Interaction of (3-Aminopropyl)triethoxysilane with Pulsed Ar-O 2 Afterglow: Application to Nanoparticles Synthesis

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ABSTRACT

The interaction of (3–Aminopropyl)triethoxysilane (APTES) with pulsed late Ar-O_2 afterglow is characterized by the synthesis of OH, CO and CO_2 in the gas phase as main by-products. Other minor species like CH, CN and C_2H are also produced. We suggest that OH radicals are produced in a first step by dehydrogenation of APTES after interaction with oxygen atoms. In a second step, the molecule is oxidized by any O_2 state, to form peroxides that transform into by-products, break thus the precursor C–C bonds. If oxidation is limited, *i.e.* a low duty cycle, fragmentation of the precursor is limited and produced nanoparticles keep the backbone structure of the precursor, but contain amide groups produced from the amine groups initially available in APTES. At high duty cycle, silicon-containing fragments contain some carbon and react together and produce nanoparticles with a non-silica-like structure.

INTRODUCTION

The elaboration of new functional coatings on inorganic surfaces by biomimetic using green technologies can be considered as a real breakthrough in the world of metallic alloys manufacturers. In particular, biological molecules strongly chemically bonded to a metallic surface could provide to a flat product interesting features such as natural lubricant, anti-biocorrosion or antifouling properties [1–5]. However, any broad development of these revolutionary "biocoatings" requires an excellent control at nanoscale of the chemical functional groups that constitute the coating.

Plasma deposition processes are well known to produce improved surface stability, controlled chemical functionality and topography [6–10]. They can be considered as the best industrial technique platform to realize, on each type of inorganic surfaces – and especially on metallic surfaces – an inorganic/organic interlayer for the covalent bonding of biomolecules [11–13]. Plasma polymerisation is a process able to produce conformal, pin-hole free, highly cross-linked polymers. The main interest of this process here is to offer the advantage of not only providing a high adhesion level with the native oxide/hydroxide layer of the metallic surface but also assisting the chemical deposition of a stable layer with highly reactive chemical groups to further promote the immobilization of biomolecules. Both features are nevertheless controlled by the choice of the precursor that must be designed so as to include suited chemical groups.

Among the different functional groups, primary amines ($-NH_2$) are interesting owing to their high reactivity towards biomolecules. Thus, NH_2-rich polymeric materials can react with biomolecules through the formation of covalent bonds with carboxyl group present in biomolecules such as peptide, enzyme or proteins [14]. Amino-alkoxysilane compounds are widely used in wet chemistry [15] for surface functionalization of oxide materials, particularly for silicon dioxide. Indeed, alkoxy groups can react with hydroxyl groups present on the

surface of many oxidized materials, and especially metallic alloys, thus forming –M–O–Si covalent bonds with the surface. Thus, a dense siloxane $SiO_xC_yH_z$ layer with a high density of –NH$_2$ groups can combine low permeation and immobilization of biomolecules.

Because of the complexity of plasma polymerization process, it is a challenging task to induce polymerisation of amino-alkoxysilane by creation of Si-O-Si bonds while keeping –NH$_2$ group as such. A strategy chosen by some groups consists in a two-step polymerisation or copolymerisation process. For instance, a two-step plasma process has been used by Jampala *et al.* [16]. They use HMDSO polymerization in order to deposit a siloxane layer followed by the deposition of pp-ethylenediamine to deposit amino-functionalised coating on stainless steel substrate. These authors noticed that, whereas pp-ethylenediamine readily delaminated from stainless steel substrate, the deposition of an intermediate pp-HMDSO layer strongly enhanced the adhesion of the amino-functionalized plasma polymer. The interest of a copolymerisation process has also been highlighted by other authors for the deposition of aminated coatings on specific polymer surfaces [17–20]. In these works, APTES, *i.e.* aminopropyltriethoxysilane ((C_2H_5O)$_3$-Si-CH_2-CH_2-CH_2-NH_2), was used as a monomer for the deposition of a siloxane layer. The group of D.E. Williams showed that the deposition of plasma polymerized ethylenediamine film on cyclo-olefin polymer had a weak adherence. This adherence, as well as the amine surface density, is significantly increased by using copolymerization of EDA and APTES [17, 18]. These authors pointed out that the siloxane network is essential to ensure an efficient amine functionalization.

Avoiding multi-step processes and/or multiple precursors use is a real challenge. Growing in a single step, a siloxane-based thin film functionalized with primary amines using a single amino-alkoxysilane precursor is still under investigation. However, only few articles deal with this issue [5, 19–22], and the majority of these studies point out a rather low amine density of the as-grown thin films and/or do not produce obvious evidences of efficient –NH$_2$

functionalization. Indeed, Gubala *et al.* [19] point out interesting results using APTES as unique precursor but they obtained a lower density compared to the APTES-EDA copolymer thin film [18]. In addition, these articles are focused on the synthesized thin films and do not deal with plasma polymerisation.

In this work, we explain how oxygen active species created in a plasma afterglow, and more precisely oxygen atoms, react with APTES.

EXPERIMENTAL METHOD

The experimental set-up is depicted in figure 1. A surface wave-driven plasma is ignited in a fused silica tube (5 mm inner diameter) by exposure to 2.45 GHz microwaves launched with a surfatron. The microwave power is set constant at 250 W. The pressure was measured upstream and downstream the flow (points E and A respectively in figure 1). All the treatments were carried out at 1500 Pa (downstream pressure in A), the upstream pressure being 2000 Pa. The maximum pumping speed of the primary pump was 40 m^3 h^{-1}. The mininum vacuum pressure was 10^{-2} mbar. An Ar–8.7vol.% O_2 mixture, flowing at 1150 sccm, is used as plasma gas. In point C (see figure 1), the plasma tube reaches a 1.2 meter long fused silica tube (16 mm inner diameter) crossed over its whole length by an infrared beam used to probe the afterglow *in situ*. Liquid APTES (99.0% purity from Sigma Aldrich) was contained in a temperature-controlled bubbler. Its vapour was transported to the afterglow by an argon flow (30 sccm) through a fused silica tube (5 mm inner diameter and 30 cm in length) connected to the main tube at point B. The APTES partial pressure was determined using the following Antoine's formula established from experimental data obtained by probing the FTIR absorption signal of pure APTES evaporated at temperatures ranging from 300 K to 390 K:

$$Log_{10}\left(\frac{P_s}{P_{ref}}\right) = 1.6 - \left(\frac{2384.4}{T(\text{K}) + 300}\right) \qquad (1)$$

where P_s is the vapour pressure (in Pa), P_{ref} the standard pressure (101325 Pa) and T the temperature (K). At 300 K, $P_s / P_{ref} = 4.23 \times 10^{-3}$ and the APTES flow rate was 0.13 sccm in the Ar-O_2 afterglow.

The light emitted by the afterglow was analysed by optical emission spectroscopy (OES). It was collected with an optical fibre connected to 550 mm focal length monochromator (Jobin–Yvon TRIAX 550) equipped with a 100 grooves mm^{-1} grating for survey spectra in the range [250–900 nm] and a 1,800 grooves mm^{-1} grating to record specific transitions with high spectral resolution. The spectrometer was coupled with a HORIBA Jobin–Yvon i-Spectrum Two iCCD detector. Each measurement was averaged over 20 spectra.

In the pulse mode, the discharge was run with a period of 100 ms, the duty cycle varying between 10 and 90%. The average velocity in the afterglow tube was easily deduced from time-resolved OES measurement. A value of 7.1 ± 0.1 m s^{-1} was found.

FTIR spectroscopy was performed with a commercial Agilent FTIR 680 spectrometer. The infrared beam leaving the spectrometer goes through lenses and collimators to get a parallel beam (16 mm in diameter). This optical arrangement decreases the intensity of the exiting beam by one order of magnitude typically. The infrared beam was next sent through a KBr window in the afterglow (single pass) and detected by a remote liquid-nitrogen cooled mercury-cadmium-telluride (MCT) detector which was protected from the afterglow by another KBr window. To minimize the influence of water and CO_2, the infrared path outside the afterglow was continuously flushed by clean, dry and carbon-free air. Absorption spectra were recorded in the range [500–4000 cm^{-1}] with a spectral resolution of 2 cm^{-1}. Reference spectra (*i.e.* either without gas and plasma or with gas but without plasma) were acquired before each acquisition to determine the absorption spectra in afterglow conditions. Each

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measurement was averaged over 100 scans in continuous mode to improve the signal-to-noise ratio.

Nanoparticles were collected on an aluminium substrate located just before the pumping unit. Transmission electron microscopy (TEM) investigation was performed on as-grown nanoparticles with a JEOL ARM 200F – Cold FEG TEM/STEM running at 200 kV (point resolution 0.19 nm) fitted with a GIF Quantum ER. Scanning Electron Microscopy (SEM) was made with a Philips XL 30. FTIR measurements on nanoparticles were performed with a commercial Agilent FTIR 680 spectrometer in attenuated total reflection (ATR) mode. Spectra were acquired in the range [500–4000 cm^{-1}] with a spectral resolution of 4 cm^{-1}.

X-ray photoelectron spectroscopy (XPS) measurements were performed with a Thermo VG Microlab350 spectrometer using a non-monochromated Al Kα and Mg Kα dual anode as X-ray source operated at 300 W and a Spherical Sector Analyser. Survey spectra to identify elements on the surface were recorded in steps of 1 eV at a 100 eV pass energy. High-resolution spectra of separate photoelectron lines (C 1s, O 1s, Si 2p and N 1s) were taken by steps of 0.05 eV at a constant pass energy of 20 eV. The normal operation pressure was 5×10^{-9} mbar. The photoelectron take-off angle (TOA) was normal to the surface of the samples. The sample surface covered by the analysis is 2×5 mm^2. Spectra processing (atomic concentrations, curve fitting, etc.) was done after the removal of a Shirley type background with the CasaXPS software. Samples were introduced in the spectrometer readily after plasma treatment in order to limit contamination from ambient air storage and directly analysed.

RESULTS AND DISCUSSION

Optical emission spectroscopy

The Ar–O_2 afterglow, run in the continuous mode, was characterized with and without APTES by OES (figure 2). Observed transitions are reported in Table 1. Without APTES, the

visible light from the afterglow is very weak and only visible in the dark with the naked eye. Besides a non-negligible contribution due to the $OH\left(A^2\Sigma^+, v'=0 \rightarrow X^2\Pi, v''=0\right)$ at 306 nm, the green line of atomic oxygen at 557.7 nm due to the $O\left(^1S \rightarrow {}^1D\right)$ transition, called the green line and the atmospheric band of O_2 due to the $O_2\left(b^1\Sigma_g^+ \rightarrow X^3\Sigma_g^-\right)$ transition at 762 nm could be also recorded with the highest iCCD gain (the OH emission saturates in these conditions – see **supplemental material 1**). Indeed, the two latter transitions arise from metastable states produced by the following mechanism:

$$O + O + O \rightarrow O(^1S) + O_2 \qquad\qquad\text{(a)}$$

$$O + O + M \rightarrow O_2\left(b^1\Sigma_g^+\right) + M \qquad\qquad\text{(b)}$$
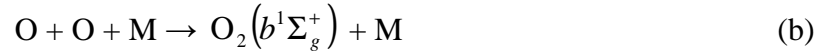
and are then very weak in intensity. Light being recorded with an optical fiber, concentrations are determined with respect to its position (defined within two mm typically), regardless of the spatial gradients that extend over much larger distances (several tens of cm).

To determine the concentration of oxygen atoms, the NO-titration method was applied. This is a two-step method described in ref. [23]: In a first step, optical calibration is performed by mixing an Ar-N_2 afterglow and a Ar–1.4vol.%NO mixture introduced instead of the Ar–APTES mixture (point B in figure 1). The same arrangement is introduced in a second step to determine the concentration of oxygen atoms at the mixing point of the Ar–1.4vol.%NO mixture with the Ar–O_2 afterglow (point B in figure 1). This method leads to [O] = $5.4\pm1.0\times10^{15}$ cm^{-3}.

Thanks to this value, the absolute concentrations of all emitting species, provided they are intense enough, could be determined with an accuracy of about 30%. Thus, the concentration of the $O_2\left(b^1\Sigma_g^+, v'=0\right)$ emitting states leading to the atmospheric band could be determined: $\left[O_2\left(b^1\Sigma_g^+, v'=0\right)\right]$ = $5.1\pm2.0\times10^{11}$ cm^{-3}. This value is consistent with the O atom
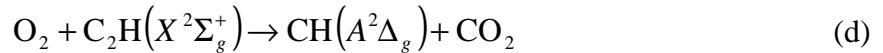
concentration reported previously. Indeed, these two concentrations are connected through process (b). To determine the O atom concentration from the $O_2\left(b^1\Sigma_g^+, v'=0\right)$ state, we used the model presented in [24]. It gives [O] = $3.2\pm1.0\times10^{15}$ cm^{-3}. We found [CH(A)] = $9.2\pm3.0\times10^6$ cm^{-3}, [CN(B)] = $2.6\pm0.8\times10^5$ cm^{-3} and [OH(A)] = $2.0\pm0.5\times10^8$ cm^{-3} when the afterglow is run in the continuous mode. The rotational spectrum of the atmospheric band was used to determine the gas temperature (as **supplemental material 2**). A temperature of 330 ± 5 K was found, confirming the low temperature of the afterglow.

When APTES is added to the afterglow, new transitions are observed. They are listed in Table 1. Besides OH emission which increases strongly (by about one order of magnitude), emissions of non-oxidized molecules, CH and CN, are found (see high-resolution spectra provided as **supplemental material 3**). The green line and the atmospheric band are no longer visible in these conditions.

CH comes naturally from APTES. The origin of the chimiluminescence band of CH(*A*) was clarified by Devriendt *et al.* [25] after the reference work by Grebe *et al.* [26]. It would be due to the following reaction:

$$O + C_2H\left(X^2\Sigma_g^+\right) \rightarrow CH\left(A^2\Delta_g\right) + CO \tag{c}$$
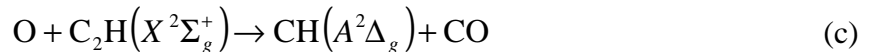
with $k_c = 1.1\times10^{-11}$ cm^3 s^{-1} at 300 K and

$$O_2 + C_2H\left(X^2\Sigma_g^+\right) \rightarrow CH\left(A^2\Delta_g\right) + CO_2 \tag{d}$$

with $k_d = 3.6\times10^{-14}$ cm^3 s^{-1} at 290 K [27]. A less probable origin of CH(*A*) is [28]:

$$C_2 + OH \rightarrow CH\left(A^2\Delta_g\right) + CO \tag{e}$$

because no emission of the Swan system due to C_2 excited states is observed in our conditions, contrary to data reported in [27].

Kinetics of CH(A) – only the ground state is considered here – is associated with the following spontaneous emission and quenching processes:

$$CH(A^2\Delta_g) \rightarrow CH(X^2\Pi_u) + h\nu \qquad \text{(f)}$$

$$CH(A^2\Delta_g) + M \rightarrow CH(X^2\Pi_u) + M \qquad \text{(g)}$$
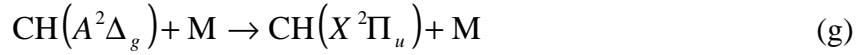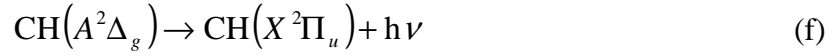
The radiative lifetime of CH(A) is 540 ns [28]. The quenching rates by Ar and O_2 are given by Chen *et al.* [29]: $k_g^{Ar} = 5.2 \times 10^{-13}$ cm^3 s^{-1} et $k_g^{O_2} = 2.2 \times 10^{-11}$ cm^3 s^{-1}. Applying the quasi-steady state approximation (processes c, d, f and g), one finds:

$$[C_2H] = \frac{\left(k_g^{Ar}[Ar] + k_g^{O_2}[O_2] + \tau_{CH(A)}^{-1}\right)[CH(A)]}{k_c[O] + k_d[O_2]} \qquad (2)$$

This leads to $[C_2H] = 1.3 \pm 3.0 \times 10^8$ cm^{-3}. The APTES concentration being 3.9×10^{13} cm^{-3}, the production of C_2H radicals is a negligible reaction pathway.

The presence of CN molecules is necessarily due to the nitrogen atom present in the amine group and to one carbon atom coming from the precursor, but not mandatorily the one attached to amine group. To the best of our knowledge, the only mechanism accounting for the emission of the CN violet system in an argon-oxygen afterglow is [30]:

$$O + O + CN(X) \rightarrow O_2 + CN(B) \qquad \text{(h)}$$
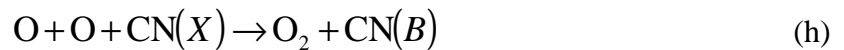
Indeed, the reaction of N with CH for example, or with C and a third body is highly unlikely, nitrogen atoms being not present as such in oxygen afterglows. The rate constant of process (h) is given with a high uncertainty: $k_h = (3 - 30) \times 10^{-31}$ cm^6 s^{-1} [30].

The loss mechanisms of the CN(B) state are the following spontaneous emission and quenching processes:

$$CN(B) \rightarrow CN(X) + h\nu \qquad \text{(i)}$$

$$CN(B) + M \rightarrow CN(X) + M \qquad \text{(j)}$$

The radiative lifetime of CN(B,v=0) is $\tau_{CN(B)}$ =66.6 ns [31]. The quenching rate by Ar is given by [32]: $k_j^{Ar} = 1.0 \times 10^{-11}$ cm^3 s^{-1}. For O$_2$, we took the same value as N$_2$ [32]: $k_j^{O_2} = 2.5 \times 10^{-11}$ cm^3 s^{-1}.

Applying the quasi-steady state approximation (processes h, i and j), one finds:

$$[CN] = \frac{\left(k_j^{Ar}[Ar] + k_j^{O_2}[O_2] + \tau_{CN(B)}^{-1}\right)[CN(B)]}{k_h[O]^2} \qquad (3)$$

This gives: [CN] = (0.56 or 5.6)×10^{11} cm^{-3}, according to the chosen value of k_h, which is at best a hundredth of the APTES concentration. Then, the production of CN radicals is a minor reaction pathway.

The origin of the OH chimiluminescence is well-known from works on acetylene combustion. It is due to the following excitation process:

$$O_2 + CH(X) \rightarrow OH(A) + CO \qquad (k)$$

Carl *et al.* [33] give for a temperature in the range [296-511] K:

$$k_k = 1.06 \times 10^{-13} \exp\left(-\frac{84}{T(K)}\right)$$

i.e. at 300 K, $k_k = 8.0 \times 10^{-14}$ cm^3 s^{-1}. The loss mechanisms of OH(A) are the following spontaneous emission and quenching processes:

$$OH(A) + M \rightarrow OH + M \qquad (l)$$

$$OH(A) \rightarrow OH \qquad (m)$$

where M=Ar or O$_2$. The rate constants of the quenching processes by these species are respectively $k_l^{Ar} = 8.8 \times 10^{-14}$ cm^3 s^{-1} [34] and $k_l^{O_2} = 8.0 \times 10^{-11}$ cm^3 s^{-1} [35]. For spontaneous emission, we consider only the v'=0 level with $\tau_{OH(A)}$=800 ns [34].

Applying the quasi-steady state approximation (processes k, l and m), one finds:

$$[CH] = \frac{\left(k_l^{Ar}[Ar] + k_l^{O_2}[O_2] + \tau_{OH(A)}^{-1}\right)[OH(A)]}{k_k[O_2]} \qquad (3)$$

This gives: $[CH] = 3.0 \times 10^{11}$ cm^{-3}, which is also a hundredth of the APTES concentration. Then, the production of CH radicals can also be considered as a minor reaction pathway.

FTIR spectroscopy

FTIR measurements were performed in the range [500–4000 cm^{-1}] without and with APTES (Figure 3). In table 2, the functional groups in gaseous APTES were identified by comparison of FTIR data with those reported in the literature [36–47].

The absorption band at 785 cm^{-1} corresponds to NH$_2$ wagging [38]. This mode has been further confirmed by isotope exchange studies where a red shift of 145 cm^{-1} was observed for the ND$_2$ wagging mode [38]. This assignment is particularly important because, together with the shoulder at 2745 cm^{-1} which is attributed by White and Tripp [40] to Si–OH…NH$_2$, these peaks are the only way to follow the NH$_2$ group through reaction processes. Indeed, other bands like ν_s (N–H) around 3305 and 3350 cm^{-1} are not detected in our case. The absorption band at 1615 cm^{-1} represents the bending vibration of aliphatic amine (N–H) groups [39] but it is extremely weak and this absorption band cannot be kept to follow the evolution of the NH$_2$ group in the APTES molecule. It is also true for the band at 3219 cm^{-1} which is due to CH$_2$-NH$_2$ [36]. The band at 785 cm^{-1} is higher (typically a tenth of the peak at 1065 cm^{-1}, for instance) and characterized by a broad profile.

The band at 878 cm^{-1} is due to CH$_3$ rocking mode [42]. Si–O–C, Si–O in APTES and Si–O–C$_2$H$_5$ contribute to bands centred at 1110, 1065 and 958cm^{-1} [40, 42, 47, 48]. CH$_2$ rocking is observed at 1176 cm^{-1} [41, 42] whereas H–C–H bending is characterized by bands at 1475 and 1449 cm^{-1} [36, 43]. The absorption band at 1241 cm^{-1} corresponds to the asymmetric stretching of SiO–CH$_2$ [42]. CH$_2$ bending is observed at 1391 cm^{-1} and corresponds to a CH$_2$

group distant from Si [38, 39]. Si–O overtones are visible at 2264 and 2116 cm^{-1} [45]. Combinations bands centred at 1938 cm^{-1} are due to Si–O and ρ CH_3 [44]. Vibrations at 2987 cm^{-1}, 2939 cm^{-1}, 2899 cm^{-1} and 2870 cm^{-1} are commonly assigned to symmetric and asymmetric stretching of CH_2 and CH_3 [41, 45]. Finally, the absorption band at 3674 cm^{-1} is assigned to free OH [45], which is likely present because of some ageing of the precursor.

When the plasma is turned on, new contributions appear (figure 3). A very broad band centred at 3234 cm^{-1} spans from 3050 to 3550 cm^{-1}. It is attributed to normal "polymeric" OH stretch (>C=O···H–O) [45, 47]. Absorption bands at 2140 cm^{-1} and 1744 cm^{-1} are attributed to CO molecules and C=O stretching respectively [42]. At 2348 cm^{-1}, the characteristic absorption of CO_2 is easily identified. The contribution at 667 cm^{-1} is clearly due to δ CO_2 [46].

The interaction of the APTES with the Ar-O_2 afterglow is characterized by the formation of ketones resulting from the interaction of oxygen with carbon atoms of the precursor and the synthesis of CO and CO_2 as gaseous by-products. In the present conditions, the APTES concentration is roughly divided by a factor of 3 in afterglow when the plasma is ignited. It is also true for the amine group.

Evolutions in pulse mode

In pulse mode, it is possible to follow the time-resolved emission of emitting species. In figure 4, the time evolutions of CH, CN and OH normalized transitions together with the control pulse are depicted. After plasma ignition, active species are transported downstream to react with APTES. The delay of ~20 ms observed before emissive species start rising is caused by this transfer step. Next, a steady state is reached within 10 ms, leading to a plateau in emission. When the plasma is turned off, an overshoot in intensity is clearly visible for all species. This phenomenon is attributed to a sudden pressure variation. When the plasma stops, the gas is no longer heated, which leads to a pressure rise until pumping overcomes this

effect. The intensity of the overshoot is all the more reduced as the pressure decreases (unreported results), which supports this assumption. Taking the value of the intensity of each transition 50 ms after the rising edge of the pulse, *i.e.* in the middle of the plateau, we observe the behaviour described in figure 5 as a function of the duty cycle. Transition intensities evolve similarly, increasing first up to DC=40-50%, reaching next a maximum from DC=40-50 to 60, and slightly decreasing beyond DC=60%.

FTIR absorption spectra were recorded as a function of the duty cycle. In figure 6, the most intense absorption bands are reported (see **Supplemental material 4** for details). The CH absorption band that characterizes the non-oxidized APTES molecule decreases when the duty cycle increases. Oxidation of organosilicon compound is known to follow such a trend, the increase in the duty cycle leading to higher oxidation level and to a transition from plasma polymer to inorganic material [48–50]. Here, the transition occurs near DC=60%, in agreement with results obtained by time-resolved emission spectroscopy.

Nanoparticles synthesis

Thin film deposition on the reactor wall is a relatively slow process in the present condition because of the low partial pressure of the precursor. The deposition rate is about 150 ng mm^{-2} min^{-1} and then, no significant drift of the afterglow parameters is observed if the inner wall of the tube is cleaned between successive experiments. A thorough description of thin film deposition will be provided in a forthcoming publication. Here, emphasis is placed on the synthesis of nanoparticles in the gas phase. The process is about one order of magnitude slower than thin film deposition and strongly dependant on the duty cycle. After a two-hour treatment, nanoparticles collected on an aluminium substrate were analysed by scanning electron microscopy, transmission electron microscopy and FTIR absorption spectroscopy. In figure 7, SEM images of micrometric powders made of agglomerates of primary nanoparticles are depicted for two duty cycles: 20% and 70%. We notice that nanoparticles collected at low

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duty cycle exhibit a highly-porous structure contrary to those collected at high duty cycle, which look like denser. Corresponding TEM images show however similar features: agglomerates have diameters of ~200 nm and are made of an assembly of amorphous nanoparticles with diameters of a few nanometres. These results are quite similar to those presented by Roth *et al.* with other organosilicon precursors [51].

FTIR analysis of nanoparticles shows also strong changes in absorption bands (figure 8 and table 3) as a function of the duty cycle. We observe in figure 9, that FTIR absorbance spectra of the nanoparticles exhibit similar features with decaying intensity when the duty cycle increases – see **supplemental material 5** for a deconvolution of the bands between 1800 and 800 cm^{-1} –. First, the Si–O–C_2H_5 absorption band in APTES appears at 950 cm^{-1}. It is a footprint of the precursor together with bands at 1169 and 1083 cm^{-1} designated hereafter as "Si–O in APTES" [42]. At 1376 cm^{-1}, δ CH_2 bonds are likely. The contribution at 1441 cm^{-1}, is assigned to δ C–O–H. δ NH_2 (the amide II band) in primary amides appears as a shoulder at 1620 cm^{-1} of the peak at 1698 cm^{-1} [46]. The contribution at 1698 cm^{-1} is attributed to ν CO (amide I band) vibration, but not to pure ketones commonly found at 1730 cm^{-1} [46]. Vibrations at 2930 cm^{-1} and 2870 cm^{-1} are convolutions of respectively symmetric and asymmetric stretching of CH_2 and CH_3 [41, 45]. The absorption band with its maximum at 3250 cm^{-1} and spanning over the range [3120–3350 cm^{-1}] is assigned to ν_s and ν_{as} NH_2. Indeed, primary amides display two strong NH_2 stretching bands, *i.e.* asymmetric stretching at 3360–3340 cm^{-1} and symmetric stretching at 3190–3170 cm^{-1} [42]. Finally, the other very broad band centred at 3500 cm^{-1} spans from 3350 to 3680 cm^{-1} is attributed to OH stretch [42]. EDX analyses of the powders (see **supplemental material 6**) confirm the presence of nitrogen at low duty cycle.

The striking feature is the change in the nanoparticle composition as the duty cycle increases. The carbon content becomes weaker but non-negligible, even at DC=70% (see **supplemental**

material 6) compared to 20%, leading to nanoparticles principally made of Si–O–C chains. XPS spectra confirm this trend (figure 10). We clearly observe the presence of nitrogen, even though the signal becomes noisy beyond DC=50%. N–C bonds (contribution at 400.1 eV) dominate but nitrogen can be also slightly oxidized (weaker contribution at 402.0 eV). Si–O–C chains are mainly found at 101.9 eV, the silica-like form of silica appearing as a shoulder at 103.6 eV. The O(1s) signal is interpreted accordingly as being the sum of 3 different peaks: -C=O at 533.1 eV, O–H at 531.8 eV and O–Si at 530.3 eV. Concerning the C(1s) signal, it is worth mentioning here that no significant C=O contribution, expected around 287 eV, was found. This fourth contribution could have been enforced, but this would have led to a minor peak. Besides, the relatively broad contribution at 288.7 eV likely contains a mixture of O–C=O and N–C=O contributions. C–O and C–(C,H) bonds are associated with peaks at 285.6 and 284.6 eV respectively. So, all in all, XPS data confirm well FTIR results.

The longer oxidation process of APTES leads to an inorganic material instead of a plasma polymer-like material, a feature that is common to most organosilicon compounds. FTIR spectra show that the C–O–H and Si–O–C absorption bands at 1441 and 950 cm^{-1} respectively are still present, but the main Si–O contributions at 1169 and 1083 cm^{-1} are barely observed. Obviously, the C–Si≡O$_3$ environment, present in APTES, is strongly affected by the duty cycle increase. Concerning the band at 950 cm^{-1}, we cannot exclude a contribution of Si–OH. However, it should be accompanied by the presence of the OH vibration band at 3500 cm^{-1}, which is not observed. So, this contribution is certainly minor. No intense contribution is found in the range [1000-1200] cm^{-1} in the present conditions, showing that nanoparticles are not silica-like.

To sum up, the synthesized nanoparticles are amorphous. They have a plasma polymer-like structure with a silicon backbone and a high amount of amide groups at low duty cycle. At high duty cycle, the nanoparticles still contain some carbon and are not silica-like.
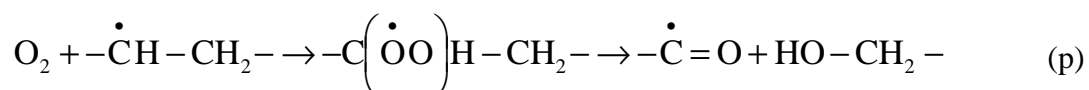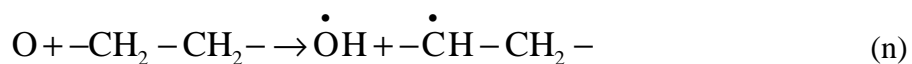
Mechanisms at stake

As described previously, the synthesis of C_2H, CH and CN radicals is very limited. As no N–O bond could be observed, the amine group is assumed to be affected to a limited extent by oxygen, which explains the formation of primary amides in the coating especially at low duty cycle.

The initiation step of APTES decomposition is attributed to atomic oxygen. To the best of our knowledge, contrary to O atoms [52-55], the singlet state $O_2(a^1\Delta_g)$ has never been identified as a reactive species with regards to dehydrogenation of organosilicon compounds. Afterglow studies with TEOS [52-54] and HMDSO [48, 53] all reinforce this conclusion.

Wrobel *et al.* [52] showed, in the case of a remote oxygen Plasma-CVD process, that the growth rate of silica film from TEOS is proportional to the concentration of atomic oxygen. The abstraction of the hydrogen atom from the TEOS molecule is a major reaction to the precursor decomposition. The value of the overall kinetic constant of oxygen atoms with TEOS molecules k_O at room temperature strongly differs according to the authors. k_O was reported to be equal to 6×10^{-15} cm^3 s^{-1} by [56], 4×10^{-14} cm^3 s^{-1} by Janča *et al.* [53] and 4×10^{-13} cm^3 s^{-1} by Sanogo and Zachariah [57]. So, these values are not helpful to evaluate the reactivity of oxygen atoms with respect to TEOS, a molecule which is relatively similar to APTES. Levedakis *et al.* [54] used a remote microwave plasma and compared the predicted oxygen atom concentrations with corresponding deposition rates. They found out that the overall SiO_2 deposition reaction is largely controlled by the concentration of oxygen atoms. Finally, Georg *et al.* [55] proposed that a major reason for thin films improvement of roughness and homogeneity is the enhanced oxidation of unwanted carbohydrogen compounds in the HMDSO molecules by atomic oxygen.

We infer from the synthesis of >CO…OH bonds (figure 3c) and the production of C=O in the precursor and/or its fragments that after OH abstraction by O atoms (process n), O_2 (in any

state, including the singlet state in this case) reacts with the as-formed radical to produce a peroxide, whose lability leads to the scission of C–C bonds according to a mechanism of the following kind:

$$O + -CH_2 - CH_2 - \rightarrow \overset{\bullet}{O}H + -\overset{\bullet}{C}H - CH_2 - \tag{n}$$

$$O_2 + -\overset{\bullet}{C}H - CH_2 - \rightarrow -C\left(\overset{\bullet}{O}O\right)H - CH_2 - \rightarrow -\overset{\bullet}{C} = O + HO - CH_2 - \tag{p}$$

Further chemical processes between APTES fragments are not accessible without time-resolved FTIR measurements. The description of the processes will require new investigations. Nevertheless, when the duty cycle increases, fragmentation of the precursor increases since oxidation proceeds longer. Then, reactions between shorter silicon radicals, still attached to carbon-containing groups, occur. This leads to a plasma-polymer material characterized by a non-negligible amount of carbon.

The fact that nanoparticles are not silica-like at high duty cycle shows the specificity of chemical reactions in the gas phase with regard to surface processes at the reactor wall. Indeed, we know from other experiments, which are not reported here, that in our conditions silica-like coatings are obtained on the reactor wall at high duty cycle. Carbon etching by oxygen is much more efficient on the growing coating than on the formed nanoparticles. The reason for this is still to be clarified.

CONCLUSION

The interaction of (3–Aminopropyl)triethoxysilane with a pulse late Ar-O_2 afterglow is characterized by the synthesis of OH, CO and CO_2 in the gas phase as main by-products. Other minor species like CH, CN and C_2H are also produced. We suggest that OH radicals are produced in a first step by dehydrogenation of APTES after interaction with oxygen atoms. In

a second step, the molecule is oxidized by any state of O_2, *i.e.* not only the ground state but also the singlet state of oxygen, for instance. The as-formed peroxides transform into by-products, break thus the C–C bonds in the precursor. If oxidation is limited, thanks to a low duty cycle, fragmentation of the precursor is limited and nanoparticles produced out of them have a composition comparable to the precursor, even though the original –NH$_2$ groups are efficiently converted into amide groups. At high duty cycle, silicon-containing fragments contain also some carbon and react together and produce nanoparticles with a non-silica-like structure at least in our experimental conditions.

It turns out that, in the present condition, a high retention level of amines in the precursor is not possible, even in pulsed conditions. Although afterglows are soft media, the behaviour of oxygen atoms toward APTES is defined once and for all and cannot be adjusted by simply changing the duty cycle. In this sense, pulsed plasmas are likely more suited to this purpose thanks to electronic collisions which should offer a more controllable way to define the dissociation pathway of APTES by acting on the electron energy.

ACKNOWLEDGMENTS

REFERENCES

1. Lecoq E., Duday D., Bulou S., Frache G., Hilt F., Maurau R., Choquet P. (2013) Plasma Polymerization of APTES to Elaborate Nitrogen Containing Organosilicon Thin Films: Influence of Process Parameters and Discussion About the Growing Mechanisms. Plasma Process Polym 10(3):250-261. doi:10.1002/ppap.201200108

2. Minier M., Salmain M., Yacoubi N., Barbes L., Methivier C., Zanna S., Pradier C. M. (2005) Covalent immobilization of lysozyme on stainless steel. Interface spectroscopic characterization and measurement of enzymatic activity. Langmuir 21(13): 5957-5965. doi:10.1021/la0501278

3. Schröder K., Finke B., Polak M., Lüthen F., Nebe J.B., Rychly J., Bader R., Lukowski G., Walschus U., Schlosser M., Ohl A., Weltmann K.-D. (2010) Gas-discharge plasma-assisted functionalization of titanium implant surfaces. Mater Sci Forum 638-642:700-705. doi:10.4028/www.scientific.net/MSF.638-642.700

4. Nebe B., Finke B., Luthen F., Bergemann C., Schroder K., Rychly J., Liefeith K., Ohl A. (2007) Improved initial osteoblast functions on amino-functionalized titanium surfaces. Biomol Eng 24(5):447-454. doi:10.1016/j.bioeng.2007.07.004

5. Borges J. N., Belmonte T., Guillot J., Duday D., Moreno-Couranjou M., Choquet P., Migeon H. N. (2009) Functionalization of Copper Surfaces by Plasma Treatments to Improve Adhesion of Epoxy Resins. Plasma Process Polym. 6(S1):S490-S495. doi: 10.1002/ppap.200931106

6. Friedrich J. F., Mix R., Kuhn G. (2003) Functional groups bearing plasma homo and copolymer layers as adhesion promoters in metal–polymer composites. Surf Coat Technol 174-175, 811-815. doi: 10.1016/S0257-8972(03)00350-5

7. Jampala S. N., Sarmadi M., Manolache S., Denes F. S. (2008) Surface functionalization by RF plasma deposition of ethylene diamine, acrylonitrile, and acetonitrile. J Appl Polymer Sci 107(3):1686-1695. doi:10.1002/app.27189

8. Kuhn G., Retzko I., Lippitz A., Unger W., Friedrich J. (2001) Homofunctionalized polymer surfaces formed by selective plasma processes. Surf Coat Technol 142-144:494-500. doi:10.1016/S0257-8972(01)01056-8

9. Rinsch C. L., Chen X., Panchalingam V., Eberhart R. C., Wang J. H., Timmons R. B. (1996) Pulsed radio frequency plasma polymerization of allyl alcohol: controlled deposition of surface hydroxyl groups. Langmuir 12(12):2995-3002. doi:10.1021/la950685u

10. Stine R., Cole C. L., Ainslie K. M., Mulvaney S. P., Whitman L. J. (2007) Formation of primary amines on silicon nitride surfaces: a direct, plasma-based pathway to functionalization. Langmuir 23(8):4400-4404. doi:10.1021/la0635653

11. Angelini E., d'Agostino R., Fracassi F., Grassini S., Rosalbino F. (2002) Surface analysis of PECVD organosilicon films for corrosion protection of steel substrates. Surf Interface Anal 34(1):155-159. doi:10.1002/sia.1273

12. Fracassi F., d'Agostino R., Palumbo F., Angelini E., Grassini S., Rosalbino F. (2003) Application of plasma deposited organosilicon thin films for the corrosion protection of metals. Surf. Coat. Technol. 174-175:107-111. doi:10.1016/S0257-8972(03)00422-5

13. Simor M., Fiala A., Kovacik D., Hlidek P., Wypkema A., Kuipers R. (2007) Corrosion protection of a thin aluminium layer deposited on polyester. Surf Coat Technol 201(18):7802-7812. doi:10.1016/j.surfcoat.2007.03.011

14.   Goddard J. M., Hotchkiss J.H. (2007) Polymer surface modification for the attachment of bioactive compounds. Prog. Polym. Sci. 32(7):698-725. doi:10.1016/j.progpolymsci.2007.04.002

15.   Witucki G. L. (1993) A silane primer: chemistry and applications of alkoxy silanes. J. Coat. Technol. 65(822):57-60.

16.   Jampala S. N., Sarmadi M., Somers E. B., Wong A. C. L., Denes F. S. (2008) Plasma-enhanced synthesis of bactericidal quaternary ammonium thin layers on stainless steel and cellulose surfaces. Langmuir 24(16):8583-8591. doi:10.1021/la800405x

17.   Gandhiraman R. P., Gubala V., Nam L. C. H., Volcke C., Doyle C., James B., Daniels S., Williams D. E. (2010) Deposition of chemically reactive and repellent sites on biosensor chips for reduced non-specific binding. Colloids Surf. B Biointerfaces 79(1):270-275. doi:10.1016/j.colsurfb.2010.04.009

18.   Gandhiraman R. P., Volcke C., Gubala V., Doyle C., Basabe-Desmonts L., Dotzler C., Toney M. F., Iacono M., Nooney R. I., Daniels S., James B., Williams D. E. (2010) High efficiency amine functionalization of cycloolefin polymer surfaces for biodiagnostics. J. Mater. Chem. 20(20):4116-4127. doi:10.1039/B925737C

19.   Gubala V., Gandhiraman R. P., Volcke C., Doyle C., Coyle C., James B., Daniels S., Williams D. E. (2010) Functionalization of cycloolefin polymer surfaces by plasma-enhanced chemical vapour deposition: comprehensive characterization and analysis of the contact surface and the bulk of aminosiloxane coatings. Analyst 135, 1375-1381. doi:10.1039/B924692D

20.   Volcke, C., Gandhiraman, R. P., Gubala, V., Raj, J., Cummins, T., Fonder, G., Nooney R.I., Mekhalif Z., Herzog G., Daniels S., Arrigan D.W.M., 2, Cafolla A.A., Williams D.E. (2010) Reactive amine surfaces for biosensor applications, prepared by plasma-

enhanced chemical vapour modification of polyolefin materials. Biosensors Bioelectron. 25(8):1875-1880. doi:10.1016/j.bios.2009.12.034

21.   Volcke C., Gandhiraman R. P., Gubala V., Doyle C., Fonder G., Thiry P. A., Cafolla A. A., James B., Williams D. E. (2010) Plasma functionalization of AFM tips for measurement of chemical interactions. J. Colloid Interface Sci. 348(2):322-328. doi:10.1016/j.jcis.2010.04.042

22.   Sakata J., Wada M. (1988) Preparation of ion exchange membranes by plasma polymerization. I. J. Appl. Polym. Sci. 35(4):875-884. doi:10.1002/app.1988.070350403

23.   Ricard A., Moisan M., Moreau S. (2001) Determination, through titration with NO, of the concentration of oxygen atoms in the flowing afterglow of Ar-O_2 and N_2-O_2 plasmas used for sterilization purposes. J. Phys. D: Appl. Phys. 34(8):1203-1212. doi:10.1088/0022-3727/34/8/311

24.   Mafra M., Belmonte T., Poncin-Epaillard F., Maliska A., Cvelbar U. (2009) Treatment of hexatriacontane by Ar–O_2 remote plasma: formation of the active species. Plasma Process. Polym. 6(S1):S198-S203. doi:10.1002/ppap.200932406

25.   Devriendt K., Peeters J. (1997) Direct Identification of the C_2H ($X^2\Sigma^+$) + O (^3P) $\rightarrow$ CH($A^2\Delta$)+ CO Reaction as the Source of the CH($A^2\Delta \rightarrow X^2\Pi$) Chemiluminescence in C_2H_2/O/H Atomic Flames. J. Phys. Chem. A 101(14):2546-2551. doi:10.1021/jp963434i

26.   Grebe J., Homann K. H. (1982) Kinetics of the Species OH($A^2\Sigma^+$), OH($X^2\Pi$) and CH ($X^2\Pi$) in the System C_2H_2/O/H. Berichte Bunsenges Phys. Chem., 86(7):581-587. doi:10.1002/bbpc.19820860702

27. Hidaka Y., Kawano H., Suga M. (1982) Additive Effect of CF_3Cl on an Oxidation Reaction in Shock Waves. I. Study with H_2–O_2. Bull. Chem. Soc. Jap. 55(2):351-357. doi:10.1246/bcsj.55.351

28. Elsamra R. M. I., Vranckx S., Carl S. A. (2005) CH ($A^2\Delta$) formation in hydrocarbon Combustion: the temperature dependence of the rate constant of the reaction $C_2H + O_2 \rightarrow$ CH ($A^2\Delta$)+ CO_2. J. Phys. Chem. A 109(45):10287-10293. doi:10.1021/jp053684b

29. Chen C., Sheng Y., Yu S., Ma X. (1994) Investigation of the collisional quenching of CH ($A^2\Delta$ and $B^2\Sigma^-$) by Ar, O_2, CS_2, alcohol, and halomethane molecules. J. Chem. Phys. 101(7):5727-5730. doi:10.1063/1.467358

30. Setser D. W., Thrush B. A. (1965) Kinetics of reactions involving CN emission. II. The reaction between oxygen atoms and cyanogen. Proc. R. Soc. Lond. A Math. Phys. Sci., 288(1413):275-291. doi:10.1098/rspa.1965.0218

31. Knowles P. J., Werner H. J., Hay P. J., Cartwright D. C. (1988) The $A^2\Pi$–$X^2\Sigma$+ red and $B^2\Sigma^+$–$X^2\Sigma^+$ violet systems of the CN radical: Accurate multireference configuration interaction calculations of the radiative transition probabilities. J. Chem. Phys. 89(12):7334-7343. doi:10.1063/1.455264

32. Tereshchenko E. N., Dodonova N. I., Photoprocesses in the gas phase (Fotoprotsessy v gazovoi faze). Leningrad, Izdatel'stvo Leningradskogo Universiteta (Leningradskii Gosudarstvennyi Universitet, Uspekhi Fotoniki, No. 8), 1983, p. 112-145. In Russian., 1, 112-145.

33. Carl S. A., Van Poppel M., J. Peeters (2003) Identification of the CH + $O_2 \rightarrow$ OH (A) + CO Reaction as the Source of OH (AX) Chemiluminescence in C_2H_2/O/H/O_2 Atomic Flames and Determination of its Absolute Rate Constant over the Range T= 296 to 511 K. J. Phys. Chem. A 107(50):11001-11007. doi:10.1021/jp035568j

34. Leblond J. B., Collier F., Hoffbeck F., Cottin P. (1981) Kinetic study of high-pressure Ar–H2O mixtures excited by relativistic electrons. J. Chem. Phys. 74(11):6242-6255. doi:10.1063/1.441015

35. Fairchild P. W., Smith G. P., Crosley D. R. (1983) Collisional quenching of $A^2\Sigma^+$ OH at elevated temperatures. J. Chem. Phys. 79(4):1795-1807. doi:10.1063/1.446025

36. Weigel Ch., Kellner R. (1989) FTIR-ATR-spectroscopic investigation of the silanization of germanium surfaces with 3-aminopropyltriethoxysilane. Fresenius Z. Anal. Chem. 335(7):663-668. doi:10.1007/BF01204067

37. Finocchio E., Macis E., Raiteri R., Busca G. (2007) Adsorption of trimethoxysilane and of 3-mercaptopropyltrimethoxysilane on silica and on silicon wafers from vapor phase: an IR study. Langmuir 23(5):2505-2509. doi:10.1021/la062972b

38. Rai V. R., Agarwal S. (2011) Mechanism of self-catalytic atomic layer deposition of silicon dioxide using 3-aminopropyl triethoxysilane, water, and ozone. Chem. Mater. 23(9):2312-2316. doi:10.1021/cm103052t

39. Rahman I.A., Jafarzadeh M., Sipaut C.S. (2009) Synthesis of organo-functionalized nanosilica via a co-condensation modification using γ-aminopropyltriethoxysilane (APTES). Ceram. Int., 35(5):1883-1888. doi:10.1016/j.ceramint.2008.10.028

40. White L. D., Tripp C. P. (2000) Reaction of (3-aminopropyl) dimethylethoxysilane with amine catalysts on silica surfaces. J. Colloid Interf. Sci. 232(2):400-407. doi:10.1006/jcis.2000.7224

41. Aissaoui N., Bergaoui L., Landoulsi J., Lambert J.-F., Boujday S. (2012) Silane layers on silicon surfaces: mechanism of interaction, stability, and influence on protein adsorption. Langmuir 28(1), 656-665. doi:10.1021/la2036778

42. Launer P. J. (1987) "Infrared analysis of organosilicon compounds: spectra-structure correlations", Silicone compounds register and review, 100

43. Okabayashi H., Shimizu I., Nishio E., O'Connor C. J. (1997) Diffuse reflectance infrared Fourier transform spectral study of the interaction of 3-aminopropyltriethoxysilane on silica gel. Behavior of amino groups on the surface. Colloid Polym. Sci. 275(8):744-753. doi:10.1007/s003960050143

44. Workman Jr J., Weyer L. (2007) "Practical guide to interpretive near-infrared spectroscopy", edited by CRC press.

45. Coates J. (2000) "Interpretation of infrared spectra, a practical approach", Encyclopedia of analytical chemistry, edited by John Wiley & Sons Ltd, Chichester.

46. Stuart B. H. (2004) "Infrared Spectroscopy: Fundamentals and Applications", Analytical Techniques in the Science, edited by John Wiley & Sons Ltd, Chichester.

47. Liakath Ali Khan F., Sivagurunathan P., Asghar J. (2008) FTIR study of hydrogen bonding interactions between alkyl esters and hexanol, p-cresol in carbon tetrachloride. Indian J. Pure Appl. Phys. 46(1):12.

48. Goujon M., Belmonte T., Henrion G. (2004) OES and FTIR diagnostics of HMDSO/O_2 gas mixtures for SiO_x deposition assisted by RF plasma. Surf. Coat. Technol. 188–189:756-761. doi:10.1016/j.surfcoat.2004.07.048

49. Bousquet A., Cartry G., Granier A. (2007) Investigation of O-atom kinetics in O_2, CO_2, H_2O and O_2/HMDSO low pressure radiofrequency pulsed plasmas by time-resolved optical emission spectroscopy. Plasma Sources Sci. Technol. 16(3):597-605. doi:10.1088/0963-0252/16/3/020

50. Supiot P., Vivien C., Blary K., Rouessac V. (2011) Organosilicon Polymers Deposition by PECVD and RPECVD on Micropatterned Substrates. Chem. Vap. Depos. 17(10-12):321-326. doi:10.1002/cvde.201106902

51. Roth C., Oberbossel G., Buitrago E., Heuberger R., Rudolf von Rohr P. (2012) Nanoparticle synthesis and growth in a continuous plasma reactor from organosilicon precursors. Plasma Process. Polym. 9(2):119-134. doi:10.1002/ppap.201100180

52. Wrobel A. M., Walkiewicz-Pietrzykowska A., Hatanaka Y., Wickramanayaka S., Nakanishi Y. (2001) Oligomerization and polymerization steps in remote plasma chemical vapor deposition of silicon-carbon and silica films from organosilicon sources. Chem. Mater. 13(5):1884-1895. doi:10.1021/cm001044+

53. Janča J., Tálský A., Zvoníček V. (1996) Kinetics of O_2 + TEOS gas-phase chemical reactions in a remote RF plasma reactor with electron spin resonance. Plasma Chem. Plasma Proc. 16(2):187-194. doi:10.1007/BF01570177

54. Levedakis D. A., Raupp G. B. (1992) "Oxygen Atom Induced Deposition of Silicon Dioxide." MRS Proceedings (Vol. 282, p. 537). Cambridge University Press.

55. Georg A., Engemann J., Brockhaus A. (2002) Investigation of a pulsed oxygen microwave plasma by time-resolved two-photon allowed laser-induced fluorescence. J. Phys. D: Appl. Phys. 35(9):875-881. doi:10.1088/0022-3727/35/9/307

56. Kawahara T., Yuuki A., Matsui Y. (1992) Reaction mechanism of chemical vapor deposition using tetraethylorthosilicate and ozone at atmospheric pressure. Jpn. J. Appl. Phys. 31(9R):2925-2930. doi:10.1143/JJAP.31.2925

57. Sanogo O., Zachariah M. R. (1997) Kinetic studies of the reaction of tetraethoxysilane with oxygen atoms. J. Electrochem. Soc. 144(8):2919-2923. doi:10.1149/1.1837918

FIGURE CAPTION

Figure 1: The experimental setup.

Figure 2: Low-resolution optical emission spectra of the Ar-O$_2$ afterglow with and without APTES.

Figure 3: Gas phase FTIR spectra of the Ar-APTES flow without (red) and with (black) the Ar-O$_2$ afterglow. Resolution: 2 cm^{-1}. a) Infrared region: from 1700 to 500 cm^{-1}. b) Infrared region: from 2600 to 1700 cm^{-1}. The signal obtained with the Ar-O$_2$ afterglow was divided by a factor of 2 to better scale the figure and show details of the other signal recorded without the afterglow. c) Infrared region: from 4000 to 2600 cm^{-1}.

Figure 4: Time evolution of CH, CN and OH normalized transitions together with the control pulse (Duty cycle = 60%, P = 15 mbar, W = 250 W, plasma gas flowrates: Ar =1050 sccm, O$_2$= 100 sccm. Precursor flowrates: Ar =30 sccm, APTES = 0.13 sccm).

Figure 5: Evolution of CH, CN and OH transitions taken 50 ms after the pulse rising edge as a function of the duty cycle (same conditions as in figure 4).

Figure 6: Evolution of the most intense FTIR absorption bands as a function of the duty cycle (same conditions as in figure 4).

Figure 7: TEM and SEM images of nanoparticles collected on an aluminium substrate at duty cycles equal to 20% and 70%.

Figure 8: FTIR spectrum of solid nanoparticles collected in the gas phase at duty cycle equal to 20%. Infrared region: from 3750 to 750 cm^{-1}. Resolution: 2 cm^{-1}.

Figure 9: FTIR spectra of solid nanoparticles collected in the gas phase at different duty cycles (from 20 to 70%). Infrared region: from 3750 to 750 cm^{-1}. Resolution: 2 cm^{-1}.

Figure 10: O(1s), N(1s), C(1s) and Si(2p) XPS spectra of nanoparticles collected on an aluminium substrate for different duty cycles.

Table 1: List of the transitions observed by optical emission spectroscopy in the Ar-O$_2$ afterglow with and without APTES.

Table 2: Vibrational groups and their corresponding frequencies observed in the APTES flow with (bold) and without Ar-O$_2$ afterglow.

Table 3: Main vibrational groups and their corresponding frequencies observed in nanoparticles produced from APTES for different duty cycles (from 20 to 70%).
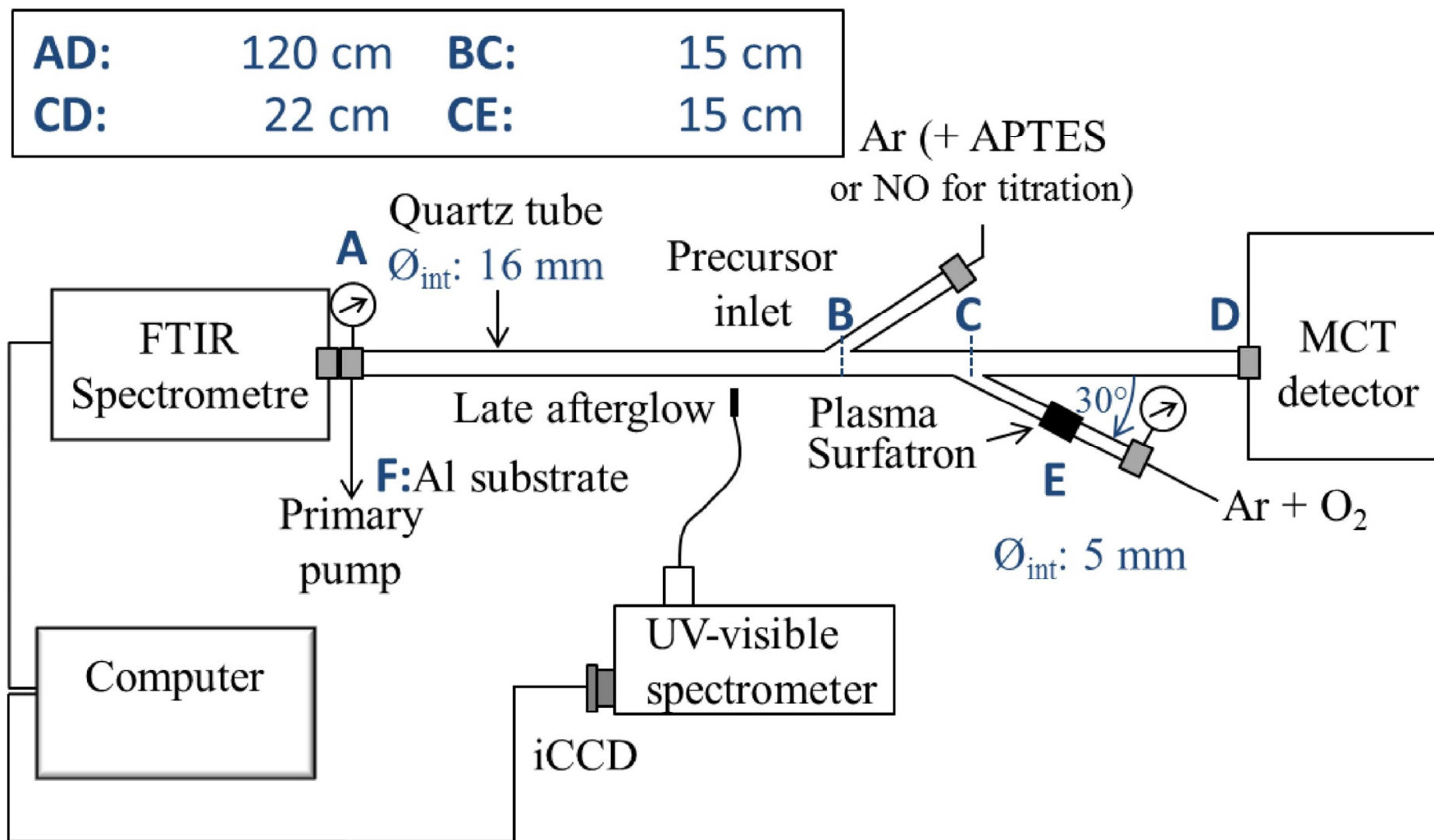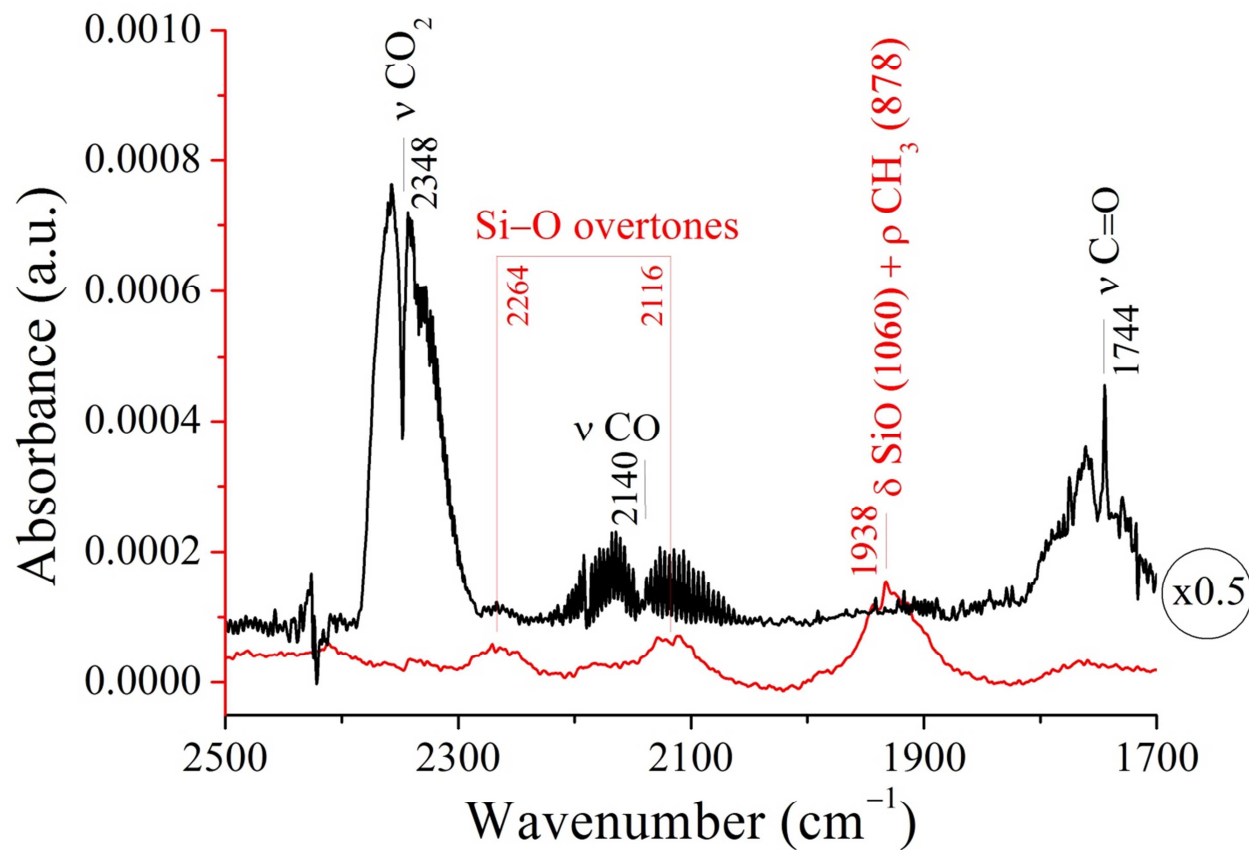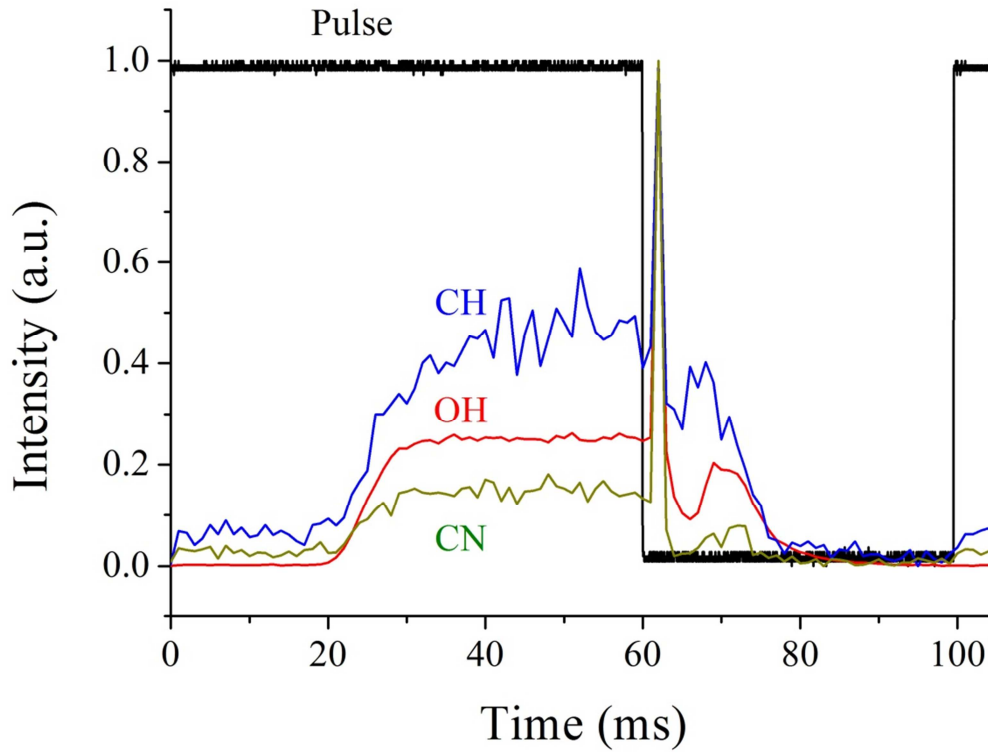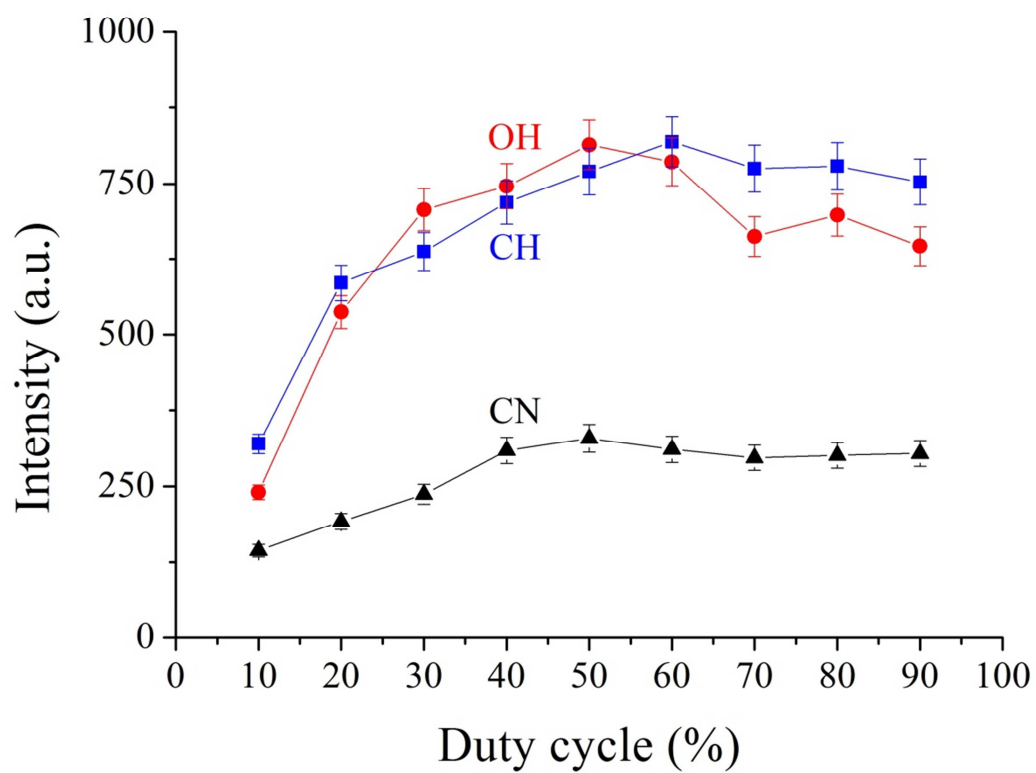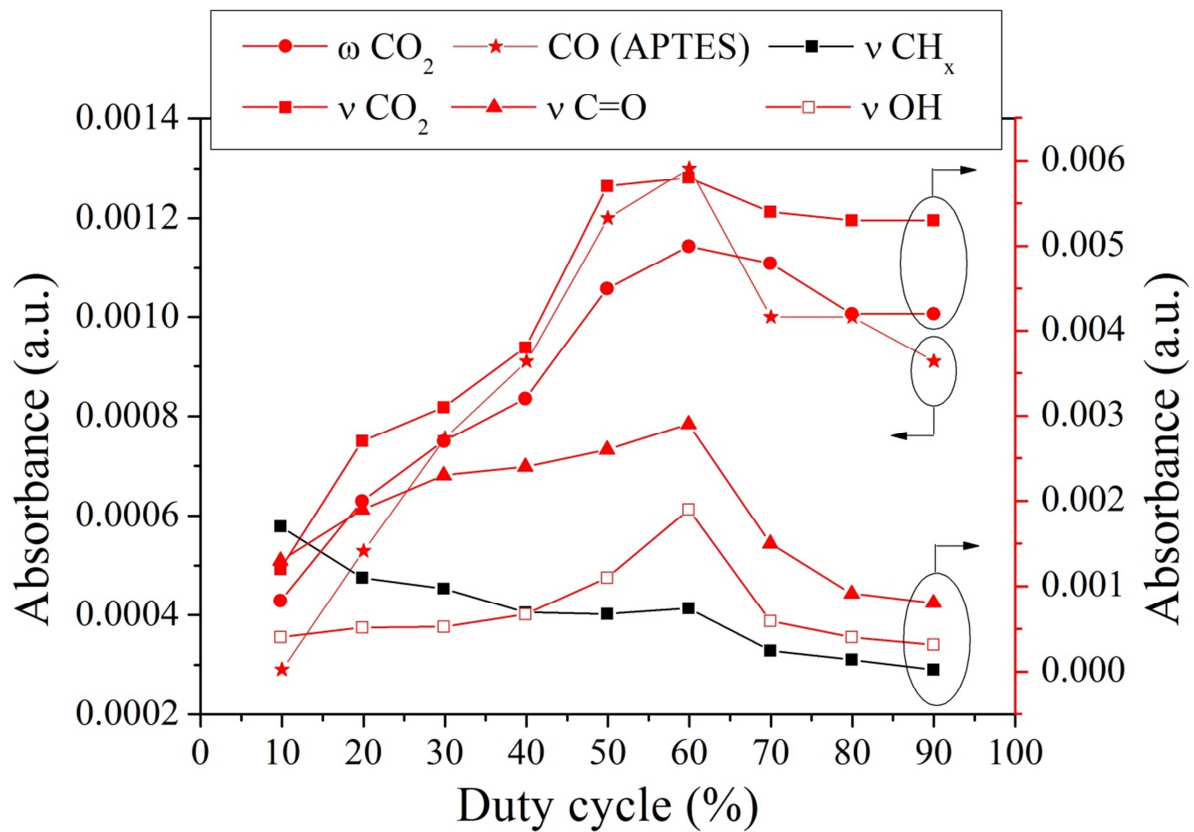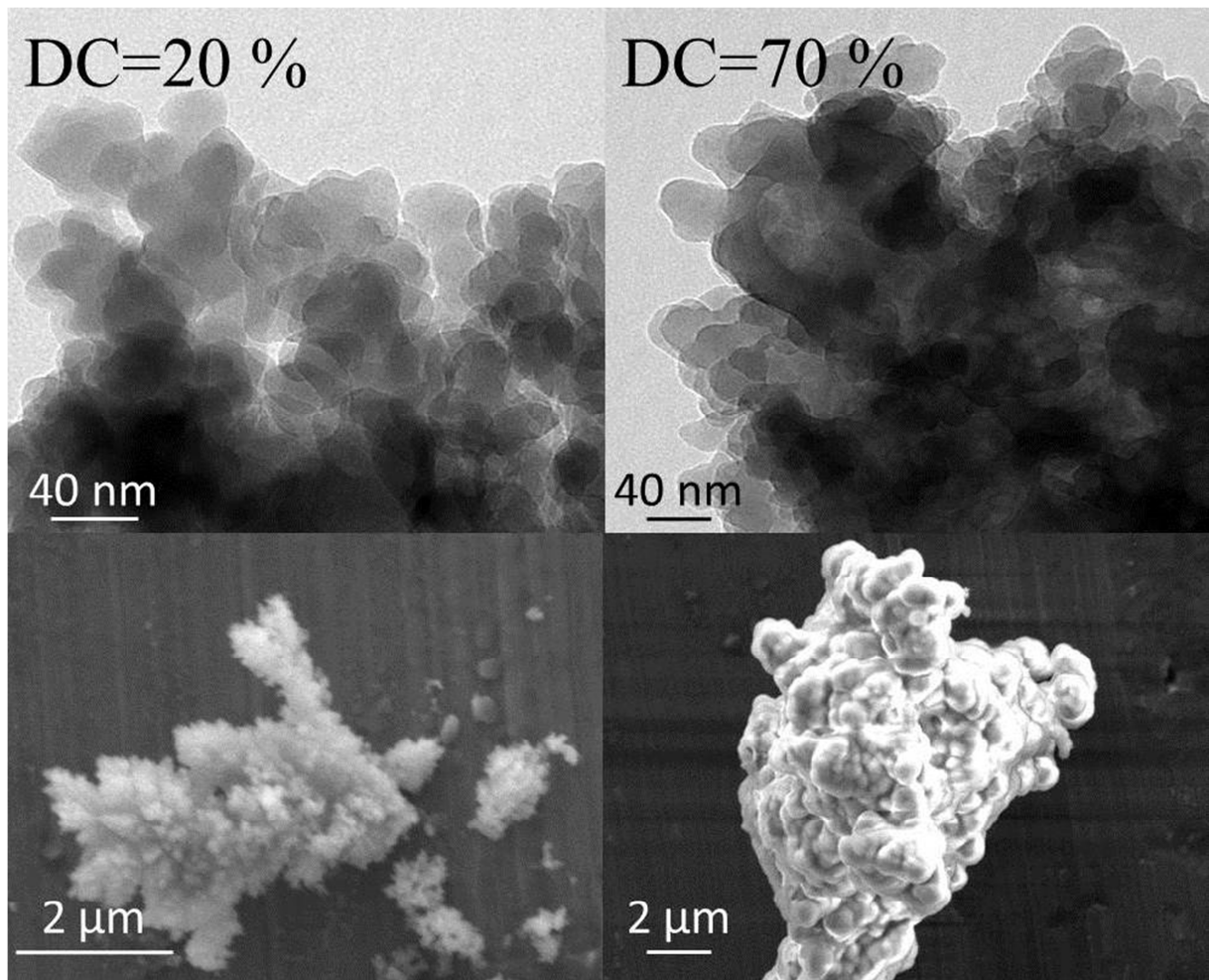
Figure 1: The experimental setup.
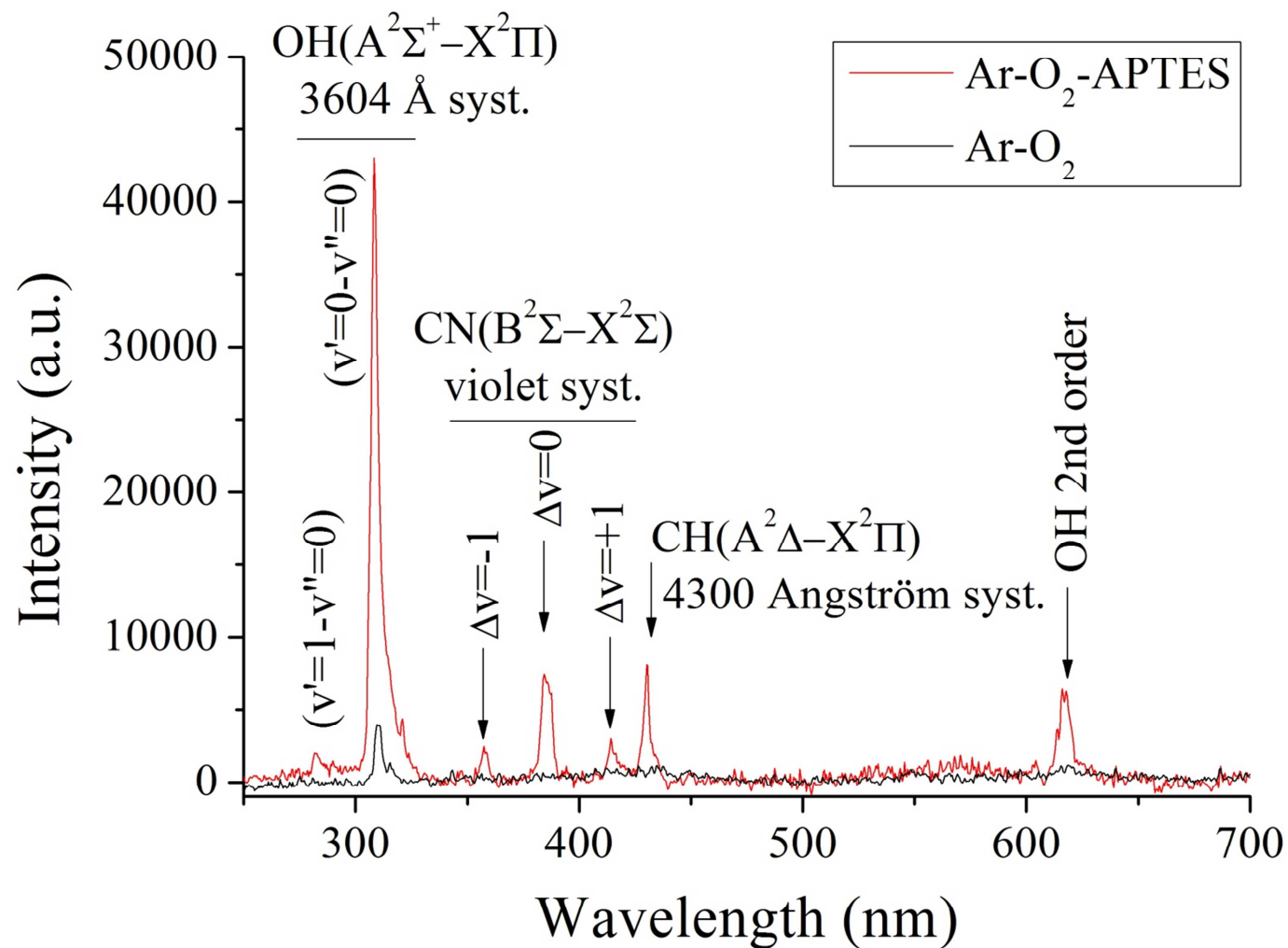
Figure 2: Low-resolution optical emission spectra of the Ar-O₂ afterglow with and without APTES.
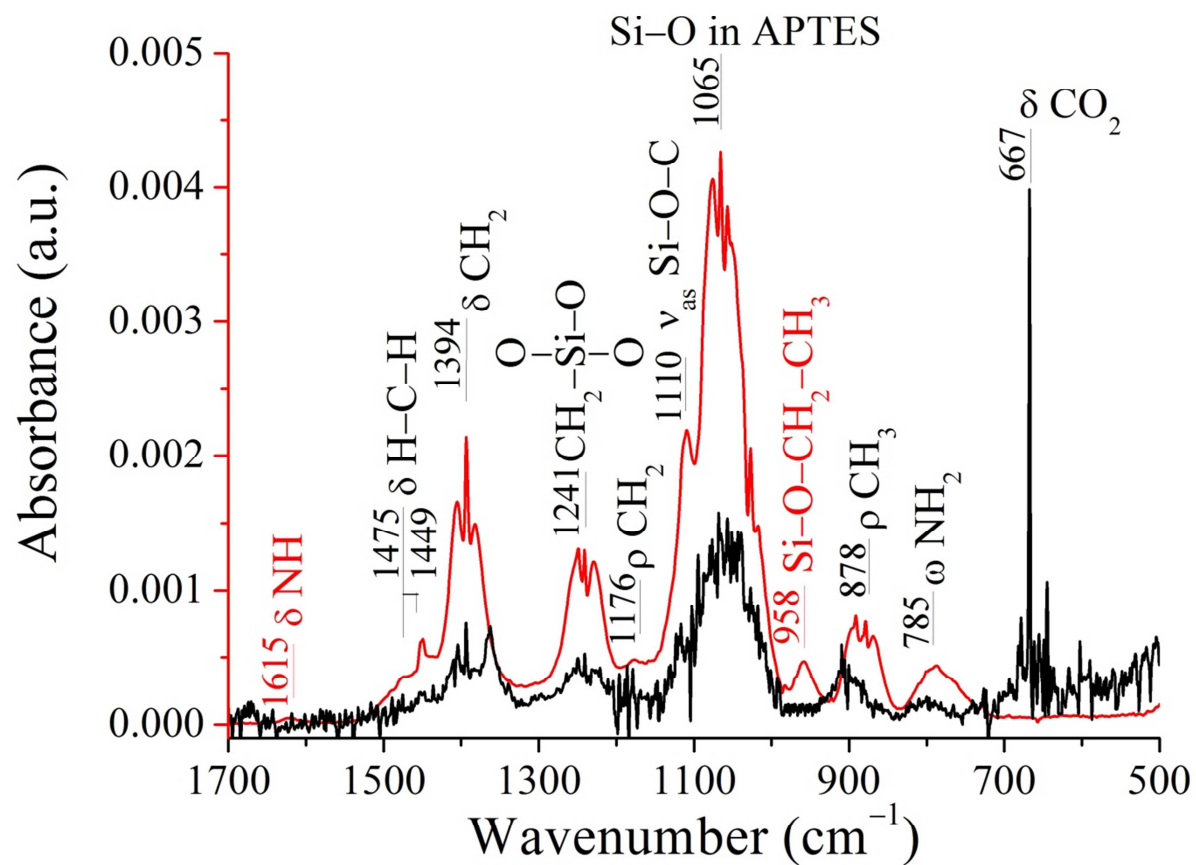
Figure 3a: Gas phase FTIR spectra of the Ar-APTES flow without (red) and with (black) the Ar-O$_2$ afterglow. Resolution: 2 cm^{-1}. a) Infrared region: from 1700 to 500 cm^{-1}.

Figure 3b: Gas phase FTIR spectra of the Ar-APTES flow without (red) and with (black) the Ar-O_2 afterglow. Resolution: 2 cm^{-1}. b) Infrared region: from 2600 to 1700 cm^{-1}. The signal obtained with the Ar-O_2 afterglow was divided by a factor of 2 to better scale the figure and show details of the other signal recorded without the afterglow.
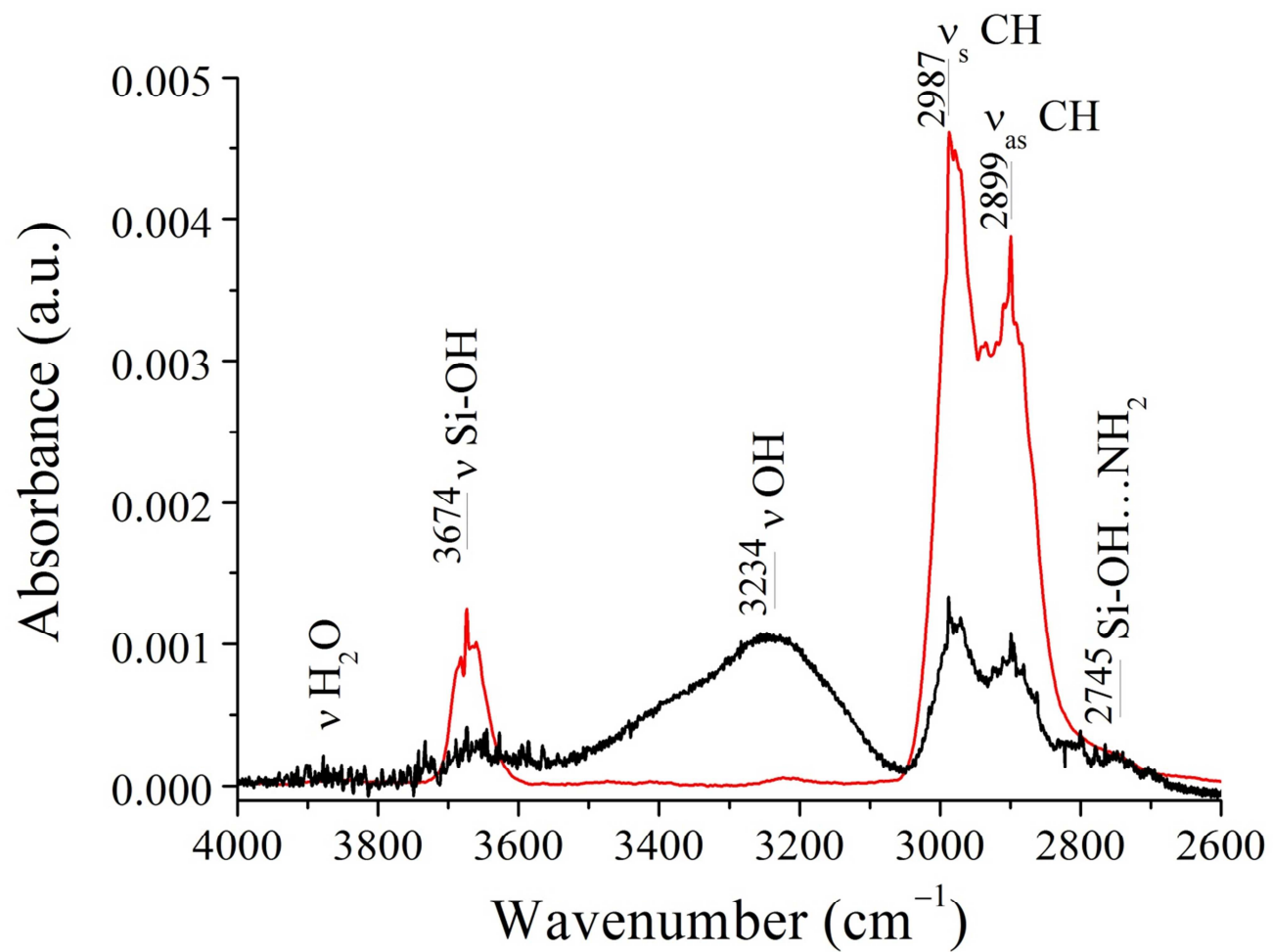
Figure 3c: Gas phase FTIR spectra of the Ar-APTES flow without (red) and with (black) the Ar-O$_2$ afterglow. Resolution: 2 cm^{-1}. c) Infrared region: from 4000 to 2600 cm^{-1}.

Figure 4: Time evolution of CH, CN and OH normalized transitions together with the control pulse (Duty cycle = 60%, P = 15 mbar, W = 250 W, plasma gas flowrates: Ar =1050 sccm, O_2= 100 sccm. Precursor flowrates: Ar =30 sccm, APTES = 0.13 sccm).

Figure 5: Evolution of CH, CN and OH transitions taken 50 ms after the pulse rising edge as a function of the duty cycle (same conditions as in figure 4).

Figure 6: Evolution of the most intense FTIR absorption bands as a function of the duty cycle (same conditions as in figure 4).

Figure 7: TEM and SEM images of nanoparticles collected on an aluminium substrate at duty cycles equal to 20% and 70%.

Figure 8: FTIR spectrum of solid nanoparticles collected in the gas phase at duty cycle equal to 20%. Infrared region: from 3750 to 750 cm^{-1}. Resolution: 2 cm^{-1}.

Figure 9: FTIR spectra of solid nanoparticles collected in the gas phase at different duty cycles (from 20 to 70%). Infrared region: from 3750 to 750 cm^{-1}. Resolution: 2 cm^{-1}.

Figure 10: O(1s), N(1s), C(1s) and Si(2p) XPS spectra of nanoparticles collected on an aluminium substrate for different duty cycles.

Condition	Species	Transition	System name	Intensity	Δv
w/o APTES	OH	OH ($A^2\Sigma^+$, $v' \rightarrow X^2\Pi$, v'')	Å system	moderate	$\Delta v = 0$
	O	O ($1S \rightarrow 1D$)	Green line	Very weak	
	O_2	O_2 ($b^1\Sigma_g^+$, $v' \rightarrow X^3\Sigma_g^-$, v'')	Atmospheric band	Very weak	$\Delta v = 0$
w/ APTES	OH	OH ($A^2\Sigma^+$, $v' \rightarrow X^2\Pi$, v'')	Å system	Strong	$\Delta v = -1, 0$
	CN	CN ($B^2\Sigma^+$, $v' \rightarrow X^2\Sigma^+$, v'')	CN violet	Moderate	$\Delta v = -1, 0, +1$
	CH	CH ($A^2\Delta_g$, $v' \rightarrow X^2\Pi_u$, v'')	CH 4300Å	Moderate	$\Delta v = 0$

Table 1: List of the transitions observed by optical emission spectroscopy in the Ar-O_2 afterglow with and without APTES.

Assignments	Frequency	Ref.	Assignments	Frequency	Ref.	Assignments	Frequency	Ref.
$\nu\ H_2O$	$3890\ cm^{-1}$	[45]	**$\nu\ CO_2$**	**$2348\ cm^{-1}$**	[42]	$\nu_{as}\ SiO–CH_2$	$1241\ cm^{-1}$	[42]
Si-OH	$3674\ cm^{-1}$	[45]	**$\nu\ CO$**	**$2140\ cm^{-1}$**	[42]	$\rho\ CH_2$	$1176\ cm^{-1}$	[41, 42]
>CO...O–H	**$3234\ cm^{-1}$**	[46, 47]	Si–O overtones	$2116\ \&\ 2264\ cm^{-1}$	[45]	$\nu_{as}\ Si–O–C$	$1110\ cm^{-1}$	[40]
$\nu_s\ CH_3$	$2987\ cm^{-1}$	[45]	Si–O + $\rho\ CH_3$	$1938\ cm^{-1}$	[44]	Si–O in APTES	$1065\ cm^{-1}$	[37, 38, 42]
$\nu_s\ CH_2$	$2939\ cm^{-1}$	[41, 45]	**$\nu\ C=O$**	**$1744\ cm^{-1}$**	[42]	$Si–O–C_2H_5$	$958\ cm^{-1}$	[42]
$\nu_{as}\ CH_3$	$2899\ cm^{-1}$	[45]	$\delta\ NH$	$1615\ cm^{-1}$	[39]	$\rho\ CH_3$	$878\ cm^{-1}$	[42]
$\nu_{as}\ CH_2$	$2870\ cm^{-1}$	[41, 45]	$\delta\ H–C–H$	$1449\ \&\ 1475\ cm^{-1}$	[36, 43]	$\omega\ NH_2$	$785\ cm^{-1}$	[38]
Si–OH…NH$_2$	$2745\ cm^{-1}$	[40]	$\delta\ CH_2$ (distant from Si)	$1394\ cm^{-1}$	[38, 39]	**$\delta\ CO_2$**	**$667\ cm^{-1}$**	[46]

Table 2: Vibrational groups and their corresponding frequencies observed in the APTES flow with (bold) and without Ar-O$_2$ afterglow.

Assignment	Comment	Frequency	Ref.	Assignment	Comment	Frequency	Ref.
Si–O–C$_2$H$_5$	strong	943 cm^{-1}	[42]	ν CO (amide I)	strong	1698cm^{-1}	[46]
Si–O in APTES	shoulder	1083 cm^{-1}	[42]	ν$_{as}$ CH$_3$		2851 cm^{-1}	[41, 45]
Si–O in APTES	strong	1169 cm^{-1}	[42]	ν$_s$ CH$_3$		2930 cm^{-1}	[41, 45]
δ CH$_2$	shoulder	1376 cm^{-1}	[36, 43]	ν$_s$ NH$_2$; ν$_{as}$ NH$_2$	very broad	3120-3350 cm^{-1}	[46]
δ C–O–H	strong	1441 cm^{-1}	[46]	ν OH	very broad	3350-3680 cm^{-1}	[46]
δ NH$_2$ (amide II)	shoulder	1620 cm^{-1}	[46]				

Table 3: Main vibrational groups and their corresponding frequencies observed in nanoparticles produced from APTES for different duty cycles (from 20 to 70%).
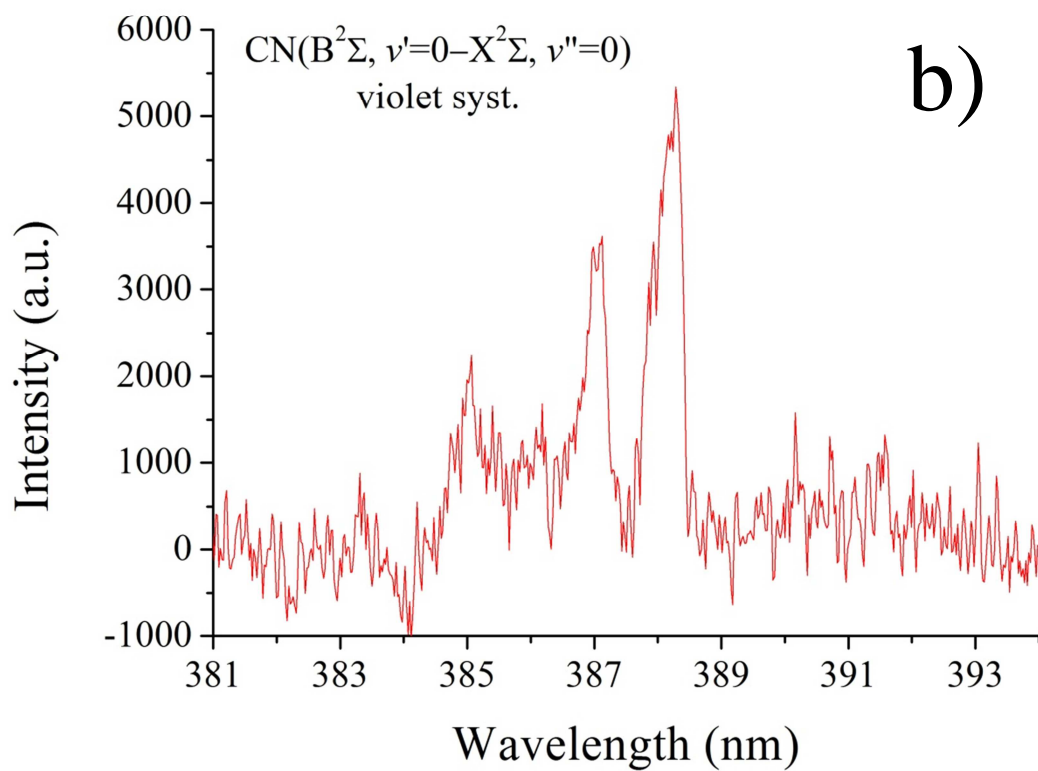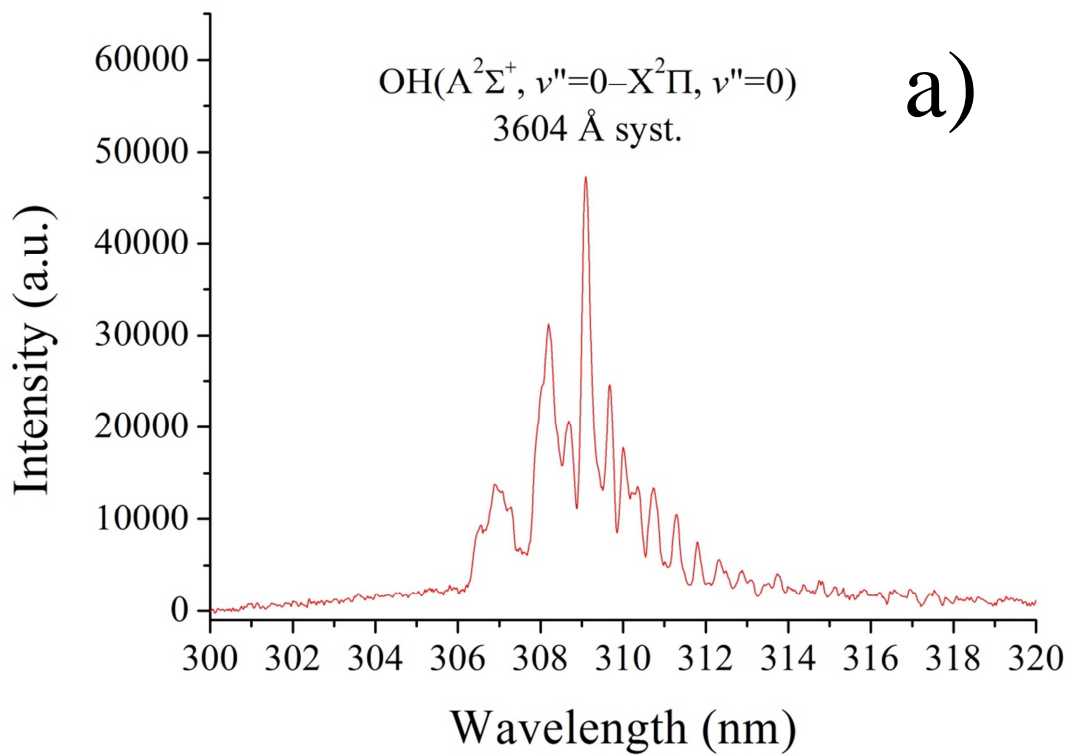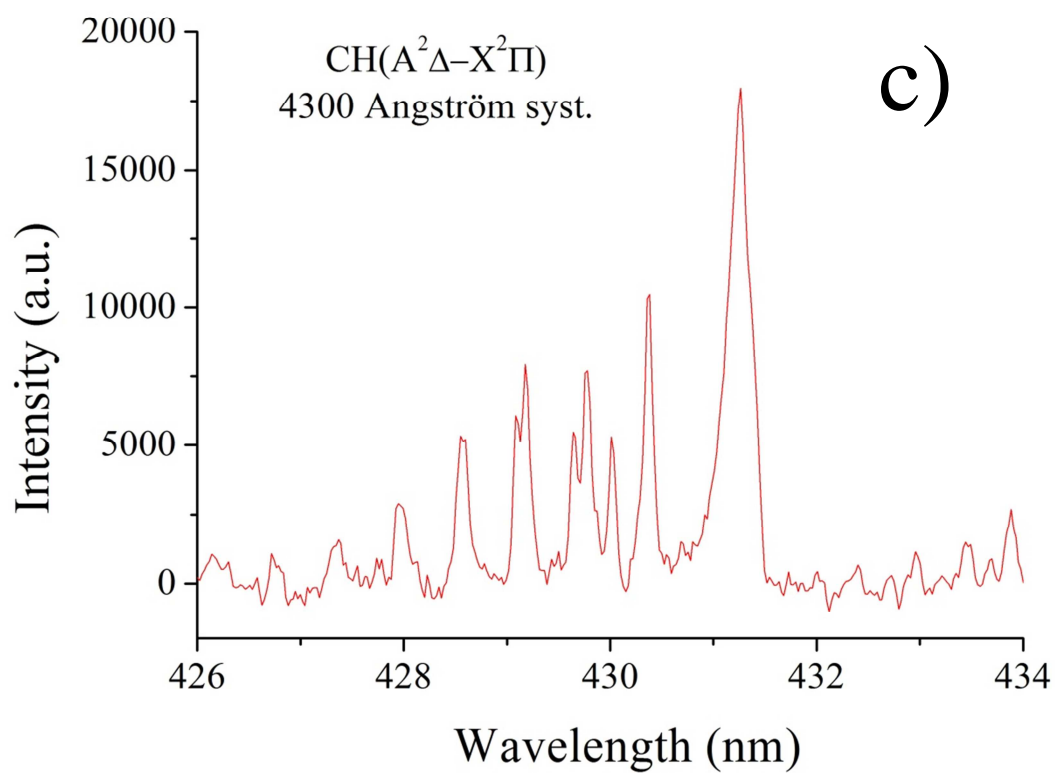
SUPPLEMENTAL MATERIALS
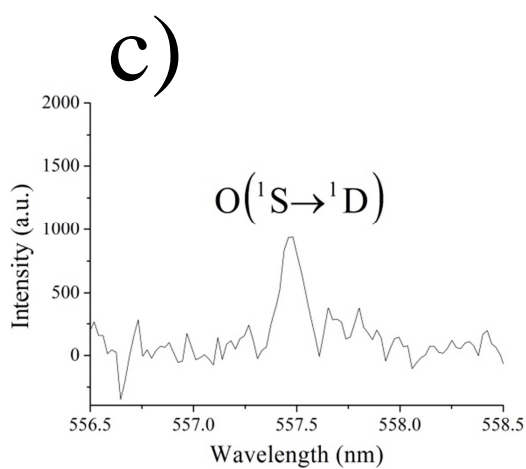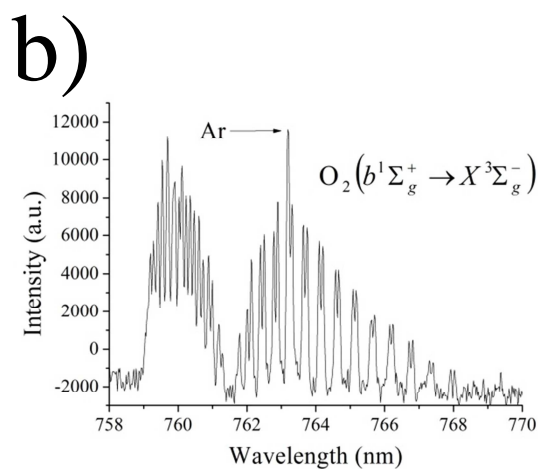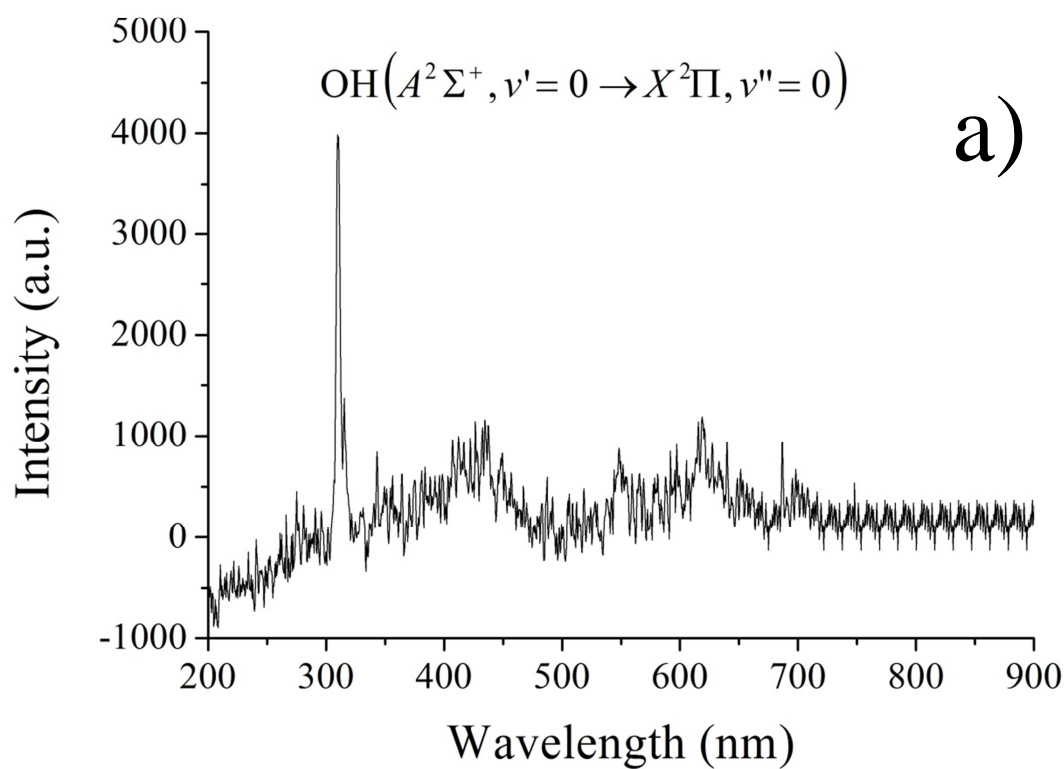
Supplemental material 1: a) OES spectrum of the Ar-O$_2$ post-discharge showing principally the OH emission at 306 nm. b) Detail of the atmospheric band (an argon line due to an unwanted reflection from the plasma is present in the middle of the spectrum and must be removed for any simulation of the rotational spectrum). c) Detail of the green line.
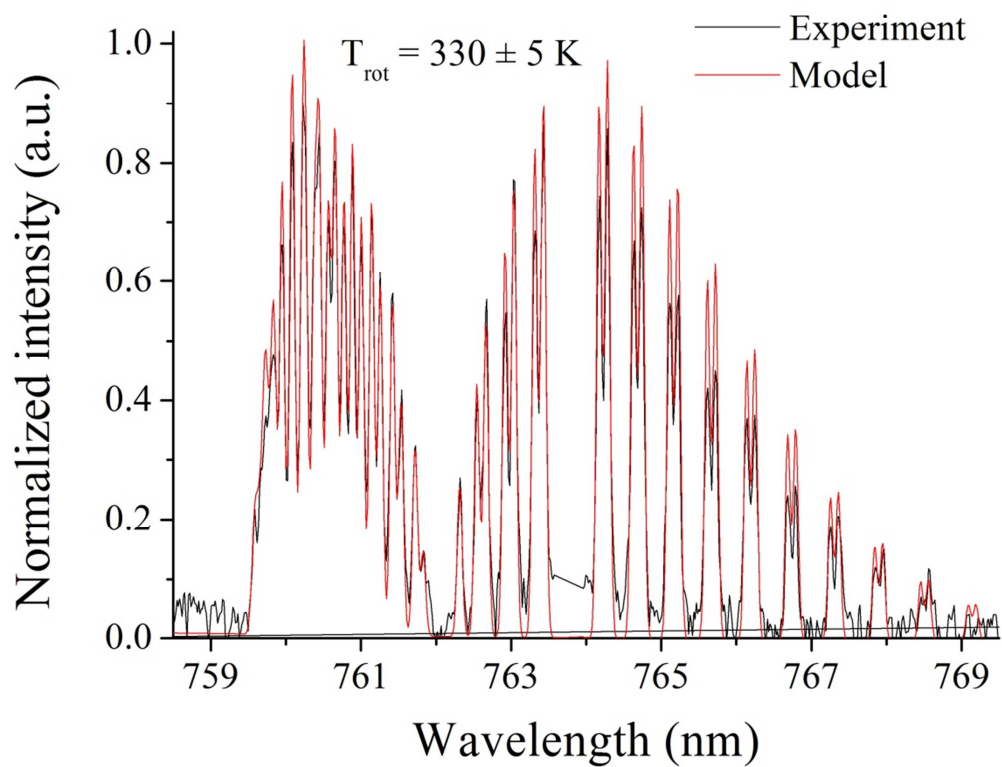
Supplemental material 2: Modelling of the rotational spectrum of the $\Delta v = 0$ transition at 762 nm of the atmospheric band to determine the gas temperature (the argon line appearing in the spectrum has been removed).

a)

OH($A^2\Sigma^+$, v''=0–$X^2\Pi$, v''=0)
3604 Å syst.

b)

CN($B^2\Sigma$, v'=0–$X^2\Sigma$, v''=0)
violet syst.

Supplemental material 3: a) OES spectrum showing the OH emission at 306 nm in the Ar-

O$_2$ post-discharge with APTES. b) Idem for CN at 388 nm. c) Idem for CH at 432 nm
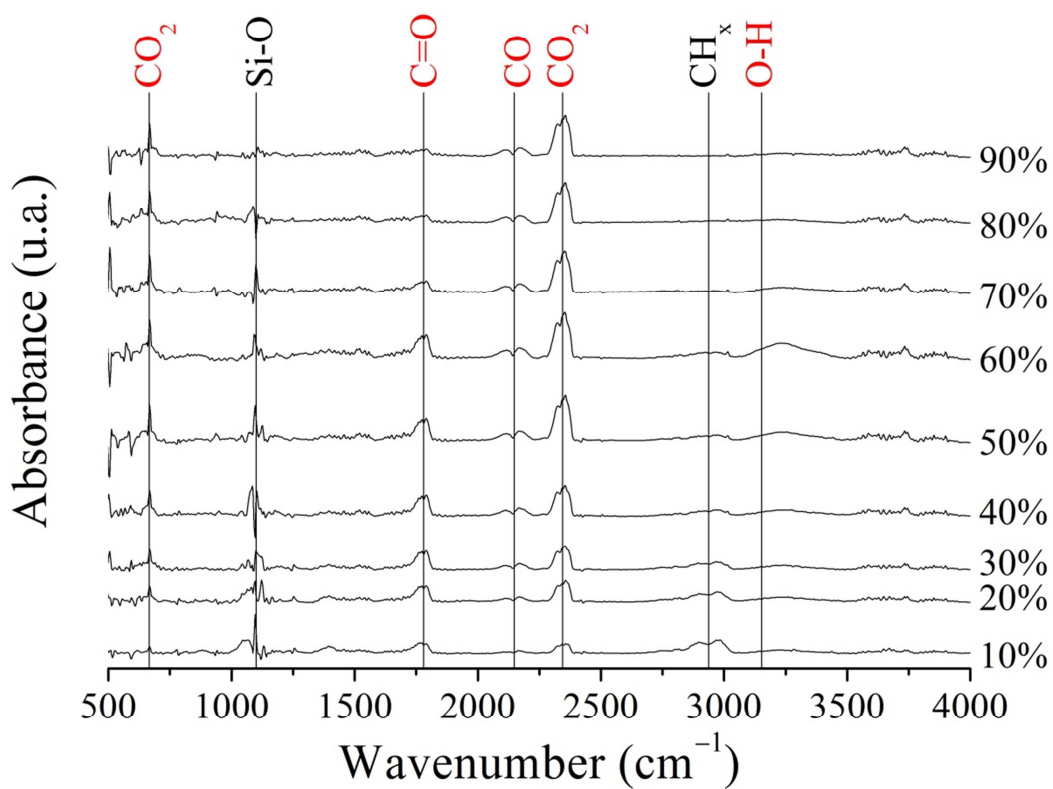
Supplemental material 4: Evolution of FTIR spectra recorded in an APTES–Ar–O_2 afterglow for various duty cycle (resolution 8 cm^{-1}, P = 15 mbar, W = 250 W, flowrates: Ar =1050 sccm, O_2= 100 sccm, Ar =30 sccm, APTES = 0.13 sccm)
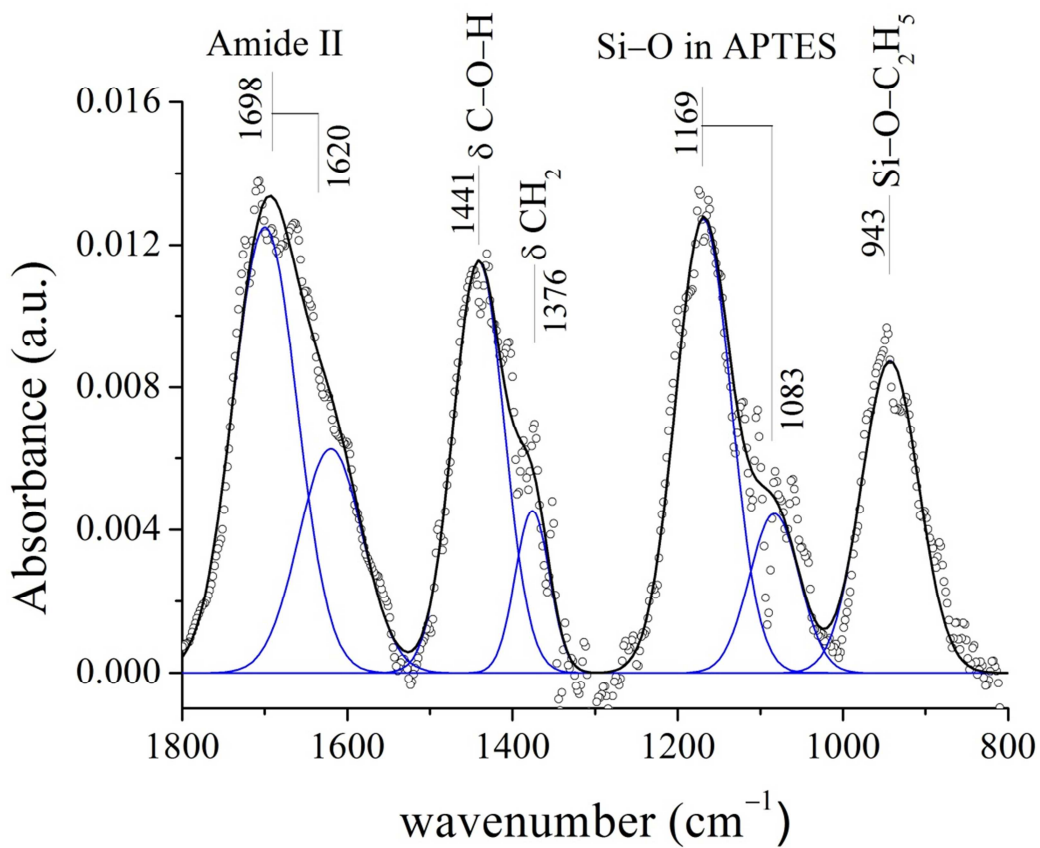
Supplemental material 5: Deconvolution of absorption bands between 1800 and 800 cm^{-1} for nanoparticles grown in an APTES–Ar–O_2 afterglow (resolution 8 cm^{-1}, DC=20%, P = 15 mbar, W = 250 W, flowrates: Ar =1050 sccm, O_2= 100 sccm, Ar =30 sccm, APTES = 0.13 sccm).
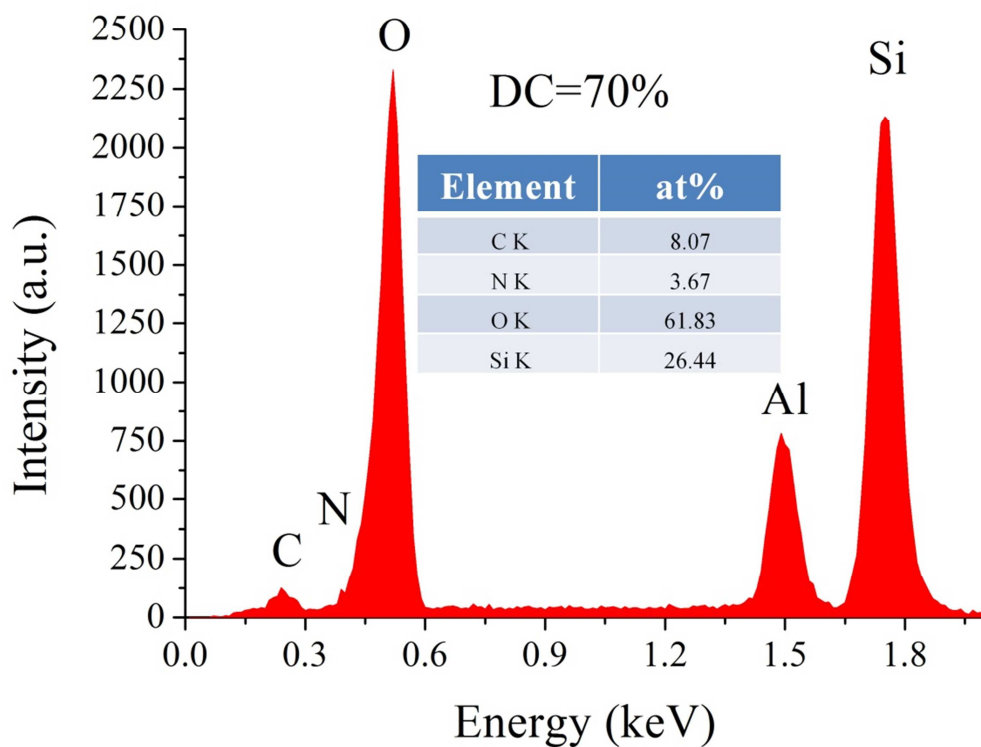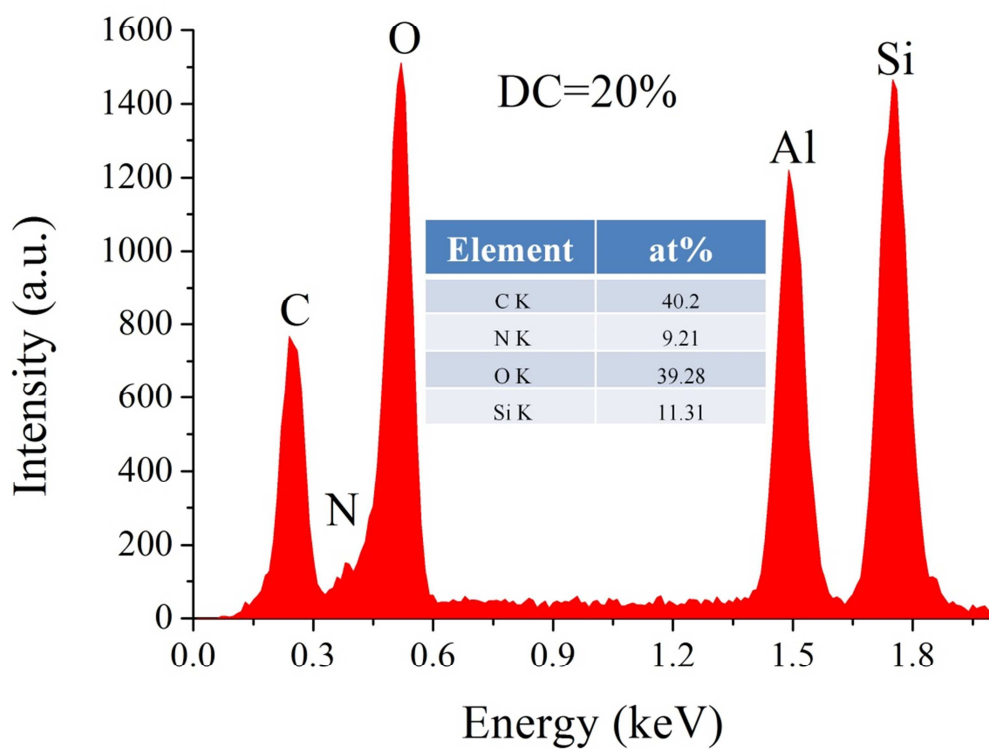
Supplemental material 6: EDX analyses of powders collected on an aluminium substrate for duty cycles equal to 20 and 70%. (P = 15 mbar, W = 250 W, flowrates: Ar =1050 sccm, O_2= 100 sccm, Ar =30 sccm, APTES = 0.13 sccm)