Supporting Information

Screening of DNA Aptamers against Myoglobin Using Positive and Negative Selection Units Integrated Microfluidic Chip and its Biosensing Application

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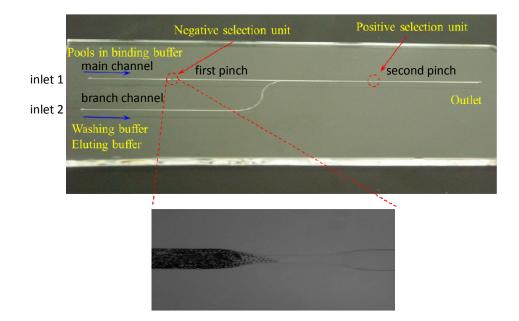


Figure S1. Structure of microfluidic device

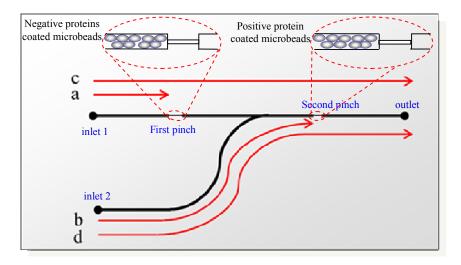


Figure S2. Sample injecting strategy. (a) Introduce the negative proteins-coated microbeads before the first pinch; (b) introduce the positive protein-coated microbeads before the second pinch; (c) inject the library in binding buffer; (d) inject washing buffer and eluting buffer in turn.

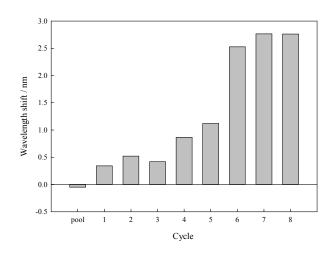
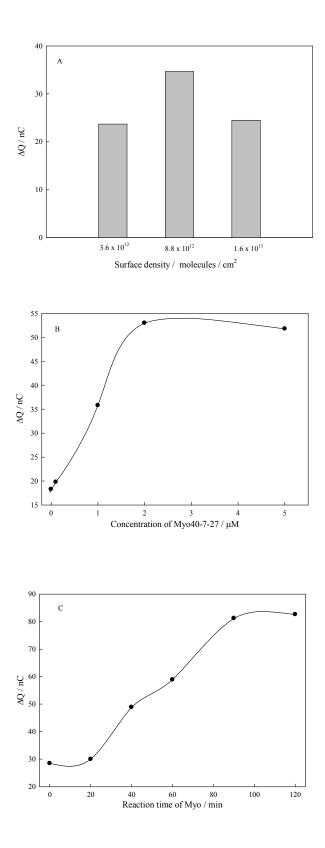


Figure S3. Efficiency of each round of selection. Original pool was used as control.



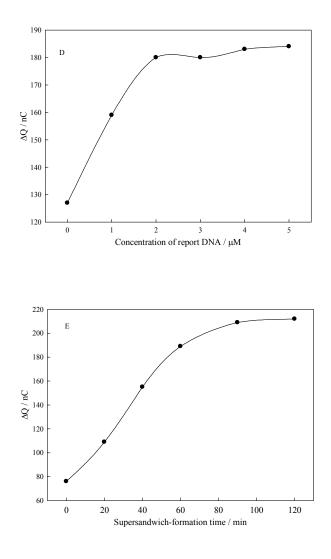


Figure S4. The effect of different factors on the detection of Myo.

(A) The effort of different surface density of capture DNA. Concentration of Myo: 10 nM; Concentration of Myo40-7-27: 5 μ M; Reaction time of Myo on the electrodes: 60 min.

(B) The effort of different Myo40-7-27 concentration. Surface density: 8.8×10^{12} molecule/cm²; Concentration of Myo: 10 nM; Reaction time of Myo on the electrodes: 60 min.

(C) The effort of different reaction time of Myo. Surface density: 8.8×10^{12}

molecule/cm²; Concentration of Myo: 10 nM; Concentration of Myo40-7-27: 2 µM.

(D) The effort of different report DNA concentration. Here, the concentration of report DNA1 equal to that of report DNA2. Surface density: 8.8×10^{12} molecule/cm²; Concentration of Myo: 10 nM; Concentration of Myo40-7-27: 2 μ M; Reaction time of Myo on the electrodes: 90 min; Supersandwich-formation time: 90 min.

(E) The effort of different supersandwich-formation time. Surface density: 8.8×10^{12} molecule/cm²; Concentration of Myo: 10 nM; Concentration of Myo40-7-27: 2 μ M; Concentration of report DNA: 2 μ M. Reaction time of Myo on the electrodes: 90 min.

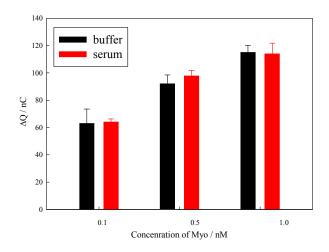


Figure S5. Detection of Myo in Human serum and buffer. The error bars represent the standard deviation of three measurements.