Enhancement of Photoconductive Detector Based on Carbon Nanotubes Decorated with Silver Nanoparticles by Adding Conductive Polymer

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Abstract

In this work, wide band range photo detector operating in UV, Visible and IR was fabricated using carbon nanotubes (MWCNTs, SWCNTs) decorated with silver nanoparticles (Ag NPs). Silicon was used as a substrate to deposited CNTs/Ag NPs by the drop casting technique. Polyamide nylon polymer was used to coat CNTs/Ag NPs to enhance the photo-response of the detector. The electro-explooding wire technology was used to synthesize Ag NPs. Good dispersion of silver NPs achieved by a simple chemistry process on the surface of CNTs. The optical, structure and electrical characteristic of CNTs decorated with Ag NPs were characterized by X-Ray diffraction and Field Emission Scanning Electron Microscopy. X-ray diffraction patterns of Ag NPs exhibited 2θ values (38.1°, 44.3°) corresponding to the Ag nanocrystal, while the XRD pattern of MWCNTs and SWCNTs/Ag NPs peaks appeared at 2θ = 26.2° corresponding to the (002) and at 2theta=44° which corresponds with miller indices (100) for CNTs and (200) for Ag NPs. The optical properties measured by UV-Vis. Spectroscopy. Broad and strong surface plasmon resonance (SPR) peak was detected at 420 nm, for Ag NPs. The absorption of CNTs/Ag NPs increased significantly from UV to near IR region (300-1000 nm). Ag NPs decorated CNTs without any impurities, according to field mission scanning electron microscopy examination, with typical particle sizes of (50-80nm) for Ag-NPs, 44nm for MWCNTs/Ag-NPs, and 30nm for SWCNTs/Ag NPs. The I-V characteristics at forward bias voltage (0.5-10) volt were studied. The figure of merits (responsivity, photocurrent gain, NEP and detectivity) after coating with polymer of the detector were measured in the dark and after illumination with UV LED (365 nm), Tungsten lamp (500-800 nm) and Laser diode (808 nm).

1. Introduction

In the past few years, there had been many significant studies on nano-crystalline materials due to their particular characteristics that bulk materials lack since nanoparticles have an extensive quantity of surface atoms than that of bulk [1]. Carbon nanotubes (SWCNTs & MWCNTs) and metal nanoparticles have various unique properties that make them useful in the field of optoelectronic and nanotechnology applications. CNTs have a wide range of thermal, electrical and structural properties [2, 3] Metal nanoparticles (NPs) have gained interest in the domains of electronics, chemistry, and biology due to their high surface-to-volume ratio and the size effect, which gives nanoparticles distinguishable features (optical, chemical, electrical, and magnetic) from their bulk material [4, 5]. CNTs decorated with metal nanoparticles...
open up new possibilities for researchers in a variety of domains, and are likely to have
different properties than individual CNTs, such as electrical, magnetic, and optical
capabilities [5]. Photodetectors play an important function in many applications of
modern life such as thermal imaging, night vision, medical and remote sensing, and
chemical testing [6]. Photoconductivity is the most generally used effect in which the
radiation incident on material changes to the electrical conductivity [7]. Previous studies
focused on the infrared (IR) response of CNTs, which explained that CNTs have
possible application in IR detector [8, 9]. To enhance the photo-response of the
photoconductive detector, the samples were coated with polyamide nylon polymer.

2. Experimental work

2.1. Synthesis of Ag-NPs by Electro-Exploding Wire (EEW)

This study developed simple and effective technologies for processing huge
counties of metal nanoparticles for wire explosions. Two electrodes were fixed in a
Teflon beaker; two electrodes were fixed in a Teflon beaker as shown Fig.1. Metal
plate was fixed on the bottom of the Teflon beaker, while a wire was placed opposite to
the metal plate. Thick copper wires were used to connect the positive terminal of the
battery 36V DC battery (welding machine) to the metal plate by an ohmic contact. The
negative terminal of the battery was connected to the wire to be exploded through
another ohmic contact by passing high current about 100A through it. (Fig.1). The
Teflon beaker was filled with 30 ml of distilled water, when just touched by silver wire
for 15 hits through precise mechanical movement, and then nominated by filtration
paper. The silver nanoparticles will be stuck in the solution.

Because of the high current density in the wire, the wire explodes in a short amount of
time. In the explosion process, silver wires with a purity of 99.99 % and silver plates
with (dimension: 2cm² area, 0.3mm thickness; purity: 99.99 %) were employed.

2.2. Decoration of carbon nanotubes (SWCNT and MWCNT) with Ag NPs

Multi-walled carbon nanotubes (MWCNTs) and single-wall carbon nanotubes
(SWCNTs) from (Nanostructured & Amorphous Materials, Inc.) were employed in this
study. The diameter of the MWCNTs ranged from 10-30 nm, length 1-2 µm and of 95%
purity, and the SWCNTs, its diameter ranged from 1-2 nm, of ~ 30 µm length and of
90% purity. Decorating CNT (SWCNT & MWCNT) with Ag NPs was carried out by
direct mixing. 0.05gm of CNTs (SWCNT & MWCNT) were dissolved in 10ml distilled
water, stirred for 1h and sonicated for half hour. 0.5 ml of Ag NPs was added to 2ml of
this solution stirred for 12h and sonicated for 1 hour.
2.3. Fabrication of photoconductive detector

Si wafer (p-type) of thickness (508+15 μm) and resistivity (1.5 Ω.cm) was used as a substrate of photoconductive detector. The samples of 1.5 x 1.5 cm² dimensions were cut from the Si wafer and rinsed in distilled water and ethanol to remove any dirt. Aluminium electrodes, the distance between the two electrodes was 0.4 mm, were made by evaporating Aluminium (Al) under vacuum with the help of a special mask deposited on surface of Si, as shown in (Fig.2).

![Figure 2: Schematic diagram for inter electrodes of photoconductive detector mask.](image)

CNTs/Ag-NPs solution was deposited on the electrodes by drop casting technique, the samples were left aside to dry at room temperature. In order to prepare polyamide nylon polymer solution (0.1 gm) of polyamide nylon polymer was dissolved in 5ml Tetrahydrofuran (THF). The solution was stirred for (10 min) then coated onto the samples by drop casting method. The samples were left in oven to dry. To determine the detector parameters mainly gain, responsivity (R), specific detectivity (D*) and NEP of the fabricated photoconductive detector on Si substrate by measuring the I-V characteristics, a suitable setup was prepared for this purpose. The system consists of: HUIER DC power supply (ps-1502DD), PC-interfaced digital multimeter (UNI-T UT803). Three sources of light were used for the illumination of the MWCNTs/Ag NPs and SWCNTs/Ag NPs by coating polymer, UV light with wavelength and power (365 nm, 10W) respectively, Tungsten lamp (500-800 nm, 250W) and laser diode (808 nm, 300 mW). The measuring circuit of the I-V characteristics circuit of the photoconductive detector is shown in (Fig.3).

3. Results and discussion

3.1. UV-Visible absorption spectroscopy

The UV-Visible absorption spectra of the pure Ag-NPs, SWCNT/Ag, and MWCNT/Ag prepared in water media were measured using Shimadzu UV-1800 spectrophotometer. (Fig.4) shows the UV-Visible absorption spectrum of Ag NPs as a function of wavelength. The spectrum reveals that Ag nanoparticles exhibit a fine structural absorption band in the UV-Visible region, with wavelengths ranging from 350 to 750 nanometers. The presence of a broad resonant absorption peak at around 420nm is due to the activation of silver nanoparticles' surface Plasmon resonance. According to (Fig.5) MWCNT/Ag and SWCNT/Ag NPs absorb light in the UV, Visible and NIR (300-1000 nm) ranges but the (SPR) peak is smaller than that of the Ag-NPs because of the induced charge transfer between (SWCNTs and MWCNTs) and Ag NPs, the number of electrons available for surface Plasmon generation in silver nanoparticles could be depleted [10, 11].
3.2. Infrared analysis spectroscopy (FTIR)

The functional group of MWCNTs/Ag NPs and SWCNTs/Ag NPs placed on glass substrates was identified using FTIR investigations in the 400 cm\(^{-1}\) to 4000 cm\(^{-1}\) range. Fig. 6 (a and b) shows the FTIR spectra MWCNTs/Ag NPs and SWCNTs/Ag NPs.

To find out how silver nanoparticles interact with carbon nanotubes (SWCNT & MWCNT) molecules. When the FTIR spectra of pure silver nanoparticles and CNTs were compared, it was discovered that the FTIR spectra of SWCNTs/Ag NPs and MWCNTs/Ag NPs contain multiple dips that are also present in the spectrum of pure Ag NPs and CNTs, but with different positions and transmission band intensities. It was slightly shifted to (2926, 1752) cm\(^{-1}\), proving the interaction between the CNTs (SWCNTs & MWCNTs) and Ag nanoparticles molecules. A broad peak appeared at the wavelength of (3461 and 3500 cm\(^{-1}\)) for MWCNTs/Ag NPs and SWCNTs/Ag NPs, respectively which were attributed to O-H stretching vibration. These results are in accordance with reported in the literature [12, 13]. The C=C bonding of aromatic rings of carbon skeleton structure were found at 1624 cm\(^{-1}\).

3.3. X-Ray diffraction

The crystalline structure of Ag nanoparticles, SWCNTs/Ag, and MWCNTs/Ag was studied using X-Ray diffraction. This approach was employed (XRD -6000 labs, supplied by SHIMADZU, with Cu K as the X-ray source), The XRD patterns of silver nanoparticles, MWCNTs/Ag NPs, and SWCNTs/Ag NPs samples are displayed in (Fig.7 (a and b)).
From Fig.7(a), it can be seen that the pristine Ag-NPs sample exhibits at 2θ=38.250° corresponding with miller indices (111), 2θ=44.353° with (200), 2θ= 64.767° with (220) and at 2θ=77.604° with miller indices (311). The crystalline planes of metallic Ag indicate that it is face-center cubic structure (F.C.C) (JCPDS No. 04-0783). After the decoration CNTs with silver nanoparticles, the XRD pattern for MWCNTs/Ag NPs and SWCNTs/Ag NPs revealed two obvious diffraction peaks at 2θ= 26.255° which correspond with miller indices (002) for CNTs (MWCNTs & SWCNTs) and 2θ=44.143° match with both miller indices (100) for CNTs and (200) for Ag NPs [14-16] as shown in (Fig.7 (b)).
3.4. Field Emission- Scanning Electron Microscopy (FE-SEM)

From (Fig. 8 (a)), the FESEM shows spherical clustered and luminous spots that correspond to the silver nanoparticles with the average particle size ranged between about (50-80 nm). The nanotubes were subsequently decorated with silver nanoparticles, (Fig. 8 (b) shows the FESEM of the Ag NPs decorated MWCNTs and SWCNTs. The luminous particles on the CNT surfaces are the silver nanoparticles. The nanoparticles on the surface of the CNTs are equally distributed, and the density of the attached nanocrystals is high, as shown in the image. Because metal nanoparticles are easily agglomerated, NPs are conjugated with carbon nanotubes to overcome problems with NP stabilization, separation, and recovery, as well as to avoid NP aggregation [5].

3.5. Hall Effect measurement

The Hall Effect was used to study the electrical properties of CNTs/Ag NPs deposited on silicon (charge concentration, conductivity, carrier mobility, Hall coefficient and resistivity). The Hall measurements parameters for CNTs/Ag-NPs deposited on Si are exhibited in Table 1. Table 1 shows that the electric conductivity of CNTs/Ag NPs become much higher than that of Ag NPs and the concentration of carriers increased when CNTs were decorated with Ag NPs. Also, it can be seen that CNTs decorated with Ag NPs were converted from n-type to – p-type when Ag NPs were added to CNTs, which results in a shift in the Fermi energy from the valence band to mid-gap. The electrical transport of CNT/Ag NPs was improved when the CNTs were decorated with Ag NPs. And when CNTs/Ag NPs deposited on silicon substrates is p-type semiconductor, the mobility and conductivity have better values.
Figure 8a: FESEM image of pure Ag NPs.

Figure 8b: FESEM images of SWCNTs/Ag NPs (right) and for MWCNTs/Ag NPs (left).

Table 1: Hall measurement parameters.

| Parameter          | Charge Concentration (1/cm³) | Conductivity (1/Ω.cm) | Mobility (Cm²/Vs) | Resistivity (Ω.cm) | Type |
|--------------------|------------------------------|------------------------|-------------------|--------------------|------|
| Ag NPs             | -1.08×10¹³                   | 2.36×10⁻¹              | 1.36×10⁵          | 4.23               | N-type |
| MWCNTs/Ag NPs      | 1.80×10²²                    | 8.83×10⁴               | 3.07×10¹          | 1.132×10⁻⁵        | P-type |
| SWCNTs/Ag NPs      | 1.99×10²²                    | 1.03×10⁴               | 3.22              | 9.708×10⁻⁵        | P-type |
| MWCNTs/Ag NPs/Si   | 9.85×10¹⁶                   | 9.91                   | 6.28×10²          | 0.1                | P-type |

3.6. I-V Characteristics for photoconductive detector

The current-voltage characteristics (I-V characteristics) of the fabricated photoconductive detector are illustrated in (Fig.9) at forward bias voltage of (0.5-10) volt. The linear I-V curve indicates the ohmic nature of the detector. The photocurrent increased under illumination with a tungsten lamp of (250 W, 500-800 nm), ultra-violet light of (10W, 365nm) and infrared light of (300 mW, 808 nm). For polyamide nylon polymer coated films the results showed that the photocurrent of the polymer coating on CNTs/Ag NPs /Si layer samples are higher than without the polymer. This polymer coating can be considered as a surface treatment of the detector film, which highly increases the photoresponse and specific detectivity of the fabricated detector. The (I-V) characterization of the fabricated photoconductive detector on silicon layer with and without coated polymer are shown in Fig. 9 (a, b and c).

Figure 9a: The variation of the UV photo-current of the fabricated photoconductive detector with coating polymer for SWCNTs/Ag NPs and MWCNTs/Ag NPs.
Figure 9b: The variation of the visible photo-current of the fabricated photoconductive detector with coating polymer for SWCNTs/Ag NPs and MWCNTs/Ag NPs.

Figure 9c: The variation of the IR photo-current of the fabricated photoconductive detector with coating polymer for SWCNTs/Ag NPs and MWCNTs/Ag NPs.

Figure of merits of the coating polymer and of the photoconductive detector for different source of light (UV, Visible and IR) are shown in Table 2. The photocurrent gain (G) which is the ratio between the photocurrent and the dark current for the same bias voltage (G = \(I_{pb}/I_{dn}\)), responsivity (R\(\lambda\) = \(I_{pb}/P_{in}\)), noise equivalent power (NEP= \(In/R_{\lambda}\)), Detectivity (D=1/NEP) and specific detectivity (D*\(=D/(A\Delta f)^{1/2}\)) were calculated. It can be observed from the results that the photocurrent of the polymer coating on SWCNTs/Ag NPs samples are higher than that of MWCNTs/Ag NPs in UV, Visible and IR radiation and have a good responsivity and photocurrent gain.

Table 2: Figure of merits photoconductive detector on silicon layer with coating polymer.

| Sample                | G  | \(R_{\lambda}\) (Amp/Watt) | NEP (Watt) | D (Watt\(^{-1}\)) | D* (Watt\(^{-1}\).Hz\(^{1/2}\).cm) |
|-----------------------|----|-----------------------------|------------|-------------------|-----------------------------------|
| **UV light**          |    |                             |            |                   |                                   |
| SWCNTs/Ag NPs        | 615| 8190×10\(^{-6}\)            | 0.019      | 52.6              | 52.6                              |
| MWCNTs/Ag NPs        | 2.6| 62.6×10\(^{-6}\)            | 4.5        | 0.22              | 0.22                              |
| **Tungsten lamp**    |    |                             |            |                   |                                   |
| SWCNTs/Ag NPs        | 654| 348×10\(^{-6}\)            | 0.06       | 16.6              | 16.6                              |
| MWCNTs/Ag NPs        | 4.2| 2.17×10\(^{-6}\)            | 95         | 0.01              | 0.01                              |
| **Laser diode (808nm)** |    |                             |            |                   |                                   |
| SWCNTs/Ag NPs        | 640.6| 12200×10\(^{-6}\)         | 0.025      | 400               | 400                               |
| MWCNTs/Ag NPs        | 6   | 1130×10\(^{-6}\)           | 0.18       | 5.5               | 5.5                               |
4. Conclusions

The wide band photoconductive detectors prepared by the drop casting technique on (p-type) silicon were fabricated and characterized to work in the UV, Vis, and IR regions. The decorating of CNTs (MWCNTs and SWCNTs) with silver nanoparticles synthesized by electro-exploding wire method was studied. It was found that the silver nanoparticles well dispersed on the surface of carbon nanotubes. The FTIR proved the interaction between CNTs and silver nanoparticles. The decorated carbon nanotubes with Ag NPs significantly improved optical absorption of CNTs. CNTs / Ag NPs absorbed light in the 300-1000 nm range better than CNTs alone, demonstrating the importance of noble metals in CNT optical absorption. The nanoparticles on the surface of the CNTs are rather evenly dispersed, according to the FESEM image. From Hall measurements, it was found that the decorating CNT surface with Ag NPs changed its characteristics from n-type to p-type. The experimental results showed that the CNTs/Ag-NPs can become essential building blocks for photoconductive detector fabricated on (P-type) silicon. The coating of the films surfaces with a polyamide nylon polymer improved the photocurrent gain, photo-responsivity, and specific detectivity in both SWCNTs/Ag NPs and MWCNTs/Ag NPs detectors. The photocurrent gain for SWCNTs/Ag NPs with coating polymer in UV, Vis, and IR light was better than photocurrent gain for MWCNTs/Ag NPs with coating polymer.

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Conflict of interest

Authors declare that they have no conflict of interest.

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تعزيز كاشف التوصيل الضوئي المعتمد على أنابيب الكاربون النانوية المزينة بجسيمات الفضة النانوية عن طريق إضافة بوليمر موصى

الخلاصة

في هذا العمل، تم تصنيع كاشف ضوئي واسع النطاق يعمل في نطاق الأشعة فوق البنفسجية والمرئية المزينة بجسيمات الفضة النانوية (SWCNTs، MWNTs) في النانوية CNTs / Ag NPs. حيث يمكن استخدام السيليكون ككبدة لترسيب (Ag NPs) النانوية CNTs / Ag NPs. تم تحضير CNTs / Ag NPs من مادة CNTs / Ag NPs الجزئية المزينة بالجسيمات الفضة النانوية باستخدام SPR (38.1°, 44.3°). المقابلة للبلازما النانوية النظيفة في حين أن قيم Ag NPs حيود الأشعة السينية من XRD (002) أظهرت قيم في 26.2° = θ تتوافق مع SXRD CNTs و MWNTs CNTs / Ag NPs من SPR (100) للنواقل.، و θ = 44° و التي تتوافق مع و (002) البريليوم (100) للنواقل.، و θ = 44°. و التي تتوافق مع العينات المزينة المزينة الضوئية CNTs / Ag NPs. والتي تفسر بالإضافة إلى الخصائص البصرية من SPR (002) التي تتوافق مع SXRD CNTs CNTs / Ag NPs في حين اثارت 220 من SPR CNTs / Ag NPs، والتي تتوافق مع SXRD CNTs و MWNTs CNTs / Ag NPs من SPR (100) البريليوم (100) للنواقل.، و θ = 44° و التي تتوافق مع و (002) البريليوم (100) للنواقل.، و θ = 44°. و التي تتوافق مع العينات المزينة المزينة الضوئية CNTs / Ag NPs. والتي تفسر بالإضافة إلى الخصائص البصرية من SPR (002) التي تتوافق مع SXRD CNTs و MWNTs CNTs / Ag NPs من SPR (100) البريليوم (100) للنواقل.، و θ = 44° و التي تتوافق مع و (002) البريليوم (100) للنواقل.، و θ = 44°. و التي تتوافق مع العينات المزينة المزينة الضوئية CNTs / Ag NPs. والتي تفسر بالإضافة إلى الخصائص البصرية من SPR (002) التي تتوافق مع SXRD CNTs و MWNTs CNTs / Ag NPs من SPR (100) البريليوم (100) للنواقل.، و θ = 44° و التي تتوافق مع و (002) البريليوم (100) للنواقل.، و θ = 44°. و التي تتوافق مع العينات المزينة المزينة الضوئية CNTs / Ag NPs. والتي تفسر بالإضافة إلى الخصائص البصرية من SPR (002) التي تتوافق مع SXRD CNTs و MWNTs CNTs / Ag NPs من SPR (100) البريليوم (100) للنواقل.، و θ = 44° و التي تتوافق مع و (002) البريليوم (100) للنواقل.، و θ = 44°. و التي تتوافق مع العينات المزينة المزينة الضوئية CNTs / Ag NPs. والتي تفسر بالإضافة إلى الخصائص البصرية من SPR (002) التي تتوافق مع SXRD CNTs و MWNTs CNTs / Ag NPs من SPR (100) البريليوم (100) للنواقل.، و θ = 44° و التي تتوافق مع و (002) البريليوم (100) للنواقل.، و θ = 44°. و التي تتوافق مع العينات المزينة المزينة الضوئية CNTs / Ag NPs. والتي تفسر بالإضافة إلى الخصائص البصرية من SPR (002) التي تتوافق مع SXRD CNTs و MWNTs CNTs / Ag NPs من SPR (100) البريليوم (100) للنواقل.، و θ = 44° و التي تتوافق مع و (002) البريليوم (100) للنواقل.، و θ = 44°. و التي تتوافق مع العينات المزينة المزينة الضوئية CNTs / Ag NPs. والتي تفسر بالإضافة إلى الخصائص البصرية من SPR (002) التي تتوافق مع SXRD CNTs و MWNTs CNTs / Ag NPs من SPR (100) البريليوم (100) للنواقل.، و θ = 44° و التي تتوافق مع و (002) البريليوم (100) للنواقل.، W. 1879(3): pp. 1-11.