

Vol. 31, No. 3, June, 2021, pp. 169-173

Journal homepage: http://iieta.org/journals/rcma

Novelty Au Nanoparticles with Different Nano Sizes as an Acidity Sensor

Zehraa Najim Abdul-Ameer

Department of Remote Sensing and GIS, College of Science, University of Baghdad, Baghdad 10001, Iraq

Corresponding Author Email: zehraanajim@gmail.com

https://doi.org/10.18280/rcma.310308	ABSTRACT
Received: 1 March 2021 Accepted: 30 April 2021 Keywords: gold, nanoparticles, size, acidity, sensor, reagent, Turkuvich method	The amazing wide range, applications of gold nanoparticles corresponding to excellent optical and electronic properties enhance utilization of Au NP as PH sensor. Gold nanoparticles were synthesized in simple and rapid Turkevich method to accomplish optical sensing technologies covalent or non-covalent, bonds formation causes, aggregation of the nanoparticles in the target substance. Au nanoparticles characterization was studied using XRD, UV-spectroscopy and SEM. XRD indicate polycrystalline. Nanoparticles with two different sizes according to reducing agent amount to obtain 20 30 nm. Optical energy gap was calculated for both particle size to find it (2.13) eV for 30nm particle size and (2.33) eV for 20nm.Scanning Electron
	Microscope showed spherical shape particles. PH test were performed to investigate

1. INTRODUCTION

Gold nanoparticles are widely used in many fields as preferred materials for their unique optical and physical properties, such as surface Plasmon oscillations for labeling, imaging, and sensing. The amazing, optical, properties of metal especially for interaction with the electromagnetic, visible light, this is due to its good resistance against the corrosion. These properties qualify Au nanoparticles to be utilized as PH sensor which is a fantastic [1, 2]. Gold nanoparticles attracted attention, due to its special, properties: the surface, Plasmon, resonance (SPR); and surface to volume ratio, besides high Surface Energy, quantum confinement and quantum dot [3, 4].

Abdulghani and Hussain [5] studied preparation of Au nanoparticles using Turkevich Method. Niu et al. studied the influence of citrate and Au concentration, reaction temperature and time, are parameters, to control, shape [6]. Sau et al. used the water phase oxidation-reduction method known as seed growth method to prepare small gold nanoparticles as seeds. Gold nano Particles can be synthesized with different sizes and, morphologies with spherical rod and triangular shape controlled by the crystal ratio to growth solution [7, 8]. Turkevich method for AuNPs synthesis was utilized for fabrication of SiO2@Au and TiO2@Au nanostructures. The nanostructures synthesized using this method were compared to SiO₂@Au and TiO₂@Au nanostructures obtained by deposition of Turkevich AuNPs on amino-functionalized SiO2 and TiO₂ corese significant red shift and broadening of the spectrum in the case of AuNPs decorated TiO₂ cores are likely related to high refractive index of TiO2 and dense packing of AuNPs on TiO₂ particles surface, resulting in strong AuNPs plasmon-plasmon coupling as studied by Yahya [9].

2. EXPERIMENTAL METHOD

efficiency of Au Np's for PH sensor and that is found to be excellent results for both sizes with more efficiency for 30 nm due to its rough surface. Which is a fantastic tool and measurement device to test the pH of soil, food, and other puncture-worthy applications.

Turkevich Method was used to synthesize Au Nano particles solution. 1 g of Chloroauric acid was dissolved in 50ml of deionized water with 58.86 mM molarity (Riedel-De Haen Ag / Seelz- Hannover, Germany) then withdraw 150 μ l of solute on the heat it up to 100°C to add 500 μ l Trisodium Citrate Dihydrate solution of 34 Mm (Panreac AppliChem / Bercelona, Spain) at once with continuous stirring. We will notice changes in the color of solution: yellow-clear white-light gray-dark gray-dark violet-red-wine, until it becomes red-wine, immediately stop heating and stirring and leave the solution to get cool.

3. TURKEVICH METHOD

One of the very well-known procedures for AuNPs synthesis is based on the diminishment of HAuCl4 by citrate in water, which was to begin with planned by Turkevich in 1951 as shown in Figure 1. In this strategy, the HAuCl4 arrangement is bubbled, and the trisodium citrate dihydrate is at that point quickly included into the bubbling arrangement under vigorous stirring. After a couple of minutes, the color of the arrangement changes from light yellow to wine ruddy. This strategy comes about in AuNPs measuring approximately 20 nm in breadth. In this procedure, citrate ions play a double part, as both stabilizing and diminishing specialists (Turkevich et al. 1951, Yahya 2017) [8, 9].

Puntes et al. have studied that a process with inverse sequence as addition of reagents (addition of HAuCl4 into a boiling sodium citrate solution) leads to the production of AuNPs with small size and a narrow size distribution as shown in Figure 2 [9, 10].



Figure 1. AuNP ion utilizing the Turkevich method in gold nanoparticle synthesis [10]



Figure 2. Direct and inverse sequence of reagent addition

4. CHARACTERIZATION OF Au NANOPARTICLES

4.1 XRD characteristics

Crystallographic information was analyzed from XRD to study structure and morphology using Shimadzu instrument, X-ray tube with target, Cukα radiation, voltage, 40.0 (kV), current 30.0 (mA) to calculate, grain, size average (D) was determined using Scherer's equation [11, 12]:

$$D=0.94\lambda/\beta COS\Theta$$
(1)

where, ($\lambda = 1.542$, Å) (CuK α), β is the full width at half, maximum, of the line (FWHM) and 2 θ scanning range was (30-80) as, in Table 1. Figure 3 shows XRD structures and for Au nanoparticles. From The pattern we can notice, that preferential, crystal at φ =38.1°, 44.3°, 64.5°, and 77.7°, corresponding, to (111), (200), (220), (311) characteristic peaks (Figure 3). Scherer's equation is used to determine average grain size (20, 30) nm.



Figure 3. XRD characteristic for Au nanoparticles

4.2 Optical properties

Shimadzu, Spectrum, of, Au, nanoparticles, of, range 190-1100 nm as shown in Figure 4, to study the optical, properties. Photoluminescence, at room, temperature was measured, using, spectrophotometer of Elico with range (300-900) nm wavelength, range. It was noticed that Energy gap values, have small difference due to agglomeration of these particles that as in Table 1. While Photoluminescence, were tested for, the, samples, using excitation wavelength of 320 nm, the spectrum. Figure 5 showed sharp peak for Au nanoparticles 20 nm with peak centered at 523 nm. Wide and broad peak noticed at (Au 30 nm) with SPR band centered at 560 nm. it is found that gold nanoparticles can be synthesized in a wide range of sizes, from 20 to 120 nm, with different characterization, taking in to account the previous work of Turkevich et al. [8], which results in numerous steps of essential and multiple steps of primary and secondary clustering leading to polycrystallites [13].

 Table 1. Structural and optical properties of Au nano, particles with two different sizes

	Eg (eV)	D (nm)	B (FWHM) (rad)	Δ	3
Au (30 nm)	2.13	22	0.32	0.002	188
(20 nm)	2.33	6	0.87	0.027	162







Figure 5. Photoluminescence intensity as function of wavelength

Maximum absorption wavelength at 523 nm was for 20 nm gold, nanoparticles, wine red color, the maximum adsorption, wavelength and, change in color (faint yellow, wine red, purple red, blue) with increase in, the particle size (30 nm) in the inter particle distance causes a red shift [14, 15].

4.3 Tunnelling Electron Microscope (TEM)

Figure 6 shows a HR Tunnelling Electron Microscope image for gold nanoparticles to measure the size of the particles. TEM images showed spherical shape gold nanoparticles and this agree with Zhao et al. [10]. TEM images of (20 nm) Au Nps with 500 μ l Trisodium Citrate Dihydrate solution.30 nm images showed more agglomeration with 300 μ l Trisodium Citrate Dihydrate causing larger particle size with moderate roughness.



Figure 6. Tunneling electron microscope: a-20 nm b- 30 nm

4.4 Acidity test procedure

Acidity measurement include several steps: setting the, PH/mV switch on the PH meter to the mV position then rinsing the electrode with (DI) water after that insert the electrode in the (PH=7) buffer for 30 minutes then record the polarity and the mV reading repeat the same steps for 4 PH buffer and the difference between the two mV readings was determined which, is the electrode span then divide the electrode's span by the theoretical span of 176.9 mV (at 250°C), then multiply by 100 and determine the slope of the electrode (Figure 7).



Figure 7. Measuring circuit

Figure 8 shows the acidity properties for both gold particle sizes. it is obvious from the figure that the Au np with 30 nm has better sensing properties [14, 15].

Recently, the chemicals internals of the sensors and the junctions that separate them from the external world come in a few variations material differences; but for the sake of application most sensors utilize silver chloride or potassium chloride and some types of ceramic junction. Increasing the robustness of these materials, or adding more junctions, would again be for more harsh applications and contaminated environments such as gold nanoparticles which is ideal with zero residues in the environment [16-18].



Figure 8. Current as function of PH: 20 nm, 30 nm



Figure 9. I-v characteristic properties of Au nanoparticles

One of the indications of efficiency of any sensor is the I-V characteristics as shown in Figure 9.

5. RESULTS AND DISCUSSION

Abdul-Ameer [13] research discussed different reasons for emission obtained from Au, nanoparticles of size 18 nm, and 13 nm. With (PLE) maximum occurs at approximately 255 nm which is about half the SPR peak position of 526 nm which is approximately close to our study. The optical properties of such materials depend on chemistry of their surfaces and on controlling of the dimensions of the crystal. Surface Plasmon (SP) is responsible in the colour of nanomaterial. As the volume, ratio increase the surface. Plasmon resonance, shift to the right or red shift occurred, the results agree with those of Tapan, K. Sau et al. [7] and become more broadening which indicates a different size or the size of Au, NPs increased while the size of Au, NPs become mono disperse when the volume ratio decreased, agree with the research results of Eustis and El-Sayed [18] and Shah et al. [19].

This research studied preparation of PH sensor which is found to have different sizes of Au Nps (20, 30 nm) with excellent optical properties for 20 nm in accordance to Au Nps with 30 nm in spite of that it was found that Au, Nps with 30 nm particle size have good sensing, properties due to roughness.

6. CONCLUSIONS

Gold nanoparticles were, synthesized in simple and rapid Turkevich method. Formation of covalent, or non-, covalent. Bonds with, the target, substance induce special optical sensing, technologies. characteristics indicate poly crystalline nanoparticles with two different sizes according to reducing agent amount to obtain 20, 30 nm with XRD, optical energy gap was calculated for both particle size to find it (2.13) eV for 30 nm particle size and (2.33) eV for 20 nm. Scanning Electron Microscope showed spherical shape particles .PH test were performed to investigate efficiency of Au Nps for PH sensor and that is found to be excellent results for both sizes with more efficiency for 30 nm due to its rough surface.

We conclude that we have, succeeded in synthesizing of Au, NPs, on the surface of, Psi using a cost-effective, and simple, method of Turkevich. The current density was calculated, of the deposition process to indicate, the size of the, formed Au NPs. These properties qualify Au nanoparticles to be utilized as PH sensor which is a fantastic tool and measurement device to test the pH of soil, food, and other puncture-worthy applications which is recorded for Au nanoparticles with 30 nm size due to agglomeration and increased roughness which is important for sensing process.

ACKNOWLEDGEMENTS

Ministry of science and technology and also laser institute for higher studies.

REFERENCES

- [1] Nagarian, R., Hatton, T.A. (2008). Nanpparticles: Synthesis, Stabilization, Passivation, and Functionalization Series. American Chemical Society. Washingtone.
- [2] Schmid, G., Corian, B. (2003). Nanoparticulated gold: Synthesis, structures, electronics, and reactivities. Eur. J. Inorg. Chem., 2003(17): 3081-3098. of https://doi.org/10.1002/ejic.200300187
- Turkevich, J., Stevenson, P.C., Hillier, J. (1951). A study [3] of the nucleation and growth processes in the synthesis of colloidal gold". Discuss. Faraday. Soc., 11: 55-75. https://doi.org/10.1039/DF9511100055
- Kimling, J., Maier, M., Okenve, B., Kotaidis, V., Ballot, [4] H., Plech, A. (2006). Turkevich method for gold nanoparticle synthesis revisited. J. Phys. Chem. B., 110(32): 15700-15707. https://doi.org/10.1021/jp061667w
- [5] Abdulghani, A., Hussain, R,K. (2014). Synthesis of gold nanoparticles via chemical reduction of Au (III) Ions by Isatin in aqueous solutions: Ligand concentrations and pH effect. Baghdad Science Journal, 11(3): 1201-1216.
- [6] Niu, W.X., Zheng, S.L., Wang, D.W., Li, X.Q., Han, S., Chen, J., Tang, Z.Y., Xu, G.B. (2009). Selective synthesis of single-crystalline rhombic dodecahedral, octahedral, and cubic gold nanocrystals. J. Am. Chem. Soc, 131(2): 697-703. https://doi.org/10.1021/ja804115r
- [7] Sau, T.K., Murph, C.J. (2004). Room temperature, highyield synthesis of multiple shapes of gold nanoparticles in aqueous solution. J. Am. Chem. Soc., 126(28): 8648-8649. https://doi.org/10.1021/ja047846d

- [8] Turkevich, J., Stevenson, P.C., Hillier J. (1951). Nucleation and growth process in the synthesis of colloidal gold. Discuss Faraday Soc., 11: 55-75.
- Yahya, Z. (2017). Plasmonic hot gold nanoparticles for [9] treatment of cancer cell line. PhD thesis, Baghdad University.
- [10] Zhao, P.X., Li, N., Astruc, D. (2013). State of the art in gold nanoparticle synthesis. Coordination Chemistry Reviews. 257(3-4): 638-665. https://doi.org/10.1016/j.ccr.2012.09.002
- [11] Chugh, H., Sood, D., Chandra, I., Tomar, V., Dhawan G., Chandra, R. (2018). Role of gold and silver nanoparticles in cancer nano-medicine. Artificial Cells, Nanomedicine, Biotechnology, and 46: 1210-1220. https://doi.org/10.1080/21691401.2018.1449118
- [12] Abdul-Ameer, Z.N., Agool, I.R. (2015). Structural and optical properties of ZnO electrodeposition method. Advances in Environmental Biology, 9(24): 445-449.
- [13] Abdul-Ameer Z.N. (2016). Effect of concentration on characterization of MgO nanoparticles using chemical bath method. Advances in Natural and Applied Sciences Journal, 10(12): 1998-1090.
- [14] Anwar, A., Minhaz, A., Khan, N.A., Kalantari, K. Afifi, A.B.M., Shah, M.R. (2018). Synthesis of gold nanoparticles stabilized by a pyrazinium thioacetate ligand: A new colorimetric nanosensor for detection of heavy metal Pd(II). Sens. Actuators B-Chem., 257: 875-881. https://doi.org/10.1016/j.snb.2017.11.040
- [15] Dong, H., Zou, F., Hu, X., Zhu, H., Koh, K., Chen, H. (2018). Analyte induced AuNPs aggregation enhanced surface plasmon resonance for sensitive detection of paraquat. Biosens. Bioelectron., 117: 605-612. https://doi.org/10.1016/j.bios.2018.06.057
- [16] Zhao, J.Y., Friedrich, B. (2015). Synthesis of gold nano particles via chemical reduction methods. NANOCON 2015. https://doi.org/10.13140/RG.2.2.28933.35049
- [17] Lin, H.Y., Chen, Y.F. (2005). Enhanced luminescence by second-harmonic surface plasmon resonance. Proc. Emerging Information Technology Conference, pp. 7803-9328. https://doi.org/10.1109/EITC.2005.1544349
- [18] Eustis, S., El-Sayed, M.A. (2006). Why gold nanoparticles are more precious than pretty gold Noble metal surface plasmon resonance and its enhancement of the radiative and nonradiative properties of nanocrystals of different shapes. Chem. Soc. Rev., 35: 209-217. https://doi.org/10.1039/B514191E
- [19] Shah, M., Badwaik, V., Kherde, Y. et al. (2014). Gold nanoparticles: Various methods of synthesis and antibacterial applications. Frontiers in Bioscience, 19(8): 1320-1344. https://doi.org/10.2741/4284

NOMENCLATURE

Eg	Enrgy band gap(eV)
D	Crystallite size(nm)

D	Crystallite siz	e(ni

Voltage(V) v

Current(A) I

Greek symbols

β	Full width at half maximum(rad)
٤	Micro strain

α	Absorption coefficient	np	nanoparticle
λ	Wavelength(nm)	XRD	Xray diffraction
θ	Diffraction angle(deg)	PL	Photoluminescence
ν	Frequency(hz)	PH	acidity
		SPR	surface Plasmon Resonance
Subscripts		HR	high resolution
-		TEM	tunneling electron microscope
Au	gold		