Supplementary Material (ESI) for Journal of Materials Chemistry This journal is (c) The Royal Society of Chemistry 2013

Supporting Information

Dinuclear iridium(III) complexes as phosphorescent

trackers to monitor mitochondrial dynamics

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Fig. S1 ¹H NMR spectra of **Ir1** in d_6 -DMSO, 500 MHz



Fig. S2 ¹H NMR spectra of Ir2 in d_6 -DMSO, 500 MHz



Fig. S3 ¹H NMR spectra of **Ir3** in d_6 -DMSO, 500 MHz



Fig. S5 ¹H NMR spectra of Ir5 in d_6 -DMSO, 500 MHz

11.5 11.0 10.5 10.0 9.5 9.0 8.5 8.0 7.5 7.0 6.5 6.0 5.5 fl (ppm)

5

8 4 503

4.0

3.5 3.0

2.5 2.0 1.5

5.0 4.5

0.5 0.0 -0

1.0



Fig. S6 ¹H NMR spectra of Ir6 in d_6 -DMSO, 500 MHz



Fig. S7 ¹H NMR spectra of Ir7 in d_6 -DMSO, 500 MHz



Fig. S8 Confocal phosphorescence image, bright field image and their overlay of living HeLa cells incubated with 5 μ M of **Ir1** in DMSO and PBS (pH = 7.4, 1:99, v/v) for 30 min at 37 °C, followed by 50 nM of MTR. (a) confocal phosphorescence images of **Ir1**; (b) bright field images; (c) confocal luminescence images of MTR; (d) overlay of a, b and c. Excitation wavelength: 405 nm for **Ir1** and 488 nm for MTG; emission filter: 580 ± 10 nm for Ir1 and 520 ± 10 nm for MTG.



Fig. S9 *In vitro* cell viability of HeLa cells incubated with 5 μ M of **Ir2-Ir7** at 37 °C for 12 h and 24 h, respectively.



Fig. S10 *In vitro* cell viability of LO2 cells incubated with 5 μ M of **Ir2-Ir7** at 37 °C for 12 h and 24 h, respectively.