Supplementary information for

Surface plasmon resonance based competitive immunoassay

for Cd²⁺

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Supplementary Figures

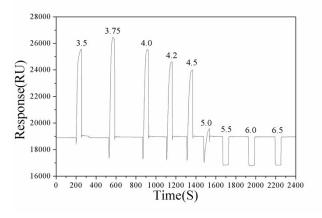


Fig. s1 The corresponding signals were produced by the antigens diluted with the acetate buffer solution of different pH values (pH 3.5, 3.75, 4.0, 4.2, 4.5, 5.0, 5.5, 6.0, 6.5 from left to right).

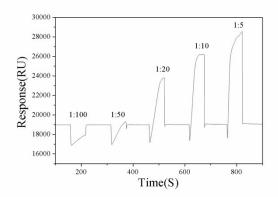


Fig. s2 The corresponding signals were produced by the antigens of different dilution rates (1:100, 1:50, 1:20, 1:10, 1:5 from left to right) with pH 3.75 acetate buffer solution.

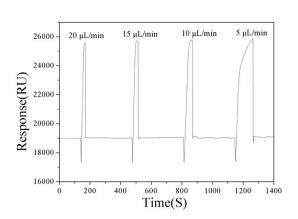


Fig. s3 The corresponding signals were produced by antigen at different flow rates (5 μ L/min, 10 μ L/min, 15 μ L/min, 20 μ L/min flow rates from left to right).

Analytical method	Operation	Linear range	Detection limit	Real sample	Ref.
		(µg L ⁻¹)	(µg L ⁻¹)		
SPR Biosensor	Simple	3.57~758.37	1.25	Water samples	This work
Carbon Paste	Complex	67.4~3372	9.0	Water samples and	1
Electrode				people hair sample	
Electrochemical Sensor	Complex	10~500	4.43	Water samples	2
ELISA	Complex	Not reported	1.95	Wheat sample	3
Fluorescence Biosensor	Simple	Not reported	56.2	Water samples	4
Naked-Eye Sensor	Simple	4.5~5001.8	4.4	Synthetic mixtures	5
				and water samples	
Nonthermal Optical	Complex	5~1000	1.5	Not reported	6
Emission Spectrometry					
Potentiometric Sensor	Simple	22.5~1.12×10 ⁶	11.2	Water samples	7
Fluorescent Sensor	Simple	112.4~1124	31.0	Cosmetic and	8
				personal care samples	

Table s1 Performance comparison of this work with other methods for Cd²⁺ detection

References

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