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Electronic Supplementary Material

Simultaneous fluoroimmunoassay of two tumor markers based on CdTe quantum dots and gold nanoclusters coated-silica nanospheres as labels

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1 Preparation of CdTe QDs

 $40~\mu L$ Thioglycolic acid (TGA) was added to 100~mL nitrogen-saturated ultrapure water containing 0.0917~g Cd(AC) $_2\cdot 2H_2O$. After adjusting to pH 10 with 0.01~M NaOH, 0.2209~g trisodium citrate and 0.0185~g Na $_2\text{TeO}_3$ were added. Then 0.01~g NaBH $_4$ was injected and bubbled by N $_2$ for 10 min. The finished CdTe precursor solution was transfered to a reaction kettle and kept at $120~^{\circ}\text{C}$ for 5 min to form yellow CdTe QDs. The CdTe QDs were precipitated in acetone and centrifuged at 10,000~rpm for 5 min, then re-dissolved in ultrapure water. They were extremely stable at $4~^{\circ}\text{C}$ for more than six months.

2 Preparation of Au NCs

Briefly, 5.0 mL of aqueous BSA solution (50.0 mg mL $^{-1}$, 37 °C) was added to an equal volume of 10.0 mM HAuCl $_4$ under vigorous stirring. After 2 min, 0.5 mL of NaOH solution (1.0 M) was introduced, and the mixture was incubated at 37 °C for 12 h. The color of the solution changed from light yellow to light brown, and then to deep brown. Finally, Au NCs solution was obtained and stored at 4 °C for use.

References

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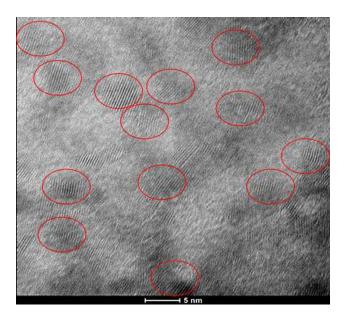


Fig.S1 HRTEM images of CdTe QDs.

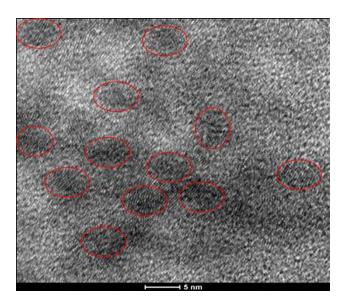


Fig.S2 HRTEM images of Au NCs.