Ag⁺ and Cysteine Detection Based on Graphene Oxide and G-Quadruplex by Ag⁺-Guanine interaction

Lin Liu^{*a*}, Tingting Hong^{*a*}, Wenting Liu^{*a*}, Xiaocheng Weng^{**a*}, Qianqian Zhai^{*a*}, Xiang Zhou^{**ab*}

^a College of Chemistry and Molecular Sciences, State Key Laboratory of Virology,
Wuhan University, Wuhan, Hubei, PR China.
Xiaocheng Weng, Tel: (+86)-27-68756663, Fax: (+86)-27-68756663, Email:
<u>xcweng@whu.edu.cn</u>
Xiang Zhou, Tel: (+86)-27-68756663, Fax: (+86)-27-68756663, Email:
<u>xzhou@whu.edu.cn</u>
^b State Key Laboratory of Natural and Biomimetic Drugs, Perking University, Hubei,

Wuhan 430072, PR China



Figure S1. (a) Fluorescence emission spectra of the fluorescent sensor in the presence of different concentrations of Ag⁺ (from 0 to 5000 nM) with the mixture of twelve metal ions together (Mg²⁺, Cu²⁺, Mn²⁺, Zn²⁺, Pb²⁺, Ni²⁺, Co²⁺, Cd²⁺, Ca²⁺, Li⁺, Hg²⁺, Fe³⁺, the concentration of each ion is 3 μ M). The concentration of AO is 500 nM while DNA is 300 nM. (b) Ag⁺ concentration-dependent change in the emission signal at $\lambda = 523$ nm. The insert figure shows the signal change in the Ag⁺ concentration range of 0-3000 nM. The red line represents a linear fit to the data.



Figure S2. (a) Fluorescence emission spectra of the Cys sensor system in the presence of different concentrations of Cys (from 0 to 5000 nM). The concentration of AO is 500 nM while DNA is 300 nM. And the concentration of Ag⁺ is 3 μ M. (b) Cys concentration-dependent change in the emission signal at $\lambda = 523$ nm.



Figure S3. The selectivity of the Cys sensor system. Red bars represent the emission signals of the sensing systems in the presence of 3 μ M of other amino acids without Cys. The green bars represent the emission signals of the sensing systems in the presence of 3 μ M Cys and 3 μ M of other amino acids together.