



# Abstract Rational Design of a Planar Junctionless Field-Effect Transistor for Sensing Biomolecular Interactions <sup>+</sup>

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- <sup>+</sup> Presented at the XXXV EUROSENSORS Conference, Lecce, Italy, 10–13 September 2023.

**Abstract:** In the ElectroMed project, we are interested in screening certain peptide sequences for their ability to selectively interact with antibodies or MHC proteins. This poses a combinatorial challenge that requires a highly multiplexed setup of label-free immunosensors. Label-free FET-based immunosensors are good candidates due to their high multiplexing capability and fast response time. Nanowire-based FET sensors have shown high sensitivity but are unreliable for clinical applications due to drift and gate stability issues. To address this, a label-free immuno-FET architecture based on planar junctionless FET devices is proposed. This geometry can improve the signal-to-noise ratio due to its larger planar structure, which is less prone to defects that cause noise and is better suited to the functionalization of different receptor molecules.

Keywords: immuno-FET; MHC proteins; label-free; planar junctionless

## 1. Introduction

Peptides are short protein sequences with diagnostic and therapeutic potential. However, the variability of the proteome between cells poses a significant challenge for screening and transducing interactions into readable signals. Label-free field effect transistor (FET)based immunosensors show promise for solving this challenge due to their high sensitivity, multiplexing capabilities, and fast response time. The nanowire-based FETs have garnered significant attention due to their 3D gating effect and faster mass transport towards the sensing area. However, the low reliability and reproducibility of the nanowire-based FET sensors has prevented these structures from reaching clinical application. In this study, a label-free immuno-FET architecture based on 2D planar junctionless FET devices is proposed. This geometry improves the signal-to-noise ratio due to the lower number of surface defects because of simpler fabrication processes compared to non-planar geometries, and it can be better suited to functionalizing different receptor molecules because of its planar structure. This study aimed to fabricate and characterize planar junctionless FETs



Citation: Shukla, R.P.; Bomer, J.G.; Wijnperle, D.; Kumar, N.; El Maiss, J.; Balakrishanan, D.; Singh, A.C.; Georgiev, V.P.; Garcia, C.P.; Krishnamoorthy, S.; et al. Rational Design of a Planar Junctionless Field-Effect Transistor for Sensing Biomolecular Interactions. *Proceedings* 2024, *97*, 121. https://doi.org/ 10.3390/proceedings2024097121

Academic Editors: Pietro Siciliano and Luca Francioso

Published: 29 March 2024



**Copyright:** © 2024 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). for pH sensing with the potential for future use in multiplexed sensing of biomolecular interactions.

### 2. Materials and Methods

Planar junctionless FETs were fabricated in a standard clean-room environment using SOI wafers from IceMOS Technology, Ltd. (Belfast, UK) To achieve a higher sensitivity suitable for detecting biomolecular interactions, a thin and lightly doped silicon device layer (thickness of 250–300 nm and resistivity of 1–10 Ohm.cm) was used [1].

#### 3. Discussion

In this study, pH sensing was performed using planar junctionless field-effect transistor sensors (FETs) that were fabricated and characterized (Figure 1a). The FET device with SiO<sub>2</sub> gate oxide displayed a voltage sensitivity of around 40 mV/pH for constant drain currents of 50 nA at a drain-to-source voltage of 0.05 V. Theoretical modeling and simulation yielded a mean value of  $3.8 \times 10^{15}$ /cm<sup>2</sup> for the surface states, with a standard deviation of  $3.6 \times 10^{15}$ /cm<sup>2</sup>. With our proposed rational design, we anticipate achieving a sensitivity that is sufficient for detecting peptide–protein interactions (Figure 1b). The performance of these planar devices can be further improved by using a thin layer of high-k gate dielectrics (e.g., HfO<sub>2</sub>) and by modifying the gate area with nanomaterials that will later be used as a multiplexed set-up of immunosensors to detect biomolecular interactions.



**Figure 1.** Microfabrication and surface functionalization: (**a**) microfabricated device with open gate area; (**b**) surface functionalization of Au/SiO2 SPR chips with APTES, peptide, and protein.

Author Contributions: Conceptualization, S.P., V.P.G., S.K., and C.P.G.; methodology, R.P.S., J.G.B., D.W., C.P.G., S.P., N.K., V.P.G., J.E.M., D.B., A.C.S., and S.K.; software, R.P.S. and N.K.; validation, R.P.S., N.K., J.E.M., D.B., A.C.S., and S.K.; formal analysis, R.P.S., S.P., C.P.G., N.K., V.P.G., J.E.M., D.B., A.C.S., and S.K.; data curation, R.P.S., N.K., and A.C.S.; writing—original draft preparation, R.P.S.; writing—review and editing, R.P.S., C.P.G., S.P., N.K., V.P.G., J.E.M., D.B., A.C.S., and S.K.; project administration, C.P.G.; funding acquisition, C.P.G.; visualization, R.P.S., S.P., N.K., J.E.M., D.B., and A.C.S. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research was funded by the European Union's Horizon 2020 research and innovation program, grant number 862539-Electromed-FET OPEN.

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

**Data Availability Statement:** Data are available on demand and in consultation with corresponding author.

Conflicts of Interest: The authors declare no conflicts of interest.

## Reference

 Shukla, R.P.; Bomer, J.G.; Wijnperle, D.; Kumar, N.; Georgiev, V.P.; Singh, A.C.; Krishnamoorthy, S.; Pascual García, C.; Pud, S.; Olthuis, W. Planar Junctionless Field-Effect Transistor for Detecting Biomolecular Interactions. *Sensors* 2022, 22, 5783. [CrossRef] [PubMed]

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