

# Dynamic Microscopy 2006

Oct. 9.-11.

**Organizers: Gregory Harms, Peter Friedl**

**Location: Lecture Hall (Hörsaal) of the  
Rudolf Virchow Center, Versbacher Str. 9**

## Symposium

- 09:00 - 09:15 Registration  
09:15 - 09:30 Introduction – Gregory Harms and Peter Friedl
- 09:30 - 13:00 Session I: Molecular, Cellular, and Intravital Imaging**
- 09:30 – 10:15 Daniel Müller (HTU, Dresden, Germany)  
"From tissues, cells to single molecules: Recent developments and applications of Atomic Force Microscopy and Spectroscopy in biology medicine"
- 10:15 – 11:00 Anne Kenworthy (Vanderbilt, Nashville, TN, USA)  
"Probing lipid raft structure and dynamics by Quantitative Fluorescence Microscopy"
- 11:00 – 11:30 Coffee break and Vendor Exhibition**
- 11:30 – 12:15 Ernst Stelzer (EMBL, Heidelberg, Germany)  
"Avoiding flat and hard surfaces with single-plane illumination microscopy"
- 12:15 – 13:00 Robert Hoffman (Anti-Cancer, Inc. San Diego, CA, USA)  
"Whole-body subcellular imaging in the live animal"
- 13:00 – 14:30 Lunch break and Vendor Exhibition**
- 14:30 – 18:15 Session II: Molecular, Cellular, and Intravital Imaging**
- 14:30 – 15:15 Jackson Egen (NIH, Bethesda, MD, USA)  
"Visualization of immune cell dynamics using intravital multiphoton microscopy"
- 15:15 - 16:00 Maxime Dahan (DNS, Paris, France)  
"Establishment of cell polarity during nerve chemotaxis: a single-molecule approach"
- 16:00 – 16:30 Coffee break and Vendor Exhibition**
- 16:30 – 17:15 David Kremers (Cal Tech, Pasadena, CA, USA)  
"Making seven dimensions into two: biting off what humans can not chew"
- 17:15 – 18:30 Keynote Speaker Lecture:**  
Zena Werb (Univ. of California, San Francisco, USA)  
"Imaging the microenvironment during tumor progression"
- 18:30 – 19:00 Reception and Vendor Exhibition**

Oct. 10, 2006

**Location: Kurssaal 1 and Microscopy Core Facility  
Rudolf Virchow Center, Versbacher Str. 9**

**Course**

- 08:00 – 08:30**      **Organization and Coffee**  
(Colloquium Room, Rudolf-Virchow-Center)
- 08:30 – 09:00**      **Tutorial I: "The roles, benefits, and limitations  
with the use of nanoparticles in single-molecule  
imaging"**  
Maxime Dahan, DNS Paris
- 09:00 - 10:45**      **First Rotating Group Session**
- 10:45 – 11:30**      **Coffee Break and Vendor Demonstrations**  
**Tutorial II: "Technical aspects of multiphoton  
microscopy: strategies and troubleshooting"**  
Jackson Egen, NIH, USA,
- 11:30 – 13:15**      **Second Rotating Group Session**
- 13:15 – 14:15**      **Lunch, Vendor Demonstrations,**  
**Tutorial III: "FRAP: special considerations,  
aspects, and models"**  
Anne Kenworthy, Vanderbilt, USA
- 14:15 – 16:00**      **Third Rotating Group Session**
- 16:00 – 16:45**      **Coffee Break and Vendor Demonstrations**  
**Tutorial IV: "Special presentation"**  
David Kremers, Cal Tech, USA
- 16:45 – 18:30**      **Fourth Rotating Group Session**
- 18:30 -**              **Workshop Dinner and Wine**

## Four Rotating Group Session Course Topics

(max. 8 people each) (105 min)

### A Single-Molecule-Imaging

(Andrey Noskov, Benjamin Klasczyk, Kira Gromova,  
Gregory Harms)

**Location:** Room 013, Rudolf-Virchow-Center, Microscopy Core Center

**System:** Zeiss, custom-built, wide-field microscopes, Roper & INTAS  
Cameras

**Topics:**

- Single-molecule properties and microscope systems
- Fluorophores for single-molecule microscopy: GFP Variants, Artificial Organics, and Nanocrystals (Quantum Dots)
- Wide-Field Single Molecule Studies: Tracking, Co-localization, Stoichiometry, and Function
- Considerations for “in vivo” situations
- Analysis

### B Dynamic confocal fluorescence and reflection microscopy

(Katarina Wolf, Stephanie Alexander, Olga Levai -  
Leica Microsystems)

**Location:** Room 125 and Kurssaal 1, Rudolf-Virchow-Center

**System:** Leica SP5

**Topics:**

- Confocal reflection imaging for tissue reconstruction
- 3D reconstruction of collagen degradation
- Principles of 4D imaging and image reconstruction
- Live invasion of cancer cells in 3D collagen lattices
- Multiple-color simultaneous imaging

### Spectral unmixing

(sponsored by INTAS GmbH/CRI)

**Location:** Kurssaal 1

- Principles of spectral unmixing
- Spectral unmixing for tumor detection in whole mice
- Multi-color unmixing from cell cultures and histopathological tissue sections

**C Multiphoton microscopy and second harmonic generation imaging  
(Eline Hahnekamp, Markus Herzberg, Volker Andresen,  
Peter Friedl)**

**Location: Room 009, Microscopy Core Center**

**System:** Laviscope Trimscope/Verdi-Miro/APE OPO

**Topics:**

- Tuning the laser
- Second harmonic generation imaging of connective tissue
- 3D reconstruction of dermis and dermis vessels
- 4D Live cell imaging of GFP
- actin cytoskeleton in melanoma cells migrating in 3D collagen
- High-speed imaging of calcium transients

**D Photoactivation and photobleaching (FRAP)  
(Geoffrey Lambright, Mike Friedrich, Wiebke Buck, Olga Levai -  
Leica Microsystems, Gregory Harms)**

**Location: Room 009 and Room 125, Rudolf-Virchow Center**

**System:** Zeiss LSM 410 and Leica SP5

**Topics:**

- The essentials for FRAP, FLIP, and Photoactivation
- Characterization through control experiments
- Qualitative vs. Quantitative FRAP with artificial membrane staining on living cells
- Qualitative vs. Quantitative FRAP with GFP fusion proteins on Living Cells
- Two-color FRAP

**Oct. 11, 2006**

**Organizers: Gregory Harms, Peter Friedl**

**Duration: 9:00 to 15:00**

**Location: Kurssaal 1, Colloquium Room and Microscopy Core Facility  
Rudolf Virchow Center, Versbacher Str. 9**

## **Demonstrator Day**

### **Vendor Showroom and Demonstrations**

- 09:00 – 15:00**
- State of the art demonstrations by industrial partners
  - Guided Tours
    - Possibility to test your samples on the demonstration units.
    - The Vendors available are:

**Chromaphor  
INTAS  
Keyence  
La Vision - BioTech  
Leica  
Nikon  
Olympus  
Perkin-Elmer  
Till Photonics  
Visitron  
Zeiss**

**The third day will be held in Kurssaal 1 and  
Colloquium Room of the Rudolf-Virchow-Zentrum.**

**Individual appointments with the individual companies are possible but  
need to be made separately with them. Please make the individual  
appointments with the vendors on or before October 9 or 10, 2006.**

**Coffee and other refreshments: 09:30 - 15:00**

**Lunch: 12:00 - 13:30**

### **Planned Specific Vendor Demonstrations and Lectures: (Tentative Schedule)**

**\*\*\* Subject to Change \*\*\***

- |                      |   |
|----------------------|---|
| <b>09:00 - 09:45</b> | <b>INTAS - Imaging Systems from INTAS with Spectral Un-mixing</b>   |
| <b>09:45 – 10:15</b> | <b>La Vision - “Ultrafast Multiphoton Microscopy”</b>   |
| <b>10:15 - 11:00</b> | <b>Leica - "Experience a new level of speed" - The new Leica Laser-<br/>Microdissection-System LMD 6000 - Intro to Leica SP5 and TIRF</b> |
| <b>11:00 – 11:30</b> | <b>Nikon - „Perfect Focusing System – Farewell to focus drift“</b>  |
| <b>11:30 – 12:00</b> | <b>Olympus - “Advanced Image-Based Screening Applications in Life Science<br/>Research”</b>   |
| <b>13:00 – 13:30</b> | <b>Perkin-Elmer- “Ultra-View ERS a focussed value for Live Cell imaging in<br/>the offering of confocal systems”</b>                      |
| <b>13:30 - 14:00</b> | <b>Visitron - “Confocal Live Cell Imaging”</b>  |
| <b>14:00 - 14:30</b> | <b>Zeiss - "CellObserver for highspeed applications and ApoTome for optical<br/>sectioning”</b>   |
| <b>14:30 - 14:45</b> | <b>Chromaphor- BIOPTECHS Cell Sample Chamber</b>  |

## **System Demonstrations by Exhibitors**

-occurring throughout the symposium and workshop and by appointment

### **1. CHROMAPHOR Analysen-Technik GmbH**

**System:** BIOPTECHS Cell Sample Chamber

### **2. INTAS GmbH/ CRI**

**System:** Nuance Microscopy Imaging System, Maestro In Vivo Imaging System

- Principles of spectral unmixing with CCD camera imaging
- Spectral unmixing for tumor detection in whole mice
- Multi-color unmixing from cell cultures and histopathological tissue sections

### **3. Keyence Deutschland GmbH**

**System:** Fluorescence Microscope BZ-8000

- Compact, fully automatic Fluorescence Microscope for Biological Research

### **4. La Vision BioTech GmbH**

**System:** TriMScope

- Ultra-fast Multi-photon Microscopy with Multiple point confocal scanning
- New light sources for Multi-photon Microscopy

### **5. Leica Microsystems GmbH**

**Systems:** Leica Laser Microdissection System LMD 6000, SP5 Confocal with FLIM, Leica AM TIRF

- laser dissection microscope for the laboratory
- Multi-spectral Confocal Microscope: FRAP, FLIM, FCS, TPE
- Fully automated TIRF microscope

### **6. Nikon GmbH**

**System:** TE2000 PFS- TIRF and TE2000 C1 confocal

- Total Internal Reflection Imaging with Perfect Focus System
- Spectral confocal imaging system on an TE2000 inverted microscope

### **7. Olympus GmbH**

**System:** *cell R*, *MVX10*, *Scan R* Microscope

- All-in-one Live Cell Imaging system
- Macro Zoom Fluorescence Microscope
- Screening Station for Life Sciences

### **8. Perkin-Elmer GmbH**

**System:** Ultra VIEW ERS

- Nipkow disc, fast live-cell confocal imaging system
- Equipped with an ultra-sensitive EMCCD camera

## **9. Till-Photonics**

**System:** iMIC imaging Setup, TILL.TIRF system

- Real Time Imaging System for Widefield, FRET, DualColor, FRAP
- flexible, versatile and PC-based
- TILL.TIRF dual-port condensor for maximal S/N-ratio for and controlable penetration depth

## **10. Visitron Systems**

**System:** VT Infinity Confocal, Cascade II EMCCD

- Low Noise Single Molecule Imaging with CascadeEMCCD
- 2D array multipoint confocal scanner with 800 frames/second and simple FRET measurements

## **11. Carl Zeiss GmbH**

**System:** Axiovert 200 ApoTome, Cell Observer

- High resolution, high contrast and simple fluorescence microscopy for 2D and 3D imaging with user friendly software
- Multi-channel, time-resolved biological imaging system