

WASTEWATER DISINFECTION BY SOLAR HETEROGENEOUS PHOTOCATALYSIS: EFFECT ON TETRACYCLINE RESISTANT/SENSITIVE *Enterococcus* strains

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ABSTRACT

Solar simulated heterogeneous photocatalysis (SSHP) with suspended TiO₂ was investigated in the inactivation of tetracycline resistant/sensitive *Enterococcus* (TRE/TSE) strains in the effluent of an urban wastewater treatment plant (UWTP). The effect of solar simulated disinfection (SSD) on the inactivation of the same *Enterococcus* strains was investigated as control. SSHP process (0.05 g l⁻¹ of TiO₂) was found to be effective in the inactivation of both *Enterococcus* strains with total inactivation (~7 log unit) observed after 60 min of irradiation. On the contrary, SSD process did not show any significant inactivation after 90 min of irradiation. The effect of both processes on the antibiotic resistance phenotypes of the surviving enterococci was also evaluated. TRE cells surviving the SSHP treatment showed that disinfection process did not affect the antibiotic resistance pattern after 45 min irradiation. The same was observed for the TSE strain. Accordingly, antibiotic resistance can spread into the receiving water body when antibiotic resistant strains survive to disinfection process.

Keywords: advanced oxidation processes, antibiotic resistant bacteria, bioindicator, urban wastewater, wastewater reuse.

1. Introduction

Because of the intensive use of antibiotics for human (domestic and hospital use), veterinary and agriculture purposes, these compounds are continuously released into the environment from different anthropogenic sources (Brown *et al.*, 2006; Kümmerer, 2009; Grassi *et al.*, 2013). In addition to the adverse effects as chemical pollutants of the environment, antibiotics are also associated with the spread of antibiotic resistant bacteria (ARB) and antibiotic resistance genes (ARGs). The contamination of the environment and the related risk for human health through the food chain is presently considered a serious public health problem as confirmed by the World Health Organization (WHO) which identified the development of antibiotic resistance as one of the major global threats to the human society (WHO, 2013).

Urban wastewater treatment plants (UWTPs) are suspected to be among the main anthropogenic reservoirs for the development of antibiotic resistance because of the simultaneous occurrence of antibiotics, ARGs and ARB (Novo and Manaia, 2010; Michael *et al.*, 2013; Rizzo *et al.*, 2013a). UWTPs are not designed to remove these contaminants and consequently they are released into the effluent

almost unaffected, possibly promoting antibiotic resistance spread in the environment (Lupo *et al.*, 2012). Moreover, wastewater reuse, especially for irrigation purposes in areas with limited rainfall, is a current practice, but the risk for human health related to the uptake of ARB is almost unknown (Fatta-Kassinos *et al.* 2011; Gatica and Cytryn, 2013).

Disinfection process (when used) should be the ultimate barrier for controlling antibiotic resistance spread before UWTP effluent disposal or reuse, but conventional disinfection processes (namely, chlorination and UV radiation) may not effectively control the release of ARB and ARGs (Rizzo *et al.*, 2013b). This drawback calls for the study of new/alternative processes, such as advanced oxidation processes (AOPs), which have been successfully investigated in water/wastewater treatment and disinfection (Rizzo *et al.*, 2008; Lofrano *et al.*, 2009; Zapata *et al.*, 2010; Rizzo *et al.*, 2013c). Among AOPs, heterogeneous photocatalysis with TiO₂ has recently emerged as an interesting water disinfection option (Dunlop *et al.*, 2011; Robertson *et al.*, 2012) because it does not result in the formation of toxic disinfection by-products compared to chemical disinfectants (Richardson *et al.*, 1996) and it can be operated with solar radiation thus saving money for energy (Malato *et al.*, 2009). Solar driven photocatalysis could be an interesting disinfection option alternative to conventional processes, in particular in arid and semi-arid areas, to improve the quality and safety of the effluent from small wastewater treatment plants (e.g., < 2000 people equivalent) before reuse.

The most studied bacteria for antibiotic resistance in wastewater belong to the common indicators of faecal contamination, namely coliforms and enterococci. Among coliforms, *Escherichia coli* have been recently used as indicators of antibiotic resistance to evaluate the effect of conventional (Templeton *et al.*, 2009; Rizzo *et al.*, 2013b; Huang *et al.*, 2013) and new disinfection processes (Rizzo *et al.*, 2014a). The effect on antibiotic resistant enterococci strains was investigated using TiO₂ photocatalysis (Tsai *et al.*, 2010) and solar photo-Fenton (Michael *et al.*, 2012), .

Tetracyclines belong to a group of broad-spectrum antibiotics generally used in the treatment of infections of the urinary and gastrointestinal tracts. Accordingly, tetracycline is one of the most frequently detected antibiotics in wastewater (Watkinson *et al.*, 2007; Michael *et al.*, 2013) and different *Enterococcus* spp. (among which *Enterococcus faecalis* and *Enterococcus faecium*) were found to be capable of acquiring resistance to tetracycline (Cauwerts *et al.*, 2007).

In the present work, two *Enterococcus* spp. strains, one tetracycline resistant (TRE) and one sensitive (TSE) were first isolated from the effluent of a biologically treated urban wastewater and used as bio-indicators to test the disinfection efficiency of solar driven TiO₂ photocatalysis process. The effect of disinfection process was evaluated based on *Enterococcus* survival and antibiotic resistance of surviving *Enterococcus* colonies.

2. Material and methods

2.1 Wastewater samples

Wastewater samples were taken from an UWTP located in the province of Salerno (Italy), from the effluent of the biological process (activated sludge). The UWTP also receives sewage from a hospital. The samples were collected in sterilized 1 l amber glass bottles and analyzed for pH, BOD₅, COD, TSS and conductivity.

2.2 Selection and identification *Enterococcus* strains

Resistant/sensitive *Enterococcus* strains were isolated from UWTP effluent sample on Slanetz Bartely Agar (SBA) culture medium (20 g l⁻¹ tryptose, 5 g l⁻¹ yeast extract, 2 g l⁻¹ glucose, 4 g l⁻¹ dipotassium phosphate, 0.4 g l⁻¹ sodium azide, 10 g l⁻¹ bacteriological agar, 0.1 g l⁻¹ TTC), using the membrane filtration method. A resistant *Enterococcus* strain was selected on SBA culture medium supplemented with 16 mg l⁻¹ of TET (the minimal inhibitory concentration according to EUCAST (2013) data base). Briefly, 5 ml of wastewater or its serial dilutions were added to 45 ml of saline solution and filtered through membranes; membranes were subsequently transferred on SBA culture medium and incubated

for 48 h at 37 °C. A confirmation test was conducted by transferring the membranes with typical red colonies onto Bile Esculin Azide (BEA) agar (17 g l⁻¹ tryptose, 3 g l⁻¹ peptone, 5 g l⁻¹ yeast extract, 10 g l⁻¹ ox bile, 5 g l⁻¹ sodium chloride, 1 g l⁻¹ esculin, 0.5 g l⁻¹ iron ammonium citrate, 0.15 g l⁻¹ sodium azide, 13 g l⁻¹ agar). Some colonies were randomly picked up and frozen in 15% glycerol Tryptone Soy Broth (TSB) at -20 °C.

2.3 Inoculum and sample preparation

Autoclaved (15 min at 121 °C) wastewater samples were inoculated with TRE and TSE strains. For disinfection experiments, fresh cultures of the selected strains were inoculated in 500 ml sterile wastewater sample at a density of 10⁷ colony forming units (CFU) 100 ml⁻¹ (0.5 McFarland). To evaluate the bacterial inactivation achieved, bacteria were enumerated by the spread plate method on SBA. Culture plates yielding corresponding to 20-200 colonies were the basis for bacterial counts.

2.4 Photocatalytic tests

Photocatalytic experiments were carried out in 2.2 l cylindrical glass batch reactor (13.0 cm in diameter) filled in with 500 ml wastewater sample (5.0 cm water height). The reactor was placed in a water bath to maintain the temperature constant (roughly 30 °C) during the experimental procedure. The wastewater solution was continuously stirred during the experiments. Solar irradiation was provided by a wide spectrum 250 W lamp equipped with a UV filter (Procomat, Italy), fixed at 40 cm from the upper water level in the reactor. A spectrometer model HR-2000 from Ocean Optics (Florida, USA), equipped with cosine corrector with Spectralon diffusing material, was used to measure irradiance spectra of UV lamp (emission spectrum 340-460 nm; average intensity: 53.46 μW cm⁻²; max intensity (375 nm): 104.0 μW cm⁻²). 0.05 g TiO₂ l⁻¹ was used in photocatalytic experiments, according to our previous results (Rizzo *et al.*, 2014b). The autoclaved wastewater sample with the TiO₂ powder (Degussa P25) was sonicated for 5 min before inoculum addition. Control tests without any photocatalyst addition (solar photolysis tests) were also performed.

2.5 Antibiotic resistance assay

The antibiotic resistance patterns of the bacterial isolates prior to and after the photocatalytic treatment was tested by Kirby–Bauer method. Briefly, the colonies that survived to photocatalytic treatment and grew on SBA medium were selected (4-5 colonies randomly selected from each one agar/irradiation time) and transferred to 10 ml physiological solutions, respectively, to achieve 10⁷ CFU 100 ml⁻¹ (0.5 McFarland). Bacterial suspensions were spread onto Mueller Hinton agar (Biolife, Italy) using a sterile cotton swab. Antibiotic-impregnated discs of ampicillin (AMP) (10 mg), ciprofloxacin (CIP) (5 mg), tetracycline (TET) (30 mg) and vancomycin (VAN) (30 mg) (all from Biolife) were placed on the surface of each inoculated plate. After 24 h of incubation at 37 °C, the diameters of antibiotic inhibition of growth were measured. The procedure was duplicated and the average values plotted. The results were compared with antibiotic resistance and inhibition diameter (mm) of *Enterococcus faecalis* for Kirby-Bauer method available in EUCAST (2013) database and summarized in Table 1.

Table 1: Summary of antibiotic resistance and inhibition diameter values (mm) of *Enterococcus faecalis* for AMP, CIP, TET and VAN (Kirby-Bauer method) available in EUCAST database (2013).

| | AMP | CIP | TET | VAN |
|------------------|------|---------|--------|------|
| Resistant (R) | ≤ 16 | ≤ 15 | ≤ 14 | < 16 |
| Intermediate (I) | - | 16 - 20 | 15 -18 | - |
| Sensitive (S) | ≥ 17 | ≥ 21 | ≥ 19 | > 16 |

3. Results and discussion

3.1 Wastewater characteristics

The wastewater characteristics of the sample taken from the effluent of the biological process meet the Italian standards (for the measured parameters) for wastewater disposal into surface water but do not meet the standard set by Italian legislator for TSS for wastewater reuse (Table 2).

3.2 Selected *Enterococcus* strains

In order to compare the effect of the solar photocatalytic process on *Enterococcus* strains, two colony types that maintained viability after treatment were characterized for the antibiotic resistance pattern (Table 1). In particular, one strain was chosen because was resistant to TET (and sensitive to CIP and VAN) (TRE) and the other one was chosen because did not show any resistance to the tested antibiotics (TSE). The occurrence of antibiotic resistant *Enterococcus* in UWTP effluents is quite common, in particular when mixed with hospital wastewater effluents. Vancomycin and ciprofloxacin resistant bacteria were detected in hospital effluent and in raw UWTP inflow, being significantly more prevalent in the hospital (Varela *et al.*, 2013); in particular, most of the vancomycin resistant enterococci isolated from the hospital effluent presented multidrug-resistance phenotypes to different antibiotics among which ciprofloxacin and tetracycline.

Table 2: Comparison of wastewater sample characteristics with Italian standards for effluent disposal and reuse

| | pH | BOD ₅ (mg l ⁻¹) | COD (mg l ⁻¹) | TSS (mg l ⁻¹) | Conductivity (μS cm ⁻¹) |
|--|---------|---|------------------------------|------------------------------|--|
| UWWTP effluent | 7.89 | 15 | 35 | 28 | 1453 |
| Italian disposal standards (D.Lgs 152/2006) | 5.5-9.5 | 25 | 125 | 35 | - |
| Italian reuse standards (D.M. 93/2006) | 6-9.5 | 20 | 100 | 10 | 3000 |

3.3 *Enterococcus* strains inactivation by solar photocatalysis

Solar simulated photocatalytic process showed a total inactivation of the two tested strains, after 60 minutes of exposure (Figure 1). In particular, a 98.627% inactivation efficiency of TRE strain was observed after 30 min irradiation time (average initial concentration as high as 1.8×10^7 CFU 100 ml⁻¹), and a 99.965% after 45 min irradiation. A similar trend was observed on the sensitive strain (average initial density 1.3×10^7 CFU 100 ml⁻¹): an inactivation of 95.106% was reached after 30 min of treatment and of 99.817% after 45 min. In parallel, solar disinfection tests did not show any significant inactivation of both strains under the same conditions of photocatalytic tests.

These results seem to support the idea that *Enterococcus* may be used as indicator of the occurrence of antibiotic resistant *Enterococcus* (in particular, TET resistant) after disinfection process. Specifically, if the disinfection process effectively inactivates *Enterococcus* indicators we can expect the simultaneous inactivation of the resistant strains.

Moreover, despite of different bacteria respond to stress conditions in different ways, it was observed that our results are in agreement with a previous work where the photocatalytic process was investigated in the inactivation of antibiotic resistant *E. coli* strain (Rizzo *et al.*, 2014b). In particular, TiO₂ photocatalytic tests under solar simulated radiation and the same operating conditions showed that total inactivation (~7 log unit) can be achieved after 60 min of irradiation. Xiong and Xu (2013), investigated the effect of a UVA/LED/TiO₂ system (TiO₂ particles coated reactor, 8000 μW cm⁻² light

intensity) on the inactivation of an antibiotic resistant *E. coli* strain (ATCC 700891) containing ampicillin and streptomycin resistance markers and they observed a 4.5 log unit inactivation (6×10^4 CFU ml⁻¹ initial bacterial density) in approximately 90 min of irradiation time. Tsai *et al.* (2010), investigated the effect of TiO₂ photocatalytic process on three clinical isolates antibiotic resistant bacteria, among which vancomycin-resistant *Enterococcus faecalis* (VRE). Differently from the present work, the authors operated the photocatalytic process under higher intensity irradiation (400-800 μW cm⁻²) and photocatalyst loadings (0.0625-0.125 g l⁻¹). According to our results, they did not observe any significant effect on VRE inactivation under light irradiation. When the catalyst was added (0.0625 g l⁻¹, 400 μW cm⁻²) inactivation drastically increased (10^3 - 10^5 CFU ml⁻¹ initial bacterial density), but total inactivation was not observed during the experiment (max irradiation time 90 min). Total inactivation was observed in 20-25 min when light irradiation intensity was increased up to 800 μW cm⁻² and photocatalyst loading was increased up to 0.125 g l⁻¹. The effect on enterococci was also investigated using a different AOP, namely solar photo-Fenton, to evaluate both inactivation rate and resistance percentage (ratio between resistant and total enterococci) (Michael *et al.*, 2012); the authors observed that total inactivation (initial density ~ 2.5-4.0 × 10² CFU ml⁻¹) took place after 180 min treatment under real solar radiation.

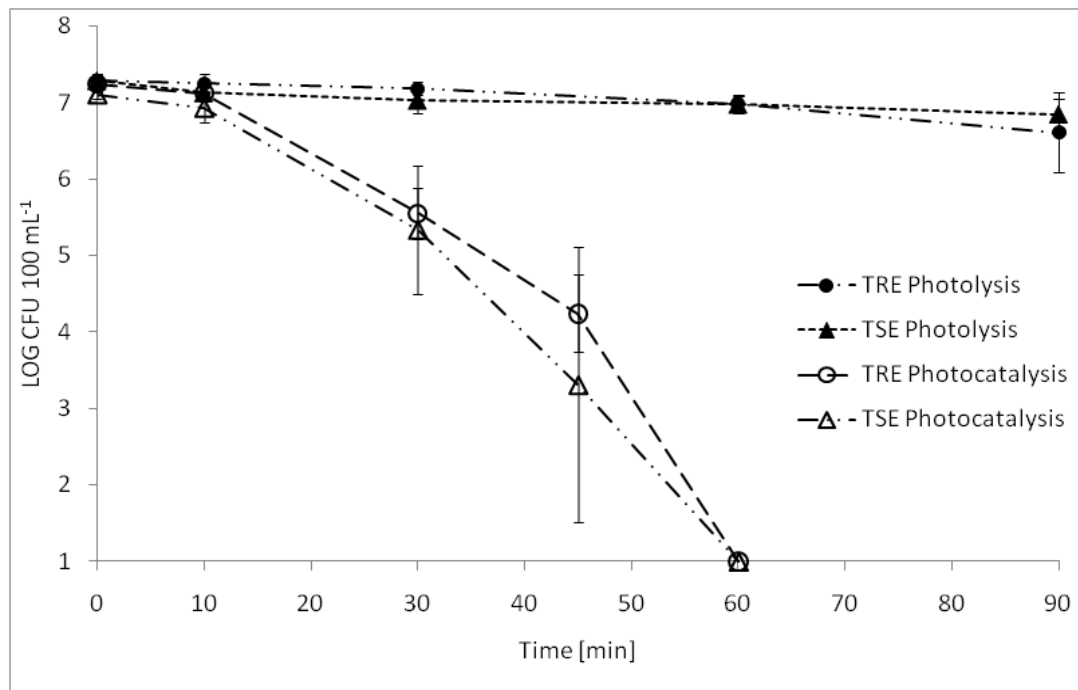


Figure 1. Inactivation rate of TRE and TSE strains under solar photolysis and photocatalytic disinfection (0.05 g TiO₂ l⁻¹).

3.4 Effect of solar photocatalysis on antibiotic resistance

The comparison of the average values of inhibition diameters for AMP, CIP, TET and VAN before disinfection treatment ($t = 0$) for TRE strain with the corresponding clinical breakpoints values for *Enterococcus* from EUCAST database (Table 1), shows that TRE strain is sensitive (S) to AMP, CIP and VAN antibiotics but resistant (R) to TET (Tables 3 and 4). The results of resistance test on the colonies that survived the photocatalytic treatment showed that disinfection process did not affect TRE strain resistance to AMP and CIP after 45 min irradiation (Table 3).

The photocatalytic treatment did not show any effect on TRE strain resistance to antibiotics tested (Table 3). Moreover, solar photolysis process did not show any effect on the inhibition diameter (no trend was observed) for all investigated antibiotics (Table 4).

Table 3. Resistance test results on TRE strain before (t=0) and after photocatalytic treatment: inhibition diameter (mm) and resistance class.

| t (min) | AMP | CIP | TET | VAN |
|---------|----------|----------|----------|----------|
| 0 | 24.3 (S) | 24.6 (S) | 10.0 (R) | 20.0 (S) |
| 10 | 23.8 (S) | 24.5 (S) | 10.0 (R) | 20.0 (S) |
| 30 | 24.0 (S) | 24.8 (S) | 10.0 (R) | 20.0 (S) |
| 45 | 31.5 (S) | 26.5 (S) | 10.0 (R) | 20.0 (S) |

Finally, both investigated processes did not affect resistance of TSE strain at all. The effect of photocatalytic process on an antibiotic resistant *E. coli* strain was investigated in a previous work (Rizzo *et al.*, 2014b). When the *E. coli* suspension was disinfected the resistance of the survived colonies to CIP and VAN decreased (inhibition diameter increased from 12.8 to 13.8 mm, and from 10.2 to 10.9 mm, respectively) as irradiation time was increased (from 0 to 30 min, respectively). In another work, the effect of solar photo-Fenton process on antibiotic resistance of enterococci was investigated in terms of resistance percentage (Michael *et al.*, 2012). The authors observed that resistance decrease in ofloxacin (initial resistance percentage ~20%) and trimethoprim (~10%) resistant enterococci took place after 180 min treatment under real solar radiation.

Table 4. Resistance test results on TRE strain before (t=0) and after photolytic treatment: inhibition diameter (mm) and resistance class.

| t (min) | AMP | CIP | TET | VAN |
|---------|----------|----------|----------|----------|
| 0 | 19.0 (S) | 23.0 (S) | 10.0 (R) | 20.0 (S) |
| 10 | 18.8 (S) | 22.5 (S) | 10.0 (R) | 20.3 (S) |
| 30 | 18.8 (S) | 22.5 (S) | 10.0 (R) | 20.0 (S) |
| 60 | 18.3 (S) | 22.3 (S) | 10.0 (R) | 20.0 (S) |
| 90 | 19.3 (S) | 22.5 (S) | 10.0 (R) | 20.5 (S) |

4. Conclusions

TiO₂ photocatalysis was found to be effective in the inactivation of TRE and TSE strains selected from the effluent of biological process of an UWTP. Remarkably, photocatalytic disinfection tests showed comparable inactivation rates for both strains. Therefore, we can expect that if *Enterococcus* strains are effectively inactivated by photocatalytic process, antibiotic resistant *Enterococcus* strains (at least some of them) can be inactivated too. Accordingly, *Enterococcus* may be used as indicators of photocatalytic process efficiency on antibiotic resistant *Enterococcus*.

Antibiotic resistance of surviving TRE colonies was not significantly affected by the photocatalytic process. This result supports the belief that antibiotic resistance can spread into the receiving water body when antibiotic resistant strains survive to disinfection process.

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