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Supporting Information for

"Electrooxidation of rhodamine B hydrazide"

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The Supporting Information contains additional LC/MS data. Figure S1 shows the mass spectrum for as-synthesized rhodamine B hydrazide (RBH). Figure S2 presents the chromatogram obtained after the electrooxidation of RBH. Figures S3–S5 demonstrate the absorption and mass spectra for the electrooxidation products of RBH: rhodamine B (compound 1), a phenanthrenone derivative (compound 2), and a phtalazinone derivative (compound 3).



4 5 6 7 8 9 Response Units vs. Acquisition Time (min) 10 Figure S2. Chromatogram for a 0.2 mM solution of RBH in the supporting electrolyte (0.1 M LiClO₄ in acetonitrile) after electrolysis at 0.9 V (vs. SCE) for 60 min. The chromatogram was recorded at 510 nm.

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3



Figure S3. Mass spectrum (a) and UV–Vis absorption spectrum (b) for peak 1 in the chromatogram of Figure S3. This peak is assigned to compound 1 (rhodamine B).



Figure S4. Mass spectrum (a) and UV–Vis absorption spectrum (b) for peak 2 in the chromatogram of Figure S3. This peak is assigned to compound 2 (a phenanthrenone derivative).



Figure S5. Mass spectrum (a) and UV–Vis absorption spectrum (b) for peak 3 in the chromatogram of Figure S3. This peak is assigned to compound 3 (a phtalazinone derivative).