Gas identification with graphene plasmons

Hai Hu^{1,#}, Xiaoxia Yang^{1,#}, Xiangdong Guo¹, Kaveh Khaliji², Sudipta Romen Biswas², F Javier García de Abajo³, Tony Low^{2*}, Zhipei Sun^{4*}, Qing Dai^{1*}.

1 National Center for Nanoscience and Technology, Beijing 100190, P. R. China.

2 University of Minnesota, Minneapolis, MN 55455, USA.

3 The Barcelona Institute of Science and Technology, 08860 Castelldefels (Barcelona), Spain.

4Aalto University Tietotie 3, FI-02150 Espoo, Finland.

daiq@nanoctr.cn;zhipei.sun@aalto.fi;tlow@umn.edu

In the present study, we identified gas molecules using araphene plasmons. The rotational-vibrational modes of gas molecules of NO₂, N₂O, NO, and SO₂, which are generally important in environmental and military monitoring applications, as well diagnostics medical as in are unambiguously detected and identified using the designed graphene nanostructures. We attribute this to the great adsorptive capacity to redistribute the gas graphene molecules to the surface (equivalent to amplifying the gas concentration), facilitating hence the interaction between ultra-confined graphene plasmons and gas molecules. Our analysis reveals theoretical that the adsorbed gas layer (about 800 zeptomole molecule per μm^2 for < 1 nm thickness) on the graphene structure, in conjunction with the strong field confinement, is critical to effectively detecting and identifying gas molecules. In addition, our araphene plasmonic sensors also successfully performed real-time monitoring of gas molecules during chemical reactions with a fast response time (< 1 min).

References

[1] Hu, H., et al. Nat. Commun. 10 (1), 1131 (2019).

[2] Hu, H., et al. Nat. Commun. 7, 12334 (2016).

[3] Rodrigo, D., et al. Science 349, 165-168 (2015).

[4] Liu, N. , et al. Nat. Mater. 10, 631-636 (2011).



Figure 1: Experimental scheme of our device. A metal chamber with a piezometer was used for precise control of gas parameters. Plasmons in a graphene ribbon array were excited using an incident infrared beam and tuned in situ by electrostatic doping through a gate voltage (Vg). The plasmon resonances were coupled with molecular excitations, thus probing the rotational-vibrational spectral fingerprints of gas molecules.



Figure 2: (a-c) Extinction spectra of graphene in the presence of N₂O, NO₂, and NO, respectively. The rotational-vibrational modes are marked with vertical lines. **(d)** Extinction spectra of graphene in the presence of two gas mixtures, one consisting of SO₂ and N₂O, and the other of SO₂, N₂O and NO₂. The graphene ribbon widths in a-c were 80, 60, and 40 nm, respectively, with a filling factor of 90%, δ VCNP of 30 V, and concentration of N₂O of 8000 ppm, and NO₂ and NO of 4000 ppm at 1 atm.