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Effect of Sodium Arsenic on the Improvement of TiO₂/Dye as Photosensitizers in Dye-Sensitized Solar Cells (DSSC)

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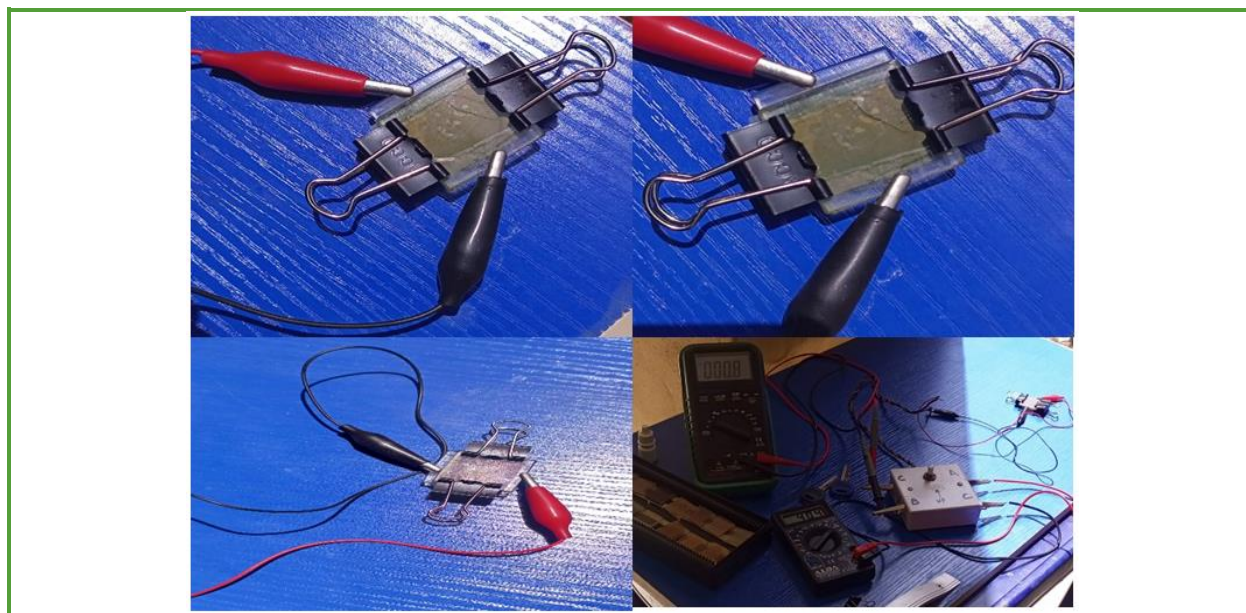
TiO₂/dye
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 Grain size
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ABSTRACT

The research presents the synthesis and fabrication of dye-sensitized solar cells (DSSCs) on the influence of sodium arsenic on the enhancement of TiO₂/dye as photosensitizers, where *Hibiscus sabdariffa* (roselle) and *Vernonia amygdalina* (bitter leaf) were used as a source of the chlorophyll, sodium arsenic (NaAs) material of different concentration (0.1-0.4 mol), was synthesized as a layer on top of TiO₂. The surface morphology study of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye revealed that the micrograph is usually defined with the granular shape of nanotubes. The grain size of TiO₂/NaAs_{0.1} is not too large and delineated by an immense sum of aggregated nanoparticles. The cells structure is polycrystalline with a most outstanding peak at 2 theta angles of 26.73° and 51.84° corresponding to hkl index numbers (111) and (202). The films have a very high absorbance from the plot, and the absorbance of the films increases as the dye molecules vary. The high absorbance of the films shows that the DSSCs will be a good material for photovoltaic applications. The fill factor of the films is 0.54, 1.24, 1.23, and 0.99 respectively while the conversion efficiency of 0.86%, 4.48%, 3.44%, and 1.81% was recorded.

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Graphical Abstract



Introduction

The world today has faced the major challenge of obtaining clean energy. Fossil fuels have been the primary energy source because they are cheap and widely accessible [1-3]. When supplies run out, it is recommended to explore and use environmentally friendly and cost-effective alternative energy sources. The sun is an eco-friendly and affordable energy source. Each hour, the sun shines on the Earth, providing more than enough energy to meet the world's energy demand for a whole year. One of several methods for converting solar energy into electrical energy is the solar cell (SC). Silicon (Si) is one of the semiconductor materials used in fabricating SCs. Si-based SC has been reported to have high efficiency ranging from 13-25%, but the major challenge limiting the use of Si as a basic material is its high cost and toxicity [4]. There are different types of solar cells, including; DSSC, organic solar cells, perovskite solar cells, and inorganic solar cells [5-7]. Among them, DSSC has outstanding properties. It provides an

economically and technically viable alternative compared to conventional solar cells. A DSSC is composed of organic dye as an absorbing layer, photoelectrode, counter electrode, and electrolyte [8]. It produces power simply by the entire lighting of organic dye in an electrochemical unit [9].

Its performance is determined by the photoelectrode component and organic dye employed. The first DSSC was initially made with a ZnO photoelectrode [10]. It had low conversion efficiency because the dye has two layers and was just able to absorb light incident on it by 1%. The DSSC performance was improved by using a porous photoelectrode, which led to the introduction of the Grätzel cell in 1991 [11], which had a 7% improved performance.

Various semiconductor materials such as gallium arsenide (GaAs), thin oxide (SnO₂), indium phosphate (InP), zinc oxide (ZnO), niobium pentoxide (Nb₂O₅), cadmium sulfide (CdS), etc. have been employed as DSSC photoelectrode [12-14], but they show low photoconversion efficiency. Ever since 14%

efficiency was achieved in DSSC using TiO_2 has been regarded as the preferred DSSC photoelectrode material [15]. TiO_2 has other unique features such as chemical inertness, eco-friendliness, abundance, and porosity. The wide band gap of TiO_2 makes the dye molecules absorb the light spectrum over a wide range [16-18].

However, the dye molecules used in DSSCs are either natural or synthetic dyes. The natural dye (e.g., chlorophyll) obtained from flowers and plants has been employed as a DSSCs photosensitizer [19, 20]. In this research, the chlorophyll obtained from *Vernonia amygdalina* (bitter leaf) and *Hibiscus sabdariffa* (roselle) would be the dye (photosensitizer) of interest. Compared to the synthetic dye molecules, the natural dye (chlorophyll) has better advantages such as easy extraction, availability, and low cost [21, 22]. The carboxylic group found in chlorophyll increases its photoconversion efficiency [23]. Its absorption peaks range from 420-660 nm, the fill factor about 47 %, and 1.96 mA and 0.585 mV I_{sc} and V_{oc} , respectively [24]. The use of chlorophyll is limited because it can easily deteriorate when exposed to sunlight. The stability of chlorophyll is mostly improved by the formation of the complex compound via interaction with metals [25, 26]. Various mordants have been used to improve the stability of chlorophyll including; alum, chalk, metals, and salts (e.g., FeCl_2 , NaAs) [27-33]. Sodium arsenic (NaAs) is preferred and is the subject of this work. The development of complex compounds between the NaAs and chlorophyll is predicted to not only enhance heat and water resistance but also boost brightness and color retention [29]. Few researches have been carried out on improving the stability of chlorophyll using a metal mordant. This would be the first time sodium arsenic would be used to improve chlorophyll performance, stability, and coloration. H.

Darmokoesoemo *et al.* [30] studied the improved properties of the composite compound of Ni (II)-chlorophyll dye sensitizer in DSSC. Ni (II)-chlorophyll has a wide range of absorbance between the wavelengths of 295 to 665 nm. The ligand-metal bond is seen in the oscillation Ni-O at 455.2 cm^{-1} wave number. The open circuit voltage of Ni (II)-chlorophyll dye sensitizer is 0.15 V, I_{sc} is 3.00 mAcm^{-2} , and the efficiency is 0.20 %. H. Darmokoesoemo *et al.* [24] also studied the improved properties of Fe (III)-chlorophyll dye sensitizer. Classic mulberry powder (CMP) serves as the source of chlorophyll pigment. Fe (III)-chlorophyll dye sensitizer showed better performance compared to Ni (II)-chlorophyll. The Fe (III)-chlorophyll dye sensitizer has the efficiency, J_{sc} and V_{oc} of the DSSCs with Fe-chlorophyll dyes are 1.3 %, 4 mAcm^{-2} , and 0.18 V respectively. Z. Arifin *et al.* [31] studied the stability and performance of Fe-chlorophyll dye sensitizer in DSSC. The chlorophyll was obtained from papaya leaf. They reported the Fe-chlorophyll dye sensitizer to have improved stability by 2 times and efficiency by 2.5 times compared to the natural chlorophyll. The efficiency, J_{sc} , and V_{oc} of the DSSCs with Fe-chlorophyll dyes are 0.16 %, 0.62 mAcm^{-2} , and 500 mV, respectively.

Herein, we present the influence of sodium arsenic on the enhancement of dye as a photosensitizer. The chlorophyll pigments were obtained from *Vernonia amygdalina* (bitter leaf) and *Hibiscus sabdariffa* (roselle). The performance of chlorophyll photosensitizer obtained from bitter leaf, roselle, and roselle-bitter leaf combined was studied.

Experimental

Materials and Methods

Ethanol, sodium arsenite, and Trixon-100 were obtained from Sigma Aldrich, p25 Degussa TiO_2 powder (99%), and FTO glass substrate

was obtained from the Institute of Chemical Education (ICE).

Experimental procedure

Hibiscus sabdariffa (roselle) and *Vernonia amygdalina* (bitter leaf) were used as a source of chlorophyll. The leaves (samples) were weighed and crushed in a blender, and ethanol was used to dissolve the particles to harvest the chlorophyll at room temperature. The extracted dyes in the beakers were filtered using the original Whatman filter paper; the filtered chlorophyll was kept in an air-tight container to avoid degradation before use. The substrate fluorine-doped tin oxide for the fabrication of the cell (FTO) went through a series of purifications before being used.

The electrode material which is titanium dioxide (TiO_2) was deposited using the Doctor Blade technique. The TiO_2 paste was prepared using 15 ml of methanol to 6 g of TiO_2 powder in a mortar with constant grinding with a pestle for proper separation of the TiO_2 particles and nonionic Triton-100 was used as a surfactant to get a good paste.

The substrate was taped to get a dimension of 2.25-2.35 cm for the deposition of paste. The

samples were annealed at 350 °C for 105 mins. The Sodium arsenic (NaAs) material of different concentrations (0.1-0.4 mol) was synthesized as a layer on top of TiO_2 using a spin coating machine for 5 sec. The spinning speed was 500 rpm and the films were dried in the oven for 15 minutes and the samples were taken for sanitization. The TiO_2 deposited samples were buried into the extracted dyes (bitter leaf dye, roselle dye, and the mixture of bitter leaf dye and roselle dye) for 24 hours and the films were properly placed for further analysis. Graphite was used to prepare the counter electrodes on fluorine-doped tin oxide conductive glass. The extracted dyes (bitter leaf dye, roselle dye, and the mixture of bitter leaf dye and roselle dye) were analyzed for their optical, structural, and elemental studies. The electrolyte solution obtained from the Institute for Chemical Education (ICE) was dropped on the cell surface and the counter electrode with graphite was applied in contact directly with the TiO_2/NaAs substrate. The 0.2 cm strip of the film without cells was noted for the crocodile clip space by joining them together at the corner. The full cell was taken for analysis to obtain the I/V characterization (see [Figure 1](#)).

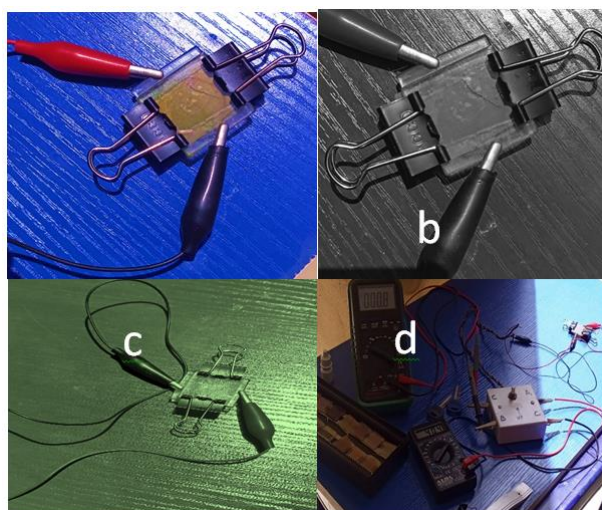


Figure 1. (a) Complete cell of $\text{TiO}_2/\text{NaAs}_{0.1}$, (b) $\text{TiO}_2/\text{NaAs}_{0.2}$ / bitter leaf dye, (c) $\text{TiO}_2/\text{NaAs}_{0.3}$ / roselle dye, and (d) $\text{TiO}_2/\text{NaAs}_{0.4}$ /the mixture of bitter leaf dye and roselle dye

Results and Discussion

The surface morphology study of $TiO_2/NaAs_{0.1}$, $TiO_2/NaAs_{0.2}$ / bitter leaf dye, $TiO_2/NaAs_{0.3}$ / roselle dye, and $TiO_2/NaAs_{0.4}$ /the mixture of bitter leaf dye and roselle dye

Figure 2a-d demonstrates the surface morphology study of $TiO_2/NaAs_{0.1}$, $TiO_2/NaAs_{0.2}$ / bitter leaf dye, $TiO_2/NaAs_{0.3}$ / roselle dye, and $TiO_2/NaAs_{0.4}$ /the mixture of bitter leaf dye and roselle dye. Figure 2a illustrates that the micrograph is usually defined with the granular shape of nanotubes. The crystallite size of $TiO_2/NaAs_{0.1}$ is not too large and delineated by an immense sum of aggregated nanoparticles $TiO_2/NaAs_{0.2}$ / bitter leaf dye as shown in Figure 2b has smaller crystallite-sized particles of which granular shapes are more gathered. The particles clustering is evident in films synthesized with the chlorophyll of bitter leaf dye [24-31]. In Figure 2c, the distributions of the particles are random as can be seen in the non-uniformity of

the particle morphology. Moreover, firm binding power and connections with wide contact areas compel the firm bonding power of the TiO_2 individual particles. The nanoparticles in Figure 2a and d suggest a better crystallite structure than the ones in Figure 5b and c.

Elemental study of $TiO_2/NaAs_{0.1}$, $TiO_2/NaAs_{0.2}$ / bitter leaf dye, $TiO_2/NaAs_{0.3}$ / roselle dye, and $TiO_2/NaAs_{0.4}$ /the mixture of bitter leaf dye and roselle dye

The spectra showing the elemental composition of $TiO_2/NaAs_{0.1}$, $TiO_2/NaAs_{0.2}$ / bitter leaf dye, $TiO_2/NaAs_{0.3}$ / roselle dye, and $TiO_2/NaAs_{0.4}$ /the mixture of bitter leaf dye and roselle dye is depicted in Figure 3. The spectrum shows the energy dispersive X-ray spectroscopy (EDX) of the samples. The EDX peaks show that titanium, sodium, arsenic, and oxygen are the major constituents of the particle composition, as shown in the plot.

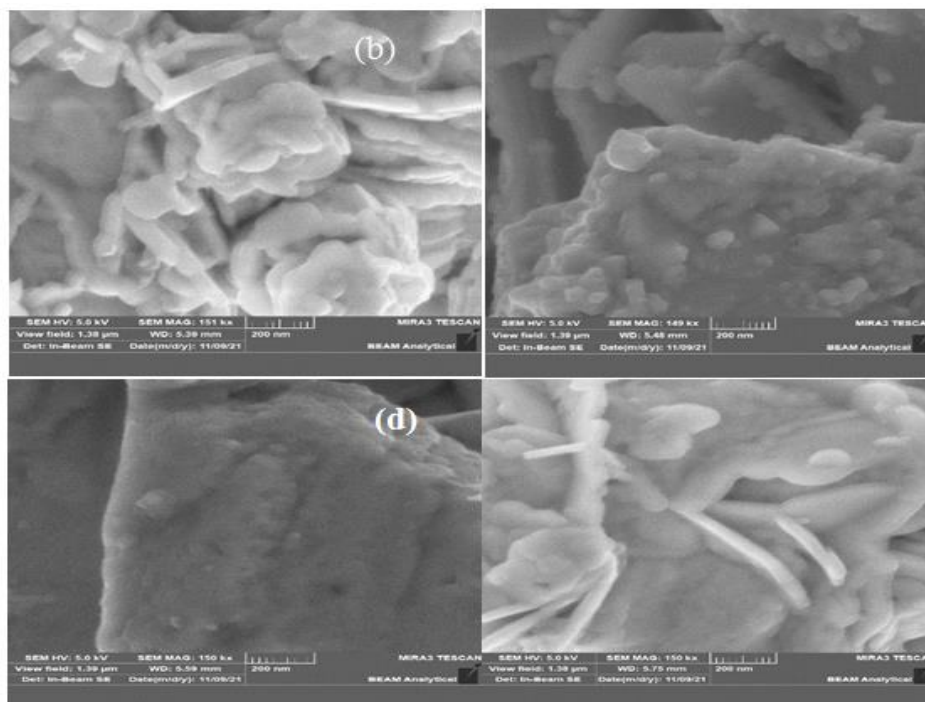


Figure 2. Surface morphology of (a) $TiO_2/NaAs_{0.1}$, (b) $TiO_2/NaAs_{0.2}$ / bitter leaf dye, (c) $TiO_2/NaAs_{0.3}$ / roselle dye, and (d) $TiO_2/NaAs_{0.4}$ /the mixture of bitter leaf dye and roselle dye

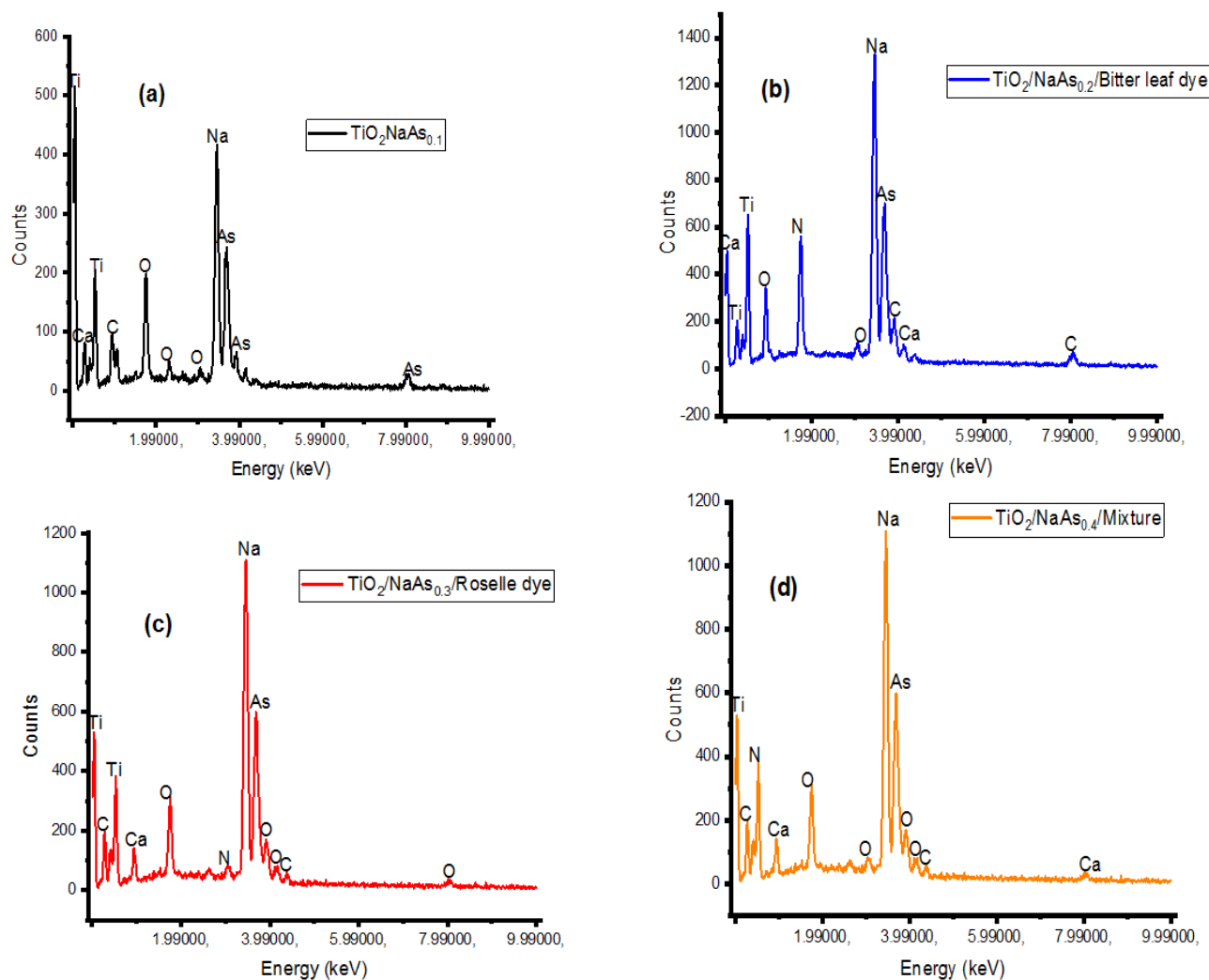


Figure 3. (a) $\text{TiO}_2/\text{NaAs}_{0.1}$, (b) $\text{TiO}_2/\text{NaAs}_{0.2}$ / bitter leaf dye, (c) $\text{TiO}_2/\text{NaAs}_{0.3}$ / roselle dye, and (d) $\text{TiO}_2/\text{NaAs}_{0.4}$ /the mixture of bitter leaf dye and roselle dye

X-ray diffraction analysis of $\text{TiO}_2/\text{NaAs}_{0.1}$, $\text{TiO}_2/\text{NaAs}_{0.2}$ / bitter leaf dye, $\text{TiO}_2/\text{NaAs}_{0.3}$ / roselle dye, and $\text{TiO}_2/\text{NaAs}_{0.4}$ /the mixture of bitter leaf dye and roselle dye

The XRD study of the synthesized $\text{TiO}_2/\text{NaAs}_{0.1}$, $\text{TiO}_2/\text{NaAs}_{0.2}$ / bitter leaf dye, $\text{TiO}_2/\text{NaAs}_{0.3}$ / roselle dye, and $\text{TiO}_2/\text{NaAs}_{0.4}$ /the mixture of bitter leaf dye and roselle dye depicted in Figure 4 are polycrystalline with a corresponding JCPDS card number 05-4562-0976, 05-4562-0979, 05-4562-0980, and 05-4562-0988 respectively. The structure of the

cells is polycrystalline with a most outstanding peak at 2 theta angles of 26.73° and 51.84° corresponding to (111) and (202). Therefore, secondary peaks were noticed at 2 theta angles of 37.94° and 65.42° which correspond to (112) and (301), subsequently. Moreover, the diffraction peak noticed raises as the molar concentration of sodium arsenic (NaAs) increases from (0.1-0.4 mol). The rise in the intensity peak is due to the enhancement of the crystal structure and energy absorption on the lattice parameters of the TiO_2 . The peak locations were not shifted due to the energy absorption of the crystals. The

structure of the lattice and single-cell distortion due to the developing increase in molar concentration of sodium arsenic (NaAs) may be concluded for the shifts in the orientation of the crystals [24-31]. The crystallite size or grain size was calculated using Equation (1).

$$D = \frac{0.9\lambda}{\beta \cos\theta} \quad (1)$$

The grain size for TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye are presented in Table 1. It is evident that the increase in molar concentration of sodium arsenic (NaAs) increases the grain sizes and enhances the crystallinity of the films (See Table 1).

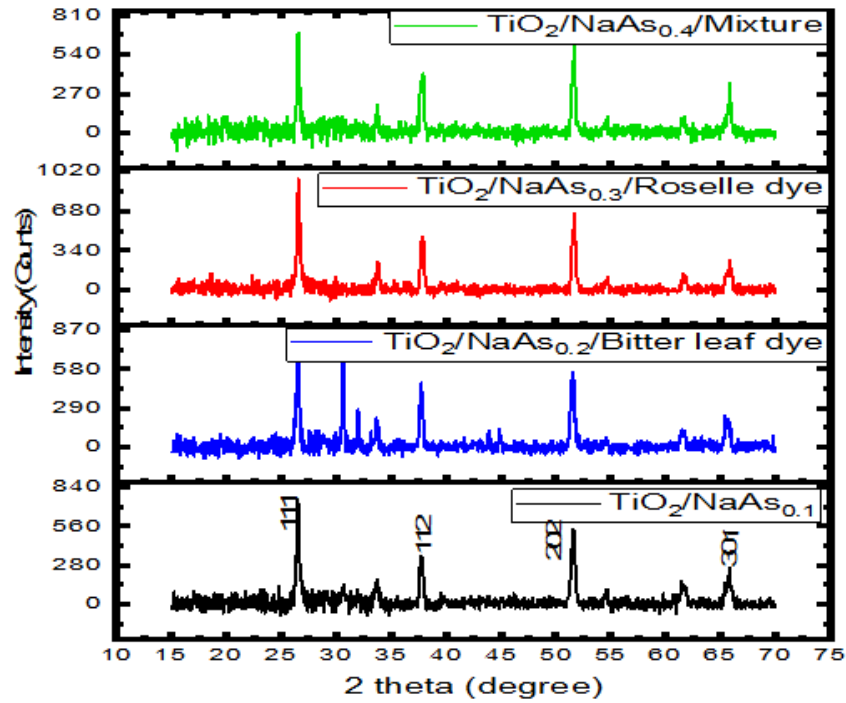


Figure 4. XRD spectrum of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye

Table 1. Structural parameter for TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye

Photoanode device	2θ (Degree)	(hkl)	d-spacing (Å)	Lattice constant (a)	FWHM (β)	Grain Size, D (nm)	Dislocation density, σ
TiO ₂ /NaAs _{0.1}	26.73	111	3.331	5.771	0.183	0.766	5.118
TiO ₂ /NaAs _{0.2} /Bitter leaf dye	37.94	112	2.369	4.738	0.209	0.779	6.205
TiO ₂ /NaAs _{0.3} /Roselle dye	51.84	202	1.762	3.524	0.148	0.784	2.799
TiO ₂ /NaAs _{0.4} /Mixture	65.42	301	1.425	3.187	0.225	0.799	5.651

The optical study of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye

The absorbance, transmittance, and reflectance of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye are displayed in Figure 5. Generally, the films have a very high absorbance from the plot, and the absorbance of the films increases as the dye molecules vary. The high absorbance of the films shows that the DSSCs will be a good material for photovoltaic applications. Figure 5 (a) presents TiO₂/NaAs_{0.1} with the lowest absorbance from the spectra and without dye molecules. It was discovered that dye molecules improve the absorbance of the films, and the *Vernonia amygdalina* (bitter leaf) dye particles have good anchoring settings on the exterior of TiO₂/NaAs_{0.2} with a better absorbance at the infrared and ultraviolet region of the spectra while *Hibiscus sabdariffa* (roselle) dye show the higher values of absorbance from the spectra at the UV region of the spectra. Both the *Vernonia amygdalina* (bitter leaf) dye molecules and *Hibiscus sabdariffa* (roselle) dye molecules give a good efficiency from the photoelectrical properties of the assembled DSSCs [24-31] which has not been reported by researchers on DSSCs. However, it clearly showed that the results obtained for TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye are suggested to be a good material for photovoltaic application for industrial purpose.

The transmittance of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye in Figure 5b reveals a very low transmittance value due to the high absorbance of the films. A high

absorbance value usually results in low transmittance. From the spectra, it was observed that TiO₂/NaAs_{0.1} has the highest transmittance from the spectra and without dye molecules [24-31]. It was discovered that dye molecules improve the transmittance due to the high absorbance value of the films, and the *Vernonia amygdalina* (bitter leaf) dye particles have good anchoring settings on the exterior of TiO₂/NaAs_{0.2} with a better transmittance at the infrared and ultraviolet region of the spectra while *Hibiscus sabdariffa* (roselle) dye show the lower transmittance values from the spectra at both region of the spectra. Both the *Vernonia amygdalina* (bitter leaf) dye molecules and *Hibiscus sabdariffa* (roselle) dye molecules give a good efficiency from the photoelectrical properties of the assembled DSSCs which have not been reported by researchers on DSSCs.

The reflectance of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye in Figure 5c revealed a negative reflectance value due to high absorbance of the films [24-31]. It is reported that dye molecules improve the absorbance of the films, and the *Vernonia amygdalina* (bitter leaf) dye particles have good anchoring settings on the exterior of TiO₂/NaAs_{0.2} with a better absorbance at the infrared and ultraviolet region of the spectra while *Hibiscus sabdariffa* (roselle) dye show the higher values of absorbance from the spectra at the UV region of the spectra, but the negative reflectance values of the films enhance the anchoring properties of the dye why perverting the films from external forces.

The energy band gap of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye in Figure 6 shows that the dye has a great effect on the band gap energy of the films.

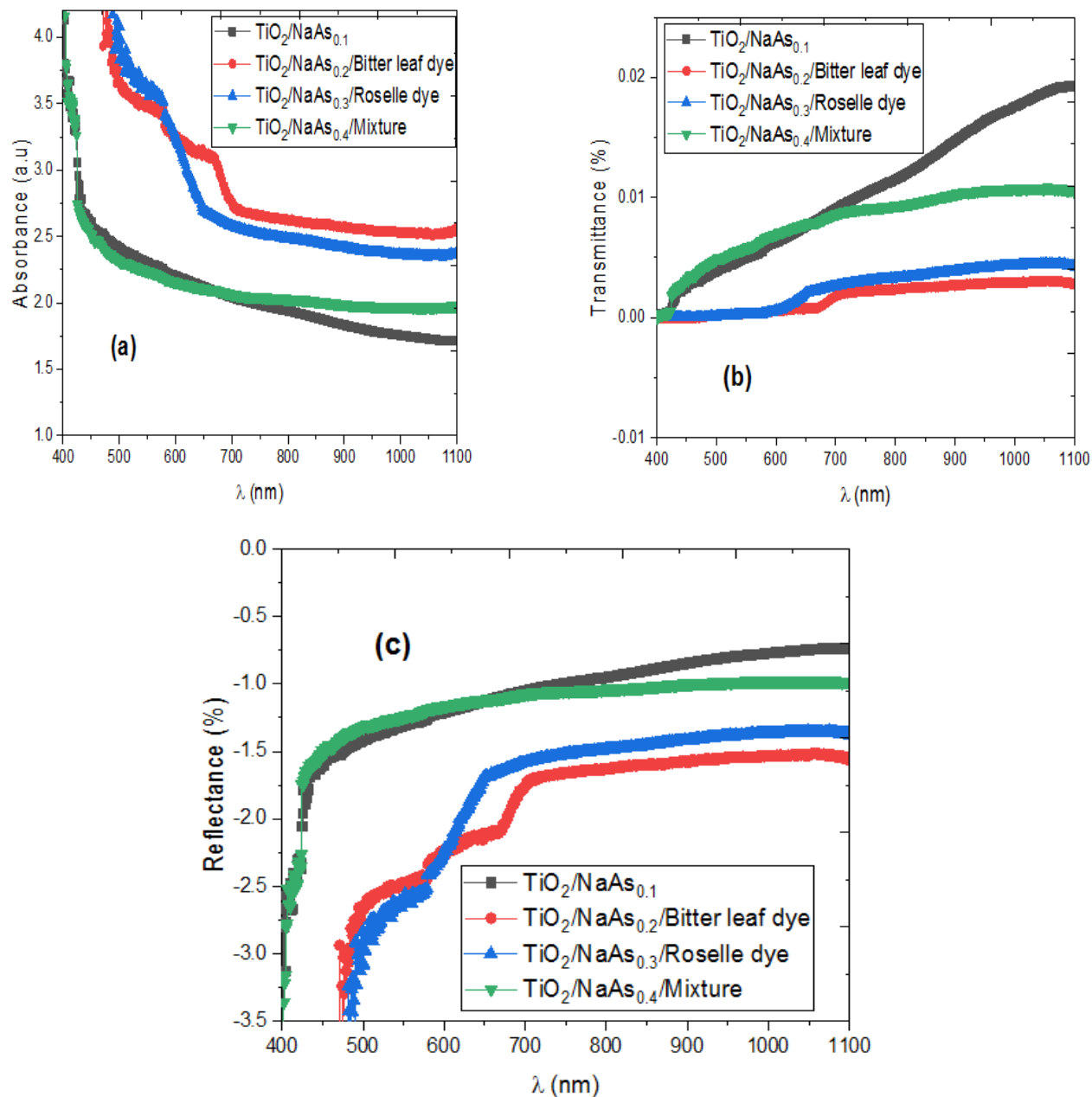


Figure 5. (a) absorbance, (b) transmittance, and (c) reflectance of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye

The *Vernonia amygdalina* (bitter leaf) dye molecules have band gap energy of 1.70 eV which is the highest in the trend, the *Hibiscus sabdariffa* (roselle) dye molecule with an energy band gap of 1.50 eV the mixture of both dye with 1.40 eV. The plot clearly shows that the mixture of *Vernonia amygdalina* (bitter leaf) dye

and *Hibiscus sabdariffa* (roselle) dye molecules has better band gap energy [24-31].

Photoelectrical properties of the assembled DSSC

Table 2 summarizes the photovoltaic performance of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and

TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye of the DSSCs. The photoelectrical properties of the assembled DSSCs without natural dye and natural dye revealed that the TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye, and roselle dye has J_{sc} of 2.62 mA/cm² and V_{oc} of 287.7 mV, J_{sc} of 4.67 mA/cm² and V_{oc} of 365.6 mV, J_{sc} of 3.72 mA/cm² and V_{oc} of 355.2 mV, and J_{sc} of 2.92 mA/cm² and V_{oc} of 296.7 mV, respectively. The

(FF) fill factor of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye is 0.54, 1.24, 1.23, and 0.99, respectively, while the conversion efficiency of 0.86%, 4.48%, 3.44%, and 1.81% was recorded for TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye, and roselle dye, subsequently.

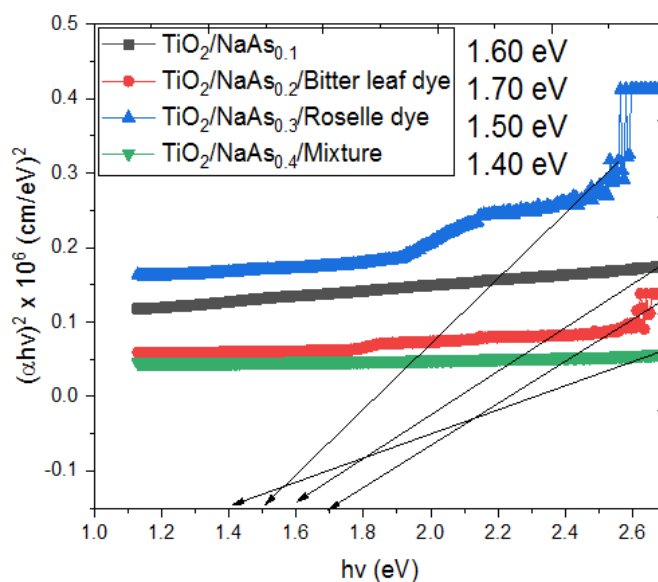


Figure 6. Energy band gap of TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye

Table 2. The photovoltaic performance TiO₂/NaAs_{0.1}, TiO₂/NaAs_{0.2}/ bitter leaf dye, TiO₂/NaAs_{0.3}/ roselle dye, and TiO₂/NaAs_{0.4}/the mixture of bitter leaf dye and roselle dye

Photoanode device	J _{sc} (mAcm ⁻²)	V _{oc} (V)	FF	η (%)
TiO ₂ /NaAs _{0.1}	2.62	287.7	0.54	0.86
TiO ₂ /NaAs _{0.2} /Bitter leaf dye	4.67	365.6	1.24	4.48
TiO ₂ /NaAs _{0.3} /Roselle dye	3.72	355.2	1.23	3.44
TiO ₂ /NaAs _{0.4} /Mixture	2.92	296.7	0.99	1.81

The photoelectrical conversion efficiency reveals that the *Vernonia amygdalina* (bitter leaf) dye particles have better anchoring properties on the TiO₂/NaAs_{0.2} surface. Thus, the performance of TiO₂/NaAs_{0.2} DSSC is considered better when compared to the performances of

TiO₂/NaAs_{0.3} with a higher molar concentration of sodium arsenic. The facts obtained here are above the range of the results obtained by different researchers on natural dyes with the characteristic performances given in the literature [24-31].

Conclusion

We have successfully synthesized and fabricated dye-sensitized solar cells (DSSCs) on the influence of sodium arsenic on the enhancement of TiO_2 /dye as photosensitizers, where *Hibiscus sabdariffa* (roselle) and *Vernonia amygdalina* (bitter leaf) were used as a source of the chlorophyll, Sodium arsenic (NaAs) material of different concentration (0.1-0.4 mol), was synthesized as a layer on top of TiO_2 . The surface morphology study of $\text{TiO}_2/\text{NaAs}_{0.1}$, $\text{TiO}_2/\text{NaAs}_{0.2}$ / bitter leaf dye, $\text{TiO}_2/\text{NaAs}_{0.3}$ / roselle dye, and $\text{TiO}_2/\text{NaAs}_{0.4}$ /the mixture of bitter leaf dye and roselle dye revealed that the micrograph is usually defined with the granular shape of nanotubes. The grain size of $\text{TiO}_2/\text{NaAs}_{0.1}$ is not too large and delineated by an immense sum of aggregated nanoparticles. $\text{TiO}_2/\text{NaAs}_{0.2}$ / bitter leaf dye as shown has smaller grain-sized particles whose shapes are granular and have more aggregated particles. The clustering of the particles is evidenced in samples synthesized with the chlorophyll of *Vernonia amygdalina* (bitter leaf). The cells structure is polycrystalline with a most outstanding peak at 2 theta angles of 26.73° and 51.84° corresponding to (111) and (202). Therefore, secondary peaks were noticed at 2 theta angles of 37.94° and 65.42° which correspond to (112) and (301) subsequently. The photoelectrical conversion efficiency reveals that the *Vernonia amygdalina* (bitter leaf) dye particles have good anchoring settings on the exterior of $\text{TiO}_2/\text{NaAs}_{0.2}$. Thus, the performance of $\text{TiO}_2/\text{NaAs}_{0.2}$ DSSC is considered superior when compared to the performances of $\text{TiO}_2/\text{NaAs}_{0.3}$ with a higher molar concentration of sodium arsenic.

Disclosure Statement

No potential conflict of interest was reported by the authors.

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Authors' Contributions

All authors contributed to data analysis, drafting, and revising of the paper and agreed to be responsible for all the aspects of this work.

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