

# Microbiology cell-staining wastewater treatment using TiO<sub>2</sub> thin films

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## Tratamiento de agua residual de microbiología usando películas delgadas de TiO<sub>2</sub>

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### ABSTRACT

Microbiology cell-staining wastewater was treated by UV/TiO<sub>2</sub> thin films photocatalysis. A simple method of applying gravity sedimentation over glass-type substrate was used for the preparation of the films. The use of Na<sub>4</sub>P<sub>2</sub>O<sub>7</sub>, microwaves, and ultrasounds were studied for decreasing the TiO<sub>2</sub> grain sizes on the films. It was established that the best method for reducing grain size resulted from a combination of Na<sub>4</sub>P<sub>2</sub>O<sub>7</sub> (0.01M) and microwave radiation (700 W, 20 min). The Films were characterized by several microscopic and spectroscopic methods. Anatase phase (gap energy of 3.2 eV) and grain sizes averaging 83 nm were achieved. Photocatalysis tests using TiO<sub>2</sub>-films showed 99.5 % of decolorization, 79 % TOC abatement, and total microbial inactivation after 14 h of treatment. No bacteria re-growth was found 48 h after the treatment was completed.

**Key words:** Bacteria, decolorization, mineralization, photocatalysis, wastewater.

### RESUMEN

La fotocatalisis con UV/TiO<sub>2</sub> usando películas delgadas fue empleada para el tratamiento de agua residual de microbiología. Se empleó un método simple de sedimentación por gravedad sobre sustrato de vidrio para la preparación de las películas. El uso de Na<sub>4</sub>P<sub>2</sub>O<sub>7</sub>, microondas y ultrasonido fue estudiado para la disminución de los granos de TiO<sub>2</sub> en las películas. Se estableció que el mejor método para disminuir los agregados resultó de una combinación de Na<sub>4</sub>P<sub>2</sub>O<sub>7</sub> (0.01M) y radiación de microondas (700 W, 20 min). Las películas fueron caracterizadas por métodos microscópicos y espectroscópicos. Se obtuvo la fase anatasa (brecha de energía de 3.2 e.V) y tamaños de grano de 83nm. Los ensayos fotocatalíticos utilizando las películas de TiO<sub>2</sub> generaron un 99.5% de decoloración, 79% de remoción de COT y una inactivación microbiana total luego de 14 h de tratamiento. No se encontró reactivación microbiana luego de 48 h de finalizado el tratamiento.

**Palabras clave:** Agua residual, bacteria, decoloración, fotocatalisis, mineralización.

## 1. Introduction

Alternative water treatment technologies such as Advanced Oxidation Processes (AOPs) have attracted worldwide attention as sustainable technologies, widely used for the removal of organic pollutants not treatable by conventional techniques due to their high chemical stability and/or low biodegradability (Bandala et al., 2008, Oller et al., 2011). Among AOPs, photocatalysis aligns with the “zero” waste scheme in water/wastewater treatments, with a number of important features that have extended their feasible applications in water treatment including mineralization of refractory organic compounds, water pathogens and disinfection by-products (Chong et al., 2010, Foster et al., 2011).  $\text{TiO}_2$  as heterogeneous semiconductor is the most active photocatalyst under the photon energy of  $300 \text{ nm} < \lambda < 390 \text{ nm}$ , with high efficiency, high physical and chemical stability, low toxicity, commercial availability and low cost (Carp et al., 2004, Chong et al., 2010, Gümüş and Akbal, 2011). The degradation of pollutants is achieved by the formation of hydroxyl radicals ( $\bullet\text{OH}$ ) highly reactive and non-selective and other reactive oxygen species (ROS) that mineralize compounds to carbon dioxide and water (Oller et al., 2011).

New achievements in  $\text{TiO}_2$  thin films have promoted its extensive application in photocatalytic water treatment mainly by avoiding agglomeration of the catalysis during operation. This methodology proved to be more efficient, practical and with better disinfection efficiency than colloidal or particulate  $\text{TiO}_2$  suspension (Benabbou et al., 2007). At present, a limited number of methods are used to prepare  $\text{TiO}_2$  thin films that include chemical and physical techniques (Carp et al., 2004, Chong et al., 2010, Gümüş and Akbal, 2011). The sedimentation technique represents a feasible way to prepare  $\text{TiO}_2$  thin films due to its simplicity and low cost. However, a non-uniform substrate coating is obtained due to aggregate formation and uncontrolled surface morphologies caused by electrostatic attraction forces, which is highly detrimental in terms of particles size preservation, surface-area reduction, its reusable lifespan and a decrease in overall efficiency (Chong et

al., 2010, Jiang et al., 2009). In view of this, a simple procedure of  $\text{TiO}_2$  thin films preparation was designed in order to reduce the formation of aggregates by combining certain parameters that could increase particle repulsion and stimulate mechanical destruction of agglomerates.

On the other hand, biological stains are used in laboratories. These chemical compounds are used continually in tests performed for the identification and diagnosis purposes. The dyes used in the staining process include triphenyl-methane and azo-dyes such as Mordants and solvents are also used in different techniques like Gram staining. However, the collected liquid residuals after staining processes result in the formation of wastewater of high toxicity, presence of microorganisms, low light transparency and high organic carbon content (Sioi et al., 2006). Some of those compounds such as crystal violet and malachite green are known for being toxic and carcinogenic, causing health disorders like mutations, cytotoxicity, neurotoxicity, and chromosomal aberrations (Oplatowska et al., 2011). Due to the high toxicity and mutagenicity of these reagents, its degradation is an indispensable step before discharging or even recycling. This study focuses on the evaluation of a novel  $\text{TiO}_2$  thin film elaborated by a simple and inexpensive method and the degradation and disinfection efficiency in photocatalysis in real cell-staining microbiology wastewater. The degradation of organic matter reflected in total organic carbon (TOC) and color removal as well as bacterial inactivation of several existing microbial populations in the wastewater and the durability of the systems in post-irradiation events are evaluated.

## 2. Experimental

### 2.1 Wastewater samples and characterization

The wastewater comes from different microbiology laboratories where cell-staining techniques are performed constantly. It might consist of a combination of different triphenyl-methane dyes such as fuchsine and crystal violet, solvents and mordants like ethanol, acetone and lugol's iodine as

Table 1. Physical and chemical parameters of water quality in cell-staining wastewater.

<i>Parameter</i>	<i>Units</i>	<i>Values</i>	<i>Method</i>
<i>Chemical Oxygen Demand (COD)</i>	<i>mg/L</i>	<i>3750 ± 214</i>	<i>Closed reflux method 5222 D</i>
<i>Biochemical Oxygen Demand (BOD5)</i>	<i>mg/L</i>	<i>1693 ± 215</i>	<i>5-day incubation method 5210 B</i>
<i>Total Organic Carbon (TOC)</i>	<i>mg/L</i>	<i>623 ± 110</i>	<i>Heated-persulfate oxidation method 5310 C</i>
<i>Color</i>	<i>Color Units</i>	<i>1891 ± 9</i>	<i>Spectrophotometric method</i>
<i>pH</i>	<i>Units</i>	<i>3.22 ± 0.03</i>	<i>Electrometric measurement 9040 C</i>
<i>Dissolved Oxygen (DO)</i>	<i>mg/L</i>	<i>7.6 ± 1.97</i>	<i>Membrane Electrode Method 4500-O G</i>
<i>Electrical Conductivity (EC)</i>	<i>mS/cm</i>	<i>0.805 ± 0.006</i>	<i>Laboratory method 2510 B</i>
<i>Settleable Solids (SS)</i>	<i>mg/L</i>	<i>0.3</i>	<i>Gravimetric method 2540 F</i>
<i>Total Suspended Solids (TSS)</i>	<i>mg/L</i>	<i>0.046</i>	<i>Gravimetric method 2540 D</i>
<i>Total Dissolved Solids (SDT)</i>	<i>mg/L</i>	<i>0.062</i>	<i>Gravimetric method 2540 C</i>

reagents of Gram staining as the most extensively performed test in microbiology. Also, other organic compounds like malachite green, Safranin O, Congo red and methylene blue might also be present. Physical and chemical characteristics of the wastewater were determined. The analyses were carried out following the procedures outlined in Standard Methods (Table 1).

Furthermore, in order to approach the bacterial community structure of the wastewater, isolations in plating media were performed. The samples were analyzed for heterotrophic and coliform bacteria in Plate Count Agar and Violet Red Bile agar (Scharlau, Spain), Gram-positive cocci in Baird-Parker (Merk, Germany), bacilli in Starch Media supplemented with polymyxin B (Oxoid, England), and *Pseudomonas* spp. in Cetrimide (Scharlau, Spain). The colonies were counted after incubation at 35 °C for 24 h and the populations were enumerated and expressed as log<sub>10</sub> (CFU/mL). In addition, to estimate the presumptive evidence of the presence of *Pseudomonas* spp., typical *Pseudomonas*-like colonies were examined under 360 nm ultraviolet radiation and a oxidase test strip was performed to the colonies that showed fluorescence (Oxoid, England). Likewise, representative colonies in

Starch Media showing hydrolysis in the plates were selected and evaluated by Gram staining. Gram-positive amyolytic colonies were estimated to potentially belong to the *Bacillus* spp. genus. Finally, an analysis of aerobic endospores was performed (Standard Methods 9218B). All samples were plated in duplicate and a microscopical characterization was carried out on the basis of Gram staining.

## 2.2 TiO<sub>2</sub> thin films preparation and characterization

TiO<sub>2</sub> USP Industrial grade was employed. In order to filter grain sizes (100 nm), this material was settled by gravity fourfold before use. In order to improve the effective surface of the films, in terms of aggregates formation (grain sizes), a number of experiments were performed establishing the best method for the films preparation (Table 2). A TiO<sub>2</sub> suspension in deionized H<sub>2</sub>O was prepared (0.05 % w/v). The suspension was adjusted to different pH values (1.6 and 9.0) and Na<sub>4</sub>P<sub>2</sub>O<sub>7</sub> (0.01 or 0.005 M) was added. Then, the suspension was either microwave irradiated for 20 min (conventional microwave, 700 W) or sonicated for 5 min (90

**Table 2.** Experimental design for the selection of the best conditions for TiO<sub>2</sub> thin films growth.

<i>Method</i>	<i>pH</i>	<i>Na<sub>4</sub>P<sub>2</sub>O<sub>7</sub> (M)</i>	<i>Equipment</i>
1	9.0	0.01	Microwave (20 min)
2	9.0	0.01	Ultrasound (5 min)
3	9.0	0.005	Ultrasound (5 min)
4	9.0	0.005	Microwave (20 min)
5	1.6	0.005	Microwave (20 min)
6	1.6	0.005	Ultrasound (5 min)
7	1.6	0.01	Ultrasound (20 min)
8	1.6	0.01	Ultrasound (5 min)

W, Fisher–Scientific FS20) (Jiang et al., 2009, Paušová et al., 2011). Films were elaborated using soda lime glasses (26 X 20 mm) as substrate, which were previously immersed in a solution of H<sub>2</sub>SO<sub>4</sub>/H<sub>2</sub>O<sub>2</sub> (3:1 v/v) in order to clean the surface. Two layers of TiO<sub>2</sub> were deposited on the substrate by simple gravity sedimentation. Films were thermally treated at 450 °C for 1 h.

The films topography was analyzed using Atomic Force Microscopy (AFM) (Nanosurf Easy Scan 2 AG) in contact mode, in order to establish the best conditions for photocatalyst preparation according to homogeneity of film surface. TiO<sub>2</sub> thin films obtained by the chosen method were characterized by Scanning Electron Microscopy (SEM) in a JEOL JSM 6490–LV electron microscope. X–ray diffraction was measured in a Siemens D–5000 diffractometer, employing an anode of Cu K $\alpha$ . UV–VIS spectra was determined using a quartz tungsten–halogen lamp (240–800 nm) coupled with a mono–chromator (Acton Research spectral pro 775), interface Lab view software® was used to process the data. FT–IR measurements were performed in Shimadzu IR prestige 21 equipment using the KBr pellets method. Raman analysis was performed in Labram equipment (Internal laser of He–Ne of 20 mW,  $\lambda = 400$  nm).

### 2.3 Description of the reactor and experimental procedure

All photocatalytic experiments were carried out in quartz glasses (24.7 X 4.7 cm) used as batch-type reactors with 0.250 L of effective volume. Two 15 W UV lamps emitting radiation at 254 nm were used as the light source. The lamps were placed 5 cm away from the glasses. Each quartz glass contained 12 films of the selected method (12 mg of TiO<sub>2</sub> each), the films were placed vertically and kept in dark for 30 min to reach adsorption-desorption equilibrium. The system operated at 120 rpm at room temperature during 14 h. Three control experiments were performed for the photocatalytic experiments: UV photolysis, TiO<sub>2</sub>/adsorption in dark and absolute control by triplicate.

### 2.4 Bacterial reappearance experiments

The disinfection durability of the photocatalysis treatment was tested post-irradiation for total bacterial inactivation. After irradiation time, the water was kept in the dark and samples were taken at 24 h and 48 h and exanimated for the presence of heterotrophic bacteria, total coliforms and Gram-positive bacilli.

## 2.5 Analysis and statistical analysis

In order to evaluate the efficiency of the treatments, the % removal of organic matter; Chemical Oxygen Demand (COD), Total Organic Carbon (TOC) and Color Units (CU) were calculated. Pseudo-first-order rate constants  $k$  in  $\text{min}^{-1}$  were calculated for CU decrease and TOC abatement. The mineralization rate was characterized qualitatively in terms of TOC removal.

Absorbance spectra were obtained to determine the level of degradation of the wastewater. A UV-Vis spectrophotometer (Thermo Scientific Evolution 60) was used with a 1 cm quartz cell. To evaluate disinfection efficiency, the bacterial count was monitored in the three bacterial populations with the highest (Colonies Forming Units) CFU/mL isolated from the wastewater: heterotrophs, total coliforms and Gram-positive bacilli. Microbial inactivation is presented as percentage of inactivation. Assuming that the bacterial inactivation efficiency followed a pseudo-first order kinetics with respect to bacterial colony count ( $N_t$ ), inactivation rate constants were analyzed.

The results obtained are presented in averages and standard deviations. A Shapiro-Wilk test for normality and Levene tests for homogeneity of variance were performed to verify data normality. Differences among measurements of removal and inactivation achieved by both treatments were assessed with One-way ANOVA (IBM SPSS® Statistics software version 20, Statistix 9.0®) and a post-hoc HSD Tukey test.  $P < 0.05$  was used to determine statistical significance of the results obtained.

## 3. Results and discussion

### 3.1 Wastewater characterization

#### 3.1.1 Physical and chemical parameters

The wastewater was intensely colored (1890.7 CU), presented a considerable high organic content: the mean COD, BOD5 and TOC in the wastewater were 3750.8 mg/L, 1693 mg/L, and

623.2 mg/L respectively. Other characteristics of the wastewater are included in Table 1. The results indicate that chemical substances such as dyes, mordants and solvents and others used in biological staining processes contribute highly to the pollution of this water. Color and COD/BOD are the most obvious indicators of water pollution and the discharge of highly colored dye effluents is aesthetically displeasing and cause severe environmental problems (Chong et al., 2010). Together, with acidic pH makes this wastewater imperative to treatment before being discharged into sewage systems or consider its reuse.

#### 3.1.2 Microbial analysis

The substances present in the wastewater are generally considered antimicrobial agents (i.e. Crystal violet, malachite green, ethanol, iodine etc.). However, a high bacterial population was found. Only four groups of bacteria were successfully identified. Overall the number of bacteria in each group was relatively high. Heterotrophic bacteria were the largest microbial population isolated (7.86 log CFU/mL), followed by total coliforms (6.56 log CFU/mL), Gram-positive bacilli (6.37 log CFU/mL) and *Pseudomonas* spp (4.71 log CFU/mL). The isolation of these microorganisms from the wastewater indicates natural adaptation to survive to the conditions of this particular wastewater, toxic bactericidal substances including dyes and solvents, and acidic pH. The higher concentration of bacteria in the wastewater could be explained either by the formation of bacterial spores or by biofilms at the surface of the containers where the water is kept (Jefferson, 2004).

However, no colonies were observed from all dilutions for spore analysis ( $< 10$  spores/mL were obtained). Gram-positive bacilli spore-forming are the species of *Bacillus* spp. and *Clostridium* spp. Even though aerobic amylolytic Gram-positive bacilli were observed, it is likely that certain compounds present in the cell-staining wastewater like ethanol, acid, iodine and dyes as methylene blue inhibited its sporulation activity (Bohin et al., 1976, Kearns et al., 2005, Setlow et al., 2002). It has shown that in the spore forming bacterium

*Bacillus subtilis* the efficiency of spore formation can be reduced more than 100-fold by alcohol stress (Bohin et al., 1976). The general stress genes and the molecular mechanisms considered to be involved in the inhibitory effect of alcohol stress on sporulation in *B. subtilis*, has been previously studied (Gottig et al., 2005, Reder et al., 2012). Likewise, spores can be photoinactivated by phenothiazinium dyes such as methylene blue, toluidine blue O and dimethylmethylene (Demidova and Hamblin, 2005).

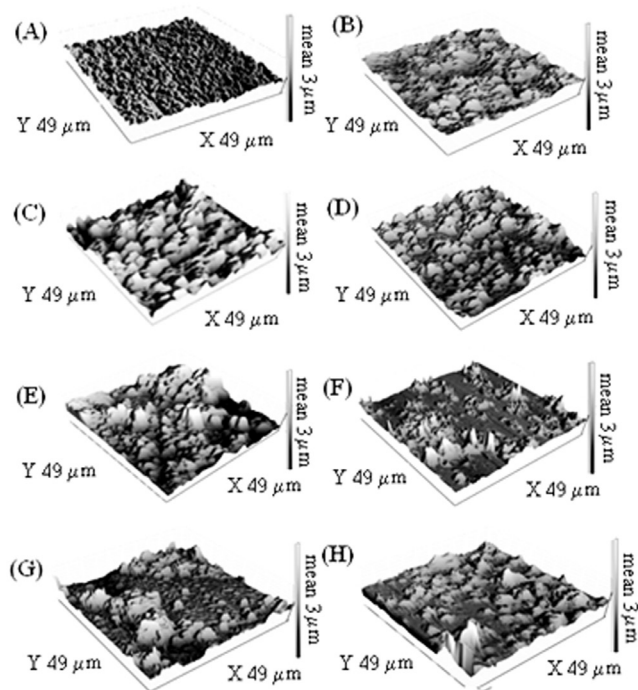
On the other hand, survival mechanisms including the formation of biofilm in Gram-positive and Gram-negative bacteria have been well studied (Andersson et al., 2008, Kearns et al., 2005). Organisms within biofilms can withstand nutrient deprivation, pH changes, oxygen radicals, disinfectants, and antibiotics making it possible to microorganisms to survive in given conditions of the cell-staining wastewater (Andersson et al., 2008, Jefferson, 2004). It has been reported that certain species can induce biofilm formation of other bacteria without out-competing them, like *B. cereus* strong EPS producers could be used

as anchors for attachment of adherence-deficit strains. Thus a wide range of species can survive in environmental challenging conditions that would not grow as pure cultures. A synergistic effect on metabolism and biosynthesis of non- or weak EPS formers such as *E. coli* can be seen when together formed a strong biofilm in wastewater medium (Andersson et al., 2008). Similarly, *Pseudomonas aeruginosa* is reported to form biofilm in dual cultures and usually found to be one of the dominating species, which could explain its relatively high concentration in the wastewater.

The profile of isolated bacteria in cell-staining wastewater must be taken into consideration as these results can serve as a tool for the design of efficient photocatalytic systems in the disinfection and/or degradation of wastewater.

### 3.2 TiO<sub>2</sub> thin film preparation and characterization

The AFM images of the evaluated combinations of variables (Fig. 1), suggest that method 1 (pH 9.0, Na<sub>4</sub>P<sub>2</sub>O<sub>7</sub> 0.01 M, 20 min of Microwave radiation)



**Figure 1.** TiO<sub>2</sub> thin films surfaces AFM images: (A) Method 1, (B) Method 2, (C) Method 3, (D) Method 4, (E) Method 5, (F) Method 6, (G) Method 7, (H) Method 8.

showed the best results in terms of substrate coating and decrease of aggregation of  $\text{TiO}_2$  particles over the substrate. This was due to the factors combined in the experiment. Because of its isoelectric point,  $\text{TiO}_2$  is charged negatively at alkaline conditions, this caused charge repulsion between  $\text{TiO}_2$  particles (Konstantinou and Albanis, 2004). Additionally, at this pH, sodium pyrophosphate is ionized, increasing the medium ionic strength, which interferes with particle aggregation (Jiang et al., 2009). Moreover, the electromagnetic field created by microwave radiation caused collisions between  $\text{TiO}_2$  aggregates leading to their fragmentation (Al-Harashseh and Kingman, 2004).

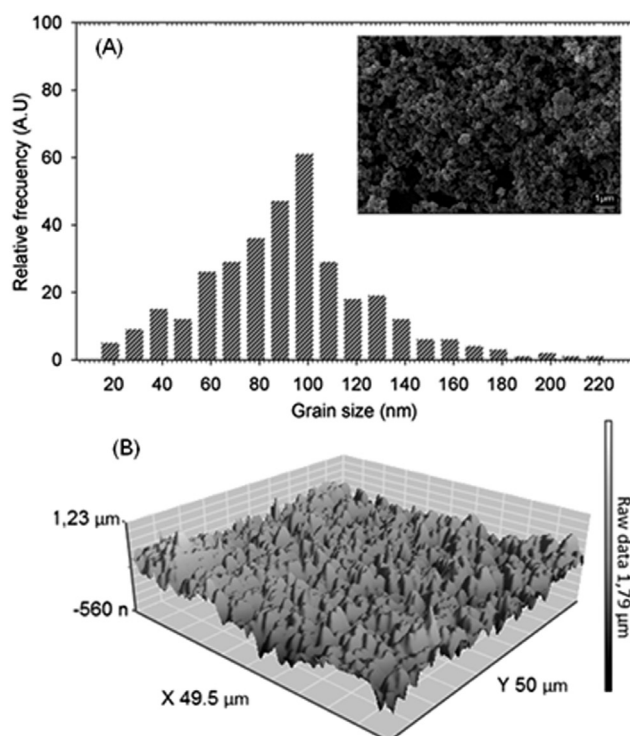
Once the method 1 for  $\text{TiO}_2$  thin films preparation was established, the films were characterized by microscopic and spectroscopic techniques. From Fig. 2-A it is observed that the higher frequencies of grain size ranged between 90 and 100 nm. The imperfection formed by the material can improve microbial inactivation since it can act as microscopic cavities, microorganisms, such as bacteria and yeast which average size are between 2 and 10  $\mu\text{m}$ ,

could be trapped and potentially inactivated by the photocatalysis process. Affirmed that material structural defects associated with forms and different grain sizes can increase the number of catalytic sites (Acosta et al., 2005).

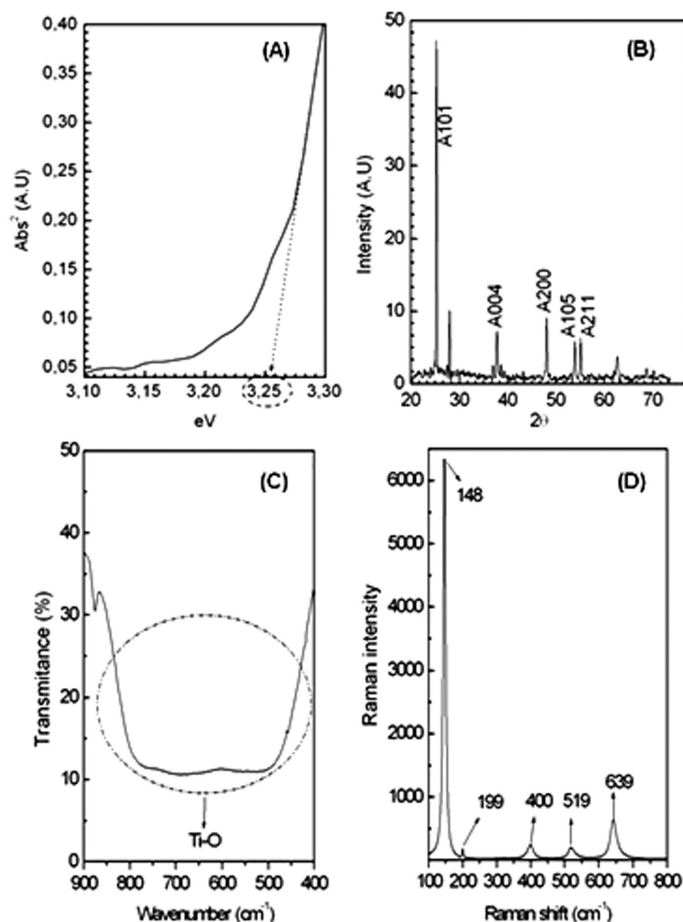
AFM high-resolution images showed that  $\text{TiO}_2$  grains formed small aggregates between 4 and 5  $\mu\text{m}$  in size. The films obtained from this methodology appear to be rough ( $136,34 \text{ nm} \pm 29,79, 39,18 \mu\text{m}^2$ ) (Fig. 2-B). This is an important aspect since it can facilitate bacteria adherence to the film allowing bacterial inactivation (Lin et al., 2008).

### 3.2.1 Spectroscopic analysis

Spectroscopic analysis was carried out in order to determine band gap energy and crystalline phase of photocatalysts. UV-Vis spectroscopy (Fig. 3-A) indicated that energy band gap corresponds to a 3.2 e.V. (387 nm), which is related to anatase band gap energy (Gaya et al., 2009, Jin-hui, 2012). From XRD profiles one can observe diffraction signals for *101*, *004*, *200*, *105* and *211* crystallographic planes,



**Figure 2.** (4a) Histogram on the grain size from SEM results, ImageJ® software was employed for measurements, (4b) AFM image of  $\text{TiO}_2$ -films growth by method 1.



**Figure 3.** Vibrational spectroscopy for  $\text{TiO}_2$ , (A) FT-IR spectroscopy and (B) Raman spectroscopy. (C) Band-Gap energy determination by UV-Vis spectra (D) XRD for  $\text{TiO}_2$  thin films.

indicating that anatase phase was predominantly found (Fig. 3-B). A broad IR signal between 500 and 700  $\text{cm}^{-1}$  has been related with Ti-O tension and associated with the presence of anatase phase (Fig. 3-C). Moreover, in Fig. 3-D, Raman Spectra showed vibrational modes, which are assigned to the previously mentioned crystalline phase, in 148, 199, 400, 519, and 639  $\text{cm}^{-1}$  (Wang et al., 2011).

### 3.3 UV/ $\text{TiO}_2$ photocatalysis tests

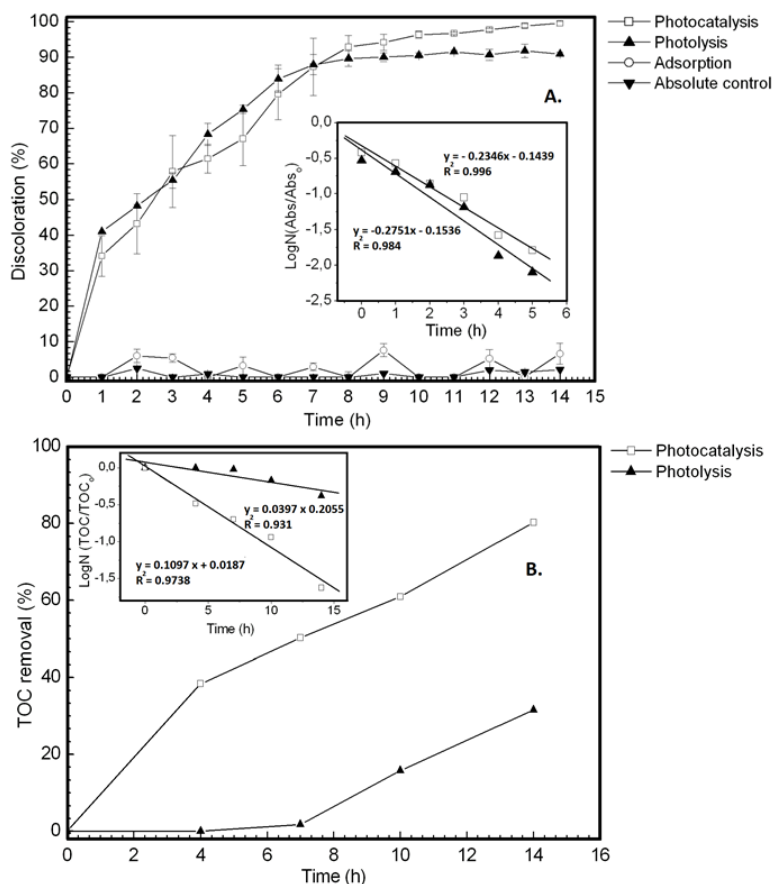
#### 3.3.1 Color removal and UV-Visible absorption changes

The photocatalysis treatment achieved 99.5 % (8.8 CU) of decolorization, while the UV photolysis control only 91.3 % (152.8 CU) and

$\text{TiO}_2$ /adsorption in dark control 6.8 % (1,623.7 CU) of color removal after 14 h. During the first 7 h of treatment decolorization percentages were similar for photocatalysis and photolysis (87.3 %). Decolorization kinetic rate constants obtained were similar as well ( $k = 0.2346 \text{ min}^{-1}$   $R^2 = 0.9963$  and  $k = 0.2751 \text{ min}^{-1}$   $R^2 = 0.9846$ , respectively) (Fig. 4-A, inlet). From Fig. 4-A, it is easy to show that photolysis is as effective as photocatalysis ( $p < 0.05$ ) and, therefore, plays an important role in the degradation of dyes in the wastewater, similar results are found in the literature (Barka et al., 2008). Adsorption control experiment showed no significant adsorption of the dyes onto the  $\text{TiO}_2$  surface

Comparing UV/ $\text{TiO}_2$  photocatalysis and photolysis from 7 h to 14 h, an increase in the





**Figure 4.** (A) Wastewater decolorization by UV/TiO<sub>2</sub> photocatalysis. Inlet: experimental data adjusted to pseudo-first-order kinetic model. (B) TOC abatement by UV/TiO<sub>2</sub> photocatalysis. Inserts represent mean values of a complex sample. Inlet: data adjusted to pseudo-first-order kinetic model.

percentage of color removal for photocatalysis can be observed. In agreement with TiO<sub>2</sub> photocatalysis systems in highly colored wastewater such as dye and textile wastewater, an efficient photocatalytic degradation and color removal can be achieved between UV/TiO<sub>2</sub> and photolysis. However, photo-oxidation and degradation efficiency in real dye wastewater depends strongly on the characteristics of the wastewater such as dye structure and other present organic compounds, irradiation time and UV lamp, etc (Al-Momani et al., 2002). It can be suggested that after 7 h of treatment, UV radiation could not penetrate effectively through the highly colored wastewater; hence, films were not successfully irradiated. Once the CU decreased and UV light reached the film surface, the wastewater was 100 % discolored

by the photocatalysis treatment. It should be emphasized, that photolytic treatment showed high percentages of decolorization. This is an interesting fact since, to the author's knowledge; triphenyl-methane dyes such as crystal violet in dilution has not been reported high decolorization rates through direct photolysis. The presence of other compounds such as solvents possibly contributed as electron donors under UV light irradiation.

### 3.3.2 TOC abatement

According to analysis of TOC abatement measurements, a clear difference between treatments and controls can be observed in Fig. 4-B. TiO<sub>2</sub> photocatalysis resulted in a significant 79.5 % of TOC mineralization, compared to 31.6

% by the use of UV light after 14 h. However, the complexity of real wastewater, as the cell-staining wastewater here studied, containing several compounds and high microbial population results in a longer treatment time. It is important to mention that TOC abatement was slower in comparison to decolorization, possibly the chromophore rupture and by-product formation could be involved in a slow TOC abatement. Similar results were reported

by Li (Li and Li, 2001), who degrade methylene blue, reaching 43.7 % of TOC abatement after 2 h of treatment. The process followed a pseudo-first-order kinetic (Fig. 4-B inlet). In photocatalysis, the  $k$  obtained value was higher ( $k = 0.197 \text{ min}^{-1} R^2 = 0.9738$ ) in comparison to photolysis ( $k = 0.0397 \text{ min}^{-1} R^2 = 0.931$ ) therefore; a faster organic matter mineralization was obtained with UV/TiO<sub>2</sub> photocatalysis.

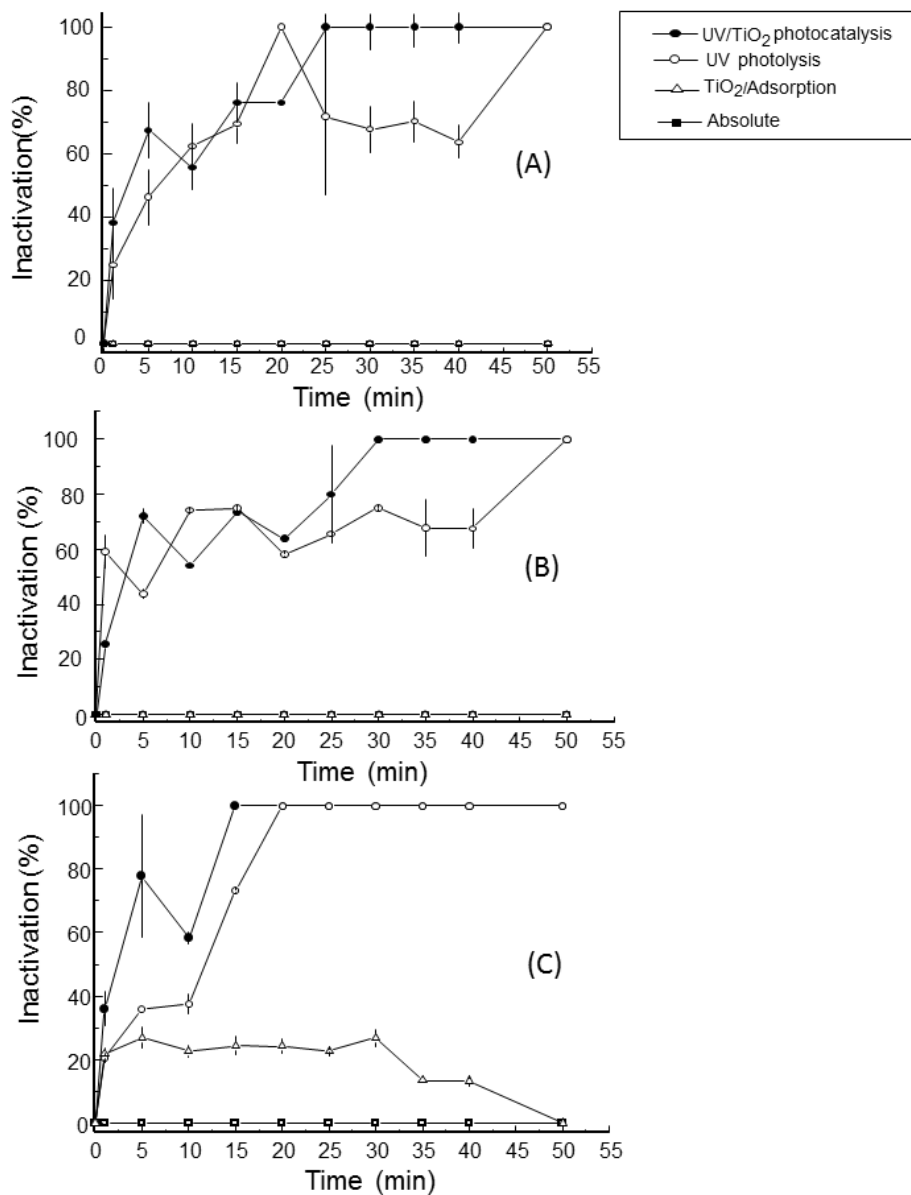


Figure 5. Microbial inactivation. A) Heterotrophic bacteria. B) Gram-positive bacilli. C) Total coliforms.

### 3.4 Bacterial inactivation and post-irradiation events

The efficiency of the treatment in the inactivation of three mayor populations found in the water was evaluated. Photocatalysis inactivated 100 % of all population after 14 h of treatment. Heterotrophic bacteria were completely inactivated after 25 min (Fig. 5-A), total coliforms after 20 minutes (Fig. 5-B) and Gram-positive bacilli were more resistant after 50 minutes of treatment (Fig. 5-C). The results showed the so-called shoulders in which bacteria repair the damage caused by reactive oxygen species (ROS). As the generation of hydroxyl radical progresses, is followed by an exponential decrease named tail, in which bacteria are no longer capable of self-repairing and eliminate intracellular ROS called "oxidative stress" leading to lysis of the cell and release of intracellular components (Chen et al., 2010).

The analysis of inactivation rate constants determined Gram-positive bacilli ( $k = 0.2677 \text{ min}^{-1}$ ,  $R^2 = 0.9240$ ) as the more resistant group to degradation by hydroxyl radicals, compared to total coliforms ( $k = 0.8744 \text{ min}^{-1}$ ,  $R^2 = 0.9946$ ). The photocatalytic inactivation of bacterial groups has been widely reported; where the inactivation rate increases with increasing cell wall thickness thus

hydroxyl radicals and other ROS are absorbed in the peptidoglycan layer lowering the number of radicals that damage the DNA (Pal et al., 2007, Rincón and Pulgarin, 2006). It has been reported that Gram-negative bacteria can be easily affected by ROS. Water disinfection studies using  $\text{TiO}_2$  affirmed that comparing the Gram-negative bacteria *E. coli* with *Staphylococcus aureus* (Gram-positive), the latter appeared to be more resistant to the photocatalysis process (Kühn et al., 2003).

Results of secondary reactivation are shown in Table 3. The photocatalysis experiment proved to be efficient in the degradation of microorganism since no bacterial growth was detected after 24 h and 48 h in dark. Reactivation occur only after photolysis due to bacteria that lost their cultivability during irradiation time but activated resistance to stressful conditions and recover their cultivability when they were introduced into more favorable conditions (Benabbou et al., 2007). This effect has been reported by other researchers and attributed to bacterial UV-induced self-defense and activation of post-treatment recovery mechanisms. Hence, a bacteriostatic effect and activation of dark repair systems could be observed, reconstitution of cellular components and DNA such as those found in *E. coli* could occur (Benabbou et al., 2007, Rincón and Pulgarin, 2004).

Table 3. Bacterial reactivation test after photocatalysis treatment.

Treatment	Microbial group	Method	
		UV/TiO2	UV
60 min of photocatalysis (Log CFU/mL)	Heterotrophic bacteria	< 1*	< 1
	Total coliforms	< 1	< 1
	Gram positive bacilli	< 1	< 1
24 h after photocatalysis (Log CFU/mL)	Heterotrophic bacteria	< 1	2.49
	Total coliforms	< 1	< 1
	Gram positive bacilli	< 1	2.35
48 h after photocatalysis (Log CFU/mL)	Heterotrophic bacteria	< 1	4.93
	Total coliforms	< 1	4.14
	Gram positive bacilli	< 1	4.90

\* < 1 means no bacterial growth detected.

#### 4. Conclusion

A simple TiO<sub>2</sub> thin films preparation based on sedimentation and a combination of pH 9.0, Na<sub>4</sub>P<sub>2</sub>O<sub>7</sub> 0,01 M, and microwave radiation for 20 min gave the best results with a uniform deposition on the substrate. Additionally, microscopic analysis showed that the higher frequencies of grain size ranged between 90 and 100 nm, and spectroscopic analysis indicates that the predominant crystalline phase in the material is anatase with gap value of 3.2 eV. The photocatalysis treatment led to 99.50 % of color removal and 79 % of TOC abatement in 14 h. As a result, mineralization was a slow process compared to decolorization, as it could be associated with the formation of transformation by-products. Likewise, the microbial population was totally inactivated in less than 1 h and no re-growth was found after 48 h of treatment. This result indicates that the oxidative species developed at the TiO<sub>2</sub> surface caused severe damage to the cells. Therefore, disinfection durability of the photocatalysis treatment was attained.

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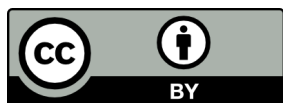
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