Supporting Information

Switching On/Off the Chemisorption of Thioctic-based Self-Assembled Monolayers on Gold by Applying a Moderate Cathodic/Anodic Potential

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Figure S1. (A) CVs of a redox-active SAM on gold ($\Gamma_{Fc} = 2.1 \times 10^{-10} \text{ mol/cm}^2 \text{ FcCO}_2\text{DT}$) recorded at different scan rate (from top to bottom: 10, 8, 6, 4, 2 and 1 V/s) and (B) plot of the anodic peak current density ($j_{p,a}$) as a function of scan rate.



Figure S2. CVs of a redox-active SAM on gold ($\Gamma_{Fc} = 2.1 \times 10^{-10} \text{ mol/cm}^2 \text{ FcCO}_2\text{DT}$) recorded at 1 V/s in 0.1 M TBABF₄/CH₃CN.



Figure S3. Normalized response (θ) of the (•) surface coverage and (•) capacitance densities plotted in Figure 2E.



Figure S4. (A) Chronoamperometric curves obtained after stepping the potential of a bare gold electrode from +0.5 to -0.4 V in (blue) a 20 μM FcCO₂DT solution and (red) in a FcCO₂DT-free solution. (B) Chronoamperometric curves obtained after stepping the potential of (blue) a saturated FcCO₂DT-SAM-coated gold electrode and (red) a bare gold electrode from -0.4 to +0.5 V in a pure solution of 0.1 M TBABF₄/CH₃CN.



Figure S5. Theoretical transient current responses deduced from the first derivation of the double-exponential fits of (A) the absorption kinetic plot shown in Figure 2E (red plot) and (B) desorption kinetic plot shown in Figure 2F.