Near-field fluorescence microscopy based on two-photon excitation with metal tips

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We present a new scheme for near-field fluorescence imaging using a metal tip illuminated with femtosecond laser pulses of proper polarization. The strongly enhanced electric field at the metal tip (∼15 nm end diameter) results in a localized excitation source for molecular fluorescence. Excitation of the sample via two-photon absorption provides good image contrast due to the quadratic intensity dependence. The spatial resolution is shown to be better than that of the conventional aperture technique. We used the technique to image fragments of photosynthetic membranes, as well as J-aggregates with spatial resolutions on the order of 20 nm. [S0031-9007(99)09142-5]

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The general aim of near-field optics is to extend optical microscopy beyond the diffraction limit. The introduction of the aperture probe for near-field microscopy [1,2] has allowed fluorescence imaging below the diffraction limit and has stimulated interests in many disciplines, especially the material and biological sciences. The widely adapted aperture approach is based on an aluminum-coated fiber tip [2] of which the foremost end is left uncoated to form a small aperture. Unfortunately, only a tiny fraction (<10⁻⁴ for a 100 nm aperture) of the light coupled into the fiber is emitted by the aperture because of the cutoff of propagation of the waveguide modes. The low light throughput and the finite skin depth of the metal are the limiting factors for resolution. Many applications require spatial resolutions that are not obtainable with the aperture technique. For example, in the effort on spectroscopic imaging of photosynthetic membranes [3], a spatial resolution of at least 20 nm was desired in order to resolve closely packed individual proteins in a lipid membrane. Moreover, the aperture technique has other practical complications: (1) It is difficult to obtain a smooth aluminum coating on the nanometric scale which introduces nonreproducibility in probe fabrication, as well as measurements; (2) the flat ends of the aperture probes are not suitable for simultaneous topographic imaging of high resolution; (3) the absorption of light in the metal coating causes significant heating and poses a problem for biological applications. To overcome these limitations, we present a new approach for high-resolution fluorescence imaging based on a sharp, laser-illuminated metal tip.

The use of laser-illuminated metal tips for near-field imaging has been discussed by many groups [4–13]. Most of the work has been limited to light scattering at the metal tip. The tip locally perturbs the fields at the sample surface; the response to this perturbation is detected in the far field at the same frequency of the incident light. The detected signals contain both near-field and far-field contributions and are generally dominated by topographic rather than spectroscopic information. The interpretation of the contrast in the recorded images is difficult.

In this Letter, we use the metal tip to provide a local excitation source for the spectroscopic response of the sample under investigation. The underlying principle of this approach has been outlined in a previous paper [11]. Excitation light of proper polarization induces a strongly enhanced field at the tip. The enhanced field consists mainly of nonpropagating (evanescent) components and is, thus, strongly confined to the tip end. The enhanced fields locally interact with the sample surface and generate a spectroscopic response that is detected in the far field at a different wavelength. The enhanced field at a gold tip is used here as a two-photon excitation source for molecular fluorescence. However, the same concept can be applied to other forms of linear and nonlinear imaging, such as imaging based on surface enhanced Raman scattering.

Rigorous electromagnetic calculations for a laser-illuminated (λ = 800 nm) gold tip with an end diameter of 10 nm have shown an intensity enhancement factor of roughly 3000 over the incident light [12]. We note that the incident wavelength is far away from the surface plasmon resonance. The field enhancement arises from a high surface charge density at the tip that is induced by the incident light polarized along the tip axis. In contrast, incident light with polarization perpendicular to the tip axis results in no field enhancement [11–13]. For a symmetrical tip perpendicular to a sample surface, the desired polarization for the field enhancement can be provided by side-on illumination. However, this leads to a large illumination area, giving rise to an overwhelming background signal. It was proposed that a tightly focused higher-order laser mode with a strong longitudinal field could be used for on-axis illumination [11]. In this work, we take the simpler approach of fabricating an asymmetrical metal tip. Figure 1(a) shows a field emission electron micrograph of a gold tip with an end diameter of 15 nm fabricated with etching and
The use of a metal tip only a few nanometers away from a chromophore raised concerns about fluorescence quenching by the metal tip. Fortunately, this problem was minimized for samples of molecular aggregates such as J-aggregates and fluorescent proteins in photosynthetic membranes. In particular, pseudoisocyanine (PIC) dye...
molecules of J-aggregates have extremely fast energy transfer along the aggregate length with an upper bound of coherent length of 50 nm [21], whereas the chlorophyll a (Chl a) and chlorophyll b (Chl b) molecules in light harvesting complex (LHCII) are separated by only a couple of nanometers according to the crystal structure [22]. Rapid energy transfer in these systems allows the fluorescence emission to take place far from the metal tip so that fluorescence quenching by the metal is reduced.

Figures 3(a) and 3(b) show the simultaneous images of topography and near-field two-photon excited fluorescence for J-aggregates of PIC dye molecules in polyvinyl sulfate (PVS). The method of sample preparation has been described by Higgins et al. [21]. The lower portions of Fig. 3 show the simultaneous cross sections taken across the aggregate strands (dotted white line). The arrows in the topography cross section indicate a feature with FWHM of 35 nm, while the corresponding feature in the emission cross section has a FWHM of 30 nm. This image demonstrates the superior spatial resolution of the present approach.

To further prove the near-field contrast, Fig. 4(a) shows the tip approach curve for the emission signal as a function of tip-sample distance at a particular point of an aggregate strand. The rapid decay of the emission signal at a tip-sample distance comparable to the tip diameter provides a clear proof of the principle of our approach.

Figure 4(b) shows the dependence of the emission signal on the average incident power while the tip was on top of an aggregate strand. The quadratic dependence proves the two-photon excited emission.

The emission spectrum of the brightest feature near the upper center of the image is shown in Fig. 5(a). With an excitation power level of \( \approx 15 \mu\text{W} \) and an acquisition time of 1 sec, the dotted line indicates the far-field contribution with no gold tip present. The solid line with a peak at 575 nm corresponds to the emission from J-aggregates with the gold tip present, which is similar to the far-field spectrum of the same sample. Although white-light continuum generation with gold tips was observed at higher power levels, no indication of such an emission occurred at the power levels at which the images were taken.

Figure 5(b) shows the near-field emission spectrum of a sample of photosynthetic membrane fragments from the chlamydomonas reinhardti strain (C2 mutant, doubly deficient in photosystem I and photosystem II [3]). The dotted line is the emission spectrum recorded when the tip was lifted. Figures 6(a) and 6(b) show the simultaneous images of topography and fluorescence of the sample, respectively. The topographic image indicates that micelles were formed when the membrane fragments were placed on the glass slip and allowed to dry. LHCII proteins embedded in lipid membrane have naturally fluorescent Chl a and Chl b molecules that are subject to two-photon excitation to the Soret band. The excitation...
light at 830 nm had an average power level of =25 μW. The dark region on the top is the region with the tip not present, resulting from only the far-field contribution with diffraction-limited resolution. When the tip was lowered to the surface, the near-field contrast appeared with many fluorescent features. The high topographic feature allowed correlation of the topographic and spectroscopic information. The high topographic feature (=400 nm) in the upper right-hand corner did not give any fluorescence signal. Most of the fluorescence features are correlated with topographic features. The cross sections below in Fig. 5(b) show four fluorescent features which all have a FWHM of =20–30 nm and similar intensities. Some features in the fluorescence image are on the order of 15 nm, with a signal-to-background ratio of better than 10:1. Transmission electron microscope micrographs as well as electron crystallography have shown that the trimer LHCII protein has a two-dimensional unit cell dimension of =13 nm [22,23], containing at least 7 Chl a and 5 Chl b per monomer [22]. We tentatively attribute the four peaks to individual units of LHCII in the micelles. The same features show up in the retrace images but reduced in emission intensity due to photobleaching. The spatial resolution of our method is approaching the limit of the size of individual protein molecules inside biological membranes.

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FIG. 6. Simultaneous topographic image (a) and near-field two-photon excited fluorescence image (b) of photosynthetic membrane fragments from *chlamydomonas reinhardtii* (C2 mutant) algae on a glass substrate. The corresponding topographic and emission cross sections along the dashed lines (A-B) are shown. The four emission peaks in the box have a FWHM of =20 nm.