

## Fast Physical-Optics Modeling of Microscopy System with Structured Illumination

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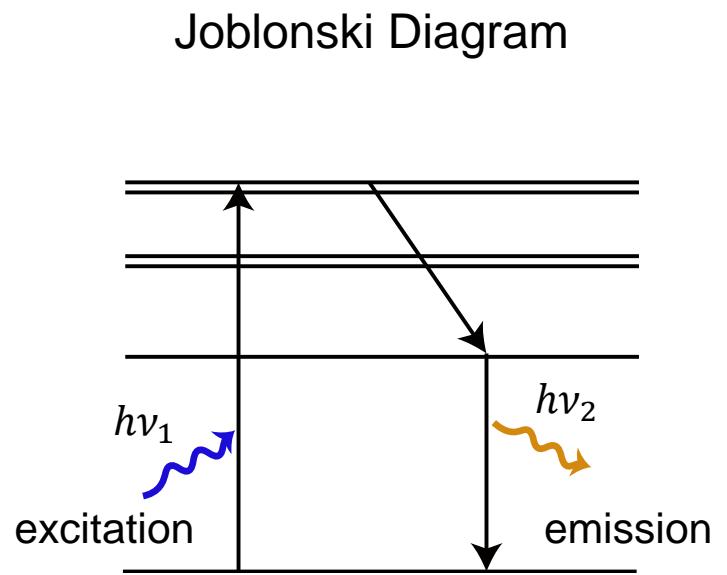
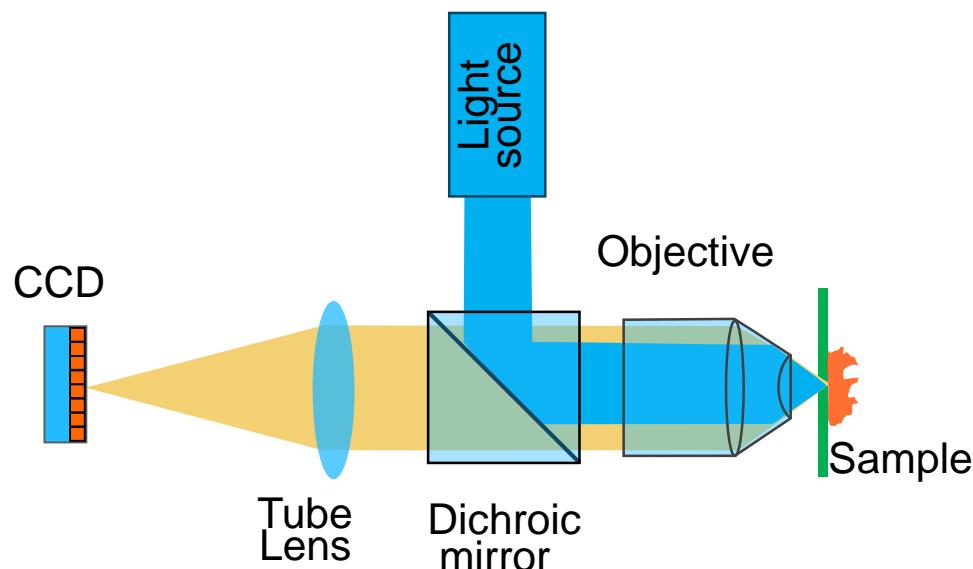
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<sup>3</sup> Wyrowski Photonics UG, Jena, Germany,

<sup>4</sup> Biomedical Imaging Group, Leibniz Institute of Photonic Technology (IPHT), Jena, Germany

## Background

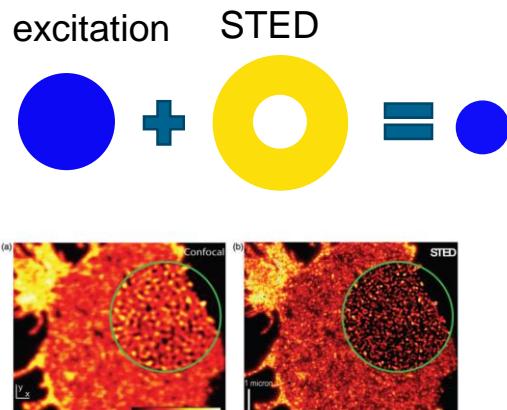
- One-photon fluorescence microscopy



# Background: Higher Resolution

## STED

[Hell et al., **Opt. Lett.** (1994)]

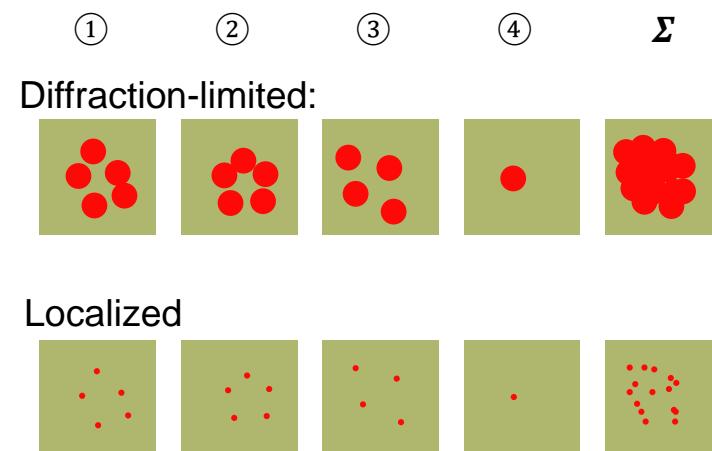


[Heintzmann et al., **Brief Funct Genomic Proteomic**(2006)]

- resolution: ~20 nm
- high power

## STORM

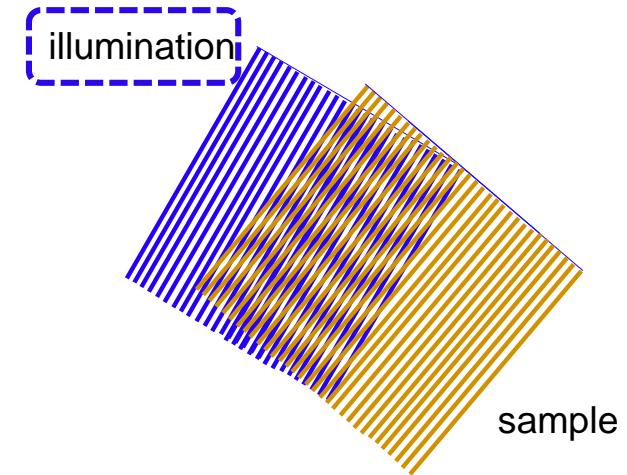
[Betzig et al., **Science** (2006)]



- resolution: ~20 nm
- low speed

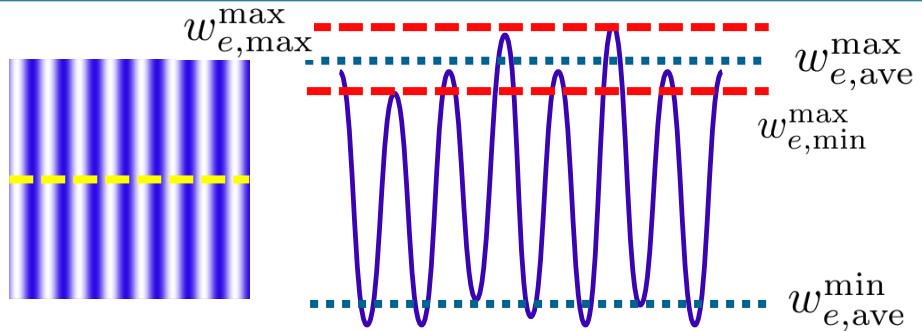
## SIM

[Heintzmann et al., **Proc. SPIE** 1998]  
[Gustafsson, **J. Microsc**(2000)]



- resolution: ~80 nm
- low power and high speed

## Motivation and Configuration



- Electric energy density

$$w_e \propto \| \mathbf{E} \|^2$$

- Contrast:

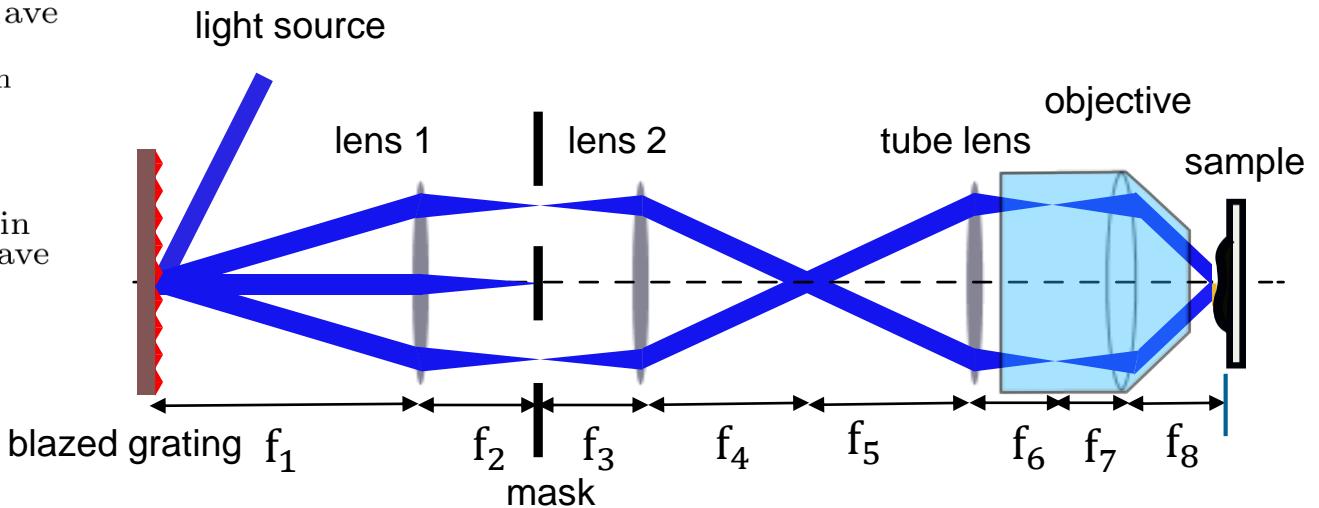
$$c = \frac{w_{e,\text{ave}}^{\text{max}} - w_{e,\text{ave}}^{\text{min}}}{w_{e,\text{ave}}^{\text{max}} + w_{e,\text{ave}}^{\text{min}}}$$

Best:  $c = 1$

- Homogeneity:

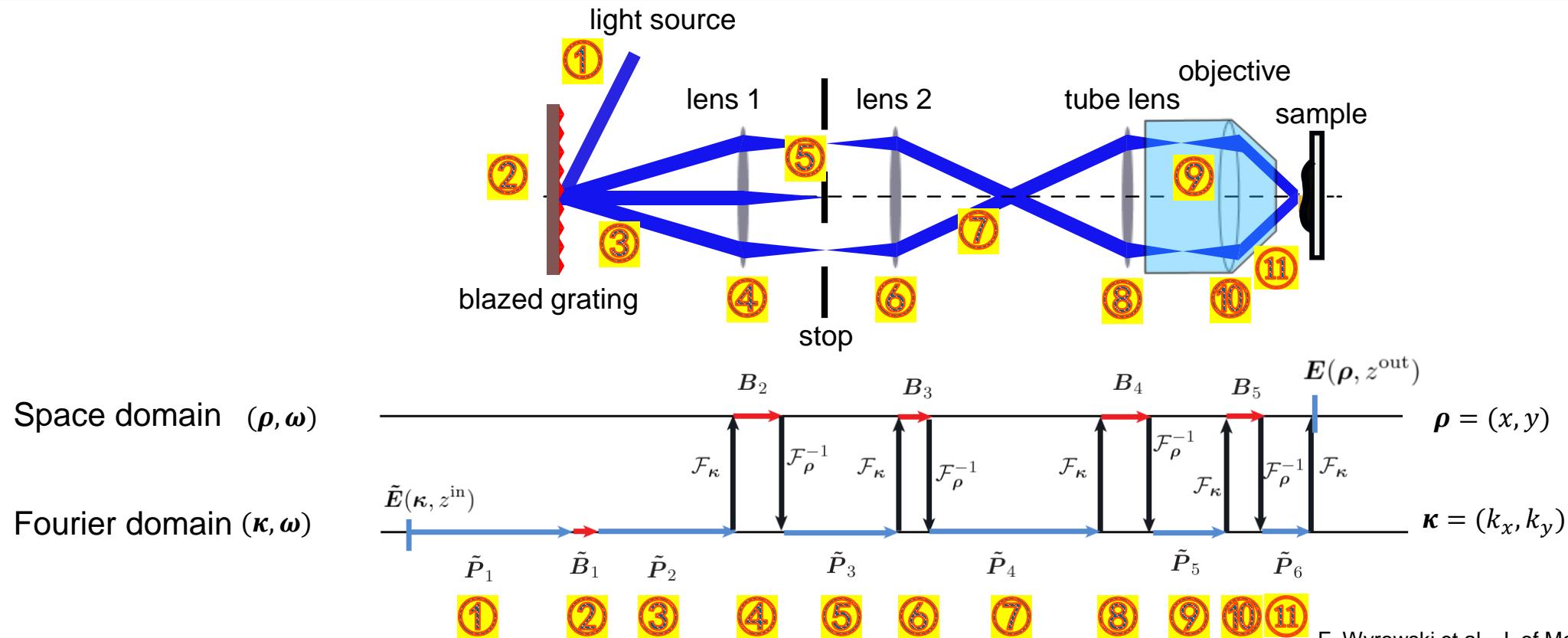
$$\sigma = \frac{w_{e,\max}^{\text{max}} - w_{e,\min}^{\text{max}}}{w_{e,\max}^{\text{max}} + w_{e,\min}^{\text{max}}}$$

Best:  $\sigma = 0$



- Polarization
- Diffraction from aperture
- Inclined illumination on blazed grating

# Theory: Field Tracing



Free space propagation

Fourier Modal Method(FMM)

Local Plane Interface Approximation(LPIA)

(1) (3) (5) (7) (9) (11)

(2)

(4) (6) (8) (10)

F. Wyrowski et al., J. of Modern Optics (2011)

S. Zhang et al. Appl. Opt. (2016)

M. G. Moharam et al. Opt. Soc. Am. A (1995)

F. Wyrowski and C. Hellmann.. Proc. DGaO (2017)

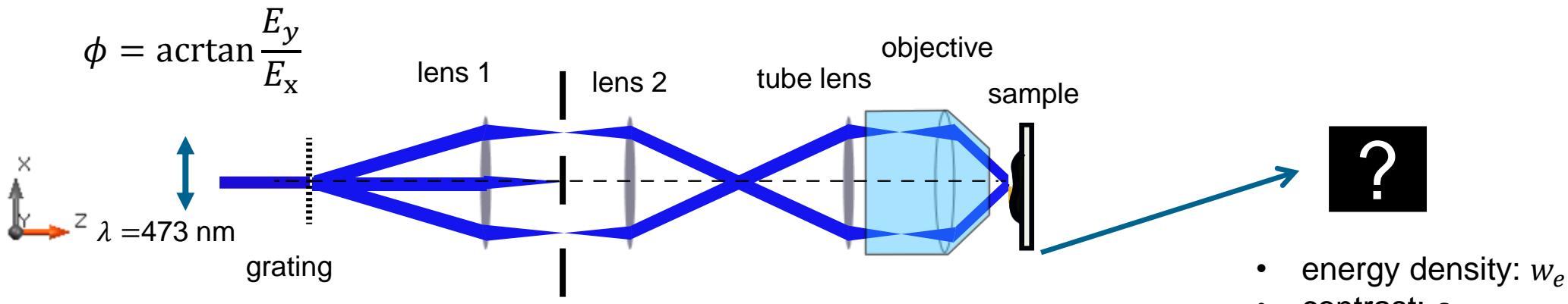
A. Pfeil et al., Appl. Opt. (2000)

The concept of **bidirectional operators** and its application to the modelling of microstructures Paper 10694-15, Prof.Frank Wyrowski

## **Simulation Results via VirtualLab Fusion**

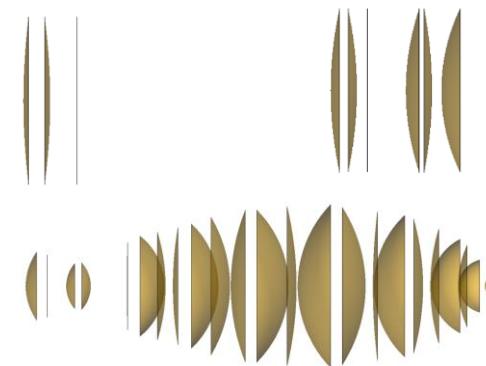
# Polarization

## Modeling Task

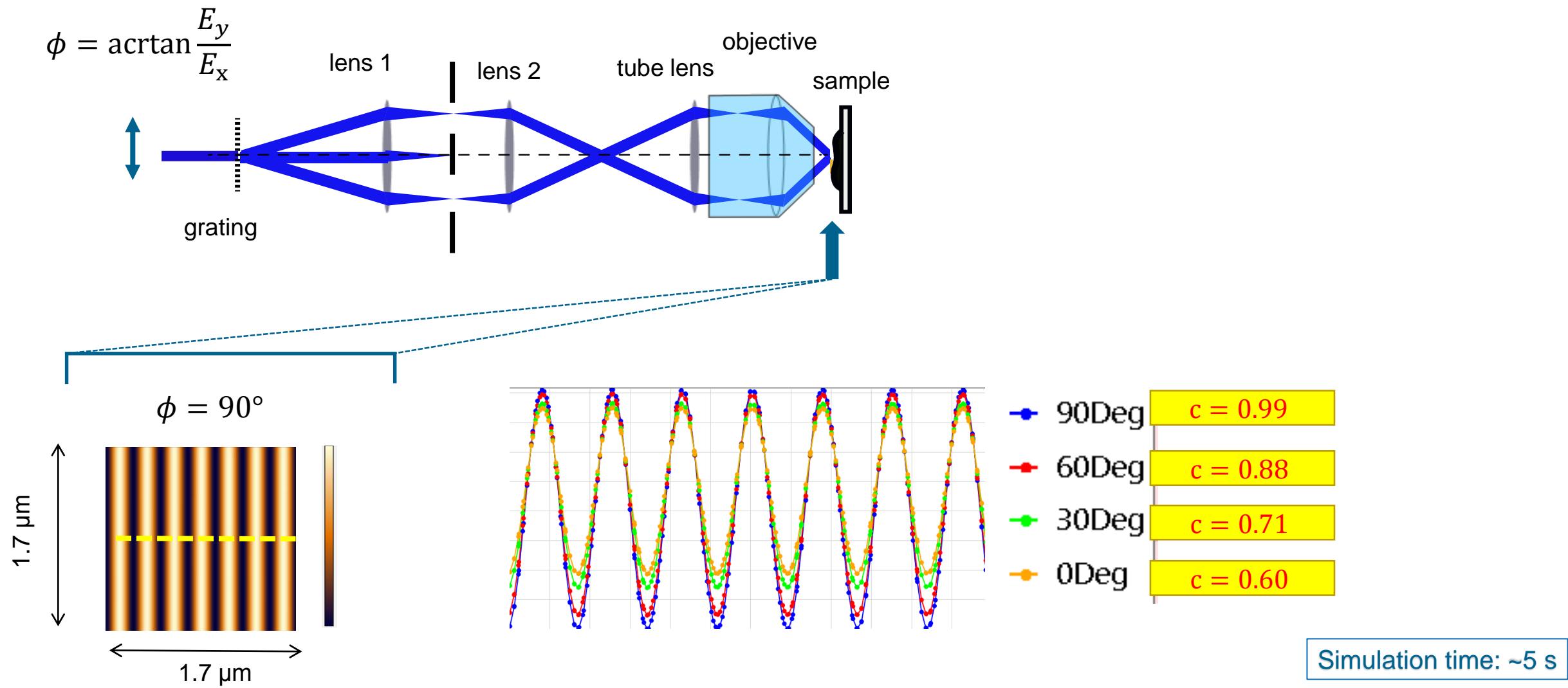


- energy density:  $w_e$
- contrast: c

Lens	Property
lens 1, 2	Thorlab AC254 double achromat
Tube lens	Nikon 200 mm
Objectives	Nikon 60X, NA=1.4, Effective NA: ~1.12 apochromatic

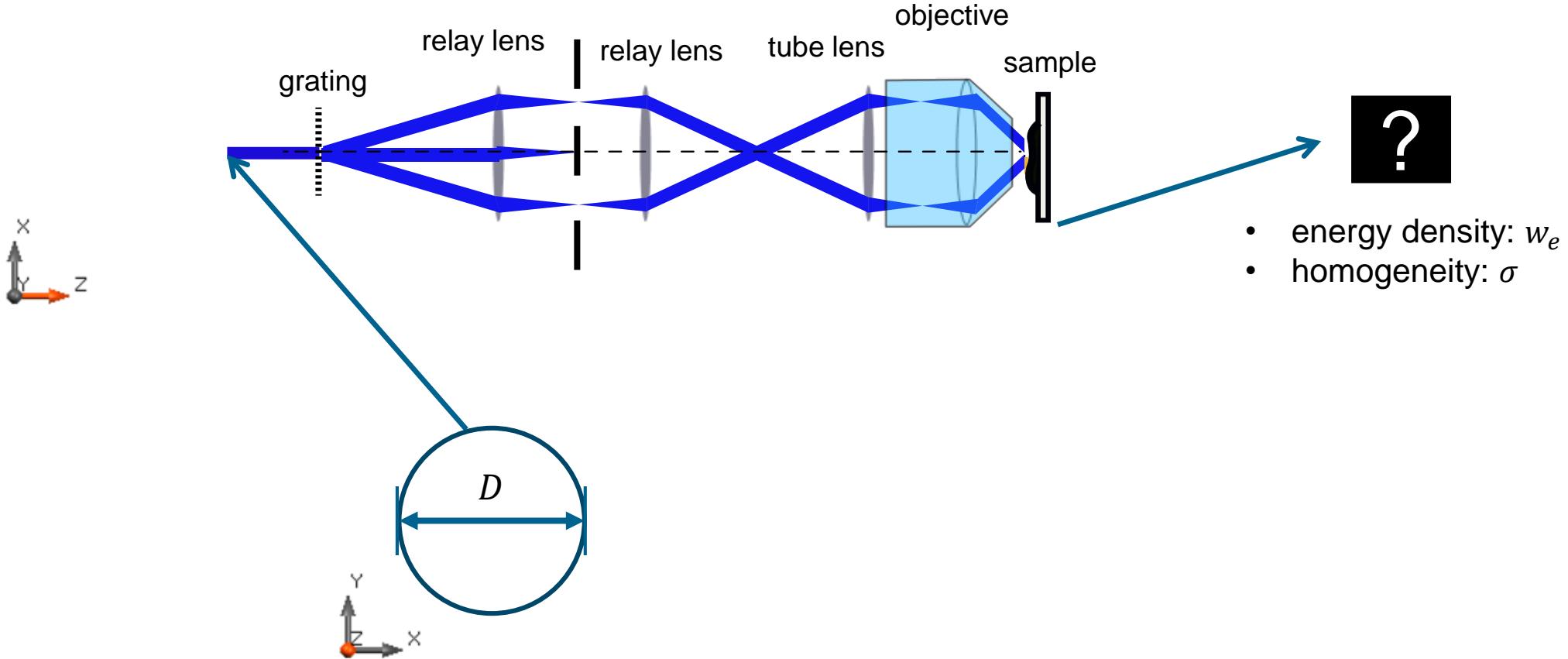


## Result: Energy Density

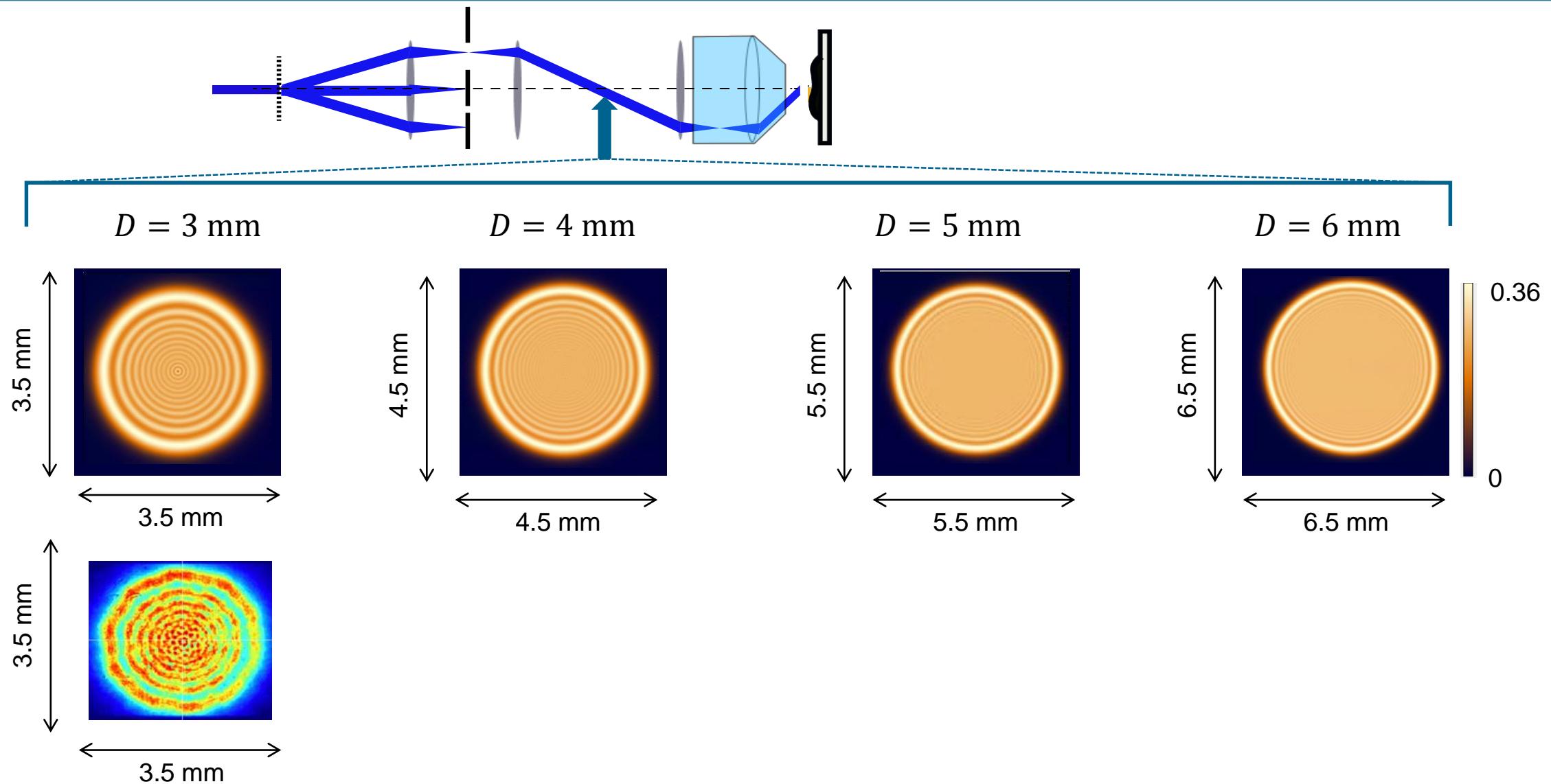


## **Diffraction from Aperture**

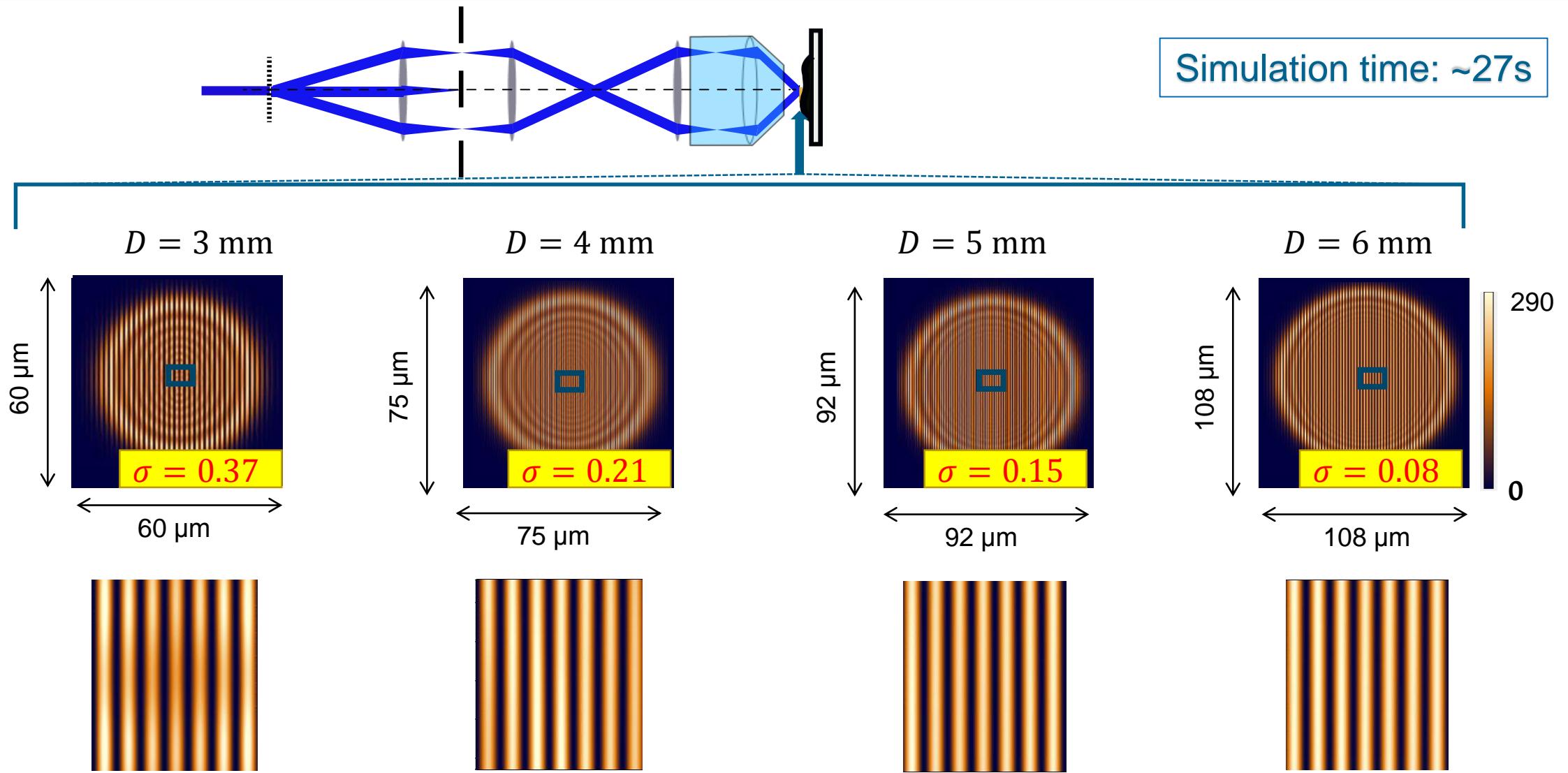
## Modeling Task



## Result: Energy Density

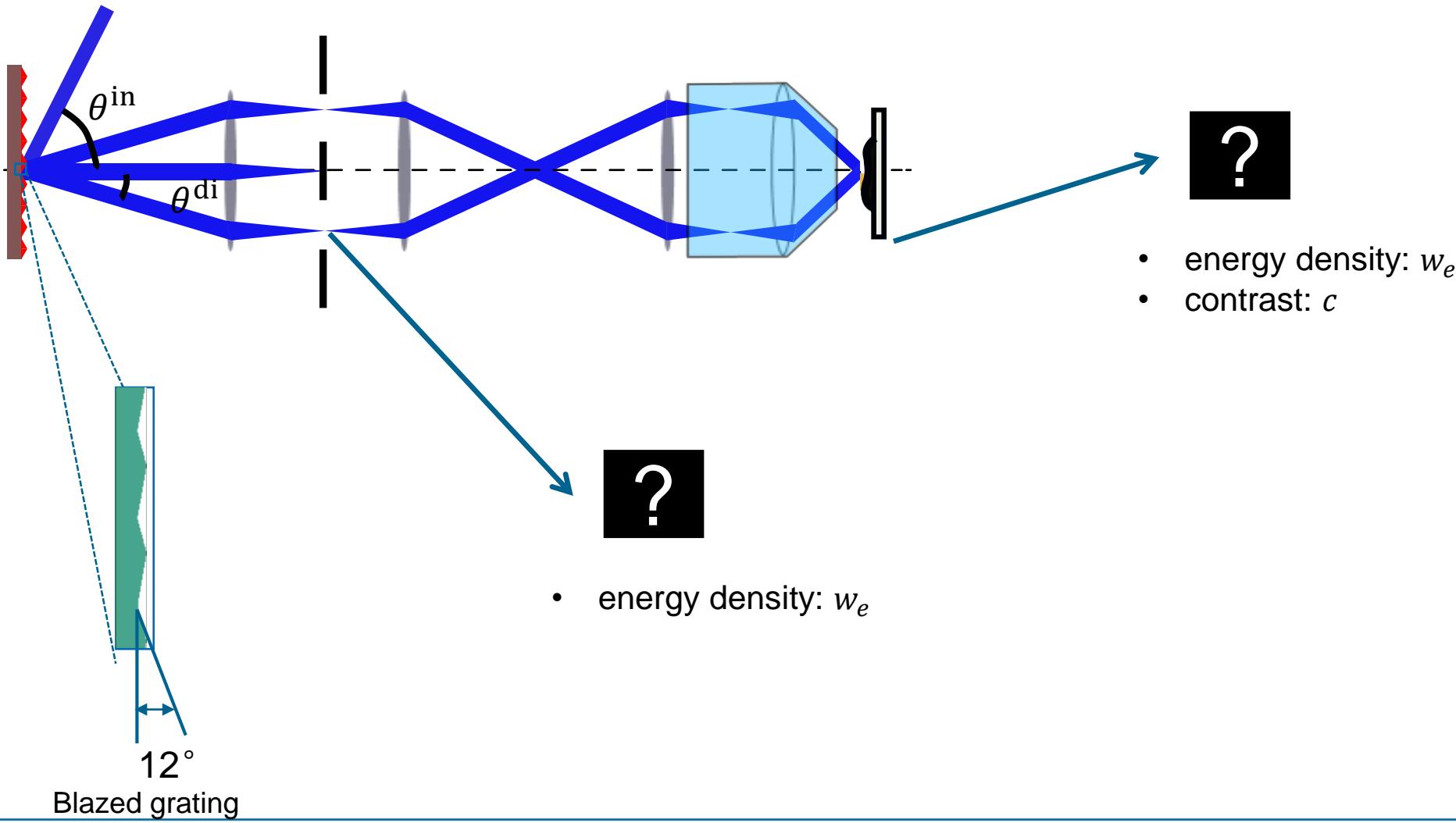


## Result: Energy Density and Homogeneity

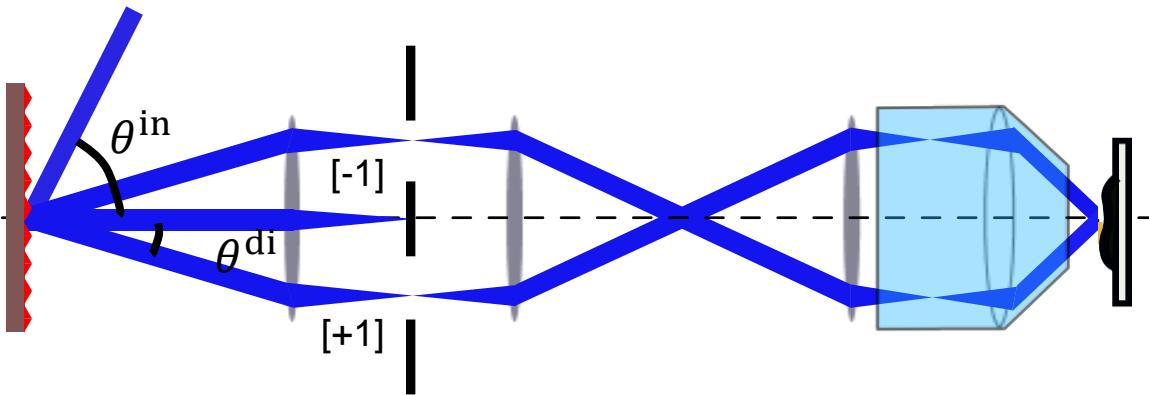


## **Inclined Illumination on Blazed Grating**

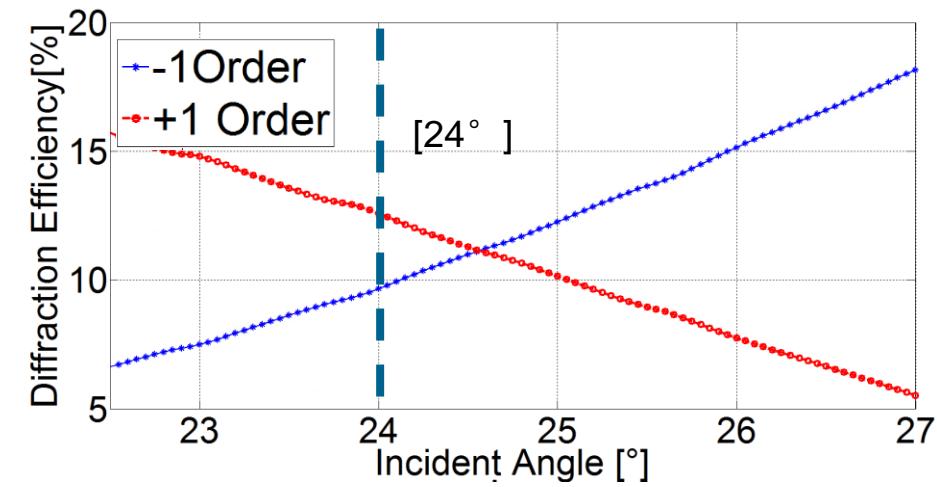
## Modeling Task



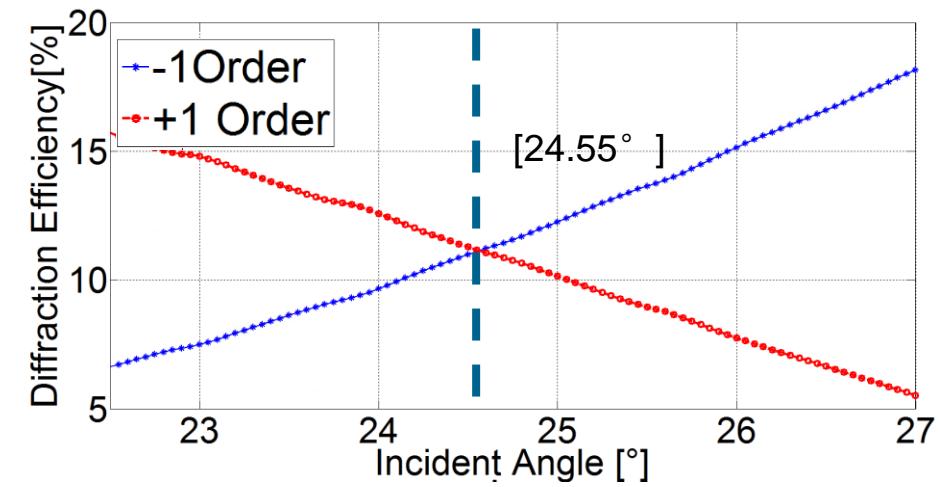
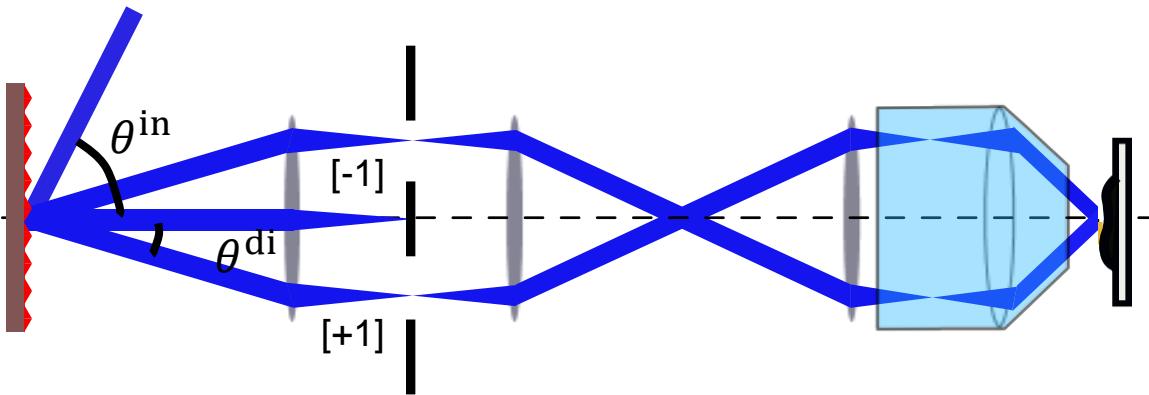
## Result: Diffraction Angle and Efficiency



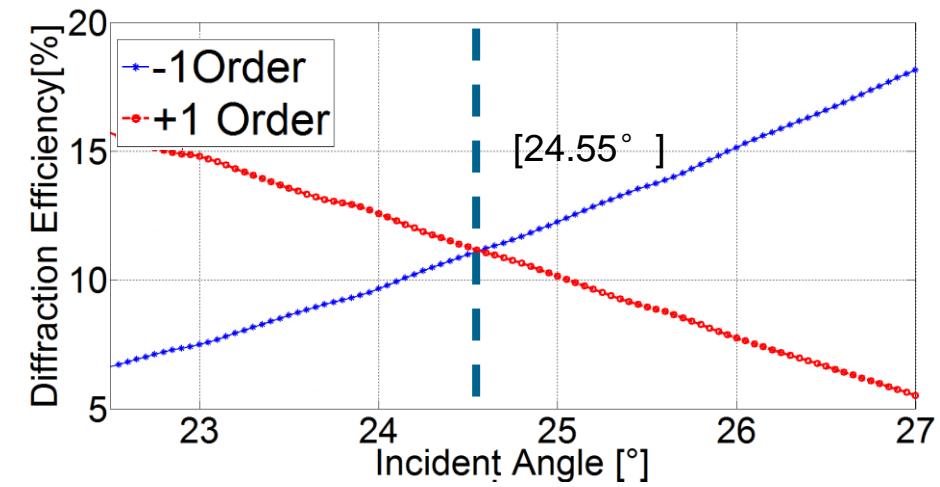
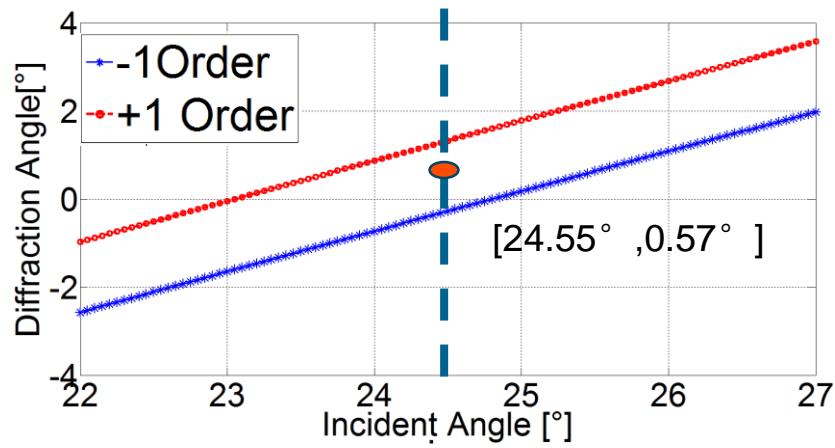
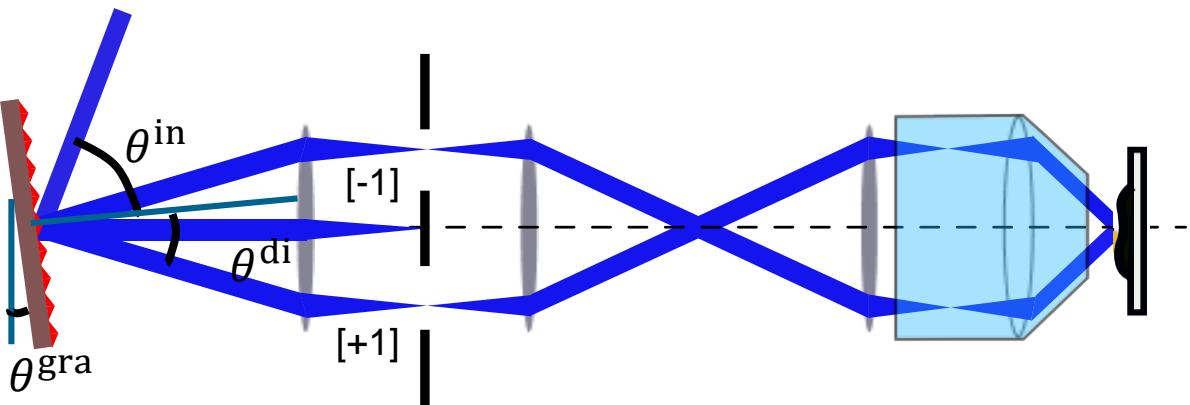
Calculated by FMM



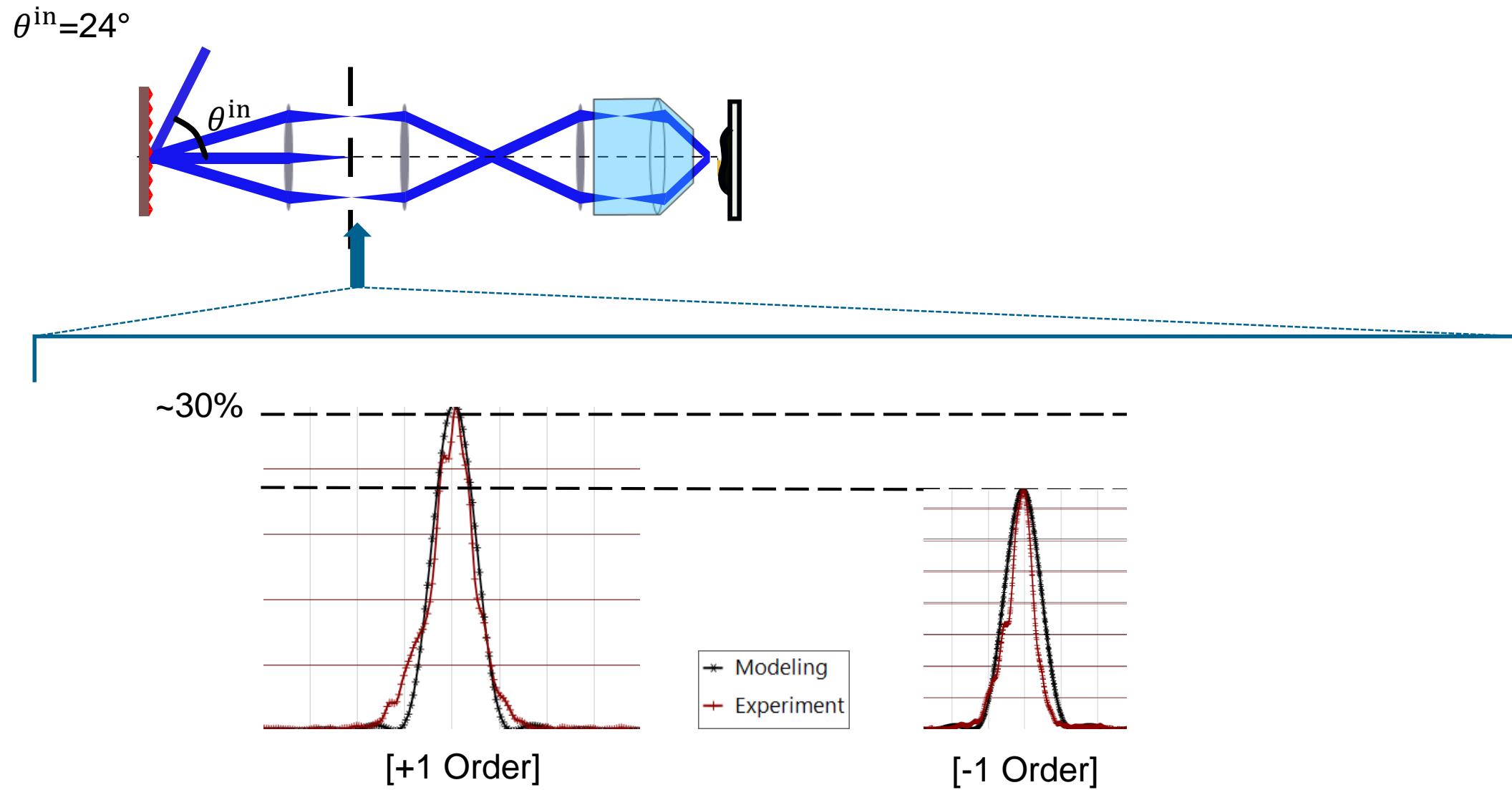
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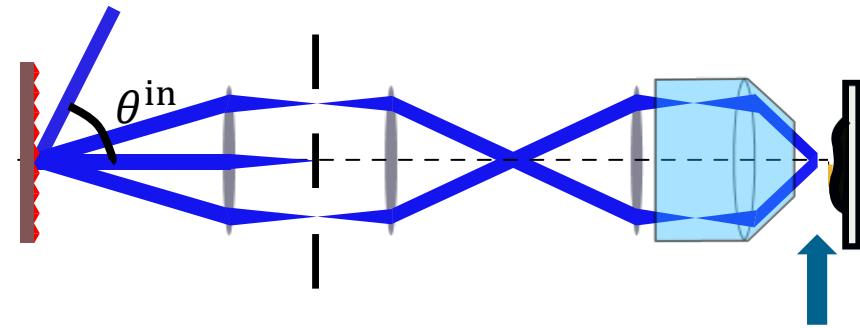


## Results: Energy Density

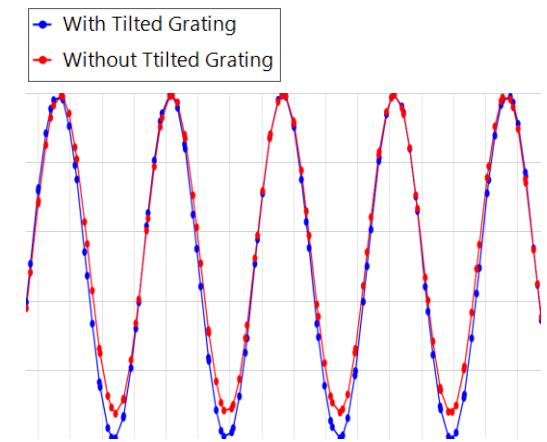
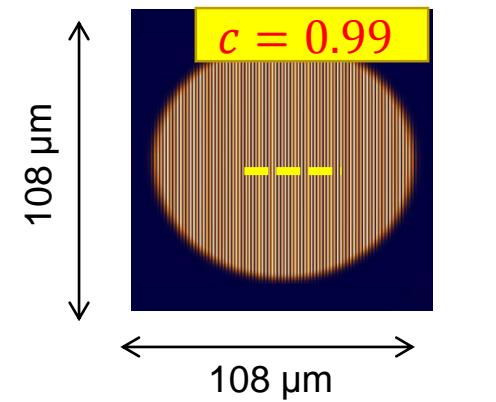
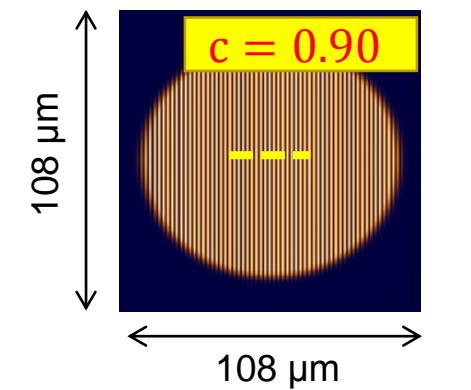
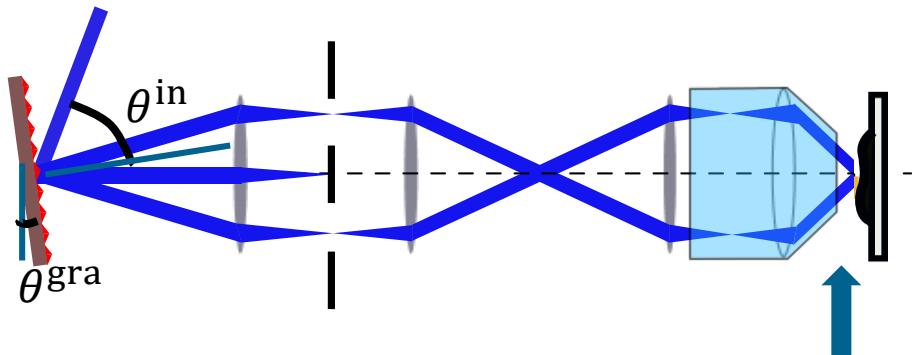


## Results: Energy Density

$$\theta^{\text{in}} = 24^\circ$$



$$[\theta^{\text{in}}, \theta^{\text{gra}}] = [24.55^\circ, 0.57^\circ]$$



## Conclusion and Outlook

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- The complexity of microscopy system with structured illumination makes it vulnerable to the undesired effects which causes the inhomogeneity and low contrast of the interference pattern.
- These effects should be analyzed and taken into account in the image reconstruction algorithm.
- In case of deep tissue imaging, adaptive optics can be applied further to compensate the undesired effects.

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Thanks!