Electronic Supplementary Information

Aptamer and Rolling Circle Amplification Involved Sandwich Assay

for Platelet-Derived Growth Factor-BB with Absorbance Analysis

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Fig. S1. The effect of template concentration on detection of PDGF-BB. Experimental condition: PDGF-BB, 50 pM; Aptamer-primer, 100 nM; Ligase, 0.02 U/ μ L; dNTPs, 0.1 mM; Polymerase, 0.4 U/ μ L; Biotinylated probe, 0.36 μ M; SA-HRP, 80-fold diluted.



Fig. S2. Effect of the diluted fold of SA-HRP on detection of PDGF-BB. Experimental condition: PDGF-BB, 50 pM; Aptamer-primer, 100 nM; Template, 56 nM; Ligase, 0.02 U/ μ L; Polymerase, 0.4 U/ μ L; dNTPs, 0.1 mM; Biotinylated probe, 0.36 μ M.



Fig. S3. Detection of PDGF-BB spiked in 1% bovine serum sample comparing with the result of detection of PDGF-BB in binding buffer solution.

Analytical method	Detection limit	Reference
Capillary electrophoresis with laser-induced fluorescence detection	0.5 nM	14
Fluorescence anisotropy assay	2 nM	16
Fluorescence resonance energy transfer assay	167 pM	32
Electrochemiluminescence assay	80 pM	33
Fluorescence analysis with structure-switching based RCA	0.4 nM	30
Chemiluminescence detection with RCA	10 fM	28
Electrochemical assay based on target binding-induced RCA	63 pM	27
Electrochemical detection with RCA	10 fM	26
Absorbance analysis with RCA	3.1 pM	this work

 Table S1 Comparison of sensitivity of aptamer-based assays for PDGF-BB