SYNTHESIS AND CHARACTERIZATION OF GRAPHENE: ITS OPTICAL PROPERTIES FOR APPLICATION IN SOLAR CELL

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Abstract

Graphene shows remarkable optical behavior, where above an incident light intensity threshold, transmission is attenuated. There is significant interest in synthesizing large-area graphene films at low temperatures by chemical vapor deposition (CVD) for optical device applications. The results obtained shows that the refractive index of graphene is \( n_g = 1.86 - 1.56 \) while that of graphene conductance is \( \sigma_r = 1.016 \) and \( \sigma_i \approx 0 \) and graphene density 0.77 mg. The modified CVD method is expected to allow the direct growth of graphene in device manufacturing processes for practical applications while keeping underlying devices intact.

Keywords: Graphene, Optical, Incident, Synthesizing and chemical vapor deposition (CVD)

1. Introduction

Graphene is a single atom, thick two-dimensional (2D) material, thus, exhibiting ~97.7% transmittance throughout the entire visible light spectrum. Additionally, graphene has a flat transmittance spectrum from the ultraviolet (UV) region to the long wave length infrared (IR) region, thus exhibiting a wide window that allows a comprehensive range of photon wave length passed through it. Unconditionally, except graphene, these combinations of remarkable optical properties are yet to be observed in any types of materials till date. Similarly, graphene has unusual electronic transport properties, which follows the characteristics of 2D Dirac fermions, quantum hall effects, Landau level quantization, and so on. Consequently, graphene’s free charges are immobile in one spatial dimension but mobile in other two dimensions, and thus, charge carrier mobility is \( \sim 10^6 \text{cm}^2/\text{v/}s \) in free-standing graphene (Yoshikawa K 2017). Similarly, graphene also exhibits excellent mechanical and thermal properties \( (k \sim 300 - 500 \text{W/mK}) \) and chemical inertness. Hence, all these properties coupled with the optical properties put together graphene a stronger candidate for applications in
transparent conducting electrodes (TCEs) flexible optoelectronics, energy harvesting devices, photo detectors, and many other optical devices (Yoshikawa 2017).

Furthermore, graphene is anticipated as an emerging alternate for conventional transparent conductive metal oxides, specifically, indium tin oxide (ITO), which contains indium as a toxic, costly, and scarce element. In particular, the graphene-based TCE for application in solar cells with enhanced efficiency is of utmost interest. To date, graphene electrodes have been applied for different types of solar cells, namely, solid-state solar cells, electrochemical solar cells, quantum dot solar cells (QDSCs), and polymer solar cells (Zang Y 2011).

2. Material and Method

The experiment setup and data measurement was done at the department of mineralogy Kaduna state Polytechnic. This research will involve Graphene films deposited on a glass substrate using spin coating. It employed a reduction protocol where a 4 Watt, 365 nm UV lamp was used for exposure of the synthesized Graphene films for different periods of time. The current-voltage (I-V) characteristics for the Graphene films were measured as a function of UV exposure time. For this, a four-probe configuration was used and current across the two inner terminals was taken as a function of the source voltage applied across the two outer terminals.

The evolution of the I-V characteristics with the UV exposure time (up to 2 hours) was analyzed. Raman and X-ray photoemission (XPS) spectra was obtained both prior to UV exposure as well as post 2 hours exposure for samples belonging to the same batch of synthesis. A comparison of the Raman spectra of the grown sample and a corresponding sample was synthesized after 2 hours exposure to UV light. The characteristic D, G and 2D peaks was labeled. A comparison of reduced GO Raman spectra with pristine graphene (CVD grown), graphene single layer ($I_{2D}/I_G = 2.7$), bilayer ($I_{2D}/I_G = 1.4$) and multilayer Raman spectra were compared to provide a recipe for estimating extent of disorder in Graphene.

A conservative estimate of the extent of disorder based on the observed $I_D/I_G$ ratio of the 2 hour reduced Graphene sample was made and were estimated.

Distilled water of 1000 cm$^3$ was measured and mixed with the graphite sample, poured into the flotation cell and agitated for 3 minutes. During the agitation process, three drops of regulator (sodium hydroxide) was added to adjust the pulp pH to 8.5 and
conditioned for seven minutes. Thereafter, 3ml of kerosene and pine oil added, immediately after the conditioning/agitation, air was allowed to pass through the pulp at a reasonable rate and froth emerged and collected till barren froth surfaced.

**Table 1.0:** Shows fixed carbon content of raw graphite (as received)

| Sample | 
|-------|---|
| MC    | Vol.7 Issue 6, June 2019 |
| VM    | 19. |
| AS    | 39. |
| FC    | 26.2 |

<table>
<thead>
<tr>
<th>Samples Content %</th>
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<tr>
<td>HeadSample</td>
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<tr>
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<tr>
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</tr>
<tr>
<td>19.</td>
</tr>
<tr>
<td>39.</td>
</tr>
<tr>
<td>5</td>
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**Table 2.0:** Shows fixed carbon content of the concentrate

<table>
<thead>
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</tr>
<tr>
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</tr>
<tr>
<td>5.4</td>
</tr>
<tr>
<td>10.</td>
</tr>
<tr>
<td>81.2</td>
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**3. Result and Discussion**

**Optical properties of graphene**

The electromagnetic properties of graphene are based on its unique atomic structure and the electronic band-gap between graphene sheet layers. As it is a two-dimensional material, the massless Dirac fermions in graphene have an amazing property: the optical response is universal and expressed only regarding to the fine-structure constant. To derive this response, we need determine the electric field of the incident light via a vector potential:

\[
\vec{A}(t) = \vec{A}_{\text{exp}} (-i\omega t) \tag{1.0}
\]

\[
\vec{E}(t) = \frac{1}{c} \frac{\partial \vec{A}}{\partial t} = i\omega \vec{A} \tag{1.1}
\]

Thus, the incident working on the Hamiltonian of Dirac electron in the presence of an electric field is:
\( \vec{H} = v \vec{\sigma} \left( \vec{p} - \frac{e}{c} \vec{A} \right) = \vec{H}_0 + \vec{H}_\text{int} \)  \hspace{1cm} (1.2)

Where:
\[ H_{\text{int}} = \frac{ve}{2c} \vec{\sigma} \vec{A} = i \frac{ev}{2\omega} \vec{\sigma} \vec{E} \]  \hspace{1cm} (1.3)

Is the Hamiltonian of the electron photon interaction, the factor \( \frac{1}{2} \) in equation (4.3) is necessary since the standard expression for the complex field is:
\[ \vec{E}(t) = \text{Re} \left[ \vec{E} \exp(-i\omega t) \right] = \frac{1}{2} \left( \vec{E} \exp(-i\omega t) + \vec{E} \exp(i\omega t) \right) \]  \hspace{1cm} (1.4)

And only the first term need to take into account. This interaction induces transitions from the occupied whole states \( \varphi_{\uparrow} (\vec{k}) \) to empty electron state \( \varphi_{\downarrow} (\vec{k}) \) with the same wave vector \( \vec{k} \), the intraband transitions being forbidden by momentum conservation.

The matrix element of the Hamiltonian equation is given by:
\[ \langle \varphi_{\uparrow} | \vec{H}_{\text{int}} | \varphi_{\downarrow} \rangle = \frac{e^2 v}{2\omega} (E_y \cos \varphi + E_x \sin \varphi) \]  \hspace{1cm} (1.5)

Where the negative and positive signs correspond to \( k \) and \( k' \) Valleys, the sign depends only on the polar angle \( \phi \) of the \( \vec{K} \) vector, not on its length. After determination the average value, the square matrix element \( |M|^2 \) over \( \phi \) is found by:
\[ |M|^2 = \langle \varphi_{\uparrow} | \vec{H}_{\text{int}} | \varphi_{\downarrow} \rangle = \frac{e^2 v^2}{8\omega^2} |\vec{E}|^2 \]  \hspace{1cm} (1.6)

Here, assuming the phone propagates perpendicular to the graphene plane, thus the electric field \( \vec{E} = (E_x, E_y, 0) \) lies within the plane. The absorption probability per unit time, to the lowest order of perturbation theory is:
\[ P = \frac{2\pi}{\hbar} |M|^2 N(\varepsilon = \frac{\hbar \omega}{2}) \]  \hspace{1cm} (1.7)

Where, \( N(\varepsilon) = \frac{2|E|}{\pi \hbar^2 v^2} \) is the density of states of single layer graphene material, the spin and valley degeneracy has been considered and the energy of the final state is \( \frac{\hbar \omega}{2} \) substituting the equation (1.6) result in:
\[ P = \frac{\sigma^2}{4\hbar^2 \omega} |\vec{E}|^2 \]  \hspace{1cm} (1.8)

Thus the optical absorption energy per unit time is:
\[ W = \rho \hbar \omega = \frac{\sigma}{4\hbar} |\vec{E}|^2 \], therefore the absorption coefficient is:
\[ A_0 \frac{W_0}{W_i} = \frac{\pi e^2}{\hbar c} \]
Optical refractive index of graphene

From Maxwell equations and Snell’s law, displacement $\vec{D}$ magnetic field $\vec{H}$ and surface current $\vec{j}$ can be expressed as:

\[
\begin{align*}
\vec{D} &= \varepsilon_0 \vec{E} + \vec{P} = \varepsilon_0 \varepsilon_r \vec{E} \\
\vec{j} &= \sigma \frac{\partial \vec{E}}{\partial t} \\
\nabla \times \vec{H} &= \frac{\partial \vec{D}}{\partial t} = \vec{j} + \varepsilon_0 \frac{\partial \vec{E}}{\partial t}
\end{align*}
\]

(1.9) (1.10) (1.11)

Where $\vec{P}$ is the dielectric polarization, $\sigma$ is the conductance, $d$ is the material thickness. $\varepsilon_0$ is the free space permittivity $(8.854 \times 10^{-12} F/m)$, and $r$ is the relative material permittivity. $\mu_0$ is the open space magnetic permeability $4\pi \times 10^{-7} H/m$, $\mu_r$ is relative permeability of the material.

The optical conductance of graphene defined from its linear dispersion relation as:

\[
\sigma = \sigma_r + \sigma_i
\]

(1.12)

Where $\sigma_r$ and $\sigma_i$ are the real and imaginary parts respectively, which can be rewritten as:

\[
\begin{align*}
\sigma_r &= \sigma_0 \left[ \frac{1}{2} + \frac{1}{72} \frac{\hbar \omega^2}{\xi^2} \times \left( \tan \frac{\hbar \omega + 2 \mu_c}{4K_B T} + \tan \frac{\hbar \omega - 2 \mu_c}{4K_B T} \right) \right] \\
\sigma_i &= \sigma_0 \left[ \frac{\mu_c}{\hbar \omega \pi} \left( 1 - \frac{2 \mu_c^2}{\xi^2} \right) - \log \left( \frac{|\hbar \omega + 2 \mu_c|}{|\hbar \omega - 2 \mu_c|} \right) \left( \frac{1}{\pi} + \frac{1}{36 \pi} \left( \frac{\hbar \omega}{\xi} \right)^2 \right) \right]
\end{align*}
\]

(1.13) (1.14)

Where $\xi \sim 3 eV$ is the energy hopping between nearest atoms in the chemical potential near Dirac point. $K_B T$ is the incident photon energy and $\sigma_0 = \frac{\pi}{2} \times \frac{e^2}{\hbar} = 6.08 \times 10^{-5} \Omega^{-1}$ is the universal graphene conductance. One theoretical work shows when the temperature $T = 300k$ under visible incidents real part and imaginary part of graphene conductance are; $\sigma_r = 1.016$ and $\sigma_i \sim 0$ respectively.

Substitute equations (1.13) and (1.14) into (1.12), we can rewrite the equation as:

\[
\frac{\sigma}{d} \frac{\partial \vec{E}}{\partial t} + i \omega \varepsilon_0 \vec{E} = i \omega \varepsilon_0 \varepsilon_r \vec{E}
\]

(1.15)

We can extract the relative permittivity $\varepsilon_r$ as:

\[
\varepsilon_r = 1 + \frac{\sigma}{j \omega \varepsilon_0}
\]

(1.16)

By definition of the refractive index of the medium is $n = \sqrt{\varepsilon_r \mu_r}$, which indicates the transmission efficiency in the medium. Regarding the linear susceptibility, the refractive

\[
1 \quad 137.04 \approx 2.3\%
\]
index is \( n_0 = \sqrt{\varepsilon_r \mu_r} = 1 \) when the material is graphene. Therefore, we can obtain the graphene refractive index as:

\[
n_g = \sqrt{\varepsilon_r} = \sqrt{1 + \frac{\sigma}{j\omega \varepsilon_0}}
\]

Where the refractive index is expressed in terms of fundamental constants and the ratio of optical wavelength to graphene thickness. We note that a universal conductance implies a refractive index with strong dispersion. For \( \lambda = 550\text{nm}, \sigma = 0.3354\text{nm} \) Therefore we find \( n_g = 1.86 - 1.56i \)

![Graphene refractive index figure](image)

**Figure 1.0:** Refractive indexes of (a) monolayer graphene and (b) bilayer graphene

4. **Density of graphene**

The unit hexagonal cell of graphene contains two carbon atoms and has an area of \( 0.053\text{nm}^2 \) and its density was found to be \( 0.77\text{mg/m}^2 \). A hypothetical hammock measuring 1m2 made from graphene would thus weigh 0.77 mg

5. **Electrical conductivity of graphene**

The sheet conductivity of a 2D material is given by. The mobility is theoretically limited to \( \mu = 200,000\text{cm}^2/\text{V}\cdot\text{s} \) by acoustic phonons at a carrier density of \( n = 1014\text{cm}^{-2} \). The 2D sheet resistivity, also called the resistance per Square, is then \( 28\Omega \). Our fictional hammock measuring 1m2 would thus have a resistance of \( 28\Omega \). Using the layer thickness we get a bulk conductivity of \( 0.86 \times 10^2\Omega^{-1}\text{cm}^{-1} \) for graphene. This is somewhat higher than the conductivity of copper which is \( 0.54 \times 10^2\Omega^{-1}\text{cm}^{-1} \).
Conclusion

In general, solar cells are not that efficient. However, recent advances in graphene-based solar cells have seen the reflectance of solar rays reduced by 20%, which provides a potential efficiency increase up to 20%, the thesis explains and demonstrates the use of Graphene for making solar panels for deriving extreme power supply using the unlimited solar energy. The results obtained shows that the refractive index of graphene is $n_g = 1.86 - 1.56i$ while that of graphene conductance is $\sigma_r = 1.016$ and $\sigma_i \sim 0$ and graphene density $0.77 \text{ mg}$ The modified CVD method is expected to allow the direct growth of graphene in device manufacturing processes for practical applications while keeping underlying devices intact.

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