The Effect of Laser Shots on Morphological and Optical Properties of Copper Oxide NPs Prepared by Nd-Yag Laser of 1064 nm Wavelengths in Distilled Water

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ABSTRACT
This study examines the synthesis of copper oxide nanoparticles (NPs) by using a Q-Switched Nd-Yag (1064 nm) laser on copper foil immersed in Distilled Water (DW). The solution color has changed to light green refers to the production of copper oxide NPs. The generated nanoparticles were studied to determine their characteristics as a function of pulse laser shots and the NPs were obtained by taking 500 and 1000 laser pulse shots on Cu target. Thin films deposited on both (glass and silicon) substrates were characterized by X-Ray Diffraction (XRD), Field Emission-Scanning Electron Microscopy (FE-SEM), and Atomic Force Microscopy (AFM) techniques. Later, regarding the colloidal nanoparticles, NPs were instantly characterized by UV-vis spectroscopy and examined by TEM microscopy. The production rate of Cu-NPs concentration in the colloidal solution was measured by atomic absorption spectra type (ICP-OES), which increased by increasing the number of laser shots in the liquid volume. The Energy Dispersion Spectroscopy (EDS) analysis resulted in the presence of copper (Cu) and oxygen (O) elements in the film structure with a nearly stoichiometry ratio. The optical energy gap was decreased to (2.44 eV) with increasing the number of laser shots in the colloidal solution.

Keywords: Copper oxide, Laser ablation, XRD, FE-SEM, TEM, Energy gap.

1. Introduction
Nanomaterials represent an important class of materials in the development of novel devices that can be used in various physical, biological, and biomedical applications, consequently. They are objects of active research in various fields of applications [1]. Recently, a number of researchers had worked on the evaluation of the biological activities of metal oxide NPs such as copper oxide from which improved biological and photocatalytic activities have been found [2, 3]. CuO NPs have attracted particular attention among various metal oxides because they are the simplest copper compounds that shows several unique physical properties such as high-temperature superconductivity and electron correlation effect [4]. The considerable antimicrobial activities of metal oxide NPs such as ZnO, Fe2O3, Fe3O4, MgO, CuO, TiO2, SiO2, have been proved at low temperature so the transition metal oxide NPs attract great interest to their notable antibacterial properties [5].

CuO NPs showed particular anticancer, antioxidant potency, and drug delivery which make them a promising tool for biomedical applications, in addition NPs prepared by the laser ablation in liquids have been lately used for manifold and inimitable applications for various fields of technology [6]. Lasers opened new physical entries into nanomaterial’s process and their characterization. Thus, this technique has various benefits with regards to other typical routes including a huge number of obtainable ablation input variables for controlling the size, shape, intrinsic stoichiometry, and the ability to generate nanomaterials with preferred chemical composition and surface free from chemical contaminations [7]. Since Pulse Laser Ablation in Liquids (PLAL) is a clean and green approach for the synthesis of metal as well as metal oxide nanostructures, it has become an increasingly popular approach for producing nanoparticles [8]. It is a promising technique for the controlled fabrication of nanomaterials via rapid reactive quenching of ablated species at the interface between the plasma and liquid with high-quality nanoparticles free from chemical reagents [9]. When the pulse laser beam is focused on the solid target immersed in DW, the laser immediately interacts with the bulk target and evaporates it into hot and dense plasma. The continuation of this process leads to nucleation and growth in the liquid and NPs will release as a

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result\(^{[10]}\). Therefore, PLAL process has received much attention as a novel NPs production technique. Regarding copper oxide, its superior nanoparticles system which inquires the size induced structured transmission and phase stability\(^{[11]}\). The simplicity of both the experimental setup and the data analysis has allowed the technique to become a widely used method for NPs generation. Copper oxide nanomaterials have attracted considerable attention among all the metal oxides due to their unique properties. In short, PLAL which combines the physical and chemical mechanism has been used as a rapid and clean approach to synthesize copper oxide NPs with different size and shapes from a pure copper foil immersed in DW. The synthesized colloidal NPs have been studied by using UV-visible spectroscopy and TEM image. Moreover, the deposited thin films were characterized by X-Ray Diffraction (XRD), and FE-SEM scanning microscopy. The size of copper oxide NPs was estimated by Scherer equation according to XRD results.

2. Experimental Procedure

2.1 Materials

Copper (Cu) foil with dimensions of 0.8 cmx 1.0 cm x 2 mm and a purity of 99.99 %, from (Alfa Aesar) company were used, and normally immersed in the bottom of a glass vessel filled with distilled water up to 8 mm of height which then irradiated with the focused output of wavelength 1064 nm from nanosecond pulsed Nd: YAG laser to generate the copper oxide nanoparticles in distilled water.

2.2 Production of Colloidal NPs

The fabrication of metal oxide NPs by Q-Switched Nd-Yag laser is an effective technique for NPs production. The copper oxide colloidal NPs of green color compared with distilled water, as in fig. (1) have been generated by letting the (500) and (1000) shots laser of energy 620 mJ to pass over the surface of bulk pure copper foil immersed in DW at the bottom of 5 ml glass vessel. The color of DW changed into light green by increasing the number of laser pulse to (500) shots, while the color of colloidal liquid turned to dark green with increasing the number of laser pulse up to (1000) shots indicating the increasing density of copper oxide NPs production in the solution. The rate of Cu NPs in the liquid was analyzed by atomic absorption spectra, type (ICP-OES, SPECTRO ACROS), for 500 and 1000 laser shots, the values were (21.35 and 31.42) mg/litter respectively. One drop of the colloidal has been dropped at the center of the piece of p-type Si (100) and left on the hot plate heater of 100°C for 5 minutes, then the prepared samples were tested for surface morphology by FE-SEM microscopy. Also, the same procedure has been applied for the synthesis of the film on a cleaned glass substrate and the films were tested by XRD. The colloidal liquid itself has been tested by TEM for both laser shots to find the shape and size of copper oxide NPs. Furthermore, optical properties have been studied by UV-Vis. 2600 spectroscopy.

![Figure 1: Copper oxide NPs: Left- dark green for (1000) shots Laser: Right -light green (500) shots laser.](image)

Results and Discussion

3. Structure Study

3.1 XRD Analysis for Thin Films Deposited on both Glass and Silicon Substrates

Copper oxide NPs synthesized by 500 laser shots have been dropped on the glass substrate at 100°C. Fig.(2) shows the XRD pattern of the film at positions of 2θ = 21.68° and 23.55° which do not belong to any copper oxide standard card, while the strongest peak is at 2θ = 43.77°, a broadening peak to be contributed to Cu (111) crystallography orientation and belongs to the reference code No.(00-001-1241). The value of full width at half maximum(FWHM) was 0.52 and by using Scherer’s formula\(^{[12]}\) the result of crystal size was 16.47 nm which refers to the poor crystalline film, due to low-temperature of film deposition.

![Figure 2: XRD pattern for Cu (111) thin film on glass substrate at 100 °C.](image)

When the film is synthesized by a drop of copper oxide colloidal NPs on p-type Si (100) at 100 oC, XRD pattern fig.(3) shows three diffraction peaks at 2θ= 25.220 which belong to Cu2O thin film according to the standard reference code (00-044-0706).
The FWHM value for the peak is 0.115° which indicates a good crystal grain size of 70.84 nm and agrees with previous studies \(^{[13]}\). The Si (100) also shows a similar XRD pattern to that grown on glass with a negligible difference 0.18°, but with a narrower diffraction peak, at a position of 2θ = 43.95°, which belongs to the CuO crystallography according to the reference code (00-044-0706). The crystal size was estimated based on FWHM value of 0.117° and from Scherer’s formula was 73.28°. This refers to a good crystallinity of the film deposited on Si substrate concerning glass when resulted in less than 17 nm for all grown peaks. The third prominent peak on Si substrate, at the position of 2θ = 77.39° belongs to Cu2O (222) thin film according to the reference code (01-078-2076) and ICSD collection code (063281), with cubic structure and lattice constant of a = 4.267Å which is connected with space group pn-3m. The results of thin films on p-type Si (100) substrate confirm the presence of both phases of copper oxide CuO and Cu2O, which refer to the presence of multi oxidation states.

While thin film deposited on glass substrate obtained under (1000) shots pulse laser and wavelength 1064 nm, fig.(4) shows pure polycrystalline CuO phase at angles 2θ = 35.32° and 38.54° in the plane directions (002) and (111) respectively, according to ICSD collection code (067850). As a result, the effect of the number of shots is nearly the same for crystalline growth on glass, while the effect of laser shots is very clear which confirms the presence of CuO phase structure with respect to the result of (500) shots pulse laser. The average crystalline size calculated was D = 13.048 nm, according to Scherer’s formula based on XRD pattern \(^{[14-16]}\).
of the NPs was produced in DW by 500 laser shots and energy of 620 mJ per laser pulse, the calculated diameters of the NPs were in the range of (10-40) nm. The result is compatible to a good approach with size of copper oxide NPs calculated by image-J program from the same TEM image which has mean size NPs of about 35.38 nm.

TEM images of the NPs grown by (1000) shots of pulse laser ablation have a corresponding size distribution of colloidal nanoparticles in distilled water as illustrated in the histogram of fig. (7). The average size of these spherical-shaped NPs is determined to be 15 nm. This reduction in the NPs size with increasing the number of shots which means elongated ablation time is attributed to the extended interaction of NPs with laser light, where higher energy absorption supports the fragmentation of bigger NPs into smaller ones [17].

Better morphology of deposited film on Si (100) might be because their surface atoms have unsaturated dangling bonds with one electron which cause the surface to be reactive and help the diffusion of Cu atoms into Si surface. Also, small agglomerated particles appear to constitute the larger particles in size.

4.2 AFM Analysis of Thin Film on Si (100) Substrate

In order to scan the thin film deposited on Si substrate, the resulting map of surface topography by AFM technique has been done by (Auto probe type, U.S.A with software: nanotech electronica) from Day Petronic company. Figure 9(a) shows the 3-D, AFM image of (5.0x5.0) µm and surface topography of the copper oxide thin film deposited on p-type Si (100) substrate at 100 °C.

4. Morphology Study and Effects of Laser Shots

4.1 (500) Shots of Pulse Laser in DW

Copper oxide NPs have been dropped on both glass and p-type Si (100) substrates at 100°C and synthesized by shooting the pure copper target in DW by (500) pulse of Nd-Yag laser with energy of 620 mJ in distilled water. The composed film was studied by FE-SEM. The surface morphology of the sample in figure 8 (a) with 2 µm scale bar shows paltry and continuous film distributions with indistinctive morphology on the glass surface and some small particles embedded on the surface. A few rounded particles ranged between 50 to 300 nm in size can be distinguished and this is maybe due to insufficient substrate temperature for film deposition. In Figure 8 (b), however, at 200 nm scale, a surface with various nanoparticles ranged approximately from 25 nm to 150 nm in size, but most of the particles happen to be identical in shape.
45.57 nm with some type of nanoparticle accumulation as can be seen from the distribution particles in figure 9(b). The particles ranged from 20 nm up to 180 nm in general, but most of the particle diameters distributed frequently in the range 60 nm to about 140 nm which is in total agreement with SEM particle size measurements for thin film deposited on Si (100) substrate.

4.3 (1000) Shots of Pulse Laser in DW

Copper oxide nanoparticles were synthesized by the same laser wavelengths of 1064 nm but doubled the number of laser shot pulses (1000) on the surface of pure copper target immersed in DW for the same pulse laser energy of 620 mJ. The colloidal NPs have been deposited on well-cleaned glass substrate at 100 °C for the sake of comparison with the surface morphology of thin film deposited on glass substrate at (500) laser shots. The SEM image in figure (10) shows perfect and neat copper oxide sphere shapes with diameters ranged approximately between 200 nm to 500 nm according to Image-J analysis. The change in morphology was found to be dependent on increasing the number of pulse laser shots up to 1000 pulses to produce copper oxide colloidal NPs. The formation of well-dispersed neat spheres and uniform crystalline structure with different diameters is extremely evident when compared with the films synthesized by 500 shots with the same laser energy on glass substrate which showed aggregated particles with round shapes. Also, the structure shows the aggregation of some nanoparticle materials.

The elemental composition of nanostructure materials for the thin film deposited on glass substrate was investigated and prepared by (1000) pulse laser shots on Cu target immersed in distilled water. EDS analysis of figure (11) shows the composition of the film which revealed the presence of copper (Cu: 76.3) wt % and oxygen element (O: 23.7) wt % respectively. The results support the atomic ratio distribution of 1:1 for CuO structure if compared with the theoretical calculated value from CuO molar mass, as shown in table (1).

![Figure 10: FE-SEM image of copper oxide deposited on glass substrate via (1000) laser shots of energy 620 mj.](image)

![Figure 11: EDS analysis of thin films deposited on glass by (1000) laser shots.](image)

**Table 1:** Elements weight percent (wt%).

<table>
<thead>
<tr>
<th>Elements</th>
<th>Cu wt %</th>
<th>O wt %</th>
</tr>
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<tbody>
<tr>
<td>Experimental value</td>
<td>76.3</td>
<td>23.7</td>
</tr>
<tr>
<td>Theoretical value</td>
<td>79.886</td>
<td>20.114</td>
</tr>
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absorption value at shorter wavelength side (UV-region). Fig. 12 spectrum(a) done by 500 laser shots shows one shoulder at 270 nm and the second position is a broadening peak extended from 640 to 720 nm. This may be due to the multi-phase of Cu-CuO-CuO in the colloidal solution. Figure 12(b) which has done by 1000 laser shots, possesses a maximum peak shifted towards a longer wavelength at 305 nm and terminates with the same behavior as curve (a), but holds an intense peak at 630 nm showing SPR band of Cu NPs, similar results reported elsewhere [18]. The SPR band is affected by particle size and in consequence redshifts it towards a longer wavelength in the visible region. The plot behavior also agrees with literature reports that Cu$_2$O strongly absorbs wavelengths below 600 nm. The absorption edge of the colloidal NPs at 360 nm for figure 12, plot (b) which is toward the shorter wavelength, indicates the higher optical energy band gap and is due to the band edge transition of Cu$_2$O [19]. Also, the NPs show the low absorption value (high transmission T, value up to 83%) on the higher wavelength side, which is a visible range and higher wavelengths up to 900 nm.

Figure 12: UV-VIS absorption spectra of copper oxide colloidal NPs by (a) 500 shots (b) 1000 shots.

The low absorption phenomenon can be explained by the energy of the incident photon which decreases at the higher wavelengths and cannot interact with atoms of the colloidal media. Thus, the photons will be transmitted instead of absorbing and the interaction of photons and the material with the absorbance of the photons can be observed [20].

5.1.3 Optical Energy Band Gap Eg Calculation

The optical band gap Eg of Cu$_2$O nanoparticles prepared by 500 and 1000 pulses of Nd-Yag laser in DW is calculated from Tauc's relation [21]. The absorption coefficient ($\alpha$) also has been calculated from the relation below which depends on the absorbance $A$:

$$\alpha = \frac{2.303A}{\lambda}$$

The optical band gap was carried out by linear extrapolation of the (a/\nu) versus (\nu) curve from the equation

$$\alpha (\nu) = A(\nu/\nu_0 - E_g)^m/2$$

Where $A$ is a constant, $\alpha$ is the absorption coefficient and $m$ equals 1 for a direct transition. Figure (13) shows the obtained energy band gap for the Cu$_2$O colloidal NPs in case of 500 pulse laser shots which is the direct optical band gap of 2.54 eV, similar results have been founded elsewhere [22].

Figure 13: (a/\nu)² vs. hv plot for Cu2O colloidal NPs by 1064 nm laser wavelengths and 500 shots.

In the case of 1000 pulse laser shots, the obtained optical energy gap for the colloidal copper oxide NPs was 2.44 eV, as shown in figure (14). It is evident that the results obtained by this method indicate that the $E_g$ of Cu$_2$O NPs is due to the direct allowed transition, and the results of optical band gap are in good agreement with the theoretical value and with previously measured values for Cu$_2$O band gap in the literature studies [23, 24]. The evaluated band gap is a little higher than the direct band gap value of bulk Cu$_2$O which is (2.0–2.17 eV). The blueshift in the value of direct band gap of the copper oxide is due to the decrease in particle size as a consequence of the appearance of quantum confinement effects [25].

Figure 14: (a/\nu)² vs. hv plot for Cu2O colloidal NPs by1064 nm laser wavelengths and 1000 shots.

6. Conclusion

Nd-Yag pulse laser technique on copper metal foil immersed in DW has been successfully used as a clean, safe and one-step physical method to produce the copper oxide colloidal NPs. It is found that the concentration of Cu NPs in the colloidal liquid increased with increasing the number of laser shots on the Cu target which produced dark green colloidal. Regarding the deposited copper oxide thin films structure and morphology, it seems to be influenced by the type of substrate (glass or silicon). TEM images of the copper oxide colloidal solution produced by (500) shots of laser show cluster NPs about (20-40) nm in size with semispherical shapes. However, in the case of (1000) shots, the TEM image and its histogram distribution shows clear change in size and shapes of NPs. The produced NPs were spherical with
an average size of about (15) nm. The size distribution and agglomeration of NPs were investigated in detail. The shape change of the NPs has also been proved from the FE-SEM images of the morphology of the thin films produced by 1000 pulses. The absorption spectrum of the colloidal NPs produced by 1000 shots results in a prominent SPR peak at a position of 630 nm which belongs to Cu metal NPs. The estimated optical band gaps were (2.54 and 2.44) eV. They decreased with increasing the number of laser shots, which most probably indicate the formation of cuprous oxide Cu$_2$O NPs. It is evident from the decreasing band gap values that the produced Cu and Cu$_2$O NPs possess bigger sizes of spherical shapes. The absorption spectrum of 1000 laser shots transpires the prominent peak at 305 nm which belongs to the inter band transition of electrons from valance band of CuO NPs. It is evident from the decreasing band gap of electrons from valance band of Cu$_2$O NPs. The results showed the possibility to control the optical energy gap as well as the particle size of the synthesized colloidal NPs by changing the number of laser shots.

References


