



# 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**

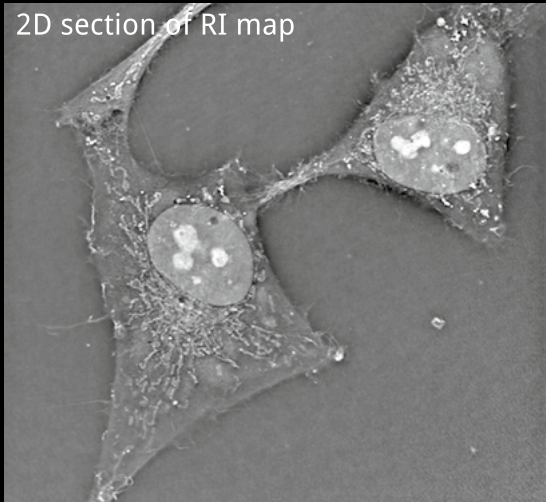
# REVOLUTIONARY TECHNOLOGY

## MULTIMODAL COMPLETE SOLUTION FOR 3D LIVE CELL EXPLORATION

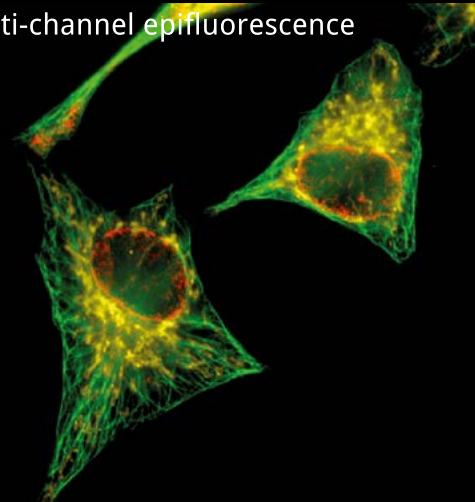
The 3D Cell Explorer measures the quantitative Refractive Index (RI) of cell organelles in seconds and 3D. This allows for biological features to be segmented based on their physical characteristics.

The 3D Cell Explorer-fluo combines 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.

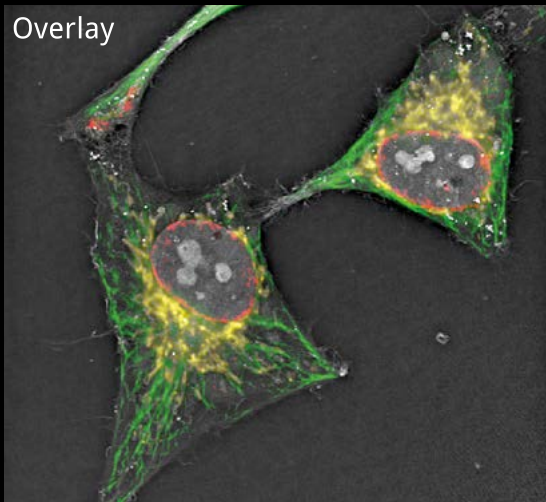
2D section of RI map



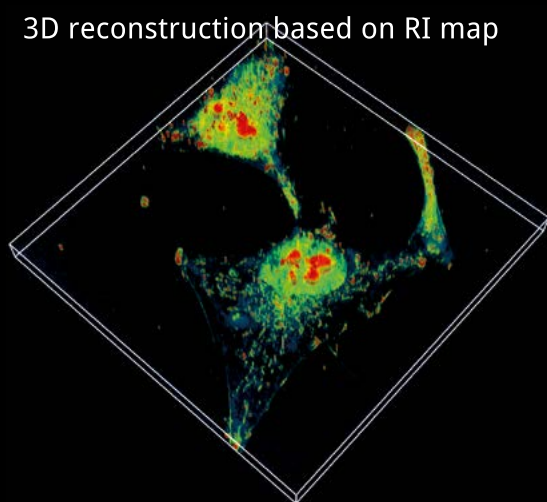
Multi-channel epifluorescence



Overlay



3D reconstruction based on RI map

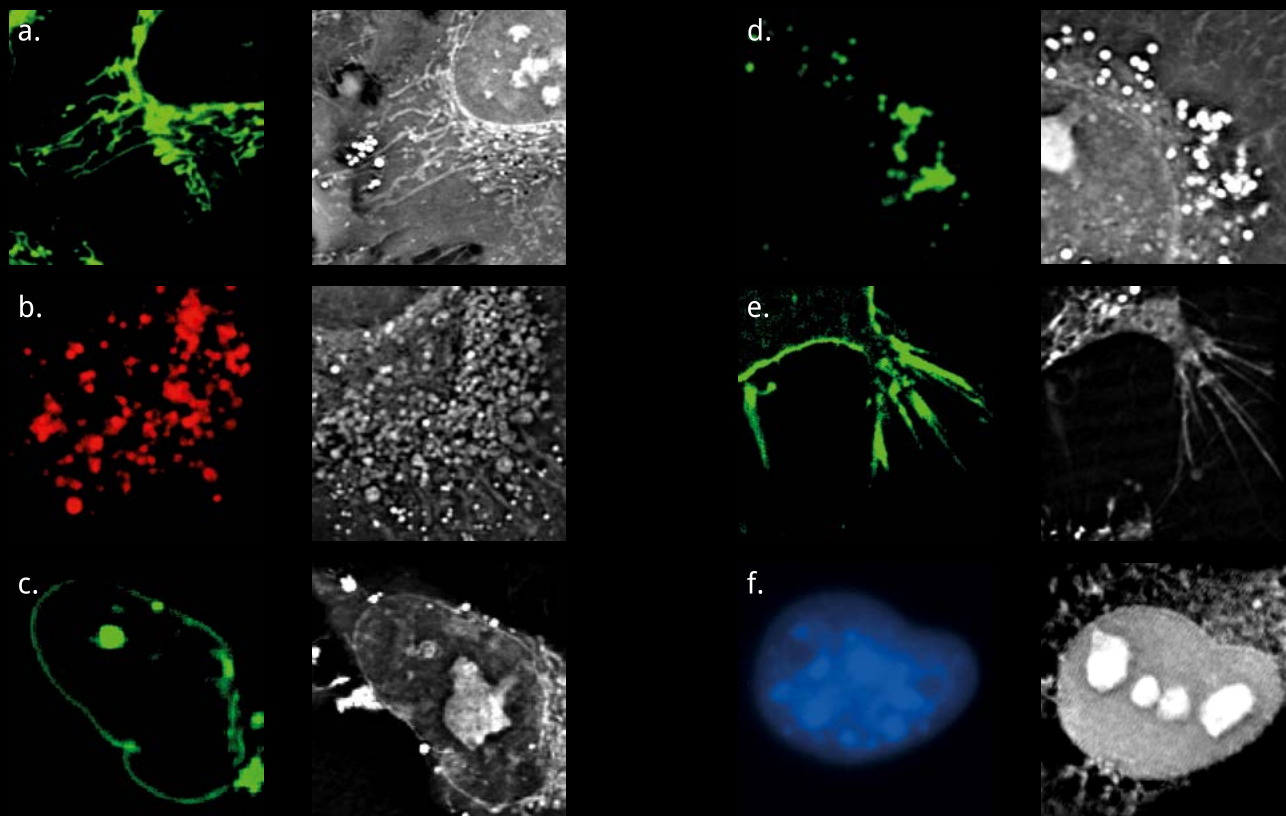


**IN A VERY SHORT PERIOD OF TIME, THE 3D CELL EXPLORER HAS BECOME VERY INTENSIVELY USED AND WE HAVE FOUND APPLICATIONS IN SEVERAL DIFFERENT DISEASE AREAS — WE WOULD NOT WANT TO BE WITHOUT THIS INSTRUMENT.**

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

## MULTIPLEXING

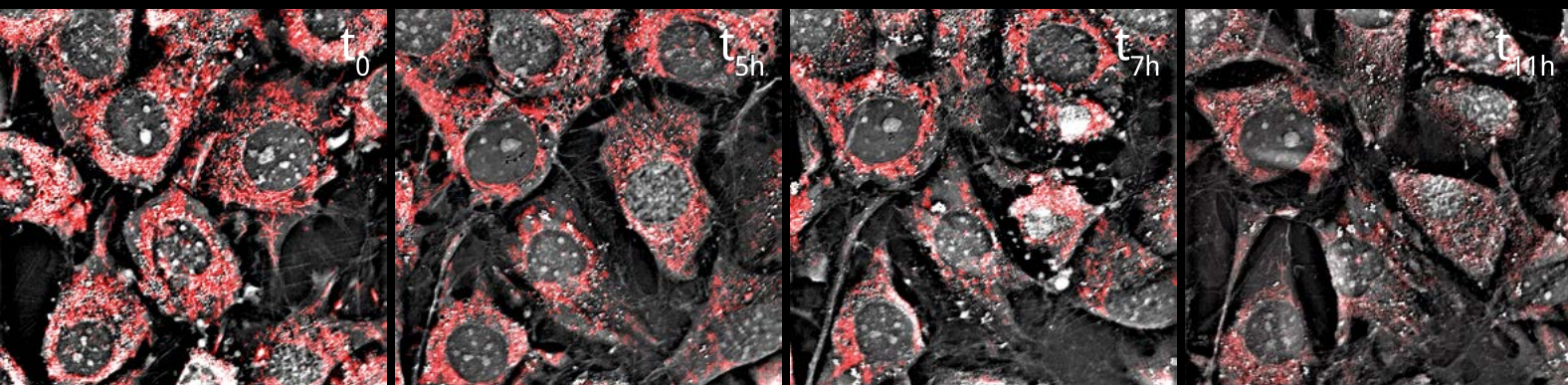
Transform 2D fluorescence into 3D cell tomography: Identify cell organelles through fluorescence and monitor non-invasively their structures & dynamics in 3D & marker-free. Explore fluorescence (3 channels) & Refractive Index (7 organelles) simultaneously.



Examples of correlation 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.



# DISCOVER MORE



## 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

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