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Photo-assisted electrodeposition of an electrochemically active polypyrrole layer on anatase type titanium dioxide nanotube arrays

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Abstract

Our purpose is here to produce a large area 3D junction between anatase type TiO₂ and polypyrrole (PPy) in a Ti/TiO₂ nanotube array (NTA)/PPy assembly with the help of electrochemical methods only. We report herein the influence of light and background salt on the electrochemical polymerization of pyrrole from aqueous electrolytic solutions on titanium oxide NTAs, a well-known large specific area n-type semiconductor substrate. As titanium dioxide is known to be poorly conducting in the anodic potential range, we confirmed that light is able to initiate a polymerization process of pyrrole, at a moderately high anodic potential. For comparison with results obtained in sodium dodecylbenzenesulfonate (SDBS) at micellar concentration, we used here also lithium perchlorate as background salt at a 0.1 M concentration, and PPy growth was found to be faster in the presence of LiClO₄. Moreover in the latter case polypyrrole could be initiated in the dark as well, yet at an enhanced deposition rate under UV light. In the two solutions. the resulting polypyrrole films were found to be

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electroactive, i.e. able to undergo electrochemically driven doping/undoping processes, yet with a higher electroactivity for the film obtained in LiClO_4 . SEM-FEG and Raman spectroscopy experiments confirmed respectively that i) the initial steps of the photo-assisted electrodeposition of polypyrrole films occur first at the mouth of the titanium dioxide nanotubes, before leading to the well-known cauliflower-like morphology for PPy and ii) PPy is systematically produced in a doped state.

Keywords : TiO_2 nanotube array, polypyrrole, impedance, photoelectropolymerisation

1. Introduction

The study of semiconductor-electrolyte interfaces is of a great interest in research as a consequence of their numerous applications. The understanding of their behaviour has an impact on numerous technologies such as environment treatment (pollutant degradation), microelectronics, sensors, photocatalysis, photovoltaics, and energy storage for example. Among the most used semiconductors, TiO_2 is well known for its low cost, chemical stability, and photoelectric conversion [1].

A way to improve metal oxide performances is to control their morphology at the nanoscale [2, 3]. Precisely, the nanostructuration of TiO_2 produced by anodization of a Ti foil enables to achieve highly ordered and vertically oriented nanotubular morphology. That leads to an increased specific area, direct transport pathways and the formation of a high density of surface states that influence the interfacial electron-transfer process [4].

Covering a nanostructured TiO_2 interface with a conducting polymer film is expected to produce innovative 3D junctions possessing interesting photoelectrochemical characteristics and/or performances [5]. Among conducting polymers, polypyrrole (PPy) electrodeposition has been widely studied on usual electrode materials (platinum, gold, glassy carbon for example), thanks in part to its relatively easy electrosynthesis process and long term chemical and electrochemical stability [6, 7]. During electropolymerization process, anions from the electrolyte [8] are incorporated in the polymer film to compensate the cationic charge resulting from the oxidation (i.e. doping) state of the growing conducting polymer film and are responsible for its ion exchange properties. It is well-known that the chemical composition of the electrolytic solution, and in particular, the type and size of the ions of the background salt present in the electrolyte are determining for the ion exchange

behaviour of conducting polymers [9] and enable to control their electrochemical properties [10]. For instance, PPy films doped with large anions such as dodecylbenzenesulfonate (DBS) anions are expected to be mainly cation exchangers as a consequence of the irreversible entrapment of DBS⁻ anions in the polymer matrix, unlike PPy/ClO₄⁻ that are mainly anion exchanger films.

Attempts to electrodeposit polypyrrole on titanium oxide nanotube arrays have been recently described [5, 12-14]. The challenge is to oxidize the pyrrole monomer in a potential range in which titanium dioxide, a n-type semiconductor, is depleted from majority carriers and only photooxidation is able to provide the holes necessary for electropolymerization. Another goal is to generate a doped and electroactive conducting polymer in tubes with a few tens of nanometers diameter. In the work published by Jin *et al.* [5], electropolymerisation was carried out under UV light (at 253.7 nm) and was shown to lead to photoelectrochemically active TiO₂-NTA/PPy junctions resulting from the electrodeposition of PPy on the outer- and inner-walls of the NTs. In spite of these convincing observations, UV light was not always used in several other works published later on and sharing the same goal, i.e. electrodeposition of PPy on TiO₂-NTAs [11, 13] which led incidentally or not in both studies to polypyrrole films whose electrochemical activity was not evidenced. On the other hand, localisation of polypyrrole electrodeposition was selected in a convincing and accurate manner by using SDBS at a micellar concentration as well as current pulses shown to favour diffusion of pyrrole into or between the TiO₂ nanotubes depending on the inner diameter of the tubes and the void between them [11]. In some yet more recent contributions, beneficial effect of light on the electrodeposition process of electronically conducting polymers on metal oxide type semiconductors was thoroughly investigated and evidenced in the course of the photoelectrochemical polymerisation of PANI and PPy on nanoporous WO₃, PANI on TiO₂-NTAs [12]. Light was used by Janaky *et al* [14] for PPy deposition on

TiO₂-NTAs in a SDBS solution at micellar concentration. The authors clearly evidenced that polymerization was photoinitiated and that electropolymerization was likely to contribute to the overall polymerization reaction, once a seed layer of PPy was formed. The important result of their work was to show the higher electroactivity of the hybrid junction in comparison with an uncovered TiO₂-NTA layer.

The present work aims at comparing pyrrole (Py) electropolymerization on a TiO₂ nanotube array in two different solutions. A particular emphasis is placed i) on the role of UV light illumination during the electrochemical generation of PPy and ii) on the influence of the electrolyte anion. The deposition is performed by successive cyclic voltammetric scans. The characterization of the obtained composite structures is carried out by field emission gun-scanning electron microscopy (FEG-SEM) and Raman spectroscopy so as to determine respectively the morphology and the oxidation state of the polymer deposit. Another important part of this contribution focuses on the analysis of electrochemical behaviour of the PPy films electrochemically grown on nanotubular TiO₂ substrates as a function of the PPy electrodeposition parameters (light, background salt) using a potentiodynamic technique (Cyclic Voltammetry, CV) performed in the dark or under UV exposure, and electrochemical impedance spectroscopy (EIS).

2. Experimental part

2.1 Synthesis of TiO₂-NTAs

Prior to anodic oxidation, titanium discs (thickness 2 mm, diameter 15 mm, purity 99.6 %, purchased from Goodfellow) were polished with silicon carbide abrasive paper (P1200), and then rinsed first with ethanol, and finally in water in an ultrasonic bath. Thereafter the Ti disc was anodized in a 3 wt% NH₄F solution in ethylene glycol containing 2 vol% deionised water using a large area platinum counter electrode in a two electrode setup.

The oxidation was performed at room temperature (21°C) under a constant potential of 20 V vs. Pt for 45 min.

Once anodized, samples were ultrasonically rinsed in ethanol and deionised water successively. At last, they were annealed at 525 °C for 2 h so as to convert the amorphous TiO₂ structure into a well crystallized anatase structure which has been thoroughly characterized [4]. Anodization of the Ti substrate led to dense and homogeneous TiO₂ nanotube arrays (NTAs). Raman spectrometry and XRD spectrometry showed that anatase was the only component of the layer. The tubes exhibited smooth walls and were highly-ordered and vertically oriented as it can be seen on the FEG-SEM pictures in Fig. 1. According to a geometric model previously used [4, 15, 16] which considers a regular network of identical and equally spaced nanotubes, the developed surface of the whole nanotube arrays is given by the following equation:

$$S_t = \frac{8\pi h R_2}{\sqrt{3}(4R_2 - 2R_1 + y)^2} \times S_{geo}$$

In our case, the flat surface of the samples (working electrode) exposed to the electrolyte, i.e. the geometric surface, S_{geo} , equals 0.78 cm². In this equation, R_1 and R_2 are respectively the internal radius and the sum of R_1 and the half thickness of the wall of the nanotubes, whereas y is the thickness of the void between the tubes and h is the average length of these latter. That leads to a developed surface (S_t) of approximately 50 cm² for all the nanotube arrays produced and used throughout this work, taking into account the values of 21 nm, 26 nm, 13 nm and 960 nm for R_1 , R_2 , y and h respectively.

2.2 Electropolymerization of Pyrrole

Fresh aqueous solutions of 0.1 M distilled pyrrole monomer were prepared in different 0.1 M supporting electrolytes, either sodium dodecylbenzenesulfonate (SDBS) or lithium

perchlorate (LiClO_4). The resulting polymers are respectively named either PPy(DBS) or PPy(ClO_4). In the case of SDBS, the selected concentration value is much higher than the value of critical micellar concentration, which has been shown to be about 1.5 mM in aqueous solution [17]. Electropolymerization was carried out on a TiO_2 -NTA used as a working electrode by cyclic voltammetry between -0.7 and 1.0 V vs. SCE at a rate of 100 mV/s and at room temperature (21 °C).

For UV assisted photoelectropolymerization, the samples were illuminated with a 125 W mercury vapour lamp (HPR 125 W (Philips), wavelength range: 300-400 nm). The UV light source was placed at a distance of 25 cm from the working electrode and the UV beam was perpendicular to the working electrode surface.

PPy(DBS) and PPy(ClO_4) thin films were also electrodeposited on Pt working electrodes ($S_{\text{geo}} = 0.78 \text{ cm}^2$) and characterized for comparison.

2.3 Characterization

The morphology of the resulting TiO_2 and polypyrrole samples was studied by FEG-SEM (Zeiss, Ultra 55). A Jobin-Yvon (Labram 010) Raman spectrophotometer equipped with a He-Ne laser (wavelength 632.8 nm) was used to confirm the structure of the polymer films as well as their oxidation state.

Electrochemical characterization of the obtained polymer films was performed in a three-electrode cell that was equipped when necessary with a UV source as described above. All the potentials are referred to a saturated calomel reference electrode (SCE). The counter-electrode was a platinum spiral wire.

An electrochemical interface (Solartron SI 1287) and a frequency response analyzer (Solartron 1250) were used to perform electrochemical impedance spectroscopy (EIS) measurements at a potential value selected inside a potential range situated between 0.5 and -1.1 V vs. SCE shown to encompass the oxidation/reduction (i.e. doping/undoping) of

polypyrrole. A 0.1 Hz to 65 kHz frequency range and a 10 mV amplitude were applied during these EIS experiments. Experimental EIS spectra were fitted with ZSimpWinTM software (Princeton Applied Research).

3. Results and discussion

3.1 Electropolymerization process

3.1.1 Preliminary experiments

Before electropolymerization, preliminary experiments were performed to characterize the substrate in the absence of pyrrole in 0.1 M SDBS (or LiClO₄) aqueous solutions. Fig. 2 shows the behaviour in SDBS, but that observed in LiClO₄ is very similar. In the dark, TiO₂-NTA is charge-depleted in the potential range positive to -0.6 V vs. SCE, whereas under UV illumination, a photocurrent is generated and first increases with increasing potential until it reaches a current plateau at 0.6 V vs. SCE. In the absence of any dissolved “oxidizable” species, this anodic photocurrent is due to water oxidation and is directly related to the exposed surface of the semiconductor.

3.1.2 Electrodeposition of PPy(DBS) on TiO₂-NTAs

Fig. 3 shows the current response versus potential during electropolymerization attempts of pyrrole in a TiO₂-NTA covered Ti substrate in the -0.7 V to 1.0 V vs. SCE potential range. One can see the clear difference between the voltammogram obtained in the dark (see Fig. 3a) and that obtained under UV light illumination (Fig. 3b).

In the dark, during the 21 first scans the anodic current measured at 1.0 V vs. SCE is close to zero, similarly to the dark current usually observed with the substrate (Fig. 2), indicating that no electrochemical oxidation of the pyrrole monomer, and therefore no electropolymerization, occurs within this potential range. From the 22nd scan, the anodic

current starts slightly increasing, reaching only 40 μA at 1 V vs. SCE after 40 scans. Nevertheless, the shape of the voltammogram does not show clearly any of the features expected for the voltammogram of the doping/undoping process of PPy films. Thus PPy electrodeposition appears to be unsuccessful in the dark in our experimental conditions in the presence of SDBS.

Under UV illumination, the behaviour is very different. During the first twelve scans, the anodic plateau reveals a photocurrent which decreases with time in the potential range from 0.0 V to 1.0 V vs. SCE, This is in close agreement with the observations made by Janáky *et al.* [14], who followed the evolution of the anodic current under illumination during the first five cycles in a similar potential range. Assuming that the anodic current was mainly a photocurrent with a possible contribution of electrooxidation, they attributed the anodic current decrease to TiO_2 shielding by the growing polymer. In the present work, further scanning was performed and the results are the following:

The anodic current measured at 1.0 V vs. SCE decreases from the first up to the 12th scan, as a result of the decrease of TiO_2 surface exposed to light, during polymerization. This anodic response drops rapidly throughout the first 5 scans (from 290 μA down to 185 μA) and then, the rate of decrease slows down. During the 12 first scans, this anodic current is assumed to be the sum of TiO_2 -NTA photocurrent on one hand and growing PPy doping/undoping and Py monomer oxidation on the other hand. Nevertheless ,according to [14], the former current is certainly the main contribution to the global anodic response hiding the latter one.

From the 13th scan, a rise of the current is observed as a function of the applied potential, and the shape of the voltammogram looks closer to the classical one observed on a conducting substrate during the electrochemical deposition of polypyrrole [18, 19]. We must admit that the contribution of the photocurrent to the total current can now be neglected and

that pyrrole oxidation and polypyrrole doping become the main contributions to the global anodic response in the 0.6 to 1.0 V vs. SCE and -0.4 to 0.6 V vs. SCE potential ranges respectively.

At 1 V vs. SCE after 40 scans, the anodic current reaches 1000 μA , which has to be compared to the 40 μA reached on Fig. 3a in similar conditions without illumination. One can point out that the cathodic wave attributed to PPy undoping never stops increasing under illumination on Fig. 3b.

In close agreement with the conclusion in Ref [14], we must admit that the PPy/TiO₂-NTA junction shows higher electroactivity in a wide potential range, than uncovered TiO₂-NTA layers.

3.1.3 Electrodeposition of PPy(ClO₄) in TiO₂-NTA

For comparison, polypyrrole was also electrodeposited in the dark and under UV illumination in a LiClO₄ aqueous electrolytic solution. Fig. 4a shows the cyclic voltammogram obtained in the dark during 40 scans. Anodic and cathodic currents measured respectively at 1 V and -0.2 V vs. SCE rise endlessly in absolute values during consecutive CV scans. Fig. 4b represents the cyclic voltammogram obtained under UV light during electropolymerization in the presence of LiClO₄. The behaviour is almost the same as that in the dark, with a clear nucleation loop appearing from the second scan. It must be noticed that the current response is twice higher in Fig. 4b by comparison with Fig. 4a, showing a higher deposition rate under lighting. It is noticeable that no photocurrent plateau is observed, indicating that the oxidation of pyrrole and doping/undoping of polypyrrole are the main contributions to the anodic and cathodic currents from the first scan of polarization.

Compared to electropolymerization of pyrrole in SDBS, it appears that the rate of electropolymerization of Py in LiClO₄ is much higher and the corresponding anodic current increases rapidly when the number of scans increases. This comparison shows that the rate of

PPy deposition does not only depend on the amount of photogenerated holes created in TiO₂ through UV excitation, but also depends on the background salt, possibly through the identity of anion and/or cation inserted during the electrochemically driven doping/undoping process and/or ion and monomer transport towards the electropolymerisation sites on/in the TiO₂ - NTA. This result will be discussed below.

In the cathodic potential range, the large cathodic current value measured at -0.2 V vs. SCE due to PPy undoping increases continuously with the increase in the scan numbers, reaching much higher values in LiClO₄ ($\approx -1400 \mu\text{A}$ for the 40th scan under UV light in Fig. 4b) than in SDBS ($-300 \mu\text{A}$ for the 40th scan under UV light in Fig. 3b).

In the two electrolytes, the vanishing of the photocurrent after 40 cycles results from incident light absorption by the polypyrrole layer. Taking an absorption coefficient value of about $2 \cdot 10^5 \text{cm}^{-1}$, (value taken from ref [20]), we can estimate that a thickness of 50 nm for the polypyrrole layer is sufficient to completely absorb the incident light.

The main conclusion from the comparison between PPy formation in the two electrolytes is that light is necessary in SDBS for deposition of a polymer seed layer, whereas in LiClO₄ polymerization already starts in the dark.

3.2 Electrochemical characterization of deposited PPy film

The electrochemical characterization was performed in the same supporting electrolyte as that used for the synthesis but without Py monomer. The aim of these experiments was i) to reveal the expected electroactivity of the polypyrrole films electrodeposited on TiO₂-NTA, and ii) to estimate the ratio between PPy covered and uncovered TiO₂ areas, assuming that the dark anodic current is mainly due to PPy oxidation, whereas the photocurrent is related to uncovered TiO₂ areas

3.2.1 PPy(DBS) in SDBS electrolyte

PPy layers synthesized in SDBS using 40 consecutive cyclic voltammograms in the dark were characterized in the dark as well as under UV light. Fig. 5a shows that the behaviour is very similar to the behaviour of uncovered TiO_2 (Fig. 2), with a dark current close to zero and a photocurrent of same amplitude as in the absence of PPy. This result confirms that PPy deposition was not initiated on TiO_2 -NTA in the dark, when SDBS was used as supporting electrolyte.

PPy layers obtained in SDBS under UV light were also characterized in the dark (Fig. 5b). During the scan towards more positive potentials, an oxidation peak observed at about -0.2 V vs. SCE can be reasonably attributed to polymer oxidation (or doping). During the reverse scan a wide reduction peak with a maximum at -0.7 V vs. SCE can be attributed to the electrochemical reduction of the doped polymer (undoping). During this reverse scan, one can predict that cations from the electrolyte, such as Na^+ and/or H^+ , are inserted in the polymeric layer to ensure electroneutrality, as a consequence of the irreversible entrapment of bulky dodecylbenzenesulfonate anions during the electrodeposition step of this polymer film. These peak attributions are confirmed by the voltammogram obtained on Pt (Fig. 5c) that shows very similar features to those observed in Fig. 5b. Nevertheless, as pointed out in [14], the distance between the anodic and cathodic peaks are higher than on Pt, indicating sluggishness of the redox reactions.

When the Ti/TiO_2 -NTA/PPy(DBS) sample was characterized under UV, the anodic current clearly increased (Fig. 5b), indicating that a photocurrent contributes to the global current. Two hypotheses can be suggested: either the PPy layer is thinner than 50 nm, or the deposit is heterogeneous and the surface of TiO_2 is only partially covered by the polymer. Fig. 6a is a FEG-SEM picture of TiO_2 -NTA/PPy synthesized under illumination in SDBS after 40 scans. In comparison with the bare TiO_2 sample (Fig. 1 or inset in Fig. 6), expansion of the

wall thickness and disappearance of nanoscopic voids between and inside the tubes can be observed. Nevertheless the deposit is non-uniform over the whole surface and some TiO₂ tubes remain uncovered, which strengthen the second of our two hypotheses raised above.

Fig. 6b is a side view of the polymeric deposit taken at the sample cracks, showing that polymerisation occurs along the whole length of the tubes.

In the dark and under UV, the cathodic currents due to PPy undoping are the same (Fig. 5b), given that PPy does not produce any photocurrent under UV. Moreover we do not expect any light induced contribution in this potential range from TiO₂ which is a n-type semiconductor

The influence of the number of scans applied during electrodeposition of PPy film deposited in SDBS was investigated, assuming that the deposition is non uniform according to the FEG-SEM images (Fig. 6a). The dark anodic current measured between -0.4 V and 1.0 V vs. SCE during the characterization is then mainly attributed to PPy oxidation, whereas the photocurrent measured in the same potential range under UV light is only related to uncovered TiO₂ areas. The ratio between PPy covered and uncovered TiO₂ areas was estimated for different PPy deposits resulting from different numbers of consecutive cyclic voltammetry scans, by comparing the dark current and the photocurrent of the composite structures to the photocurrent measured for uncovered pristine nanotubes. The measurements were performed for composite structures obtained after respectively 50, 70 and 100 scans.

The percentage of uncovered nanotubes surface S% is then given by the following relation:

$$S\% = \frac{I_a^{uv} - I_a^{dark}}{I_a^{uv}(pristineTiO_2)} * 100 \quad (1)$$

From Table 1, one can see that this percentage is almost the same (~25%) whatever the number of scans. This means that the surface coverage of TiO₂-NTA by the polypyrrole does not increase anymore beyond the 50th scan in spite of the increase in the current attributed to the doping/undoping of the growing polypyrrole film. It seems that beyond the

50th scan, the polypyrrole growth implies a thickening of the film without any further spreading out on the TiO₂ substrate, the TiO₂/PPy junction providing a preferential path for electron transfer.

3.2.2 PPy(ClO₄) in LiClO₄ electrolyte

After synthesis of the organic polymer film in TiO₂-NTA in LiClO₄ aqueous electrolyte, the electrochemical activity of the composite structure was characterized in the same supporting electrolyte using cyclic voltammetry, firstly in the dark, then under UV light. When PPy was deposited in the dark in LiClO₄ electrolyte (Fig. 4a), the cyclic voltammograms obtained after transfer in a blank electrolytic solution respectively in the dark and under UV light strongly differed (see Fig. 7a). It is interesting to notice that the CV obtained in the dark clearly shows the electroactivity of PPy, by the presence of an anodic peak at 0.2 V vs. SCE and a cathodic peak at -0.15 V vs. SCE corresponding respectively to the electrochemically driven doping and undoping of polypyrrole. One can thus conclude that when the background salt chosen for the electrodeposition step is LiClO₄, PPy can be electrochemically grown on a TiO₂-NTA array in the absence of light. This is very different from SDBS solution, in which no PPy film was electrodeposited in the dark. The characterization of PPy(ClO₄) currents under light exposure clearly shows the contribution of a photocurrent to the total anodic current (Fig. 7a). This result indicates either that the PPy layer is very thin (<50 nm) or that the coverage of the surface by PPy is non homogeneous. The FEG-SEM images (not shown) indicate the same mechanism of deposition as in Fig. 6a, with an enlargement of some of the tube walls and an overall heterogeneous deposition.

When PPy was previously deposited under UV light in LiClO₄ electrolyte (Fig. 4b), a typical voltammogram corresponding to electrochemical doping/undoping of PPy was observed. Moreover, no difference was observed between the voltammogram obtained in the dark or under UV light (Fig. 7b). The absence of photocurrent indicates that the surface of

TiO₂-NTA is fully covered with an electrodeposited polypyrrole whose thickness is higher than 50 nm. Accordingly, the corresponding FEG-SEM picture in Fig. 6c shows the well-known cauliflower-like morphology over the whole surface of the TiO₂-NTA.

At this stage, one can conclude that polypyrrole deposition on TiO₂-NTA is easier (faster) in the presence of lithium perchlorate than in that of SDBS used as a supporting electrolyte. This observation confirms the detrimental character of SDBS in the efficiency of electrodeposition of polypyrrole on anatase type TiO₂-NTAs from aqueous solutions. Nevertheless a higher deposition rate does not necessarily lead to a better coverage rate of the tube walls by the polymer, given the low rate of mass transport of the reactants which control the electrochemical process. Moreover direct comparison between the two media can hardly be performed, since in SDBS surfactant aqueous solution, the pyrrole monomer is likely to be included and therefore transported inside SDBS micelles, which is not the case in LiClO₄. The efficiency of monomer supply is therefore expected to be very different in the two solutions.

For the need of comparison, polypyrrole films electrodeposited on platinum electrodes in similar conditions to those used in Fig. 4 were electrochemically characterized in LiClO₄ electrolyte (Fig. 7c). In Fig. 7a, b and c, two cathodic peaks appear in the same potential range on the three CVs, namely at about -0.85 V and -0.15 V vs. SCE respectively, whereas the oxidation peak is slightly more positive on TiO₂-NTA/PPy(ClO₄) compared to that observed on Pt-PPy(ClO₄) (at about 0.2 V vs. SCE in the former case and 0.1 V vs. SCE in the latter case). This slight difference can be attributed to a difference in conductivity between the two substrates, TiO₂ being more resistive than Pt. It is noticeable that the anodic currents measured in Fig. 7b and c are comparable, though the specific area of TiO₂-NTA is close to 50 cm² whereas that of platinum is about 0.78 cm². The similarity between the current values in the two cases can be due to a lower thickness of the PPy film on TiO₂-NTA in comparison

with Pt, or to the fact that the surface of TiO₂-NTA covered by a PPy film is close to the geometric surface, and that deposition only occurred at the upper part of the tubes. The voltammogram corresponding to the electrochemical characterization of PPy(ClO₄) in LiClO₄ electrolyte under UV light is the same as that obtained in the dark (Fig. 7b). For such PPy films, the photo-activity of the TiO₂-NTA substrate is totally hidden by the electroactivity of PPy deposit, indicating that UV illumination is strongly beneficial during electropolymerization. Moreover, it has to be noted that electrochemical deposition of PPy seems to have electrically activated the surface of TiO₂-NTA, since the TiO₂-NTA/PPy(ClO₄) junction behaves like a conductor in the dark, in spite of the underlying TiO₂-NTA that is insulating in the anodic potential range in these conditions.

A comparison between the voltammograms obtained in SDBS (Fig 5b) and those obtained in LiClO₄ (Fig 7b) shows that the shape of the voltammograms of the hybrid structure obtained in LiClO₄ is very close to the shape observed on Pt (Fig 7c) in the same medium, whereas a clear difference between the shapes on the two substrates is observed in SDBS (fig 5b and b). Though the deposited polymer amount is different in the two cases, it seems that the hybrid structure formed in LiClO₄ shows higher electroactivity in comparison with the structure formed in SDBS.

3.3 Electrochemical Impedance Spectroscopy

For a deeper insight into the electrochemical behaviour of Ti/TiO₂-NTA/PPy junctions EIS measurements were performed in the dark in order to avoid the contribution of the charge transfer resistance related to photocurrent. The electrolyte was the supporting electrolyte used for the synthesis (aqueous SDBS or LiClO₄ solutions). Nyquist diagrams were plotted in the potential range -1 V to 0.5 V vs. SCE, by 100 mV steps, but the results are only reported for three potential values: -0.9 V, -0.6 V and -0.3 V vs. SCE. (see Fig. 8 and 9 for PPy films obtained in SDBS and LiClO₄ solutions respectively).

3.3.1 EIS investigations in the dark and under UV of junctions produced in SDBS

EIS analyses of Ti/TiO₂-NTA/PPy(DBS) were performed in SDBS (0.1 M) and the response was compared to that of a bare TiO₂-NTA. The sample with a PPy film synthesized in the dark (Fig. 8, black squares) exhibits a capacitive behaviour similar to that of the pristine sample (Fig. 8, blue discs) in the whole potential range, confirming that no significant amount of PPy was deposited in the dark at the surface of the tubes. In both cases, the interface could be represented by a R_{par}(CPE) circuit in series with an electrolyte resistance R_s. R_{par} is the resistance in parallel with a CPE element characterized by Q (modulus) and n (exponent). This element was used instead of a pure capacitance, but the equivalent capacitance C could be easily calculated using Brug's equation [21]:

$$C = (RQ)^{1/n} / R \quad (2)$$

where R is defined by

$$\frac{1}{R} = \frac{1}{R_s} + \frac{1}{R_{par}} \quad (3)$$

An n value of about 0.95 was found in the whole potential range under study, and the capacitance value decreased when the potential increased, which is the usual behaviour for a space charge layer of an n-type semiconductor (Fig. 10a, black squares and blue discs). The R_{par} value was very high and increased from 4 kΩ at -1.1 V vs. SCE to 0.3 MΩ at 0.5 V vs. SCE, as a consequence of progressive depletion of the space charge layer.

The situation is more complex in the presence of PPy(DBS) films electrodeposited under UV light (Fig. 8, red triangles). Between -0.6 V vs. SCE and 0.2 V vs. SCE, they exhibit the typical shape of the EIS diagram reported in [18, 22, 23] in the case of PPy deposited on a conducting surface, with a semi-circle in the high frequency range and a diffusion feature in the low frequency range. The capacitance associated with the HF semi-circle was roughly estimated using the same equivalent circuit as for PPy electrodeposited in dark, and its evolution as a function of applied potential is also reported in Fig. 10a (red

triangles). It appears that for the 3 samples (bare TiO₂-NTA, PPy(DBS) electrodeposited in dark and under UV light) this capacitance decreases in the same way with increasing potential, confirming that the semiconducting behaviour of TiO₂-NTA prevails. This result is in agreement with the assumption that the surface of TiO₂ is only partially recovered by PPy. Indeed the capacitance of the polymer layer is then in series with that of the semiconductor, and the capacitance response is consequently dominated by the smaller value, i.e. the TiO₂ capacitance. In previous works [4, 16], it has been shown that this capacitance actually includes the space charge layer capacitance and the surface state capacitance which are in parallel. Nevertheless at first approximation, the experimental capacitance values were reported in a Mott Schottky representation, leading to a rough estimation of either the flat band potential value (between 0V and -0.1 V vs. SCE in Fig. 10b) or to the position of a localized energy state in this potential range.

The parallel resistance (R_{par}), which equals the HF semicircle diameter is much lower in the case of PPy(DBS) electrodeposited under UV light (Fig. 11). For the deposit obtained in SDBS under UV, a minimum value for R_{par} is observed at -0.5 V vs. SCE ($R_{\text{cp}} = 420 \Omega$), in agreement with the cathodic peak in Fig. 5b due to insertion of cations into the PPy film (the difference between the potentials at the maximum of current in Fig. 5b and the minimum of resistance in Fig. 11 can be due to the lack of stationarity during CV experiments). In the potential range positive to -0.2 V vs. SCE, the increase of the resistance is due to incomplete coverage of the surface of TiO₂ by PPy. In this case the blocking behaviour of TiO₂ prevails in the anodic range.

3.3.2 EIS investigations in the dark and under UV of junctions produced in LiClO₄

EIS analyses were also performed in LiClO₄ (0.1 M) in the dark and the response was compared to that of the corresponding TiO₂ NTAs layer without polypyrrole.

The Nyquist diagrams obtained for Ti/TiO₂-NTA/PPy(ClO₄) junctions synthesized in the dark (Fig. 9, black squares) are close to those of the bare TiO₂-NTA (Fig. 9, blue discs) in the cathodic potential range, but they show a less capacitive behaviour when the potential gets more anodic than -0.5 V vs. SCE, due to increasing doping of PPy when the potential increases.

The diagrams obtained for TiO₂-NTA/PPy(ClO₄) synthesized under UV light (Fig. 9, red triangles) exhibit the typical EIS diagram of PPy deposited on platinum [22-24] in the whole potential range under study. For the two PPy(ClO₄) samples, a simple R_{par}(CPE) circuit can no more be used to fit the experimental results, indicating that the PPy deposit has significantly modified the electrochemical behaviour at the interface. The resistance (R_{par}) which equals the HF semicircle diameter is reported in Fig. 11 for the four samples obtained in the two media for comparison.

The physical meaning of R_{par} is not simple and can hardly be attributed to a charge transfer resistance at the PPy/aqueous electrolytic solution interface. In [25] an accurate model has been developed in order to describe the mixed ionic /electronic conduction of a PPy membrane or of a PPy film deposited on a Pt substrate. In the present work the model is moreover complicated by the presence of a semiconducting substrate and will be the subject of a separate future work. Nevertheless for simplicity R_{par} was considered here at first approximation to compare the conductivities of the four layers.

In Fig. 11, it clearly appears that hybrid structures formed in LiClO₄ shows lower R_{par} values than the layer obtained in SDBS, confirming the higher electroactivity of junctions formed in LiClO₄.

In the two media, hybrid structures obtained under UV light show lower R_{par} values than the layer obtained in the dark in the same electrolyte. In both cases, the maximum of R_{par} is located around -0.6 V vs. SCE, whereas the resistance drastically decreases in the anodic

range. This behaviour is in agreement with the oxidation current increase observed above -0.6 V vs. SCE, in the voltammogram of Fig. 7.

EIS measurements confirm that PPy deposition on TiO₂-NTA is favoured by UV light which provides the photo-holes necessary for pyrrole electrochemical oxidation and electropolymerization. From our results it clearly appears that the rate of photo-electropolymerisation depends on the background electrolyte and that it is much higher in LiClO₄ than in SDBS solutions. Moreover, the deposition process starts right from the first scan in LiClO₄ solution, whereas an activation step seems to be necessary in SDBS solution. These observations are likely to be correlated to difference in mass transport and TiO₂ surface activation in the two solutions. The initiation of electropolymerization in LiClO₄ solution in the dark is less obvious to explain. Since during the synthesis, alternative anodic and cathodic scans are applied on the substrate, hydrogen insertion into the TiO₂ nanotubes can be promoted, increasing progressively the doping level of the semiconductor and consequently its conductivity [16, 26-28]. A similar explanation has been given in the case of the WO₃/PPy system, for which the electroactivity in the dark in the anodic range has been attributed to H⁺ uptake [12]. A slight dark deposition has also been reported for the TiO₂/PANI hybrid but was not clearly explained [12].

Our result is different from that of Kowalski *et al.* [11], who suggest that SDBS at micellar concentration favours PPy deposition in or between the tubes, even though the experimental conditions for PPy formation in their work (anodic current pulses with very high amplitude) were very different from ours.

The impedance and voltammetric results show that once the coverage rate is complete, the TiO₂-polypyrrole junction behaves like a conductor in the dark, although the underlying TiO₂ substrate was initially insulating in the anodic potential range. This suggests that the

contact between PPy and TiO₂ is ohmic. From energy level diagrams established for PPy by Miyauchi *et al.* [29], from reference [28], and from the estimation of the energy of the conduction band for TiO₂-NTA (Fig. 10b), the electron transfer process between the oxide and the polymer is schemed in Fig. 12. It appears that the highest occupied molecular orbital (HOMO) of PPy is very close in energy to the position of the conduction band of TiO₂, leading to the possibility of direct charge transfer between the two materials. The initiation of polymerisation on TiO₂ under lighting, with a well adapted light source providing the holes necessary for oxidative polymerization can easily be understood. But the initiation of electropolymerisation in LiClO₄ solution in the dark is less obvious to explain. and further experiments are necessary for a clear understanding.

3.4 Characterization of Ti/TiO₂-NTA/PPy junctions by Raman spectroscopy

A Raman spectrum recorded in the air is reported as an example in Fig. 13. It corresponds to the sample synthesized in SDBS under illumination, after 40 scans of polarization. Similar spectra were also obtained on hybrid samples synthesized in LiClO₄. According to several detailed interpretations of Raman spectra of polypyrrole films [31-33], the band at 926 cm⁻¹ corresponds to the CH out-of-plane bending, the band at 987 cm⁻¹ is assigned to the ring deformation, the band at 1050 cm⁻¹ reflects the CH in-plane bending, bands at 1323 cm⁻¹ and 1329 cm⁻¹ are linked to the C-C stretch and the band at 1605 cm⁻¹ corresponds to the C=C backbone stretch. Moreover, the spectrum does not display a peak at 1680 cm⁻¹ usually assigned to C=O stretching vibration, and implying overoxidation of the polymer film. The ratio between peak intensities at 926 cm⁻¹ and 987 cm⁻¹ is superior to the unit. This fact combined to the presence of a peak at 1605 cm⁻¹ confirm the oxidized state of the polymer [34].

4. Conclusion

This work confirms the beneficial role of UV light illumination on the conductivity of anatase type TiO_2 NTAs in the anodic potential range and therefore on the electrodeposition of polypyrrole on such substrates. This positive influence of lighting during electropolymerisation, which has already been evidenced by Janaky and co-workers[14], when PPy was deposited in sodium dodecylbenzenesulfonate, was also observed in the present work in lithium perchlorate containing aqueous electrolytic solutions, yet with a higher deposition rate in the second case.

Nevertheless two main differences have been evidenced between the two media:

- Deposition of PPy onto TiO_2 NTAs in the dark is observed from the first scan in the presence of LiClO_4 , but is hardly detected in SDBS, in which PPy deposition must be photoinitiated..
- Once the films are formed, hybrid structures obtained in LiClO_4 showed higher electroactivity than those formed in SDBS.

The overall low deposition efficiency in SDBS was mainly attributed to anion and monomer transport in and between the TiO_2 NTs in micellar and therefore visqueous SDBS aqueous solutions. In the two media, once the PPy deposit was initiated, the film kept on growing mainly at the covered sites, which are likely to constitute preferential paths for charge transfer. This result, together with the energy level diagram, suggests that the contact between PPy and TiO_2 is ohmic, leading to an electroactive composite 3D junction. Indeed, whatever the electrodeposition conditions are, the resulting PPy films were found to be generated in their oxidized (and not overoxidized) state, according to Raman investigations and thus able to undergo electrochemically driven redox (doping/undoping) reactions, as shown by cyclic voltammetry and electrochemical impedance spectroscopy.

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Number of electropolymerization cycles under UV [1 cycle: from -0.7 V to 1 V VS. SCE at 100 mV/s]	Characterization in SDBS [From -1.09 to 0.5 V VS. SCE at 100 mV/S]		Uncovered TiO ₂ -NTA surface (%)
	$I_a^{uv} - I_a^{darkness}$ at 0.5 V VS. SCE (μA)	Average I_a^{uv} (TiO ₂ pristine) at 0.5 V VS. SCE (μA)	
50	66	260	25.4
70	65	260	25.0
100	67	260	25.8
Pristine	260	260	100

Table 1. Percentage of uncovered TiO₂-NTA surface.

Fig. 1. FEG-SEM images of a pristine TiO_2 -NTA : a) top view and b) side view.

Fig. 2. Cyclic voltammetry analyses of pristine TiO_2 -NTA in an aqueous electrolytic solution containing SDBS (0.1 mol.L^{-1}) at 10 mV/s in the dark and under UV illumination.

Fig. 3. Consecutive CVs (40 scans) allowing electropolymerization of pyrrole on a $\text{TiO}_2\text{-NTA}$ in an aqueous electrolytic solution containing SDBS (0.1 mol.L^{-1}) and Py (0.1 mol.L^{-1}) at 100 mV/s between -0.7 V and 1 V/SCE , a) in the dark and b) under UV illumination.

Fig. 4. Consecutive CVs (40 scans) allowing electropolymerization of pyrrole on a TiO_2 -NTA in an aqueous electrolytic solution containing LiClO_4 (0.1 mol.L^{-1}) and Py (0.1 mol.L^{-1}) at 100 mV/s between -0.7 V and 1 V/SCE a) in the dark and b) under UV illumination.

Fig. 5. CV characterization of $\text{TiO}_2\text{-NTA/PPy(DBS)}$ junction in an electrolytic aqueous solution containing SDBS (0.1 mol.L^{-1}) at 10 mV/s under UV illumination or in the dark corresponding to electrodeposition of PPy for a) in the dark, b) under UV and c) on Pt, a metal on which electrodeposition of PPy is not influenced by illumination.

Fig. 6. FEG-SEM images of TiO₂-NTA/PPy junctions obtained in SDBS, under UV and potentiodynamically between -0.7 V and 1V at 100 mV/s: a) Top view after 40 scans, TiO₂ nanotubes are partially filled by PPy (Inset: FEG-SEM image of a pristine TiO₂-NTA) (b) Side view over the sample cracks after 20 scans (c) cauliflower-like structure in LiClO₄ after 40 scans.

Fig. 7. CV characterization of $\text{TiO}_2\text{-NTA/PPy}(\text{ClO}_4)$ junction in an aqueous electrolytic solution containing LiClO_4 (0.1 mol.L^{-1}) at 10 mV/s under UV illumination or in the dark corresponding to electrodeposition of PPy for a) in the dark, b) under UV illumination and c) on Pt, a metal on which electrodeposition of PPy is not influenced by illumination.

Fig. 8. Nyquist representation of EIS spectra of a TiO_2 -NTA/PPy(DBS) junction synthesized in the dark (square) or under UV illumination (triangle). EIS analyses were performed in an electrolytic aqueous solution containing SDBS (0.1 mol.L^{-1}) in the dark. EIS spectrum corresponding to pristine TiO_2 -NTA is also represented (disc).

Fig. 9. Nyquist representation of EIS spectra of a $\text{TiO}_2\text{-NTA/PPy}(\text{ClO}_4)$ junction synthesized in the dark (square) or under UV illumination (triangle). EIS analyses were performed in an electrolytic aqueous solution containing LiClO_4 (0.1 mol.L^{-1}) in the dark. EIS spectrum corresponding to pristine $\text{TiO}_2\text{-NTA}$ is also represented (disc).

Fig. 10. a) Variation of the space charge layer capacitance as a function of applied potential in SDBS in the presence of PPy (TiO₂-NTA is measured in SDBS without PPy) b) in a Mott Schottky representation. The plots correspond to TiO₂ NTA/PPy(DBS) junctions synthesized in the dark (black squares) or under UV illumination (red triangles). Evolution of the capacitance corresponding to pristine TiO₂-NTA is also represented (blue discs).

Fig. 11. Evolution of the R_{par} value as a function of potential: a) overall view and b) Enlargement of the low resistance domain. $\text{TiO}_2\text{-NTA/PPy(DBS)}$ junction synthesized in the dark is represented by squares. $\text{TiO}_2\text{-NTA/PPy(DBS)}$ junction synthesized under UV illumination is represented by triangles. $\text{TiO}_2\text{-NTA/PPy(ClO}_4\text{)}$ junction synthesized in the dark is represented by stars. $\text{TiO}_2\text{-NTA/PPy(ClO}_4\text{)}$ junction synthesized under UV is represented by pentagons. Pristine $\text{TiO}_2\text{-NTA}$ is represented by discs.

Fig. 12. Band structure of PPy* in comparison with flat band potential of TiO₂ (*From ref [27]). The conversion of the vacuum scale to the potential scale is made using $E_{\text{SCE}} = -4.70 \text{ eV}$ [28].

Fig. 13. Raman spectrum of a TiO₂-NTA/PPy(DBS) junction.