Comparative Observation of Silver Nano and Microstructures Deposited from Aerosol and Fog

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Abstract. A comparative study of the structure and fractal properties of arrays of the silver nano-/micro-particles deposited on the silicon substrate both from the aerosol and fog showed that the form of the silver individual particles and nano-/microstructures greatly depends on the deposition conditions. By passing an aerosol through isopropyl alcohol, the formation of fractal aggregates of the silver nano-/micro-particles both in the air and in alcohol was observed. Deposition of the silver nano-/micro-particles in the atmosphere of the saturated isopropyl alcohol vapours led to formation of fog. Micro-droplets of the silver colloidal solution were deposited on the substrate. The further evaporation of alcohol created the silver nano-/microstructures in the form of annular layers. It was found that the concerned annular layers contained silver particles of the same shape in the form of a Crescent (or Janus-nano-/microparticles). The nature of discovered effects is discussed.

1 Introduction

It is known [1], that anomalous optical properties of the metal nano-/microparticle (NMP) arrays can be determined by the interaction of light with the surface plasmon polaritons and localized plasmons. The structure and form of the metallic NMP are the important factors determining the efficiency of the concerned “plasmon-photon” interactions. Theoretical analysis [2] [3] [4] showed that the sickle shaped metallic NMP can enhance the interaction of localized plasmon with light photons, compared to the NMP having the spherical shape. However, the proposed method of the sickle metallic NMP synthesis by means of the colloidal microlithography method [5] is multi-stage, and requires complicated hardware design. So the studies aim to simplify the method of the sickle metallic NMP synthesis.

The disadvantage of the newly proposed method of the metallic nano-/microstructures (NMS) layers synthesis by the “dry” aerosol deposition [6] consists in the random ballistic mechanism of coatings formation on the substrate surface. These coatings contain both the NMP of different sizes and the NMS of chain and dendritic shapes that can lead to significant scatter of the optical and other properties of the resulting porous metal films [7]. Therefore it is desirable to develop a method of the metallic NMS formation in the controlled conditions which could provide an opportunity for receiving the NMS of the same shape and, preferably,

containing the arrays of the metallic NMP having the same geometric shape. Adherence to conditions allowed achieving the stability and reproducibility of the optical properties of the metallic NMS.

A promising approach to solve this problem may be the creation of the metallic NMS precipitates formed during the evaporation of the micro-droplets from the metallic NMP colloidal solutions in various solvents [8], [9].

In the given case, depending on the conditions of the evaporation process [10], the metallic NMP deposits can exist either in the form of solid spots (this situation is considered in the “coffee stains” model [8]), or form of ring structure.

According to the research [8] [9] [10], the concerned method of the metallic NMS preparation can have many practical applications such as production of plasmon resonance sensors, catalytic surfaces, etc. The different forms of the received metallic NMS are due to the occurrence of the sedimentation processes in the highly non-equilibrium conditions which lead to the development of the self-organization. In that case the shape of the NMS deposits is determined by the deviation degree of the evaporation conditions from the equilibrium [10]. Usually the colloidal NMP solutions can be considered as the micro-emulsions which can be formed and stabilized by the surfactants. Then it is necessary to wash off the NMP sludge from the surfactant and emulsifier admixtures that can result in the deformation and the NMP damage. This problem can

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be solved if the deposition of the aerosol metallic particles on the silicone surface has been carried out in the atmosphere of the solute saturated vapors. Metallic NMP is the center where the condensation of the solute molecules and micro-droplet formation is taking place. Finally, the initial aerosol of the metallic particles transforms into fog [11]. After that, the evaporation of micro-droplets deposited on the silicone surface forms the metallic NMS, which do not contain the surfactant and emulsifier admixtures.

The aim of present study consisted in:

a) the experimental confirmation of the possibility to form the ring metallic NMS synthesized by the evaporation of micro-droplets deposited on the substrate surface from the fog formed as a result of the interaction between the aerosol NMP and the solvent molecules;

b) the study of the structure and shape of metal NMP contained in the NMS, which formed either after the micro-droplet deposition and evaporation, or the passing of aerosol NMP through the solvent media;

c) conducting the fractal analysis of images that show the metal NMP shape and the structure of the metal NMS synthesized in both modes, obtained by transmission and scanning electron microscopy (TEM and SEM, respectively), and atomic force microscopy (AFM).

2 Experimental

To obtain the colloidal solutions of silver NMP in isopropyl alcohol (IPA) without the admixture of the emulsifier and surfactant molecules the self-made experimental facility [6] has been used (Fig.1(a)). The scheme of the concerned facility includes an aerosol spark generator, a diffusion aerosol spectrometer (DAS), the ultrasonic generator of the liquid and the aerosol flow stabilizer. After mixing the silver aerosol and the IPA saturated vapour the condensation processes were developed in the volume of the aerosol flow stabilizer and the micro-droplets of the colloidal solution appeared. Then these micro-droplets were deposited on the surfaces of three silicon wafers located at the output of the aerosol flow stabilizer. IPA vapours were obtained using the ultrasonic generator. For that purpose an IPA media with the chemical grade "pure for analysis" (State Standard 9805-84) was placed in ultrasonic bath before switching on the ultrasonic generator. Experiments were carried out at room temperature and at the atmospheric pressure. The IPA saturated vapour pressure was ~ 25.3 mm Hg [12], however not measured or fixed specifically. The silicon plates with dimensions 10×10 mm and the thickness 0.5 mm (State Standard 8.592-2009) were used in research. The duration of the separate experiment did not exceed one hour.

The silver NMP aerosol was created with the help of the spark discharge between two silver electrodes and an air stream passed at the speed of 1 cm/s through the working chamber of the aerosol spark generator [6]. The frequency of the spark breakdown was 50 min⁻¹. Then the silver aerosol was transported to the ultrasonic generator and to the DAS measuring system, which

determined the size distribution and concentration of the silver NMP in the aerosol (Fig. 1(b)).

During the experiment the composition of the silver aerosol particles was controlled every minute. It has been established that the largest part of the aerosol composition was about 45 - 50 nm (Fig. 1(b)-1). The total number of the silver aerosol particles received in the volume of the working chamber during the experiment was about $\sim 5.2 \times 10^6$. As it is shown in Fig. 1(b)-2, the particle size distribution can be satisfactorily approximated by lognormal function [11], [13].

Fig. 1. (a) Technological scheme of the silver NMP micro-droplets deposition: 1—micro-pump; 2—high efficiency particulate air filter; 3—aerosol spark generator; 4—diffusion aerosol spectrometer; 5—ultrasonic generator; 6—aerosol flow stabilizer; 7—silicon substrate. Black dots represent aerosol particles of silver; (b) Histogram (I) and the aerosol silver particle size distribution function (II) obtained by the approximation of experimental data by the lognormal distribution.

The colloidal solutions of silver NMP were also prepared by passing the aerosol through the IPA medium. Before carrying out the TEM and SEM measurements the colloidal solutions of silver NMP in IPA were treated in an ultrasonic bath for 10 minutes. Then the sample of solution (about ~ 0.1 ml) was deposited on the copper mesh modified by carbon. Investigation of sludge formed on the silicon surface and carbon substrate after evaporation of the colloidal solution of micro-drops was conducted using TEM and SEM electronic microscopes FEI Tecna G2 F20 S-Twin TMP and Quanta-3D FEG (Fig. 2 and Fig. 3 respectively). TE-microscope FEI Tecna G2 F20 S-Twin

TMP worked at an accelerating voltage of 200 kV in bright field. The device resolution was 0.14 nm (line resolution).

Fig. 2. Images of the obtained silver nano-/microstructures on the copper-carbon substrate at different magnifications: 9900 $\times$ (a); 38,000 $\times$ (b); 97,000 $\times$ (c); and 450,000 $\times$ (d).

It should be noted that the study of sludge which formed after evaporation of the micro-drops received by the second method was carried out during half an hour after it had been placed on a copper-carbon substrate. When the micro-drops were received using the first method, the TEM- and SEM-measurements were performed after the storage of the precipitates in the air and at the room temperature within two weeks.

AFM investigation of the silicon surface roughness after deposition of the silver NMP arrays was carried out by Multimode 8 scanning probe microscope, manufactured by Bruker (USA). The concerned array of AFM data was further processed using Image J 1.49 [14] software with the help of the Fraclac plug-in [15].

3 Results and Discussion

3.1 Silver Nano-/Microstructures Deposited from Aerosol

As follows from the TEM images obtained in the first case the silver aerosol passage through IPA media resulted in intense NMP coagulation and formation of the different NMS types (Fig. 2). In that case the formation of the numerous silver NMS resulted in the appearance of the porous silver film [6], that is an array of the randomly linked chains consisting of several (at least 10) silver NMP with sizes in the range of 10 - 200 nm (Fig. 2(a)-(c)). The facets of silver nano/microcrystals can be observed on the NMP surface (Fig. 2(d)).

It is known [16], that the chain structure of NMS occurs as a result of the silver NMP coagulation in the air. Possibly, coagulation can take place in the IPA medium. The secondary structure observed in Fig. 2(a) and Fig. 2(b), consists of chaotically intertwined chains

containing the numerous silver NMP's. Such structure occurs both at the stage of the storage in alcohol, and the evaporation of nano-/micro-drops. The emergence and growth of the secondary structures (or fractal aggregates [17]) is a result of the aggregation of the numerous silver particles in chains, and it can be described in terms of different models of aggregation [18]. The estimation of the fractal dimension for layers of the silver NMS located in the regions with high and low NMP concentrations (Fig. 2(a) and Fig. 2(b)) gives the values of the surface fractal dimension $D = 1.85$ and 1.99 which correspond to the case of Brownian motion and/or the motion of NMS's on a linear trajectory in the model of the three-dimensional "cluster-cluster association" [18]. Therefore, at the formation of the silver NMS layers which is described within of the "cluster-cluster association" model, the different mechanisms of silver particles fusion and structures can be implemented.

3.2. Silver Nano-/Microstructures Deposited from Fog
 The observed ring shape of the silver NMS, synthesized by deposition from the fog, repeats the profile of evaporated micro-droplet (Fig. 3). At the same time it was found that the silver NMS's with sizes from 0.5 to 20 nm are uniformly distributed on the substrate surface that is consistent with the ideas of the "coffee stains" model [8],[9],[10].

Considering the arrays of the silver NMS, which are formed by the micro-droplets of colloidal silver solution in isopropyl alcohol on the silicon plate surfaces after deposition from fog (Fig. 3(a) and Fig. 3(b)).

Fig. 3. SEM-images of the silver nano-/microstructures on the silicon surface formed by evaporation of the micro-drops of colloidal solution. Measurement conditions are listed on the insert.

Despite the significant time interval between the deposition of the micro-droplets on the substrate and SEM-measurements, the border of drops can be clearly observed on the silicon surface. This may be due to the sorption of the alcohol molecules by the silicon substrate. The numerous silver NMSs with the size from <0.1 to $3 \mu\text{m}$ can be observed on the SEM images also. It should be noted that a pronounced ring structure is characteristic for silver NMSs with the size from <0.1 to $1 \mu\text{m}$.

According to [9] [10] [19], capillarity leads to one micron particle dynamics in evaporated drop. In this case, mass transfer processes can move the silver NMP either to the drop surface, or in the opposite direction depending on the ratio of the speed of the drop shape relaxation and the evaporation rate. The formation of the ring structure (or self-organization) may be considered a result of the mass transfer processes.

Unfortunately, due to the high level of the electron field emission the origin of which is associated with the adsorbed solvent molecules, it was impossible to carry out the SEM-studies of the ring silver NMS at the higher magnifications. Further investigation of the structure and distribution of silver NMPs in the ring boundary layer was carried out by AFM (Figure 4). As follows from the AFM-images (Figure 4(a) and Figure 4(b)), the size of the observed numerous silver NMP is about from 0.01 to $0.5 \mu\text{m}$. All the silver particles considered have the same Crescent shape as well as the same orientation. All the observed silver NMPs are oriented by thickened side towards the outer border of the ring, while the Crescent is oriented towards the center of the vaporized micro-droplet.

It is possible to result the following arguments in favour of the fact that with the help of AFM method exactly the silver particles have been registered. First, in the AFM profiles the silver NMP revealed itself as the fluctuations of the spatial distribution of the surface height with the magnitude from 30 to 35 nm (Fig. 4(a) and Fig. 4(c)), that is much higher than the surface roughness of the silicon substrate ($2.5 - 5.0 \text{ nm}$). Second, the maximum height of the discovered sickle silver NMP is 44 nm that coincides with the maximum of the size distribution ($40 - 45 \text{ nm}$) in the aerosol containing the silver NMP which has been determined previously (Fig. 1(b)).

The most part of the sickle silver NMP's, shown in the Fig. 4, has a two-or three-layer structure (Fig. 5(a)). At least two aerosol particles participated in the formation of the considered silver NMP, which originally had a spherical shape (Fig. 2(d)). Possibly, the deformation of the considered two-or three- particle clusters tends towards its boundaries away from the center. Probably the mass-transfer processes have been moving in that direction during the evaporation of the IPA molecules.

Fig. 4. 2D- (a) and 3D-images (b) of the silver NMP arrays on the surface of the silicon support recorded by AFM-method in the field of the ring structure appeared in the place of the vaporized micro-droplets; (c) the AFM-profile of the silicon surface containing the silver nano- and micro-particles (shown by the white line on the image (a)). An enlarged image of the silver particle, selected by white rectangular frame, is shown in the Fig. 5(a).

An assessment of geometric characteristics of the Crescent-shaped silver NMP has been done. For this purpose, the shape of the particles has been represented in the form of a Crescent (Fig. 5(b)) and the geometric parameters of the silver NMP have been calculated using the following formulas [20] [21]:

$$\begin{cases} h_i = h_2 + \Delta h; s_i = \alpha_i r_i; r_i = h_i + d_i = \frac{h_i}{2} + \frac{c^2}{8h_i} \\ c = \frac{2r_i \sin \alpha_i}{2} = 2\sqrt{h_i(2r_i - h_i)}; \alpha_i = \frac{2\arccos d_i}{r_i} \\ A_i = \frac{1}{2} r_i^2 (\alpha_i - \sin \alpha_i) \end{cases} \quad (1)$$

In the system of the Equation (1) $i = 1, 2$; $h_i = h_1, h_2$ is the height of the circle segments, the arc lengths s_1, s_2 are part of these segments, correspondingly, limiting the considered particle with external and internal sides; Δh is the thickness of the Crescent; d_1, d_2 are the distances from the centres of circles O_1 and O_2 to the middle of the chord AB , which have the lengths c ; r_1, r_2 are the radii of the circles with centres O_1 and O_2 ; α_1, α_2 are the angles AO_1B and AO_2B ; A_i is the area of segments. The area of the Crescent is determined as the difference of the squares of segments.

According to the calculations, the radii of the circles O_1 and O_2 are practically identical (Table 1). Consequently, the Crescent-shaped silver particle is formed by two deformed aerosol NMP (Fig. 5(a)). On the other hand, the estimated values of the circle radii O_1 and O_2 are $4 - 5$ times higher than the most probable

silver NMP size in the initial aerosol (Fig. 1(b)). Probably, the obtained difference is due to the presence of a layer of the adsorbed isopropyl alcohol molecules, i.e. the cluster “silver NMP–isopropyl alcohol molecules” can be seen as a direct micelle. Possibly, the evaporation of micro-droplets is accompanied by the coagulation of silver particles existing in the solution in the form of direct micelles. Further, clusters containing two or three NMP’s, which surrounded by a layer of the IPA molecules, have been created.

Table 1. The geometrical characteristics of the Crescent modelling the silver sickle nano-/micro-particles.

№	Name, designation	Value	
		Arc1	Arc2
1*	The thickness of the Crescent in the center of the figure Δh , μm	0,18	
2*	The length of the chord c , μm	0,36	
3*	The height of the segment in the center h , μm	0,29	0,11
4	The radius of the circle r , μm	0,201	0,205
5	The length of the arc s , μm	0,82	0,44
6	Angle α , degree	232,6 9	125,72
7	The area of the segment A_i , μm^2	0,10	0,03
8	The area of the Crescent (particles), μm^2	0,07	

Note: (*) The values are determined by measuring the Fig. 5(a).

The processes of spinodal decomposition [22] of the considered clusters and the formation in the cluster volume of the two-phase system containing the layer of the deformed Crescent shape silver NMPs have been developed during further drying and storage of the sample. Subsequent evaporation of the IPA molecules leads to the formation of the observed silver Crescent-shaped NMP. It is known [23], that this phenomenon is often observed in the study of the metastable heterogeneous colloidal systems as a result of the complex thermo- and mass -transfer processes [24] and it is called “the syneresis”.

According to [25] [26] [27] the driving force the effect of syneresis is concentration gradient of the alcohol molecules in the cluster and on the substrate surface during the evaporation process. The value of the occurring stress concentration is sufficient for cluster splitting, and deformation of the original spherical silver NMP’s.

The value of the fractal dimension for silver Crescent-shaped NMP (Fig. 5(a)) is about of $D \sim 1.69$, that is significantly lower than the established similar values for the silver NMP fractal dimension obtained by passing the silver aerosol through the IPA medium [18]. This value of the fractal dimension D corresponds to the model of the “cluster–cluster association in the regime of Brownian motion model” [18].

Thus, experimentally synthesized silver NMS with sickle NMP’s was obtained by means of controlled evaporation of the colloidal solutions droplets deposited from the fog on the surface of the silicone support. It is possible that the arrays of the silver Crescent-shaped

NMP’s may be associated with the processes of silver nanoparticles self-organization, which develop during the micro-drops evaporation. In conclusion, it should be noted that the detected silver Crescent shaped NMP belongs to the recently discovered kind of specific asymmetric nanoparticles—the so called Janus silver nanoparticles [25],[26] [27].

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References

1. V.V. Klimov, *Nanoplasmonics*, 480 (2009)
2. A. Aubry, Y.L. Dang, S.A. Maier, J.B. Pendry, *A Physical Review*, 82B (2010)
3. N. Bocchio, M. Alvarez, F. Gaul, H. Rochholz, K. Koynov, M. Kreiter, *Molecular Plasmonics. International Symposium, Abstracts. Jena* (2007)
4. V. Krapek, A.L. Koh, L. Brinek, M. Hrton, O. Tomanec, R. Kalousek, S.A. Maier, T. Sikola, *Optics Express*, 23, 11855 (2015)
5. H. Rochholz, N. Bocchio, M. Kreiter, *New Journal of Physics*, 9, 53, (2007) <https://doi.org/10.1088/1367-2630/9/3/053>
6. A.S. Smolyanskii, V.A. Zagaynov, Yu.G. Biryukov, E.P. Magomedbekov, O.G. Stepanova, L.I. Trakhtenberg, *Composites and Nanostructures*, 6, 137-147 (2014)
7. S.V. Gaponenko, N.N. Rosanov, E.L. Ivchenko, A.V. Fedorov, A.V. Baranov, A.M. Bonch-Bruevich, T.A. Vartanyan, S.G. Przybelski, *Nedra*, 326 (2005)
8. M. Layani, M. Gruchko, O. Milo, I. Balberg, D. Azulay, S. Magdassi, *ACS Nano*, 3, 3537-3542 (2009)
9. V.V. Vysotskaya, O.J. Uryupin, I.N. Senchikhin, V.I. Roldugin, *Colloidal Journal*, 75, 161-169 (2013)
10. I.V. Vodolazsky, Yu.Yu. Tarasevich, O.P. Isakova, *Nonlinear World*, 8, 142-150 (2010)
11. P. Reist, *Aerosols* (Mir, Moscow, 1997)
12. S.S. Kutateladze, V.M. Borishanskii, *Handbook of Heat Transfer* (2012)
13. K.W. Lee, H. Chen, *Aerosol Science and Technology*, 3, 327-334 (1984)
14. T. Ferreira, W. Rasband, *ImageJ User Guide-IJ 1.46r*. (2012)
15. A.L. Karperien, *FracLac for ImageJ*, 1-36, (2013)
16. A.E. Aloyan, V.O. Arutyunyan, A.A. Lushnikov, V.A. Zagaynov, *Journal of Aerosol Science*, 28, 67-85 (1997)
17. A.V. Kolesnichenko, M.Y. Marov, *Modeling of Processes of Formation of Dust Fractal Clusters as the Basis of Loose Preplanetesimals in the Solar Deplanatum the Cloud* (Preprint IPM, Moscow, 2014)

18. B.M. Smirnov, *Uspekhi Fizicheskikh Nauk*, 149, 177-219 (1986)
19. I.P. Suzdalev, *Nanostructures and Nanomaterials* (2006)
20. G. Korn, and T. Korn, *Mathematics Handbook (for Science Officers and Engineers)* (1977)
21. Online Calculator. <http://planetcalc.ru>
22. A.A. Rempel, A.A. Valeeva, *Materials and Methods of Nanotechnology*, (2015)
23. D.A. Fridrikhsberg, *Course of Colloid Chemistry*, (1995)
24. J.E. Geguzin, *Physics of Sintering* (Nauka, Moscow, 1984)
25. A.Y. Olenin, T.R. Nizamov, G.V. Lisichkin, *Russian Nanotechnology*, 9, 19-24 (2014)
26. Y. Song, S.W. Chen, *Chemistry—An Asian Journal*, 9, 418-430 (2014)
27. M. Lattuadaa, T.A. Hattonb, *Nano Today*, 6, 86-308, (2011)