Electronic Supplementary Information

A series of iridophosphors with tunable excited states for hypoxia monitoring via time-resolved luminescence microscopy

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Table S1 HOMOs and LUMOs distributions of Ir1–Ir4 at T_1 and S_1 states



Fig. S1 Plots of average phosphorescence intensity as a function of oxygen contents in HeLa cells incubated with **Ir1** (10 μ M).



Fig. S2 (a) Confocal luminescence images in Hela cells incubated with **Ir3** (10 μ M) under different oxygen level; (b) Photoluminescence lifetime images in Hela cells incubated with **Ir3** (10 μ M) under different oxygen level.

NMR and MALDI-TOF spectra



Fig. S3 ¹H NMR spectrum of 9-(5'-bromo-[2,2'-bipyridin]-5-yl)-9*H*-carbazole.



Fig. S4 ¹³C NMR spectrum of 9-(5'-bromo-[2,2'-bipyridin]-5-yl)-9*H*-carbazole.



Fig. S5 ¹H NMR spectrum of ligand **1**.



Fig. S6 ¹³C NMR spectrum of ligand 1.



Fig. S7 ¹⁹F NMR spectrum of ligand 1.



Fig. S8 ¹H NMR spectrum of Ir1.



Fig. S9¹⁹F NMR spectrum of Ir1.



Fig. S10 ¹H NMR spectrum of Ir2.



Fig. S11¹⁹F NMR spectrum of Ir2.



Fig. S12 ¹H NMR spectrum of Ir3.



Fig. S13 ¹⁹F NMR spectrum of Ir3.



Fig. S14 ¹H NMR spectrum of Ir4.



Fig. S15¹⁹F NMR spectrum of Ir4.



Fig. S16 MALIDI-TOF spectrum of Ir1.



Fig. S17 MALIDI-TOF spectrum of Ir2.



Fig. S18 MALIDI-TOF spectrum of Ir3.



Fig. S19 MALIDI-TOF spectrum of Ir4.



Fig. S20 The corresponding Stem–Volmer plots of **Ir2–Ir4** (a-c) under the quenching by oxygen (red points stand for the values of I_0/I and corresponding $K_{SV} = 0.129$ (a), 0.032 (b) and 0.017 (c), respectively; black points stand for the values of τ_0/τ and corresponding $K_{SV} = 0.137$ (a), 0.031 (b) and 0.018 (c), respectively.).