



Plasmonically enhanced lasing by different size silver nanoparticles-silver film hybrid structure



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ABSTRACT

We reported on the enhanced lasing in organic dyes based on plasmonic hybrid structure of Ag nanoparticles (NPs)-Ag film, the diameters of Ag NPs ranged from 20 nm to 100 nm. The lowest lasing threshold was achieved by the optimal size Ag NPs-Ag film hybrid structure, which was reduced by 5.2 times than that of the neat gain medium. Comparing to the separate Ag NPs or Ag film, the hybrid structure presented the more intense local electric field due to the plasmonics coupling between the localized surface plasmons of Ag NPs and the surface plasmon polariton of Ag film, and the stronger scattering due to the reinjection of the leaking photons by external feedback of Ag film. The effects of different sizes Ag NPs-Ag film hybrid structures on lasing were investigated. It found that when the Ag NPs in hybrid structure is small (diameter ≤ 40 nm), the enhanced localized electric field plays a major role on enhanced lasing; with the increase of Ag NPs size, the enhanced electric field and scattering have comparable contribution on enhancing lasing; for the larger size Ag NPs-Ag film (diameter ≥ 80 nm), the scattering effect is the dominant mechanism for random lasing. Then the lowest threshold was dominated by the balance of enhanced localized electric field and scattering effect. Our results could provide us a unique idea to effectively enhance the lasing of organic dyes, and realize the lower pumped threshold and stronger lasing.

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1. Introduction

Localized Surface Plasmons (LSPs) can be excited on metal nanoparticles (NPs) whereas delocalized surface plasmon polariton (SPP) can be excited on metallic film, which both can induce the field enhancement in the near-field region [1–4]. Various groups have already used metallic NPs to enhance lasing. For example, O. Popov et al. utilized gold NPs to enhance lasing from a polymer film doped with Rhodamine 6G [5,6]. X. Meng et al. achieved the enhanced emission of coherent random lasing in polymer films by introducing Ag NPs [7,8]. T. Zhai et al. demonstrated an enhanced random laser based on gold nano-island structures with a layer of dye-doped polymer [9]. E. Heydari et al. reported the emission enhancement for the gold NP-based waveguided random laser [10]. However, for the metallic film in planar waveguide structure, it

would increase the lasing threshold due to the quenching effect of metallic film on lasing of dyes, which is a major obstacle to obtain stimulated emission under electrical pumping [11,12]. Substantial efforts have been made to restore and optimize the optically pumped lasers in the presence of metallic film, such as polymer film deposited on distributed feedback metallic structure [13,14], adopting in low loss metal cladding [15–17], or introducing the spacer between polymer film and metallic film [18]. Despite all these attempts, it is still desired to further reduce the lasing threshold of optically pumped laser in the presence of metallic film.

Here, a particularly interesting plasmonic system which has received somewhat less attention, metallic NPs-metallic film hybrid structure was explored. We systematically investigated the effects of different sizes Ag NPs-Ag film hybrid structures on the lasing of gain medium, the diameters of Ag NPs ranged from 20 to 100 nm. The lowest lasing threshold of the gain medium deposited on 60 nm Ag NPs-Ag film hybrid structure was found, which greatly reduced 5.2 times than that of the gain medium deposited on glass. For comparing, the devices that gain medium deposited on Ag film,

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silver island film (SIF) were also prepared. It found that the Ag NPs-Ag film coupled system could better enhance lasing, which is because that the Ag NPs-Ag film hybrid structure presents the more intense local electric field due to the plasmonics interaction between LSPs of the Ag NPs and the SPP of Ag film, and the stronger scattering effect due to the re-injection of emitted light into the organic gain media of planar waveguide structure by the external feedback of Ag film.

At meanwhile, the effects of different sizes Ag NPs-Ag film hybrid structures on lasing properties were studied. It was noting that the enhanced localized electric field and scattering effect of Ag NPs-Ag film all work on the reduction of lasing threshold. However, according to the experiment and theoretical analysis, it was found that when the Ag NPs in hybrid structure is small (diameter ≤ 40 nm), the enhanced localized electric field plays a more significant role on the enhanced lasing of organic dyes; with the increase of Ag NPs size for hybrid structure, the enhanced localized electric field and scattering play the comparable role on the lasing; when the Ag NPs size is larger (diameter ≥ 80 nm), the scattering effect of Ag NPs-Ag film mainly contributes to the enhanced lasing. Then we concluded that the optimal size Ag NPs-Ag film could provide a balance between enhanced local electric field and scattering to realize maximum lasing. At meanwhile, the negative effect of metallic film is avoided to realize the lower pumped threshold than that of the metal-free device.

2. Experiment

2.1. Synthesis of different size Ag NPs

Different sizes Ag NPs were synthesized according to the seed-mediated growth method by citrate reduction of silver nitrate (AgNO_3) with NaBH_4 as strong reducing agent [19]. Firstly, small Ag NPs were synthesized under chemically reducing AgNO_3 in aqueous solution by a rapid nucleation-growth-ripening principle, the resulting Ag NPs were used as starter seeds. Then, slowly adding proper portions of Ag salt and citrate reducer into the starter

seeds solution obtained in the first step. In this way, we prepared Ag NPs with diameters about: 20, 40, 60, 80, 100 nm.

2.2. The devices based on different sizes Ag NPs-Ag film hybrid structure

In order to systematically study the properties of different sizes Ag NPs-Ag film hybrid structures, we developed the devices that gain media deposited on different sizes Ag NPs-Ag film, the diameters of Ag NPs were about 20, 40, 60, 80, and 100 nm, respectively. The device structures were Glass/Ag film (50 nm)/ SiO_2 (10 nm)/Ag NPs/LiF (5 nm)/PS:BMT-TPD (shown in Fig. 1). For comparison, the gain medium deposited on glass was also prepared as reference. The preparation process of devices was following manner:

Firstly, a continuous Ag film of thickness 50 nm was evaporated onto a glass substrate. The 10 nm- SiO_2 layer was radio frequency sputtered over the silver surface, which could serve to protect the metal surface from chemical attack, and to provide a means to chemically attach silver colloids. SIFs were deposited according to the procedure described elsewhere [20]. The Ag film coated with SIF was prepared. Fig. 2 shows the atomic force microscopy (AFM) images of the Ag NPs with different sizes.

Then the planar waveguide structures were fabricated. N,N'-bis(3-methylphenyl)-N,N'-diphenyl-[1,1':4,1''-terphenyl]-4,4''-diamine (BMT-TPD) was used as gain medium [21]. Polystyrene (PS) and BMT-TPD were dissolved in chloroform solution (PS: BMT-TPD = 4:1, wt%). The gain medium films were obtained by spin coating the solutions onto LiF layers at speed 4000 rpm, and were annealed at 110 °C for 10 min to remove the solvent. The 5 nm-LiF layer was deposited over the SIF surface to prevent quenching caused by direct contact between Ag NPs and gain medium. The thickness of gain medium layer is approximately 250 nm.

2.3. Characterization

The absorption and photoluminescence (PL) spectra were

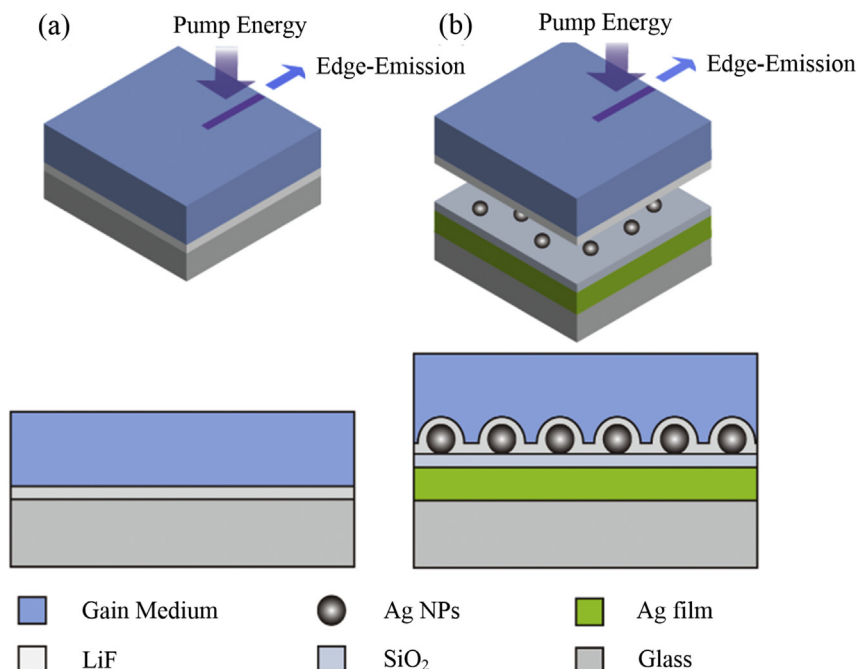


Fig. 1. The schematic illustrations of different devices. (a) Glass/LiF (5 nm)/PS:BMT-TPD (reference device); (b) Glass/Ag film (50 nm)/ SiO_2 (10 nm)/Ag NPs/LiF (5 nm)/PS:BMT-TPD.

obtained by UV–Vis spectrophotometer (HITACHI U-3010, Japan) and Fluorescence Spectrometer (Fluoromax-4 spectrofluorometer). The thicknesses of polymer layers were measured with Stylus Profiler (Dektak 6M, USA). The devices were photopumped at normal incidence with a Nd: YAG laser (355 nm, 10 Hz repetition rate, and 5.5 ns pulse duration). An adjustable slit and a cylindrical lens were used to shape the beam into a stripe with the size of $7\text{ mm} \times 1\text{ mm}$. For the devices shown in Fig. 1, they present the waveguide scheme. When the devices are pumped, the light is partially confined into the waveguide and amplified by the gain medium as it is reflected (or scattered) by the internal surfaces of waveguide and propagates along the path of optical gain. As a result, the light emitted from the edge of the waveguide could undergo the longer light path and more gain process in gain

medium, which leads the lower lasing threshold and stronger emission intensity according to the optical confinement of the waveguide. Therefore, the edge emission spectrum was measured to study the lasing properties of devices [22,23]. And The edge emission spectra were collected by Fiber Optic Spectrometer (Ocean Optics SpectraSuite, USB2000).

3. Results and discussion

Fig. 3 shows the localized surface-plasmon resonance (LSPR) bands of different sizes Ag NPs, together with the absorption and emission spectra of BMT-TPD. It exhibits that the LSPR peaks of these different sizes Ag NPs undergo a red shift from 410 to 449 nm as the diameters of Ag NPs range from 20 nm to 100 nm, which lead

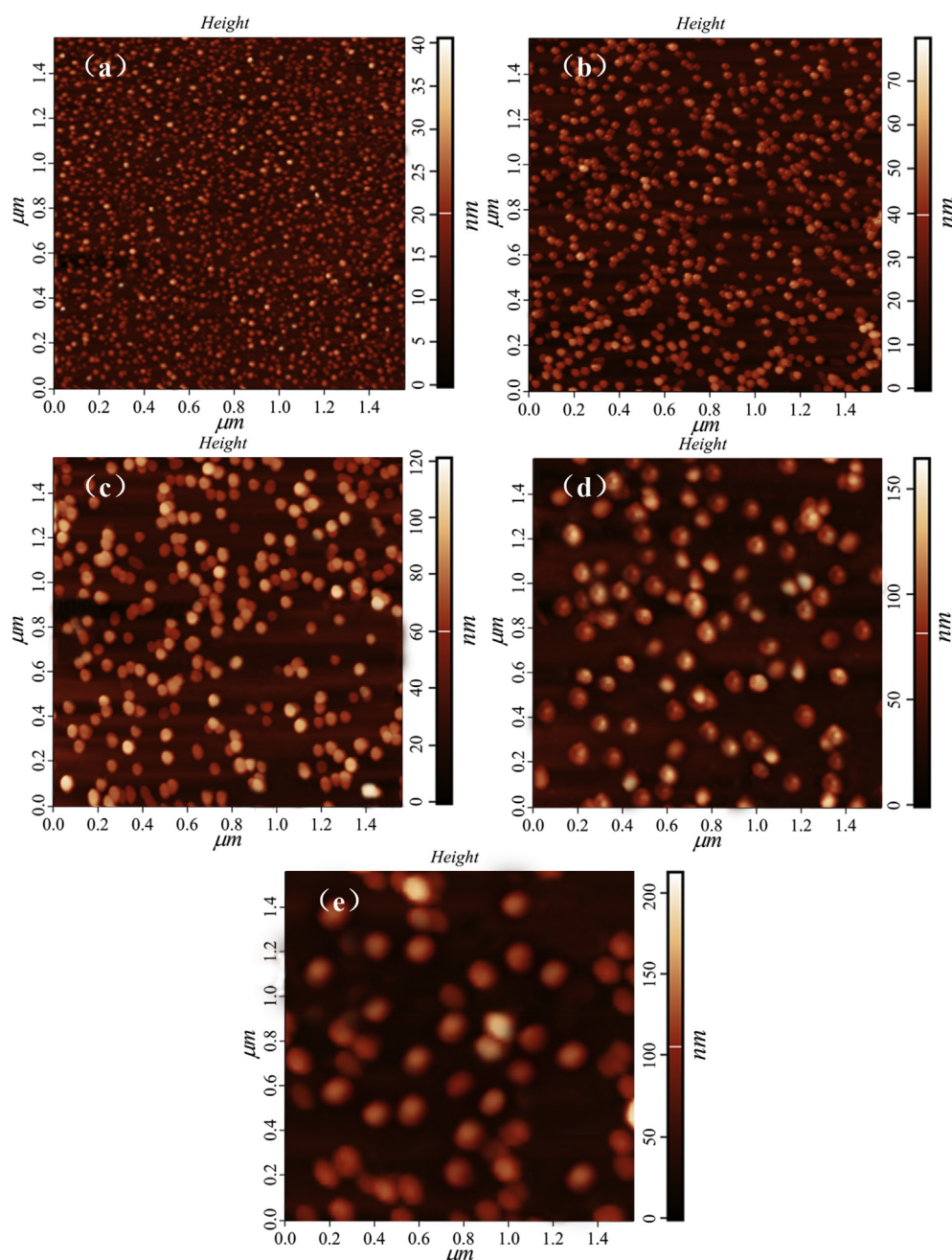


Fig. 2. The AFM images of different sizes Ag NPs, the diameters are about (a) 20 nm, (b) 40 nm, (c) 60 nm, (d) 80 nm, and (e) 100 nm.

the different effect on surface plasmon mediated emission enhancement.

For comparison, the device that gain medium deposited on glass was fabricated as reference (shown in Fig. 1(a)). Fig. 4(a) depicts the emission spectra of reference device at different pump intensities, which exhibits an obvious amplified spontaneous emission (ASE) behavior [24,25]. At first, the emission intensity increases slowly with the increase in pump intensity, the broad emission spectrum is shown. With further increase in the pump intensity, the emission spectrum becomes narrow with a steep increase of the emission intensity. The rapid decrease in linewidth of emission spectra above the threshold is shown in the inset of Fig. 4(a), which exhibits the threshold of $42.9 \mu\text{J}/\text{cm}^2$.

To investigate the effect of different sizes Ag NPs-Ag film hybrid structures on lasing properties, the different sizes Ag NPs-Ag film hybrid structures were introduced into the gain media (the device structure is shown in Fig. 1(b)) and the optimal size Ag NPs-Ag film hybrid structure was found. Fig. 4(b)-(d) show the emission spectra of devices that gain media deposited on Ag NPs-Ag film hybrid structures, the diameters of Ag NPs in hybrid structures are 20, 60 and 100 nm, respectively. The sharp spikes with full width at half maximum (FWHM) of about 1 nm are observed, which indicate the occurrence of coherent random lasing. For the waveguide-plasmonic scheme shown in Fig. 1(b), a large part of the scattered light may be totally reflected back at the gain medium/air interface to propagate within the waveguide and scattered further by the Ag NPs, then the scattered light is enhanced or amplified by the gain medium through the confinement of the scattered light into the waveguide. Therefore, the multiple scattering of Ag NPs and waveguide confinement mechanisms provide effective gain channels [9,26,27]. The random lasing could occur. The intensities and FWHMs of corresponding emission spectra are shown in the inset of Fig. 4(b)-(d). It shows that when the pump energy is higher than the lasing threshold, the emission intensity increases rapidly and the FWHM decreases to about 1 nm. Fig. 4(e) illustrates the emission intensity as a function of pump energy for devices that gain medium deposited on different sizes Ag NPs-Ag film hybrid structures, the Ag NPs diameters are 20, 40, 60, 80 and 100 nm, respectively. The corresponding lasing thresholds are exhibited in Fig. 4(f) and Table 1. It shows that with the increase of Ag NPs size, the lasing threshold exhibits a profound reduction at first, and turns to increase. The lowest lasing threshold of $8.3 \mu\text{J}/\text{cm}^2$ is found for the device with 60 nm Ag NPs-Ag film hybrid structure, about a

factor of 5.2 times lower than that of neat gain medium.

For further presenting the advantages of Ag NPs-Ag film hybrid structure on enhanced lasing, we developed the devices that gain media deposited on Ag film, SIFs, respectively, the corresponding device structures are Glass/Ag film (50 nm)/SiO₂ (10 nm)/LiF (5 nm)/PS:BMT-TPD, Glass/Ag NPs (60 nm)/LiF (5 nm)/PS:BMT-TPD, respectively. Fig. 5 (a)-(c) show the emission spectra of the devices that gain medium deposited on Ag film, 60 nm Ag NPs, and 60 nm Ag NPs-Ag film with 100 nm SiO₂ spacer, respectively. Fig. 5 (b) and (c) show the sharp spikes in emission spectra above the lasing threshold, which indicate the random lasing behaviors due to the existing of Ag NPs. Fig. 5(d) illustrates the emission intensity as a function of pump energy for the devices that gain medium deposited on glass, Ag film, 60 nm Ag NPs, 60 nm Ag NPs-Ag film with 100 nm and 10 nm SiO₂ spacers. The corresponding lasing thresholds are 42.9, 37.4, 28.3, 21.4 and $8.3 \mu\text{J}/\text{cm}^2$, respectively. We can find that the lasing thresholds of device based on Ag NPs-Ag film are lower than that based on other substrates, suggesting that the Ag NPs-Ag film hybrid structure could more effectively enhance the lasing properties than independent Ag NPs and Ag film. In addition, for the device with Ag NPs-Ag film hybrid structure, comparing to the Ag NPs-Ag film with 100 nm SiO₂ spacer, there is lower lasing threshold for that with 10 nm SiO₂, which indicates that the interaction of LSP and SPP plays the important role on enhanced lasing.

As we known, the metallic nanostructures are constantly used to enhance the lasing due to the both underlying mechanisms of enhanced localized electric field and enhanced scattering strength [6]. On the one hand, the enhanced localized electric field could increase pump light density and excitation rate, the more dye molecules could be excited to the higher energy levels. Meanwhile, the quantum yield and radiative decay rate of dye molecule could be increased. On the other hand, the metallic nanostructures could also scatter the emitter energy with the greater scattering strength, which could further enhance the lasing properties.

In order to determine which mechanism of hybrid structure takes effect on enhanced lasing properties, the devices based on different sizes Ag NPs-Ag film hybrid structures with 50 nm-LiF spacer between SIFs and gain media were prepared, the thickness of LiF spacer of device shown in Fig. 1(b) was changed to 50 nm [28]. For the device with 5 nm-thick LiF spacer, the quenching effect of dyes could be avoided, both the enhanced localized electric field and scattering of Ag NPs-Ag film contribute to enhanced lasing; and for the device with 50 nm-thick LiF spacer, the enhanced localized electric field effect of Ag NPs-Ag film is avoided, only the scattering effect of Ag NPs-Ag film hybrid structure works [28].

The lasing thresholds of devices with 5 nm or 50 nm LiF spacer layers between gain medium and SIFs are shown in Table 1. In Table 1, E_{th-5nm} ($E_{th-50nm}$) is the lasing threshold of devices with 5 nm (50 nm) LiF spacer; E_{th} is the ASE threshold of reference device; D is the diameter of Ag NPs. E_{th-5nm}/E_{th} ($E_{th-50nm}/E_{th}$) is the ratio of threshold for device with 5 nm (50 nm) LiF spacer and the reference device, which exhibits the degree of threshold reduction for the devices with 5 nm (50 nm) LiF spacer comparing to the reference device. E_{th-5nm}/E_{th} reveals the contribution of both enhanced localized electric field and scattering effect of Ag NPs-Ag film hybrid structure on threshold reduction, and $E_{th-50nm}/E_{th}$ reflects the contribution of only scattering effect of corresponding Ag NPs-Ag film on reduced threshold.

In Table 1, it shows that the devices with 5 nm LiF spacer exhibit the superior performance and lower lasing threshold comparing to corresponding devices with 50 nm LiF spacer. And we found that the effects of enhanced localized electric field and scattering of hybrid structure all contribute to the reduced lasing threshold. When the Ag NPs is small in hybrid structure (the diameter of Ag

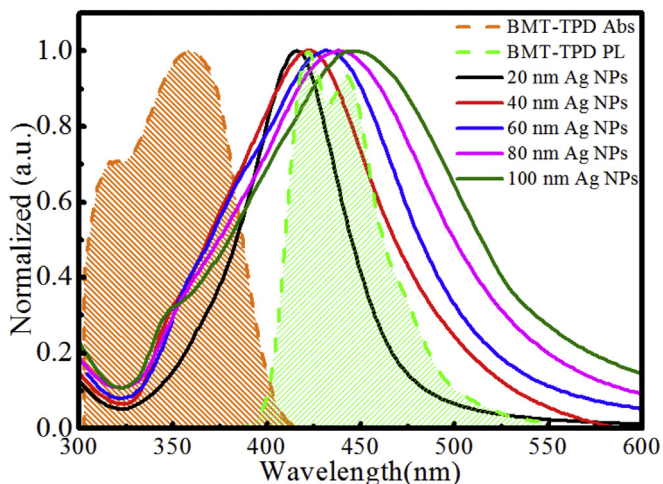


Fig. 3. The LSPR spectra of different sizes Ag NPs, together with the absorption and emission spectrum of BMT-TPD.

NPs ≤ 40 nm), the enhanced localized electric field is the major mechanism to enhance lasing. For example, for the device with 40 nm Ag NPs-Ag film, the $E_{th-5nm}/E_{th} \sim 0.25$, the corresponding $E_{th-50nm}/E_{th} \sim 0.74$, suggests that comparing to the scattering effect of 40 nm Ag NPs-Ag film, the contribution of enhanced localized electric field of 40 nm Ag NPs-Ag film on the lasing can be more, which plays the main role on reducing lasing threshold. With the increase of Ag NPs size, the scattering effect is stronger, for the device with 60 nm Ag NPs-Ag film, the $E_{th-5nm}/E_{th} \sim 0.2$, the corresponding $E_{th-50nm}/E_{th} \sim 0.62$. It shows that the enhanced localized electric field and scattering have comparable contribution to enhance lasing. In the same way, when the size is larger in hybrid structure (the diameter of Ag NPs ≥ 80 nm), such as device with 100 nm Ag NPs-Ag film, the $E_{th-5nm}/E_{th} \sim 0.33$, the corresponding $E_{th-50nm}/E_{th} \sim 0.43$, suggests that comparing to the enhanced localized electric field effect, the scattering effect of 100 nm Ag NPs-Ag film plays the main role on reducing lasing threshold.

In order to investigate the reason of experiment results above, two mechanisms of enhanced localized electric field and scattering are discussed respectively.

On the one hand, for the enhanced localized electric field of Ag NPs-Ag film hybrid structure. We know that the light can directly excite the LSPR of metallic NPs, but it cannot directly excite the SPP of metallic film due to the inherent wave vector mismatch between the SPP and the photons traveling in the air region. In the Ag NPs-

Table 1

The thresholds of devices with 5 nm or 50 nm LiF spacer layers between gain medium and SIFs. E_{th} is the ASE threshold of reference device; E_{th-5nm} , $E_{th-50nm}$ are the lasing threshold of devices with 5 nm or 50 nm LiF spacer; D is the diameter of Ag NPs. E_{th-5nm}/E_{th} ($E_{th-50nm}/E_{th}$) is the ratio of lasing threshold for device with 5 nm (50 nm) LiF spacer and the reference device.

D (nm)	Device with 5 nm LiF spacer		Device with 50 nm LiF spacer	
	E_{th-5nm} ($\mu\text{J}/\text{cm}^2$)	E_{th-5nm}/E_{th}	$E_{th-50nm}$ ($\mu\text{J}/\text{cm}^2$)	$E_{th-50nm}/E_{th}$
20	24.6 ± 0.98	0.57	36.5 ± 0.96	0.85
40	10.7 ± 0.91	0.25	31.8 ± 0.99	0.74
60	8.3 ± 1.18	0.2	26.6 ± 1.1	0.62
80	13.1 ± 1.36	0.31	22.9 ± 0.84	0.53
100	14.1 ± 1.1	0.33	18.6 ± 1.04	0.43

Ag film hybrid structure, the Ag NPs could provide an indirect way to excite the SPPs of the Ag film by the diffraction effect, thus the SPP can be excited [3,4].

In order to investigate the localized electric field of different sizes Ag NPs-Ag film hybrid structures, the electric field intensity distributions of different sizes Ag NPs-Ag film hybrid structures were calculated using the COMSOL software. Fig. 6 (a)-(e) show the electric-field distribution of different sizes Ag NPs-Ag film hybrid structures with 10 nm SiO_2 spacer at the incident wavelength of 435 nm which is wavelength of emission light, the diameters of Ag NPs are 20, 40, 60, 80, and 100 nm, respectively. We can find that

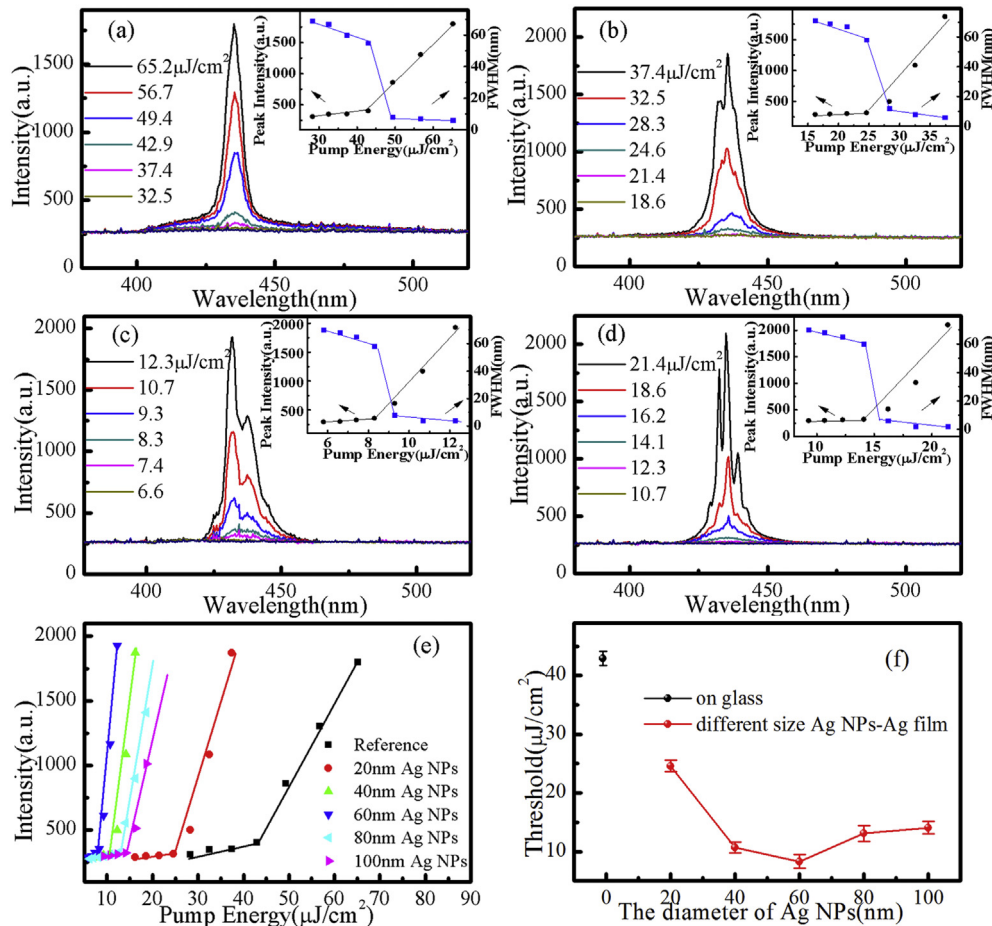


Fig. 4. The emission spectra of the devices that gain media deposited on (a) glass, (b) 20 nm, (c) 60 nm, (d) 100 nm Ag NPs-Ag film hybrid structures. The insets show the dependence of the emission intensity and the FWHM of the emission spectra on the pump intensity. For the sharp spikes of spectra appear in (b)–(d), the FWHMs above the lasing threshold were estimated from those of sharp spikes, instead of that of the broad background. (e) Dependences of the emission intensity on the pump energy intensity for devices with different structures. (f) The lasing thresholds of the gain media deposited on glass and different sizes Ag NPs-Ag film hybrid structures.

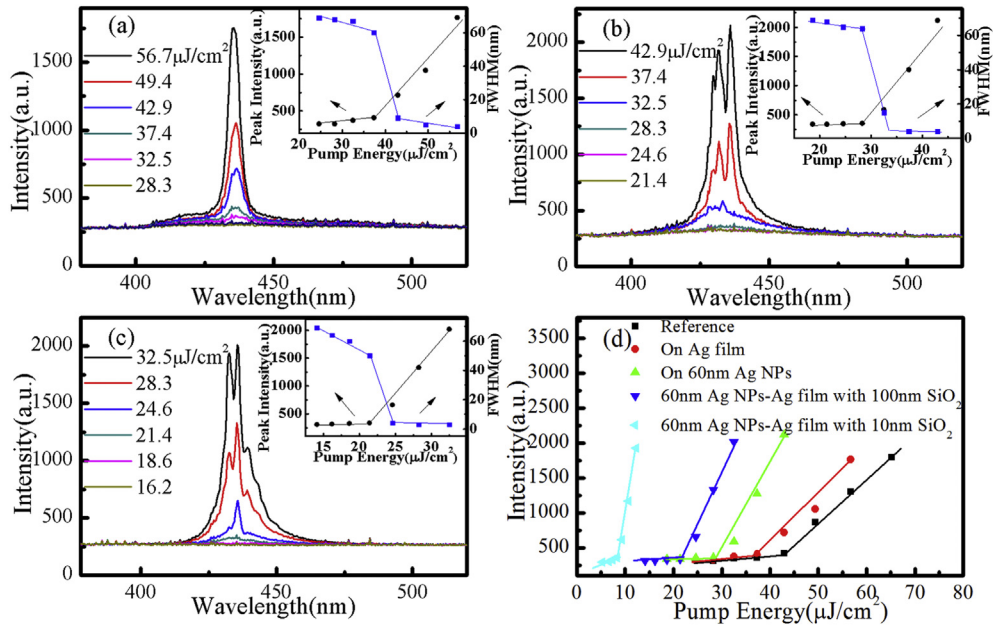


Fig. 5. The emission spectra of devices that gain media deposited on (a) Ag film, (b) 60 nm Ag NPs, (c) 60 nm Ag NPs-Ag film with 100 nm SiO_2 . The insets show the emission intensity and the FWHM of the emission spectra. (d) Dependences of the emission intensity on the pump energy for different devices.

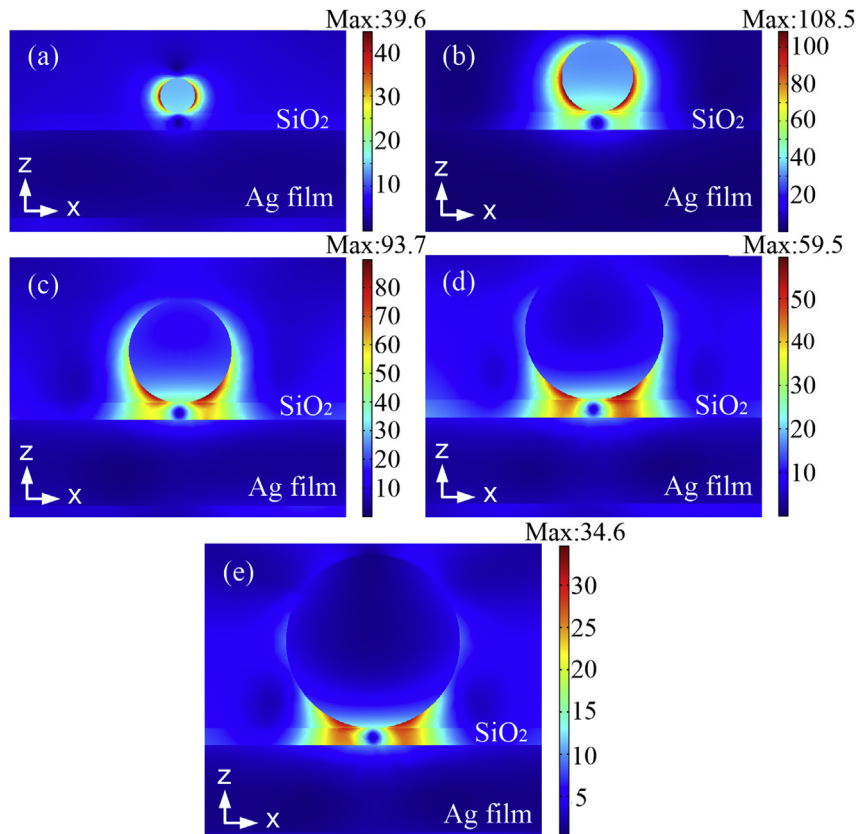


Fig. 6. The electric-field distribution of (a) 20 nm, (b) 40 nm, (c) 60 nm, (d) 80 nm, and (e) 100 nm Ag NPs-Ag film with 10 nm SiO_2 spacer.

with the increasing of Ag NPs size, the electric field intensity of hybrid structure increases at first, and then reduces, there is the strongest electric field when the diameter of Ag NPs is 40 nm. In addition, for illustrating the advantage of Ag NPs-Ag film hybrid

structure on enhanced localized electric field and proving the existing of the plasmon coupling interaction between Ag NPs and Ag film, the electric field intensities of independent 60 nm Ag NPs (Fig. 7(a)) and the 60 nm Ag NPs-Ag film with 100 nm SiO_2 spacer

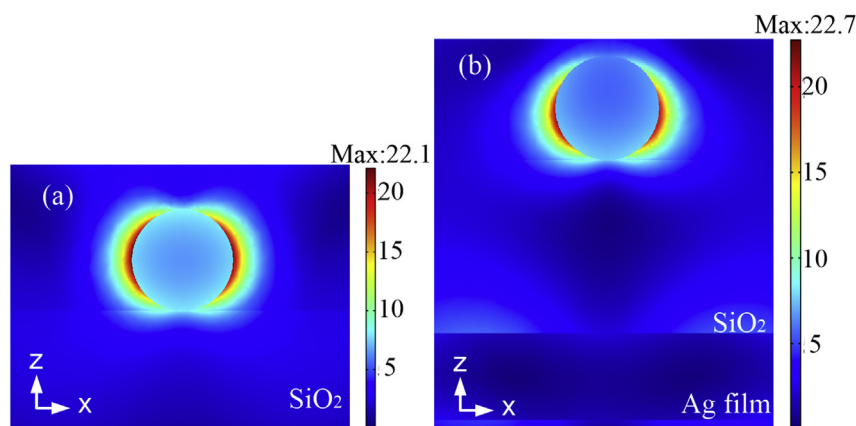


Fig. 7. The electric-field distribution of (a) 60 nm Ag NPs on SiO₂ without Ag film, (b) 60 nm Ag NPs-Ag film with 100 nm SiO₂ spacer.

(Fig. 7(b)) were also simulated. Comparing with those two structures shown in Fig. 7, we found the 60 nm Ag NPs-Ag film with 10 nm SiO₂ spacer has stronger electric field. Fig. 7 also shows that when the distance between 60 nm Ag NPs and Ag film changes to 100 nm, the electric field intensity is similar as that of independent 60 nm Ag NPs, which indicates that there is no plasmon coupling when the distance between Ag NPs and Ag film is too large. Therefore, the lasing threshold of device based on Ag NPs-Ag film hybrid structure with 100 nm SiO₂ spacer is higher than that with 10 nm SiO₂ spacer shown in Figs. 4 and 5. And the stronger localized electric field of hybrid structure due to the plasmon coupling interaction between LSPR of Ag NPs and SPP of Ag films is identified in Figs. 6 and 7.

On the other hand, to evaluate another mechanism of scattering effect of different size Ag NPs-Ag film, the scattering cross sections (σ_s) of different sizes Ag NPs at $\lambda = 435$ nm were calculated using Mie theory [29]. For comparison, $\sigma_s/V_{particle}$ was calculated as normalized scattering cross section, $V_{particle}$ is the particle volume [5,6]. Fig. 8 shows that, with the Ag NPs diameter increasing from 20 nm to 100 nm, the $\sigma_s/V_{particle}$ increases. Moreover, comparing to the Ag NPs only, the Ag NPs-Ag film has stronger scattering effect. Theoretical and experimental have already verified that the leaking photons escaping from gain medium can be reflected and re-injected into the gain media by the external feedback of metallic film

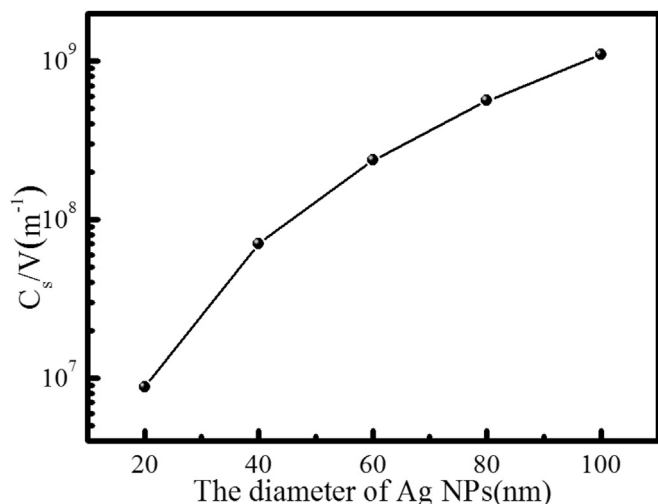


Fig. 8. The normalized scattering cross section at $\lambda = 435$ nm versus the Ag NPs size.

[30–32]. Fig. 5 shows that the gain media deposited on hybrid structure with 100 nm SiO₂ spacer has lower lasing threshold than that deposited on separate Ag NPs, which is due to the stronger scattering by external feedback of Ag film (the electric field intensities of separate Ag NPs and hybrid structure with 100 nm SiO₂ are about the same shown in Fig. 7). And Table 1 shows that with increase of Ag NPs, the $E_{th-50nm}$ is lower, which also suggests that with the increasing of Ag NPs size, the scattering effect of hybrid structure increases.

According to the discussion of two mechanisms about enhanced localized electric field and scattering effect of different sizes Ag NPs-Ag film hybrid structures, we can know that comparing with the independent Ag NPs or Ag film, the two mechanisms of enhanced localized electric field and scattering effect of Ag NPs-Ag film hybrid structure are all stronger, which lead to the significantly enhanced lasing. When the Ag NPs is small in hybrid structure (diameter ≤ 40 nm), comparing to the scattering effect, the enhanced localized electric field is the dominant role to enhance lasing. With the increase of Ag NPs diameter, the scattering effect increases, the enhanced localized electric field and scattering have comparable contribution on enhancing lasing. When the size is larger (diameter ≥ 80 nm), the scattering effect is the major mechanism for enhanced lasing. And it found that there is the lowest lasing threshold in device with 60 nm Ag NPs-Ag film hybrid structure, about a factor of 5.2 times than that of reference device, both the enhanced localized electric field and scattering effect play the important role on reduction of lasing threshold. At meanwhile, the negative effect of Ag film is avoided in gain medium to realize the lower pumped threshold than that of the neat gain medium.

4. Conclusion

In summary, avoiding the negative effect of metallic film on the lasing has been an important issue for the electrical pumped laser. In this article, we investigated the lasing properties based on different sizes Ag NPs-Ag film hybrid structures, the Ag NPs diameters ranged from 20 nm to 100 nm. The lowest lasing threshold was achieved by introducing 60 nm Ag NPs-Ag film hybrid structure. Comparing to the separate Ag NPs and Ag film, the Ag NPs-Ag film hybrid structure presented the more intense local electric field due to the plasmonics coupling between the Ag NPs and the Ag film. In the meantime, the re-injection of emitted light into the organic gain media by the Ag film could also enhance scattering strength. By studying the enhanced localized electric field and the scattering effect of Ag NPs-Ag film hybrid structure in experiment and Mie theory, we found that the enhanced localized electric field

and scattering all contribute to the enhanced lasing. When the Ag NPs is small in hybrid structure (diameter ≤ 40 nm), enhanced localized electric field is the major role for enhanced lasing; with the increase of Ag NPs size, the enhanced electric field and scattering all play the important roles on the enhanced lasing; when the size is larger (diameter ≥ 80 nm), compared with enhanced localized electric field, the scattering effect plays a main role on the enhancement of lasing properties. The lowest lasing threshold was achieved due to both the enhanced localized electric field and scattering effect. Our results could provide us a unique idea to effectively avoid the negative effect of metallic film and realize the lower pumped threshold. This method is expected to be a potential metal-modified technology for improving the lasing performance.

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