

Cite this: *J. Mater. Chem. C*, 2023,
11, 7982Received 21st December 2022,
Accepted 17th February 2023

DOI: 10.1039/d2tc05444b

rsc.li/materials-c

The design of novel organic electrochemical transistor (OECT) channel materials that can be controlled by a whole range of external stimuli is key towards the emergence of unprecedented technologies in bioelectronics. Like the established multiresponsive field-effect transistors, multiresponsive OECTs can in principle be realised *via* blending, by combining multiple components with each one imparting a specific function to the device. Here we report the first example of an optically switchable OECT which is capable of undergoing a reversible modulation of its ON current by up to 30% upon irradiation with UV and visible light. By investigating the electrical characteristics of the channel material, in conjunction with the electronic characterisation performed by a macroscopic Kelvin probe technique and photoemission yield spectroscopy in air, we gained distinct insight into the electrochemical doping process occurring within the blend upon light irradiation. Such a proof-of-concept work opens perspectives towards the implementation of complex neuromorphic operations and algorithms in OECTs.

Introduction

Functional diversification in organic materials represents a major research challenge with high potential for disruptive technological applications in tomorrow's optoelectronics and nanoelectronics.^{1–3} The motion of ionic charge carriers at a solid–liquid (or semisolid) electrolyte interface brings extraordinary functionalities to traditional organic semiconductors (OSCs), thus enabling the future development of these technologies.^{4,5} Iontronics is a discipline that exploits such concepts to control the ionic–electronic transport in mixed conductors,⁶ with its most promising application being device interfacing with biological tissues and living systems, *i.e.*,

A photo-responsive organic electrochemical transistor†

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10th Anniversary Statement

The *Journal of Materials Chemistry C* has established itself as a high quality, interdisciplinary platform for reporting innovative concepts in which materials are brought to function through their integration in optical, magnetic and electronic devices. All in all, 10 years of success and a bright future ahead!

bioelectronics.^{7,8} In the field of organic bioelectronics, organic electrochemical transistors (OECTs) have concurrently emerged as prototypical devices whose operation, mediated by ionic motion, relies on the electrochemical (EC) doping of an OSC channel material covered by an electrolyte in which the gate electrode is immersed.⁹

Among OSCs, conjugated polymers and polyelectrolytes exhibiting mixed conductivity represent the current state-of-the-art materials for bioelectronics,^{10,11} and they are thus extremely interesting for the realisation of integrated organic/bio-organic functional systems.¹² Recently developed materials have already enabled the sensing of cell parameters,¹³ the actuation of functions in living cells *in vitro*,⁸ and the efficient drug delivery in living tissues as rooted by electrochemical signaling driven by OECTs.¹⁴ Such proofs of concept opened the door to new therapeutical approaches, *e.g.* by locally monitoring disease states.¹⁵ The development of biocompatible systems that are fueled by or respond to multiple inputs under physiological conditions such as electronic stimuli, ionic gradients, light pulses and mechanical stresses is a prerequisite for the full integration of man-made electronics in the human body.^{16,17} During the past decade, the bioelectronics endeavor has mainly been focused on the figures of merit of OECTs, such as transconductance,¹⁸ reaching performances that make such

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2tc05444b>



devices suitable for functional applications.¹⁹ Molecular doping strategies in organic materials have been continuously improved towards the development of reliable electronics.²⁰ Yet, device stability needs further optimisation,²¹ due to the special environment of living systems, and more testing both in bioelectronic circuits²² and for *in vivo* applications is required.²³

The integration of an optoelectronic control in organic bioelectronics, however, remains relatively unexplored. Preliminary work has been recently reported on photocapacitive and photofaradaic processes in photoelectrodes based on OSC materials,²⁴ the use of photocapacitors to control the activity of living cells by acting as electrodes for extracellular stimulation,²⁵ or the interfacing of traditional light-based devices such as organic LEDs and solar cell photodetectors with living systems.²⁶ These bioelectronics and neuromorphic devices may then be self-powered by transparent photovoltaics, and implemented, for example, as an artificial skin.²⁷ The use of optical tools represents a viable strategy combining minimal invasiveness with the potential to yield fast and low-power photonic devices.²⁸ Light-based smart healthcare for diagnostics and therapeutics is already widespread in clinics, although the realisation of functional tools that can be directly worn by or implanted in patients is still very limited.¹ Hence, there are several opportunities to develop new (photo)chemical approaches for the realisation of switchable bioelectronic devices based on OSC materials,²⁹ aiming to mimic and control biological systems with the aid of external stimuli.³⁰ Photoswitchable biomaterials, pioneered by Willner,^{31,32} can be designed to integrate molecular photoswitches³³ into biological systems³⁴ and/or interact with bioelectronic materials as stimuli-responsive blocks to generate dynamic interfaces.³⁰

The use of light to modulate the electrical response of OECTs can provide an additional tool to remotely control the device operation in living systems with minimum invasiveness. Some OSC materials possess bandgap energies of 1–4 eV, thus they are prone to absorbing visible or UV light. This optical absorption determines an increase in conductivity, whereas its lowering occurs much more rarely.³⁵ One possible strategy to achieve a light-triggered bidirectional conductivity modulation consists of the blending of molecular photochromic additives into OSC polymer matrices.^{36,37} For applications in bioelectronics, the blending can be made by combining molecular photoswitches with an electrochemically active polymer, which is an unprecedented strategy for controlling OECT operation. Among photoswitching molecules that can be used for imparting an additional light responsiveness to the OSC polymer, spiropyran (SP) is a first choice for the design and tuning of dynamic materials.³⁸ SP derivatives exist in two relatively stable isomers under ambient conditions: the closed ring non-polar spiropyran, and the ring-opened conjugated, metastable zwitterionic merocyanine (MC) form.³⁹ It is possible to trigger the isomerisation from SP to MC upon irradiation with UV light ($\lambda \approx 365$ nm), whereas the opposite process can be accomplished by illumination with visible light ($\lambda \approx 530$ nm) or activated thermally.

On the other hand, an OSC polymer matrix is chosen based on its ability to efficiently transport charges and ions as well as its affinity with the side chains of the chosen spiropyran derivative to promote efficient blending. Since a common choice for the side chains of OECT polymers is either oligo(ethylene glycol) (OEG)⁴⁰ or alkyl chains,⁴¹ due to the advantages of the easy polymer synthesis, solution processability and structural arrangement of the polymer, we focused our attention on poly(2-(4,4'-bis(2-methoxyethoxy)-5'-methyl-[2,2'-bithiophen]-5-yl)-5-methylthieno[3,2-*b*]thiophene) (**pgBTTT**),⁴² a semiconducting polythiophene derivative whose structure comprises a BTTT backbone decorated by OEG chains in a regiosymmetrical arrangement. To ensure structural similarity for optimal mixing, a SP derivative exposing OEG chains of comparable length (**OEG-SP**) was also synthesised for improved blending (ESI† – Section S1). Optically switchable OECTs were fabricated by casting solutions containing a mixture of **pgBTTT** with SP molecular additives to form thin blend films onto Au prepatterned quartz substrates. By coupling a mixed ion-electron conductor with a light-sensitiser, we fabricated OECTs that can toggle between two states of current upon UV or visible light irradiation.

Results and discussion

Multifunctionality in multicomponent molecular films, such as those obtained by blending a polymer with a small molecule additive, is boosted *via* the subtle control over the interface between the different components.³⁷ The process of blending is aimed at avoiding phase segregation while favoring intermixing towards a maximal degree of intercomponent interfacing, as inspired by van der Waals heterostructures.⁴³ In our photo-switchable blend, the **pgBTTT** polymer imparts mixed ionic-electronic conductivity whereas the **OEG-SP** additive confers a photoresponsive nature to the hybrid material. These two components have been mixed in a 20% m/m proportion of **OEG-SP** in **pgBTTT** and deposited as drop-cast thin films from CHCl₃ solutions at various concentrations. The responsiveness of the **OEG-SP** and **OEG-SP/pgBTTT** blends to light stimuli at different wavelengths is investigated by UV-vis absorption spectrophotometry both in solutions and in thin films. Monocomponent films of **OEG-SP** reveal efficient and reversible photoisomerisation when exposed to UV (365 nm) and visible (530 nm) light (Fig. 1a). The **OEG-MC** state is characterised by a strong absorption band centred at 569 nm resulting from its conjugated zwitterionic structure. Although partially hidden by the main absorption band of the **pgBTTT** polymer, this band was also observed in the bicomponent film (red shifted by about 20 nm due to matrix effects)^{44,45} thus corroborating the efficient photoisomerisation also in the blend (Fig. 1b). Additionally, a weaker band associated with electronic transitions of higher energy⁴⁶ was *ca.* 40 nm red shifted in the **OEG-MC/pgBTTT** blend (from 352 nm of the neat **OEG-MC** film to *ca.* 395 nm) further supporting the existence of intermolecular interactions between the **MC** form and the **pgBTTT** matrix. The photoisomerisation kinetics of **OEG-MC** has been studied



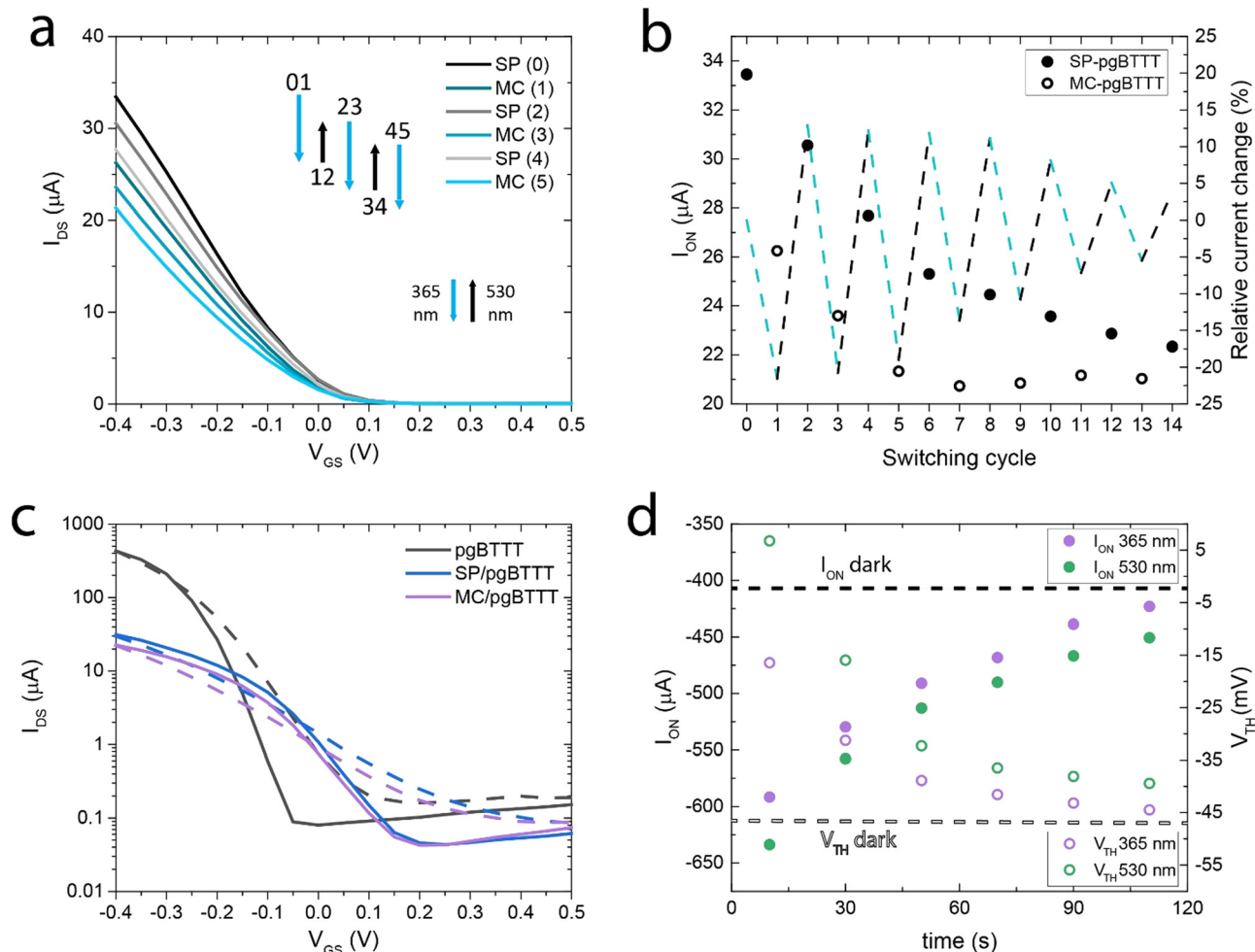


Fig. 3 (a) Photoswitching properties of OECT devices based on the **OEG-SP/pgBTTT** blend. The transfer curves were sequentially collected by performing three complete photoisomerisation cycles. Even and odd numbers refer to the spiropyran (SP) and merocyanine (MC) state, respectively. The blue (downward, 365 nm LED) and black (upward, 530 nm LED) arrows indicate the optical switching between the two states. The length of the arrows is proportional to the change in current caused by the SP switching. (b) I_{ON} data for 14 switching operations (full, **SP-pgBTTT**, and empty, **MC-pgBTTT**, circles). The relative change in I_{ON} after each switching operation is indicated by a dashed line (light blue: conversion to MC, black: conversion to SP). (c) Transfer curves of neat **pgBTTT**, **SP/pgBTTT** and **MC/pgBTTT** bottom contact Au OECTs. The curves (forward, solid, and backward, dashed, sweep) are reported on a logarithmic scale. (d) Recovery of neat **pgBTTT** device characteristics (I_{ON} and V_{TH}) after turning off the irradiation by 365 nm and 530 nm LEDs. The initial I_{ON} and V_{TH} values (dashed lines) are prior to light irradiation. All transfer curves were recorded at $V_{DS} = -0.5$ V, 0.5 M NaCl electrolyte, voltage sweep rate: 100 mV s^{-1} , W/L: 10.

7.2% absolute I_{ON} modulation) is comparable to a 57% loss in the initial signal observed in the literature.⁴⁷ Overall, the I_{ON} of the OECT decreases by 40% due to photo-fatigue, with the concomitant gradual loss of the **OEG-SP** photoswitching capability. Compared to neat **pgBTTT** OECTs (Fig. 3c) with an identical device configuration and a similar channel depth (60 nm vs. 30 nm for the blend), the product of charge carrier mobility and volumetric capacitance (μC^*), extracted from the transfer curves at an overdrive voltage ($V_{OV} = V_{GS} - V_{TH} = -0.35$ V) is a factor ≈ 15 lower for the blend (7 vs. 113 F cm^{-1} V^{-1} s^{-1}). The I_{ON}/I_{OFF} ratio is also reduced by a factor of ≈ 10 and V_{TH} is shifted towards more positive values (from -50 mV to $+220$ mV). To compensate for these performance losses in a commercial device application, V_{TH} and the absolute I_{ON} value can both be tuned by controlling the substrate wettability and the

thickness of the film, hence providing a tool for further optimisation (see ESI† – Section S3). Remarkably, both **pgBTTT** and the blend can be processed *via* spin-coating diluted (0.05 mg mL^{-1}) solutions in $CHCl_3$ to yield continuous ultrathin films whose electrical and interfacial properties can be easily investigated (Fig. S11 and S12, ESI†).

Another important difference between **pgBTTT** and the **OEG-SP/pgBTTT** OECTs is that the neat polymer devices exhibit a unidirectional increase in photoconductivity upon both UV and visible light irradiation as a result of the photogeneration of charge carriers (Fig. 3d and Fig. S13, ESI†). In fact, continuous illumination of the thin film causes an increase of I_{ON} by about 30% of the photocurrent contribution relative to I_{ON} measured in dark conditions, irrespective of the wavelength used for irradiation. A reversible V_{TH} shift of about $+50$ mV is also



observed upon irradiation. While this photoconductive state is transient (Fig. S14, ESI†), the presence of photo-responsive elements in the blend causes the quenching of the photoconductivity and promotes a prolonged subsistence of the electrical properties in one or in the other switching state. Conversely, in the neat **pgBTTT** devices, both I_{ON} and V_{TH} quickly decay as a function of time after irradiation with either UV (365 nm) or visible (530 nm) light (Fig. 3d). The photoexcitation dynamics of the neat **pgBTTT** system are discussed in more detail in the ESI.†

The optoelectronic properties of neat **pgBTTT** thin films are sensitive to the EC doping.⁴² The UV-vis spectrum of these monocomponent films is characterised by a strong absorption due to a $\pi-\pi^*$ optical transition (maximum at about 602 nm) with an absorption onset at 717 nm, resulting in an optical gap of $E_{OPT} = 1.73$ eV. The reversible doping of **pgBTTT** in a NaCl electrolyte solution is characterised by a colour change from an electric blue tint to a pale shade that tends to transparency, as the maximum absorption of the polymer redshifts to about 900 nm.⁴² To understand the operation of our OECTs, macroscopic Kelvin probe (KP) and photoelectron yield spectroscopy in air (PYSA) measurements have been made aiming to investigate the electrochemically doped (and dedoped) states of the active material.

The ionisation energy (IE) and the Fermi level (E_F) of the neat **pgBTTT** polymer on ITO substrates were measured before and after the processes of doping, and subsequent dedoping (Table 1). No significant variation of IE is observed upon the doping and dedoping processes, suggesting that the presence of Cl^- ions within the bulk of **pgBTTT** is not distorting the energetic landscape of the π -conjugated system in the neat polymer. Conversely, the Fermi level is shifted towards the HOMO by almost 0.3 eV when Cl^- ions are intercalated, indicating that the material undergoes p-doping. The incorporation of negative ions *via* EC intercalation generates hole carriers in the **pgBTTT** backbone that increase the overall conductivity of the material, enabling the OECT operation.

The illumination of **pgBTTT** *via* a UV LED does not significantly affect the energy levels of the polymer, indicating its stability towards 365 nm light irradiation (Table S2, ESI†). The spiropyran itself has IE and E_F values higher than the polymer, but, on the contrary, the merocyanine form (**OEG-SP** after UV, **OEG-MC**) is more n-doped by about 0.2 eV compared to **OEG-SP**. Both EC and the UV photoswitching experiments were carried out on the polymer blend (Table 1).

The primary effect of blending is to reduce the IE of the non-EC-doped blend compared to the neat polymer by about 0.3 eV.

Since **OEG-SP** possesses an IE higher than **pgBTTT**, the lowering of IE in the blend cannot be simply ascribed to electron transfer effects arising from the 20% m/m concentration of the photochrome. The addition of such a compound would not directly affect the minimum IE, but would rather change the doping state of the material. Aside from this major change, the $IE - E_F$ difference (related to the doping character of the material) remains around 0.5 eV before and after EC doping of the blend, but also when the blend is phototuned *via* UV irradiation, indicating that there is already an accumulation of charge carriers in the material at 0 V, in agreement with the V_{TH} values determined previously.

The effects of UV photoswitching are instead more evident in the EC-doped blend. In fact, the 0.15 eV shift in the IE (towards a more oxidised material) observed when **SP-pgBTTT** undergoes EC doping is compensated by the conversion of the **OEG-MC** form in the blend (having the EC reduction effects on the channel material). The 0.15 eV change in IE causes a variation in the charge injection (different injection barrier at the Au/OSC interface) that is responsible for the photomodulation observed in the **OEG-SP/pgBTTT** devices.⁴⁹ Note here that due to the moderate solubility of the **OEG-MC** form in the water electrolyte, accurate IE and E_F values were collected when the photoswitching experiment was performed after the EC doping step (and not *vice versa*).

Conclusions and outlook

Optically switchable OECT devices have been fabricated by blending a well-known mixed conductor, **pgBTTT**, and a versatile photochromic derivative based on a spiropyran core decorated with OEG chains, **OEG-SP**. Bottom contact devices based on a 20% m/m blend of **pgBTTT** and **OEG-SP** exhibited a modulation of the OECT I_{ON} by a factor of 30% upon illumination with 365 nm UV light. At the same time, due to the additional capability of the device to respond to light stimuli, we observed a decrease in the OECT mobility and I_{ON}/I_{OFF} ratio of about one order of magnitude compared to the neat **pgBTTT** device. The blending is accompanied by a notable change in the electrical behaviour of the OECT when exposed to light since the neat polymer exhibited solely photocurrent enhancements for both UV and visible light. The UV photocurrent quenching in the blend might suggest that more fundamental processes can be involved in the accumulation/depletion of charge carriers. The response to light stimuli decays after about 6 switching cycles due to photochemical fatigue, causing the irreversible I_{ON} negative drift over time which stabilises after

Table 1 IE and E_F of **pgBTTT** and the photoswitching **SP-pgBTTT** blend on ITO before and after doping ($V_{GS} = -0.5$ V) experiments in 0.5 M NaCl. ITO substrates underwent the cleaning procedure specified in the ESI. Data for dedoping ($V_{GS} = +0.5$ V) of **pgBTTT** and EC doping of **MC-pgBTTT** are also reported. The experimental parameters of substrate (ITO) and **pgBTTT** thin films were measured in ambient air and were averaged for $n \geq 3$ samples

Property	ITO	pgBTTT V_{GS} : 0 V	pgBTTT V_{GS} : −0.5 V	pgBTTT V_{GS} : +0.5 V	SP-pgBTTT V_{GS} : 0 V	SP-pgBTTT V_{GS} : −0.5 V	MC-pgBTTT V_{GS} : 0 V	MC-pgBTTT V_{GS} : −0.5 V
IE (eV)	5.04 ± 0.03	5.29 ± 0.05	5.32 ± 0.05	5.28 ± 0.03	4.96 ± 0.05	5.11 ± 0.01	4.99 ± 0.04	4.95 ± 0.03
E_F (eV)	4.81 ± 0.02	4.50 ± 0.01	4.79 ± 0.06	4.54 ± 0.03	4.45 ± 0.03	4.59 ± 0.02	4.48 ± 0.03	4.46 ± 0.01



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