

ORIGINAL RESEARCH ARTICLE

Influence of illumination on eco-friendly titanium dioxide/epoxy nanocomposite-based triboelectric nanogenerators

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Abstract

Triboelectric nanogenerators (TENGs) based on titanium dioxide (TiO₂)/epoxy nanocomposites have attracted increasing interest as self-powered systems capable of harvesting mechanical friction and light energy for sustainable electricity generation. In this study, a photoinduced TENG was developed using TiO₂/epoxy nanocomposites with a polyethylene terephthalate charge collector. Scanning electron microscopy and X-ray diffraction analyses were conducted to evaluate the structural and morphological properties of the nanocomposite, revealing the presence of TiO₂ nanoparticles with an anatase crystal structure. Ultraviolet-visible spectroscopy was employed to investigate the optical properties of the nanocomposite, identifying an optical band gap energy of 3.52 eV. In addition, Fourier transform infrared spectroscopy was used to determine chemical bonds and functional groups. Epoxy serves as the matrix material, binding the TiO₂ nanoparticles and providing mechanical integrity to the composite film. It ensures uniform dispersion of TiO₂ nanoparticles and offers a flexible yet robust substrate essential for the repeated contact-separation cycles of TENG operation. The TiO₂/epoxy nanocomposite, used as the electrode for the TENG device, was created by incorporating TiO₂ nanoparticles into the epoxy resin. Without ultraviolet illumination, the TENG devices generated output voltages of 25, 45, 50, and 55 V at TiO₂ concentrations of 0, 2, 3, and 4 wt%, respectively. Under ultraviolet illumination, the corresponding voltages increased to 65, 75, 77, and 80 V. The highest voltage of 80 V was obtained for the device containing 4 wt% TiO₂. These findings provide valuable insights into the structural, morphological, optical, and chemical characteristics of the nanocomposite, supporting the development of effective and sustainable power sources.

Keywords: Triboelectric nanogenerator; Photoinduced triboelectric nanogenerator; Titanium dioxide/epoxy nanocomposite; Solution blending technique; Power sources

1. Introduction

Mechanical energy, such as motion or friction, can be converted into electrical energy using an energy-harvesting device known as a triboelectric nanogenerator (TENG). The term “triboelectric effect” refers to the movement of charges between two materials when they come into contact and subsequently separate.^{1–3} Utilizing nanomaterials in the design of TENGs can improve their performance. Among these materials, titanium dioxide (TiO₂) is widely used in many applications due to its unique properties.^{4–6} When integrated as a functional layer within the TENG structure, TiO₂-based composites can further improve energy-conversion efficiency.^{7,8} Electron-hole pairs are produced when light interacts with TiO₂ nanoparticles; these charge carriers can then facilitate the triboelectric charging process if friction is applied to the composite material.^{9–12} The interaction between light and friction enables the simultaneous harvesting of energy from both sources. The TiO₂/epoxy nanocomposite-based TENG offers several benefits for producing sustainable energy, as it can capture ambient light energy, making it suitable for interior settings or places with low light levels.^{13–15} Parimalam *et al.*¹⁶ investigated the organic-inorganic coatings utilizing bisphenol A diglycidyl ether, incorporating silica and other fillers via solution casting. TiO₂ and silicon dioxide nanoparticles enhanced the chemical resistance over 21 days, while the epoxy/silica/TiO₂/zinc oxide/latex nanocoating demonstrated outstanding heat resistance (efficiency of 90–99%) across various substrates. It is also possible to effectively transform the frictional energy produced by diverse mechanical actions into electrical energy. This capacity supports a wide range of applications, including harvesting energy from mechanical vibrations or human body movements.¹⁷

Through the provision of a self-powered system, nanocomposite-based TENGs can achieve energy self-sufficiency.^{18–22} Although TiO₂/epoxy nanocomposites show strong potential for TENG applications, several challenges remain, including enhancing the durability and long-term stability of the materials, fine-tuning the nanocomposite composition, and improving the overall energy conversion efficiency.^{23–25} Related studies demonstrate the multifunctional advantages of composite fillers. For example, epoxy/silica/TiO₂/zinc oxide coating demonstrated the highest bacterial reduction (>95%) after 4 h.²⁶ The combined filler effect enhances stress, viscosity, and torque compared to individual fillers, suggesting potential industrial and medical applications in reducing microbial growth. TENGs employ TiO₂/epoxy nanocomposites to harness light and friction energy, providing a sustainable approach to electricity generation. Renewable reinforcements based on hydroxyapatite

have been reported to improve the performance of biocomposite materials, and treated carbon black particles can increase the glass transition temperature and thermal decomposition resistance by 7°C.²⁷ Ongoing research continues to advance the performance and usability of these nanocomposite-based TENGs.^{1,7,28}

The ability of TiO₂/epoxy nanocomposite-based TENGs to harvest energy simultaneously from light and mechanical friction makes them particularly promising. While frictional motion generates triboelectric charging, light energy absorbed by the TiO₂ nanoparticles contributes to the total charge generation.^{29–31} These TENGs can generate electrical energy for low-power electronic devices without the need for external power sources by effectively harnessing ambient light sources, such as indoor illumination.^{32–34} A previous study examined epoxy-based composite coatings containing 20 wt% colloidal nanosilica and varying amounts of micro-sized TiO₂ (5, 10, 15 wt%).³⁵ Fourier transform infrared (FTIR) spectroscopy confirmed the presence of functional groups, indicating a good interaction. Scanning electron microscopy (SEM) showed well-dispersed TiO₂ at lower loading. A higher percentage of TiO₂ resulted in enhanced absorption of ultraviolet (UV) light. In comparison to pure epoxy, nanocomposites showed an enhanced oxygen transport rate. To improve the total energy conversion efficiency of TENGs, current studies focus on optimizing electrode layout, device design, and the properties of nanocomposites. In addition, scaling up the production of TiO₂/epoxy nanocomposites and TENG devices in a cost-effective manner is receiving increasing attention to enable broader deployment.^{36–38}

This work aims to develop polycondensed epoxy nanocomposites incorporating nitrogen-doped titanium dioxide (n-TiO₂) nanoparticles. The innovative aspect of this study lies in the creation of a TiO₂/epoxy nanocomposite deposited on a polyethylene terephthalate (PET) substrate for a photoinduced TENG.³² This design expands the potential applications of nanogenerators and enables efficient harvesting of environmental energy. The epoxy matrix, enhanced with TiO₂ nanoparticles, was combined with the PET substrate, endowing the TENG with improved performance and unique functional properties. The device performance was examined at various TiO₂ concentrations (0 wt%, 2 wt%, 3 wt%, and 4 wt%), revealing that optimizing the TiO₂ content increased the voltage output, leading to a maximum voltage of 80 V.

2. Materials and methods

2.1. Materials

The materials used in this study included Kapton tape, double-sided copper tape, PET sheet (Techinstro, MH,

India), and double-sided tape. The epoxy matrix was based on the ED-20 resin. Diaminopentane (GOST 10587-84; Rearus LLC, Russia) was used as a hardener in the preparation of the TENG devices, with TiO₂ particles incorporated at 0, 2, 3, or 4 wt%. In the synthesis of the nanocomposite films, 4,4'-diaminodiphenylmethane (DDM; Sigma-Aldrich, United States; molar mass 198 g/mol, purity 97%) was used as the curing agent. The synthesized TiO₂ nanoparticles consisted of 75% anatase and 25% rutile with an average particle diameter of 46 nm. Dimethyldichlorosilane (HIMMED Trading House LLC, Russia) was employed as the surface treatment agent.

2.2. Synthesis procedure

Nanocomposites were prepared by sequentially curing ED-20 diene-epoxy resin at 90°C and 160°C in the presence of an aromatic hardener, with titanium(IV) dioxide nanoparticles synthesized via a plasma-chemical approach incorporated into the resin. Ultrasonic dispersion was employed to distribute the nanoparticles throughout the polymer matrix using a SONOREX Digital 10p ultrasonic bath (Bandelin, Sigma-Aldrich, United States). The ED-20 epoxy resin had a molecular weight of 385 and an epoxy group content of 22.6 wt%. DDM, produced by microwave-assisted synthesis, was used as the hardener. According to the process outlined by Bukichev *et al.*,³⁹ films of n-TiO₂/ED-20 epoxy resin can be prepared with TiO₂ concentrations ranging from 0.2 to 5 wt%. In this work, films containing 0, 2, 3, and 4 wt% TiO₂ were fabricated. To ensure correct stoichiometry, the TiO₂ nanoparticles were added to a mixture of ED-20 and DDM in an equimolar ratio of epoxy and amine groups, and dispersed for 20 min in an ultrasonic bath at 35 kHz. The mixture was treated with a toluene-based solution of dimethyldichlorosilane, which served as an anti-adhesive, and then cast into glass molds. A temperature-

controlled oven was used to cure the samples using a stepped temperature regime of 90°C and 160°C each (3 h at each temperature), which, according to calorimetric data, ensured 95–97% curing. The resulting films had a thickness of 80–100 µm.

2.3. Device fabrication and experimental setup

The fabrication procedure for the TENG device is illustrated in Figure 1A, and the experimental setup is depicted in Figure 1B. The TENG structure was composed of PET sheets, copper tape (2 mm thick), Kapton tape (1 mm thick), double-sided foam tape, and an epoxy resin (ED20) cured with diaminopentane, with TiO₂ nanoparticles added at 0, 2, 3, or 4 wt%. Two copper tapes were attached to the PET sheets, serving as electrodes, with one electrode secured in place using Kapton tape. The surface of this Kapton tape was roughened using abrasive sandpaper to enhance the triboelectric effect. The second electrode was coated with the TiO₂/epoxy nanocomposite, where the TiO₂ heterostructure can strengthen the internal electric fields at the nanoparticle surfaces and increase the charge transfer rate between nanoparticles. The two contact layers were separated by double-sided foam tape spacers, each with an effective surface area of 24 cm². A Keithley 6517B electrometer (Tektronix/Keithley, USA) was used to measure variations in voltage.

2.4. Characterization methods

2.4.1. Scanning electron microscopy

Scanning electron microscopy was used to examine the surface morphology of the nanocomposite films. Before imaging, samples were sputter-coated with a thin gold layer to improve surface conductivity. SEM observations were performed using a JSM 6490 LV scanning electron microscope (JEOL, Japan) operated at an accelerating voltage of 20 kV. Micrographs were acquired at multiple magnifications.

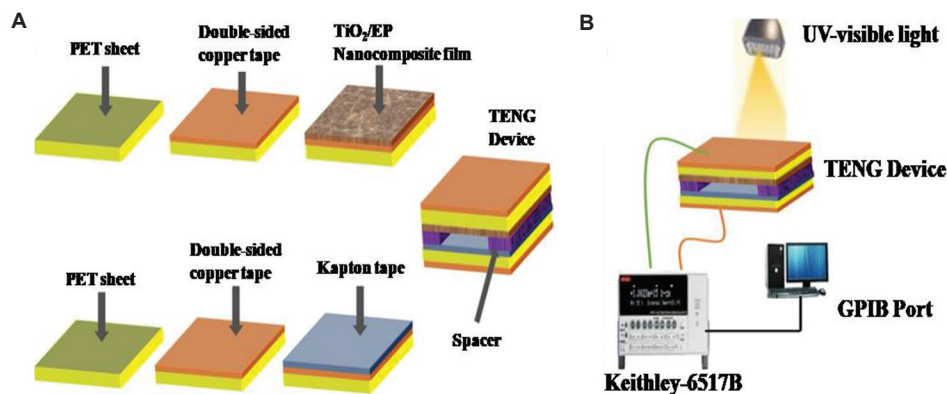


Figure 1. Fabrication of a TENG. (A) Fabrication steps of the TENG. (B) Experimental setup for testing the effect of ultraviolet-visible light. Abbreviations: EP: Epoxy; GPIB: General Purpose Interface Bus; PET: Polyethylene terephthalate; TENG: Triboelectric nanogenerator.

2.4.2. Energy-dispersive X-ray spectroscopy (EDS)

EDS was used to determine the elemental composition of the films. EDS analysis was performed using an INCAx-act EDS system (Model 51-1385-007, Oxford Instruments, Japan) integrated with the SEM, and spectra were processed using the INCA software package. Elemental spectra and point analyses (and/or elemental mapping, if performed) were collected to confirm the presence and distribution of constituent elements.

2.4.3. X-ray diffraction (XRD)

XRD patterns were collected using a Bruker D8 Advance diffractometer (Bruker, Germany) with Cu K α radiation ($\lambda = 1.5406 \text{ \AA}$). Data were acquired over a 2θ range of $10\text{--}80^\circ$ with a step size of 0.02° and a scan rate of 2° min^{-1} . Phase identification was performed by comparison with standard reference data (JCPDS/ICDD).

2.4.4. Ultraviolet–visible (UV–Vis) spectroscopy

UV–Vis absorbance spectra of the TiO₂/epoxy films were recorded using a Thermo Scientific Evolution 201 UV–Vis spectrophotometer (Japan) in the wavelength range of 200–800 nm with a data interval of 1 nm. Measurements were performed on film samples (thickness 80–100 μm), using distilled water as the reference/baseline.

2.4.5. FTIR spectroscopy

FTIR spectra were acquired using a Thermo Scientific Nicolet 6700 FTIR spectrometer (USA). Spectra were recorded in the range of 200–800 nm at a resolution of 0.09 cm^{-1} with 36 scans per sample. Measurements were performed for TiO₂ nanoparticles and epoxy films containing 0, 2, 3, and 4 wt% TiO₂.

3. Results and discussion

3.1. Surface morphology

TiO₂ nanoparticles were dispersed throughout the epoxy matrix volume, both as individual particles and aggregates, as determined by SEM measurements (Figure 2). The displayed histograms of the particle size distribution show that even at a relatively low concentration of nanofillers (0.5 wt%), the average diameter of particles in the original n-TiO₂ powder increased from 40 to 83 nm upon incorporation into the TiO₂/epoxy nanocomposite. Figure 2A shows the SEM image of pure epoxy resin, whereas Figure 2B–D shows the SEM images of 2 wt% n-TiO₂, 3 wt% n-TiO₂, and 4 wt% n-TiO₂ incorporated in the matrix of the epoxy resin samples. These SEM images clearly demonstrate the incorporation of TiO₂ nanoparticles into the epoxy matrix. Figure 2E and F depict the SEM images of the pure n-TiO₂ nanoparticles, which are spherical and range in size from

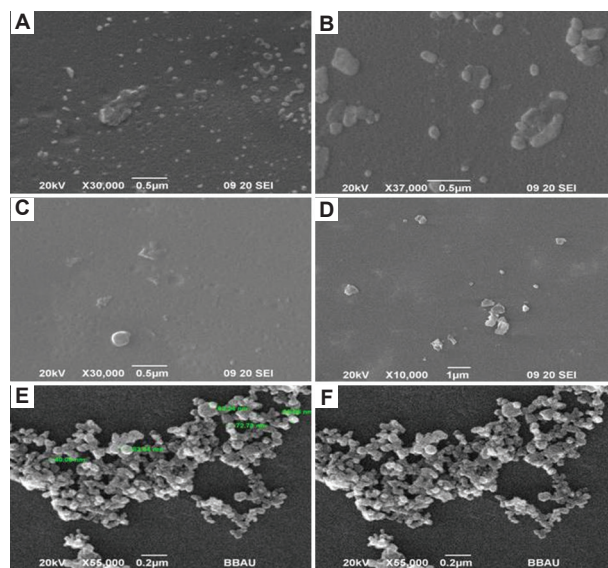


Figure 2. Scanning electron microscopy images of TiO₂/epoxy nanocomposite films. Images of films incorporated with (A) 0 wt%, (B) 2 wt%, (C) 3 wt%, and (D) 4 wt% n-TiO₂. Scale bars: 0.5 μm and 1 μm ; magnifications: 10K \times , 30K \times , and 37K \times . (E and F) Pure n-TiO₂ nanoparticles. Scale bar: 0.2 μm ; magnification: 55K \times . Abbreviations: n-TiO₂: Nitrogen-doped titanium dioxide; TiO₂: Titanium dioxide.

10 to 100 nm. During curing, nanoparticle aggregation can lead to an increase in the effective size of the n-TiO₂ particles.

3.2. Energy dispersive spectroscopy analysis

Figure 3 presents the EDS spectra of epoxy resin with and without n-TiO₂, confirming the presence and weight percentage of titanium and oxygen. EDS was used alongside field-emission SEM to examine the chemical composition of the material. Figure 3A shows the EDS spectrum of 0 wt% TiO₂ epoxy, indicating the absence of titanium peaks. Figure 3B–D shows the EDS spectra of epoxy resin with 2 wt%, 3 wt%, and 4 wt% TiO₂, respectively, indicating the presence of titanium and carbon. According to the EDS analysis, the prepared TiO₂ exhibited distinct titanium and oxygen peaks while showing fewer contaminants, such as chlorine.

3.3. Structural analysis

Figure 4 depicts the X-ray diffraction pattern of the created TiO₂/epoxy composite film. According to Chen *et al.*,⁴⁰ the absence of spurious diffraction peaks indicates good crystallographic purity. The experimental data agree with the JCPDS card no. 00-015-0875, confirming the anatase phase of TiO₂. The 2θ at a peak of 25.4° confirms the TiO₂ anatase structure,¹³ supported by additional strong peaks at 30.04° and 48° . No extraneous diffraction peaks were

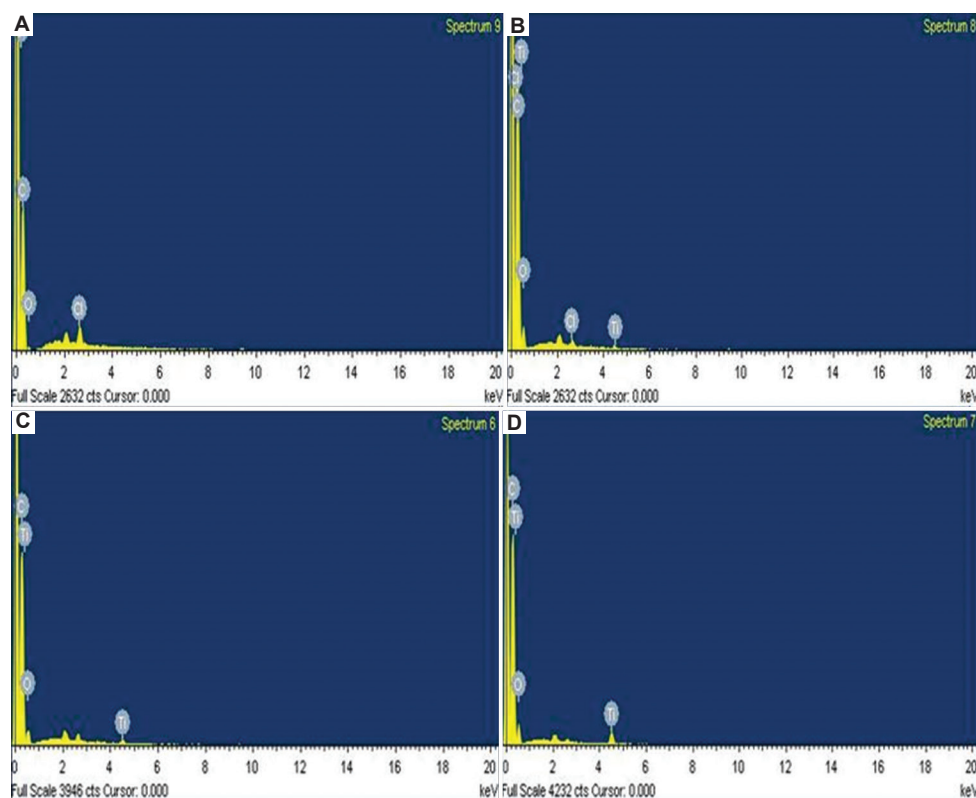


Figure 3. Energy dispersive spectroscopy spectrum and elemental compositions. Spectrum of epoxy with (A) 0 wt%, (B) 2 wt%, (C) 3 wt%, and (D) 4 wt% titanium dioxide.

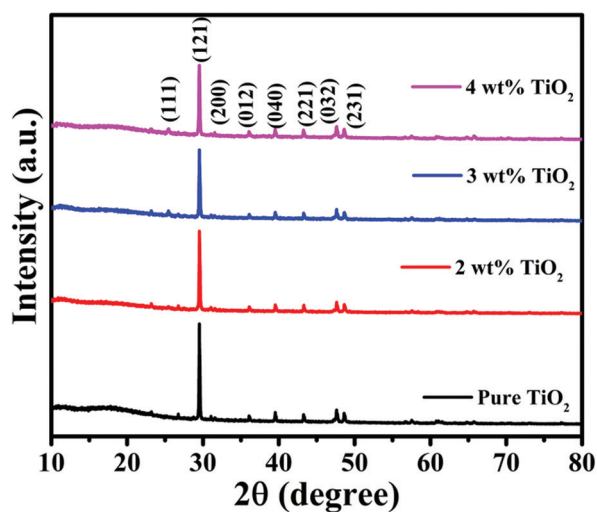


Figure 4. X-ray diffraction pattern of pure TiO₂ and epoxy composites of 2 wt%, 3 wt%, and 4 wt% TiO₂, respectively
Abbreviation: TiO₂: Titanium dioxide.

observed, indicating phase purity. The broad nature of the diffraction peaks suggests an extremely small crystallite size, while the overall peak intensities reflect the crystalline nature of the material.

3.4. UV-Vis absorption analysis

Figure 5A depicts the UV-Vis (300–1,200 nm) absorbance spectrum for the TiO₂/epoxy nanocomposite. The main absorption edge was observed at around 380 nm in the UV region. Tauc's plot was used to analyze the optical energy band gap value for the TiO₂/epoxy nanocomposite (inset in Figure 5B). The optical band gap (E_g) was calculated using Tauc's relation, as given by Equation 1:

$$\alpha = \frac{\alpha_0 (h\nu - E_g)^n}{h\nu} \quad (1)$$

where α_0 is a characteristic parameter, α is the absorption coefficient, h is Planck's constant, and n is the power factor. Depending on the type of transition that occurred, the value of n may change. Since the transition in this nanocomposite material was direct, n is presumed to be 1/2. For the TiO₂/epoxy nanocomposite, the calculated value of E_g was 3.52 eV.

3.5. FTIR spectroscopy analysis

FTIR analysis of TiO₂ nanoparticles confirmed the formation of high-purity products. FTIR spectra (Figure 6) of these nanoparticles primarily display peaks associated

with TiO₂. A peak at 499.9 cm⁻¹ is attributed to Ti–O stretching vibrations. The final product appeared to have Ti–O bonds and lacked OH or peroxy groups, as indicated by the FTIR spectrum. The produced TiO₂ nanoparticles' infrared spectra, which can be used to identify chemical bonds and functional groups in a product, were in the range of 400–4,000 cm⁻¹. Low-wavelength peaks at 585.6, 586.4, 566.9, and 565.1 cm⁻¹ were assigned to C–I stretching vibrations. The highest peak observed for the TiO₂/epoxy nanocomposite in 0, 2, 3, and 4 wt% TiO₂ was 3,420.5, 3,403.3, 3,387.9, and 3,384.2 cm⁻¹, respectively, due to O–H stretching. In the low-energy region (<1,000 cm⁻¹), broad absorption bands associated with Ti–O stretching were

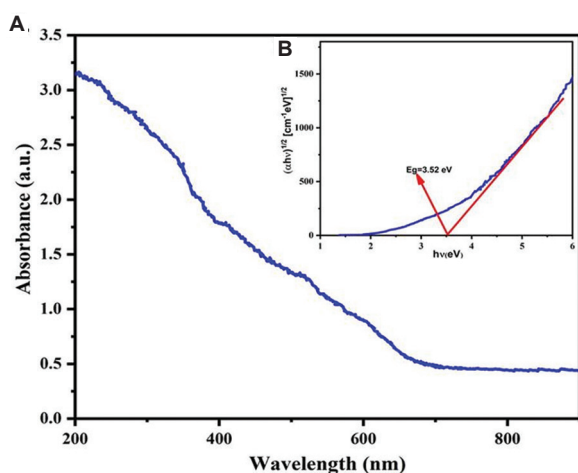


Figure 5. (A) Ultraviolet–visible spectrum of titanium dioxide/epoxy nanocomposite and (B) the Tauc's plot in the inset

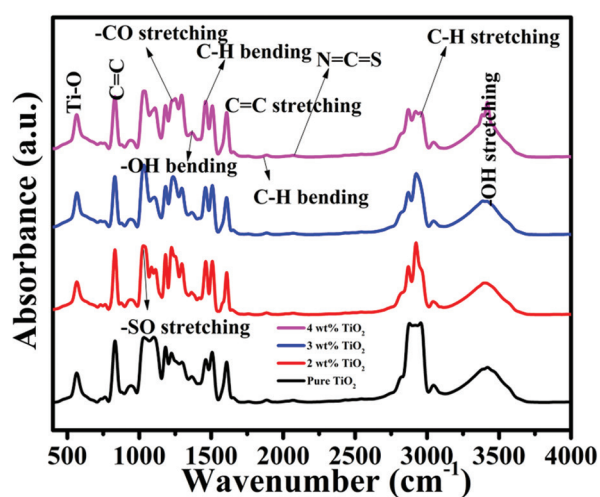


Figure 6. Fourier transform infrared analysis of TiO₂/epoxy nanocomposite of pure TiO₂ and epoxy of 2 wt%, 3 wt%, and 4 wt% TiO₂, respectively
Abbreviation: TiO₂: Titanium dioxide.

observed. For the TiO₂/epoxy nanocomposite containing 0, 2, 3, and 4 wt% TiO₂, characteristic peaks appeared near 586 and 566 cm⁻¹.

3.6. Experimental analysis for TENG devices

A TENG converts mechanical energy into electrical energy through the combined action of the triboelectric effect and electrostatic induction. It typically comprises two triboelectric layers made from materials with differing electron affinities. When these materials come into contact and then separate, one surface tends to gain electrons while the other loses them. Nylon, polytetrafluoroethylene, and polydimethylsiloxane are a few examples of frequently used triboelectric materials.³¹ As seen in Figure 7A, electrodes on each side of the triboelectric layers collect the electrical charges generated during contact or separation. These electrodes are usually made of conductive materials, such as conductive polymers or metal foils. One electrode is attached to each triboelectric surface, enabling efficient charge collection.⁷ The layers are often supported by a flexible substrate, such as a polymer film or a flexible printed circuit board, which provides both flexibility and structural stability. The electrical output of TENG may be utilized to charge batteries or power other devices. The electrodes are linked to an external load, such as a resistor or energy storage device, completing the electrical circuit and enabling practical energy utilization. To enable the creation of mechanical energy, the TENG device may incorporate various mechanical structures. The overall fabricated TENG device is shown in Figure 7A, while the photographic images of the fabricated TENG and

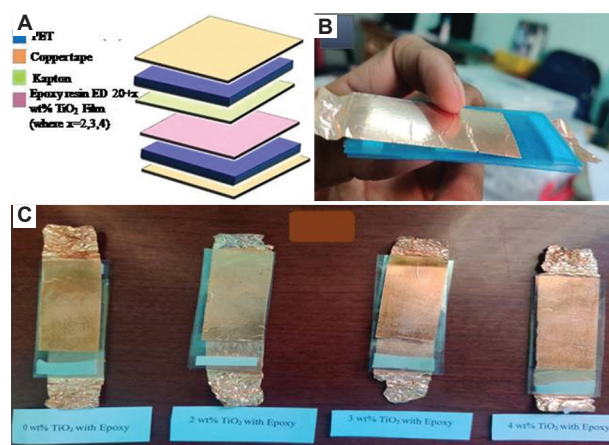


Figure 7. Device fabrication. (A) Constituent layers in epoxy resin TiO₂-based TENGs. (B) Image of fabricated TENG. (C) Images of fabricated TENG with TiO₂/epoxy nanocomposite films incorporated with 0 wt%, 2 wt%, 3 wt%, and 4 wt% n-TiO₂, respectively.
Abbreviations: n-TiO₂: Nitrogen-doped titanium dioxide; TENG: Triboelectric nanogenerator; TiO₂: Titanium dioxide.

TiO₂/epoxy nanocomposite-based devices with varying TiO₂ concentrations are presented in Figures 7B and C, respectively.

The optimized output voltage characteristics of the TENG device without UV light irradiance are presented in Figure 8. The TENG device produced a voltage of 25 V using 0 wt% TiO₂ nanoparticles, as shown in Figure 8A. Under triboelectric operation, this device exhibits the smallest electrical potential difference between its electrodes. Nonetheless, even at this lower voltage, the device is capable of generating a measurable charge flow. The TENG device containing 2 wt% TiO₂ nanoparticles generated a higher voltage of 45 V compared to the sample containing 0% TiO₂ (Figure 8B). This implies that the inclusion of TiO₂ nanoparticles increased the electrical potential difference. In comparison to the two prior samples, the TENG device with 3 wt% TiO₂ nanoparticles produced a voltage of 50 V (Figure 8C). This suggests that increasing the concentration of TiO₂ nanoparticles leads to a larger electrical potential difference and enhances charge transfer within the system. The TENG device with 4 wt% TiO₂ nanoparticles generated a voltage of 55 V (Figure 8D), the highest among all the samples. This indicates that higher TiO₂ loading is associated with a larger electrical potential difference and a greater charge flow. Although it is slightly lower than the earlier samples, it remains in the same ballpark, indicating a sizable flow of electrical charges. Voltages usually increase as TiO₂ nanoparticle concentration rises, indicating an increase in electrical potential differences.^{40,41} TiO₂ nanoparticles possess unique properties that enhance the performance of nanogenerators. They are wide-bandgap semiconductor materials that effectively transform light energy into electrical energy. Their inclusion expands the surface area,

creating a network of connections that improves electrical conductivity, electron mobility, and charge transfer. When materials with various electron affinities come into contact, TiO₂ nanoparticles enhance the triboelectric effect by acting as electron acceptors and donors. This makes charge transfer easier, which raises the output voltage. In addition, TiO₂ nanoparticles strengthen the epoxy matrix, improving its flexibility, toughness, and resistance to mechanical stress. TiO₂/epoxy nanocomposite-based photoinduced TENGs exhibit improved performance and durable power production capabilities due to these combined features.

The optimized output voltage characteristics of the TENG device under UV light irradiance are presented in Figure 9. Table 1 compares the output voltage characteristics of the TENG device with and without UV light, showing a drastic increase in the output voltage of the TENG devices. When exposed to UV light, the TENG device with 0 wt% TiO₂ nanoparticles produced a higher voltage of 65 V (Figure 9A) compared to the corresponding device operated without UV illumination, as presented in Figure 9B. The electrical potential difference in this sample appears to be amplified by UV light irradiation. Under UV light irradiation, the TENG device with 2 wt% TiO₂ nanoparticles generated a voltage of 75 V (Figure 9B). This implies that the electrical potential difference was higher than it was in the UV-free sample. With 3 wt% TiO₂ nanoparticles and UV light irradiation, the TENG device generates a voltage of 77 V (Figure 9C). The voltage in this sample was slightly higher than previously reported ones.⁴² When subjected to UV light, the TENG device with 4 wt% TiO₂ nanoparticles produced the highest voltage of 80 V among all the samples (Figure 9D). This indicates that the largest electrical potential difference

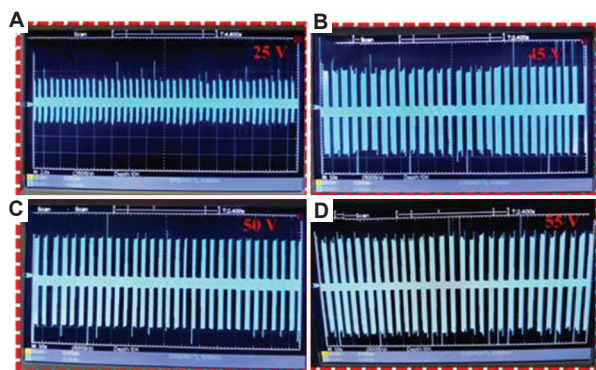


Figure 8. Characteristics of TiO₂/epoxy nanocomposite-based triboelectric nanogenerators without ultraviolet light exposure. Output voltage for (A) 0 wt%, (B) 2 wt%, (C) 3 wt%, and (D) 4 wt% TiO₂, respectively.

Abbreviation: TiO₂: Titanium dioxide.

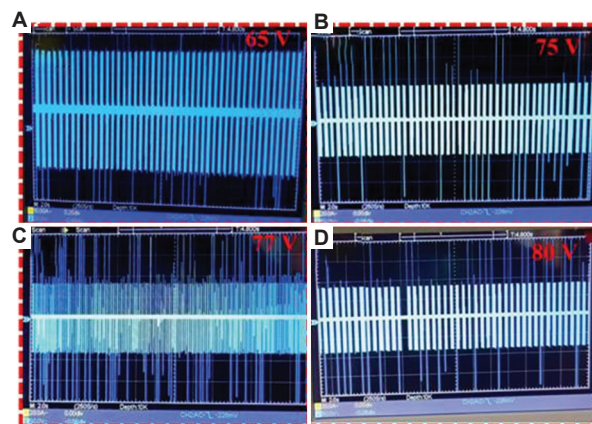


Figure 9. Characteristics of TiO₂/epoxy nanocomposite-based triboelectric nanogenerators with ultraviolet light exposure. Output voltage for (A) 0 wt%, (B) 2 wt%, (C) 3 wt%, and (D) 4 wt% TiO₂, respectively.

Abbreviation: TiO₂: Titanium dioxide.

occurred when UV light irradiation and TiO₂ nanoparticle concentration are combined. UV light illumination led to an increased generation of electrical charges. The TENG device performed better when UV light irradiance was added together with different TiO₂ nanoparticle concentrations. For all compositions, UV illumination increased the voltage output. The current output remained relatively constant for most samples, except for the device containing 4 wt% TiO₂, which exhibited a substantial increase in current under UV light. These findings suggest that UV light, particularly when combined with precise concentrations of TiO₂ nanoparticles, can enhance the generation of electrical energy in TENG devices. The triboelectric voltage produced in the samples is greatly influenced by UV radiation. When UV light irradiates TiO₂ nanoparticles, it activates the photovoltaic effect. Due to its wide bandgap, TiO₂ can absorb photons in the UV region, leading to the generation of electron-hole pairs. Upon photon absorption, electrons are excited to higher energy levels, leaving behind positively charged holes. This charge separation creates an internal electric potential, resulting in a built-in voltage across the material.⁴³ This built-in voltage influences the total triboelectric voltage measured. The TiO₂ nanoparticles can be used to produce electron-hole pairs, which can enhance the triboelectric effect. The triboelectric effect occurs when two materials with differing electron affinities, such as the opposing substance and the TiO₂ nanoparticles, come into contact. The opposing material may absorb the excited electrons from the TiO₂ nanoparticles, thereby enhancing the charge density and triboelectric voltage.

3.7. Mechanism of TENGs

The mechanism can be understood as follows: when two dielectric materials, Kapton and a TiO₂/epoxy nanocomposite, come into contact, electrification occurs, resulting in the formation of opposite static charges on the inner surfaces of the triboelectric layers. Upon separation of these layers, a potential difference is created, causing electrons to flow under short-circuit conditions.⁴⁴ A charge imbalance develops between two materials that have

different electron affinities when they come into contact and then separate, creating an electric potential.⁴⁵ TiO₂ nanoparticles are well known for their superior electrical and optical characteristics, which make them suitable for use in energy-related applications. The PET substrate serves as the foundation material for the nanocomposite, providing mechanical flexibility, toughness, and insulation. In this instance, inherent motion or deformation of the device provides the mechanical energy needed for the triboelectric effect. The photoinduced mechanism indicates that light can activate or enhance the performance of the nanogenerator.²³ TiO₂ is a semiconductor with photoactive characteristics, meaning that when exposed to light, it can produce electron-hole pairs. The photoactive characteristics of TiO₂ nanoparticles play a role in the reaction of TENG to UV radiation. Because TiO₂ nanoparticles in the anatase phase are photocatalytic, they produce electron-hole pairs when exposed to UV light. Figure 10 illustrates the role of UV light on the enhanced characteristics of TiO₂/epoxy nanocomposite-based TENG. On the TiO₂/epoxy composite, these electron-hole pairs improve charge separation and increase surface charge density. The TENG produces more voltage as a result of the enhanced triboelectric effect caused by this greater charge density. Higher concentrations of TiO₂ produce higher voltage output as they enhance the material's capacity to capture and transmit charges more effectively when exposed to UV light. By enhancing charge transfer during contact and separation, these charge carriers can support the triboelectric effect and increase the total electrical output of the nanogenerator.⁴⁶ Moreover, the incorporation of TiO₂ nanoparticles into the epoxy matrix can also influence the surface morphology and interfacial properties of the nanocomposite. This can lead to improved contact electrification and charge transfer efficiency, further enhancing the voltage output of the TENG device. During the deformation process, the TiO₂/epoxy nanocomposite layer and the PET substrate

Table 1. Output voltage of the triboelectric nanogenerator devices with and without UV light

Sample wt% of titanium dioxide	Without UV light	With UV light
0	25 V	65 V
2	45 V	75 V
3	50 V	77 V
4	55 V	80 V

Abbreviation: UV: Ultraviolet

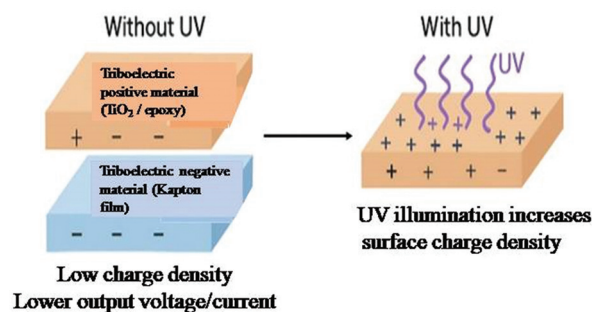


Figure 10. Impact of UV light on the charge density of TiO₂/epoxy nanocomposite-based triboelectric nanogenerator
Abbreviations: TiO₂: Titanium dioxide; UV: Ultraviolet.

come into contact and eventually separate due to the subsequent application of pressure. The triboelectric effect is produced by the contact and separation process. The TiO₂/epoxy nanocomposite layer and the PET substrate transmit electrons due to the triboelectric phenomenon, and between the two materials, this charge transfer produces an electric potential. Through the use of proper electrodes and electronics that are attached to the device, the created electric potential can be harnessed and collected.³⁴ By applying UV light, TiO₂ nanoparticles in the nanocomposite layer can absorb photons and produce additional charge carriers. The triboelectric effect is enhanced by this photoinduced action, resulting in increased electrical output.^{47,48} The triboelectric interaction between the TiO₂/epoxy nanocomposite layer and the PET substrate can be exploited in a self-powered, photoinduced TENG to convert mechanical energy into electrical energy.⁴⁹ The charge transfer process during contact and separation can therefore be facilitated by these photoexcited charge carriers, increasing the total electrical output of TENG. The operating range of TiO₂-based TENGs can be extended by UV light, which acts as a supplementary energy source.^{50,51} The TENG device can capture energy from both mechanical motion and UV illumination in its surroundings, utilizing visible or UV light, enabling it to function autonomously even with modest mechanical stimulation. It is a prospective contender for several applications, including wearable technology, self-powered sensors, and energy harvesting from ambient light sources due to its dual-mode energy-collecting capacity. Comparison with previously reported devices indicates that most earlier studies achieved higher output voltages⁴⁸⁻⁵⁰ than those obtained here, with the exception of Alam *et al.*⁵² However, to the best of our knowledge, none of these studies systematically examined the effect of UV illumination on TENG performance, which is the key innovation of the present work.

4. Conclusion

This study used a TiO₂/epoxy nanocomposite-blended PET substrate to create a photoinduced TENG. Structural and morphological investigations revealed the presence of TiO₂ nanoparticles with anatase crystal structure in the nanocomposite. The performance of the TENG device was examined under the effect of UV light, and the findings indicated that varying TiO₂ concentrations had an impact on the output voltage. The device containing 4 wt% TiO₂ produced the highest voltage output of 80 V under UV illumination. For the device without TiO₂ (0 wt%), the voltage increased from 25 V (without UV) to 65 V under UV illumination. These results provide important new insights into the structural, morphological, optical, and chemical

properties of the blended TiO₂/epoxy nanocomposite PET substrate. The development of such self-powered, photoinduced TENGs may significantly advance sustainable power-source technologies. Environmental stimuli, such as UV light, can thus be exploited to generate energy, paving the way for self-sufficient and portable energy-generating systems. Future studies can focus on maximizing the TiO₂ concentration and exploring other materials to enhance TENG performance.

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Conflict of interest

The authors declare that they have no competing interests

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Not applicable.

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Availability of data

Data will be available from the corresponding authors upon reasonable request.

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