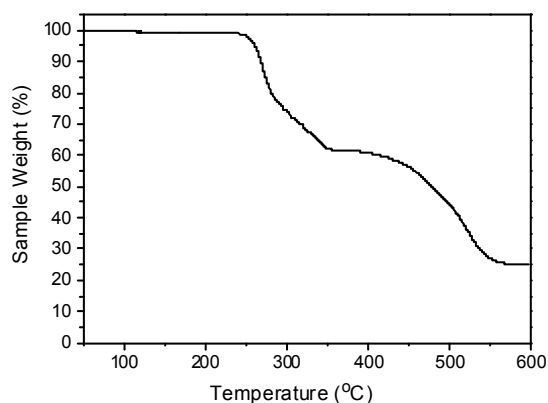


## Supporting Information for Manuscript ID JM-ART-01-2012-030599

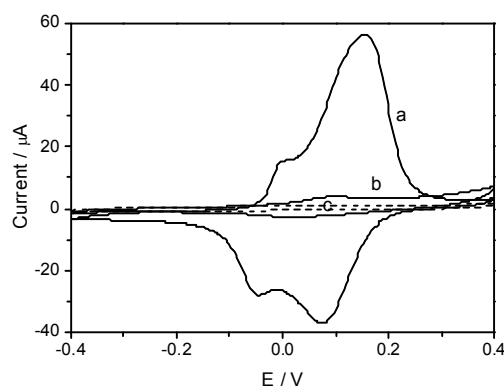
### Electrochemical sensing by surface-immobilized poly(ferrocenylsilane) grafts†

Xiaofeng Sui,<sup>+a</sup> Xueling Feng,<sup>+a</sup> Jing Song,<sup>b</sup> Mark A. Hempenius<sup>a</sup> and G. Julius Vancso<sup>\*\*a</sup>

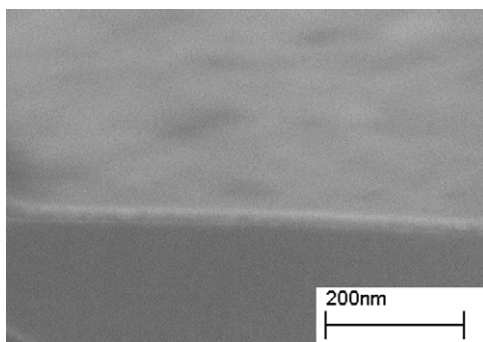
DOI: 10.1039/b000000x



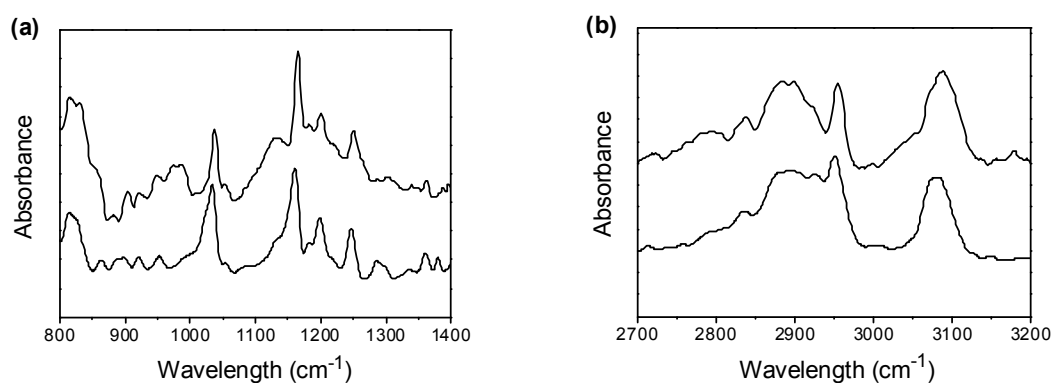
**Fig S1.** Thermogravimetric analysis of PFS 1.



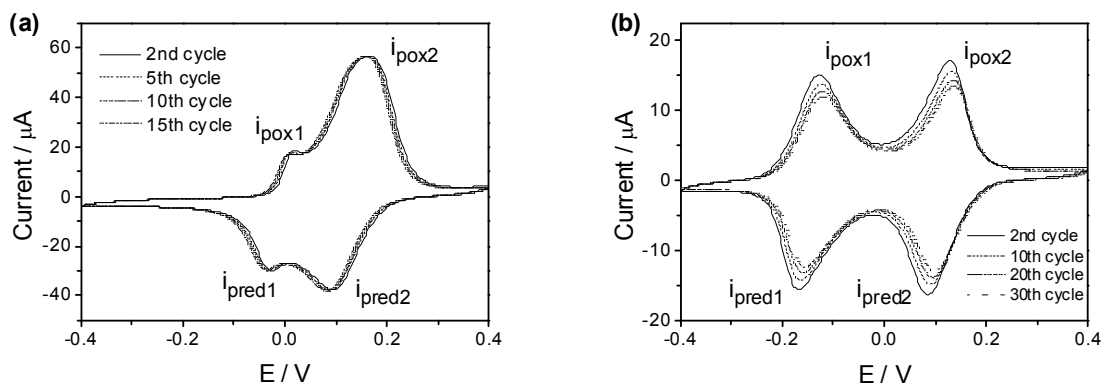
**Fig S2.** Cyclic voltammogram of a) PFS 1 covalently bound to an amine-terminated SAM on gold; b) PFMDS on the gold substrate with the cysteamine SAM; c) bare gold in 0.1 M NaClO<sub>4</sub> with Pt wires as the reference and counter electrode. All the substrates were soaked in THF overnight before measurements.



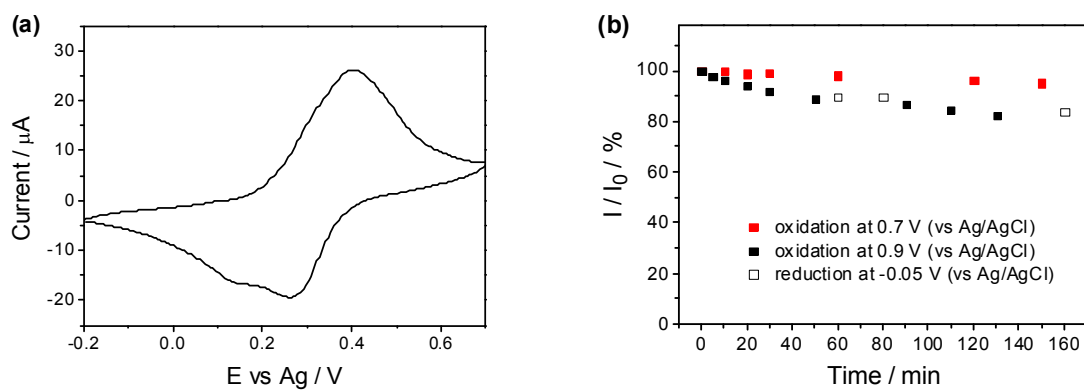
**Fig S3.** SEM image of the covalently anchored PFS film on silicon.



**Fig S4.** FTIR spectra of PFS **1** in bulk (bottom) and PFS brushes (top) on gold; (a) low energy region; (b) high energy region.



**Fig S5.** Cyclic voltammograms of PFS chains on gold at different scan rates, Pt reference and counter electrode. (a) In aqueous  $\text{NaClO}_4$  (0.1 M); (b) in  $\text{CH}_2\text{Cl}_2$  containing  $\text{NBu}_4\text{PF}_6$  (0.1 M).



**Fig S6.** (a) Cyclic voltammogram and (b) remaining percentage of the CV current intensity in the oxidation process relative to time = 0 as a function of accumulated holding time at the specified potential. Experiments were carried out in aqueous  $\text{NaClO}_4$  (0.1 M) using a Ag/AgCl reference electrode and a Pt counter electrode.