

Cite this: *Chem. Sci.*, 2024, 15, 2055

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# Infrared photoinduced electrochemiluminescence microscopy of single cells†

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Electrochemiluminescence (ECL) is evolving rapidly from a purely analytical technique into a powerful microscopy. Herein, we report the imaging of single cells by photoinduced ECL (PECL;  $\lambda_{em} = 620$  nm) stimulated by an incident near-infrared light ( $\lambda_{exc} = 1050$  nm). The cells were grown on a metal-insulator-semiconductor (MIS) n-Si/SiO<sub>x</sub>/Ir photoanode that exhibited stable and bright PECL emission. The large anti-Stokes shift allowed for the recording of well-resolved images of cells with high sensitivity. PECL microscopy is demonstrated at a remarkably low onset potential of 0.8 V; this contrasts with classic ECL, which is blind at this potential. Two imaging modes are reported: (i) photoinduced positive ECL (PECL+), showing the cell membranes labeled with the [Ru(bpy)<sub>3</sub>]<sup>2+</sup> complex; and (ii) photoinduced shadow label-free ECL (PECL-) of cell morphology, with the luminophore in the solution. Finally, by adding a new dimension with the near-infrared light stimulus, PECL microscopy should find promising applications to image and study single photoactive nanoparticles and biological entities.

Received 8th November 2023

Accepted 7th December 2023

DOI: 10.1039/d3sc05983a

rsc.li/chemical-science

## Introduction

Electrochemiluminescence (ECL) is a well-known phenomenon of light emission generated by an electrochemical reaction in the vicinity of the electrode surface.<sup>1</sup> In an aqueous environment, it involves a highly exergonic electron-transfer reaction between a luminophore and the electrogenerated radicals of a sacrificial co-reactant, populating the excited state of the luminophore that relaxes to the ground state by emitting a photon.<sup>2</sup> In simple terms, in ECL technologies, an electrical signal is converted into an optical signal. Thanks to its high sensitivity, spatial and temporal resolution, and near-zero background, ECL has become an important tool in medical diagnosis or clinical immunoassays.<sup>3–6</sup> The most efficient and most used ECL system comprises the tris(2,2'-bipyridine)ruthenium(II) complex ([Ru(bpy)<sub>3</sub>]<sup>2+</sup>) and tri-*n*-propylamine (TPrA), which act as the luminophore and the co-reactant, respectively.<sup>7–10</sup> With this tandem system, bright ECL emission is generated during oxidation in a neutral aqueous solution (pH = 7.4).<sup>11,12</sup> Since ECL provides an optical readout, in the last decade, it has evolved into a powerful technique for microscopy

applications, allowing for the study of micro/nanoparticles and biological entities such as cells or organelles.<sup>13–23</sup> Two main imaging approaches have been developed: (i) positive ECL (ECL+) with a light-emitting object, which directly generates ECL or is labeled with an ECL tag, on a dark background; and (ii) shadow or negative label-free ECL (ECL-), which is based on a non-ECL emissive object hindering the ECL reagents diffusion and thus shadowing the ECL background.<sup>18,24</sup> Both provide complementary information about the imaged object. In the latter configuration, Su and co-workers described the ECL imaging of fingerprints and cell motion.<sup>25–28</sup> To pursue the quest for higher sensitivity, single photons and single biomolecules were imaged at the level of cells, organelles or bacteria by developing original approaches and specifically tailored nanomaterials.<sup>29–31</sup> Feng and co-workers reported a direct optical method for imaging single ECL photons generated by individual electron-transfer reaction events in solution.<sup>32</sup> They extended their original approach to image cells and nanoparticles with remarkable optical resolution.<sup>33–35</sup> Liu and co-workers reported the imaging of single biomolecules at the cell membrane level with ECL nano-emitters and their motion tracking.<sup>36–38</sup>

A different ECL approach has been developed first in organic solvents<sup>39,40</sup> and more recently in water<sup>41–45</sup> by photo-stimulating the ECL emission at depleted semiconductors (SCs). It can be considered counter-intuitive because, contrary to classical ECL, where the experiments are performed in the dark, this approach, named photoinduced electrochemiluminescence (PECL), requires an incident light ( $\lambda_{exc}$ ).<sup>46–48</sup> The photogenerated charges react with the ECL reagents dissolved in the solution

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3sc05983a>







Fig. 1 Schematic representation of two PECL microscopy modes for imaging single cells with back-illumination configuration. (a) Direct imaging by PECL+ of cell membrane labeled with the streptavidin-modified [Ru(bpy)<sub>3</sub>]<sup>2+</sup> luminophore (SA@Ru) in TPrA solution; and (b) label-free imaging of cells by PECL- with cells immersed in a PBS solution containing both [Ru(bpy)<sub>3</sub>]<sup>2+</sup> and TPrA.



Fig. 2 (a) Cyclic voltammograms and (b) corresponding ECL signals of n-Si/SiO<sub>x</sub>/Ir in the dark (black curve) or under 1050 nm infrared illumination (red curve) compared to the p<sup>+</sup>-Si/SiO<sub>x</sub>/Ir electrode in the dark (blue curve) in a PBS solution (pH 7.4) containing 30 μM [Ru(bpy)<sub>3</sub>]<sup>2+</sup> and 0.1 M TPrA. Scan rate: 50 mV s<sup>-1</sup>.

These results show that potentials higher than 0.7 V are sufficient to generate PECL emission. Thus, we selected a potential of 0.8 V for further PECL microscopy experiments on cells.

This data demonstrates that the excitation stimulus at λ<sub>exc</sub> = 1050 nm triggers PECL emission (λ<sub>em</sub> = 620 nm) at n-Si/SiO<sub>x</sub>/Ir. The anti-Stokes shift of -430 nm is large enough to allow for efficient filtering of the near-infrared excitation wavelength, thus easily discriminating the PECL signal from the incident near-infrared light (Fig. 3). Therefore, it enables microscopy



Fig. 3 Spectra of the 1050 nm LED excitation (purple curve), the resulting PECL emission (red curve), and the filter transmission (green curve).

experiments on cells using the PECL emission from [Ru(bpy)<sub>3</sub>]<sup>2+</sup>-TPrA. Moreover, the stability of these electrodes demonstrated in a recent report, with a bright PECL emission for at least 35 h, makes them suitable for microscopy applications.<sup>47</sup> Next, the n-Si/SiO<sub>x</sub>/Ir photoanodes were tested for imaging CHO-K1 cells using both PECL+ and PECL- and compared to the PL and ECL images recorded with their corresponding but non-photoactive p<sup>+</sup>-Si/SiO<sub>x</sub>/Ir anodes.

CHO-K1 cells were grown on both n-Si/SiO<sub>x</sub>/Ir and p<sup>+</sup>-Si/SiO<sub>x</sub>/Ir photoanodes. Cell growth on the sputtered Ir thin film was comparable to a glassy carbon electrode or a glass substrate, in good agreement with previously reported results.<sup>58</sup> The cells were imaged on an inverted epifluorescence microscope in two ways: PECL+ and PECL-, with a three-electrode set-up mounted on a commercially available chamber (Idylle tech transfer platform, Fig. S1†). For the first set of imaging experiments, we tested the PECL+ mode (Fig. 1a) with cells that were labeled using the [Ru(bpy)<sub>3</sub>]<sup>2+</sup> luminophore bearing a streptavidin (SA@Ru).<sup>59</sup> The cells were fixed and permeabilized with Triton X-100. This step enables the TPrA to diffuse through the membranes and to initiate the electrochemical steps over the whole cell and not only on the cell periphery. As reported previously, the entire cell became visible by ECL with this procedure.<sup>60</sup> Then, the cells were incubated with biotin X, which reacts with the primary amino groups of proteins. Biotinylation is a classical method to label cell proteins.<sup>61</sup> After incubation with SA@Ru, the cell membranes were decorated with the ECL emitter. PECL+ microscopy was performed in a commercially available ProCell buffer solution, which contains 0.18 M TPrA (Fig. 4). The direct oxidation of TPrA at the electrode surface generates the cation radical TPrA<sup>•+</sup>. This is followed by its fast deprotonation reaction, which gives the neutral radical TPrA<sup>•</sup>.<sup>62</sup> The SA@Ru labels located on the cell membranes react with both radicals, as demonstrated previously, and generate the excited state [Ru(bpy)<sub>3</sub>]<sup>2+\*</sup>, which decays by emitting ECL locally.

Fig. 4 shows the PECL+ study of the cells labeled with SA@Ru (Fig. 1a). The microscopic images (green false color; Fig. 4a and d) correspond to the PL of the [Ru(bpy)<sub>3</sub>]<sup>2+</sup> and so highlight the SA@Ru label localization. The entire cell is visible using PL. The pictures on the right (red false color; Fig. 4b and e) are the ECL



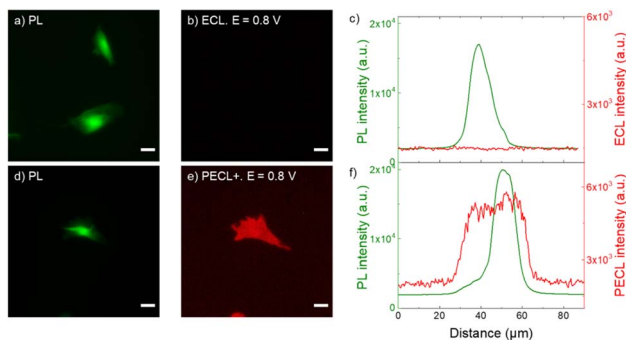


Fig. 4 (a and d) PL micrographs (green color) of CHO-K1 cells labeled with SA@Ru. (b) ECL and (e) PECL+ images (red color) of the corresponding labeled cells recorded in ProCell under near-infrared ( $\lambda_{\text{exc}} = 1050 \text{ nm}$ ) back-illumination at 0.8 V on (b)  $\text{p}^{++}\text{-Si/SiO}_x/\text{Ir}$  and (e)  $\text{n-Si/SiO}_x/\text{Ir}$  electrodes. (c and f) Comparison of the luminescence intensity profiles of the cells in PL. (c) ECL and (f) PECL+. The axis along which the profiles were extracted is shown in Fig. S4.† Green and red are false colors coding the luminescence intensity. Cells were grown on the Ir surfaces, fixed, permeabilized with Triton X-100 and then labeled with SA@Ru. Scale bar: 20  $\mu\text{m}$ .

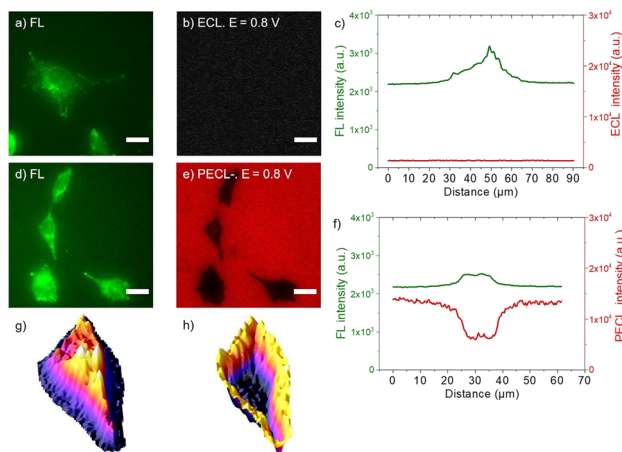
and PECL+ images recorded at 0.8 V. Green and red false colors have been selected to code the PL and ECL intensities of the images, respectively, but the same wavelength (*i.e.*, 620 nm) is emitted by the SA@Ru in the PL, ECL or PECL modes. At this potential, the cells are not visible by classic ECL because TPrA and SA@Ru are not oxidized on the  $\text{p}^{++}\text{-Si/SiO}_x/\text{Ir}$  electrode (Fig. 2). The labeled cells become visible using ECL only at more anodic potentials, typically 1.2 V (Fig. S2†). In the PECL+ mode, one can see the whole cell on the  $\text{n-Si/SiO}_x/\text{Ir}$  electrode at 0.8 V under near-infrared illumination (Fig. 4e). On the one hand, this highlights an important advantage of the reported PECL+ approach, since it shows that lower potentials are required in comparison to classic ECL experiments. On the other hand, illumination of the biological samples is required. This is not an issue in the reported conditions since the cells were fixed, but potential phototoxicity effects should be considered in future research on PECL imaging, even if the illumination wavelength is in the near-infrared range. The control experiments of the  $\text{n-type}$  electrode recorded without near-infrared light at 0.8 V and under near-infrared light at open circuit potential did not show any PECL+ images in these conditions (Fig. S3†). The cells remain invisible in these experimental conditions. This demonstrates that the PECL+ images correspond to photo-generated  $\text{h}^+$  that oxidize TPrA and populate the excited state of the SA@Ru labels. Indeed, there is no PECL emission without the synergetic effect of incident near-infrared illumination and the imposition of a 0.8 V potential, meaning that both stimuli are required to produce PECL+ images. This is confirmed by examining the PL, ECL (Fig. 4c) and PECL+ (Fig. 4f) intensity profiles that were extracted along the axis depicted in Fig. S4.† The luminescence intensity profiles plotted in green for PL and in red for ECL (Fig. 4c) and PECL+ (Fig. 4f) confirm the sharp contrast between ECL and PECL+ for these conditions. As observed previously on a glassy carbon electrode in classic ECL mode, Fig. 4d and e shows that a significantly larger spatial

extension of the cell membrane is visible by PECL+ when compared to PL. This confirms the surface-confined features of the PECL+ microscopy, which depends on the concentrations of both ECL reagents and their respective reactivity.<sup>62,68,69</sup> This thin emitting layer improves the contrast in comparison to PL images. Finally, this set of microscopy experiments also proved the possibility of back-illumination with an MIS structure to record PECL+ images of the cells.

Finally, to complete this PECL study, we tested the second ECL imaging mode, *i.e.*, the shadow PECL (PECL−) imaging of the cells on the  $\text{n-Si/SiO}_x/\text{Ir}$  electrode (Fig. 1b). The negative or shadow ECL mode is a label-free imaging method where ECL reagents (*i.e.*,  $[\text{Ru}(\text{bpy})_3]^{2+}$  and TPrA) are not covalently bonded to the cell and are dissolved in solution. This allows imaging of the morphology of the objects under investigation and the transport properties of the ECL reagents through it.<sup>60,63–65</sup> The object, which is captured on the electrode surface, locally hinders the diffusion of both the luminophore and/or the co-reactant and thus partially or totally inhibits the initial electrochemical step and, thus, the eventual ECL generation. Therefore, the object is visible by a negative optical contrast: it appears dark on a bright luminescent background because stronger ECL intensity is produced at the bare electrode.

Herein, the cells were labeled with the calcein-AM dye. This is a standard procedure to discriminate between live and dead cells.<sup>66</sup> Indeed, the hydrophobic AM (acetomethoxy) group makes it enter viable cells. Then, the acetomethoxy group is removed by the intracellular esterases, and the resulting green dye is trapped within the cell and fluoresces strongly. This labeling step simplifies the imaging process since it enables us to see them first by fluorescence (FL) and to compare the FL and the PECL− images. In addition, calcein-AM ( $\lambda_{\text{abs}} = 494 \text{ nm}$ ,  $\lambda_{\text{em}} = 517 \text{ nm}$ ) did not interfere with the electrochemical reactions involved in ECL or with the near-infrared excitation and ECL wavelengths. The fixed and calcein-labeled cells were immersed in the PBS solution containing the freely diffusing  $[\text{Ru}(\text{bpy})_3]^{2+}$  and TPrA (Fig. 1b). FL images of the calcein-AM labels define the cells' shapes (Fig. 5a and d). The images depicted in Fig. 5b and e correspond to the ECL and PECL− signals, respectively. Here again, no ECL light was emitted at 0.8 V, showing that the potential was too low to enable ECL generation at the  $\text{p}^{++}\text{-type}$  anode. In contrast, PECL occurred at 0.8 V on a  $\text{n-Si/SiO}_x/\text{Ir}$  back-illuminated electrode, and it revealed the cells in a negative (or shadow) contrast. The intensity profiles of the cells (Fig. S6†) were extracted from the FL and ECL images (Fig. 5c) and PECL− (Fig. 5f). There was no signal in ECL, whereas a local decrease in intensity was observed in the PECL− configuration. The correlation between the positive FL and the negative PECL− information is better emphasized on the 3D images. This is shown by comparing Fig. 5g with Fig. 5h, which demonstrates that the 3D FL images of a typical single cell have the same but inverted shape as the PECL intensity. As in the PECL+ mode, the control experiment of the  $\text{n-type}$  electrode without near-infrared back-illumination at 0.8 V and under near-infrared light at open circuit potential did not show any PECL in these conditions (Fig. S7†). However, the cells' morphology remains visible by classical ECL on a  $\text{p}^{++}\text{-type}$  electrode at 1.2 V (Fig. S7†). This





**Fig. 5** (a and d) Fluorescence micrographs (green color) of CHO-K1 cells labeled with calcein-AM. (b) ECL and (e) PECL– images (red color) of the same calcein-labeled cells recorded in PBS with 30  $\mu\text{M}$   $[\text{Ru}(\text{bpy})_3]^{2+}$  and 0.1 M TPrA under near-infrared ( $\lambda_{\text{exc}} = 1050 \text{ nm}$ ) back-illumination at 0.8 V on (b)  $\text{p}^{++}\text{-Si/SiO}_x/\text{Ir}$  and (e)  $\text{n-Si/SiO}_x/\text{Ir}$  electrodes. (c and f) Comparison of the luminescence intensity profiles of the same cells in FL, (c) ECL and in (f) PECL–. (g and h) 3D imaging of the same cell in (g) PL and (h) PECL– modes. The axis along which the profiles were extracted is shown in Fig. S6.† Green and red are false colors coding the luminescence intensity. Cells were grown on the Ir surfaces, labeled with calcein-AM and then fixed. Scale bar: 20  $\mu\text{m}$ .

highlights the interest in the PECL approach, which requires a lower potential. In addition, since ECL intensity decreases when recording successive ECL images of cells or other entities in TPrA solution due to a progressive lower TPrA oxidation current,<sup>67</sup> local photo-addressing different regions of the electrode may constitute an efficient and highly beneficial approach to avoid this general drawback in ECL microscopy.

## Conclusions

Herein, we reported the PECL microscopy of single cells with the anodic  $[\text{Ru}(\text{bpy})_3]^{2+}/\text{TPrA}$  system, which generates a visible emission at 620 nm under an incident non-visible light at 1050 nm in back illumination. This corresponds to an anti-Stokes shift of  $-430 \text{ nm}$  that allows easy separation of both wavelengths and thus makes the approach suitable for microscopy applications. The strategy is based on an n-type Si semiconductor coated by Ir into a MIS structure,  $\text{n-Si/SiO}_x/\text{Ir}$ , which provides high stability and bright PECL emission. CHO-K1 cells were grown on the Ir surface of the photoanode and were imaged by PECL, exploiting the photogenerated charges. PECL microscopy was demonstrated at a remarkably low potential of 0.8 V, where cells cannot be visualized in classic ECL with a non-photoactive  $\text{p}^{++}$ -type anode. Positive PECL (PECL+) of the cells was reported first by labeling the plasma membrane with the  $[\text{Ru}(\text{bpy})_3]^{2+}$  luminophore. Then, the cell morphology was imaged by an alternative label-free shadow PECL (PECL–) by immersing  $\text{n-Si/SiO}_x/\text{Ir}$  in the electrolyte containing free diffusing luminophore and co-reactant. The local control of imaging with the incident near-infrared light has

a strong potential in the microscopy field considering the large penetration depth of these wavelengths in biological samples. Finally, PECL microscopy adds a new dimension to the ECL toolbox with the tunable incident illumination, opening new possibilities for local photo-addressing of the electrode and for imaging biological samples.

## Data availability

Additional data can be obtained from the corresponding author upon request.

## Author contributions

J. Descamps: investigation, methodology, analysis, writing – original draft. Y. Zhao: investigation, methodology, writing – original draft. B. Goudeau: investigation, methodology, analysis. D. Manojlovic: conceptualization, methodology, writing – review & editing. G. Loget: conceptualization, supervision, writing – review & editing. N. Sojic: conceptualization, supervision, writing – review & editing.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by Agence Nationale de la Recherche (LiCORN, ANR-20-CE29-0006 and ELISE, ANR-21-CE42-001) and the PHC “Pavle Savic”.

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