



HAL
open science

Affinity and Distribution of Silver Nanoparticles Within Plasma Polymer Matrices

Ali Mansour, Fabienne Poncin-Epaillard, Dominique Debarnot

► **To cite this version:**

Ali Mansour, Fabienne Poncin-Epaillard, Dominique Debarnot. Affinity and Distribution of Silver Nanoparticles Within Plasma Polymer Matrices. *Journal of Materials Science*, 2019, 54 (19), pp.12972-12987. 10.1007/s10853-019-03772-6 . hal-03016391

HAL Id: hal-03016391

<https://hal.science/hal-03016391>

Submitted on 20 Nov 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Affinity and Distribution of Silver Nanoparticles Within Plasma Polymer Matrices

A. Mansour, F. Poncin-Epaillard, D. Debarnot*

Institut des Molécules et Matériaux du Mans (IMMM) - UMR 6283 CNRS, Le Mans
Université, Avenue Olivier Messiaen, 72085 Le Mans Cedex 9, France

*Corresponding author: Dominique.Debarnot@univ-lemans.fr

J. Mat. Sci., 54(19), 12972-12987 (2019)

Abstract

The affinity and repartition of silver nanoparticles in different plasma polymer matrices are studied according to the polymer functional groups. The polymer layers are synthesized by plasma polymerization of allyl alcohol (AAI), aniline (ANI), or heptadecafluoro-1-decene (HDFD). The matrix is then impregnated in a metallic salt solution and a reducing one. When plasma polymerized (pp) ANI is used as matrix, a higher amount of silver nanoparticles (AgNPs) is introduced compared to the other two polymer matrices. Otherwise, with pp-AAI, smaller AgNPs with narrower size distribution are inserted. Finally, different plasma polymer/AgNPs interaction mechanisms are proposed according to the polymer functional groups. Moreover, the sensing behavior of pp-ANI, well-known as sensitive layer towards ammonia, is markedly enhanced by the presence of AgNPs.

Keywords: hybrid nanocomposites; plasma polymer; silver nanoparticles; metal distribution; polymer/metal interaction.

1. Introduction

Polymer/metal nanocomposites have attracted considerable attention as they combine the advantages of both polymer and metal offering interesting properties in many fields such as electronics, photonics, sensing, catalysis, biology and medicine.¹⁻⁵

Among the several approaches to elaborate such hybrid nanocomposite materials, the generation of metal nanoparticles from a metal precursor in the polymer, considered as a simple approach, is carried out by mixing the polymer with the metal precursor followed by different physical (photoirradiation, thermal treatment) or chemical treatments (chemical reduction by hydrazine, sodium citrate, or sodium borohydride).^{2,6} In a lot of studies, the polymer matrix is synthesized by conventional ways (chemical or electrochemical ones).⁷⁻¹⁰

Recently, plasma polymerization is gaining more and more attention for its ecofriendly process (e.g., solvent free). Indeed, plasma polymerization is initiated by collisions of the monomer molecules (many organic and inorganic vapors) with free electrons accelerated by the electric field. It results in the formation of excited species, radicals and ions. Different reactions (recombination, addition...) will then be held between these active species to form the plasma polymer leading to a more irregular structure compared to conventional polymers. However, a great resemblance to conventional polymers can be obtained after optimizing plasma parameters such as discharge power, duty-cycle, frequency, etc. A large range of polymers with various functional groups can be obtained by plasma polymerization.

Different plasma polymer/metal composite materials have been synthesized by the approach mentioned before, using n-heptylamine,¹¹ acrylic acid,¹² allyl alcohol,¹³ or maleic anhydride¹⁴ as monomers, silver nitrate (AgNO_3) as metallic solution, and sodium borohydride (NaBH_4) solution as silver reducing agent. The purpose of these studies concerns the antibacterial application, and no intensive investigation on the role of the polymer nature on the silver nanoparticles (AgNPs) formation was realized. However, the properties of the composites are

highly dependent on the distribution of the metal particles in the polymer matrix. Therefore, the objective of this work is to study the silver nanoparticles formation: amount, size, size distribution and oxidation state according to the chemical composition of the polymer matrix, in order to better understand the interaction mechanisms between metal ions and the plasma polymer. For this purpose, different plasma polymer layers with different functional groups are chosen containing nitrogen, oxygen or fluorine functions using aniline (ANI), allyl alcohol (AAl) or heptafluoro-1-decene (HDFD) as precursors respectively. The formation of silver nanoparticles into these matrices is carried out by impregnation in AgNO_3 metallic solution followed by a chemical reduction in NaBH_4 .

Ammonia (NH_3) sensing is one of the most attractive applications due to its large production and toxicity.¹⁵⁻¹⁷ Indeed, numerous studies have recently been devoted to the development of polymer-based gas sensors for NH_3 detection, specially polyaniline well-known for its selectivity to ammonia.¹⁸⁻²¹ Moreover, an improved sensing performance of conventional polymers towards different toxic gases is shown by incorporating metals such as Cu, Ag, Pt, Pd etc.²²⁻²⁵ Polyaniline/AgNPs composites have then been prepared for the detection of ammonia both in solution based on the change in color of the films²⁶ and in gas phase based on the change of electrical or optical properties of the layers^{27,28}. In these works, polyaniline is synthesized by conventional ways and no real investigation is carried out in order to understand how these metals could improve sensing performance. In our work, polyaniline is elaborated by the plasma technique with the purpose to demonstrate how the metal nanoparticles can improve the ammonia gas sensing. Therefore, the final part of this work concerns the study of the ammonia sensitivity, evaluated by absorbance variation, of plasma polymerized aniline (pp-ANI) containing silver nanoparticles compared to bare pp-ANI.

2. Experimental

2.1. Materials and methods

Aniline (mol. wt = 93.13 g.mol⁻¹, purity ≥ 99.5%), allyl alcohol (mol. wt = 58.08 g.mol⁻¹, purity ≥ 98.5%), and heptadecafluoro-1-decene (mol. wt = 446.1 g.mol⁻¹, purity = 99%) monomers were obtained from Sigma-Aldrich and used as received. The structure of the monomers is given in Figure 1. Silver nitrate (AgNO₃) (mol. wt = 169.87 g mol⁻¹, Fisher), sodium borohydride (NaBH₄) (mol. wt = 37.83 g mol⁻¹, purity = 96%, Sigma Aldrich), and doubly distilled water (Millipore, resistivity 18 MΩ.cm) were used to prepare the aqueous solution of the metal salt and the reducing agent, respectively.

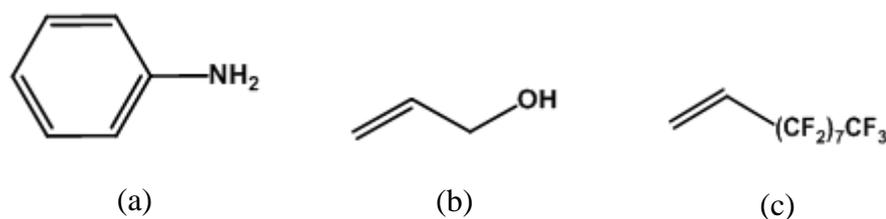


Figure 1. Chemical structures of the monomers (a) aniline, (b) allyl alcohol and (c) heptadecafluoro-1-decene.

2.2. Incorporation of silver nanoparticles into the different plasma polymer films

The plasma polymers' synthesis and deposition were achieved in a cylindrical aluminum reactor capacitively coupled with two parallel circular electrodes with a R.F. (13.56 MHz) generator. The adhesion of the plasma polymer layers to the different types of substrates was successfully enhanced by argon plasma pre-treatment ($P = 100$ W, $F = 10$ sccm, $t = 1$ min). The plasma polymerization of ANI, AAl and HDFD was realized in pulsed mode using plasma parameters optimized to a power (P) equal to 30 W, a duty cycle ($D.C.$) of 50% (defined as $D.C. = \text{discharge time } (t_{on}) / (\text{discharge } (t_{on}) + \text{post-discharge } (t_{off}) \text{ times})$), and a frequency of 30 kHz. In order to avoid thickness dependence, different deposition times (8, 20, and 5 min, respectively for the polymerization of ANI, AAl and HDFD) were judiciously selected to obtain the same thickness

of about $\sim 285 \pm 5$ nm. Then, all these polymers were successively dipped in 0.1 M AgNO₃ for 3 h and 0.1 M NaBH₄ aqueous solutions for 1 h at room temperature under continuous and constant stirring in a dark place. The concentrations and dipping times of the silver salt and the reducing agent have been optimized in a previous work.²⁹ Between each step, the composites were rinsed with distilled water and dried with air.

2.3. Characterization techniques

UV–Vis spectroscopy is known to be the most used technique for metal nanoparticle characterization. UV-Vis absorption measurements were performed with Varian Cary 100, in the wavelength range of 200–900 nm, using glass slides as substrates. The spectra were measured in a double beam mode and the reference was carried out with a glass slide.

The silicon wafers, used for determination of the layer thickness via atomic force microscopy (AFM, Bruker-Innova), were preventively cleaned and partially masked with a stable adhesive tape during the plasma deposition. AFM operating in the tapping mode was also used to examine the topography and roughness of the polymer and composite thin films. The root mean square roughness (Rq) values were derived from (10x10 μm) AFM images using the Gwiddion software.

The metal particle size, size distribution, and metal diffraction patterns were determined by the transmission electron microscopy (TEM) micrographs coupled with selected area electron diffraction (SAED). These characterizations were carried out on a JEOL (STEM 2100) microscope operating at 200 kV. The samples must have a thickness lower than 100 nm and deposited on nickel grids (Formvar/carbon film, 200 mesh, Ni, USA / Canada).

The FTIR-ATR spectra were recorded in the absorbance mode in the range of 400–4000 cm^{-1} with a resolution of 2 cm^{-1} using 32 scans. The spectrometer was a Bruker-vertex 70V apparatus with a globar light source and a DTGS detector. Polypropylene (PP) (Goodfellow Production,

England) was used as substrate for the deposition of plasma polymers and plasma polymer/metal nanocomposites. All the spectra were normalized with respect to thickness. The spectra were measured in the Attenuated Total Reflectance (ATR) technique using diamond. The sampling depth was in the range of about 1.66 μm at 1000 cm^{-1} and 45° as angle of incidence.

XPS determination of surface structures deposited onto silicon wafers (100) (Neyco s.a., France) was carried out on an Axis Nova (Kratos) instrument at the Institut des Matériaux de Nantes, France. The emission was analyzed at a take-off angle of 90° relative to the sample surface, yielding a sampling depth of maximal 10 nm due to the mean free path of the electrons. The curve fitting was performed using CasaXPS software (Casa Software Ltd.). The peak shape was chosen with Gaussian (70%)/Lorentzian (30%) curve fitting and quantification was reliable to $\pm 5\%$.

2.4. Gas sensing

The *ex-situ* (out of the plasma reactor) doping with iodine has been used in this work to obtain NH_3 sensitive thin films. It consists of introducing the polymer and nanocomposite films into a chamber saturated with iodine vapours (I_2 , Aldrich) during 24 h.

The optical ammonia gas sensing system (Figure 2) is composed of a sealed measuring chamber inserted into the UV-Vis spectrophotometer, a gas dilution system with flowmeters and gas cylinders, and a computer for data collection. The gas sensitive films (doped pp-ANI or doped pp-ANI/AgNPs) deposited onto glass substrates were introduced into the sealed chamber. A stream of nitrogen gas containing ammonia was then passed through the sensitive film and spectroscopic measurements were performed. The concentration of NH_3 was varied by mixing different flows of NH_3 and 500 sccm of N_2 . The concentration of NH_3 (ppm) was defined as the ratio of the flow rate of NH_3 to the total flow rate of NH_3 and N_2 . In this work, the ammonia

concentration varied between 92 and 4618 ppm. The ammonia concentrations lower than 92 ppm were not measured due to experimental set-up limitation. The interaction between NH_3 and the sensitive film leads to its optical absorbance variation. When the absorbance variation tends to a constant value depending on time, the NH_3 gas was turned off and stream of pure N_2 was passed through the sensor to purge completely the NH_3 molecules in the measuring chamber and to regenerate the sensitive film. The wavelength of the light source used in this work was fixed at 430 nm. All experiments were performed at room temperature. Sensitivity (S) was calculated as $(A-A_0)/A_0$ ratio, where A_0 is the initial optical absorbance of the sensor under N_2 and A, the absorbance of the sensor when exposed to ammonia gas. The sensitivity value given for one concentration is the average of at least three measurements.

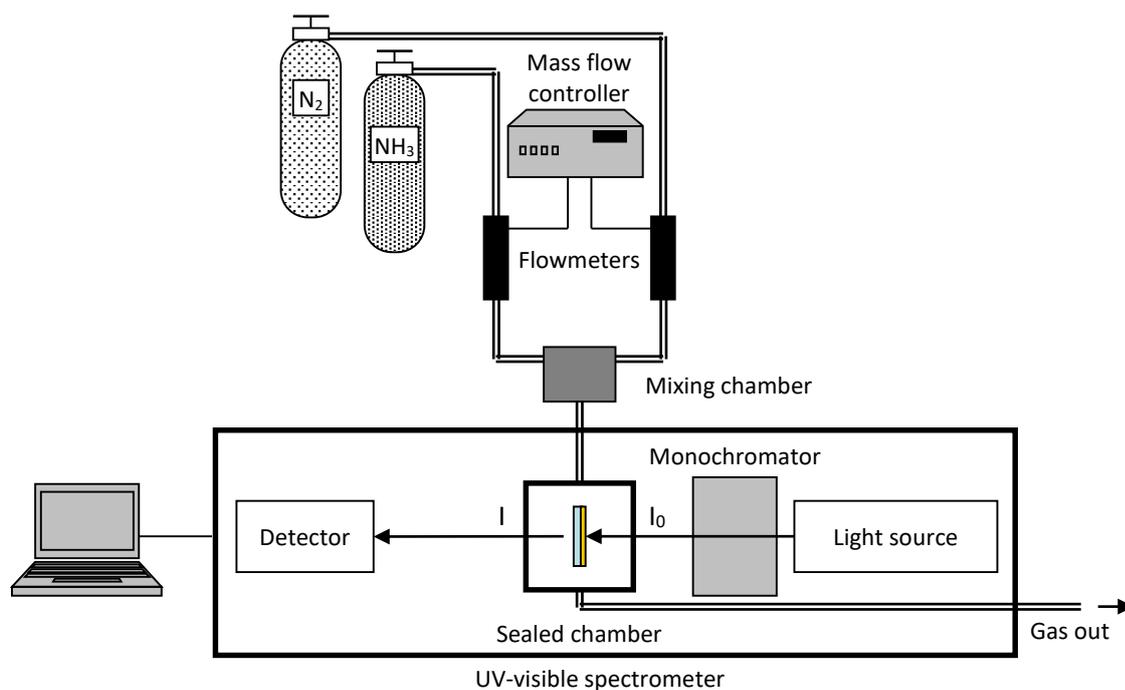


Figure 2. Experimental set-up of ammonia gas sensing based on absorbance variation measurement.

3. Results and discussion

In order to describe the plasma polymer/silver interaction mechanisms, the silver chemical affinities towards plasma polymers, and the polymer chemical functions involved with, are first

investigated. In a second time, the silver particle size, and particle size distribution are examined according to the polymer matrix. We are interested in three polymer matrices with different chemical functions: pp-ANI, pp-AAI and pp-HDFD. The interest of such composite materials is finally demonstrated through the study of its gas sensitivity using pp-ANI as matrix, which is well-known to be sensitive to ammonia.

3.1. Silver chemical affinity towards plasma polymers

Table 1 shows the XPS quantification of the different elements in the plasma polymer and composite materials.

Table 1. XPS quantification of the different elements in the plasma polymers and nanocomposites.

	Atomic percentages (%)				
	C	N	O	F	Ag
Bare pp-ANI	87.1±4.3	9.2±0.5	3.7±0.2	-	-
pp-ANI after impregnation+reduction	82.7±4.1	6.6±0.3	5.8±0.3	-	4.9±0.2
Bare pp-AAI	81.2±4.1	-	18.8±0.9	-	-
pp-AAI after impregnation+reduction	81.1±4	-	17.9±0.9	-	0.9±0.04
Bare pp-HDFD	50.2±2.5	-	0.4±0.02	49.4±2.5	-
pp-HDFD after impregnation+reduction	52.6±2.6	-	1.4±0.07	45.9±2.3	0.1±0.01

The presence of oxygen in plasma polymers deposited from monomers which do not contain oxygen (like ANI and HDFD) is due to the fact that the plasma polymer surfaces remain active and react with atmospheric oxygen after the films are removed from the reactor chamber.³⁰⁻³² As well as, the lower experimental ratios than the theoretical ones expected from the starting monomer N/C (0.1 vs. 0.16), O/C (0.23 vs. 0.33), and F/C (1 vs. 1.7) of pp-ANI, pp-AAI and

pp-HDFD respectively are mostly ascribed to fragmentations of the monomer in the R.F. glow discharge followed by a loss of nitrogen, oxygen, and fluorine during the polymerization.^{33,34} Moreover, the surface rearrangement phenomena of plasma polymers can also contribute to these theoretical and experimental differences.^{35,36}

All the composites are shown to contain Ag, C, and O near their surfaces, in addition to N for pp-ANI matrix and F for pp-HDFD one. A decrease in nitrogen and fluorine percentages is observed for the pp-ANI and pp-HDFD composites, with at the same time, an increase in the oxygen percentage. We can suppose that nitrogen and fluorine groups move from the surface to the bulk of the polymers when dipped in the aqueous solutions. The presence of AgNPs leaves this organization fixed once the composite is formed. The increase of the O percentage for the pp-ANI and pp-HDFD composites may be explained by the higher reactivity of the latter for which active species are mainly diradicals, whereas monoradicals species are principally created in pp-AAI.³⁷

The characteristic doublet at 368.6 eV and 374.6 eV of Ag3d_{5/2} and Ag3d_{3/2} with a difference of 6 eV on the wide-scan XPS spectra (not shown here) indicates that silver particles have been successfully synthesized in all the composites.^{13,38} However, the amount of silver depends on the type of the polymer matrix, since pp-ANI shows the most significant amount of silver (4.9%), then pp-AAI (0.9%) and pp-HDFD (0.1%). One of the main reasons for this difference is the type of interaction that exists between the polymer functional groups and the metal ions. In order to identify which chemical functions are involved through plasma polymer/silver interactions, XPS studies of N1s, and O1s are presented in Table 2.

Table 2. Atomic percentages of the different components in the high resolution XPS spectra of (a) N1s of pp-ANI, and (b) O1s of pp-AAI after impregnation+reduction.

	-N= (398.7±0.2 eV)	-NH- (399.7 eV)	⁺ • -NH- (400.6 eV)	-NH-/-N= (400.6 eV)	⁺ • -NH-/-N= (400.6 eV)
Bare pp-ANI (fwhm = 1.4 eV)	6.2±0.3	82.7±4.1	11±0.5	13.3±1.3	1.8±0.2
pp-ANI after impregnation (fwhm = 1.3 eV)	6.8±0.3	76.9±3.8	16.3±0.8	11.3±1.1	2.4±0.2
pp-ANI after impregnation+reduction (fwhm = 1.4 eV)	4.4±0.2	80.5±4	15.1±0.7	18.2±1.8	3.4±0.3

(a)

	C-OH/R, O-C-O (532.8±0.1 eV)	C=O (533.9±0.1 eV)
Bare pp-AAI (fwhm = 1.5 eV)	91.4±4.6	8.6±0.4
pp-AAI after impregnation (fwhm = 1.8 eV)	65.6±3.3	34.4±1.7
pp-AAI after impregnation+reduction (fwhm = 1.7 eV)	98±4.9	2±0.1

(b)

Concerning pp-ANI, the N1s spectra can be decomposed into three peaks at 398.7 and 399.7 eV, attributed to the imine (-N=) and amine (-NH-) groups of polyaniline, respectively, and at higher binding energy (400.6 eV) due to the positively charged nitrogen groups (-NH⁺•).³⁹⁻⁴² The decrease of amine functions of pp-ANI after adding metallic salt indicates that this last is oxidized by the silver ions to increase the level of doping (-NH⁺•), showing that Ag⁺ ions play the role of oxidizing dopant. The FTIR-ATR spectra (Figure 3-a) also reveal that the amines are the main chemical groups that interact with silver ions, by the decrease of the band at 3370 cm⁻¹ (νN-H). This result shows that the interaction of the (-NH⁺•/-NH-) redox couple of pp-

ANI with the silver ions is favored compared to the (-N=/⁺NH-) one whose standard potential ($E^\circ(-N=/⁺NH-) = 0.7-0.75 \text{ V}$)⁴³ is too close to that of silver ions ($E^\circ(\text{Ag}^+/\text{Ag}) = 0.80 \text{ V}$)^{44,45}.

To get a complete reduction of Ag^+ , the NaBH_4 solution ($E^\circ(\text{H}_2\text{BO}_3^-/\text{BH}_4^-) = -1.24 \text{ V}$) induces not only the reduction of the remaining metallic ions but also that of pp-ANI, since we observe a decrease in the imine (-N=) functions and an increase in the amine ones (-NH-) (Table 2-a),

whereas the ⁺NH - concentration remains unchanged. The fact that NaBH_4 is not able to reduce the ⁺NH - groups into amine ones suggests that the redox standard potential of ⁺NH -/⁺NH-, not found in the literature, is close to that of NaBH_4 . The standard potentials of pp-ANI, silver and NaBH_4 are summarized in Table 3-a.

Regarding pp-AAI, the polymer chain of pp-AAI is also oxidized by silver ions, as shown by the O1s decomposition⁴⁶ where a remarkable decrease of C-OH/R, O-C-O, with an increase of C=O is observed (Table 2-b). This result is also well observed on the FTIR-ATR spectra of pp-AAI (Figure 3-b), by the decrease of the characteristic band at 3380 cm^{-1} assigned to νOH , and which indicates the interaction between OH functions and Ag^+ . Thereafter, to get a complete reduction of Ag^+ , the NaBH_4 solution induces also the reduction of pp-AAI where an important increase of the hydroxyl functions is observed after reduction (Table 2-b and Figure 3-b). All these results are consistent with the standard potentials of pp-AAI ($E^\circ(\text{C}=\text{O}/\text{C}-\text{O}) = -0.16 \text{ V}$)⁴⁷, silver and NaBH_4 which are summarized in Table 3-b.

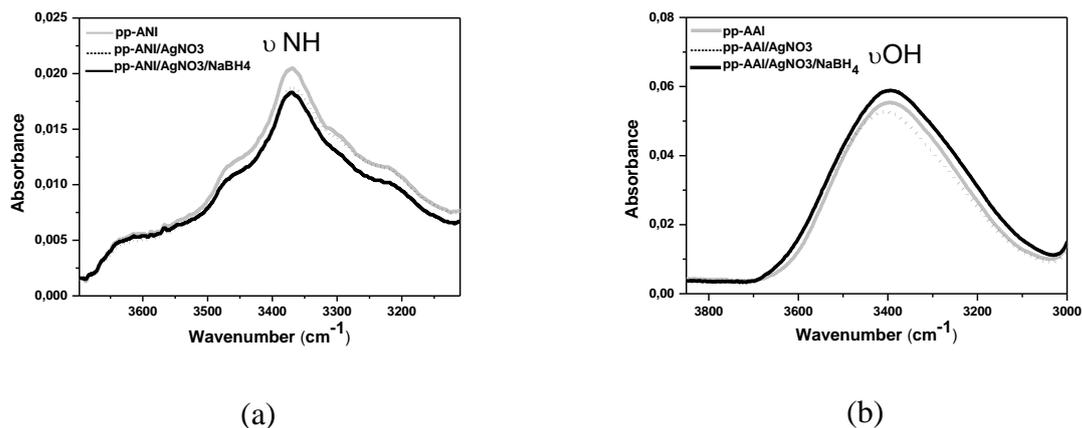


Figure 3. FTIR-ATR spectra of (a) pp-ANI, and (b) pp-AAI after impregnation+reduction.

Table 3. Redox standard potentials of (a) pp-ANI and (b) pp-AAI with those of silver and the reducing agent.

	E° (V)		E° (V)
Ag^+/Ag	0.80	Ag^+/Ag	0.80
pp-ANI (-N=/ NH -)	0.70 – 0.75	pp-AAI (C=O/C-O)	-0.16
pp-ANI (-NH^{\bullet} -/ NH -)	< -1.00 (estimated value)	$\text{H}_2\text{BO}_3^-/\text{BH}_4^-$	-1.24
$\text{H}_2\text{BO}_3^-/\text{BH}_4^-$	-1.24		

The oxidation of the polymer chains of pp-ANI and pp-AAI after impregnation in the metallic salt solution, must therefore be accompanied by a reduction of Ag^+ ions. For pp-ANI (Figure 4-a,b) and pp-AAI (Figure 4-c,d) matrices containing silver, the Ag $3d_{5/2}$ peak envelop is decomposed into two component peaks at 366.9 eV and 368.9 eV, which correspond to two different oxidation states of silver, namely silver oxide (Ag^{+I}) and metallic silver (Ag°), respectively. It is known that a metal ion having a higher standard potential than a polymer, once in contact, can be spontaneously reduced by this last to form zero-valent metal. This is the

case of pp-ANI used as a polymeric matrix and as silver reducing agent. The incomplete reduction even after adding NaBH_4 , may be explained by the strong interaction between Ag^+ and nitrogen functions, and the possible formation of a complex as shown by XPS (Figure 4-a,b). It has already been reported the formation of coordination bonds of nitrogen-containing polymers such as polyaniline,⁴⁸ poly(1,8-diaminonaphthalene),⁴⁹ polyacrylonitrile⁵⁰ with silver ions due to the nitrogen free electron pair that interacts with the metal ions.

Based on the standard potential of pp-AAI, Ag^+ is reduced to its metallic form (97.2 % of Ag°) by the latter (Figure 4-c,d). In fact, the use of alcohols as reducing agents is well-known in the literature.⁵¹ In contrast to pp-ANI, no silver ions are observed after reduction, and all the remaining Ag^+ ions are converted to Ag° by NaBH_4 which is a stronger reducing agent (lower standard potential) than the polymer matrix. Therefore, the only type of interaction between Ag^+ and OH functions is the electrostatic one.

It has been shown in the literature, a decrease of the free electrons pair ability to form a coordinating bond with a metal, when the electronegativity of atoms increases.^{52,53} The electron donating ability of nitrogen in pp-ANI is then higher than that of oxygen in pp-AAI since the electronegativity of nitrogen is lower than that of oxygen. Additionally, the pp-ANI structure is rich in π electrons (aromaticity) which further promotes the coordination between Ag^+ and N atoms, thus explaining this higher affinity of Ag^+ towards pp-ANI (4.9% of Ag) compared to pp-AAI (0.9% of Ag).

For pp-HDFD (Figure 4-e,f), the $\text{Ag}3d_{5/2}$ high resolution XPS spectra decomposition shows that silver is present in its metallic state. Due to the weak interaction between silver ions and the fluorine components (0.1% Ag), it can be predicted that the oxygen functions present in a very small quantity (0.4%) could interact with the silver ions and reduce them.

Therefore, the plasma polymers play the role of reducing agents, contributing to the formation of metallic particles.

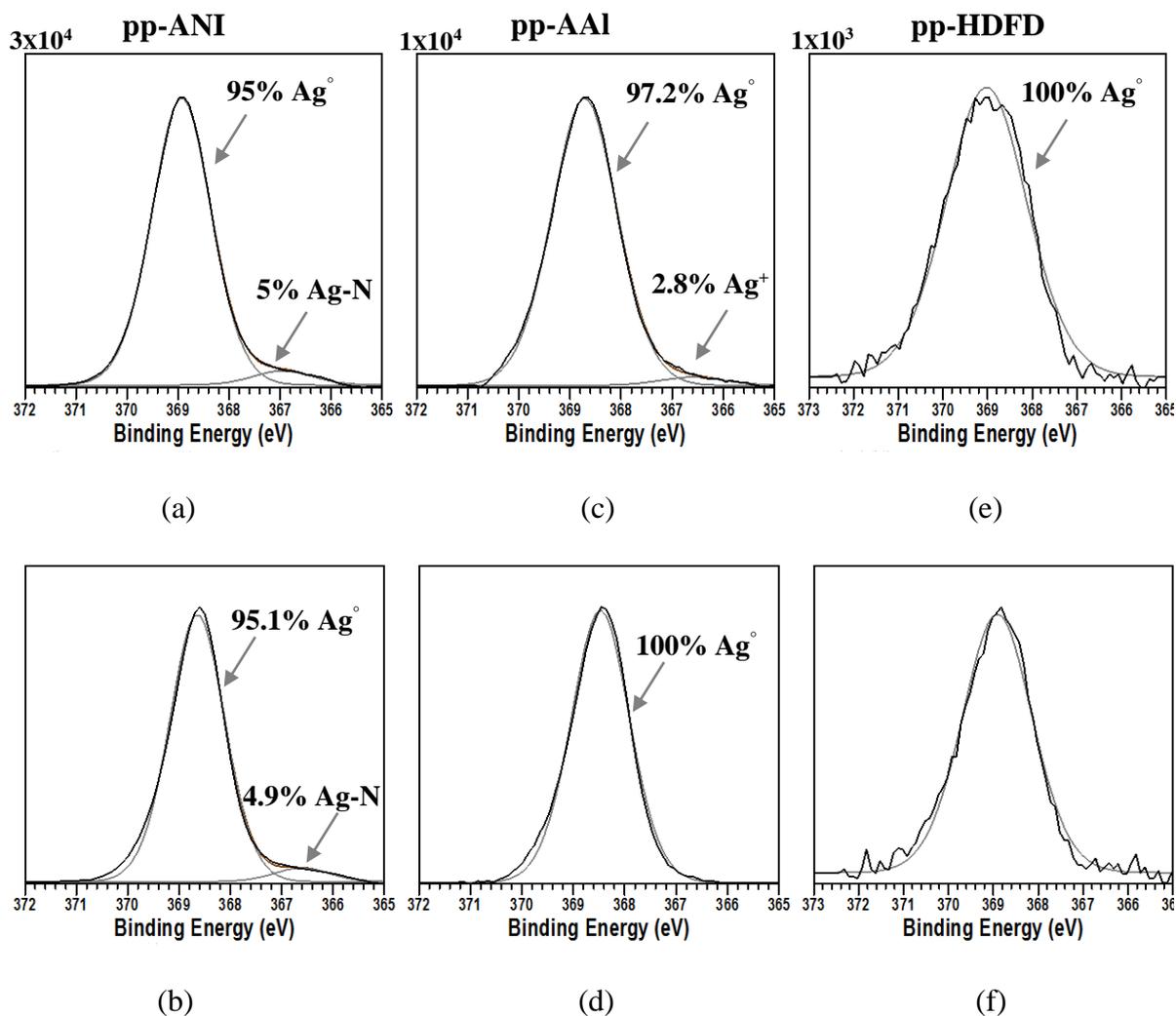
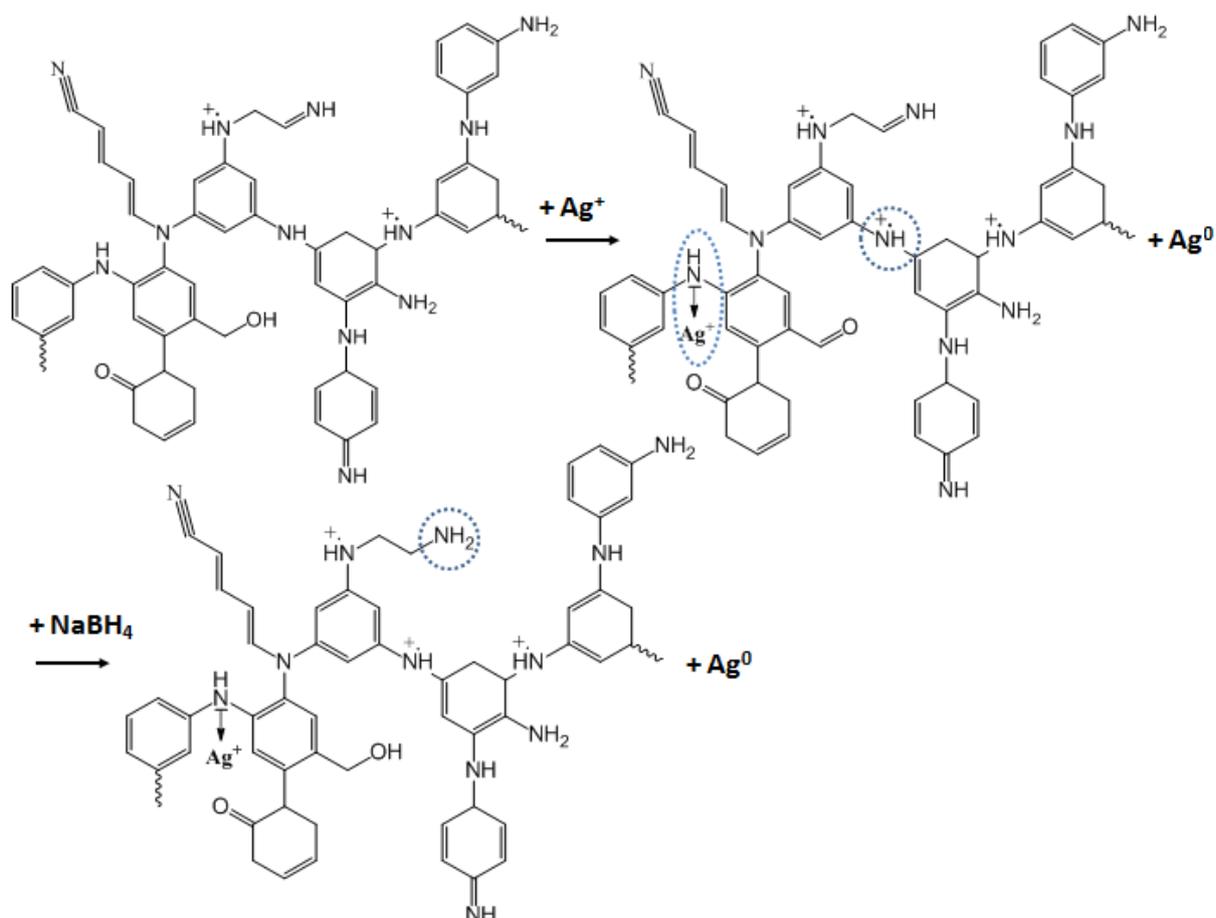


Figure 4. Ag $3d_{5/2}$ high resolution XPS spectra of (a) pp-ANI, (c) pp-AAI, (e) pp-HDFD after impregnation, and (b) pp-ANI, (d) pp-AAI, (f) pp-HDFD after impregnation+reduction.

Following these results, the proposed interaction mechanism between pp-ANI and silver salt on the one hand, and the reducing agent on the other hand is presented in scheme 1, according to the results observed by FTIR-ATR and XPS. The chemical structure of pp-ANI is non-linear due to the fragmentation and the recombination of the monomer in the plasma phase. This fragmentation can give rise to a loss of aromaticity either by the aromatic ring opening or by the π -bond scission in the aromatic ring. pp-ANI mainly contains amine functions, with some

imines and positively charged nitrogen groups, in addition to oxygen functions following the post-oxidation of the polymer.

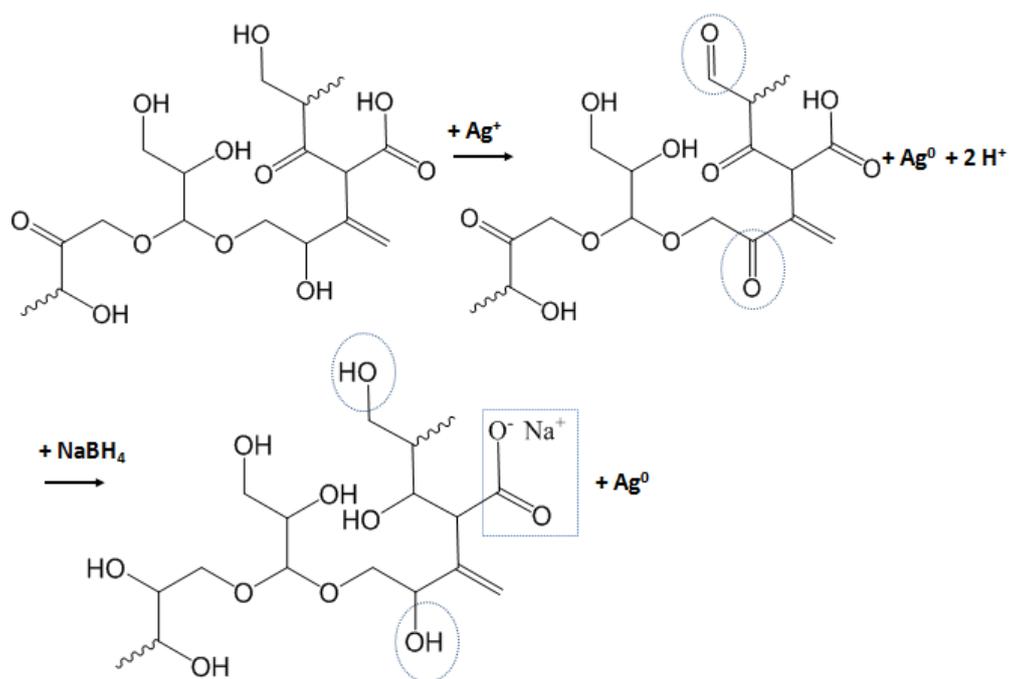
There are two different reactions that occur during the pp-ANI/AgNPs formation. One of them is the electrostatic interaction between Ag^+ and pp-ANI followed by the reduction of Ag^+ leading to the metallic nanoparticles Ag^0 , where at the same time, pp-ANI is oxidized (increase of $-\text{NH}^+$). The other one is the complexation of Ag^+ with the amine groups. Thus, the interactions are of electrostatic and coordination type,⁵⁴ taking into consideration that the electrostatic interaction followed by the reduction is more predominant than that of the complexation. After reduction, the imines functions are reduced by the NaBH_4 solution into amines.



Scheme 1. Complexation and electrostatic interactions between Ag^+ and pp-ANI.

Concerning pp-AAI, its chemical structure is also non-linear due to the fragmentation and the recombination of the monomer in the plasma phase. pp-AAI mainly contains hydroxyl functions, with some carbonyl and carboxyl groups.

The mechanism that can be involved in the formation of Ag° in the pp-AAI matrix is presented in scheme 2, where only the electrostatic interaction mechanism is involved in the nanocomposite formation, followed by the reduction of Ag^+ . At the same time, hydroxyl groups of pp-AAI are oxidized to form $-\text{C}=\text{O}$ groups corresponding either to a ketone, an aldehyde or a carboxyl group. After reduction, the $-\text{C}=\text{O}$ groups are reduced to hydroxyl ones, and a new bond is formed (COO^-Na^+) as a result of the interaction of pp-AAI with the NaBH_4 solution.



Scheme 2. Electrostatic interactions between Ag^+ and pp-AAI.

3.2. Repartition and size distribution of silver nanoparticles

AFM studies are carried out to examine the surface roughness of the different polymers and composite layers. The root mean square roughness values of the surfaces (R_q) are presented in Table 4. R_q is about 0.3 ± 0.1 and 0.9 ± 0.1 nm for pp-ANI and pp-AAI respectively, in comparison to about 1.3 ± 0.1 nm for pp-HDFD, indicating that the latter exhibits the highest

surface roughness, associated with a degradation and aggregation behavior, induced by fluorinated radicals. After silver particles addition, an increase of Rq for all the polymers is observed. For pp-ANI, Rq increases from 0.3 to 2.8 ± 0.2 nm, for pp-AAI it increases from 0.9 to 2.9 ± 0.2 nm, and from 1.3 to 5.3 ± 0.2 nm for pp-HDFD. The high quantity of silver particles inserted in pp-ANI compared to pp-AAI, explains the more pronounced increase in Rq. Although the amount of silver inserted in pp-AAI is more important than in the case of pp-HDFD, the increase in Rq is lower. Following these results, Rq values are then not only attributed to the amount of silver particles but also to their size and which will be studied afterwards.

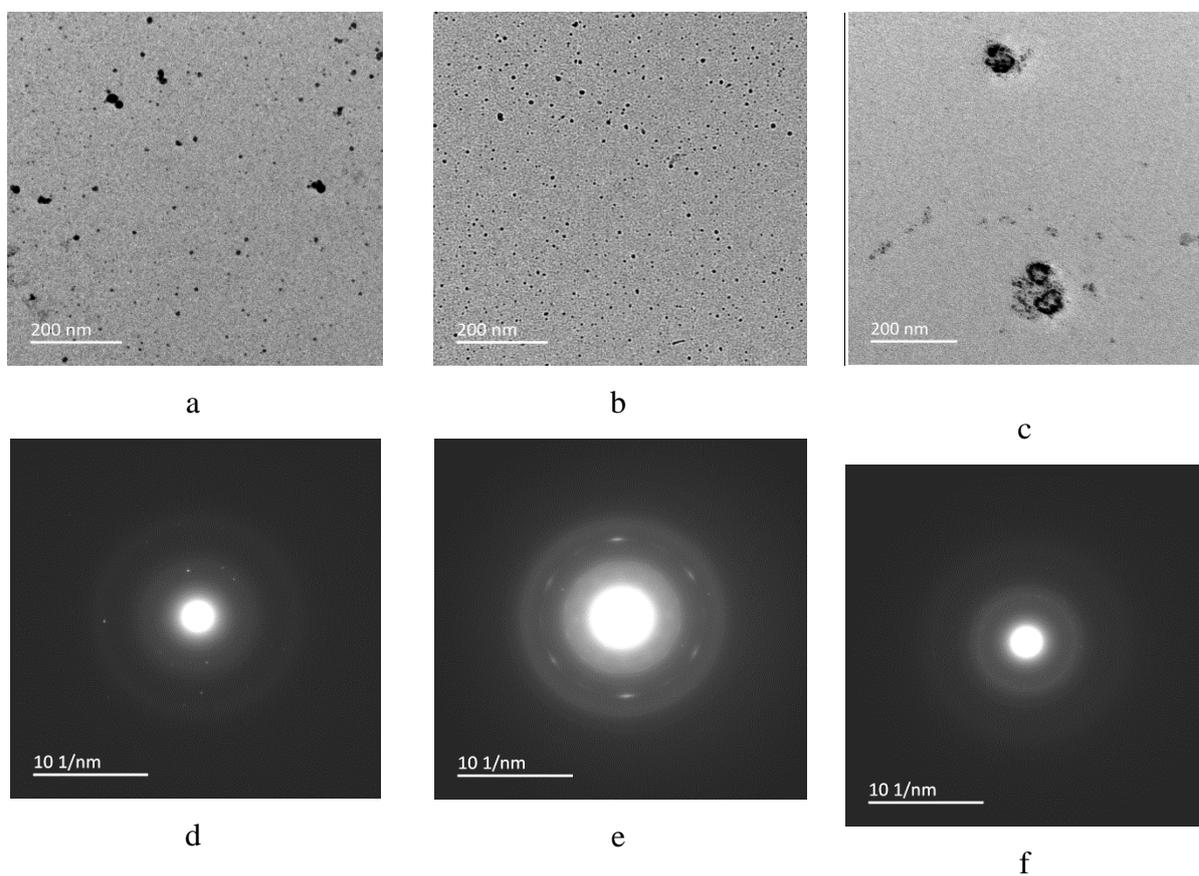
Table 4. Root mean square roughness values of the different polymer and composite layers.

Samples	Rq (nm)
Bare pp-ANI	0.3±0.1
pp-ANI after impregnation+reduction	2.8±0.2
Bare pp-AAI	0.9±0.1
pp-AAI after impregnation+reduction	2.9±0.2
Bare pp-HDFD	1.3±0.1
pp-HDFD after impregnation+reduction	5.3±0.2

The samples were investigated by transmission electron microscopy (TEM) to visualize the size and shape of silver particles. Spherical nanoparticles of 7.4 ± 0.4 nm and 12 ± 1.2 nm mean diameter are obtained for pp-AAI and pp-ANI as matrices, respectively (Figure 5). The smallest metal inclusions with the narrowest size distribution are produced by using pp-AAI (Figure 5-b,h). Moreover, for pp-HDFD, the weak chemical affinity between silver and the fluorocarbon film contributes to aggregate silver nanoparticles, inducing a broader size distribution in the range of 6 to 40 nm (Figure 5-c,i).^{55,56} Accordingly, we can say that silver nanoparticles likely

prefer to interact with each other rather than with the pp-HDFD matrix, thus the R_q value obtained for the fluorinated composite is well related to the aggregation of silver particles.

The electron diffraction pattern coupled to TEM makes it possible to study the structure of the crystallized materials. The SAED patterns (Figure 5-d,e,f) reveal that the diffraction rings from inner to outer, with d -spacings of 2.4, 2, 1.4, and 1.2 Å could be indexed as (111), (200), (220), and (311) reflections, respectively, corresponding to face-centered cubic silver structure (fcc). These values are consistent with diffraction data of fcc silver metal. These results confirm the crystallinity of silver nanoparticles with typical crystalline dots, in contrast to the amorphous structure of polymers.



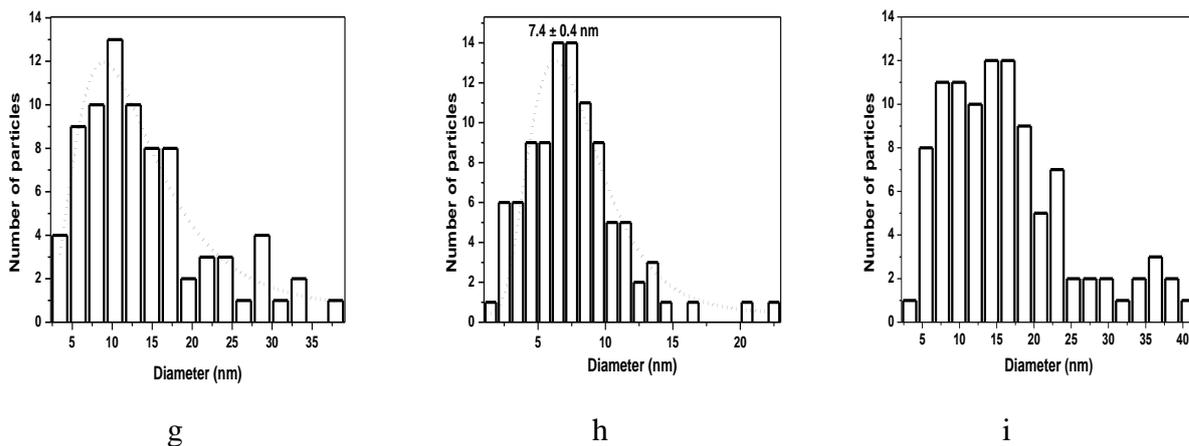


Figure 5. TEM images, SAED pattern and Ag particle size distribution in (a,d,g) pp-ANI, (b,e,h) pp-AAI, and (c,f,i) pp-HDFD after impregnation+reduction.

To confirm the formation of AgNPs, UV-Vis absorption spectroscopy has been demonstrated to be an effective tool, which depends on the Surface Plasmon Resonance (SPR).^{12,14,57,58} Figure 6 shows the UV-Visible absorbance spectra of plasma polymers before and after incorporation of AgNPs. pp-ANI shows one peak at 260 nm assigned to π - π^* transition of benzene rings with a shoulder at 290 nm assigned to polaron- π^* .^{35,59} pp-AAI shows a peak at 253 nm assigned to C-O, C=O, C=C,¹³ and for pp-HDFD, a peak at 251 nm attributed to CF, CF₂ is observed.^{60,61} Previous studies have shown that spherical AgNPs contribute to the absorption band at around 400 nm in the UV-visible spectra.⁶²⁻⁶⁴ A SPR band characteristic of AgNPs is detected around 426±5 nm for pp-ANI, and pp-AAI after impregnation, confirming the formation of spherical AgNPs in the plasma polymer matrices. However, after reduction, a slight increase in the peak intensity is observed in the case of pp-ANI, in contrast to pp-AAI, where the peak position, shape and intensity are changed. Taking into account that the peak position is very dependent on the nanoparticle size,^{65,66} the nanoparticles in pp-AAI (SPR band at 399 nm) are then smaller than those in pp-ANI (SPR band at 423 nm), and the size distribution is narrower, due to the narrower shape of the peak, thus supporting the results obtained by TEM. However, in the case of pp-HDFD, the absence of a characteristic peak of silver nanoparticles in the range of 400-

430 nm, confirms the formation of larger silver particles that tend to agglomerate,⁵⁷ in addition to the low incorporated amount (0.1%). All these observations show the effect of the nature of the polymer matrix on the size and particle size distribution of AgNPs.

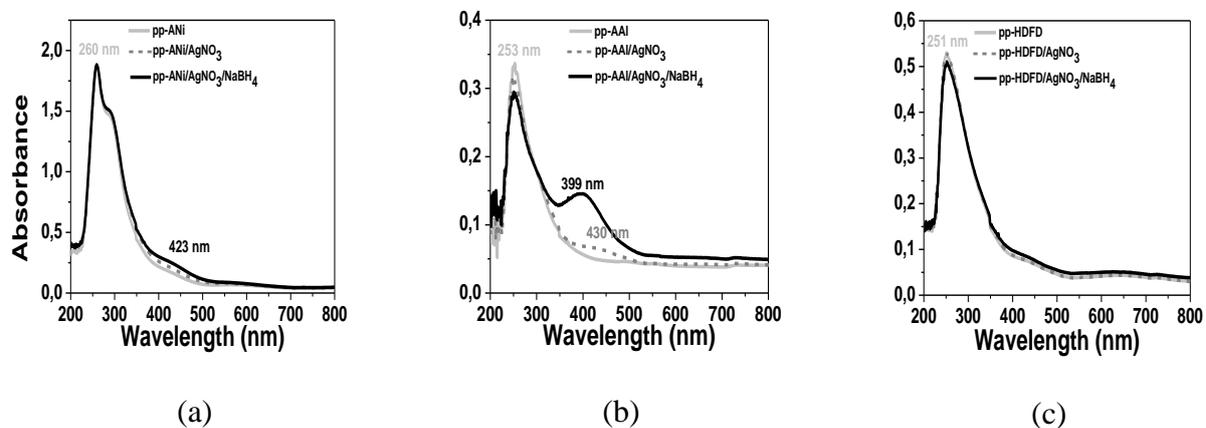


Figure 6. UV-Vis spectra of (a) pp-ANI, (b) pp-AAI, and (c) pp-HDFD after impregnation+reduction.

In conclusion, the results obtained by AFM and TEM are in good agreement with the results of UV-Vis absorption measurements. A uniform size distribution of AgNPs (narrower plasmon band) and smallest particle size of 7.4 ± 0.4 nm (SPR wavelength centered at 399 nm) are obtained by using pp-AAI as matrix. In contrast, the fluorocarbon film (pp-HDFD) exhibits particular behavior, where a low affinity for silver is observed with a tendency to aggregate.

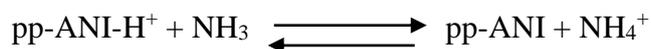
3.3. Gas sensing properties

In this part, we will show how the sensitivity to ammonia detection is improved when silver nanoparticles are inserted into pp-ANI matrix.

pp-ANI must be doped in order to create gas absorption sites, which are the positively charged nitrogen groups (including polarons). The doped form (pp-ANI-H⁺) is obtained by oxidative or acidic doping. In this work, the oxidative doping by iodine is used for bare pp-ANI and pp-

ANI/AgNPs composite to create more absorption sites,^{19,20} taking into account that pp-ANI already shows the presence of some absorption sites -NH⁺ - as analyzed by XPS (Table 2).

The interaction of ammonia gas with doped polyaniline corresponds to the deprotonation (dedoping) of the polymer and can be schematically expressed by the following equation:^{20,67,68}



The wavelength of 430 nm has been chosen to analyze the absorbance variation after ammonia absorption since it corresponds to a polarons absorption band of I₂-doped pp-ANI. The sensitivity of the layers containing or not AgNPs as a function of ammonia concentration is illustrated in Figure 7. The layers' sensitivity increases in a non-linear way when the ammonia concentration increases and tends to reach a plateau. This behaviour is explained by the saturation of the absorption sites at a certain ammonia concentration. Nevertheless, the ammonia vapour-sensing response of the layer with AgNPs is found to be higher than that without AgNPs, for all ammonia concentrations. The introduction of metal nanoparticles onto the polymer matrix is then effective in promoting the sensing response to ammonia. In order to explain the above observations, the chemical and morphological structures of these sensing layers are studied as follows.

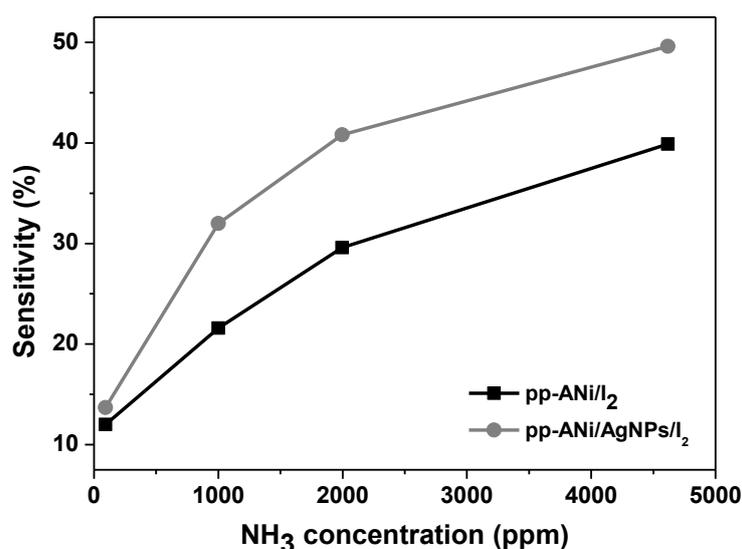


Figure 7. Sensitivity of doped pp-ANI and doped pp-ANI/AgNPs composite as a function of ammonia concentration.

On the one hand, the N1s XPS spectra (Table 2) show higher absorption sites (-NH^{\bullet} at 400.6 eV) for pp-ANI containing AgNPs (thanks to Ag^+ oxidative doping agents) than for bare pp-ANI, thus inducing in the end a better sensitivity. This increase in absorption sites is also observed by UV-Vis spectroscopy, where the shoulder at 290 nm assigned to polaron- π^* increases after adding AgNPs (Figure 6-a).

On the other hand, I/N XPS atomic ratio obtained after 24 h of I_2 doping, shows a higher value for pp-ANI containing AgNPs (0.6 ± 0.06) than for bare pp-ANI (0.5 ± 0.05), which means that the presence of AgNPs seems to provide a better interaction between pp-ANI and I_2 . Moreover it is worth to note that the morphological structure of these sensitive layers cannot be neglected. A rougher doped pp-ANI/AgNPs layer ($R_q = 5.8 \pm 0.2$ nm) is obtained compared to doped pp-ANI ($R_q = 0.5 \pm 0.1$ nm) (Figure 8). Thus, the presence of AgNPs leads to a higher surface-to-volume ratio that facilitates the diffusion of doping agents and ammonia molecules inside pp-ANI, hence facilitating their interactions.^{68,69}

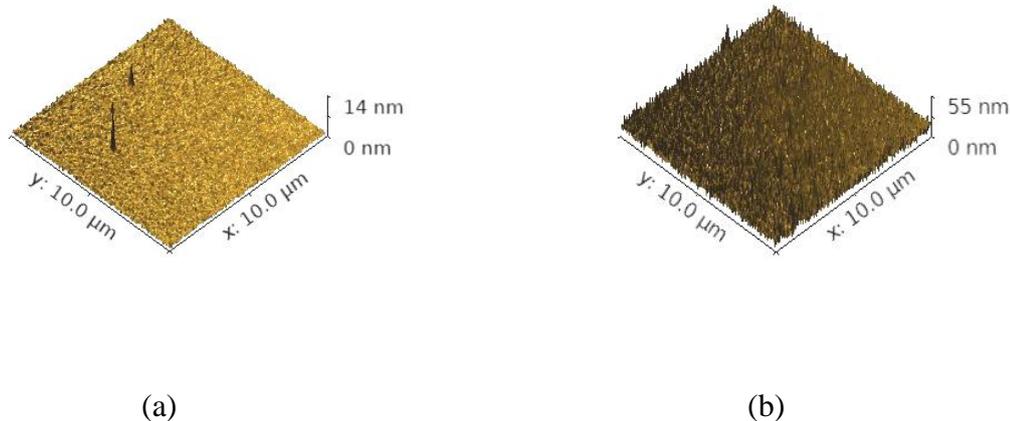


Figure 8. AFM images of (a) bare doped pp-ANI, and (b) doped pp-ANI/AgNPs composites.

4. Conclusion

To summarize, we succeeded in the formation of silver nanoparticles in the different plasma polymer layers but in different amounts and distribution according to the plasma polymer. pp-AAI is the most suitable matrix for the formation of the smallest particle size and the narrowest particle size distribution. But in terms of silver amount, pp-ANI with amine functions shows a pronounced affinity towards AgNPs, in comparison to the other polymer matrices explained by the structure of pp-ANI rich in π electrons, as well as the lower electronegativity of N atom compared to O and F ones. We have also demonstrated that AgNPs inserted into pp-ANI result in the improvement of sensitivity towards NH_3 , due to a change in the chemical and morphological structure of the polymer after adding AgNPs.

Conflict of Interest

The authors declare that they have no conflict of interest.

References

1. A. L. Stepanov, S. N. Abdullin, Y. V. Petukhov, Y. N. Osin, R. I. Khaibullin, I. B.

- Khaibullin, Formation of metal-polymer composites by ion implantation, *Philosophical Magazine B* 80 (2000) 23–28.
2. G. V. Ramesh, S. Porel, T. P. Radhakrishnan, Polymer thin films embedded with in situ grown metal nanoparticles, *Chem. Soc. Rev.* 38 (2009) 2646-2656.
 3. Q. Chen, M. Zhou, Y. Fu, J. Weng, Y. Zhang, L. Yue, F. Xie, C. Huo, Magnetron sputtering synthesis silver and organic PEO nanocomposite, *Surf. Coatings Technol.* 202 (2008) 5576–5578.
 4. K. Gupta, P. C. Jana, A. K. Meikap, Optical and electrical transport properties of polyaniline-silver nanocomposite, *Synth. Met.* 160 (2010) 1566–1573.
 5. J. Shan, Z. Ma, A review on amperometric immunoassays for tumor markers based on the use of hybrid materials consisting of conducting polymers and noble metal nanomaterials, *Microchim. Acta* 184 (2017) 969–979.
 6. S. Prakash, T. Chakrabarty, A. K. Singh, V. K. Shahi, Polymer thin films embedded with metal nanoparticles for electrochemical biosensors applications, *Biosens. Bioelectron.* 41 (2013) 43–53.
 7. M. Trchová, J. Stejskal, The reduction of silver nitrate to metallic silver inside polyaniline nanotubes and on oligoaniline microspheres, *Synth. Met.* 160 (13–14) (2010) 1479–1486.
 8. P. Xu, X. Han, B. Zhang, N. H. Mack, S. H. Jeon, H. L. Wang, Synthesis and characterization of nanostructured polypyrroles: Morphology-dependent electrochemical responses and chemical deposition of Au nanoparticles, *Polymer (Guildf)* 50 (2009) 2624–2629.
 9. N. Cioffi, L. Torsi, I. Losito, L. Sabbatini, P. Zambonin, T. Bleve-Zacheo, Nanostructured palladium–polypyrrole composites electrosynthesised from organic solvents, *Electrochim. Acta* 46 (2001) 4205–4211.

10. P. Pascariu, A. Airinei, M. Grigoras, L. Vacareanu, F. Iacomi, Metal-polymer nanocomposites based on Ni nanoparticles and polythiophene obtained by electrochemical method, *Appl. Surf. Sci.* 352 (2015) 95–102.
11. K. Vasilev, V. Sah, K. Anselme, C. Ndi, M. Mateescu, B. Dollmann, P. Martinek, H. Ys, L. Ploux, H. J. Griesser, Tunable antibacterial coatings that support mammalian cell growth, *Nano Lett.* 10 (2010) 202–207.
12. V. Kumar, C. Jolivalt, J. Pulpytel, R. Jafari, F. Arefi-Khonsari, Development of silver nanoparticle loaded antibacterial polymer mesh using plasma polymerization process, *J. Biomed. Mater. Res. A* 101 (2013) 1121–1132.
13. A. Fahmy, J. Friedrich, F. Poncin-Epaillard, D. Debarnot, Plasma polymerized allyl alcohol/O₂ thin films embedded with silver nanoparticles, *Thin Solid Films* 616 (2016) 339–347.
14. A. Airoudj, L. Ploux, V. Roucoules, Effect of plasma duty cycle on silver nanoparticles loading of cotton fabrics for durable antibacterial properties, *J. Appl. Polym. Sci.* 132 (2015).
15. E. Kosenko, Y. Kaminsky, E. Grau, M. D. Miñana, S. Grisolia, V. Felipo, Nitroarginine, an inhibitor of nitric oxide synthetase, attenuates ammonia toxicity and ammonia-induced alterations in brain metabolism, *Neurochem. Res.* 20 (1995) 451–456.
16. G. Sarala Devi, V. Bala Subrahmanyam, S. C. Gadkari, S. K. Gupta, NH₃ gas sensing properties of nanocrystalline ZnO based thick films, *Anal. Chim. Acta* 568 (2006) 41–46.
17. Q. Chang, K. Zhao, X. Chen, M. Li, J. Liu, Preparation of gold/polyaniline/multiwall carbon nanotube nanocomposites and application in ammonia gas detection, *J. Mater. Sci.* 43 (2008) 5861–5866.
18. A. Joshi, S. A. Gangal, S. K. Gupta, Ammonia sensing properties of polypyrrole thin

- films at room temperature, *Sensors Actuators, B Chem.* 156 (2011) 938–942.
19. A. Airoudj, D. Debarnot, B. Bêche, F. Poncin-Epaillard, Development of an optical ammonia sensor based on polyaniline/epoxy resin (SU-8) composite, *Talanta* 77 (2009) 1590–1596.
 20. T. Mérian, D. Debarnot, V. Rouessac, F. Poncin-Epaillard, Ammonia absorption study of pulsed-plasma polyaniline by quartz crystal microgravimetry and UV/vis spectrometry, *Talanta* 81 (2010) 602–608.
 21. X. Ma, G. Li, H. Xu, M. Wang, H. Chen, Preparation of polythiophene composite film by in situ polymerization at room temperature and its gas response studies, *Thin Solid Films* 515 (2006) 2700–2704.
 22. S. Cui, H. Pu, G. Lu, Z. Wen, E. C. Mattson, C. Hirschmugl, M. Gajdardziska-Josifovska, M. Weinert, J. Chen, Fast and selective room-temperature ammonia sensors using silver nanocrystal-functionalized carbon nanotubes, *ACS Appl. Mater. Interfaces* 4 (2012) 4898–4904.
 23. X. Yang, L. Li, F. Yan, Polypyrrole/silver composite nanotubes for gas sensors, *Sensors Actuators, B Chem.* 145 (2010) 495–500.
 24. U. V. Patil, N. S. Ramgir, N. Karmakar, A. Bhogale, A. K. Debnath, D. K. Aswal, S. K. Gupta, D. C. Kothari, Room temperature ammonia sensor based on copper nanoparticle intercalated polyaniline nanocomposite thin films, *Appl. Surf. Sci.* 339 (2015) 69–74.
 25. T. Sen, N. G. Shimpi, S. Mishra, R. Sharma, Polyaniline/ γ -Fe₂O₃ nanocomposite for room temperature LPG sensing, *Sensors Actuators, B Chem.* 190 (2014) 120–126.
 26. E. Detsria, J. Popanyasak, Fabrication of silver nanoparticles/polyaniline composite thin films using Layer-by-Layer self-assembly technique for ammonia sensing, *Colloids and Surfaces A: Physicochem. Eng. Aspects* 467 (2015) 57–65.

27. A.A. Athawale, P.P. Katre, Ag dispersed conducting polyaniline nanocomposite as a selective sensor for ammonia, *J. Metastable Nanocrystalline Mater.* 23 (2005) 323–326.
28. X. Li, Y. Gao, J. Gong, L. Zhang, L. Qu, Polyaniline/Ag composite nanotubes prepared through UV rays irradiation via fiber template approach and their NH₃ gas sensitivity, *J. Phys. Chem. C* 113 (2008) 69–73.
29. A. Mansour, F. Poncin-Epaillard, D. Debarnot, Distribution of metal nanoparticles in a plasma polymer matrix according to the structure of the polymer and the nature of the metal, *Thin Solid Films*, <https://doi.org/10.1016/j.tsf.2019.04.036>.
30. T. R. Gengenbach, Z. R. Vasic, S. Li, R. C. Chatelier, H. J. Griesser, Contributions of restructuring and oxidation to the aging of the surface of plasma polymers containing heteroatoms, *Plasmas Polym.* 2 (1997) 91–114.
31. X. Gong, L. Dai, A. W. H. Mau, H. J. Griesser, Plasma-polymerized polyaniline films: Synthesis and characterization, *J. Polym. Sci. Part A Polym. Chem.* 36 (1998) 633–643.
32. R. Hernandez, A. F. Diaz, R. Waltman, J. Bargon, Surface characteristics of thin films prepared by plasma and electrochemical polymerizations, *J. Phys. Chem.* 88 (1984) 3333–3337.
33. P. A. F. Herbert, L. O'Neill, J. Jaroszyńska-Wolińska, Soft Plasma polymerization of gas state precursors from an atmospheric pressure corona plasma discharge, *Chem. Mater.* 21 (2009) 4401–4407.
34. P. A. F. Herbert, L. O'Neill, J. Jaroszyńska-Wolińska, C. P. Stallard, A. Ramamoorthy, D. P. Dowling, A comparison between gas and atomized liquid precursor states in the deposition of functional coatings by pin corona plasma, *Plasma Process. Polym.* 8 (2011) 230–238.
35. D. Debarnot, T. Mérian, F. Poncin-Epaillard, Film Chemistry Control and Growth Kinetics of Pulsed Plasma-Polymerized Aniline, *Plasma Chem. Plasma Process.* 31

- (2011) 217–231.
36. A. Fahmy, R. Mix, A. Schönhals, J. F. Friedrich, Structure–Property Relationship of Thin Plasma Deposited Poly(allyl alcohol) Films, *Plasma Chem. Plasma Process.* 31 (2011) 477–498.
 37. H. Yasuda, M. O. Bumgarner, J. J. Hillman, Polymerization of organic compounds in an electrodeless glow discharge. V. Amines and nitriles, *J. Appl. Polym. Sci.* 19 (1975) 1403–1408.
 38. I. Medina-Ramirez, S. Bashir, Z. Luo, J. L. Liu, Green synthesis and characterization of polymer-stabilized silver nanoparticles, *Colloids Surfaces B Biointerfaces* 73 (2009) 185–191.
 39. X. Li, A. Rafie, Y. Y. Smolin, S. Simotwo, V. Kalra, K. K. S. Lau, Engineering conformal nanoporous polyaniline via oxidative chemical vapor deposition and its potential application in supercapacitors, *Chem. Eng. Sci.* (2018) DOI: [10.1016/j.ces.2018.06.053](https://doi.org/10.1016/j.ces.2018.06.053).
 40. M. M. Mahat, D. Mawad, G. W. Nelson, S. Fearn, R. G. Palgrave, D. J. Payne, M. M. Stevens, Elucidating the deprotonation of polyaniline films by X-ray photoelectron spectroscopy, *J. Mater. Chem. C* 3 (2015) 7180–7186.
 41. Z. F. Li, E. T. Kang, K. G. Neoh, K. L. Tan, Effect of thermal processing conditions on the intrinsic oxidation states and mechanical properties of polyaniline films, *Synth. Met.* 87 (1997) 45–52.
 42. S. Kumar, F. Gaillard, G. Bouyssoux, A. Sartre, High-resolution XPS studies of electrochemically synthesized conducting polyaniline films, *Synth. Met.* 36 (1990) 111–127.
 43. P. Xu, X. Han, B. Zhang, Y. Du, H.-L. Wang, Multifunctional polymer–metal nanocomposites via direct chemical reduction by conjugated polymers, *Chem. Soc. Rev.*

- 43 (2014) 1349–1360.
44. T. Zhao, R. Sun, S. Yu, Z. Zhang, L. Zhou, H. Huang, R. Du, Size-controlled preparation of silver nanoparticles by a modified polyol method, *Colloids Surfaces A Physicochem. Eng. Asp.* 366 (2010) 197–202.
 45. N. V. Blinova, J. Stejskal, M. Trchová, I. Sapurina, G. Ćirić-Marjanović, The oxidation of aniline with silver nitrate to polyaniline-silver composites, *Polymer (Guildf)* 50 (2009) 50–56.
 46. R. Zeng, M. Z. Rong, M. Q. Zhang, H. C. Liang, H. M. Zeng, Interfacial interaction in Ag/polymer nanocomposite films, *J. Mater. Sci. Lett.* 20 (2001) 1473–1476.
 47. T. Yonezawa, N. Toshima, Mechanistic consideration of formation of polymer-protected nanoscopic bimetallic clusters, *J. Chem. Soc. Faraday Trans.* 91 (1995) 4111.
 48. G. Neshar, G. Marom, D. Avnir, Metal-polymer composites: Synthesis and characterization of polyaniline and other polymer@silver compositions, *Chem. Mater.* 20 (2008) 4425–4432.
 49. X. G. Li, M. R. Huang, S. X. Li, Facile synthesis of poly(1,8-diaminonaphthalene) microparticles with a very high silver-ion adsorbability by a chemical oxidative polymerization, *Acta Mater.* 52 (2004) 5363–5374.
 50. Y. Wang, Q. Yang, G. Shan, C. Wang, J. Du, S. Wang, Y. Li, X. Chen, X. Jing, Y. Wei, Preparation of silver nanoparticles dispersed in polyacrylonitrile nanofiber film spun by electrospinning, *Mater. Lett.* 59 (2005) 3046–3049.
 51. Y. Sun, Y. Xia, Large-scale synthesis of uniform silver nanowires through a soft, self-seeding, polyol process, *Adv. Mater.* 14 (2002) 833–837.
 52. S. Ahrland, J. Chatt, N. R. Davies, The relative affinities of ligand atoms for acceptor molecules and ions, *Q. Rev. Chem. Soc.* 12 (1958) 265–276.
 53. H. Wang, X. Qiao, J. Chen, X. Wang, S. Ding, Mechanisms of PVP in the preparation

- of silver nanoparticles, *Mater. Chem. Phys.* 94 (2005) 449–453.
54. F. J. Liu, L. M. Huang, T. C. Wen, A. Gopalan, J. S. Hung, Interfacial synthesis of platinum loaded polyaniline nanowires in poly(styrene sulfonic acid), *Mater. Lett.* 61 (2007) 4400–4405.
 55. X. Wang, J. Zuo, P. Keil, G. Grundmeier, Comparing the growth of PVD silver nanoparticles on ultra thin fluorocarbon plasma polymer films and self-assembled fluoroalkyl silane monolayers, *Nanotechnology* 18 (2007) 265-303.
 56. V. Mankad, R. K. Kumar, P. K. Jha, Investigation of blue-shifted plasmon resonance: An optical properties study of silver nanoparticles, *Nanosci. Nanotechnol. Lett.* 5 (2013) 889–894.
 57. N. T. Nguyen, J. H. Liu, A green method for in situ synthesis of poly(vinyl alcohol)/chitosan hydrogel thin films with entrapped silver nanoparticles, *J. Taiwan Inst. Chem. Eng.* 45 (2014) 2827–2833.
 58. S. Ashraf, A. Z. Abbasi, C. Pfeiffer, S. Z. Hussain, Z. M. Khalid, P. R. Gil, W. J. Parak, I. Hussain, Protein-mediated synthesis, pH-induced reversible agglomeration, toxicity and cellular interaction of silver nanoparticles, *Colloids Surfaces B Biointerfaces* 102 (2013) 511–518.
 59. Q. Huang, G. Chen, J. Liu, One-pot synthesis of polyaniline doped with transition metal ions using H₂O₂ as oxidant, *Polym. Adv. Technol.* (2014) 1391-1395.
 60. F. Fanelli, Optical emission spectroscopy of argon-fluorocarbon-oxygen fed atmospheric pressure dielectric barrier discharges, *Plasma Process. Polym.* 6 (2009) 547–554.
 61. I. Vinogradov, A. Lunk, Spectroscopic diagnostics of DBD in Ar/fluorocarbon mixtures - Correlation between plasma parameters and properties of deposited polymer films, *Plasma Process. Polym.* 2 (2005) 201–208.
 62. K. Shameli, M. Bin Ahmad, S. D. Jazayeri, S. Sedaghat, P. Shabanzadeh, H. Jahangirian,

- M. Mahdavi, Y. Abdollahi, Synthesis and characterization of polyethylene glycol mediated silver nanoparticles by the green method, *Int. J. Mol. Sci.* 13 (2012) 6639–6650.
63. Y. Sun, Y. Xia, Gold and silver nanoparticles: A class of chromophores with colors tunable in the range from 400 to 750 nm, *Analyst* 128 (2003) 686-691.
64. K. Stamplecoskie, J. Scaiano, Light Emitting Diode Irradiation Can Control the Morphology and Optical Properties of Silver Nanoparticles, *J. Am. Chem. Soc.* 132(6) (2010) 1825-1827.
65. L. Balan, R. Schneider, D. J. Loughnot, A new and convenient route to polyacrylate/silver nanocomposites by light-induced cross-linking polymerization, *Prog. Org. Coatings* 62 (2008) 351–357.
66. G. A. Martinez-Castanon, N. Niño-Martínez, F. Martínez-Gutierrez, J. R. Martínez-Mendoza, F. Ruiz, Synthesis and antibacterial activity of silver nanoparticles with different sizes, *J. Nanoparticle Res.* 10 (2008) 1343–1348.
67. H. T. Hien, H. T. Giang, N. Van Hieu, T. Trung, C. Van Tuan, Elaboration of Pd-nanoparticle decorated polyaniline films for room temperature NH₃ gas sensors, *Sensors Actuators, B Chem.* 249 (2017) 348–356.
68. S. Mikhaylov, N. Ogurtsov, Y. Noskov, N. Redon, P. Coddeville, J. L. Wojkiewicz, A. Pud, Ammonia/amine electronic gas sensors based on hybrid polyaniline-TiO₂ nanocomposites. the effects of titania and the surface active doping acid, *RSC Adv.* 5 (2015) 20218–20226.
69. S. Virji, J. Huang, R. B. Kaner, B. H. Weiler, Polyaniline nanofiber gas sensor: examination of response mechanisms, *Nano Lett.* 4 (2004) 491–496.