

Supplementary Information for

**Point of care biosensor for uric acid based on the target induce changing of  
photothermal effect of Gold nanostars using thermometer as readout**

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## **1. Experimental section**

### **1.1 Instruments and equipment.**

Transmission electron microscopy (TEM) images were acquired on a Tecnai G2 F20 S-TWIN microscope. UV-vis absorption spectra were acquired by Microplate spectrophotometer (Multiskan GO, Thermo Scientific, USA). The portable digital thermometer used for the tests in this work was CIE 305P. The commercial hand-held NIR laser pointer (808 nm, 1.5 W) was purchased from Changchun Laser Technology Co., Ltd. (Changchun, China).

### **2.2 Synthesis of AuNSs**

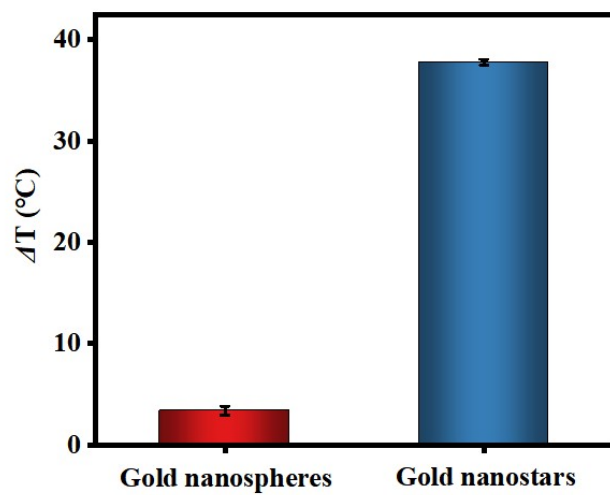
Gold seeds was firstly prepared by using sodium citrate reduction method. The mixture of  $\text{HAuCl}_4$  solution (1%, 1 mL) and ultrapure water (99 mL) was heated along with stirring and refluxing. Then, trisodium citrate solution (38.8 mM, 10 mL) was rapidly added to the above mixture, followed by further stirring and refluxing at 120°C for 20 minutes. When the color of solution gradually changed to wine red, the reaction solution was cooled down to room temperature. Then, the above gold seed solution (750  $\mu\text{L}$ ) was added to an aqueous solution (38.5 mL) containing hydroxylamine hydrochloride (40 mM, 750  $\mu\text{L}$ ) and HEPES (100 mM, 18.75 mL, pH = 8.5). The solution was mixed evenly by stirring at 1450 rpm. Subsequently,  $\text{HAuCl}_4$  solution (1 mM, 5 mL) was added dropwise, followed by continued stirring at 700 rpm for 15 minutes. The color of solution gradually turned to blue, whose UV-vis absorption wavelength was around 800 nm. After the reaction is completed, centrifuge three times at 8°C (4000 rpm, 10 minutes). Finally, the solution is

redispersed in HEPES and stored at 4°C for later use.

### **1.3 Serum sample preparation**

To verify the practical application potential of this photothermal biosensor, the UA levels of serum samples from patients at the First Affiliated Hospital of Fujian Medical University (Fujian, China) were tested. The serum samples were firstly centrifuged (4000 rpm/min, 10 min) to obtain supernatant, and then stored at -20°C before utilization. Serum samples were diluted 100-fold for the detection of the concentration of UA according to this photothermal biosensor testing procedures as mentioned above.

## 2. Supplementary figures



**Figure S1** Comparison of photothermal effects between spherical and star shaped gold nanoparticles.