about

microscope optics



RESOLVING POWER

A human eye cannot distinguish two points when these are closer than approximately 0.3 mm or 300 μ m to each other. That is what is called "resolving power"; the capacity to resolve two points which are close together or "optical resolution", the ability of an imaging system to resolve detail

MAGNIFICATION

If one needs to detect details of 10 μ m, one needs at least 300 μ m / 10 μ m = 30x magnification in order to resolve this detail. Knowing that conventional stereo microscopes can magnify - without the help of additional lenses and without introducing additional optical aberrations – up to approximately max. 100 times, the smallest detail that a stereo microscope can resolve is 300 μ m / 100 = 3 μ m

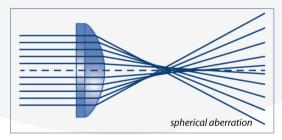
For details smaller than approximately 3 µm, one could better choose an optical compound microscope to resolve these details

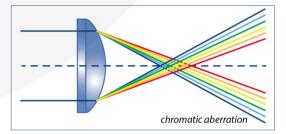
OPTICAL RESOLUTION

The resolution of an optical microscope can be defined as the shortest distance between two points that the microscope can reveal for a human eye or camera

The theoretical maximum resolution is limited by the diffraction laws of physics and is equal to

$$\label{eq:r} \begin{split} r = 1.22 \, \lambda \, / \, &(N.A. \, of \, objective + N.A. \, condenser) \\ \text{with } r = resolution, \, N.A. = numerical \, aperture, \\ \lambda = imaging \, wavelength \end{split}$$





ABERRATIONS

In practice, the resolution of a microscope is lower than the theoretical resolution due to optical aberrations, improper illumination or bad alignment of the optics and components of the microscope. Despite corrections for chromatic and spherical aberrations, coma, astigmatism and distortion, the real maximum resolution of a microscope with a 100x objective is about 0.25 µm for 550 nm light rays

Chromatic aberration
Spherical aberration
Astigmatism

Coma aberration

Distortion

Difference in focus of different colors caused by difference in refractive indices for each wavelength

Difference in focus caused by defects or irregularities on the surface of a lens Rays traveling through two perpendicular planes having different focal planes

Variation in magnification over the entrance pupil

Stretching of an image as a variation in magnification across the lens field