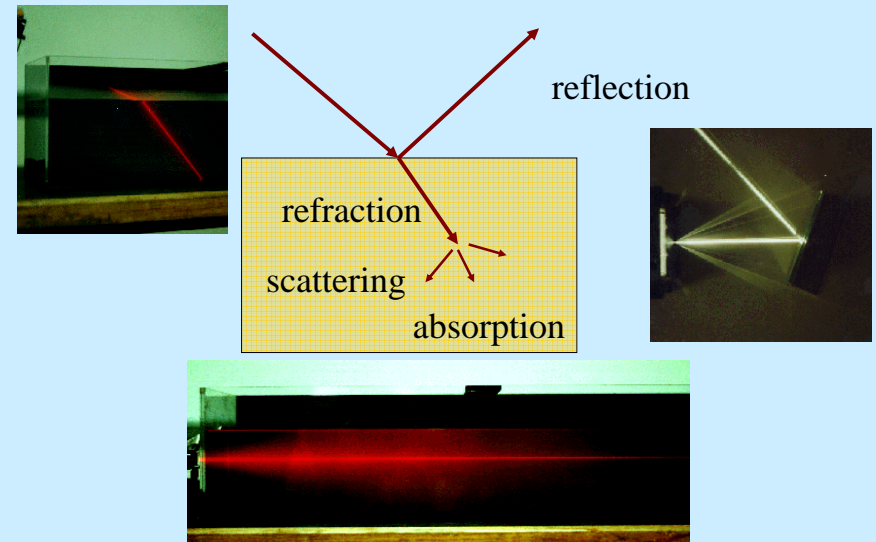


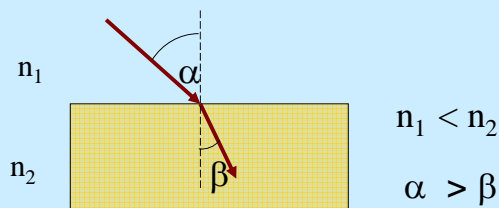
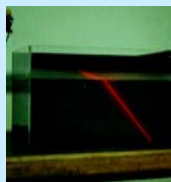
Interaction of light with matter 1.

Interaction of light with matter



Refraction of light

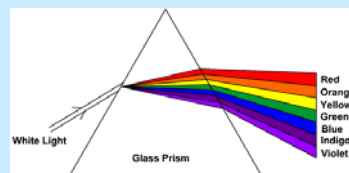
Fermat's Principle: Light follows the path of least time



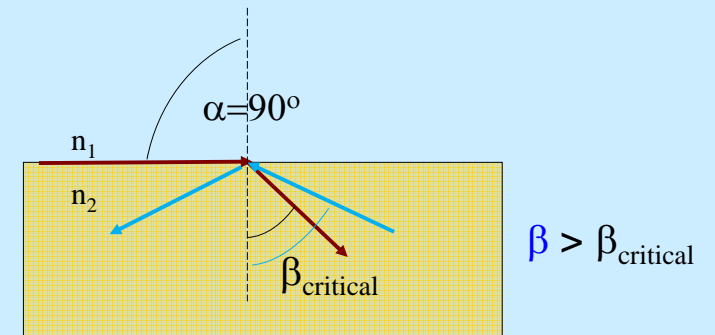
Snell's Law

$$\frac{\sin \alpha}{\sin \beta} = \frac{c_1}{c_2} = \frac{n_2}{n_1} = n_{21} \quad \text{The index of refraction}$$

Dispersion of light



Critical angle – total internal reflection



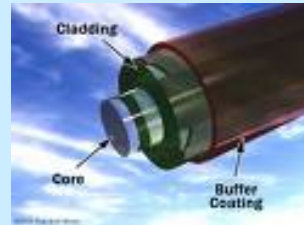
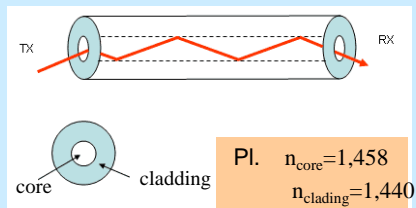
Medical application

Determination of concentration – refractometry

Concentration of solutions is proportional
with their index of refraction .



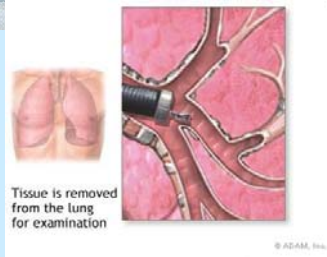
Optical fibers



Application in dentistry



Other medical applications



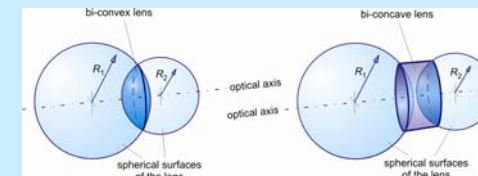
Bronchoscopy



Colonoscopy

Image formation occurs, when light rays emerging from one point converge at another point.

Image formation by thin lenses – Geometrical optics



Optical lenses and their interpretation by spherical surfaces

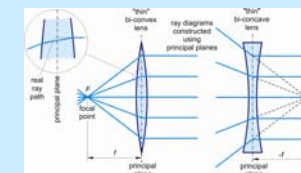
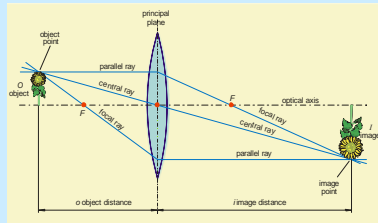


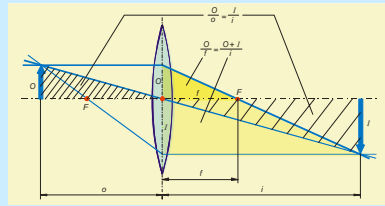
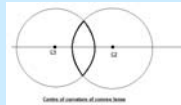
Image formation (thin lens approximation)

Image construction by principal rays



$$M = \frac{I}{O} = \frac{i}{o}$$

the radii of curvature

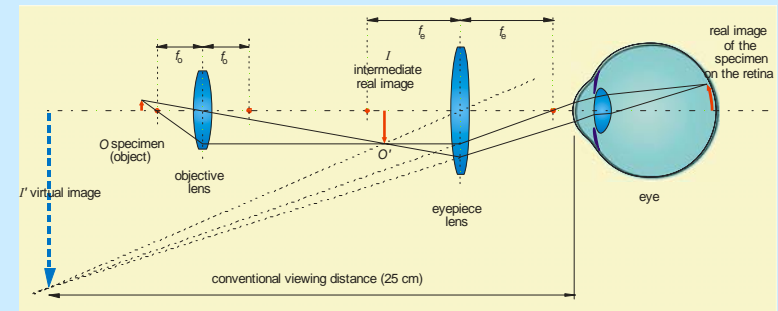


lensmaker's formula.

$$D = \frac{1}{f} = \frac{1}{o} + \frac{1}{i} = (n-1) \left(\frac{1}{r_1} + \frac{1}{r_2} \right)$$

D - diopter: measure of the optical power of a lens, which is equal to the reciprocal of the focal length measured in meters

Image formation – compound microscope

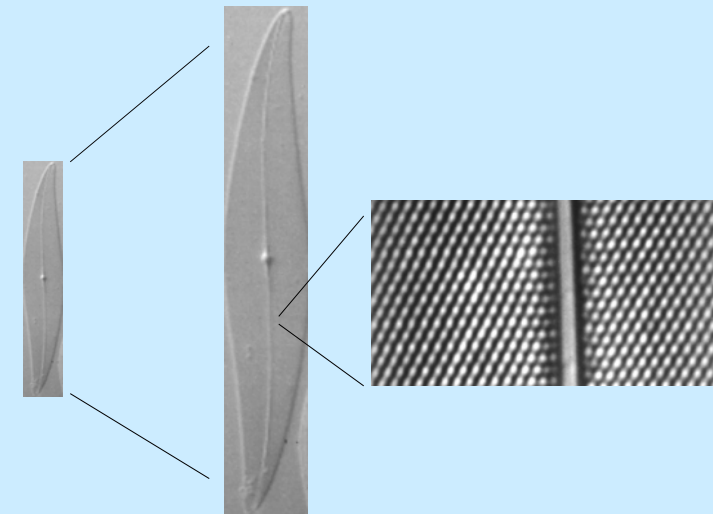


The image is magnified
reversed
virtual

Magnification vs Resolution



Magnification vs Resolution

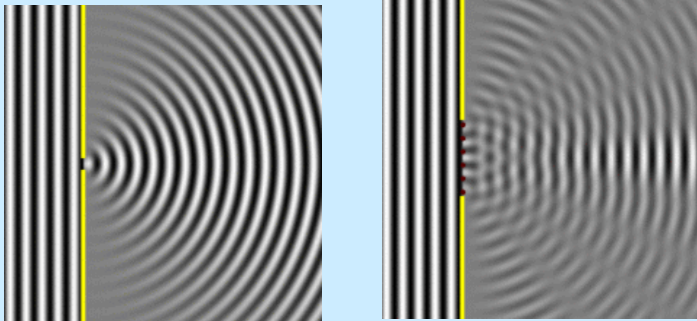


diatome

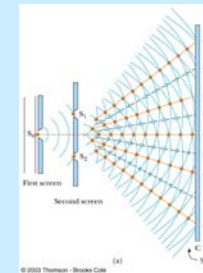
Limited resolution of microscopy

Wave nature of light

Huygens-principle



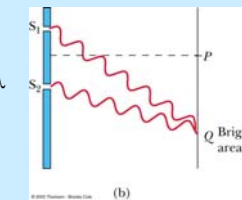
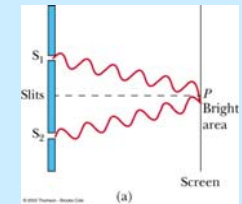
Young experiment



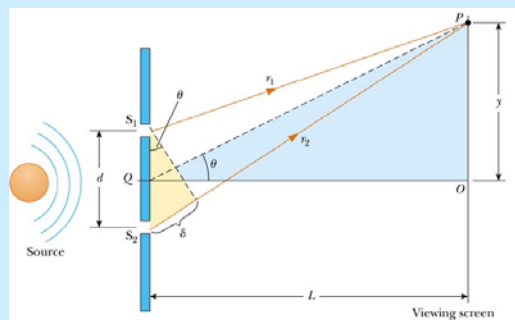
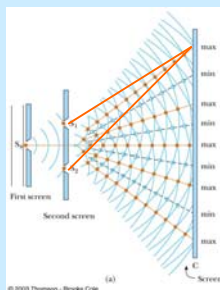
Where are the bright areas?

Constructive interference

If the distances from the slits
- are equal
or
- their difference is equal $n \cdot \lambda$



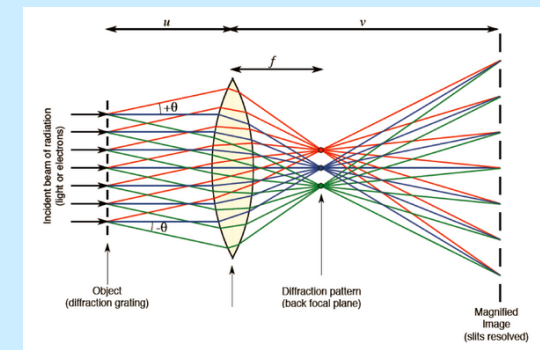
Young experiment



$$\delta = d \cdot \sin \Theta = k \cdot \lambda$$

$$d = \frac{\lambda}{n \sin \Theta}$$

Diffraction pattern in the microscope



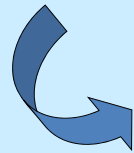
$$d = 0,61 \frac{\lambda}{n \sin \Theta}$$

ABBE'S PRINCIPLE: An optical system can resolve only those details of the specimen, which diffract light rays in a way that **besides the principal maximum at least the first order diffraction rays** are allowed to contribute to the image formation.

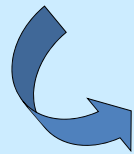
To decrease the limit of resolution – shorter wavelength – ¿matter wave?

Electron microscope

$$\lambda = h / mv$$



$$v = \sqrt{\frac{2eU}{m_e}}$$



U : 10 – 100 kV

$\lambda \sim 2 \text{ pm}$

Limit of resolution

Light microscope

Electron microscope

$$\lambda \sim 400 \text{ nm}$$

$$\lambda \sim 2 \text{ pm}$$

$$d = 0,61 \frac{\lambda}{n \sin \Theta}$$



$$d = \lambda / NA$$

NA ~ 2

NA ~ 10⁻³

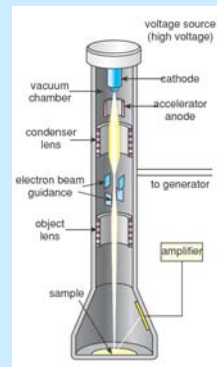
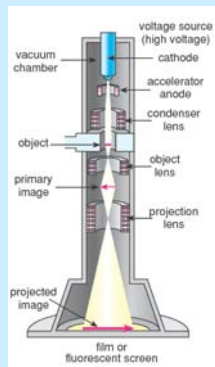
Limit of resolution ~ **200 nm**

Limit of resolution ~ **0,2 - 0,5 nm**

Structure of electron microscopes

Transmission electron microscope
TEM

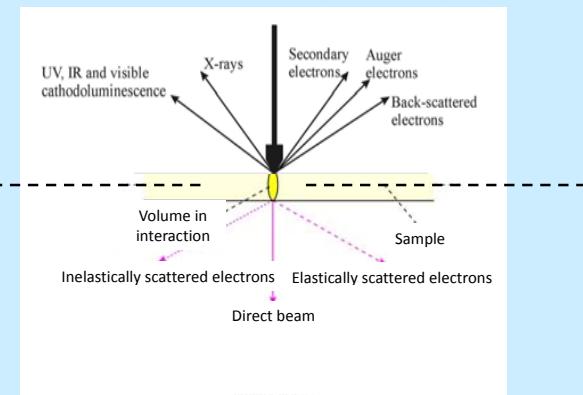
Scanning electron microscope
SEM



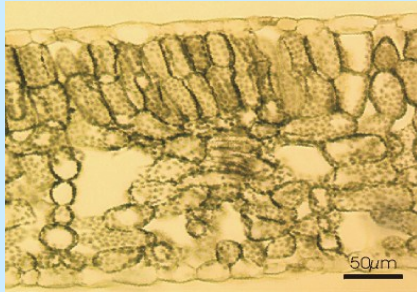
Interactions of electron beam

SEM

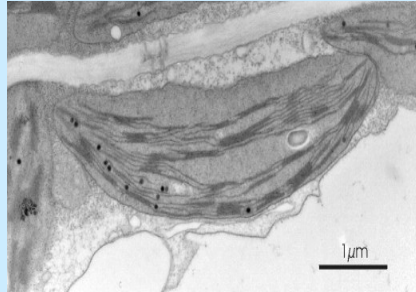
TEM



Light microscope vs Electron microscope

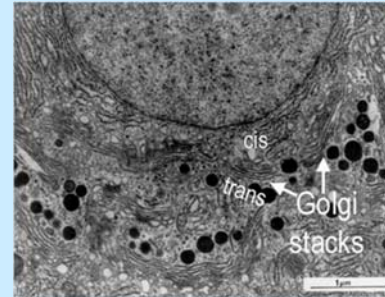


Semi-thin section of a spinach leaf in the light microscope.

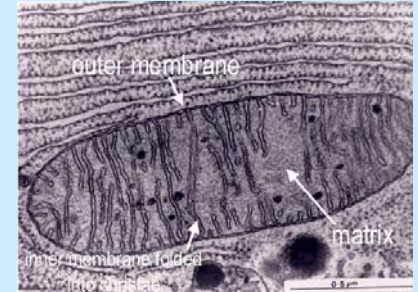


TEM micrograph: ultra-thin section of a spinach cell (chloroplast).

TEM

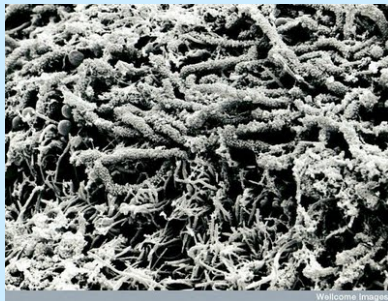


An electron micrograph showing golgi stacks



An electron micrograph showing mitochondrion

SEM



Brush your teeth often because this is what the surface of a tooth with a form of plaque looks like.

SEM

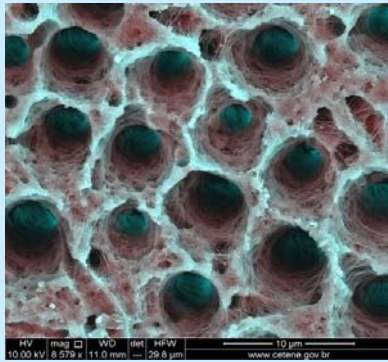


Scanning Electron Microscope image of bacteria in dental plaque magnified 30000 times !



Photograph © Mr. Steve Gschmeissner

SEM



Dentin is found in teeth and comprises tiny channels called dentinal tubules. This image shows those tubules.

Optics of the eye

$$D = \frac{n' - n}{R}$$

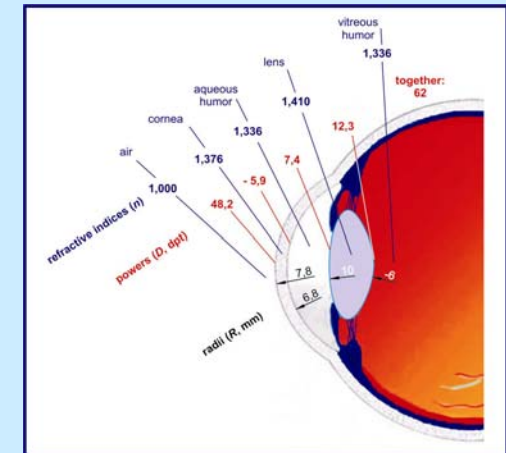
Power of the human eye

D : power (dpt)

n : refr. index of the 1. medium

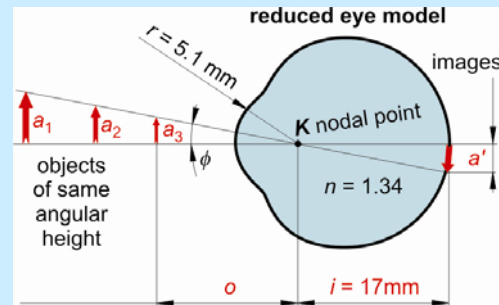
n' : refr. index of the 2. medium

R : radius (m)
+ convex
- concave



Optics of the eye

Image formation of the reduced eye



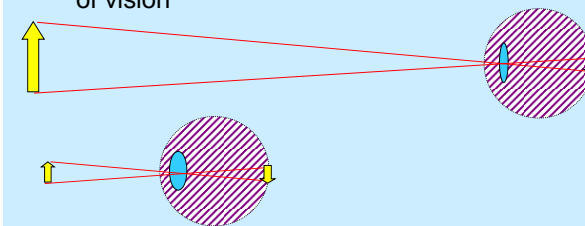
Power of the eye: sum of the powers of all refractive surfaces of the eye
unit: 1 dpt = 1/m, $D_{\text{eye}} = 59 - 72$ dpt

The image:

- real
- diminished
- inverted

Accommodation power

far point of vision



$$D_r = \frac{n'}{i} + \frac{n}{o_r}$$

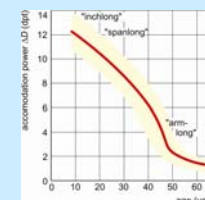
near point of vision

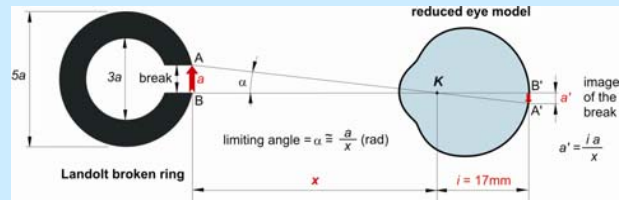
$$D_p = \frac{n'}{i} + \frac{n}{o_p}$$

Accommodation power: the difference of the largest and the smallest power of the eye

$$\Delta D = D_p - D_r = \frac{1}{o_p} - \frac{1}{o_r}$$

Near point o_p o_r Far point





Limiting angle of view (α):

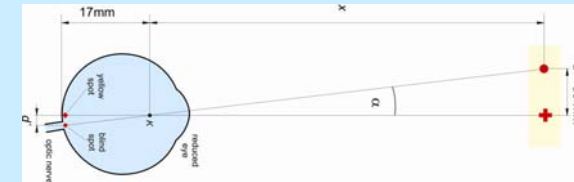
the smallest angular view of separated points A and B that can be just distinguished

Resolution of the eye or visual acuity (visus):

$$\text{visus} = \frac{1(^{\circ})}{\alpha(^{\circ})} \cdot 100\%$$

normal limiting angle ($1'$)

individual limiting angle



Measurement of the distance of the blind spot from the yellow spot

Question of the week

What is (are) the precondition(s) of total internal reflection in the core of optical fiber?

Related chapters

Damjanovich, Fidy, Szöllősi: Medical Biophysics

- II. 1.1.
 - 1.1.1
 - 1.1.3
- II. 2. 1.
 - 2.1.1
 - 2.1.2
 - 2.1.3
 - 2.1.4
 - 2.1.5
 - 2.1.8
- VI. 2.
 - 2.1.
 - 2.2.
- X.5.