Electronic Supplementary Material (ESI) for Analyst.

Supporting Information

A label-free and time-resolved luminescent strategy for the detection of protein based on DNA-Tb^{3+} luminescence quenched by graphene oxide

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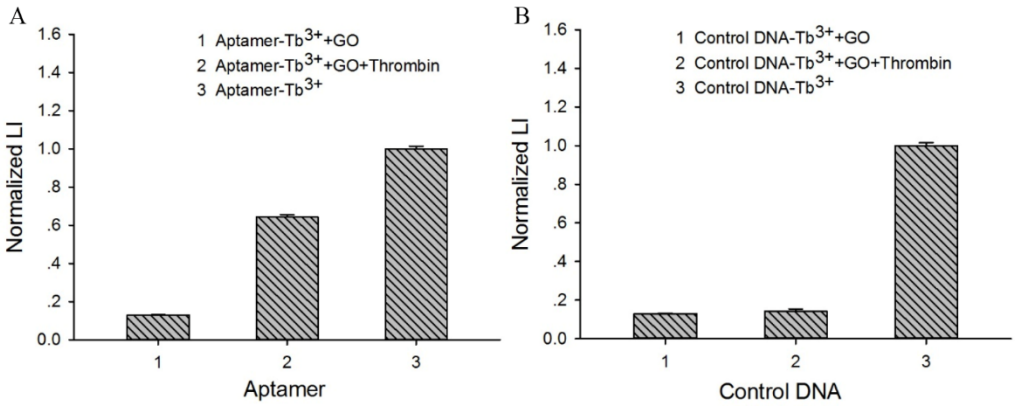

Figure S1. (A) Anti-thrombin aptamer binding with thrombin to desorb from the GO. (B) The control DNA interacting with thrombin.

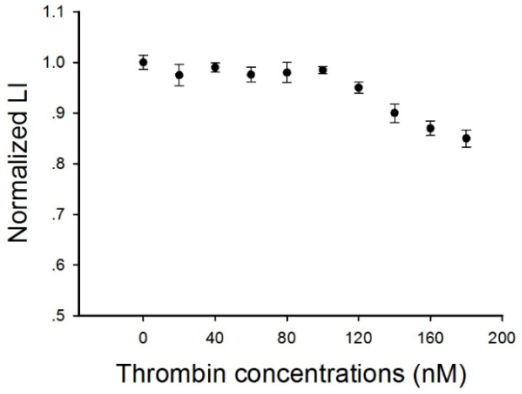

Figure S2. The influence of different concentrations thrombin to aptamer-Tb^{3+} complex.

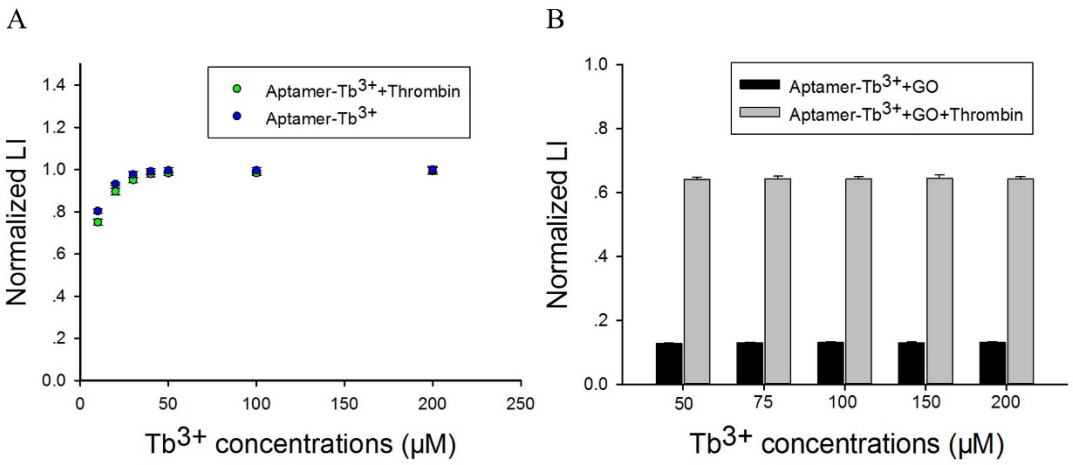

Figure S3. (A) The influence of thrombin to the luminescence of Aptamer-Tb^{3+} complex for different Tb^{3+} concentrations (green dots), luminescence of different concentration of Tb^{3+} binding

aptamer complex (blue dots). Thrombin, 100 nM. (B) The influence of Tb^{3+} concentration to the recovery experiments.

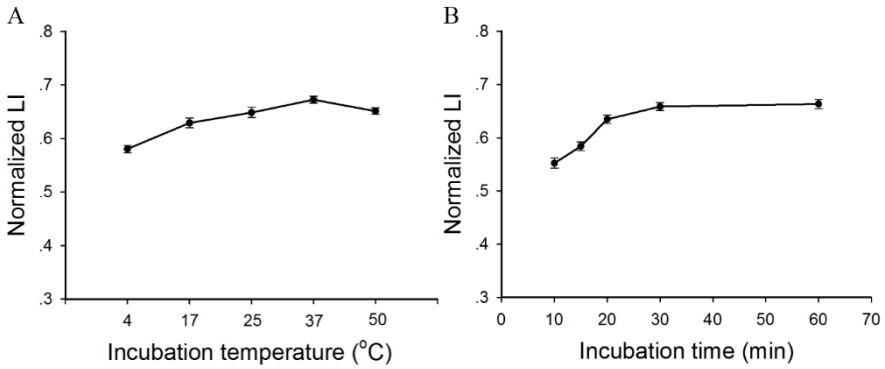

Figure S4. (A) The optimization of incubation temperature. (B) The optimization of incubation time.