Self-assembled monolayer of push-pull chromophore on gold: towards the modulated polarization for the controlled detection of biomolecules

Junlong Wang^a, Virginie Gadenne^a, Jean-Manuel Raimundo^b, Lionel Patrone^a

^aAix Marseille Univ., Université de Toulon, CNRS, IM2NP UMR 7334, ISEN Yncréa Méditerranée,

Maison du Numérique et de l'Innovation, Place G. Pompidou, 83000 Toulon, France,

lionel.patrone@im2np.fr

^bCINaM UMR CNRS 7325, Aix-Marseille Université, 13288 Marseille cedex 09, France

Corresponding author email: junlong.wang@etu.univ-amu.fr

The specific detection of antibodies using antibody/antigen binding interactions has been the subject of numerous studies in the field of biosensors [1-3]. Among the different approaches, a promising functional ON/OFF system has been proposed based on the reversible modification of the conformation of a charged oligopeptide by the application of an electrical potential on the surface, thus making it possible to control the antibody-antigen interaction [4]. So far, this system can only respond to a single electrical stimulus, thus limiting its use to a single antibody/antigen pair. To overcome this scientific obstacle, we propose to develop an alternative system allowing the detection of the binding interaction of several antibody/antigen pairs within a single platform.

For this purpose, we are considering an original approach based on the use of "push-pull" chromophores to control the potential allowing the conformation of the oligopeptide to be switched between the OFF state (antibody-antigen interaction not allowed) and the ON state (interaction permitted). This structure of the push-pull chromophores forms an electrical dipole depending on the nature of the donor and attractor groups, and the π -conjugated bridge. The idea is to add push-pull chromophores presenting different electrical dipole moments - each one being associated to a specific electrical potential value - between the surface and the oligopeptide supporting a given antigen, and to insert these systems within an inert matrix of molecular compound of oligoethylene glycol (OEG).

References:

- [1] Casalini, S.; Dumititru, A.C.; Leonardi, F.; et al., ACS Nano 2015, 9, 5051.
- [2] Marquette, C. A.; Blum, L.J. *Biosens. Bioelectron.* **2006**, *21*, 1424.
- [3] Wen, W.; Yan, X.; Zhu, C. et al., Anal. Chem. 2017, 89, 138.
- [4] Santos Gomes, B.; Cantini, E.; Tommasone, S.; et al., ACS Appl.Bio.Mater., 2018, 1, 738.