



# Emission enhancement of plasmonically modified YAG structures co-doped by $\text{Ce}^{3+}$ , $\text{Bi}^{3+}$ and $\text{Yb}^{3+}$ ions at low temperature

M. Kushlyk<sup>1</sup>, V. Tsiurma<sup>1,2</sup>, Ya. Zhydachevskyy<sup>2,3</sup>, V. Haiduchok<sup>3,4</sup>, I.I. Syvorotka<sup>3,4</sup>, D. Sugak<sup>3,4</sup>, M. Baláž<sup>5</sup>, A. Suchocki<sup>2</sup>



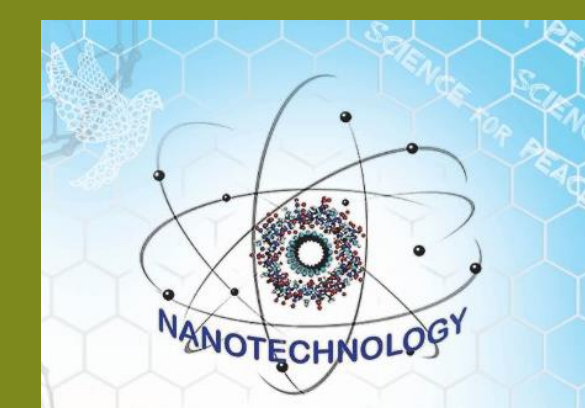
<sup>1</sup> Ivan Franko National University of Lviv, 79017, 107 Tarnavskoho Str., Ukraine

<sup>2</sup> Institute of Physics PAS, Al. Lotników 32/46, Warsaw 02-668, Poland

<sup>3</sup> Lviv Polytechnic National University, 79646, 12 Bandera, Lviv, Ukraine

<sup>4</sup> SRC „Carat“, 202 Stryjska Str., Lviv 79031, Ukraine

<sup>5</sup> Institute of Geotechnics SAS, Watsonova 45, 04001 Košice, Slovakia



## Abstract

In recent years, much attention has been paid to the plasmonic effect of metal nanoparticles formed on the surface of the down-converting phosphors [1, 2]. Combination of thin conversion layer, with silver plasmonic nanostructures leads to increase donor absorption and emission capacity.

High enhancement factors are caused by a close matching of the maximum of plasmon extinction and  $\text{Bi}^{3+}$  and  $\text{Ce}^{3+}$  emission bands. The enhancement in PL intensity can be explained by the interaction between the spontaneous recombination and surface plasmon in metal nanoparticles.

According to the FDTD simulation, near surface of NPs exhibits dipole and quadrupole resonance modes LSPR which high enhancement factor can lead to the high PL efficiency. The position of dipole mode of Ag NPs is in close vicinity to  $\text{Ce}^{3+}$  luminescence and leads to 300% of enhancement [3]. However, experimental research shows negligible changes in its PL enhancement with temperature decreasing. On the other hand, quadrupole mode of Ag plasmon resonance could be used for  $\text{Bi}^{3+}$  emission enhancement. However, because the relatively short penetration depth of SPR electric field of NPs gives rise to the limitations of luminescence enhancement efficiency.

## Methodology

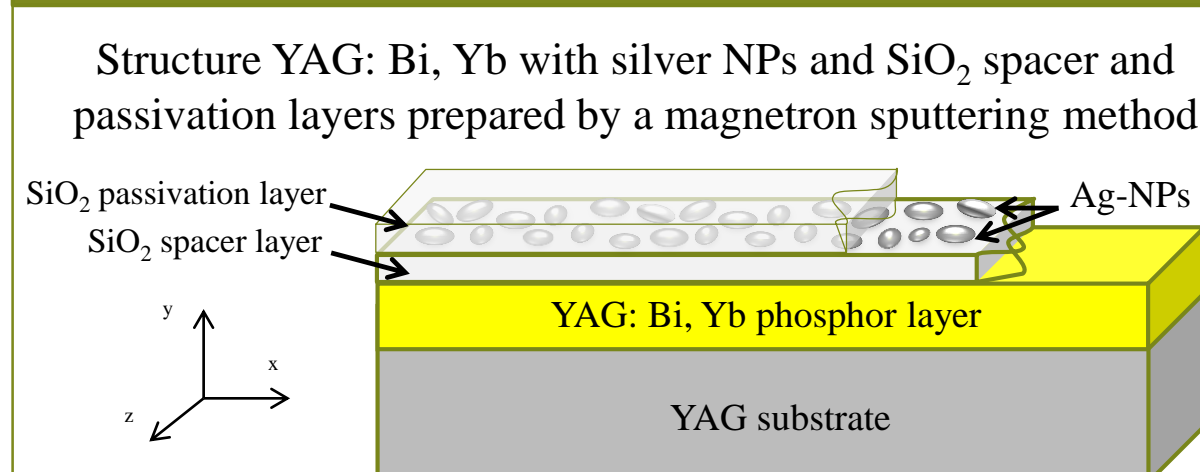
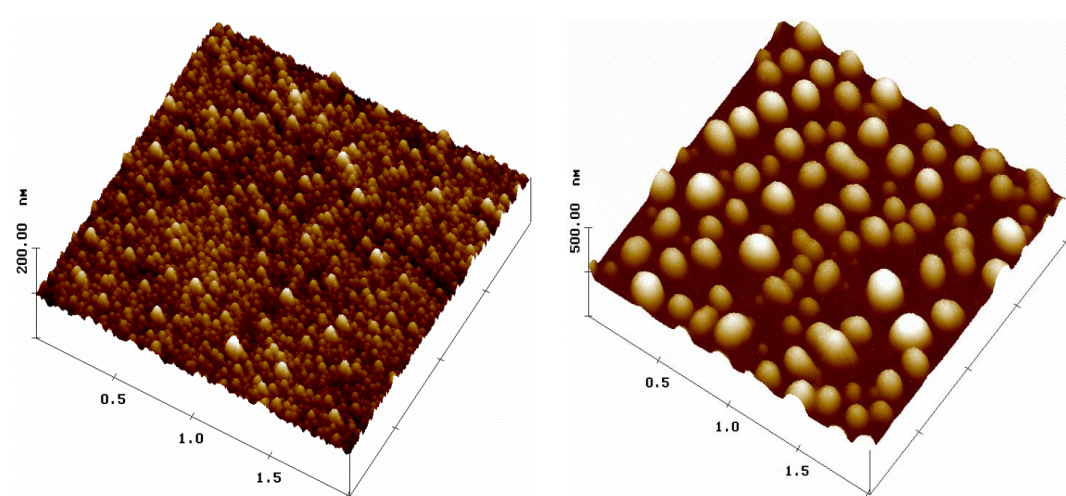


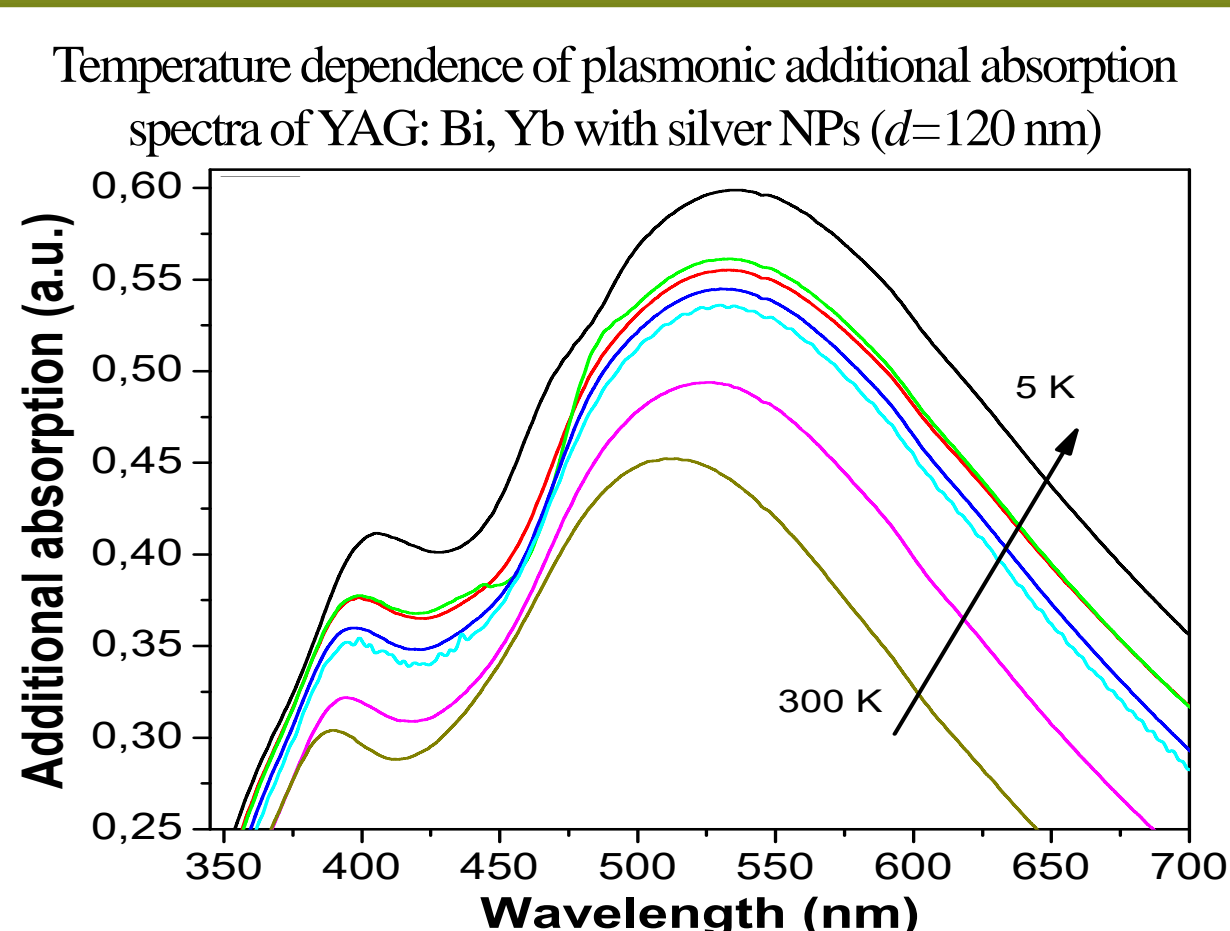
Table 1. Preparation parameters of samples

Sample	$\text{SiO}_2$ thickness, nm	Ag mass thickness, nm	Annealing parameters	$\text{SiO}_2$ passivation	Mean diameter, nm
3YYb-1-1	6	10	T=500 °C, t = 15 min	No	38.43
2YYb-2-1	10	50	T=400 °C, t = 40 min	Yes, 3 nm	70.8
2YYb-2-2	3	100	T=500 °C, t = 40 min	Yes, 3 nm	123.5
3YYb-2-2	15	150	T=500 °C, t = 40 min	Yes, 3 nm	181.6

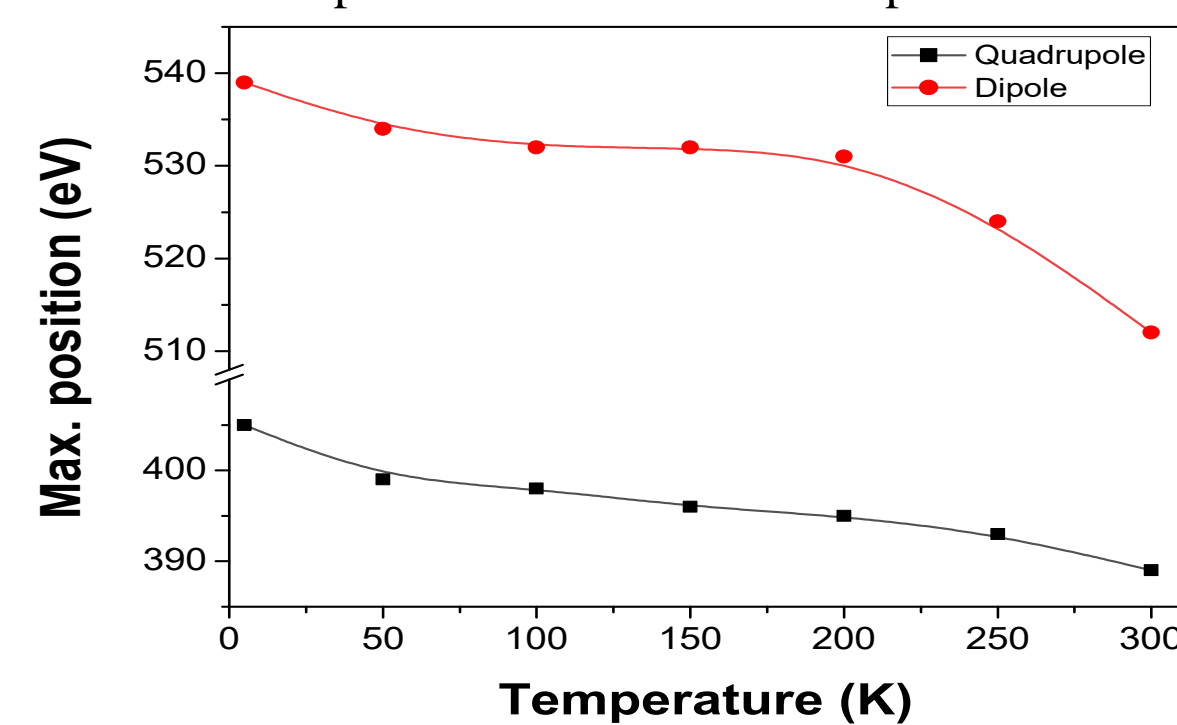
Morphology of silver NPs on surface of YAG: Bi, Yb with  $\text{SiO}_2$  spacer layer for “small” NPs ( $d < 50\text{nm}$ , left image) and “large” NPs ( $d > 100\text{nm}$ , right image)



## Optical Properties of SPR

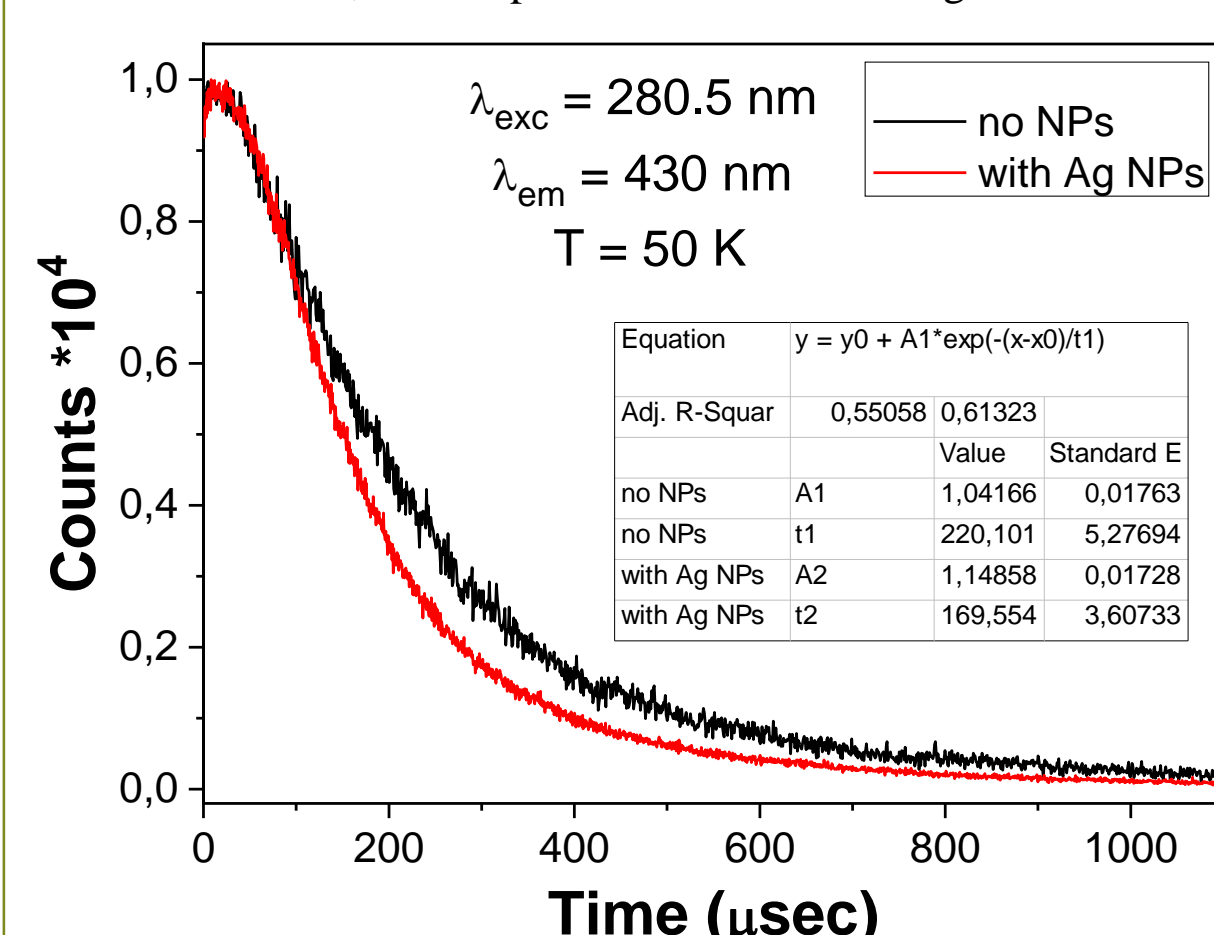


Position of maximums of dipole and quadrupole modes of SPR peaks as a function of temperature

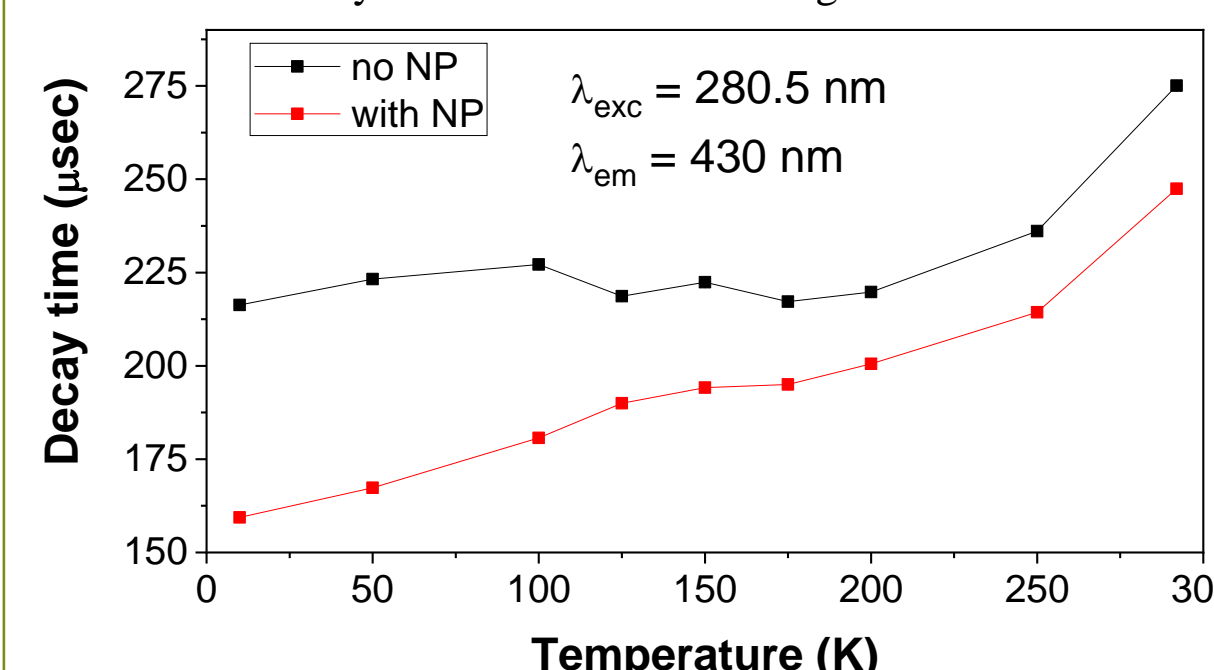


## Decay Kinetics

Decay kinetics temperature dependence of  $\text{Bi}^{3+}$  for two types of YAG: Bi, Yb samples with and without Ag NPs.

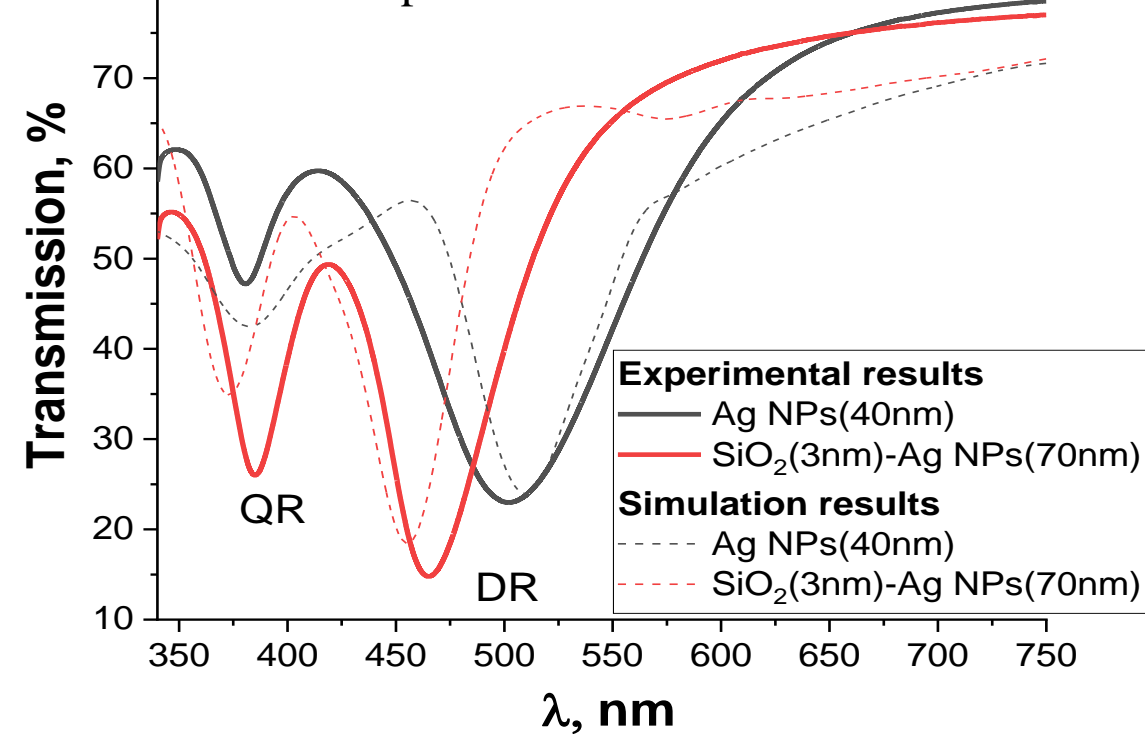


Temperature dependence of decay time of  $\text{Bi}^{3+}$  emission in YAG crystals with and without Ag NPs.

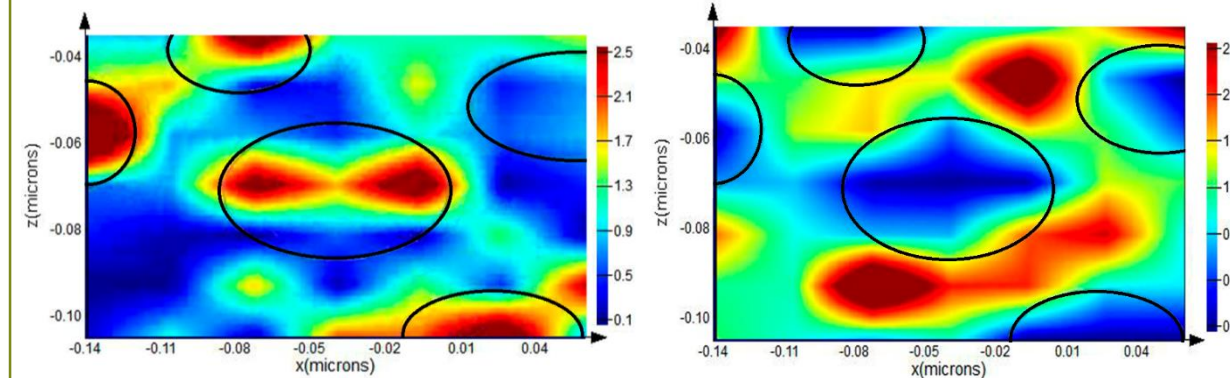


## Simulation Results

Transmission spectra samples with silver NPs that demonstrate QR – quadrupole resonance and DR – dipole plasmon resonance

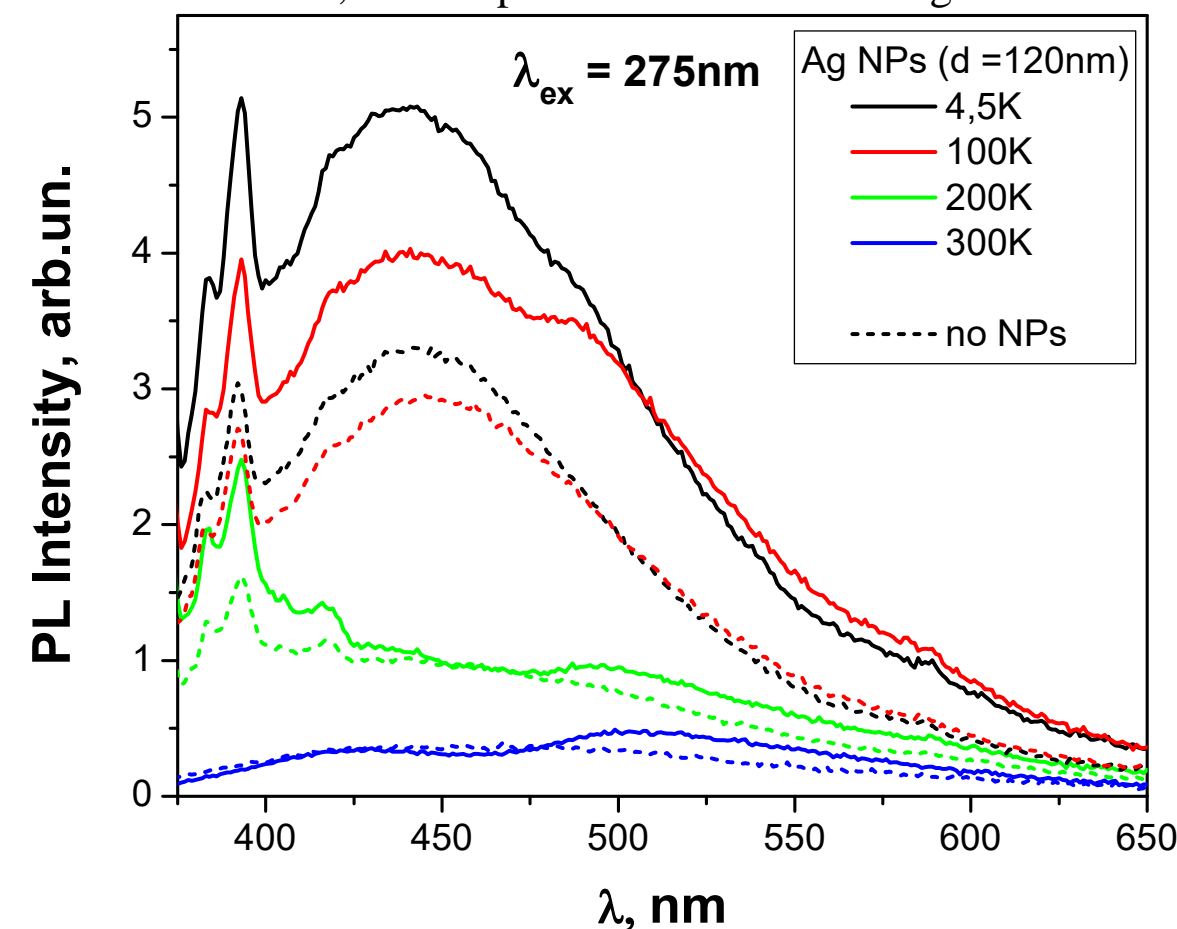


Map of the local electrical field on the samples surfaces for dipole ( $\lambda = 510\text{ nm}$ , left image) and quadrupole ( $\lambda = 375\text{ nm}$ , right image) resonances respectively obtained by finite-difference time-domain (FDTD) simulation

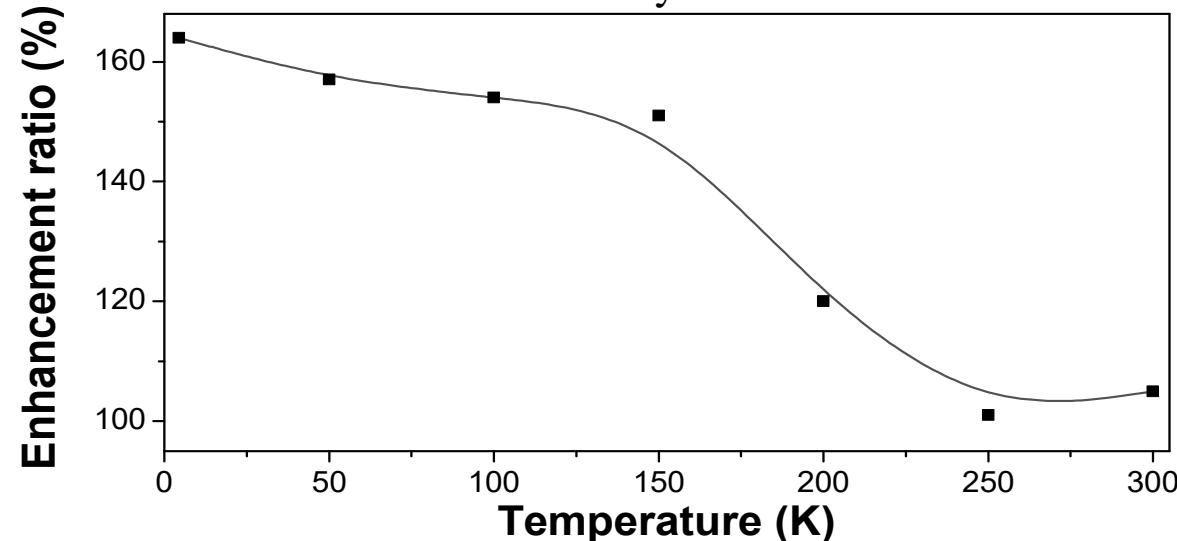


## PL Enhancement

Temperature dependence of PL spectra of  $\text{Bi}^{3+}$  for two types of YAG: Bi, Yb samples with and without Ag NPs.



Temperature dependence of enhancement ratio of  $\text{Bi}^{3+}$  emission in YAG crystals



## Conclusions

According to the FDTD simulation, near surface of NPs exhibits dipole and quadrupole resonance modes LSPR which high enhancement factor can lead to the high DC efficiency. Quadrupole mode of Ag plasmon resonance could be used for  $\text{Bi}^{3+}$  emission enhancement. However, because the relatively short penetration depth of SPR electric field of NPs gives rise to the limitations of luminescence enhancement efficiency. Two competing processes of enhancement and quenching of PL were obtained as a function of NPs mean size. Enhancement process prevails in case of NPs size larger than 50 nm.

Enhancement of  $\text{Bi}^{3+}$  emission in YAG epitaxial films with Ag NPs during cooling the samples below 200 K was observed. We obtained maximum value of enhancement factor near 170% at 4 K temperature. Such enhancement arises as the result better matching of quadrupole mode of SPR maximum with  $\text{Bi}^{3+}$  emission wavelength. The observed redshift of quadrupole peak is suggested to be caused by temperature changes of Ag,  $\text{SiO}_2$  and YAG dielectric constants.

## Acknowledgements:

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## Contact information:

Dr. Markiyana Kushlyk, Email: [markiyana.kushlyk@lnu.edu.ua](mailto:markiyana.kushlyk@lnu.edu.ua), [kushlykmarik@gmail.com](mailto:kushlykmarik@gmail.com)

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