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# Dynamics of VOCs degradation and bacterial inactivation at the interface of $\text{Ag}_x\text{O}/\text{Ag}/\text{TiO}_2$ prepared by HiPIMS under indoor light

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## Abstract

$\text{Ag}_x\text{O}/\text{Ag}/\text{TiO}_2$  photoactive thin films were produced by different modes of magnetron sputtering (namely, DCMS and HiPIMS). The spectral and microstructural properties of the  $\text{Ag}-\text{Ag}_x\text{O}/\text{TiO}_2$  thin films were studied by Diffuse Reflection Spectroscopy (DRS) and X-Ray Photoemission Spectroscopy (XPS). The atomic deposition of  $\text{TiO}_2$  and Ag in the thin sputtered film were studied by TEM.  $\text{Ag}_x\text{O}/\text{Ag}/\text{TiO}_2$  exhibited fast photocatalytic volatile organic compounds (VOCs) degradation and bacterial inactivation under low visible light intensity. The Langmuir Hinshelwood model was applied to highlight the photocatalytic performance of the  $\text{Ag}_x\text{O}/\text{Ag}/\text{TiO}_2$  catalyst. The oxidative states of  $\text{Ag}-\text{Ag}_x\text{O}/\text{TiO}_2$  were studied by XPS. Deconvolution of the Ag3d peak of the  $\text{Ag}_x\text{O}/\text{Ag}/\text{TiO}_2$  thin films showed the presence of AgO and metallic Ag. It is suggested that the mechanism involving the degradation of VOCs on  $\text{AgO}/\text{Ag}/\text{TiO}_2$  catalysts is due to the transfer of holes from AgO to  $\text{TiO}_2$  via the  $\text{AgO}/\text{Ag}/\text{TiO}_2$  heterojunction. The electrostatic interaction between both semiconductors allows this charge transfer. In addition, the contribution of reactive oxygen species (ROS) in bacterial inactivation at the interface of  $\text{AgO}/\text{Ag}/\text{TiO}_2$  thin film was also examined. Light-induced interfacial charge transfer (IFCT) between AgO and  $\text{TiO}_2$  leading to pollutants oxidation appears to require low photons' energy and can oxidize pollutants even at high concentrations. The sputtered coatings have also been studied for repetitive reuse, showing the sustainable activity of the prepared materials.

## Keywords

HiPIMS coating; Photocatalysis; indoor air treatment; Redox catalysis; Bacterial inactivation.

## 1. Introduction:

As it is largely connected to human health, indoor air quality has become a very important topic for the social, economic, and scientific community. It has been shown that modern society spends between 70% and 90% of their time indoors [1], such as

home, offices, cars, hospitals, and healthcare buildings... Besides, the world health organization declared that in 2012, 7 million premature deaths are due to air pollution. Amongst these 7 million deaths, 4.3 million were related to indoor contamination [2]. This is a complex problem containing different types of pollutants, such as particles (dust, and smoke...), radon, asbestos, biological agents (bacteria, viruses...), and gaseous contaminants such as ammonia [3-11]. Different sources are emitting VOCs such as outside pollution (food industry, swimming pool water treatment, hospital cleaning room....) and from everyday activities (e.g., cooking, smoking, or even using cleaning agents, and office equipment) [12-20]. To reduce the VOCs from air, various techniques were used, such as adsorption [21], condensation [22], plasma combined with catalysis [5-7,23-25], and photocatalysis [20,24]. Notwithstanding their significant ability in the VOCs removal, these proposed techniques were shown to produce high amounts of dangerous intermediates, by-products, and waste. Furthermore, these processes exhibit a temporary pollution transfer from gaseous to solid phase or pollution allocation from one area to another [27].

Heterogeneous photocatalysis is deemed as a promising method for air purification owing to its ability to remove VOCs even at relatively elevated concentrations [5, 7, 28-30]. This technique was able to degrade a wide range of pollutants and their by-products [31-33]. Titanium dioxide ( $\text{TiO}_2$ ) is widely employed photocatalyst for pollutants' removal/abatement due to its high stability, excellent activity, inexpensive to build and operate, and its non-toxicity [40, 41].  $\text{TiO}_2$  supported on glass fiber tissue [34, 35], cellulosic paper [36], polymer support [34], and  $\text{TiO}_2$  on PES fabrics [20, 37]. However, the main limiting factor affecting the photocatalytic application of  $\text{TiO}_2$  in real settings is its wide band gap (3.2 eV) allowing only the absorption of UV photons [38, 39]. To extend the spectral absorption of  $\text{TiO}_2$  in the visible light range, ionic (cationic or anionic) doping was seen to tighten its band gap allowing solar/visible light absorption [42-45]. Other researchers adopted the engineering of tailored heterojunctions with other metal oxides allowing visible light absorption [39, 46-49].

In this study, we chose to combine  $\text{TiO}_2$  with  $\text{Ag}_x\text{O}$  to enhance the photocatalytic activity in the visible spectral range.  $\text{Ag}_x\text{O}$  was chosen due to its strong plasmon resonance occurring under visible light excitation. The choice of  $\text{Ag}_x\text{O}$  is favorable for energy levels' alignments that authorize the silver plasmonic electrons to be

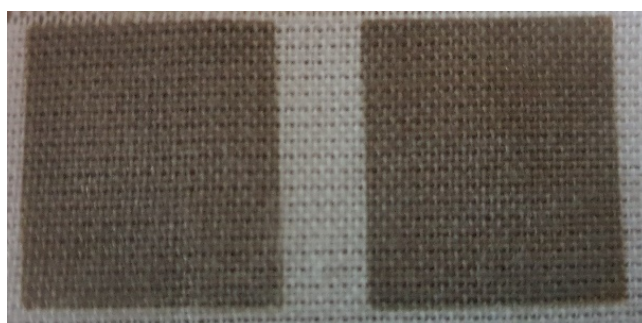
transferred into the  $\text{TiO}_2$ . Both oxides were sputtered on textile fabrics to overcome the labor and time required to recover the catalyst after use.

This work addresses the synthesis of  $\text{Ag}_x\text{O-TiO}_2$  on polyester cloth by two sputtering modes (DCMS and HiPIMS). Odorous compounds such as chloroform (Trichloromethane,  $\text{CHCl}_3$ ) and Diacetyl (butane-2,3-dione,  $(\text{CH}_3\text{CO})_2$ ) were chosen as indoor VOCs models. Chloroforms are model pollutants commonly encountered in hospital settings. Diacetyl is normally emitted during food processing and preparation; they are dangerous to health [20]. In addition, the *E. coli* bacteria was chosen as a microorganism model. The effect of the applied sputtering mode on the kinetics of COVs removal (chloroform and butadione) and bacterial inactivation was studied and is presented in details in this work. The surface properties of the sputtered samples were examined by up-to-date surface science techniques. The prepared films will open the doors to potential applications in hospital settings to reduce bacterial infections and to remove malodors.

## 2. Experimental

### 2.1. Catalyst synthesis: $\text{Ag/TiO}_2$ sputtered PES

Ag and Ti were sputtered on PES using magnetron sputtering in the (HiPIMS and DCMS modes). The HiPIMS deposition was assisted with using a HiP3 5 kW Solvix generator at a time-average power of 100 W ( $5 \text{ Wcm}^{-2}$ , pulse-width of 50  $\mu\text{s}$  and a frequency of 1 kHz, gas phase 95% Ar - 5%  $\text{O}_2$ ). The applied power to the Ti-target was 90 W for the first 30 s then at 50 W to regulate the relative content of Ti in the film as it will be shown later by transmission electron microscopy (TEM). The distance between the sputtered target and the sample holder was 10 cm as previously optimized [58]. The visual perception of the sputtered PES fabrics is shown in Figure 1.



**Figure 1.** Visual perception of the  $\text{Ag}_x\text{O-TiO}_2$  sputtered on PES.

## 2.2. Thickness of the sputtered Ag<sub>x</sub>O-TiO<sub>2</sub>-PES, TEM, spectral absorption and XPS

The thicknesses of TiO<sub>2</sub>, Ag, and AgO-TiO<sub>2</sub> deposited on Si-wafer we measured using a Profilometer (Alphastep-500, Tencor, USA). To carry out TEM observations, the PES cloths were embedded in epoxy resin (Embd 812) and cross-sectioned with an ultra-microtome (Ultracut E) and then imaged using FEI Osiris operated in 200kV. With Spot size of 5, dwell time 50 μs, real time 600 s.

Diffuse reflectance spectroscopy (DRS) was carried out by means of a Perkin Elmer Lambda 900 UV-VIS-NIR spectrometer within 250-700 nm wavelength range.

In order to determine the existing species on the sputtered Ag<sub>x</sub>O-TiO<sub>2</sub> film, X-ray photoelectron spectroscopy (XPS) was performed using an AXIS NOVA photoelectron spectrometer (Kratos Analytical, Manchester, UK, monochromatic Al-K<sub>α</sub> with  $h\nu = 1486.6$  eV). In order to correct the charging effect at the interface of the sputtered samples, the carbon C1s line having position at 284.6 eV was used as reference. The surface atomic concentration was determined from the peak areas considering the sensitivity factors for each element. The XPS spectral peaks were deconvoluted using the CasaXPS V2 software.

## 2.3. Target compounds and tested bacteria

Two target organic compounds were used, Butadione (C<sub>4</sub>H<sub>6</sub>O<sub>2</sub>) and Chloroform (CHCl<sub>3</sub>), existing particularly in the food and hospital sectors, respectively. Butadione (99% purity) and Chloroform (99.4% purity) were used as VOCs probes. *Escherichia coli* (DSM 10198-0307-001) was used as a model microorganism to test the antibacterial activity. The preparation protocol and the testing method have been detailed before [20].

## 2.4. Reactor design for VOC removal

The used batch reactor has approximately 10 mm as upper diameter and 50 mm of working height contains two septa that are able to contain up to 50 mL of solution. The first one is used for samples extraction and the second one is used for VOCs injection. The catalytic samples were fixed to the internal wall of the batch reactor to be in direct contact with the studied VOC and light radiation. For all experiments, 20 cm<sup>2</sup> of coated Ag<sub>x</sub>O-TiO<sub>2</sub>-PES were used. A fluorescent visible lamp (Sylvania CF-L 24 W/840) with emission in the visible (400 – 720 nm) has been inserted inside the reactor. This light is a typical indoor light like those used for houses, hospitals and schools. To prevent extra-light from, the hall reactor was covered with Al-foil during the

photocatalytic experiments. The homogenization of the gas phase was allowed by a magnetic stirrer and barrel.

## 2.5. VOCs analysis system and antibacterial testing

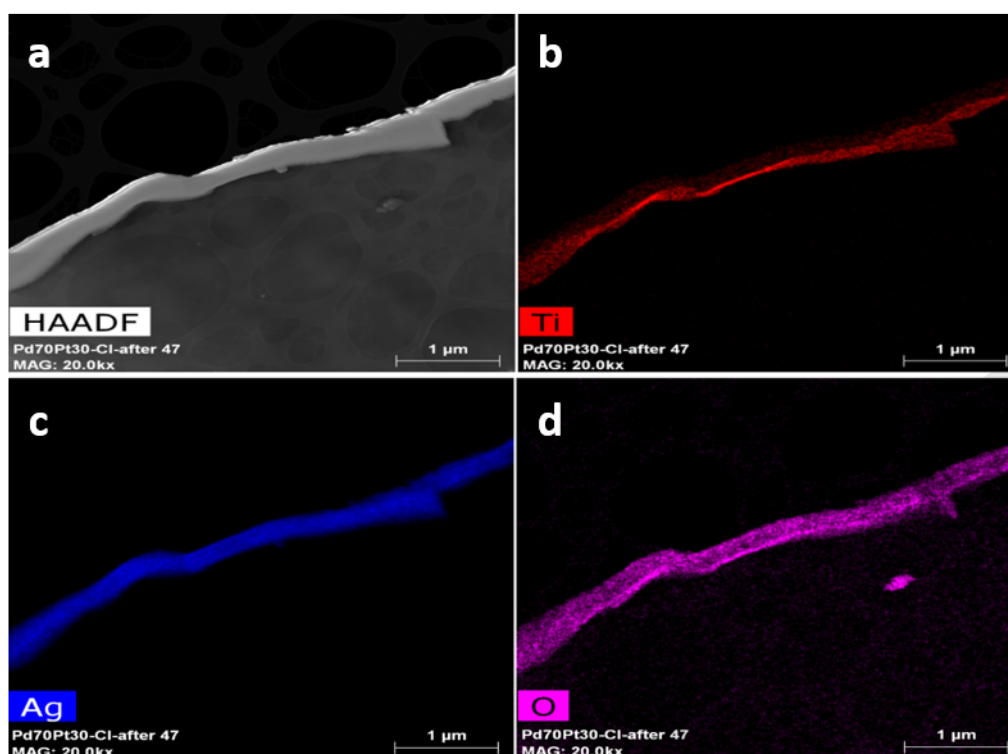
To examine the performance of our catalysts for VOCs removal  $C_4H_6O_2$  and  $CHCl_3$  analysis was carried out using gas chromatography (Fisons GC9000) equipped with a FID. The mobile phase (carrier gas) was Nitrogen. We used a syringe of 500  $\mu$ L for all injection. All injections were carried out in triplicates. Once the flow inside the reactor gets stabilized (at equilibrium), the pollutants analysis starts.

The antibacterial activity of  $Ag_xO-TiO_2-PES$ , *Escherichia coli* (*E. coli*) was studied as reported in our previous study [20]. In this study, the initial bacterial concentration is  $\sim 2.6 \cdot 10^6$  colony-forming unit per milliliter (CFU/mL).

## 3. Results and discussion

### 3.1. Thickness calibration and TEM of sputtered $Ag_xO-TiO_2-PES$

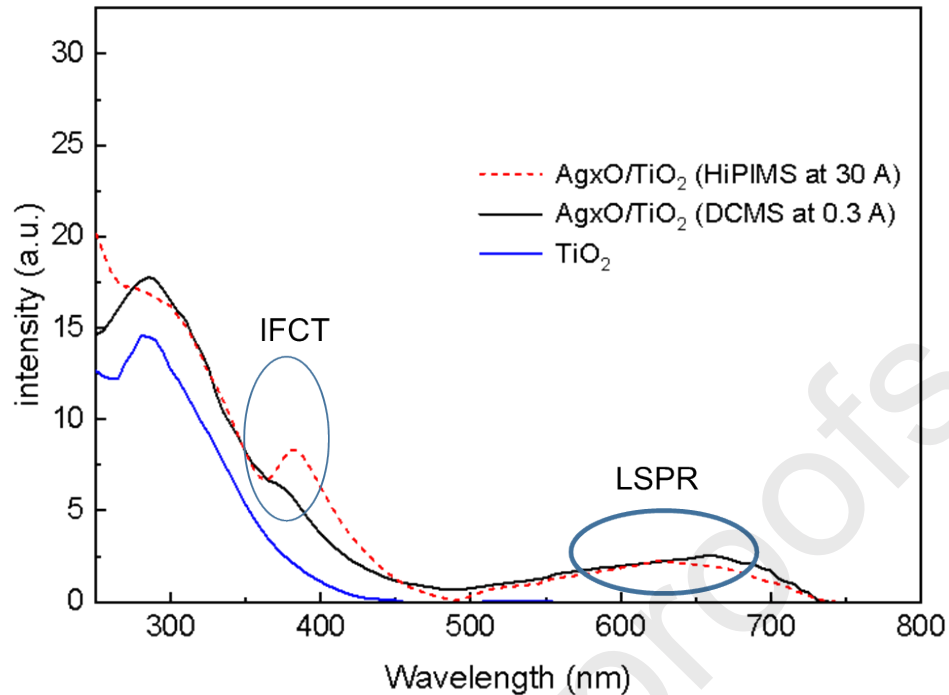
Figure 2a presents the high-angle annular dark field (HAADF) mapping of elements in the  $Ag_xO-TiO_2-PES$  samples sputtered by HiPIMS at 30 A. Figures 2b, 2c and 2d show the mapping of Ti-, Ag-, and O-element, respectively. The Ag-atoms distribution (red image) appears to be dense and uniform. This likewise provides evidence for the presence of different oxidation states of Ag-NPs (AgO) in the topmost layers. In addition, many kinds of research studied in a detailed way the properties of Ag-coated PES [64-68]. The EDX mapping of oxygen clearly shows (Fig. 2d) relatively large amounts of oxygen species (purple image). This is due to  $TiO_2$  (EDX mapping of Ti is shown in figure 2b) and AgO nanoparticles, which exhibit a higher O-content compared to the content of oxygen species in the film bulk. In the context of a comparison between the samples sputtered by DCMS and those sputtered by HiPIMS, Rtimi et al reported that the distribution of oxygen species in the samples sputtered by DCMS shows more oxidized copper (in this case Ag) species in the bulk than the samples prepared by HiPIMS, while the latter has shown more oxidized copper species at the upper layers in contact with bacteria [69].



**Figure 2.** TEM images of  $\text{Ag}_x\text{O-TiO}_2\text{-PES}$  as sputtered by HiPIMS at a current of 30 A for 20s: a) High-angle annular dark-field imaging (HAADF), b) EDXS mapping of Ti, c) EDXS mapping of Ag, and d) EDXS mapping of O.

### 3.2. Surface properties of the sputtered $\text{Ag}_x\text{O-TiO}_2\text{-PES}$

Figure 3 shows the spectral absorption of the sputtered  $\text{Ag}_x\text{O-TiO}_2\text{-PES}$  by HiPIMS (at 30 A) and DCMS (at 0.3 A). Analysis of the absorption spectra show that  $\text{TiO}_2$  thin film do not absorb in the visible and presented an absorption onset ( $\lambda$ ) at 260 nm. In contrast, the light absorption of  $\text{Ag}_x\text{O/TiO}_2$  sputtered by HiPIMS and DCMS extends to the visible region. In addition, a significant phenomenon was observed related to the interaction of light with The Ag incorporated into Titanium dioxide near the absorption edge. This was related to the interfacial charge transfer (IFCT) between  $\text{Ag}_x\text{O}$  and  $\text{TiO}_2$  semiconductors [38]. A wide absorption band from 500 to 750 nm is noted for  $\text{Ag}_x\text{O/TiO}_2$  photocatalysts sputtered by HiPIMS at 30A and DCMS at 0.3A. These bands can be ascribed to surface plasmon resonance (SPR) of the Ag species [38,44].



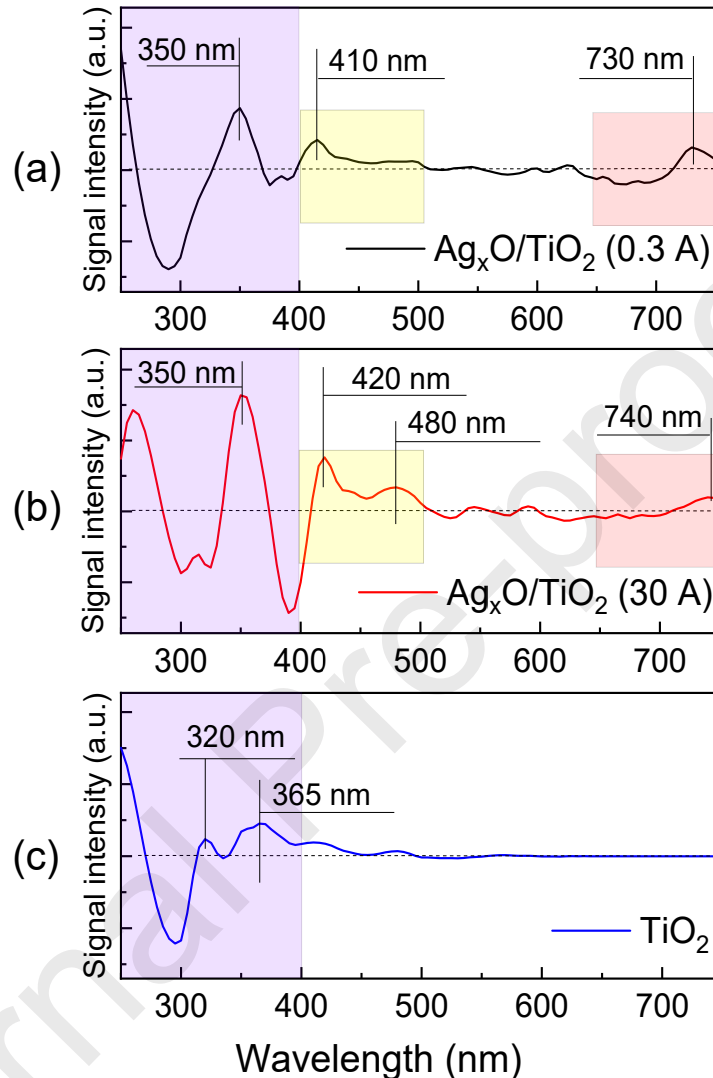
**Figure 3.** Diffuse reflectance spectroscopy (UV-Vis) of sputtered TiO<sub>2</sub> (using DCMS at 0.3 A) and Ag<sub>x</sub>O-TiO<sub>2</sub> (using DCMS at 0.3 A and HiPIMS at 30 A).

In order to better analyze the light absorption of our coatings, we applied a spectral analysis based on high derivatives. The method of higher derivatives has been known for decades [70], but it has not previously been applied to the analysis of photocatalytic materials' spectra. Analysis of extrema in the representation of the second and fourth derivatives allows the determination of the UV-Vis absorption maxima giving a clear idea about the appropriate wavelengths to be used with this system. The signal-to-noise ratio in modern spectrometers is high enough to reliably determine higher derivatives. The fourth derivative of the absorption spectra allows to reliably determining the position of significant peaks. If these peaks overlap slightly, the second derivative becomes enough to locate the significant peaks. Figure 4 shows the second derivatives of the absorption spectra of (a) TiO<sub>2</sub> (using DCMS at 0.3 A) and Ag<sub>x</sub>O-TiO<sub>2</sub> (using DCMS at 0.3A and HiPIMS at 30 A).

It is readily seen from the 2<sup>nd</sup> derivative of the Figure 4a (Ag<sub>x</sub>O/TiO<sub>2</sub> (0.3 A)) that the thin film is absorbing at 350 nm and at 410 and 730 nm with a lower intensity. However, the thin Ag<sub>x</sub>O/TiO<sub>2</sub> film prepared by HiPIMS exhibited higher spectral absorption peak at 350 nm than the film prepared by DCMS. In addition, the HiPIMS prepared film showed peaks at 420, 480 and 740 nm. The TiO<sub>2</sub> thin film showed only spectral

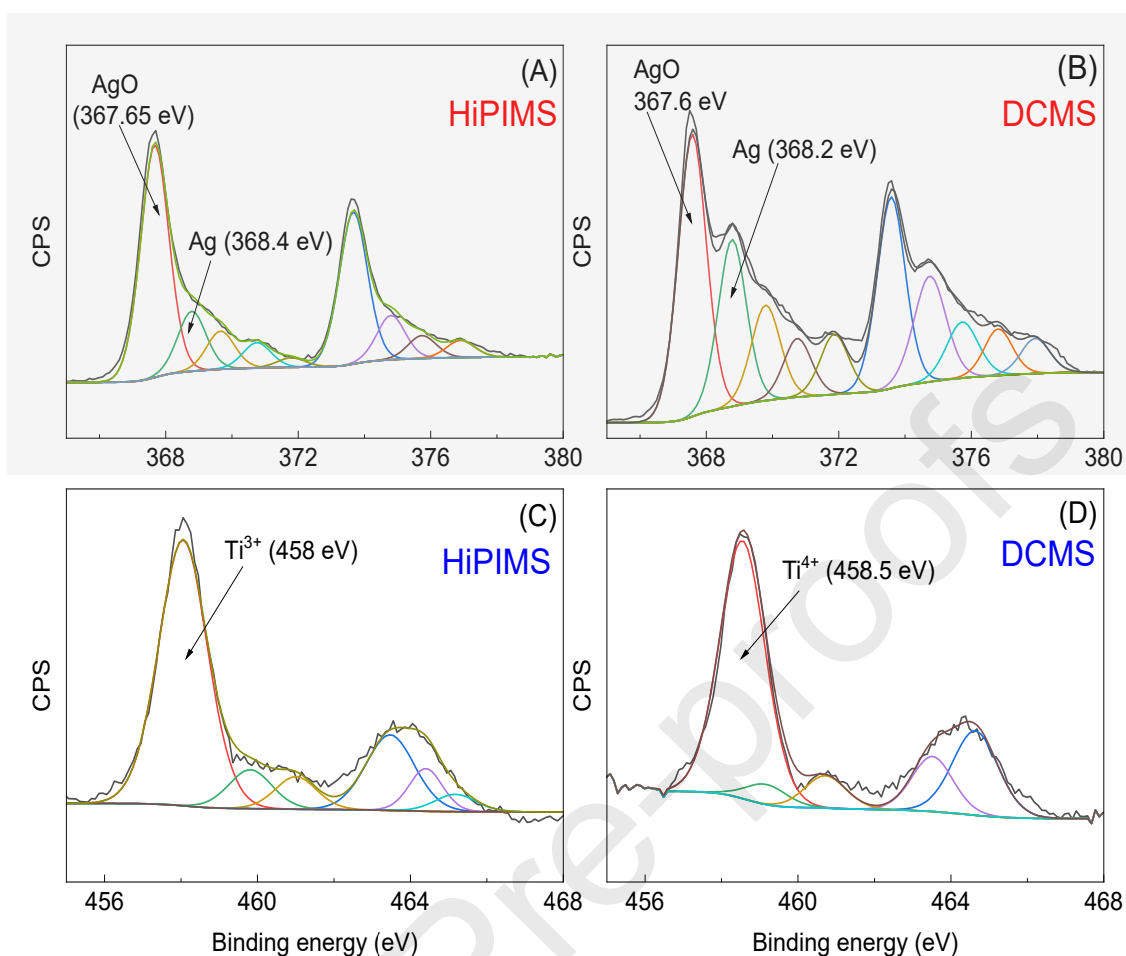


absorption in the UV range (320 and 365 nm). From Figure 4b, the HiPIMS thin film absorb in the UVA, violet and blue. This is attributed to the Ag surface plasmon. The weak peak at 730-740 nm refers to the far-red absorption.



**Figure 4.** Analysis of spectral absorption extrema through second derivatives of light absorption of (a)  $Ag_xO/TiO_2$  (0.3 A), (b)  $Ag_xO/TiO_2$  (30 A) with respect to (c) sputtered  $TiO_2$ .

Figure 5 shows the oxidative states of Ag and Ti sputtered on the polyester cloth as determined by XPS. Figure 5 shows the  $Ti2p$  and  $Ag3d$  peaks from  $Ag_xO-TiO_2$  deposited by HiPIMS and DCMS. The found peaks were asymmetric.  $AgO$  peaks at 367.6 eV were detected in both samples (HiPIMS and DCMS). The spectra showed also the presence of lower peaks at 368 eV characteristics of metallic Ag. The presence of  $Ti^{3+}$  in the HiPIMS samples could arise from the  $AgO$  incorporation on the  $TiO_2$ . The peak at 367.6 eV is characteristic of Ag-ions adsorbed on hydroxylated  $TiO_2$  [71].



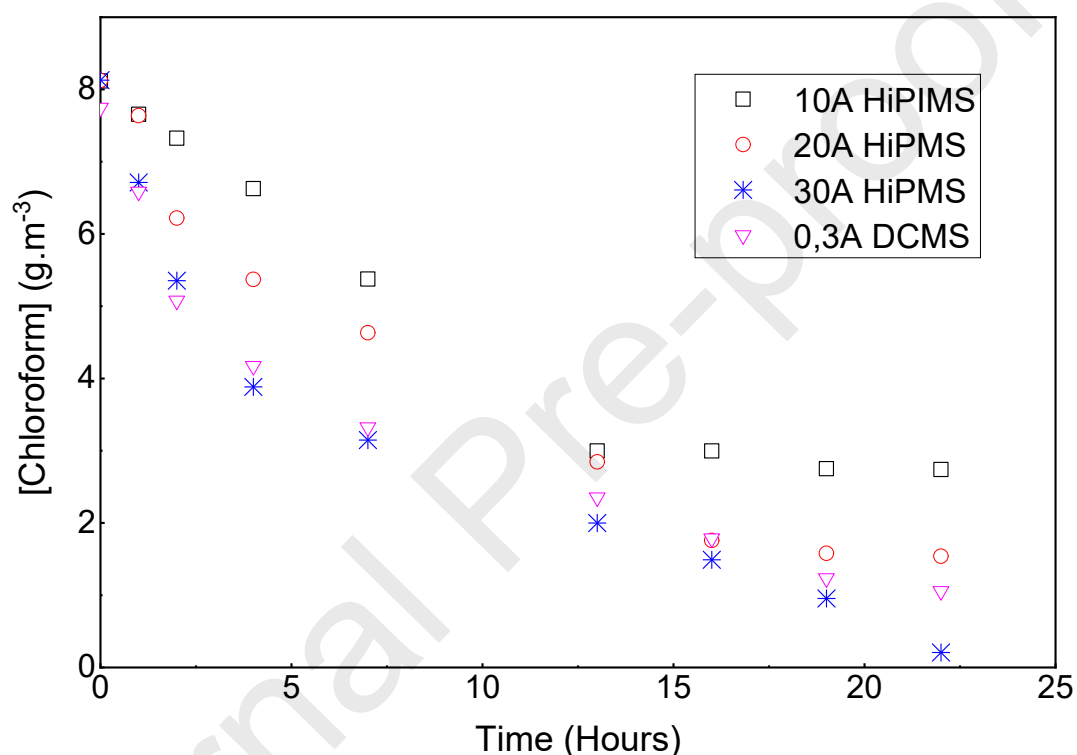
**Figure 5.** XPS deconvolution of Ag3d and Ti2p peaks from AgxO-TiO<sub>2</sub> deposited by HiPIMS and DCMS.

### 3.3. Photocatalytic experiments: COVs removal and Bacterial inactivation tests

#### 3.3.1. Effect of deposition of Ag-NPs TiO<sub>2</sub>-PES by HiPIMS and DCMS on the photocatalytic activity of AgxO-TiO<sub>2</sub>-PES photocatalyst

The photocatalytic performances of AgxO-TiO<sub>2</sub>-PES catalysts were evaluated by photocatalytic removal of COVs (Chloroform) under visible light irradiation at ambient temperature. Figure 6 shows the chloroform removal using AgxO-TiO<sub>2</sub>-PES sputtered by HiPIMS for 10, 20, 30 A and sputtered by DCMS at 0.3A. Figure 6 shows that the chloroform removal is pertinently affected by the different sputtering current. In this test, it is readily seen that the initial concentrations of Chloroform decreased over time under visible light irradiation. Also, an increase in sputtering current from 10A until 30A increases the chloroform removal to about 98% of the initial concentration of 10 g.m<sup>-3</sup>. This enhancement is due to the availability of active sites generated at the interface of the sample deposited at 30A [59]. However, at low current values the chloroform degradation reaches a limit; in fact, these current values are rather small to form a

catalyst layer leading to perfect and complete removal of chloroform. Previously S. Rtimi et al reported [69] that the HiPIMS deposition current generates different ions and charge species, leading to distinguish charges at the interfaces of the thin film. In addition, compared to the AgxO-TiO<sub>2</sub>-PES sputtered by direct current magnetron sputtering (DCMS) at 0.3A, AgxO-TiO<sub>2</sub>-PES (HiPIMS at 30A) exhibits a high photocatalytic activity under visible light. This is due to the generation of a higher number of Ag-ions at the catalysts surface when using HiPIMS (30A) than DCMS [64, 69].



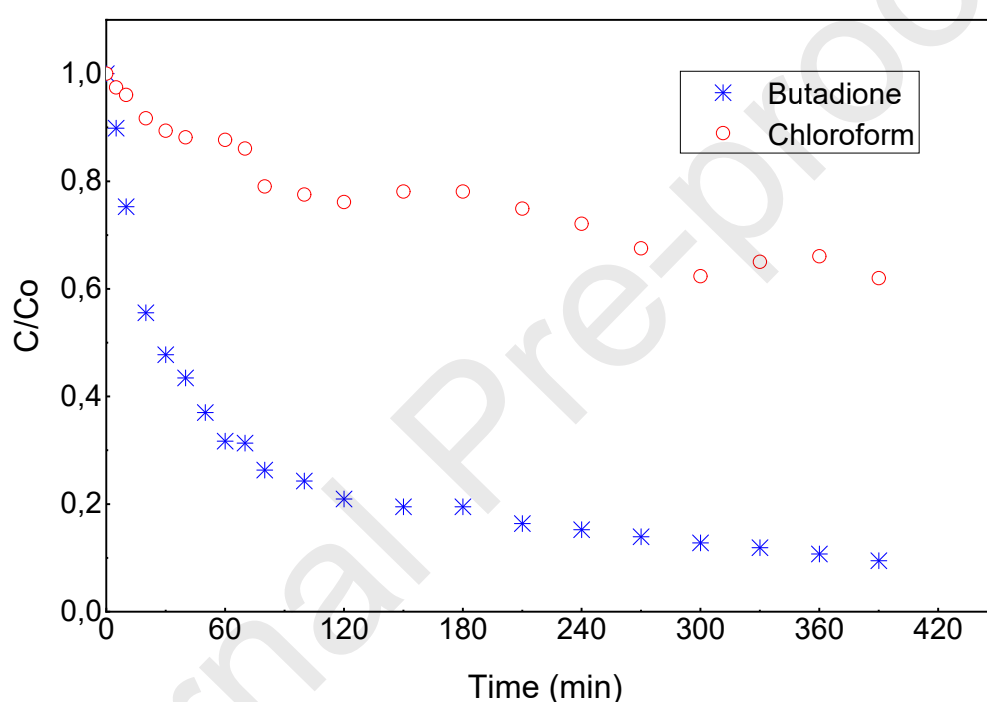
**Figure 6.** Removal of chloroform with catalyst AgxO-TiO<sub>2</sub>-PES at different HiPIMS sputtering currents deposited for 20 s. Light source: fluorescent visible lamp (Sylvania CF-L 24 W/840; 400 – 720 nm)

### 3.3.2. Mixtures of pollutants (VOCs and bacteria)

#### 3.3.2.1. Effect of the presence of two Kinds of VOCs on AgxO-TiO<sub>2</sub>-PES performance

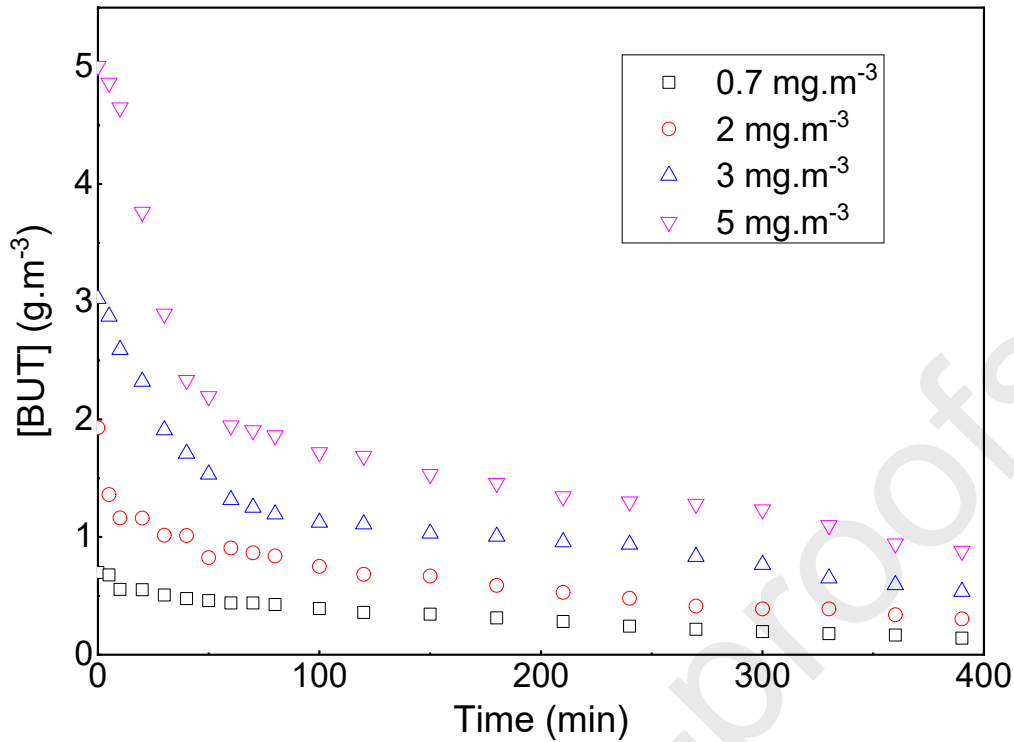
The removal performance of two kinds of indoor air pollutants, with equimolar concentrations, was investigated. Figure 7 shows that at 30A HiPIMS and low visible light illumination, the degradation of Chloroform is quite similar to that of Butadione

(BUT). Firstly, before switching on the lamp and during the adsorption stage, it was found that the adsorbed Butadione amount is slightly greater than that of chloroform. Once adsorption of both species reaches saturation, the lamp is switched on. We found (Fig. 7) that the photocatalytic degradation kinetics of BUT is faster than that of chloroform. This may be attributed to the chemical composition of the pollutants (e.g., carbon number), their affinities to the catalyst and the amounts of their intermediate by-products on the catalyst surface, which in turn leads to a competitive effect of the pollutants towards the active catalytic sites. This result is similar to our previous work [20] where we used  $\text{Cu}_x\text{O}/\text{TiO}_2$ -PES for the removal of VOCs.



**Figure 7.** Two kinds of Indoor air pollutants removal on  $\text{Ag}_x\text{O}-\text{TiO}_2$ -PES HiPIMS-sputtered 30 A for 20 s under a fluorescent visible lamp (Sylvania CF-L 24 W/840; 400 – 720 nm)

Figure 8 shows the BUT degradation in the presence of the  $\text{Ag}_x\text{O}-\text{TiO}_2$ -PES photocatalyst under light starting with different initial concentrations. The degradation rate decreases when the BUT initial concentration increases. This is owing to the availability of active catalytic sites at low concentrations [58, 70]. Indeed, the amount of the molecules involved in the reaction is not increased in the same ratio than that of the inserted initial concentration resulting in a decrease in the degradation efficiency.



**Figure 8.** BUT degradation versus inlet BUT concentration under visible light irradiation (Sylvania CF-L 24 W/840; 400 – 720 nm) based on HiPIMS sputtered Ag<sub>x</sub>O-TiO<sub>2</sub>-PES (30 A for 20 s).

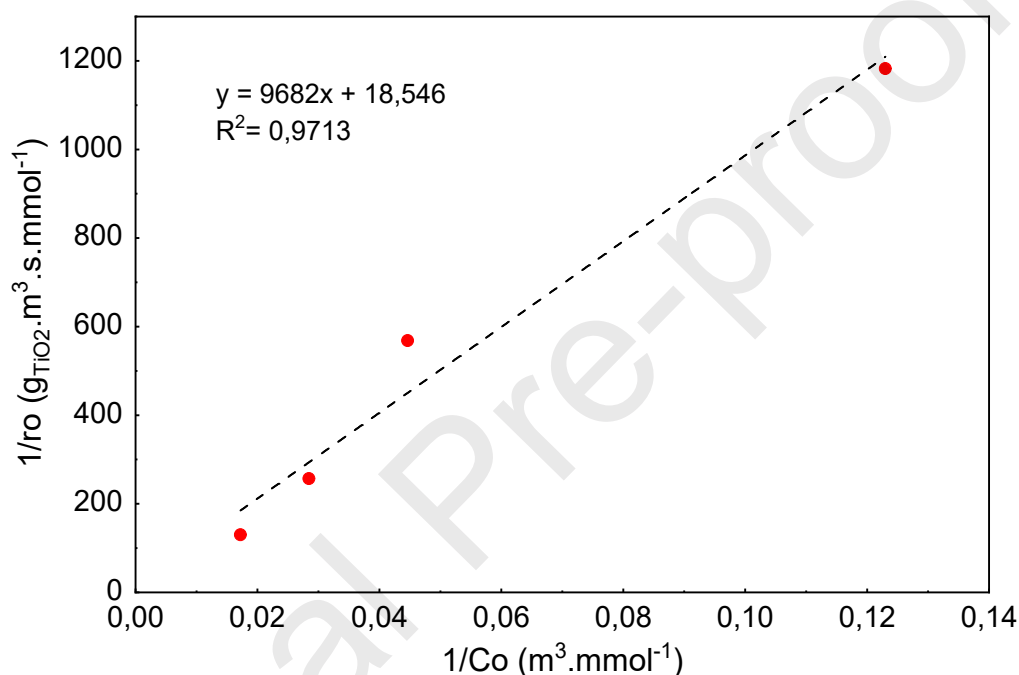
To further understand the chemical transformation of the heterogeneous photocatalytic degradation of Butadione over Ag<sub>x</sub>O-TiO<sub>2</sub>-PES, the Langmuir-Hinshelwood (L-H) model was employed to fit the experimental results [58, 71]. The L-H reaction rate equation shown in the following equation is employed to explain the adsorption and degradation of Butadione:

$$r_0 = -\frac{d[VOC]}{dt} = k_c \frac{K[VOC]_0}{1 + K[VOC]_0}$$

Where  $r_0$  (mmol/g<sub>cat</sub> m<sup>3</sup> s) is the initial photocatalytic degradation rate,  $[VOC]_0$  is the initial BUT concentration (mmol/m<sup>3</sup>),  $K$  is the adsorption constant (m<sup>3</sup>/mmol) and  $k_c$  is the kinetic constant (mmol.m<sup>-3</sup> s g<sub>TiO2</sub>) at maximum coverage for the experimental conditions.

The plot of  $1/r_0$  versus  $1/[\text{COV}]_0$  enables determining the values of  $k_c$  and  $K$ . According to Figure 9, we estimate the values of the kinetic and adsorption constants (Table 1) using the linearized L-H equation ( $\frac{1}{r_0} = \frac{1}{k_c K} \times \frac{1}{[\text{VOC}]_0} + \frac{1}{k_c}$ ).

Table 1 shows Butadione shows a slow degradation rate regarding the use of AgxO-TiO<sub>2</sub>-PES sputtered at 30A; this may be due to the physicochemical properties of the pollutant and to the interfacial charge transfer mechanism (IFCT), which will be shown in a later section.



**Figure 9.** Linear regression using L-H model for the BUT degradation on AgxO-TiO<sub>2</sub>-PES sputtered at 30A for 20s.

**Table 1.** L-H constants ( $k_c$  and  $K$ ) at AgxO-TiO<sub>2</sub>-PES photocatalyst

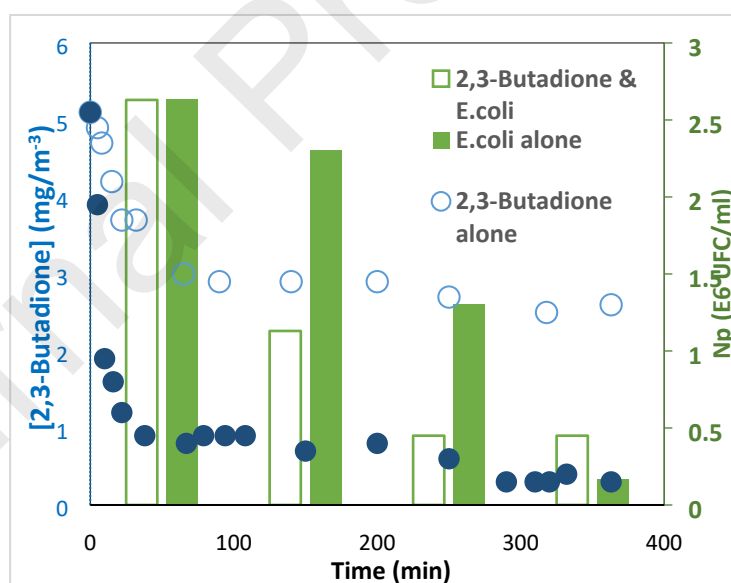
Catalyst tested	$k_c$ : Kinetic constant of L-H ( $mmol \cdot m^{-3} \cdot s \cdot g_{TiO_2}$ )	$K$ : Adsorption Constant of L-H ( $m^3 \cdot mmol^{-1}$ )
AgxO-TiO <sub>2</sub> -PES	$54 \cdot 10^{-3}$	$2 \cdot 10^{-3}$

Table 1 shows that AgxO-TiO<sub>2</sub>-PES leads to a more efficient degradation of butadione compared to PES-TiO<sub>2</sub>, which in turn leads to an almost twice faster degradation of butyraldehyde under UV irradiation [59]. This result may be due to the Ag amount

present on the photocatalyst surface and to the chemical properties of the target pollutants.

### 3.3.2.2. Simultaneous removal of butadione and bacteria

After studying the effect of the existence of two kinds of indoor air pollutants on AgxO-TiO<sub>2</sub>-PES performance, we investigate the bacterial (*E. coli*) inactivation activity of AgxO-TiO<sub>2</sub>-PES together with simultaneous removal of BUT and *E. coli*. Many experiments were done with our composites with initial bacterial concentration and inlet BUT concentration of  $\sim 2.6 \times 10^6$  (CFU/mL) and 5 mg/m<sup>3</sup>, respectively. In fact, the goal of mixing target molecules with bacteria is to validate the effectiveness of the advanced oxidation process (AOP) in a case closer to real conditions where large amounts of pollutants must be removed simultaneously, and to see that there is always a degradation of pollutants during mixing. Figure 10 shows the bacterial inactivation performance of AgxO-TiO<sub>2</sub>-PES during separate and simultaneous removal of BUT and *E. coli* under visible light irradiation.

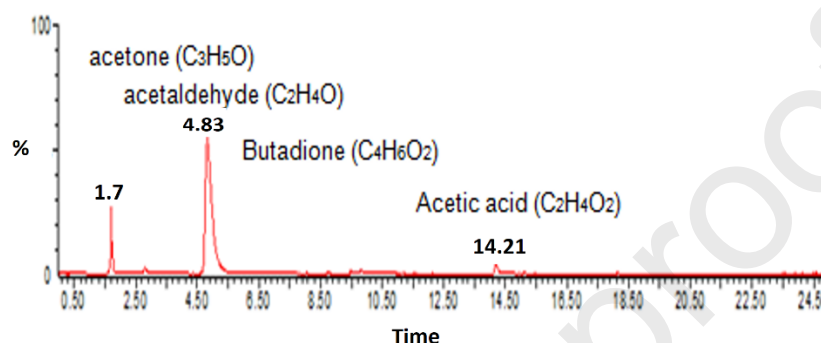


**Figure 10.** Simultaneous removal of *E. coli* and BUT on AgxO-TiO<sub>2</sub>-PES catalyst sputtered at a peak current of 30A. The initial concentration of *E. coli* is  $\sim 2.6 \times 10^6$  CFU/mL. (Light source: Sylvania CF-L 24 W/840; 400 – 720 nm).

### 3.4. Identification of the VOCs' degradation by-products

The by-products emanating from the photocatalytic degradation of butadione in presence of bacteria using the AgxO-TiO<sub>2</sub>-PES catalyst were determined by Gaseous Chromatography-Mass Spectroscopy (GC-MS). Figure 11 shows GC-MS spectra of

by-products originating from AgxO-TiO<sub>2</sub>-PES – based Butadione photo degradation. The detected degradation by-products are: (i) acetone (C<sub>3</sub>H<sub>5</sub>O), (ii) acetaldehyde (C<sub>2</sub>H<sub>4</sub>O) and (iii) acetic acid (C<sub>2</sub>H<sub>4</sub>O<sub>2</sub>). Butadione (C<sub>4</sub>H<sub>6</sub>O<sub>2</sub>) is also present after 4.83 minutes. These results are similar to those reported by Abou Saoud et al. [5], subsequent to the use of TiO<sub>2</sub>-GFT (Glass Fiber Tissue) as photo-catalyst for butadione degradation under photocatalytic/plasma processes.

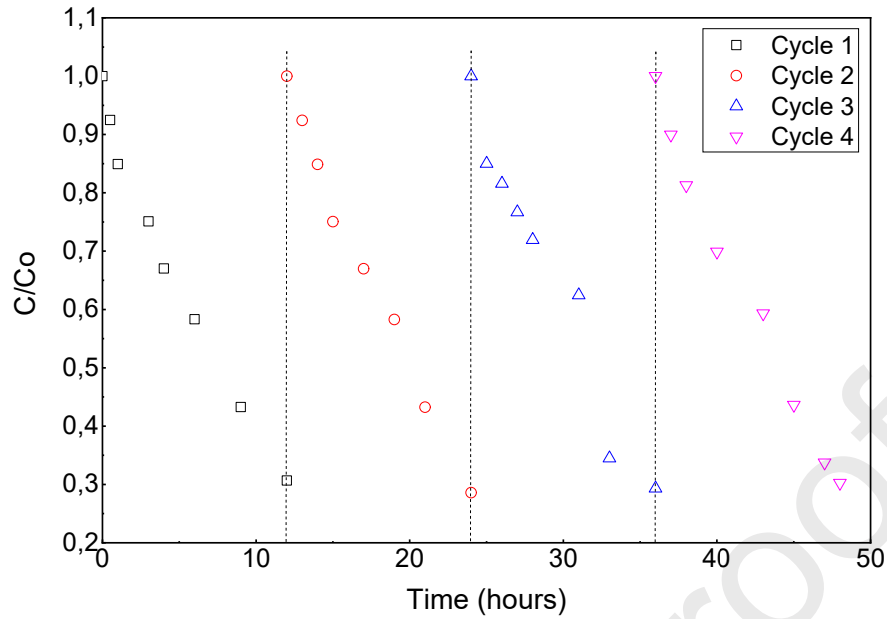


**Figure 11.** GC-MS of products observed during Butadione photo degradation by AgxO-TiO<sub>2</sub>-PES.

### 3.5. Catalyst recyclability

The reusability of the catalyst is an important factor to take into account since it concerns the possibility of its potential application in environments polluted by toxic gases [20]. The re-use of the catalyst was examined for four successive cycles. The AgxO-TiO<sub>2</sub>-PES was washed well after each photodegradation cycle and reused. Figure 12 shows no loss of sample photoactivity after four cycles of use. These results prove the overall potential of the AgxO-TiO<sub>2</sub>-PES to be used as photocatalyst for air treatment applications. Furthermore, the same slope is noted in each cycle; these results can confirm the non-deactivation of the surface sites of the catalyst and its excellent catalytic stability.

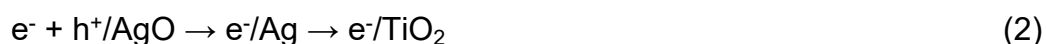




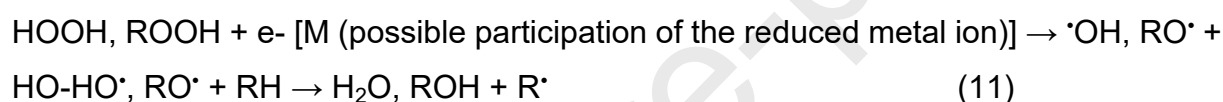
**Figure 12.** Repetitive degradation of Butadiene on Ag<sub>x</sub>O-TiO<sub>2</sub>-PES. VOC concentration = 10g/m<sup>3</sup>.

### 3.6. Photocatalytic degradation mechanism

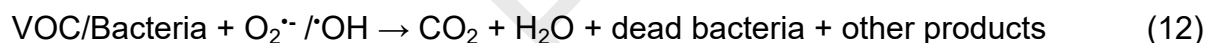
To understand the possible photocatalytic mechanism of VOC removal and *E. coli* inactivation in Ag<sub>x</sub>O/TiO<sub>2</sub> composite, Figure 13 was proposed. Under UV-vis light, electron-hole pairs generate in the valence bands (VB) of AgO and TiO<sub>2</sub>, then migrate to the conduction bands (CB). The AgO conduction band potential was 0.72 eV and the Ag Fermi level was 0.99 eV [72]. As a result, and because the Fermi level of Ag was more positive than the conduction band potential of AgO, the electrons that are in the AgO conduction band can move to the Fermi level of Ag. At the same time, electrons on the Fermi level of Ag can be rapidly transferred to the TiO<sub>2</sub> semiconductor interface, which has been reported in Ag/ZnO and Ag/TiO<sub>2</sub> systems [73,74]. This could be due to the fact that the heterojunction in Ag-AgO/TiO<sub>2</sub> assisted the transfer of photo-generated electrons and holes. Thus, these electrons can quickly be transferred into the TiO<sub>2</sub> interface and then undergo a transformation as described below:



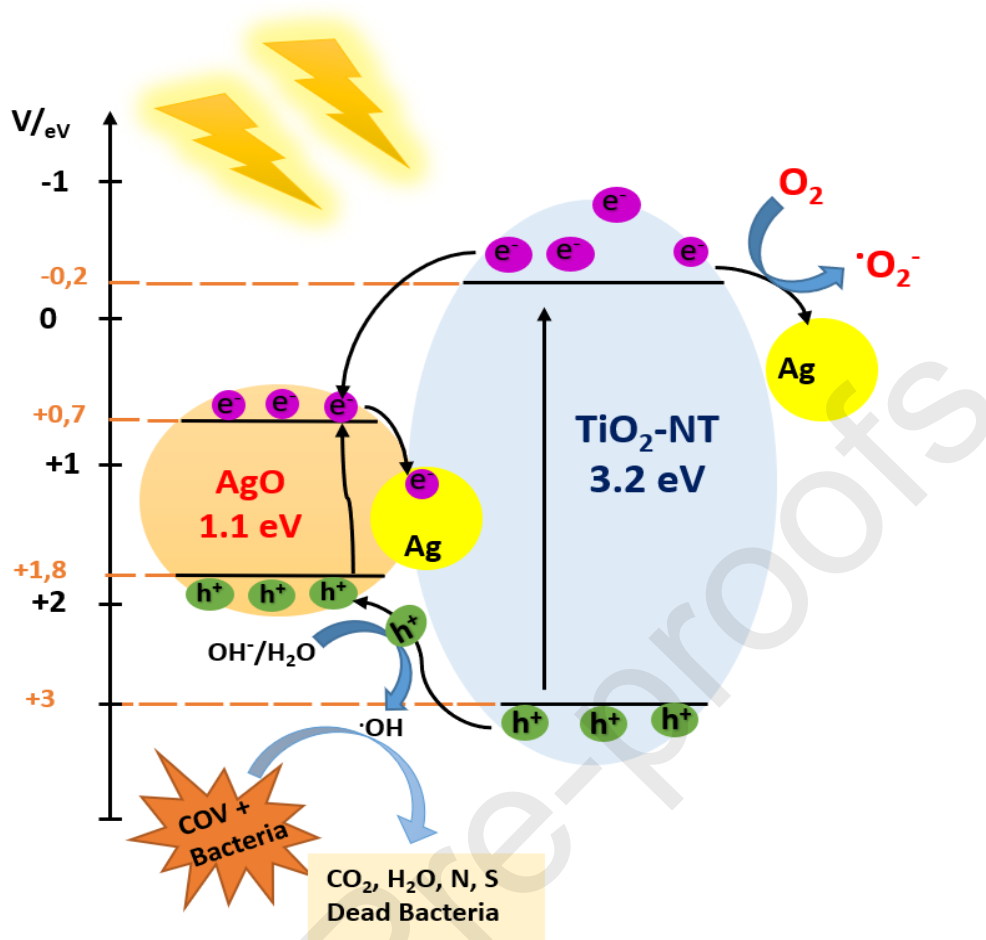
In reaction (4), one can describe it as a multi-electron transfer process. However, the activation energy of such reactions, as follows from the Marcus theory [75], is directly proportional to the square of the number of transferred electrons. The proposed mechanism is then described as the classical chain degenerated-branched oxidation of organic materials (eqs.5-11) [76-78]:



Reaction can be simplified as seen in eq.12:



At the same time, the photo-generated holes in the Valence band of  $TiO_2$  are easily transferred to the Valence band of  $AgO$ . This result could be due to the fact that the Valence band of  $AgO$  (1.84 eV) is more negative than that of  $TiO_2$  (~3eV). Therefore, while the photo-generated electrons flow on the  $TiO_2$  surfaces, the holes are gathered in the  $AgO$  Valence band. Therefore, it was constructed an inter-semiconductor transfer flows for electrons ( $AgO \rightarrow Ag \rightarrow TiO_2$ ) and holes ( $TiO_2 \rightarrow AgO$ ), which significantly limited the charge recombination and improved the charge separation efficiency. The fact that there was effective charge separation resulted in an increase in the lifetime of photogenerated charges, which increased the photocatalytic performance of our photocatalyst. Indeed, adsorbed oxygen ( $O_2$ ) and water ( $H_2O$ ) reacted with electrons ( $e^-$ ) and holes ( $h^+$ ) to produce the superoxide anion ( $O_2^-$ ), hydroxyl radical ( $\cdot OH$ ),  $OH$  and  $H^+$ . The superoxide anion  $O_2^-$  and the hydroxyl radical ( $\cdot OH$ ) were considered to be highly oxidizing reactive oxygen species (ROS). They can decompose VOCs, bacteria and other organic materials adsorbed on the active sites of the nanocomposites into  $CO_2$ ,  $H_2O$  and other products, as shown in eq. (1-12).



**Figure 13.** Proposed mechanism of photocatalysis by Ag-AgO/TiO<sub>2</sub> nanocomposite under light.

#### 4. Conclusion

This paper reports the testing of a novel, uniform and stable Ag<sub>x</sub>O/TiO<sub>2</sub>-PES photocatalyst prepared at low temperature by high power pulsed magnetron sputtering. This is the first study where uniform Ag/Ag<sub>x</sub>O coatings on PES were used for the degradation of indoor air VOCs as well as for bacterial deactivation. We have studied the degradation of two types of concentrated VOCs (butadione and chloroform) using the Ag<sub>x</sub>O/TiO<sub>2</sub>-PES catalyst. We have shown that this catalyst has stable performance during the removal of butadione by Ag<sub>x</sub>O/TiO<sub>2</sub>-PES. Likewise, this catalyst gives almost reproducible results and good recyclability when used under visible irradiation. It also exhibits excellent catalytic stability and has promising potential for application in air treatment. The Ag<sub>x</sub>O/TiO<sub>2</sub>-PES photocatalyst showed high antibacterial activity, which reached almost complete bacterial inactivation within 6 hours for Ag<sub>x</sub>O sputtered at 30 A.

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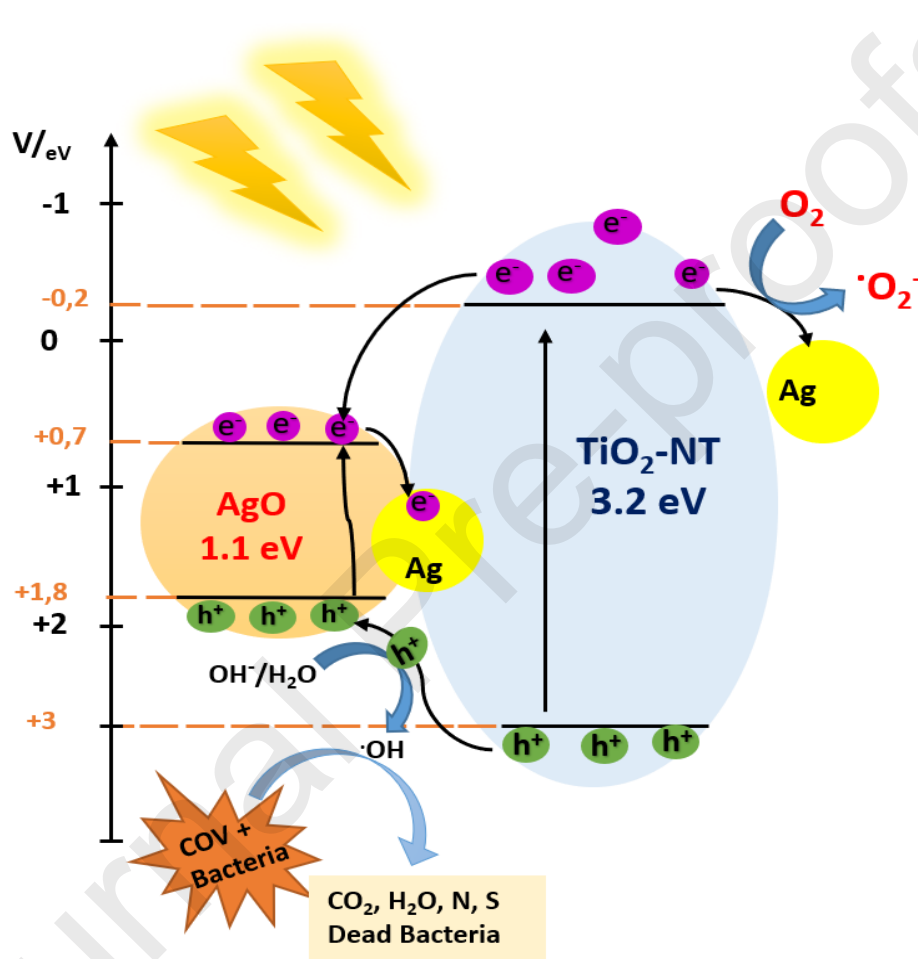
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### Graphical abstract



[79]

### Highlights

- Indoor/outdoor air pollution and bacteria were degraded on sputtered  $Ag_xO/Ag/TiO_2$  on textile.
- Photocatalytic kinetics are carried out in the dark and under visible light.
- $Ag/Ag_xO$  nanoparticles show a significant effect on the activity of  $TiO_2$ .
- A possible mechanism of photocatalytic processes is suggested.

[80]

Journal Pre-proofs