



Improving the Functional Properties of Egyptian Cotton Fabrics Using Tio₂ Nanoparticles



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Abstract

Titanium dioxide nanoparticles (TiO₂-NPs), which are optically active metal nanoparticles that give cotton textiles excellent self-cleaning, UV blocking, and antibacterial performance, are prepared using the sol-gel method to investigate the novel qualities of cotton fabric. Using succinic acid 5 g/L/SHP as an across-linking agent, TiO₂-NPs were prepared at two distinct concentrations (1% and 3% OWF) and applied to cotton fabrics (Giza 88 and Giza 94) to investigate the mechanical characteristics, UV-blocking abilities, microorganisms, and self-cleaning capabilities of textile. The characterization of TiO₂-NPs was investigated using UV–VIS spectroscopy, Transmission Electron Microscopic (TEM), Energy Dispersive X-ray spectra (EDX) and Zeta potential and particle size. The elemental analysis and surface morphology of treated cotton fabrics are characterized by Fourier transform infrared spectroscopy (FTIR), X-ray diffraction (XRD), and scanning electron microscopy (SEM). Tensile strength and elongation of the treated fabrics are also evaluated. The results showed that the average size of TiO₂ nanoparticles was 65.7 nm which has excellent UV protection value (UPF 50+), self-cleaning property, the inhibition zone increased at concentration of 3% for negative and positive bacteria (*Escherichia coli* and *Staphylococcus aureus*). The tensile strength decreased using 1% TiO₂-NPs but with increasing the concentration to 3%, the mechanical properties were improved. The results of this investigation demonstrated that the cotton fabric treated with TiO₂-NPs exhibited superior general characteristics when compared to the untreated samples. Giza88 was superior to Giza 94, due to the genetic characteristics of the variety.

Keywords: Titanium dioxide nanoparticles, UV protection, Self-cleaning, Antimicrobial, Textiles, Cotton fabrics, X-ray diffraction (XRD), Fourier transform infrared spectroscopy (FTIR).

1. Introduction

In recognition of its special qualities such as its softness, affinity for skin, biodegradability, absorbency, and moisture comfort cotton is the most widely used natural fiber in the textile industry [1]. Durably functionalizing cellulosic textiles to make them stain-repellent, air-permeable, hydrophobic, electrically conductive, photo-luminescent, selfcleaning, antimicrobial, UV protecting, and pollutant-removing while maintaining comfort is now possible. Novel approaches utilizing a range of functionalizing agents derived from metal salts can accomplish these characteristics such as metalorganic compounds, metal oxides, and metal nanoparticles [2]. The process of giving textile materials useful qualities is being revolutionized by nanotechnology [3]. Nanotechnology is needed to give cotton, the king of natural fibers, properties like flame retardancy, antimicrobial properties, UV protection, and extreme hydrophobicity. These qualities are also imparted with the aid of nanofinishing, which preserves cotton's natural comfort level. Furthermore, because of its nano-size range, comparatively little material is needed to impart the functional feature [2,4]. Due to its remarkable qualities, nanotechnology and nanoscience have become extremely important in recent years [5]. Nanoparticles (NPs) have garnered interest in a variety of sectors, including electronics, research, medicine, and environmental cleaning [6]. Since of their high surface area-to-volume ratio (SVR) and high surface energy, NPs are becoming more and

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more in demand since they may be used in a variety of ways [7]. The NPs can be divided into two groups according to the elements they contain: metallic and non-metallic [8].

While metal NPs primarily consist of gold (Au), silver (Ag), Titanium (Ti), platinum (Pt), copper (Cu), and Fe (0), metallic NPs also include metal oxides. Titanium dioxide (TiO₂) [9], zinc oxide (ZnO) [10], iron oxide (Fe₂O₃/Fe₃O₄) [11], magnesium oxide (MgO) [12], copper oxide (CuO) [13], alumina (Al₂O₃), and numerous other metal oxide nanoparticles are among the metal oxide nanoparticles [14]. The synthesis of TiO₂-NPs can be achieved by three distinct methods: chemical, physical, and biological. The physical methods include the chemical gas-phase atomic layer deposition (ALD) technique, reactive DC magnetron sputtering, thermal evaporation, pulsed discharge plasma, and pulsed laser deposition (PLD). TiO₂-NPs recently employing a pressurized argon atmosphere and high-voltage discharge plasma [15].

Sol-gel synthesis was used by Jongprateep et al. [16] to create TiO₂-NPs with a size range of 48-85 nm. TiO₂-NPs were synthesized in this study using titanium (IV) isopropoxide (TTIP) as a precursor. TiO₂-NPs have several advantages over other metal oxides, including photo-catalysis, low cost, great abundance, self-cleaning properties, strong oxidizing power, and improved chemical stability [17]. Additionally, it has antibacterial, flame-retardant and UV protection qualities when used in textile finishing. TiO2-NPs have a wide surface area, excellent surface morphology, and inherently non-toxic [2,18]. Because of the oxygen vacancies, which encourage the growth of positive electrons or Ti³⁺ centers and, as a result, excess edonors in the electronic structure of titanium, it is an N-type semiconductor [19]. According to Žerjav et al. [20], the two main disadvantages of using undoped TiO₂ as a photo-catalyst are its broadband gap, which ranges from 3.00 to 3.30 eV depending on the polymorph of TiO₂ utilized, and high charge carrier recombination rate. Therefore, since TiO2 exhibits photo-catalytic action in the presence of a UV light source, this problem could be resolved by using a UV source. The purpose of this research is to coat cotton fabrics with titanium dioxide nanoparticles, which will enable the fabrics to selfclean and become anti-microbial as well as protect against damaging UV radiation.

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Sol-gel synthesis was used by Jongprateep et al. [22] to create TiO_2 -NPs with a size range of 48-85 nm. TiO₂-NPs were synthesized in this study using titanium (IV) isopropoxide (TTIP) as a precursor. TiO₂-NPs have a number of advantages over other metal oxides, including photo-catalysis, low cost, great abundance, self-cleaning properties, strong oxidizing power, and improved chemical stability [23]. Additionally, it has antibacterial, flame-retardant and UV protection qualities when used in textile finishing. TiO2-NPs have a wide surface area, excellent surface morphology, and are inherently non-toxic [3,24]. Because of the oxygen vacancies, which encourage the growth of positive electrons or Ti³⁺ centers and, as a result, excess edonors in the electronic structure of titanium, it is an N-type semiconductor [25]. According to Zheng et al. [26] and Žerjav et al. [27], the two main disadvantages of using undoped TiO₂ as a photocatalyst are its broad band gap, which ranges from 3.00 to 3.30 eV depending on the polymorph of TiO₂ utilized, and high charge carrier recombination rate. Therefore, since TiO₂ exhibits photo-catalytic action in the presence of a UV light source, this problem could be resolved by using a UV source. The purpose of this research is to coat cotton fabrics with titanium dioxide nanoparticles, which will enable the fabrics to self-clean and become antimicrobial as well as protect against damaging UV radiation.

2. Materials and Methods

2.1. Materials

Unbleached long stable Egyptian cotton fabrics were purchased from Miser-El-Mhala Company for Spinning and Textile, Egypt, which were created from Giza 88 and Giza 94. The textiles are plan weaved, with a warp of 36 yarns per cm² and a weft of 30 yarns per cm², weighing 150 grams. Every experimental material was kept in a controlled atmosphere with a temperature of $25^{\circ}C \pm 2^{\circ}C$ and a relative humidity of $65^{\circ}C \pm 2\%$.

2.2. Chemicals

Titanium (IV) isopropoxide,+97% (C₁₂H₂₈O₄Ti) was obtained and purchased from Thermo Fisher (Kandel), Germany. Succinic acid 99%, NaOH and

 H_2O_2 were purchased from PIOCHEM. Sodium hypophosphate (SHP) was obtained from Central Drug House (P) LTD, New Delhi. Reactive (drimen) dye blue 2 ($C_{29}H_{20}ClN_7O_{11}S_3$ was purchased from ICN Biomedical INC., Germany.

2.3. Pretreatment processes

2.3.1. Scouring

The procedure of scouring was in accordance with Hebeish *et al.* **[21]**. Giza 88 and Giza 94 cotton fabrics were immersed in one liter of an aqueous solution containing 40g/L NaOH (1M) and 1% mercerol (wetting agent) at 100°C for ninety minutes while being constantly stirred. The liquor total ratio will remain at 1:30. Following scrubbing, the samples underwent a thorough washing under running water and then samples were dried.

2.3.2. Bleaching

The following bleaching procedure was to bleach fabric in the slack state according to Hebeish *et al.* **[21]** method: The scoured materials were bleached for one hour at 90°C using the silicate (Na₂SiO₃.9H₂O) 3.5%, Na₂CO₃ 1% and H₂O₂ 4%. At a liquor ratio of 20:1, every desiring, scouring and bleaching experiment was run in triplicate.

2.3.3. Dyeing process

The altered textiles were colored using reactive (drimen) dye with 6% owf (on fabric weight) at a material-to-liquor ratio of 1:50 using 1% owf. The samples were first submerged in the water, and the dyeing bath was heated to 50°C. The prepared salt was added to the dyeing bath three times, and the dye solution was added. After 15 minutes of boiling, the dyeing was allowed to continue for 45 minutes, at which point it was stopped and the dyeing bath was cooled. Following a thorough rinsing with running water, the dyed samples were soap-soaped for 15 minutes at 70°C in a solution containing 5g/L nonionic detergent (Hostapal CV-ET). After being cleaned, the samples were finally rinsed with water and allowed to dry in air [22].

2.4. Treatment Processes

2.4.1. Preparation of TiO₂-NPs

Titanium tetra isopropoxide (6 mL) and 1% acetic acid (2 mL) were mixed using a magnetic stirrer. After five minutes, two milliliters of strong HCl were added to the solution to adjust its pH, after 56 milliliters of ethanol had been added dropwise while stirring continuously. After 45 minutes of vigorous magnetic swirling of the combination, a viscous solution was created, signifying the formation of TiO₂-NPs sol-gel. Abou-Okeil *et al.* **[23]** followed this procedure with two hours of calcination at 450°C and twenty-four hours of drying at 110°C in a muffle furnace.

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2.4.2 Cotton treatment with TiO₂-NPs

Succinic acid (5 g/L) and sodium hypophosphite (4% wt/wt) were dissolved in a 100 mL finishing bath to create TiO₂-NPs sol-gel cross-linked on cotton fabric samples. To create a dispersed solution, the prepared solution of TiO₂-NPs (1% and 3% owf) was sonicated for 20 minutes. The cotton fabrics were submerged in the prepared solution at room temperature. Ultimately, the treated materials underwent three minutes of drying at 120°C and one minute of curing at 150°C. These procedures completed twofold drying and curing [24].

2.5. Characterization of nanoparticles

2.5.1. UV – VIS spectroscopy

A PG Instruments Ltd. T80 UV-Vis spectrophotometer was used at the Cotton Research Institute (CRI), Agriculture Research Centre (ARC), Egypt, to study the optical characteristics of TiO₂-NPs. UV-VIS spectra were recorded between 300 and 700 nm in wavelength.

2.5.2. Transmission Electron Microscopy (TEM)

Utilizing transmission electron microscopy (JEM - 1400, JEOL model), TiO₂-NPs were morphologically examined. Cairo University Research Park's (CURP) electronic microscope lab captured the images. TiO₂-NPs were put on a carbon grid that had been glow-discharged. The TEM operating system then determined the diameter and shape of the nanoparticle specimens.

2.5.3. Fourier transforms infra-red spectroscopy studies (FTIR)

Using a Shimadzu FTIR 8400 Spectrophotometer (National Institute for Standards (NIS)), the FTIR spectrum was acquired to determine the functional groups contained in the samples.

2.5.4. Particle size and zeta potential

Particle size and zeta potential of the nanomaterials measured by photon interconnection were spectroscopy and Laser Doppler intensity measuring, respectively, using a Zetasizer® 3000 particulate size description analyzer (Malvern Instruments) in the central lab of NIS, a complementary scientific search institute with analytical services. The mean hydrodynamic diameter was determined by advance analysis after size modification was carried out three times at a 25°C scattering angle and confined for three minutes each time. The automatic water dip cell mode was used to verify the zeta potential adjustment.

2.6. TiO_2 -NPs coated with cotton fabrics: characterization.

2.6.1. Scanning Electron Microscopy (SEM studies)

SEM Quanta 250 FEG (Field Emission Gun) combined with an EDX unit (Energy Dispersive X-ray Analyses) at 30 KV accelerating voltage and 14x to 1 million magnification was used to capture images of the nanoparticles.

2.6.2. X-ray diffraction (XRD)

The nanoparticle on the textiles was characterized using X-ray diffraction (XRD). Using CuK radiation and λ =1.5406A, XRD patterns were obtained with a Bruker D8 advanced rotaflex diffraction meter.

Crystallinity of cellulose: The XRD crystallinity index (CIXRD) was determined from the height ratio given below using the peak height method for native cellulose, which was created by Segal *et al.* [25].

CI _{XRD} (%) =
$$\frac{I_{002} - I_{am}}{I_{002}} \times 100$$

Where I_{002} was the intensity of the crystalline and I_{am} the height of the minimum.

2.7. Application of TiO₂-NPs on cotton fabrics

2.7.1. UV protection

Using a Cary 50 solar screen transmission spectrophotometer, the transmissions of UV radiation were measured by AS/NZS 4399(1996)-Sun protective apparel - Evaluation and Classification. According to Sharma *et al.* **[26]**, UPF values were automatically calculated and classified using the Australian Standards Association of Australia standard SAAS/NZS 43399, which stands for Australian / New Zealand Standards, Home bush, Australia, 1996. The amount of UV light that can pass through a substance determines its LIPE. When compared to an

substance determines its UPF. When compared to an unprotected person, UPF indicates how long a wearer of a specific clothing may stay outside in the sun before suffering skin reddening [28].

Table 1: UV protection and classification [27]

UV protection	UPF classification	Transmitted UV radiation		
Excellent	40,45,50,50+	≤2.5%		
Very good	25,30,35	4.1-2.6%		
Good	15,20	6.7-4.2%		
Non-ratable	0,5,10	>6.7%		

2.7.2. Anti-bacterial screening test

Gram-positive *Staphylococcus aureus* (ATCC 6538) and Gram-negative *Escherichia coli* (ATCC 8739) were the two microorganisms employed in the agar plate method as a qualitative test to determine the antibacterial impact of the materials **[29]**.

2.7.3. Agar plate method

The bacteriostatic activity was applied in sterile Petri plates. The test organisms, 24-hour broth cultures of *S. aureus* and *E. coli*, were used as inoculums. Using a sterile cotton swab, the test organisms were swabbed all over the surface of the agar plates. The test textiles (fabrics coated with nanoparticles) and the control fabrics were gently pressed against the core of the mat culture. The plates were incubated at 37°C for 18 to 24 hours.

2.7.4. Self-cleaning property

Self-cleaning finished goods and experiments The untreated and TiO₂-loaded fabric samples were dried and conditioned after being dipped for 15 minutes in a basic blue dye solution (1% owf) for staining. The samples were subjected to a 24-hour exposure to UV light. Fabric samples that had not been treated or that had been loaded with TiO₂ were assessed for color strength or K/S value. Degradation of the dye stains directly causes a drop in K/S values [**23,30,31**]. It is determined according to the following relation:

Extent of discoloration (%) = $(K/S)a \ge 100 (K/S)b$ where (K/S)a is the color strength after exposure to daylight and (K/S)b is the color strength before exposure to daylight.

2.7.5. Strength and Elongation

The tensile stress is expressed as the force per unit linear density of the unstrained specimen. The tensile strength (Kg f) and elongation (%) were measured according to ASTM 5035 [**32**] using Shimadzu Testing Machino-Japan at Textile Metrology Department- National Institute for Standard (NIS-Egypt).

2.8 Statistical analysis

All experiments were independently repeated in triplicates. Results were expressed as mean \pm standard error (SE). The statistical analysis was performed in MS Excel and Origin 6.1 software.

3. Results and Discussion

The main task of this work is to enhancement of the cotton fabric properties such as UV-protective ability, self-cleaning, and antimicrobial properties by preparing TiO_2 -NPs and succinic acid as cross-linking agents. Results discuss the size and morphology of the TiO_2 nanoparticles and effect on the cotton fabrics.

3.1. Characterization of TiO₂-NPs.

3.1.1. UV – VIS spectroscopy of TiO₂-NPs

The absorption (A) spectrum of TiO₂-NPs was measured by Lambada spectrophotometer to find the wavelength peak (lambda max \approx 316 nm) the absorption spectrum shown in Fig. (1). These results are in agreement with Yang **[33]** which found that the wavelength range of TiO₂-NPs between 300-400nm and the optimum particle size of spherical TiO₂ is 50-120 nm.



Figure 1: UV-VIS spectrum of TiO2-NPs.

3.1.2. Transmission Electron Microscope (TEM) of TiO₂-NPs

TEM is a useful technique for the analysis of the size and shape of ultrafine particles. Fig. (2) shows the TEM of the prepared TiO_2 -NPs by sol-gel method. The size and shape of the prepared TiO_2 -NPs are homogeneous mono-dispersed with some agglomeration. TiO₂-NPs were observed to be irregular and hexagonal and the nanoparticle's size displays arrange of 65.7nm [**34**].



Figure 2: Electron micrographs of TiO₂-NPs.

3.1.3. Energy Dispersive X-Ray Spectra (EDX) of TiO_2 -NPs

EDX analysis was carried out to verify the composition of the grown nanoparticles. The standard EDX analysis of the grown TiO_2 -NPs is shown in Fig. (3). The EDX analysis confirms that the grown nanoparticles consist solely of titanium and oxygen, with a percentage of atomic Ti:O of 58.69% compared to 41.31%. Based on EDX and quantitative analysis data, the molecular ratio of Ti:O of the grown nanoparticles was calculated to

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be nearly that of the bulk. The fact that the spectrum only shows peaks for Ti and O and no other element further indicates that the nanoparticles are made entirely of TiO_2 [35].



Figure 3: Energy Dispersive X-Ray Spectra of TiO₂-NPs.

3.1.4. Fourier Transform Infrared studies (FTIR).

Fourier Transform Infrared Analysis of TiO_2 -NPs was shown in Fig. (4). The broad band centered at 500-600 cm⁻¹ is assigned to the bending vibration (Ti-O-Ti) bonds in the TiO_2 lattice. The broad band centered at 3600-3400 cm⁻¹ was assigned to the intermolecular interaction of the hydroxyl group for water molecule with TiO_2 surface. The peak at 1650 cm⁻¹ refers to the characteristic bending vibration of OH⁻ [**36**].



Figure 4: Fourier Transform Infrared of TiO2-NPs.

3.1.5. Zeta potential and particle size.

Zeta analysis, which is dependent on surface charge, is used to determine the stability of suspended nanoparticles. Particle size and zeta potential of produced TiO₂-NPs were investigated according to Rasmussen *et al.* [**37**]. The type of hydrodynamic size (diameter) of produced TiO₂-NPs in distilled water was investigated. Fig. (5a) displays the zeta potential for a TiO₂-NPs suspension in distilled water. A distinct stability index was observed on zeta potential of -3.14 mV, with an electrophoretic movement (mean) of 0.00253 cm²/Vs. while Fig. (5b) displays the average hydrodynamic diameter of 150 nm for the particle size, These results in agreement with Albukhaty *et al.* **[38]**, who found that the DLS of TiO₂-NPs is 150 nm with zeta potential (A) of the opposite value of -44,6mV with electrophoretic movement (mean): 0,000 346cm²/Vs and these findings in agreement with Li *et al.* **[39]**, who found that the prepared suspension confirmed the potential general parameters of zeta negatively to improve stability.



Figure 5: (a) Zeta potential and (b) Size distribution by number of TiO₂ -NPs.

3.2. Characterization of Nano finishing in textiles 3.2.1. Fourier Transform Infrared studies (FTIR) of cotton fabrics (Esterification with succinic acid)

Fig. (6a) shows the *FTIR* spectrum of succinic acid. It has been demonstrated that the carboxylic group of succinic acid and the hydroxyl group of cellulose polymer chains can form an ester bond (Fig. 6b). Sodium hypophosphite (SHP) has also been claimed to have the ability to speed up ester formation reactions. It first reacts with an intermediate cyclic anhydride before reacting with cellulose.



Figure 6: (a) Synthesis of TiO_2 -NPs, cellulose, and dicarboxylic acids (b) Ester covalent and electrostatic bonds, and cyclic anhydride intermediates are formed during the esterification of cellulose with succinic acid [40].

Figs. (7 and 8) display the recorded FTIR spectra for samples (G88 and G94) treated with 5 g/L of

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succinic acid and 3% of TiO₂-NPs. The carbonyl group's (C=O) absorbance peak for esters can be seen between 1700 cm⁻¹ and 1750 cm⁻¹. The antisymmetric stretching vibrations of ester carbonyl in carboxyl groups are responsible for the absorption band at 1725 cm⁻¹. Regarding the TiO₂ spectrum (586 cm⁻¹), the peak at 569 cm⁻¹ belongs to the Ti–O stretching band. This is explained by the Ti–O covalent connections that develop between cotton fibers and TiO₂. It follows that TiO₂-NPs interact with cotton fibers [**40**]. This band is visible in every sample that has had carboxylic acid treatment. This suggests that cotton fabric and succinic acid have established ester bonds.



Figure 7: FTIR spectrum of cotton fabrics samples (G88) coated with TiO_2 -NPs (a) Untreated cotton (b) Treated with TiO_2 -NPs.



Figure 8: FTIR spectrum of cotton fabrics samples (G94) coated with TiO₂-NPs (a) Untreated cotton (b) Treated with TiO₂-NPs.

3.2.2. Scanning electron microscope (SEM)

The surfaces of the treated fabrics were examined using SEM microscopy. The surface morphology of cotton fabric samples (Fig. 9b) treated with TiO_2 -NPs and bleached cotton fabric (Fig. 9a) is shown in Fig. 9 (a,b). We can see a picture of the delicate and smooth untreated fiber surface in Fig. (9a). However, Fig. (9b) makes it possible to distinguish the TiO_2 -NPs on the surface of each sample in the treated fabrics. TiO_2 -NPs attach to the fibers efficiently, adhering to the surface as well as penetrating textile flaws. Additionally, the nanoparticles have a homogeneous distribution throughout the fiber surface, even



though some aggregated nanoparticles are still visible. The treated cloth appeared brilliant and white in the SEM images, indicating the presence of white TiO_2 -NPs on the fabric surface.

The size of the particles affects their ability to adhere to the fibers. The findings are in agreement with those of Saad [34] and Azeem *et al.* [41] and imply that while the smaller particles will penetrate deeper and adhere strongly to the fabric matrix, the bigger particle agglomerates will be easily removed from the fiber surface. Albukhaty, *et al.* [38] explain that sol-gel is one of the most efficient methods for creating TiO_2 nanopowder based on their investigation and analysis.



Figure 9: SEM of cotton fabric samples coated with $TiO_2 - NPs$ (a) Untreated cotton (b) Treated with $TiO_2 - NPs$.

3.2.3. X-Ray Diffraction Studies

The substance applied to the fabrics throughout the investigation was examined for crystallinity using XRD studies. Both the applied powder and the substrate's crystallinity are revealed by the XRD pattern. The XRD of cotton fabric G94 and G88 are displayed in Figs. 10 (a,b) and 11 (a,b), respectively. Figures displaying the cotton fabric treated with titanium dioxide XRD graph. Two major peaks can be seen in the graph. The peak at 17 is from cellulose found in all figures, while the peak at 25 is from TiO₂-NPs found only in the treated fabrics. This peak confirms the presence of nanoparticles on the fabrics. These results are consistent with Azeem et al. [41]. The crystallinity index was increased from 76.40 % to 80.88% in the case of G88 and crystallinity index of G94 was increased from 75.42 % to 76.84% in concentration 3% TiO₂-NPs which confirmed by the results of tensile strength at concentration 3% these due to the filling of inter yarn spaces in fabric due to this the friction between yarns produced and they resist to elongation and These findings are in agreement with Azeem et al. [41].



Figure 10: XRD studies of cotton fabric samples (G94) coated with TiO2–NPs (a) Untreated cotton (b) Treated with TiO_2 –NPs.



Figure 11: XRD studies of cotton fabric samples (G88) coated with TiO_2 -NPs (a) Untreated cotton (b) Treated with TiO_2 -NPs.

3.3. UV protection properties

The main aim of the current work was to use TiO_2 -NPs to create a highly UV-absorbing technology that could be used on fabrics. The application of TiO_2 -NPs with succinic acid/SHP as a cross-linking agent at varying concentrations (1% and 3%), as demonstrated by the results in Table (2).

When compared to untreated cellulose fabrics, treated cellulose fabrics (G88 and G94) showed lower UV transmission. This led to an increase in the automatically calculated UPF (Ultraviolet Protection Factor) value, which indicates how long a person can stay in the sun with special clothing before their skin starts to redden. Table (2) displays the UVA radiation transmitted for cotton fabrics in the UVA (320–400 nm) and UVB (280–320 nm) zones (abbreviated as TUVA and TUVB).

According to the findings, G88 and G94 both suggest a higher UPF value. Excellent UV protection is provided by all treated cotton fabrics. The findings showed that TiO_2 -NPs treated fabrics could be used for various technological purposes as well as for protecting the skin from sun radiation. Abbas *et al.* [42], and Prorokova *et al.* [43] are in agreement with these results of TiO_2 -NPs.

Table 2:	Effect of	TiO ₂ -NPs	with	succinic	acid o	on UV	protection
of cotton	fabrics						

	G94			G88		
Verities	UPF value	TUVA	TUVB	UPF value	TUVA	TUVB
Control	33.33±3.179797	7.29	3.03	35.86±2.194944	5.6	3.15
TiO ₂ -NPs 1%	57.54±3.539371	1.71	1.65	72.04±3.308294	2.56	1.05
TiO ₂ -NPs 3%	97±3.997528	1.85	0.76	112.12±4.246667	1.18	0.68

Transmitted UVA (320-400 nm) and UVB (280 -320 nm) regions which were expressed as TUVA and TUVB. UPF: UV protection factor TUV Transmitted UV radiation %. Transmitted UVA (320-400 nm) and UVB (280 -320 nm) regions which were expressed as TUVA and TUVB. UPF: UV protection factor TUV Transmitted UV radiation %. *Each value represents the mean \pm standard error, n=3.

3.4. Antibacterial properties

The results of the antibacterial test are reported in Tables (3 and 4). The tables show the comparison of antimicrobial activity of the treated fabrics as zone inhibition of bacteria (for the two representative types *S. aureus* and *E.coli*) for the fabric samples (G88 and G94) treated with two different concentration levels of TiO₂-NPs concerning fabric weight such as 1% and 3%. It is clear from the results there is no reduction of bacterial found on the bleaching cotton fabrics but the treated fabric with TiO₂-NPs with 1% and 3% show the highest antimicrobial activity against both *S.aureus* and *E.coli* bacterial and response of Giza 88 Variety.

The antibacterial activity of TiO_2 -NPs in textiles is related to its crystal structure, shape and size. It is based on degrading organic materials by photo-catalytic reaction. Strong chemical reactivity is exhibited by the oxygen and hydroxyl radicals that are formed. Specifically, atomic oxygen reacts with the majority of organic compounds to produce carbon dioxide and water when it reacts with organic compounds in bacteria. Consequently, the bacteria can be eliminated quickly, and textile odors and oil stains can be removed [**38**].

 Table 3: Effect of TiO₂-NPs with different concentration on zone inhibition of *E.coli*

Verities	G88	G94			
Inhibition					
	Zone (mm)				
Control	0±.000	0±0.00			
TiO ₂ -NPs 1%	10±1.04083	9±1.4433			
TiO ₂ -NPs 3%	13±1.2583	11±1.5275			

*Each value represents the mean \pm standard error, n=3.

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 Table 4: Effect of TiO₂-NPs with different concentration on zone inhibition of *Staphylococcus aureus*

Verities	G88	G94			
Inhibition					
Control	0±.000	0±0.000			
TiO ₂ -NPs 1%	12±1.607275	11±1.322876			
TiO ₂ -NPs 3%	15±1.154701	14±1.040833			

*Each value represents the mean \pm standard error, n=3

3.4.1. Mechanism of antimicrobial activity of TiO₂-NPs

O₂-NPs exhibit negative charges on their surface at the point of zero charge (pzc) at pH = 6.2, which indicates a reduced bactericidal effect in neutral and alkaline solutions. This is so because, in these conditions, TiO₂-NPs repel microorganisms with a negative charge in the absence of light [44]. The TiO₂-NPs are positively charged and exhibit significant bacterial cell-to-cell interactions in acidic pH conditions, the bacterial membrane is penetrated, leading to oxidative damage [45]. A thorough sequence implicated in TiO2-NPs toxicity to microorganisms is shown in Fig. 12. By adhering to TiO_2 -NPs on the surface of the microorganism, TiO_2 inhibits or eliminates the microbe. Reactive oxygen species (ROS) (O^{2-}, OH^{\cdot}) are formed and initially interact with the lipids on the microorganisms' membrane surface [46]. Lipid peroxidation results from the interaction of ROS with the lipids on the membrane. Furthermore, there is an increase in the lipid membrane's permeability. After causing damage to the membrane lipids, the ROS enters the microorganism's cytoplasm and begins to destroy its DNA and ribosomal proteins as well as the nucleus and other cellular organelles. Denaturation of proteins and DNA is caused by ROS oxidizing them. Finally, the microorganisms are killed when the damaged lipid membrane allows all of the damaged cytoplasmic contents to flow out. Prior research has also demonstrated the toxicity of TiO2-NPs on a variety of microorganisms. Kiwi et al. [9] demonstrated that TiO2-NPs had a bactericidal impact on Escherichia coli when they came into close contact with the bacteria in a dark condition. The minus-charged bacterial cell wall at a pH near but below the point of zero charge is attracted to the TiO₂-NPs electrostatically, which damages the cell wall during this process. According to Vatansever et al. [47], the fundamental source of TiO₂-NPs antibacterial action is the formation of ROS when UV light is present. This implies that the bactericidal activity is caused by UV light rather than TiO₂-NPs. TiO₂-NPs have also shown promise in the destruction of multidrug-resistant bacteria by generating reactive radicals via electron-hole pairs

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during UV irradiation [48]. The power and duration of UV-A light irradiation are critical factors in the inactivation of drug-resistant bacteria by photocatalytic materials like TiO₂-NPs [49]. Because of this, the disinfection approach requires a high-power UV source to excite TiO₂-NPs, and because of their poor photo-excitation in visible light, there are fewer bactericidal applications. Because of this, TiO₂-NPs effectiveness against microbes is restricted in indoor environments with low UV light levels. Therefore, one of the most pressing needs in the medical and industrial sectors is the development of such TiO₂-NPs that, in addition to their remarkable antibacterial capabilities, may be activated by visible light.



Figure 12: Schematic diagram of the toxicity of TiO₂NPs to microorganisms [50].

3.5. Self-cleaning properties

Cotton fabrics coated with TiO2-NPs were analyzed using the photo-catalytic degradation methylene blue under UV-visible irradiation. Fig. 13 shows the degradation methylene blue of TiO₂-NPs on fabrics (G88 and G94) with different concentrations (1% and 3 with respect to the weight fabrics) under UV-visible irradiation at 24 h the data shows that the degradation rate increases with the increasing of TiO₂ concentration and irradiation time these results in agreement with Kale et al. [51]. who found that all TiO₂ coated samples show the photo-catalytic properties and capable of stain degradation. Also, the results were in agreement with Alvarez-Amparán et al. [52], who found that the TiO₂-NPs functionalized cotton fabrics degraded 99% of methylene blue in 60 min by photo-catalysis under UV and solar irradiation conforming selfcleaning properties. Abou-Okeil et al. [23] found that 2 % of TiO₂ nanoparticles are almost sufficient to give enough self-cleaning.



Figure 13: Image of methylene blue degradation by $\rm TiO_{2}\mathchar`-NPs$ on cotton fabrics.

3.5.1. Self- cleaning mechanism of TiO₂-NPs

The fundamental mechanisms of TiO2-NPs photocatalysis have been the subject of numerous studies [53-56]. As previously mentioned in the introduction, heterogeneous photo-catalysis is based on the theory that photon energy either equals or exceeds the band gap ($hv \ge Eg$) of a semiconductor, typically TiO₂-NPs, upon absorption. As shown in Fig. 14, highly reactive oxygen species (ROS) are generated when the newly formed electron-hole pairs take part in a series of oxidation-reduction events with species adsorbed on the TiO₂-NPs surface to generate highly reactive oxygen species (ROS). These organisms cause hazardous pollutants like CO₂ and H₂O to break down when they interact with the bacteria or adsorbed organic compounds on the TiO₂-NPs surface. However, the shorter halflifetimes of electrons and holes usually restrict their availability for participation in redox reactions. Moreover, the recombination event is indicated by energy emitted as waste heat or light and a decrease in process efficiency.



Figure 14: TiO2-NPs semiconductor photo-catalysis principle.

3.6. Mechanical properties of cotton fabric

3.6.1. Tensile elongation and strength Tensile strength is measuring of fiber strength which depends on the crystal region of the cellulose unit which is observed as a parallel unit that makes a block that makes strength. The mechanical data of the treated and untreated samples were obtained in Table (5). TiO₂-NPs treatment with two different concentrations and succinic acid caused increments in fabric strength accompanied by a decreasing in concentration recorded from 420 N in the control sample to 392 N under a concentration 1% then followed slightly increased by concentration of 3% recorded of 423.63 N these in Giza 88.

The same matter in Giza 94 also decreased in concentration recorded from 365.46 N in the control sample to 317 N in concentration of 1% then followed by increasing under concentration of 3%. The decreasing value of tensile strength due to the acidity of succinic acid in the impregnation bath. This leads to an irreversible de-polymerisation of the cellulose molecules **[24]**.

TiO₂-NPs exhibit strong photo-catalyst ability; the fiber macromolecules are readily broken, increasing the number of molecular chains in the amorphous portion of the fiber and weakening its supermolecular structure. In the end, this causes the fibers' macro-mechanical properties to decrease. Therefore, during the development and use of nanoscale composite multifunctional textiles, it will be essential to make the nanometer particle equally spread onto the textile and firmly combine it with nanometer material and fiber [57]. However, Rahman et al. [58] discovered that a 2% concentration of TiO₂ increased the tensile strength. Further improves the tensile strength because the fabric between spaces are filled, creating friction between the yarns that prevents elongation. These findings concur with those of Azeem et al. [41], who discovered that tensile strength improves along with increasing TTIP concentration.

On the other hand Fabric elongation is the increase in length of specimen expressed as a percentage of the original fiber length. Elongation is a property against the strength in which every increase in strength opposite decrease in elongation as shown in Table (5).

TiO₂-NPs treatment with two different concentrations and succinic acid 5 g/L leads to an increase in fabric elongation % in Variety G94 which was increased by increasing the concentration relative to control that recorded 19.40 mm and increased in concentration 1% to 25.77 mm then decrease gradually to 24.14 mm by concentration of 3% TiO₂ these concentrations have a highest tensile strength.

On the other hand, the elongation % of G88 in the control (untreated cotton fabric) was recorded at 18.33 mm and the treatment of TiO_2 -NPs with succinic acid led to an increase in concentration 1% recorded at 22.24 mm and then decreased with increased concentration to 3% which recorded 21.02

mm. Sadr and Montazer [59] found that The shrinkage that may occur during sonication, causing a slight increase in breaking elongation from 5 to 6%, could be linked to the loss of tensile strength fabric and/or to the cleavage of the cellulosic chains by acid hydrolysis. The results showed that *in situ* sono-synthesis of TiO₂-NPs on cotton fabric did not significantly damage the cotton structure. These results are in agreement with Saad [34] which found that all treated fabrics with TiO₂-NPs showed a decrease in TiO₂ nanoparticles can be attributed to those spaces between fibers after treatment decreases.

 $\label{eq:constraint} \begin{array}{c} \textbf{Table 5:} \\ \textbf$

	T.S (N)		Elongation (mm)		
Treatment	G94	G88	G94	G88	
Control	365.46±19.1796	420±22.1435	19.40±2.34109	18.33±2.3784	
TiO ₂ NPs 1%	317 ±22.3960	392±23.4591	25.77±1.2850	22.24±2.9834	
TiO ₂ NPs3%	375 ±22.8983	423.63±23.3225	24.14±2.3140	21.02±2.5027	

*Each value represents the mean \pm standard error, n=3.

4. Conclusion

Significant functional qualities including antibacterial activity, self-cleaning activity using the methylene blue stain method, and ultraviolet protection factor were evaluated for the TiO_2 -NPs with succinic acid treated fabric surfaces. All findings demonstrated the best performance at concentrations (3% TiO_2 -NPs). TiO_2 -NPs reduced the tensile strength by 1%, but the mechanical characteristics increased when the concentration was raised to 3%. The variety's genetic traits demonstrated that Giza 88 is superior to Giza 94.

5. Conflicts of interest

There are no conflicts to declare.

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