THE STRUCTURAL, MORPHOLOGICAL AND OPTICAL STUDY OF PURE AND W-DOPED TiO2 NANO PARTICLES AND ITS APPLICATION TO ANTIMICROBIAL ACTIVITY

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ABSTRACT

In this research work, the effect of tungsten-doping on the crystal structure, morphology and antimicrobial of titanium dioxide nanoparticles were studied. The pure and different weight % of tungsten doped TiO2 nanoparticles were synthesized by sol-gel method and calcinated at 600°C for 5 h. The synthesised products have been characterized by X-ray Diffraction studies (XRD), Field Emission Scanning Electron Microscopy (FESEM), Energy Dispersive X-ray analysis (EDXA), Ultraviolet-Visible Diffuse Reflectance Spectroscope (UV-Vis), Photoluminescence Spectra (PL), High Resolution Transmission Spectroscopy (HRTEM) and Fourier Transform Infra Red Spectroscopy (FT-IR). XRD pattern of pure TiO2 and 1 wt % W-doped TiO2 nanoparticles confirms the anatase structure and increase in the W-doping changes the phase of TiO2 to rutile. Average crystallite size of synthesized nanoparticles was determined using the Debye-Scherrer formula. The crystallite size obtained for pure TiO2 is 37 nm and W-doped TiO2 is 28 nm, 34 nm and 33 nm. The FESEM images show the agglomerated particles of spherical-like morphology. Optical property and direct bandgap of pure and W-doped TiO2 nanoparticles also further characterised by UV–Vis Spectroscopy. The images of HRTEM clearly confirm that particles present in the W-doped TiO2 powdered sample is nanosized particles. The Kirby Bauer Agar Well Diffusion Assay method was employed to explore antimicrobial activity of nanosized pure and W-doped TiO2 colloidal suspension against the test microorganisms two Gram positive bacteria (Staphylococcus aureus, Bacillus subtilis), two Gram negative Bacteria (Escherichia coli, Pseudomonas aeruginosa) and two fungi (Candida albicans, Aspergillus niger). It shows that the W-doped TiO2 nanoparticles inhibited the multiplication and growth of the above mentioned test bacteria and fungi. Antimicrobial activity was found against all tested microorganisms which confirmed that W-doped TiO2 nanoparticles possess high antimicrobial activity compared to pure TiO2 nanoparticles.

Keywords: Nanoparticles, Sol–gel method, XRD, HRTEM, Antimicrobial activity.

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. INTRODUCTION

Titanium dioxide is used as an eco friendly photo catalyst. It is an inexpensive, easily available, nontoxicity and chemically stable one. However, TiO2 is used other than photocatalyst and this is used in catalyst support, photoconductors, solar cells, gas sensors, coatings etc. Now the scientist are focused in antimicrobial activity of TiO2 due to its rapid recombination of photo activated electrons and positive holes. Titanium dioxide present in three different crystalline phases: rutile, anatase, and brookite. Out of these three phases, Rutile phase is stable one compared to other two phases and are in metastable. Anatase and rutile systems have tetragonal unit cells. The rutile phase possesses two TiO2 molecules per unit cell having lattice constant a=4.5937Å and c=2.9587 Å and the anatase phase has four TiO2 molecules per unit cell having lattice constant a=3.7842 Å and c=9.5146 Å. Generally, the temperature of TiO2 nano crystal is increased to above 450°C [1, 2] phase has been changed from anatase to rutile structure. Anatase and rutile phases of TiO2 nanocrystals are the two important photoactive polymorphic phases with the band-gap energy of 3.20 eV and 3.02 eV respectively [3]. Band gap value of the anatase phase is larger than that of the rutile phase, so the rutile phase properties are slightly better than the anatase phase properties in semiconducting performance [4]. In recent years, researcher’s showed interest on the antimicrobial activity of doping of titanium dioxide with transition metals like tungsten, cobalt etc.

In this present study, synthesis of W-doped TiO2 nanoranged particles by sol-gel method is to study the highest possible antimicrobial activity and it can be compared with pure TiO2 also prepared by solgel method. There are several methods such as sol-precipitation [5], ion-impregnation [6], hydrothermal synthesis [7], sol-gel synthesis [8] to available for obtain homogeneous doping of W in TiO2. But, Sol-gel is the most simple and sophisticated method among the various methods for producing nanoparticles. The preparation of thin film W-doped TiO2 nanoparticles in a colloidal state by sol-gel method which is used in antimicrobial coatings in water filters, leathers, textiles and medical devices. The prepared W-doped TiO2 nanoparticles were characterised by different techniques like X-ray diffraction (XRD), Field emission scanning electron microscopy (SEM), Energy dispersive X-ray Spectroscopy (EDXA), UV–Visible Diffuse Reflectance Spectroscopy (UV-Vis), Photoluminescence analysis (PL), Fourier Transform Infra Red spectroscopy (FTIR), and High resolution Transmission Electron Microscopy (HRTEM). The Kirby Bauer Agar Well Diffusion Assay method was employed to explore antimicrobial activity of nanosized pure and W-doped TiO2 colloidal suspension against the test microorganisms two Gram positive bacteria (Staphylococcus aureus, Bacillus subtilis), two Gram negative Bacteria (Escherichia coli, Pseudomonas aeruginosa) and two fungi (Candida albicans, Aspergillus niger). It shows that the W-doped TiO2
nanoparticles induced the multiplication and the growth of the above mentioned test bacteria and fungi. Antimicrobial activity was found against all tested microorganisms at a concentration range of 25 µg, 50 µg, 75 µg and 100 µg of pure TiO$_2$, 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ solutions.

2. EXPERIMENTAL

2.1. Materials

Titanium isopropoxide (Sigma-Aldrich > 97 % pure), sodium tungstate dihydrate (Merk, 98 % pure), ethanol (Hayman (German) 99.99 % pure), hydroxylamine hydrochloride (Sigma-Aldrich 99 % pure) were used as precursor and is used without any further purification. Doubly distilled water was used for the whole synthesis process. Bacterial cultures such as Staphylococcus aureus, Bacillus subtilis, Escherichia coli, Aspergillus niger and fungus cultures such as Pseudomonas aeruginosa, Candida albicans were obtained from Eumic analytical Lab and Research Institute, Tiruchirappalli. Bacterial strains were maintained on nutrient agar slants (Hi media) at 4°C.

2.1.1 Physicochemical characterization

The X-ray diffraction pattern analysis for pure TiO$_2$ and doped TiO$_2$ nanoparticles was recorded by Lab X XRD6000 Shimadzu model with Cu-Kα radiation. The structure and morphology of the nanoparticles were investigated by Field Emission Scanning Electron Microscope (FESEM) using FEI Quanta FEG 200-High Resolution Scanning Electron Microscope. The absorption spectra and optical band gap of the TiO$_2$ nanoparticles were measured by using UV-Vis Spectrophotometer (JASCO U-670 Spectrometer) and the alcohol as a solvent. The spectrum was recorded between 200 – 800 nm. A Photoluminescence spectrum was recorded between 370-770 nm and it was carried out by using Horiba Jobynvcon model spectrophotometer and the alcohol is used as a solvent. FTIR absorption spectrum was recorded by JASCOFP8200 spectrophotometer. The particle size and lattice structure of the nanoparticles were investigated by JASCO XRD6000 Shimadzu model with CuKa radiation. The structure and morphology of the nanoparticles were investigated by Field Emission Scanning Electron Microscope. The absorption spectra and optical band gap of the TiO$_2$ nanoparticles were measured by using UV-Vis Spectrophotometer (JASCO U-670 Spectrometer) and the alcohol as a solvent. The spectrum was recorded between 200 – 800 nm. A Photoluminescence spectrum was recorded between 370-770 nm and it was carried out by using Horiba Jobynvcon model spectrophotometer and the alcohol is used as a solvent. FTIR absorption spectrum was recorded by JASCOFP8200 spectrophotometer. The particle size and lattice structure of the nanoparticles were investigated by JASCO XRD6000 Shimadzu model with CuKa radiation. The structure and morphology of the nanoparticles were investigated by Field Emission Scanning Electron Microscope. The absorption spectra and optical band gap of the TiO$_2$ nanoparticles were measured by using UV-Vis Spectrophotometer (JASCO U-670 Spectrometer) and the alcohol as a solvent. The spectrum was recorded between 200 – 800 nm. A Photoluminescence spectrum was recorded between 370-770 nm and it was carried out by using Horiba Jobynvcon model spectrophotometer and the alcohol is used as a solvent. FTIR absorption spectrum was recorded by JASCOFP8200 spectrophotometer. The particle size and lattice structure of the individual crystal was visualised by using High Resolution Transmission Electron Microscopy JE2100 (JEOL-200KV, LB6 filament) and EDX/A analysis was carried out to find the composition of pure and doped TiO$_2$ samples by using the detector attached with the same instrument.

2.2. Synthesis of pure TiO$_2$ and W-doped TiO$_2$ nanoparticles

Pure titanium dioxide nanoparticles and 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles were prepared by sol gel method [9]. For the preparation of TiO$_2$ nanoparticles, aqueous solution of titanium (IV) isopropoxide was used as starting material. The sol was prepared by mixing titanium isopropoxide (3 ml) with 24 ml of ethanol and dissolved 1000 ml of doubly distilled water at room temperature. The molar ratio of titanium isopropoxide and alcohol is 1:8 respectively. Hydroxylamine hydrochloride 0.694 g was dissolved in 100 ml of deionised water and added gradually to the titanium isopropoxide sol. After stirring, an aqueous solution was centrifuged by using centrifuge machine. The precipitate obtained was dried at 105°C in hot air oven. It was then calcinated at 600°C in a muffle furnace for 5 h at a constant temperature rise of 2°C/minute. For the preparation of 1 wt %, 3 wt %, 5 wt % W-doped TiO$_2$ nanoparticles, aqueous solution of titanium (IV) isopropoxide was used as starting material. The sol was prepared by mixing titanium isopropoxide (3 ml) with 24 ml of ethanol and dissolved 1000 ml of doubly distilled water at room temperature. The molar ratio of titanium isopropoxide and alcohol is 1:8 respectively. Hydroxylamine hydrochloride 0.694 g was dissolved in 100 ml of deionised water and added gradually to the titanium isopropoxide sol. For tungsten doping 1 wt % of sodium tungstate dihydrate solution was added to the TiO$_2$ sol. The mixture of titanium (IV) isopropoxide and sodium tungstate dihydrate solutions were stirred for 3 h in the magnetic stirrer. After stirring, an aqueous solution was centrifuged by using centrifuge machine. The precipitate obtained was dried at 105°C in hot air oven. It was then calcinated at 600°C in a muffle furnace for 5 h at a constant temperature rise of 2°C/minute. Similarly 3 wt % and 5 wt % W-doped TiO$_2$ powders were prepared by the same procedure as mentioned in the above method.

2.3. Antimicrobial activity

2.3.1. Preparation of culture media

2.3.1.1 Nutrient agar medium

Nutrient agar medium of pH 7 is one of the most commonly used medium for several routine bacteriological purposes. It was prepared by dissolving 5 g of peptone, 3 g of beef extract, 15 g of agar, 5 g of sodium chloride, 1.5 g of yeast extract in 100 ml of distilled water, it is boiled to dissolve the medium completely and sterilized by autoclaving at 15 lb psi pressure (121°C) for 15 min.

2.3.1.2 Inoculum preparation

Bacterial cultures were subcultured in liquid medium (Nutrient broth) at 37°C for 8 h and further used for the test (10$^2$-10$^8$CFU/ml). The suspensions were prepared before the test was carried out.

2.3.1.3 Assay of antimicrobial activity

The nutrient broth was prepared, and bacterial and fungal colonies were inoculated into the broth culture and are used to assay the antimicrobial activity.
2.3.2 Inhibition zone assay (Kirby Bauer Agar Well Diffusion Assay)

The nutrient agar medium was prepared and sterilized by autoclaving at 121°C 15 lbs pressure for 15 min then aseptically poured the medium into the sterile petriplates and allowed to solidify the bacterial and fungal broth culture was swabbed on each petriplates using a sterile buds. Then, wells were made by well cutter. 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles containing solutions were prepared dissolving 100 mg of each in 100 ml of DMSO solvent and from this stock solution, different concentrations of 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles calcinated at 600°C (25 µg, 50 µg, 75 µg and 100 µg) solutions were taken for assay.

The antimicrobial activity of 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles calcinated at 600°C were assayed against Staphylococcus aureus, Bacillus subtilis, Escherichia coli, Pseudomonas aeruginosa, Aspergillus niger and Candida albicans at 37°C for 24 h. After incubation the plates were observed for the zone of inhibition.

The bacterial and fungal viable count was determined after 24 h by plating pure TiO$_2$, 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles calcinated at 600°C on nutrient agar plates and the number of colony forming units (CFU) which were counted by a viable count method. To control this, the bacterial and fungal cultures were incubated with Gentamicin. The sample material which has antimicrobial activity was identified by inhibited growth of the microorganisms and it is clearly seen as distinct zone of inhibition. The diameter of zone of inhibition was measured and expressed in millimetre [10]. The prepared 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles calcinated at 600°C C nanoparticles with the sizes varied from 28 to 34 nm. Kirby Bauer Agar Well Diffusion Assay test was conducted using 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles calcinated at 600°C C nanoparticles and common antibiotic Gentamicin. The diameter of zone of inhibition for 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles calcinated at 600°C C was compared to this antimicrobial agent [11]. The diameter of zone of inhibition for pure TiO$_2$ at 600°C was also compared to this antimicrobial agent. To analyse the antimicrobial activity of 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles calcinated at 600°C C was obtained by the microemulsion method.

3. Results and discussion

3.1. X-ray diffraction analysis (XRD)

![XRD patterns of pure TiO$_2$ (a) and W-doped TiO$_2$ at different concentrations (b) 1 wt % (c) 3 wt % (d) 5 wt % W-doped TiO$_2$ calcinated at 600°C](image_url)

Figure 1 XRD pattern of (a) pure TiO$_2$ (b) 1 wt % W-doped TiO$_2$ (c) 3 wt % W-doped TiO$_2$ and (d) 5 wt % W-doped TiO$_2$ calcinated at 600°C
X Ray Diffraction analysis was used to determine the crystalline structure and phase of the synthesised nanoparticles. Figure 1(a-d) shows the X Ray Diffraction patterns of pure and W-doped TiO$_2$ nanoparticles. The results shows that most of the peaks to pure and 1 wt % W-doped TiO$_2$ nanoparticles calcinated at 600° C confirms the formation of anatase phase by the existence of strong diffraction peaks at 2θ values of 25.3°, 38.44°, 48°, 54°, 55.07°, 63°, 69.23°, 70.89°, and 75.38° corresponding to the crystal planes of (101), (112), (200) (105), (211) (204), (116), (220), and (215), respectively [JCPDS card no 21-1272] belonging to the tetragonal structure. The XRD diffraction patterns obtained for 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles are shown in the Figure 1(c and d). It showed the presence of characteristic rutile peaks at 2θ values of 27.4°, 36.0°, 39.1°, 54.3°, and 64.0° 2θ to the crystal planes of (110), (101), (200), (211), and (002) respectively [JCPDS: 88-1175] being a mix of anatase and rutile phase. The fineness peaks indicates that the nanosized materials were well crystallized. There are several studies reported [12, 13, 14] that sol-gel sample of TiO$_2$ should undergo a phase transformation from anatase to rutile during the higher calcinations (above 500° C) temperature. Here, the results showed that major phase transformation from anatase to rutile takes place with the increase of wt % of WO$_3$, new peaks appeared at 2θ =20.45° and 22.83° (marked as W), for the W-doped TiO$_2$.The new peaks may be attributable to a new component of W$_x$Ti$_{1-x}$O$_2$. It has been observed by other researchers that, At high wt % of tungsten doping (5 wt %), retarded the phase transformation (up to 900° C) [15, 7]. But this experiment is not found in any retardant at the current level of doping (3 wt % and 5 wt %) [16] is due to W ions in TiO$_2$ can either replace titanium ions to form W-O-Ti bonds or locate at interstitial sites.

The average particle size of pure TiO$_2$ is 37 nm and 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ powders are about 28 nm, 34 nm and 33 nm respectively. The average particle sizes was calculated using the full-width at half maximum measurement at 2θ of the maximum diffraction peaks using the following Debye-Scherrer’s formula,

$$D = \frac{K \lambda}{\beta \cos \theta}$$

In this equation, D is the crystallite size, K the Scherrer constant usually taken as 0.89, λ the wavelength of the X-ray radiation (0.15418nm for Cu Kα). Comparing the XRD patterns W-doped TiO$_2$, it appears that W loading does not influence the crystalline structure of TiO$_2$.

3.2. Field Emission Scanning Electron Microscope (FESEM)

![Figure 2](https://www.cirworld.com)
The FESEM images of pure and W-doped TiO$_2$ calcinated at 600° C are shown in Figure 2 (a-d). From the images, it can be confirmed that the average agglomerated particle size is nearly spherical and homogeneous particles. FESEM micrograph of W-doped TiO$_2$ nanoparticles shows that the surface morphology of the particles is nearly spherical with uniform sized particles and coherent together. However, the individual spherical particles are clearly seen due to the nano-clusters formed during the growth for 1 wt %, 3 wt % W-doped TiO$_2$. But it can be seen from the 5 wt % W-doped TiO$_2$ nanoparticles, nanorod type of nanoparticles were obtained. It can be seen that the average agglomerated particle size of 1 wt%, 3 wt % W-doped TiO$_2$ prepared by sol-gel method has no influence on the particle except 5 wt % W-doped TiO$_2$.

3.3. Energy Dispersive Analysis by X-Rays (EDXA)

Analysis of EDXA is used to analyze the chemical composition of the prepared material. It is clear that from the figure 3(a) TiO$_2$ is in pure form and free from any observable impurities. Figure 3 (b-d) also shows the EDXA of 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ samples, prepared by sol-gel method. EDXA shows only peaks of titanium, tungsten and oxygen elements. From the figure, it is clear that W-doped TiO$_2$ is free from impurities.

3.4. UV–Visible diffuse reflectance spectra (UV-Visible spectra)

![Graph showing UV-Vis diffuse reflectance spectra](image-url)

Figure 4 UV-Vis diffuse reflectance spectra of pure TiO$_2$, 1 wt %, 3 wt % and 5 wt % of W-doped TiO$_2$ calcinated at 600° C
UV-Visible spectroscopy measurement was performed to explore the absorbance and band gap of pure and 1 wt %, 3 wt% and 5 wt% W-doped TiO$_2$ nanoparticles. The optical absorbance spectra of pure and W-doped TiO$_2$ samples were recorded by a UV-Vis spectrophotometer in the range of 200-800 nm are shown in figure 4. From the figure, it indicates that blue shift was observed for pure TiO$_2$ compared with W-doped TiO$_2$ samples. The band edge absorption for pure TiO$_2$ blue shifted with tungsten doping indicates that widening of the optical band gap of pure TiO$_2$[17]. The obtained results can be confirms the band gap value of W-doped TiO$_2$ (4.47 eV, 3.96 eV and 3.41 eV) larger than the pure TiO$_2$ (2.86 eV).

But in the case of UV-Vis absorbance spectra of doped TiO$_2$, 1 wt %W-doped TiO$_2$ compared with other doped (3 wt% and 5 wt %) TiO$_2$ samples observed redshift when increasing the doping concentration of tungsten. It can be seen that from the figure 4. Optical absorption edges are shifted to higher wavelength region (red shift) with increasing dopant tungsten [18]. This redshift may create consistent of the tungsten onto the TiO$_2$ crystal lattice which creates impurity in the band gap [19] leading to reduction in the band gap energies [20]. The band gap of the samples can be determined by extrapolation of the absorption edge onto the x-axis and using the Planck’s equation

$$E_g = \frac{hC}{\lambda}$$

Where $E_g$ is the energy gap of pure and doped TiO$_2$ at absorption wavelength $\lambda$,

- $h$ is the Planck’s constant,
- $C$ is the velocity of light.

The calculated bandgap values for pure TiO$_2$ is 2.86 eV and W-doped nanoparticles values are 4.47 eV, 3.96 eV and 3.41 eV. From the bandgap values of W-doping, decreases when increase the doping of W content which shift to the longer wavelength. This may be attributed to the new electronic states and are introduced in the middle of the TiO$_2$ bandgap after doping the W atoms [18].

3.5 Photoluminescence study (PL)

![Figure 5 Photoluminescence spectrum of pure TiO$_2$, 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ samples calcinated at 600$^\circ$C](image)

Photoluminescence spectroscopy (PL) is a useful method to find the efficiency of trapping of charge carrier, transfer and circumstances of electron-hole pairs in semiconductor particles. The excitation wavelength of the pure TiO$_2$ is found to be at 325 nm. Similarly, for other excitation wavelength of W-doped TiO$_2$ nanoparticles is observed at 325 nm, 325 nm and 355 nm which indicates that trapped electrons and oxygen vacancies. Photoluminescence spectra of pure TiO$_2$ is greater than the PL intensity of 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$, which is due to a lower recombination rate of electrons and holes [21, 22] in the presence of light irradiation by the transition of energy levels between WO$_3$ orbital and TiO$_2$ orbital. It takesplace by photogenerated electrons and are transferred to WO$_3$ conduction band from TiO$_2$ conduction band and the holes accumulate in the TiO$_2$ valence band which results in photogenerated electrons and holes are separated. The figure 5 shows the photoluminescence spectrum of pure and 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles which show that the position of the peaks is almost similar except the 5 wt % W-doped TiO$_2$ nanoparticles and the photoluminescence intensity of pure TiO$_2$ is greater than the PL intensity of W-doped TiO$_2$. It indicates that the recombination of charge carriers is effectively reduced by the doping tungsten metal [23]. The peaks for the doped TiO$_2$ shift to red direction and also this shift of emission peak towards longer wavelengths further supports the lowering of the band gap of TiO$_2$ due to the tungsten doping treatment.
3.6 Fourier Transform Infrared Spectroscopy (FTIR)

![FTIR Spectra Diagram](image)

**Figure 6** Fourier transforms infrared spectra of (a) pure TiO$_2$, (b) 1 wt% W-doped TiO$_2$, (c) 3 wt% W-doped TiO$_2$, and (d) 5 wt% W-doped TiO$_2$ samples calcinated at 600° C.

IR spectra of pure and W-doped TiO$_2$ are shown in Figure 6. The absorption bands are shown in the region of 3300-3400 cm$^{-1}$ for vibrations of O-H and 1620-1630 cm$^{-1}$ in the region indicates the bending vibrations of the molecules. The little difference of band around 450 to 800 cm$^{-1}$ indicates the presence of Ti-O-W, compared to pure TiO$_2$ [24].

3.7 High Resolution Transmission Electron Microscopy (HRTEM)

![HRTEM Image](image)

**Figure 7** High Resolution Transmission Electron Microscopy image of W-doped TiO$_2$

The High Resolution TEM is used to study the morphology, distribution pattern, growth, of sample and it is also used to confirm the size of the particles. From the images clearly confirms that particles present in the W-doped TiO$_2$ powdered sample is nanosized particles. HRTEM images shows, all the particles are in irregular shapes and agglomerated. The figure (b) shows presence of fringes which are identified by lattice d-spacing values of 1.694 nm which correspond to the lattice spacing values (220) by using the standard JCPDS data.
3.8 Antimicrobial activity

(a) *Staphylococcus aureus* (Gram positive)

(b) *Bacillus subtilis* (Gram positive)

(c) *Escherichia coli* (Gram negative)

(d) *Pseudomonas aeruginosa* (Gram negative)

(e) *Candida albicans* (fungus)

(f) *Aspergillus niger* (fungus)
Figure 7(a-f) Inhibition zone of different microorganisms by media subjected to pure TiO$_2$, 1 wt% W-doped TiO$_2$, 3 wt% W-doped TiO$_2$, 5 wt% W-doped TiO$_2$ calcinated at 600°C
The antimicrobial activities of the samples are identified from the zone of inhibition. The diameter of zone of inhibition was measured and expressed in millimeter. The results showed that for all of the four samples at different concentration showed some good zone of inhibition against all the pathogens used. In addition, it could be observed that the doped samples are having much higher zone of inhibition compared to the pure TiO$_2$ sample. It shows that there is a significant effect of the tungsten induced antimicrobial activity for all the other doped samples. Also, the zone of inhibition increases as the concentration of the samples increases for almost all the samples against all the pathogens. However, none of the samples were able to reach the zone of inhibition of the standard Gentamicin except for the two samples which showed some higher zone of inhibition against the gram positive bacteria. The 1 wt % W-doped TiO$_2$ at higher concentration (100 mg) showed a higher zone of inhibition against the gram positive bacteria Staphylococcus aureus compared to all other pure and doped samples. However the 3 wt % and 5 wt % W-doped TiO$_2$ showed the zone of inhibition to that of the standard Gentamicin. Similar results were observed for the 3 wt % W-doped TiO$_2$ against the gram positive bacteria Bacillus subtilis. 1 wt % W-doped TiO$_2$ at higher concentration showed a higher zone of inhibition compared to the standard Gentamicin against the gram negative bacteria Pseudomonas aeruginosa. These results proved that the effect of the W doping on TiO$_2$ enhances the antimicrobial activity due to modified surface area, morphology and the reactivity of the samples.

CONCLUSION

Pure and 1 wt %, 3 wt % and 5 wt % W-doped TiO$_2$ nanoparticles were successfully synthesized by sol-gel method using hydroxylamine hydrochloride as a hydrolysis catalyst. The prepared nanoparticles are calcinated at 600° C for 5 h. According to the XRD pattern, 1 wt % W-doped TiO$_2$ was in an anatase crystalline form and it may be due to the smaller amount of W in TiO$_2$. It did not affect the crystalline structure. Whereas 3 wt % and 5 wt % W-doped TiO$_2$ shows rutile crystalline structure which confirms the phase transformation due to the tungsten doping. The average particle sizes of pure TiO$_2$ powder is approximately 37 nm. The average particle size of 1 wt %, 3 wt %, and 5 wt % W-TiO$_2$ powders are about 28 nm, 34 nm and 33 nm respectively. FESEM images of 1 wt % 3 wt % and 5 wt % W-doped TiO$_2$ which confirms the spherical with uniform sized particles and coherent together of all these nanoparticles. EDXA analysis shows that no impurites were present in the prepared pure and W-doped samples. UV-Vis absorbance spectra of doped TiO$_2$, 1 wt % W-doped TiO$_2$ compared with other doped 3 wt % and 5 wt % TiO$_2$ samples observed redshift when increasing the doping concentration of tungsten. The bandgap values of W-doping, decreases when increase in the doping of W content which shift to the longer wavelength. This may be attributed to the new electronic states and are introduced in the middle of the TiO$_2$ bandgap after doping the W atoms PL spectrum indicates that the recombination of charge carriers is effectively reduced by the doping tungsten metal. From the HRTEM images spherical shaped particles are shown. But all the spherical shaped nanoparticles are agglomerated together to form a larger particle present in the nanostructure domain and particles size are approximately 30 to 35. Broadening of peak of pure TiO$_2$ in FTIR spectra is due to increasing the W doping. Antibacterial activity of the W doping on TiO$_2$ enhances the antimicrobial activity due to modified surface area, morphology and the reactivity of the samples.

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