

Electrochemical characterization of graphene gated field effect transistors: route for smart biological sensors

Juliette Simon, Adrien Hugo, Pascal Mailley, Fabienne Blanc, Thomas Alava

▶ To cite this version:

Juliette Simon, Adrien Hugo, Pascal Mailley, Fabienne Blanc, Thomas Alava. Electrochemical characterization of graphene gated field effect transistors: route for smart biological sensors. Graphene2020, Jun 2020, Grenoble (E-meeting), France. hal-04815256

HAL Id: hal-04815256 https://hal.inrae.fr/hal-04815256v1

Submitted on 2 Dec 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Electrochemical characterization of graphene gated field effect transistors: route for smart biological sensors

Juliette SIMON

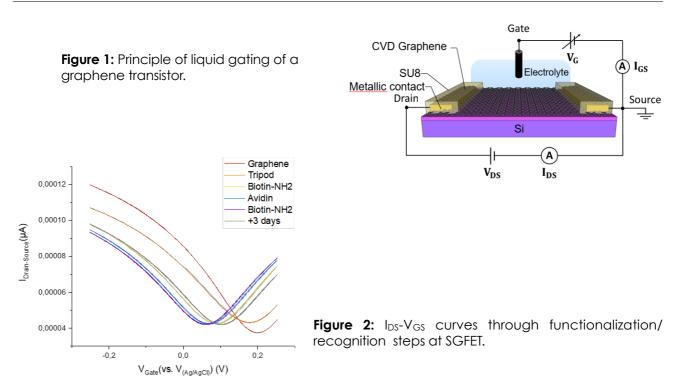
Adrien HUGO, Pascal MAILLEY, Fabienne BLANC, Thomas ALAVA CEA, 17 avenue des martyrs, Grenoble, France Juliette.simon@cea.fr

Highly conductive, lightweight, flexible, transparent, mechanically robust, CVD graphene has sparkled large interest in the biosensors community. Indeed graphene ultimate surface-to-volume ratio, associated with exceptional electron mobility is particularly appealing for the detection of charged biological species. Previous work in our group demonstrated a unique fabrication protocol for Graphene Solution Gated Field Effect Transistor (SGFET) on silicium substrates (Figure 1). This study first introduces an extended electrochemical characterization of these devices exploring their limits in robustness. Next, results from our first biological detection campaign are presented. In Phosphate Buffer Saline (PBS) 0.01X, V_{GS} was swept from -0.25V to 0.25V with a scan rate of 0,01V/s to characterize the l_{DS} vs. V_{GS} evolution. Graphene was functionalized with either positively or negatively charged proteins. A unique tripodal molecular compound was immobilized on the graphene surface to support a biotin probe [1]. Consecutively, avidin-class proteins were specifically captured on the sensor [2]. Post Data processing gave us access to leakage current evolution, sensitivity of the sensor and V_{Dirac} evolution. Figure 2 shows the obtained responses of the graphene transistors I_{DS} vs. V_{GS} after injection of the different biological molecules involved in our experiment. The direction and magnitude of the shift of the Dirac peak from one functionalization/recognition step to another helps at understanding the inner mechanisms involved in the modification of the electrical double layer at the interface between graphene and the liquid media and their influence on graphene conductivity.

References

- [1] J. A. Mann et al., Angew. Chem. Int. Ed. 11, 2013, "Preservation of antibody selectivity on graphene by conjugation to a tripod monolayer".
- [2] T. Alava et al., Anal. Chem., 2013, "Control of the Graphene–Protein Interface Is Required To Preserve Adsorbed Protein Function".





Graphene2020