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## COMMUNICATION

## Recognizing Conductive Islands in Polymeric Redox Surface Using Electrochemical-Coupled Vibrational Spectromicroscopy

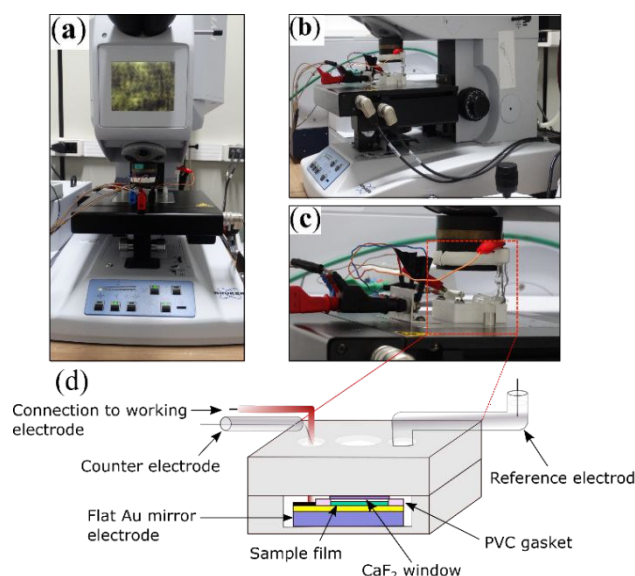
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We introduce a set up by coupling the multiplex FTIR microscopy to electrochemistry through a home-made spectroelectrochemical cell to observe real time changes in the electronic states of polymeric islands by monitoring the oxidation states of polyaniline (PANI). The resultant technique called as electrochemical-coupled vibrational spectromicroscopy (EVSM) enables the measurements of structural changes in the conductive islands of PANI with the spatial resolution as high as 2.5  $\mu\text{m}$ . Unique 2D and 3D chemical maps obtained by the integration of the spectral bands in the subtractively normalized interfacial infrared (SNIFTIR) spectra reveal electrochemical heterogeneity, showing promising topological properties control for conducting polymer-based electronic devices.

In flexible electronics and bioelectronics, it is still a great challenge to get an unambiguous correlation between the topological structural order and electrical conductivity. For instance, polyaniline (PANI) thin films and its composites have useful applications in flexible electronics and have revolutionized the actual generation of high-tech devices, such as molecular imprinted polymer-based chemical sensor<sup>1</sup>, organic electronics<sup>2</sup>, and communication devices<sup>3</sup>. Generally, Atomic Force Microscopy (AFM)-based techniques are used to reveal the out-of-plane elastic moduli and charge mobilities<sup>4</sup>, which is important to obtain conductive films with ordered topology. However, AFM requires long times and has poor spatial resolution in localizing real-time topological distortion within larger areas of the electrodes. Here we introduce an Electrochemical-Coupled Vibrational Spectromicroscopy (EVSM) technique<sup>5</sup>, that is about 100 times faster than AFM, with the possibility of analysing large surfaces in few minutes, and with the advantage of obtaining both spectroscopic and electronic information of the conductive surfaces with micrometre ( $\mu\text{m}$ ) range spatial resolution. Among conducting polymers, PANI has been extensively studied and employed in the fabrication of several devices owing to its higher electrical conductivity and easier synthesis<sup>2</sup>. However, a common issue



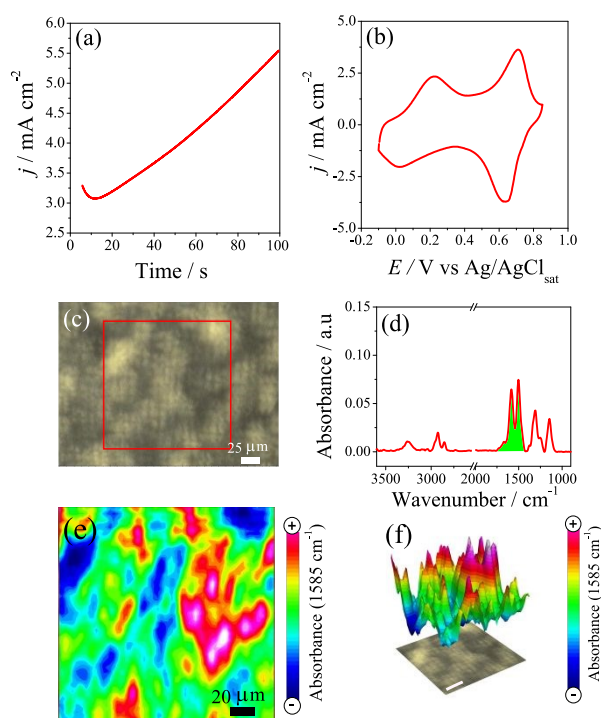
**Fig. 1. EVSM instrumental set up.** Photographs of the FTIR microscope coupled to electrochemistry through a home-built electrochemical cell mounted to the microscope stage (a) frontal view (b) lateral view (c) microscope objective and spectroelectrochemical cell (d) sketch of the cell exhibiting the connections of the reference and counter electrodes (top) and the position of working electrode, PVC, electrolyte, and  $\text{CaF}_2$  window (bottom).

with the PANI-based electrodes is the formation of disordered aggregates, resulting in the non-uniform structure of the polymer. As structural disorder is associated to the localization of charges within the polymer matrix, this leads to the development of electronic traps, which limits the charge transfer within the polymeric film<sup>6</sup>, a phenomenon called charge transfer hindrance. This has been previously observed for the PANI through electrochemical impedance spectroscopy (EIS), where the charge transfer resistance ( $R_{ct}$ ) value increased by several folds in magnitude for PANI/Au as compared to Au electrode due to the charge transfer hindrance inherent to the disordered polymer structure<sup>7</sup>.

Elucidating the impact of changes in the structural disorder, particularly under *operando* conditions, on the electronic and

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**Fig. 2. Electrodeposited PANI and micro-FTIR.** (a) Chronoamperometric curve of the electrodeposition of PANI on Au mirror electrode at applied potential of 0.8 V (aniline solution, 0.1 mol L<sup>-1</sup>) and H<sub>2</sub>SO<sub>4</sub> (0.5 mol L<sup>-1</sup>) during 100 s. (b) Cyclic voltammogram of the resultant electrode in H<sub>2</sub>SO<sub>4</sub> (0.5 mol L<sup>-1</sup>). Scan rate: 50 mV s<sup>-1</sup> (c) Optical microscopic image showing the PANI on Au-coated glass substrate. The area highlighted in red is selected for the extraction of the spectra and 2D/3D chemical maps, (d) Micro-FTIR spectrum of the PANI recorded without any potential control exhibiting the characteristic vibrational modes of the polymer structure, (e) 2D chemical map showing the distribution of the spectral bands highlighted by green in d and (f) 3D chemical map obtained in the similar fashion as 2D. The scale bar in f is 40 μm.

conducting properties of PANI electrodes by an easy-to-handle, fast and real-time spectroscopic measurements is urgently needed. The molecular structure of PANI may be conveniently studied by the Fourier Transform Infrared (FTIR) spectroscopy, as revealed by several studies reported in the literature<sup>1,8</sup>. Besides, the PANI exhibits distinctive redox properties in addition to its distinct electrochromic behaviour, that may be easily monitored by the application of FTIR spectroelectrochemical technique. The EVSM has gotten all the accessories to follow simultaneously the changes in these different kinds of properties when the external potential is modulated.

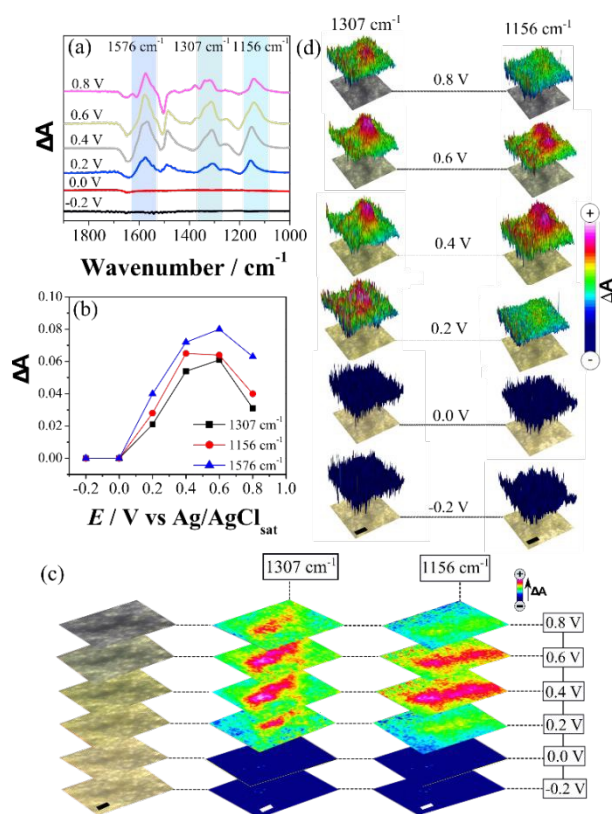
EVSM technique was applied in this work under two different perspectives: i) PANI thin film as a redox polymer and showing real time changes on the surface morphology as a function of applied electrochemical potential despite of the spectral interferences from the absorption modes of the water molecules in the studied region and ii) PANI as a conducting polymer and showing the topological structural changes in term of optical microscopic and spectromicroscopic chemical images. EVSM is introduced here as a powerful technique to explore

topological effects. Briefly, EVSM comprise a multiplex FTIR chemical imaging from focal plane array (FPA) detector connected to an inverted electrochemical cell, as shown in Fig. 1. Specific vibrational modes are selected using FPA, allowing the collection of 4096 IR spectra from the scanned area of the sample (micro-FTIR) in a single detection, which then allows the construction of the chemical maps showing the surface changes in colour patterns, providing useful information regarding the charge transfer processes and its impact on the chemical composition of the probed material in specific spatially resolved 2D regions<sup>9</sup>.

A home-made three electrodes spectroelectrochemical cell containing the PANI modified flat Au mirror as the working electrode, Pt wire sealed in a glass tube as counter electrode and Ag/AgCl<sub>sat</sub> as a reference electrode (see Fig. 1d) is used for EVSM analysis. The spectroelectrochemical cell was positioned in the motorized stage of the FTIR microscope coupled to an FTIR spectrometer, which provides the facility of moving the electrochemical cell and hence the measurement can be performed at any desired position of the working electrode by initially focusing the visible and then IR light through CaF<sub>2</sub> window when the measurement is performed (see Fig 1a-c). The CaF<sub>2</sub> window also controls the thickness of the electrolyte layer above the working electrode. The electrodeposition of PANI on flat Au mirror electrode was obtained through chronoamperometry (Fig. 2a, see SI for further detail) and characterized through cyclic voltammetry experiments, representing the characteristic redox couples (Fig. 2b), which are consistent to the interconversion of different forms of the PANI. Prior to performing the EVSM, the PANI electrode was characterized through micro-FTIR spectroscopy without coupling to the electrochemistry (for experimental detail see SI). The optical image of the sample film was initially recorded with microscope objective, as shown in Fig 2c. The FTIR spectra were extracted from the scanned area (marked by the red square in c), out of which a representative spectrum is shown in Fig. 2d, which exhibits the characteristic vibrational modes of the polymeric backbone of the PANI, such as asymmetric and symmetric C-C ring vibrations located at 1590 and 1504 cm<sup>-1</sup>, respectively. Besides the bands located at 1305, 1236 and 1150 cm<sup>-1</sup> arise from the N-H, C-C, and in plane C-H bending vibrations modes, respectively<sup>8</sup>. Fig. 2e shows the 2D chemical map obtained by the integration of the C-C ring vibrational modes, highlighted in green in the FTIR spectrum. This map shows the distribution of these two bands by the colour contrast, where the blue indicates the lowest concentration and the red suggests the highest concentration<sup>10</sup>. A visual comparison with optical microscopic image confirms that the darker regions, highly concentrated with PANI content shows a similar trend in the chemical image, where these regions show the highest concentration (red in colour) as compared to the lighter regions, which are blue in colours. The 3D chemical map (Fig. 2f) was obtained in a similar fashion and contrast similar to 2D map was obtained. The FTIR chemical maps suggest that the deposition of the polymer occurs inhomogeneously in the form of islands in the μm range.

The surface-confined electrochemical information from these conductive islands is then obtained by using FPA detector. A typical 64×64 elements FPA detector can collect 4096 spectra from the selected region of the sample with a pixel resolution of ~2.5 μm (see Fig. S1, SI) in a single detection within 2 minutes, thus making possible the spatially resolved measurements in a





**Fig. 3. EVSM of PANI modified flat Au mirror electrode recorded with FPA detector** (a) Spectral changes of PANI recorded at different applied potentials in acidic medium ( $\text{H}_2\text{SO}_4$ ,  $0.5 \text{ mol L}^{-1}$ ), (b) Changes in the intensities of the spectral bands measured from SNIFTIR spectra as a function of applied potential (c) 2D chemical maps obtained by integrating the specific bands at  $1307 \text{ cm}^{-1}$  and  $1156 \text{ cm}^{-1}$  from the SNIFTIR spectra shown in b, and (d) 3D chemical maps obtained in the similar fashion as in c. The scale bar is  $40 \mu\text{m}$ .

short duration of time. This also allows the direct construction of chemical maps from the spectra collected via a mathematical treatment, which shows the chemical distribution of vibrational mode from the selected sample area, as shown previously. EVSM system was initially evaluated by performing the cyclic voltammetry to verify whether the electrodes are properly connected, and that the system is suitable for carrying out the spectroelectrochemical measurements (Fig. S2, S1). Subtractively normalized interfacial infrared (SNIFTIR) spectra of the PANI film, where the spectrum recorded at each applied potential is subtracted from that obtained at the initial potential<sup>11</sup>, are initially recorded with the microscope and FPA detector in the reflectance mode. The electrochemical potential range used for monitoring these spectra and hence the observed changes in the PANI electrode is the same as used in cyclic voltammetry measurements, namely  $-0.2$  to  $+0.8 \text{ V}$ . Fig. 3a exhibits the SNIFTIR spectra, recorded at the different applied potential. With increasing potential, some of the peaks increased in intensity, while others decreased, suggesting that the polymer chain undergoes structural changes during the potential sweep. For example, the vibrational peaks at  $1576$ ,  $1307$  and  $1156 \text{ cm}^{-1}$  gained an increase in their intensity,

whereas the intensity of the absorption peaks at  $1510$  and  $1202 \text{ cm}^{-1}$  are decreased. The absorption band observed near  $1500 \text{ cm}^{-1}$  is assigned to the stretching vibration of the benzenoid ring of the polymer and the decrease in the intensity of this band suggests symmetry breaking and loss of benzenoid ring structure.

On the other hand, the absorption bands observed at  $1576$  and  $1487 \text{ cm}^{-1}$  arise from the semi-quinoid ring structure of the polymer chain<sup>12</sup> and an increase in their intensities with the increasing potential are consistent with the conversion of benzenoid ring (i.e.  $(-\text{B}-\text{NH}-\text{B}-\text{NH}-\text{B}-\text{NH}-\text{B}-\text{NH})$ ) into semi-quinoid ring or polaron lattice structure (i.e.  $(-\text{B}-\text{NH}-\text{B}-\text{NH}^+-\text{B}-\text{NH}-\text{B}-\text{NH}^+)$ ). Similarly, the increase in the intensity of the vibrational peaks at  $1307$  and  $1156 \text{ cm}^{-1}$  are correlated to the protonation processes of the leucoemeraldine form, resulting in the conversion to the following structure:  $(-\text{B}-\text{NH}-\text{B}-\text{NH}-\text{B}-\text{NH}^+=\text{Q}=\text{NH}^+)$ . The changes in the intensity of specific bands of the SNIFTIR spectra are followed as a function of applied potential (Fig. 3b) and are then used as reference bands for the evolution of chemical maps. All the three bands centred at  $1156$ ,  $1307$  and  $1576 \text{ cm}^{-1}$  initially gain an increase in their intensity, when the potential is increased from  $0.0 \text{ V}$  to  $+0.6 \text{ V}$ , which are consistent to the formation of semiquinoid ring and protonation processes during the conversion of reduced form (non-conducting) into partially oxidized form (conducting). Similar vibrational modes have been observed in the difference spectra of PANI, recorded as a function of a slow potential sweep ( $1 \text{ mV s}^{-1}$ ).<sup>13</sup> In general, the strengthening in the intensity of these bands indicates an increase in the degree of doping of the polymeric backbone. Alternatively, this could be due to the increase in the dipole moment by the development of the positive charge on the polymer chain, which increases in the intensity of these absorption bands. With the further increase in the potential namely at  $+0.8 \text{ V}$ , a decrease in the absorption intensity of these bands are observed and may be related to the loss of the conductivity of the polymer during its conversion into fully oxidized form. These results are in close agreement with the in situ FTIR spectroscopic studies of the PANI recorded by sweeping the potential of the working electrode in two different steps: first oxidation step ranging from  $-200 \text{ mV}$  to  $+400 \text{ mV}$  and second oxidation step ranging from  $+400 \text{ mV}$  to  $+800 \text{ mV}$ .<sup>14,15</sup> The absorption behaviour of the various absorption bands observed during the first step is identical to what we observed here in the SNIFTIR spectra up to  $0.4 \text{ V}$ , which means that during the first potential sweep, an increase in the intensity of the vibrational absorptions at  $1564$ ,  $1481$ ,  $1304$ ,  $1250$  and  $1140 \text{ cm}^{-1}$  was observed. Meanwhile, during the second oxidation step, the absorption bands at  $1629$ ,  $1571$ ,  $1507$  and  $1376 \text{ cm}^{-1}$  gained an increase in the absorption intensities, which is consistent to the appearance of some new peaks, particularly  $1627$  and  $1376 \text{ cm}^{-1}$  in the SNIFTIR spectra recorded at  $0.6$  and  $0.8 \text{ V}$ , as can be seen in Fig. 3a. The appearance of the band at  $1627 \text{ cm}^{-1}$ , which is assigned to N-ring vibration, confirm the transformation of polaron lattice into original quinoid structure ( $(-\text{B}-\text{N}=\text{Q}=\text{N}-\text{B}-\text{N}=\text{Q}=\text{N}-)$ ).

Since, PANI exhibits a distinctive electrochromic behaviour, we followed the changes in colour of the PANI by recording the optical microscopic images with the increasing potential, while dealing with chemical maps and we find a trend similar to that observed in the spectra. At the initial applied potentials of  $-0.2$  to  $0.0 \text{ V}$ , it exists in the leucoemeraldine base form and the colour of the polymer is yellow (Fig. 3c). With the increase in the

applied potential from 0.0 V - +0.6 V, the electrode colour changes from yellow to light green and then dark green, corresponding the conversion into emeraldine form. As the potential exceeds the +0.6 V, the onset potential for the conversion to pernigraniline form, the polymer adopts a dark blue colour, as can be seen in the microscopic image recorded at +0.8 V.

We monitor the chemical maps as a function of increasing potential, for which the areas under the vibrational peaks in the SNIFTIR spectra are integrated. Fig 3c shows the multiplex FTIR chemical images in the 2D format corresponding to two distinct vibrational modes at 1307 and 1156  $\text{cm}^{-1}$ . At initial applied potentials ca. -0.2 V and 0.0 V, the respective chemical maps show no contrast and are not only consistent with the spectra but with optical images at these two applied potentials, where no apparent spectroelectrochemical processes are observed. Since the electrochemical redox activity of the PANI starts only after 0.0 V, as we have seen in the cyclic voltammogram, the first SNIFTIR spectrum with distinct vibrational modes appeared only at the +0.2 V. The chemical maps were obtained by integrating the bands at 1307 and 1156  $\text{cm}^{-1}$ , where the obtained contrast is related to the chemical distribution of these bands along the sample surface. Similar procedure was adopted at +0.4 and +0.6 V. The 2D chemical images follow a trend like the spectra for these two absorption bands. A higher concentration of these bands at the centre of the sample area, particularly in the potential interval from 0.2 – 0.6 V, indicates that the polymeric molecules are not evenly distributed; instead they appear in the form of dispersed islands, when the potential on the electrode is changed. Finally, when the intensity of the absorption bands drops suddenly at +0.8 V, the chemical maps reveals the same effect by showing a decrease in the intensity of the polymeric islands at this higher potential. 3D chemical maps were obtained in a similar fashion, as shown in the Fig. 3d, where a similar tendency was observed in the colour contrast of the chemical images. All these results suggest that the electrochemical heterogeneity of the conducting polymer electrode can be successfully controlled and monitored in real time by the EVSM, showing the applicability of the technique towards the elaboration and understanding of the mechanism behind the complex electrochemical properties changes, which can help in their proper exploitation in practical applications.

In summary, the surface morphology of PANI modified flat Au mirror electrode was evaluated by using micro-FTIR spectroscopy and then underwent through in-situ monitoring of the surface redox reaction in real time through micro-FTIR spectroscopy coupled to the electrochemistry. Consistent results between the changes in the vibrational modes of the SNIFTIR spectra and the consequent 2D and 3D FTIR chemical maps were obtained at different applied potential. The results obtained are promising in the sense, that regardless of the restrictions imposed by the experimental conditions, such as the interference of absorption mode of the water (OH bending), in the region of 1590  $\text{cm}^{-1}$ , the changes in the spectral properties of the polymeric islands are successfully monitored, which gave an insight into their topological disorder along the entire electrode area and may pave the way for the better understanding of the charge transfer hindrance due to poor chain connectivity.

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## Conflicts of interest

There are no conflicts to declare.

## Notes and references

- 1 M. B. Regasa, T. R. Soreta, O. E. Femi, P. C. Ramamurthy and S. Kumar, *J. Mol. Recognit.*, 2020, 1–11.
- 2 T. Zhang, H. Qi, Z. Liao, Y. D. Horev, L. A. Panes-Ruiz, P. S. Petkov, Z. Zhang, R. Shivhare, P. Zhang, K. Liu, V. Bezugly, S. Liu, Z. Zheng, S. Mannsfeld, T. Heine, G. Cuniberti, H. Haick, E. Zschech, U. Kaiser, R. Dong and X. Feng, *Nat. Commun.*, 2019, **10**, 1–9.
- 3 M. F. Preeti, B. Vasisth, S. Singhal, L. Gaur, V. Verma, S. P. Gairola and S. K. Dhawan, in *Proceeding of International Conference on Intelligent Communication*, 2016, pp. 979–989.
- 4 P.-Y. Chen, H.-L. Hung, C.-C. Han and H.-C. Chiu, *Langmuir*, 2020, **36**, 4153–4164.
- 5 BR 10 2019 001293 5, 2019.
- 6 R. Noriega, J. Rivnay, K. Vandewal, F. P. V. Koch, N. Stingelin, P. Smith, M. F. Toney and A. Salleo, *Nat. Mater.*, 2013, **12**, 1038–1044.
- 7 A. Dey, A. Kaushik, S. K. Arya and S. Bhansali, *J. Mater. Chem.*, 2012, **22**, 14763–14772.
- 8 M. Tanzifi, S. H. Hosseini, A. D. Kiadehi, M. Olazar, K. Karimipour, R. Rezaie Mehr and I. Ali, *J. Mol. Liq.*, 2017, **244**, 189–200.
- 9 L. J. A. Macedo and F. N. Crespillo, *Anal. Chem.*, 2018, **90**, 1487–1491.
- 10 L. J. A. Macedo, F. C. D. A. Lima, R. G. Amorim, R. O. Freitas, A. Yadav, R. M. Iost, K. Balasubramanian and F. N. Crespillo, *Nanoscale*, 2018, **10**, 15048–15057.
- 11 Z. Su, V. Climent, J. Leitch, V. Zamlynnny, J. M. Felio and J. Lipkowski, *Phys. Chem. Chem. Phys.*, 2010, **12**, 15231–15239.
- 12 G. A. Planes, J. L. Rodríguez, M. C. Miras, G. García, E. Pastor and C. A. Barbero, *Phys. Chem. Chem. Phys.*, 2010, **12**, 10584–10593.
- 13 Z. Ping, G. E. Nauer, H. Neugebauer, J. Theiner and A. Neckel, *J. Chem. Soc. - Faraday Trans.*, 1997, **93**, 121–129.
- 14 N. S. Sariciftci, H. Kuzmany, H. Neugebauer and A. Neckel, *J. Chem. Phys.*, 1990, **92**, 4530–4539.
- 15 Z. Ping, H. Neugebauer and A. Neckel, *Electrochim. Acta*, 1996, **41**, 767–772.

## TOC

Micro-FTIR spectroscopy through its coupling with electrochemistry was used to evaluate the formation of conductive polymeric islands of polyaniline film on Au mirror electrode.

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