

Cells

Microscopes

Optical

Principles

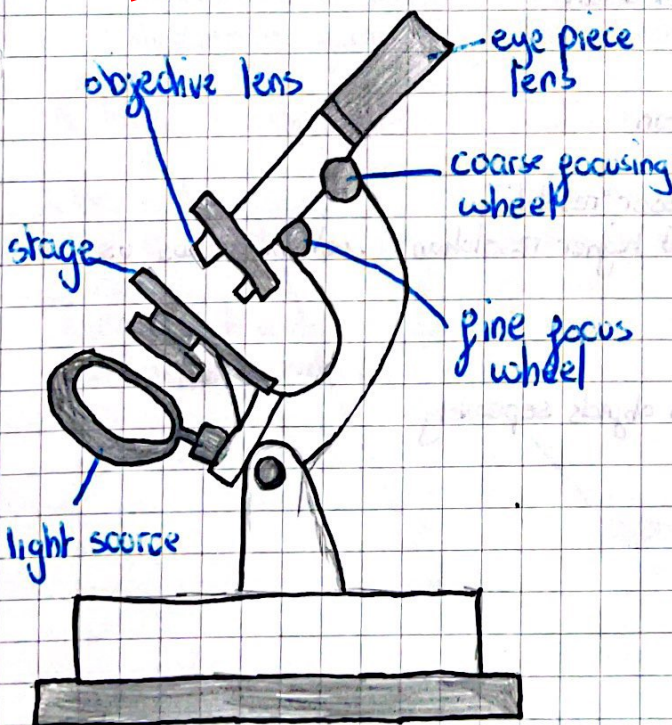
- ↳ The prepared specimen must be thin so it is only one layer of cells
- ↳ This will enable enough light to pass through
- ↳ Light waves pass through (or onto) the specimen

Limitations

- ↳ Lighter (not the 'microscope') has a longer wavelength
- ↳ So a lower resolution
- ↳ Lower magnification

Advantages

- ↳ Can view live specimens
- ↳ Coloured images
- ↳ Preparations rarely damage the specimen



Scanning Electrons

Principles

- ↳ Specimen is preserved & stained
- ↳ Place in a vacuum
- ↳ Uses an electron wave length
- ↳ Electron wavelength is shorter ... higher resolution (than optical)
- ↳ Electrons bounce off the surface (then are detected to create an image)

Limitations

- ↳ Electrons scattered (by molecules in air), so electrons are in a vacuum ... so there is no oxygen, so no respiration, so cannot examine living organisms
- ↳ More complex & time consuming specimen preparation procedure ... which may damage the specimen - alteration of the specimen's appearance are called 'artefacts'
- ↳ Black & white image (no colour)

Advantages

- ↳ Electron wavelength is shorter
- ↳ ... so higher resolution (than optical)
- ↳ ... so can study more detail (than optical)
- ↳ Can study 3D structures
- ↳ Higher magnification (than optical)