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ESTIMATION OF SIZE AND CONCENTRATION OF SILVER NANOPARTICLES IN AQUEOUS SUSPENSIONS FROM EXTINCTION SPECTRA

S.Z. Malynych

V.E. Lashkaryov Institute for Physics of Semiconductors NAS of Ukraine,
41, Nauky Avenue, 03028, Kyiv, Ukraine
E-mail: s.malynych@gmail.com

The size and concentration of silver nanoparticles suspended in water can be easily determined from their extinction spectra. This is due to the strong correlation of the localized surface plasmon resonances which are excited by electromagnetic radiation inside nanoparticles with size and shape of metal nanoparticles. Calibration curve obtained from Mie theory calculations for silver nanoparticles suspended in water is presented. A comparison with published experimental data is also discussed.

Keywords: SILVER NANOPARTICLES, COLLOID, EXTINCTION SPECTRA, LOCALIZED SURFACE PLASMON RESONANCE.

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1. INTRODUCTION

Growing interest in the physical properties of metal nanoparticles observed over the last decades caused by their wide potential applications in modern science and technology. These applications embrace single molecule surface enhanced Raman scattering [1], nonlinear optics [2], and photovoltaics [3]. The unique optical properties of noble metal nanoparticles, especially gold and silver make them an appealing platform for biosensing [4] and fluorescence-based biochips [5]. Recently, a colorimetric method with gold nanoparticles in DNA-dispersed solution, in which the plasmon absorption band of gold nanoparticles determined DNA hybridization, has been reported [6]. For most biomedical applications it is essential to determine the size and concentration of the nanoparticles dispersed in a carrying substance. Direct measurement of the nanoparticles size using TEM is expensive and time-consuming although it provides high accuracy in particle sizing. The methods based on dynamic light scattering are widely used but they are limited by a model of Brownian motion for spherical particles. At the same time there is no well-developed technique for concentration measurements of nanoparticles. In every-day laboratory practice it is desired to have low-cost, simple and fast method for the determination of nanoparticle size and concentration. Since the optical properties of metal nanoparticles are governed by its size and shape as well as the dielectric function of the particle and surrounding medium it is highly expected to use optical spectroscopy as an express tool for monitoring the particles size and concentration.

N. Khlebtsov in his thorough paper considered the correlations between particle size and their spectral properties based on published experimental data and his own T-matrix simulations [7]. He presented calibration curves

to determine the size and concentration of gold nanoparticles from the extinction spectra. It should be mentioned here that most papers considering analytical tools for the characterization of metal nanoparticle suspensions are mainly devoted to gold nanoparticles. This is due to low chemical activity of gold metal and well-developed nanoparticle synthesis technique. Among noble metals silver possesses the highest interband threshold that amounts 3,9 eV, while for Au and Cu it occurred at 2,4 and 2,1 eV respectively [8]. Thus, contribution of free electrons dominates in the optical response of silver nanoparticles. It means that the excitations in Ag nanoparticles have lower damping rates in compare to other noble metals, which makes them more favorable for photonic applications. Despite vast of works on physical properties and synthesis of silver nanoparticles there is lack of papers focusing on the experimental determination of size and concentration from their optical spectra. Systematic optical studies of Ag NPs ranging in size from 20 to 200 nm were performed by Chumanov and co-workers [9]. Here, we compare published experimental data on spectral properties of Ag nanoparticles depending on size and concentration with Mie theory simulations.

2. THEORETICAL CONSIDERATIONS

Optical response of metal nanoparticles results from the excitation of the conduction electrons coupled to the electromagnetic field. These modes are non-propagating and thus are termed Localized Surface Plasmon Resonances (LSPR). On the other hand, LSPR arise naturally from the scattering problem of a small, sub-wavelength conductive nanoparticle in an oscillating electromagnetic field. Surprisingly enough, a complete theory of the scattering and absorption of electromagnetic radiation by a sphere developed by G. Mie in 1908 does not use the notion of surface plasmons, but completely describes the distribution of electromagnetic radiation in far-field. Nowadays that theory is widely used and is treated in a variety of books.

According to Mie theory the extinction, scattering, and absorption cross-sections of spherical particles of arbitrary size can be calculated by series expansion of the internal and scattered fields into a set of partial waves described by vector harmonics:

$$\begin{aligned}
 C_{ext} &= \frac{2\pi}{|k|^2} \sum_{L=1}^{\infty} (2L+1) \operatorname{Re} \left\{ |a_L| + |b_L| \right\} \\
 C_{sca} &= \frac{2\pi}{|k|^2} \sum_{L=1}^{\infty} (2L+1) \left(|a_L|^2 + |b_L|^2 \right) \\
 C_{abs} &= C_{ext} - C_{sca}
 \end{aligned} \tag{1}$$

with

$$\begin{aligned}
 a_L &= \frac{m\psi_L(mx)\psi_L'(x) - \psi_L(x)\psi_L'(mx)}{m\psi_L(mx)\xi_L' - \xi_L(x)\psi_L'(mx)} \\
 b_L &= \frac{\psi_L(mx)\psi_L'(x) - m\psi_L(x)\psi_L'(mx)}{\psi_L(mx)\xi_L' - m\xi_L(x)\psi_L'(mx)},
 \end{aligned}$$

where $m = n/n_m$, n and n_m are the complex refractive index of the nanoparticle and real refractive index of the surrounding medium respectively, \mathbf{k} is the wave vector, $x = |\mathbf{k}|R$ is the size parameter, ψ_L , ξ_L are Riccati-Bessel cylindrical functions. The prime indicates differentiation with respect to the argument. The summation index L gives the order of the partial wave, $L = 1$ corresponds to dipole fields, $L = 2$ to quadrupole, $L = 3$ to octupole fields and so on. From the analysis of Eq. (1) follows that spectral position of plasmon resonance strongly depends on the size of a particle. Namely, Mie theory predicts a shift of LSPR towards longer wavelength with increasing the diameter of a particle. The red shift is a consequence of retardation effect when particle dimensions become comparable with the wavelength of light.

Two different processes take place when light travels through a suspension of small particles. They are absorption of light which leads to thermal losses and scattering or re-radiation of the incident light. Total losses that account for both components are termed the extinction of light. Optical transmittance defined as the ratio of the transmitted light intensity to the incident one is a usual measured quantity in most UV-vis spectroscopic experiments. Sometimes it is convenient to plot transmittance in a semi-logarithmic scale then the extinction would be a linear function of the optical path and/or concentration of a solution. For an ensemble of N non-interacting particles transmitted intensity follows Beer's law $I = I_0 \exp(-C_{ext}lN)$, where C_{ext} is the extinction cross-section of individual particle in cm^2 , l is the path length in cm, N is the particle concentration in cm^{-3} . Thus, the extinction of a suspension of nanoparticles can be defined as

$$E = 0,4343C_{ext}lN. \quad (2)$$

Hence, Mie theory provides a fundamental basis for the experimental determination of the size and concentration of nanoparticles from their extinction spectra.

3. RESULTS AND DISCUSSION

The calculations of the extinction cross sections of Ag nanoparticles where performed according to the Eq. (1) with the use of modified version of Mie solution code [10]. The dielectric functions of silver used in calculations were taken from [11]. Fig. 1 depicts the position of the dipole plasmon resonance as a function of particle diameter. The peak positions were determined from the calculated extinction spectra for the set of spherical nanoparticles of different size suspended in pure water ($n = 1,33$). The experimental data from [9b, 12, 13] are also shown. One can see that experimental peak positions for nanoparticles of the same size are shifted to the longer wavelengths in compare to those obtained from calculations. The shift is more notable for the particles smaller than 60 nm. The disagreements between calculated and experimental data can be explained by deviations of the particle shape from ideal sphere. Indeed, Mie theory deals with spherical particles while the nanoparticle shape obtained by chemical synthesis is rather polyhedral [9a, 12, 13]. Those shape deviations are more pronounced for very small particles and diminish with increasing the particle size. The influence of particle shape on their optical properties was studied in [14].

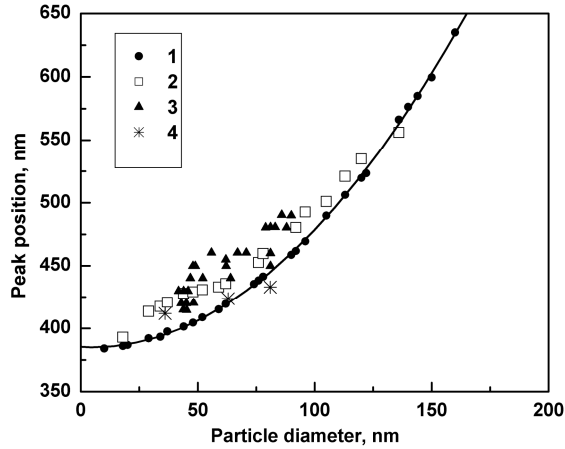


Fig. 1 – Position of the LSPR peak as a function of particle diameter for Ag nanoparticles in water. 1 – calculated data from Mie theory; 2, 3, 4 – experimental data from Refs. 9b, 12, 13 respectively. Solid line represents polynomial fit to the calculated data.

Decreasing of the nanoparticles size constrains the applicability of Mie theory due to additional collisions of free electrons with the boundary of the particle. As a result, dielectric function and plasmon peak positions for metal nanoparticles differ from those for bulk metal. Usually, only the imaginary part of the dielectric function is being modified to take into account size-limiting effects. According to [15, 16] small-size regime should be considered for the particles less than 10 nm in diameter. To avoid size and shape limitations several numerical methods have been developed. Discrete dipole approximation (DDA) is one of the most frequently used so far. In this method the particles of interest are divided into a cubic array of finite elements. The response of the whole array of elementary dipoles to an applied electromagnetic field is then described by self-consistently determining the induced dipole moment in each element, i.e. the dielectric properties of nanoparticle is directly related to the polarizability of the individual element. Jensen et al. have shown a good agreement of DDA and Mie theory results for silver nanoparticles with diameters ranging from 40 to 200 nm [17].

Polydispersity of silver nanoparticle suspensions embarrasses precise measurement of plasmon peak position introducing additional error to the determination of the particle size from the extinction spectra. Another plausible reason of the deviations of experimental data from calculated ones is the presence of chemical reaction products used during silver nanoparticle synthesis [12, 13]. The reaction products form a thin layer on the nanoparticles surface and can substantially alter their optical response.

Theoretical calibration curve is described by following empirical expression

$$d = \sqrt{24,01 + 100(\lambda_{\max} - 385)} + 4,9 ,$$

where d , λ_{\max} are the nanoparticle diameter and the LSPR wavelength, respectively, expressed in nanometers. From the discussion above follows that the calibration can be successfully used for the determination of the diameter of Ag nanoparticles in aqueous suspension when the latter exceeds

60-70 nm. For smaller nanoparticles the calibration allows only rough estimation of the particle diameter.

Evanoff and Chumanov developed a method to determine the concentration of silver nanoparticles which they named as standard subtraction [9b]. The method is based on the measurement of extinction spectra of Ag nanoparticle aqueous suspension prior and after the removal of a known number of particles. Clean glass slides modified with poly(vinylpyridine) were used to remove the particles from a suspension. The slides were immersed into a measured volume of colloid and exposed about 70 hours with gentle stirring. Due to the strong affinity of pyridyl group to metals the particles self-assembled on the surface forming a single layer with an average interparticle distance of 200-300 nm [18]. Total number of particles self-assembled on a glass surface was determined employing scanning electron microscopy. Then the concentration of particles in a suspension was calculated from the decrease of the extinction spectra. Extinction cross sections of individual nanoparticles were determined using expression analogous to the Eq. (2). Corresponding plot is shown in Fig. 2 along with the cross sections calculated from Mie theory.

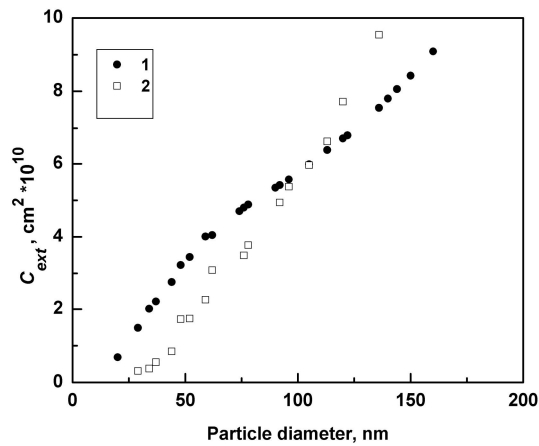


Fig. 2 – Extinction cross section of silver nanoparticles in water as a function of particle diameter obtained from Mie theory simulations (1), experimental data from Ref. 9b (2).

As it is seen from the figure that experimental cross section almost linearly depends on the particle diameter while the dependence obtained from Mie theory consists of two parts. Calculated dependence changes its slope at 60 nm which qualitatively corresponds to the experimental findings of Evanoff and Chumanov [9a, b]. They observed a maximum on the extinction efficiency curve for the nanoparticles of 50-60 nm in diameter. This is the size when phase retardation significantly contributes to the optical response of a nanoparticle and dominates over the increase of conduction electrons with increasing the particle size. Phase retardation results in reducing of the effective number of electrons that participate in plasmon resonance thus decreasing the incline of the extinction cross section curve. Reynolds and Stilwell [19] have shown that the mean free path of electrons in silver is about 52 nm which may also result in mentioned changes.

4. CONCLUSIONS

In this paper, we have shown that optical spectroscopy provides a simple analytical tool for the estimation of the size and concentration of silver nanoparticle in aqueous suspension. Calibration curve for the LSPR position and the extinction cross section vs. particle size are also presented along with corresponding empirical expression. Deviation of the nanoparticle shape from ideal sphere and extraneous chemicals on the nanoparticles surface causes disagreements between experimental data and the results obtained by Mie theory simulations. The disagreements become substantial for the particles less than 60 nm in diameter allowing rough estimation of size and concentration of Ag nanoparticles. Nevertheless, simplicity and express character of the described method makes it useful in the laboratory practice.

ОЦІНКА РОЗМІРУ ТА КОНЦЕНТРАЦІЇ НАНОЧАСТИНОК СРІБЛА У ВОДНОМУ ЗАВИСІ ЗІ СПЕКТРІВ ЕКСТИНКЦІЇ

С.З. Малинич

Институт фізики напівпровідників ім. В.Є. Лашкарьова НАН України,
просп. Науки, 41, 03028, Київ, Україна
E-mail: s.malynych@gmail.com

Розмір та концентрацію наночастинок срібла, завислих у воді, можна легко оцінити за їх спектрами екстинкції. Це можливо завдяки сильній кореляції локалізованих поверхневих плазмонних резонансів, що збуджуються у наночастинках електромагнітним випромінюванням, із розміром та формою металевих наночастинок. Представлено калібрувальну криву для наночастинок срібла у воді, отриману із розрахунків згідно з теорією Мі. Проведено порівняння з опублікованими експериментальними даними.

Ключові слова: НАНОЧАСТИНКИ СРІБЛА, КОЛОЇД, СПЕКТРИ ЕКСТИНКЦІЇ, ЛОКАЛІЗОВАНИЙ ПОВЕРХНЕВИЙ ПЛАЗМОННИЙ РЕЗОНАНС.

ОЦЕНКА РАЗМЕРА И КОНЦЕНТРАЦИИ НАНОЧАСТИЦ СЕРЕБРА В ВОДНОЙ ВЗВЕСИ ИЗ СПЕКТРОВ ЭКСТИНКЦИИ

С.З. Малинич

Институт физики полупроводников им. В.Е. Лашкарева НАН Украины,
просп. Науки, 41, 03028, Киев, Украина
E-mail: s.malynych@gmail.com

Размер и концентрацию взвешенных в воде наночастиц серебра можно легко оценить из спектров экстинкции. Это возможно благодаря сильной корреляции локализованных поверхностных плазмонных резонансов, возбуждаемых электромагнитным излучением, с размером и формой металлических наночастиц. Представлена калибровочная кривая для оценки размера наночастиц серебра, полученная из расчетов согласно теории Ми. Проведено сравнение с опубликованными экспериментальными данными.

Ключевые слова: НАНОЧАСТИЦЫ СЕРЕБРА, КОЛЛОИД, СПЕКТРЫ ЭКСТИНКЦИИ, ЛОКАЛИЗОВАННЫЙ ПОВЕРХНОСТНЫЙ ПЛАЗМОННЫЙ РЕЗОНАНС.

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