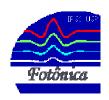
Fabrication and optical connection of 3D microstructures









outline

introduction

fs-micromachining

superhydrophobic surfaces production of nanoparticles in glass scaffolds for biological applications

fs-laser microfabrication

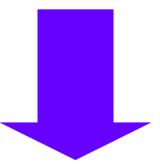
two-photon polymerization birefringent microstructures fluorescent microstructures connection with silica nanowires micro structres with ZnO nanowires micro- environment for bacterial growth

introduction

short pulse duration \rightarrow high intensity

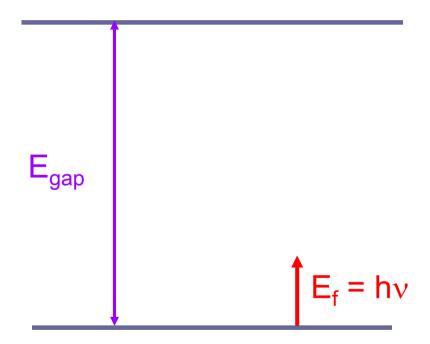
(even at low energy)

photon energy < bandgap

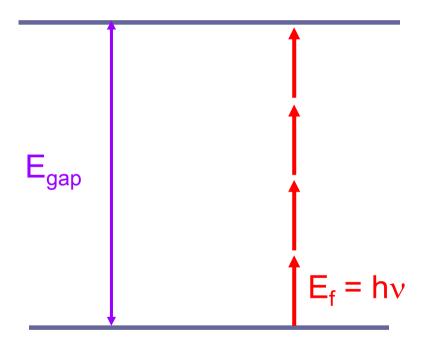


nonlinear interaction

nonlinear interaction



nonlinear interaction

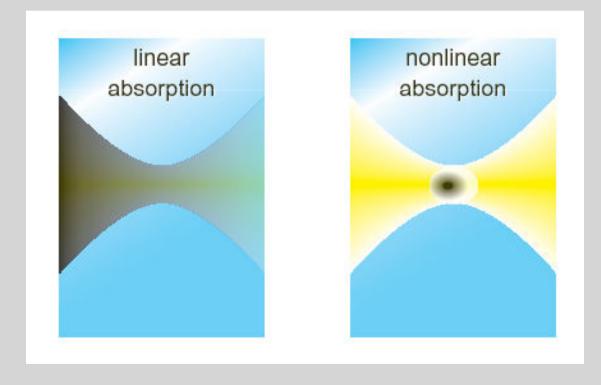


multiphoton absorption

Two-photon absorption

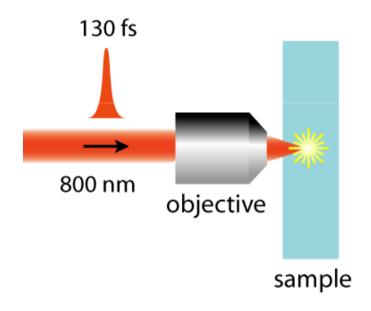
Nonlinear interaction provides spatial confinement of the excitation

fs-microfabrication



$$\alpha = \alpha_0 \qquad \alpha = \alpha_0 + \beta I$$

focus laser beam inside material



femtosecond pulses

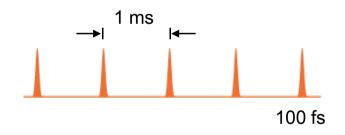
Ti:Sapphire lasers



$$1 \text{ fs} = 10^{-15} \text{ s}$$

Repetition rate

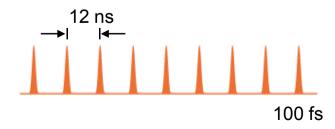
1 KHz



Energy

mJ

86 MHz

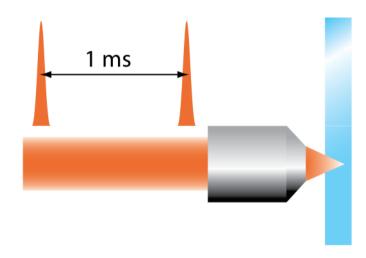


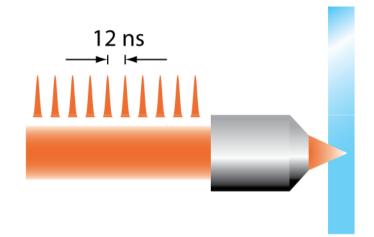
nJ

femtosecond pulses

amplified laser

oscillator





repetitive

cumulative

fs-micromachining

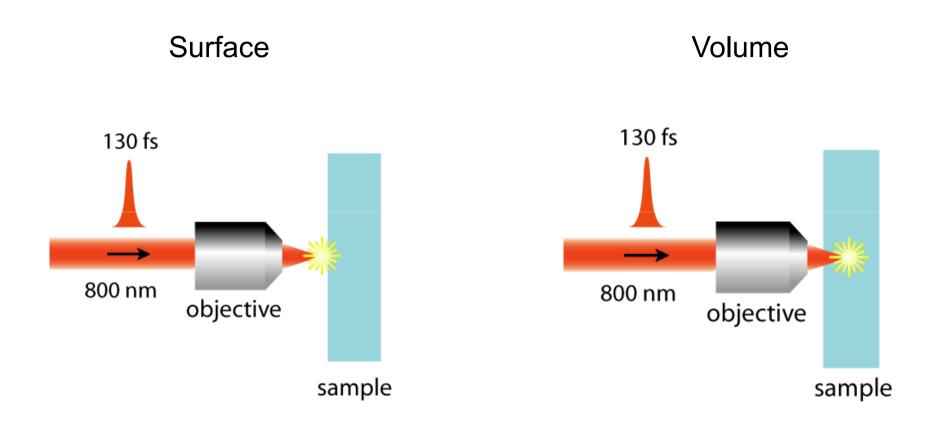
microfabrication can be controlled by

- objective NA
- number of pulses scanning speed
- pulse energy

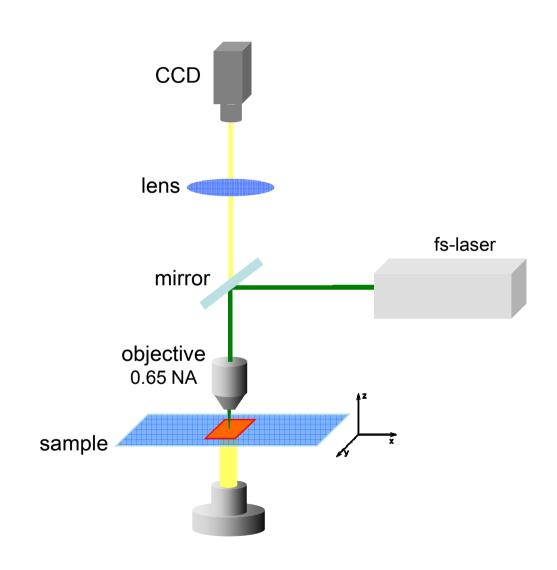
two main techniques

- fs-laser micromachining
- microfabrication via two-photon polymerization

fs-laser micromaching

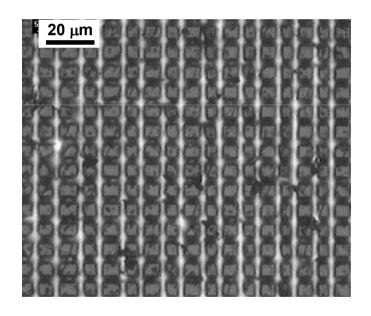


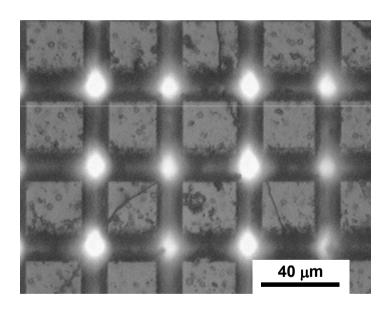
fs-laser microstructuring experimental setup



laser microfabrication: super hydrophobic surface

examples of fabricated surfaces





laser microfabrication: super hydrophobic surface

Superhydrophobic surfaces

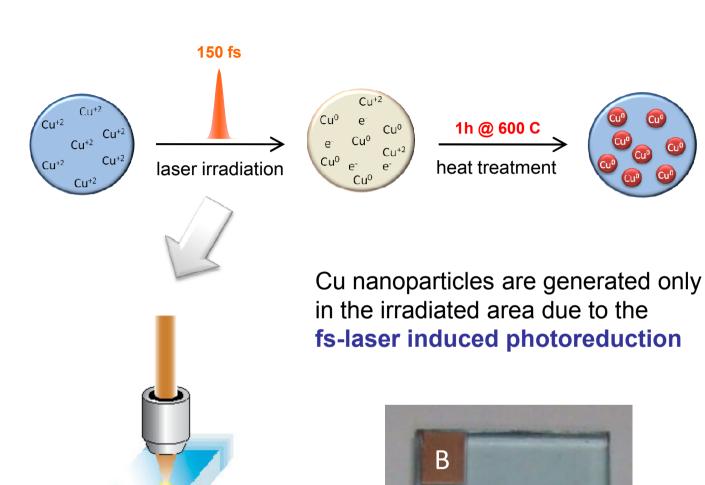




flat surface

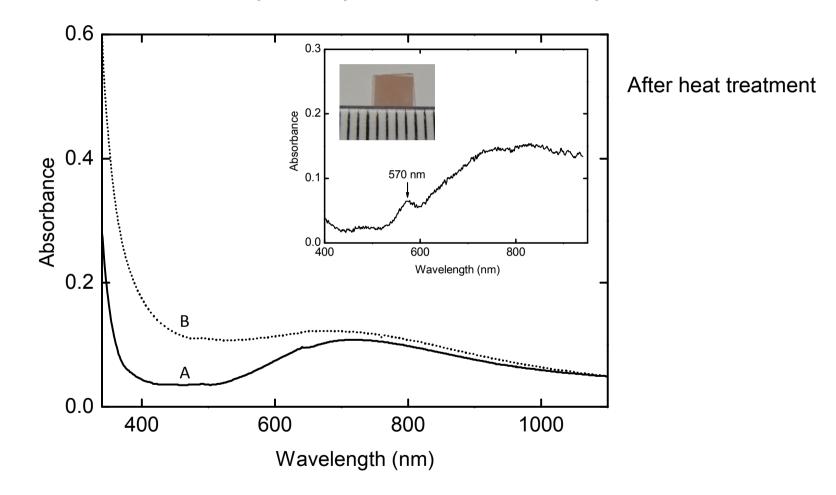
microstructured surface

Generation of Cu nanoparticles



Generation of Cu nanoparticles

Absorption spectrum of the sample



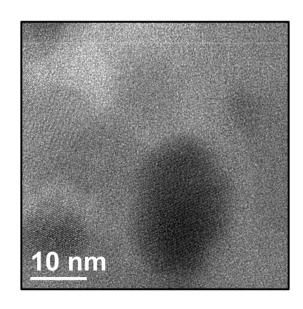
A: as-prepared sample

B: sample after fs-laser irradiation

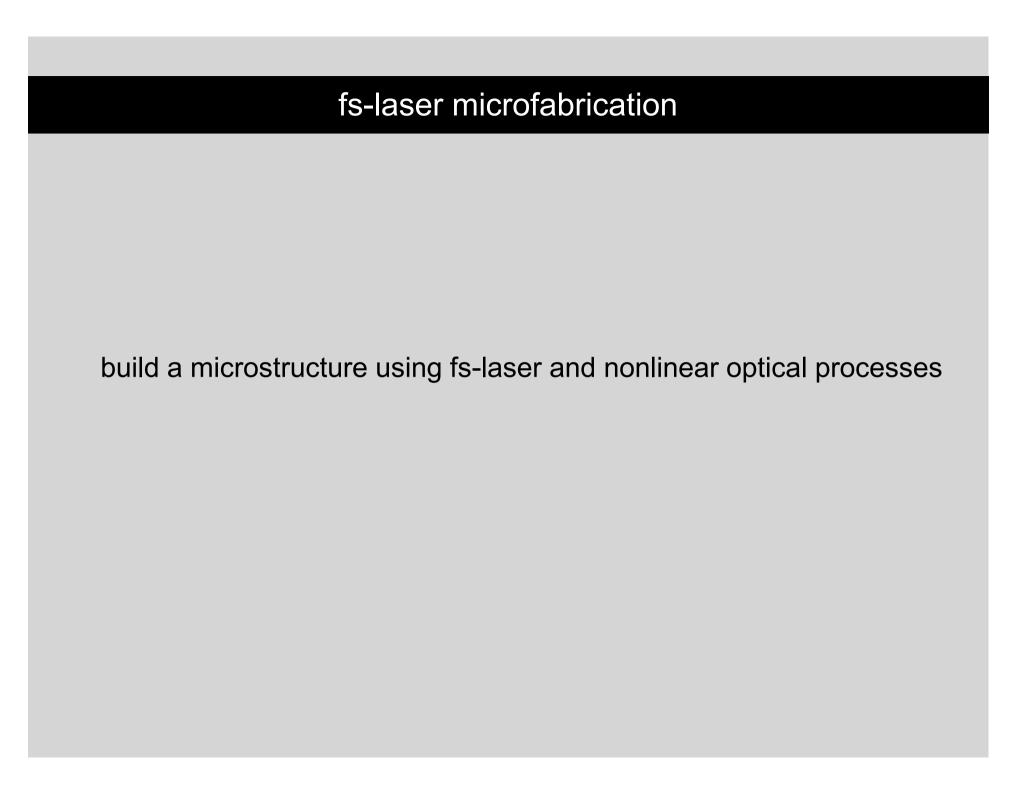
Generation of Cu nanoparticles

Optical microscopy showing the regions in the bulk where Cu nanoparticles were produced





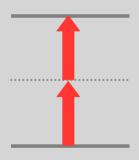
TEM image of the nanoparticles



Monomer + *Photoinitiator* → *Polymer*



Photoinitiator is excited by *two-photon absorption*

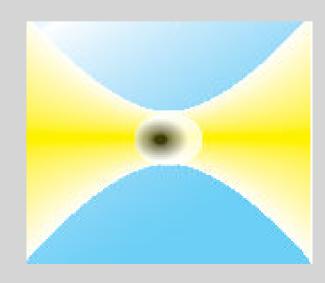


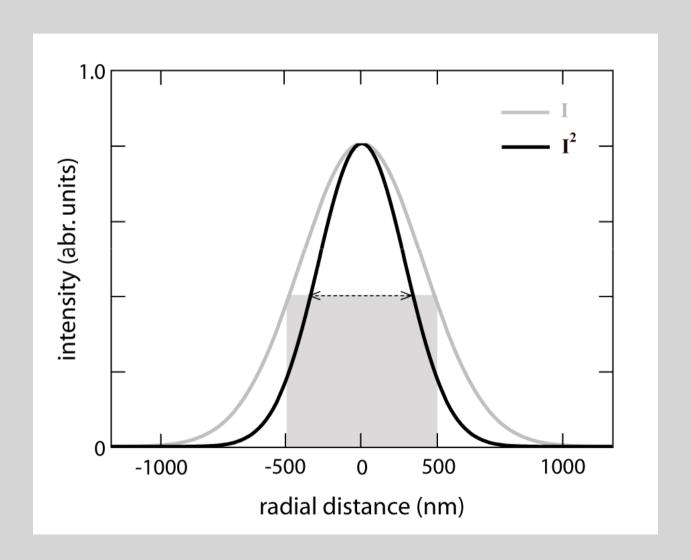
$$R_{2PA} \propto I^2$$

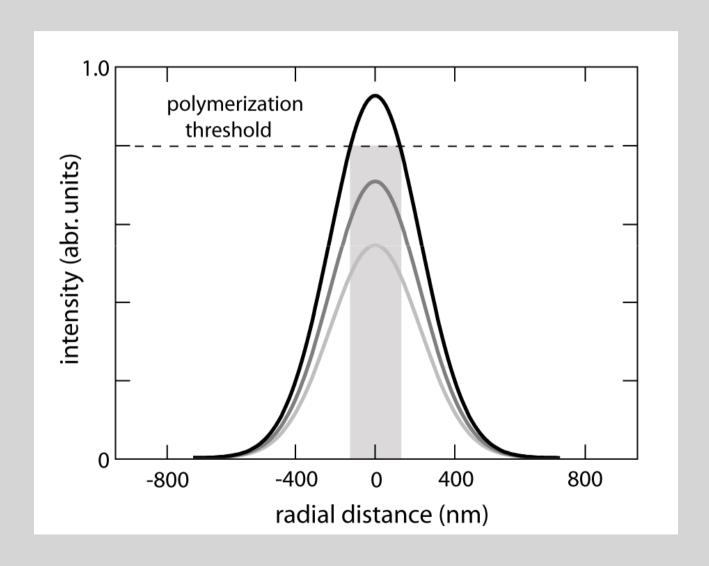
The polymerization is confined to the focal volume.



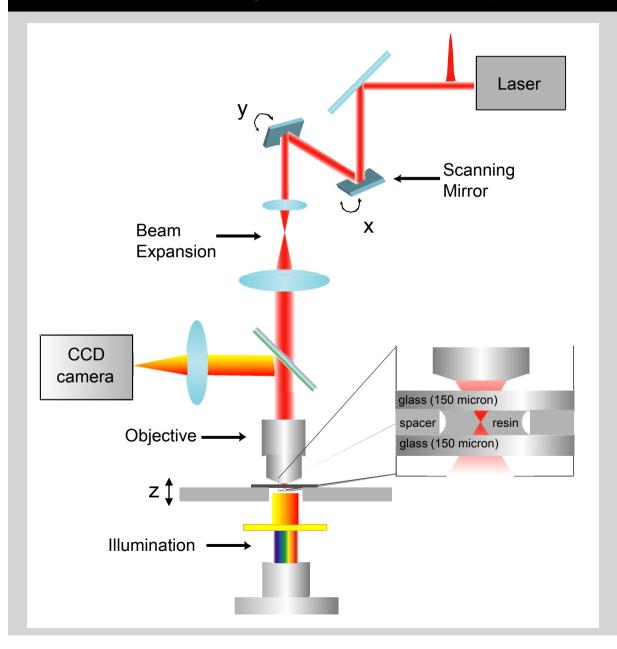
High spatial resolution







Two-photon polymerization setup



Ti:sapphire laser oscillator

- 130 fs
- 800 nm
- 76 MHz
- 20 mW

Objective

40 x 0.65 NA

Resin preparation

Monomers

Monomer A

$$CH_{2} \leftarrow CH_{2} - CH_{2} - CH_{2} - CH = CH_{2}$$

$$CH_{3} - CH_{2} \quad CH_{2} \leftarrow CH_{2} - CH_{2} - CH_{2} - CH = CH_{2}$$

$$CH_{3} - CH_{2} \quad CH_{2} \leftarrow CH_{2} - CH_{2} - CH_{2} - CH = CH_{2}$$

reduces the shrinkage upon polymerization

Monomer B

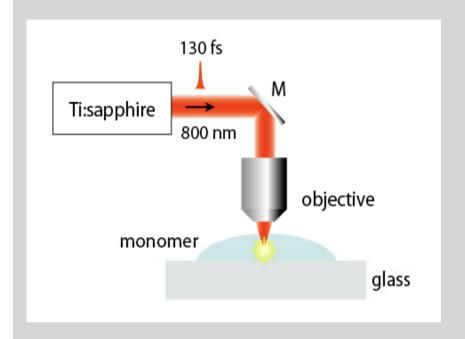
gives hardness to the polymeric structure

Photoinitiator

Lucirin TPO-L

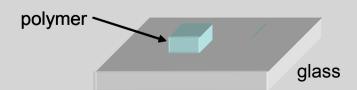
$$H_3C$$
 CH_3
 CH_3
 CH_3
 CH_3
 CH_3
 CH_3

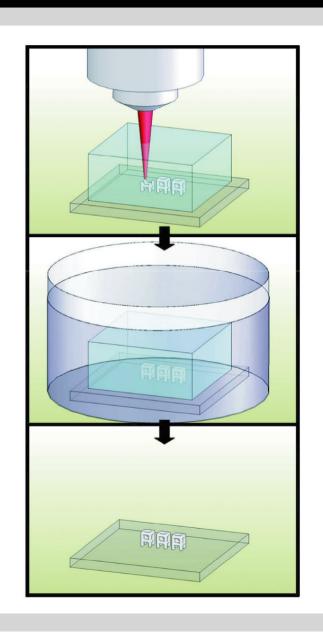
Appl. Phys. A, 90, 633-636 (2008)





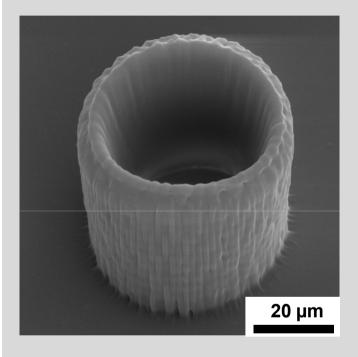
30 μ m x 30 μ m x 12 μ m cube

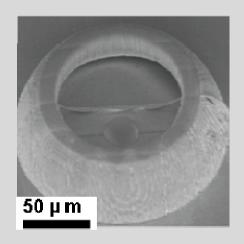


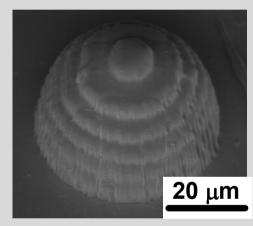


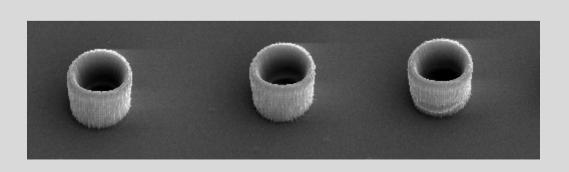
After the fabrication, the sample is immersed in ethanol to wash away any unsolidified resin and then dried

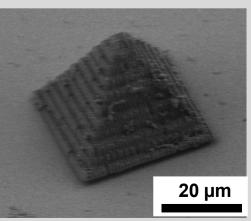
Microstructures fabricated by two-photon polymerization



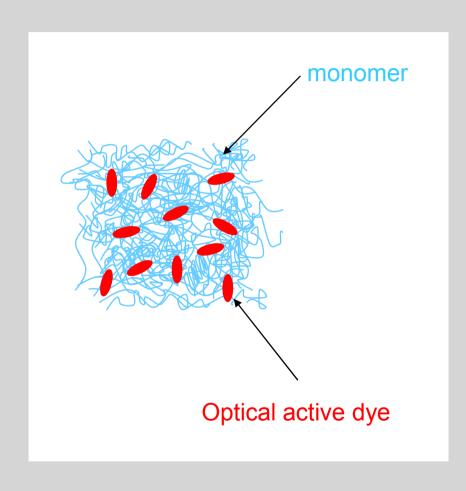


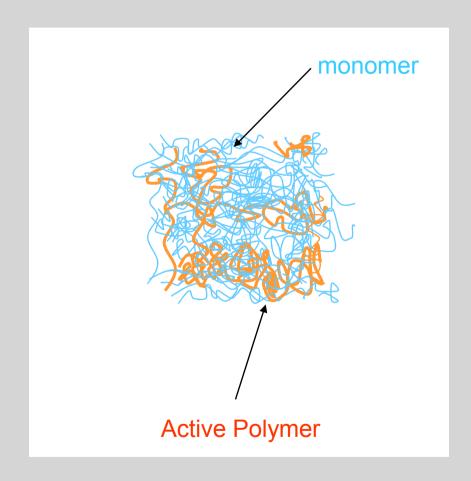






Microstructures containing active compounds





Applications of two-photon polymerization

Optics and Photonics

Doping microstructures with organic molecules and metals

fluorescence birefringence conductivity

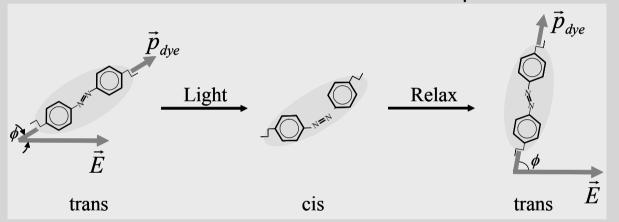
Bio-applications

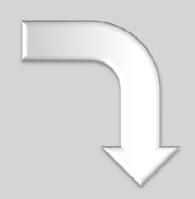
Fabrication using bio-compatible resins to biological applications

tissue engineering scaffolds fabrication of microneedle cell study

microstructures for optical storage

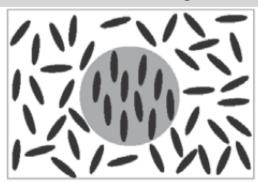
orientation mechanism in azochromophores



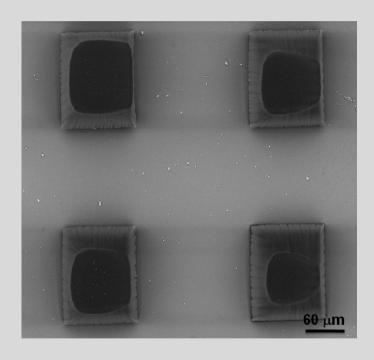


after writing

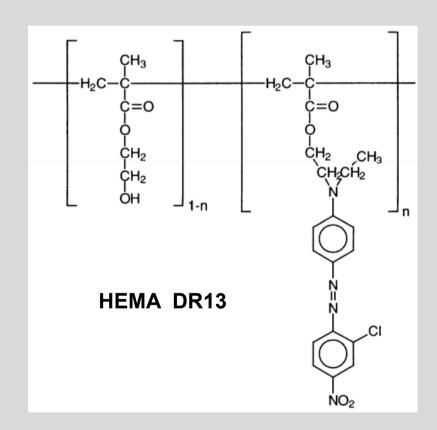
optically induced **birefringence**



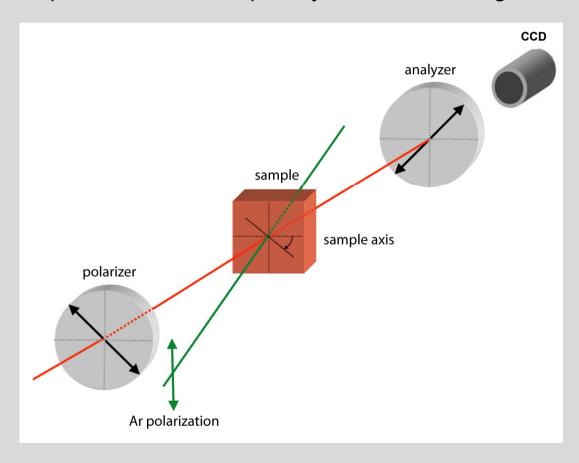
Resin composition: acrylate resin + HEMA DR13



microstructure fabricated by twophoton polymerization containing HEMA DR13



setup to measure the optically induced birefringence



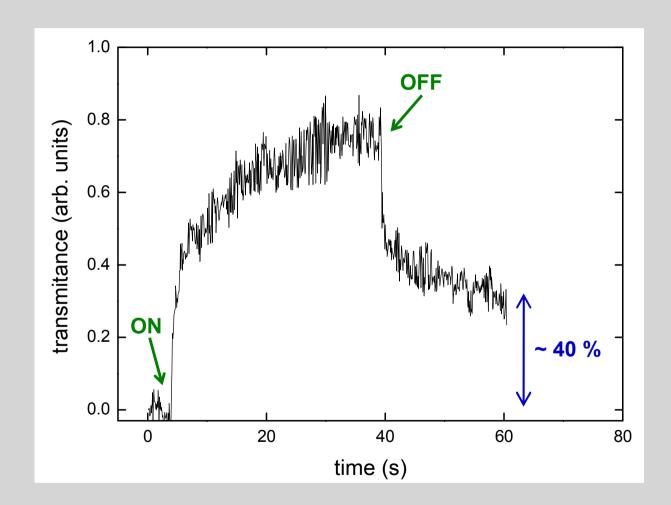
WRITING

Pump beam: Argon ion at 514 nm

READING

Probe beam: HeNe at 632.8 nm

Result demonstrating the induction of birefringence in the microstructure



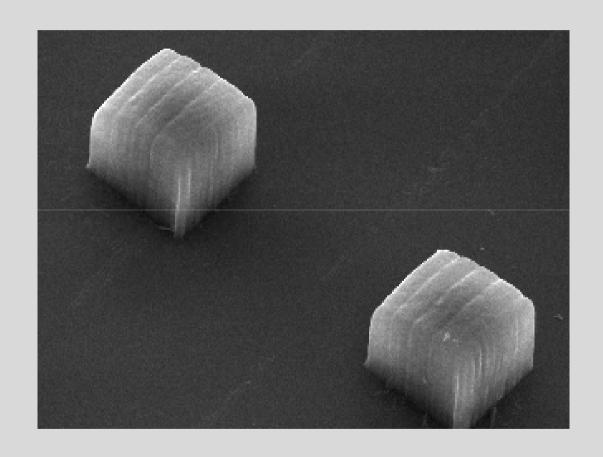
Microstructures containing Rhodamine

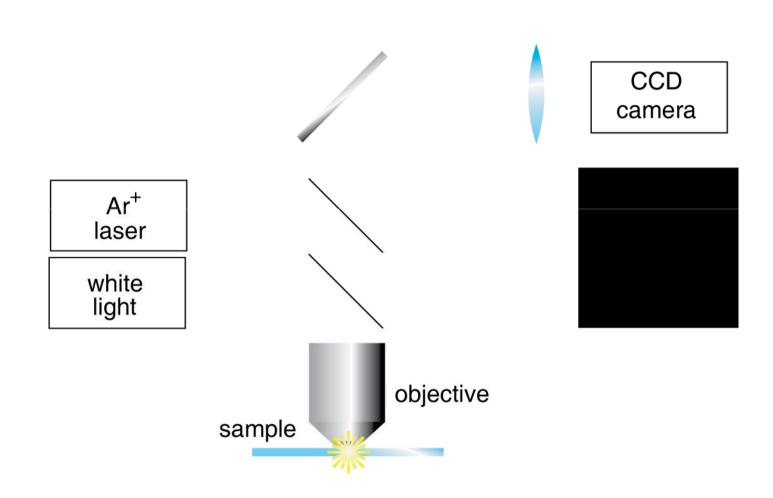
Rhodamine 6G

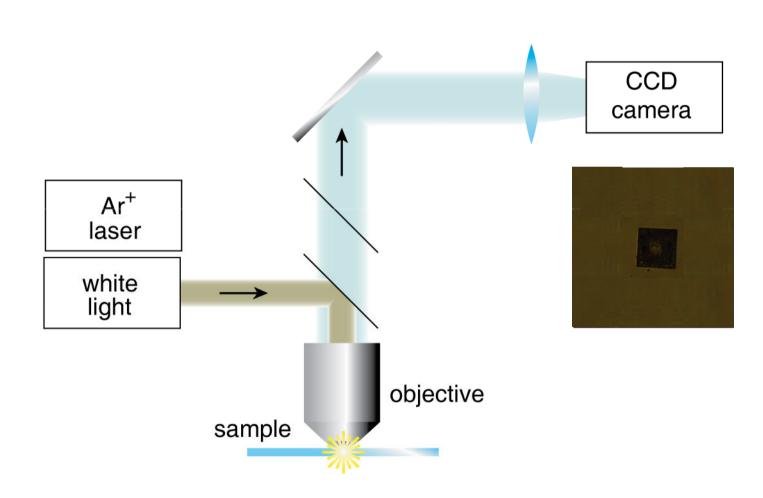
$$H_3C$$
 O
 O
 CH_3
 O
 CH_3
 O
 CH_3

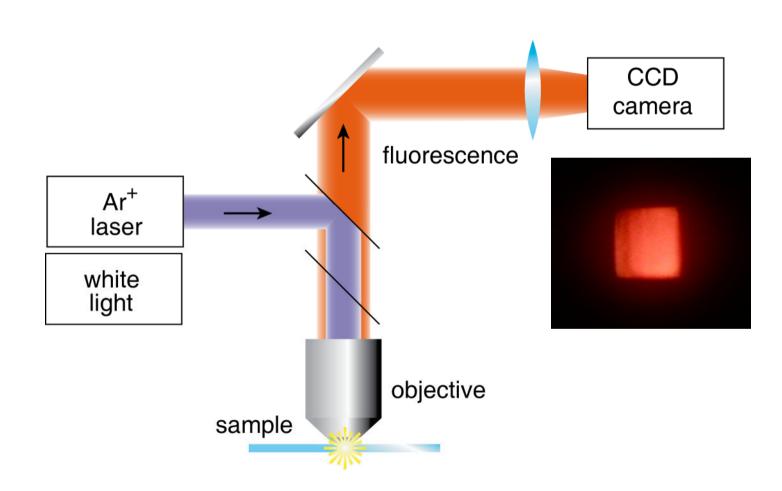
- High luminescence
- Used as dye laser gain medium

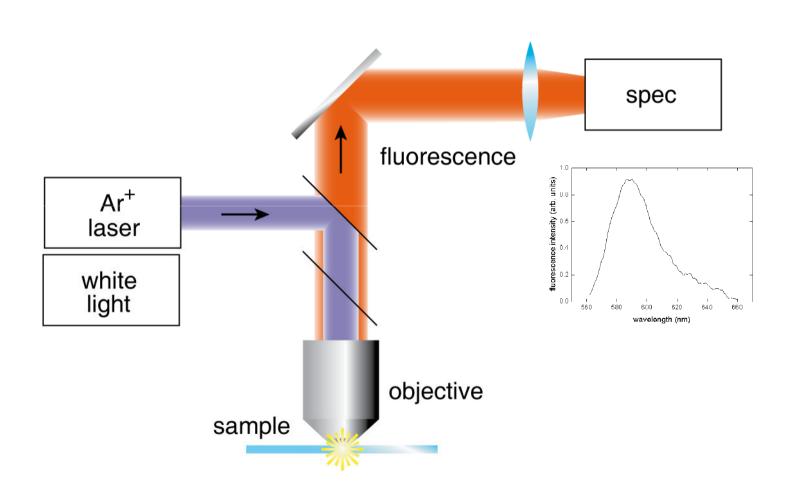
Microstructure containing Rhodamine

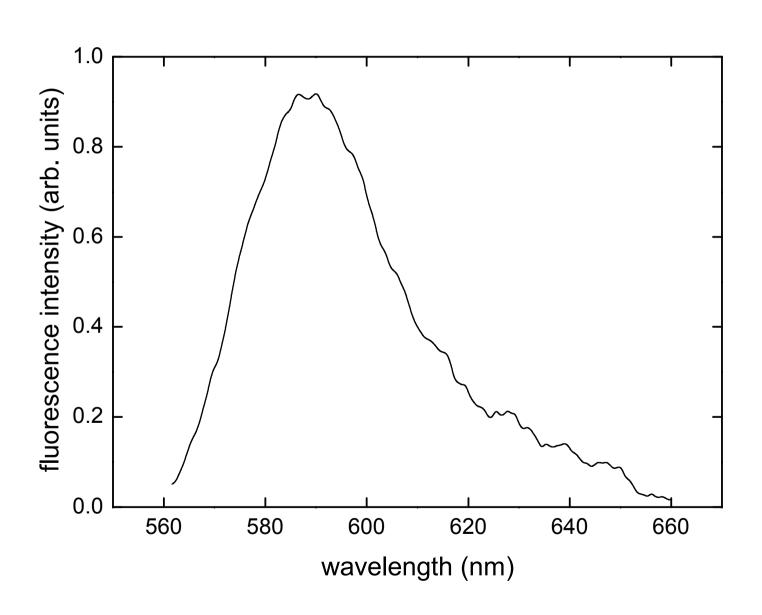




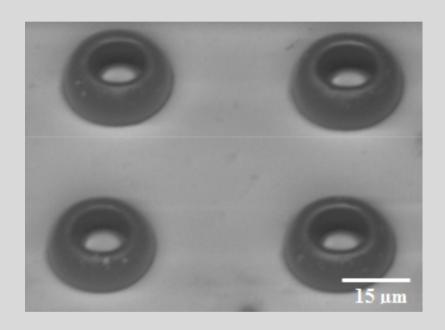




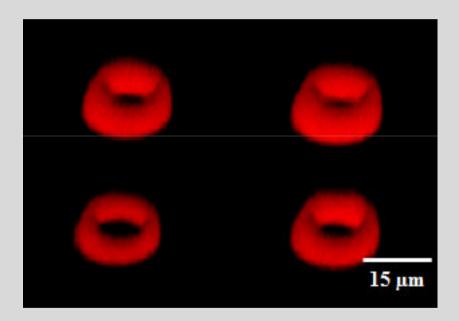




fabrication of array of doped microstructures

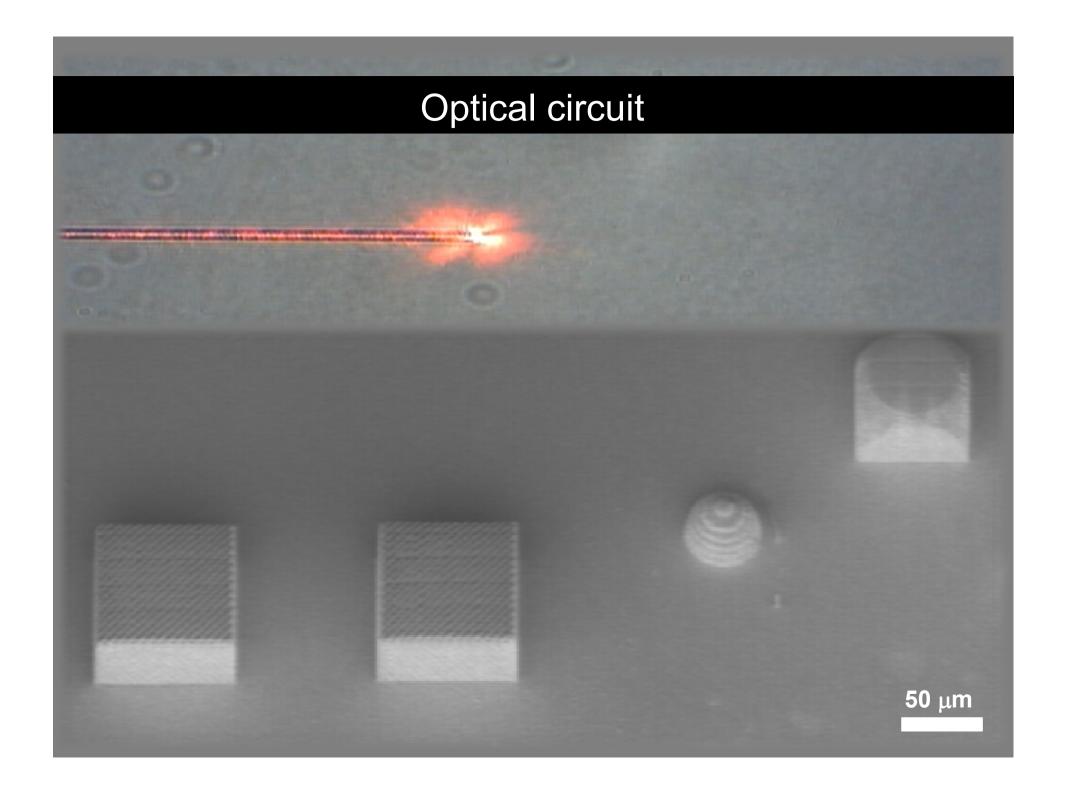


SEM



Confocal microscopy

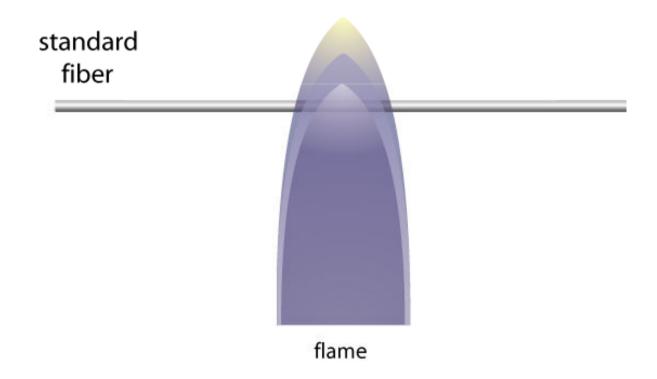
Optical circuit



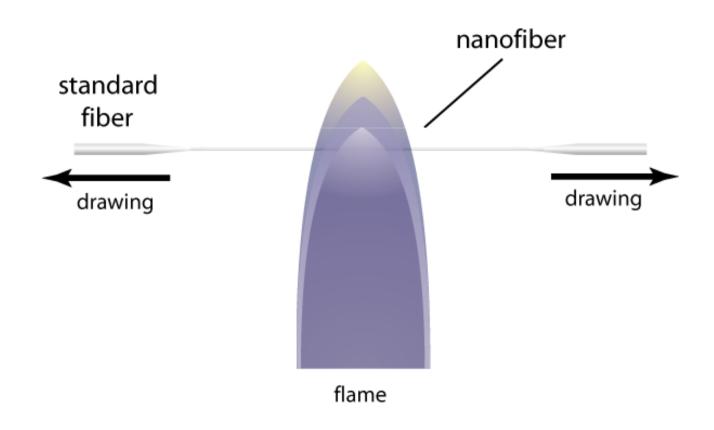
nanowires fabrication process

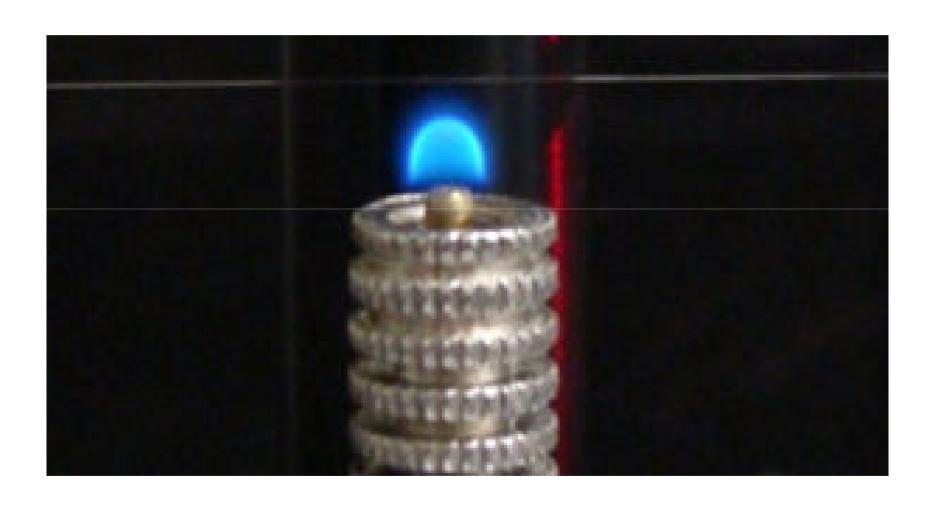
standard fiber

nanowires fabrication process

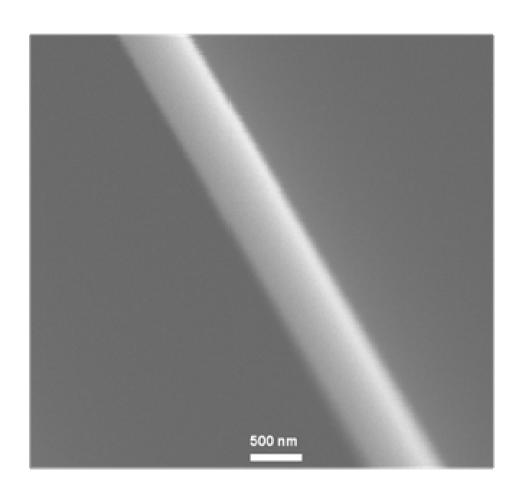


nanowires fabrication process





Silica nanowires 70 μm 1 μm

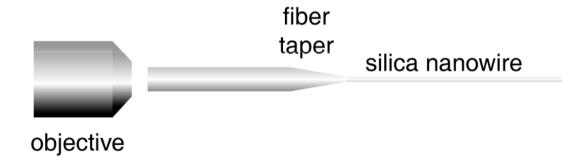


coupling light into nanowires

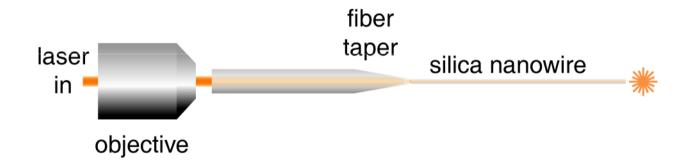
fiber taper

silica nanowire

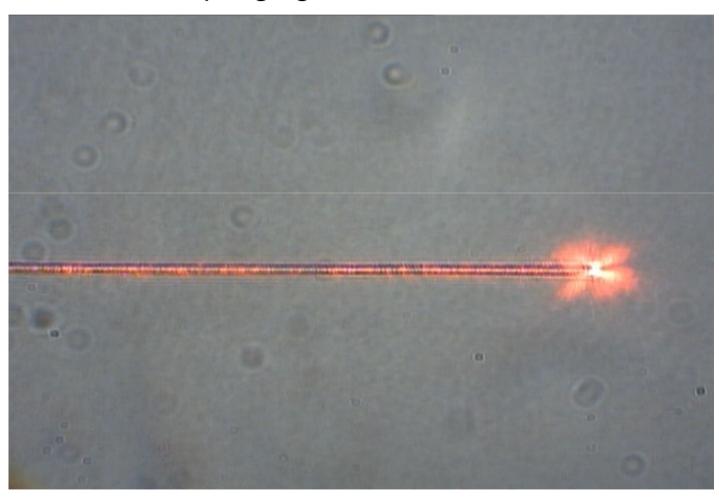
coupling light into nanowires

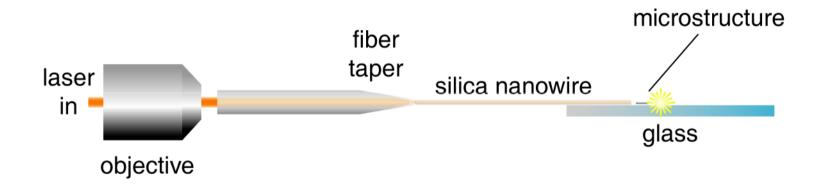


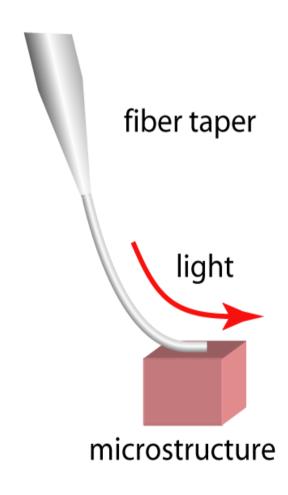
coupling light into nanowires



coupling light into nanowires

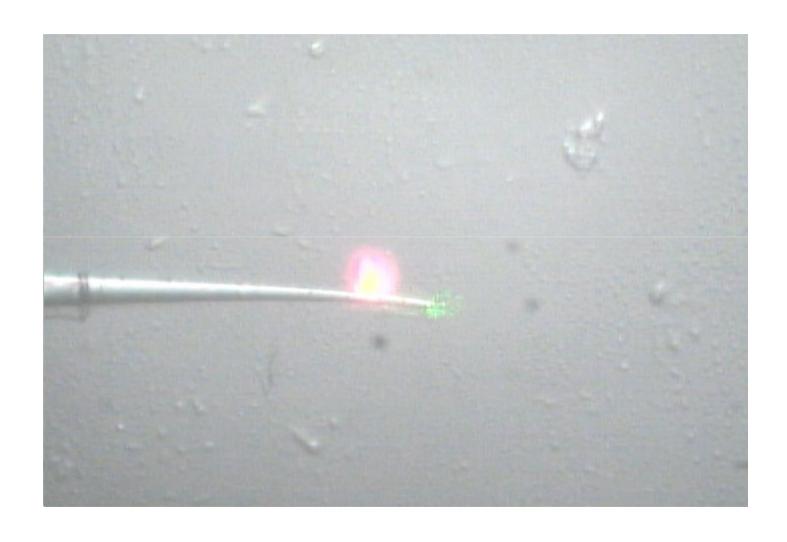


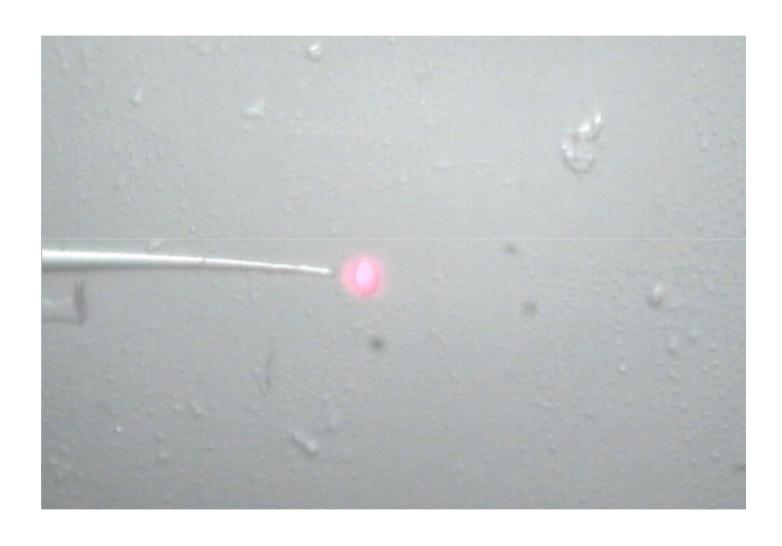


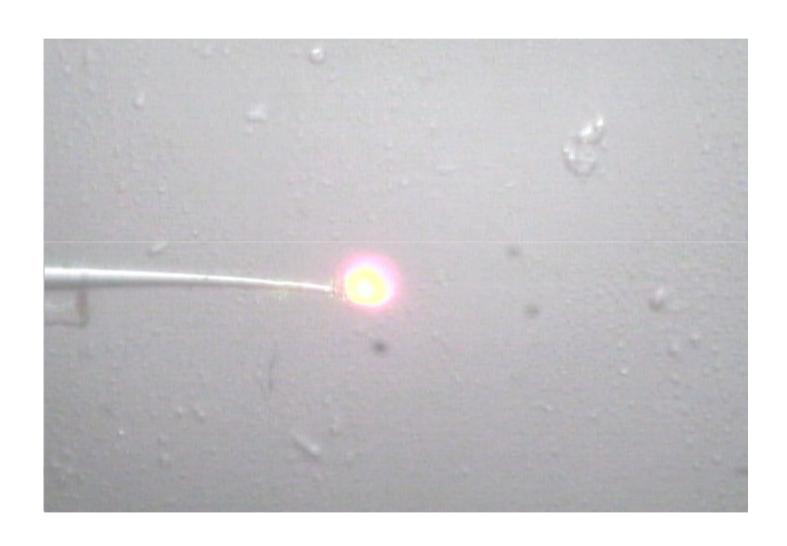








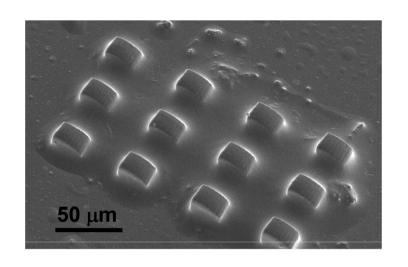


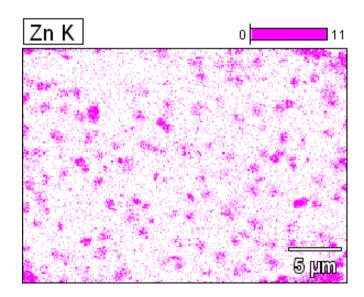


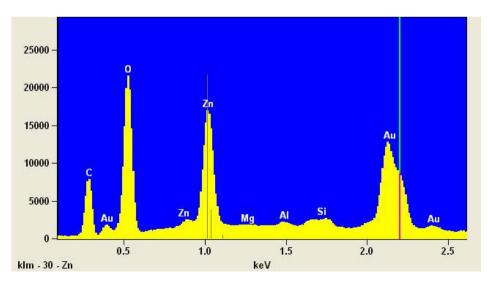


Microstructures with ZnO nanowires

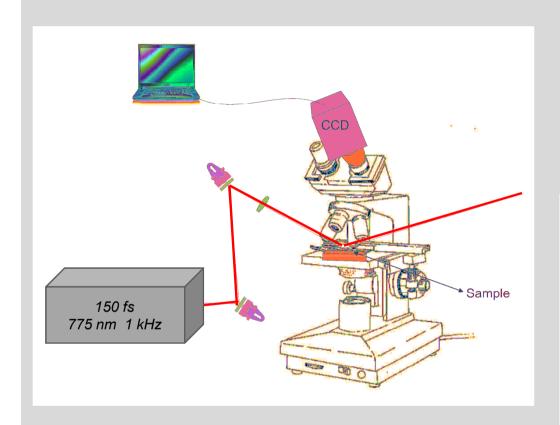
Microstructures with ZnO nanowires

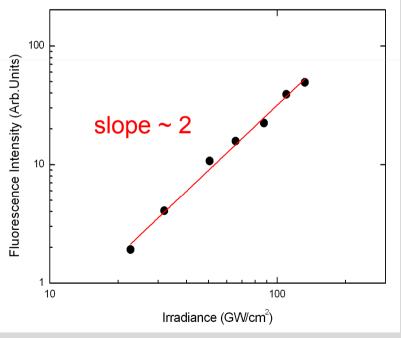






Microstructures with ZnO nanowires

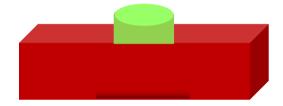




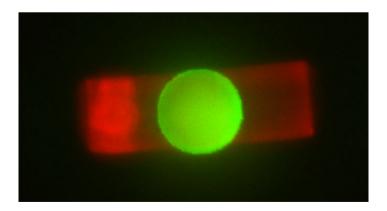
Guiding bacterial growth in a micro-environment microfabrication of multidoped microstructures

Guiding bacterial growth in a micro-environment

to study bacterial growth it was needed to develop double doped microstructures

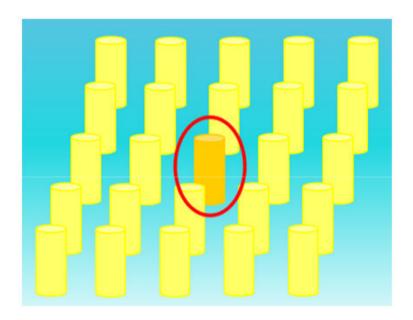


microstructure containing Fluorescein and Rhodamine



Guiding bacterial growth in a micro-environment

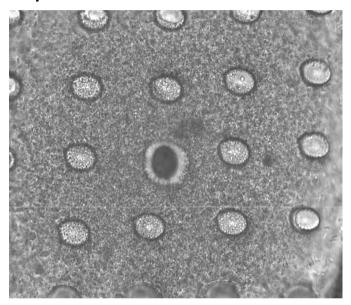
Study the development of E. coli in micro-environments:



micro-environment in which the central structure contains antibiotic.

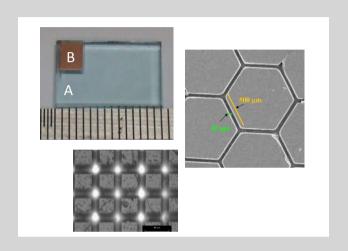
Guiding bacterial growth in a micro-environment

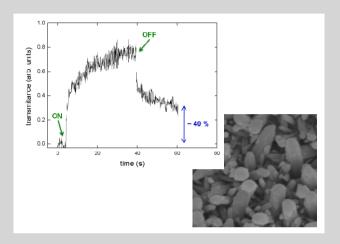
Study the development of E. coli in micro-environments:

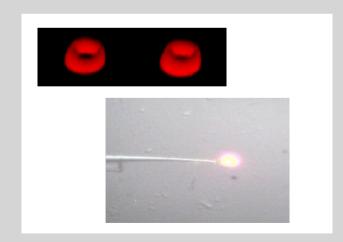


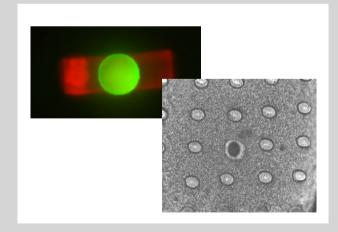
after 3 hours, we observed that a small region around the doped structure does not show bacterial growth.

Summary









Acknowledgments

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Gustavo F. B. Almeida

Nathalia Tomazio

FAPESP

CAPES

CNPq

www.fotonica.ifsc.usp.br

