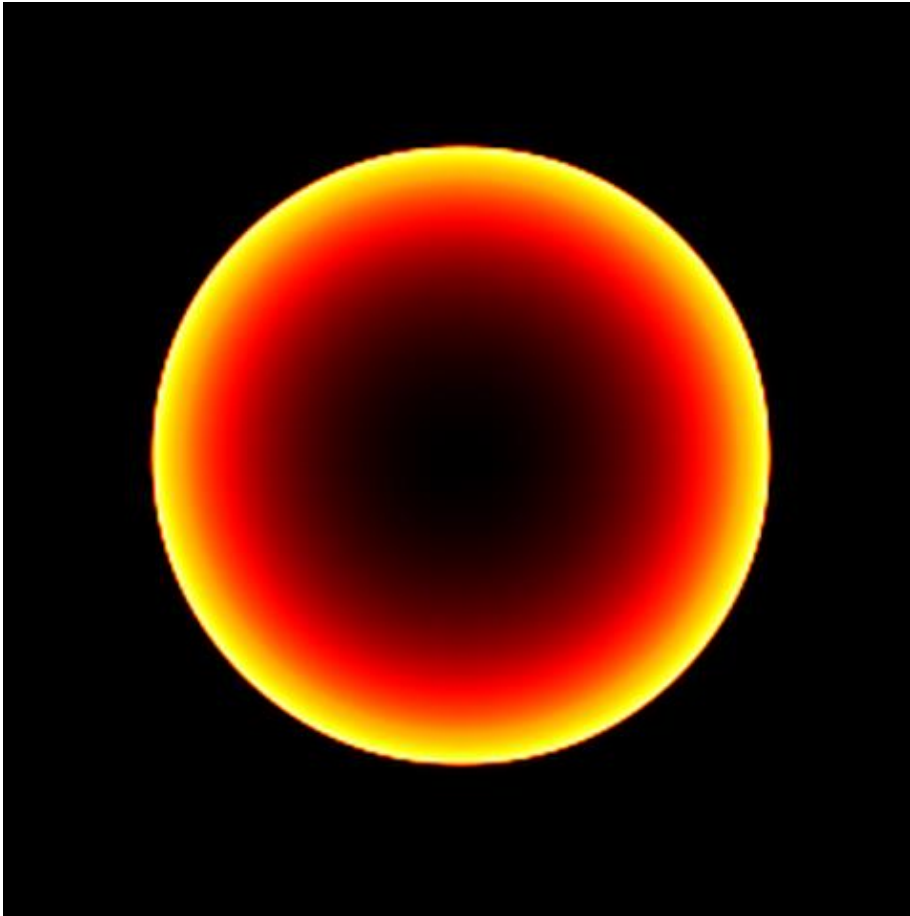


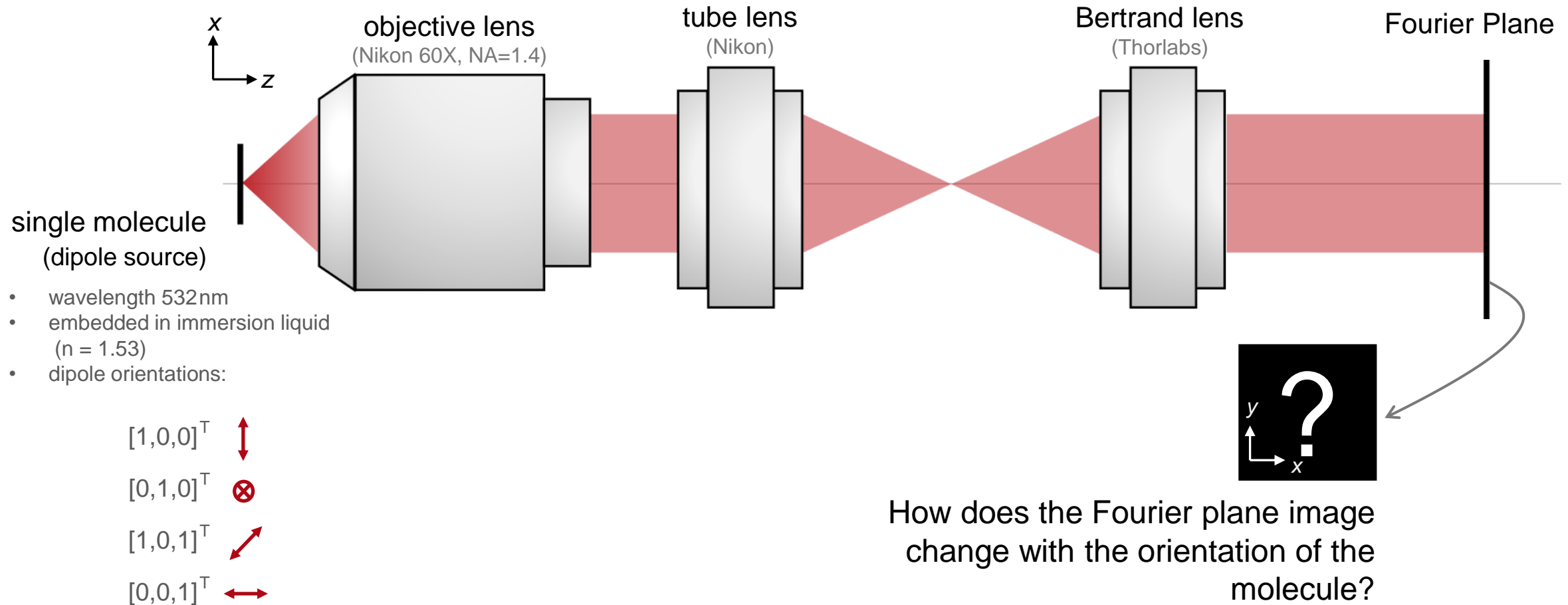
# Single-Molecule Imaging with High-NA Fourier Microscope

# Abstract

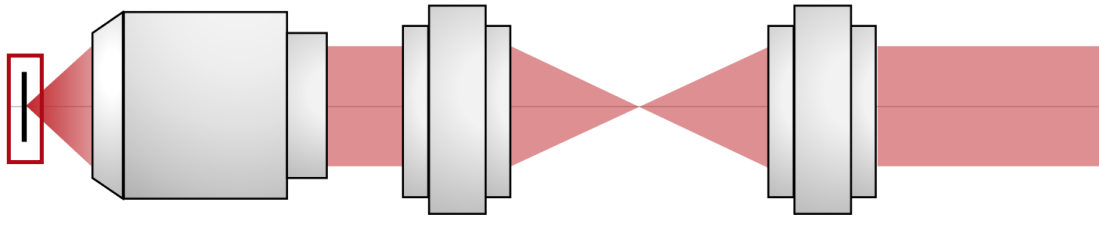


Fourier microscopy is widely used for single-molecule imaging, surface plasma observation, photonic crystal imaging, etc. It enables the direct observation of the spatial frequency distribution. Different effects in the high-NA Fourier microscope (angle-dependent Fresnel losses at each lens surface, diffraction, etc.) can affect the final image quality obtained for the single molecule. The fast physical optics software VirtualLab Fusion can model the entire system with its powerful Field Tracing engine, including the Fresnel losses and aperture diffraction effects. An example is presented, and we compare the simulation results with experimental results from literature.

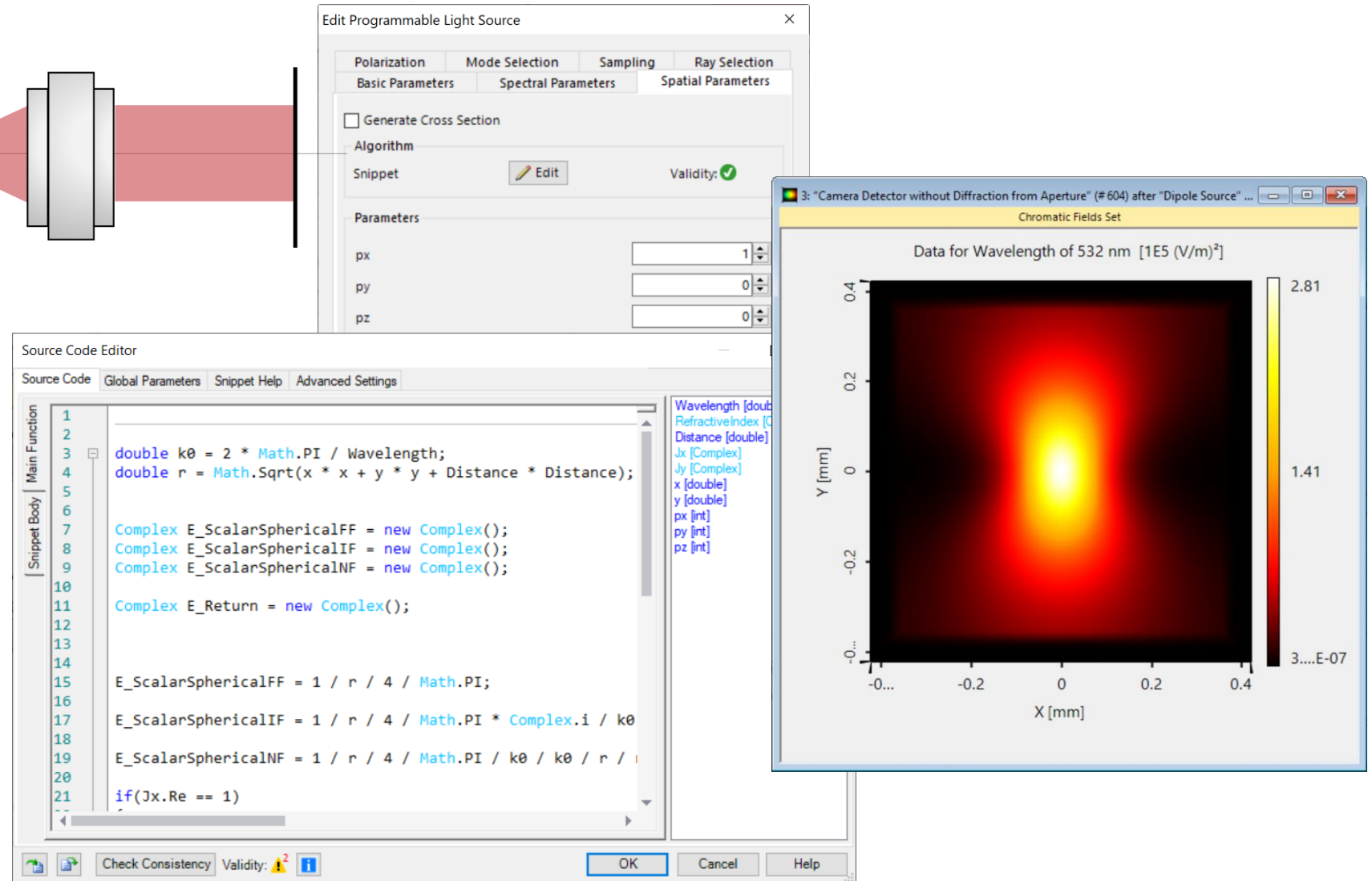
# Modeling Task



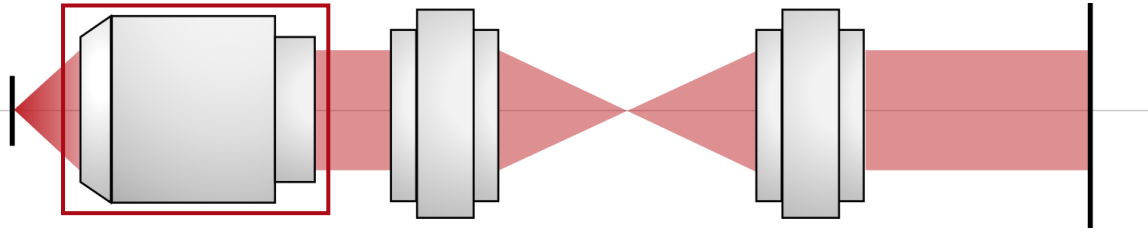
# System Building Blocks: Dipole Source



- The *Programmable Light Source* allows for the specification of an arbitrary lateral field distribution. In our case, we specify the field generated by a dipole.
- The dipole source emits a locally polarized field (meaning that the spatial profile of the  $E_x$  and  $E_y$  components at the source plane is fundamentally different and therefore cannot be expressed with a single function).
- In order to accurately model the polarization characteristics, we employ a *Multiple Light Source*, that allows us to define the different profiles for the different components.

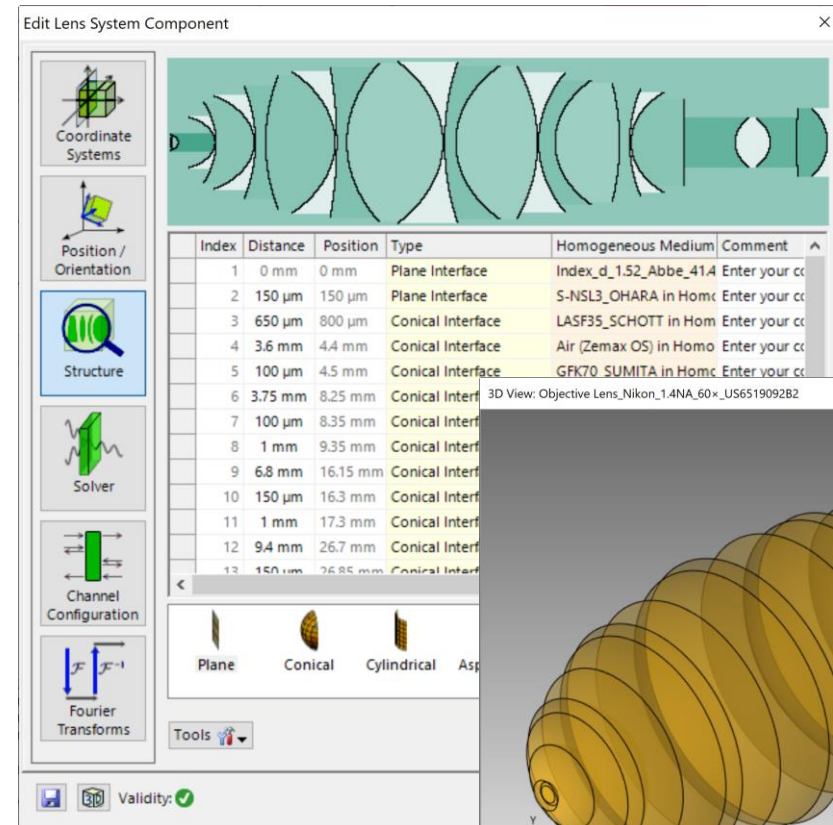


# System Building Blocks: Objective Lens

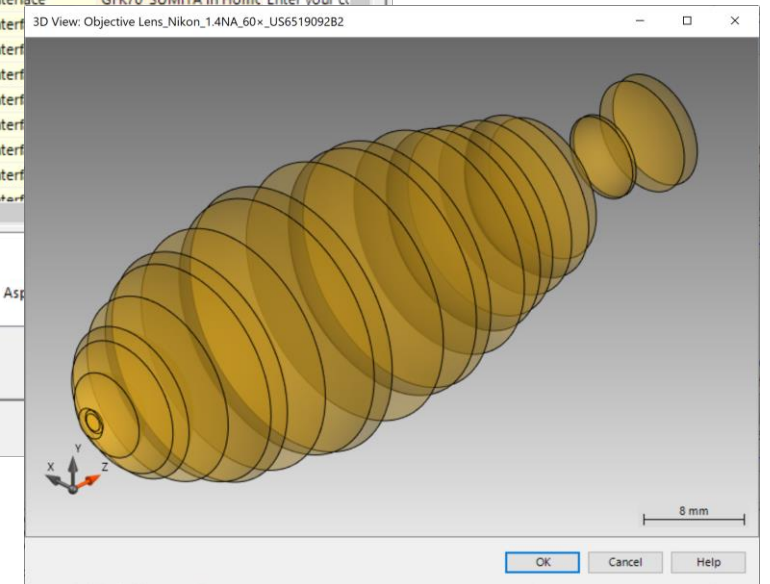


Objective lenses, such as the one used in this system, are usually quite complex structures, containing many interfaces and dispersive materials.

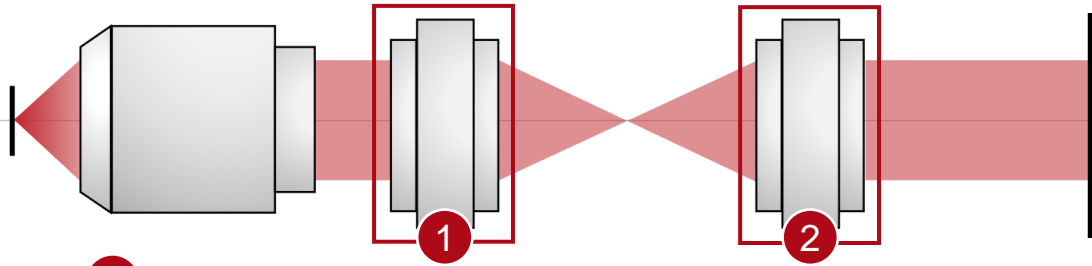
In VirtualLab Fusion, this can be modeled using the *Lens System Component*. There, the optical engineer can build up a component from a sequence of interfaces and materials.



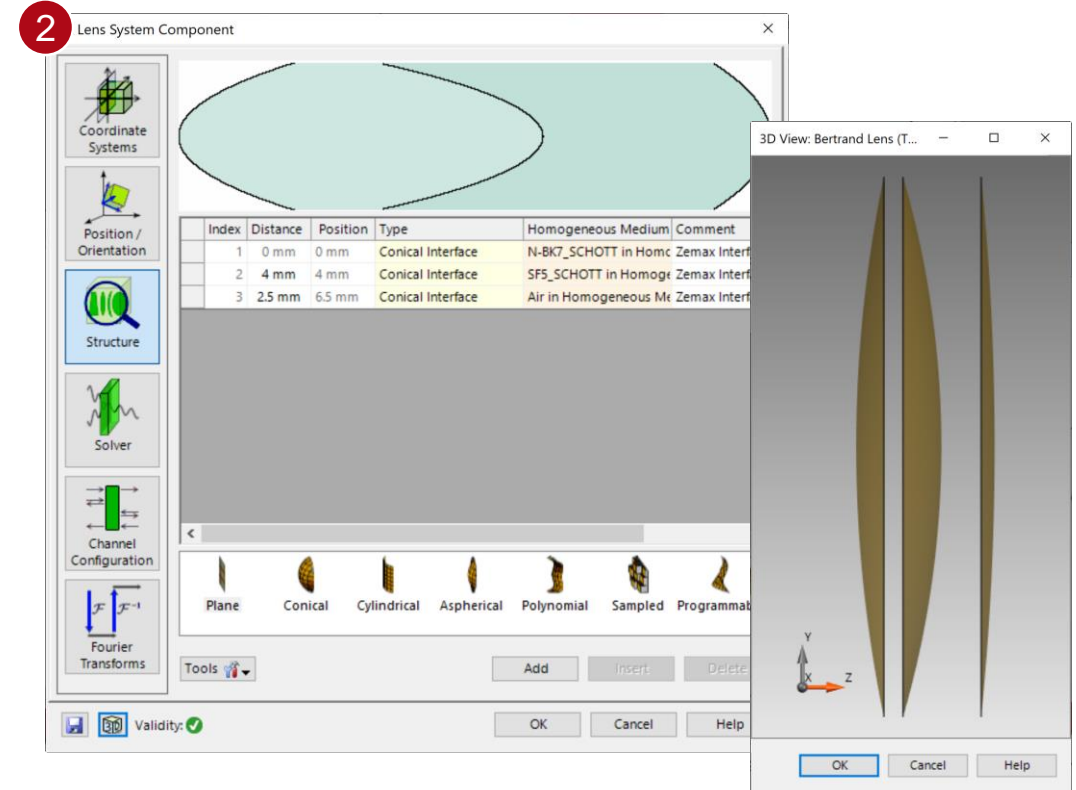
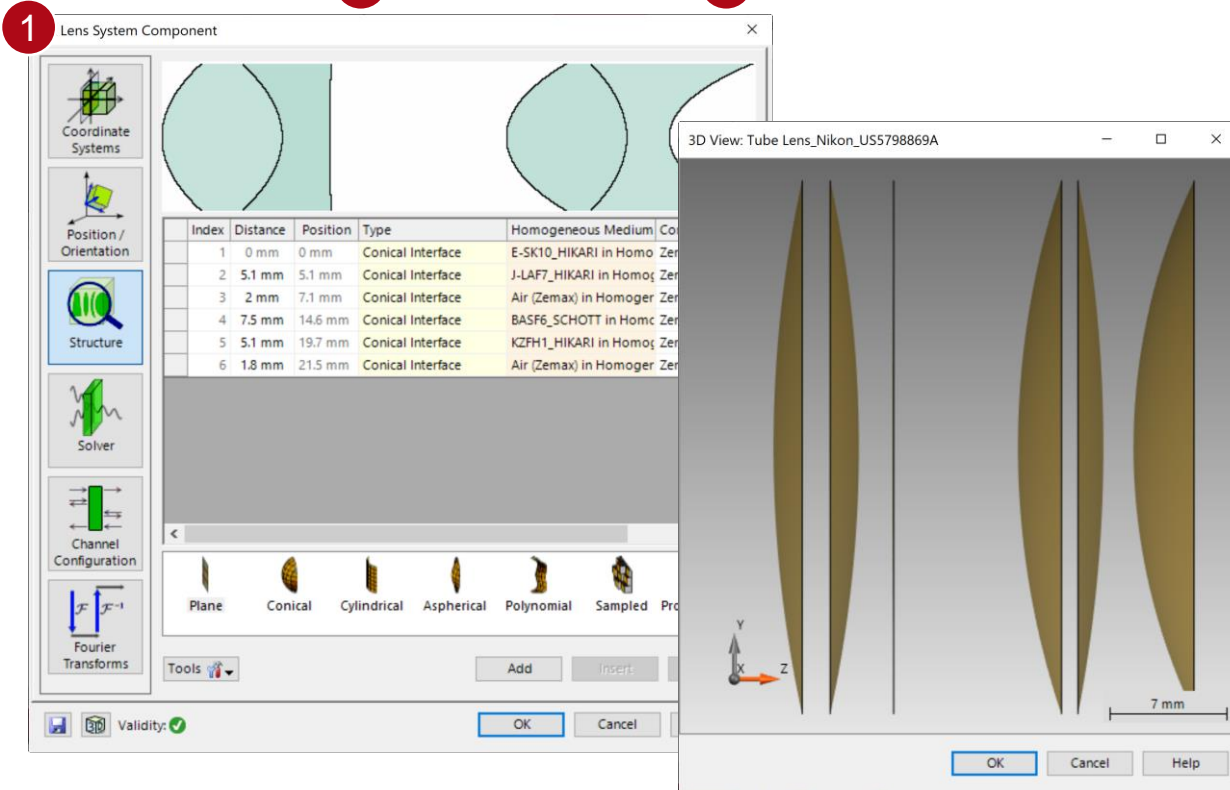
*Note: An immersion liquid is used to fill the gap between source and objective. Hence the surrounding material of the lenses is the immersion liquid.*



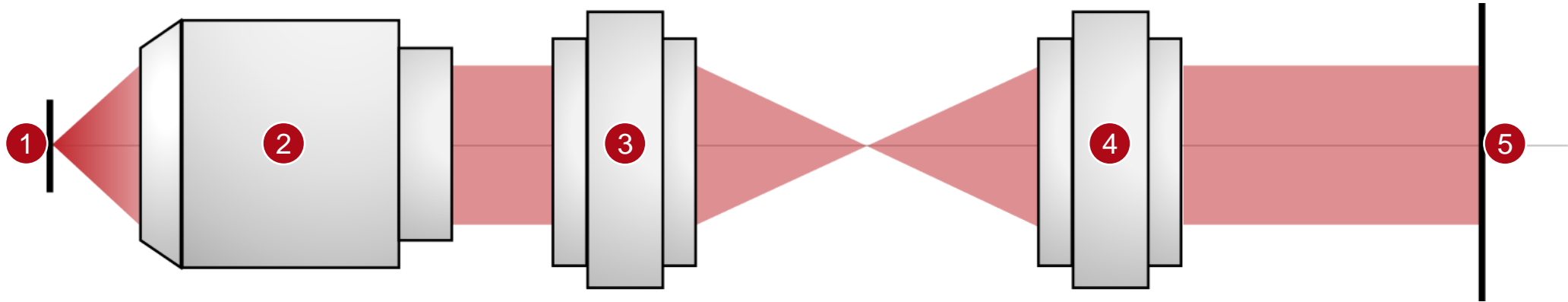
# System Building Blocks: Tube & Bertrand Lenses



Tube lenses and Bertrand lenses are commonly used in microscopes to reproduce the image from the objective lens to the eye-piece. They can be imported into VirtualLab Fusion. We use the same modeling strategy as for the objective lens.



# Summary of Model

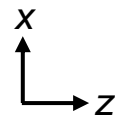


Optical System	Elements in VirtualLab Fusion	Model/Solver/Detected Value
1. dipole source	<i>Programmable &amp; Multiple Light Source</i>	lateral field distribution
2. objective lens	<i>Lens System Component</i>	Local Plane Interface Approximation
3. tube lens	<i>Lens System Component</i>	Local Plane Interface Approximation
4. Bertrand lens	<i>Lens System Component</i>	Local Plane Interface Approximation
5. detector	<i>Camera Detector</i>	energy density measurement



# Image at the Fourier Plane

Dipole moment  
 $[p_x, p_y, p_z]^T$



$$[1, 0, 0]^T$$



$$[0, 1, 0]^T$$



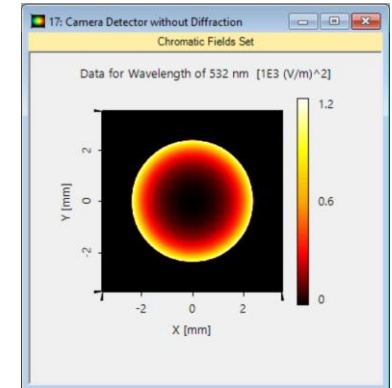
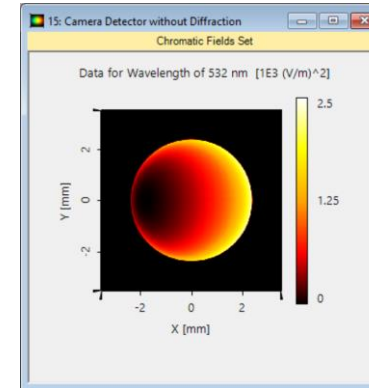
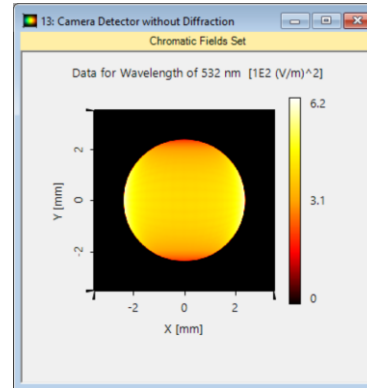
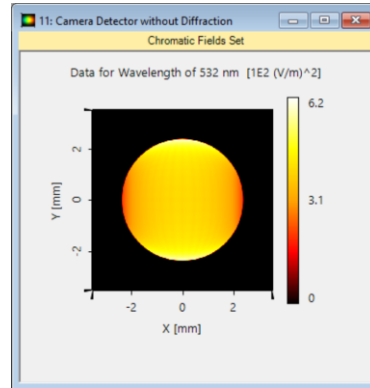
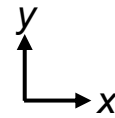
$$[1, 0, 1]^T$$



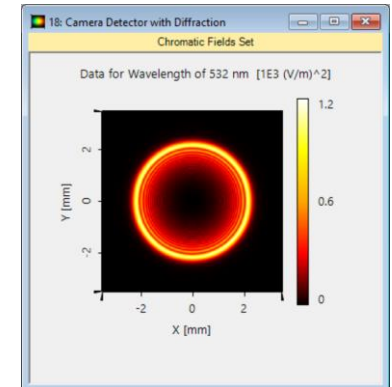
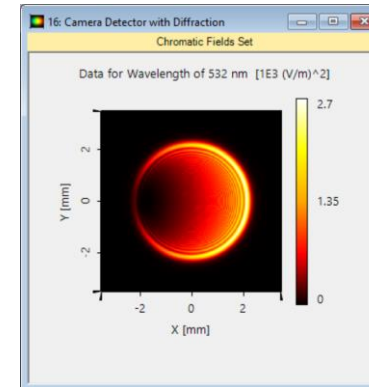
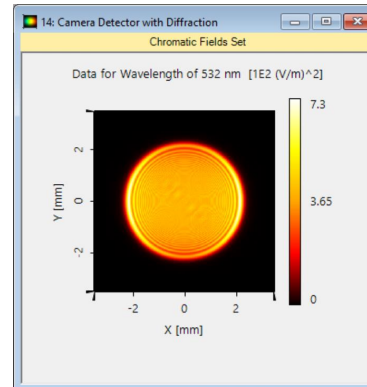
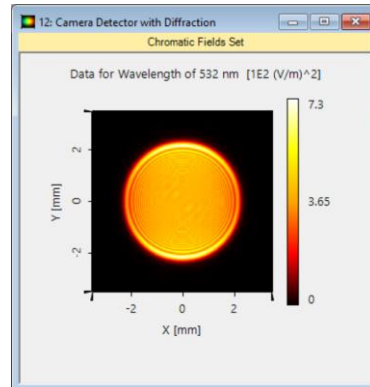
$$[0, 0, 1]^T$$



- without diffraction from aperture

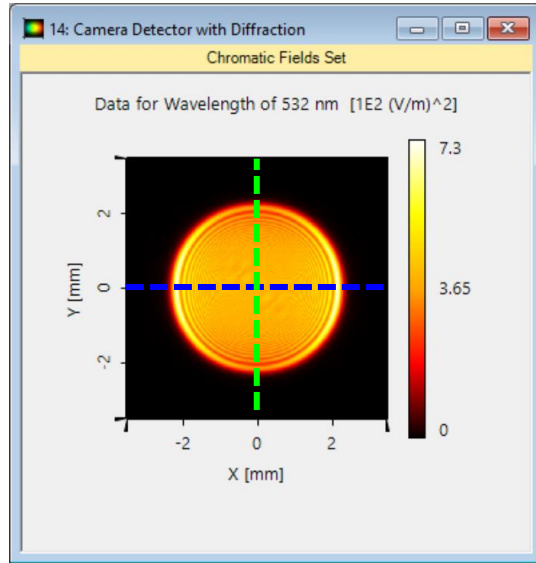


- with diffraction from aperture

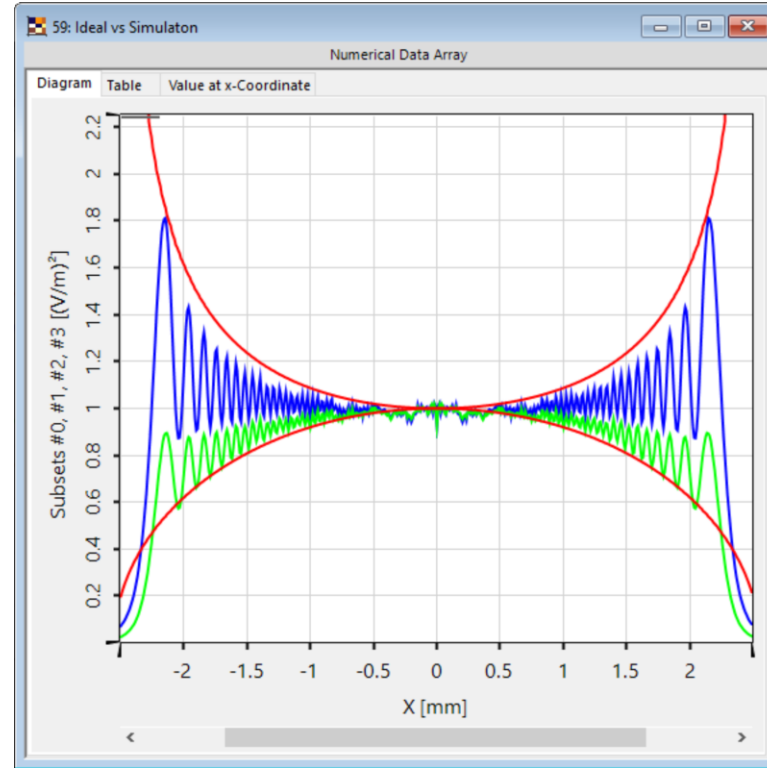




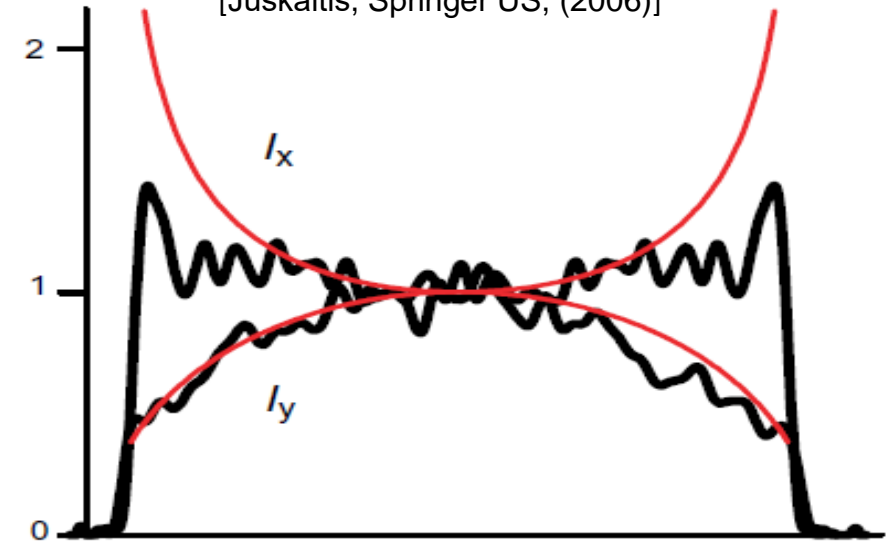
# Simulation Comparison for Orientation $[0,1,0]$



$[0,1,0]^T \otimes$

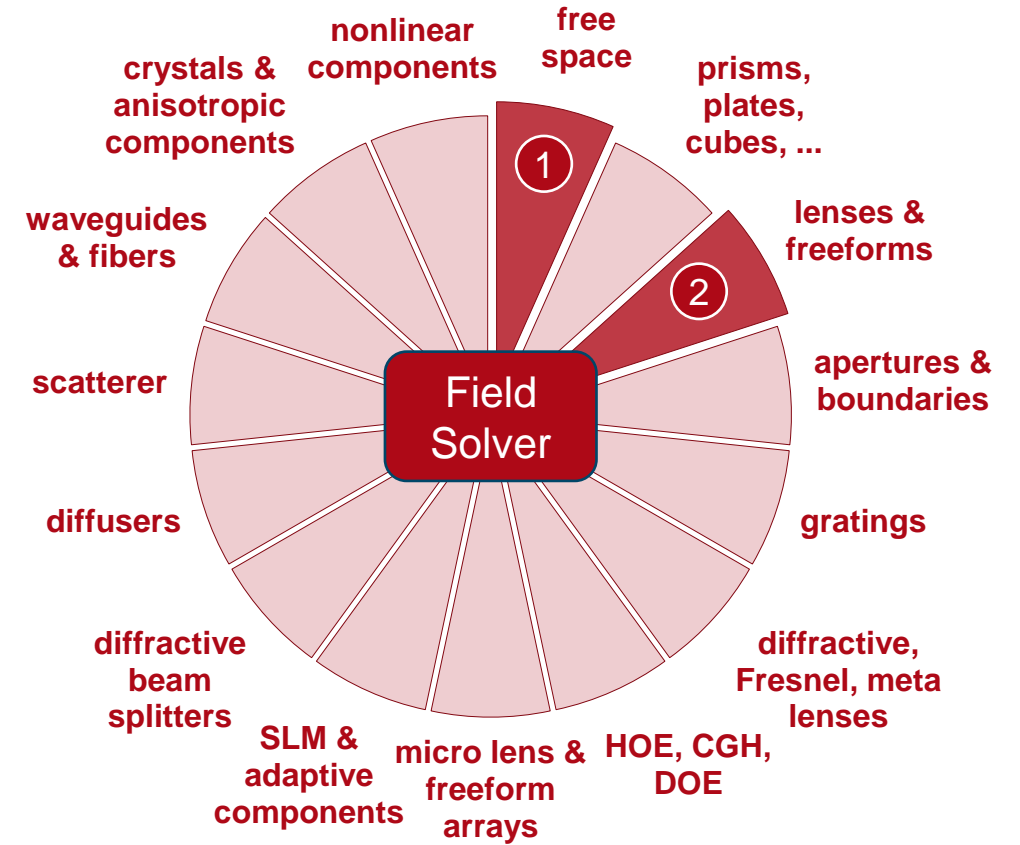
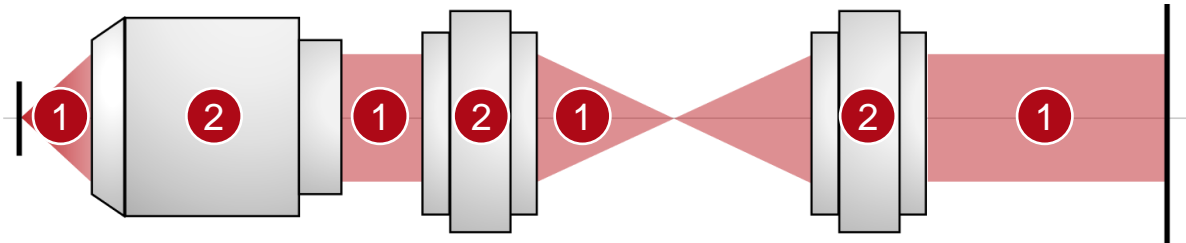


Reference  
[Juškaitis, Springer US, (2006)]



In order to further investigate the physical effects, we took dipole orientation  $[0,1,0]$  and compared the obtained results to an experimental measurement [Juškaitis, Springer US, (2006)]. The blue and green curves are taken from the corresponding 1D cross-sections of the simulation result. Cross-sectional references of an idealized case (diffraction neglected) are depicted in red. The data of the reference curves was calculated analytically by applying formulas given in the reference publication and finally imported into VirtualLab Fusion.

# VirtualLab Fusion Technologies



# Document Information

title	Single-Molecule Imaging with High-NA Fourier Microscope
document code	MIC.0008
document version	1.1
software edition	VirtualLab Fusion Basic
software version	2021.1 (Build 1.180)
category	Application Use Case
further reading	<ul style="list-style-type: none"><li>- <a href="#">Analyzing High-NA Objective Lens Focusing</a></li><li>- <a href="#">Resolution Investigation for Microscope Objective Lenses by Rayleigh Criterion</a></li><li>- <a href="#">Reflecting Microscope System with Very High Numerical Aperture</a></li></ul>