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Abstract: Recently, titanium dioxide (TiO₂) nanomaterials have gained increased attention because of their cost-effective, safe, stable, non-toxic, non-carcinogenic, photocatalytic, bactericidal, biomedical, industrial and waste-water treatment applications. The aim of the present work is the synthesis of electrospun TiO₂ nanofibers (NFs) in the presence of different amounts of air–argon mixtures using sol-gel and electrospinning approaches. The physicochemical properties of the synthesized NFs were examined by scanning and transmission electron microscopies (SEM and TEM) coupled with energy-dispersive X-ray spectroscopy (EDX), ultraviolet-visible spectroscopy and thermogravimetric analyzer (TGA). The antibacterial and antibiofilm activity of synthesized NFs against Gram-negative Pseudomonas aeruginosa and Gram-positive methicillin-resistant Staphylococcus aureus (MRSA) was investigated by determining their minimum bacteriostatic and bactericidal values. The topological and morphological alteration caused by TiO₂ NFs in bacterial cells was further analyzed by SEM. TiO₂ NFs that were calcined in a 25% air-75% argon mixture showed maximum antibacterial and antibiofilm activities. The minimum inhibitory concentration (MIC)/minimum bactericidal concentration (MBC) value of TiO₂ NFs against P. aeruginosa was 3 and 6 mg/mL and that for MRSA was 6 and 12 mg/mL, respectively. The MIC/MBC and SEM results show that TiO₂ NFs were more active against Gram-negative P. aeruginosa cells than Gram-positive S. aureus. The inhibition of biofilm formation by TiO₂ NFs was investigated quantitatively by tissue culture plate method using crystal violet staining.
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violet assay and it was found that TiO$_2$ NFs inhibited biofilm formation by MRSA and $P$. aeruginosa in a dose-dependent manner. TiO$_2$ NFs calcined in a 25% air-75% argon mixture exhibited maximum biofilm formation inhibition of 75.2% for MRSA and 72.3% for $P$. aeruginosa at 2 mg/mL, respectively. The antibacterial and antibiofilm results suggest that TiO$_2$ NFs can be used to coat various inanimate objects, in food packaging and in waste-water treatment and purification to prevent bacterial growth and biofilm formation.

**Keywords:** TiO$_2$ nanofibers; electrospinning; biofilm prevention and control; multidrug-resistant bacteria; biomedical application

1. Introduction

Titanium dioxide (TiO$_2$) is among the investigated photocatalytic nanomaterials and is used extensively in diverse applications and for diverse purposes [1]. TiO$_2$ nanomaterials are widely used in waste-water treatment and purification, air-pollutant decomposition, implantable devices, air-conditioning coatings, hydrophilic coatings, self-cleaning and self-disinfecting devices, pesticide degradation (e.g., herbicides, insecticides and fungicides) and in the production of hydrogen fuel [2,3]. TiO$_2$ is usually non-toxic, highly durable with a high refractive index, high absorption of light and a lower-cost production with antibacterial activity [4,5]. Because of its strong stability, TiO$_2$ materials can be applied easily on inanimate items, e.g., metal, glass and biomedical implants [5]. Recently, TiO$_2$ nanoparticles (NPs) have attracted increased interest in the scientific and industrial community because of their extensive applications in biological and pharmaceutical areas, purification of environmental sources, electronic system, solar energy cells, photocatalysts, photo-electrodes and gas sensors. TiO$_2$ NPs are proven to be employed in food technology, drugs, cosmetics, paint pigment, ointments and toothpaste [6,7]. Because of their cost-effective, safe, stable, non-toxic, non-carcinogenic, photo-induced super-hydrophobicity and antifogging properties, TiO$_2$ NPs have been used to kill bacteria, remove toxic and harmful organic elements from water and air and for self-sterilize glass surfaces [8–11].

However, it is difficult to separate TiO$_2$ NPs after a photochemical reaction, which limits their practical applications [12]. TiO$_2$ NFs aggregate easily in solution, which reduces their photocatalytic efficacy because of the decreased surface area. These limitations can be overcome by preparing TiO$_2$ nanofibers (NFs) using simple, rapid and cost-effective electrospinning (ES) methods [13–18]. TiO$_2$ NFs have gained increased attention because of their mesoporous structure [19], stability in solution, little or no aggregation, high surface to volume ratio that enhances photocatalytic reactions and their ease in separation and collection from solution after photochemical reactions [20,21]. However, the photocatalytic efficacy of TiO$_2$ NFs is comparatively low and is effective only under ultraviolet (UV) light because of their relatively large band-gap energy and low-ordered crystalline structure [22]. An exceptional feature of TiO$_2$ nanoparticles (NPs) is their photocatalytic activity that enhances the bacterial killing when exposed to UV light [7,23]. TiO$_2$ NPs tend to exist in three principal forms, namely brookite, rutile and anatase, and it has been reported that the anatase form has a high photocatalytic and antibacterial activity [23–26]. A major biomedical application of TiO$_2$ NPs is to prevent biofilm formation on medical devices that is related to infections and sepsis [3,27,28]. Several researchers have focused on the antibacterial and antibiofilm activities of TiO$_2$ NPs under UV light against standard bacterial strains, e.g., ATCC, MTCC and NCIM. However, limited work has been published on the antibacterial and antibiofilm activities of TiO$_2$ NFs without application of UV light against drug resistant isolates. The objective of present investigation is to explore the antibacterial and antibiofilm efficacies of TiO$_2$ NFs in dark against two major human pathogenic drug resistant bacteria i.e., Gram-positive *methicillin-resistant Staphylococcus aureus* (MRSA) and Gram-negative *Pseudomonas aeruginosa* by using different methods.
2. Experimental Methodology

2.1. Electrospinning and Heating Protocol

Both the sol-gel and electrospinning approaches were used to synthesize electrospun TiO\(_2\) NFs. Briefly, Titanium isopropoxide (IV), acetic acid and ethanol were mixed and stirred with respect to volume ratio of 3:1:3. After that, 12\% by weight of polyvinylpyrrolidone (PVP) was dissolved in the obtained TiO\(_2\) solution. This mixed TiO\(_2\)/PVP sol-gel was then placed within a plastic syringe for electrospinning experiment. Additional details are provided in a preliminary study [15]. Thermal gravimetric analysis (TGA) and differential scanning calorimetry (DSC) for the non-isothermal heating of electrospun TiO\(_2\) NFs were performed on a Mettler Teledo thermal gravimetric analyzer TGA/DSC. The samples were heated from ambient temperature to 900 °C at a rate of 10 °C/min with an argon protective gas of 20 mL/min in various mixtures of air and argon. The thermal experiments were carried out by utilizing alumina crucibles that were charged with 25 mg of sample and in mixtures of 50\% air-50\% argon, 25\% air-75\% argon and 100\% argon. It is worth noting that the argon shielding gas is included in the relative percentage of air to argon gas. For safety reasons, samples that were contacted with 100\% air were heated in an oven under the same conditions [29].

2.2. Characterization of Electrospun TiO\(_2\) NFs

The morphological and structural properties of as-prepared NFs were characterized by SEM (FEI Inspect S50) and TEM (FEI Morgagni 268). The elemental composition was determined by energy-dispersive spectroscopy (EDX). A strong correlation can be established from the initial microstructure images. The TiO\(_2\) grains were structured as microspheres and a complete description of the microstructure is provided. The microstructure relates to monitoring by three-dimensional imaging of the evolution of internal porosity as a function of annealing temperature. A Jasco V-670 UV–visible diffuse reflectance spectrophotometer (DRS) under a wavelength ranging between 200 and 750 nm was used to estimate the band gap energy (\(E_g\)) of various TiO\(_2\) NFs.

The values of band gap energy (\(E_g\)) were calculated from the absorption spectra versus wavelength using the following expression:

\[
E_g = \frac{hC}{\lambda_0}
\]  

In this expression, \(h\) is Planck’s constant (6.626 \times 10^{-34} \text{ J.s}) and \(C\) is the speed of light (3 \times 10^8 \text{ m/s}). \(\lambda_0\) (expressed in nm) is the cut off wavelength obtained from the absorption spectra [30]. Accordingly, \(\lambda_0\) denotes the absorption edge wavelength, obtained from the offset wavelength derived and extrapolated from the low energy absorption band.

2.3. Evaluation of Antibacterial Activity of Electrospun TiO\(_2\) NFs

2.3.1. Bacterial Culture

The laboratory strain of Gram-negative \textit{Pseudomonas aeruginosa} PAO1 and Gram-positive methicillin resistant \textit{Staphylococcus aureus} (MRSA) ATCC 33591 used in this study was obtained from Molecular Microbiology Laboratory, Institute for Research and Medical Consultations, Imam Abdulrahman Bin Faisal University, Dammam, Saudi Arabia. The bacterial strains preserved in glycerol cultures (~80 °C) were cultivated on Tryptic soy broth (TSB) at 37 °C in a shaker incubator before being used for microbial studies.

2.3.2. Investigation of Minimum Inhibitory and Minimum Bactericidal Concentration (MIC/MBC) Values of Electrospun TiO\(_2\) NFs

The MIC values of TiO\(_2\) NFs against \textit{P. aeruginosa} and MRSA was estimated by serial two-fold dilutions of TiO\(_2\) NFs from 32 to 1 mg/mL as described previously [31,32]. The determination of MBC values was also investigated as method described in previous studies [32,33].
2.4. Effect of TiO$_2$ NFs on Biofilm Formation

The antibiofilm potential of TiO$_2$ NFs against *P. aeruginosa* and MRSA biofilm was examined quantitatively in a sterilized 96-well polystyrene (flat bottom) microtiter tissue culture plate using crystal violet assay as described in our previous study [31,33].

2.5. Effect of TiO$_2$ NFs on the Morphology of *P. aeruginosa* and MRSA: SEM Analysis

Further, the effects of TiO$_2$ NFs on the morphological features of *P. aeruginosa* and *S. aureus* cells were analyzed by SEM. In Brief, $\sim$10$^6$ CFU/mL of *P. aeruginosa* and *S. aureus* cells treated with 1 mg/mL of TiO$_2$ NFs for 18 h were incubated at 37 °C [33,34]. After incubation, the treated and untreated samples were centrifuged at 10,000 rpm for 15 min. The obtained pellets were washed with PBS (1×) three times and fixed with primary fixative (i.e., 2.5% glutaraldehyde) for 6 h at 4 °C and then further fixed with secondary fixative (i.e., 1% osmium tetroxide) for 1 h. After fixation, the samples were dehydrated by a series of ethanol [34,35]. The cells were then fixed on the aluminum stubs, dried in a desiccator and coated with gold. Finally, the treated and untreated samples were examined by SEM.

3. Results and Discussion

3.1. Effects of the Calcining Atmosphere on TiO$_2$ Colour

Figure 1 presents a gradual color change from white to dark grey after heat treatment in 100% air and in different mediums of air–argon compositions up to 100% argon medium. This change is likely because of oxygen vacancy defects. The change and the intensification of the color are mainly a result of defects associated with oxygen vacancies that rise from an increase in argon content [36].

![Figure 1](Image)

Figure 1. Color changes in electrospun titanium oxide (TiO$_2$) nanofibers (NFs) in argon-air mixtures.

3.2. Microstructure Analysis of the Prepared NFs

Figures 2 and 3 show the typical SEM and TEM micrographs of the as-spun TiO$_2$ and calcined NFs. The electrospinning process could produce good quality TiO$_2$ NFs, possibly without nodes and defects. The diameter of the as-spun fibers varied between 80 and 600 nm, whereas the estimated average thickness was ~400 nm (Figure 2a). Upon annealing in different mediums of air/argon (100%-0%, 50%-50%, 25%-75%, and 0%-100%), the fibers shrank, and their morphology changed slightly from smooth to rough. This figure also shows the presence of a heterogeneous matrix made up of agglomerated grains for the initial microstructure and leads to faster granular growth. The fibers size was between 50 and 300 nm (Figure 2b–e). Several thin-fibers of about 50 nm were perceived in specimens annealed under 25-75% air-argon. The quality and shape of fiber mats were preserved after calcination as clarified by the TEM images (Figure 3b–e) unlike the electrospun fibers that are often
composed of oxide nanoparticles (Figure 3a) [37]. The as-spun fibers showed organic species, whereas the annealed fibers exhibited a solid morphology with high-quality individual particles in the range of 100 nm. The annealing of TiO$_2$ NFs at 900 °C in 50%-50% air-argon led to pure TiO$_2$ fibers formation, which was proven by EDX and TGA characterization techniques. In Figure 4, the EDX spectrum illustrates high-intensity O and Ti peaks and a small Pt peak from the platinum coating on the TiO$_2$ NFs heated in 50-50% air-argon, which is mainly similar to those observed in specimens annealed in 100% air, 25% air-75% argon, and 100% argon. Figure 5 shows the TGA result for samples heated under 50-50% air-argon medium. The PVP polymer and organic material are completely removed from the electrospun TiO$_2$ NFs at ~450 °C, and ~100 °C, respectively.

Figure 2. Scanning electron microscopy (SEM) of TiO$_2$ NFs calcined in different air and argon mixture. (a) As-spun TiO$_2$, (b) 100% Air, (c) 50% Air and 50% Argon, (d) 25% Air and 75% Argon and (e) 100% Argon.
Figure 3. Transmission electron microscopy (TEM) of TiO\textsubscript{2} NFs calcined in different air and argon mixture. (a) As-spun TiO\textsubscript{2}, (b) 100% Air, (c) 50% Air and 50% Argon, (d) 25% Air and 75% Argon and (e) 100% Argon.

Figure 4. An energy-dispersive X-ray spectroscopy (EDX) spectrum of electrospun TiO\textsubscript{2} NFs prepared in 50% air-50% argon mixture.
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3.3. Wide-Band Gap Analysis of Calcined Electrospun TiO$_2$ NFs

Figure 6 shows the UV-vis DRS spectra of as-electrospun TiO$_2$ NFs calcinated in air-argon media at 900 °C and cooling to ambient temperature. Table 1 shows the values of band-gaps at room temperature for various TiO$_2$ NFs. The band-gap value reduced from 3.33 eV for as-spun and non-calcinated samples to about 3.09 eV for the ones calcinated in 100% air. Under various air-argon environments, the value of $E_g$ decreased from about 3.09 to 2.18 eV with an increase in argon content. A previous study on similar specimens revealed that the growth of vacancies was minimal and the reduction of $E_g$ value was ascribed to the increase in crystallinity [38]. The measured difference agrees with that weighted according to the concentration of pure anatase and rutile phases [38–40]. Alterations in levels and phase mixing and gradual development of oxygen vacancies are two factors that can reduce the band-gap energy with argon introduction. The measured energy gap was 2.18 eV for sample heated in 100% argon and for the phase composition for which the difference according to the concentration would be 3.05 eV. The difference of 0.87 eV is assigned to the development of oxygen vacancies and allows a greater density of charge carriers. The development of oxygen vacancies leads to the creation of Ti$^{3+}$ centers or unpaired electrons that generate vacant states under the conduction band [41,42]. The development of oxygen vacancies for different argon concentrations has been previously discussed [38]. When the specimen is annealed in argon, oxygen disappears and the non-stoichiometric anatase (TiO$_{2-x}$) forms [43]. The formation of oxygen vacancy defects in titanium oxide is induced from the occurrence of new localized states of oxygen vacancies between the conduction and valence bands. The excitation of electrons from the valence band to the vacant oxygen states can be done in visible light. With rising argon amount, the effective $E_g$ moves thoroughly to the red region, the specimen is being active under visible light and thus the $E_g$ is reduced. So, the mutual effects of the formation of oxygen vacancies and crystallinity treatment have prolonged the excitation of light of electrospun TiO$_2$ NFs from ultraviolet to visible light range without the need of chemical doping.

Table 1. Band gap energies for as-electrospun TiO$_2$ nanofibers (non-calcinated), and TiO$_2$ NFs obtained after calcination at 900 °C in various air-argon media.

| Calcination Conditions       | $E_g$ (eV) |
|-----------------------------|------------|
| As-electrospun              | 3.33       |
| 100% Air                    | 3.09       |
| 50% Air and 50% Argon       | 2.94       |
| 25% Air and 75% Argon       | 2.91       |
| 100% Argon                  | 2.18       |
3.4. Antibacterial and Antibiofilm Activity of TiO₂ NFs

3.4.1. MIC and MBC

The microbiocidal activities of TiO₂ photocatalysis were reported for the first time by Matsunaga and co-workers in 1985 [44]. They investigated the killing of bacteria and yeast cells in water by employing TiO₂-Pt photocatalysts in near-ultraviolet radiation. They reported that the inhibition of respiratory activity was the mechanism for cell death.

In this research work, the antibacterial property (MIC/MBC) of TiO₂ NFs calcined with different ratios of air-argon mixtures (i.e., 100% air, 50% air-50% argon, 25% air-75% argon, and 100% argon) has been investigated against P. aeruginosa and MRSA (Supplementary Figure S1). The MIC/MBC values of TiO₂ NFs heated with different ratios of air-argon mixtures against P. aeruginosa and MRSA are presented in Table 2. TiO₂ NFs heated in the presence of 25% air-75% argon showed a maximum antibacterial activity and MIC/MBC values against P. aeruginosa were 3 and 6 mg/mL and for MRSA it was 6 and 12 mg/mL, respectively (Table 2). Based on the MIC and MBC results, it was observed that Gram-negative P. aeruginosa was more susceptible to TiO₂ NFs than Gram-positive MRSA. These results agree with results from previous studies [45,46], and may occur owing to differences in their cell wall structures and to bacterial strain growth rate [45–47]. Pigeot-Rémy and co-workers [48] investigated the effects of TiO₂ particles against E. coli K-12 in the dark and reported that the attachment of NPs to bacterial surfaces causes membrane damage and perturbation, which may increase the permeability of the outer cell membrane and the resultant damage to the envelope of bacterial cells leads to bacterial cells death.

| Electrospun TiO₂ Nanofibers Code | Calcination Conditions | Methicillin Resistant S. aureus | P. aeruginosa |
|----------------------------------|------------------------|-------------------------------|--------------|
| (a)                              | 100% Air               | 7                             | 7            |
| (b)                              | 50% Air and 50% Argon  | 7                             | 7            |
| (c)                              | 25% Air and 75% Argon  | 6                             | 3            |
| (d)                              | 100% Argon             | >16                           | >16          |

Table 2. Minimum inhibitory concentration (MIC) and minimum bactericidal concentration (MBC) (mg/mL) values of tested electrospun TiO₂ nanofibers against Methicillin resistant S. aureus and P. aeruginosa.
3.4.2. Effects of Electrospun TiO₂ NFs on the Morphology of Bacterial Cells

Morphological alterations in Gram-negative P. aeruginosa (Figure 7) and Gram-positive MRSA (Figure 8) after exposure to TiO₂ NFs were further examined by SEM. The untreated P. aeruginosa had a normal, rod-shaped structure and regular, smooth and intact cell surface (Figure 7). However, the morphology of P. aeruginosa cells was altered considerably, and cells were damaged to different extents after treatment with TiO₂ NFs. After 18 h of treatment, the cell envelope and cell wall were rough, irregular, abnormal in form and main damage was categorized by the creation of “pits” and depressions that probably lead to a loss of bacterial cell membrane integrity (Figure 7). Similarly, the untreated Gram-positive MRSA was normal with smooth and regular cell surfaces (Figure 8). However, MRSA cells treated with TiO₂ NFs exhibited noticeable alterations and damage and the clusters of NFs were linked and anchored on the surface of bacterial cells (Figure 8). Irregularities, shallows and depressions on the cell envelopes and cell walls of certain MRSA cells suggest that bacterial damage occurred (Figure 8). SEM analysis showed that TiO₂ NFs were more effective against P. aeruginosa bacterial cells in comparison with MRSA and were severely injured compared with Gram-positive MRSA. The obtained results may be due to morphological dissimilarities in the cell walls of bacteria. Gram-negative bacterial cells display thin layers of peptidoglycan that facilitate the mobility of metal-ion NPs within cells and facilitate the interaction among NPs and walls of bacterial cells. Gram-negative bacteria exhibit a negative charge due to their high content of lipopolysaccharides. This negative charge attracts and interacts with positive metal ions, which may lead to the NP penetration, intracellular damages and protein and DNA destruction [46]. It was suggested that the interaction of TiO₂ NPs with bacterial cells in the dark caused bacterial membrane integrity destruction, especially of lipopolysaccharides [48]. TiO₂ NPs form pores in bacterial cell walls and membranes, which increases the permeability and leads to cell death [10]. However, other published work has shown that the contact among metal oxides and bacterial cells provokes oxidation and formation of reactive oxygen groups including O₂•⁻, •OH, and H₂O₂. These free radicals attack bacteria cell walls and alter the membrane integrity and permeability, which leads to bacterial cell death [48–51]. It has been reported that the destruction of cell envelope by incorporation of TiO₂ NPs inside the cells damages bacterial DNA and RNA, which could provoke cell death [48]. The antimicrobial activity of TiO₂ in the absence of photoactivation has been also reported. Nakano and co-worker [51] stated that TiO₂ deactivates bacterial DNA and enzymes via coordination of electron-donor groups, like hydroxyls, indoles, carbohydrates, amides, and thiols in the absence of light. Pit formation in bacterial cell walls and envelopes that enhanced the permeability lead to bacterial cell death [51,52]. It has been reported that there is proportional relationship between the light and the antimicrobial activity of TiO₂. Senarathna et al [53] and Lee et al [54] reported that the presence of sunlight enhanced the antimicrobial activity of TiO₂ against S. aureus might be due to generation of free radicals [53,54].
Figure 7. Effect of electrospun TiO$_2$ NFs on the morphological aspects of $P. aeruginosa$ as examined by scanning electron microscopy: (A) control without any treatment and treated with TiO$_2$ calcined in (B) 100% Air, (C) 50% Air and 50% Argon, (D) 25% Air and 75% Argon; and (E) 100% Argon.

3.4.3. Inhibition of Biofilm Formation by TiO$_2$ NFs

The antibiofilm potential of TiO$_2$ NFs heated under different air-argon environments was evaluated at various amounts of 0.25, 0.5, 1.0 and 2.0 mg/mL against MRSA and $P. aeruginosa$ biofilms using crystal violet microtiter assays in a 96-well flat-bottom polystyrene plate at OD$_{595}$ nm. Plots in Figure 9A,B show that TiO$_2$ NFs inhibit the biofilms formation by MRSA and $P. aeruginosa$ in a dose-dependent manner. It was reported that a rise in TiO$_2$ concentration provoked a reduction in the cultivability of bacteria [48]. As shown in Figure 9A,B, TiO$_2$ NFs heated in a 25% air-75% argon mixture exhibited the highest biofilm inhibition of about 75.2% for MRSA and 72.3% for $P. aeruginosa$, respectively at 2 mg/mL of TiO$_2$ NFs. These results agree with those reported in previous studies [55,56]. In a previous study, epoxy/Ag-TiO$_2$ nanocomposites were found to inhibit biofilm creation of $S. aureus$ ATCC 6538 and $E. coli$ K-12 by 67% and 77%, respectively [56].
Figure 8. Effect of electrospun TiO$_2$ NFs on the morphological aspect of *S. aureus* as examined by scanning electron microscopy: (A) control without any treatment and treated with TiO$_2$ NFs calcined in (B) 100% Air; (C) 50% Air and 50% Argon; (D) 25% Air and 75% Argon, and (E) 100% Argon.
4. Conclusions

This study focuses on the heat treatment of TiO$_2$ NFs to develop photoactive titanium photocatalysis in the visible spectrum and to evaluate their antibacterial and antibiofilm potential against Gram-negative bacteria *P. aeruginosa* and Gram-positive MRSA. The $E_g$ value was 3.09 eV for specimens heated in 100% air and 2.18 eV for the ones heated in 100% argon. The value of $E_g$ decreased systematically with rising argon amount in the various air-argon mixtures. The increase in the amount of argon brings the state under the TiO$_2$ conduction band. TiO$_2$ NFs calcined in a 25% air-75% argon environment showed maximum antibacterial and antibiofilm activities. The MIC/MBC and SEM results show that TiO$_2$ NFs were more operative against Gram-negative *P. aeruginosa* than Gram-positive *S. aureus*. The inhibition of biofilm formation by TiO$_2$ NFs shows that TiO$_2$ NFs inhibit the biofilms formation by MRSA and *P. aeruginosa* in a dose-dependent manner. From the obtained data on antibacterial antibiofilm analysis, it has been concluded and suggested that TiO$_2$ NFs can be used in hydrophilic coatings, coating of various inanimate object surfaces, such as metals, glass, medical devices and equipment to prevent biofilm formation on medical devices or medical device-related infections and sepsis, and also can be applied in food packaging, wastewater treatment and purification, self-cleaning and self-disinfecting, killing of bacteria and the removal of toxic and damaging organic compounds from water and air.
Supplementary Materials: The following are available online at http://www.mdpi.com/2079-6382/9/5/572/s1, Figure S1: represents MHA plates showing MBC values of tested electrospun TiO$_2$ NFs in various air–argon environments.

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References
1. Fujishima, A.; Rao, T.N.; Tryk, D.A. Titanium dioxide photocatalysis. J. Photochem. Photobiol. C: Photochem. Rev. 2000, 1, 1–21. [CrossRef]
2. Mills, A.; Le Hunte, S. An overview of semiconductor photocatalysis. J. Photochem. Photobiol. A: Chem. 1997, 108, 1–35. [CrossRef]
3. Gupta, S.M.; Tripathi, M. A review of TiO$_2$ nanoparticles. Chin. Sci. Bull. 2011, 56, 1639–1657. [CrossRef]
4. Kamat, P. Photophysical, Photochemical and Photocatalytic Aspects of Metal Nanoparticles. J. Phys. Chem. B 2002, 106, 7729–7744. [CrossRef]
5. Mantravadi, H.B. Effectivity of Titanium Oxide Based Nano Particles on E. coli from Clinical Samples. J. Clin. Diagn. Res. 2017, 11, DC37–DC40. [CrossRef]
6. Awati, P.; Awate, S.; Shah, P.; Ramaswamy, V. Photocatalytic decomposition of methylene blue using nanocrystalline anatase titania prepared by ultrasonic technique. Catal. Commun. 2003, 4, 393–400. [CrossRef]
7. Othman, S.; Salam, N.R.A.; Zainal, N.; Basha, R.K.; Talib, R. Antimicrobial Activity of TiO2Nanoparticle-Coated Film for Potential Food Packaging Applications. Int. J. Photoenergy 2014, 2014, 1–6. [CrossRef]
8. Hashimoto, K.; Irie, H.; Fujishima, A. TiO2Photocatalysis: A Historical Overview and Future Prospects. Jpn. J. Appl. Phys. 2005, 44, 8269–8285. [CrossRef]
9. Schmidt, H.; Naumann, M.; Muller, T.; Akarsu, M. Application of spray techniques for new photocatalytic gradient coatings on plastics. Thin Solid Film. 2006, 502, 132–137. [CrossRef]
10. E Teo, W.; Ramakrishna, S. A review on electrospinning design and nanofibre assemblies. Nanotechnology 2006, 17, R98–R106. [CrossRef] [PubMed]
11. Albetran, H.; Low, I.M. Crystallization kinetics and phase transformations in aluminium ion-implanted electrospun TiO$_2$ nanofibers. Appl. Phys. A 2014, 116, 161–169. [CrossRef]
12. Albetran, H.; O’Connor, B.H.; Prida, V.M.; Low, I.M. Effect of vanadium ion implantation on the crystallization kinetics and phase transformation of electrospun TiO$_2$ nanofibers. Appl. Phys. A 2015, 120, 623–634. [CrossRef]
13. Albetran, H.; Low, I. Parameters controlling the crystallization kinetics of nanostructured TiO$_2$—An overview. Mater. Today Proc. 2019, 16, 25–35. [CrossRef]
19. Zhang, X.-W.; Xu, S.; Han, G. Fabrication and photocatalytic activity of TiO2 nanofiber membrane. Mater. Lett. 2009, 63, 1761–1763. [CrossRef]
20. Chandrasekar, R.; Zhang, L.; Howe, J.Y.; Hedin, N.E.; Zhang, Y.; Fong, H. Fabrication and characterization of electrospun titania nanofibers. J. Mater. Sci. 2009, 44, 1198–1205. [CrossRef]
21. Li, Q.; Satur, D.J.G.; Kim, H.; Kim, H.G. Preparation of sol–gel modified electrospun TiO2 nanofibers for improved photocatalytic decomposition of ethylene. Mater. Lett. 2012, 76, 169–172. [CrossRef]
22. Xu, S.; Li, A.; Poirier, G.; Yao, N. In Situ Mechanical and Electrical Characterization of Individual TiO2 Nanofibers Using a Nanomanipulator System. Scanning 2012, 34, 341–346. [CrossRef] [PubMed]
23. Ahmad, R.; Sardar, M. TiO2 nanoparticles as an antibacterial agents against E. coli. Int. J. Innov. Res. Sci. Eng. Technol. 2013, 2, 3569–3574.
24. Albetran, H.; Low, I. Crystallization kinetics study of In-doped and (In-Cr) co-doped TiO2 nanopowders using in-situ high-temperature synchrotron radiation diffraction. Arab. J. Chem. 2020, 13, 3946–3956. [CrossRef]
25. Albetran, H.; Vega, V.; Prida, V.M.; Low, I.-M. Dynamic Diffraction Studies on the Crystallization, Phase Transformation, and Activation Energies in Anodized Titania Nanotubes. Nanomaterials 2018, 8, 122. [CrossRef]
26. Albetran, H.; Low, I.-M. Effect of indium ion implantation on crystallization kinetics and phase transformation of anodized titania nanotubes using in-situ high-temperature radiation diffraction. J. Mater. Res. 2016, 31, 1588–1595. [CrossRef]
27. Ravishankar, R.V.; Jamuna, B.A. Nanoparticles and their potential application as antimicrobials. In Science against Microbial Pathogens: Communicating Current Research and Technological Advances; Mendez-Vilas, A., Ed.; University of Mysore: Mysore, India, 2011; pp. 197–209.
28. Arora, H.; Doty, C.; Yuan, Y.; Boyle, J.; Petras, K.; Rabatic, B.; Paunesku, T.; Woloschak, G.E. Titanium Dioxide Nanoparticles and their potential application as antimicrobials. In Nanomaterials for Life Sciences; Kumar, C.S.S.R., Ed.; Wiley-VCH Verlag GmbH & Co KGaA: Weinheim, Germany, 2010; pp. 1–42.
29. Albetran, H.; O’Connor, B.; Low, I.-M. Effect of calcination on band gaps for electrospun titania nanofibers heated in air-argon mixtures. Mater. Des. 2016, 92, 480–485. [CrossRef]
30. Bhadwal, A.S.; Tripathi, R.; Gupta, R.K.; Kumar, N.; Singh, R.P.; Shrivastav, A. Biogenic synthesis and photocatalytic activity of CdS nanoparticles. RSC Adv. 2014, 4, 9484. [CrossRef]
31. Balasamy, R.J.; Ravinayagam, V.; Alomari, M.; Ansari, M.A.; Almofty, S.A.; Rehman, S.; Dafalla, H.; Marimuthu, P.R.; Akhtar, S.; Hamad, M. Cisplatin delivery, anticancer and antibacterial properties of Fe/SBA-16/ZIF-8 nanocomposite. RSC Adv. 2019, 9, 42395–42408. [CrossRef]
32. Sultan, A.; Khan, H.M.; Malik, A.; Ansari, A.; Azam, A.; Perween, N. Antibacterial activity of ZnO nanoparticles against ESBL and Amp-C producing gram negative isolates from superficial wound infections. Int. J. Curr. Microbiol. App. Sci. 2015, 1, 38–47.
33. Baig, U.; Ansari, M.A.; Gondal, M.; Akhtard, S.; Khan, F.A.; Falathaf, W.S. Single step production of high-purity copper oxide-titania nanocomposites and their effective antibacterial and anti-biofilm activity against drug-resistant bacteria. Mater. Sci. Eng. C 2020, 113, 110992. [CrossRef] [PubMed]
34. Ali, F.A.A.; Alam, J.; Shukla, A.K.; Alhoshan, M.; Ansari, M.A.; Al-Masry, W.A.; Rehman, S.; Alam, M. Evaluation of antibacterial and antifouling properties of silver-loaded GO polysulfone nanocomposite membrane against Escherichia coli, Staphylococcus aureus, and BSA protein. React. Funct. Polym. 2019, 140, 136–147. [CrossRef]
35. Shukla, A.K.; Alam, J.; Ansari, M.A.; Alhoshan, M.; Alam, M.; Kaushik, A. Selective ion removal and antibacterial activity of silver-doped multi-walled carbon nanotube/polyphenylsulfone nanocomposite membranes. Mater. Chem. Phys. 2019, 233, 102–112. [CrossRef]
36. Gamboa, J.A.; Pasquevich, D.M. Effect of Chlorine Atmosphere on the Anatase-Rutile Transformation. J. Am. Ceram. Soc. 1992, 75, 2934–2938. [CrossRef]
37. Szilágyi, I.M.; Santala, E.; Heikkilä, M.; Pore, V.; Kemell, M.; Nikitin, T.; Teucher, G.; Firkala, T.; Khriachtchev, L.; Rasanen, M.; et al. Photocatalytic properties of WO3/TiO2 core/shell nanofibers prepared by electrospinning and atomic layer deposition. Chem. Vap. Depos. 2013, 19, 149–155. [CrossRef]
38. Albetran, H.; O’Connor, B.H.; Low, I.-M. Activation energies for phase transformations in electrospun titania nanofibers: Comparing the influence of argon and air atmospheres. Appl. Phys. A 2016, 122. [CrossRef]
39. Scanlon, D.O.; Dunnill, C.W.; Buckeridge, J.; Shevlin, S.A.; Logsdail, A.J.; Woodley, S.M.; Catlow, C.R.A.; Powell, M.J.; Palgrave, R.G.; Parkin, I.P.; et al. Band alignment of rutile and anatase TiO2. Nat. Mater. 2013, 12, 798–801. [CrossRef]  
40. Natoli, A.; Cabeza, A.; Torre, Ángeles, G.; Aranda, M.A.G.; Santacruz, I.; Cabeza, A. Colloidal Processing of Macroporous TiO2 Materials for Photocatalytic Water Treatment. J. Am. Ceram. Soc. 2011, 95, 502–508. [CrossRef]  
41. Nakamura, I.; Negishi, N.; Kutsuna, S.; Ihara, T.; Sugihara, S.; Takeuchi, K. Role of oxygen vacancy in the plasma-treated TiO2 photocatalyst with visible light activity for NO removal. J. Mol. Catal. A: Chem. 2000, 161, 205–212. [CrossRef]  
42. Seo, H.; Baker, L.R.; Hervier, A.; Kim, J.; Whitten, J.L.; Somorjai, G.A. Generation of Highly n-Type Titanium Oxide Using Plasma Fluorine Insertion. Nano Lett. 2011, 11, 798–801. [CrossRef]  
43. Andersson, S.; Collén, B.; Kuylenstierna, U.; Magnéli, A.; Pestmalis, H.; Åsbrink, S. Phase Analysis Studies on the Titanium-Oxygen System. Acta Chem. Scand. 1957, 11, 1641–1652. [CrossRef]  
44. Matsunaga, T.; Tomoda, R.; Nakajima, T.; Wake, H. Photoelectrochemical sterilization of microbial cells by semiconductor powders. FEMS Microbiol. Lett. 1985, 29, 211–214. [CrossRef]  
45. Nagalakshmi, M.; Karthikeyan, C.; Anusuya, N.; Brundha, C.; Basu, M.J.; Karuppuchamy, S. Synthesis of TiO2 nanofiber for photocatalytic and antibacterial applications. J. Mater. Sci. Mater. Electron. 2017, 28, 15915–15920. [CrossRef]  
46. Azizi-Lalabadi, M.; Ehsani, A.; Divband, B.; Alizadeh-Sani, M. Antimicrobial activity of Titanium dioxide and Zinc oxide nanoparticles supported in 4A zeolite and evaluation the morphological characteristic. Sci. Rep. 2019, 9, 1–10. [CrossRef] [PubMed]  
47. Vardanyan, Z.; Gevorkyan, V.; Ananyan, M.; Vardapetyan, H.; Trchounian, A. Effects of various heavy metal nanoparticles on Enterococcus hirae and Escherichia coli growth and proton-coupled membrane transport. J. Nanobiotechnol. 2015, 13, 1–9. [CrossRef]  
48. Pigeot-Remy, S.; Simonet, F.; Errazuriz-Cerda, E.; Lazzaroni, J.; Atlan, D.; Guillard, C. Photocatalysis and disinfection of water: Identification of potential bacterial targets. Appl. Catal. B: Environ. 2011, 104, 390–398. [CrossRef]  
49. Ranjan, S.; Ramalingam, C. Titanium dioxide nanoparticles induce bacterial membrane rupture by reactive oxygen species generation. Environ. Chem. Lett. 2016, 14, 487–494. [CrossRef]  
50. Wang, D.; Zhao, L.; Ma, H.; Zhang, H.; Guo, L.-H. Quantitative Analysis of Reactive Oxygen Species Photogenerated on Metal Oxide Nanoparticles and Their Bacteria Toxicity: The Role of Superoxide Radicals. Environ. Sci. Technol. 2017, 51, 10137–10145. [CrossRef]  
51. Chakra, C.S.; Mateti, S. Structural, Antimicrobial and Electrochemical Properties of Cu/TiO2 Nanocomposites. J. Nanosci. Technol. 2018, 4, 331–334. [CrossRef]  
52. Nakano, R.; Hara, M.; Ishiguro, H.; Yao, Y.; Ochiai, T.; Nakata, K.; Murakami, T.; Kajioka, J.; Sunada, K.; Hashimoto, K.; et al. Broad Spectrum Microbicidal Activity of Photocatalysis by TiO2. Catalysts 2013, 3, 310–323. [CrossRef]  
53. Senarathna, U.L.N.H.; Fernando, N.; Gunasekara, C.; Weerasekera, M.M.; Hewageegana, H.G.S.P.; Arachchi, N.D.; Siriwardena, H.D.; Jayaweera, P.M. Enhanced antibacterial activity of TiO2 nanoparticle surface modified with Garcinia zeylanica extract. Chem. Cent. J. 2017, 11, 7. [CrossRef] [PubMed]  
54. Lee, W.S.; Park, Y.-S.; Cho, Y.-K. Significantly enhanced antibacterial activity of TiO2 nanofibers with hierarchical nanostructures and controlled crystallinity. Analyst 2015, 140, 616–622. [CrossRef] [PubMed]  
55. Alavi, M.; Karimi, N.; Valadbeigi, T.; Valadbeigi, T. Antibacterial, Antifogging, Antiquorum Sensing, Antimottility, and Antioxidant Activities of Green Fabricated Ag, Cu, TiO2, ZnO, and Fe3O4 NPs via Protoparmeliopsis muralis Lichen Aqueous Extract against Multi-Drug-Resistant Bacteria. ACS Biomater. Sci. Eng. 2019, 5, 4228–4243. [CrossRef]  
56. Santhosh, S.M.; Natarajan, K. Antibiofilm Activity of Epoxy/Ag-TiO2 Polymer Nanocomposite Coatings against Staphylococcus Aureus and Escherichia Coli. Coatings 2015, 5, 95–114. [CrossRef]