## **Electronic Supplementary Information**

## Highly sensitive and selective detection of Pb<sup>2+</sup> ions using a novel and simple DNAzyme-based quartz crystal microbalance with dissipation biosensor

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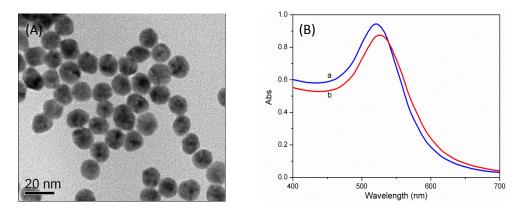


Fig. S1 (A) TEM image of AuNPs and (B) UV-Vis absorption spectra of AuNPs before (a) and after (b) immobilization with thiol-substrate.

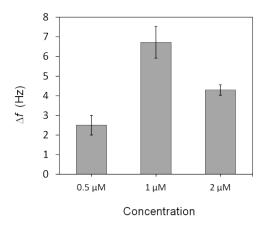


Fig. S2 The frequency response of 500 nM of Pb<sup>2+</sup> ions with different concentration of DNAzyme probe on the sensor surface. The surface density of the sensor was controlled by changing the concentration of DNAzyme from 0.5 to 2  $\mu$ M. With increasing amount of DNAzymes bound to the sensor surface, the frequency change increased as more substrate strands were hybridized with DNAzymes to form the Pb<sup>2+</sup>-active structure. The frequency change decreased after the concentration of DNAzyme increase to 2  $\mu$ M due to steric hindrance effect. Therefore, 1  $\mu$ M DNAzyme was chosen as the optimum condition for sensor fabrication according to the result.

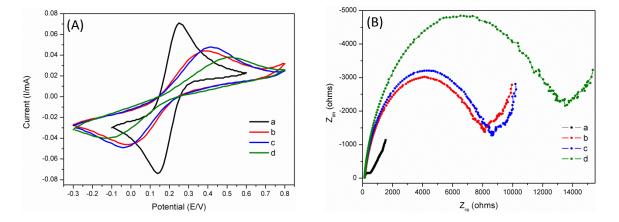


Fig. S3 Characterization of the QCM-D surface during sensor fabrication by (A) cyclic voltammetry (CV) and (B) electrochemical impedance spectroscopy (EIS). Conditions: working buffer 25 mM HEPES including 100 mM NaCl in the presence of 10 mM  $K_3$ Fe(CN)<sub>6</sub> and 10 mM  $K_4$ Fe(CN)<sub>6</sub>, CV: scan rate 50 mV/s, EIS: applying 5 mV alternative voltage in the frequency range of 100 mHz to 100 kHz. (a) bare gold; (b) after immobilization of GR5-E; (c) after backfilled with MCH; (d) after hybridization with substrate-functionalized AuNPs.