Domino-fluorination-protodefluorination enables decarboxylative cross-coupling of α -oxocarboxylic acids with styrene via photoredox catalysis

Muliang Zhang,† Junwei Xi,† Rehanguli Ruzi,† Nan Li,† Zhongkai Wu,† Weipeng Li,† Chengjian Zhu*†‡

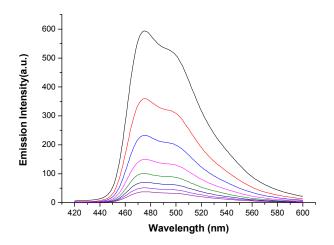
†State Key Laboratory of Coordination Chemistry, Jiangsu Key Laboratory of Advanced Organic Materials, School of Chemistry and Chemical Engineering, Nanjing University, Nanjing 210093, China †‡State Key Laboratory of Organometallic Chemistry, Shanghai Institute of Organic Chemistry, Shanghai 200032, China. E-mail: cjzhu@nju.edu.cn

Table of Contents

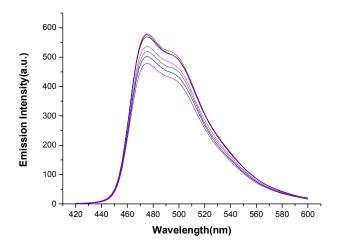
1. Luminescence quenching experiment	S2
2. Copies of ¹ H NMR, ¹³ C NMR, ¹⁹ F NMR Spectra	S3

1. Luminescence quenching experiment

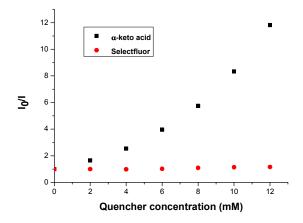
The luminescence quenching experiment was taken using a Cary Eclipse fluorescence spectrophotometer (Varian, USA). The experiments were carried out in 2.5 x 10^{-5} mol/L of [Ir{dF(CF₃)ppy}₂{dtbbpy}]PF₆ in CH₃CN-H₂O (V/V, 1:1) at 25 °C. The excitation wavelength was 350 nm and the emission intensity was collected at 475 nm. The concentrations of quenchers (α -keto acid 1 and Selectfluor) in CH₃CN-H₂O were 0, 2, 4, 6, 8, 10, 12, 14mM.



Luminescence quenching of [Ir{dF(CF₃)ppy}₂{dtbbpy}]PF₆ by α -keto acid



Luminescence quenching of [Ir{dF(CF $_3$)ppy} $_2$ {dtbbpy}]PF $_6$ by Seclectfluor



 $Ir[dF(CF_3)ppy]_2(dtbbpy)PF_6\ emission\ quenching\ with\ Selectfluor\ and\ benzoylformic\ acid.$

2. Copies of ¹H NMR, ¹³C NMR spectra

