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Light Emission Property from Organic Dye Thin Films due to Excitation of Multiple Surface Plasmons

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1. INTRODUCTION

Surface plasmon resonance (SPR), that is, attenuated total reflection (ATR) has attracted much attention for measurements and sensing, since surface plasmons (SPs) resonantly excited at ultrathin metal surfaces are strongly influenced by conditions of the surfaces [1]. The ATR measurements utilizing SP excitations have been used to evaluate structure and optical properties of organic ultrathin films on metal ultrathin films, to estimate orientations of liquid crystal molecules [2-5] and as one of sensing methods [6,7]. The ATR methods have been also investigated for device applications, because of strong optical absorption and strong electric fields due to SP excitations [8, 9]. Recently, emission light at a resonant angle region of SP excitations was observed through the prism in the ATR Kretschmann configuration, when metal ultrathin films on the prism or organic ultrathin films on metal ultrathin films were directly irradiated from air by a laser beam [10,11]. The emission light depended upon resonant conditions of SPs in the Kretschmann configuration, and it is thought that multiple SPs were excited by means of the direct excitation of organic dye films by a laser beam, that is, reverse irradiation [11-13].

In this study, the emission light properties have been investigated in details for merocyanine (MC) and crystal violet (LB) films on silver (Ag) films using the conventional ATR method and the reverse irradiation in the Kretschmann configuration.

2. EXPERIMENTS

2.1 Sample preparation

Merocyanine (MC) and crystal violet (CV) were used as organic dye molecules and these Langmuir-Blodgett (LB) films were prepared by the LB method. MC is one of cyanine dyes and shows photoelectric properties [9] and p-type conduction. CV is one of triphenylmethyl groups and shows n-type conduction. The MC or CV molecules were mixed with arachidic acid (C20) for good depositions and the molar ratios of these mixtures were [MC or CV]: [C20] = 1:2.

In this study, two types of LB films were prepared. One was the MC LB film with 16 monolayers. Another was an alternate LB film with 16 monolayers of MC and CV LB monolayers (MC/CV LB film). The both LB thin films were deposited on the C20 LB films with 2 monolayers on Ag thin film. The Ag thin film of which thickness was about 50 nm was used as SP active layer.

The Ag thin film was deposited on a microscopic cover glass by a vacuum evaporation method.

2.2 A measuring system of emission light by the reverse irradiation in the Kretschmann configuration

Figure 1 shows a system to detect emission light through the prism when the sample was excited by the reverse irradiation in the Kretschmann configuration. The sample was located on the bottom of a half-cylindrical prism (BK-7, $n=1.512$ at $\lambda = 488$ nm) as shown in Fig. 1. The sample was irradiated at the vertical incident angle, $\theta_i=0^\circ$, by a p-polarized Ar⁺ laser beam at 488nm. The emission light was observed through the prism in the Kretschmann configuration [11-13]. The spectra of the emission light were measured at various emission angles where the light was observed. The emission light was detected with a photo-multiplier (PM) through an optical fiber and a spectrometer, and a photo-counting unit mounted on the computer.

3. RESULTS AND DISCUSSIONS

Figure 2 shows the emission light spectra from the prism/Ag/LB films, ((a): MC LB film, (b): MC/CV LB film) at various emission angles in the reverse irradiation measurements. The spectra of the emission light strongly depended on the emission angle. The peak wavelengths of the emission light became shorter as the emission angles increased. The relation between the wavelength and the emission angle agreed with the resonant condition of excitations of SPs in the Kretschmann configuration of the ATR method. It was thought that multiple SPs were simultaneously excited and the light was emitted through

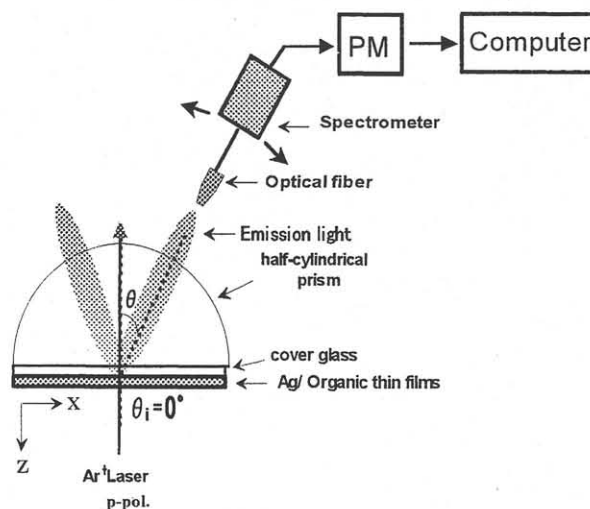


Fig. 1 Sample configuration and a measuring system.

the prism. The spectra in Figs. 2 (a) and (b) distributed around about 600nm and 700nm, respectively. These peaks almost corresponded with the PL peaks of MC and CV, respectively. Especially in Fig.2 (b), the emission light from MC was not observed. It was thought that these

and the spectra remarkably changed with the emission angles. It was concluded that the emission light was caused by simultaneous excitations of multiple SPs at the Ag/organic dye thin film.

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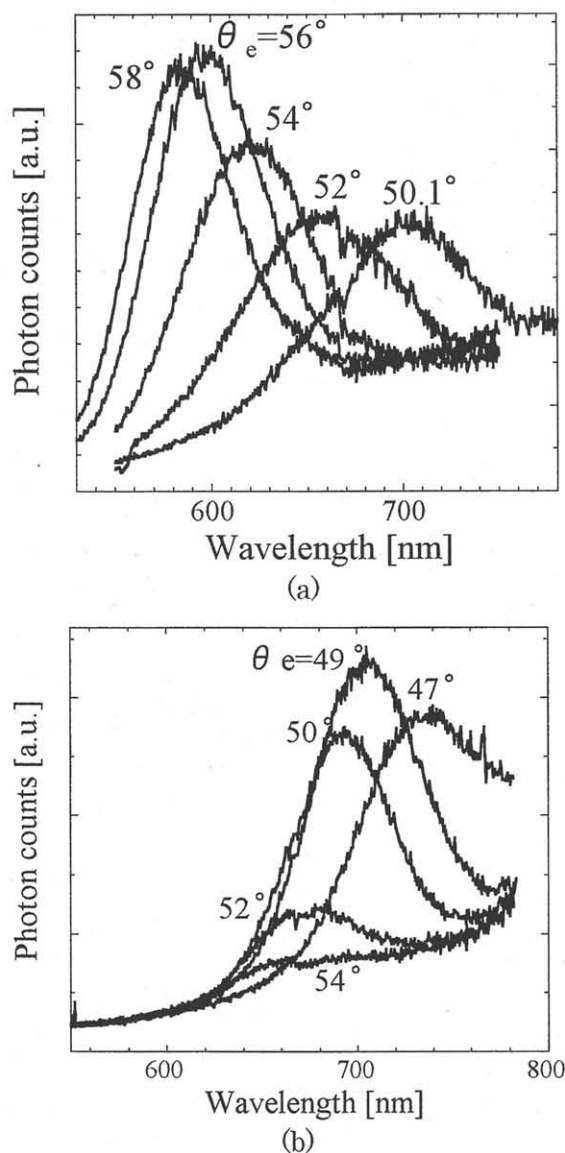


Fig.2. Emission light spectra at various emission angles due to the reverse irradiation. (a); MC LB films, (b); MC/CV LB films.

phenomena were caused by the energy transfer from MC to CV. It is thought that these properties may be used as a device with light emission of selectable colors and new sensing techniques.

4. CONCLUSION

The light emission property from Ag/organic dye thin films due to surface plasmon (SP) excitations was investigated using the reverse irradiation of the laser beam in the Kretschmann ATR configuration. By the reverse irradiation, emission light was observed through the prism,