Fourier-transform coherent anti-Stokes Raman scattering microscopy.
Jennifer Ogilvie, Emmanuel Beaurepaire, Antigoni Alexandrou, Manuel Joffre

To cite this version:
Jennifer Ogilvie, Emmanuel Beaurepaire, Antigoni Alexandrou, Manuel Joffre. Fourier-transform coherent anti-Stokes Raman scattering microscopy.. Optics Letters, Optical Society of America, 2006, 31 (4), pp.480-2. 10.1364/OL.31.000480 . hal-00094226

HAL Id: hal-00094226
https://hal.archives-ouvertes.fr/hal-00094226
Submitted on 3 Oct 2013

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L’archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d’enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.
Fourier-transform coherent anti-Stokes Raman scattering microscopy

Jennifer P. Ogilvie,* Emmanuel Beaurepaire, Antigoni Alexandrou, and Manuel Joffre

Laboratoire d’Optique et Biosciences, Centre National de la Recherche Scientifique Unité Mixte de Recherche 7645, Institut National de la Santé et de la Recherche Médicale U696, Ecole Polytechnique, 91128 Palaiseau Cedex, France

Received September 13, 2005; accepted October 14, 2005; posted November 10, 2005 (Doc. ID 64777)

We report a novel Fourier-transform-based implementation of coherent anti-Stokes Raman scattering (CARS) microscopy. The method employs a single femtosecond laser source and a Michelson interferometer to create two pulse replicas that are fed into a scanning multiphoton microscope. By varying the time delay between the pulses, we time-resolve the CARS signal, permitting easy removal of the nonresonant background while providing high resolution, spectrally resolved images of CARS modes over the laser bandwidth (~1500 cm\(^{-1}\)). We demonstrate the method by imaging polystyrene beads in solvent. © 2006 Optical Society of America

OCIS codes: 170.5660, 020.4180, 180.6900, 020.1670, 300.6410, 300.6450.

Multiphoton microscopy is an important tool that is increasingly used in biological research owing to its ability to produce three-dimensional images of complex samples. To probe particular species, fluorescent labels are most commonly used but have several disadvantages, including photobleaching and possible modification of the system under study. Coherent anti-Stokes Raman scattering (CARS) microscopy offers the advantage of providing endogenous contrast based on the inherent vibrations of different chemical species. Because of the nonlinear nature of the involved light–matter interaction, CARS microscopy shares many of the benefits of multiphoton microscopy while avoiding problems associated with fluorescent labeling. In this Letter we present a Fourier-transform-based method of CARS microscopy. The technique employs a single broadband laser source and provides a simple approach to obtaining high-resolution spectrally resolved CARS images. We demonstrate the method by spectrally resolving images of polystyrene beads immersed in solvent.

CARS is a nonlinear scattering process in which light of different frequencies interacts through the third-order susceptibility of the medium. Two photons, of frequencies \(\omega_p\) and \(\omega_s\), such that the difference frequency \(\omega_p - \omega_s\) matches a vibrational resonance in the sample, cause the scattering of a third photon, of frequency \(\omega_p\), to produce a higher-energy anti-Stokes photon of frequency \(\omega_p - \omega_s + \omega_p\), as shown in Fig. 1(a). Other nonlinear processes that are not related to any vibrational resonances can also occur, producing a nonresonant background that can obscure the CARS signal [see Fig. 1(b)].

Since its first demonstration in 1982, most CARS microscopy implementations to date have employed picosecond pulses and have reduced the nonresonant background through the use of near-IR excitation and polarization selection. While using picosecond pulses reduces the nonresonant background, it also necessarily restricts the spectral range of the CARS method, such that single CARS modes are imaged at a time. Obtaining CARS spectral images over the entire fingerprint region, where molecular spectra are most distinguishable, is important for simultaneous imaging of multiple species and thereby significantly enhances the capabilities of CARS microscopy. Picosecond implementations of CARS microscopy can produce broadband CARS spectra by the time-consuming process of tuning one of the excitation frequencies. A more rapid approach has been to combine picosecond and femtosecond excitation, providing CARS images with up to ~200 cm\(^{-1}\) of bandwidth.

The use of femtosecond pulses provides yet larger spectral bandwidth but requires more elaborate methods for dealing with the nonresonant background.
ground, which increases with decreasing pulse duration. Recently CARS images were obtained using broadband femtosecond pulses, which can potentially excite simultaneously all vibrational modes of frequencies smaller than the laser bandwidth, as in impulsive stimulated Raman scattering. In this approach, mode selectivity is achieved using various techniques such as phase-only pulse shaping, spectral focusing with chirped pulses, and selecting a narrowband pump beam by use of amplitude-only pulse shaping associated with spectral broadening of the Stokes beam and frequency-resolved detection of the CARS emission. In the case of phase-only pulse shaping, a sinusoidal spectral phase was applied such that the exciting pulse actually consisted of a quasi-periodic pulse sequence that allows mode selectivity as in multiple-pulse impulsive stimulated Raman scattering. Frequency-resolved Raman spectra were then obtained through a Fourier transform of the data measured as a function of the sequence pseudoperiod. However, although multiple pulse sequences are ideal for driving a given Raman mode, it is well known that the entire Raman spectrum can also be obtained in a more straightforward way through a Fourier transform of time-domain impulsive stimulated Raman scattering data. Our approach to single-laser CARS microscopy is thus to use a broadband laser source to record the CARS emission as a function of the time delay between only two identical collinear excitation pulses and then to obtain the CARS spectrum through a simple Fourier transform. Furthermore, windowing out the contribution around zero time delay allows a straightforward removal of the nonresonant background, similarly to a previous time-domain CARS experiment performed at a fixed time delay and an interferometric CARS experiment that demonstrated a difference in time evolution between resonant and nonresonant signals.

In the research reported in this Letter we generate spectrally resolved CARS images over a bandwidth of $\sim 1500 \text{ cm}^{-1}$, using a single broadband Ti:sapphire oscillator. In our time-domain Fourier-transform CARS (FTCARS) method, two broadband pulse replicas are created in a Michelson interferometer. The first pulse provides the pump and Stokes fields to impulsively excite a vibrational coherence, while the second, time-delayed pulse probes this coherence. The bandwidth obtainable is determined by the spectral width of the excitation source, which must impulsively excite the Raman modes to be imaged.

We demonstrate FTCARS microscopy by using a home-built laser-scanning microscope and a $\sim 20 \text{ fs}$ Ti:sapphire oscillator (Ti:sapph; center wavelength, 820 nm) excitation source. Before it reached the microscope, the 100 MHz pulse train was sent into a Michelson interferometer, as shown in Fig. 1, producing two collinear pulses that could be variably time delayed with respect to each other. This delay was accurately monitored with a He–Ne laser (633 nm). Precompensation of the microscope dispersion was performed with a fused-silica prism compressor, which was optimized via second-order autocorrela-

![Figure 2](image-url)

**Fig. 2.** (Color online) (a) Two-photon fluorescence image of a 20 $\mu$m polystyrene bead immersed in 2-propanol. (b) Spectrally resolved FTCARS image of the line in (a) [see (c) for scale]. (c) FTCARS spectra at positions 1 and 2 in (b).
816 cm\(^{-1}\) is clearly resolved,\(^{21}\) as is the dominant mode in polystyrene at 1005 cm\(^{-1}\), as well as smaller-amplitude modes at 783 and 620 cm\(^{-1}\) [Fig. 2(c)].\(^{21,22}\) The peak near 800 cm\(^{-1}\) in the polystyrene spectrum may have contributions from both the 2-propanol and the polystyrene as a result of limited axial resolution (~30 \(\mu\)m). The CARS signal exhibits attenuation at the solvent–bead interface, in agreement with other reports.\(^{23}\) The spectral line image in Fig. 2(b) was obtained by slowly scanning the time delay while simultaneously scanning the microscope’s galvanometric mirrors at ~200 Hz. A time series was then reconstructed for each pixel in the line, from which the CARS spectrum for each pixel was determined after averaging over 100 time-delay scans. Figure 3 shows the direct time-domain CARS signal that corresponds to location 1 in Fig. 2(b), where the ~1000 cm\(^{-1}\) mode in polystyrene dominates the signal, as shown in the inset of Fig. 3. The peak at zero delay reflects the nonresonant response, which is easily windowed out before the Fourier transform, yielding the CARS spectra shown in Fig. 2(c). The maximum time delay actually used was 2.5 ps, yielding a spectral resolution of 13 cm\(^{-1}\), a value that can easily be reduced by use of longer delays. Furthermore, scanning the time delay more rapidly and improving data processing should significantly reduce the total acquisition time (currently 10 min).

In summary, we have proposed a Fourier-transform technique for CARS microscopy that employs a single broadband laser source and a Michelson interferometer. We have demonstrated the technique by spectrally imaging a polystyrene bead in 2-propanol. This approach offers a straightforward method for removing the nonresonant background from CARS images while sharing the advantages of other Fourier-transform spectroscopies (no need for a grating spectrometer, high throughput, arbitrarily high spectral resolution limited only by the maximum time delay). Compared with methods based on pulse shaping, FTCARS microscopy offers a more compact optical setup and higher spectral resolution. Furthermore, when the possibility of recording two-photon absorption spectra as well is taken into account,\(^{13}\) Fourier-transform microscopy should make possible the simultaneous acquisition of frequency-resolved two-photon and CARS images.

We thank the Délégation Générale pour l’Armement for supporting this research. J. P. Ogilvie’s e-mail address is jogilvie@umich.edu.

*Present address, Departments of Physics and Biophysics Research Division, University of Michigan, Ann Arbor, Michigan 48109-1040.

References