

DOI: 10.1002/((please add manuscript number))

Article type: Communication

High Performance Vertical Organic Electrochemical Transistors

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Keywords: electrochemical transistor, organic bioelectronics, transconductance, device density, vertical

Since the initial demonstration of a working transistor in 1947, reduction in physical size (downscaling or miniaturization) of various inorganic transistors has led to better utilization of available substrate space while improving transistor performance and lowering power

requirements and device cost.^[1] Downscaling of transistors has enabled the advancement of a wide range of commonly used electronic devices.^[2] Compared to traditional inorganic transistors, the development of transistors based on organic materials is at a relatively young phase. Downscaling of organic field-effect transistors (OFETs) has been primarily driven by active-matrix display technologies,^[3] with much effort focused on understanding and optimizing organic semiconductor (OSC) morphology and charge transport properties.^[4] An alternative organic transistor structure to the OFET, the organic electrochemical transistor (OECT), has also benefitted from OSC materials research, progressing significantly in the last decade. Whereas OFET current modulation relies on the field-effect, responding to charge accumulation at the dielectric interface of the channel, the current in OECTs is controlled by the injection of ions from an electrolyte into the bulk of the channel. This makes them particularly suitable for biointerfacing applications where documented advantages of organic technologies are numerous.^{[5],[6]} Examples include neural implants,^{[7]-[14]} *in vitro* arrays for biological assays,^{[15]-[21]} and neuromorphic devices.^{[22]-[25]} OECTs have reached a significant level of maturity and a major contemporary challenge is to pursue miniaturization for increasing density and improving performance.

Various vertical transistor orientations have been previously investigated in OFETs.^{[26]-[30]} However, the majority of these devices suffer from low mobilities, undesired leakage currents, and rather high driving voltages. In contrast, OECTs are inherently suitable for miniaturization. This results from the different principle of OECT operation, good-quality contacts to the channel due to the high levels of doping, and simplicity of the fabrication process. A first attempt to make use of a vertical architecture in combination with poly(3,4-ethylenedioxythiophene) doped with poly(styrenesulfonate) (PEDOT:PSS) deals with screen printed devices. Vertical vias in plastic or paper are exploited to make use of both sides of the substrate.^[31] In this communication we demonstrate vertical organic electrochemical transistors (vOECTs) with PEDOT:PSS channels, fabricated using photolithography. The

fabrication process takes advantage of the unique ionic-to-electronic transducing capability of the OECT, while maintaining low-voltage operation, compatibility with flexible substrates, and amplification with high signal-to-noise ratio. For biointerfacing applications this provides a means to considerably increase the number of possible recording sites (transistor density) while significantly improving transistor performance. Indeed, the geometry-normalized intrinsic transconductance shown here reaches values greater than 800 S/m, the highest reported for OECTs to date.^[32]

The vOECT fabrication process is interesting because it does not require exotic methods. It relies on a sacrificial layer lithography process described previously^[32] to pattern PEDOT:PSS channels. The key difference between the vOECT and planar organic electrochemical transistors (pOECTs) lies in the definition of the channel length. Previously, the pOECT channel length was defined by the distance between source and drain gold contacts formed in the same plane. This 2D arrangement of transistor contacts limits the number of transistors which may be fabricated in a given area, i.e. device density. Additionally, the channel length is limited by the fabrication process, typically to around 5 μm . The vOECT channel length, L , is defined by the thickness of the parylene-C (PaC) dielectric layer separating the vertically-stacked source and drain contacts, allowing for sub-micron resolution (here L was varied down to 450 nm). This is illustrated in **Figure 1**. As seen in Figure 1b, the exposed gold contact width defines the channel width, W , while the intermediate PaC layer thickness defines L .

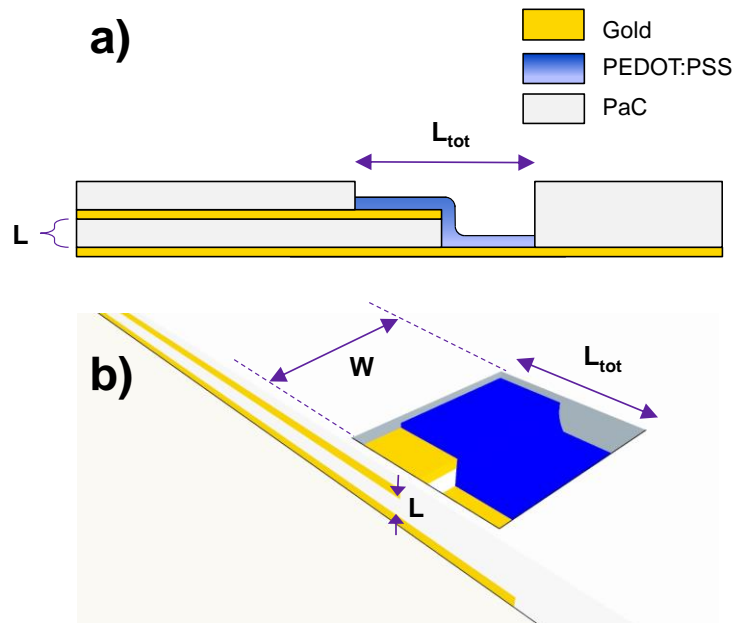


Figure 1. vOECT structure. (a) Cross-sectional view demonstrating how the PaC layer thickness defines the channel length (L). (b) Top view with PEDOT:PSS shown partially filled in, to aid in the visualization of L . W and L_{tot} are defined by the PaC opening, L_{tot} is specified for calculation of the total PEDOT:PSS layer volume.

Fabricated vOECTs exhibit very high transconductance as well as good cutoff frequency. This was characterized through steady-state (direct current, d.c.) and transient (alternating current, a.c.) measurements, performed on all vOECT and pOECT device variations. Steady-state characterization was performed by measuring the output (sweeping the drain-source bias, V_D , from 0 to -0.6 V and stepping the gate-source bias, V_G , from -0.2 to 0.6 V) and transfer (sweeping V_G from -0.2 to 0.6 with $V_D = -0.6$ V) characteristics. The transfer characteristics were subsequently utilized to extract the transconductance of various OECT channel geometries. As demonstrated by Rivnay, et al, OECTs store charge in the bulk of PEDOT:PSS rather than at an interface, and their transconductance depends on Wd/L , where d is the thickness of the PEDOT:PSS film and W and L are the channel dimensions.^[11] The inversely proportional relationship between transconductance g_m and channel length L holds

true for other transistor technologies and has been extensively studied in both inorganic^[33] and organic field effect transistors (FETs),^[3] where the thickness of the semiconductor layer, d , does not play a role. This inverse proportionality with L led to the concept of the vertical transistor where L can be substantially reduced, benefiting from increased inherent transistor amplification. To demonstrate these benefits, pOECTs and vOECTs were fabricated with the same channel widths ($W = 70 \mu\text{m}$) to allow for direct comparison. **Figure 2a** illustrates the dramatic increase in g_m for the vOECT (up to 24 mS for $L = 450 \text{ nm}$) while using less than half of the substrate area necessary for the pOECT. Figures 2b,c,d,e show SEM images of the pOECT and vOECT and corresponding schematics illustrating the W and L of the transistors.

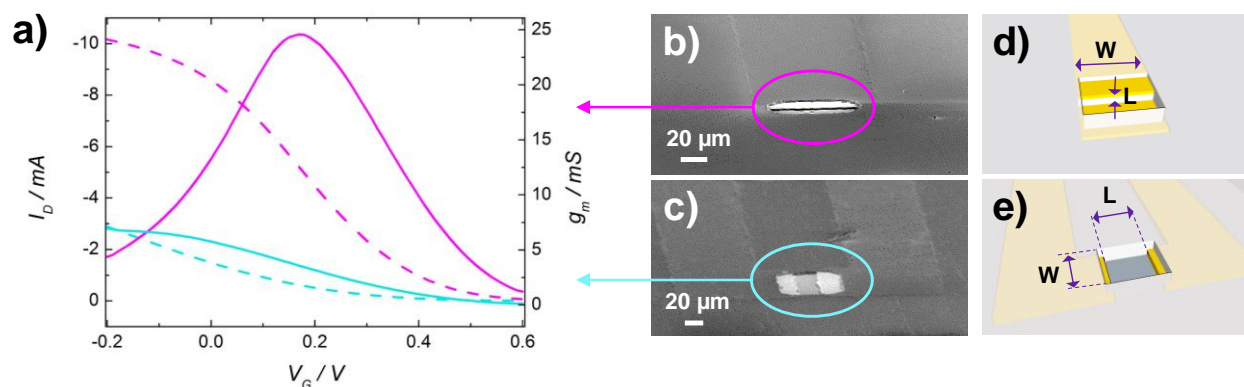


Figure 2. vOECT versus pOECT. (a) Demonstration of increased vOECT amplification through transfer characteristics (dashed lines) measured at $V_D = -0.6$ and their derivatives (transconductance, g_m , solid lines). Magenta and cyan lines correspond to the vOECT and pOECT, respectively. The vOECT in (b), (d) utilizes less substrate space than the pOECT in (c), (e) as seen by SEM and sketched in 3D.

The vOECT transfer characteristics were measured for various channel lengths ($L = 450 \text{ nm} - 1.4 \mu\text{m}$) and two PEDOT:PSS thicknesses ($d = 70 \text{ nm}, 180 \text{ nm}$). The transconductance was obtained by taking the derivative of the transfer characteristic. The scaling of g_m with decreasing L (PaC thickness) is shown in **Figure 3a, b**. To correlate vOECT and pOECT geometries, the peak transconductance values for each Wd/L are shown in Figure 3c. Here, the vOECT peak g_m values (star symbols) are overlaid with past pOECT data.^[11] The

vOECT geometry leads to higher values of Wd/L , previously unattainable with the pOECT layout and standard lithography. However, slight deviation from the fit is also observed. This effect becomes particularly evident as the channel length is shortened and/or the PEDOT:PSS thickness is increased, thus leading to higher values of Wd/L . This deviation results from the parasitic series resistance which in our case is mainly due to the contact lines, namely the source-drain series resistance, R_{SD} . This effect and the resulting difference between measured (g_m) and intrinsic (g_{mi}) transconductance has been investigated previously for short channel FETs.^[34] For instances of non-zero drain conductance (due to short-channel effects or the transistor not being in saturation mode) the intrinsic transconductance, g_{mi} , was found to be

$$g_{mi} = g_m^0 / (1 - R_{SD}g_d(1 + R_{SD}g_m^0)) \quad \text{Eq (1)}$$

where g_d is the drain conductance ($g_d = \partial I_D / \partial V_D$), and g_m^0 is:

$$g_m^0 = g_m / (1 - R_{SD}g_m) \quad \text{Eq (2)}$$

with g_m representing the measured transconductance ($g_m = \partial I_D / \partial V_G$). Eq (2) is accurate only when $g_d = 0$, and it may be seen that Eq (1) reduces to Eq (2) when this is the case.

Contact line resistances of the vOECT devices used in this study were both calculated and measured. The drain conductance was found from the OECT output characteristics for $V_D = -0.6$ V. The drain conductance and source/drain series resistance were utilized to calculate the intrinsic transconductance of vOECTs according to Eq (1) (**Figure S1**). The obtained peak g_{mi} values are represented by open star symbols in Figure 3c. These intrinsic transconductance values follow the previous trend, while reaching values of Wd/L previously unseen in OECTs. Although a potential additional issue could result from the contact resistance of the devices (i.e. the PEDOT:PSS/Au contact), the scaling of g_m as expected with reduced device dimensions indicates negligible effects due to this interface. Intrinsic transconductance values

of up to 57 mS were found, which translates to 814 S/m when normalized to the channel width (g_m/W), the highest reported value for OECTs to date.^[32] This geometry-normalization represents a method to compare transistor amplification properties while considering the device footprint.

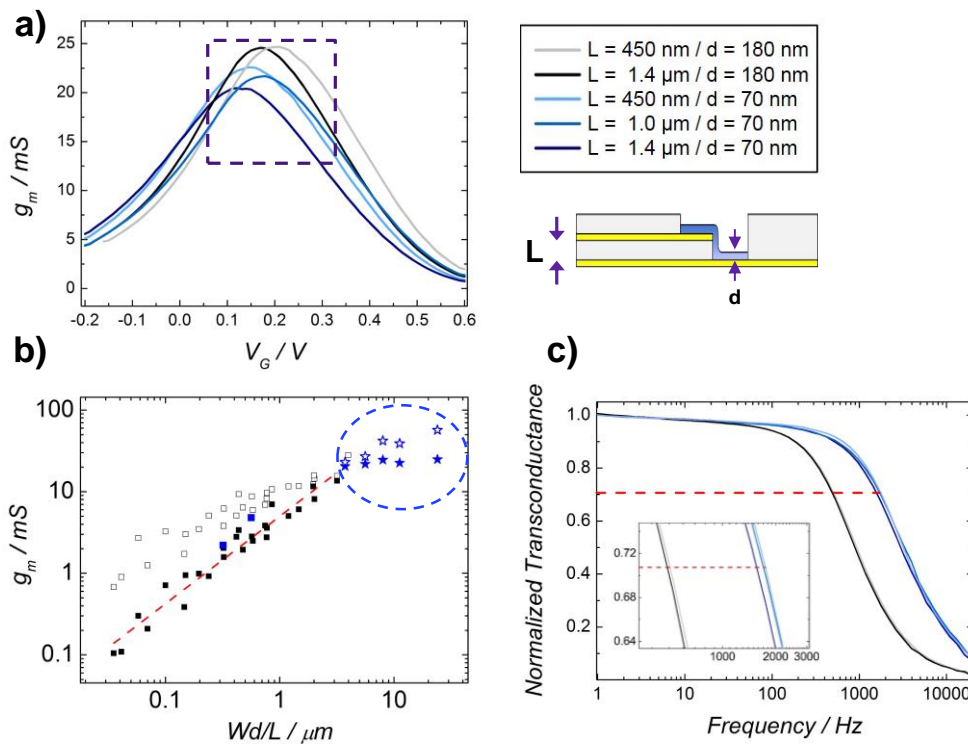


Figure 3. Scaling of transistor transconductance with channel length (L) (i.e. PaC thickness).

The increase in peak transconductance (g_m) and corresponding legend are shown in (a). (b) Peak vOECT g_m values are overlaid on previous pOECT data.^[11] Blue stars represent vOECT devices, with filled and empty symbols indicating measured (g_m) and intrinsic (g_{mi}) transconductance, respectively. Blue squares represent pOECT data obtained in this study. (c) Frequency response of vOECTs with the zoomed-in cut-off frequency in the inset.

Good transient response of vOECTs is shown through the normalized transconductance as a function of a.c. frequency, compared for different transistor geometries. In a past study the

cut-off frequency (f_T) of pOECTs was determined to depend on the PEDOT:PSS layer

volume,^[11] namely

$$\frac{1}{2\pi f_T} \sim d \cdot (WL_{tot})^{1/2} \quad \text{Eq (3)}$$

where d is the PEDOT:PSS thickness and W and L_{tot} are the PEDOT:PSS channel dimensions.

This means that the entire PEDOT:PSS film (thickness, as well as planar dimensions – i.e. the surface area in contact with the electrolyte) affects the device performance. Typically the

overlap of PEDOT:PSS on the gold electrodes of pOECTs has been negligible in comparison

to the channel length and L_{tot} may be approximated by L . Consequently, reduction in the

PEDOT:PSS thickness and/or transistor dimensions (L and W) leads to an increased cut-off

frequency. In the case of vOECTs, the polymer/gold electrode overlap must be taken into

account and the entire PEDOT:PSS volume is calculated using L_{tot} as indicated in Figure 1b.

Our vOECTs exhibited cut-off frequencies between 1.3 and 1.7 kHz for $d = 70$ nm and

approximately 550 Hz for $d = 180$ nm (Figure 3d). These results fall short of the values

obtained utilizing Eq (3) by a factor of 3 to 4 (f_T values estimated from previous data and Eq

(3) are ~ 5.5 kHz and 2.3 kHz for the 70 nm and 180 nm PEDOT:PSS films, respectively).^[11]

This is likely due to additional parasitic resistance/capacitance as a result of interconnects and

polymer/electrode overlap. However, source/drain series resistance and some amount of

overlap (i.e. additional capacitive effects) are ultimately unavoidable, particularly in situations

of limited substrate space. Thus, vOECT transient device performance indicates that higher

cutoff frequencies may be obtained as the channel dimensions are reduced (thereby reducing

the total volume), while conserving high amplification. These enhancements of the OECT are

ideal for implantable devices.

A 3D fabrication process for OECTs has been demonstrated which, when seen from a top

down perspective, consolidates all three terminals of a transistor (source/drain and local gate

if desired) into a single line perpendicular to the substrate. This dramatically decreases the spatial footprint required by each transistor and provides a pathway for transistor miniaturization. Additionally, the reduced channel length improves transconductance compared to planar OEECTs. We have demonstrated vertical OEECTs with geometry-normalized intrinsic transconductance greater than 800 S/m and with cut-off frequencies exceeding 1.5 kHz. These values represent a significant improvement over planar OEECTs and can be further enhanced by reducing parasitic effects such as the overlap between the electrolyte and source/drain contacts. Ultimately, we have shown a path toward high-density, high-performance OEECT arrays that will enable spatially and temporally resolved neural interfacing and high-throughput biological assays.

Experimental Section

vOEECT/pOEECT Fabrication: The fabrication process is based on methods reported previously.^[32] Metal interconnects were photolithographically patterned on glass microscope slides using a positive photoresist (Shipley 1813), a SUSS MJB4 UV broadband contact aligner and MF26 developer (refer to the fabrication schematic in **Figure S3**). A 2 nm chromium adhesion promoting layer and 100 nm of gold were thermally evaporated onto the substrates and interconnects were then defined through lift-off. A parylene-C (PaC) film was deposited on the glass slides to thicknesses of 450 nm – 1.4 μm (defining the channel length, L) using a SCS Labcoater 2 with 3-(Trimethoxysilyl)propylmethacrylate present in the chamber to act as an adhesion promoter. For the vOEECTs a second set of interconnects were patterned on top of the first PaC layer, defined analogously to the first metal lines and overlapping these to create the vertically-stacked source and drain contacts. Insulating and sacrificial PaC layers were subsequently deposited, both to thicknesses of 2 μm . Prior to the sacrificial PaC layer, a dilute solution of Micro-90 industrial cleaner was spin coated onto the insulating PaC. This allows the sacrificial layer to later be peeled from the substrate, defining

the OECT channel. The PaC layers were etched by reactive ion etching (RIE) in an Oxford 80 Plasmalab plus with an O_2/CHF_3 plasma to open contact pads and to create an opening down to the source/drain electrodes. These are overlapped in the case of the vOECT and this etching step simultaneously exposes the source/drain electrodes, while establishing the channel length, L . The exposed width of the gold contacts defines the channel width, W . AZ9260 photoresist was utilized as an etch mask, using the UV contact aligner and AZ developer. A dispersion of PEDOT:PSS (CleviosTM PH 1000 from Heraeus Holding GmbH), 5 volume % ethylene glycol, 0.1 volume % dodecyl benzene sulfonic acid, and 1 wt % of (3-glycidyloxypropyl)-trimethoxysilane was spin coated onto the substrates to attain a thickness of 70 nm or 180 nm, coating the vOECT vertical source/drain step or the planar channel area of the pOECT. The sacrificial PaC layer was peeled off removing superfluous PEDOT:PSS and defining the OECT channels. The devices were baked for 1 h at 140 C to crosslink the film. Finally, the devices were immersed in deionized water to remove low molecular weight compounds.

vOECT/pOECT Characterization: All OECT electrical characterization was carried out in 0.1 M NaCl aqueous solution with a Ag/AgCl pellet (Warner Instruments) gate electrode. A National Instruments PXIe-1062Q system with a PXIe-4145 source measure unit (SMU) was used along with custom LabVIEW software to carry out steady-state and transient measurements. This provided OECT output (sweeping the drain-source bias, V_D , from 0 to -0.6 V and stepping the gate-source bias, V_G , from -0.2 to 0.6 V) and transfer (sweeping V_G from -0.2 to 0.6 with $V_D = -0.6$ V) characteristics. A Keithley 2612A dual SourceMeter was utilized for the measurement of drain conductance used in the intrinsic transconductance calculation.

Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

Acknowledgements

This work was supported by Fondation pour la Recherche Médicale. H.G. was supported by the Engineering and Physical Sciences Research Council (EPSRC) of U.K. (grant EP/P511420/1). J.F. and R.M. acknowledge the National Science Foundation, Grant No. 1509909.

Received: ((will be filled in by the editorial staff))

Revised: ((will be filled in by the editorial staff))

Published online: ((will be filled in by the editorial staff))

References

- [1] S. E. Thompson and S. Parthasarathy, *Mater. Today*, **2006**, *9*, 20–25.
- [2] P. M. Zeitzoff and J. E. Chung, *IEEE Circuits Devices Mag.*, **2005**, *21*, 4.
- [3] H. Siringhaus, *Adv. Mater.*, **2005**, *17*, 2411.
- [4] H. Siringhaus, *Adv. Mater.*, **2014**, *26*, 1319.
- [5] M. Berggren and A. Richter-Dahlfors, *Adv. Mater.*, **2007**, *19*, 3201.
- [6] T. Someya, Z. Bao, and G. G. Malliaras, *Nature*, **2016**, *540*, 379.
- [7] S. J. Wilks, S. M. Richardson-Burns, J. L. Hendricks, D. C. Martin, and K. J. Otto, *Front. Neuroengineering*, **2009**, *2*.
- [8] D. Khodagholy, T. Doublet, P. Quilichini, M. Gurfinkel, P. Leleux, A. Ghestem, E. Ismailova, T. Hervé, S. Sanaur, C. Bernard, and G. G. Malliaras, *Nat. Commun.*, **2013**, *4*, 1575.
- [9] J. Reeder, M. Kaltenbrunner, T. Ware, D. Arreaga-Salas, A. Avendano-Bolivar, T. Yokota, Y. Inoue, M. Sekino, W. Voit, T. Sekitani, and T. Someya, *Adv. Mater.*, **2014**, *26*, 4967.
- [10] A. Williamson, M. Ferro, P. Leleux, E. Ismailova, A. Kaszas, T. Doublet, P. Quilichini, J. Rivnay, B. Rózsa, G. Katona, C. Bernard, and G. G. Malliaras, *Adv. Mater.*, **2015**, *27*, 4405.
- [11] J. Rivnay, P. Leleux, M. Ferro, M. Sessolo, A. Williamson, D. A. Koutsouras, D. Khodagholy, M. Ramuz, X. Strakosas, R. M. Owens, C. Benar, J.-M. Badier, C. Bernard, and G. G. Malliaras, *Sci. Adv.*, **2015**, *1*, e1400251.
- [12] D. Khodagholy, J. N. Gelinas, T. Thesen, W. Doyle, O. Devinsky, G. G. Malliaras, and G. Buzsáki, *Nat. Neurosci.*, **2015**, *18*, 310.
- [13] D. Khodagholy, J. N. Gelinas, Z. Zhao, M. Yeh, M. Long, J. D. Greenlee, W. Doyle, O. Devinsky, and G. Buzsáki, *Sci. Adv.*, **2016**, *2*, e1601027.
- [14] I. Uguz, C. M. Proctor, V. F. Curto, A.-M. Pappa, M. J. Donahue, M. Ferro, R. M. Owens, D. Khodagholy, S. Inal, and G. G. Malliaras, *Adv. Mater.*, **2017**, *29*.
- [15] L. H. Jimison, A. Hama, X. Strakosas, V. Armel, D. Khodagholy, E. Ismailova, G. Malliaras, B. Winther-Jensen, and R. Owens, *J. Mater. Chem.*, **2012**, *22*.
- [16] M. Sessolo, D. Khodagholy, J. Rivnay, F. Maddalena, M. Gleyzes, E. Steidl, B. Buisson, and G. Malliaras, *Adv. Mater.*, **2013**, *25*, 2135.
- [17] P. Lin and F. Yan, *Adv. Mater.*, **2012**, *24*, 34.
- [18] A. Williamson, J. Rivnay, L. Kergoat, A. Jonsson, S. Inal, I. Uguz, M. Ferro, A. Ivanov, T. A. Sjöström, D. T. Simon, M. Berggren, G. G. Malliaras, and C. Bernard, *Adv. Mater.*, **2015**, *27*, 3138.
- [19] A. Romeo, G. Tarabella, P. D'Angelo, C. Caffarra, D. Cretella, R. Alfieri, P. G. Petronini, and S. Iannotta, *Biosens. Bioelectron.* **2015**, *68*, 791.
- [20] A. Jonsson, S. Inal, Ilke Uguz, A. J. Williamson, L. Kergoat, J. Rivnay, D. Khodagholy, M. Berggren, C. Bernard, G. G. Malliaras, and D. T. Simon, *Proc. Natl. Acad. Sci.*, **2016**, *113*, 9440.

- [21] X. Strakosas, M. Huerta, M. J. Donahue, A. Hama, A.-M. Pappa, M. Ferro, M. Ramuz, J. Rivnay, and R. M. Owens, *J. Appl. Polym. Sci.*, **2017**, *134*,.
- [22] F. Alibart, S. Pleutin, D. Guérin, C. Novembre, S. Lenfant, K. Lmimouni, C. Gamrat, and D. Vuillaume, *Adv. Funct. Mater.*, **2010**, *20*, 330.
- [23] P. Gkoupidenis, N. Schaefer, B. Garlan, and G. G. Malliaras, *Adv. Mater.*, **2015**, *27*, 7176.
- [24] G. Tarabella, P. D'Angelo, A. Cifarelli, A. Dimonte, A. Romeo, T. Berzina, V. Erokhin, S. Iannotta, *Chem. Sci.* **2015**, *6*, 2859.
- [25] Y. van de Burgt, E. Lubberman, E. J. Fuller, S. T. Keene, G. C. Faria, S. Agarwal, M. J. Marinella, A. Alec Talin, and A. Salleo, *Nat. Mater.*, **2017**, *16*, 414.
- [26] N. Stutzmann, R. H. Friend, and H. Sirringhaus, *Science*, **2003**, *299*, 1881.
- [27] R. Parashkov, *Appl. Phys. Lett.*, **2003**, *82*, 4579.
- [28] M. Uno, *Appl. Phys. Lett.*, **2008**, *93*, 173301.
- [29] A. Fischer, R. Scholz, K. Leo, and B. Lüssem, *Appl. Phys. Lett.*, **2015**, *101*, 213303.
- [30] B. Lüssem, A. Günther, A. Fischer, D. Kasemann, and K. Leo, *J. Phys. Condens. Matter*, **2015**, *27*, 443003.
- [31] J. Kawahara, P. A. Ersman, K. Katoh, and M. Berggren, *IEEE Trans. Electron Devices*, **2013**, *6*, 2052.
- [32] D. Khodagholy, J. Rivnay, M. Sessolo, M. Gurfinkel, P. Leleux, L. H. Jimison, E. Stavrinidou, T. Herve, S. Sanaur, R. M. Owens, and G. G. Malliaras, *Nat. Commun.*, **2013**, *4*.
- [33] S. M. Sze, K. Ng. Kwok. *Physics of semiconductor devices*. John wiley & sons, **2006**.
- [34] S. Y. Chou and D. A. Antoniadis, *IEEE Trans. Electron Devices*, **1987**, *34*, 448.

Vertical organic electrochemical transistors are demonstrated with decreased channel length down to 450 nm, while retaining a simple fabrication process. These devices exhibit an intrinsic transconductance of up to 57 mS and a geometry-normalized transconductance of 814 S/m. The reduced spatial footprint and improved device performance are particularly suited for biointerfacing applications.

Electrochemical transistor

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High Performance Vertical Organic Electrochemical Transistors

