

Supporting Information

**Post-imprinting Modification: Electrochemical and Scanning Electrochemical
Microscopy Studies of Semi-Covalently Surface Imprinted Polymer**

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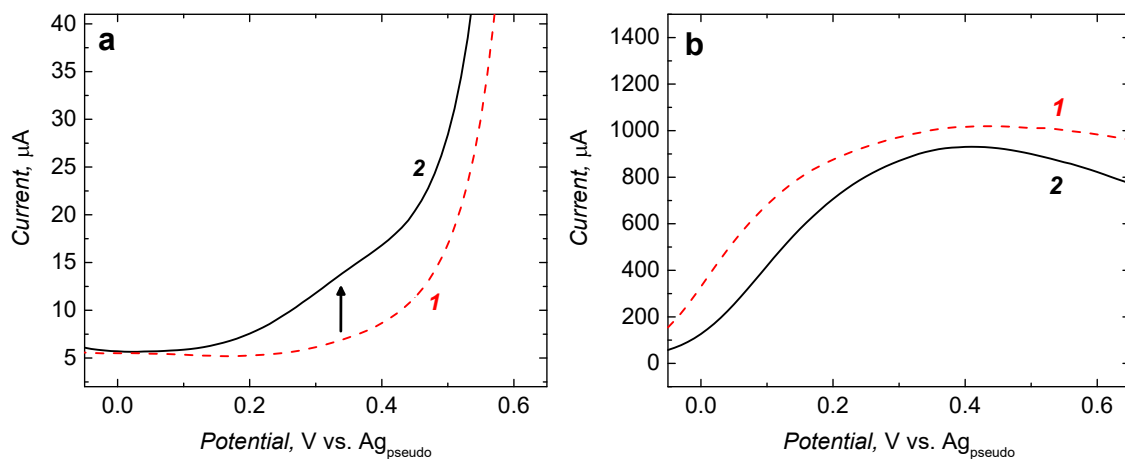


Figure S1. SWV measurement performed in 0.1 M $\text{LiClO}_4/\text{ACN}$ for co-polymers of (a) 3,3'-bithiophene (**CR2**) and 2,2'-bithiophene-5-carbonyl chloride (**FM1**) and (b) 4H-cyclopenta[1,2-b:5,4-b']dithiophene (**CR4**) and 2,2'-bithiophene-5-carbonyl chloride (**FM1**) (1) before and (2) after covalently immobilized ferrocene modification. Measurement conditions include pulse height: 25 mV, pulse width: 50 ms, and step height: 10 mV.

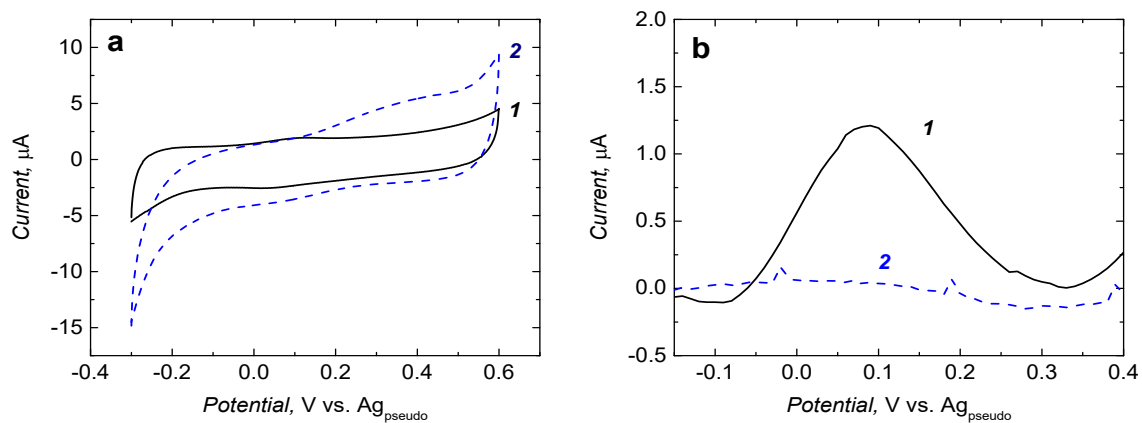


Figure S2. (a) Cyclic voltammetry and (b) square wave voltammetry measurements of BSA post-imprinted modified electrode. Semi-covalent surface imprinted polymers were prepared with (1) **FM2** and **CR2** (2) **FM1** and **CR3** combinations. Electrolyte: 0.5 mM H₂SO₄, CV: 50 mV/s, SWV: pulse height: 25 mV, pulse width: 50 ms, step height: 10 mV.

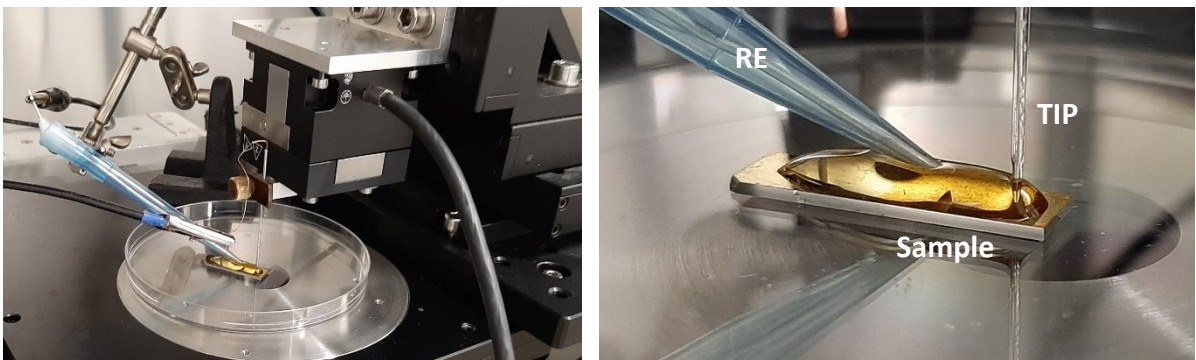


Figure S3. SECM measurement setup for polymer surface characterization. RE – Ag pseudo reference and counter electrode, TIP – platinum microelectrode (ca. 25 μm diameter), Sample - post-imprinting polymer modified electrode or NIP polymer.

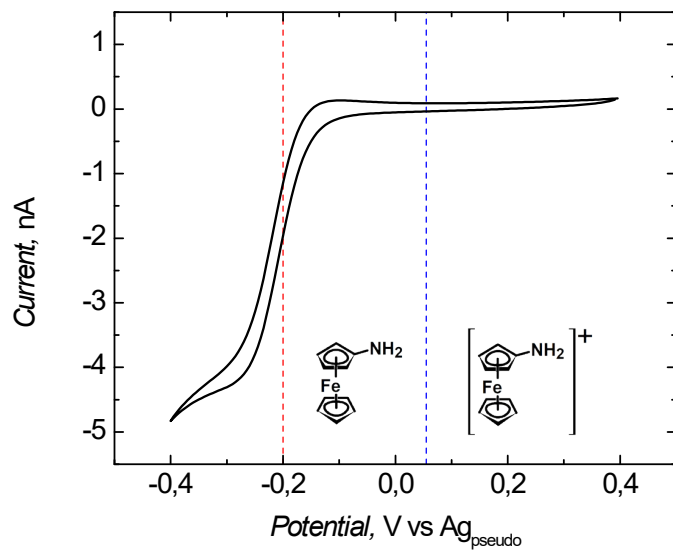


Figure S4. Voltammogram recorded on the SECM Pt probe tip in the bulk electrolyte, containing 1 mM Ru(NH₃)₆Cl₃ mediator in 0.1 M KCl. Redox potentials of Ru(NH₃)₆³⁺ (red dotted line) and aminoferrocene (blue dotted line) are marked. The potential scan rate was 50 mV/s.