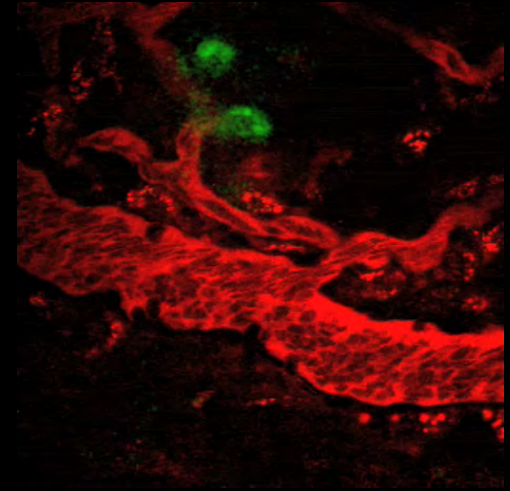
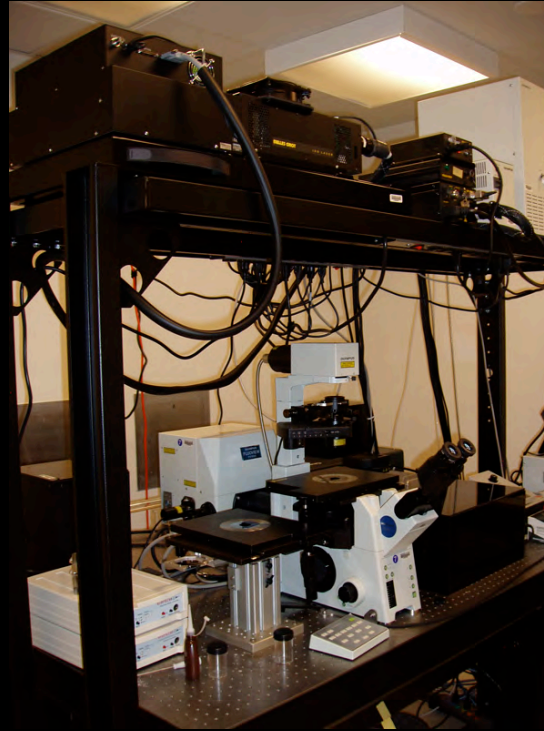
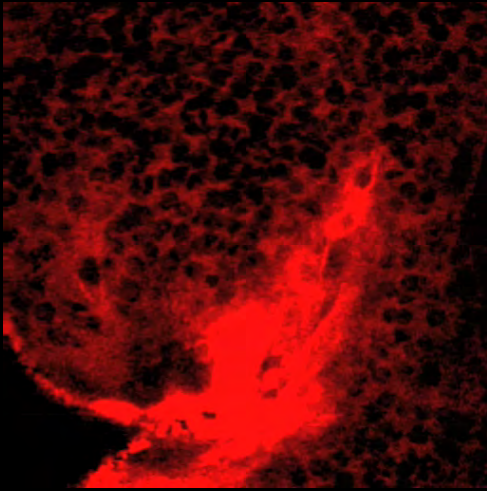


# Building Your Own 2-Photon Microscope: Challenges, Advantages and Limitations



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**Oral and Pharyngeal Cancer Branch**  
**NIDCR-NIH**



# Building Your Own 2-Photon Microscope: Challenges, Advantages and Limitations

## How did we manage to build a 2-photon microscope?

12% Ethanol (Preferably Italian)



Caffeine (Strictly Italian Espresso)



Being a pain in the neck.....

# Building Your Own 2-Photon Microscope

Turn Key System

Why?

Build your own

More expensive

~~\$400K-500K~~

\$150K

\$150K Laser

1) Budget

1) Initial expenses

2) Maintenance

2) Flexibility

3) Upgrades

Buy Confocal microscope



Convert to a 2-photon

First step (first major decision)

Which Platform?

Thorlabs  
Labvision

Olympus

1) We copied the system built here

2) Flexibility

3) Support

# Upright vs. Inverted

Flexibility:

- 1) 2-photon
- 2) Confocal

- 1) Intravital imaging
- 2) Live Cell imaging

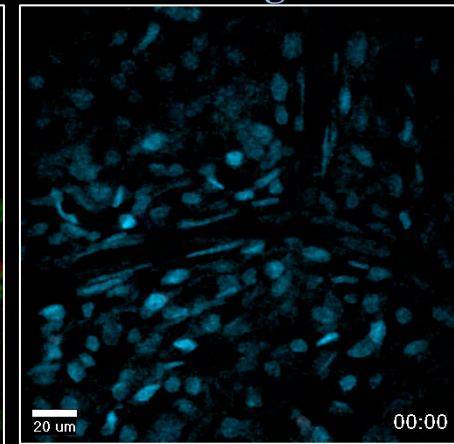
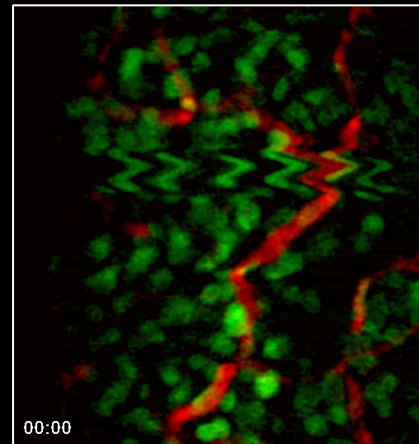
- 1) All the organs

## Motion artifacts

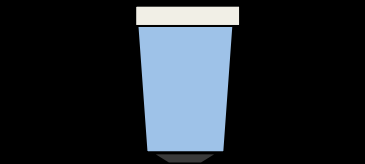
**“Custom made holding device specifically designed for the organ of interest”**

w/o holding device

with holding device

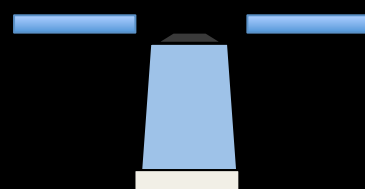


Upright



All the organs

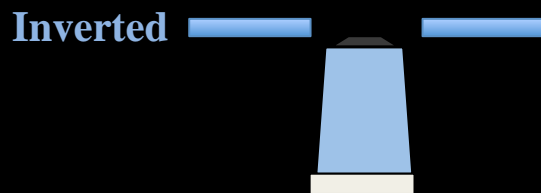
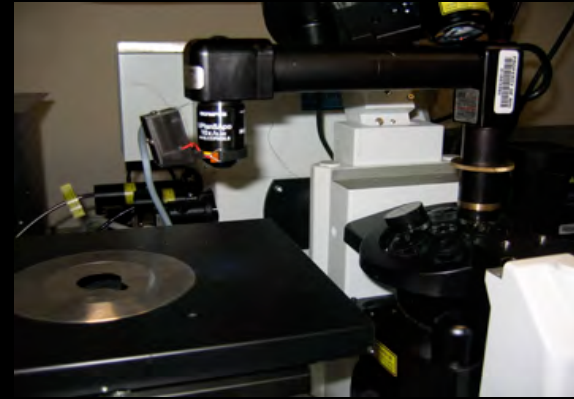
Inverted



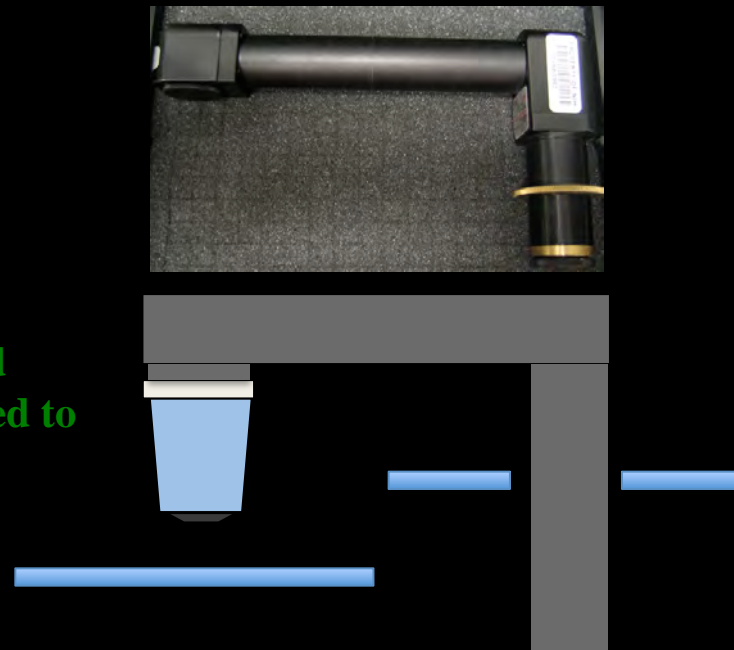
All the organs but the brain  
Live Cell Imaging

**“Positioning and securing the organ to the coverslip”**

# Upright vs. Inverted

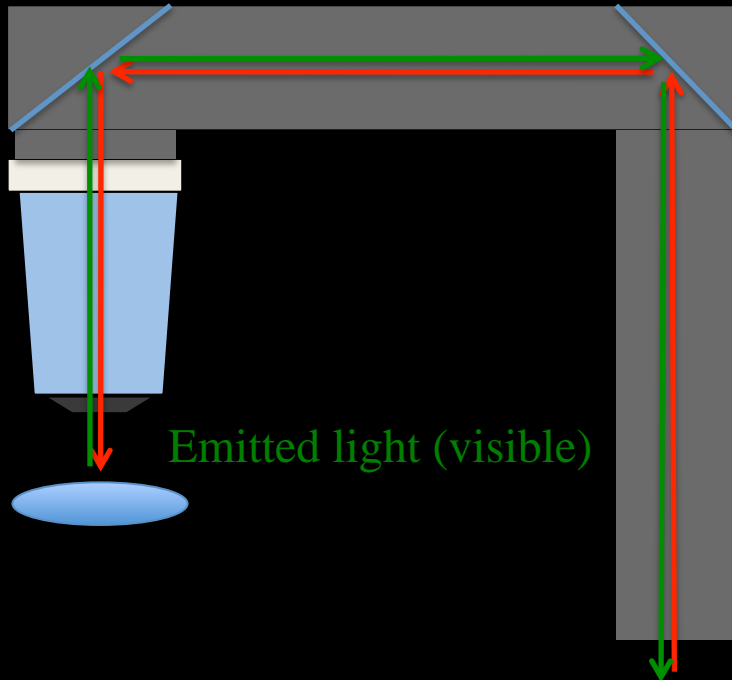


Inverted  
converted to  
upright



All the organs but the brain  
Live Cell Imaging

# Upright vs. Inverted



Excitation beam (IR)

- 1) Optimized for visible light
- 2) Increase the light path
  - 1) Model available with PMT on top
- 3) Loss of power (5-10%)
- 4) No effects on laser pulse width
- 5) Requires extra stage
- 6) Head can be rotated
- 7) Adaptors for lenses



# Upright vs. Inverted



Upright



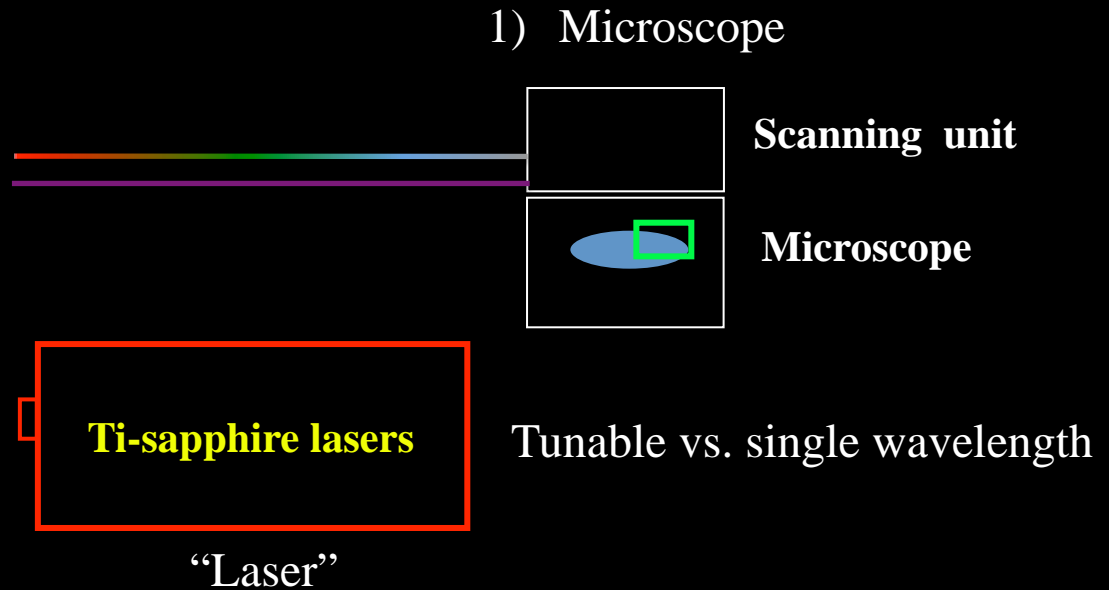
All the organs

Upright  
converted to  
inverted



# Laser

Laser combiner **488 nm**  
**561 nm**  
**633 nm**  
UV laser **405 nm**



**High power lasers (3-4 W)**  
**Repetition rates: 80-100 Hz**

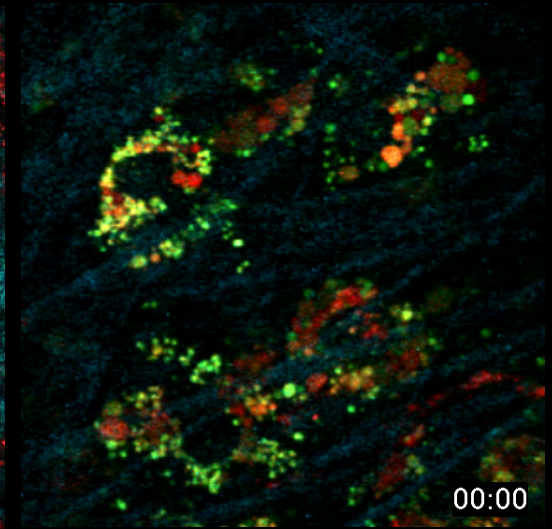
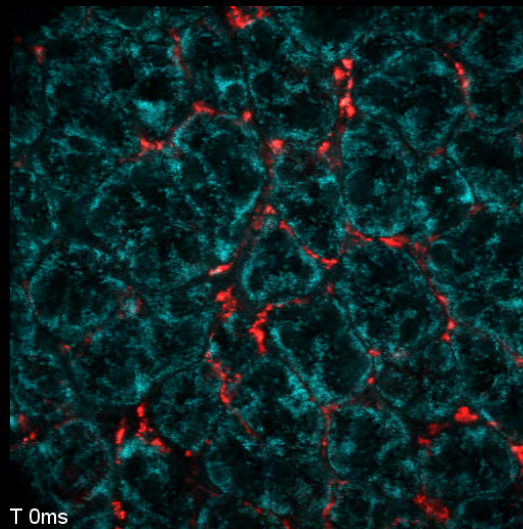
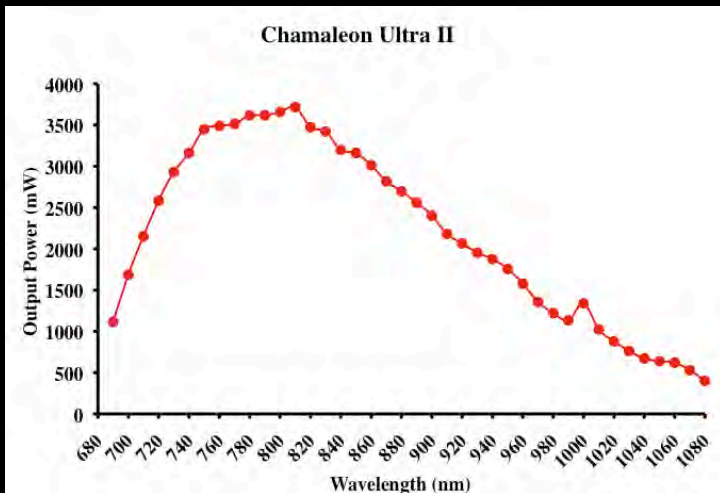
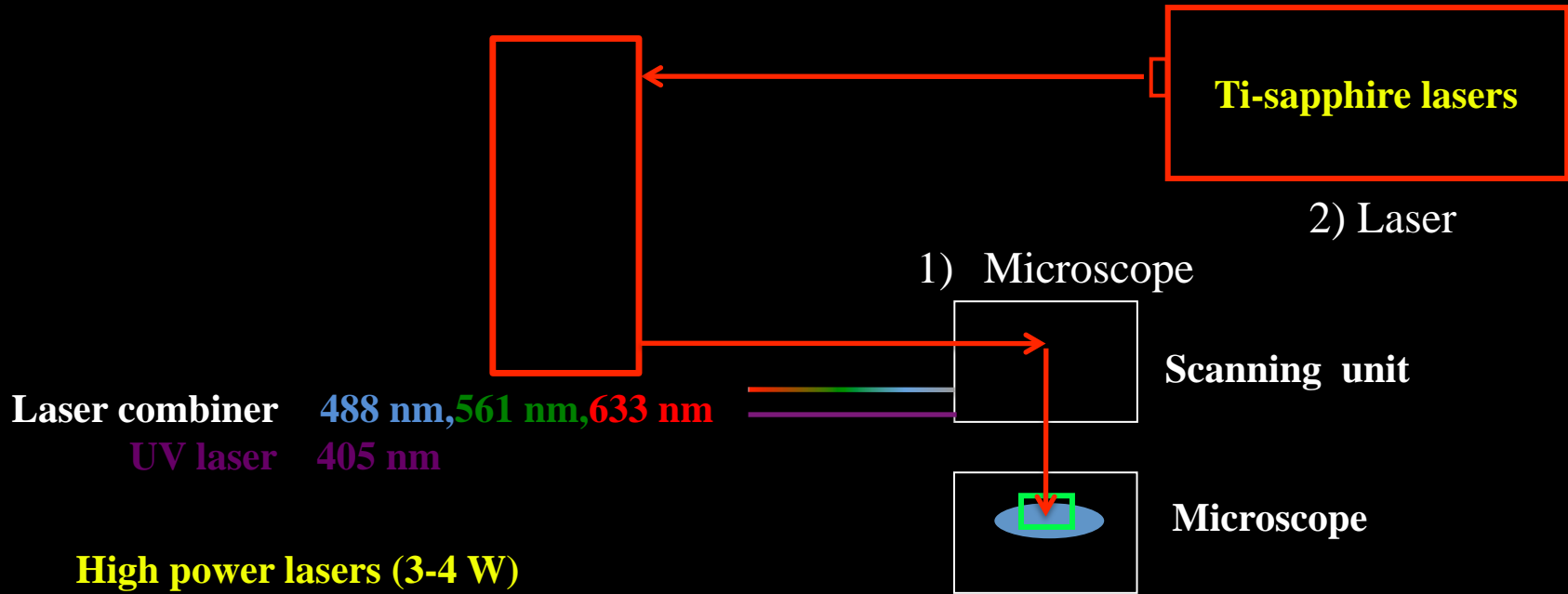
**Pulses: 100-150 fs**

**Beam diameter: 1.2 +/- 0.2 mm**

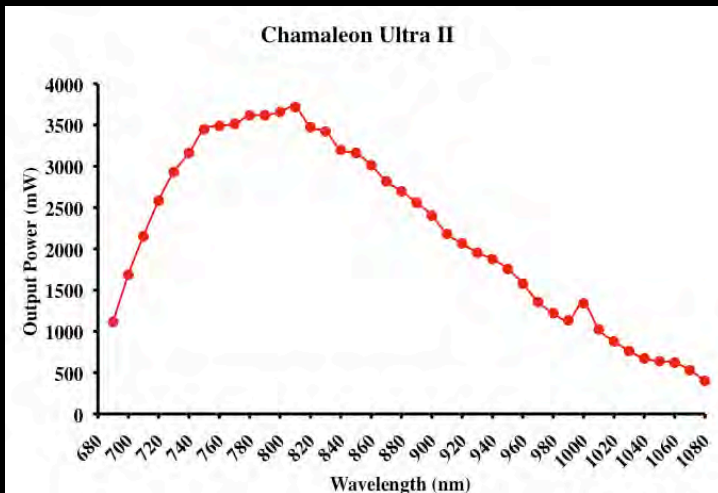
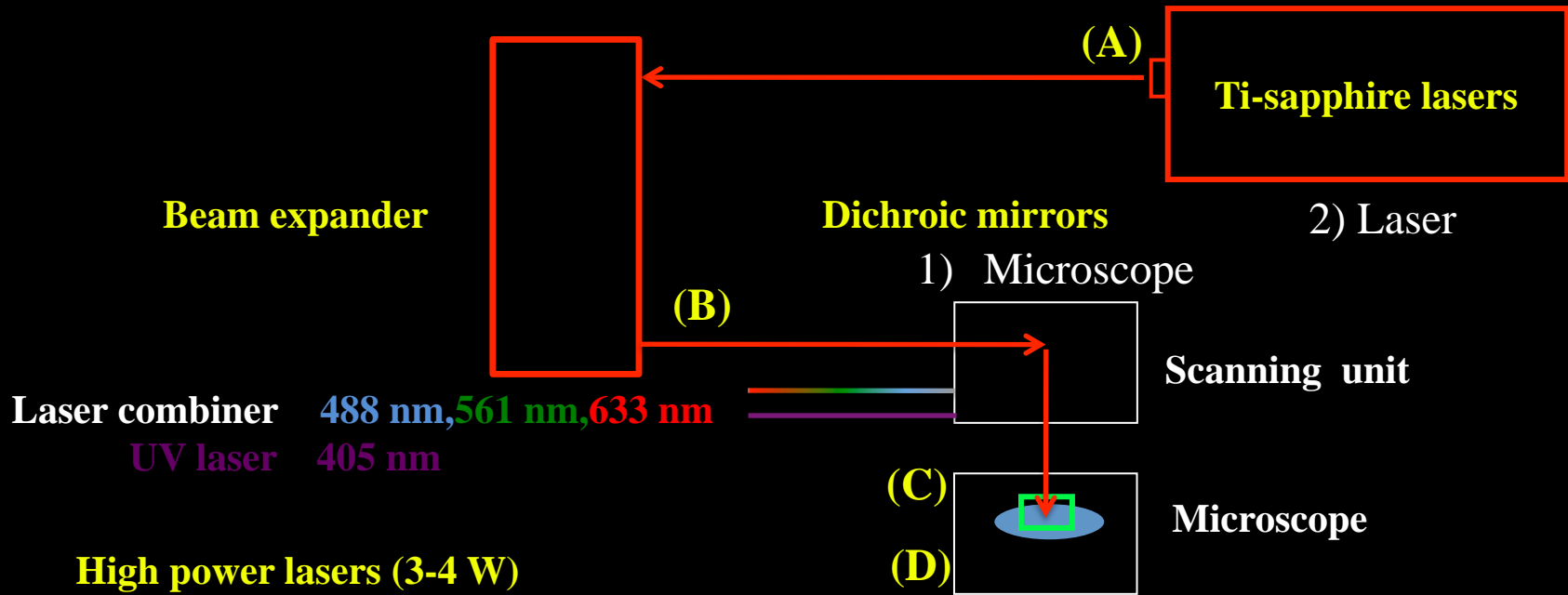
**Tunable: 680-1080 nm**



# Laser output power

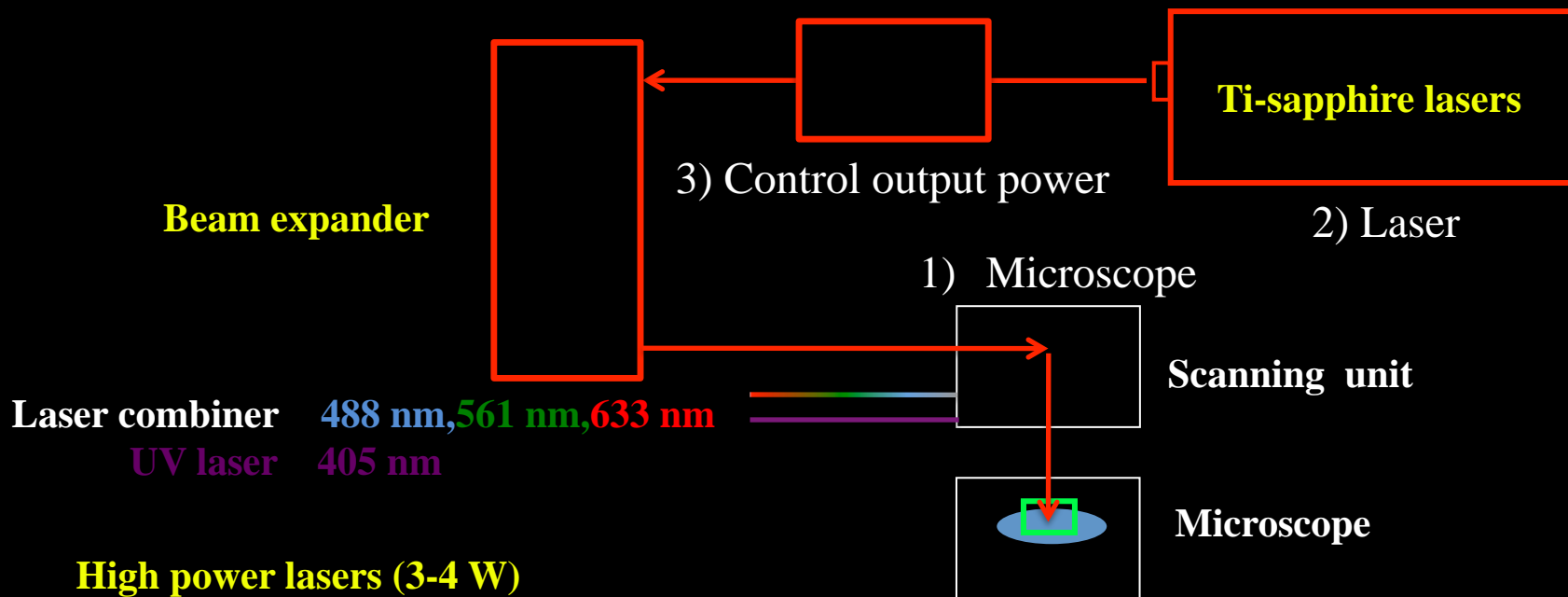


# Loss of power throughout the optics



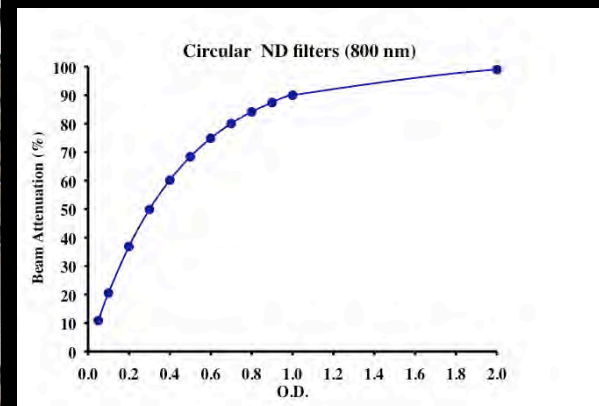
- (A) 800 nm – 3520 mW (100%)**
- (B) 3280 mW (93%)**
- (C) 864 mW (25%) – S.H. cuts 20%**
- (D) 340 mW (10%) – 60x N.A 1.2**

# Control the power at the specimen

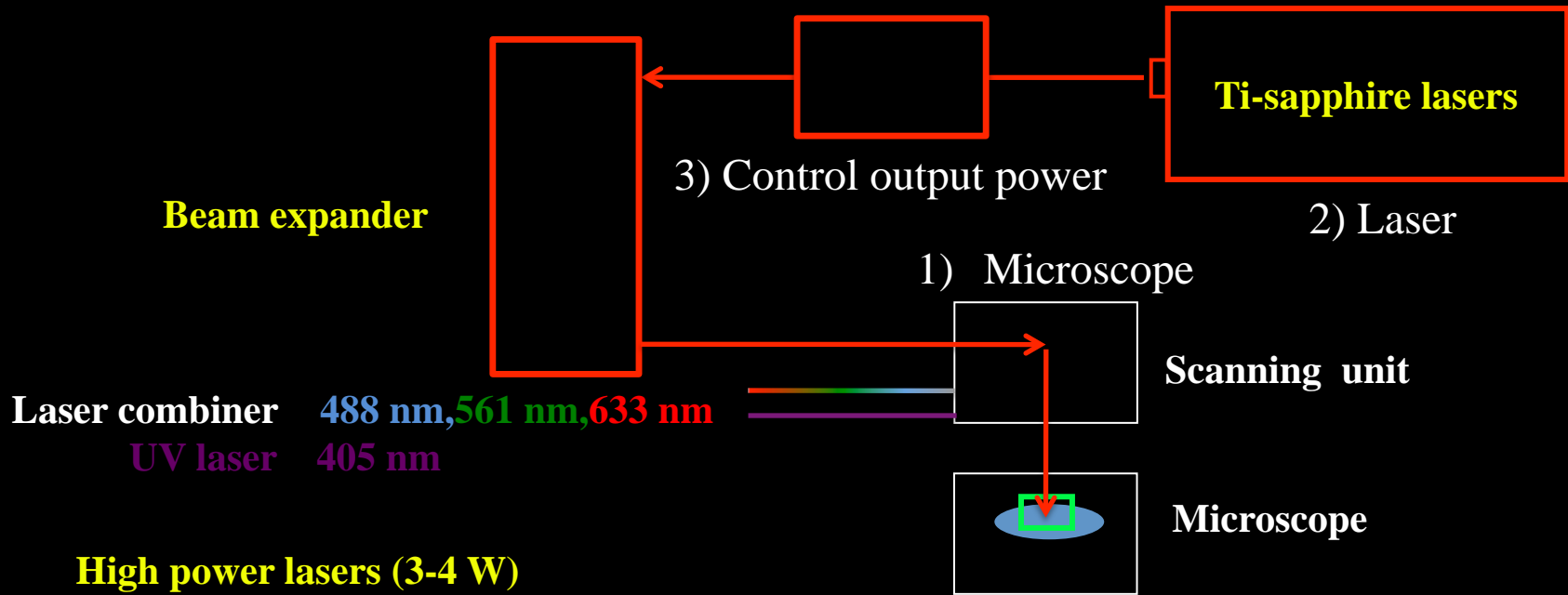


- 1) ND filters
  - a) Single
  - b) Carousel with multiple filters (8-10)
    - a) Manually or software controlled

- 2) ND continuous filter wheel



# Control the power at the specimen



1) ND filters

2) ND continuous filter wheel

3) AOM (Acousto-optic modulator)

4) EOM (Electro-optic modulator)

a) Easy integration with the software

b) Size of the beam matching the aperture of the AOM

c) Significant pulse broadening (up to 600 fs)

a) Need for a pre-chirping system

d) Deflection of the beam

a) Not practical if different wavelengths are needed

b) Need for an automatic realignment set up (expensive)

# Broadening of the pulse width

(a)



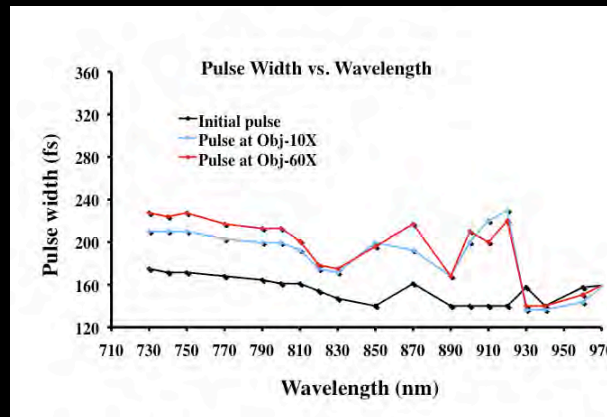
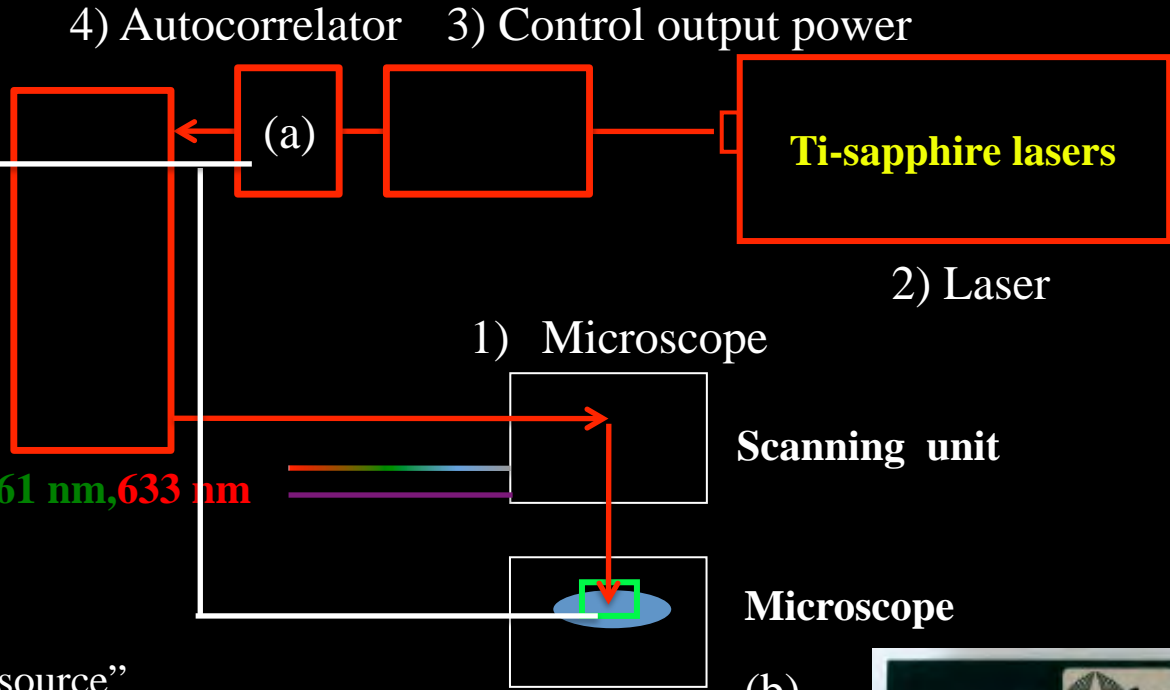
**Beam expander**

Laser combiner 488 nm, 561 nm, 633 nm  
 UV laser 405 nm

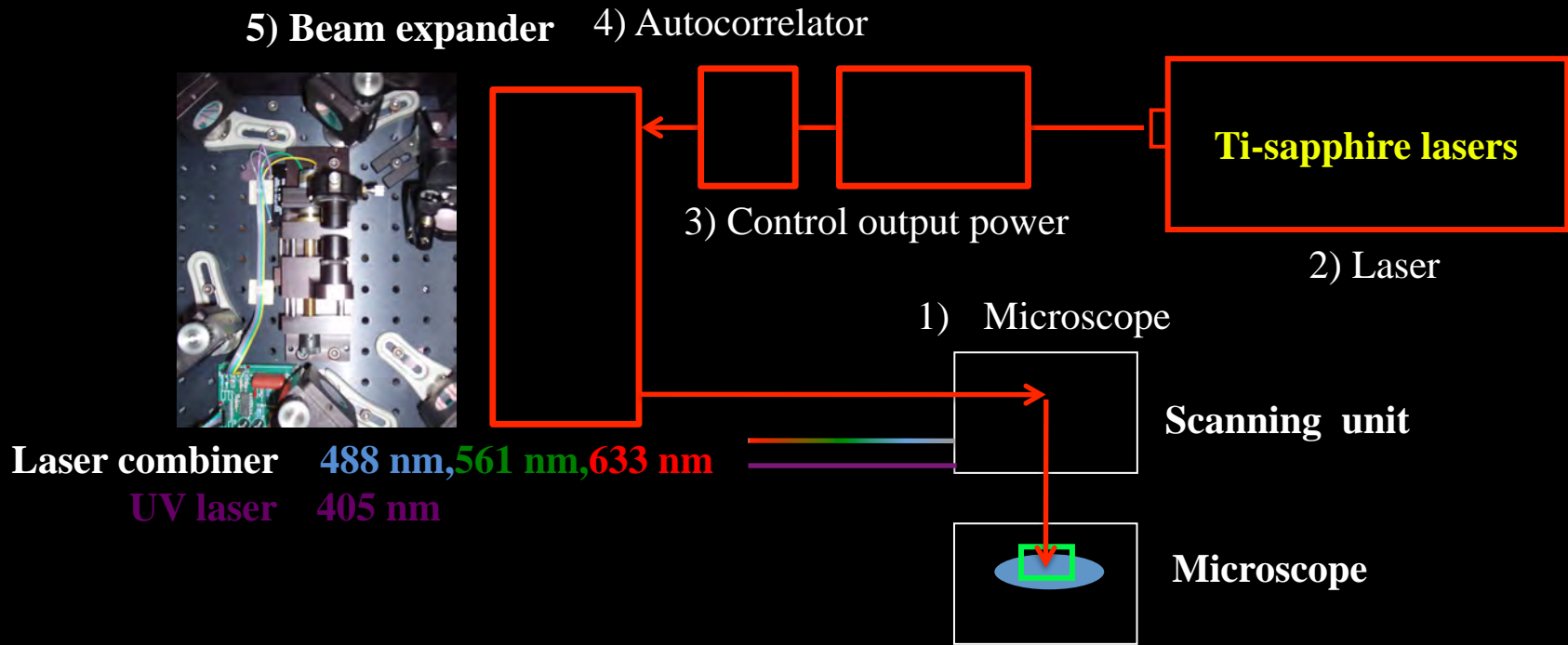
**Pulses: 100-150 fs**

(a) Measure the pulse at the “source”

(b) Measure the pulse and the power at the “specimen”



# Size of the laser beam

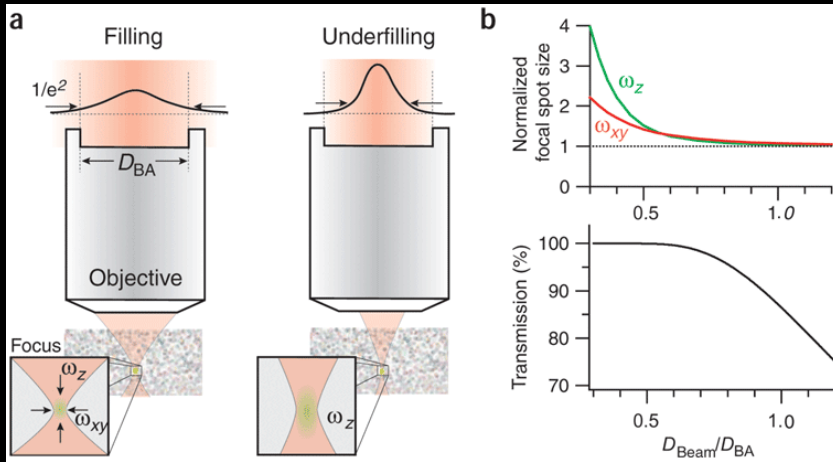


1) Control the size of the beam

2) Control the power at the specimen

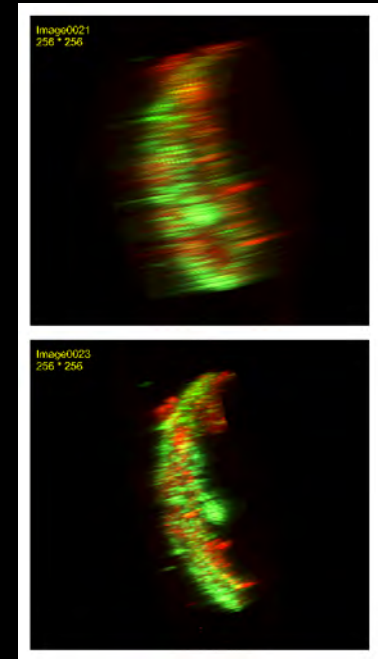
# Filling the backaperture of the lens

## Filling the back aperture of the lens



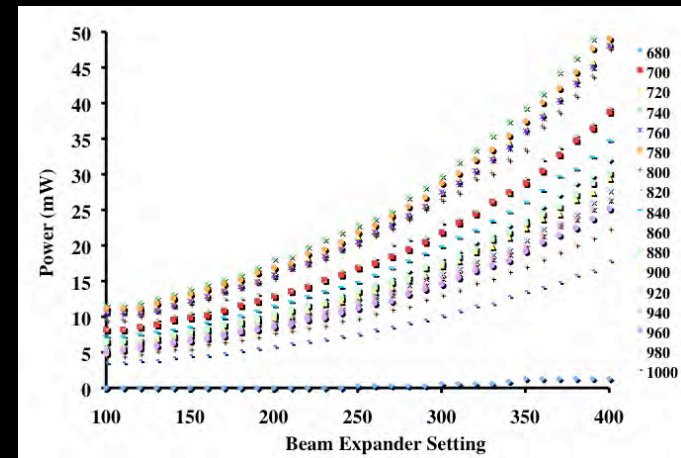
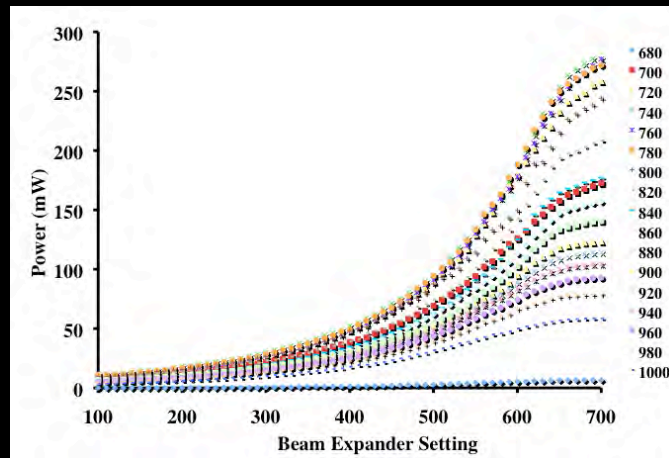
Essential for large lenses such as the 20X

## Underfilling



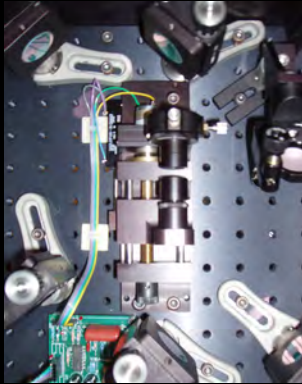
## Filling

## Control the power at the specimen by overfilling

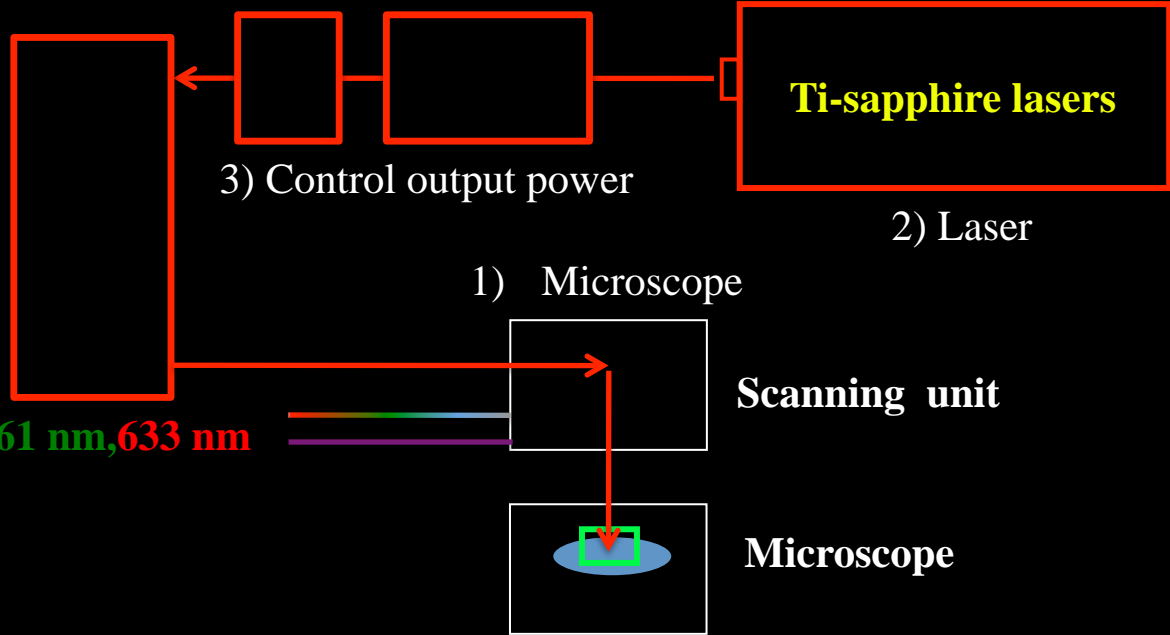


# Challenge: alignment of the beam

5) Beam expander    4) Autocorrelator

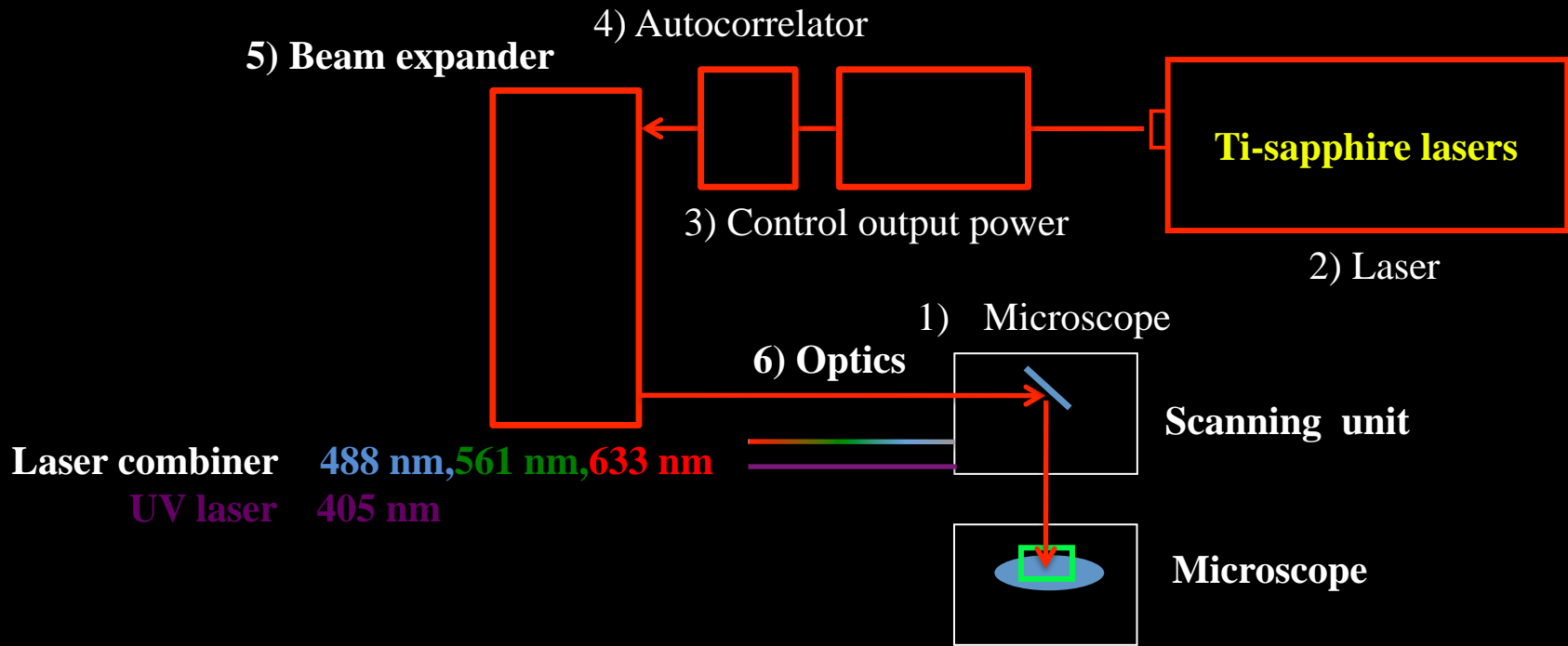


Laser combiner    488 nm, 561 nm, 633 nm  
UV laser    405 nm



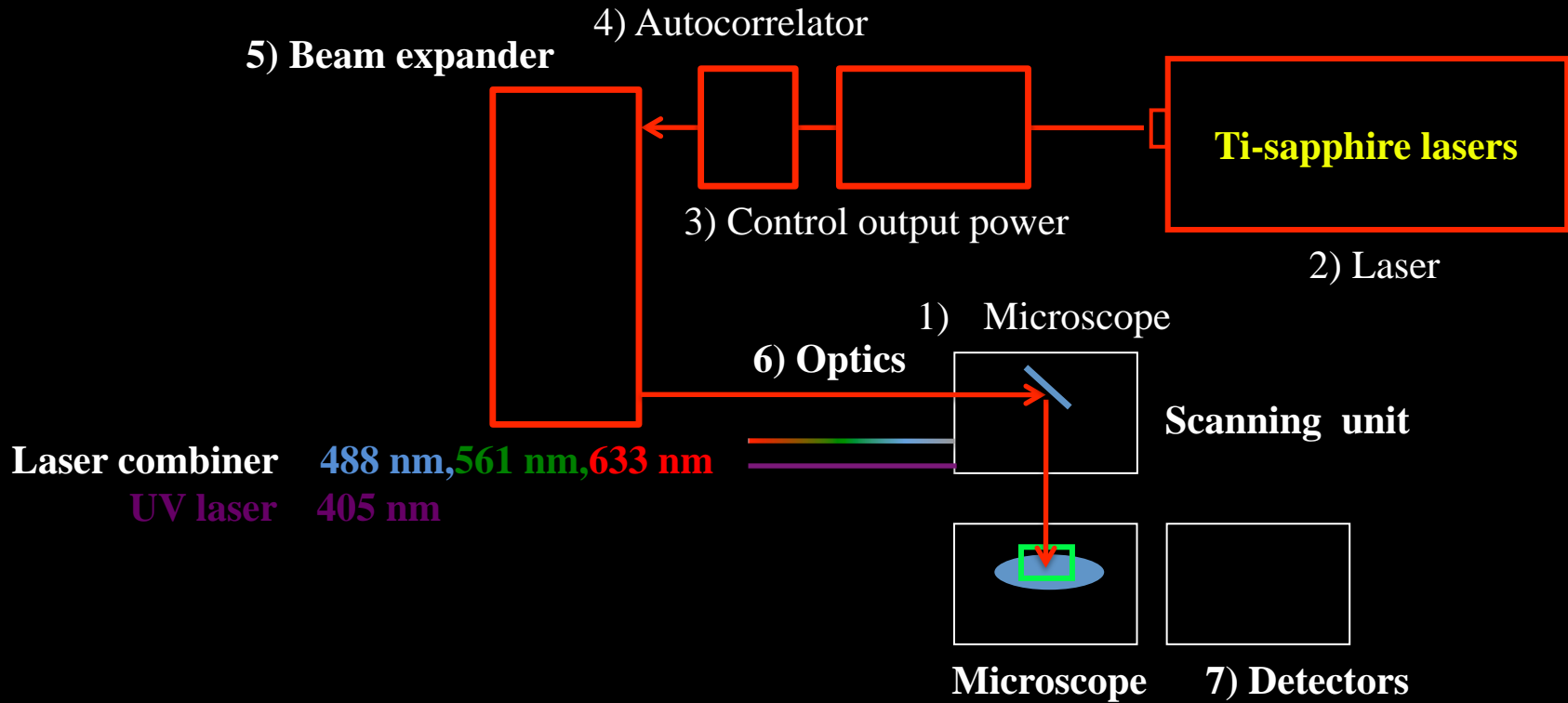


# Proper optics



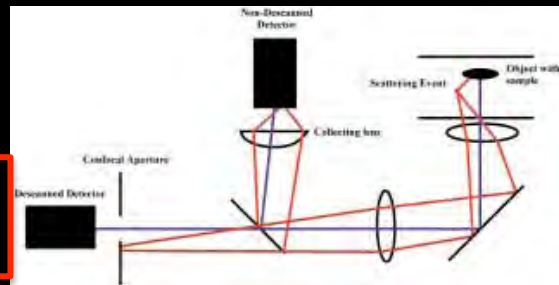
6) Excitation Dichroic mirror – reflect above 675-680 nm

# Non-descanned detectors



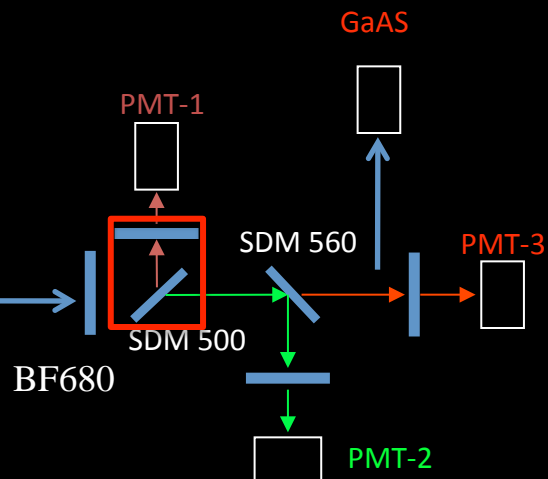
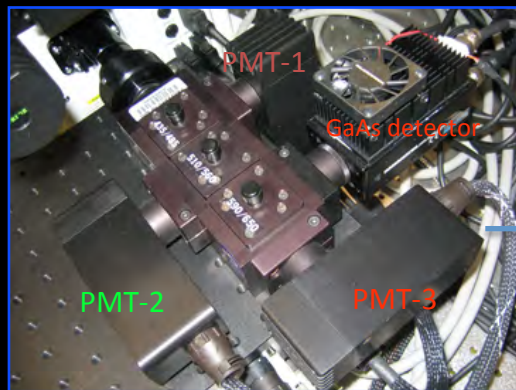
A) Descanned detectors

B) Non-descanned detector



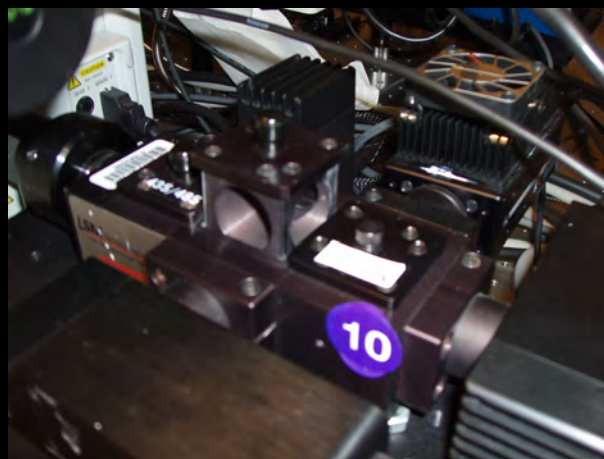
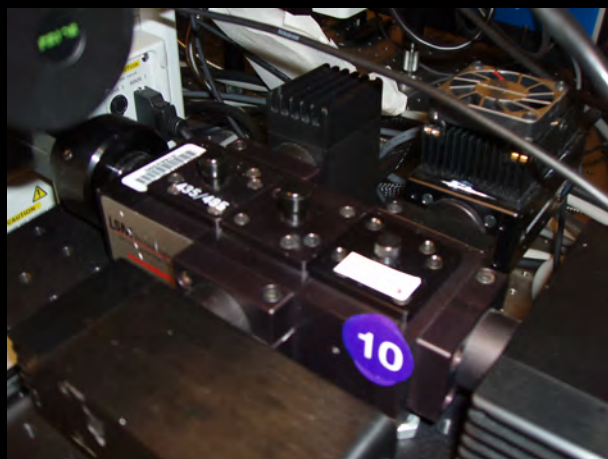
Positioning

# Non-descanned detectors



3 Cooled PMT from Hamamatsu R6060-11

1 Gallium Arsenide PMT



20x  
excitation 750  
emission < 510

Cooled PMT



GaAs detector



GaAs det gradient

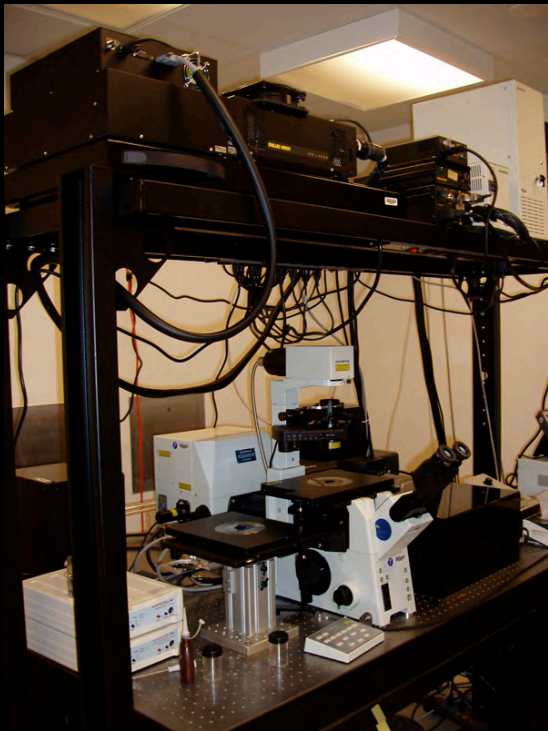
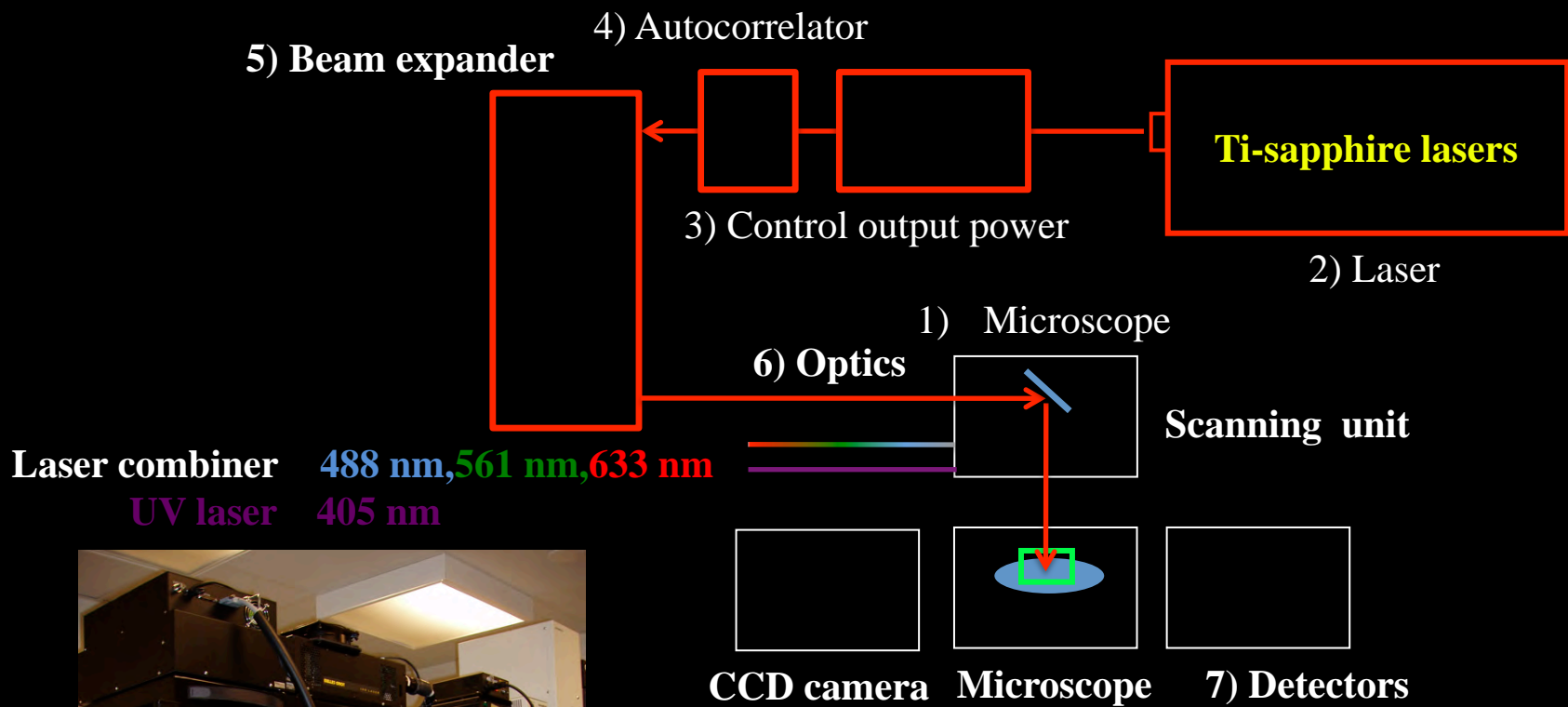


0 300  $\mu\text{m}$

# Non-descanned detectors

## Objective inverter with PMT





Confocal microscopy

Two-photon microscopy