



### 3D CELL EXPLORER-fluo



# COMBINE THE BEST OF TWO WORLDS

#### **COMPLETE 3D SOLUTION**

Combine high quality tomographic data with fluorescent markers

#### **MULTIPLEXING**

Explore up to 10 markers in parallel

# EXTENDED LIVE CELL IMAGING

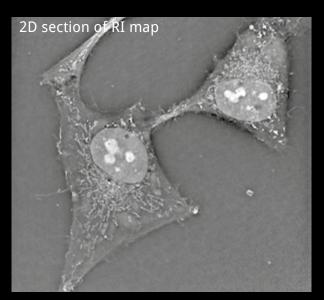
Limit cell damages caused by fluorescent markers, bleaching and phototoxicity

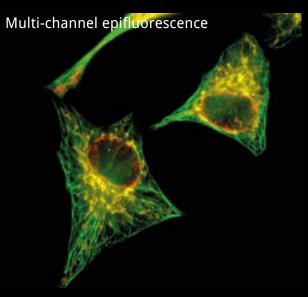
#### THE HOLOTOMOGRAPHIC FLUORESCENCE MICROSCOPE

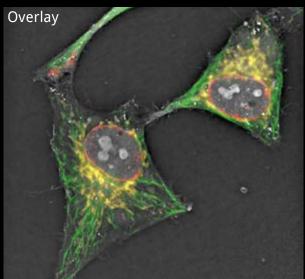
## REVOLUTIONARY TECHNOLOGY

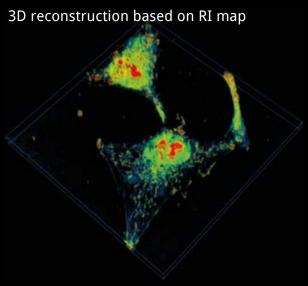
## MULTIMODAL COMPLETE SOLUTION FOR 3D LIVE CELL EXPLORATION

Combine 3D refractive index analysis with a fully integrated 3 channel fluorescence module to image your live cells as they are and as long as you want. Put chemical information into structural context for new biological insights.









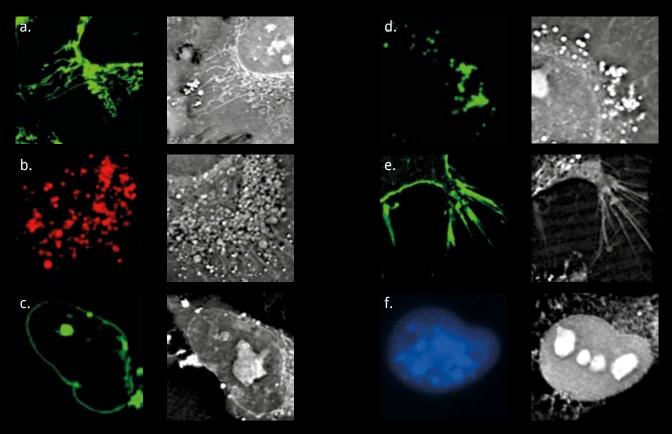


IN A VERY SHORT PERIOD OF TIME, THE 3D CELL EXPLORER HAS BECOME VERY INTENSIVELY USED AND WE HAVE FOUND APPLICATIONS IN SEVERAL DIFFERENT DISEASE AREAS — WE WOULD NOT WANT TO BE WITHOUT THIS INSTRUMENT.

Oliver Nayler, PhD Senior Director, Head Cardiovascular & Fibrosis Biology Idorsia Pharmaceuticals Ltd, Allschwil, Switzerland

#### **MULTIPLEXING**

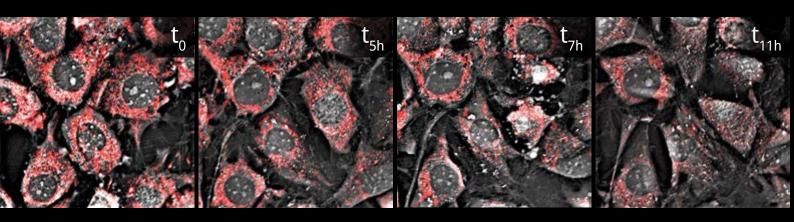
Transform 2D fluorescence into 3D cell tomography: Identify cell organelles through fluorescence and monitor non-invasively their structures & dynamics in 3D & marker-free. Explore fluorescence (3 channels) & Digital Stains (8 channels) simultaneously.



Examples of co-localization for a. mitochondria, b. lysosomes, c. nuclear membrane, d. lipid droplets, e. plasma membrane, f. nucleus & nucleoli.

#### **EXTENDED LIVE CELL IMAGING**

Image your live cells as long as you need. Limit cell damages caused by fluorescent markers, bleaching and phototoxicity.



Long-term imaging (11hrs) of mouse pre-adipocytes. Mitochondria were labeled with mitoTracker. A holotomographic image was taken every 15 seconds and a fluorescence image every 5 minutes.

# DISCOVERMORE



#### **EXPLORE A NEW VISION**

Long observation time New space for discoveries

#### IMPROVE KNOWLEDGE

Combine fluo and RI tomography Up to 10 markers in parallel

#### **PROCESS NEW DATA SETS**

Unique organelle segmentation Quantitative data analysis

#### **SAVE EXPERIMENTAL TIME**

No preparation Short setup time Fast & easy acquisition

#### **TECHNICAL SPECIFICATIONS**

Illumination Source	Holotomography: Class 1 low power laser ( $\lambda$ =520 nm, sample exposure 0.2 mW/mm <sup>2</sup> ) Fluorescence: High speed switchable <100 $\mu$ m, Lifetime > 20'000 hours each channel
Resolution	Holotomography: x,y: 200 nm; z: 400 nm (3D image) Fluorescence: rx,y: ~ 400 nm (2D image)
Field-of-view	Holotomography: $90 \times 90 \times 30 \mu m$ Fluorescence: $90 \times 90 \mu m$
Microscope Objective	Dry objective / 60× magnification / NA 0.8
Channels	Holotomography: Up to 7 simultaneous Fluorescence: DAPI + FitC + TritC   FitC + TritC + Cy5   DAPI + FitC + TritC / Cy5
Imaging	Holotomography: 3D Fluorescence: 2D 4D time lapse: (RI + fluo)
Time resolution	Holotomography: 0.5 fps 3D RI frame Fluorescence: 3 fps each channel
Camera	USB 3.0 CMOS Sony IMX174 sensor / Quantum Efficiency (typical) 70% (at 545 nm) / Dark Noise (typical) 6,6 e <sup>-</sup> / Dynamic Range (typical) 73,7 dB
Dimensions (width × depth × height in mm)	3D Cell Explorer-fluo: 380 × 170 × 445 Fluorescence module: 77 × 186 × 162
Weight	12 kg