

# Synthesis and Characterization of Au/Fe<sub>3</sub>O<sub>4</sub> NCs Decorated Graphene Oxide via Pulsed Laser Ablation Technique

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The demand for nanocomposites (NCs) of graphene and magnetite materials decorated with gold nanoparticles is increasing on account of their applications in science and technology. First, graphene oxide (GO) was made with the pulsed laser ablation technique in liquid (PLAL) and then loaded with magnetite (Fe<sub>3</sub>O<sub>4</sub>) that was prepared by the same technique to form Fe<sub>3</sub>O<sub>4</sub>/GO nanocomposite (NCs). The Au/Fe<sub>3</sub>O<sub>4</sub> NCs were also synthesized and added by volume ratio to the GO solution to form Au/Fe<sub>3</sub>O<sub>4</sub>/GO NCs. We investigated the functional groups in the three samples using Fourier transform infrared spectroscopy (FTIR). We measured the optical absorption spectra of the samples using UV-visible light. From these optical spectra, we calculated the optical conductivity ( $\sigma$ ), absorption coefficient ( $\alpha$ ), and other related optical properties. We improved the optical absorption and other optical parameters while decreasing the band gap ( $E_g$ ) from 4.1 eV to 3 eV for the three samples. These materials are suitable for photonic applications, drug delivery applications, and water treatment applications.

## Keywords

graphene oxide (GO), magnetite (Fe<sub>3</sub>O<sub>4</sub>), gold/magnetite NCs, and the pulsed laser ablation technique.

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## Introduction

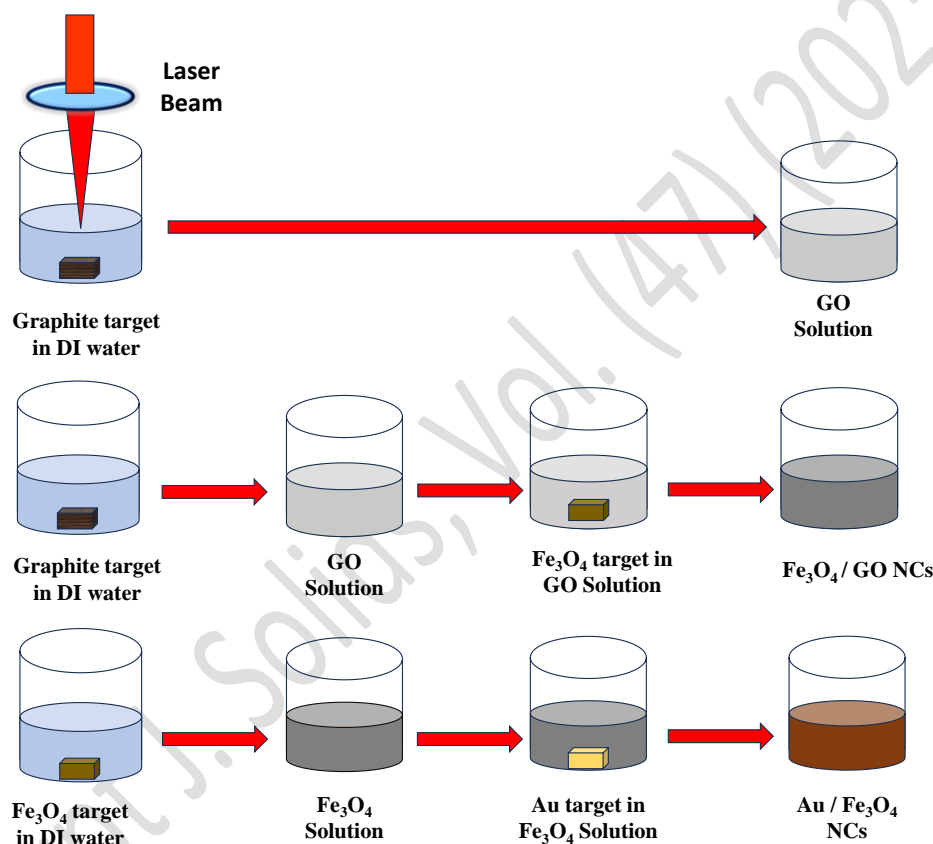
A layered carbon structure with functional groups including oxygen is called graphene oxide (GO). Graphene oxide (GO) contains various types of oxygen groups, including epoxide (=O) and hydroxyl (-OH) on its flat surface, and carbonyl (-C=O) and carboxyl (-COOH) on its edges, which enable it to form connections with a variety of organic and inorganic materials [1]. This happens through non-covalent, covalent, and ionic interactions, creating new hybrids and composites with special properties. The possibility of bandgap modification in GO is of interest for its execution in photonic and electronic devices [2]. Researchers found that GO nanosheets, prepared by pulsed laser ablation in

liquid (PLAL), are photoluminescent and biocompatible, making them suitable for bioimaging applications.  $\text{Fe}_3\text{O}_4$  is gaining a lot of attention because of its unique structural characteristics. The distribution of cations between its sites and their exchange interaction provides superior optical, electrical, electronic, and magnetic properties [3].  $\text{Fe}_3\text{O}_4$  is the most magnetic of all the naturally occurring metals on Earth; it has garnered a lot of interest; also, it is non-toxic and biocompatible. Moreover, noble metal nanoparticles, like gold (Au), have drawn a lot of interest due to their unique physical characteristics, especially their pronounced plasmon absorption peaks in the visible spectrum. The size, shape, and stability of the particles over time all affect the resonance frequencies. The demonstrated absorbance demonstrates a promising strategy to boost the efficacy of diverse cancer treatments. Researchers have invested significant effort in developing a multifunctional nanomaterial, aiming to achieve multiple properties in a single material [4]. These materials consist of at least two unique materials, each possessing specific properties from their respective segments [5]. There are several chemical methods available for obtaining nanomaterials. However, the purity of the finished nanomaterials—that is, their absence of impurities and precursor residues—is crucial for many applications [2]. Researchers have recently been actively exploring high-energy physical methods for nanoparticle production. Among these techniques is pulsed laser ablation in a liquid (PLAL). This method enables the direct production of particles in the form of colloids in pure liquids with either no precursors or very few [9]. Furthermore, it is an approach for micro-/nanostructure generation directly from bulk materials [7]. When the laser beam focuses on the surface of a solid target material in the surrounding media (gas or liquid), it rapidly raises the temperature of the irradiated spot, vaporizing the target material. When evaporating species (atoms and clusters) collide with nearby molecules, they create a laser-induced plasma plume by exciting the electron state, emitting light, and generating electrons and ions [3]. The target material, surrounding media, ambient pressure, and laser settings all influence the plasma structures, including the size of the plume and its emission spectrum. The process of laser ablation in liquid confines the plasma plume in a small region, thereby directly producing nanoparticles in the liquid phase [8]. In this work, the NGOs were prepared by Nd:YAG pulsed laser in 20 ml of deionized water for 60 min. Au/ $\text{Fe}_3\text{O}_4$ NCs were also prepared the same way and then loaded on NGOs to obtain Au/ $\text{Fe}_3\text{O}_4$ /GO NCs. The prepared samples were characterized by transmission electron microscopy (TEM), Fourier transform infrared spectroscopy (FTIR), and UV-visible absorption spectroscopy. The absorption curve we got was used to analyze the optical features of the samples. This includes the bandgap ( $E_g$ ), optical conductivity ( $\sigma$ ), absorption coefficient ( $\alpha$ ), Urbach energy ( $E_U$ ), electronegativity ( $\Delta x$ ), refractive index ( $n$ ), steepness parameter ( $\sigma_s$ ), electron-phonon interaction ( $E_{e-p}$ ), electronic polarizability ( $\alpha_e$ ), molar volume ( $V_m$ ), molar refraction ( $R_m$ ), and molar electronic polarizability ( $\alpha_{me}$ ).

## 1. Experimental

### a. Synthesis of GO, Fe<sub>3</sub>O<sub>4</sub>/GO, and Au/Fe<sub>3</sub>O<sub>4</sub>/GO NCs

We first put the graphite rod from Sigma Aldrich in an ultrasonic bath for 10 min and then we used deionized water to wash it. The rod was fixed at the bottom of 20 ml of distilled water, and then the Q-switched Nd:YAG laser with a wavelength of 1064 nm, pulse width (8-12) ns, repetition rate of 10 Hz and 450 mj was focused on the graphite target by a converging lens of 10 cm focal length for 60 min; a solution of GO nanosheets was simply formed.



*Fig.1. Shows the schematic diagram of the laser ablation technique.*

Fe<sub>3</sub>O<sub>4</sub> cube was supplied from (SAMMLUNG VON), and a high-purity Au slab (99.99%) from Sigma Aldrich was also cleaned in the same way as graphite. Fe<sub>3</sub>O<sub>4</sub>/GO has been obtained by immersing Fe<sub>3</sub>O<sub>4</sub> cubes in GO solution and ablating them by laser for 30 min. Also, Au/Fe<sub>3</sub>O<sub>4</sub> NCs were prepared by the same technique; first, Fe<sub>3</sub>O<sub>4</sub> cubes were immersed at 20 ml below the distilled water surface and radiated for 5 min, then Au slabs were fixed at the bottom of the Fe<sub>3</sub>O<sub>4</sub> solution and ablated for 45 min, resulting in the production of Au/Fe<sub>3</sub>O<sub>4</sub> NCs. The Au/Fe<sub>3</sub>O<sub>4</sub> and GO solutions were mixed by volume ratio 1:3 and sonicated for 30 min to accelerate the reaction to form Au/Fe<sub>3</sub>O<sub>4</sub>/GO.

Finally, the three samples were obtained: GO NPs,  $\text{Fe}_3\text{O}_4/\text{GO}$  NCs, and  $\text{Au}/\text{Fe}_3\text{O}_4/\text{GO}$  NCs. The experimental setup is shown in Fig. 1. All samples were prepared at ambient temperature and pressure.

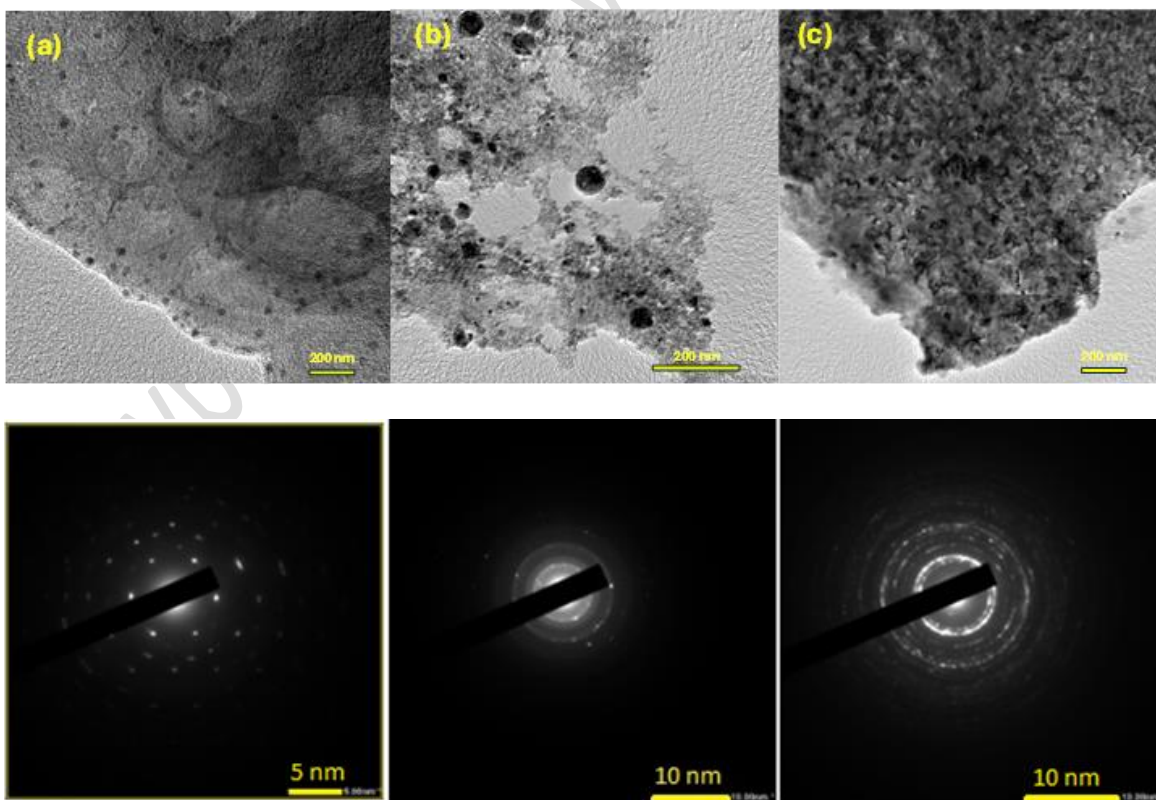
### b. Characterization tools

Nd:YAG laser from The Quanta-Ray® Lab series, the used model is (Quanta Ray) Lab-150. The size and morphology of NPs were identified using a high-resolution transmission electron microscope (HRTEM) (JEOL-JEM-2100 Plus, Japan). Scan images were recorded at an acceleration voltage of 200 kV. The functional groups present in the samples were identified using Fourier transform infrared (FT-IR) (Bruker ALPHA II, Germany), and the optical absorption spectrum of the samples was measured in the 200–600 nm wavelength range using quartz cells with a 1 cm optical path length using a JASCO 670 UV-vis spectrophotometer (Japan). Sonicator type EQ-VGT-1613QTD from MTI.

## 3. Results and discussion.

### a) Transmission electron microscopy

TEM images of prepared GO nanosheets,  $\text{Fe}_3\text{O}_4/\text{GO}$  NCs, and  $\text{Au}/\text{Fe}_3\text{O}_4/\text{GO}$  NCs are shown in Fig. 2 (a, b, and c), respectively.

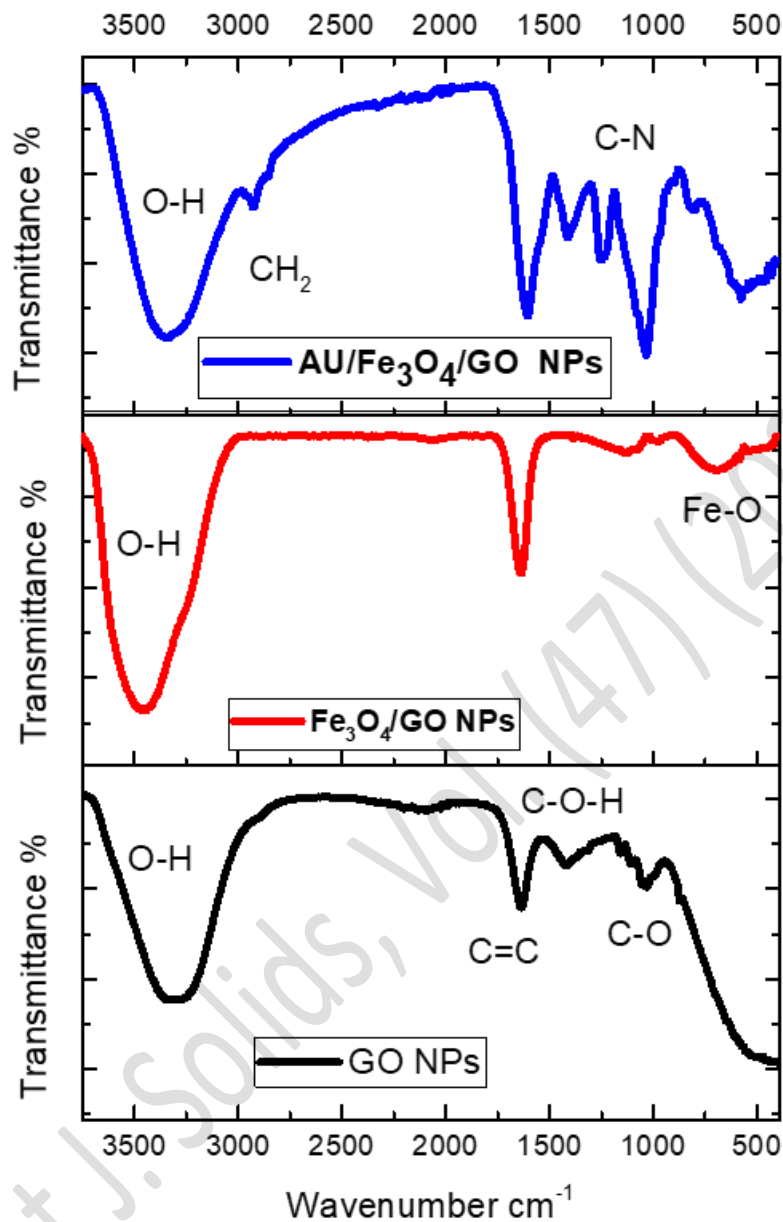


**Fig.2** Shows the TEM images of GO,  $\text{Fe}_3\text{O}_4/\text{GO}$  and  $\text{Au}/\text{Fe}_3\text{O}_4/\text{GO}$  NCs and SAED Patterns.

They reveal the crumpled nature; it's possible that some agglomeration happened when the samples were drying on the TEM grid. It is observed that GO was formed as an irregular thin layer. The non-deformed nanosheets help the charge carriers to transform fast with less energy, leading to a high interaction with the surrounding [4]. It is easily observed that some dots were found at Fig. 2(a) that were not observed at lower resolution, our result matching with [6]. In their work, laser ablation of a graphite target led to obtaining porous graphene sheets as primary products and graphene quantum dots (GQDs) as secondary products. We estimate the lattice spacing to be 0.29 nm. Our result is in good agreement with [7], where laser-ablated GQDs have a lattice spacing larger than chemically oxidized GQDs, which is nearly in the range of 0.3 nm. They attribute the grater lattice spacing of the laser-ablated GQDs [29]. The increased hydroxyl group relative concentration causes the carbon backbone gap to widen. Fig. 2(a1) shows the SAED pattern of GO which reveals the mono crystal structure. Fe<sub>3</sub>O<sub>4</sub>/GO NCs with different resolution are shown in the Fig.2(b). Fe<sub>3</sub>O<sub>4</sub> nanoparticles in a spherical shape have been loaded on the GO surface. A wide range of particle size distributions with average particle sizes nearly 51.64 nm are shown in Fig 2(b). Fig 2(c) illustrates that Au/Fe<sub>3</sub>O<sub>4</sub>/GO NCs, where Au/Fe<sub>3</sub>O<sub>4</sub> NCs diffused completely in GO sheets, and its shape is no longer recognized, which reveals the effect of sonication that accelerates the combination. This acceleration is due to either physical or chemical effects of cavitation [8]. Where Fig 2(b1, c1) show the SAED patterns for Fe<sub>3</sub>O<sub>4</sub>/GO and Au/Fe<sub>3</sub>O<sub>4</sub>/GO NCs multi crystal structure nature.

#### b) Fourier transform infrared spectroscopy (FTIR)

The FT-IR spectrum of GO, Fe<sub>3</sub>O<sub>4</sub>/GO, and Au/Fe<sub>3</sub>O<sub>4</sub>/GO is shown in the Fig.3. The three curves clearly display similar absorption bands. The broad band that starts from (3683) up to (2980) cm<sup>-1</sup> could be assigned to the absorbed water molecules over the nanoparticles [9]. GO Spectrum confirmed the presence of carboxyl COOH (1422 cm<sup>-1</sup>); the band at 1636 cm<sup>-1</sup> is assigned to the bending mode of the unsaturated carbon double bond (C=C), and the one at 1038 cm<sup>-1</sup> belongs to the stretching vibration of alkoxy C-O functional groups. For the Fe<sub>3</sub>O<sub>4</sub>/GO composite, it differs from that of GO by the disappearance of some oxygen groups, indicating some reduction that took place in graphene oxide sheets [10]. Moreover, the peak at (582 cm<sup>-1</sup>) assigned to the Fe-O stretching modes of the magnetite lattice and two peaks at (425 cm<sup>-1</sup>) and (682 cm<sup>-1</sup>) suggest a cation vacancy arrangement that is creating a reduced symmetrical structure, most likely because of surface oxidation [11]. These peaks suggest that the Fe<sub>3</sub>O<sub>4</sub> NSs were effectively decorated on the surface of GO. For Au/Fe<sub>3</sub>O<sub>4</sub>/GO, besides the GO bands and Fe-O bands, two peaks at (2938 cm<sup>-1</sup>) and (2852 cm<sup>-1</sup>) indicate C-H stretching alkane and C-H stretching aldehyde, respectively [22]. The weak bands at (1248 cm<sup>-1</sup>) and (1038 cm<sup>-1</sup>) show C-N stretching of aliphatic amines and C-O stretching of carboxylic amines, respectively [12]. It confirmed the successful attachment of Au nanoparticles to the Fe<sub>3</sub>O<sub>4</sub>/GO NCs.

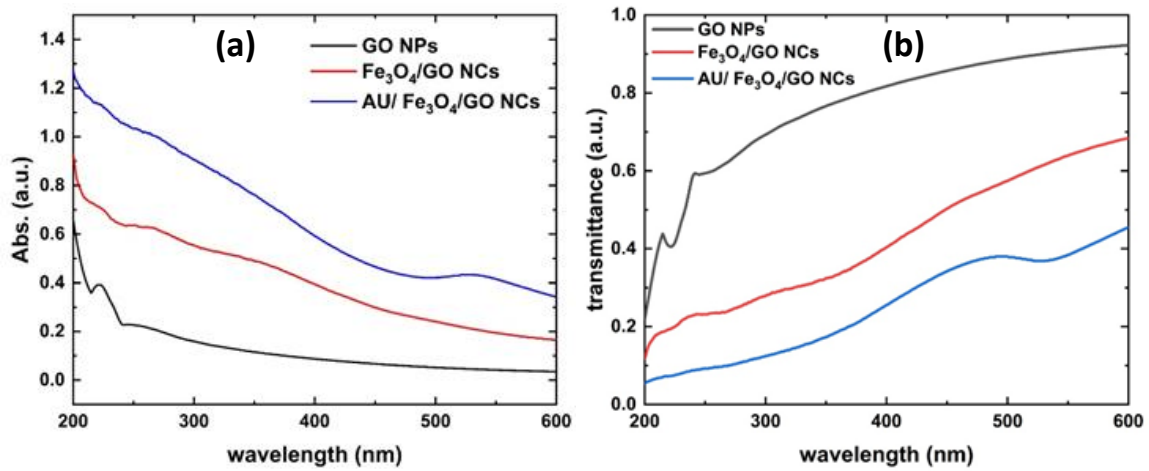


**Fig.3.** The FT-IR spectra of GO, Fe<sub>3</sub>O<sub>4</sub>/GO and Au/ Fe<sub>3</sub>O<sub>4</sub>/GO NCs.

### c) Optical properties studies

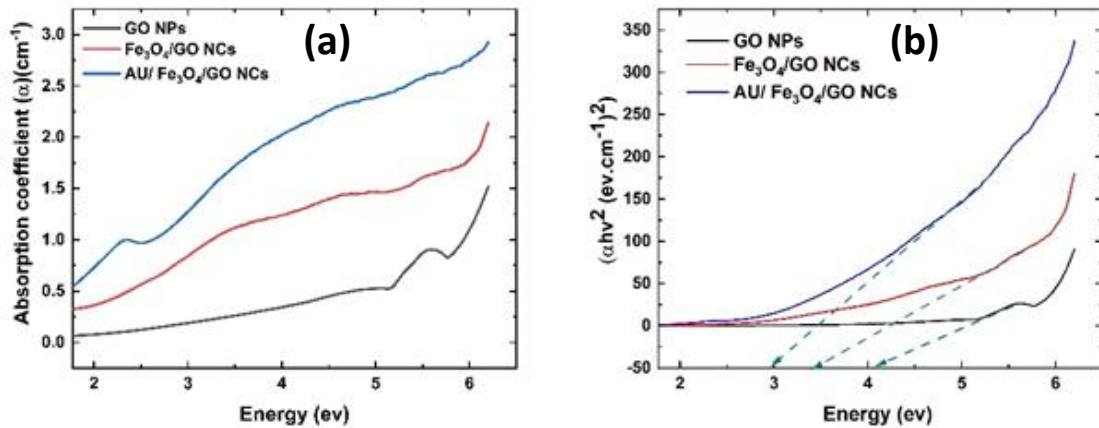
**Fig 4-a** represents the optical absorption intensity of GO, Fe<sub>3</sub>O<sub>4</sub>/GO and Au/ Fe<sub>3</sub>O<sub>4</sub>/GO NCs as a function of the wavelength.





**Fig.4, a) Absorption spectra of GO, Fe<sub>3</sub>O<sub>4</sub>/GO and Au/ Fe<sub>3</sub>O<sub>4</sub>/GO NCs b) Transmission spectra of GO, Fe<sub>3</sub>O<sub>4</sub>/GO and Au/ Fe<sub>3</sub>O<sub>4</sub>/GO NCs.**

It is observed that in the case of GO spectra, the characteristics peak at 222 nm due to the aromatic transition of  $\pi$ - $\pi^*$  belonging to the C-C bond, and another peak at nearly 265 nm corresponds to the  $n$ - $\pi^*$  transition [7]. Adding Fe<sub>3</sub>O<sub>4</sub> to GO and irradiating it with a laser again, the GO peak started to make a red shift of the absorption peak relative to the  $\pi$ - $\pi^*$  transitions, meaning a repairing of the  $sp^2$  hybridization, which means some reduction of oxygen groups [13], as well as the interaction between Fe<sub>3</sub>O<sub>4</sub> and GO. The absorption spectrum of Au/Fe<sub>3</sub>O<sub>4</sub>/GO NCs shows an increase in the absorption intensity; this is because under light excitation, localized surface plasmon resonance (LSPR) of gold dominates. The observed peak at 530 nm corresponds to surface plasmon of Au-decorated Fe<sub>3</sub>O<sub>4</sub> [14]. Fig 4(b) shows the transmission curve of GO, Fe<sub>3</sub>O<sub>4</sub>/GO, and Au/Fe<sub>3</sub>O<sub>4</sub> NCs; the figure reveals that the transmission decreases by adding Fe<sub>3</sub>O<sub>4</sub> NPs and Au/Fe<sub>3</sub>O<sub>4</sub> NCs. The absorption coefficient ( $\alpha$ ) of GO, Fe<sub>3</sub>O<sub>4</sub>/GO, and Au/Fe<sub>3</sub>O<sub>4</sub>/GO NCs as a function of incident light energy were calculated and given in Fig. 5(a) [15].



**Fig.5, a) Absorption coefficient curve of GO, Fe<sub>3</sub>O<sub>4</sub>/GO and AU/ Fe<sub>3</sub>O<sub>4</sub>/GO NCs. b) Variation of  $(\alpha hv)^2$  with the photon energy (hv) of GO, Fe<sub>3</sub>O<sub>4</sub>/GO and AU/ Fe<sub>3</sub>O<sub>4</sub>/GO NCs.**

It shows that the decorated GO nanosheets have a much higher optical absorption coefficient than pure GO nanosheets, which indicates the high absorption nature of the nano  $\text{Fe}_3\text{O}_4$  and  $\text{Au/Fe}_3\text{O}_4$ .  $\alpha$  and photon energy ( $h\nu$ ) are related to each other by Tauc's equation as [2]:

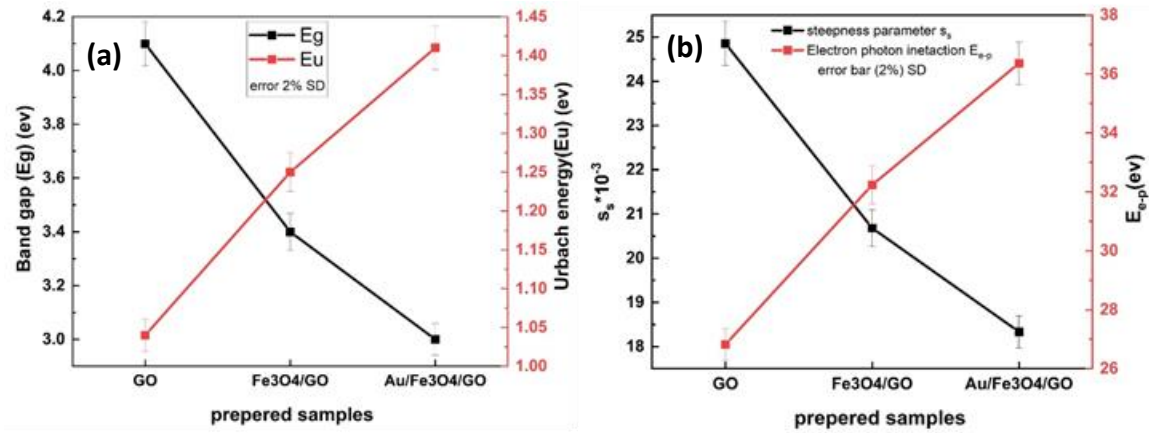
$$(\alpha h\nu)^{\frac{1}{n}} = \beta(h\nu - E_g) \quad (1)$$

Since  $\beta$  is a band tailing parameter,  $E_g$  is optical band gap energy, and  $n$  is the power factor of the transition mode. The range of  $n$  (0.3-0.8) indicates the electronic permitted direct transition [17]. The bandgap values are established by analyzing the Tauc-plot curve of the  $(\alpha h\nu)^2$  value on  $(h\nu)$ . By intercepting this linear part's extrapolation to the energy axis, we obtain the energy bandgap  $E_g$  [18]. Plotting  $(\alpha h\nu)^2$  against photon energy ( $h\nu$ ) for all the samples is displayed in Figure 5-b. The values of  $E_g$  for GO,  $\text{Fe}_3\text{O}_4/\text{GO}$ , and  $\text{Au/Fe}_3\text{O}_4/\text{GO}$  NCs are 4.1 eV, 3.4 eV, and 3 eV, respectively. The presence of different oxidation states of iron in the magnetite ( $\text{Fe}^{2+}$  and  $\text{Fe}^{3+}$ ) could also introduce specific impurity states, particularly in the electronic structure of the material, altering the conductivity or introducing new energy levels within the band gap. Also, interaction between the graphene oxide, magnetite and gold could lead to charge transfer between the three materials, creating localized states in the graphene's conduction or valence bands. The iron atoms in magnetite can also interact with the carbon atoms in graphene, leading to hybridized electronic states at the interface, which could behave as impurity states. Because of this, the doped material's bandgap energy narrows, and it absorbs photons with bandgap energies between 3 and 4.1 eV, which causes more excited electrons to remain inside the doped sheets' conduction band. Urbach energy is calculated by the following equation: [16][19].

$$\ln \alpha = \ln \alpha_o + \frac{h\nu}{E_U} \quad (2)$$

$\alpha_o$  is a constant and  $E_U$  is the band tail width (Urbach energy) of the localized states in the optical energy gap. We have determined the  $E_U$  by calculating the equation's inverse slope (2). Table 1 tabulates the value of the band tailing. The relation between  $E_g$  and band  $E_U$  is shown in Fig 6 (a) for the three samples. It is clearly observed that by the addition of nanoparticles to GO sheets, the weak bonds of oxygen are replaced by decorating material. Increasing in the  $E_U$  value ashore that a broadening in localized states in the forbidden gap led to a decrement in it [22].





**Fig.6.** a) Variation of the optical band gap ( $E_g$ ) and Urbach energy ( $E_u$ ) of GO, Fe3O4/GO and Au/ Fe3O4/GO NCs. b) Variation of both steepness parameter ( $\sigma_s$ ) and electron-phonon interaction ( $E_{e-p}$ ) of GO, Fe3O4/GO and Au/ Fe3O4/GO NCs.

The steepness parameter ( $\sigma_s$ ), which demonstrates the widening of the absorption band, can be calculated empirically using this equation: [20] [22].

$$\sigma_s = \frac{K_B T}{E_U} \quad (3)$$

We define  $k_B$  as the Boltzmann constant and  $T$  as the absolute temperature. Furthermore,  $\sigma_s$  determines how strongly the electron-phonon interacts ( $E_{e-p}$ ), and they related to one another by the relationship: [21]

$$E_{e-p} = \frac{2}{3\sigma_s} \quad (4)$$

The  $\sigma_s$  and  $E_{e-p}$  values were calculated using equations (3) and (4), respectively, and tabulated in Table 1. The plot of the three prepared samples versus the three prepared samples is shown in Fig. 6. The  $s$  decreases as the  $E_{e-p}$  increases. The total electronegativity difference ( $\Delta\chi$ ) is an important parameter to understand the chemical bonding. You can obtain the value of ( $E_g$ ) by substituting it into the Duffy relation: [21]

$$\Delta X = 0.2688 E_g \quad (5)$$

Where ( $\Delta\chi$ ) value decreased from 1.04 to 0.806 as calculated in table 1. Finding the refractive indices of materials is crucial, particularly for those that can be utilized to make any kind of optical device, such as switches, filters, modulation, etc. Many relations are presented between the refractive index ( $n$ ), ( $E_g$ ) and ( $\Delta\chi$ ) shown in the following relations [22][23]: The value decreased from 1.04 to 0.806 in Table 1.

**Table 1** contains the calculated values for the refractive index ( $n$ ). The average refractive index ( $n_{avg}$ ) was increased, assures the inverse proportional relation between  $E_g$  and  $n$ .

Also, materials with small  $n$  are transparent in the visible region where the sample's colour becomes gradually darker.

$$n = -\ln(0.027E_g) \quad (6)$$

$$n = \sqrt{\left(\sqrt[6]{\frac{5}{E_g}} - 2\right)} \quad (7)$$

$$n = -\ln(0.102\Delta X) \quad (8)$$

**Table 1:** Bandgap ( $E_g$ ), Urbach energy ( $E_u$ ), electronegativity ( $\Delta X$ ), refractive index ( $n$ ), steepness parameter ( $\sigma_s$ ), electron phonon interaction ( $E_{e-p}$ ) of GO, Fe<sub>3</sub>O<sub>4</sub>/GO and Au/ Fe<sub>3</sub>O<sub>4</sub>/GO NC respectively.

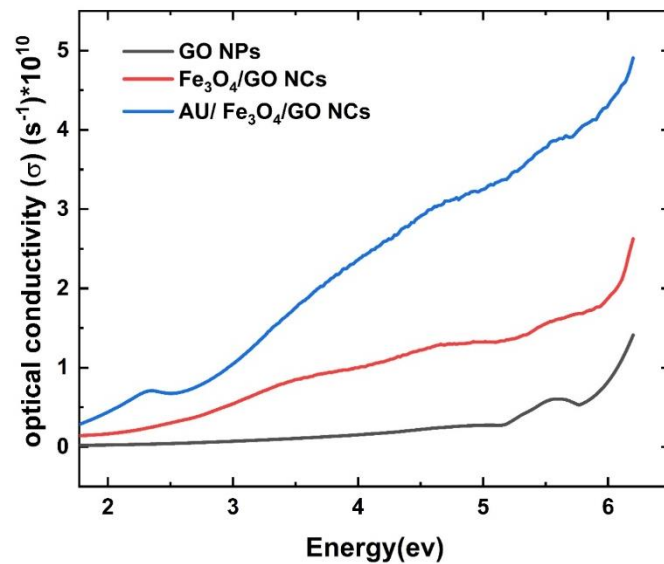
	$E_g$ (ev)	$E_u$ (ev)	$\Delta X$	Equation (n)			Average(n) ( $n_{avg}$ )	$\sigma_s$	$E_{e-p}$
Samples				Eq (6)	Eq (7)	Eq (8)			
GO	4.1	1.04	1.102	2.2	2.15	2.18	2.17	24.8	26.8
Fe <sub>3</sub> O <sub>4</sub> /GO	3.4	1.25	0.913	2.38	2.29	2.37	2.39	20.6	32.2
Au/ Fe <sub>3</sub> O <sub>4</sub> /GO	3	1.41	0.806	2.51	2.39	2.49	2.46	18.33	36.36

The optical response is studied for the samples in terms of optical conductivity calculated using the following relation:[3]

$$\sigma = \frac{\alpha n c}{4\pi} \quad (9)$$

Since ( $c$ ) is the speed of light in vacuum, **Fig.7** gives optical conductivity as a function of the photon energy. The optical conductivity is higher at larger photon energy due to higher electron excitation at this energy, and the conductivity of decorated GO sheets is higher than pure GO sheets, which reveals the high photo-response [24]. Using the values of  $n_{avg}$  obtained in **Table 1**, the electronic polarizability can be calculated by using in the following equation: [22] [25]

$$\alpha_e = \frac{3(n_{avg}^2 - 1)}{4\pi N_A(n_{avg}^2 + 2)} \quad (10)$$



**Fig.7,** optical conductivity spectrum of GO, Fe<sub>3</sub>O<sub>4</sub>/GO and Au/ Fe<sub>3</sub>O<sub>4</sub>/GO NC.

$V_m$  is the molar volume, can be evaluated by the electronic polarizability ( $\alpha_e$ ) and by substitution in the following relation: [4]

$$\alpha_e = 0.395 \left( \frac{n_{avg}^2 - 1}{n_{avg}^2 + 1} \right) V_m \quad (11)$$

$R_m$  is the molar refraction, which calculated based on the value of the average refractive index and molar volume by the Volf and Lorentz-Lorenz formula: [5][6]

$$R_m = \left( \frac{n_{avg}^2 - 1}{n_{avg}^2 + 1} \right) V_m \quad (12)$$

$\alpha_{me}$  is the molar electronic polarizability, which given by: [5]

$$\alpha_{me} = \frac{R_m}{2.52} \quad (13)$$

The obtained values of ( $\alpha_e$ ), ( $V_m$ ), ( $R_m$ ) and ( $\alpha_{me}$ ) are present in **Table 2**.

**Table 2:** Electronic polarizability ( $\alpha_e$ ), molar volume ( $v_m$ ), molar refraction ( $R_m$ ) and molar electronic polarizability ( $\alpha_{me}$ ) of GO, Fe<sub>3</sub>O<sub>4</sub>/GO and Au/ Fe<sub>3</sub>O<sub>4</sub>/GO NC respectively.

	$\alpha_e * 10^{-25}$	$V_m * 10^{-25}$	$R_m * 10^{-25}$	$\alpha_{em} * 10^{-25}$
Samples				
GO	2.20	8.55	4.75	1.88
Fe <sub>3</sub> O <sub>4</sub> /GO	2.38	8.71	4.83	1.92
Au/ Fe <sub>3</sub> O <sub>4</sub> /GO	2.49	8.80	4.89	1.94

It's clearly seen that by the addition of  $\text{Fe}_3\text{O}_4$  NPs and  $\text{Au/Fe}_3\text{O}_4$  NCs, the electronic polarizability increased. There is a general thought by which the electronic polarizability of samples containing oxide ion increases with increasing refractive index and decreasing energy gap [27]. The ( $V_m$ ), ( $R_m$ ), and ( $\alpha_{me}$ ) are proportionally related to ( $\alpha_e$ ), so they also increased.

#### 4. Conclusion

GO,  $\text{Fe}_3\text{O}_4/\text{GO}$ , and  $\text{Au/Fe}_3\text{O}_4/\text{GO}$  NCs were successfully prepared by a powerful green PLAL technique. The TEM image showed that the samples had uneven, flat layers of GO, which were evenly covered with  $\text{Fe}_3\text{O}_4$  particles that are about 51.64 nm in size. Finally, by adding  $\text{Au/Fe}_3\text{O}_4$  NCs to the GO solution, they diffused completely with the aid of sonication. The FT-IR spectrum confirms the functional groups of GO,  $\text{Fe}_3\text{O}_4$ , and  $\text{Au/Fe}_3\text{O}_4$ , as well as the bonds that have been built between them. The UV-visible spectrum indicates a red shift in the absorption edge of each of the three samples. The ( $E_g$ ) decreased from 4.1 to 3 eV and ( $\alpha$ ,  $E_U$ ,  $n$ ,  $\sigma$ ,  $\alpha_e$ ,  $V_m$ ,  $R_m$ ,  $\alpha_{em}$ , and  $E_{e-p}$ ) all increased for the three samples, respectively due to introducing new impurity energy levels, where ( $\sigma_s$  and  $\Delta x$ ) decreased. This shows that our structure is applicable for different applications, such as photonic applications, drug delivery applications, and water treatment applications [28].

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