

# Hard X-Ray Scanning Microscopy with Fluorescence and Diffraction Contrast

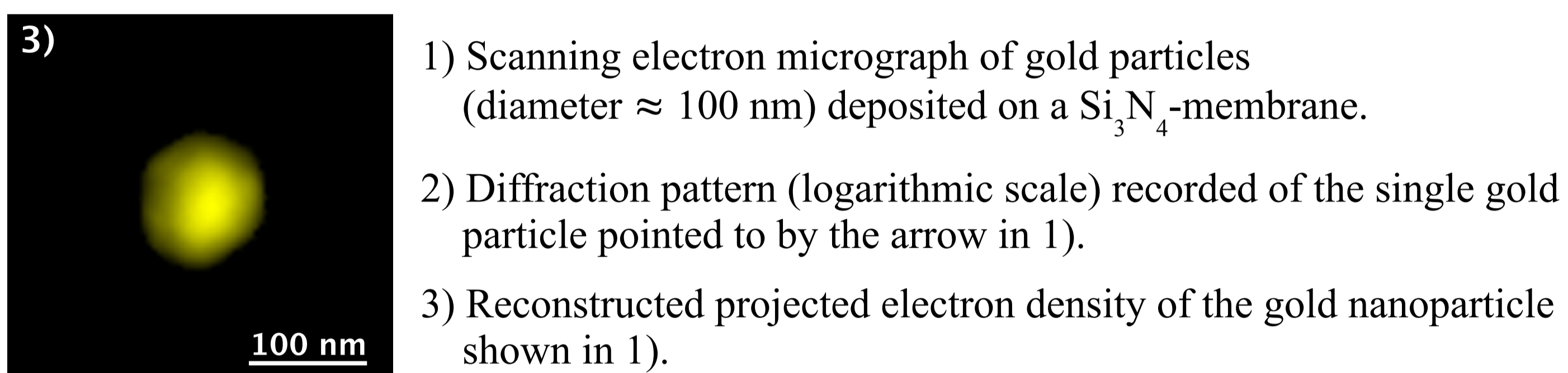
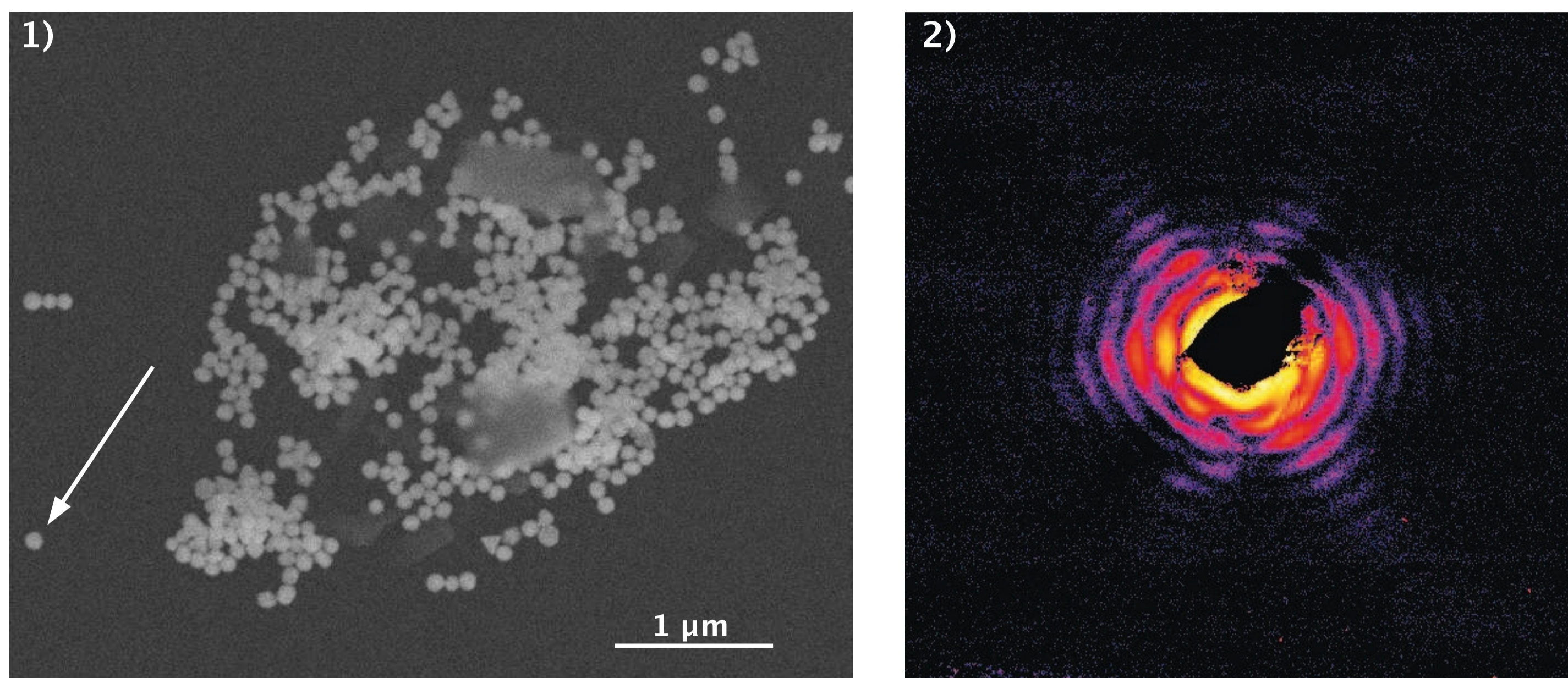


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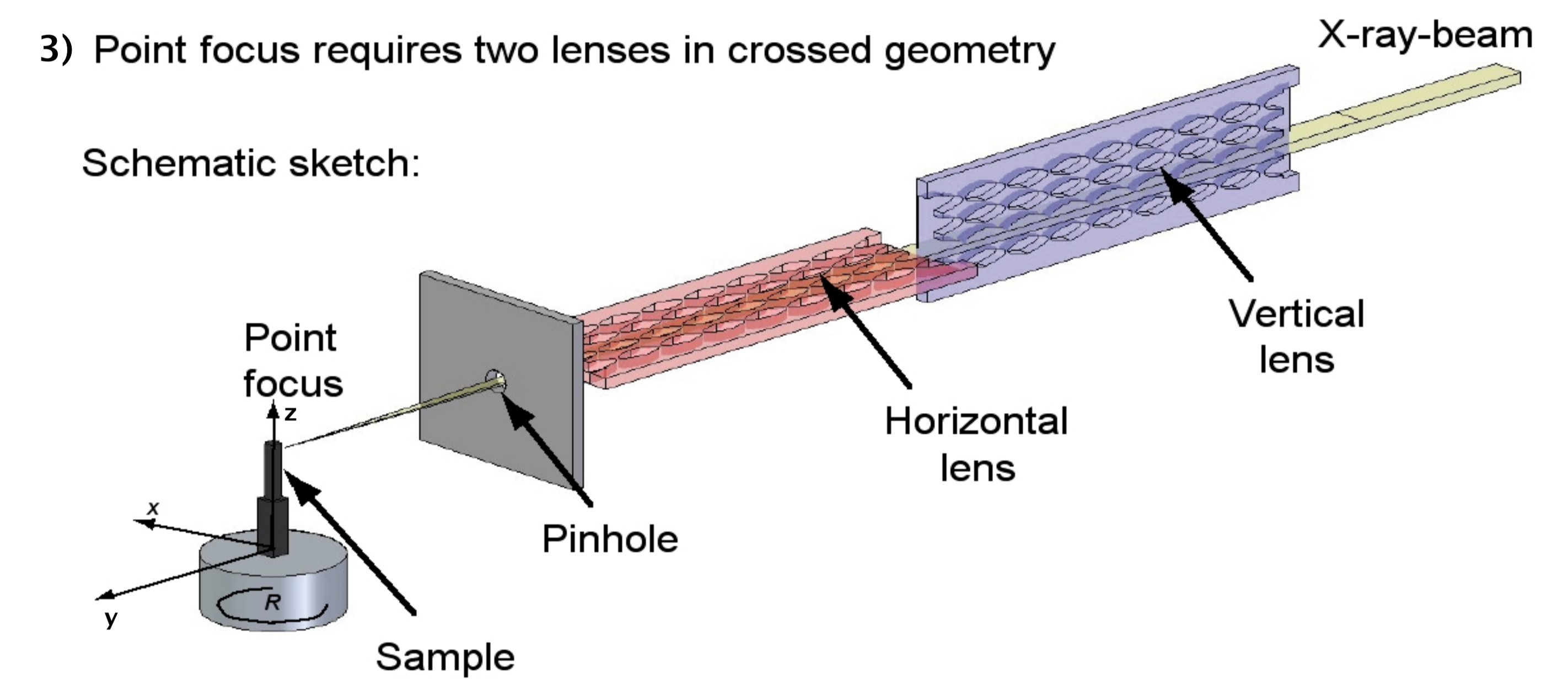
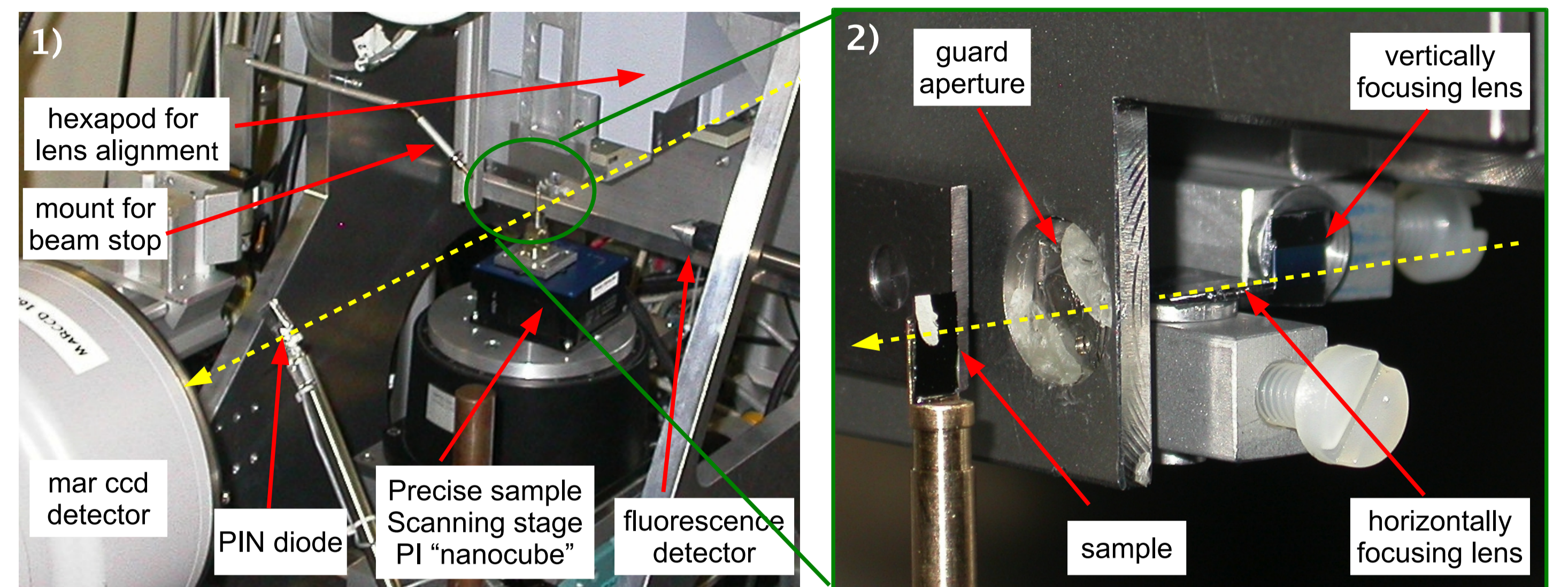


## Coherent Diffraction Imaging



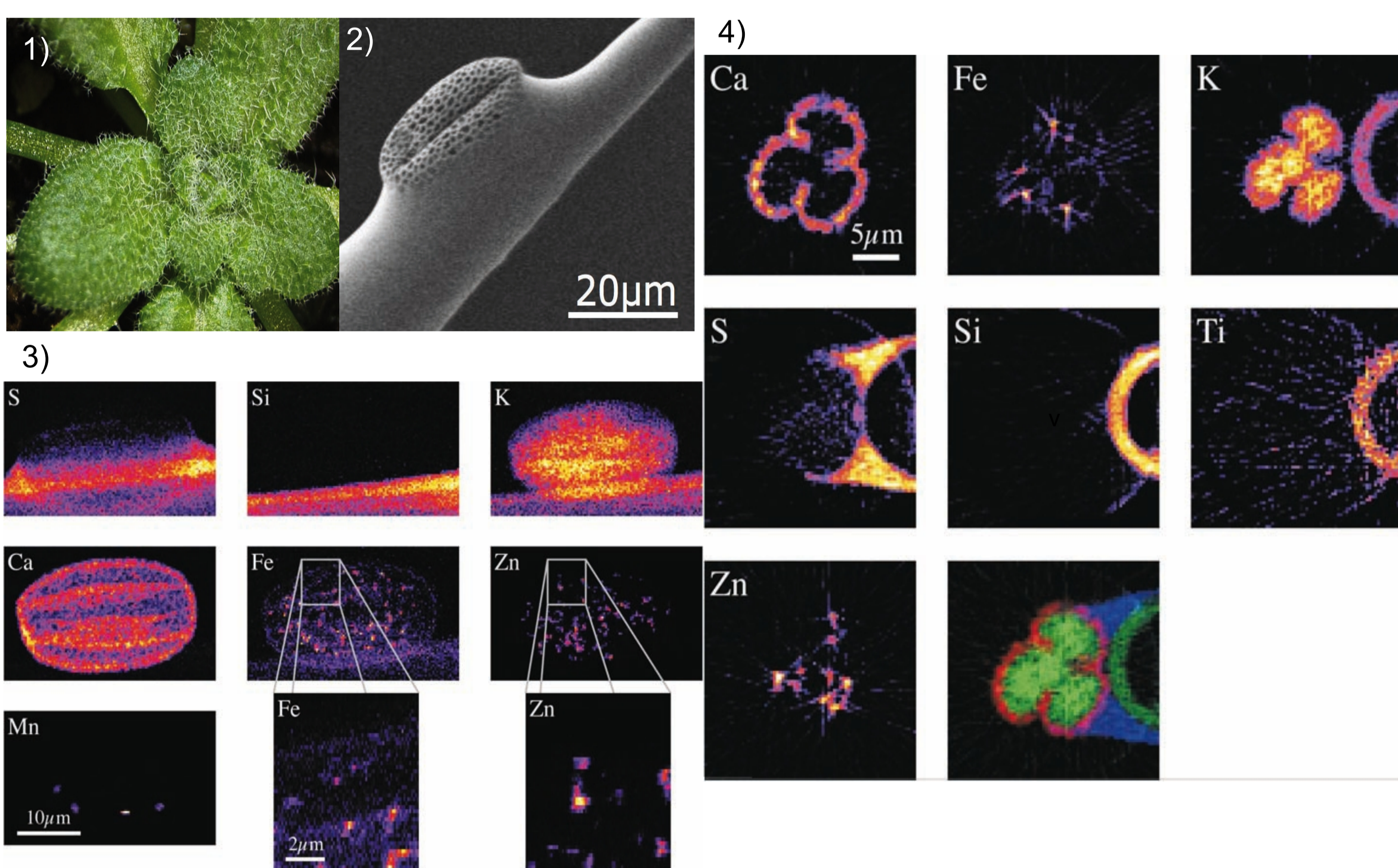
- Illumination with a nanobeam of  $100 \times 100 \text{ nm}^2$  size in focus, an energy of  $E = 15.25 \text{ keV}$ , and a flux of  $10^8 \text{ ph/s}$  (exposure time  $10 \times 60 \text{ s}$ ).
- Reconstruction by using the diffraction pattern shown in 1) and the hybrid input-output (HIO) method together with a so-called shrink wrap algorithm.
- 200 independent reconstructions.
- 191 converged to similar enantiomorphs of the gold particle. These were combined to an average reconstruction shown in 3).
- Spatial resolution between 3.8 nm and 7.6 nm ( $\sim 5 \text{ nm}$  by phase retrieval transfer function).

## Experimental Setup



- 1)2) Foto of the experimental setup as realized at ID 13, ESRF.
- There are several detectors such as a mar ccd camera, a high resolution x-ray camera, an energy dispersive detector and a PIN diode.
- The x-ray beam (yellow dashed line) is focused in two directions by crossed nanofocusing lenses shown in 3). The horizontal lens is fixed to the setup and can be aligned using a hexapod table on which the whole experiment is set up. The vertical lens is then adjusted by means of a mini hexapod.
- The sample is mounted on a high-precision piezo stage and can be scanned through the beam.
- A SiLi-detector detects the fluorescence light from the sample and a mar ccd detector records the diffraction pattern.

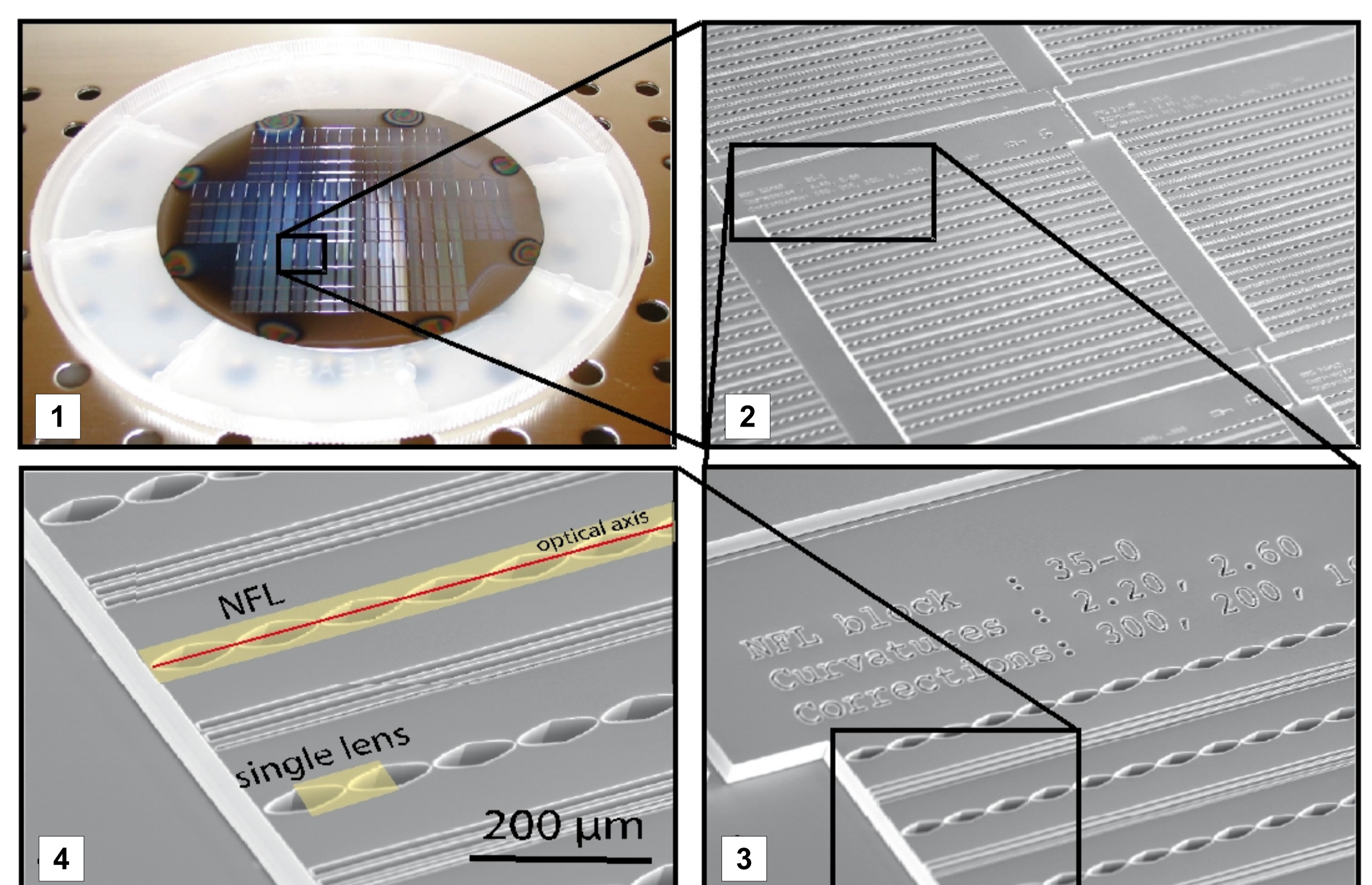
## Fluorescence Mapping and Tomography



Fluorescence experiment on a pollen of *Arabidopsis thaliana*.

- 1) Photograph of *Arabidopsis thaliana*.
- 2) Scanning electron microscope image of the pollen used in the experiment. The pollen was glued to a glass capillary.
- 3) Two-dimensional element distribution mapping with a resolution of 100 nm.
- 4) Tomographic reconstruction of the element distribution within the pollen. The lateral resolution was 300 nm.

## Focusing Device: NFL



- 1) Lenses made by 4-inch silicon wafer technology. There are 14 identical chips of a set of lenses arranged on the wafer shown in 1). Each chip holds more than 40000 single structures with 16 blocks of lenses.
- On each block there are 14 lenses with 2 different radii of curvature and 7 different corrections for under-etching. The different blocks contain different numbers of single lenses in a row, 2)/3).
- 2) Scanning electron microscope image of one block.
- 3) An individual bar code is structured into the wafer in between the lenses to ease the process of alignment of the lenses in the beam.

## References

- C. G. Schroer et al, Coherent X-Ray Diffraction Imaging with Nanofocused Illumination, *Phys.Rev.Lett.*, **101**(9), 090801, 2008.
- C. G. Schroer et al, Hard x-ray nanoprobe on refractive x-ray lenses, *Appl.Phys.Lett.*, **87**, 124103, 2005.

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