

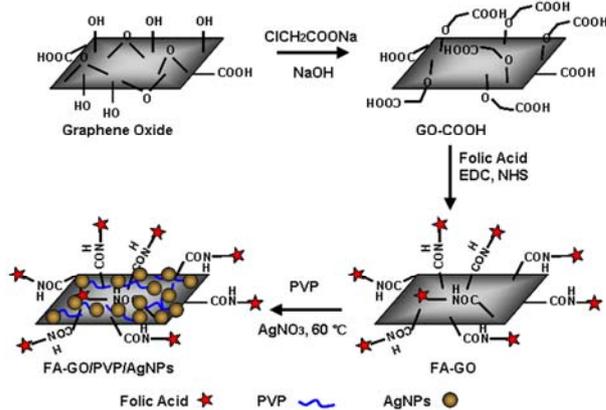
Supporting Information

Graphene oxide based surface-enhanced Raman scattering probes for cancer cell imaging

Zhiming Liu, Zhouyi Guo,* Huiqing Zhong, Xiaochu Qin, Mingming Wan and Biwen

Yang

MOE Key Laboratory of Laser Life Science & SATCM Third Grade Laboratory of Chinese
Medicine and Photonics technology, College of Biophotonics, South China Normal University,
Guangzhou 510631, China.



Scheme S1 Illustration of the fabrication of FA-GO/AgNPs.

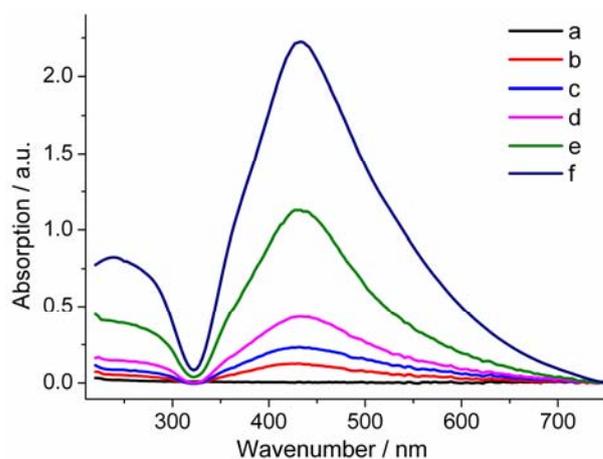


Fig. S1 UV-Vis absorption spectra of GO/AgNPs with identical graphitic carbon (5 µg/mL) prepared with a various weight ratio between the AgNO₃ and GO: 0 (a), 48 (b), 96 (c), 192 (d), 384 (e), 768 (f).

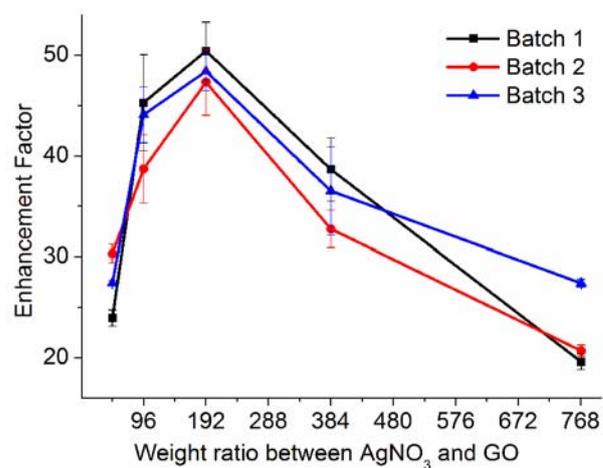


Fig. S2 Batch-to-batch reproducibility of the as-prepared GO/AgNPs samples. The uniform trends of EFs in response to the AgNO₃/GO ratio indicate a good reproducibility of the synthesis process. Errors bars are based on standard deviations (SD) of three parallel measurements of each sample.

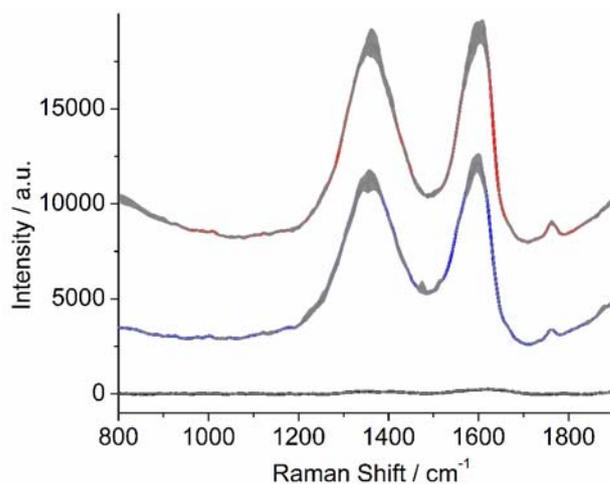


Fig. S3 The stability of the GO/AgNP hybrids. Black line represents the Raman spectrum of GO (25 µg/mL). Red line represents the SERS spectrum of as-prepared GO/AgNPs (192) with 25 µg/mL graphitic carbon. Blue line represents the SERS spectrum of GO/AgNPs (192) with 25 µg/mL graphitic carbon stored for two months. The SERS EF of GO/AgNPs (192) is measured as ~43.3 after two months storage, which is similar to the value of as-prepared hybrids (~48.4). The grey regions show the SD of three parallel measurements of each sample.

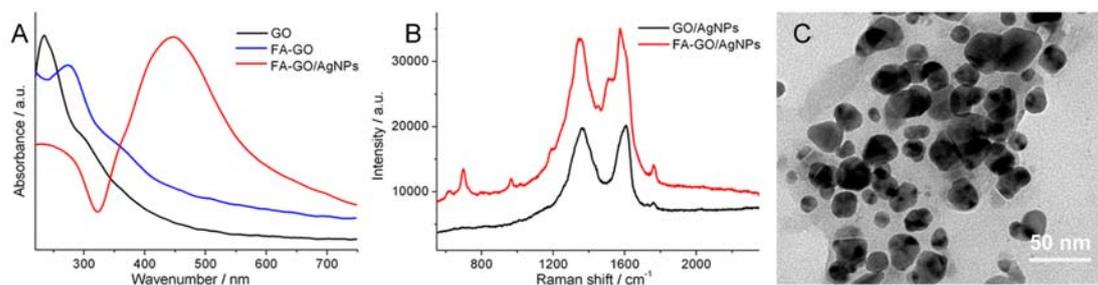


Fig. S4 Characterization of FA-GO/AgNPs. (A) UV-Vis absorption spectra. (B) SERS spectra. (C) TEM image of FA-GO/AgNPs.

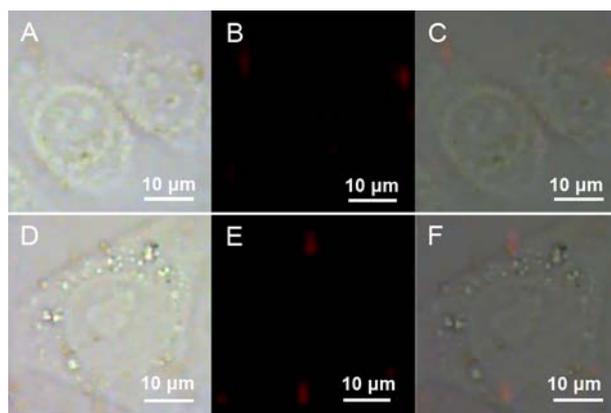


Fig. S5 Typical bright field images (A and D), SERS images (B and E) and the overlap images (C and F) of A549 cells (A-C) and HeLa cells (D-F) incubated with 1 µg/mL GO/AgNPs for 2h.