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Velocimetric third-harmonic generation microscopy: micrometer-scale quantification of morphogenetic movements in unstained embryos

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We demonstrate the association of third-harmonic generation (THG) microscopy and particle image velocimetry (PIV) analysis as a novel functional imaging technique for automated micrometer-scale characterization of morphogenetic movements in developing embryos. Using a combined two-photon-excited fluorescence and THG microscope, we characterize the optical properties of Drosophila embryos and show that sustained THG imaging does not perturb sensitive developmental dynamics. Velocimetric THG imaging provides a quantitative description of the dynamics of internal structures in unstained wild-type and mutant embryos. © 2004 Optical Society of America

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The development of animal embryos exhibits a complex ensemble of cell movements that are highly regulated in time and space. Investigating these complex dynamic processes remains a challenge in biology. In particular, control of morphogenetic movements involved in Drosophila melanogaster embryo development serves as a major model for developmental genetics. However, Drosophila embryos exhibit fast development dynamics and are highly scattering for visible light at early stages, which limits direct imaging of dynamic processes. In addition, fluorescent labeling can introduce unwanted perturbations and might be difficult to obtain in complex mutants. As a consequence, the mechanical behavior of specific cells and tissues is not completely understood in many cases. In this context, novel noninvasive techniques for visualizing and quantifying morphogenetic movements in vivo should have a significant effect on developmental biology.

Third-harmonic generation (THG) microscopy has been proposed as a novel general-purpose technique for obtaining structural images of transparent objects with micrometer three-dimensional resolution comparable with that of two-photon-excited fluorescence (2PEF) microscopy and was demonstrated recently in biological samples such as cells and zebrafish embryos. In this Letter we extend THG microscopy to the quantitative measurement of tissue velocity fields inside unstained opaque embryos by combining THG microscopy and particle image velocimetry (PIV) analysis techniques adapted from hydrodynamics. First we use a combined 2PEF–THG approach to characterize the optical properties of Drosophila early embryos. Then we demonstrate that sustained THG embryo imaging does not perturb the dynamics of sensitive developmental processes. Finally, we illustrate velocimetric THG imaging by micrometer-scale quantification of morphogenetic movements in live unstained wild-type and mutant embryos.

We characterized early Drosophila embryo properties (Fig. 1) by using a custom-built combined 2PEF–THG microscope incorporating a femtosecond titanium:sapphire oscillator (Coherent, Inc.), an optical parametric oscillator (APE), a galvanometer mirrors (GSI Lumonics), water-immersion objectives (Olympus), and photon-counting photomultiplier modules (Electron Tubes). 2PEF was epidetected when green fluorescent protein- (GFP-) expressing embryos were excited at 820–920 nm. Alternatively, THG was detected in the transmitted direction when embryos were excited at 1180 nm. In either case, the signal was selected by use of appropriate filters (Chroma). Combined 2PEF–THG imaging of embryos with nuclear GFP labeling illustrates the complementarity of these imaging modalities (Fig. 1). At the onset of gastrulation (the universal primary morphogenetic process in animal development), Drosophila embryos comprise a single cell layer surrounding a heterogeneous medium (yolk), which is totally opaque in transmitted light microscopy. As expected, GFP-labeled nuclei produce a strong 2PEF signal when they are excited at 820 nm. THG is enhanced for
micrometer-sized optical heterogeneities. When embryos are illuminated at 1180 nm, lipid droplets that are present around the nuclei and at the periphery of the central yolk region produce a readily detectable THG signal \((5-10 \times 10^6 \text{ detected photons/s in our imaging conditions})\). These droplets have typical sizes of the order of \(0.5 \mu\text{m}\), which contribute to THG signal enhancement.

The key parameter for nonlinear microscopy of tissues is the scattering mean free path of excitation light \(l_s^{(\text{ex})}\) because it defines the increase in laser power that is necessary to compensate for the scattering of excitation light with depth. When absorption is neglected and with the assumption of constant collection efficiency, the detected signal for an \(n\)-th-order process locally scales as \(\exp\left[-z/l_s^{(\text{ex})}\right]^{-n}\) for a given incident power, where \(z\) is the imaging depth. We estimated \(l_s^{(850)}\) (850-nm excitation) at different depths within embryos at the onset of gastrulation and found that three regions with distinct optical properties could be defined: (i) the peripheral region (nuclei) for which \(l_s^{(850)} \sim 100 \mu\text{m}\), (ii) a highly scattering region at the yolk surface \(l_s^{(850)} \sim 25 \mu\text{m}\), and (iii) the internal yolk \(l_s^{(850)} \sim 100 \mu\text{m}\). As is evident from THG images, the highly scattering nature of the yolk's surface is related to the density of lipid droplets in this region, which strongly hampers inner tissue imaging by conventional techniques. We estimated an average scattering length \(l_s^{(1180)}\) for THG imaging over the entire yolk region (see Fig. 1). We found that \(l_s^{(1180)} \sim 90 \mu\text{m}\), indicating that THG microscopy performs well deep within embryos owing to the reduced scattering of excitation wavelengths in the 1100–1300-nm range.

Rapid THG imaging of embryos with \(~80\)-MHz pulse trains requires instantaneous intensities in the TW/cm\(^2\) regime (90-mW average power with a 0.8-N.A. objective). Although THG microscopy is believed to be noninvasive because harmonic generation processes do not involve energy deposition in the sample, the possibility of laser-induced perturbation with such intensities cannot be excluded without experimental evidence. As a matter of fact, we observed bubble formation in isolated cells with slightly higher pulse energy in the same wavelength range (data not shown). To establish the validity of THG microscopy for measuring the dynamics of developmental processes we monitored the process of cellularization, a critical and temperature-sensitive dynamic event of embryonic cells. Cellularization involves oocyte plasma membrane folding between nuclei, resulting in the formation of individual cells. The rate and the completion of cellularization front invagination (CFI) are sensitive indicators of the integrity of cytoskeleton dynamics. We found that CFI can be followed in vivo by THG microscopy because it induces changes in the distribution of lipid droplets about the nuclei. We compared the CFI rate in control cytoskeleton-labeled GFP (sGMCA) embryos by using 2PEF microscopy and in unlabeled wild-type embryos by THG microscopy. As illustrated in Fig. 2, we found similar dynamics and phase durations for the two imaging modalities. These data also corroborate transmitted-light measurements (not shown). We conclude that for these imaging conditions (1180 nm, 80 MHz, 0.9 nJ, 0.8 N.A., 120-μm/ms scan speed, 20 s between successive images), sustained THG imaging does not induce significant perturbation. Moreover, we observed a normal survival rate (85 ± 3%) after long-term imaging, which further confirms the validity of THG microscopy for measuring embryo developmental dynamics.

To provide a micrometer-scale quantitative description of morphogenetic movements, we estimated instantaneous velocity fields from THG image sequences of developing embryos by use of PIV, a technique commonly used in hydrodynamics. PIV relies on correlation calculations and can extract velocimetric data with subpixel resolution even when moving structures are not clearly defined in the images, in contrast to standard tracking methods. THG microscopy is sensitive to micrometer-sized optical inhomogeneities and provides rich structural information throughout entire unstained embryos. As a consequence, THG image sequences are ideally suited for correlation analysis (Fig. 3). PIV analysis was performed with the MatPIV software package with \(64 \times 64\) pixel interrogation windows. As shown in Fig. 3, velocimetric analysis of THG data provides information about tissue...
THG microscopy and PIV as a functional imaging technique to conventional imaging techniques. Our experiments establish the combination of 2PEF–THG data to provide insight about optical properties of embryos. We showed that this approach is applicable to early Drosophila embryos, a major model in developmental genetics that is difficult to observe with conventional imaging techniques. Our combined 2PEF–THG data provide a simultaneous description of tissue and internal structure dynamics and reveals their coupling.

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