RAMAN OPTICAL TRAPPING OF GRAPHENE

M. G. Donato^a, <u>G. Faggio^{b*}</u>, F. Bonaccorso^c
D. Alfieri^b, S. Santangelo^b, C. D'Andrea^a
B. Fazio^a, P. G. Gucciardi^a, A. C. Ferrari^c
O. M. Maragò^a and G. Messina^b

^aCNR-IPCF, Istituto per i Processi Chimico-Fisici, Messina, Italy ^bDipartimento di Ingegneria Elettronica, Università "Mediterranea", Reggio Calabria, Italy ^cDepartment of Engineering, University of Cambridge, Cambridge, UK *Corresponding author: *faggio@unirc.it*

Dimensionality plays a special role in nature. From phase transitions to transport phenomena, two-dimensional (2d) systems often exhibit a strikingly different behavior from those with higher or lower dimensionality. Graphene [1] is the prototype 2d material and, as such, has unique mechanical, thermal, electronic, and optical properties, already proven outstanding for both fundamental research and applications [2,3]. Here we demonstrate optical trapping of individual graphene flakes in aqueous dispersion [4]. This enables the investigation of their Raman spectra in the optical trap. The modification of the Raman spectra due to colloidal metallic nanoparticles [7] of different shapes [8] added to the dispersion are investigated both by micro-Raman and Raman tweezers analysis.

Graphite is exfoliated by ultrasonication in a water-surfactant solution, followed by ultracentrifugation. (Fig. 1). We do not use any functionalization nor oxidation, in order to retain the electronic structure of pristine graphene in the exfoliated layers [9]. We use dihydroxy sodium deoxycholate (SDC) as surfactant. We then place 75 microlitres of dispersion in a chamber attached to a piezo-stage with 1nm resolution. Optical trapping (Fig. 2a) is obtained by focusing a helium-neon (633 nm) laser through a 100X oil immersion objective (NA=1.3). The latter is coupled to a spectrometer through an edge filter. This allows us to use the same laser light both for optical trapping and for Raman scattering, realizing a Raman optical tweezers to directly probe the structure of the trapped flake. A typical Raman spectrum of trapped flakes measured at 633 nm is plotted in Fig. 2b. Besides the G and 2D peaks, this has significant D and D' intensities, and the combination mode D+D'~2950 cm⁻¹. The large intensity of the D peak in Fig. 2b is assigned to the edges of our submicrometer flakes [10]. We note that the 2D band, although broader than in pristine graphene [5], is well fitted by a single Lorentzian lineshape. Thus, even if the flakes are multilayers, they are electronically almost decoupled.



Figure 1: Graphite exfoliation. (a) Molecular structure of SDC. (b) Schematic illustration of the graphite exfoliation process. (c) A mild ultrasonication produces exfoliated mono- and few-layer graphene encapsulated by SDC. (d) Photograph of the as-prepared dispersion.



Figure 2: (a) Experimental setup. The same optics is used to trap graphene in dispersion and collect the Raman signal that is sent to a spectrometer. (b) Raman spectrum of an optically trapped flake for 633 nm trapping and excitation wavelength. (c) Micro-Raman spectra of graphene dispersions with and without gold nanoparticles for 514.5nm excitation.

References

- [1] K. S. Novoselov, et al., Science, 306 (2004) 666669.
- [2] A. K. Geim, K. S. Novoselov, Nat. Mater., 6 (2007) 183191.
- [3] F. Bonaccorso, et al., Nat. Photonics, 4 (2010) 611622.
- [4] O. M. Maragò, et al., ACS Nano, 4, (2010) 7515-7523.
- [5] A. C. Ferrari, et al., Phys. Rev. Lett., 97 (2006) 187401.
- [6] A. C. Ferrari, et al., Phys. Rev. B, 61 (2000) 1409514107.
- [7] F. Schedin, et al., ACS Nano, 4 (2010) 56175626.
- [8] P.H. Jones, et al., ACS Nano, 3 (2009) 30773084.
- [9] Y. Hernandez, et al., Nat. Nanotechnol., 3 (2008) 563568.
- [10] C. Casiraghi, et al., Nano Lett., 9 (2009) 14331441.