



Synthesis, Structural and Optical Characterization of TiO₂ Nanoparticles and its Assessment to Cytotoxicity Activity

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ABSTRACT

In this research work, TiO₂ nanoparticles was synthesised by solgel method using titanium (IV) isopropoxide and calcinated at 500 °C, 600 °C and 700 °C for five hours. The synthesised nanoparticles are characterized by using X-ray Diffraction studies (XRD), Field Emission Scanning Electron Microscopy (FESEM), Energy Dispersive X-ray Spectroscopy (EDXA), UV-Vis Diffuse Reflectance Spectroscopy (UV-Vis), Photoluminescence Spectra (PL), High Resolution Transmission Spectroscopy (HRTEM) and Fourier Transform Infra-Red Spectroscopy (FTIR). The synthesized nanoparticle of TiO₂ calcinated at 500 °C and 600 °C for 5 h shows anatase nature crystal structure and temperature increased to 700 °C, they undergo structural changes to rutile structure at the same duration of calcination. Average crystallite size can be determined by Debye-Scherrer formula for synthesized nanoparticles shows the particle sizes range from 30 nm to 47 nm. EDXA analysis confirms no impurities present in the sample. The band gap value decreases from 2.95 eV to 2.79 eV when calcination temperature increases in the UV-Visible analysis. The photoluminescence study of TiO₂ shows the direct recombination between electrons in the conduction band and holes in the valence band. FTIR analysis shows the bending and stretching mode of Ti-O-Ti. HRTEM of TiO₂ nanoparticles shows nearly spherical with approximately particles size of 35 nm to 50 nm. The cytotoxicity activity of TiO₂ nanoparticles calcinated at 500 °C, 600 °C and 700 °C has been explored.

Keywords: Cytotoxicity activity; HRTEM; SEM; Solgel method; TiO₂; XRD.

1. INTRODUCTION

TiO₂ is naturally occurring in oxide of Titanium. The high refractive index, non-toxic property, high chemical stability are some of the important properties of titanium dioxide and it has relatively low-cost of production (Kamat *et al.* 1999; Rammal *et al.* coworkers 2002; Rubing Zhang *et al.* 2004). Generally, Titanium dioxide has been used in paints as a pigment and filler (Krishnamurthy Prasad *et al.* 2010), ointments, toothpaste etc (Shuai Yuan *et al.* 2005; Juergen, Braun *et al.* 1992; Zallen and Moret 2006). Figure 1 shows the three forms TiO₂, rutile (Tetragonal, a=b=3.78Å, c=9.5Å), anatase (Tetragonal, a=b=3.78Å, c=9.5Å) and brookite (Rhombohedral, a=5.43Å, b=9.16Å, c=5.13Å) forms. All crystalline forms TiO₂ consist of [TiO₆]²⁻ octahedral structure. Anatase form of TiO₂ possess a wider optical band gap (3.2 eV), a higher fermi level, a smaller electron effective mass, and high mobility of charge carriers. But rutile phase is highly stable phase at high temperature (Mogyorosi *et al.* 2003). The edges and corners of this structure have different manners, but the overall stoichiometry as TiO₂ (Mark *et al.* 1983; Weast *et al.* 1984). Anatase phase has changed to rutile phase is usually occurs at 600 °C to 700 °C (Czanderna

et al. 1958; Yoganarasimhan and C.N.R.Rao 1962). This phase transition to rutile is non-reversible. Because, rutile form has greater thermodynamic stability compared to anatase phase (Reidy *et al.* 2006; Navrotsky and Kleppla, 1967).

A nanoparticle of titanium dioxide plays an important role in the fields of photo catalyst material for degradation of organic contaminants (Desong Wang *et al.* 2011) and as a self disinfecting material for surface coating in many applications. It is used in environmental purification, photo-electric chemical conversions in solar cells (Alam Khan *et al.* 2010), electronic devices, photo electrodes and gas sensors. The properties of Titanium dioxide such as photo induced super-hydrophobicity, nontoxicity and antifogging effect have been used in removing bacteria and harmful organic materials from water and air. The size, the crystalline structure, and the morphology of the TiO₂ nanoparticles strongly affect the performance of the TiO₂ (Kavan *et al.* 1996; Krol *et al.* 1997; Hadjipanayis *et al.* 1994;). So, there are variety of researchers focusing on the preparation of nanoparticles of TiO₂ using ethanol along with different acid and bases.

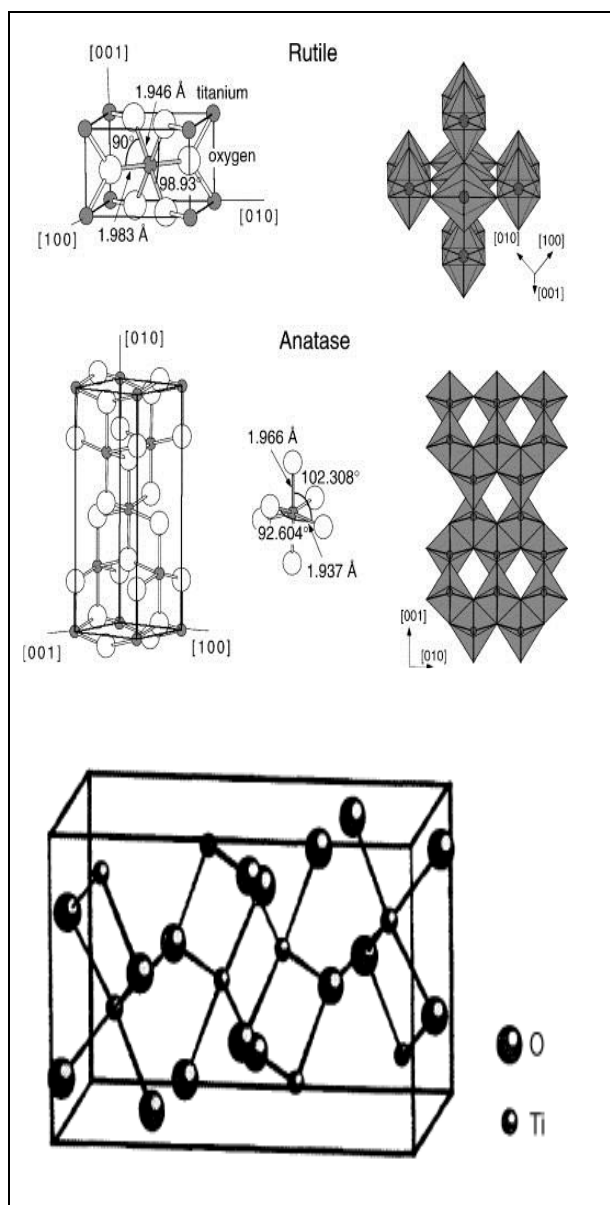


Fig. 1: Crystal structures of TiO₂ rutile, anatase and brookite phase.

In this work, an easy and new way of synthesizing nano particles of TiO₂ using titanium (IV) isopropoxide, ethanol and hydroxylamine hydrochloride has been discussed. Synthesised TiO₂ nanoparticles have been employed for the characterization such as X-ray Diffraction (XRD), Field Emission Scanning Electron Microscopy (FESEM), Energy Dispersive X-ray Spectroscopy (EDXA), Ultraviolet-Visible Diffuse Reflectance Spectroscopy (UV), Photoluminescence Analysis (PL), Fourier Transform Infra-Red Spectroscopy (FT-IR), and High Resolution Transmission Electron Microscopy (HRTEM).

2. EXPERIMENTAL METHODS

2.1 Material

Titanium isopropoxide (Sigma-Aldrich > 97 % pure), ethanol (Hayman (German) 99.99 % pure), hydroxylamine hydrochloride (Sigma-Aldrich, 99 % pure) were used as precursors and is used without any further purification. Doubly distilled water was used for the whole synthesis process.

2.2 Physicochemical Characterization

The X-ray diffraction pattern analysis for TiO₂ 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 TiO₂ nanoparticles samples 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 Jobnyvon model spectrophotometer and the alcohol is used 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 EDXA analysis was carried out to find the composition of TiO₂ samples by using the detector attached with the same instrument.

2.3 Synthesis of Pure TiO₂ Nanoparticles

Aqueous solution of titanium (IV) isopropoxide was used as starting material. Solgel is the most simple and sophisticated method proposed by Byun *et al.* 2000 among the various methods for producing nanoparticles. The sol was prepared by mixing titanium iso propoxide and ethanol having molar ratio 1:8 and dissolved in 1000 ml of deionised water at room temperature. Hydroxylamine hydrochloride 0.694 g was dissolved in 100 ml of deionised water and added gradually to the titanium iso propoxide sol. After 45 min of vigorous stirring, the suspension was centrifuged and precipitate obtained was washed with single step deionised water. After centrifugation, the precipitate was dried at 105° C till the samples were converted into dry powder. The prepared samples were calcinated at 500° C, 600° C and 700° C for five hours at a constant temperature of 2° C/min.

2.3 Cytotoxicity Activity of TiO₂

2.3.1 Methodology

2.3.1.1 Cell Line

The human embryonic kidney cell line (HEK 293) was obtained from National Centre for Cell Science (NCCS), Pune and grown in Eagles Minimum Essential Medium containing 10 % fetal bovine serum (FBS). The cells were maintained at 37° C, 5 % CO₂, 95 % air and 100 % relative humidity. Maintenance cultures were passaged weekly, and the culture medium was changed twice a week.

2.3.1.2 In vitro Cytotoxic Activities for TiO₂ Nanoparticles

The monolayer cells were detached with trypsin-ethylenediamine tetra acetic acid (EDTA) to make single cell suspensions and viable cells were counted using a hemocytometer and diluted with medium containing 5 % FBS to give final density of 1x10⁵ cells/ml. One hundred microlitres per well of cell suspension were seeded into 96-well plates at plating density of 10,000 cells/well and incubated to allow for cell attachment at 37° C, 5 % CO₂, 95 % air and 100 % relative humidity. After 24 h the cells were treated with serial concentrations of the test samples. They were initially dispersed in phosphate buffered saline (PBS) and an aliquot of the sample solution was diluted to twice the desired final maximum test concentration with serum free medium. Additional four serial dilutions were made to provide a total of five sample concentrations (0.25 µg, 2.5 µg, 25 µg, 50 µg, and 100 µg/ml). Aliquots of 100 µl of these different sample dilutions were added to the appropriate wells already containing 100 µl of medium, resulting in the required final sample concentrations. Following sample addition, the plates were incubated for an additional 48 h at 37° C, 5 % CO₂, 95 % air and 100 % relative humidity. The medium containing without samples were served as control and triplicate was maintained for all concentrations.

2.3.1.2 MTT Assay

3-[4,5-dimethylthiazol-2-yl]2,5-diphenyltetrazolium bromide (MTT) is a yellow water soluble tetrazolium salt. A mitochondrial enzyme in living cells, succinate-dehydrogenase, cleaves the tetrazolium ring, converting the MTT to an insoluble purple formazan. Therefore, the amount of formazan produced is directly proportional to the number of viable cells.

After 48 hours of incubation, 15 µl of MTT (5 mg/ml) in phosphate buffered saline (PBS) was added to each well and incubated at 37° C for 4 h. The medium with MTT was then flicked off and the formed formazan crystals were solubilised in 100 µl of DMSO and then measured the absorbance at 570 nm using micro plate reader.

- The percentage cell growth was then calculated with respect to control as follows

$$\% \text{ Cell growth} = [\text{A}] \text{ Test} / [\text{A}] \text{ control} \times 100$$

- The % cell inhibition was determined using the following formula.

$$\% \text{ Cell Inhibition} = 100 - \text{Abs}(\text{sample}) / \text{Abs}(\text{control}) \times 100.$$

3. RESULTS & DISCUSSION

3.1 X-Ray Diffraction Study (XRD)

The XRD patterns of TiO₂ samples calcinated at 500° C, 600° C and 700° C shown in the figure 2. It is clear that the TiO₂ present in its anatase form at 500° C, 600° C. Phase transformation to rutile form is identified in XRD results when calcination temperature increased to 700° C. The obtained 2θ values and corresponding (hkl) planes are 25.3° (101), 38° (004), 48° (200), 54° (105), 63° (204), 69.23° (116), 70.89° (220), 75.38° (215) respectively for TiO₂ nanoparticles calcinated at 500° C, 600° C (JCPDS Card No.21-1272). When calcination temperature increases to 700° C, shows the shifting of peaks at 27.3°, 36°, 41.2°, 54°, 69° (in figure 2(c) corresponding to the hkl values (110), (101), (111), (210), (112) (JCPDS Card No.88-1175) indicate that phase has been changed from anatase into rutile. The preferred peak for 2θ value 25.3° was observed with corresponding plane (101), which is the strongest peak among other peaks for all the TiO₂ nanoparticles calcinated at temperatures of 500° C, 600° C which is not available when the calcination temperature increased to 700° C, supports the changes in phase from anatase to rutile. The peaks of the graph are in good agreement with the literature report by Akarsu *et al.* (2006). The average size of the particles was calculated using Debye-Scherrer's formula.

$$\text{Crystallite size} = 0.9\lambda / \beta \cos \theta,$$

Where β is the full width at half-maximum (FWHM_{hkl}) of an hkl peak at 2 θ value,

θ is the half of the scattering angle.

The calculated particle size of TiO₂ nanoparticles is approximately about 30, 37 and 47nm respectively.

3.2 Field Emission Scanning Electron Microscope (FESEM)

FESEM images provide information about the surface morphology. From the images shown in the figure 3, confirms the grains of nanoparticles of TiO₂ appear to be nearly spherical and uniform sized particles and coherent together. However, the individual spherical

particles are not clearly seen due to the nano-clusters formed during the growth.

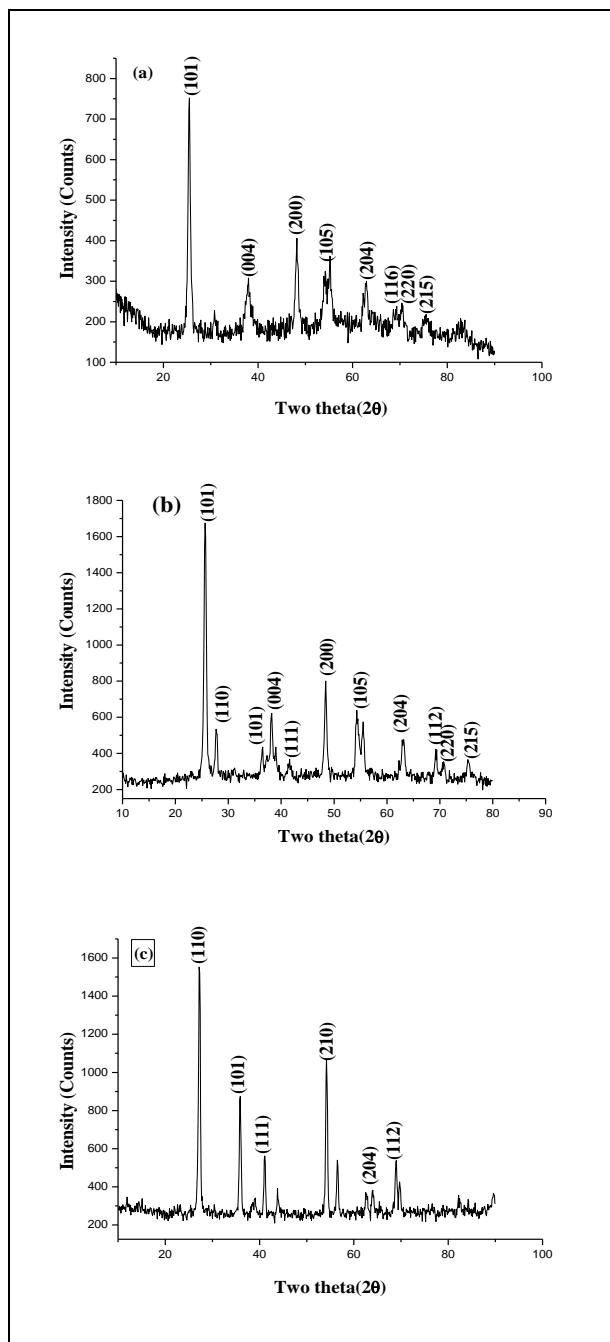


Fig. 2: XRD patterns of TiO₂ nanoparticles calcinated at (a) 500 °C (b) 600 °C (c) 700 °C.

3.3 Energy Dispersive Analysis by X-Rays (EDXA)

EDXA is used for chemical composition of a material. Figure 4 shows the EDXA of TiO₂ nanoparticles which contains only peaks of titanium and oxygen. From the figure 4 (a, b and c), it is clear that TiO₂ is free from impurities.

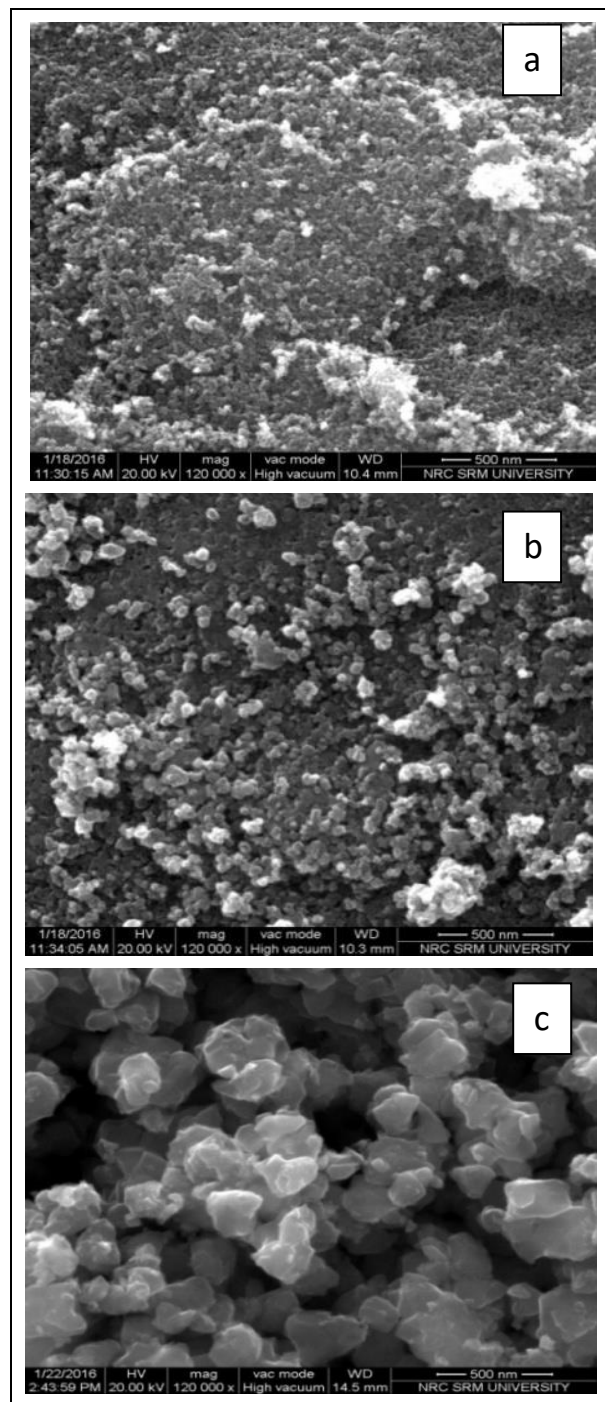


Fig. 3: FESEM images of TiO₂ nanoparticles calcinated at (a) 500 °C (b) 600 °C (c) 700 °C.

3.4 UV–Visible Diffuse Reflectance Spectroscopy (UV-Vis)

Fig. 5 shows the UV-Visible spectra of the TiO₂ at different temperatures of 500°C, 600 °C and 700 °C. It shows the absorbance can be decreased at wavelength after 350 nm for intrinsic band gap absorbance of TiO₂ due to electron transition from the valence band to the conduction band. Red shift was observed for TiO₂

nanoparticles in the absorbance edge is due to this change in surface morphology, particles size (Senthil *et al.* 2014) and also variation in the absorbance may be due to variation in the annealing temperature (Elangovan *et al.* 2015). The band gap is determined by analysing the absorption of incident light by the nanomaterials. The band gap energy can be determined by extrapolation of the absorption edge onto the x-axis and by using the Planck's equation

$$E_g = hc/\lambda$$

Where, E_g is the energy gap of pure TiO_2 at absorption wavelength λ , h is the Planck's constant, C is the velocity of light.

The calculated band gap value for TiO_2 was calculated at 500°C , 600°C and 700°C are 2.95 eV, 2.86 eV and 2.79 eV respectively. The band gap energy decreased with the increase of calcinated temperature. At higher calcinated temperature 700°C shows lowest band gap (2.79) is due to rutile phase is formed which has a bigger particle size (Swapan *et al.* 2010).

3.5 Photoluminescence Study (PL)

The photoluminescence spectrum was recorded for TiO_2 nanoparticles which are shown in figure 6. The emission of first peak in the photoluminescence spectra obtained at three calcination temperature of TiO_2 nanoparticles between 360 nm to 420 nm corresponds to the direct recombination between electrons in the conduction band and holes in the valence band (Liqiang *et al.* 2004). The presence of emission peaks in the visible region is due to the presence of defect levels below the conduction band. Similar peak has been observed in earlier work on TiO_2 nano particles (Vijayalakshmi and Rajendran 2012).

The presence of broad peak in the visible region of PL spectrum indicates the presence of defect levels below the conduction band and the electronic transition takes place by defect levels such as oxygen vacancies in the band gap (Zhao *et al.* 2007).

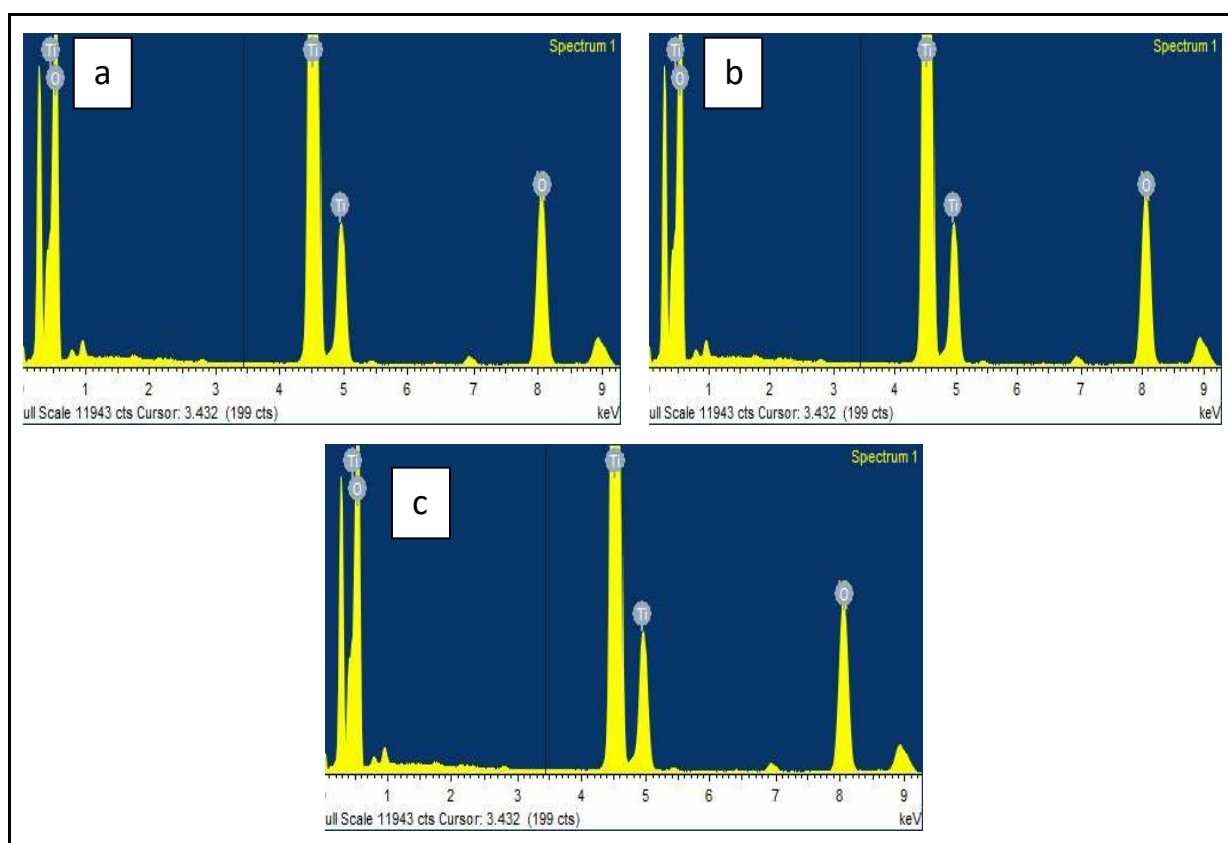


Fig. 4: EDXA pattern of TiO_2 nanoparticles calcinated at (a) 500°C (b) 600°C (c) 700°C .

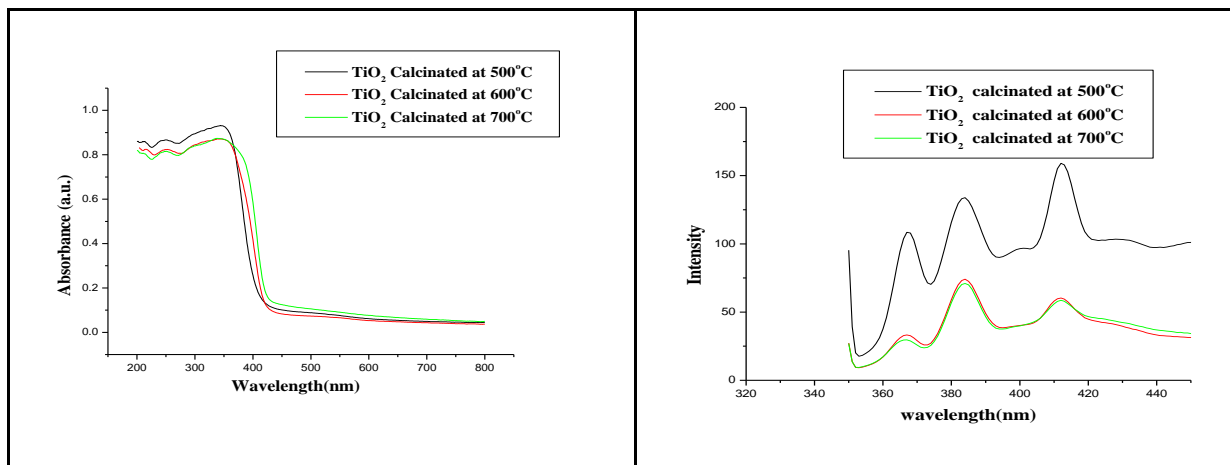


Fig. 5: UV-Vis diffuse reflectance spectra of TiO₂ nanoparticles calcinated at 500 °C, 600 °C and 700 °C.

Fig. 6: Photoluminescence spectra of TiO₂ nanoparticles calcinated at 500 °C, 600 °C and 700 °C.

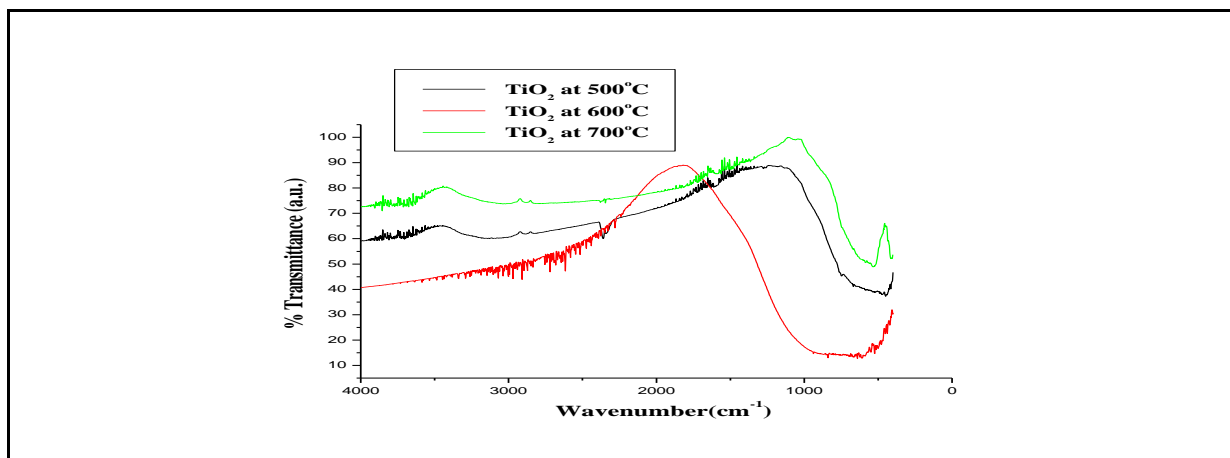


Fig. 7: FTIR spectra of TiO₂ nanoparticles calcinated at 500 °C, 600 °C, and 700 °C.

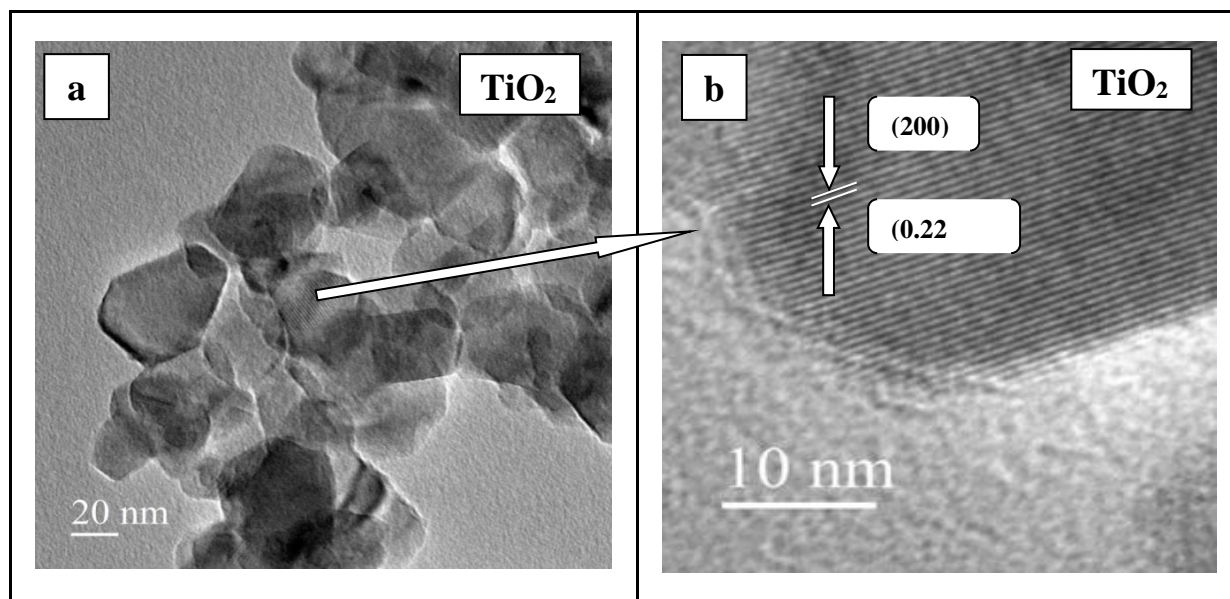


Fig. 8: High Resolution Transmission Electron Microscope image of TiO₂ nanoparticles prepared by sol-gel method.

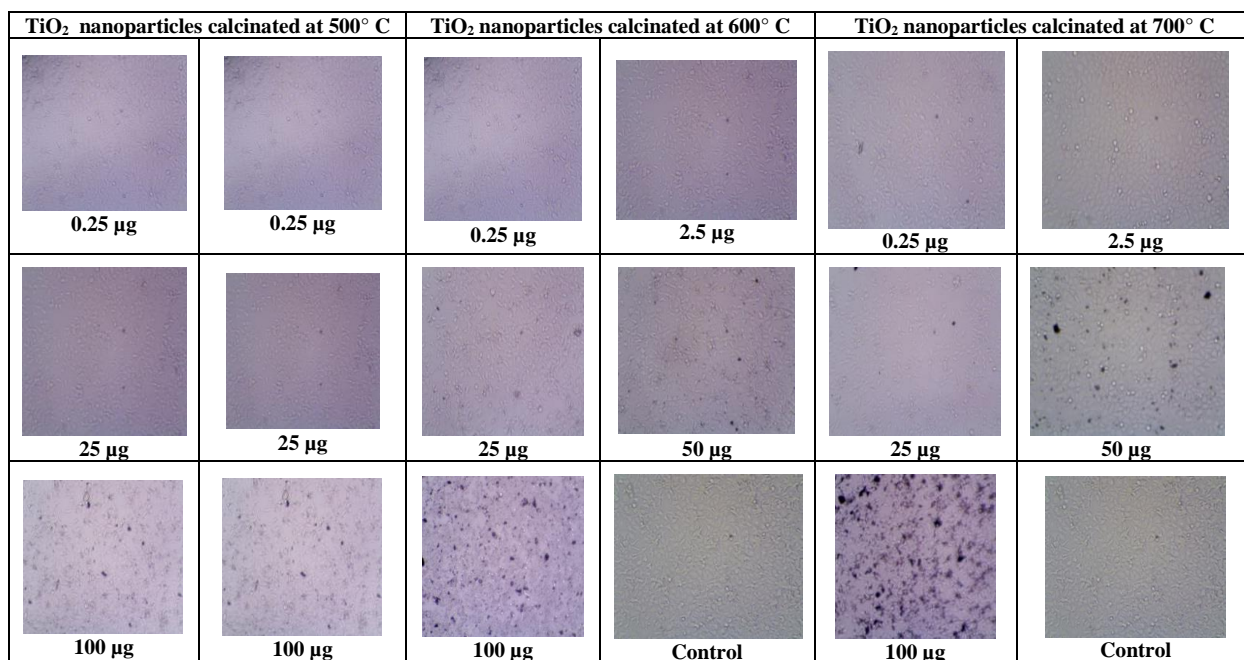


Fig. 9: Percentage inhibition of TiO₂ nanoparticles calcinated at 500 °C, 600 °C, 700 °C.

Table 1. Percentage of inhibition and IC50 value for different calcinated TiO₂ nanoparticles at various concentrations.

Percentage inhibition of TiO ₂ nanoparticles calcinated at 500 °C			Percentage inhibition of TiO ₂ nanoparticles calcinated at 600 °C			Percentage inhibition of TiO ₂ nanoparticles calcinated at 700 °C		
Conc (µg/ml)	% cell inhibition	IC50 value (µg)	Conc (µg/ml)	% cell inhibition	IC50 value (µg)	Conc (µg/ml)	% cell inhibition	IC50 value (µg)
0.25	0	>100	0.25	3.14	>100	0.25	1.33	>100
2.5	1.2		2.5	2.31		2.5	4.54	
25	1.57		25	3.21		25	3.34	
50	3.59		50	4.06		50	5.02	
100	7.67		100	13.4		100	12.03	
Control	0.337		Control	0.337		Control	0.337	

3.7 High Resolution Transmission Electron Microscope (HRTEM)

The HRTEM was used to confirm the distribution pattern, growth of prepared nanoparticles and also confirm the size of nano particles by directly measured from the ruler of the images. In general, TiO₂ nanoparticles have strong tendency to agglomerates to form a larger particle. The image 8 shows that most of the TiO₂ nanoparticles aggregates together and also spherical or nearly spherical and expand shaped particles with approximately uniform size of 35 nm to 50 nm. This

exhibit lattice fringes which was used to calculate d-spacing and found to be 0.222 nm. This d-spacing value has been compared with standard JCPDS data corresponds to the plane (200) of anatase.

3.8 Cytotoxicity Activity of TiO₂

The synthesised TiO₂ nanoparticles assessed cytotoxic effects after 48 h exposure and cisplatin was used as standard Mosmann *et al.* 1983; Monks *et al.* 1991;. Figure 8 shows the percentage inhibition of TiO₂ nanoparticles calcinated at 500 °C, 600 °C, and 700 °C. The IC50 value is found greater than 100 µg/ml against

human embryonic kidney cell line (HEK 293) for tested TiO₂ nanoparticles. This result indicates TiO₂ nanoparticles did not show any cytotoxicity activity at the tested concentrations. Table 1 shows the % of inhibition and IC₅₀ value for different calcinated TiO₂ nanoparticles at various concentrations. The tested TiO₂ nanoparticles low or did not show cytotoxicity activity against human cell line.

4. CONCLUSION

Titanium dioxide nanoparticles have been successfully synthesized by sol-gel method with using of hydroxylamine hydrochloride as a hydrolysis catalyst. From the XRD analysis, anatase phase was identified for TiO₂ nanoparticles calcinated at 500° C and 600° C for 5 h and converting it into rutile form when it is calcinated at 700° C for 5 h. TiO₂ nano particle size determined from Debye-Scherrer's formula is 30 nm, 37 nm and 47 nm. EDXA analysis shows that no impurities are present in the prepared TiO₂ samples. UV-Visible spectrum of TiO₂ nanoparticles at three temperatures shows red shift in the absorbance edge and is due to the change in surface morphology, particles size. The band gaps of nanoTiO₂ measured by graphical method are 2.95 eV, 2.86 eV and 2.79 eV. Photoluminescence study indicates that the strong emission peaks obtained in 383 nm, 385 nm and 385 nm region confirm direct recombination between electrons in the conduction band and holes in the valence band. High Resolution Transmission Electron Microscope of TiO₂ nanoparticles shows aggregate together and also spherical or nearly spherical with approximately uniform size of 35 to 50 nm. FTIR spectrum analysis shows the peaks at 433 cm⁻¹, 514 cm⁻¹ to 700 cm⁻¹ shows bending and stretching mode of Ti-O-Ti. The cytotoxicity activity results of TiO₂ at different temperatures show very small or no cytotoxicity activity.

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CONFLICTS OF INTEREST

The authors declare that there is no conflict of interest.

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