

“ I prefer to take the train.”

*Joris Kaal (a week ago)*



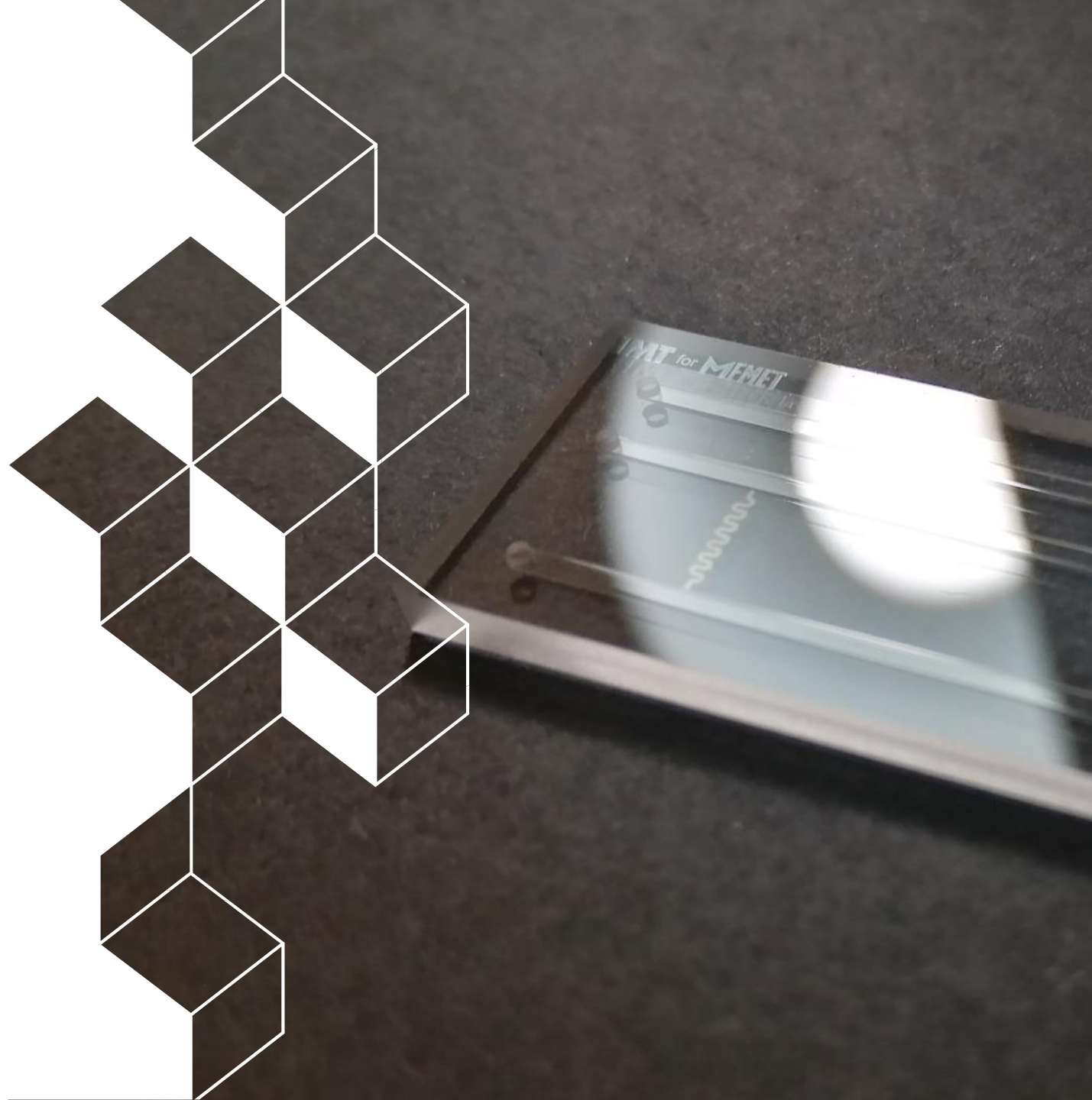




# Dimensional characterization of microfluidic chips

What we can and can't do

Joris Kaal, CEA-Leti



# Why?

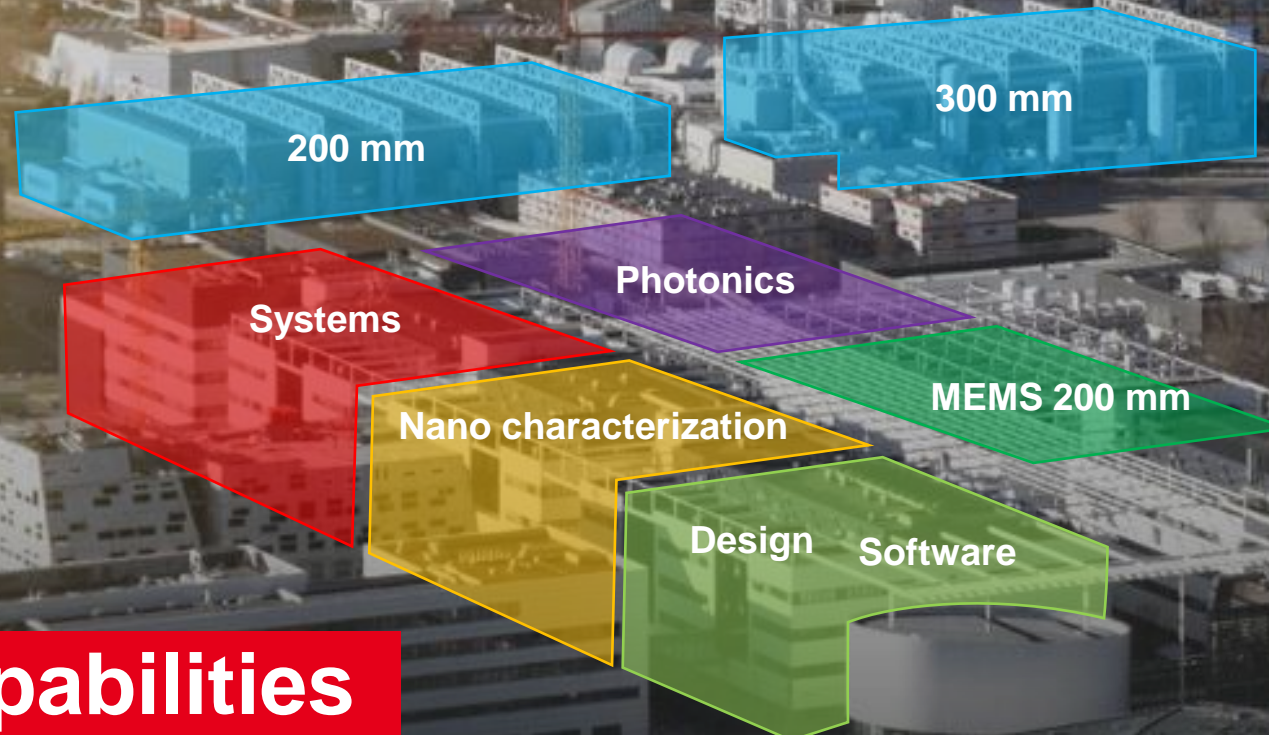


**01**

**GENERAL NEED  
MEASUREMENT  
METHODS**

**02**

**DIMENSIONS ARE  
BASIS OF OTHER  
MEASUREMENTS**



## CEA-Leti's capabilities

- > 11 000 m<sup>2</sup> of cleanroom dedicated to nanofabrication
- > 800 people divided in 4 shifts
- > Supported by a **nanocharacterization platform**

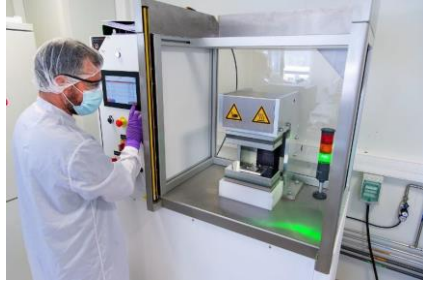


# PRISM

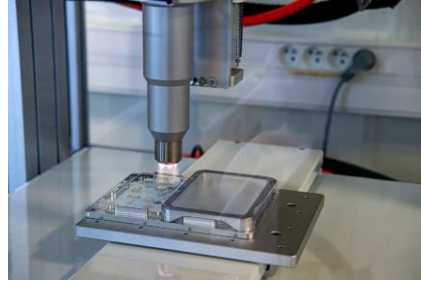
## Research Platform for Integration, Surface functionalization and Microfabrication



Polymer micro-machining and aluminium moulds



Thermoforming and heat sealing



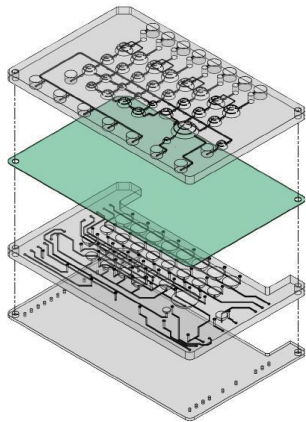
Atmospheric plasma



Silicone film lamination



Electrochemical sensors and surface functionalisation



Generic library for standardised and ISO-conform building blocks



Cleanroom environment

# How are we measuring?



Protocol  
(CEA)  
**1**

Optical  
profilometry

Protocol  
(LNE)  
**2**

Optical  
microscopy

Protocol  
(UofG)  
**3**

Tiled digital  
imagery

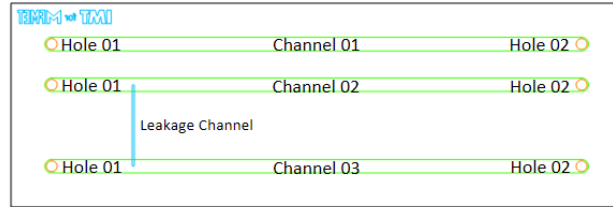
Protocol  
(UofG)  
**4**

Confocal  
microscopy

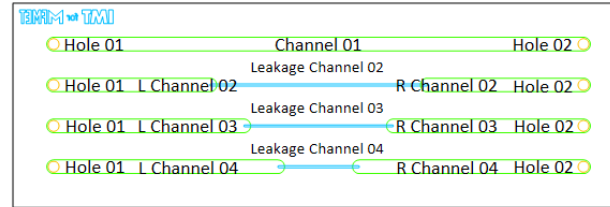
# What are we measuring?



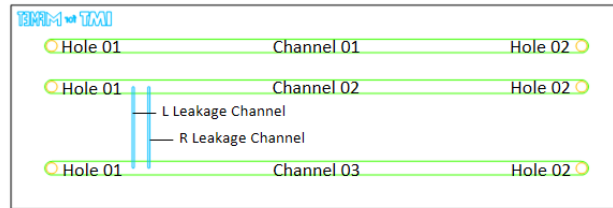
Design 01



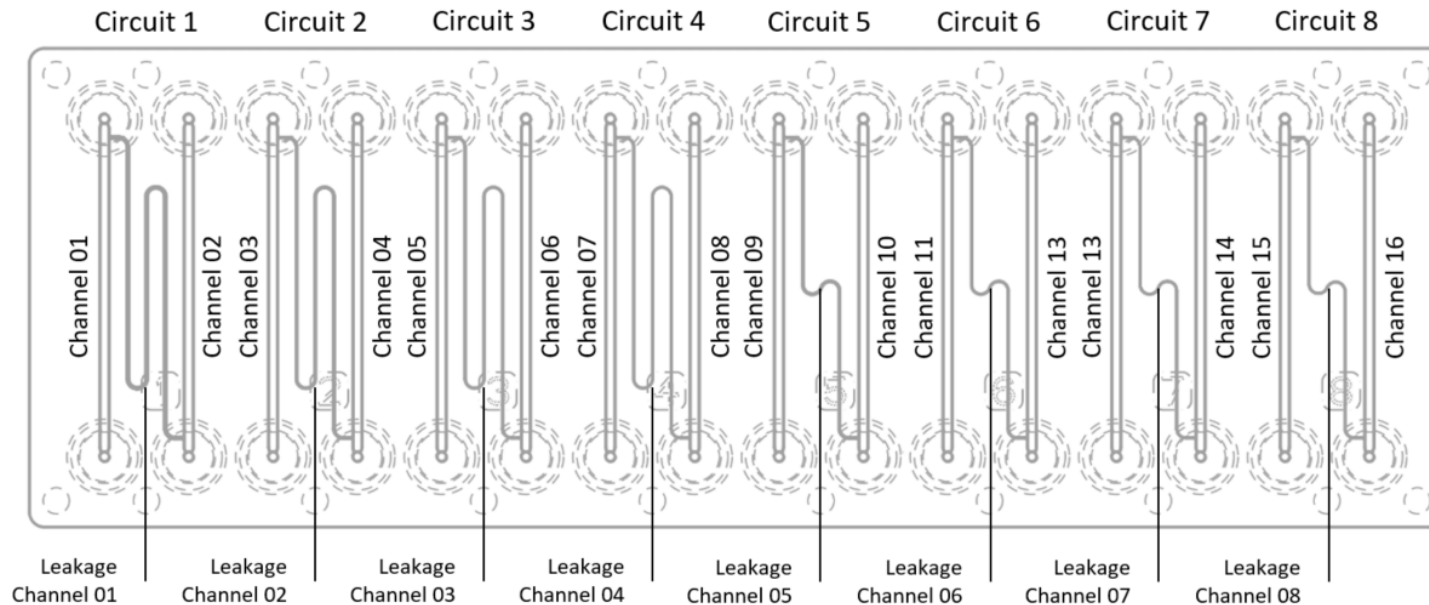
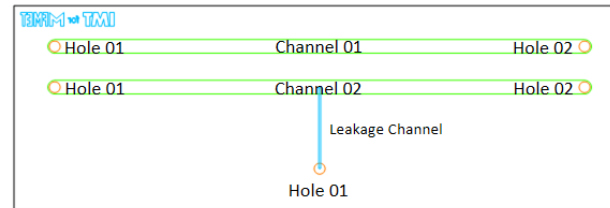
Design 05



Design 02



Design 06

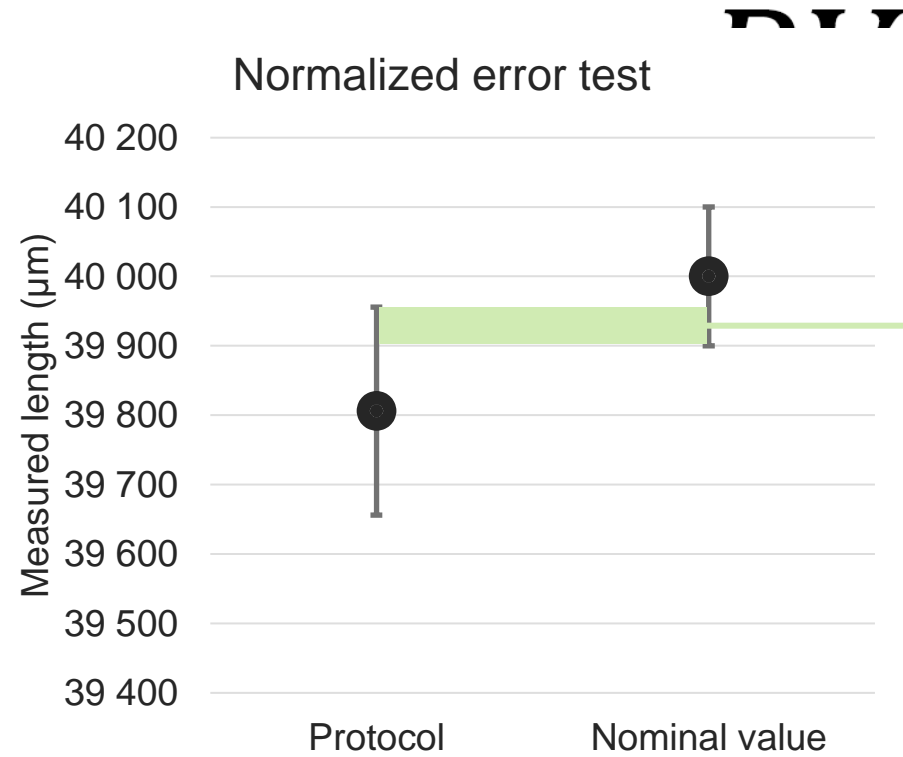


263 © Bio glass  
POPAS © COC

# How do we compare?



$$E_n = \frac{\sigma}{\mu} \sqrt{\frac{1}{n} \sum_{i=1}^n (x_i - \mu)^2}$$



$$\frac{|E_n|}{1} \leq 1$$

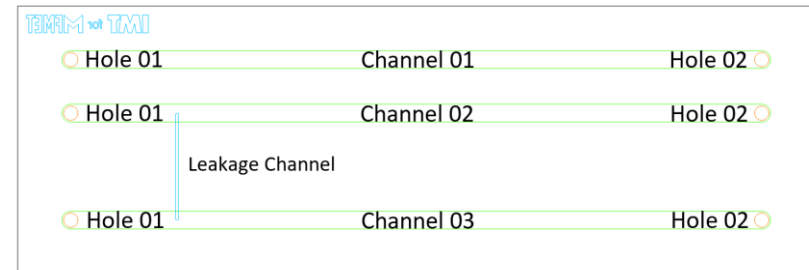


# Normalized error values in glass



	<u>Diameter Hole 01</u>	<u>Diameter Hole 02</u>	<u>Width</u>	<u>Depth</u>	<u>Length</u>
Protocol 1	0.16	0.33	-0.93	<b>-7.81</b>	0.02

Design 01



# Material influence (xy-plane)



## Optical profilometer vs. Optical profilometer

on TOPAS® COC

Width

Average  
measured

Standard  
deviation

Channel 01	503.1 $\mu\text{m}$	0.1 $\mu\text{m}$ (0.0 %)
Leakage Channel 01	108.5 $\mu\text{m}$	0.1 $\mu\text{m}$ (0.1 %)
Channel 02	508.1 $\mu\text{m}$	0.1 $\mu\text{m}$ (0.0 %)

On D263® Bio glass

Width

Average  
measured

Standard  
deviation

Channel 01	991.3 $\mu\text{m}$	0.2 $\mu\text{m}$ (0.0 %)
Channel 02	992.1 $\mu\text{m}$	0.2 $\mu\text{m}$ (0.0 %)
Channel 03	992.4 $\mu\text{m}$	0.2 $\mu\text{m}$ (0.0 %)
Leakage Channel	143.0 $\mu\text{m}$	0.5 $\mu\text{m}$ (0.3 %)

# Material influence (depth)



## Optical profilometer vs. Optical profilometer

### on TOPAS® COC

Depth

	<u>Average measured</u>	<u>Standard deviation</u>
Channel 01	358.0 $\mu\text{m}$	1.7 $\mu\text{m}$ (0.5 %)
Leakage Channel 01	123.6 $\mu\text{m}$	1.8 $\mu\text{m}$ (1.4 %)
Channel 02	372.9 $\mu\text{m}$	2.7 $\mu\text{m}$ (0.7 %)

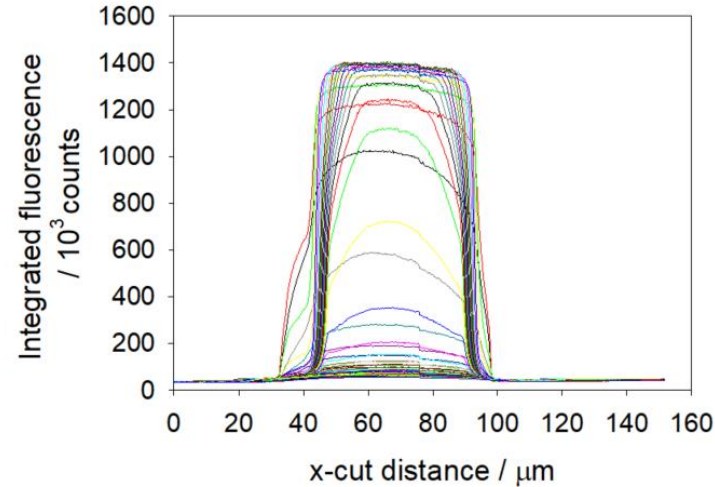
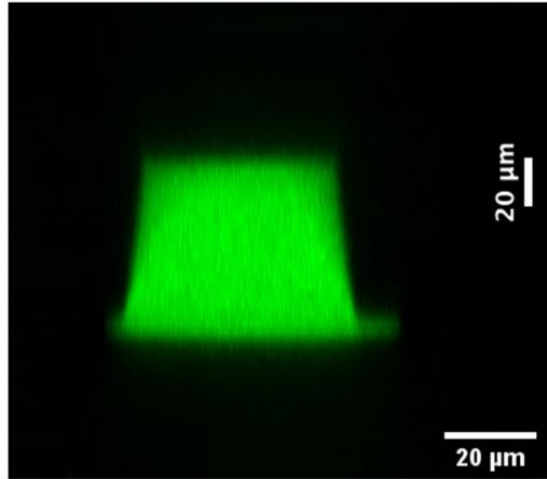
### On D263® Bio glass

Depth

	<u>Average measured</u>	<u>Standard deviation</u>
Channel 01	13.8 $\mu\text{m}$	4.6 $\mu\text{m}$ (33.0 %)
Channel 02	17.9 $\mu\text{m}$	1.3 $\mu\text{m}$ (7.6 %)
Channel 03	20.2 $\mu\text{m}$	3.3 $\mu\text{m}$ (16.3 %)
Leakage Channel	N/A	N/A

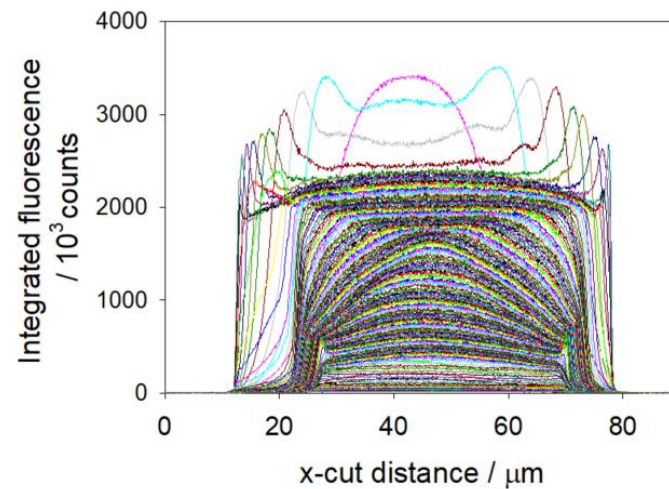
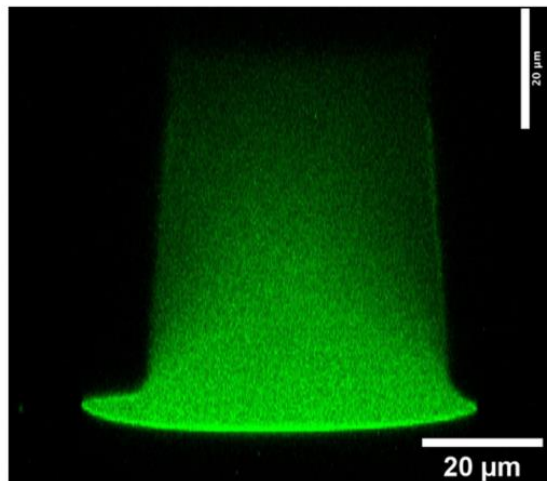


# A way to solve depth measurements



Zeiss LSM 510  
x20 NA = 0.5 objective  
resolution in Z of 2.07 μm

Estimated channel depth:  $74.5 \pm 4 \mu\text{m}$



Zeiss LSM 510  
x100 NA = 1.3 objective  
resolution in Z of 0,44 μm

Estimated channel depth:  $73.0 \pm 0.8 \mu\text{m}$

# Conclusion

- All protocols are consistent
- D263© Bio glass and TOPAS© COC are comparably compatible with measurements
- Depth measurements are a challenge

# Outlook

- How to do depth measurements?
  - Improve optical profilometry?
  - Zeta plane imagery?
  - Confocal microscopy?
- Compare more materials







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