a gas phase diffusion of diethyl ether into an acetone/methanol solution of 1 at room temperature to give yellow-green solid (510 mg, 87%). ESI-MS (m/z): [M+] 608.08 (calcd 608.08). IR (CH3CN, ν(CO), cm−1): 2041s, 1954s, 1928s. 1H NMR (acetone-d6, 298 K; δ): 9.17 (dt, JHH 5.2 and 1.2 Hz, 2H, 2,9-H phen), 9.13 5.2 Hz, 2H, 2,9-H phen), 8.67 (d, JHH 8.2 Hz, 2H, 3,8-H phen), 8.09 (s, 2H, 4,7-H phen), 7.85 (dd, JHH 5.2 and 8.2 Hz, 2H, 5,6-H phen), 7.78 (d, JHH 9.7 Hz, 6H, TPPTS), 7.31 (dd, JHH 8.0 and 7.1 Hz, 3H, TPPTS), 7.13 (dd, JHH 8.0 Hz, 3H, TPPTS, 7.13 (d, JHH 9.4 Hz, TPPTS). 31P{1H} NMR (acetone-d6, 298 K; δ): 9.21. Anal. Calcd for C40H28N2Na5O20PReS2.4H2O: C 37.97; H 2.61; N 1.69; S 11.96. Found: C 37.97; H 2.61; N 1.69; S 11.96.
(ddd, $J_{HH}$ 8.3, 1.2 and 0.9 Hz, 2H, 3,8-H phen), 8.46 (s, 2H, 4,7-H phen), 8.27 (dd, $J_{HH}$ 5.2 and 8.3 Hz, 2H, 5,6-H phen), 4.34 (d, $J_{HH}$ 13.2 Hz, 3H, PTA), 4.25 (d, $J_{HH}$ 13.2 Hz, 3H, PTA), 3.82 (s, 6H, PTA). $^{31}P$ NMR (acetone-d$_6$, 298 K; δ): 82.0 (s). Anal. Calcd for C$_{27}$H$_{32}$F$_3$N$_8$O$_5$P$_2$ReS: C 36.61; H 3.64; N 12.65; S 3.62. Found: C 36.67; H 3.69; N 12.71; S 3.59.

**Bioanalytical Study:**

1) **In vitro Cytotoxicity Assay:** To assess the cytotoxicity of the complexes 1 and 2, we treated them on the HepG2 cells and analyzed the cell viability by a colorimetric assay agent, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT, Roche). The HepG2 cells were seeded in a 96-well plate with 5 × 10$^4$ cells per well in a 90% Dulbecco’s Modified Eagle Medium supplemented with 10% heat-inactivated fetal bovine serum, penicillin, and streptomycin. To contrast the control differences in the concentrations of the complex, the cells were added to each well. After 15 h of incubation, the cells were washed twice with PBS buffer and then incubated with 200 μL of the culture medium with 10% MTT agent per well. After 4 h of reaction time, the culture medium was removed and refilled with 200 μL of dimethyl sulfoxide (Sigma-Aldrich) per well to dissolve the purple MTT formazan crystal. The optical density of these samples was measured at 595 nm. All measurements were done with three replicates using an ELISA reader (VersaMax Microplate Spectrophotometers; Molecular-Devices). The uptake of probes in HepG2 cells was also validated via two-photon luminescence microscopy and spectroscopy under the illumination of a 800-nm femtosecond laser.

2) **Ex vivo Calibration:** To calibrate the phosphorescence quenching rates of collagen-bound 2, we measured the TP-PLIM of the complex 2 administered to the collagen tissue in vivo (Figure S9, Supporting Information). The fresh collagen tissue was excised from a sacrificed mouse’s ear and then bathed in the normal saline with 100 μm complex 2 at 33 °C. The prepared sample in saline was held on a bottom-glass petridish for 800 nm excited TP-PLIM. The oxygen mole fraction in the measurement chamber was adjusted by O$_2$–N$_2$ gasning system and monitored by an oxygen meter (500DL, Unisense) during the experiment. All the experiments were done within an hour. Finally, we generated the lifetime histogram of each PLIM image (Figure S9c, Supporting Information) and chose the peak lifetime as the representative one at different oxygen mole fraction (Figure S9b, Supporting Information).

3) **Phosphorescence lifetime of probes in biological substances and Matrigel:** In order to make sure it’s the collagen that stabilizes phosphorescence probes and achieves a longer phosphorescence lifetime, we mixed (100 μm) complex 2 in PBS buffer, albumin solution (0.2 mg mL$^{-1}$), and FBS (10%, v/v, solvent: saline). Under normoxia and 0% O$_2$, we measured the phosphorescence decay traces and fit the corresponding lifetimes.

Furthermore, the phosphorescence lifetime of complex 2 in Matrigel was also studied. 0.25 mL Matrigel containing 10 ng mL$^{-1}$ LPS, 20 ng mL$^{-1}$ IFN-γ, and 30% L929 cell-conditioned medium was injected into the abdomen of C57BL/6-C2J mouse. Seven days later, the mouse was euthanized with CO$_2$, and the Matrigel was taken out. The Matrigel was filled with biosynthesized collagen fibers. Then the Matrigel was incubated with complex 2 solution (100 μM) and observed with PLIM system under 20% Oxygen ambient condition and depleted oxygen with a nitrogen purge. As a control, the PLIM of pure Matrigel was also characterized.

4) **In vivo Studies:** The animal study was approved by the Institutional Animal Care and Use Committee (IACUC) of National Taiwan University (the assigned approval number of the laboratory: NTU107-EL-00135). The C57BL/6-c2J (c2J) mice were provided by the National Taiwan University College of Medicine. About 10$^3$ melanoma cells (cell lines: B16F1 mCherry) were injected subcutaneously in 4-week-old mice’s ears. Tumor environments were imaged by two-photon microscopy and PLIM as it grew to 1 mm in diameter. A 2% of isoflurane mixed with 98% O$_2$ was used to initiate anesthesia. During the experiment, the anesthesia was achieved by injecting Avertin through IP with the proper dosage (0.25 mg g$^{-1}$).

For the preparation of the stock solution, complex 2 was dissolved in sterilized normal saline and then pass through a syringe filter with 0.22 μm pore size. A 0.2-mL solution was injected through the tail vein of mice. The final concentration of 2 in mice was estimated to be 100 μM. The ear was mounted on cover glasses and then observed by the multiphoton microscope. To provide two extreme oxygen levels in mice ear, on the one hand, we supplied pure oxygen gas via inhalation; on the other, we blocked the vessels at the root of the ear. Cessing the blood circulation was done by sandwiching a bent 26G needle, and the mice ear between a coverslip and a cover glass. The needle was put upon and cross over the major vessel of the mice ear. The additional force was applied on both sides of the glass to ensure the blockage. The color of tissue and blood was turned into white and purple, respectively, during this procedure. For the reperfusion studies, we simply increased the space between two glasses by inserting a tweezer and removed the needle simultaneously. A high tissue oxygen level was reached by the inhalation of pure oxygen (760 mm Hg). Gas was warmed to 33 °C and then delivered via a tube, which connected a mask at the end. The mask covered the entire snout of mice during the experiment. To prevent irreversible damages to the tissues, we only held the ischemic condition for 10 min. For depth-dependent examinations, the focal plane of images was shifted 5 μm per step by a motorized objective stage. The environment was kept around 33 °C to maintain the body temperature of mice. Their vital sign was monitored during the experiment.

**Multiphoton Luminiscence and PLIM Imaging System:** Multiphoton luminescence imaging was performed on a commercial laser-scanning microscope (Observer.Z1 and LSM 710, Zeiss) equipped with a micro-incubator and a fluorescence/phosphorescence lifetime imaging system (SPC-150, Becker & Hickl). An 80-MHz Ti: Sapphire laser (Mai Tai DeepSee, Spectra-Physics) was used as the excitation source. The 800 nm excitation beam was focused on samples by a 40X water immersion objective (C-APO, Zeiss), and the average power was kept below 10 mW for in vitro study and 20 mW for in vivo study. To measure the phosphorescence decay, we turned the laser excitation on for 5 μs at a 20-kHz gating rate by the DGC-200 module (Becker & Hickl). This resulted in a 50 μs pixel dwell time for PLIM, which is much longer than the phosphorescence lifetime of complex 2. During the PLIM imaging, the frame time was 3.3 s.
and the overall image acquisition time was 5 min in each area to collect high enough photon counts (>100 photons) for correct curve fitting. Data analysis was performed by the software SPCI mage 6.0 (Becker & Hickl).

Retention of Complex 2 In Vivo: 1.5 mM of complex 2 (solution: saline) was injected intravenously into C57BL/6-C2J mouse via tail vein injection. Then the ear of the mouse was mounted with cover glass and observed with Nikon Inverted Multiphoton Microscope Eclipse (Nikon instrument Inc) equipped with a fluorescence/phosphorescence lifetime imaging system (Becker & Hickl) under a 40× NA = 1.15 water-immersion objective at different time points (0, 5, 10, 15, 20, 25, 30 min, and 6 h). During the experiment, the anesthesia was achieved by injecting Avertin through IP with a proper dosage (0.25 mg g

Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

Acknowledgements

C.-H. W. and K.S. K. contributed equally to this work. Financial support from the Academy of Finland (grant 317903, I.O.K.; Flagship Programme, Photonics Research and Innovation PREIN, decision 320166), Russian Science Foundation (grant 19-73-20055, J.R.S., synthesis of rhenium complexes); the Academy of Finland (grant number 317903, I.O.K.; Flagship Programme, MYRG2018-00070-FHS and MYRG2019-00022-FHS), from The Science and Technology Development Fund, Macau SAR (File no. 112/2016/A3, 018/2017/A1, 0011/2019/AKP, and 0120/2020/A3), and from National Health Research Institutes, Taiwan (grant number NHRI-EX105-10427EI). The authors would like to thank Prof. Russell A. Prough in the University of Louisville for his plasmid with ALDH1A1 promotor. Pi-Tai Chou thanks the Ministry of Science and Technology, Taiwan for financial support.

Conflict of Interest

The authors declare no conflict of interest.

Data Availability Statement

Research data are not shared.

Keywords

phosphorescence lifetime imaging microscopy, phosphorescent oxygen sensors, Re\textsuperscript{II} diimine carbonyl complexes, tissue ischemia, tumor hypoxia, two-photon phosphorescence


Received: July 1, 2021          Published online:


