

Supplementary Information

Evidence of multiple pathways capable of emitting peroxyoxalate chemiluminescence using a charge coupled device spectrometer

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Table 1 Comparison of background emission characteristics generated in ODI derivative CL reactions

	<i>ODI^d</i>	<i>OD2MI^d</i>	<i>OD2EI^d</i>	<i>OD4MI^d</i>	<i>OD2MI + OD4MI^d</i>
I_{\max}^a	0.60	0.05	0.02	1.00	0.80
τ_{\max}^b	0.34	0.40	0.52	0.30	0.30
τ_{half}^c	0.90	2.09	3.87	0.45	0.45

^a Relative maximum background intensity for each experimental condition normalized by that observed for the OD4MI CL reaction. ^b Time to reach the maximum emission (sec). ^c half-life of the decay (sec). ^d ODI derivatives formed from the reaction between TCPO and ImH derivatives for 90 seconds at room temperature (21 ± 2.0 °C). Reaction condition: [TCPO] = 1.0 mM, [ImH derivative] = 10.0 mM, [H₂O₂] = 0.1 M in the mixture of 50% acetonitrile + 50% ethyl acetate. Background emissions were measured with PTI with photo-multiplier tube.

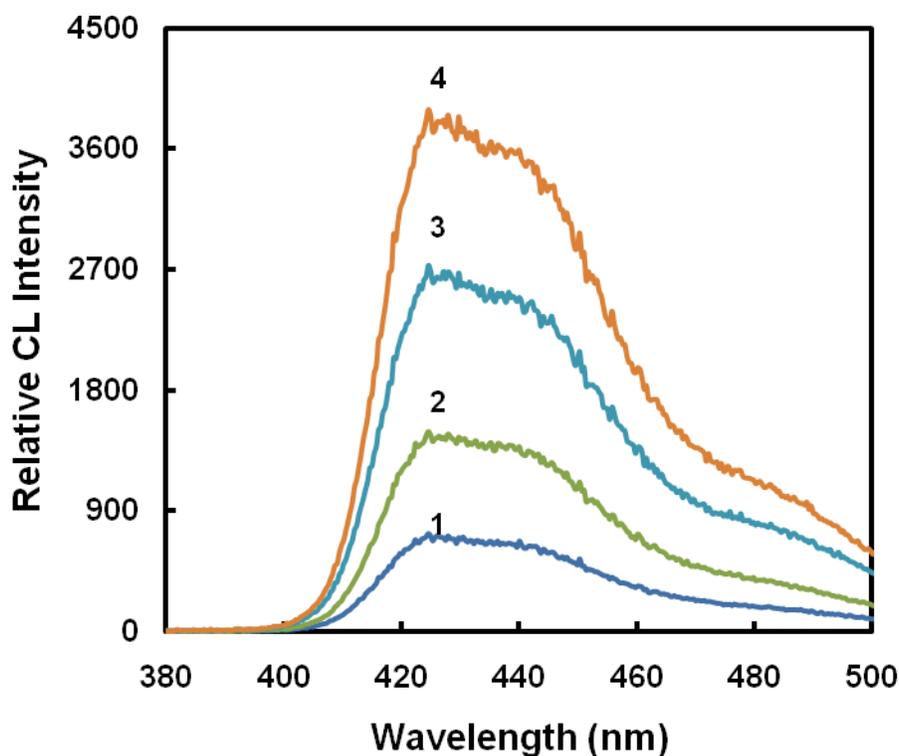


Fig. S-1. CL spectra observed in ODI derivative CL reactions.
1. OD2EI-CL, 2. OD2MI-CL, 3. ODI-CL, 4. OD4MI-CL.

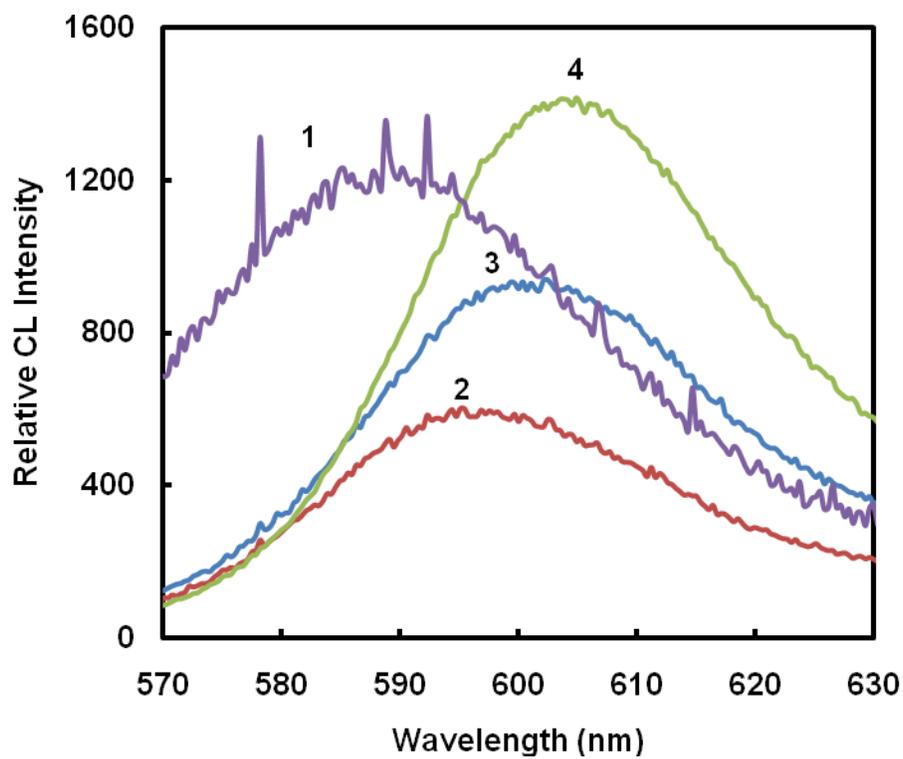


Fig. S-2. Fluorescence and CL spectra of rhodamine 101 in mixture (50 % ethyl acetate, 25 % water, and 25 % isopropyl alcohol). 1. Fluorescence, 2. ODI-CL, 3. TCPO-CL, 4. DNPO-CL