

Potential of local material SiO₂ water Hyacinth for semiconductor materials

Sony Hidayat^{1*}, Ahmad Ziyen Nafis¹, Muhammad Noorman Perdana¹, Upik Nurbaiti¹

¹Department of Physics, Faculty of Mathematics and Natural Sciences, Semarang State University, Indonesia

* Corresponding author's e-mail: sonyhidayat777@students.unnes.ac.id

ABSTRACT

This study aims to examine the potential of local SiO₂ material synthesized from water hyacinth biomass waste as a candidate for semiconductor materials. Synthesis was carried out through a calcination process at temperatures above 600 °C to remove cellulose, lignin, and other impurities. Material characterization was carried out using UV-Vis spectroscopy, X-ray Diffraction (XRD), and Raman Spectroscopy. The results of UV-Vis spectroscopy showed that the water hyacinth derived SiO₂ exhibited an unusually reduced optical band gap of approximately 2.3 eV, likely influenced by impurity phases.. XRD tests indicated the presence of two crystal phases, namely the trigonal structure of SiO₂ and the monoclinic structure of the CaH₁₂O₁₇Si₂U₂ compound. Raman analysis confirmed the presence of functional groups such as Si, amorphous SiO₂, CH₂/CH₃ and vibration signals from irregular sp² carbon. This combination of structures is thought to cause a lower band gap value compared to pure SiO₂. This finding indicates that SiO₂ from water hyacinth has potential as an intermediate semiconductor material, although further purification is still needed to increase the purity of the SiO₂ phase.

Keywords:

Band Gap; Local Water Hyacinth Biomass; Semiconductor; SiO₂.

Introduction

The development of semiconductor materials has become increasingly advanced in various fields of technology (Nurhayati et al., 2023). These applications include LEDs, computer chips, and renewable energy technologies. Semiconductors are materials that have unique properties, the energy gap is not too large, between metal and insulator materials (Rasyidi, 2015). One of the materials that is widely used in the manufacture of semiconductors is silicon (Si) because it has a fairly small band gap, around 1.4 eV, and 3-4 eV for silicon dioxide or silica (SiO₂) (Rai et al., 2014; Ücker et al., 2023). Research in the field of semiconductors has been widely carried out, but research that utilizes local organic materials is still not widely studied, especially biomass waste. Therefore, investigating biomass waste as a potential source of advanced materials is important for identifying locally derived alternatives.

According to Hidayat et al (2024) and Junaudi et al (2022), water hyacinth is one of the biomass waste materials that can be synthesized into silica. Water hyacinth is an aquatic weed that grows very quickly and is invasive if its population is not controlled (Kalsum, 2015) (Figure 1). Excessive water hyacinth populations will cause quite serious environmental impacts, such as shallowing of the water ecosystem, reduced water oxygen levels, excessive methane gas in the water ecosystem due to water hyacinth decay, and blockage of water flow due to excessive water hyacinth numbers (Ningsih et al., 2019; Ratnani et al., 2024; Zevriawan et al., 2023). Therefore, the existence of water hyacinth in nature needs to be controlled and utilized wisely. The processing of water hyacinth into handicrafts has been widely carried out by creative economy activists, but other potentials need to be further developed, one of which is for semiconductor materials.

The purification of SiO₂ from water hyacinth is carried out using the calcination method, calcination is a material purification process using a heating temperature in the range of 500°C-1000°C (Usman et al., 2020; Pratama et al., 20230. In this case, calcination is intended to purify

SiO₂ from cellulose, lignin and unnecessary impurities, so that it is expected to produce SiO₂ with high purity. In addition, the calcination process is intended to form a more uniform SiO₂ crystal structure.



Figure 1. (a) Illustration of Over Population Water Hyacinth on Bengawan Solo River (KOMPAS.COM); (b) Illustration of Over Population Water Hyacinth on Martapura River (KOMPASIANA.COM).

This study focuses on determining the band gap value of SiO₂ synthesized from water hyacinth, its crystal structure and functional groups. However, this study does not cover the direct application of SiO₂ material, this is also a limitation in the study.

Methods

Synthesis of water hyacinth SiO₂



Figure 2. Extraction Process SiO₂ from Water Hyacinth.

Freshly obtained water hyacinth is cleaned from impurities and then dried under the hot sun. The dried water hyacinth is then burned using conventional methods to obtain its ash (Figure 2). Next, 150 mL of citric acid solution (C₆H₈O₇) is added to 48 g of the water hyacinth ash. Soaking is carried out for several minutes with a total mixture temperature of around 50°C using a stirrer heater. Water hyacinth that has been soaked in citric acid solution is then washed with hot water 3 times to remove the citric acid content in it. The residue from washing in the form of water hyacinth ash is then ovened at a temperature of 100°C for 25 minutes. The next step is the purification of SiO₂ using a furnace at a temperature above 600°C. (Hutomo, 2017). Then the synthesis results were tested using UV-VIS to determine the band gap of the material, XRD to determine the crystal structure of the synthesized SiO₂ material, and Raman Spectroscopy to determine the functional groups of the material.

Results and Discussions

Water hyacinth has been synthesized into powder suspected to be SiO₂. The powder obtained was then subjected to XRD characterization to see the crystals and validate the presence of SiO₂, then strengthened with Raman spectroscopy, then to see the semiconductor properties of the band gap, UV-Vis characterization was carried out (Figure 3).



Figure 3. Results of Extraction SiO₂ from Water Hyacinth.

UV-VIS Analysis

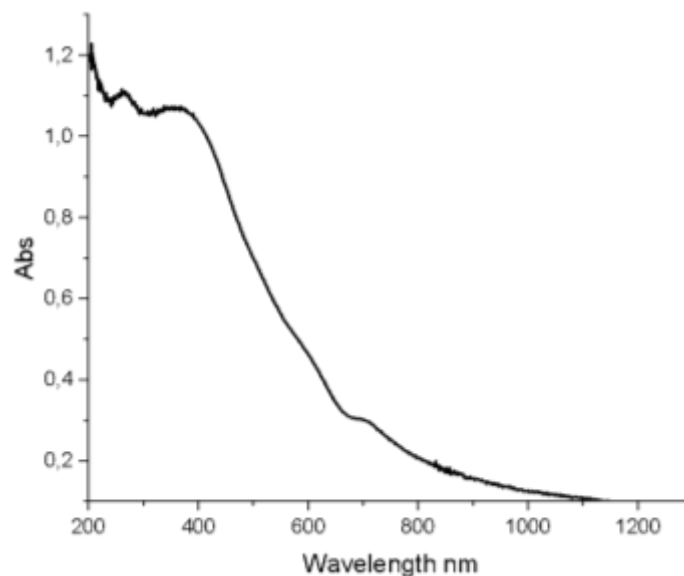


Figure 4. Graph of SiO₂ Absorbance Value of Water Hyacinth

The results of the synthesis of SiO₂ water hyacinth powder were then tested for optical absorbance using UV-VIS Spectroscopy in the testing wavelength range of 200nm-1200 nm. It can be seen that there is one fairly sharp absorbance peak, which indicates an electronic transition from the valence band to the conduction band (Figure 4). The phenomenon of the emergence of one fairly sharp absorbance peak indicates that the SiO₂water hyacinth material has direct band gap characteristics, because indirect band gap materials generally occur two or more energy transitions accompanied by tiered absorbance due to phonon participation (Sharma et al., 2012). In direct band gap materials, electron transitions from the valence band to the conduction band can occur directly without any change in momentum, resulting in one sharp peak in the absorbance spectrum (Muttaqin, 2015). Thus, the water hyacinth-derived SiO₂ appears to exhibit direct band gap behavior (Babyszko et al., 2023). To find out the band gap value, further analysis can be carried out using the tauc plot method with the equation:

$$(\alpha h\nu)^n = A(h\nu - E_g)$$

Where α is the absorption coefficient, $h\nu$ is the photon energy, E_g is a constant, n is the band gap energy, $n = \frac{1}{2}$ is the value for direct transitions, and $n = \frac{1}{2}$ is for indirect transitions.

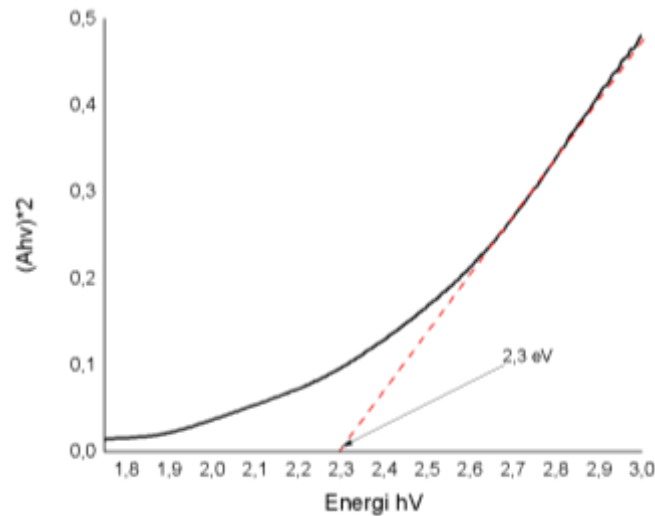


Figure 5. Energy Gap Graph SiO₂ Water Hyacinth Tauc Plot Method

The Tauc plot analysis yielded a band gap value of approximately 2.3 eV (Figure 5). The band gap value of 2.3 eV is included in the category of semiconductors with an intermediate band gap, which is too small for an insulator and too large for a metal conductor, but is ideal for various light-active semiconductor applications such as red-orange LEDs ($\lambda \sim 540\text{--}560\text{ nm}$) which require materials with a band gap of around 2 eV (Telefunken, 1999). When compared with research by Seguíni et al (2011); Hussin (2017), SiO₂ has a band gap of around 3 eV–5 eV, while SiO₂ synthesized from water hyacinth has a band gap of around 2.3 eV. This is likely due to the presence of other elements or compounds that are still present in the SiO₂ from the synthesized water hyacinth. The fairly flat graph curve indicates that the SiO₂ synthesized from water hyacinth is not completely pure, in contrast to the band gap graph of SiO₂ which tends to be almost vertical. Therefore, further analysis is needed with XRD to determine the structure of the crystals in the material, as well as Raman Spectroscopy analysis to determine the presence of other functional groups in the synthesized material.

X-Ray Drifraction XRD Analysis

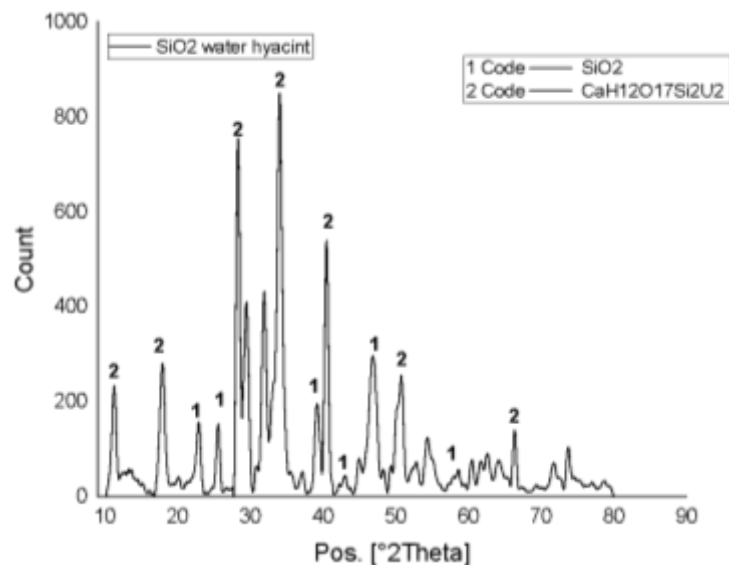


Figure 6. XRD Graph of SiO₂ Water Hyacinth

The results of the synthesis of SiO₂ water hyacinth will then be characterized by XRD (target Cu, $\lambda K\alpha = 1.54060 \text{ \AA}$). Testing using XRD aims to determine the crystal phase that appears and to determine the crystal shape formed from the sample.

After the XRD characterization test was carried out, the data obtained were then processed using Match Software with the Crystallography Open Database (COD) 2025 comparative database (Figure 6). The diffraction peaks shown (2θ) (code 1): 23; 25.99; 39.02; 43; 46.9; 58.71, are diffraction peaks owned by the SiO₂ crystalline structure, with a trigonal crystal structure, and COD id code: 8103513. The peaks at $2\theta = 11.25^\circ, 17.88^\circ, 28.35^\circ, 34.2^\circ, 40.8^\circ, 47.02^\circ, 50.84^\circ$, and 66.4° (code 2) correspond to a monoclinic phase indexed in the COD as CaH₁₂O₁₇Si₂U₂ (COD ID 9014766). The emergence of crystal structures other than SiO₂ is a by-product of imperfect material synthesis. Basically, the elements Ca, H, and O are indeed found in the structure of organic materials, especially plants, so the emergence of the CaH₁₂O₁₇Si₂U₂ crystal system is very common if the purification process is not perfect. To visualize the trigonal SiO₂ and monoclinic CaH₁₂O₁₇Si₂U₂ structures, the corresponding COD files were processed using VESTA software (Figure 7).

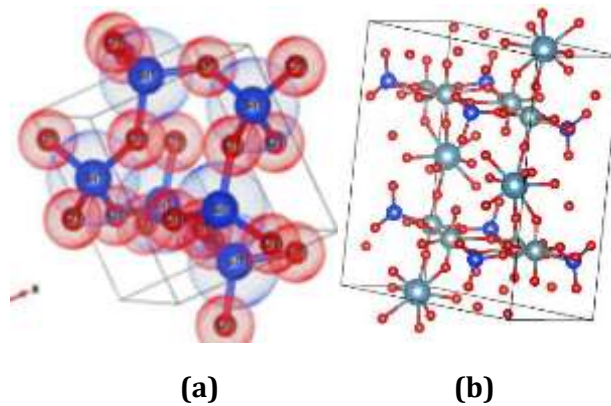


Figure 7. (a) Crystal structure of SiO₂ , (b) Crystal structure of CaH₁₂O₁₇Si₂U₂

Raman Spectroscopy Analysis

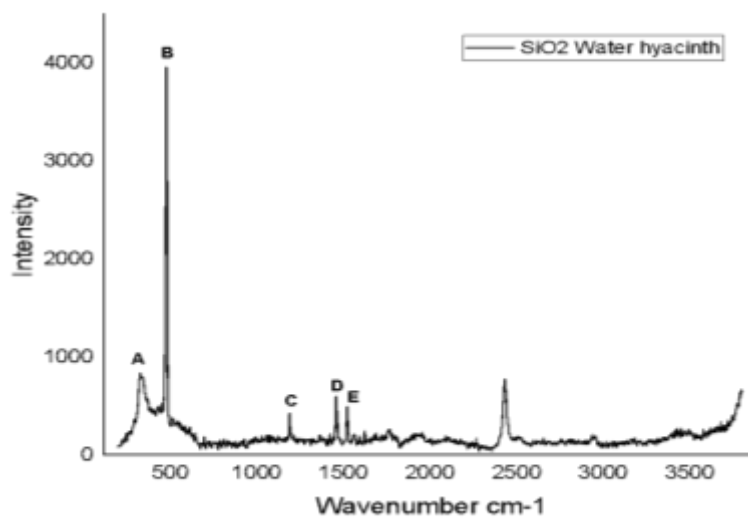


Figure 8. Raman Spectroscopy Graph of SiO₂ Water Hyacinth

The SiO₂ water hyacinth sample was then subjected to Raman Spectroscopy characterization test to see the phase of the compound or other elements in the sample (Figure 8). The Raman characterization test was carried out in the wave number range of 100-4000 cm⁻¹. The Raman spectroscopy graph shows several sharp peaks and quite a lot of noise at several wave number points (Table 1). The appearance of noise is likely due to the camber of the sample used and the appearance of fluorescence phenomena in the sample when exposed to laser light. Fluorescence is a phenomenon where the outermost electrons in the sample compound are excited, and emit light with a higher wavelength than the wavelength of the exciting light. In Raman Spectroscopy testing this often occurs, especially in organic samples, usually this phenomenon appears in the low wave number range of 300-450 cm⁻¹ depending on the sample.

Table 1.Functional Group of SiO₂ Water Hyacinth

Code	Wave Number cm ⁻¹	Intensity	Bond
A	336.24	664.68	Fe-S
B	471.89	3962.64	Si-a
C	1100	401	SiO ₂
D	1458.33	455.85	CH ₂ /CH ₃
E	1529.06	408.74	G

According to Galuskina et al (2025) The spectrum peak at wave number 336 cm⁻¹ is related to the symmetric stretching vibration of Fe-S. The appearance of the Fe element in the sample is most likely due to contaminants during the sample refining process. The spectrum peak that appears at wave number 471 cm⁻¹ is a peak indicating the presence of vibrations of the Si-a element which is an amorphous Si compound, as reported by Abed et al (2020). The noise spectrum that appears in the wavelength range of 1100 cm⁻¹ is a spectrum peak indicating the presence of SiO₂ compounds (Abed et al., 2020). Meanwhile, the spectrum peak that appears at wave number 1458 cm⁻¹ is a peak due to deformation of the CH₂/CH₃ compound as reported by research conducted by Zyubin et al in 2018. Peak D at wave number 1529 cm⁻¹ according to research conducted by Galaburda et al (2019); Tan et al (1999); Cho et al (2015) can be associated with the spectrum due to irregular sp² carbon Graphene (G) vibrations. The emergence of graphen elements or carbon bonds (C) is very natural if the purification process is not perfect, the C element appears from the burning process of water hyacinth which has not completely disappeared during calcination. The results of Raman Spectroscopy testing support XRD and UV-VIS test data showing compounds other than SiO₂, causing a decrease in the band gap value compared to pure SiO₂.

UV-VIS characterization shows that the SiO₂ material synthesized from water hyacinth has direct band gap properties and has a band ap value of 2.3 eV using the touch plot method. The results of the XRD test showed that the synthesized material had 2 types of crystal phases, namely Trigonal SiO₂ and monoclinic CaH₁₂O₁₇Si₂U₂. Raman Spectroscopy testing shows the presence of Si, amorphous SiO₂, CH₂/CH₃ and Vibrations of the G element.

Conclusion

The results of the study showed that the SiO₂ compound synthesized from water hyacinth has a type of electron transfer from the conduction band to the valence band of the direct band gap type. In addition, the SiO₂ compound from water hyacinth also has an energy gap of 2.3 eV. This indicates that the synthesized water hyacinth-derived SiO₂ can be classified as an intermediate-band-gap semiconductor. XRD analysis shows 2 crystal structure phases, namely trigonal SiO₂ and CaH₁₂O₁₇Si₂U₂ monoclinic. CH₂/CH₃ and Graphene (G) compounds are also still present in the synthesized SiO₂ material. This combination has justified the value of the SiO₂ band gap from water hyacinth 2.3 eV is influenced by other compounds, not purely from SiO₂.

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Conflicts of interest

The authors declare that there are no conflicts of interest.

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