

Thermodiffusion of latex beads studied with a microfluidic cell

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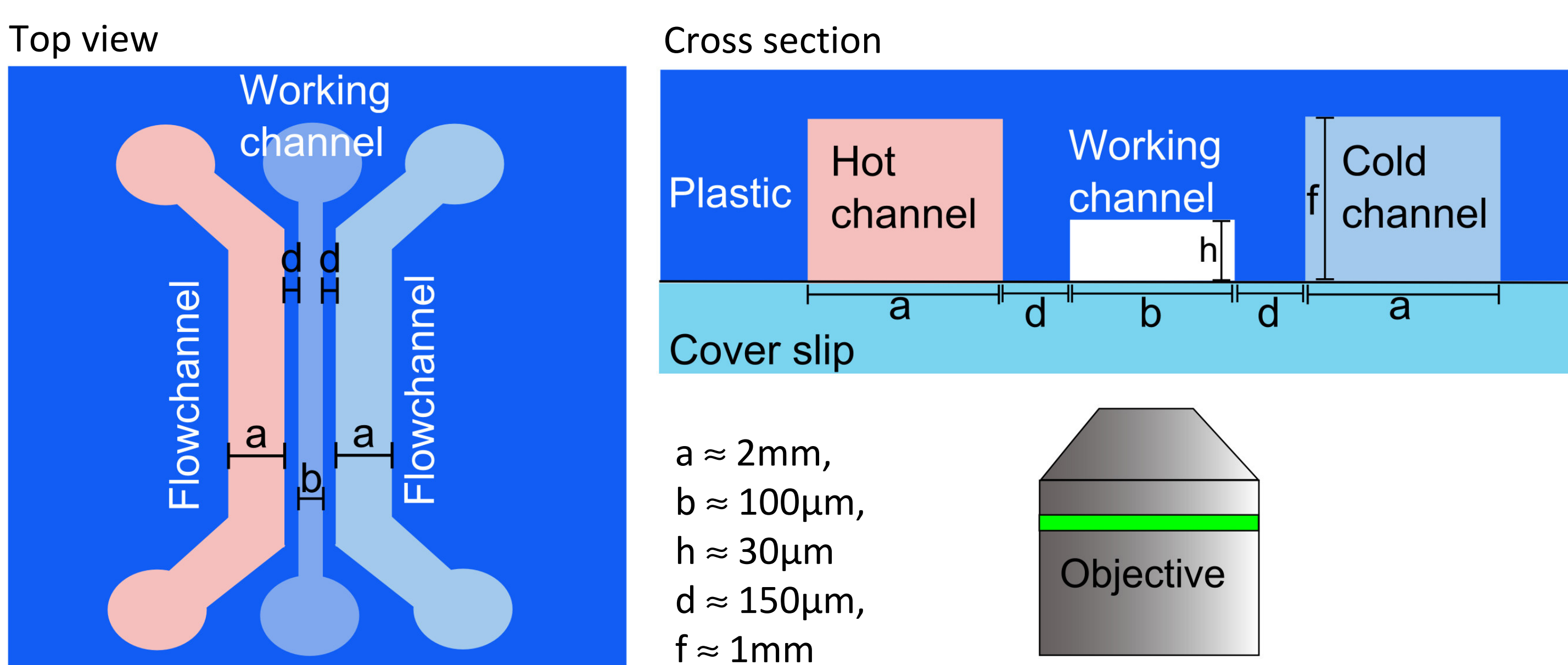
Introduction

Challenges in thermophoresis

- Current understanding of thermophoresis in liquids is not complete.
- Both theory and experiments are required.
- Most of currently used methods (optical, based on beam deflection or diffraction) are not suitable for studying thermophoresis of big colloidal particles ($d > 100\text{nm}$) as well as for complex mixtures
- Application of thermophoresis for biosensors

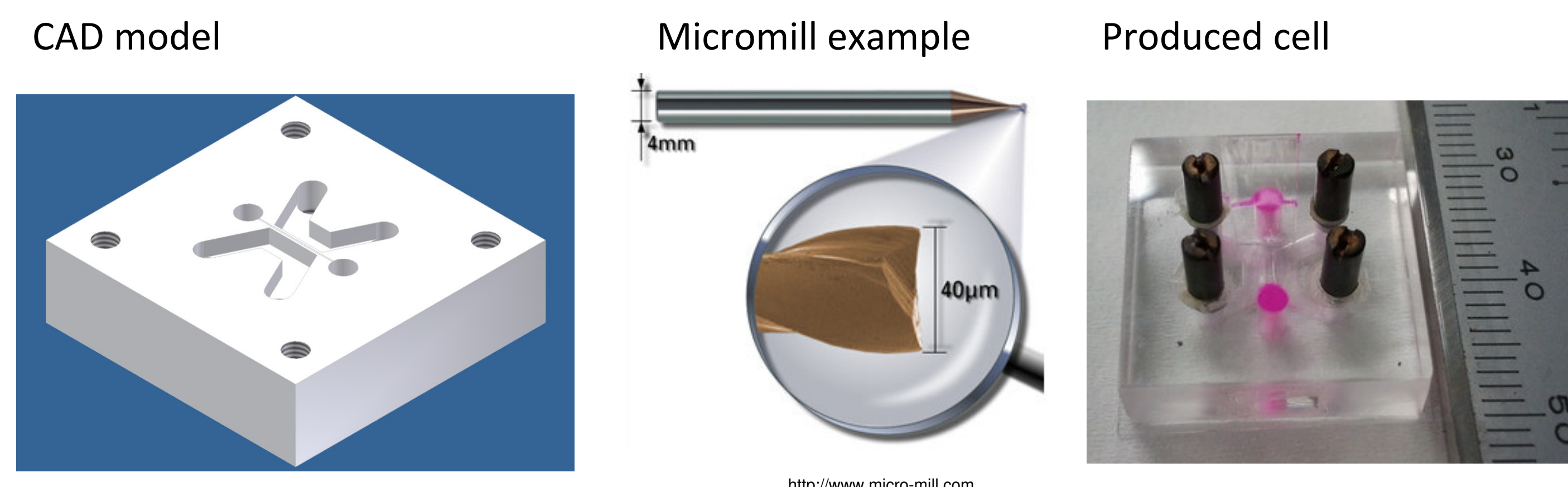
Microfluidic cell

Design



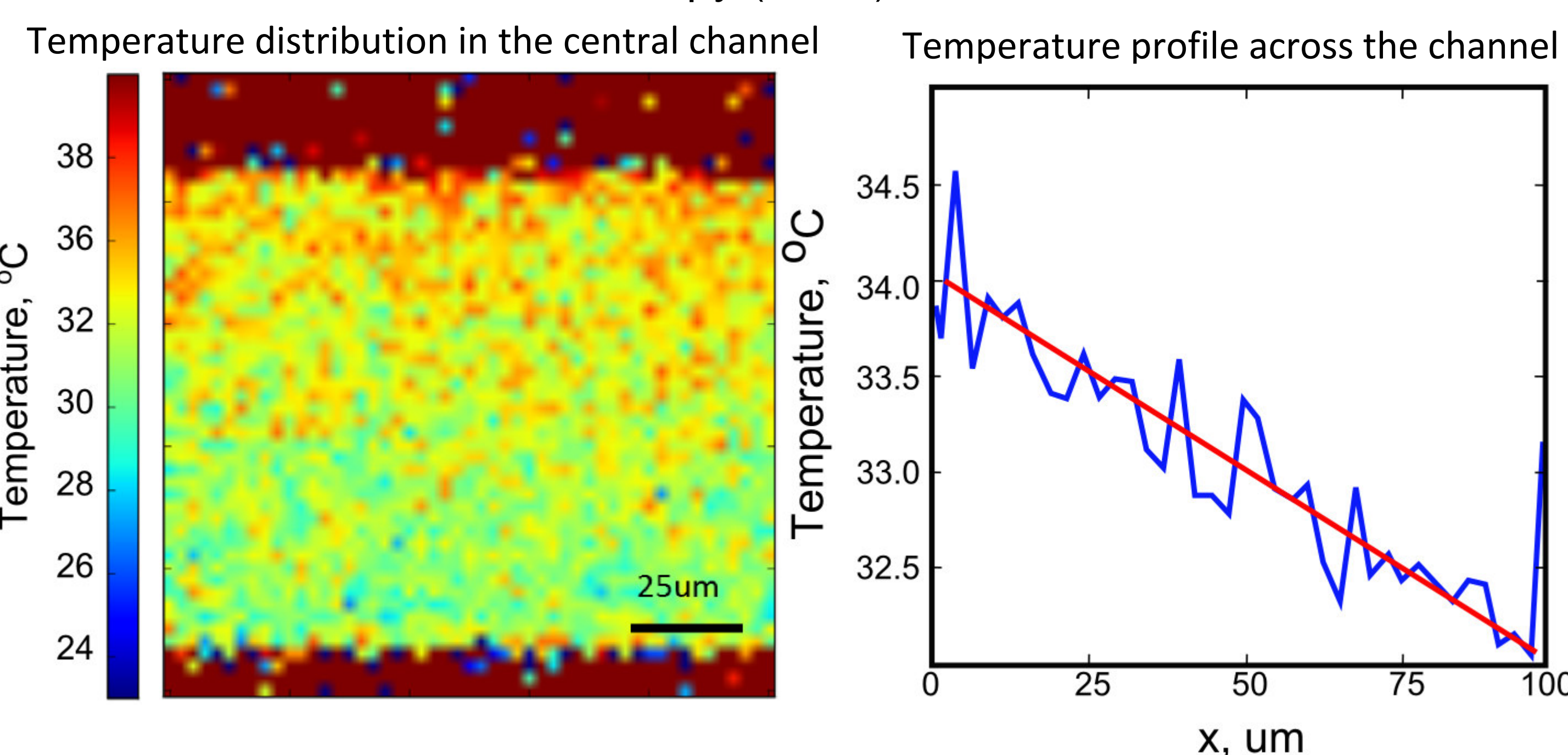
Production

Made of Plexiglas by micromilling



Characterization

Fluorescence life time microscopy (FLIM) with Rhodamine B



Advantages

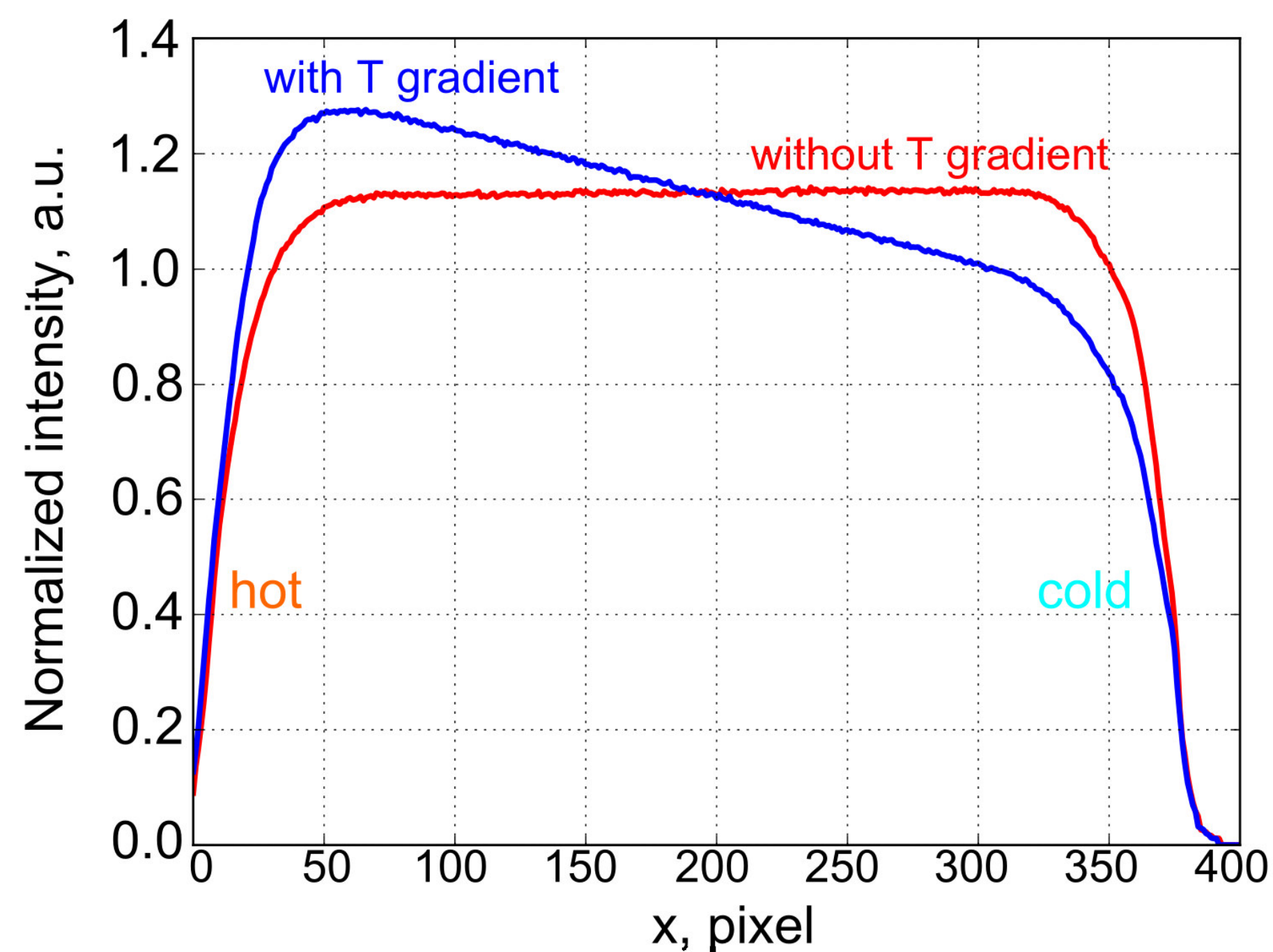
- Investigation in buffer solutions is possible
- Large colloids can be studied ($>100\text{nm}$)
- Complex mixtures can be investigated
- Large temperature gradient 10^4 K/m
- Single particles can be tracked (studied)

Small particles

Latex nanobeads (25 nm) 1% in water

Thermo Scientific™ Fluoro-Max green fluorescent internally dyed polystyrene particles in water.

Particles cannot be distinguished in the microscope. Fluorescence intensity is taken as a measure of concentration



At the equilibrium when $c \ll 1$:

$$\vec{J} = -D\vec{\nabla}c - cD_T\vec{\nabla}T = 0$$

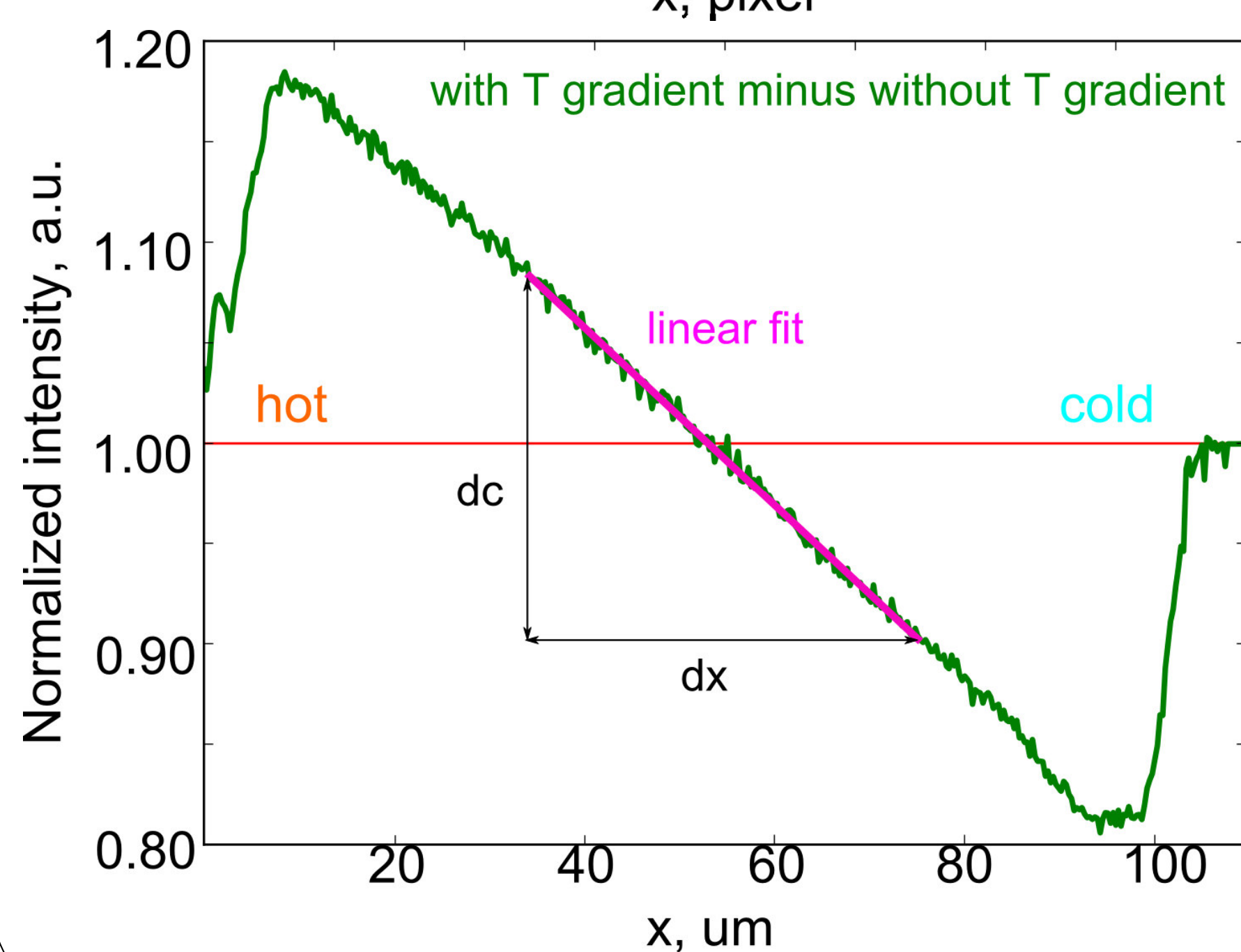
$$\downarrow$$

$$S_T = -\frac{D_T}{D} = -\frac{1}{c} \frac{|\vec{\nabla}c|}{|\vec{\nabla}T|} = -\frac{dc/dx}{c dT/dx}$$

$$S_T \approx -0.18\text{ K}^{-1}$$

for $T = 25^\circ\text{C}$

Validation against TDFRS measurements is planned

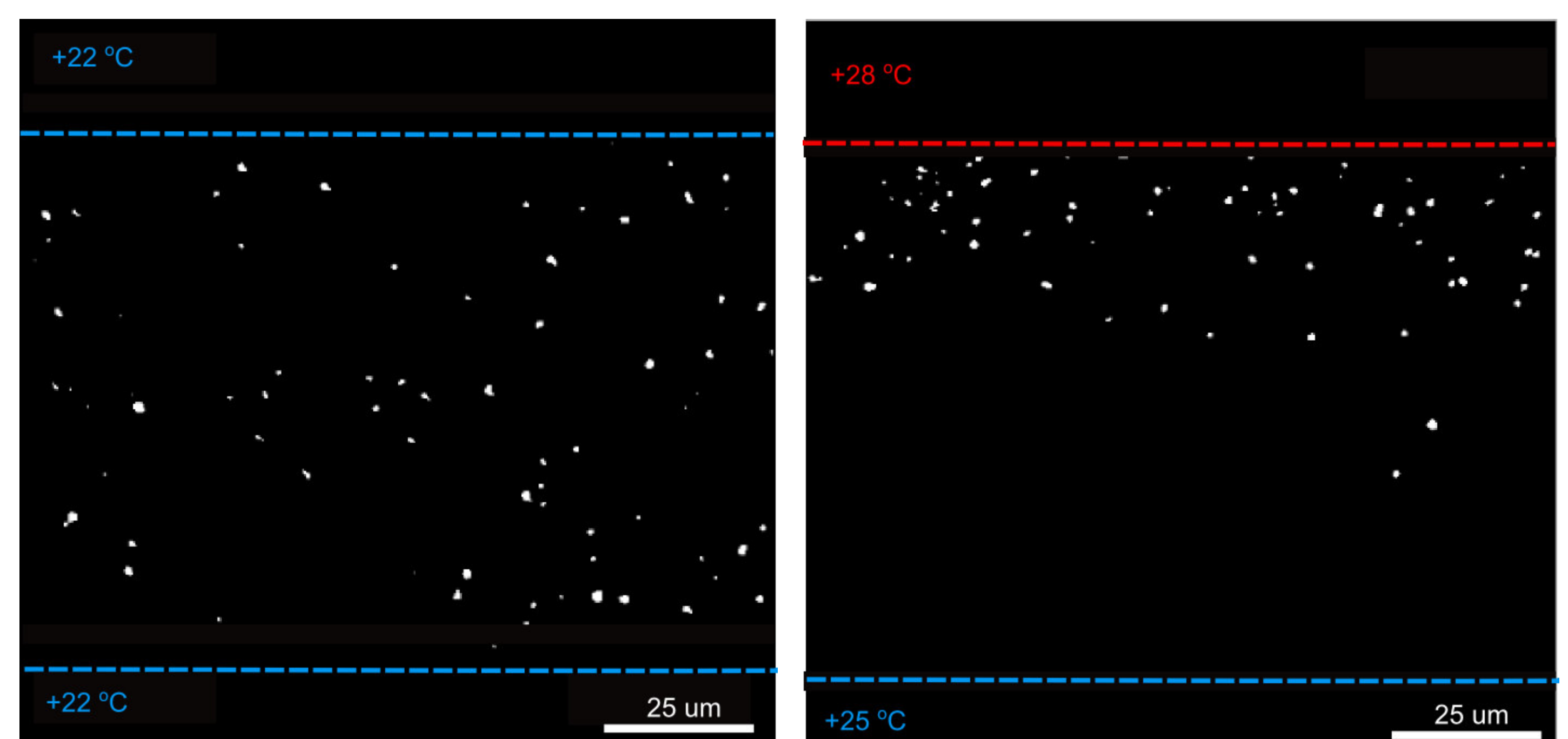


Large particles

Latex microbeads (0.5 μm) 0.01% in water

Thermo Scientific™ Fluoro-Max green fluorescent internally dyed polystyrene particles in water.

Single particles can be distinguished in the microscope and counted



At the equilibrium when $c \ll 1$:

$$\vec{J} = -D\vec{\nabla}c - cD_T\vec{\nabla}T = 0$$

$$\downarrow$$

$$c = c_0 \exp\left(\frac{S_T \Delta T}{L} x\right)$$

$$S_T \approx -3.0\text{ K}^{-1}$$

fitting

