Electronic Supplementary Information for

Aptamer aided plasmonic nano-urchin for reporter free surfaceenhanced Raman spectroscopy analysis of cortisol

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1. Supplementary experimental and ethical approval

1.1. Characterization methods

Transmission electron microscopy (TEM) imaging was performed on a JEM 2100 (JOEL, Japan). Fourier transform infrared (FT-IR) spectra were generated on a Nicolet iS 10 Fourier spectrophotometer (USA). Zeta potential was measured by a Nano ZS90 Zeta analyzer. Ultraviolet–visible (UV–vis) absorption spectra were tested with a UV-2700 UV-Vis spectrophotometer (Shimadzu, Japan).

1.2. Ethical approval

Bio-fluids of serum and urine samples were donated by healthy volunteers in Zhongshan Third People's Hospital. All the experimental protocols involved in this work were approved by the committees on human subject research and ethics of the Zhongshan Third People's Hospital and Yunnan University. In accordance with the Helsinki Declaration, written informed consents were offered by all participants to ascertain their participation in the study and approve the utilization of their biological samples for analysis.

2. Supplementary Figures



Fig. S1. TEM images of the Au plasma.



Fig. S2 Zeta potentials of α -FeOOH, α -FeOOH@Au and α -FeOOH@Au-aptamer. The error bars were calculated as the relative standard deviation of three measurements.



Fig. S3 Raman spectra of cortisol solution (276.0 μ mol·L⁻¹) being directly dropped onto a silicon wafer for Ramman analysis and treating with the α -FeOOH@Au-aptamer based SERS protocol.



Fig. S4 SERS spectral intensities of cortisol analyzed by the same α -FeOOH@Auaptamer nanomaterials under different circles. (The peak at 1645 cm⁻¹ was taken as representative.)



Fig. S5 Raman spectra of cortisol standard solutions with various concentrations analyzed by α -FeOOH@Au-aptamer.



Fig. S6 Raman spectra of cortisol captured and analyzed by α -FeOOH@Au-aptamer from different cortisol spiked urine.