



Imaging Basics: with photons/electrons

“Small is beautiful...”

Electron Microscopy ETH Zurich (EMEZ), Roger.Wepf@emez.ethz.ch

- Question to audience:
- Did you use any imaging techniques?
- LM..., EM..., MRI..., PET..., X-ray....?
- Who uses Image processing?
- Who is “comfortable” with contrasting/staining technique?
- Who is familiar with labeling techniques?
- Who can explain the duality of the probes?
- Who is familiar with the resolution range of any mode?
- Do you know the difference between frequency and wavelength...?

Content:

- why imaging - why 3D...
- *Imaging modes...*
- *Imaging “space”...*
- *Imaging principle*
- *Waves or particle - Duality*
- *Photon - no mass high speed*
- *Electron - mass and fast*
- *Imaging modalities e.g in LM and EM...& Applications*
- *Resolution...*
- *other Imaging modes..... X-ray Tomo, Imaging, PET, MRI...*

Aim of Imaging is primarily:

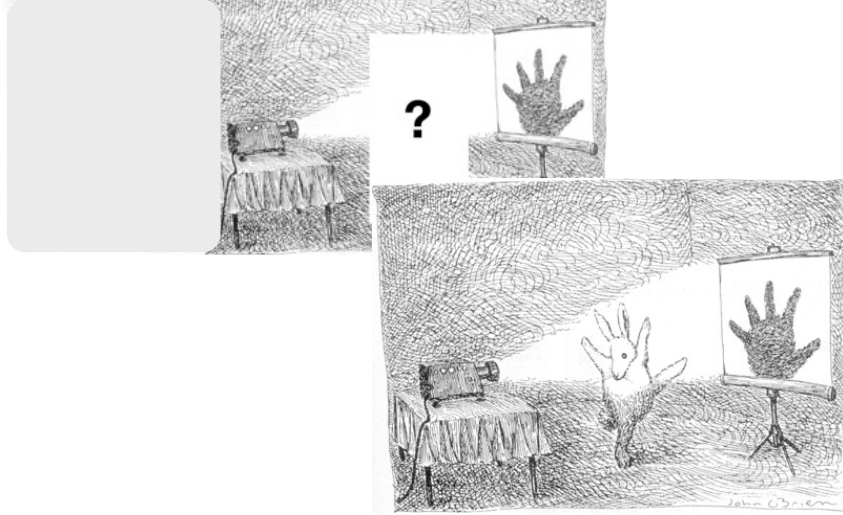
Get information on:

- **Structure/ Morphology**
- **Chemical composition/molecular composition**
- **Regional function**

As fast as possible, non-invasive and as natural as possible along all length scale to understand structure-function and time domain of life.

?? ?? ?? ??

Front View



Description of the "whole"....

- A question of perspective and methodic approach...

Perspectives and the truth:
 Is there **one** only Image of a object?



A View - A series of views...:



Vor etwa 100 Jahren versuchte Claude Monet die Gesamtheit (hier z.B. eines Heuschobers) durch mehrere Bilder aus unterschiedlichsten Perspektiven, Lichtverhältnissen und Jahreszeiten zu erfassen und zu beschreiben und so den Betrachter zum „Gesamten“ zu führen....

Art: "Imagination - One Image..."

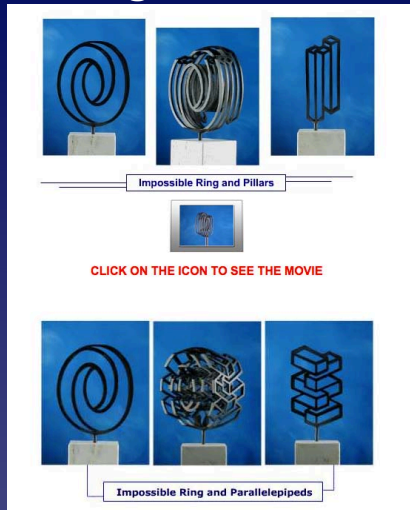


Do you trust your view-point...

-> Guido Morettis Body-Worlds....

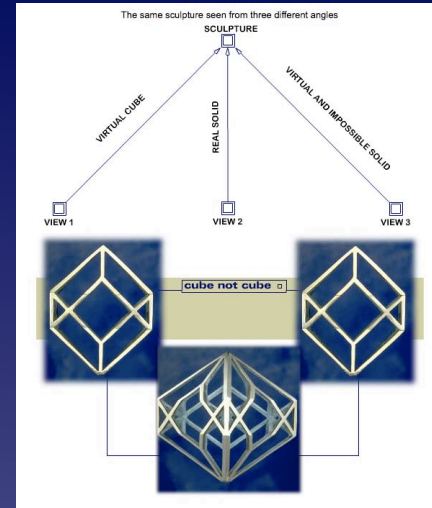
<http://www.coolopticalillusions.com/artists/guido-moretti/floating-3d-cube.htm>

Art: "Imagination - One Image..."



-> Guido Morettis Body-Worlds...
http://www.guidomoretti.it/E_illusione.htm

Art: "Imagination - One Image..."



<http://www.coolopticalillusions.com/artists/guido-moretti/floating-3d-cube.htm>

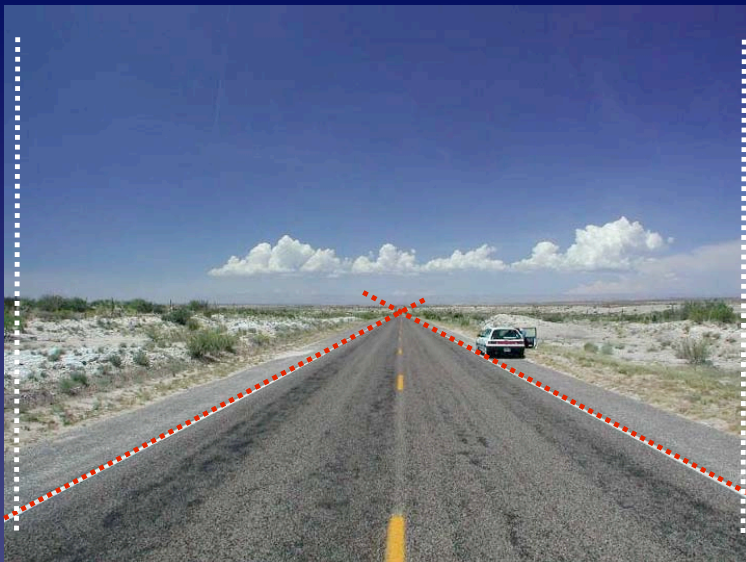


You need more than one
 view point -
 more than one image

Do you trust your view-point...
 -> Guido Morettis Body-Worlds....

http://www.guidomoretti.it/E_illusione.htm

The real world fools us day by day...The "parallel-paradox" -
 "two parallel lines never meet...".....



3D Imaging.....Microscopy:

- Essential to make the "Nano-Word" -
 - "visible, tangible & concrete"
- 3D required because single images may miss-lead you..
 (Viewpoint, Overlap, Irritation, Illusions, miss association....)
- 3D because the "small world" is translated to our "Thinking-Dimension"you can turn it and hold it...
 - Visual perception and brain power.....

A virtual experiment: let's enlarge the world by 1 Million times:

Than the real size of X becomes experienceable in our world

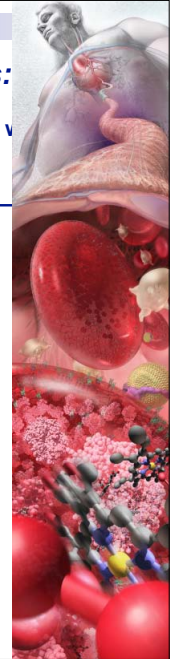
What:	Real Size:	Corresponding Size:
Atom	0,1-0,3nm	0,1-0,3mm (diameter body hair)
Protein	2-5nm	2-5mm (Rice corn)
Zytoskelett Fibers	9-25nm	9-25mm (robe)
Ribosom	25nm	25mm (2€ piece)
Nanoparticle	30-200nm	30-200mm (Walnut - CD)

If we enlarge the world by 1 Million times:

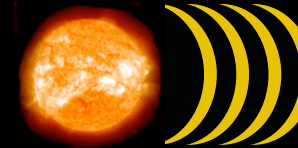
Than the real size of X becomes experienceable in our world

What:	Real Size:	Corresponding Size:
Atom	0,1-0,3nm	0,1-0,3mm (diameter body hair)
Virus	30-400nm	30-400mm (upto a glass bottle)
Bacteria	500-2000nm	0,5-2m
Cell nucleus	5 μm	5m
Human hair	60-100 μm	60-100m (Diameter)
Human eye	4 cm	40km
Human	1,7-1,8 m	1700-1800km

Your feet in Rom - your head in Hamburg
if an Atom is just 0,1mm large



How does an image emerge or how is it formed ?



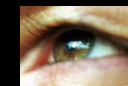
Self emitter



Eye/
Detector...



Remission:
Scattering/
Absorption/
Conversion



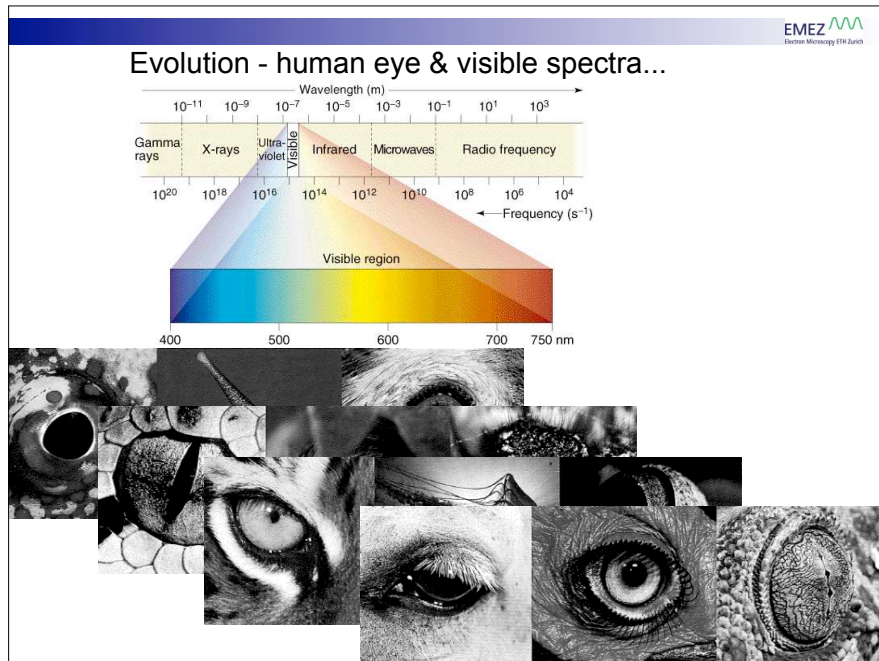
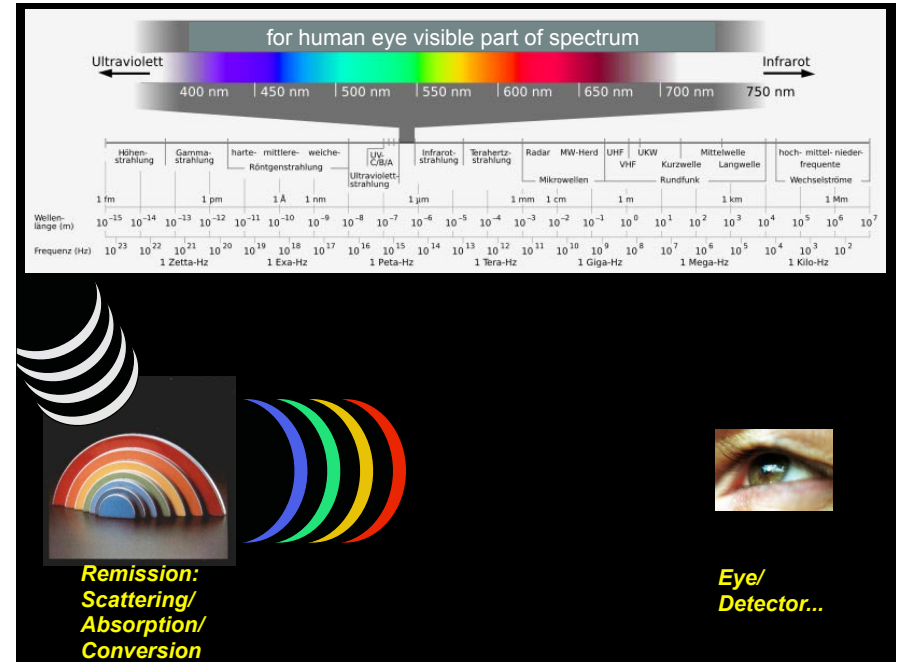
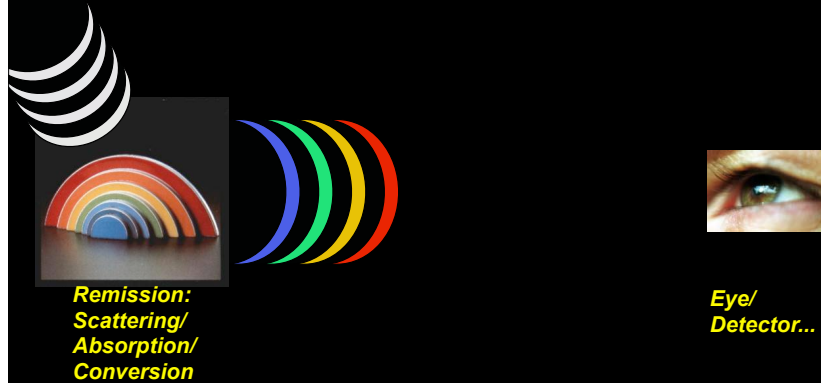
Eye/
Detector...

Rays (Waves):

Light (Photons - lampe, laser...)

Particle (Electrons, Ions,...)

Sound (Ultrasound, electromagnetic...)



Imaging Modalities in Life-Science

MRI, MEG, EEG, fMRI

IR, PET

„CERN“

X-ray, NMR

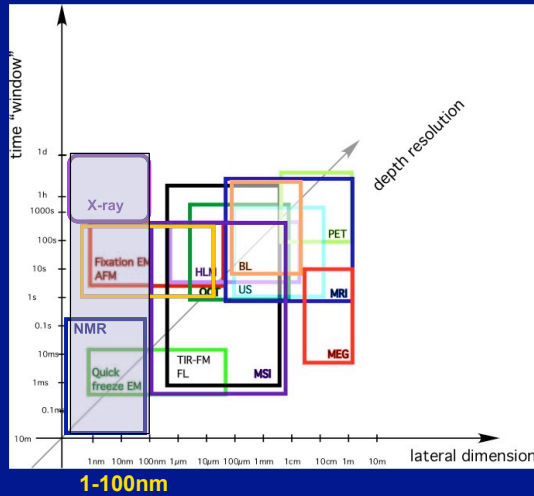
EM, IM, SPM

LM, CLSM

CIMST

www.neuroscience.cam.ac.uk

The "Imaging Space"



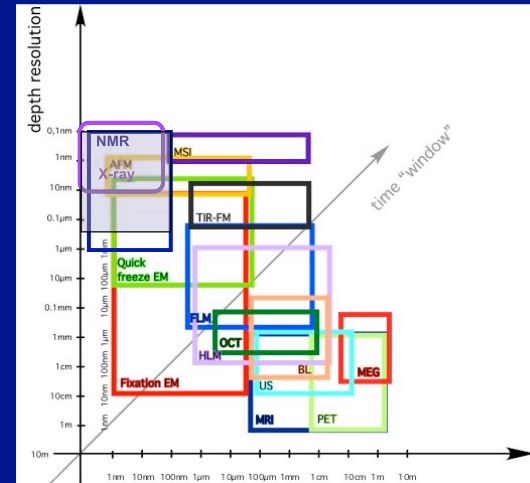
1-100nm

Aim: enlarge samples to make them visible for the human eye or processable in a PC (Pixel/Voxel)

im:ag:ing

Date 1967: the action or process of producing an image especially by means other than visible light

The "Imaging Space"



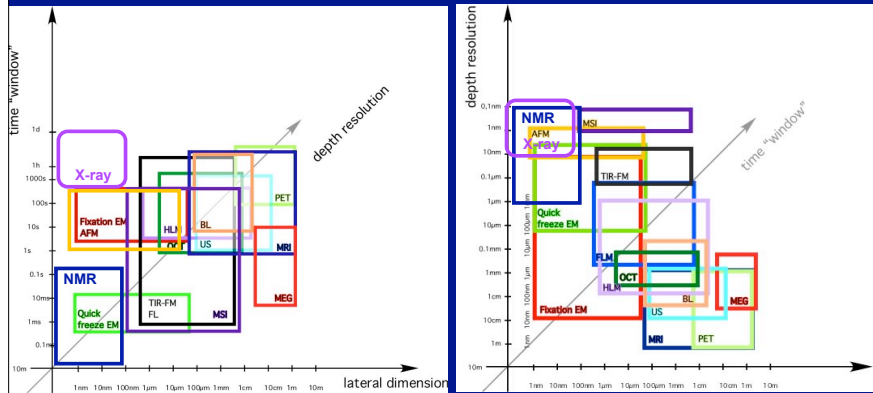
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The "Imaging Space"



1-100nm

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Aim: enlarge samples to make them visible for the human eye or processable in a PC (Pixel/Voxel)

im:ag:ing

Date 1967: the action or process of producing an image especially by means other than visible light

Imaging: Principle

"Information transfer chain"

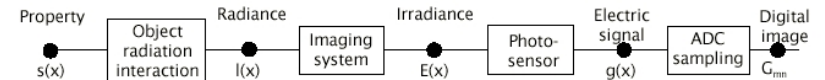


Figure 1.1: Chain of steps linking an object property to the signal measured by an imaging system.

What is Electromagnetic Radiation?

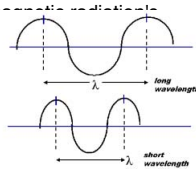
Electromagnetic radiation is energy – we describe it as a wave – visible light is only a small portion

The characteristics which distinguish different types of light are the electron

- wavelength λ
- frequency f
- energy

frequency

$$f = c / \lambda = \frac{v}{\lambda}$$



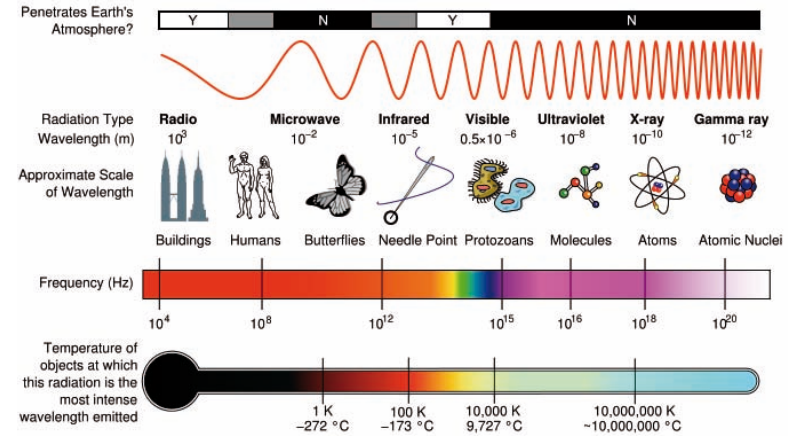
Wavelength: – the distance between two peaks (or two troughs) of the wave.
Frequency: – the number of wavelengths passing a given point in one second.

->The longer the wavelength, the more time it takes for a full wave to pass a given point (or the fewer waves pass the point in a given time).

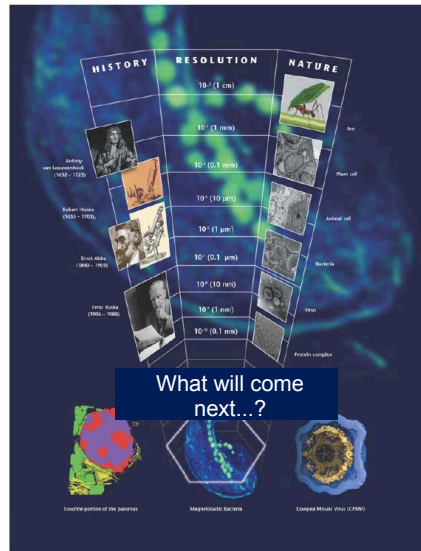
Since frequency is wavelengths per second, as the wavelength becomes longer the frequency decreases, and vice versa.

Energy: - is directly proportional to the frequency-- if the frequency increases, so does the energy of the radiation and vice versa.

The Electromagnetic Spectra.....



Wikipedia....



CLASS	FREQUENCY	WAVELENGTH	ENERGY
γ	300 EHz	1 pm	1.24 MeV
HX	30 EHz	10 pm	124 keV
SX	3 EHz	100 pm	12.4 keV
	300 PHz	1 nm	1.24 keV
EUV	30 PHz	10 nm	124 eV
NIR	3 PHz	100 nm	12.4 eV
MIR	300 THz	1 μ m	1.24 eV
FIR	30 THz	10 μ m	124 meV
	3 THz	100 μ m	12.4 meV
EHF	300 GHz	1 mm	1.24 meV
SHF	30 GHz	1 cm	124 μ eV
UHF	3 GHz	1 dm	12.4 μ eV
VHF	300 MHz	1 m	1.24 μ eV
HF	30 MHz	1 dam	124 neV
MF	3 MHz	1 hm	12.4 neV
LF	300 kHz	1 km	1.24 neV
VLF	30 kHz	10 km	124 peV
VF	3 kHz	100 km	12.4 peV
ELF	300 Hz	1 Mm	1.24 peV
	30 Hz	10 Mm	124 feV

FEI; Wikipedia....

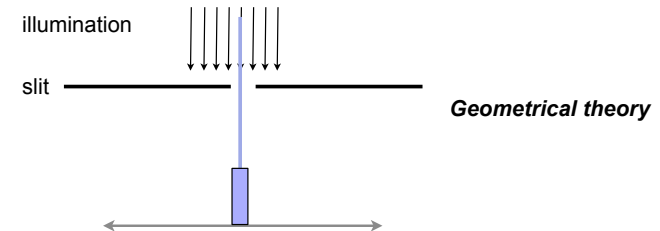
Duality of "light" - particle or waves ?

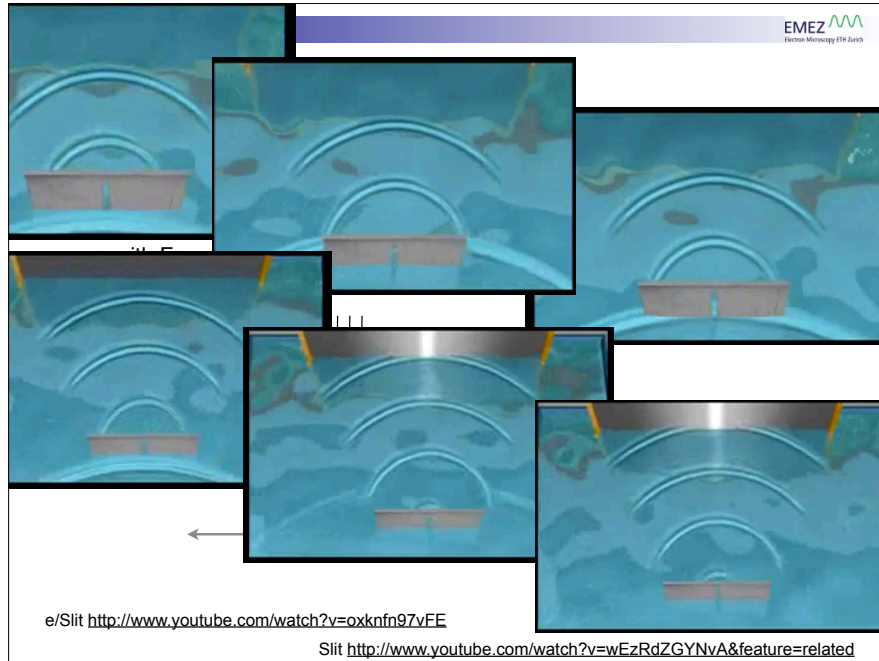
The particle concept:

an energy concept with Electronvolt (eV) - 1eV is the kinetic energy of a charged particle which is accelerated by an electric field of 1 Volt.Newton (1642- 1727)

The wave concept:

waves with Frequenz (f), Wellenlänge (l) Wavenumber (wz) represent energy-proportional units... Huygens (1629-1695)&Hooke (1638-1703)





EMEZ
Dietrich Hertz

- Duality of "light" - particle or waves ?

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illumination

slit

Geometrical theory

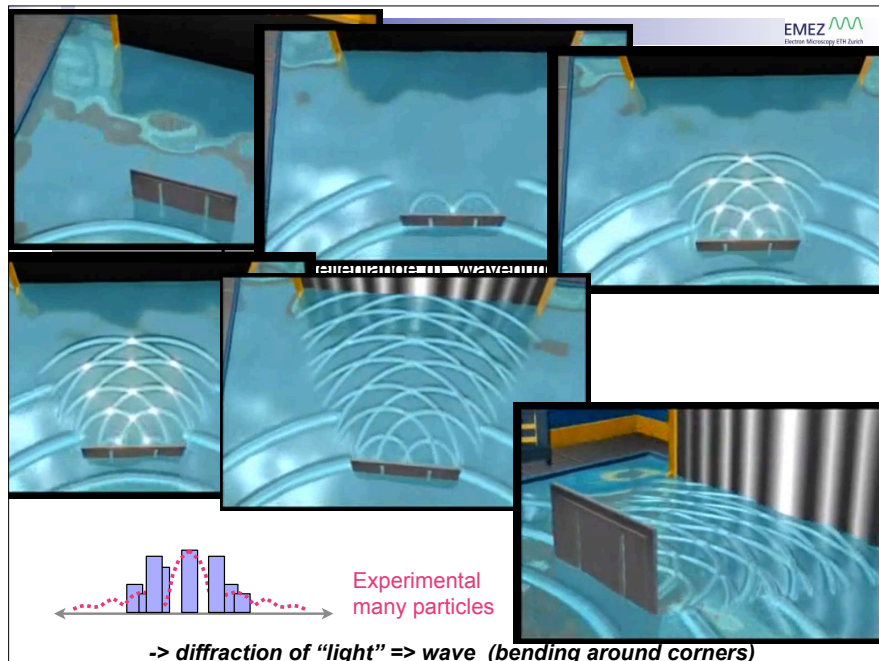
Experimental x particles

Experimental many particles

<http://www.youtube.com/watch?v=oxknfn97vFE>

-> diffraction of "light" => wave (bending around corners)

e/Slit <http://www.youtube.com/watch?v=oxknfn97vFE>
Slit <http://www.youtube.com/watch?v=oxknfn97vFE>



EMEZ
Dietrich Hertz

Unbelievable.....

....also particles e.g. electrons/ He-ions do it if not observed

<http://www.youtube.com/watch?v=oxknfn97vFE>

■ Duality of “light” - particle or waves.....

The particle concept:

a energy concept with Electronvolt (eV) - 1eV is the kinetic energy of a charged particle which is accelerated by an electric field of 1 Volt.

-> “classical ray optics”

The wave concept:

waves with frequency (f), wavelength (λ) wave-number (wz) represent energy-proportional units.....

-> “wave optics” (wave front and Schrödinger eq.)

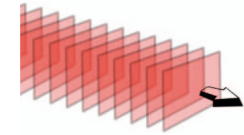
$$E = h \cdot f$$

(h...PLANCK’s constant) directly relates the energy with the frequency of a “ray” Quantum Theory (Planck & Einstein)

The relation between the wavelength (λ) of a particle of mass, m, moving at a velocity, v, is given by the **DeBroglie** wave equation:

$$\lambda = \frac{h}{p}, \quad \lambda = \frac{h}{mv} \quad (1)$$

■ Not one wave but plane waves are used for imaging



In the physics of wave propagation, a **plane wave** is a constant-frequency wave whose **wavefronts** (surfaces of constant **phase**) are infinite parallel planes of constant **amplitude** normal to the **phase velocity** vector.

Mathematically, a plane wave is a wave of the following form:

$$u(\mathbf{x}, t) = A e^{i(\mathbf{k} \cdot \mathbf{x} - \omega t)}$$

where *i* is the **imaginary unit**, **k** is the **wave vector**, **ω** is the **angular frequency**, and **A** is the **(complex) amplitude**.

-> two dimensional sinusoids have a frequency, phase, amplitude and direction!

(This becomes of interest when Fourier Transformations (FFT) are used to analyze images (Amplitude in real part, Phase in imaginary part))

■ Conversion of energy-> wavelength-> wave-number.....

$$E = h \cdot f$$

The wavelength of light is via the speed of light (c) linked to the frequency (general: $\lambda = \frac{v}{f}$,) => $\lambda = c / f$

The so called wave-number is the reciprocal of the wavelength (wz, n)
 $n = 1 / \lambda$ (n Wave number usually in cm-1)

-> $E = h \cdot c \cdot n$

Conversion from one to the other unit:

$\lambda [\mu\text{m}] = 10'000 / n [\text{cm}^{-1}]$; $\lambda [\text{nm}] = 10'000'000 / n [\text{cm}^{-1}]$

$f [\text{Hz}] = 3 \cdot 10^{10} \cdot n [\text{cm}^{-1}]$

$E [\text{eV}] = 1 / 8065,5 \cdot n [\text{cm}^{-1}]$

<http://www.cactus2000.de/de/unit/masswav.shtml>

Imaging: Principle

“Information transfer chain”

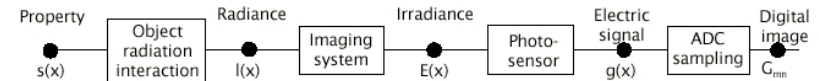


Figure 1.1: Chain of steps linking an object property to the signal measured by an imaging system.

To image a certain structure the used wavelength should be in the range of the structure detail to be imaged... if not the wavelength can not interact linearly with the object...

- Conversion in our head.....
 - Visual perception and brain power.....

Watch out what your brain does!

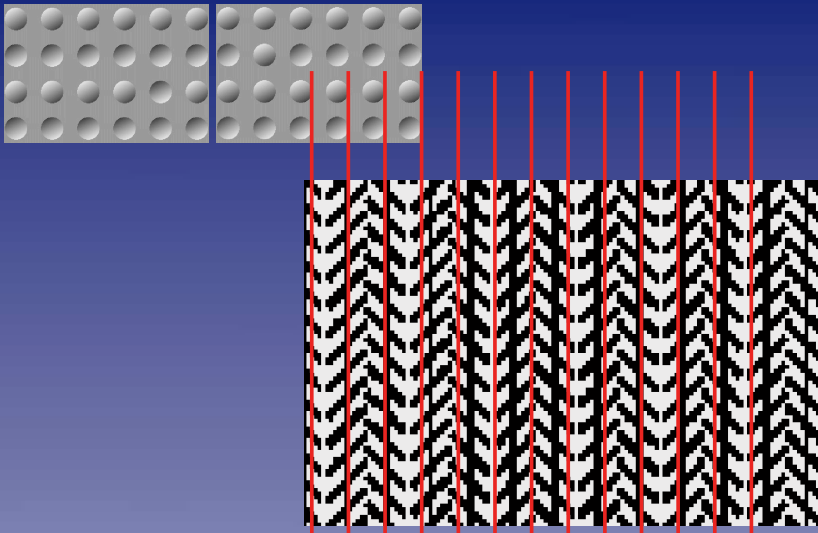
Just start to read (engl. or german).....

Aoccdrnig to a rscheearch at an Elingsh uinervtisy, it deosn't mtttaer in waht oredr the ltteers in a wrod are, the olny iprmoetnt tihng is taht frist and lsat ltteer is at the rghit pclae. The rset can be a toatl mses and you can sitll raed it wouthit porbelm. Tihs is bcuseae we do not raed ervey lteter by it slef but the wrod as a wlohe. ceehiro.

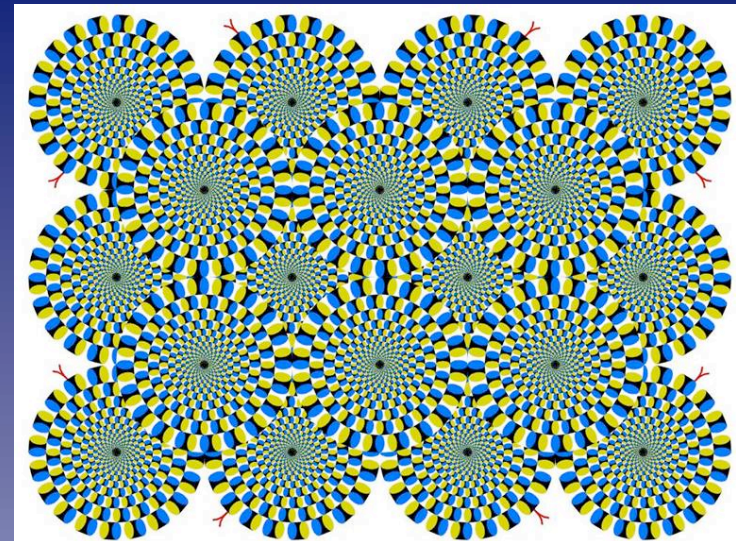
David Harris 2003; David R. Perrott and Caltech senior research fellow Kourosh Saberi, Nature 1999 on speech "Cognitive Restoration of Reversed Speech."

Afugnmud enier Sduite an enier Elingshcn Unvirestiät ist es eagl, in wlehcer Rienhnefoge die Bcuhtsbaen in eniem Wrot sethen, das einzig woihitge dbaei ist, dsas der estre und lzete Bcuhtsbae am rcihgiten Paltz snid. Der Rset knan ttolaer Bölsdinn sien, und du knasnt es torztedm onhe Porbelme lseen. Das ghet dseahlb, wiel wir nchit Bcuhtsbae für Bcuhtsbae enizlen lseen, snodren Wröetr als Gnaezs.

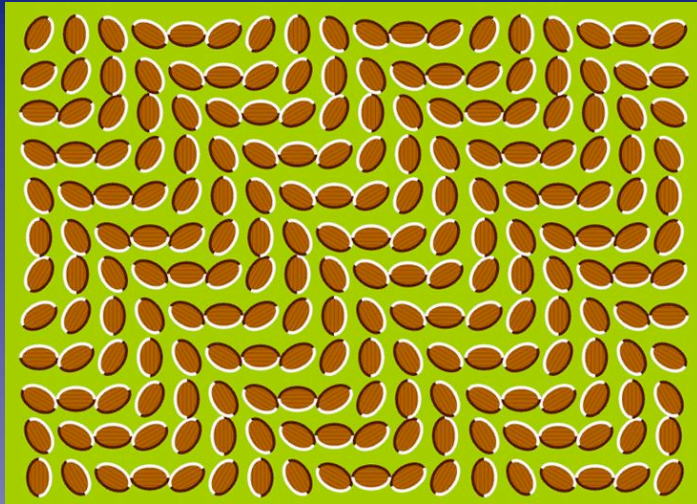
Can you trust what we see...



Nothing is turning - or...?



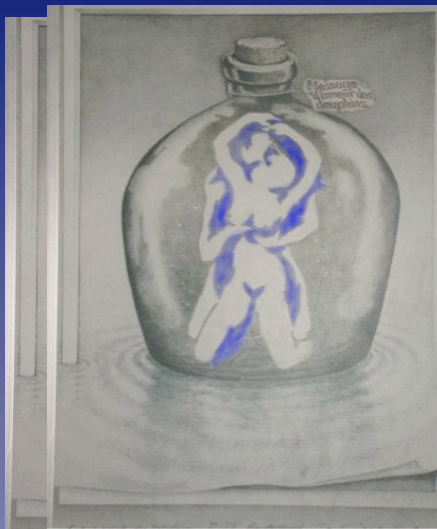
.... an nothing is moving!



Do we trust what we see... Illusionen!



What do you see in this image.....?



Children see only
Dolphins
...what did you see???

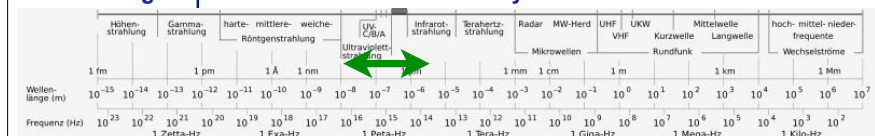
Sandro Del-Prete

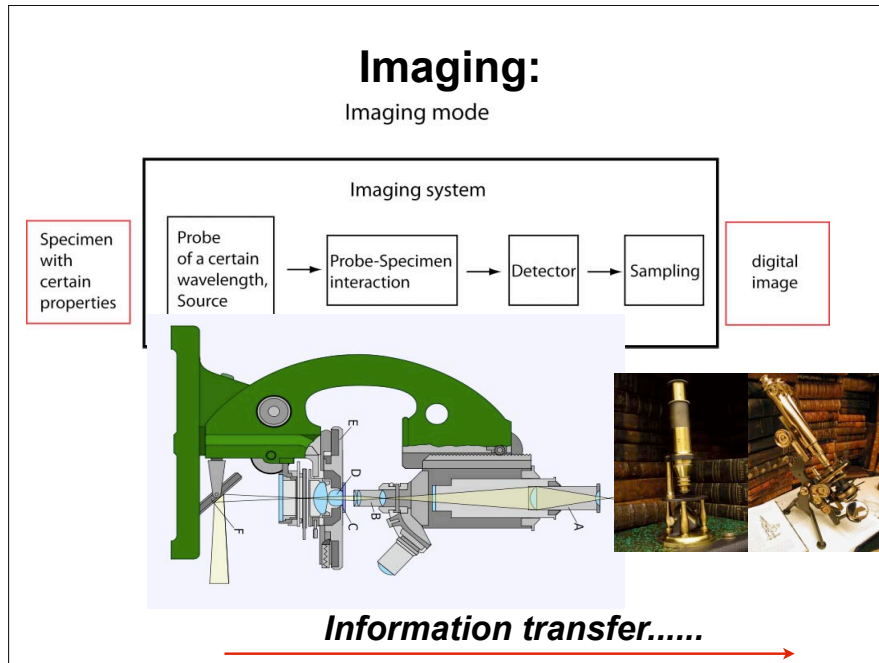
"A world were your
eyes deceive you"

<http://www.illusoria.com>

Photon..... Properties:

	Photon (Boson/Eichboson)	Electron (Fermion/Lepton)
Charge:	0	
Mass:	0	
Resting Energy:	eV	
Compton Wavelength:	-	
Spin:	1	
Radius:	-	
Wavelength	300nm-1600nm in use today...	





Imaging Mode: Light Microscopy (LM)

- **Full field illumination** (Bright field, Phase contrast, Differential Interference Contrast, Fluorescence, Polarisation, Spectroscopy...)
 - -> parallel light is exposed to the whole specimen
 - -> thickness of specimen is limiting resolution
 - -> fast light exposure possible
 - -> projection/reflexion images of the exposed area
- **Scanning mode** (Reflexion, Fluorescence, Spectroscopy - Raman)
 - -> focused light spot is scanned through the specimen
 - -> scanning in x,y and z possible
 - -> thickness not so critical
 - -> slow for large image area
 - -> pixel by pixel images

Light Microscope (LM) Illumination types....

Wide field & confocal illumination

TIRF illumination

2007; Jyoti K Jaiswa & Sanford M Simon

Motile cells (Fish keratocyte) - Fluorescence LM... (Resolution or Seeing....?)

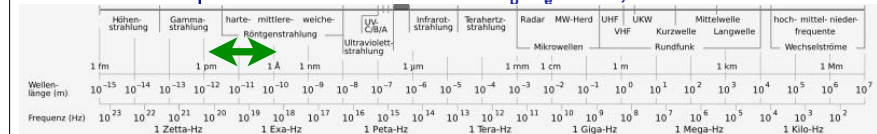
F-actin
(ϕ 9nm)

Microtubules
(ϕ 25nm)

- **Imaging Mode: Light Microscopy (LM)**
- **Probe:** bundle of light or focused spot of light
- **Wavelength:** 300-1000nm
- **Probe-Specimen Interaction:** Absorption, elastic and inelastic scattering, conversion (fluorescence, Raman), phase shift...
- **Modalities:**
 - full-field illumination/Interference, Phase contrast, Scattering & Absorption, Emission
 - Scanning probe imaging/ Absorption, Emission, Scattering...
- **Resolution:**
- **Application:**
- **Advantage:**
- **Limitation:**

Photon/Electron Properties:

	Photon (Boson/Eichboson)	Electron (Fermion/Lepton)
Charge:	0	-1e=-1,602x10 ⁻¹⁹ C
Mass:	0	5,485x10 ⁻⁴ u (1 m _e) 9,109x10 ⁻³¹ kg
Resting Energy:	eV	0,5109 MeV
Compton Wavelength:	-	2,426x10 ⁻¹² m
Spin:	1	+/- 1/2
Radius:	-	2,817x10 ⁻¹⁵ m m _p /m _e = 1836,1



Electron: Energy & Wavelength

The dualism „wave - particle“ is quantified by the De Broglie equation:

$$\lambda = h/p = h/mv$$

λ : wavelength; h : Planck constant; p : momentum

The energy of accelerated electrons is equal to their kinetic energy:

$$E = eV = \frac{m_0v^2}{2}$$

V : acceleration voltage

$e / m_0 / v$: charge / rest mass / velocity of the electron

$$p = m_0v = (2m_0eV)^{1/2}$$

$$\lambda = h / (2m_0eV)^{1/2} (\approx 1.22 / V^{1/2} \text{ nm})$$

At the acceleration voltages used in TEM, relativistic effects have to be taken into account $\lambda = h / [2m_0eV (1 + eV / 2 m_0c^2)]^{1/2}$

Wavelengths of Electrons

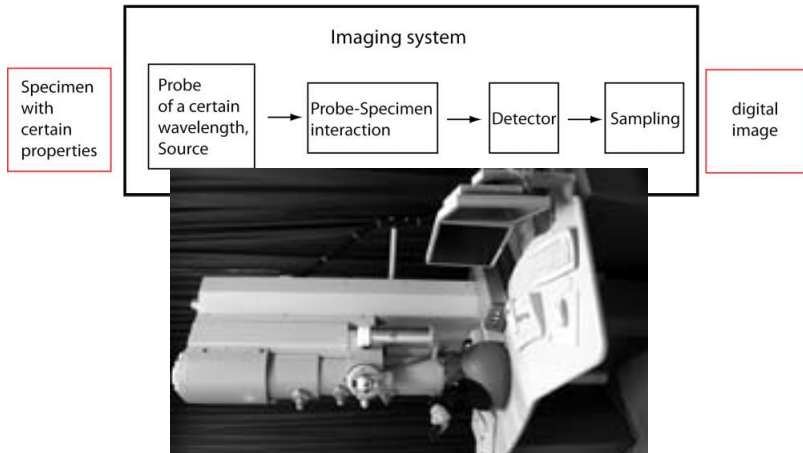
Accelerating voltages: SEM 0.5 – 30 kV
TEM 100 – 1'000kV

V _{acc} / kV	Nonrelativistic wavelength [nm]	Relativistic wavelength [nm]
1	0.0388	0.0388
40	0.00613	0.0060
100	0.00386	0.00370
300	0.00223	0.00197
1000	0.00124	0.00087

(Atomic distances: ~ 0.1 nm (Å))

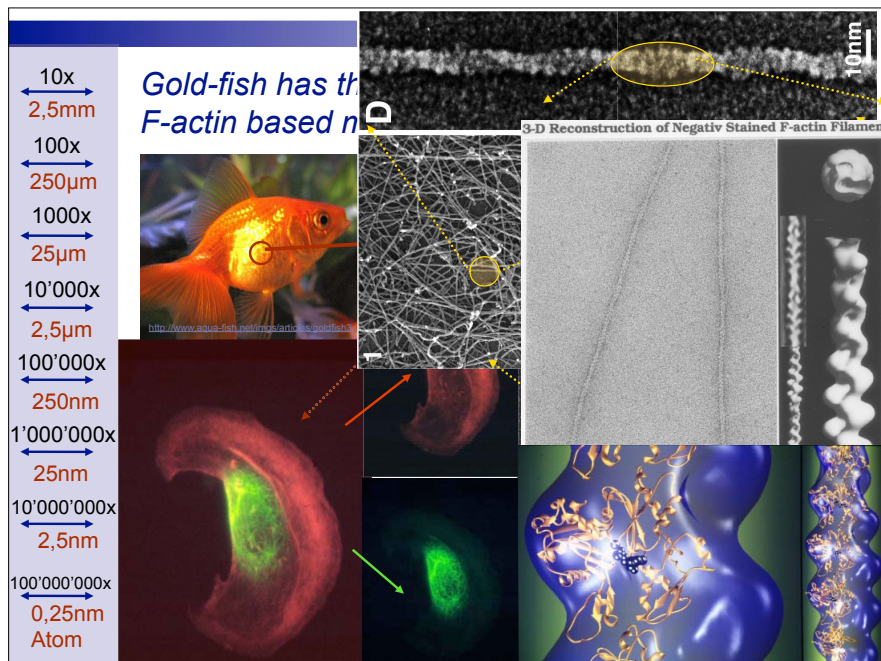
Imaging: “Information transfer chain”

Imaging mode



Imaging Mode: Electron Microscopy (EM)

- Probe: parallel beam or focused spot of electrons
- Wavelength: 10^{-3} - 10^{-6} nm
- Probe-Specimen Interaction: Absorption, elastic and inelastic scattering, conversion (fluorescence, luminescence), phase shift, ...
- Modalities:
 - full-field illumination; Interference, Phase contrast, Scattering & Absorption, Emission
 - Scanning probe imaging; Absorption, Emission, Scattering...
- Resolution:
- Application:
- Advantage:
- Limitation:



Imaging Modalities in Life-Science

MRI, MEG,
EEG, fMRI

IR,
PET

X-ray,
MR

Thanks for your attention...

LM, CLSM

EM,
IM
SPM

