

Electronic supplementary information

A graphene oxide-based fluorescent aptasensor for the turn-on detection of epithelial tumor marker mucin 1

Yue He, Yi Lin, Hongwu Tang* and Daiwen Pang

* To whom correspondence should be addressed. Prof. Hongwu Tang, E-mail:
hwtang@whu.edu.cn (H.-W. T.), Tel: 86-27-68756759, Fax: 86-27-68754685.

*Corresponding author:

*Key Laboratory of Analytical Chemistry for Biology and Medicine (Ministry of Education),
College of Chemistry and Molecular Sciences, Research Center for Nanobiology and
Nanomedicine (MOE 985 Innovative Platform), and State Key Laboratory of Virology, Wuhan
University, Wuhan, 430072, P. R. China.*

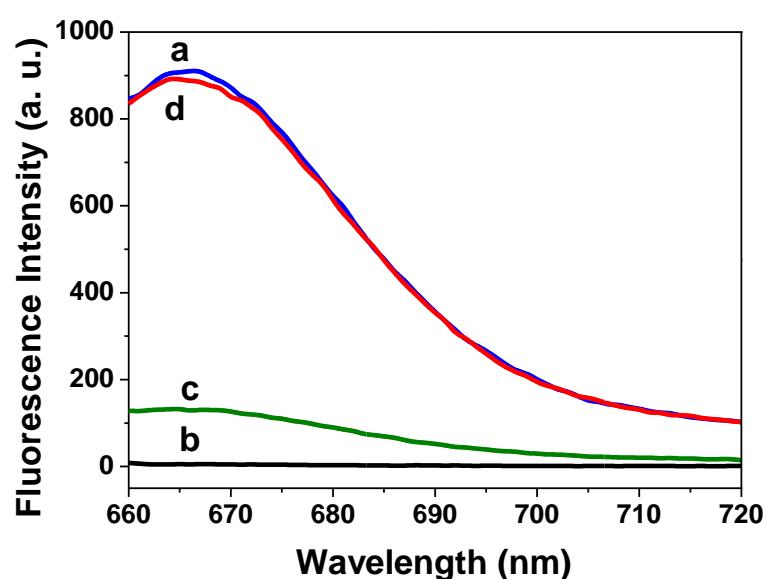


Figure S1

Figure S1. Fluorescence emission spectra of Cy5-labeled MUC1 aptamer (20 nM) at different conditions: (a) Cy5-labeled MUC1 aptamer in Tris–HCl buffer; (b) Cy5-labeled MUC1 aptamer + GO; (c) Cy5-labeled MUC1 aptamer + GO + 5 μ M MUC1; (d) Cy5-labeled MUC1 aptamer + 5 μ M MUC1. Excitation: 645 nm.

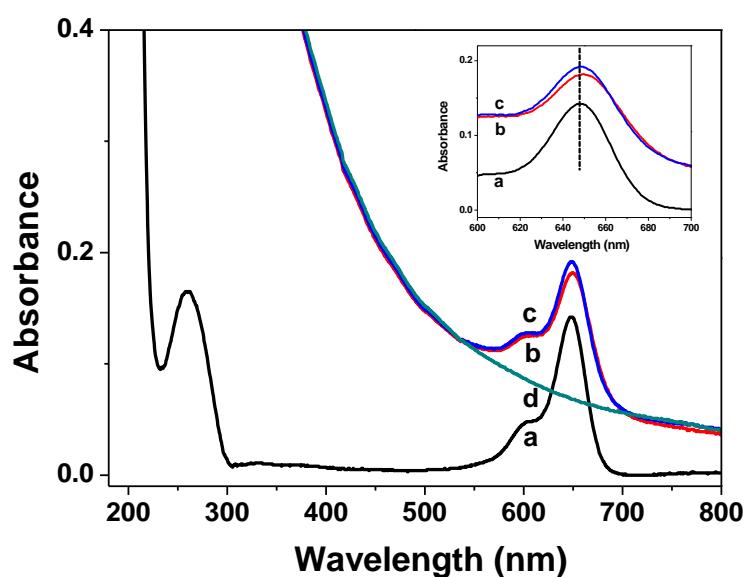


Figure S2

Figure S2. UV-visible absorbance spectra of free (a) P_0 ; (b) $P_0 + \text{GO}$; (c) $P_0 + \text{MUC1} + \text{GO}$; (d) GO in Tris–HCl.

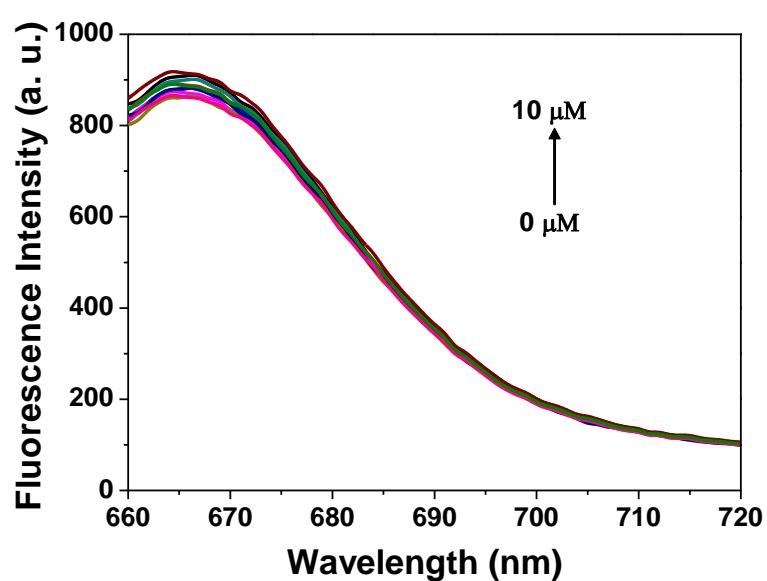


Figure S3. Fluorescence emission spectra of P_0 (20 nM) upon the addition of MUC1 at different concentrations. Excitation: 645 nm.

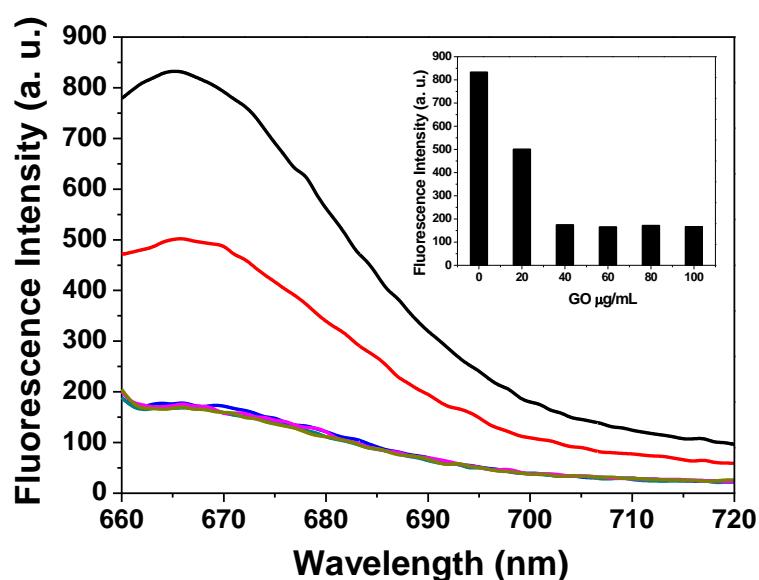


Figure S4. Fluorescence emission spectra of P_0 (20 nM) in 2% serum solution upon the addition of different concentrations of GO. Excitation: 645 nm. Inset: fluorescence intensity versus concentration of GO. Excitation: 645 nm.

Table S1. Fluorescence anisotropy changes of P₀ (20 nM) at different conditions: (a) P₀ in Tris–HCl bufferr; (b) P₀ + 10 μM MUC1; (c) P₀ + GO; (d) P₀ + GO + 5 μM MUC1; (e) P₀ + GO + 100 μM MUC1. Excitation: 645 nm.

		r ^a	RSD (%)
(a)	P ₀	0.4261	2.5
(b)	P ₀ +MUC1 (10 μM)	0.4415	2.9
(c)	P ₀ +GO	0.6647	2.4
(d)	P ₀ +GO+MUC1 (5 μM)	0.4854	3.0
(e)	P ₀ +GO+ MUC1 (10 μM)	0.4700	2.7

^a Average of three measurements.