

Engineering optical fibres for nonlinear optical endoscopy

Min Gu

Centre for Micro-Photonics, Faculty of Engineering and Industrial Sciences
Swinburne University of Technology, VIC 3122, Australia

mgu@swin.edu.au

Telephone: (61) 3 92148776, Fax: (61) 3 9214 5435

Abstract: This paper is to review the recent development of fibre-optical nonlinear optical microscopy for 3D endoscope tissue imaging. The new compact probes are designed with a double-clad photonic crystal fibre or a double-clad fibre.

Introduction

One of the implementations toward a compact nonlinear optical microscope is the adoption of fibre optics. The invention of nonlinear optical microscopy based on multiphoton fluorescence and harmonic generation has provided a powerful tool for 3D imaging through tissue. The development of compact nonlinear optical microscopy or endoscopy is motivated by three novel applications. First, it provides high-resolution *in vivo* cellular imaging under conditions in which conventional bulk microscopy cannot be used. The examples of these conditions include imaging hollow tissue, imaging within solid organs, imaging freely-moving animals, and remote delivery and collection in a minimally invasive manner. Second, nonlinear optical endoscopy is a critical tool for long term imaging studies such as the investigation of the cellular effect of aging and the development of new *in vivo* assays for testing of drugs and therapeutics. Third, it facilitates the development of minimally invasive clinical diagnostics and surgical procedures.

Various efforts have been made to produce a compact probe based on single mode fibre optics [1-3]. It has been shown that the use of a single-mode fibre does not

efficiently facilitate both the delivery of near-infrared pulsed laser light and the collection of visible nonlinear optical signal [2, 4-6]. Here we report on the new development of a fibre-optical nonlinear optical endoscope probe towards 3D high-resolution tissue imaging. The signal of this new probe is increased by two orders of magnitude.

Nonlinear optical micro-probe with a double-clad photonic fibre

The probe comprises of a double-clad photonic crystal fibre (PCF), a micro-prism, a two-dimensional (2D) microelectromechanical system (MEMS) mirror, and a gradient-index (GRIN) lens [3, 7-11]. A pulsed laser beam of wavelength 800 nm generated from a turnkey Ti:Sapphire (Spectra Physics, MaiTai) is launched into the double-clad PCF through a prechirp unit consisting of a grating pair. The endoscope probe allows side-view imaging at a working distance of approximately 200 μm . The back-scattering nonlinear optical signals propagating through the fibre and the MEMS mirror are collected by a photomultiplier tube.

The imaging capability of the nonlinear optical endoscope is demonstrated by the two-photon fluorescence imaging with 10 μm diameter fluorescent microspheres. It shows that the 2D MEMS mirror enables the efficient delivery of the light beam over the broadband wavelength range and the smooth response for image acquisition. *In vitro* three-dimensional (3D) images of internal organ tissue and cancer tissue have been achieved with a penetration depth of up to

100 μm and axial resolution of approximately 10 μm . Epithelial cells and intestinal crypts in 3D images from rat large intestine tissue are clearly identified. Morphological details in the breast cancer tissue (human u-87 MG glioblastoma cells) are also provided by the visualisation of cell nuclei (Fig. 1) [11].

The capability of the nonlinear endoscopic probe can be further extended to have 3D functional imaging [12]. This can be achieved by the use of supercontinuum generation in nonlinear photonic fibre.

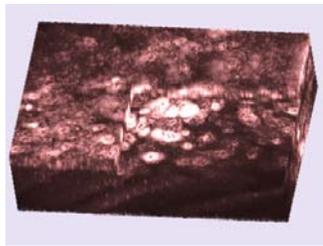


Fig. 1 Three-dimensional visualization of *in vitro* images of the human u-87 MG glioblastoma tissue.

Nonlinear optical micro-probe with a double-clad photonic fibre

Although the nonlinear optical endoscope that uses the double-clad photonic crystal can improve the signal strength significantly, its weak mechanical strength has limited the probe in endoscope applications. An alternative way of designing a high efficiency nonlinear optical endoscope is the adoption of a double-clad fibre [13] that has an improved mechanical property for fast scanning. In addition, the double-clad fibre does not reduce the overall system efficiency due to the large size of the inner clad region that can transmit efficiently the visible signal from a 3D sample.

Acknowledgements

The author thanks the Australian Research Council for its support and acknowledges the contributions from Ling Fu, Wei Tao, Dru Morrish and Hongchun Bao from Swinburne University and Huikai Xie and Lei Wu from University of Florida.

References

- [1] B. A. Flusberg, E. Cocker, W. Piyawattanametha, J. Jung, E. Cheung and M. Schnitzer, "Fiber-optic fluorescence imaging", *Nat. Methods* **2**, 941-950 (2005).
- [2] D. Bird and Min Gu, "Fibre-optic two-photon scanning fluorescence microscopy", *J. Microscopy* **208**, 35-48 (2002).
- [3] Ling Fu and Min Gu, "Fibre-optical nonlinear optical microscopy and endoscopy", *J. Microscopy* **226** 195-206 (2007).
- [4] D. Bird and Min Gu, "Compact two-photon fluorescence microscope using a single-mode optical fibre coupler", *Opt. Lett.* **27**, 1031-1033 (2002).
- [5] Damian Bird and Min Gu, "Two-photon fluorescence endoscopy with a micro-optic scanning head", *Opt. Lett.* **28**, 1552-1554 (2003).
- [6] Ling Fu, Xiaosong Gan and Min Gu, "Use of a single-mode fiber coupler for second harmonic generation microscopy", *Opt. Lett.* **30**, 385-388 (2005).
- [7] L. Fu, X. Gan and Min Gu, "Nonlinear optical microscopy based on double-clad photonic crystal fibers", *Opt. Express* **13**, 5528-5534 (2005).
- [8] Ling Fu, Xiaosong Gan and Min Gu, "Characterization of the GRIN lens-fiber spacing toward applications in two-photon fluorescence endoscopy", *Applied Optics*, **44**, 7270-7274 (2005).
- [9] L. Fu, A. Jain, H. Xie, C. Cranfield and Min Gu, "Nonlinear optical endoscopy

- based on a double-clad photonic crystal fiber and a MEMS mirror”, *Opt. Express* **14**, 1027-1032 (2006).
- [10] Ling Fu and Min Gu, „A double-clad photonic crystal fiber coupler for compact nonlinear optical microscopy imaging”, *Opt. Lett.* **31**, 1471-1473 (2006).
- [11] Ling Fu, Ankur Jain, Charles Cranfield, Huikai Xie and Min Gu, “Three-dimensional nonlinear optical endoscopy”, *J. Biomedical Opt.* **12**, 040501 (2007).
- [12] Ling Fu and Min Gu, “Polarization anisotropy in fiber-optic second harmonic generation microscopy”, *Opt. Express* **16**, 5000-5006 (2008).
- [13] Hongchun Bao, John Allen, Robert Pattie, Rod Vance, and Min Gu, “A fast handheld two-photon fluorescence micro-endoscope with a $475\mu\text{m}\times 475\mu\text{m}$ field of view for in vivo imaging”, *Opt. Lett* **33**, (2008), in press.