

A comparative study of two different media effect on the electropolymerization of 2-(9-ethylcarbazol-3-yliminomethyl)phenol by cyclic voltammetry, impedance spectroscopy, XPS, UV-visible measurements and DFT calculation.

Abstract

A carbazole-based monomer, 2-(9-ethylcarbazol-3-yliminomethyl)phenol (SIC), was first synthesized and characterized. It was afterward polymerized by potentiostatic methods. Oxidative polymerization of SIC was carried out in two media. The first one (*i.e.* Bu₄NBF₄ (0.1M)/MeCN) gives the poly(2-(9-ethylcarbazol-3-yliminomethyl)phenol). However, the second one (LiClO₄/MeCN + 35mM HClO₄), leads to another polymer, the poly(2-(9-ethylcarbazol-3-yliminomethyl)cyclohexa-2,5-diene-1,4-dione). These two novel polymers will be abbreviated hereafter as PSIC and PDIC, respectively. If the formation of PSIC was expected, the formation of PDIC in acidic medium isn't. It was explained by a hydroxyl function oxidation of SIC. The obtained polymers were then characterized by cyclic voltammetry, electrochemical impedance spectroscopy, XPS, IR and UV-Vis spectroscopies. Furthermore, the corresponding electrochemical and optical band gap values were calculated in order to get an idea about the conductive properties. The related results show that PDIC and PSIC are good conductive polymers.

Keywords: Conductive polymer; Electropolymerization; Cyclic voltammetry; Polyimine; Band gap, DFT calculation.

1. Introduction

Electronic polymers are the subject of much research due to their reversible switchability between insulating and conducting state. This propriety allows them to be used in several applications, such as energy storage materials and electronic devices. Among organic polymers, polycarbazoles exhibit an interesting optoelectronic and photochemical properties [1,2] due to the high electron-donating character of the carbazole ring. Owing to their proton doping [3], polycarbazole films show a fair conductivity and stable redox properties in protic medium [4].

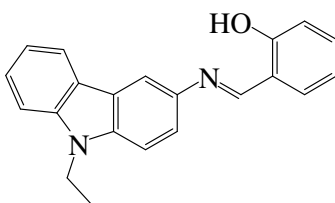
In the present work, we investigated the electrosynthesis of a polyimine based on a carbazole unit. The monomer of this class of polymers is easily obtained. Its synthesis consists on a simple condensation between an amine and an aldehyde. We reported the electrochemical behaviour and the

anodic electropolymerization of the compound 2-(9-ethylcarbazol-3-yliminomethyl)phenol (**SIC**) in two different electrolytic solutions (Bu_4NBF_4 (0.1M)/MeCN and $\text{LiClO}_4/\text{MeCN} + 35\text{mM HClO}_4$).

2. Experimental

2.1. Materials

3-amino-9-ethylcarbazole and salicylaldehyde were obtained from *Aldrich*, and used without further purification. Tetrabutylammonium tetrafluoroborate obtained from *FLUKA*, was recrystallized in a mixture of water/methanol and dried at 100°C for 24h. Aluminium oxide was dried at 300°C under vacuum for at least 24h prior to use. The monomer (**SIC**) was prepared by condensation between 3-amino-9-ethylcarbazole and salicylaldehyde in ethanolic medium. A yellow product was obtained with a yield of 65% (m.p. 114°C).



2-(9-ethylcarbazol-3-yliminomethyl)phenol (**SIC**)

2.2. Electrochemical measurements

All electrochemical experiments were carried out using a computer-controlled VoltaLab PGZ 301 potentiostat/galvanostat. All data were collected and analyzed using Voltamaster software. A three-electrode cell consisting of glassy carbon disk (GCE) as working electrode (1cm^2), an Ag/AgCl electrode as reference and a platinum wire as counter electrode were used. A 0.1M solution of tetrabutylammonium tetrafluoroborate (Bu_4NBF_4) in acetonitrile was used as supporting electrolyte. Anhydrous Al_2O_3 was added for drying, into the electrolytic solution. All solutions were deaerated by bubbling N_2 gas for a few minutes prior to electrochemical measurements. The potential of ferrocene/ferrocinium redox couple in the supporting electrolyte-solvent system was measured and found to be 0.435V vs. Ag/AgCl. Square wave parameters were pulse half-peak-to-peak $\Delta E_{\text{SW}} = 25\text{mV}$, staircase step height $\Delta E_{\text{S}} = 5\text{mV}$ and scan rate $v = 100\text{mV/sec}$. The electrochemical impedance measurements on the modified electrode were done over the frequency range from 0.1 to 100kHz with 10mV as amplitude. For large-scale electrolysis, the working potential was controlled by a Tacussel potentiostat PRT 20-2X, and the charge consumption was measured with a Tacussel IG 6-N integrator.

2.3. Optical measurements

UV-Vis measurements were performed on a JASCO V-660 spectrophotometer. The absorption spectra of the compounds were recorded on ITO glass transparent film (solid phase). The infrared absorption spectra were recorded on a JASCO FT/IR-6300 spectrometer. The XPS analyses were performed with a photoelectron spectrometer ESCALAB 250 (Thermo VG).

2.4. Theoretical calculations

Theoretical calculations were performed to estimate the energy values of the frontier molecular orbitals (HOMO and LUMO) of the studied molecules. The molecular geometries were optimized with the DFT method using the Gaussian 09 software. The used functional was B3LYP (Becke 3-parameter-Lee Yang-Parr) with the basis set of 6-311G(d,p). The figures were generated with GaussView 5.0.

3. Results and discussion

3.1. Electrochemical polymerization of SIC in $Bu_4NBF_4(0.1M)/MeCN$

The electrochemical behaviour of SIC was investigated on a GCE electrode by cyclic voltammetry (CV) and square wave voltammetry (SWV). Using CV, the voltammogram obtained during electrooxidation of 5×10^{-3} M of SIC in Bu_4NBF_4 (0.1M) / MeCN (Figure 1), showed two peaks at 1.12 and 1.45V vs. Ag/AgCl reference electrode. These two peaks appeared also in the SWV's voltammogram at slightly lower potentials, at 1 and 1.3V, (Figure 2). This difference in peaks potentials may be due to the high sensitivity of SWV techniques [5].

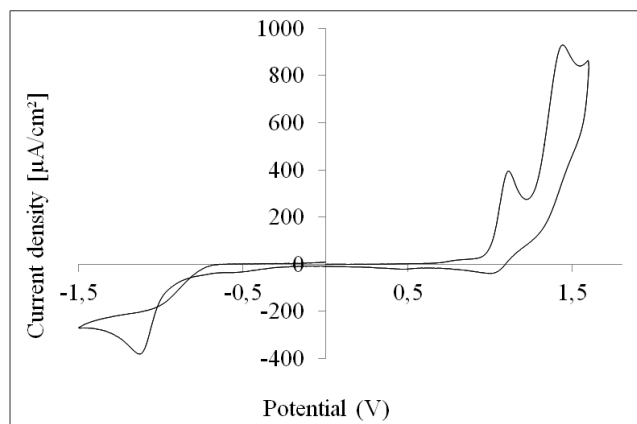


Figure 1. Cyclic voltammogram of SIC (5×10^{-3} M) in Bu_4NBF_4 (0.1 M)/MeCN; first cycle, $v = 100$ mV/s; working electrode: GCE; reference electrode: Ag/AgCl

The first peak probably corresponds to the oxidation of hydroxyl group which reacts easily due to the presence of n electrons into the oxygen atom [6]. At this potential, no polymerization of SIC

was observed. The second peak corresponds to the formation of the radical cation of the monomer. In the cathodic scan region, SIC showed only one irreversible reduction peak related to the reduction of imine moiety [7].

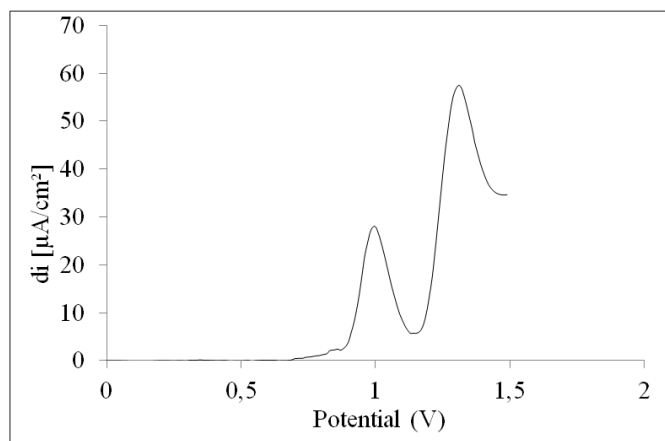


Figure 2. Square wave voltammogram of SIC (5×10^{-3} M) in Bu_4NBF_4 (0.1 M)/MeCN; $\nu = 100$ mV/s; working electrode: GCE; reference electrode: Ag/AgCl

The electrochemical polymerisation of SIC performed by recurrent potential scans between 0 and 1.4 V in Bu_4NBF_4 (0.1M)/MeCN (Figure3), led to the gradual development of a new redox system centred at lower potential than the monomer (around 1.1V). The current densities of the new system increased during the successive cycles, indicating that the formed polymer film has a good adherence on the electrode [8].

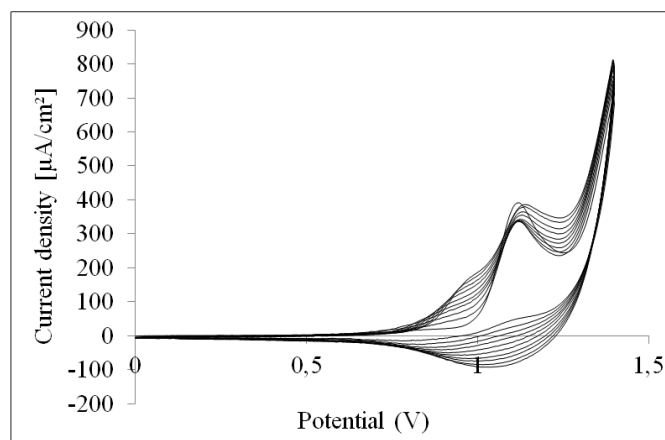


Figure 3. Electropolymerization of SIC (5×10^{-3} M) in Bu_4NBF_4 (0.1 M)/MeCN; 10 sweeps between 0V and 1.4V; $\nu = 100$ mV/s; working electrode: GCE; reference electrode: Ag/AgCl

3.2. Electrochemical polymerization of SIC in LiClO_4 (0.1M)/MeCN in the presence of HClO_4

In an acidic medium, the imine bond undergoes a protonation which commonly enhances the planarity of the polymeric material and subsequently, improves its electrical properties. A preliminary study of the **SIC** electropolymerization was carried out to specify the appropriate concentration of HClO_4 to add in the electrolytic solution. The current densities values at p-doping/undoping potential of **PDIC** polymer ($E_{1/2} = 1.02\text{V}$) versus HClO_4 concentration (Figure4) showed a maximum at 35mM. Beyond this value, the current densities decreased, proving that a stronger acid concentration would limit the growth of **PDIC** polymer [9].

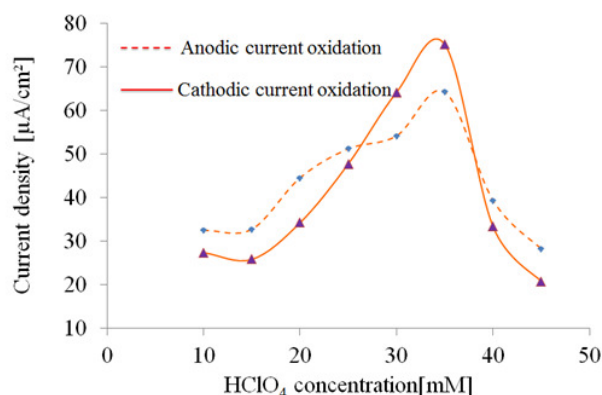


Figure 4. Influence of HClO_4 concentration on current density measured at p-doping/undoping potential of **PDIC**. (----) Anodic current density, (—) Cathodic current density.

The species mobility which increases with temperature increasing is another factor which can improve a potentiodynamic polymerization. However, this increasing is not always favourable, and can influence the electropolymerization process providing short, no-structured and less conductive chains.

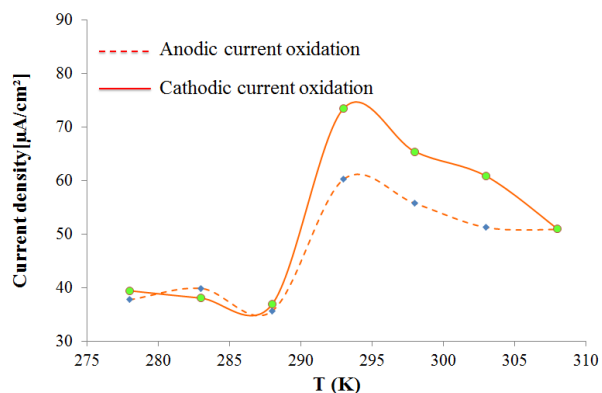


Figure 5. Influence of temperature on current density values measured at p-doping/dedoping potential of **PDIC**. (----) Anodic current density, (—) Cathodic current density.

Figure 5, represents the current densities values recorded at doping-undoping potential ($E_{1/2} = 1.02\text{V}$) versus the temperature, varied from 5 to 30°C. The anodic and cathodic currents intensities

increase simultaneously and reach a maximum at 20°C. Beyond this value, the current densities decreased, and the polymer becomes less electroactive. The figure 6 presents the cyclic voltammogram obtained during electrooxidation of **SIC** in LiClO₄ (0.1M)/MeCN + 35mM HClO₄. It exhibits two peaks with unequal intensities. The first one was recorded at 0.98V and was reversible, while the second one which was observed at 1.32V was irreversible. On another hand, SWV's voltammogram (Figure 7) shows three oxidation peaks at 0.93, 1.24 and 1.44V. The first peaks, as on CV's voltammogram, was assigned to the oxidation of the hydroxyl function, while the second and third ones, combined in one peak in CV (the second peak), were related to the oxidation of imine and carbazole moieties.

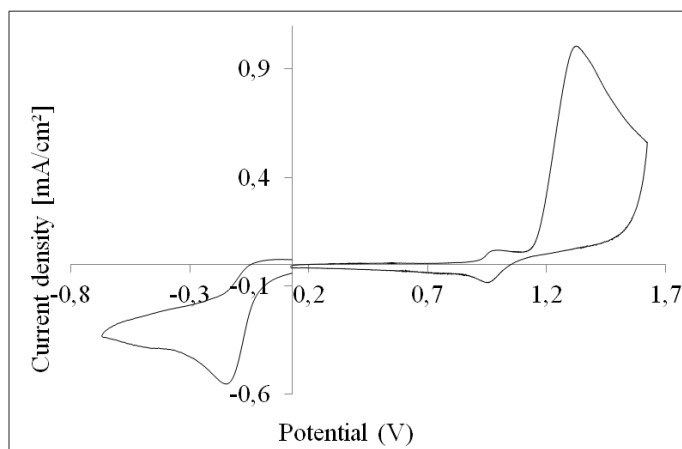


Figure 6. Cyclic voltammogram of **SIC** 5×10^{-3} M in LiClO₄ (0.1 M)/MeCN + 35mM HClO₄; $v = 100$ mV/s; working electrode: GCE; reference electrode: Ag/AgCl

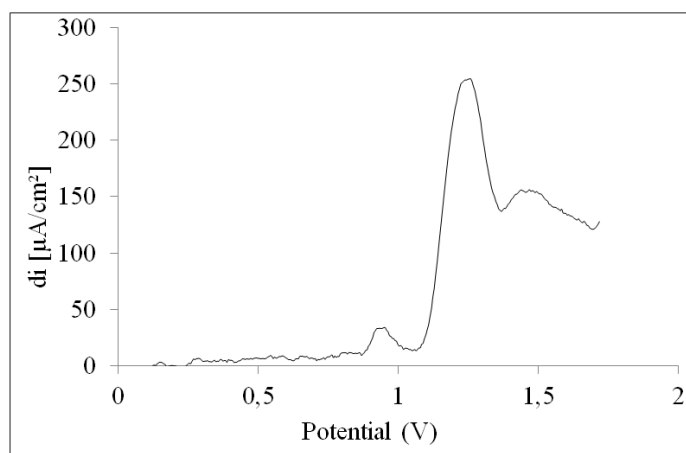


Figure 7. Square wave voltammogram of **SIC** 5×10^{-3} M in LiClO₄ (0.1 M)/MeCN + 35mM HClO₄; $v = 100$ mV/s; working electrode: GCE; reference electrode: Ag/AgCl

The electrochemical polymerisation of **SIC** performed in protic medium (Figure 8a) led to the growth of a new redox system centred at lower potentials (around 1.02 V) comparing to the monomer. The electrochemical behaviour recorded in this medium was different from the one recorded in aprotic solvent. The oxidation of the phenol moiety in acidic medium occurs at 0.98V, and

leads to its hydrolysis resulting in the formation of two products, *para*-quinone and *ortho*-quinone (scheme 2)[6]. The both compounds showed a reversible electrochemical behavior with two electrons and two protons transfer (scheme 3)[6]. On the first negative scan (Figure 8b), the two very weak reduction peaks recorded at 0.39 and 0.67V were attributed to the reduction of the two 2-substituted phenol oxidation products, while the peak at 0.85V corresponding to the OH oxidation. In figures 8a and 8b, E_{Red2} corresponds to the reduction of *ortho*-(2-substituted)quinone in *ortho*-(2-substituted)catechol, whereas E_{Red1} corresponds to the reduction of *para*-(2-substituted)quinone in *para*-(2-substituted)hydroquinone. On the second positive scan (Figure 8b), two anodic peaks were recorded at $E_{Ox1} = 0.5V$ and $E_{Ox2} = 0.72V$, showing the reversibility of the two reductive peaks of quinones.

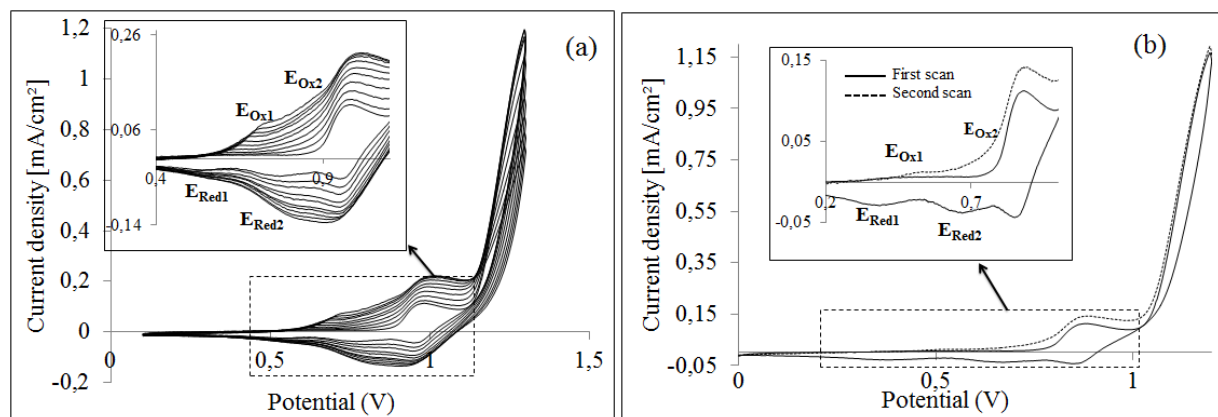


Figure 8. Electropolymerization of SIC (5×10^{-3} M) in $LiClO_4$ (0.1 M)/MeCN + 35mM $HClO_4$; $v = 100$ mV/s; between 0 and 1.29 V; working electrode: GCE; reference electrode: Ag/AgCl a) 10 sweeps; b) First and second sweep.

Therefore, in the protic medium, the monomer (SIC) undergoes hydrolysis around 1V, which modifies its structure before its polymerization. This reaction which is very fast, leads to the formation of two isomers, *para*-quinone and *ortho*-quinone which oxidizes at the same potential to give PDIC. Therefore, the theoretical calculations showed that the two isomers *para*-quinones and *ortho*-quinones, have practically the same HOMO value (-5.46 and -5.4eV, respectively), and a very near LUMO values (-3.41 and -3.53eV, respectively) (Figure 9).

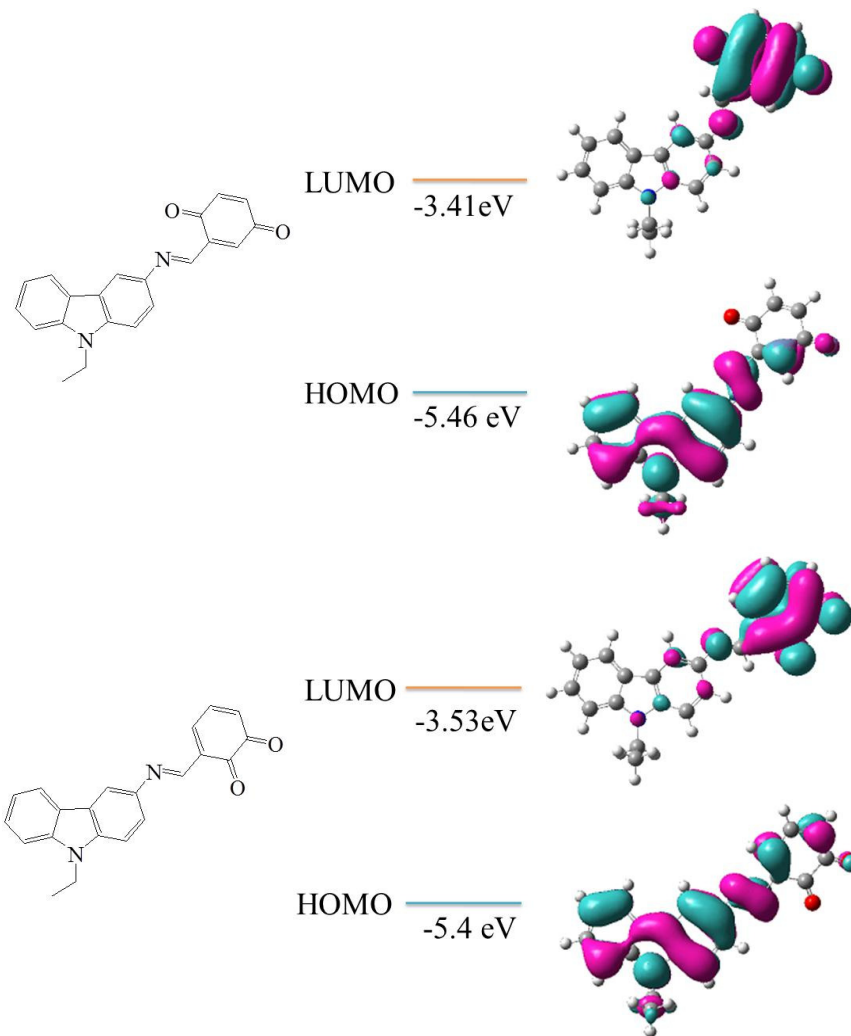


Figure 9. The contour plots and energy values of HOMO and LUMO orbitals of *para*-quinone and *ortho*-quinone compounds.

The variation of the first peak potential of SIC with the logarithm of sweep rate : $E_{pa} = f(\log v)$ (Figure 10) provides a straight line with zero slope value, confirming that the oxidation of hydroxyl function occurred rapidly.

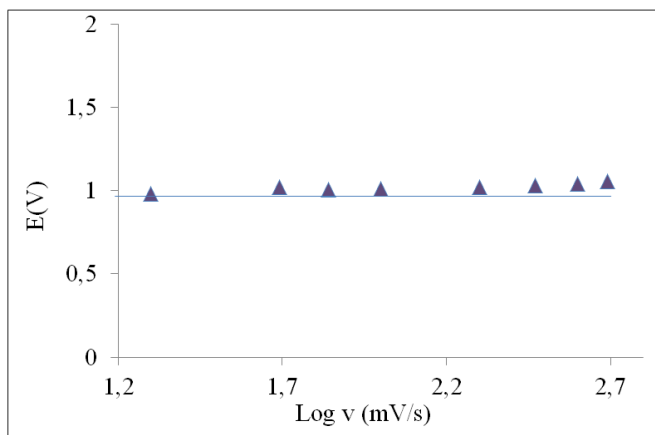
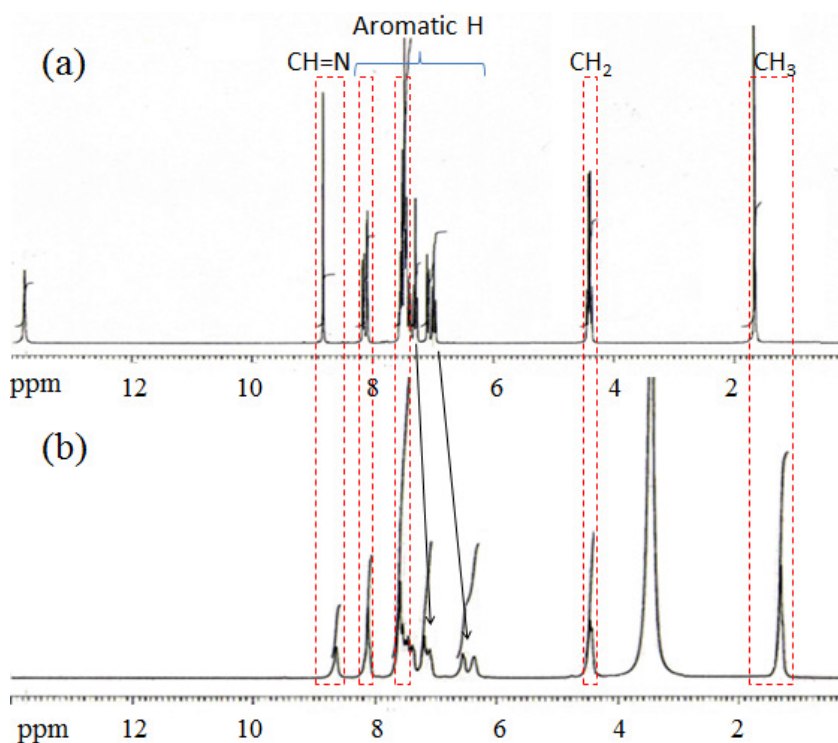


Figure 10. Variation of the potential of the first oxidation peak of **SIC** as a function of the logarithm of sweep rate: $E_{pa}=f(\log v)$, on GCE in LiClO_4 (0.1 M)/MeCN + 35mM HClO_4 . Scan rates from 20 to 500 mV/s.

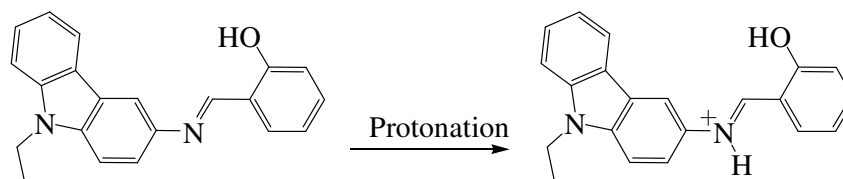
To confirm that the nature of the compounds formed was due to the hydrolysis of hydroxyl function, a large-scale electrolysis of **SIC** at controlled potential value of 0.98V was carried out. A nacre white product was formed immediately, which once again **proves** the velocity of the reaction. The ^1H NMR spectrum (Figure 11) of the obtained precipitate, denoted DIC, is in accordance with the disappearance of the OH function after oxidation. Moreover, the two peaks of phenolic ring protons recorded between 6.98 and 7.37ppm before oxidation, have moved to smaller values namely 6.3 to 6.5ppm, reflecting therefore the **environment** change of the corresponding protons.



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Figure 11. ^1H NMR spectrum of a) **SIC**, b) **DIC**.

176 Furthermore, the color change that occurred when we dissolve **SIC** in MeCN with
 177 $\text{LiClO}_4(0.1\text{M})/35\text{mM HClO}_4$, allowed us to think that **SIC** underwent a chemical protonation. Treated
 178 with a solution of HClO_4 (35mM) in MeCN, **SIC** changes its color from yellow to orange and the FTIR
 179 analysis showed the appearance of new bands (Figure 12) of which the signal recorded at 1229 cm^{-1}
 180 was assigned to the vibration of the group C-N^+ [10]. Thus, we can presume that **SIC** underwent a
 181 protonation before its electropolymerization (Scheme.1).



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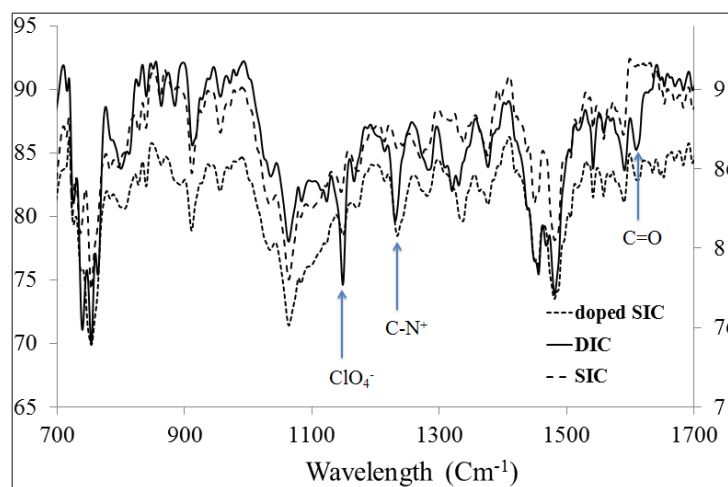
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Scheme.1 Protonation of **SIC**

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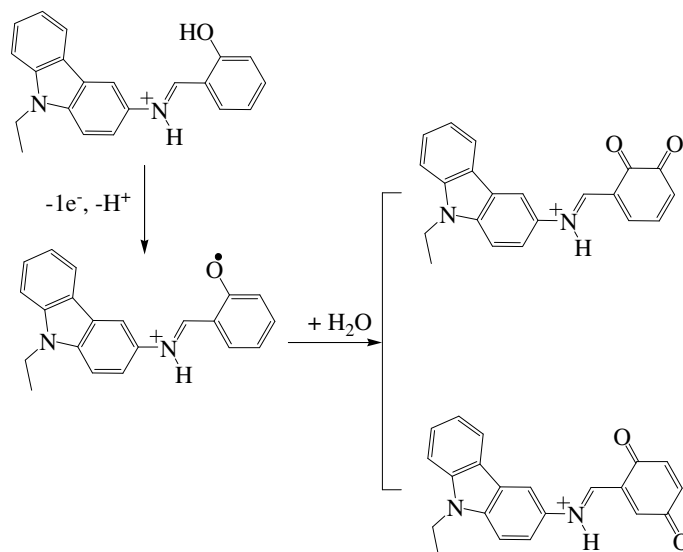
From these data, we can say that the **SIC** monomer changes its structure in the protic medium and becomes a mixture of two isomers *para*-quinone and *ortho*-quinone (**DIC**).



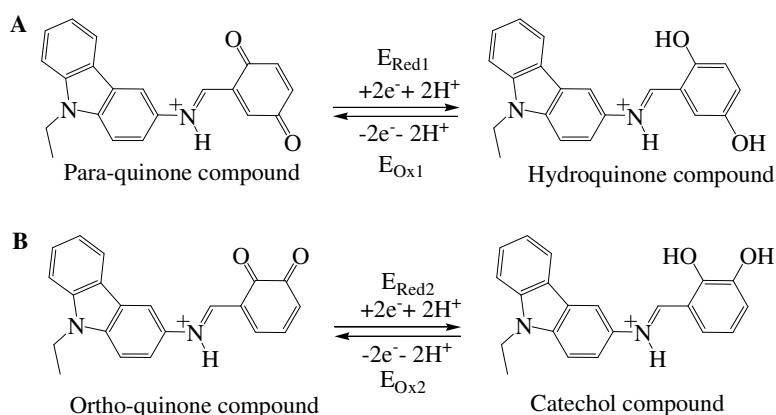
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Figure 12. IR spectra of **SIC** (---), **DIC** (—) and protonated **SIC** (·····).



Scheme 2.First oxidation mechanism of SIC.



Scheme 3. Redox behaviour of: A) *Para*-quinone compound, B) *Ortho*-quinone compound.

3.3. Polymers electroactivity

The study of electrodeposited films of **PSIC** and **PDIC** polymers in both electrolytic media was done in another cell containing only the electrolytic solutions, without monomer. During the positive scans, in the two media (Figure13), voltammograms show that the redox behaviour corresponds to the positive doping/undoping process, due to the participative accumulation of anionic charges, BF_4^- and ClO_4^- respectively in the polymeric matrices, pointing out the electroactive nature of the formed materials[11]. Otherwise, in the protic medium, the redox peaks appearing at $E_{1/2} = 0.61\text{V}$ was related to the redox moiety of *para*-quinone / hydroquinone [6] in the polymer matrices. However, the redox system corresponding to *ortho*-quinone/catechol moiety was overlapped with the one of **PDIC**. The electrochemical behaviours of **PSIC** and **PDIC** affirm the difference between their structures.

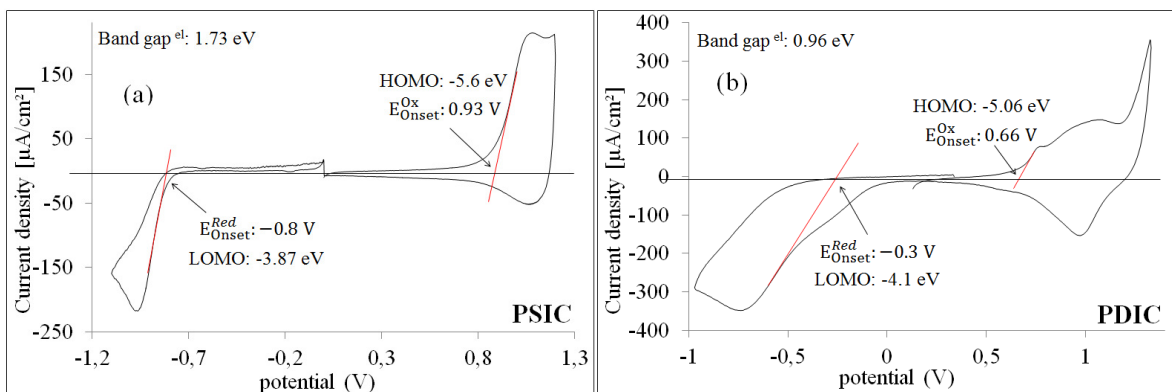


Figure 13. Cyclic voltammograms for oxidation of: a) **PSIC** coated GCE in Bu_4NBF_4 (0.1 M)/MeCN; b) **PDIC** coated GCE in LiClO_4 (0.1 M)/MeCN + 35mM HClO_4 . $v = 100\text{mV/s}$.

The thin layer film of both polymers was analyzed by XPS, a much more sensitive measurement than IR. The XPS spectra of the two polymers coated ITO, are represented on figure 14. Either for **PSIC** or **PDIC**, the band at 399.9eV corresponds to the N_{1s} signal, while the one at 532.5eV corresponds to the signal of O_{1s} [12]. In the spectrum of **PSIC**, there are two bands in the C region (Figure15), corresponding to C-C (284.7eV) and C-OH (286.5eV) [13]. However, in the spectrum of **PDIC**, we have the band of the C-C at 284.7eV, accompanied by another one at 288.2eV corresponding to the C=O energy (Figure16) [14].

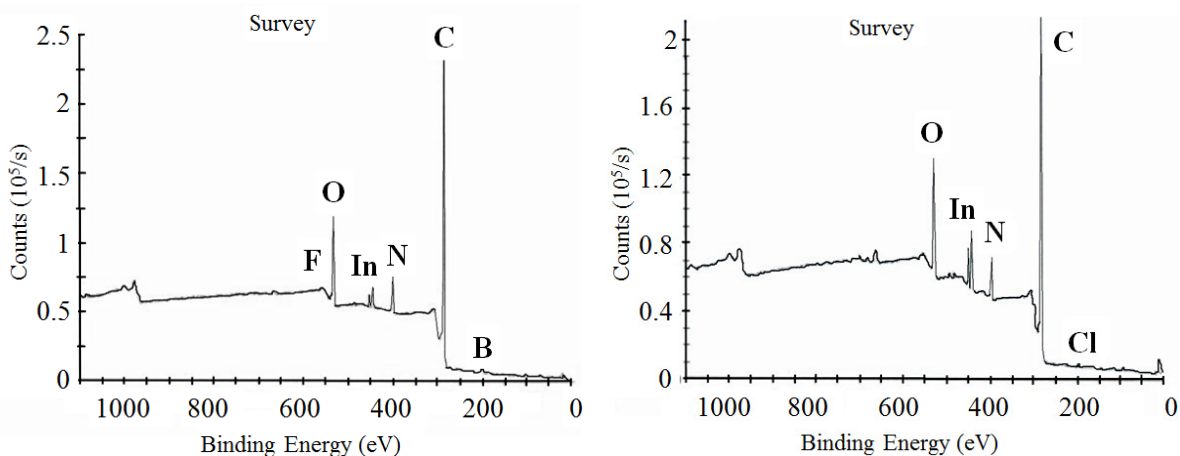


Figure 14. Survey spectra of : a) **PSIC** coated ITO, b) **PDIC** coated ITO.

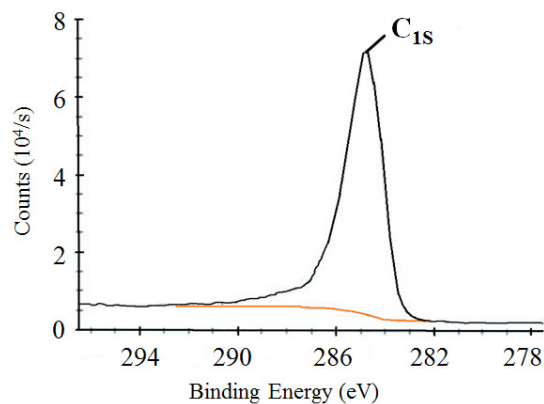


Figure 15. Survey spectra of PSIC-coated ITO.

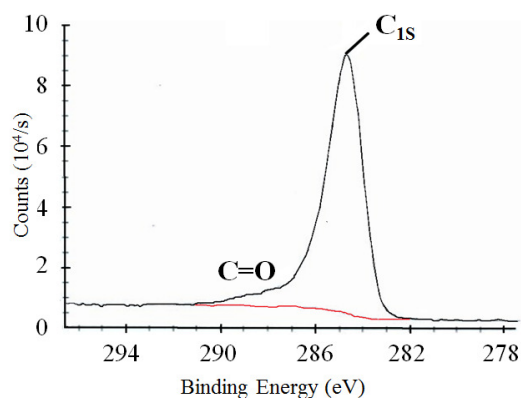


Figure 16. Survey spectra of PDIC-coated ITO.

3.4. Electrochemical impedance measurements (EIS)

The electrochemical impedance spectroscopy permits to understand the charge transport mechanism of electrolyte ions during the redox reaction. Several works have been focused on the mass transport in conducting polymers. However, due to the complexity of these phenomena, their complete understanding is not achieved yet. The impedance measurements were conducted at a potential corresponding to the oxidation state of each polymer (around 1V) while varying the frequency from 0.1 to 100kHz. From the impedance response, the thin film coated electrode can be represented from electrical point of view by a Randles-type equivalent circuit [15].

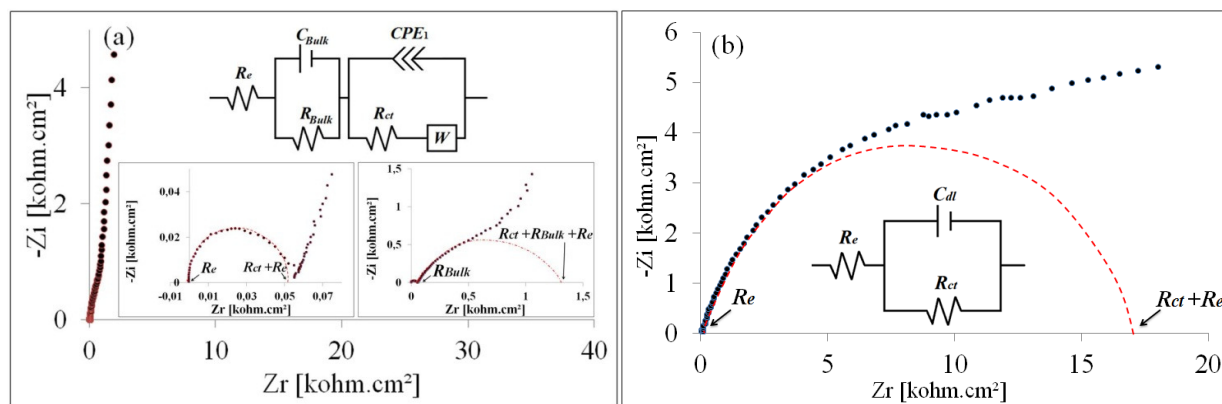


Figure 17. Nyquist plots and Randles-type equivalent circuits of: a) **PDIC**, b) **PSIC**.

Figure 17a shows the nyquist plot of **PDIC** coated GCE in LiClO_4 (0.1 M)/MeCN + 35mM HClO_4 , while Figure 17b shows **PSIC** coated GCE in Bu_4NBF_4 (0.1 M)/MeCN. For **PDIC** coated GCE, there is three characteristic parts of the diagram. At high frequencies, a semicircle characteristic for charge transfer in polymer bulk is observed, and represented in equivalent circuit by parallel combination of film resistance R_{Bulk} and capacitance C_{Bulk} in series with solution resistance R_e [16]. At intermediate frequencies, a beginning of a second semicircle is recorded. The impedance response depends on the shapes of pores in the surface (theory of porous electrodes developed by *de LEVIE*) [17]. The more the pore is easily accessible, the more the semicircle tends to disappear and to be replaced by a straight line. Compared to the first one, the second semicircle is flattened, because the film-electrolyte interface is not smooth and uniform [18]. Consequently, the double layer capacitance in the equivalent circuits is replaced by a constant phase element (CPE) [18]. The impedance response is represented in equivalent circuit, by a parallel combination of charge transfer resistance R_{ct} (between film and solution) and pseudo-capacitive charging of the film CPE_1 [16]. At low frequencies, a straight line representing Warburg-type impedance due to a finite diffusion of counter-ions species within the film is observed. This line is recorded with a slope $\theta \neq 45^\circ$, as often found in the presence of inhomogeneities in a coated material, or under non-uniform diffusion conditions [19].

For the **PSIC** coated GCE, the diagram showed only a beginning of a semicircle, characteristic for a charge transfer between polymer and solution, and represented in equivalent circuit by parallel combination of the charge transfer resistance R_{ct} and the interfacial capacitance of the film C_{dl} . Thus, there is a pure charge transfer control and no diffusion limitations [20,21].

The charge transfer resistance recorded for **PDIC** film was much lower than the one recorded for **PSIC** film, which implies that **PDIC** is more conductive than **PSIC** [22].

3.5. Optical and electrochemical bandgap

The electrical properties of conjugated polymers depend on their structures and on the delocalization of π electrons that is related to the planarity of the chain. Thus, due to its protonated state, **PDIC** would be more planar than **PSIC** and then, would exhibit better electrical properties.

Otherwise, the bandgap is another important factor determining the electrical conductivity. The bandgap can be measured in two different ways, electronic bandgap determined from cyclic voltammetry and optical bandgap estimated from UV-Visible spectroscopy.

By cyclic voltammetry, the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) energy levels could be estimated from the onsets of oxidation and reduction potentials, respectively, according to the following equations [23]:

$$E_{\text{HOMO}} (\text{eV}) = - [E_{\text{onset}}^{\text{ox}} + 4.4]$$

$$E_{\text{LUMO}} (\text{eV}) = - [E_{\text{onset}}^{\text{red}} + 4.4]$$

The difference between HOMO and LUMO energies, give the electrochemical bandgap value (E_g^{el}). The obtained results were presented in Figures 13a and 13b and summarized in Table 1.

The electrochemical bandgap recorded for **PDIC** is very smaller than the one recorded for **PSIC** (0.96 and 1.73 eV, respectively). This value which reflects the conductivity of the material was related to the doping level. Indeed, when a polymer is oxidized, by analogy of the admitted results, an electron is removed from the HOMO and a positive polaron or radical cation is created. The energy level associated with that positive polaron, partially relocated on monomer units of the polymer chain, is represented by a destabilized bonding orbital of which the energy is higher than that of the HOMO. Furthermore, the protonation of imine moiety, which was preferably compared with that of the amine sites [24], increases planarity of polyazometines [25], which decreases the LUMO level and increases the HOMO level. Subsequently, the bandgap is decreased. Hence, it is clear that in the medium containing LiClO_4 (0.1M) / MeCN + 35mM HClO_4 , **PDIC** underwent protonation and electrochemical doping, which can explain its low energy bandgap.

Optical properties of the polymer films on ITO glass were observed by recording the changes in the absorption spectra under voltage at 0 and 1V. The electronic absorption of neutral films exhibit an absorption band around 355 and 305 nm for **PSIC** and **PDIC**, respectively. These bands which were due to $\pi \rightarrow \pi^*$ transition lose intensity, during oxidation.

Polarized at 1V in LiClO_4 (0.1M)/MeCN/ HClO_4 (35mM), **PDIC** thin layer presents a modification in the optical spectrum with the appearance of a broad band around 605nm (Figure 18). In

a similar way, when **PSIC** thin layer is polarized at 1V, the optical spectrum shows a new absorption band at 477nm (Figure19).

The optical bandgap is calculated according to $E_g^{opt} = 1240/\lambda_{onset}$, where λ_{onset} is obtained from the intersection between the baseline and the tangent of the UV-Vis band-end[23].

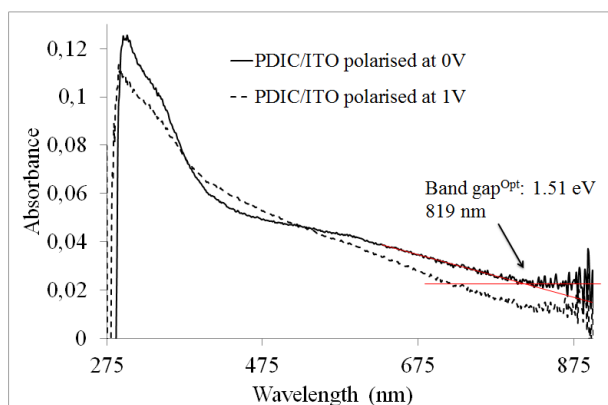


Figure 18. The absorbance versus the wavelength registered with **PDIC**coated ITO polarized in the LiClO_4 (0.1M) / MeCN + (35mM) HClO_4 , at: 0V (—) and 1V (---).

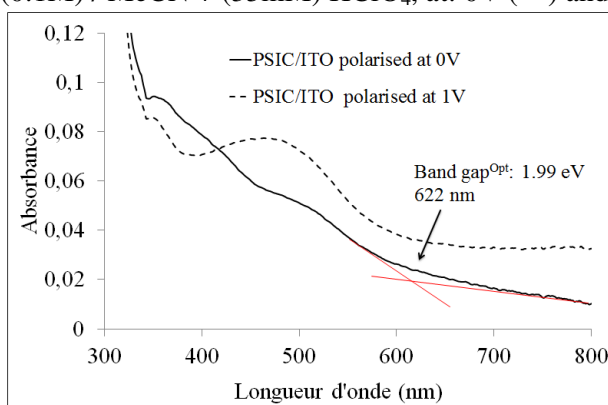


Figure 19. The absorbance versus the wavelength registered with **PSIC**coated ITO polarized in the LiClO_4 (0.1M) / MeCN + (35mM) HClO_4 at: 0V (—) and 1V (---).

Table 1. Electrochemical bandgaps.

	E_{onset}^{ox} (vs. Ag/AgCl) ^a	E_{onset}^{red} (vs. Ag/AgCl) ^a	E_{HOMO} (eV)	E_{LUMO} (eV)	E_g^{el} (eV) ^b	E_g^{opt} (eV) ^c
SIC	1.20	-0.96	-5.87	-3.71	2.16	2.84
PSIC	0.93	-0.80	-5.60	-3.87	1.73	1.99
DIC	1.14	-0.01	-5.54	-4.39	1.15	-----
PDIC	0.66	-0.30	-5.06	-4.10	0.96	1.51

^a The onset potentials are determined from the intersection of the two tangents drawn at the rising current and the baseline charging current of the CV traces. ^b Energy of the bandgaps calculated from the difference between the energy of the HOMO and the LUMO. ^c The optical bandgap is calculated according to $E_g^{opt} = 1240/\lambda_{onset}$

The optical bandgap of **PDIC** (1.51eV) is lower than that of **PSIC** (1.99eV). The bandgap value can be affected by several factors such as an intra-chain charge transfer and substituents effect [26]. Figure 21 shows the (DFT) theoretical calculation for the distributions of the frontier molecular orbitals of **SIC**. Compared to **SIC**, *para*-quinone and *ortho*-quinone compounds are formally donor-acceptor (D-A) type moieties. Phenol oxidation in **SIC** leads to an electronic deficient state of the quinone moiety of the formed molecules (*para*-quinone and *ortho*-quinone), while their carbazole moieties are electrons donor. Indeed, in a D-A system, a charge transfer for the donor to the acceptor increases the length of the π -system through resonance. The increasing in the conjugation length of a molecule, causes the decrease of its bandgap[27].

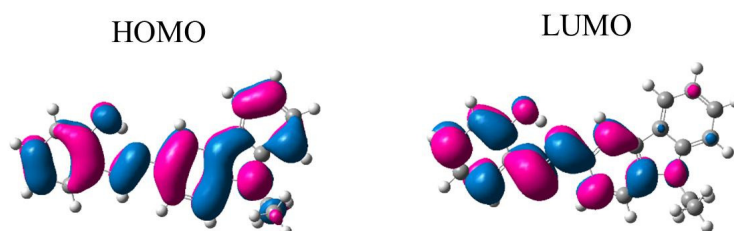


Figure 20. The contour plots and energy values of HOMO and LUMO orbitals of **SIC**.

The difference in the surface topography was observed by the atomic force microscopy working in the contact mode (Figure21). The 3D view shows that there are numerous smaller surface undulations with a precipitation of different shapes [28]. However, we notice that precipitations in **PDIC** are not as numerous as in **PSIC** thin film.

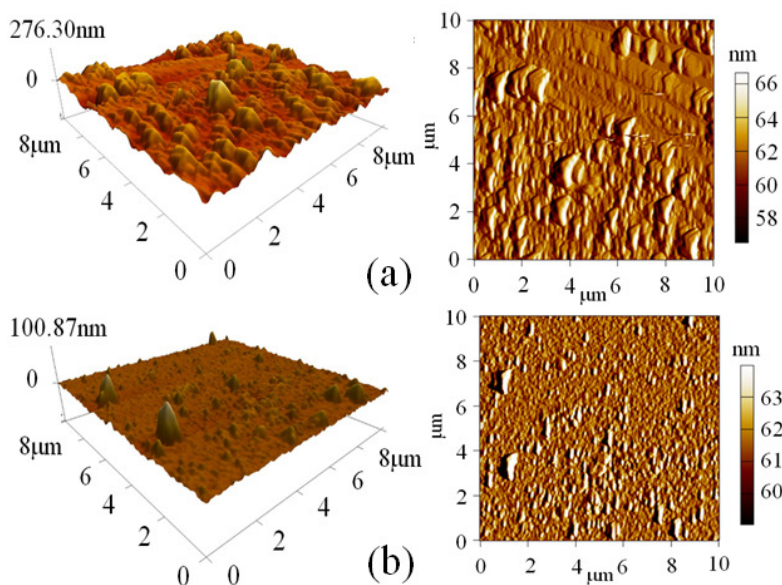


Figure 21. AFM image of: a) **PSIC** thin layer, b) **PDIC** thin layer.

Conclusion

The electrochemical polymerisation of **SIC** is performed in both protic and organic medium. During the anodic oxidation in protic medium containing $\text{LiClO}_4/\text{MeCN} + (35\text{mM}) \text{HClO}_4$, **SIC** undergoes hydrolysis around 1V, which modifies its structure before its polymerization. This reaction is very fast, and leads to the formation of two isomers, *para*-quinone and *ortho*-quinone (**DIC**) which oxidizes at the same potential and form **PDIC**. In the organic medium, the **SIC** retains its initial structure, and leads after oxidation to the corresponding polymer **PSIC**. The electrochemical behaviour of **PSIC** and **PDIC** supports the conductive character of both polymers and the difference between their structures. In an acidic medium, the imine bond undergoes a protonation which increases commonly the planarity of the polymeric material and subsequently, improves its electrical properties. The charge transfer resistance recorded for the **PDIC** film was much lower than the one recorded for the **PSIC** film. The electrochemical and the optical bandgaps of **PDIC** are lower than those of **PSIC**. The difference in structures of both polymers could explain these results. Compared to **SIC**, *para*-quinone and *ortho*-quinone compounds are formally electron donor-acceptor (D-A) type moieties. Phenol oxidation in **SIC** leads to an electronic deficient state of the quinone moiety of the formed molecules (*para*-quinone and *ortho*-quinone), while their carbazole moieties are electron donor. Indeed, in a D-A system, a charge transfer for the donor to the acceptor increases conjugation length through resonance. The increasing of the conjugation length of a molecule causes the decrease of its bandgap. Therefore, these findings indicate that **PDIC** is more conductive than **PSIC**.

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