

DNA hybridization Detection by Charge Perturbation  
Through DNA at poly(thionine)-Modified Glassy Carbon  
And Gold Electrodes

Mohammad Mahbubur Rahman<sup>1</sup>, and Jae-Joon Lee<sup>1,2\*</sup>

<sup>1</sup> Department of Advanced Technology Fusion, Konkuk  
University, Seoul 143-701, Korea

<sup>2</sup> Nanotechnology Research Center & Department of  
Applied Chemistry, Konkuk University, Chungju 380-  
701, Korea

\* Author to whom correspondence should be addressed;  
Tel.: +82-43-840-3580; Fax: +82-43-851-4169  
([jjlee@kku.ac.kr](mailto:jjlee@kku.ac.kr))

Simple and label-free electrochemical sensors for DNA hybridization detection were developed based on poly(thionine) [PTH] modified glassy carbon and gold electrodes. Probe ssDNA was immobilized on the PTH film via covalent linkage between pendant amine (-NH<sub>2</sub>) group of the PTH and the phosphate (PO<sub>4</sub><sup>-</sup>) group of the ssDNA. The hybridizations were examined with different target ssDNA sequences. DPV showed a significant decrease of Fe<sup>2+</sup> oxidation peak current density ( $J_{peak}$ ) when hybridized with complementary and 1-base miss match ssDNA sequences. 3-base miss match and non-complementary ssDNA sequences showed the negligible changes of Fe<sup>2+</sup> oxidation  $J_{peak}$ . EIS demonstrated an increased charge transfer resistance ( $R_{ct}$ ) and decreased charge transfer rate constant ( $K_a^0$ ) after hybridization of complementary sequence. The PTH/GCE and PTH/GE sensors showed the excellent sensitivity of 1.44 and 50  $\mu$ A/cm<sup>2</sup>/nM with the detection limit of 0.14 and 0.36 nM, respectively for sensing complementary ssDNA hybridization.