

Resistivity Scaling Transition in Ultrathin Metal Film at Critical Thickness and Its Implication for the Transparent Conductor Applications

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Understanding of ultrathin metal film's electrical and optical properties at sub-10 nm thickness may provide important engineering insight on its application as a transparent conductor. Here, a rapid change is observed in the ultrathin metal film's electrical and optical scaling properties as the thickness shrinks to below a certain critical thickness d_c . Below this thickness, the metal film's electrical properties are shown to be strongly influenced by the inhomogeneity of the film which can be modeled via general effective media theory by incorporating size-effect contribution. As a result, below d_c , carrier's scattering time rapidly decreases with a reduced mean free path leading to a rapid rise in resistivity. Also, the film's optical loss increases while the optical transmission plateaus below d_c . As one promising application of thin metal film is transparent conductor where the film's electrical and optical properties are equally important, its maximum theoretical figure-of-merit is shown, which is determined at this d_c serving as an important engineering metric.

1. Introduction

Understanding the electrical and optical properties of thin metal films serves as the foundation for its application in electronic and photonic devices.^[1] Recently, metal film with thickness in the ultrathin (<10 nm) scale has caught great interest because metals such as silver (Ag) or gold is known to have excellent optoelectronic or plasmonic property due to their low resistivity and optical loss.^[2,3] To obtain continuous and ultrathin films of these metals, efforts were made to lower the percolation threshold thickness of these materials by overcoming their intrinsic de-wetting problem.^[4,5] These efforts enabled numerous applications such as transparent conductor,^[2,6,7] low-loss plasmonic waveguide,^[8] EMI shielding,^[9] or other nanophotonic applications.^[10,11] In particular, ultrathin metal film as a transparent conductor^[3,12] has gained great interest over the past years exhibiting advantages including excellent electrical/optical

properties with mechanical flexibility and simple fabrication procedure compared to counterparts like metal-oxide films,^[13] carbon-based materials,^[14] or metal nano-wires.^[15] Additionally, our recent results show that ultrathin metal film as a transparent conductor can completely eliminate waveguide mode in organic light-emitting devices to achieve better device efficiency, an outstanding benefit over metal-oxide films.^[16] As the apparent electrical resistivity becomes strongly dependent on the film's thickness in this regime, understanding the resistivity change at the ultrathin regime may provide insight and guidance in engineering applications. Electrical properties of metal films become a sophisticated problem especially when the thickness is close to extremely thin (≈ 5 nm) regime where not only the size effect

affecting the resistivity but also the morphological changes of the films starts to strongly influence its electrical property.^[17,18] Despite its significance, research on ultrathin metal films so far mostly involve experimental observation of resistivity scaling^[6,19] with little effort on rigorously investigating how understanding its electrical and optical properties can better serve to guide or solve engineering problems.

Several studies were conducted in the past to understand the morphological evolution of metal film growth and how it impacts the electrical and optical properties of the film at the ultrathin regime.^[17,18,20–22] For example, Zhang et al.^[17] discussed the resistivity change of ultrathin Ag films near critical thickness where the transition of growth mode happens, though the basis of defining critical thickness is not clear. Maaroo et al.^[18] introduced the term “critical thickness” near the percolation threshold for Pt and Ni thin films by modeling the electrical resistance associated with morphological change of the film. Hovel^[22] observed the optical property change of ultrathin gold film near the percolation threshold where metal-to-insulator transition occurs. These studies use the term “critical thickness” or “percolation threshold” to better identify the morphological transition of film growth, but they did not provide further insight or practical implications in addressing engineering applications. As a step closer to bridging the gap, Ghosh et al. experimentally demonstrated how the term “percolation threshold thickness” of metal film can be associated with transparent conductor's optimum figure-of-merit.^[20,21] However, the work is based on experimental observation only and

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is lacking in theory. Although these works help us to understand the film's property at ultrathin thickness regime, it is difficult to find a direct correlation on how resistivity modeling or terminologies can help us better design electronic or photonic devices.

We provide a detailed study on the implication of critical thickness d_c at which the sharp transition of electrical resistivity occurs in ultrathin metal films. This transition of resistivity near d_c is explained by using general effective media theory, treating the film as an inhomogeneous medium composed of metal film with air voids, which provides an excellent fit to the measured resistivity scaling behavior. Based on this analysis, the scientific and engineering implication of d_c is discussed. First, below d_c , a drastic increase in electrical resistivity is attributed to the rapid decrease in scattering time, a regime where size effect scattering models no longer dominate. This is explained via the film's morphology change that impacts the electron mean free path. Second, the optical absorption of the film increases while optical transmission reaches a plateau. Finally, when using such an extremely thin metal film as a transparent conductor, its figure-of-merit reaches a maximum value at d_c . This interesting correlation becomes a useful design guideline in designing an ultrathin Ag film transparent conductor to improve the efficiency of light emitting devices^[23] as our recent work shows that waveguiding in OLED can be eliminated.^[16]

2. Results and Discussions

2.1. Observation of Critical Thickness

Ultrathin copper-seeded thin silver film, denoted as Ag (Cu), was prepared according to our previous work.^[16] The measured resistivity of Ag (Cu) thin metal films shows an exponential increase with decreasing average film thickness d . **Figure 1a** shows the log-log plot of the measured Ag (Cu) film's resistivity

(open symbol) as a function of d . Clearly the resistivity curve shows two distinct linear regimes on a log-log scale (two dashed-lines are drawn as a guide only), in which the rapid resistivity scaling at lower thickness can be easily overlooked if plotted under linear scale. This double-sloped behavior is quite universal, as it was also observed in Ag films with germanium as a seed-layer or even without any seed layer (Figure S1, Supporting Information); and can be seen in other metal films like Au,^[5] Cu,^[24] or NiCr;^[25] though none of these works closely examined this phenomenon and its implications. It is worth paying attention to the film's thickness at which two slopes in Figure 1a intersect which is marked in vertical red dashed line and will be referred to as the critical thickness d_c , which is empirically obtained as 5 nm for Ag (Cu) film.

2.2. Electrical Resistivity Approximation

A better understanding of the rapid increase in resistivity below d_c is important in designing metal film based transparent conductors because divergence in resistivity should be avoided. First, we start by using widely accepted size effect models to describe the resistivity scaling of the film. Simply put, size effect models describe the increase in resistivity of the metal film as the film's size (thickness or grain size) decreases which is primarily attributed to the surface or grain boundary scattering of the conduction electrons.^[26] As the details of these models can be found elsewhere,^[26,27] we will not go into details but simply use them to model our film (calculations in Section 3 of supporting information) using Matthiessen's rule by treating each scattering mechanism independently:^[26]

$$\rho_{size} = \rho_{Surf} + \rho_{GB} - \rho_i \quad (1)$$

where ρ_{Surf} and ρ_{GB} are resistivity contribution due to surface and grain boundary scattering, respectively, and ρ_i is the bulk resistivity. The blue dotted line in Figure 1b is the approximated

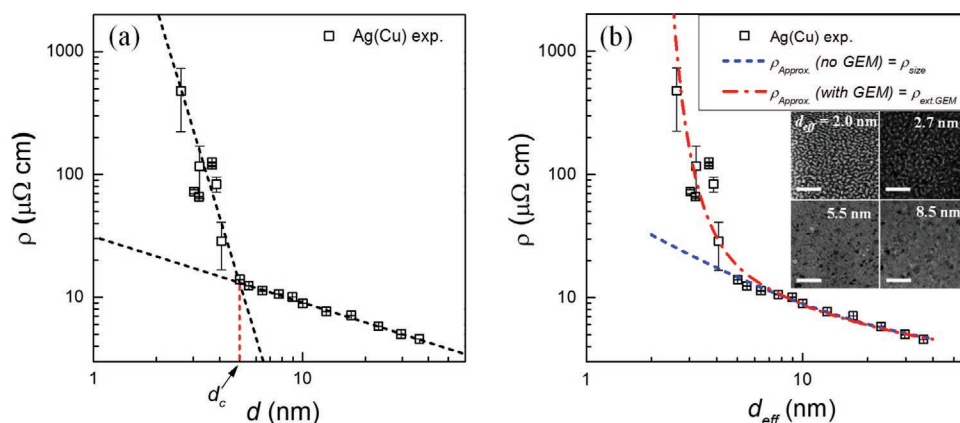


Figure 1. a) Resistivity ρ of Ag (Cu) film as a function of film thickness d plotted in log-log scale. The open symbols are the measured data while dotted lines extrapolating double-slope behavior of resistivity scaling are not based on any physical model. The critical thickness d_c is indicated as a vertical red dotted line which is 5 nm for Ag (Cu) film. b) ρ versus d approximated by using size effect model only (ρ_{size}) is plotted in blue dashed line and that by extended GEM model ($\rho_{ext.GEM}$) is plotted in a red dashed-dot line. Measured experimental resistivity values are plotted in open symbols for reference. Inset images are the top-down TEM images of Ag (Cu) films with thickness d ranging from 2.0 nm to 8.5 nm where corresponding thickness for each image is indicated. Regions appear as light or dark color each correspond to region with air (void) or metal, respectively. All images have a scale bar of 50 nm. The images are processed to show better contrast between the projected area of metal and air.

resistivity ($\rho_{approx.}$) of Ag (Cu) thin metal film calculated by size effect models called ρ_{size} which shows good agreement with experimental data (symbol) for a thickness of $d > 5$ nm but fails for thickness below $d < 5$ nm. It can be inferred that different scattering mechanisms may be limiting the electron conductivity below 5 nm. As top-down TEM images for selected film thickness show as inset of Figure 1b, film's morphology below 5 nm can be treated as an inhomogeneous medium comprised of metal film with air voids. Effective thickness d_{eff} will be used from now to define the nominal thickness of metal films. Resistivity of such inhomogeneous medium is known to scale as the fraction of air (insulator) in the medium is increased, which can be modeled using effective medium approaches.^[28,29] We extend the general effective media (GEM) theory as an empirical model to fit our data because this model entails both the aspects of effective medium approximation and percolation model, which fully describes the rapid resistivity change for a wide range of metal-air composite.^[30] This is appropriate for our case as we are dealing with rapid resistivity change in the vicinity of the percolation threshold with different resistivity scaling behaviors. The resistivity as determined by the GEM (ρ_{GEM}) model can be calculated from its conductivity σ_{GEM} ($=1/\rho_{GEM}$) as:^[29]

$$(1-\phi) \cdot \frac{\sigma_m^{1/t} - \sigma_{GEM}(\phi)^{1/t}}{\sigma_m^{1/t} + A \cdot \sigma_{GEM}(\phi)^{1/t}} + \phi \cdot \frac{\sigma_0^{1/t} - \sigma_{GEM}(\phi)^{1/t}}{\sigma_0^{1/t} + A \cdot \sigma_{GEM}(\phi)^{1/t}} = 0 \quad (2a)$$

$$A = \frac{1-\phi_c}{\phi_c} \quad (2b)$$

where ϕ is the metal fraction, σ_0 is the bulk conductivity of metal, t is the critical exponent of conductivity, ϕ_c is percolation threshold fraction, A is a constant, σ_m is the conductivity of the medium which can be approximated as ≈ 0 (for insulator). For our case, $t = 3.06 \pm 0.2$ with a percolation threshold value ϕ_c of 0.59 were empirically extracted from Ag (Cu) film's physical parameters (Figure S5a for details). In its original form Equation (2a) does not have any dependence on the film thickness, but only the metal fraction with its bulk conductivity

σ_0 . To obtain explicit film thickness dependence required in this study, we extend the GEM model by choosing σ_0 to be σ_{size} ($=1/\rho_{size}$), where σ_{size} is determined by the size effect model which is a function of the film thickness. By doing so, we can dynamically capture the change in resistivity as film size shrinks to an ultrathin regime. This is important because the conduction of the electron at such a thin regime cannot be described in a piecewise manner, but two mechanisms are interlinked to determine the total resistivity. Resistivity value approximated by this extended GEM model $\rho_{ext.GEM} = 1/\sigma_{ext.GEM}$ is plotted as red dashed line in Figure 1b. Surprisingly, it shows that using the GEM model with the simple substitution of ρ_0 by ρ_{size} provides an excellent fit for the experimental data throughout the entire thickness range, including the $d_{eff} < 5$ nm regime. Also note that for the range of film thickness sufficiently large where the film is free of voids ($\phi = 1$), $\rho_{ext.GEM}$ naturally converges to ρ_{size} . The discrepancy arises if we assume a constant resistivity ρ_0 ($=1/\sigma_0$) throughout the entire range of $\phi(d_{eff})$ which may not be correct because the scattering at grain boundary or surfaces still plays a role in this regime (Section 6 of supporting information for details). Resistivity contribution by other mechanisms such as tunneling at discontinuous films^[27,31] have been ruled out because these models yield resistivity at least a few orders of magnitude higher.

2.3. Implication of Critical Thickness

Despite divergence of resistivity at metal-insulator transition regime is a widely studied topic in the thin film community,^[22,29,32] less efforts were made to study the resistivity change near critical thickness. Here we want to pay attention to the change in electrical and optical properties near critical thickness d_c to gain some engineering insight, which will be helpful in utilizing the ultra-thin metal film as a transparent conductor and for other optoelectronic applications.

First, the carrier conduction mechanism undergoes a transition with respect to d_c . This is shown in Figure 2a where left axis is the modeled resistivity and the right axis is the contribution

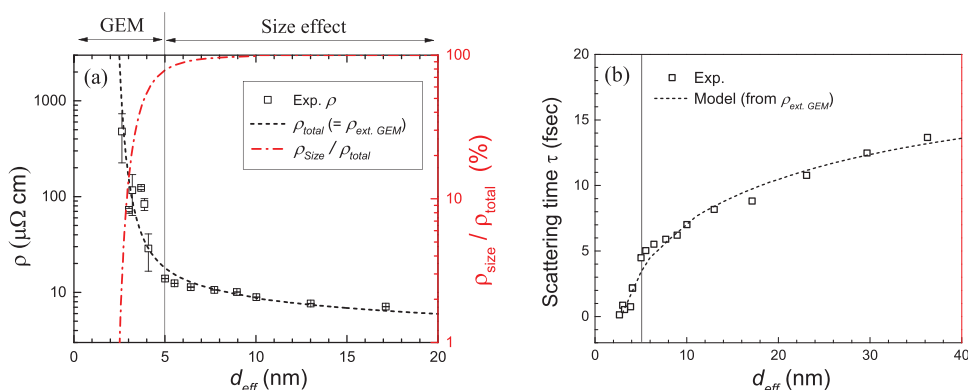


Figure 2. a) Ag (Cu) film's total resistivity ρ_{total} ($=\rho_{ext.GEM}$, dashed black line) on the left axis and contribution of size effect model over total resistivity, ρ_{size}/ρ_{total} (dashed-dot red line) on the right axis both as a function of film thickness d_{eff} . Symbols on the left axis are measured experimental resistivity values. Transport mechanism above and below the critical thickness d_c ($=5$ nm) are each governed by size effect theory and GEM, respectively. b) Experimental (symbol) and modeled (dashed line) carrier scattering time τ of Ag (Cu) film as a function of d_{eff} . In both figures, Ag (Cu) film's d_c is indicated as a vertical solid line.

of size effect over total resistivity shown as ρ_{size}/ρ_{total} (where $\rho_{total} = \rho_{ext.GEM}$) as a function of d_{eff} . In the graph, d_c is represented as a solid vertical line. Note that, log scale plot of ρ_{size}/ρ_{total} (%) shows a rapid decrease below d_c , conversely indicating that the electron conduction is strongly influenced by the inhomogeneity of the film. In fact, the measured electron scattering time τ ($=\mu \cdot m_e/q$, μ is mobility, q is charge, m_e is effective electron mass in silver where $1.03 \times m_0$ was used^[33]) rapidly decays below d_c which is plotted in Figure 2b with symbol.

This is attributed to the increase in metal-air interfaces due to the inhomogeneity of film, in which electron transport is impeded at these interfaces causing mean scattering time to decrease and resistivity to increase. For the film thickness above d_c , τ gradually decays as thickness shrinks which is expected per size effect theory with the value of τ being consistent with those reported elsewhere.^[34] If we assume constant carrier concentration N_c and electron mass (N_c of $5.85 \times 10^{22} \text{ cm}^{-3}$ ^[33]) within the metal phase, the overall trend of τ can be approximated by using extended GEM model $\tau^{-1} = \rho_{ext.GEM} \cdot N_c \cdot q^2/m_e$ and is plotted as black dotted line in Figure 2b. Interestingly, the Drude model (in size effect model) remains valid even for an inhomogeneous film when taking into account the reduced scattering time.

To better understand how film's morphological change affect resistivity near d_c , effective mean free path of electron l ($=\tau \cdot v_F$) was calculated using τ from $\rho_{ext.GEM}$ and Fermi velocity v_F of $14.5 \times 10^5 \text{ m sec}^{-1}$ ^[35] and plotted in Figure 3 with symbol. If we simplify our problem by assuming that grain boundaries dominate the electron scattering in the size effect model (see Figure S3), the effective mean free path will be determined by the average grain size D because electron will scatter at each grain boundary as it travels. Also, as D is known to proportionally scale with film's thickness for a physical vapor deposited metal films,^[36] we will also assume this proportionality relation ($d_{eff} \sim D$) to be valid for our case, which is an important premise for resistivity scaling due to the size effect (i.e., grain boundary scattering) model. If so, it would be the point at which l becomes smaller than D (or d_{eff}) where the size effect theory no longer becomes the dominant contributor of scattering events. As shown in Figure 3, l coincides with $d_{eff} \sim D$ relation (red dotted line) for above d_c , which implies that the scattering event is governed by size effect model. This is illustrated in top-right schematic of Figure 3 which shows a top-down view of film densely packed with metal-clusters with l determined by the grain size. However, l starts to deviate from $d_{eff} \sim D$ relation for below d_c . This is due to the increased metal-air (insulator) boundaries perpendicular to the direction of electric field causing diffusive reflection of electron, thus reduce l to below the size of grain boundary (top-left schematic of Figure 3). Therefore, it is easy to see that d_c is simply the thickness at which a film transitions from a quasi-continuous to a continuous state. Resistivity scaling transitions at d_c is the result of the inhomogeneity of metal film leading to more rapid increase of the resistivity when its thickness decreases below d_c . Moreover, the resistivity below d_c tends to have large sample-to-sample variation due to the randomness of quasi-continuous film. This could raise a practical concern on the uniformity of the sample as it would mean that resistivity becomes uncontrollable as the film gets thinner. Thus,

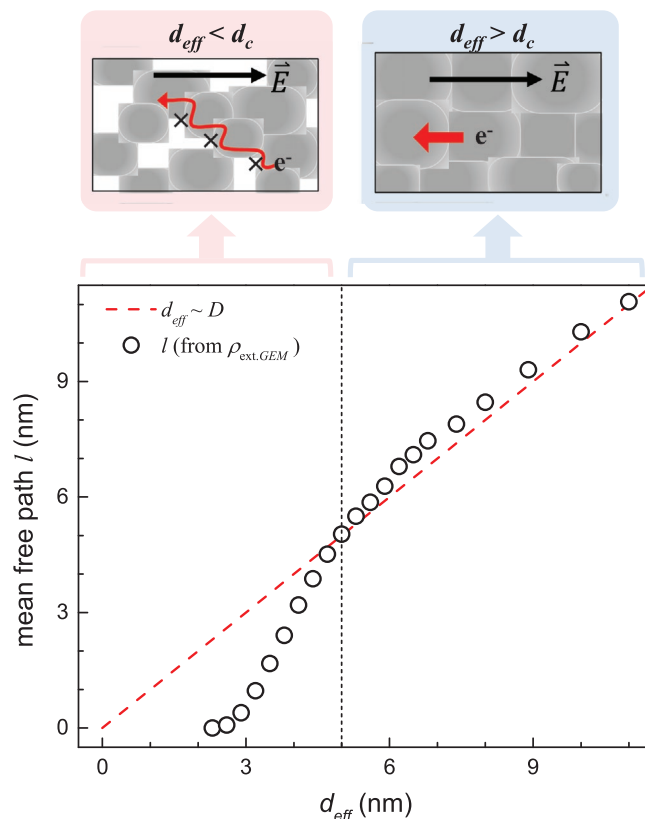


Figure 3. Film's predicted effective mean free path l (symbol) as a function of d_{eff} calculated from $\rho_{ext.GEM}$. Proportionality relation of grain size D with d_{eff} is plotted (red dashed line) as well to illustrate the size effect theory. Ag (Cu) film's critical thickness d_c ($=5 \text{ nm}$) is indicated as a vertical dotted line. Schematics above represent a top-down nanoscopic view of film's morphology in relation with the l and D for the thickness regime below (left, red) and above (right, blue) d_c . Granular shapes represent metal grains and a red arrow indicates the path of traveling electrons under the presence of electric field \vec{E} (black arrow). The left schematic illustrates the co-existence of metal clusters and air (void) depicting the inhomogeneity of the film when d_{eff} is below d_c . Each air-metal boundary (marked as cross) acts as a strong scattering site. The right schematic illustrates the homogeneous metal film free of voids when d_{eff} is above d_c .

one needs to consider metal films with thickness above d_c for practical electronic applications.

Another important implication of critical thickness is that the film's optical property also changes its trend at d_c . Figure 4a shows the average absolute transmission (T_{AVE}) and absorption (A_{AVE}) of the Ag (Cu) film on the glass substrate in the visible wavelength range (380–780 nm). Note the film's optical transmission gradually increases as the film's thickness is reduced to d_c . This is anticipated because metal film can transmit electromagnetic waves when its thickness is less than the skin depth at visible frequency ($\approx 30 \text{ nm}$ for Ag^[37]). As the film's thickness is further reduced to below d_c , T_{AVE} reaches a plateau followed by an increase in the film's absorption. This may be due to the increased absorption (and scattering) of light by a metal cluster network due to the excitation of localized surface plasmon resonance. For most photonic or optoelectronic applications of metal films, it is desirable to suppress optical loss and maximize transmission, in which d_c may be used as

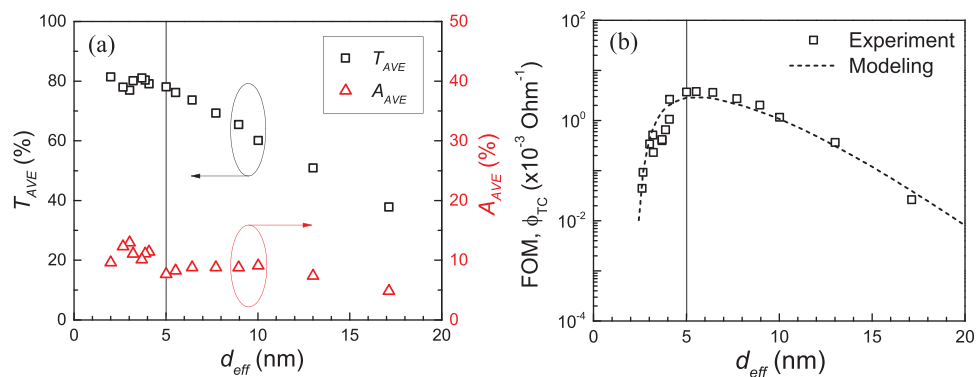


Figure 4. a) Film's measured average transmission (T_{AVE}) and absorption (A_{AVE}) over visible wavelength (380–780 nm) plotted as a function of d_{eff} . b) Haacke's figure-of-merit ϕ_{TC} ($= T^{10}/R_s$) of Ag (Cu) film as a function of d_{eff} where T is transmission at 550 nm wavelength and R_s is sheet resistance. Symbols and dotted lines are experimental and modeled ϕ_{TC} , respectively. For modeling of ϕ_{TC} , R_s was calculated from approximation (using $\rho_{ext, GEM}$) used in Figure 1b and T is calculated from measured Ag (Cu) film's optical constants using transfer-matrix method. In both figures, Ag (Cu) film's critical thickness d_c ($=5$ nm) is indicated as a vertical solid line.

a design criterion in determining the metal film's thickness to maximize the performance.

The performance of a transparent conductor is typically characterized by Haacke's figure-of-merit (FoM) ϕ_{TC} ($= T^{10}/R_s$)^[38] where T is the transmittance at a wavelength of 550 nm and R_s ($=\rho/d$) is the sheet resistance. From the above discussion, one can expect an interesting aspect of d_c for the Ag (Cu) film: its FoM will reach maximum value at this thickness. Figure 4b plots the measured (symbol) and modeled (dotted line) ϕ_{TC} of Ag (Cu) film on a substrate as a function of film thickness. To model ϕ_{TC} for Ag (Cu) film, the electrical resistivity was calculated from the resistivity approximated from the red dashed line in Figure 1a using $\rho_{ext, GEM}$ while optical transmission T was simulated from the measured refractive index of the film by using transfer matrix method. As the results in Figure 4b show, the measured ϕ_{TC} of the film shows a bell-shaped curve as a function of film's thickness in which the curve is well approximated by the modeled curve. Indeed, the film's ϕ_{TC} reaches its maximum value roughly at a thickness of d_c ($=5$ nm). This is anticipated because below this critical thickness, the resistivity rapidly increases while the film's optical transmittance reaches a plateau. For the film thickness greater than d_c , though resistivity decreases but the film becomes too thick to effectively transmit visible light, leading to a decrease in ϕ_{TC} . Therefore, it is desirable for metal film to be as thin as possible to maximize optical transmission but right before it enters a quasi-continuous film at which both electrical and optical properties start to deteriorate.

A similar observation of associating percolation threshold with optimum ϕ_{TC} thickness was made elsewhere,^[20] but we think the use of the term "percolation threshold" may not be accurate in this case. Percolation threshold is an important terminology to describe the change in morphology of the film growth, and it was calculated to be 2.4 nm in our case (calculation in Section 4 of supporting information). Clearly, such a percolation threshold is different from the thickness at which optimal FoM for a transparent conductor is obtained. Strictly speaking, the percolation threshold is the point at which metal-to-insulator transition occurs which should be lower than d_c where transition from continuous to quasi-continuous state

occurs. Ideally, it is important to make this d_c as low as possible to achieve maximum performance of metal film as a transparent conductor. However, making d_c too small will increase the sheet resistance of the film, which can deteriorate transparent conductor performance. For example, even when we assume an ideal case of Ag (Cu) film with d_c below 2 nm, our calculation shows that ϕ_{TC} still peaks at a film thickness of around 4–5 nm which decreases below this thickness due to high electrical resistance. At this film thickness regime, electrical resistance plays a dominant role in determining the transparent conductor performance and so it is critical to engineer the film to have low electrical resistance even at extremely thickness.

3. Conclusion

In summary, ultrathin Ag (Cu) film's electrical and optical properties were studied with respect to the critical thickness d_c . For film's thickness below d_c , its electrical resistivity exponentially increases which is attributed to the rise in morphological inhomogeneity of the film. Good approximation of electrical resistivity near d_c using a slightly modified GEM model by treating the film as metal-air composite medium indicates that the film's resistivity is strongly influenced by its morphological change. First, below d_c , the conduction of electron was shown to be strongly influenced by the inhomogeneous nature of metal film which increases the scattering time. Second, the film's optical transmittance plateaus while absorption increase below d_c . Finally, as one promising application of ultrathin metal films, Ag (Cu) film's FoM as a transparent conductor reaches maximum value at d_c , which serves as an important engineering and design metric.

4. Experimental Section

In this study, Ag film was deposited on a fused silica substrate where 5 Å Cu was used as a seed-layer to promote the wetting of Ag film, which will be referred to as Ag (Cu) film throughout this paper. This seeded growth of Ag (Cu) film enables smooth and continuous

Ag film down to an extremely thin regime. The optimum thickness of Cu was chosen that guarantees Ag film with low electrical and optical loss with minimum surface roughness. Silica with a size of 2 cm by 2 cm was used as a substrate for depositing Ag (Cu) film. The Ag (Cu) films were deposited using physical vapor deposition under base pressure of 10^{-7} Torr at room temperature (Kurt J. Lesker Co, LAB 18 & PVD-75). The film's resistivity was measured using the 4-point-probe method (Miller Design & Equipment FPP-5000) and was cross-checked with Hall measurement (Ecopia HMS-3000) where the values were consistent within 10% of range. For Hall measurement, samples were measured with and without applying InGa Eutectic on four corners of the sample which showed a negligible difference. Also, the ohmic contact of each sample for Hall measurement was verified by checking the linearity of all four configurations of current-voltage characteristics including films near critical thickness. For the thickness range dealt in this work, all showed metallic behavior while those thickness near 2 nm was not considered in the data as these films showed non-ohmic behavior exhibiting resistivity values that are a few orders of magnitude higher than what is dealt with here. A minimum of three samples were prepared for a given thickness where the resistivity value of each sample was obtained from the average of five different measurement data. The film's thickness was measured via ellipsometer (Woollam M-2000) which was then cross-checked using X-ray reflectivity (XRR, Rigaku SmartLab), showing consistency in the result. Film's transmission and reflection spectra were obtained using a spectroscopic ellipsometer and a reflectometer (F20, Filmetrics), respectively. For transmission electron microscopy (TEM) analysis, Ag film samples were directly deposited on a Silicon Dioxide Support Films TEM grid (PELCO, 18 nm, $60 \times 60 \mu\text{m}$ apertures (24) on $0.5 \times 0.5 \text{ mm}$ Window, $\varnothing 3 \text{ mm}$). Then, top-down images were taken under TEM bright field image mode (JEOL 2010F) with 200 kV high voltage condition. In the TEM image of Ag films, the dark contrast is diffraction contrast due to strong electron diffraction from Ag grains, indicating regions covered with Ag. Region appearing as a bright spot is a sign of amorphous behavior, in which region above cutoff value was treated as a void region with no Ag. Images were obtained and processed to extract the projected area fraction ratio of metal-insulator composite films for an extremely thin film regime. For each thickness value, images were taken from at least eight different locations of the film and the area fraction value was averaged over these images.

Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

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Conflict of Interest

L.J.G. declare financial interest in the ultra-thin metal based transparent conductor technology.

Data Availability Statement

The data that support the findings of this study are available in the supplementary material of this article.

Keywords

general effective media, resistivity scaling, size effect theory, ultrathin metal film

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- [1] a) S. M. Rossnagel, T. S. Kuan, *J. Vac. Sci. Technol., B: Microelectron. Nanometer Struct. –Process., Meas., Phenom.* **2004**, *22*, 240; b) M. Guilmain, T. Labbaye, F. Dellenbach, C. Nauenheim, D. Drouin, S. Ecoffey, *Nanotechnology* **2013**, *24*, 245305; c) E. Ando, M. Miyazaki, *Thin Solid Films* **2008**, *516*, 4574.
- [2] C. Ji, D. Liu, C. Zhang, L. J. Guo, *Nat. Commun.* **2020**, *11*, 3367.
- [3] C. Zhang, C. Ji, Y.-B. Park, L. J. Guo, *Adv. Opt. Mater.* **2021**, *9*, 2001298.
- [4] a) A. Anders, E. Byon, D.-H. Kim, K. Fukuda, S. H. N. Lim, *Solid State Commun.* **2006**, *140*, 225; b) V. J. Logeeswaran, N. P. Kobayashi, M. S. Islam, W. Wu, P. Chaturvedi, N. X. Fang, S. Y. Wang, R. S. Williams, *Nano Lett.* **2009**, *9*, 178.
- [5] Y.-G. Bi, J. Feng, J.-H. Ji, Y. Chen, Y.-S. Liu, Y.-F. Li, Y.-F. Liu, X.-L. Zhang, H.-B. Sun, *Nanoscale* **2016**, *8*, 10010.
- [6] X. Yang, P. Gao, Z. Yang, J. Zhu, F. Huang, J. Ye, *Sci. Rep.* **2017**, *7*, 44576.
- [7] G. Zhao, W. Shen, E. Jeong, S.-G. Lee, S. M. Yu, T.-S. Bae, G.-H. Lee, S. Z. Han, J. Tang, E.-A. Choi, J. Yun, *ACS Appl. Mater. Interfaces* **2018**, *10*, 27510.
- [8] C. Zhang, N. Kinsey, L. Chen, C. Ji, M. Xu, M. Ferrera, X. Pan, V. M. Shalae, A. Boltasseva, L. J. Guo, *Adv. Mater.* **2017**, *29*, 1605177.
- [9] a) H. Wang, C. Ji, C. Zhang, Y. Zhang, Z. Zhang, Z. Lu, J. Tan, L. J. Guo, *ACS Appl. Mater. Interfaces* **2019**, *11*, 11782; b) N. Erdogan, F. Erden, A. T. Astarlioglu, M. Ozdemir, S. Ozbay, G. Aygun, L. Ozyuzer, *Curr. Appl. Phys.* **2020**, *20*, 489.
- [10] F. Moresco, M. Rocca, T. Hildebrandt, M. Henzler, *Phys. Rev. Lett.* **1999**, *83*, 2238.
- [11] G. Fahsold, M. Sinther, A. Priebe, S. Diez, A. Pucci, *Phys. Rev. B* **2002**, *65*, 235408.
- [12] Y.-G. Bi, Y.-F. Liu, X.-L. Zhang, D. Yin, W.-Q. Wang, J. Feng, H.-B. Sun, *Adv. Opt. Mater.* **2019**, *7*, 1800778.
- [13] J.-W. Park, G. Kim, S.-H. Lee, E.-H. Kim, G.-H. Lee, *Surf. Coat. Technol.* **2010**, *205*, 915.
- [14] D. S. Hecht, L. Hu, G. Irvin, *Adv. Mater.* **2011**, *23*, 1482.
- [15] B. Bari, J. Lee, T. Jang, P. Won, S. H. Ko, K. Alamgir, M. Arshad, L. J. Guo, *J. Mater. Chem. A* **2016**, *4*, 11365.
- [16] C. Jeong, Y.-B. Park, L. J. Guo, *Sci. Adv.* **2021**, *7*, eabg0355.
- [17] D. Zhang, H. Yabe, E. Akita, P. Wang, R.-i. Murakami, X. Song, *J. Appl. Phys.* **2011**, *109*, 104318.
- [18] A. I. Maarroof, B. L. Evans, *J. Appl. Phys.* **1994**, *76*, 1047.
- [19] a) N. Formica, D. S. Ghosh, A. Carrilero, T. L. Chen, R. E. Simpson, V. Pruneri, *ACS Appl. Mater. Interfaces* **2013**, *5*, 3048; b) S. Jeong, S. Jung, H. Kang, D. Lee, S.-B. Choi, S. Kim, B. Park, K. Yu, J. Lee, K. Lee, *Adv. Funct. Mater.* **2017**, *27*, 1606842.
- [20] D. S. Ghosh, T. L. Chen, V. Pruneri, *Appl. Phys. Lett.* **2010**, *96*, 091106.
- [21] D. S. Ghosh, T. L. Chen, N. Formica, J. Hwang, I. Bruder, V. Pruneri, *Sol. Energy Mater. Sol. Cells* **2012**, *107*, 338.
- [22] M. Hövel, B. Gompf, M. Dressel, *Phys. Rev. B* **2010**, *81*, 035402.
- [23] C. Zhang, Q. Huang, Q. Cui, C. Ji, Z. Zhang, X. Chen, T. George, S. Zhao, L. J. Guo, *ACS Appl. Mater. Interfaces* **2019**, *11*, 27216.
- [24] E. V. Barnat, D. Nagakura, P. I. Wang, T. M. Lu, *J. Appl. Phys.* **2002**, *91*, 1667.

- [25] N. Chuang, J. Lin, T. Chang, T. Tsai, K. Chang, C. Wu, *IEEE J. Electron Devices Soc.* **2016**, 4, 441.
- [26] T. Sun, B. Yao, A. P. Warren, K. Barmak, M. F. Toney, R. E. Peale, K. R. Coffey, *Phys. Rev. B* **2010**, 81, 155454.
- [27] M. A. Angadi, *J. Mater. Sci.* **1985**, 20, 761.
- [28] a) M. H. Cohen, J. Jortner, I. Webman, *Phys. Rev. B* **1978**, 17, 4555; b) P. M. Kogut, J. P. Straley, J. C. Garland, D. B. Tanner, *AIP Conf. Proc.* **1978**, 40, 382; c) R. Landauer, *J. Appl. Phys.* **1952**, 23, 779.
- [29] D. S. McLachlan, M. Blaszkiewicz, R. E. Newnham, *J. Am. Ceram. Soc.* **1990**, 73, 2187.
- [30] M. Taya, *Electronic Composites: Modeling, Characterization, Processing, and MEMS Applications*, Cambridge University Press, Cambridge **2005**.
- [31] a) G. Dittmer, *Thin Solid Films* **1972**, 9, 317; b) P. Sheng, B. Abeles, *Phys. Rev. Lett.* **1972**, 28, 34.
- [32] A. L. Efros, B. I. Shklovskii, *Phys. Status Solidi B* **1976**, 76, 475.
- [33] C. Kittel, *Introduction to Solid State Physics*, Wiley, Hoboken, NJ **2004**.
- [34] G. Ding, C. Clavero, D. Schweigert, M. Le, *AIP Adv.* **2015**, 5, 117234.
- [35] D. Gall, *J. Appl. Phys.* **2016**, 119, 085101.
- [36] a) A. F. Mayadas, R. Feder, R. Rosenberg, *J. Vac. Sci. Technol.* **1969**, 6, 690; b) K. N. Tu, A. M. Gusak, I. Sobchenko, *Phys. Rev. B* **2003**, 67, 245408; c) M. Philipp, *Fakultät Mathematik und Naturwissenschaften*, Technische Universität Dresden, Dresden **2011**.
- [37] Y. Li, *Plasmonic Optics: Theory and Applications*, SPIE Press, Bellingham, Washington **2017**.
- [38] G. Haacke, *J. Appl. Phys.* **1976**, 47, 4086.