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# Magneto-Optical Kerr Effects of Magnetic Garnet Thin Films Incorporating Gold Nanoparticles: *A Possible Coupling between Localized Surface Plasmons and Magneto-Optical Effects*

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# Introduction

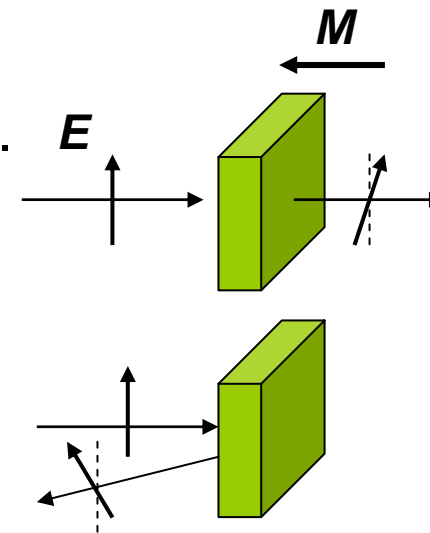
Magneto-Optical (MO) effects:

Linearly polarized light interacts with magnetized materials

The plane of polarization of light is rotated.

Transmission: MO Faraday effects

Reflection: MO Kerr effects



Coupling between MO Kerr effects and surface plasmon polariton (SPP)

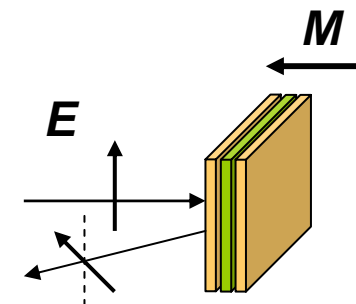
Au/Co/Au multi-layers in ATR geometry

Enhancement of MO Kerr FOM

Traveling SPP coupled to MO effect

Surface-enhanced MO media ?

*V. I. Safarov et al., PRL 73, 3584 (1994).*



# Motivation

MO medium + noble metal nanoparticles (MO inactive) , e.g., Au

Little is known about

MO Kerr effects coupled with localized SPP in the visible region

*Theoretical: V.A.Kosobukin, SPIE Proceeding 2535, 9 (1995).*

*Theoretical: M.Abe and T.Suwa, PRB 70, 235103 (2004).*

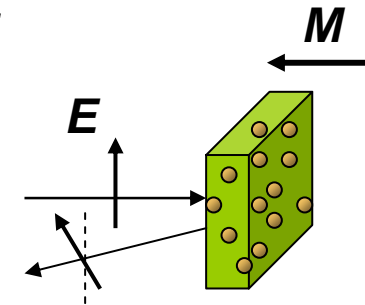
Localized SPP

Direct excitation by traveling light wave

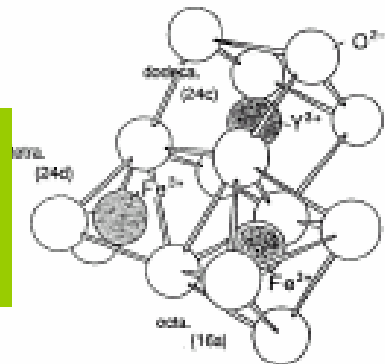
Large specific surface

Localized SPP influence MO effects

Novel MO effects of composite materials



$Y_3Fe_5O_{12}$ : yttrium iron garnet (YIG)  
Ferrimagnet  
Transparent in visible region



**Present study:**

**Preparation and MO Kerr study of  
YIG films incorporating Au nanoparticles**

# Preparation

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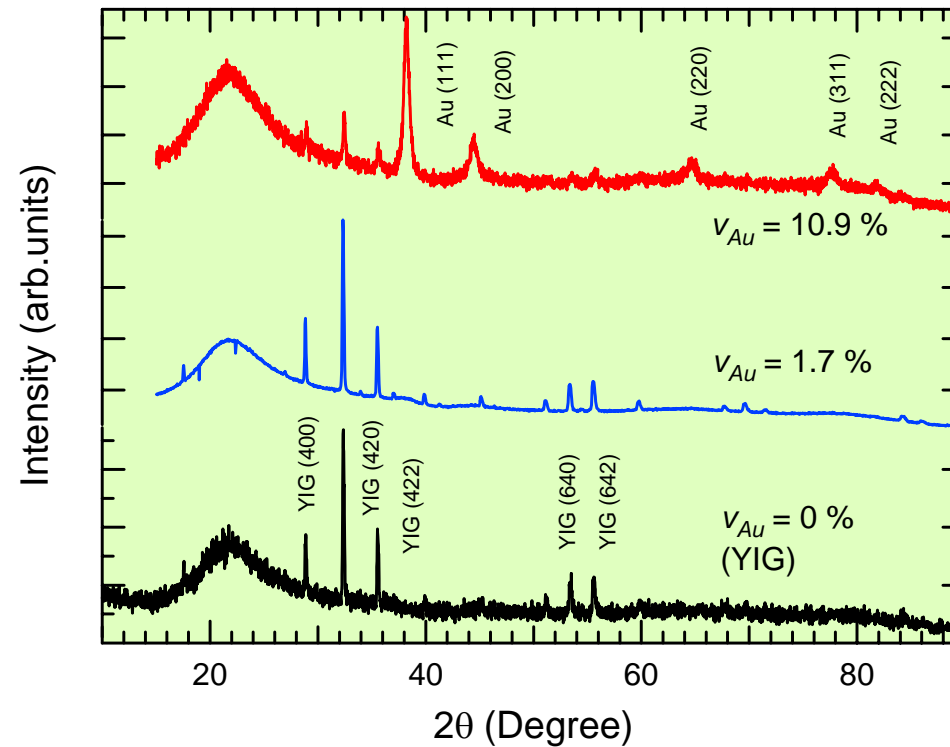
Film deposition: Co-sputtering method  
4 inch  $\phi$  YIG target, Au chips  
Ar gas 20mTorr  
Deposition rate  $\sim$ 6nm/min  
Film thickness: 200-500nm  
on quartz substrates

Au volume fraction ( $v_{Au}$ ) evaluated by EPMA  
 $v_{Au} = 0, 1.7, 10.9$  %

Post annealing: 900°C 30min in N<sub>2</sub> gas

# X-ray diffraction profiles

\*  $\lambda_{\text{CuK}\alpha} = 1.54 \text{ \AA}$   
\* after annealing



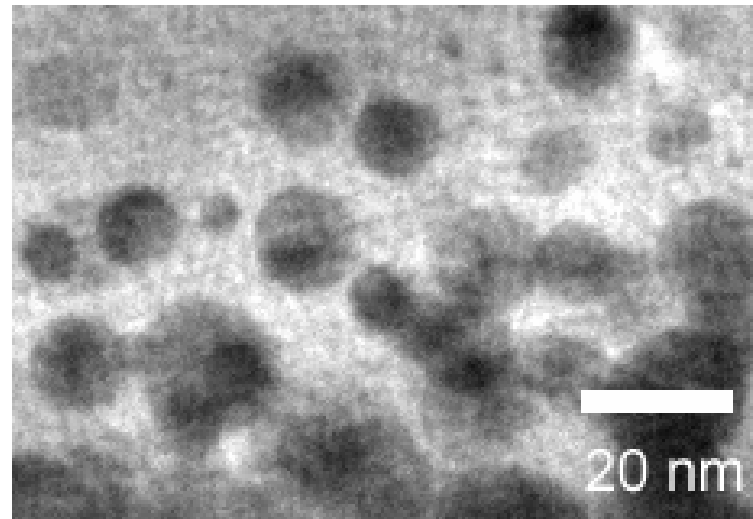
fcc-Au is grown and coexist with crystalline YIG in a film with  $v_{Au} = 10.9\%$

# TEM observation

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\* at 200 kV

Cross-sectional transmission electron microscopy (TEM)

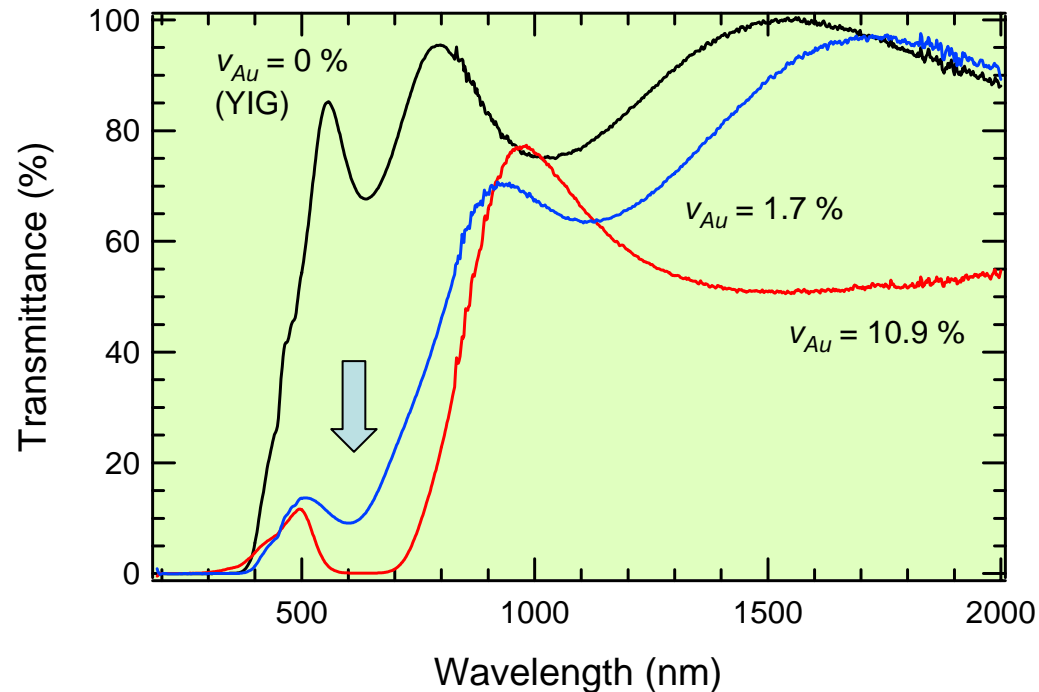


$V_{Au} = 10.9\%$

Spherical Au nanoparticles 12 nm in average diameter are embedded randomly in YIG matrices

# UV-Vis. transmission spectra

\* using a double-beam-type spectrometer



Below 500nm: Strong absorption due to CT-type transition in YIG

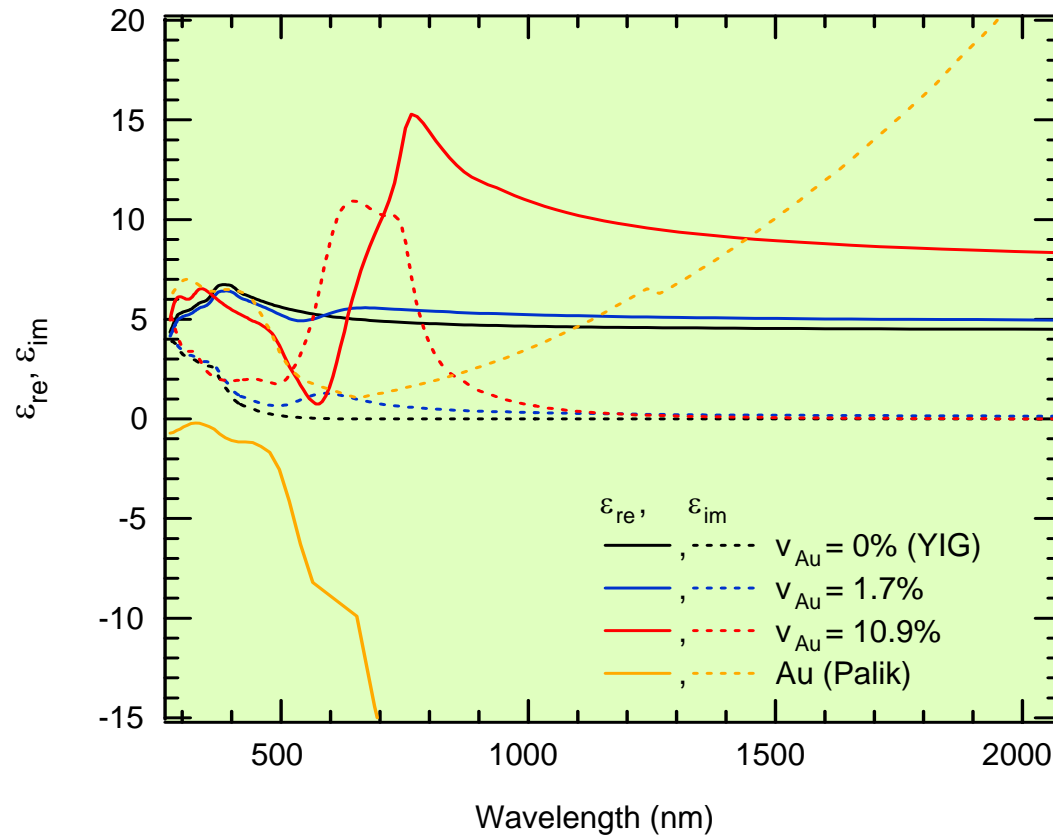
Above 500nm: Almost transparent for YIG

Incorporation of Au results in a strong absorption around 600nm

SPP of Au nanoparticles

# Spectroscopic ellipsometry

\* using Horiba UVISSEL NIR

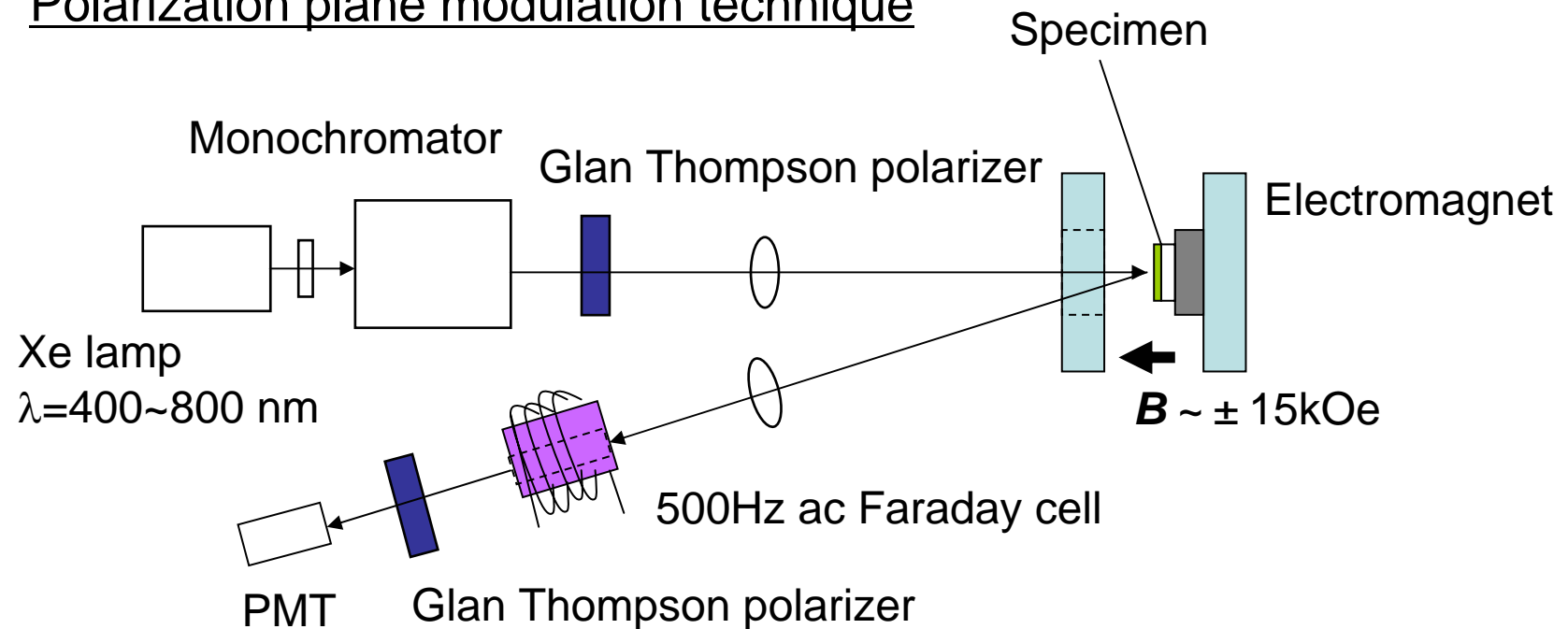


We observe dispersion of  $\epsilon$  due to SPP of Au nanoparticles in YIG



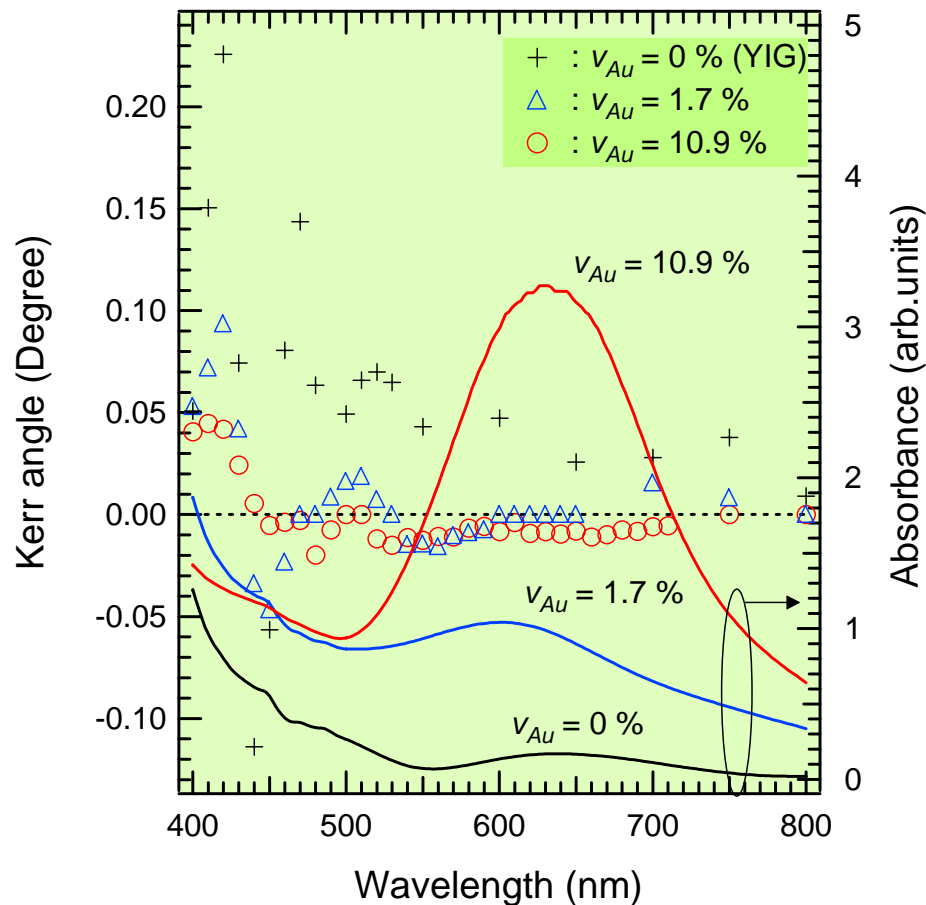
# Polar MO-Kerr effect measurements

## Polarization plane modulation technique



Faraday cell between two polarizers compensates  
the rotation of polarization plane due to Kerr effects of the specimen  
High angle resolution about  $0.001^\circ$

# Polar MO-Kerr spectra



$v_{Au} = 1.7\%$ :  
With Au nanoparticles  
**Negative Kerr rotation angle**  
around 550nm

$v_{Au} = 10.9\%$ :  
Increase in  $v_{Au}$   
Red shift of the region of  
Anomalous negative Kerr rotation

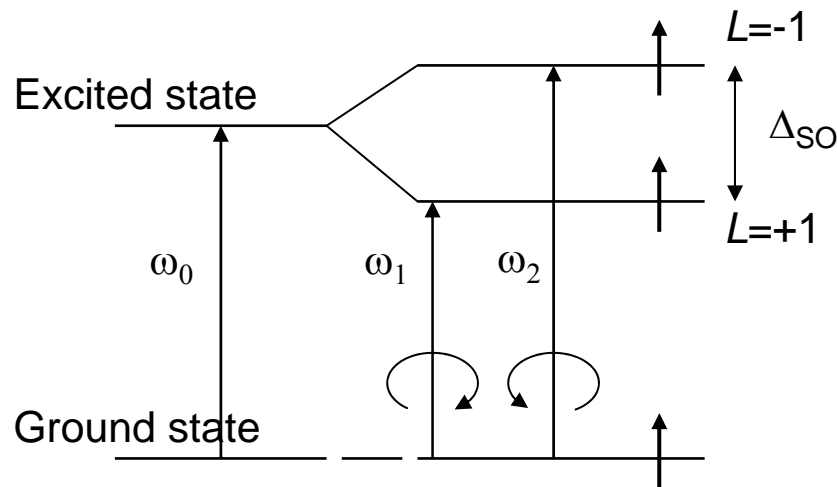
*Correlation between the region of  
negative Kerr rotation and  
SPP wavelength of Au nanoparticles*  
**A possible coupling  
between SPP and MO effects**

# CT-type transition

*F. J. Kahn et al., Phys.Rev.* **186**, 891 (1969).

Optical transition related to  $d$  electron in YIG:  
 Charge transfer (CT) type transition, crystal field (CF) type transition

1. CT-type transition: transition from  $p$  level to  $d$  level



$$\varepsilon'_{xy} = \frac{Ne^2 f_0 \Delta_{SO}}{2m\varepsilon_0 \omega \tau} \cdot \frac{\omega_0 - \omega}{\{(\omega_0 - \omega)^2 + \gamma^2\}^2}$$

$$\varepsilon''_{xy} = -\frac{Ne^2 f_0 \Delta_{SO}}{4m\varepsilon_0 \omega} \cdot \frac{(\omega_0 - \omega)^2 - \gamma^2}{\{(\omega_0 - \omega)^2 + \gamma^2\}^2}$$

MO effects:

Oscillator strength  $f_0$ , spin-orbit splitting  $\Delta_{SO}$ , frequency  $\omega$

# Possible origin

1. Charge transfer (CT) type transition around 490 nm is dominant in MO property of YIG in visible region

The **effective SO coupling** of excited state in CT transition is modified by localized SPP accompanied by an enhanced electric field

$$H_{so} = \left[ \frac{\hbar}{4m_0c^2} \right] \{ \nabla V \times \vec{p} \} \cdot \vec{\sigma}$$

Negative Kerr rotation angle  
cf: Modification of SO coupling constant in  $\text{Bi}_x\text{:Gd}_{3-x}\text{Fe}_5\text{O}_{12}$   
*Takeuchi, JJAP 14, 1903 (1975).*

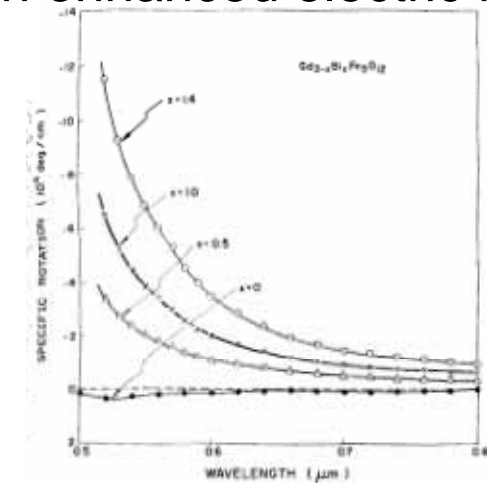


Fig. 1. Faraday rotation spectra of  $\text{Gd}_{3-x}\text{Bi}_x\text{Fe}_5\text{O}_{12}$ .

2. Enhancement of oscillator strength of crystal field (CF) type transition in YIG at 610 nm by relieving **the parity constrain**

# Summary

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- ◆ We have prepared YIG thin films containing small amount of Au nanoparticles.
- ◆ UV-visible transmission spectra shows strong absorption around 600 nm, which is caused by SPP of Au nanoparticles.
- ◆ Polar MO-Kerr spectra shows negative Kerr angle around 600nm, indicating the near-field coupling between MO Kerr effect and SPP.
- ◆ Possible origin for negative Kerr angle:
  1. Modification of spin-orbit coupling in CT-type transition by SPP with an enhanced electric field,
  2. Enhancement of oscillator strength of CF-type transition in YIG by SPP as an electric dipole