Supplementary Information

Combining a Loop-Stem Aptamer Sequence with Methylene Blue: A simple assay for Thrombin Detection by Resonance Light Scattering Technique

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Fig. S1 RLS spectra of the reaction system upon different concentration of H-eTBA (1 μ L, 1, 2, 4, 8, 10, 30 and 50 μ M). Conditions: MB: 50 μ M; pH value: 7.40. Error bars were the standard deviation of three repetitive measurements.



Fig. S2 RLS spectra of the reaction system upon different concentration of CTBA (1 μ L, 1, 2, 4, 8, 10, 30 and 50 μ M). Conditions: MB: 50 μ M; H-eTBA: 1 μ L, 10 μ M; pH value: 7.40. Error bars were the standard deviation of three repetitive measurements.



Fig. S3 RLS spectra of the reaction system upon the different incubation time. Conditions: MB: 50 μ M; H-eTBA: 1 μ L, 10 μ M; CTBA: 1 μ L, 10 μ M; thrombin: 4.91 nM; pH value: 7.40. Error bars were the standard deviation of three repetitive measurements.



Fig. S4 RLS intensity comparison of the protein interferences toward human thrombin. All interferences (BSA, Trypsin and α -Chymotrypsin) were tested at the concentration of 27.25 nM. The RLS signal of thrombin was tested at 17.18 nM. Other conditions: MB: 50 μ M; H-eTBA: 1 μ L, 10 μ M; CTBA: 1 μ L, 10 μ M; pH7.40. Error bars were the standard deviation of three repetitive measurements.