



Shining a **light** on synchrotron **light**

Sylvain RAVY

Laboratoire de Physique des Solides
CNRS, Université Paris-Saclay
91405 Orsay
France

OUTLINE

Brilliance, emittance... coherence

Synchrotron light: waves and photons

Scattering and absorption

Diffraction, Spectroscopy, Imaging
beamlines

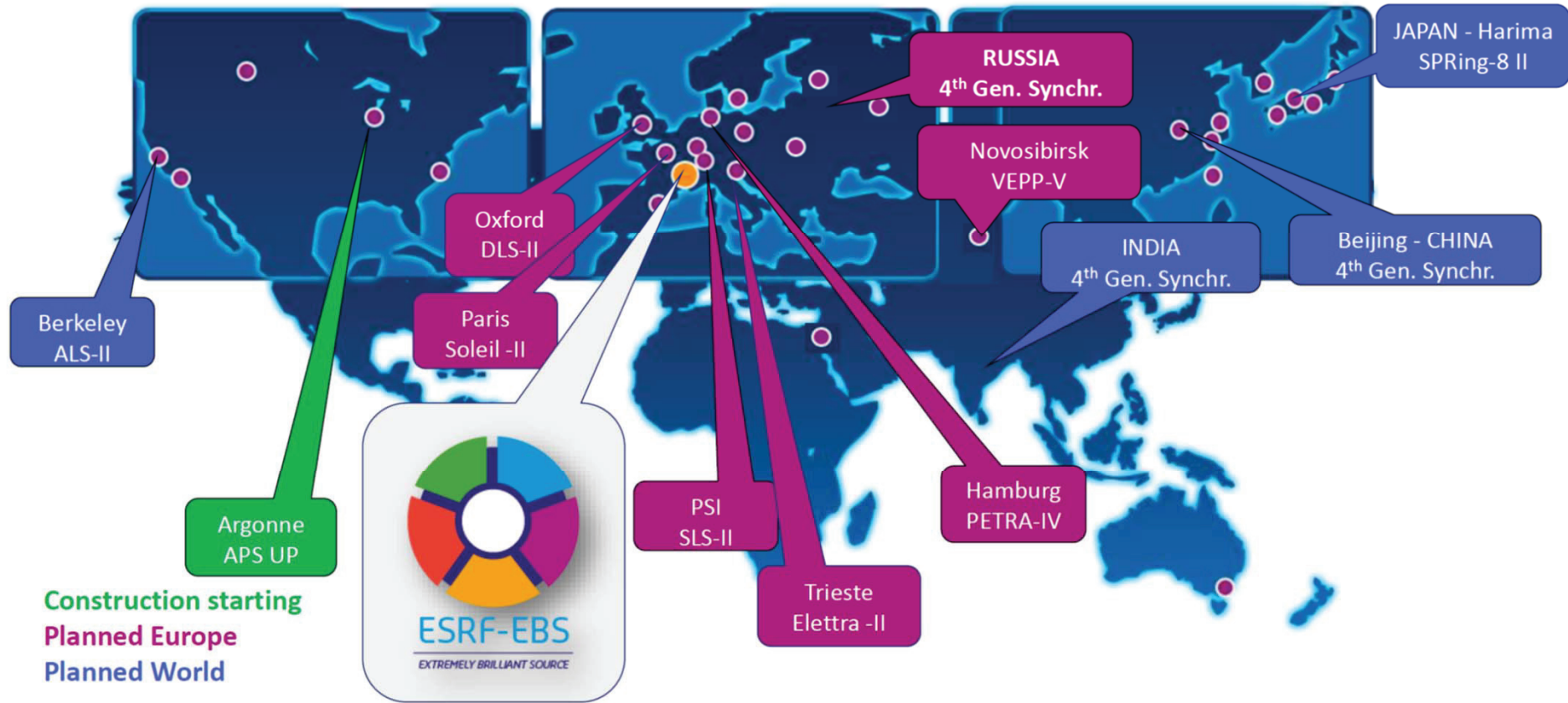
Conclusion



Why?
Because its light is
brilliant

There are now
more than 50 synchrotron light sources
In the world.

At least 13 upgrade programs in the world... ...to increase brilliance



Brilliance

Emittance

Coherence

Brilliance (or brightness) of a light Source

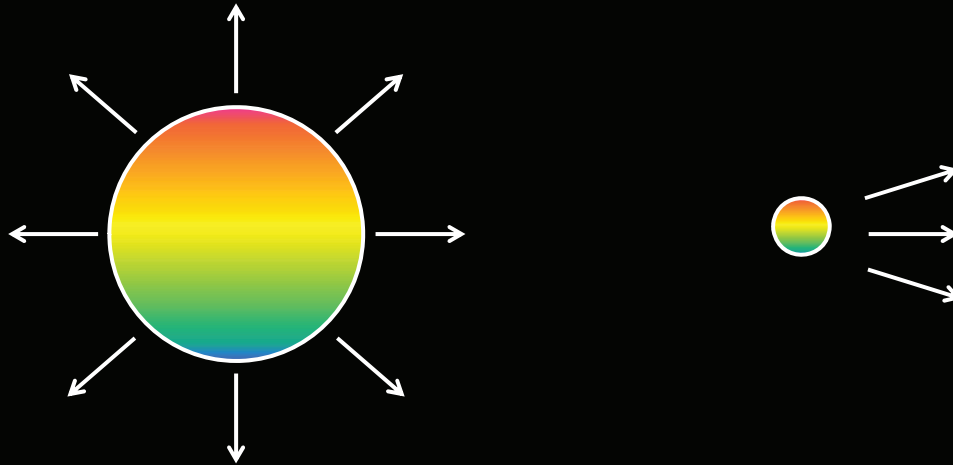
The more a source is :

Intense

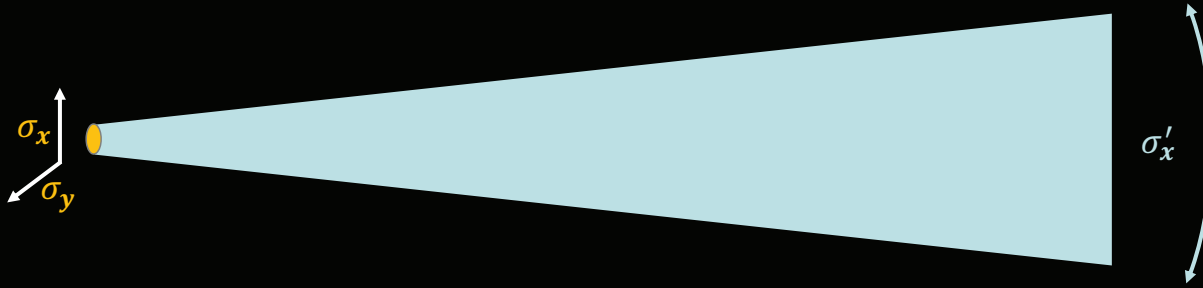
Small

Parallel

The more it is **brilliant**



Light source (and spectral brilliance)



Flux : in ph/s in $\frac{1}{1000}$ bandwidth

Source size σ_x, σ_y : in mm

Divergence σ'_x, σ'_y : in mrad

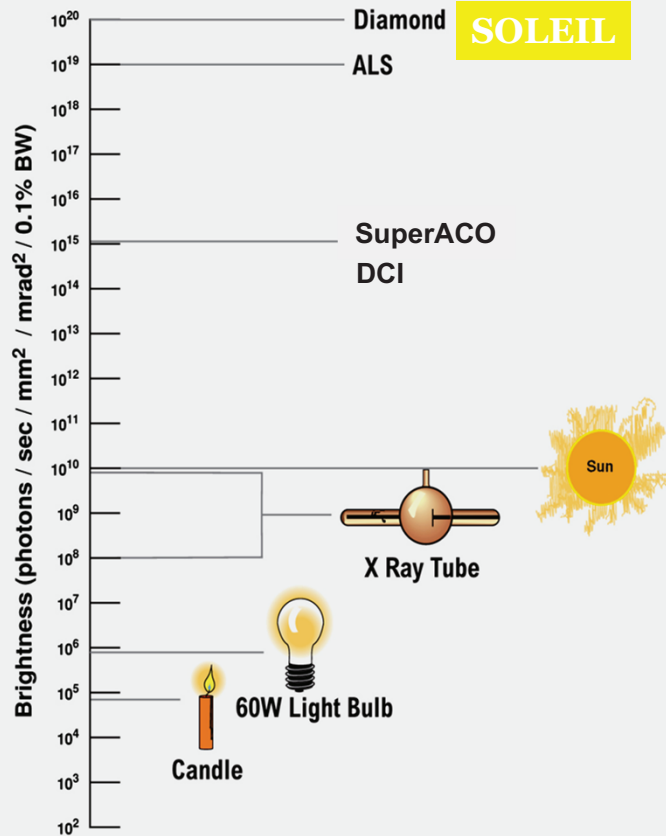
Spectral brilliance:

Flux divided by size and divergence of the source

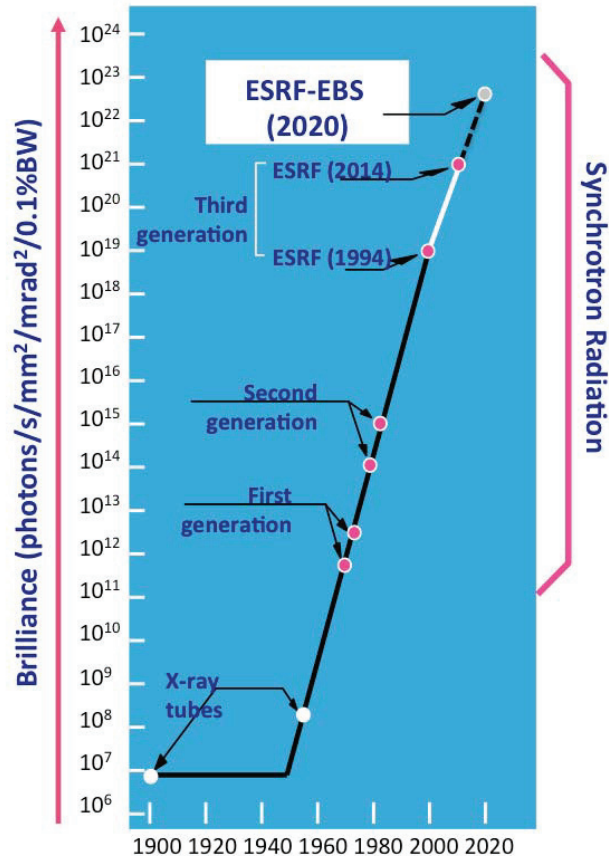
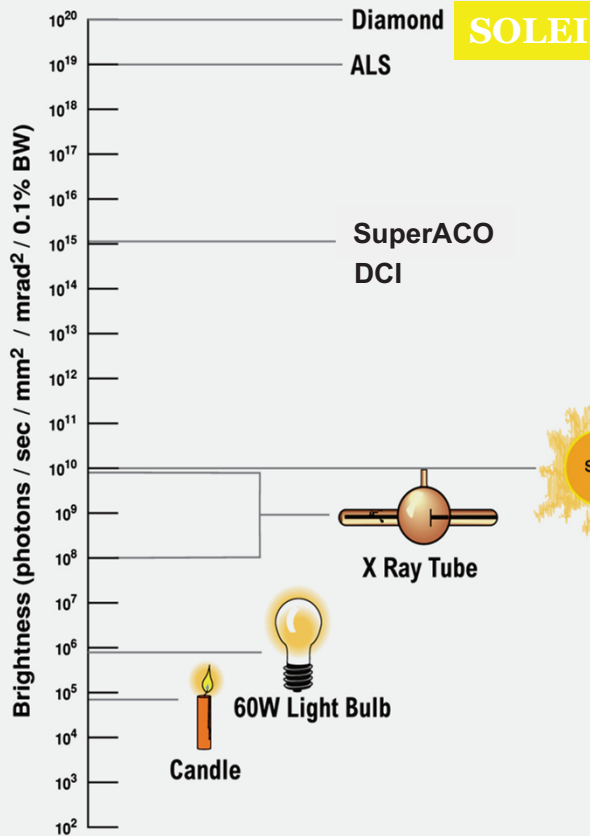
$$B = \frac{F}{\sigma_x \sigma'_x \sigma_y \sigma'_y}$$

$$\frac{[\text{ph/s}]}{[\text{mm}]^2 [\text{mrad}]^2 [0,1\% \text{ bandwidth}]} (= \text{UB})$$

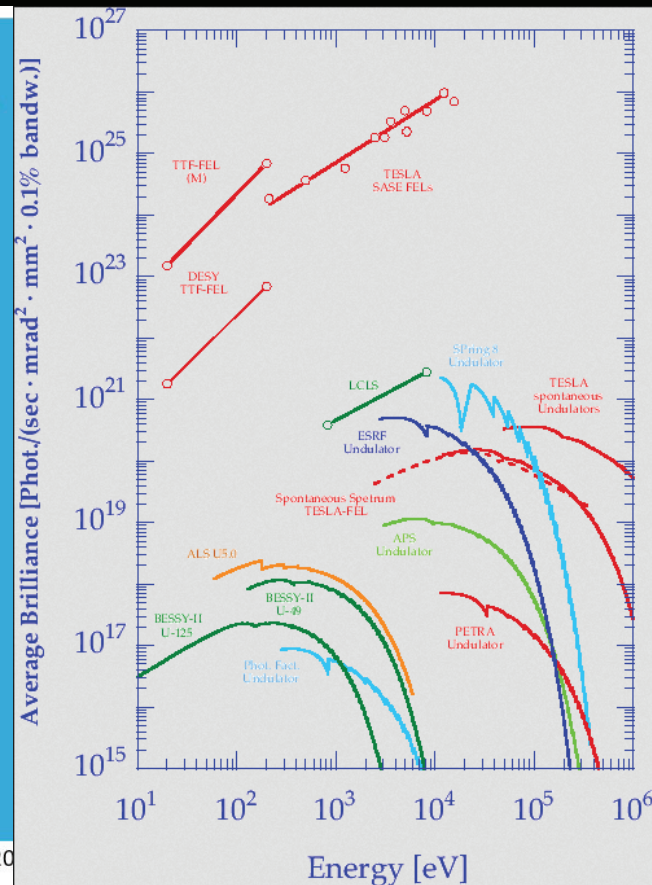
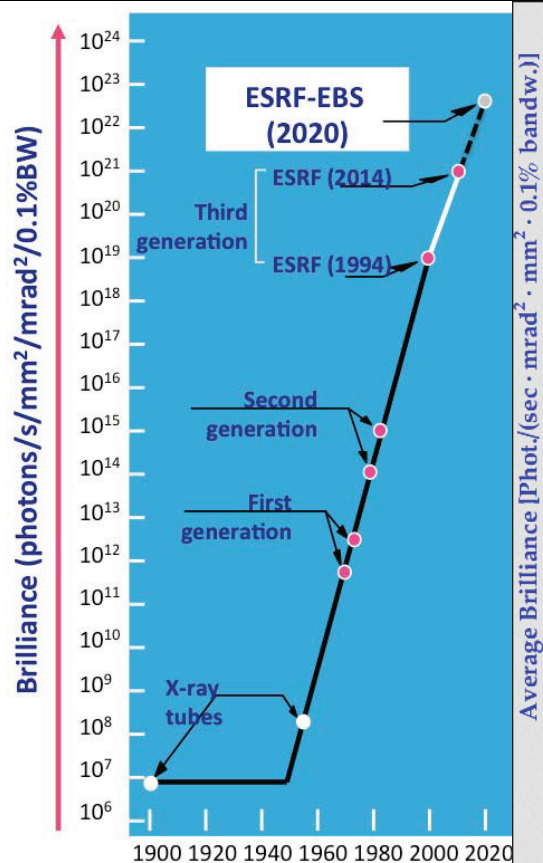
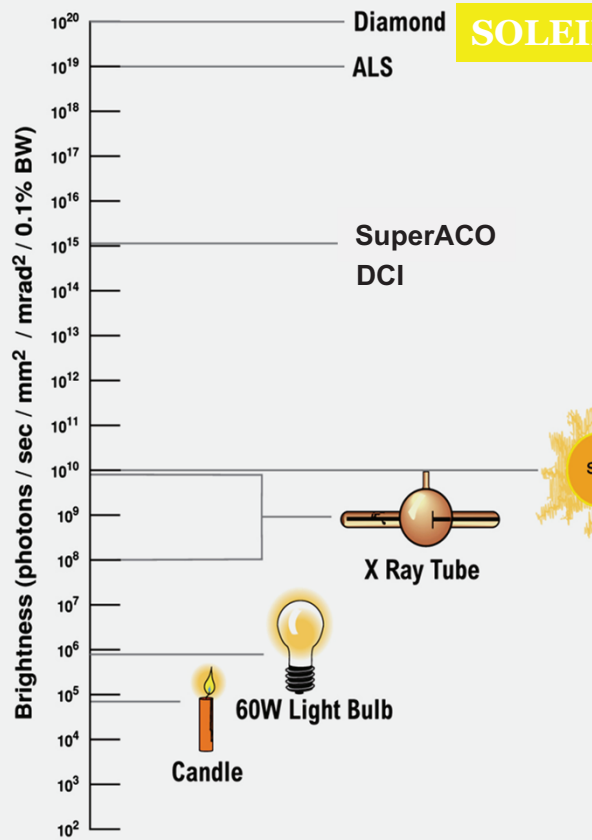
Brilliance: a figure of merit



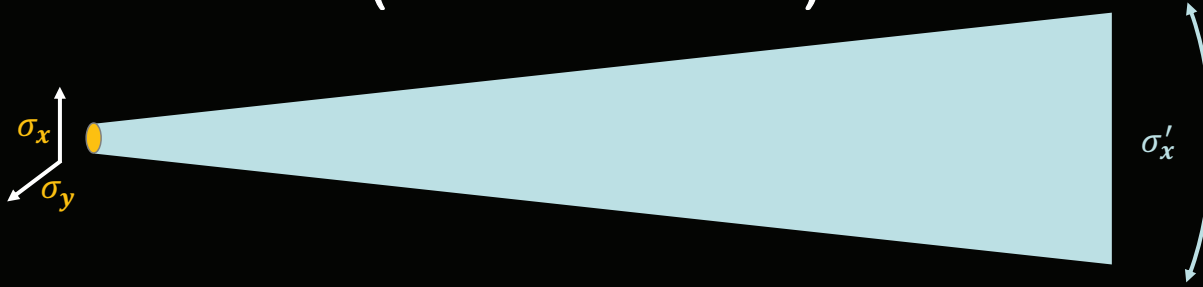
Brilliance: a figure of merit



Brilliance: a figure of merit



Light source (and emittance)



$$B = \frac{F}{\sigma_x \sigma'_x \sigma_y \sigma'_y}$$

Emittance: $\varepsilon_x = \sigma_x \sigma'_x$
Vertical x and horizontal y

$$B = \frac{F}{\varepsilon_x \varepsilon_y}$$

The smaller the emittance
The higher the brilliance

Brilliance and emittance

$$B = \frac{F}{\varepsilon_x \varepsilon_y}$$

Increasing the brilliance consists in decreasing the emittance
Yes, but there is a lower limit for the emittance:

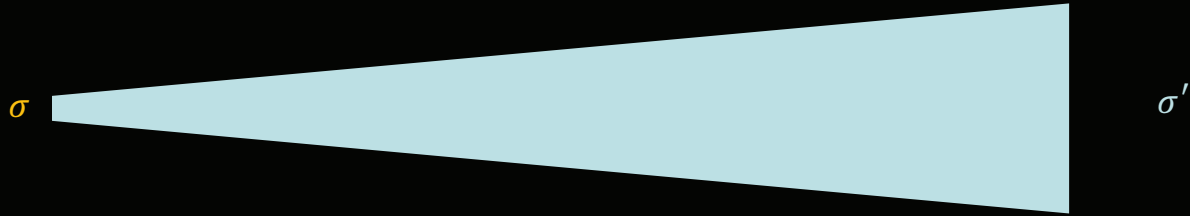
The limit of diffraction

Brilliance and emittance

$$B = \frac{F}{\varepsilon_x \varepsilon_y}$$

Increasing the brilliance consists in decreasing the emittance
Yes, but there is a lower limit for the emittance:

The limit of diffraction

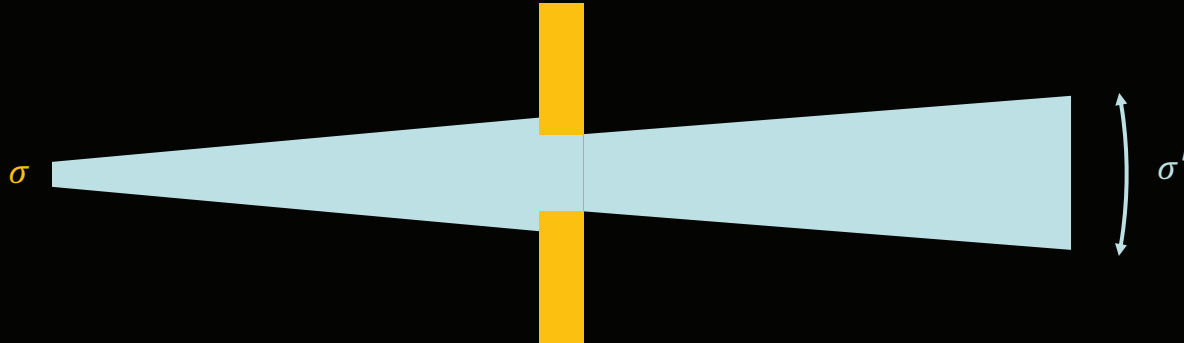


Brilliance and emittance

$$B = \frac{F}{\varepsilon_x \varepsilon_y}$$

Increasing the brilliance consists in decreasing the emittance
Yes, but there is a lower limit for the emittance:

The limit of diffraction

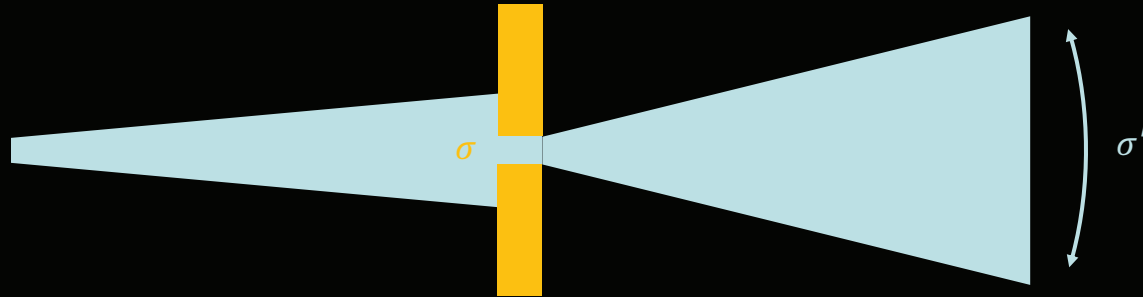


Brilliance and emittance

$$B = \frac{F}{\varepsilon_x \varepsilon_y}$$

Increasing the brilliance consists in decreasing the emittance
Yes, but there is a lower limit for the emittance:

The limit of diffraction

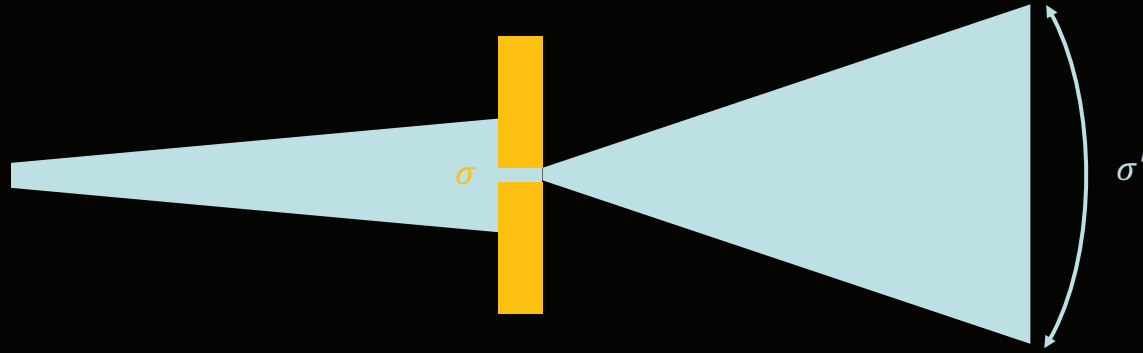


Brilliance and emittance

$$B = \frac{F}{\varepsilon_x \varepsilon_y}$$

Increasing the brilliance consists in decreasing the emittance
Yes, but there is a lower limit for the emittance:

The limit of diffraction

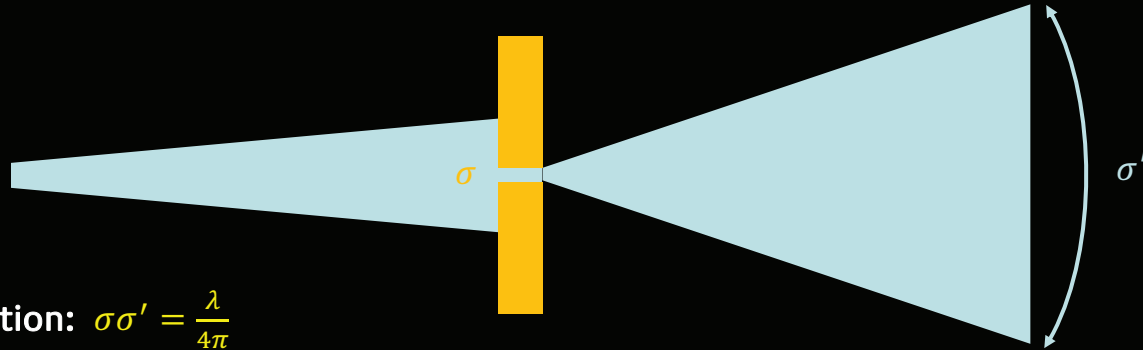


Brilliance and emittance

$$B = \frac{F}{\varepsilon_x \varepsilon_y}$$

Increasing the brilliance consists in decreasing the emittance
Yes, but there is a lower limit for the emittance:

The limit of diffraction



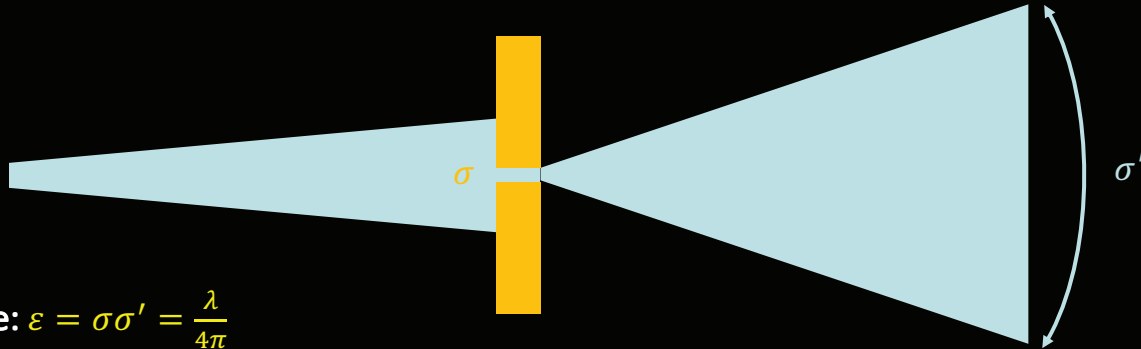
Theory of diffraction: $\sigma \sigma' = \frac{\lambda}{4\pi}$

Brilliance and emittance

$$B = \frac{F}{\varepsilon_x \varepsilon_y}$$

Increasing the brilliance consists in decreasing the emittance
Yes, but there is a lower limit for the emittance:

The limit of diffraction



Minimum emittance: $\varepsilon = \sigma \sigma' = \frac{\lambda}{4\pi}$

Rule of the thumb: **10 keV $\Leftrightarrow$ 10 pm.rad**

Diffraction limited or Extremely brilliant

Reaching this value would give **fully coherent** beam
And an extremely brilliant beam

Currently ESRF : 3800 pm.rad; APS 3000 pm.rad; SOLEIL 3700 pm.rad

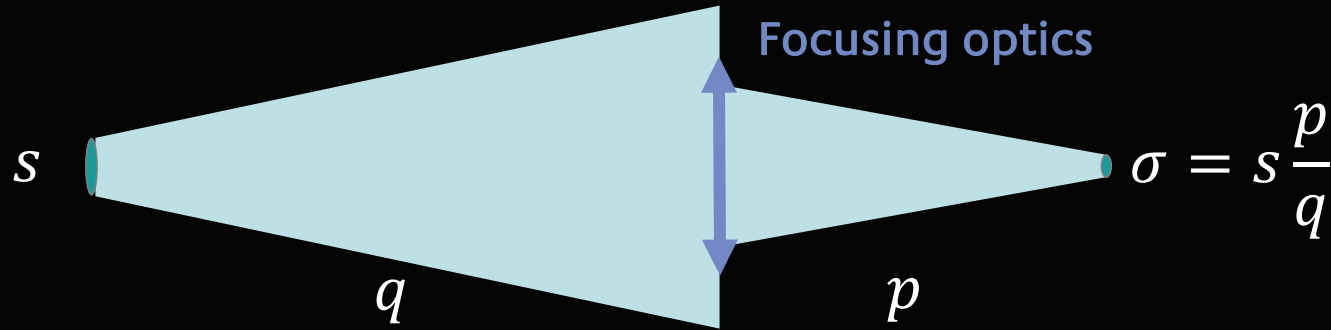
New possibilities using multi-bend achromat lattice
MAX IV 330 pm.rad; EBS-ESRF 134 pm.rad
Projects: SOLEIL 70 pm.rad; PETRA IV: 10 pm.rad

Increasing the brilliance: why?

Nanobeams

Coherence

Why increasing the brilliance?



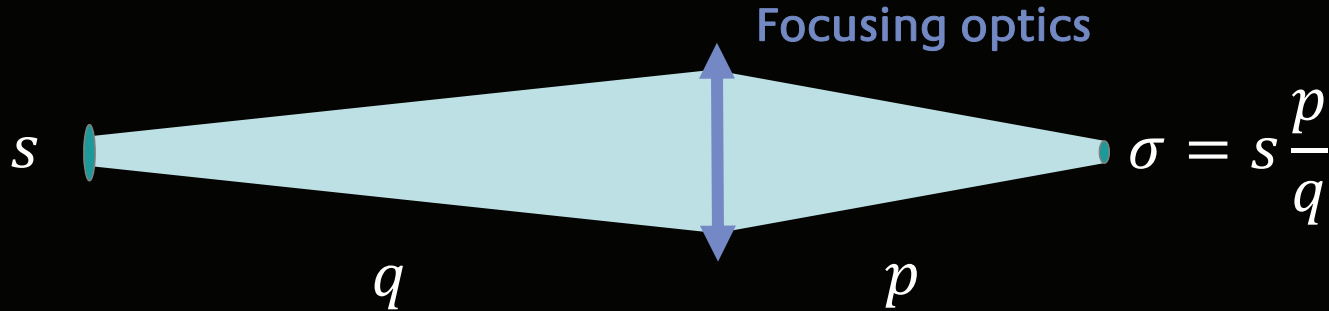
Beam is easier to **focus** (few 10 nm)

S is small (brilliance)

Adapted to small numerical aperture of optics (brilliance)

p/q is small (long beamlines);

Why increasing the brilliance?



Beam is easier to **focus** (few 10 nm)

s is small (brilliance)

Adapted to small numerical aperture of optics (brilliance)

p/q is small (long beamlines);

Diffraction micro- and nano-crystals ; extreme conditions

Scanning imaging (sub-micron, STXM)

Parallel beam (powder diffraction, SAXS)

Coherence

Characteristics of synchrotron light



Thomas Young

1773-1829



Augustin Fresnel

1788-1827

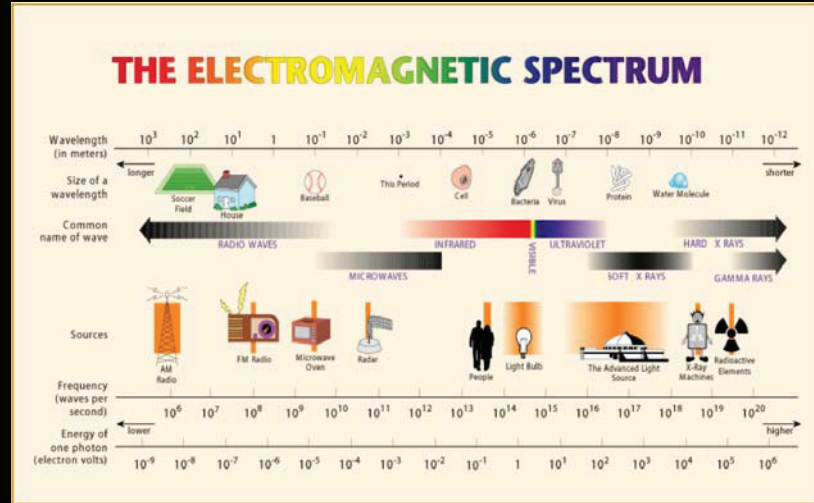
~ 1800

Waves are characterized by their:



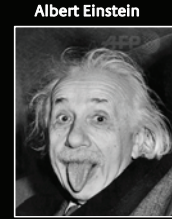
Wavelengths λ (Å, nm), frequencies ν (Hz)

Needed to interpret phenomena
where **scattering** is involved



Max Planck

1858-1947



Albert Einstein

1879-1955

~ 1900

Photons are characterized by their:



Energy $h\nu$ (eV)

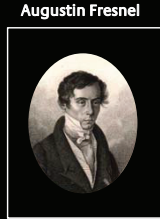
Needed to interpret phenomena
where **absorption** is involved

Characteristics of synchrotron light



Thomas Young

1773-1829



Augustin Fresnel

1788-1827

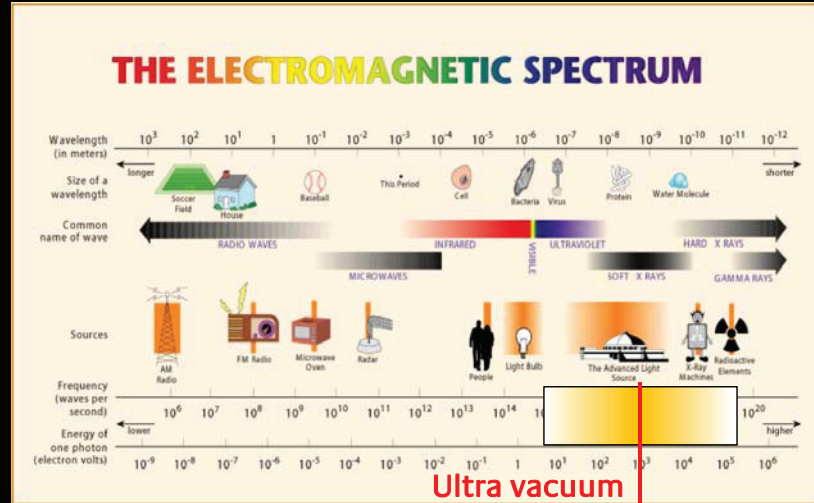
~ 1800

Waves are characterized by their:



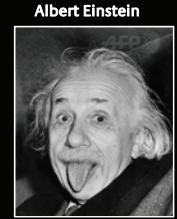
Wavelengths λ (Å, nm), frequencies ν (Hz)

Needed to interpret phenomena
where **scattering** is involved



Max Planck

1858-1947



Albert Einstein

1879-1955

~ 1900

Photons are characterized by their:



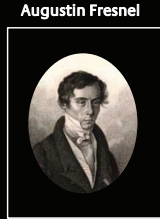
Energy $h\nu$ (eV)

Needed to interpret phenomena
where **absorption** is involved

Characteristics of synchrotron light

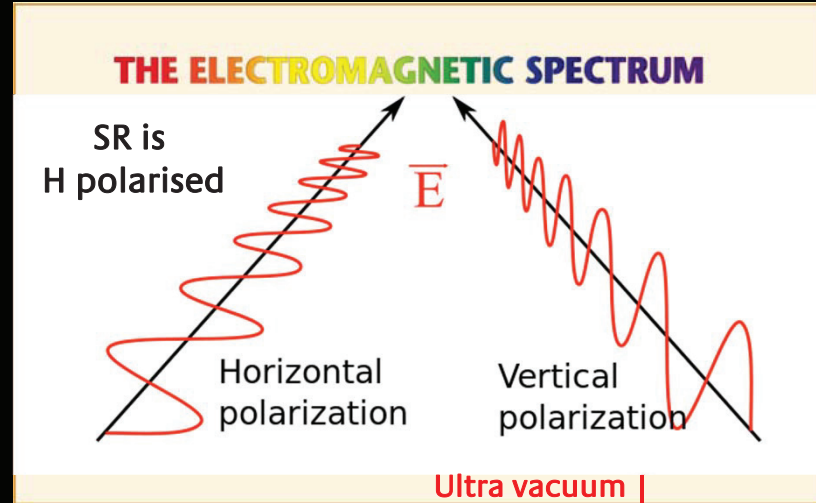


1773-1829

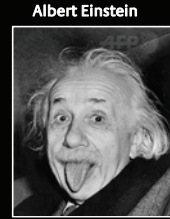


1788-1827

~ 1800



1858-1947



1879-1955

~ 1900

Waves are characterized by their:



Wavelengths λ (Å, nm), frequencies ν (Hz)

Needed to interpret phenomena
where **scattering** is involved

Photons are characterized by their:



Energy $h\nu$ (eV)

Needed to interpret phenomena
where **absorption** is involved

Complementarity

Two families of beamlines

Beamlines using scattering (and diffraction)

- Diffraction beamlines (MX)
- Small angle scattering (SAXS)
 - Surface diffraction

Beamlines using absorption (and associated phenomena)

- X-ray Absorption Spectroscopy
(XAS-EXAFS-XANES)
- Photoelectron spectroscopy
(PES-ARPES)

Imaging beamlines

Can use both!

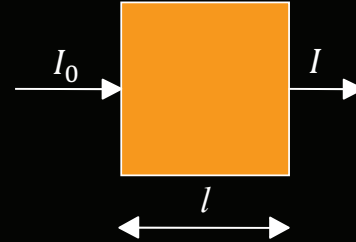
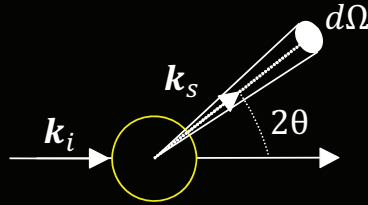
XPEEM, STXM, Tomography

Coherent diffraction imaging, Ptychography

Two processes of interaction between x-ray and matter

Scattering and Absorption

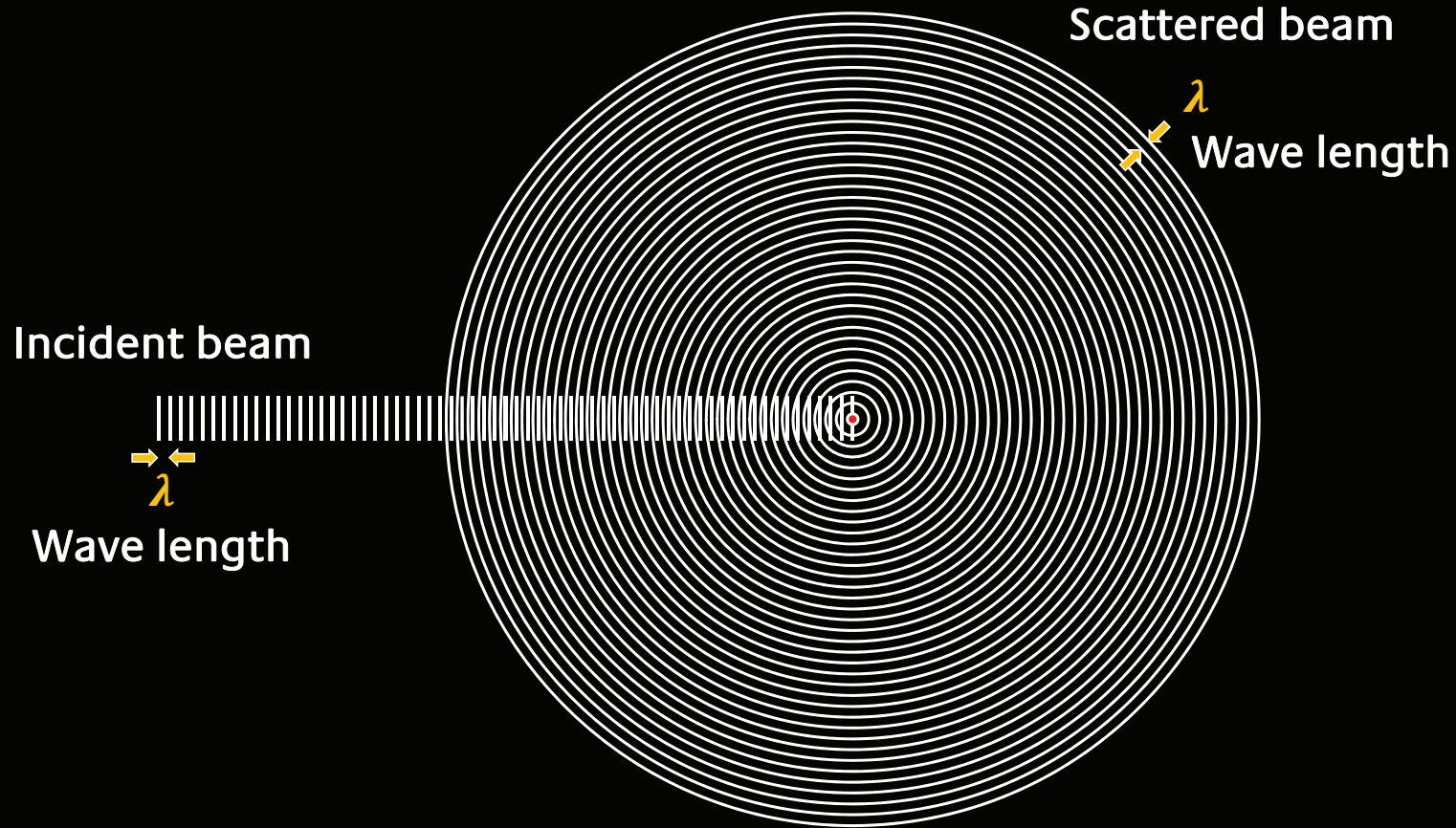
$$k = \frac{2\pi}{\lambda}$$



$$I = I_0 e^{-\mu l}$$

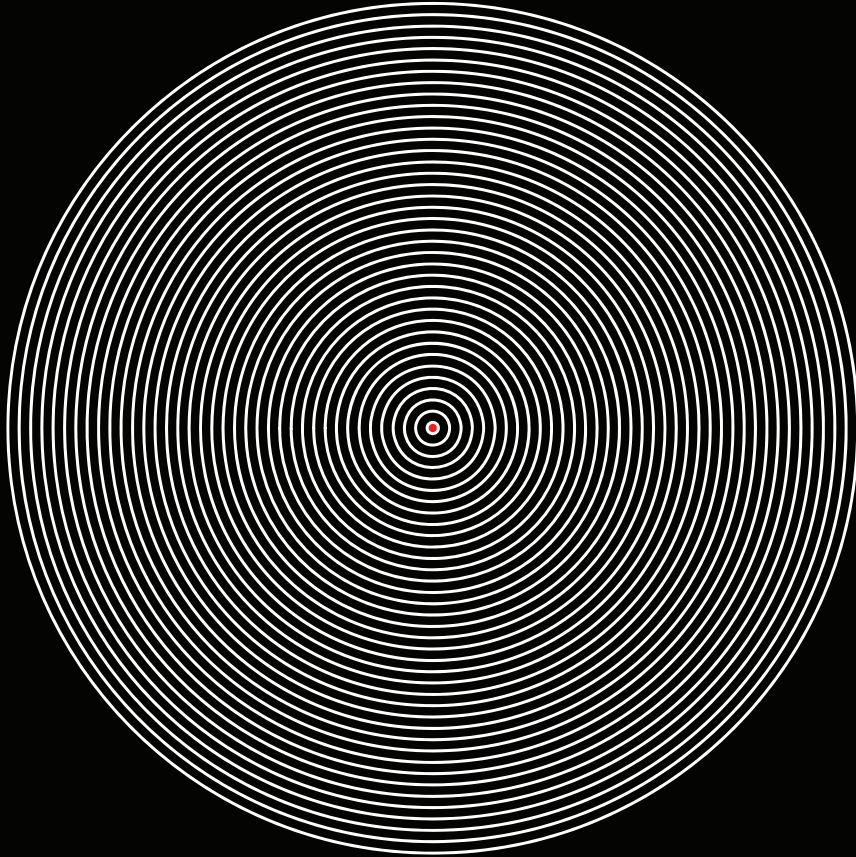
Scattering and diffraction

Scattering



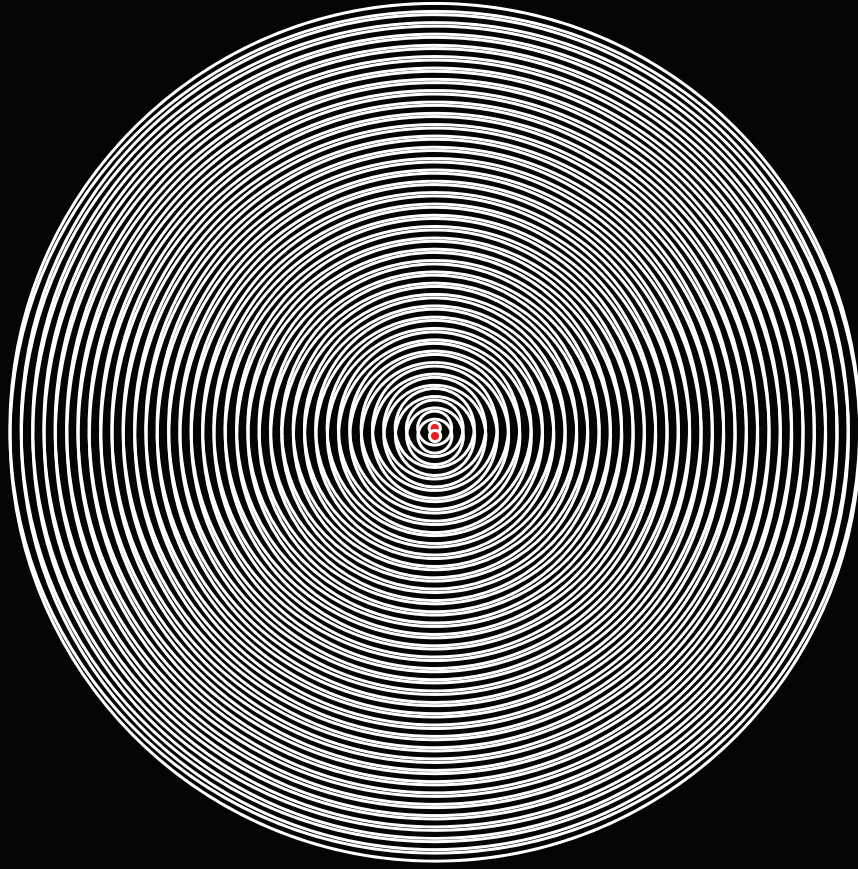
Diffraction: principle

Interference



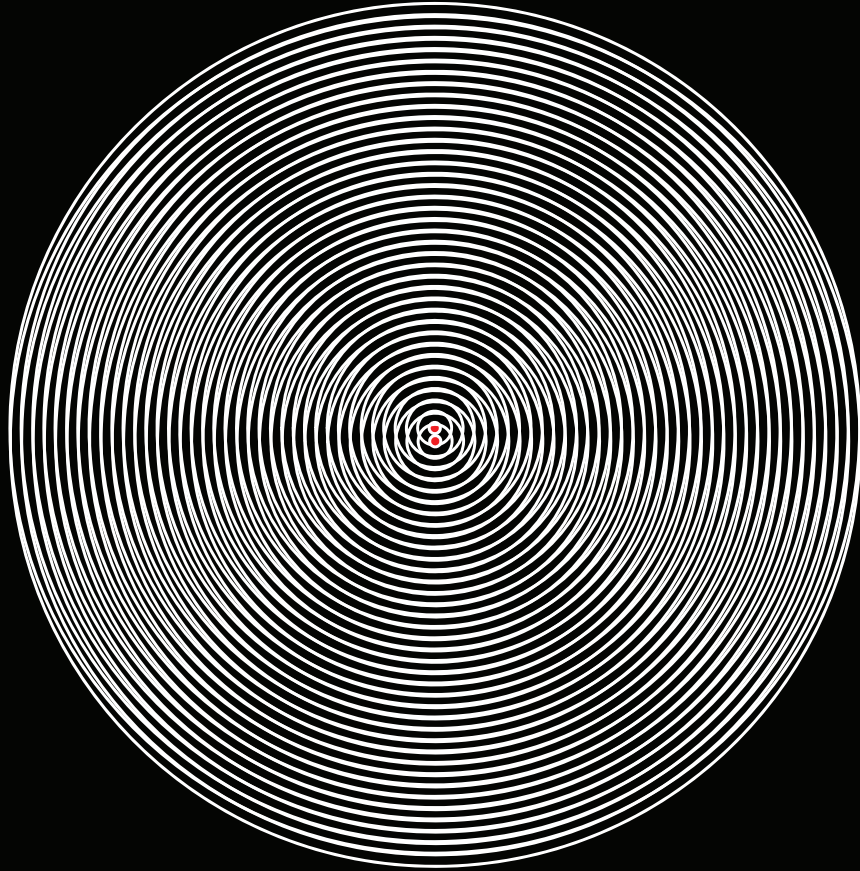
Diffraction: principle

Interference



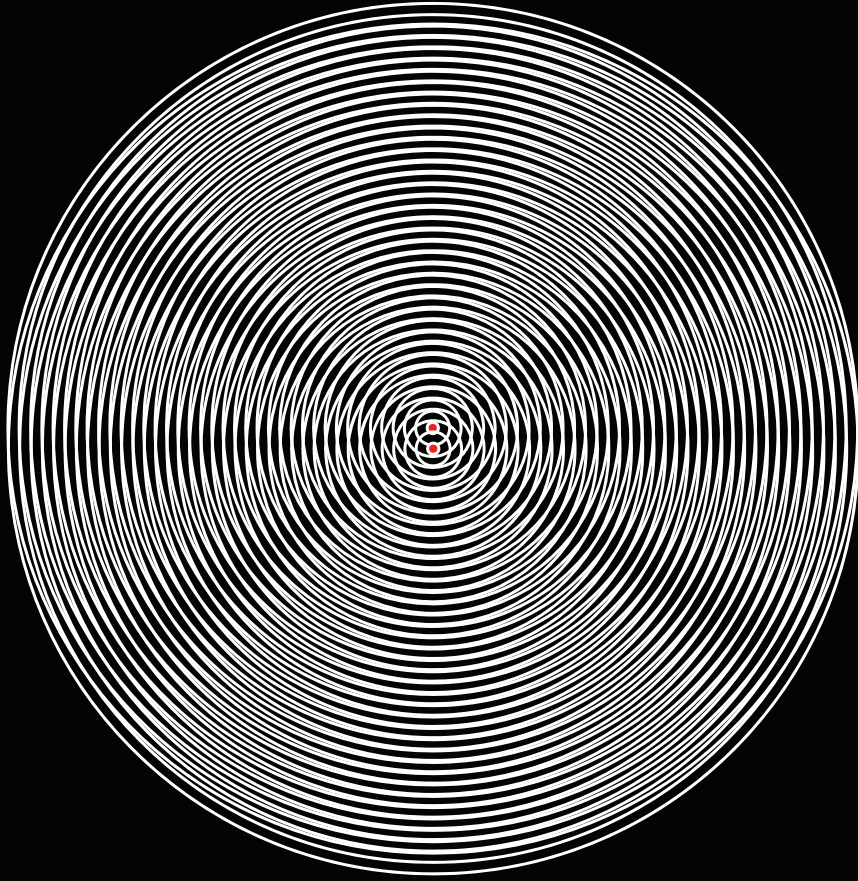
Diffraction: principle

Interference



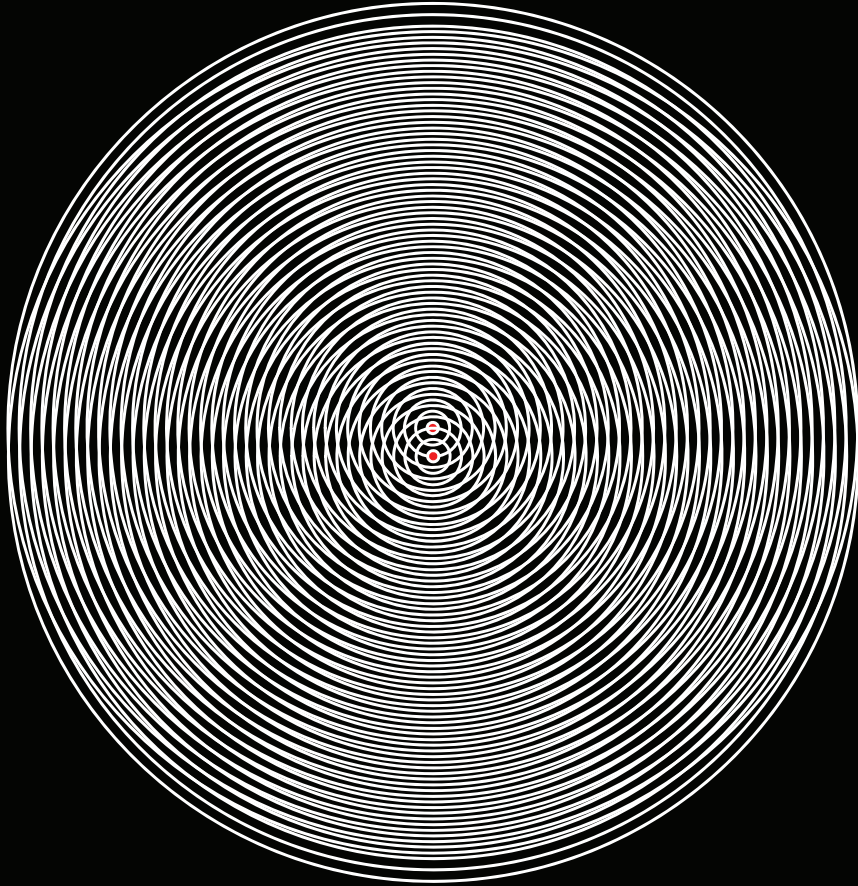
Diffraction: principle

Interference



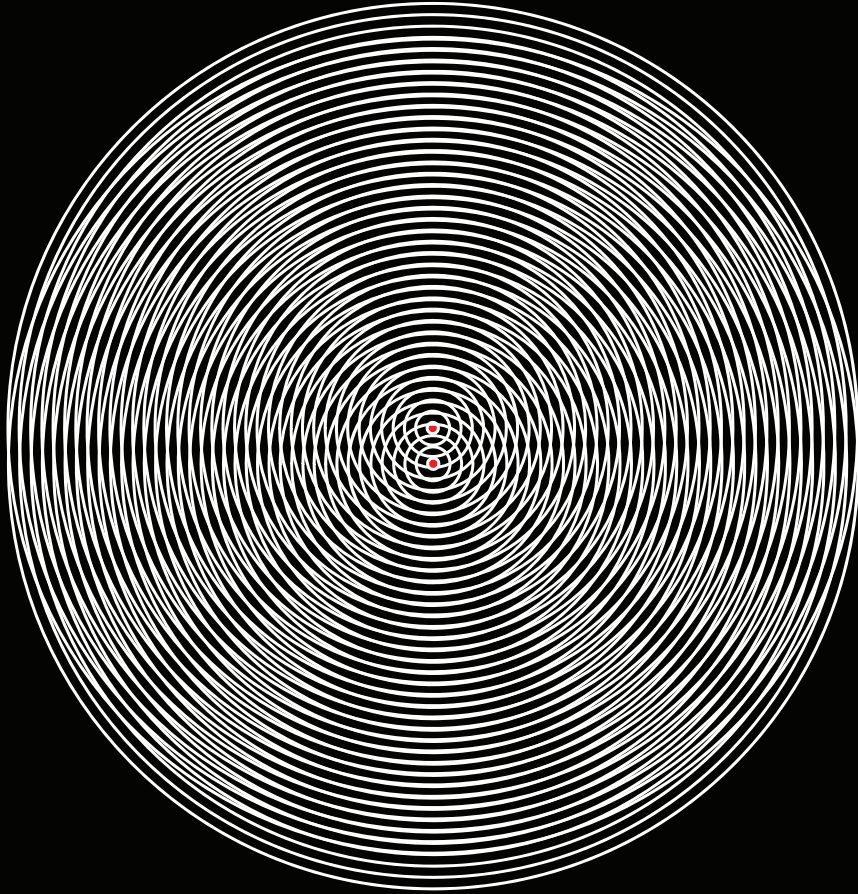
Diffraction: principle

Interference



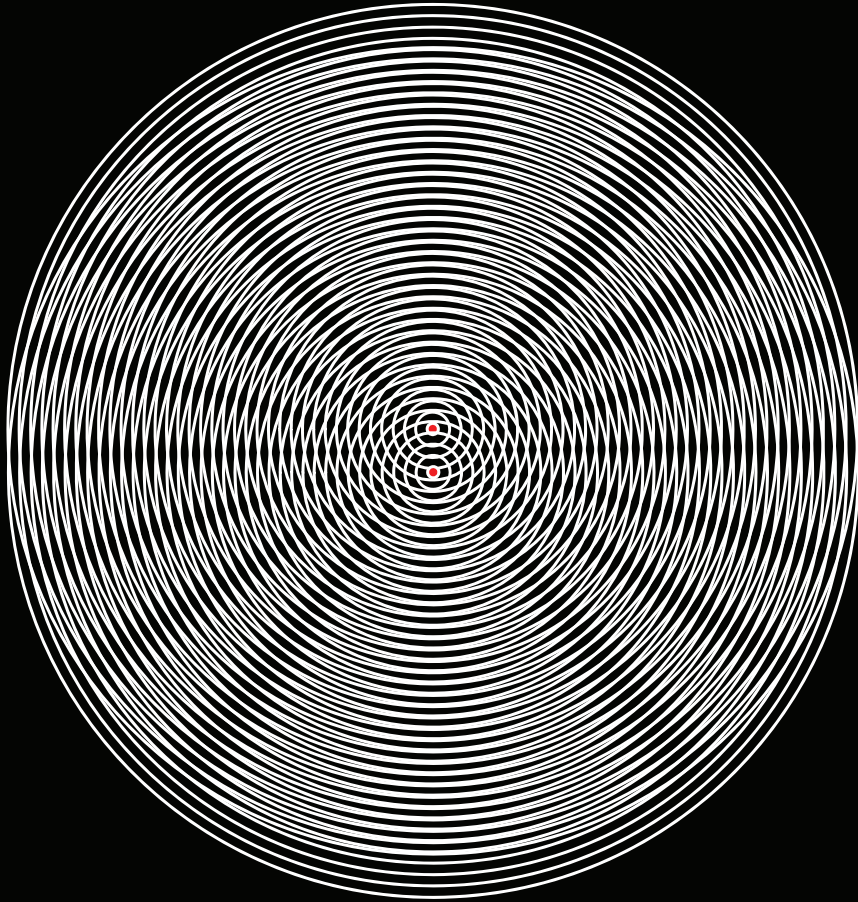
Diffraction: principle

Interference



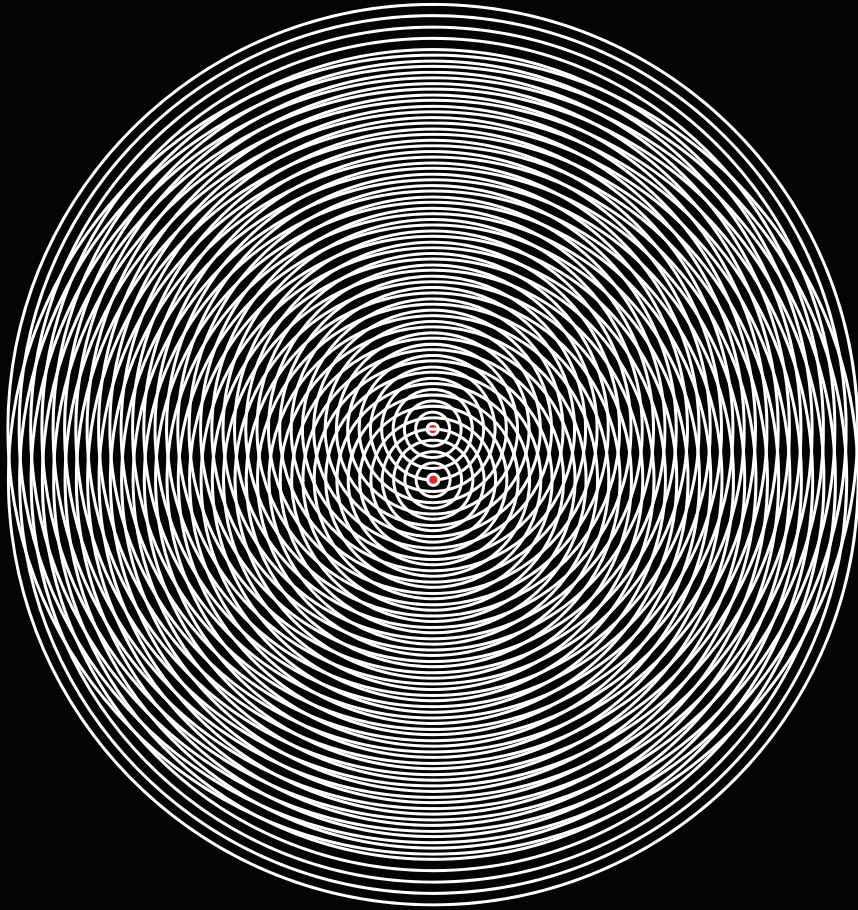
Diffraction: principle

Interference



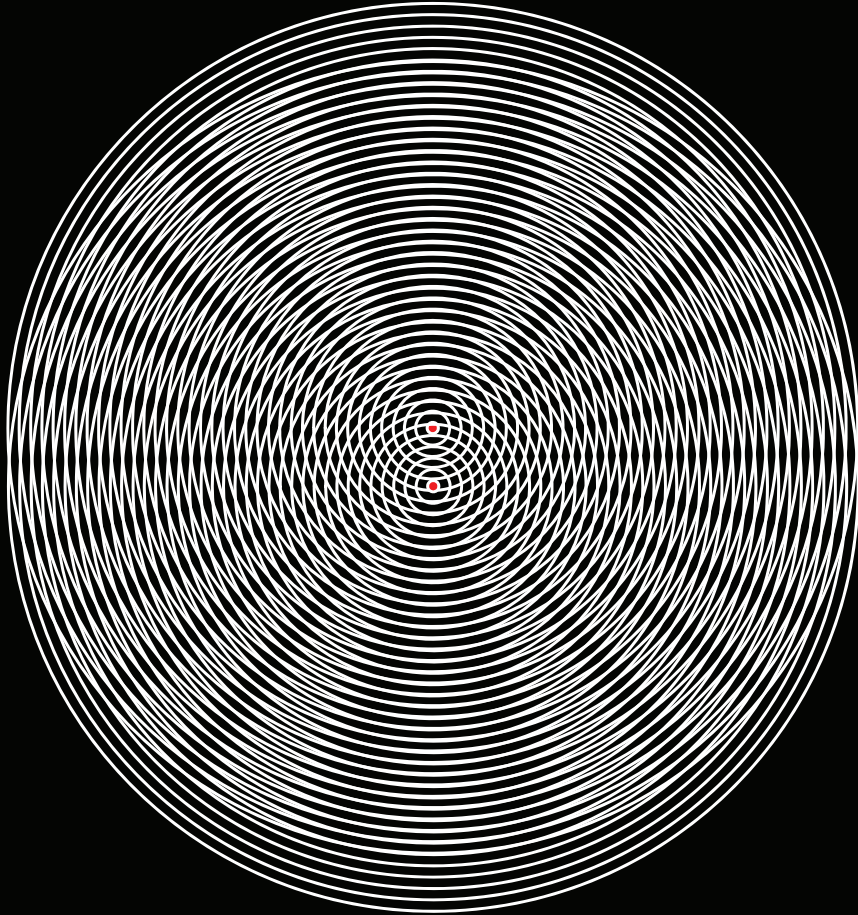
Diffraction: principle

Interference



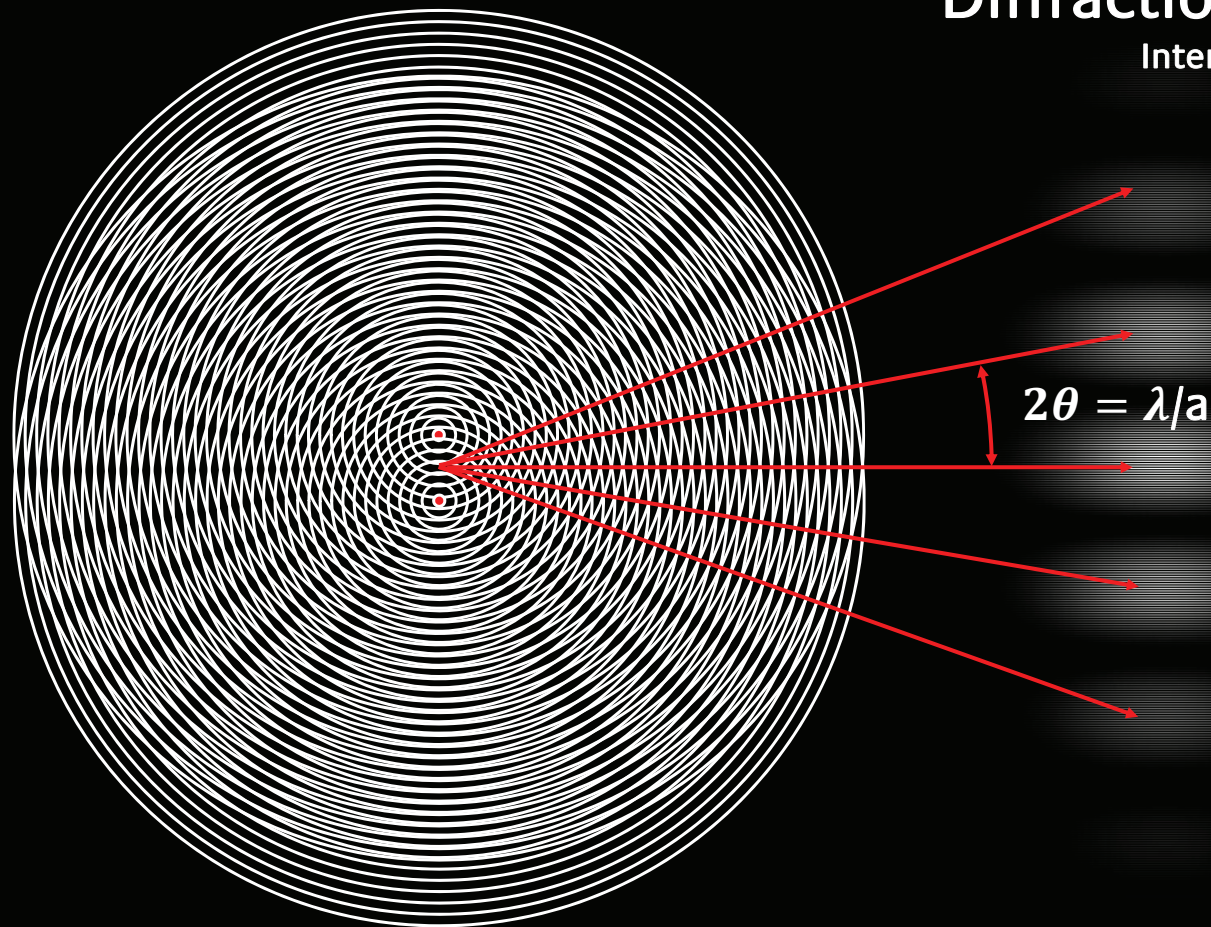
Diffraction: principle

Interference



Diffraction: principle

Interference

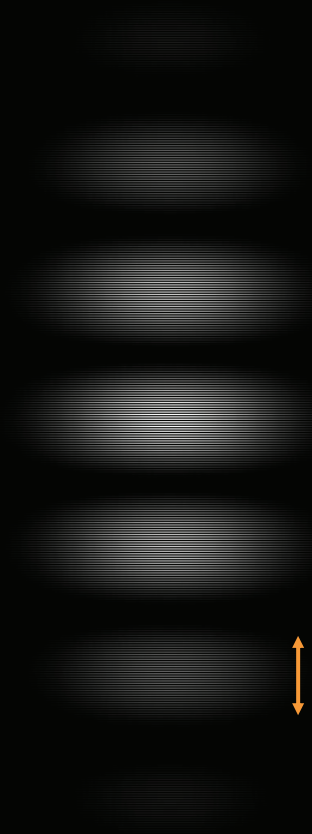


To measure
distance: a
measure
angle: 2θ

Diffraction
is essentially
Goniometry
(and detection)

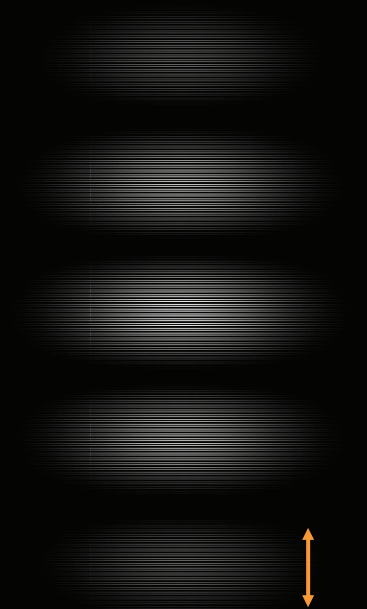
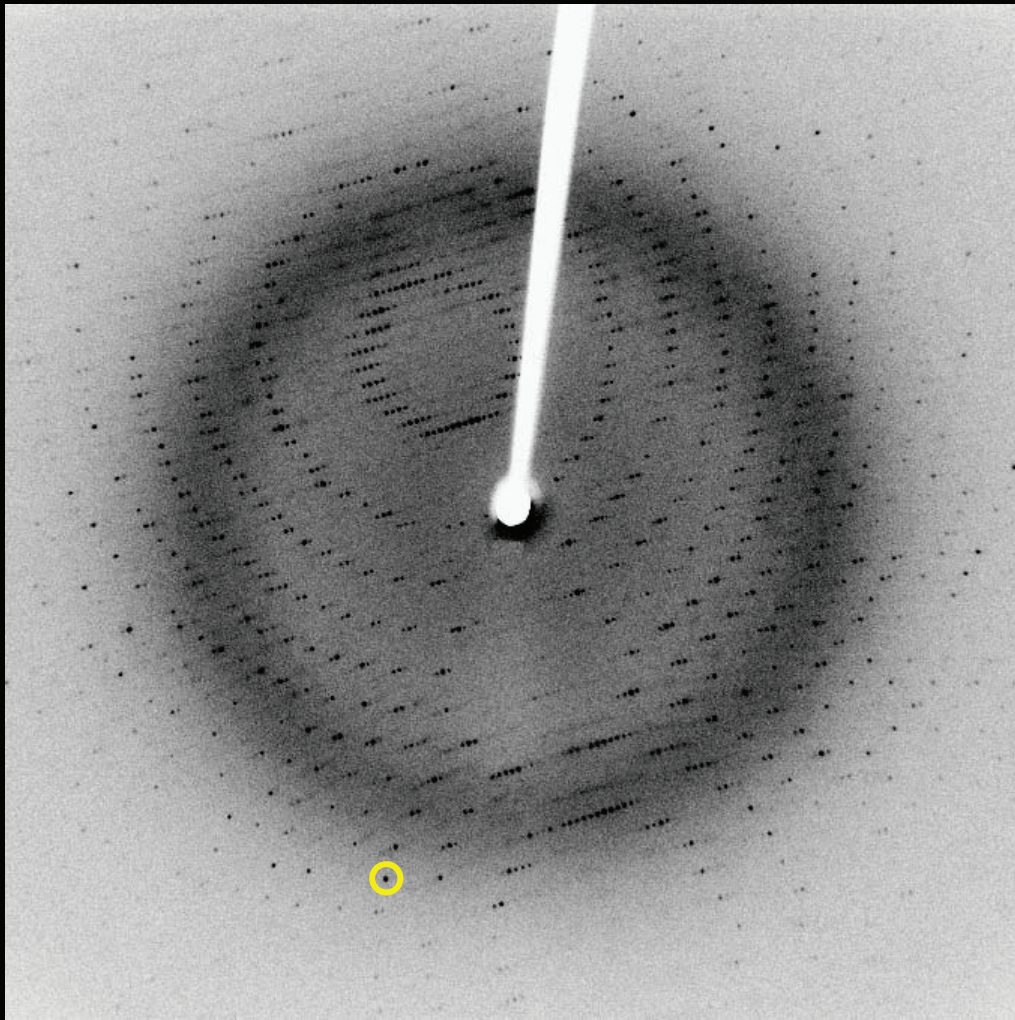
To measure
angles
**as precisely
as possible**

Diffraction: principle



With 2 atoms
spots **are broad**

Diffraction: principle



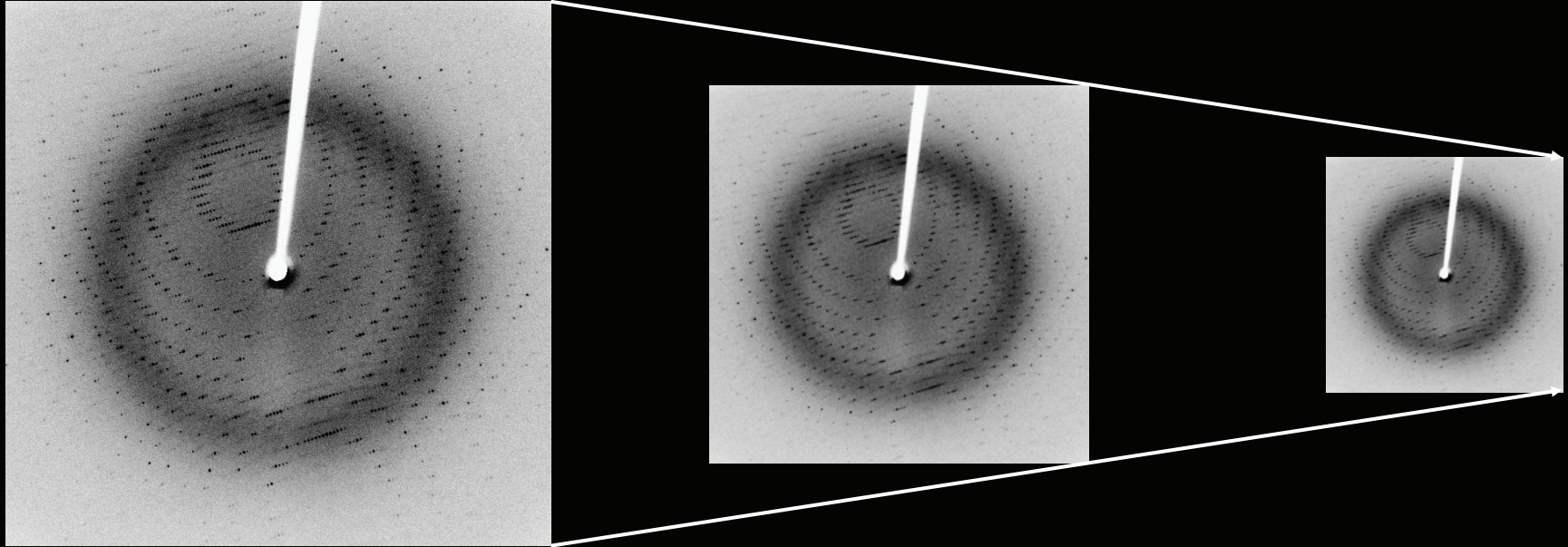
With 2 atoms
spots **are broad**

With many as in a crystal
very narrow ($5 \mu\text{rad} = \text{arc s}$)

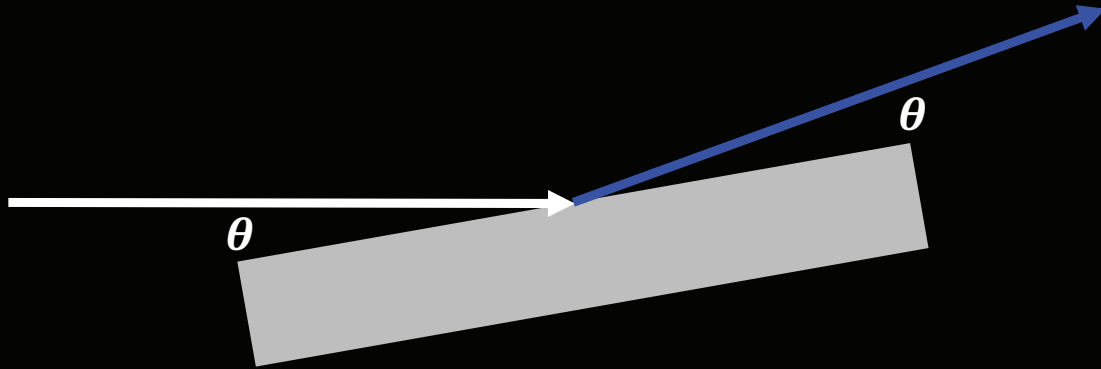
Diffraction: principle

Because $2\theta = \lambda/a$

Diffraction is **more difficult** at high energy (small λ)



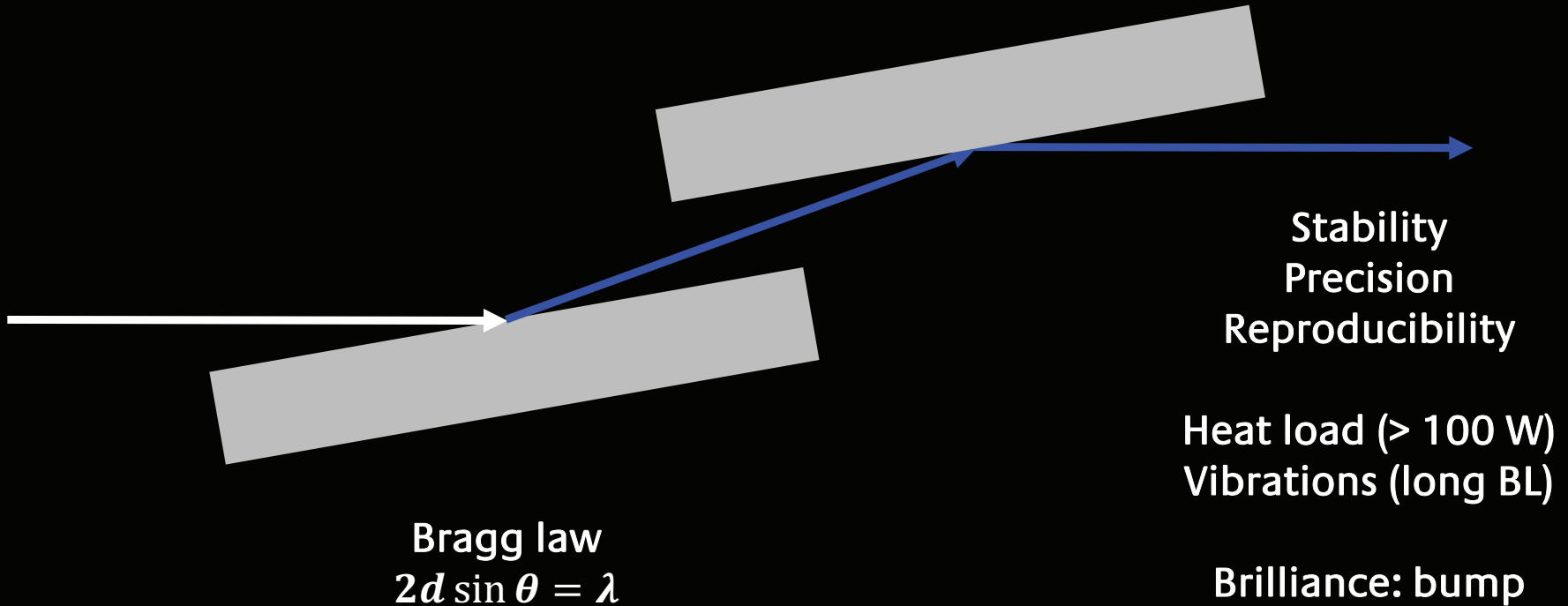
Monochromator: principle



Bragg law
 $2d \sin \theta = \lambda$

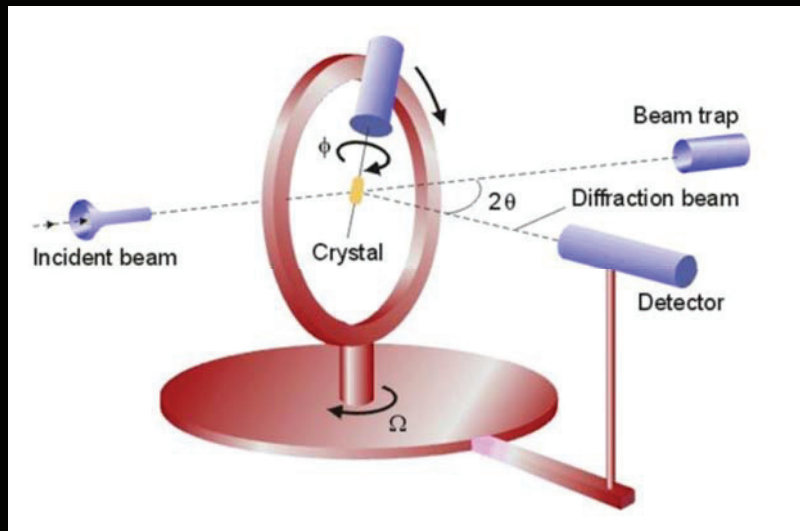
Monochromator: principle

Double Crystal Monochromator



Diffraction: the 4-circle diffractometer

An assembly of 4 goniometers to adjust crystal (3) and detector (1)



High **precision** goniometers (1 arc sec)
Sphere of Confusion (5 μm or better)
or Cylinder of Confusion

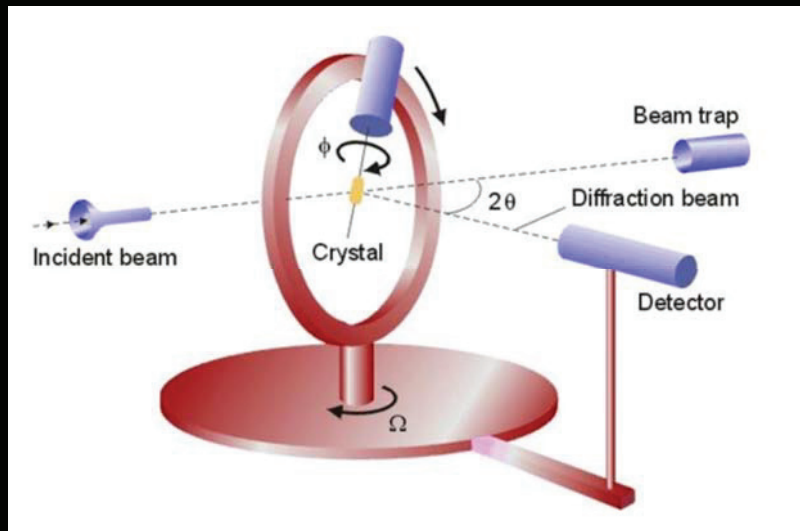
Euler
Vs
Kappa



More circles for complex **sample environments**
Nano beams, very small SOC
Detectors are more and more heavy (arms)
Because of H polarization, better to work in vertical plane

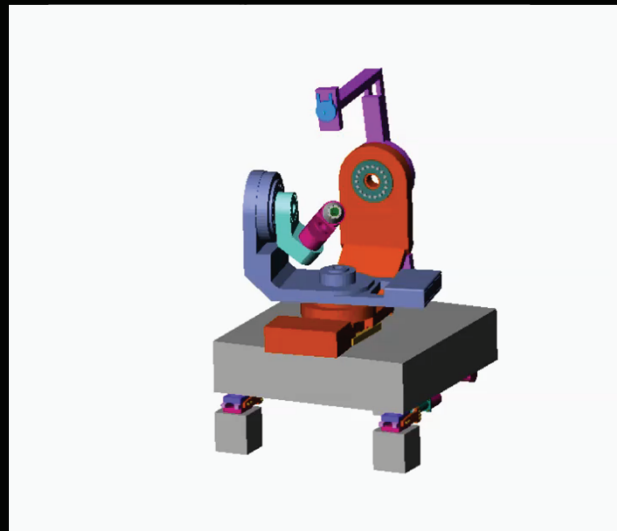
Diffraction: the 4-circle diffractometer

An assembly of 4 goniometers to adjust crystal (3) and detector (1)



High **precision** goniometers (1 arc sec)
Sphere of Confusion (5 μm or better)
or Cylinder of Confusion

Euler
Vs
Kappa



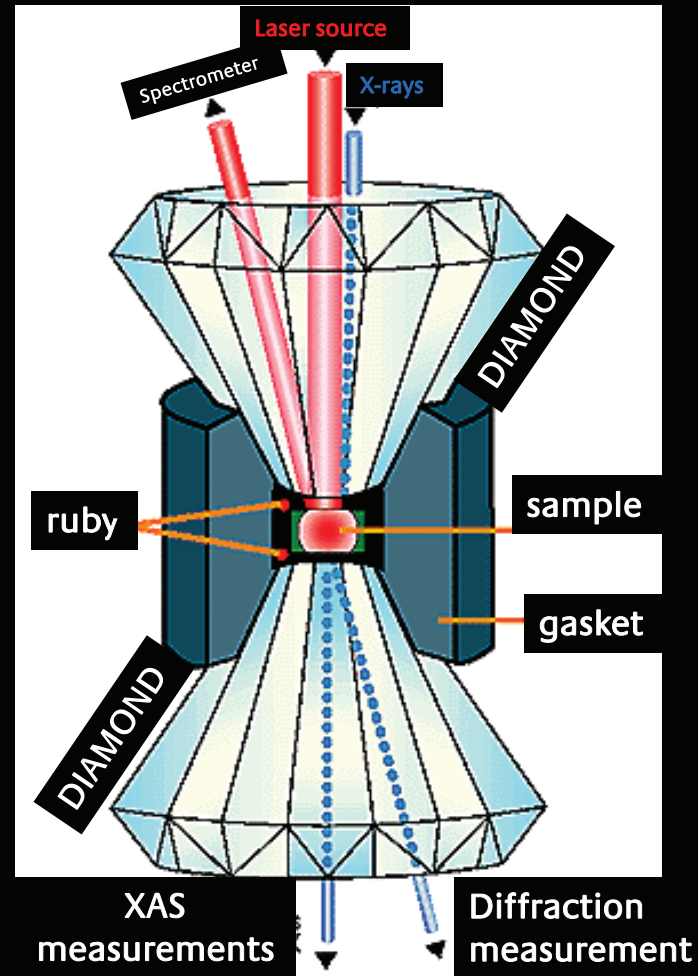
More circles for complex **sample environments**
Nano beams, very small SOC
Detectors are more and more heavy (arms)
Because of H polarization, better to work in vertical plane

Nano-beams are necessary to reach
Ultra high pressure
In Diamond anvil cells

The higher the pressure
The smaller the aperture
The smaller the beam

Mechanical challenge:

Low (high) temperature
High pressure



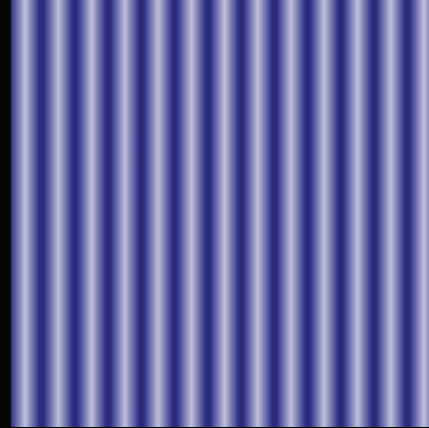
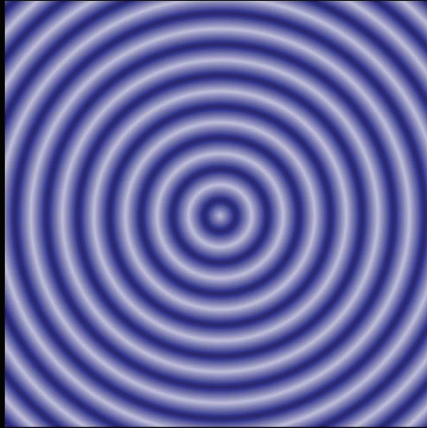
Brilliance

Emittance

Coherence

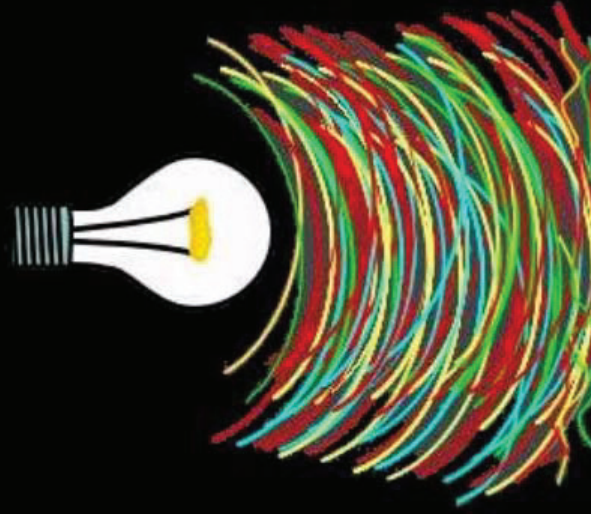
What is coherence?

Light wave is coherent if it looks like a perfect spherical or plane wave



What is coherence?

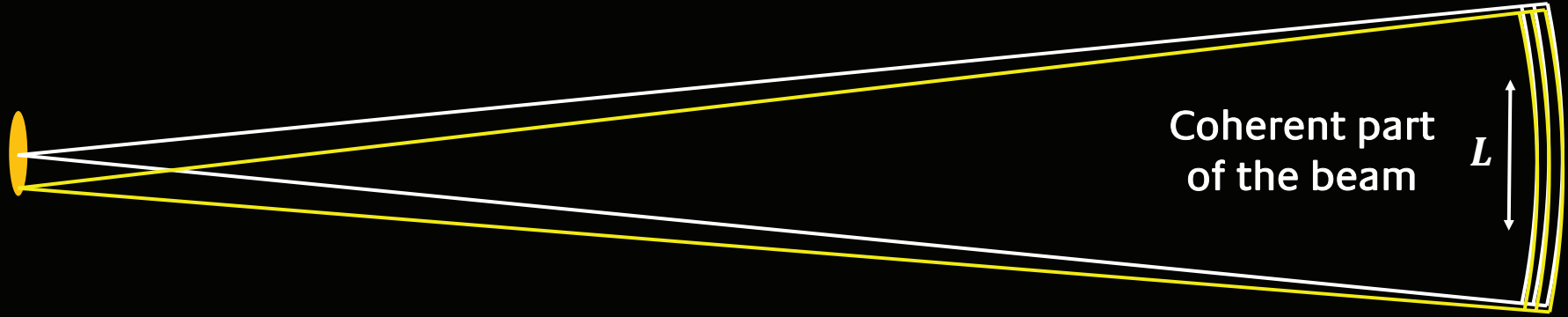
Unfortunately, light (even synchrotron light) looks like that:



The longitudinal coherence is given by the **wave length purity** (monochromator)

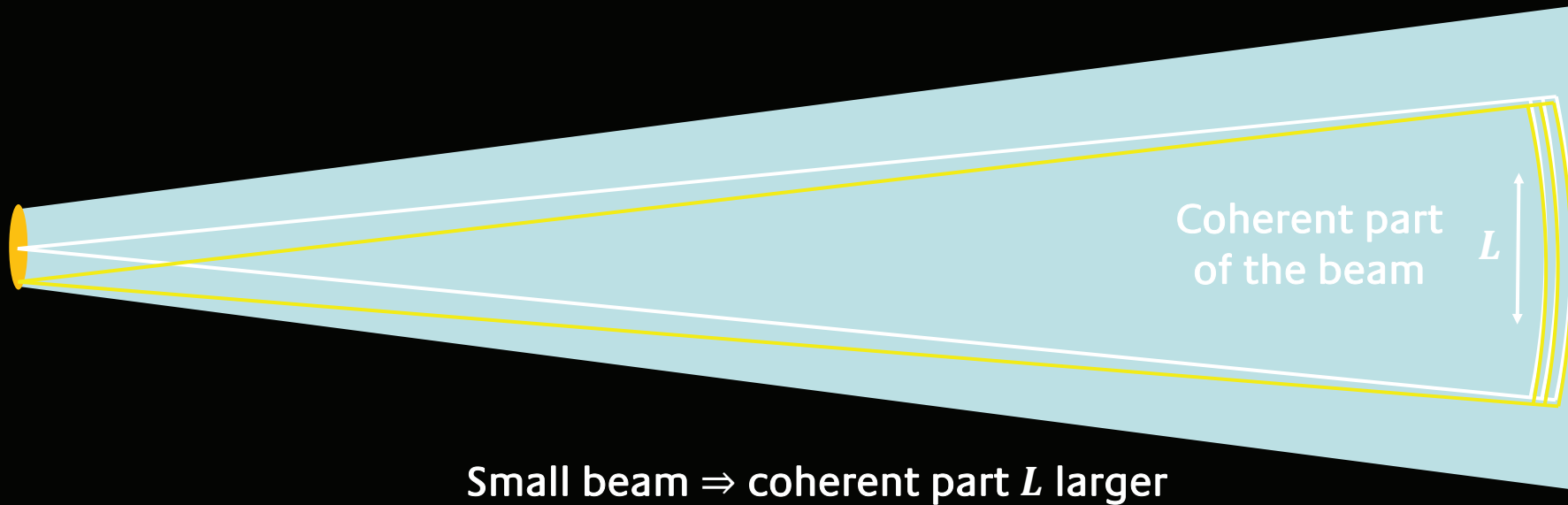
The transverse coherence depends on the **size of the source**

Brilliance vs Coherence

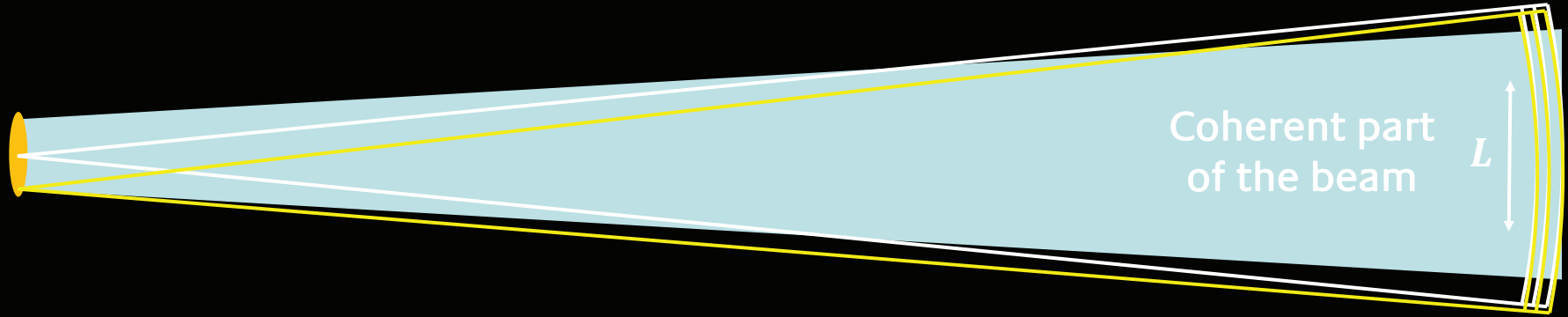


Small beam $\Rightarrow$ coherent part L larger

Brilliance vs Coherence

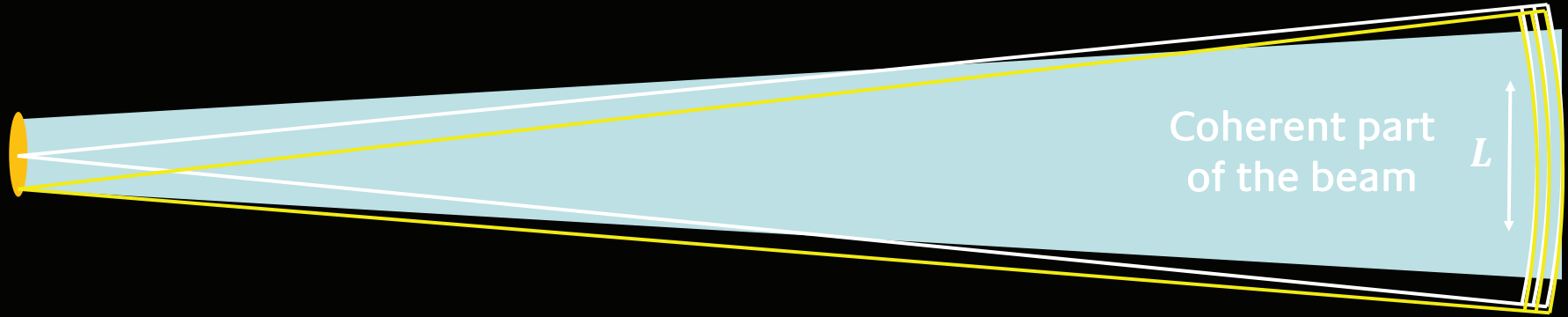


Brilliance vs Coherence



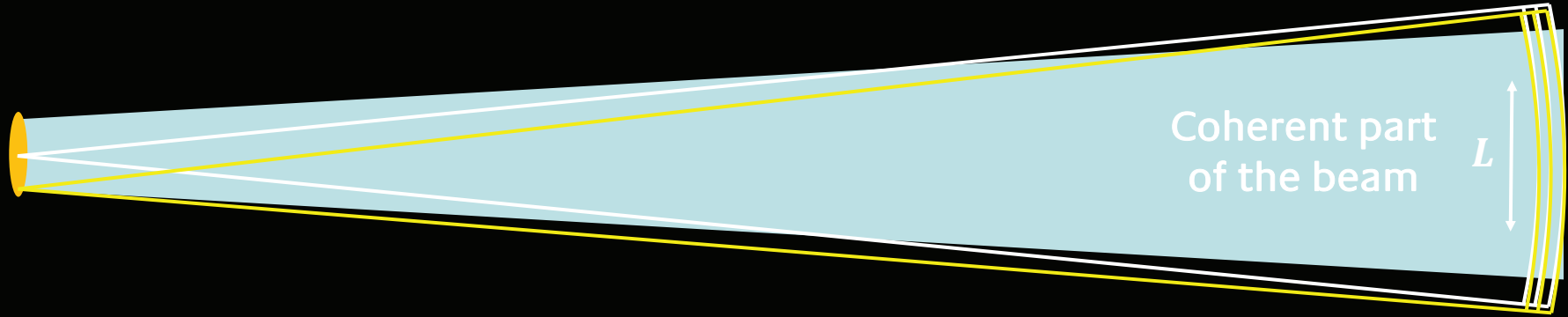
Small beam $\Rightarrow$ coherent part L larger

Brilliance vs Coherence



Small beam $\Rightarrow$ coherent part L larger
Small divergence $\Rightarrow$ more flux in the coherent part

Brilliance vs Coherence



Small beam $\Rightarrow$ coherent part L larger

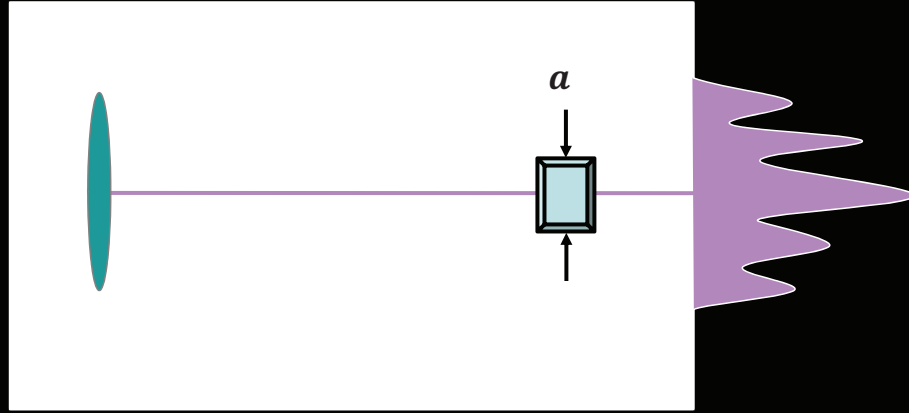
Small divergence $\Rightarrow$ more flux in the coherent part

More brilliance $\Rightarrow$ More coherence!

$$F_{coh} \propto B\lambda^2$$

Why coherence?

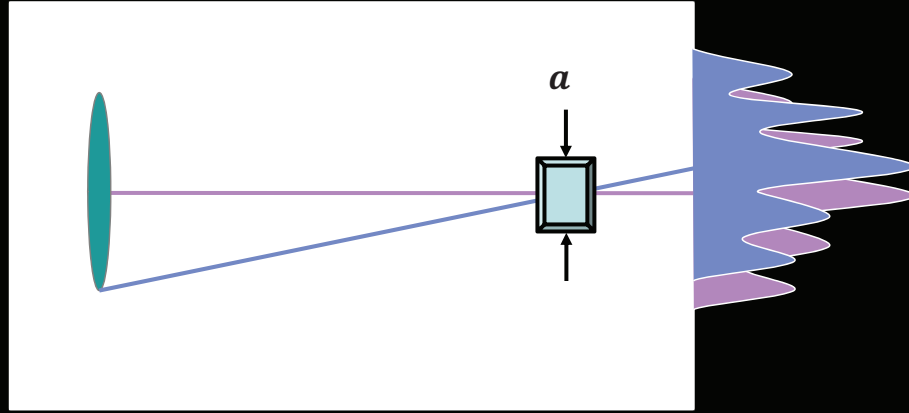
To measure the speckles



Fringes (or speckles) are visible iff $a < L$

Why coherence?

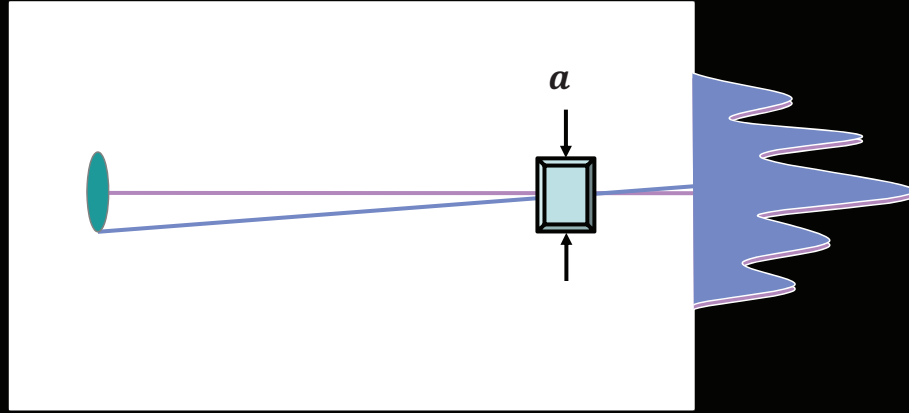
To measure the speckles



Fringes (or speckles) are visible iff $a < L$

Why coherence?

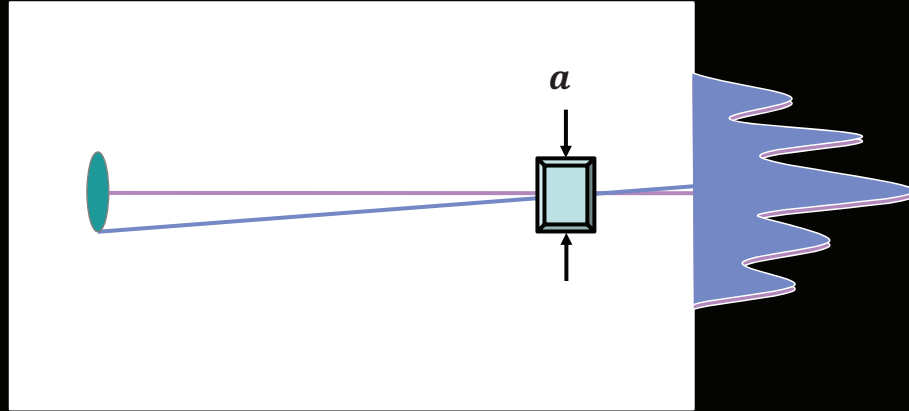
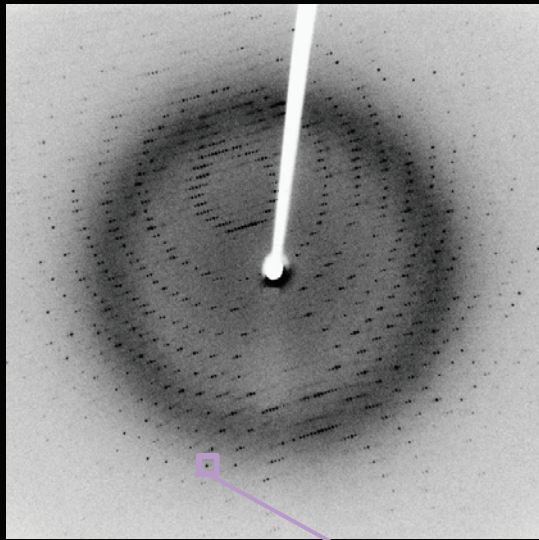
To measure the speckles



Fringes (or speckles) are visible iff $a < L$

Why coherence?

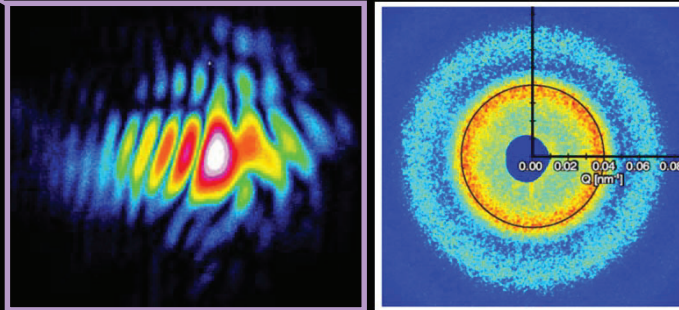
To measure the speckles



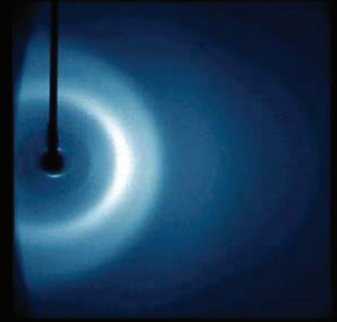
Fringes (or speckles) are visible iff $a < L$

New imaging
Techniques: CDI, Ptycho.

(Much) more demanding
of stability ($10 \mu\text{rad}$)



P. Wochner et al., *PNAS* 106, 11511 (2009).

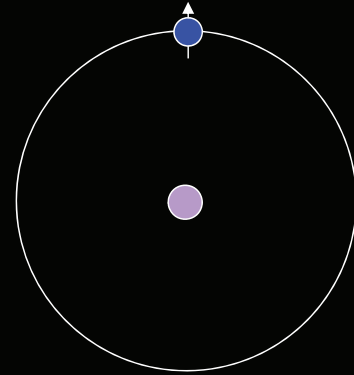


T. Head-Gordon et al., *Chem. Rev.* 102, 2651 (2002).

Absorption (and related techniques)

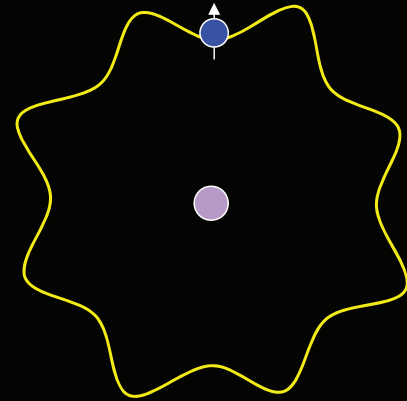
Absorption: energy levels in atoms

Where are the orbits of electron?



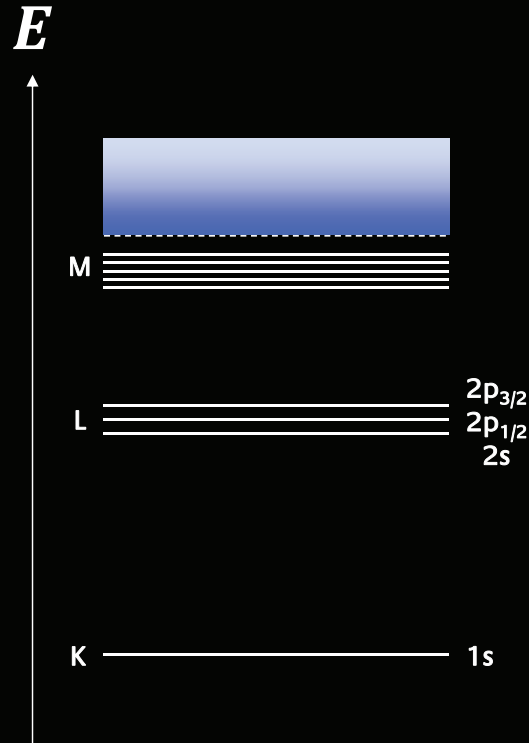
Absorption: energy levels in atoms

Where are the orbits of electron?
Because they are also waves...

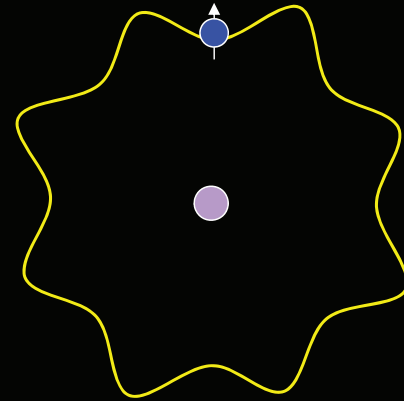


A few possibilities
called
Atomic orbital

Absorption: energy levels in atoms

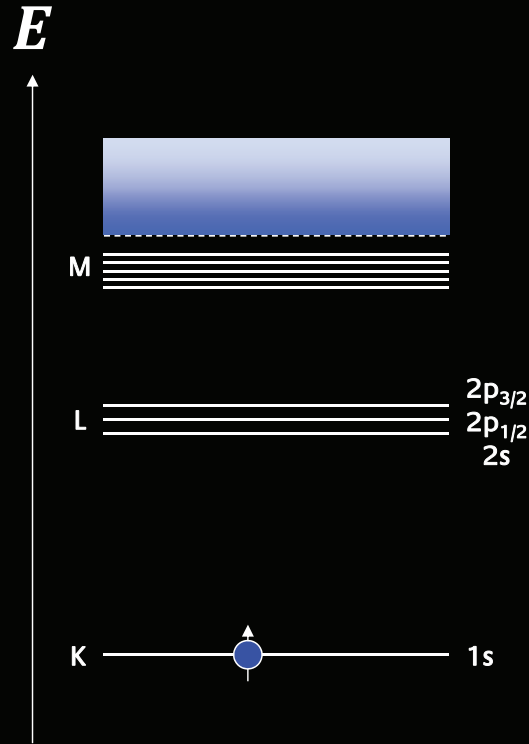


Where are the orbits of electron?
Because they are also waves...

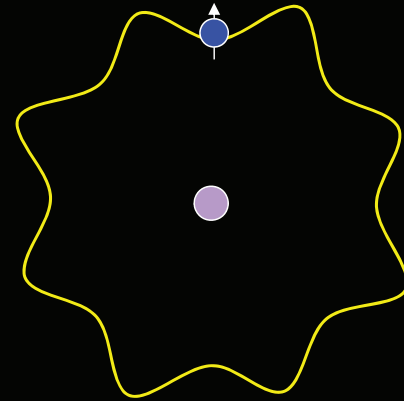


A few possibilities
called
Atomic orbital

Absorption: energy levels in atoms

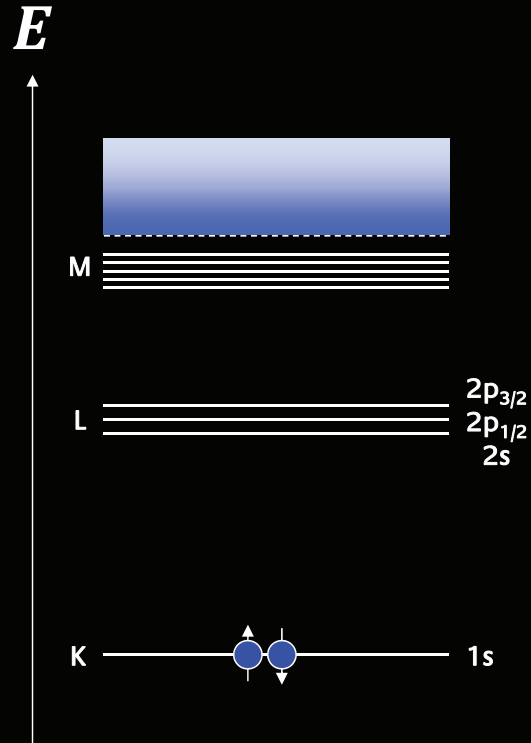


Where are the orbits of electron?
Because they are also waves...

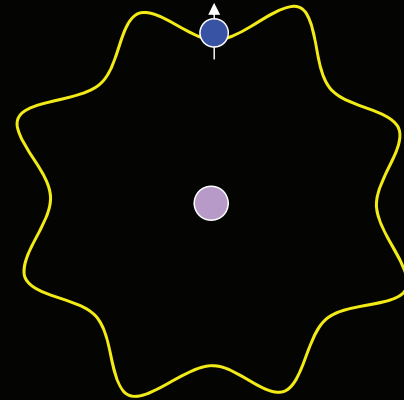


A few possibilities
called
Atomic orbital

Absorption: energy levels in atoms

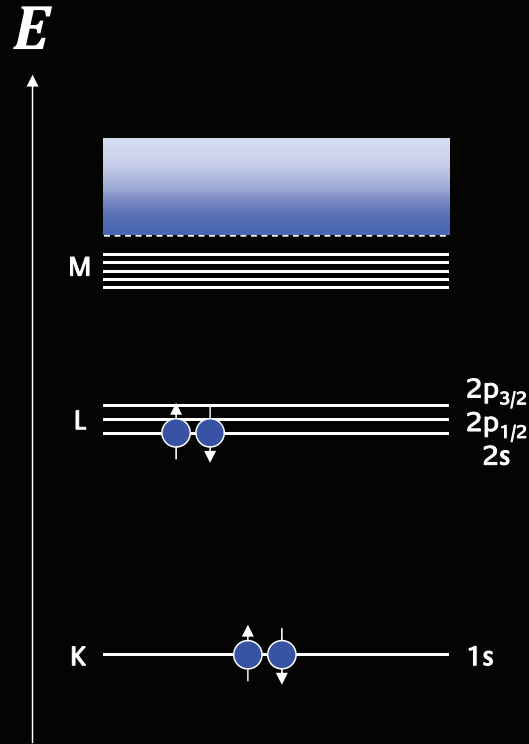


Where are the orbits of electron?
Because they are also waves...

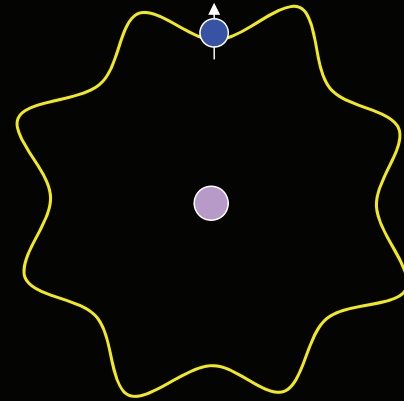


A few possibilities
called
Atomic orbital

Absorption: energy levels in atoms

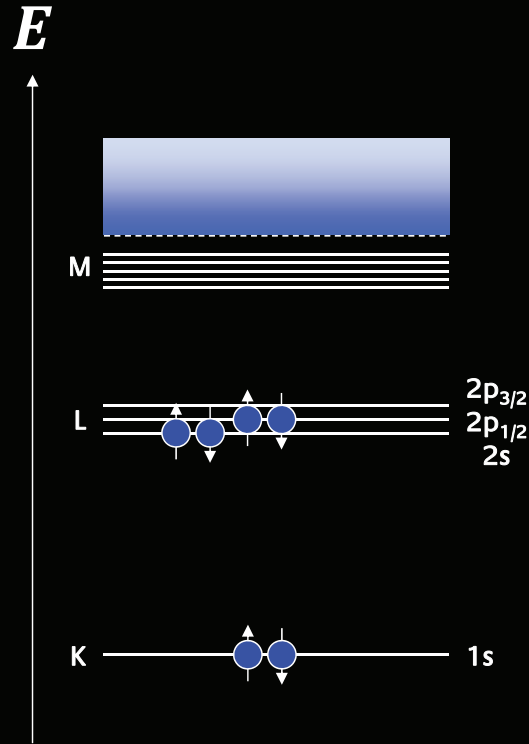


Where are the orbits of electron?
Because they are also waves...

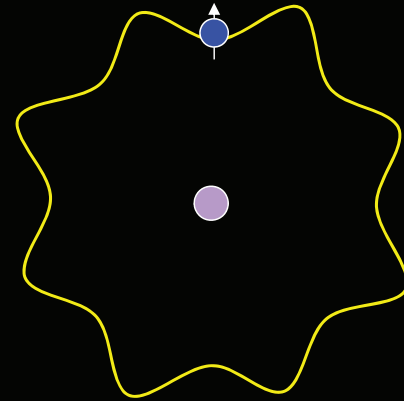


A few possibilities
called
Atomic orbital

Absorption: energy levels in atoms

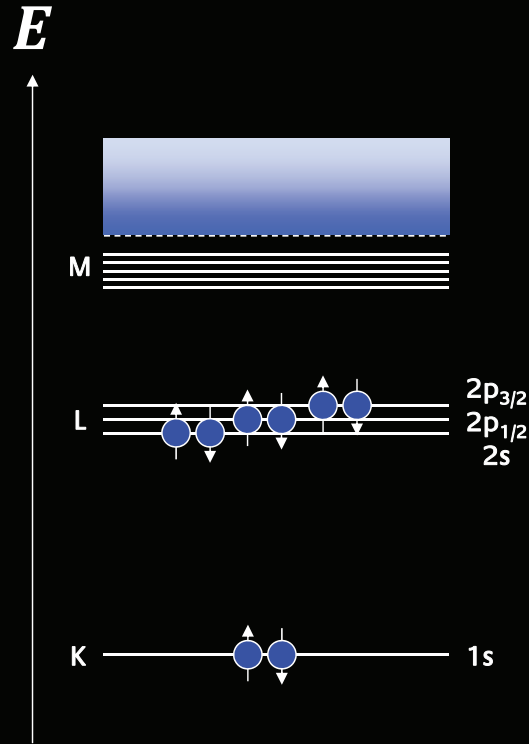


Where are the orbits of electron?
Because they are also waves...

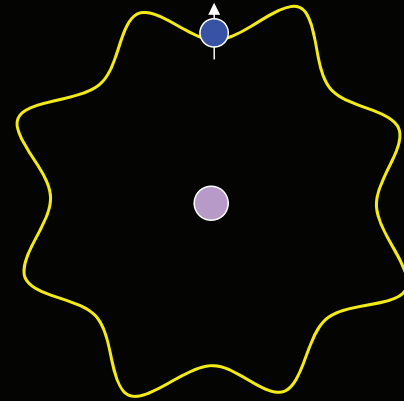


A few possibilities
called
Atomic orbital

Absorption: energy levels in atoms

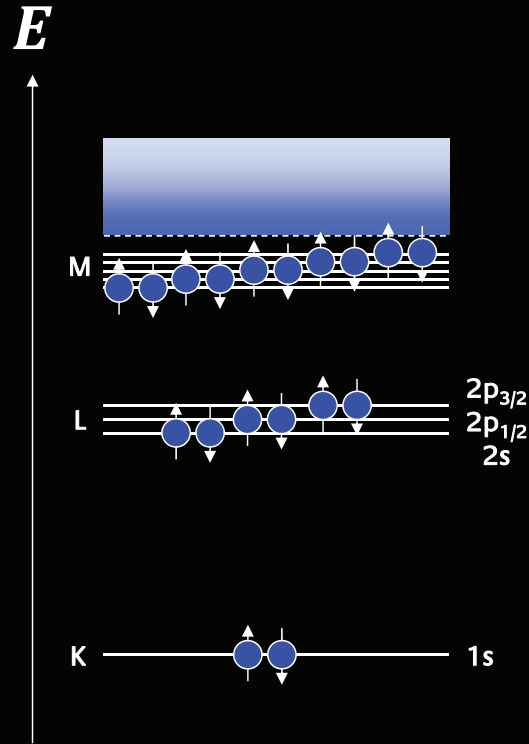


Where are the orbits of electron?
Because they are also waves...

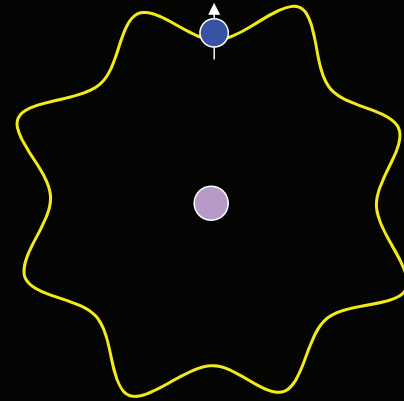


A few possibilities
called
Atomic orbital

Absorption: energy levels in atoms

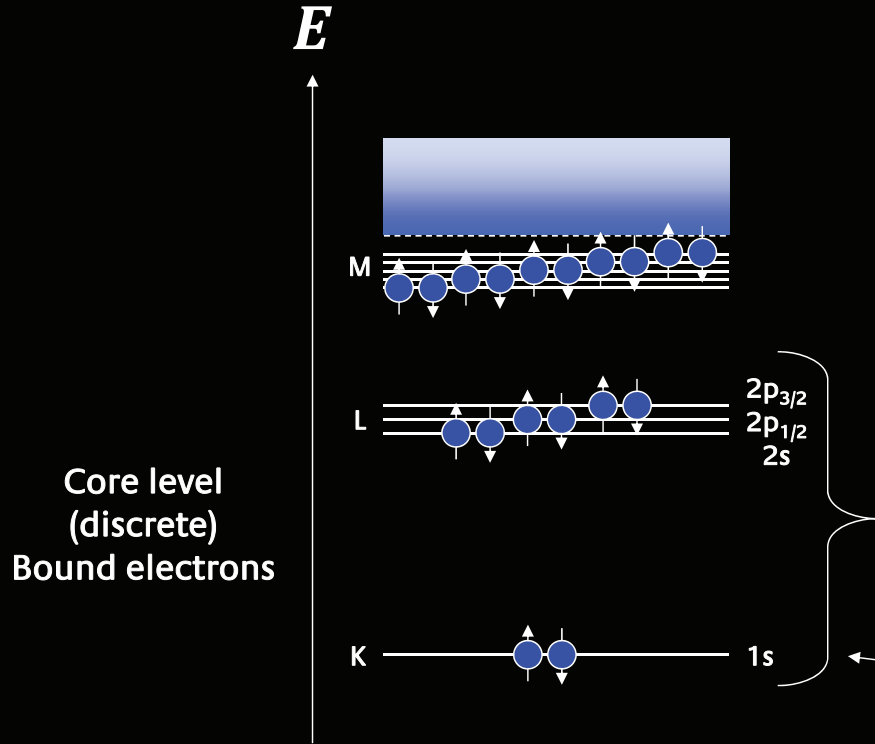


Where are the orbits of electron?
Because they are also waves...

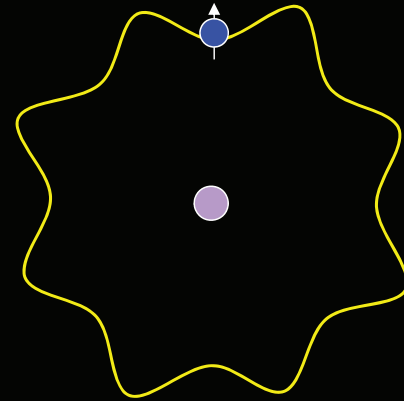


A few possibilities
called
Atomic orbital

Absorption: energy levels in atoms

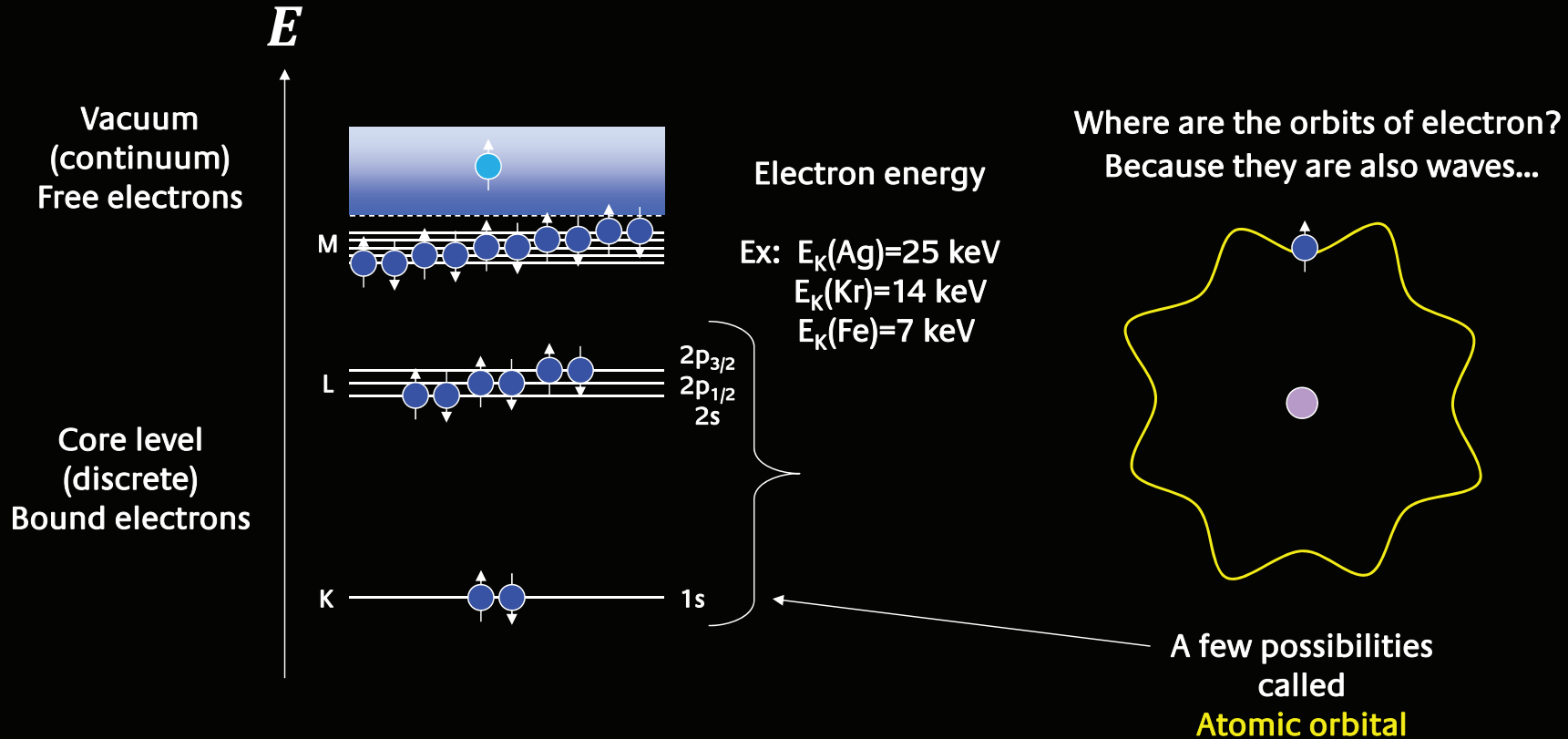


Where are the orbits of electron?
Because they are also waves...

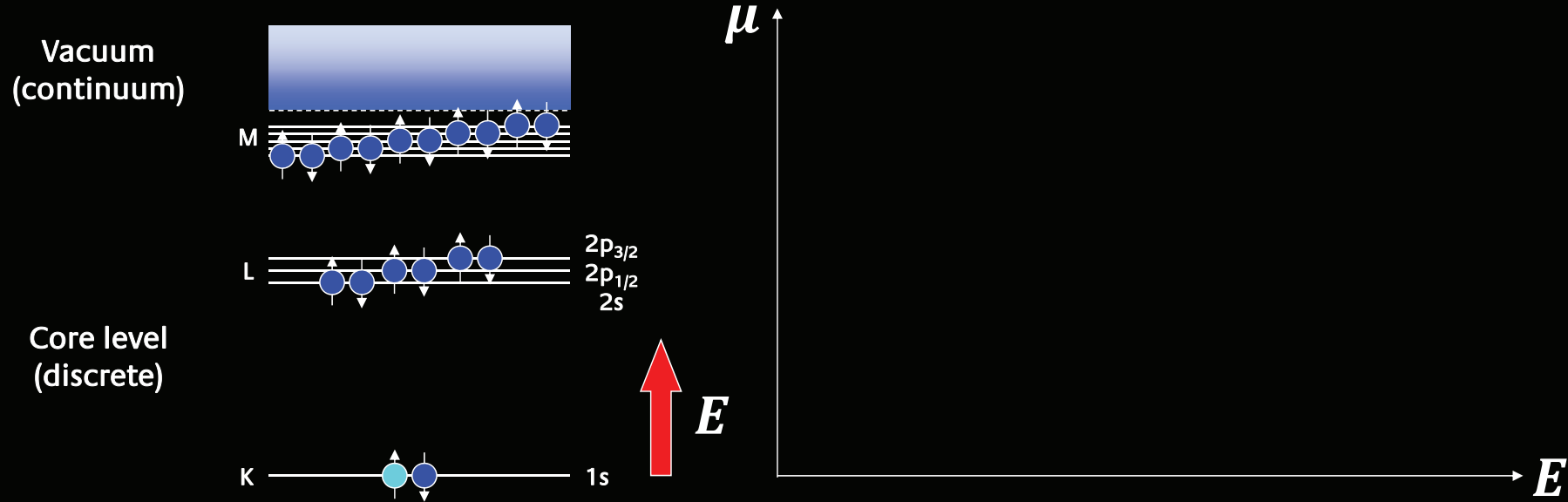


A few possibilities
called
Atomic orbital

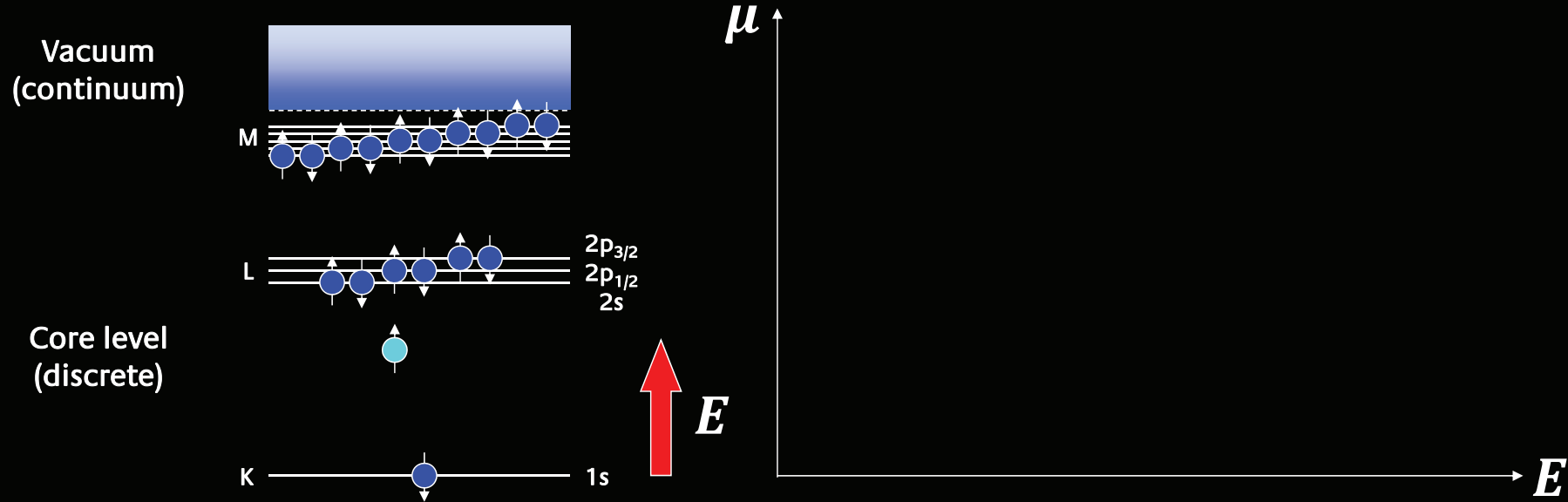
Absorption: energy levels in atoms



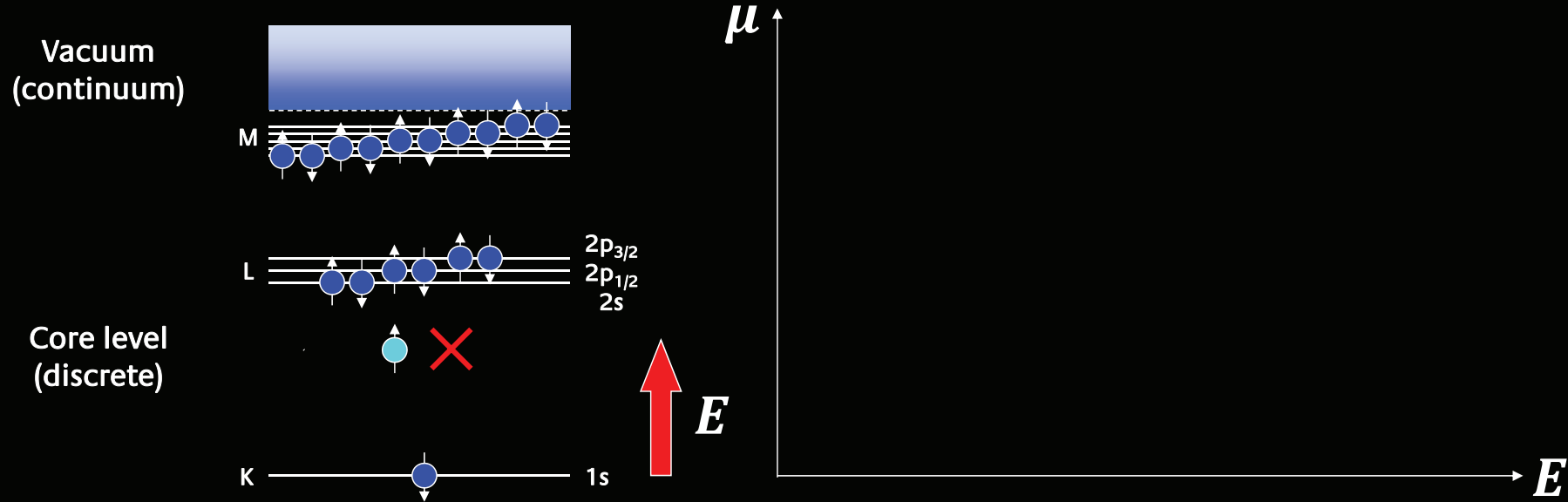
Absorption: increase of coefficient μ



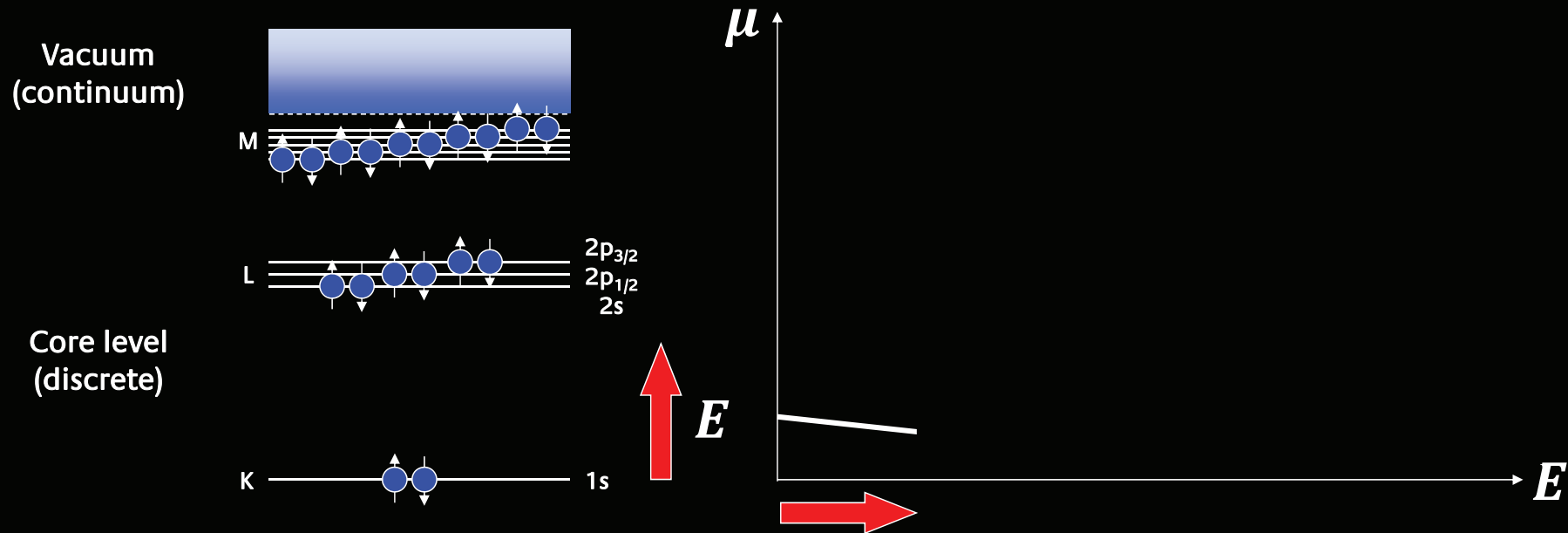
Absorption: increase of coefficient μ



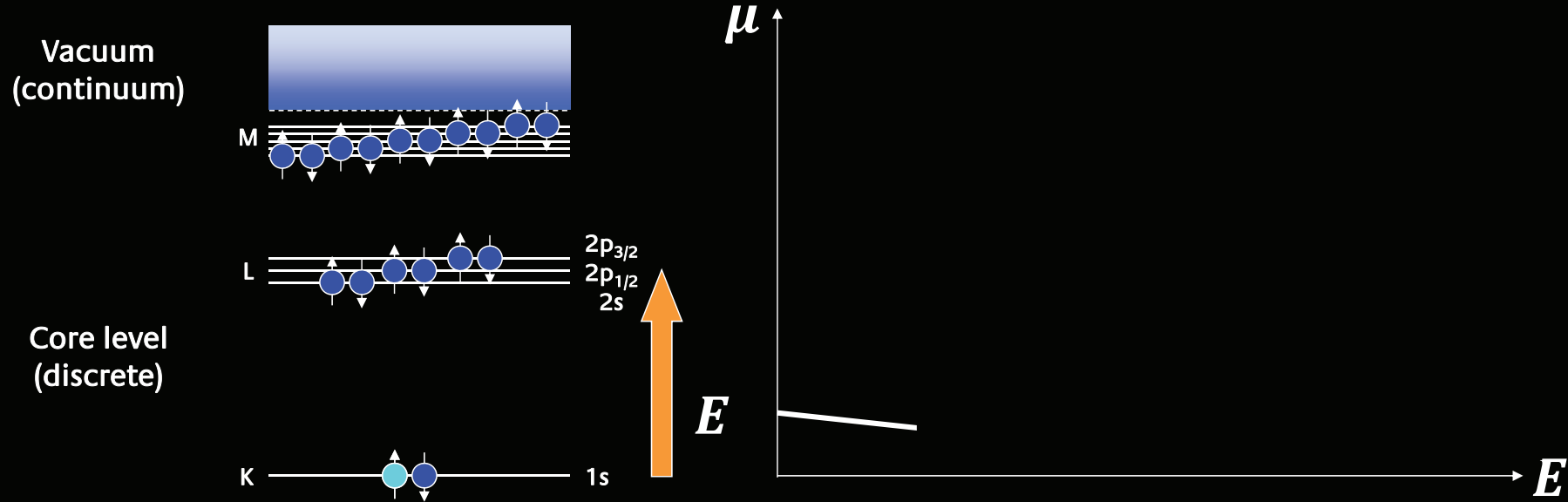
Absorption: increase of coefficient μ



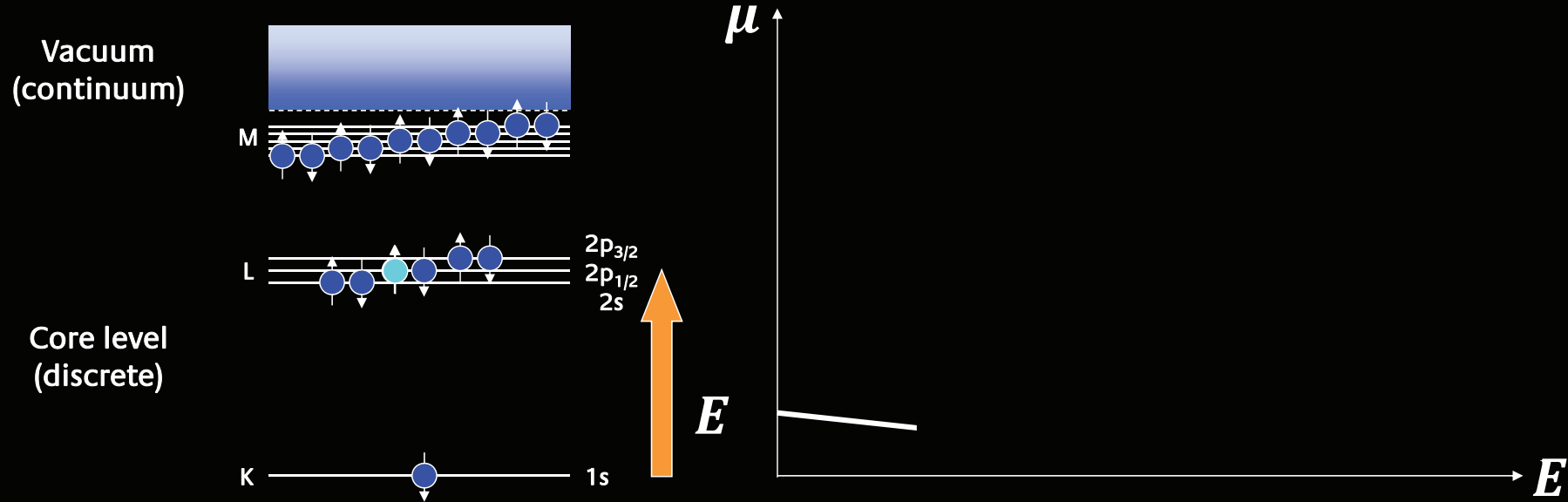
Absorption: increase of coefficient μ



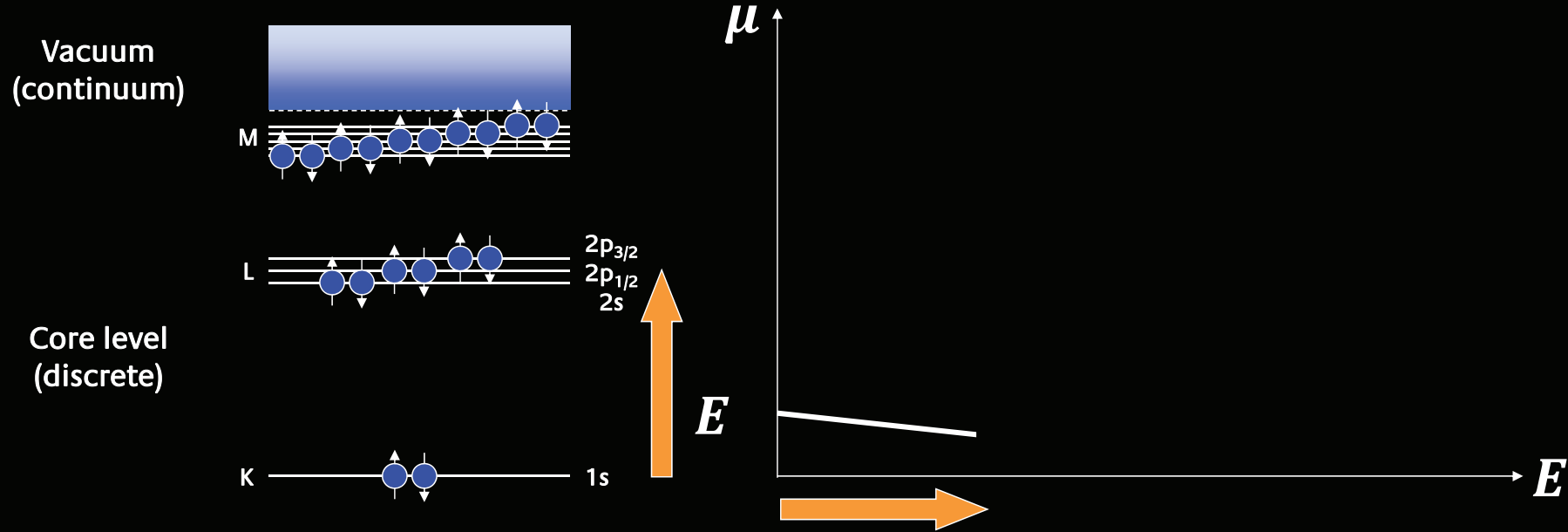
Absorption: increase of coefficient μ



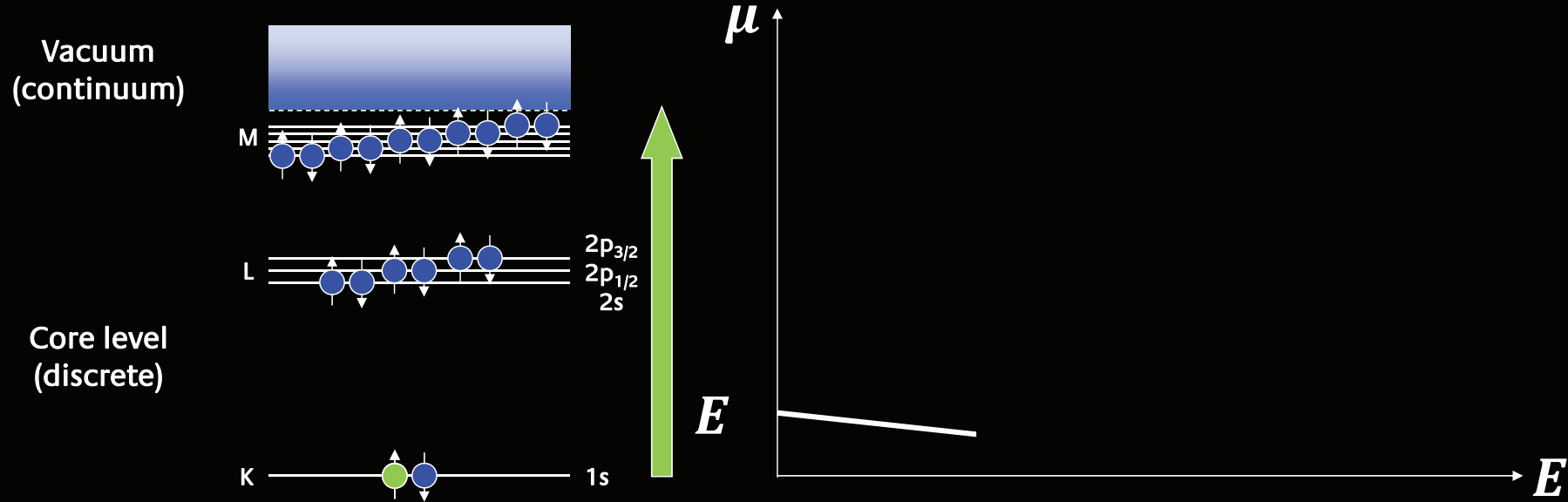
Absorption: increase of coefficient μ



Absorption: increase of coefficient μ



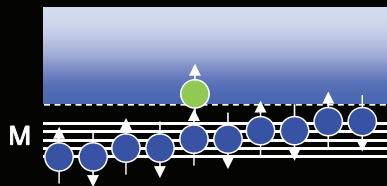
Absorption: increase of coefficient μ



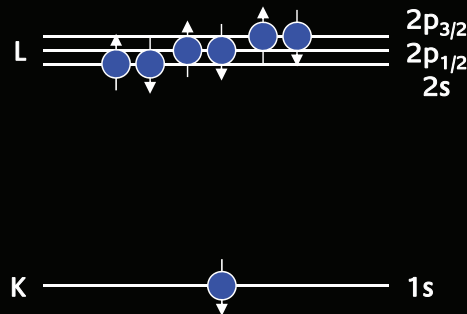
Absorption: increase of coefficient μ

Yes!

Vacuum
(continuum)



Core level
(discrete)



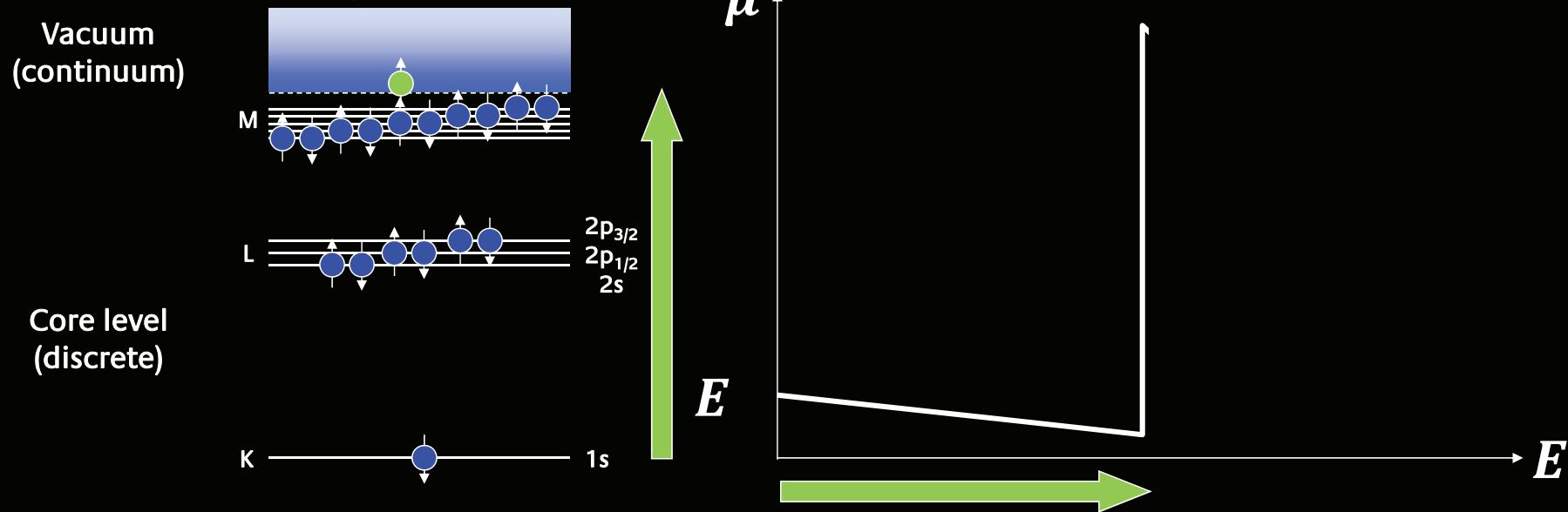
μ

E

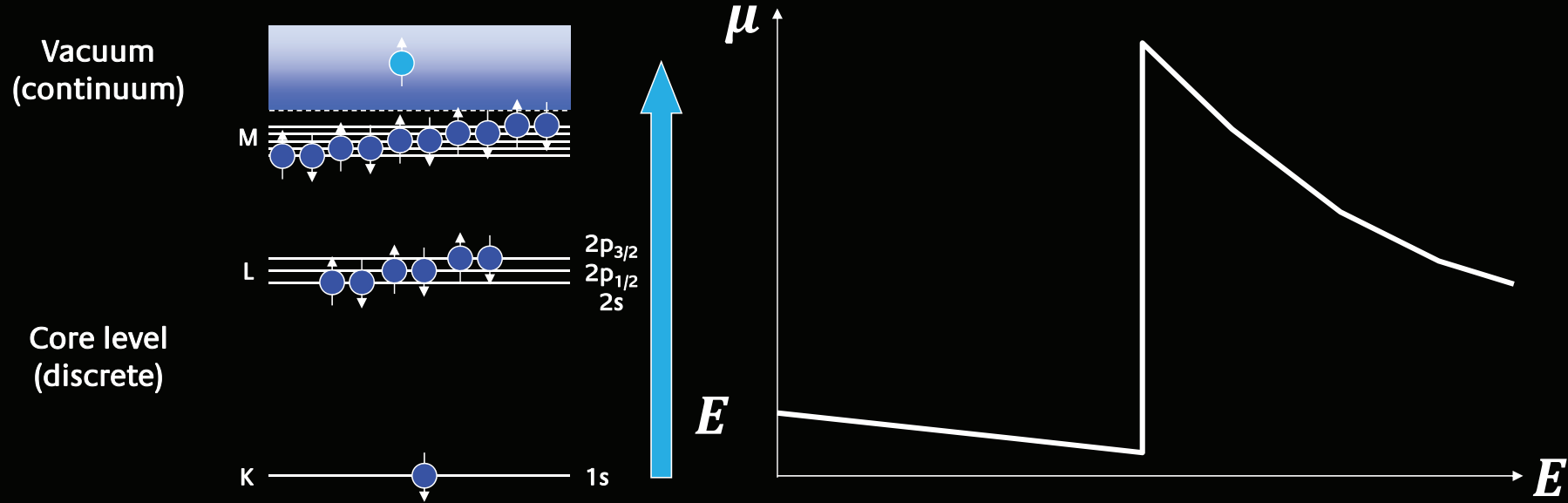
E

Absorption: increase of coefficient μ

Yes!



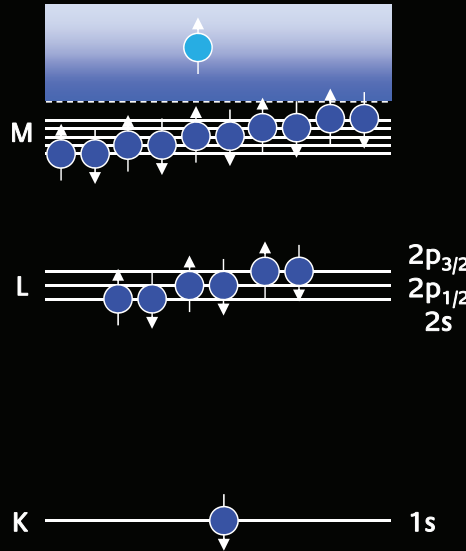
Absorption: increase of coefficient μ



Absorption: increase of coefficient μ

Photo-electron

Vacuum
(continuum)



Core level
(discrete)

μ

X-ray
Absorption
Spectroscopy

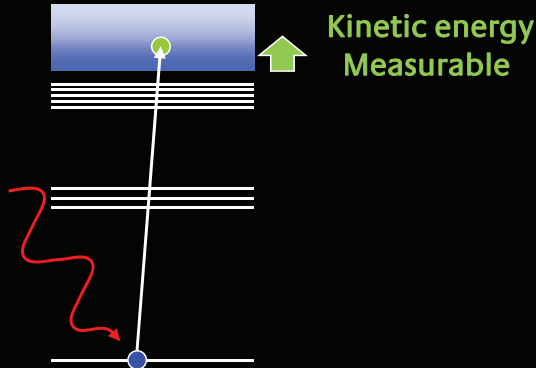
E

E

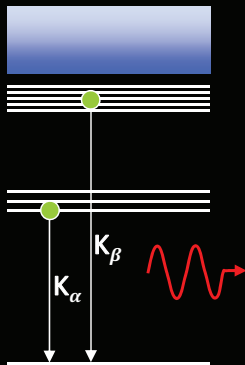
Element specific
K-edge, L-edge, etc.

Absorption: processes of relaxation

Photoelectric effect

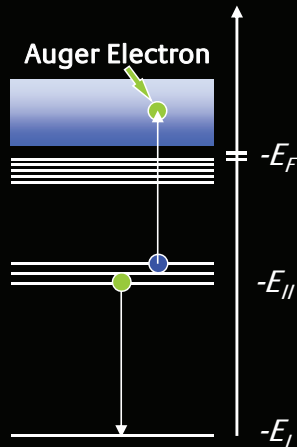


Fluorescence



Fluo and Auger
Are element
specific

Auger Electron



Atomic excitation:

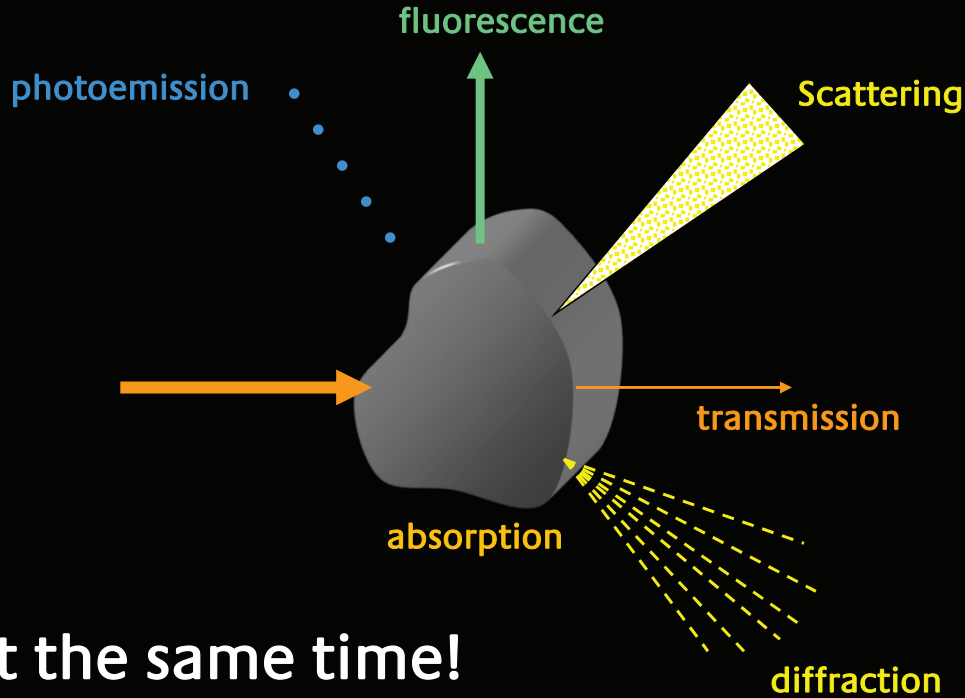
Absorption of photon (bulk)
Emission of Photo-electron
(surface ~10 nm, UHV)

Atomic relaxation:

Release of the excess of energy

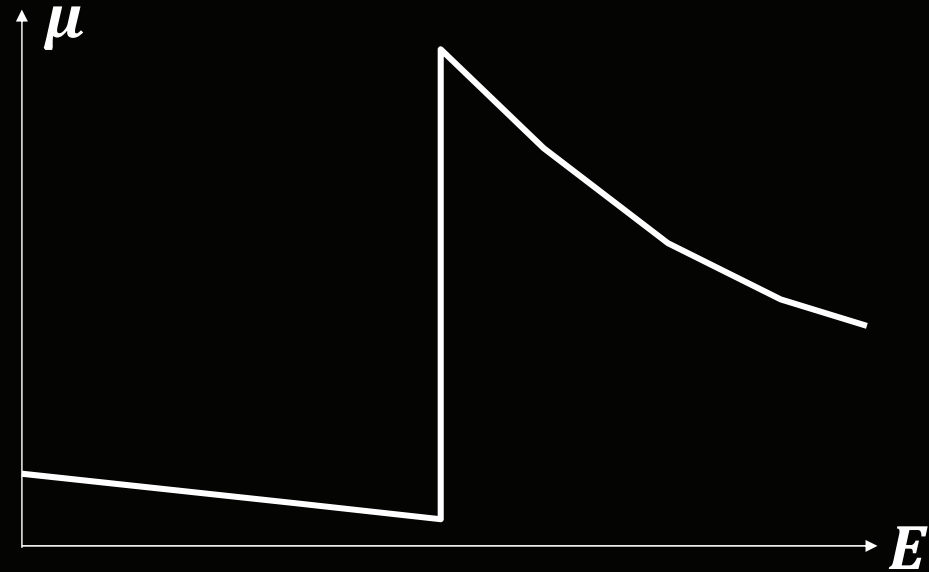
Emission of fluorescence photon
Emission of Auger electron

Interactions light/matter

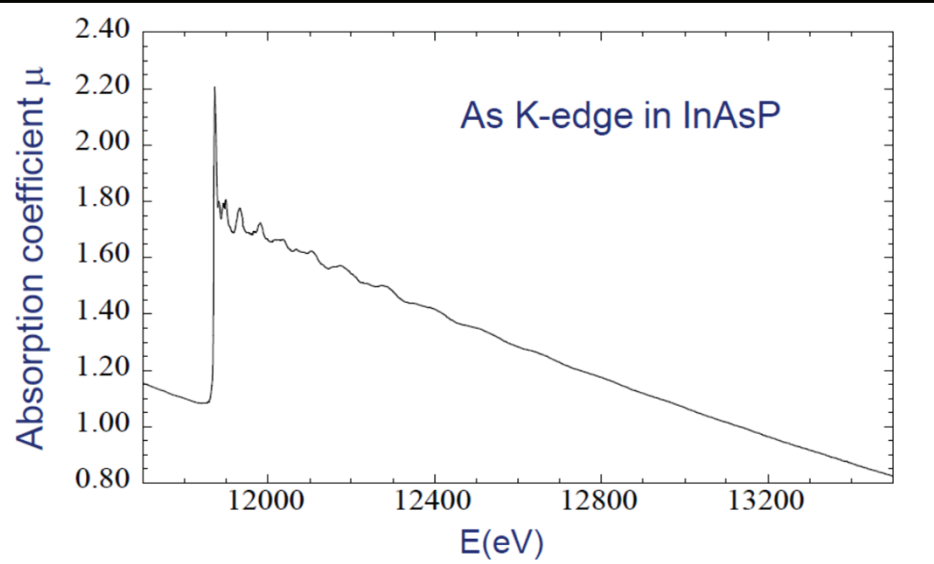


All (almost) at the same time!

Absorption: XAS, EXAFS, XANES

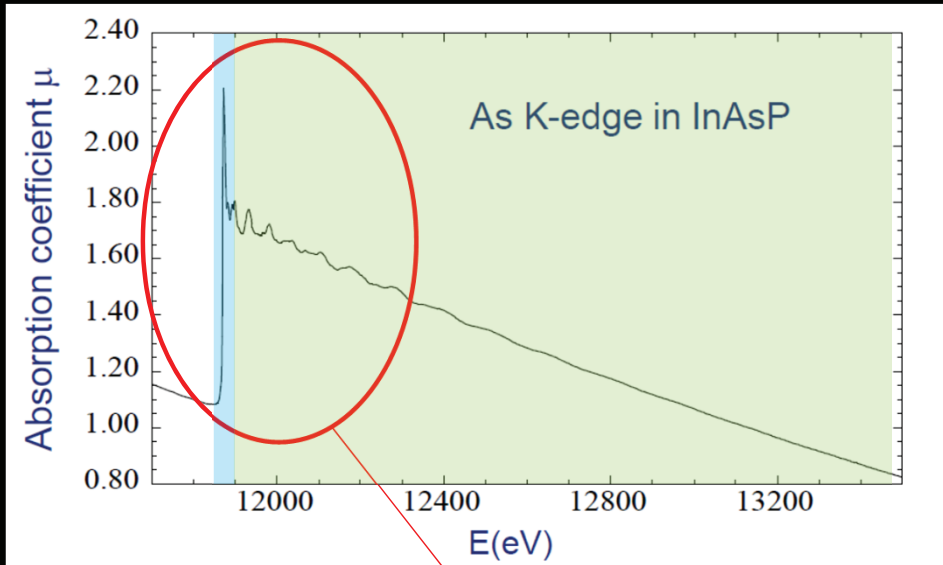


Absorption: XAS, EXAFS, XANES



Credit Sakura Pascarelli @ ESRF

Absorption: XAS, EXAFS, XANES



Credit Sakura Pascarelli @ ESRF

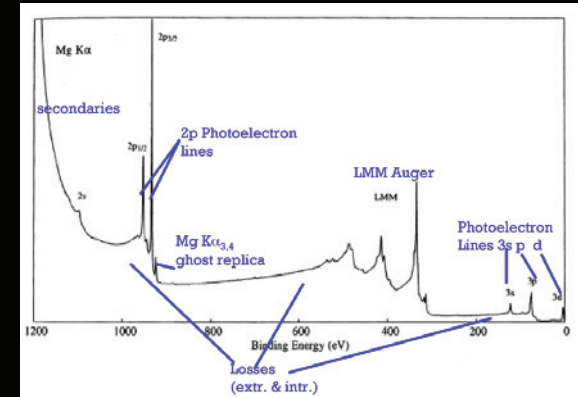
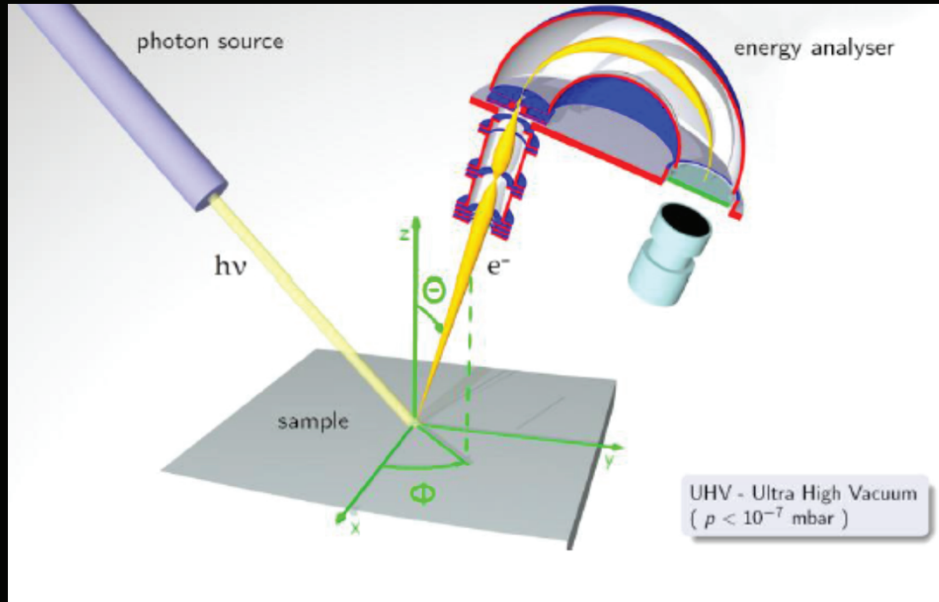
XAS measurements:

Absorption directly measured
Measurements of fluorescence
as a function of energy!

Constraints on monochromator
Reproducibility, precision, fixed exit
Dispersive XAS, quick EXAFS

Oscillations give information on atomic (**EXAFS**) and electronic (**XANES**) structure
Measurement of oscillations **as precisely as possible**

Photoemission spectroscopies

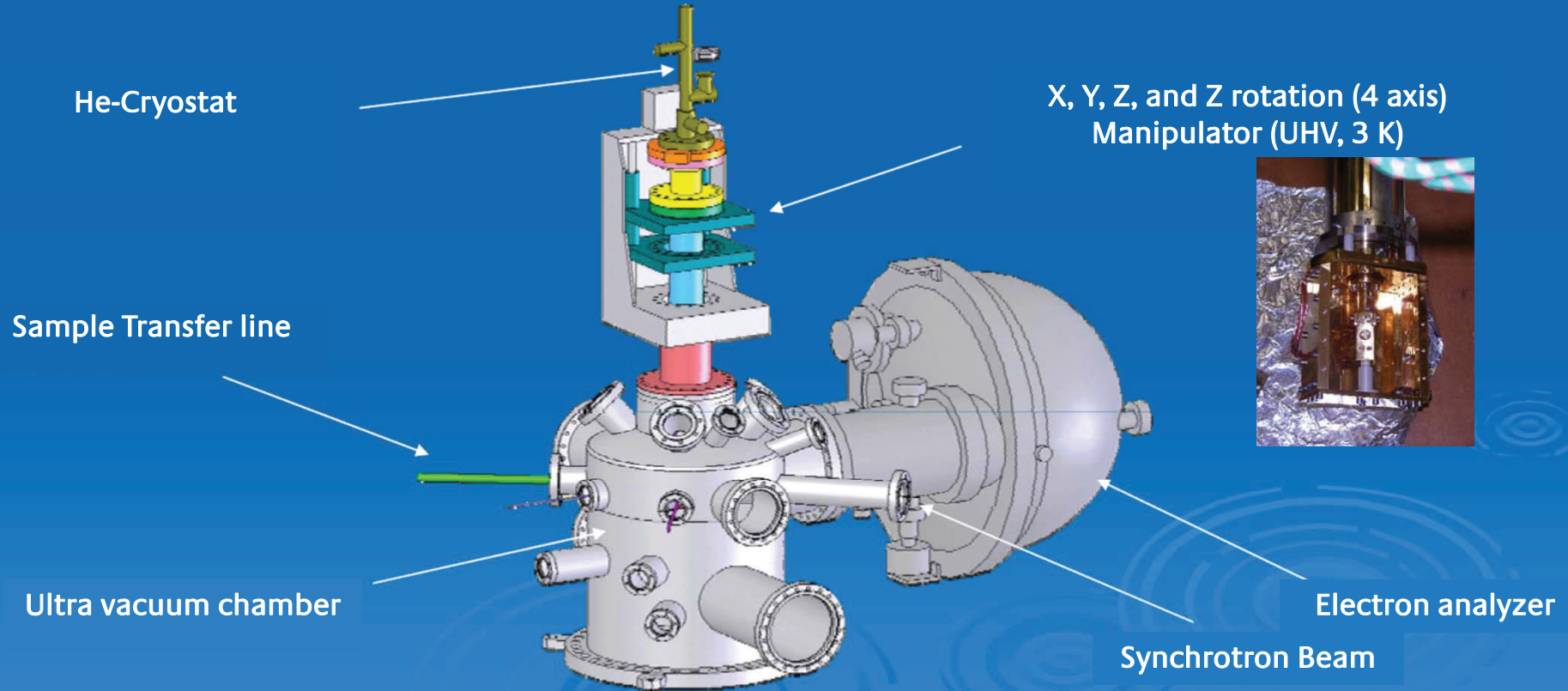


XPS: Measuring the photoemitted electrons as a function of electron energy

Angle Resolved Photo-Electron Spectroscopy (ARPES) as a function of angles (θ)
Temperature broadens the lines: low T

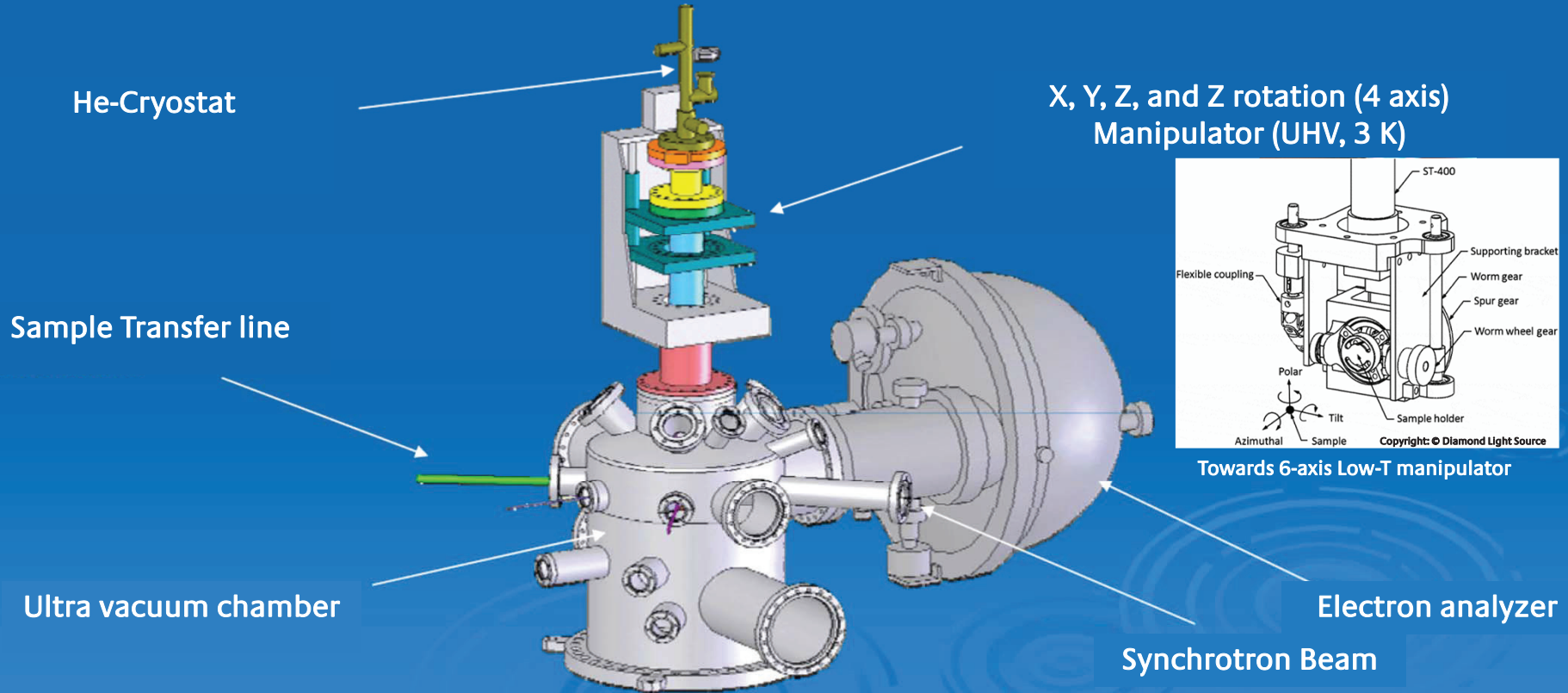
ARPES manipulators

Rotating the sample at very low T



ARPES manipulators

Rotating the sample at very low T

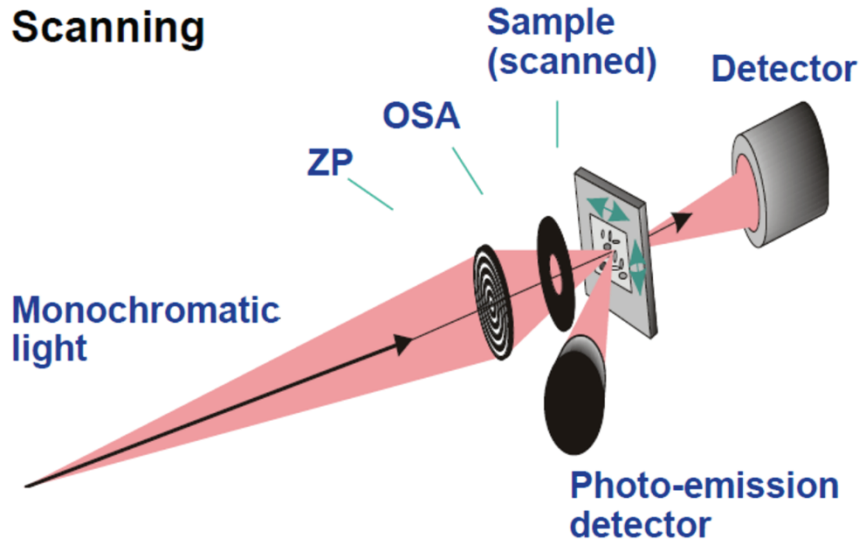


Imaging

Two main classes of 2D imaging techniques

Contrast and resolution

Scanning

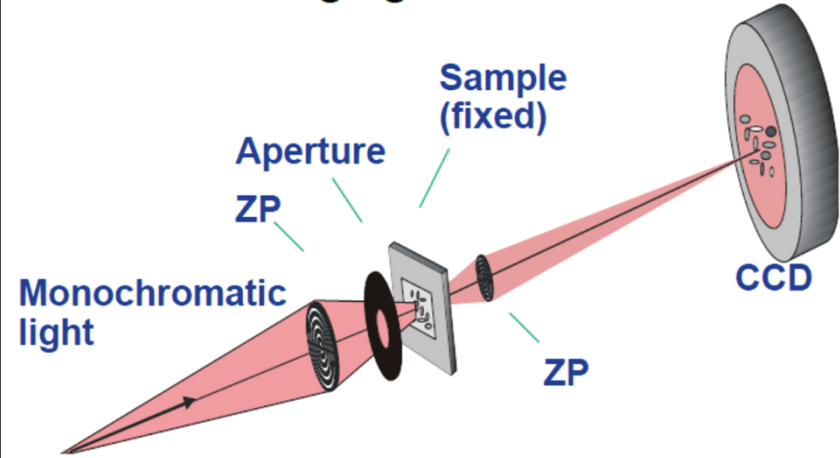


Photon efficiency

Different channels of measurements (XRD, XRF...)

Low speed; sensitive to drift

Full-field imaging



High speed: whole image at once
(Time-resolved expts)

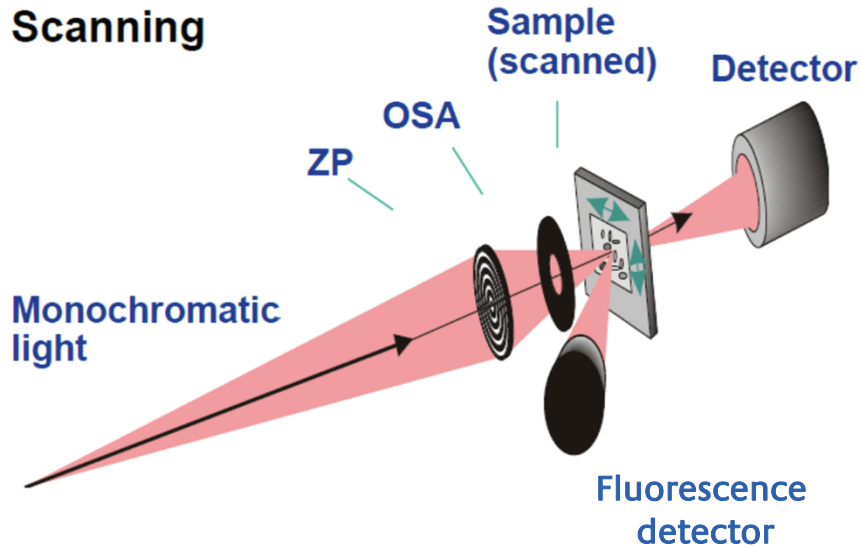
Low photon efficiency

Background from other channels

Two main classes of imaging techniques

Examples using energy absorption to contrast

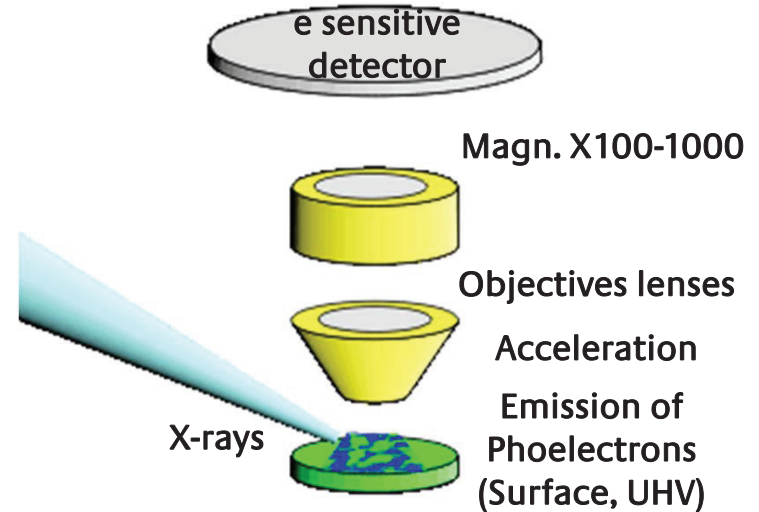
Scanning



Scanning transmission X-ray spectroscopy (STXM)

Resolution a few tens of nm.

All scanning techniques need
very precise, fast and reproducible actuators



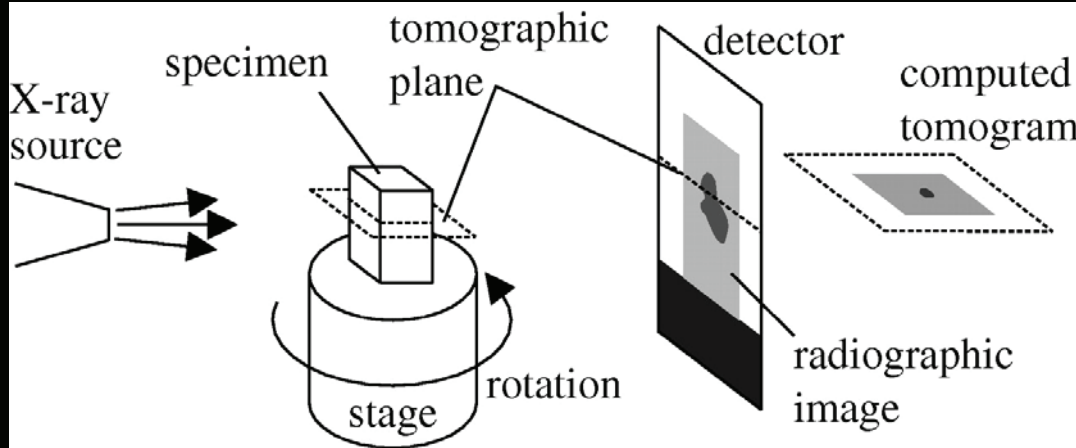
Credit A. Scholl ALS

X-ray Photoemission Electron Microscopy (XPEEM)

Same principle as electron microscope.

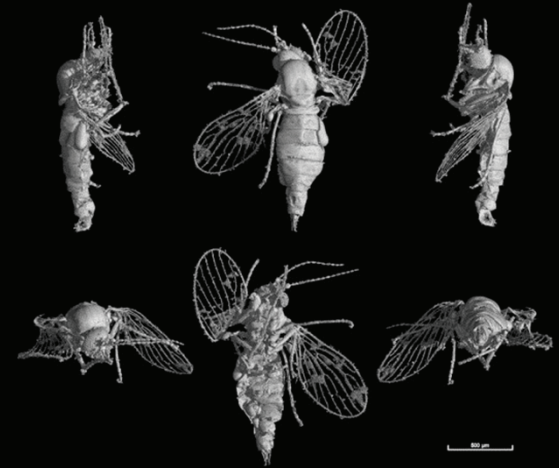
3D imaging: tomography

Based on absorption



Measurement of absorption image, as a function of angle
Computation of the 3D object

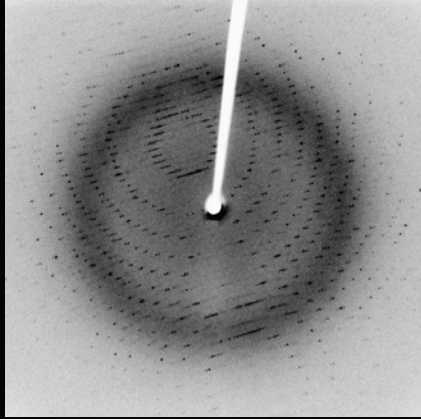
Need of **high precision**,
heavy load rotation stage (>50 kg, 100 nm, 60 rpm)



CREDIT P. Tafforeau ESRF

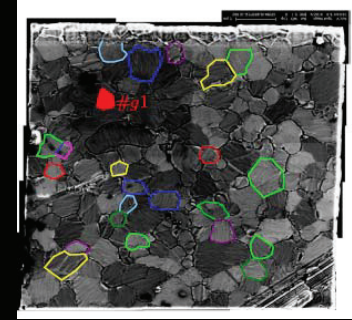
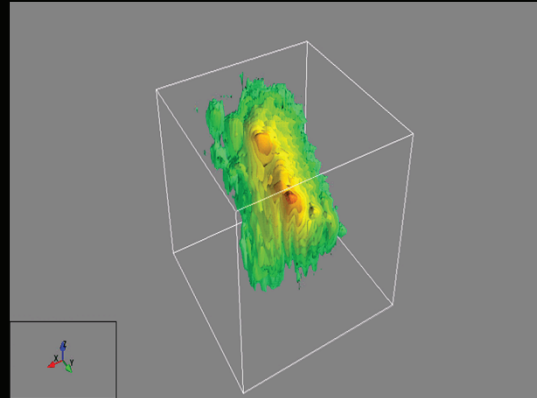
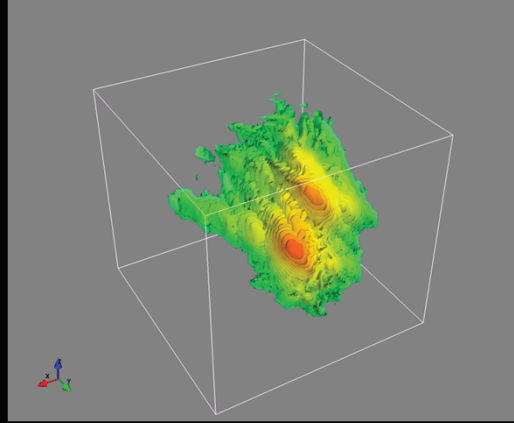
New 3D technique using coherence

Coherent Diffraction Imaging CDI



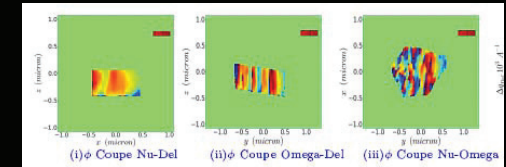
Principle:

Measurement of a **Bragg spot**
of a micrometric crystal
as a function of Bragg angle
In coherent conditions



CREDIT: N. Vaxelaire, S. Labat IM2NP

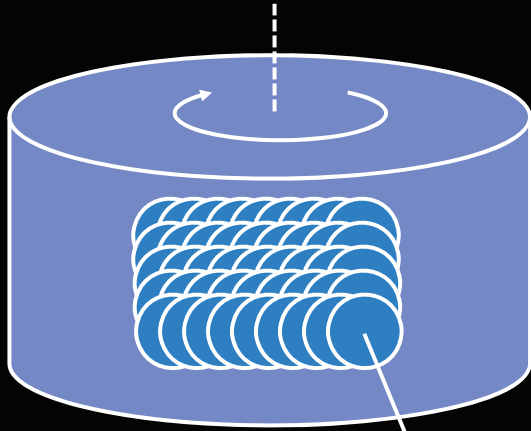
Reconstruction of strain
In a sub- μ gold grain



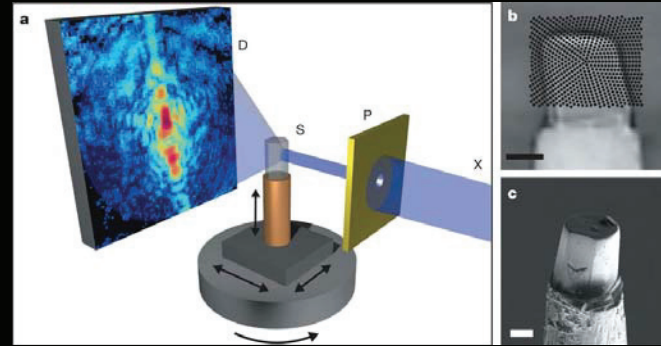
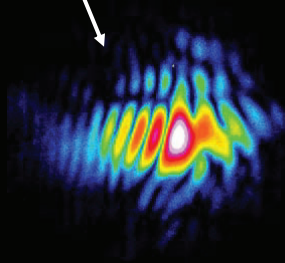
Very demanding expts:
Stability (coherence)
Precision of rotation stages
Sample environment (low T)

New 3D technique using coherence

Ptychography: raster scan with overlap, and rotation



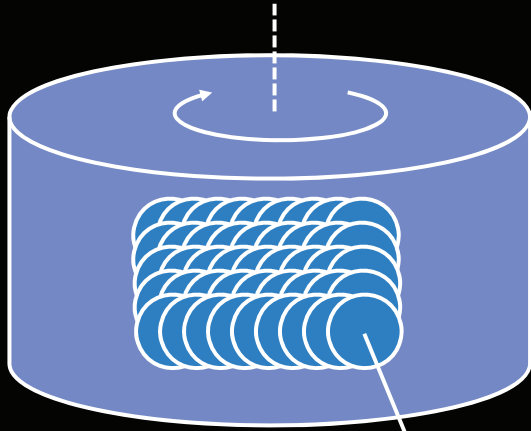
Measurement of diffraction
Pattern at each point



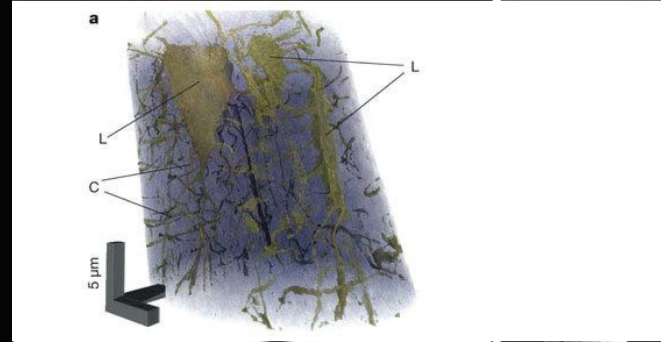
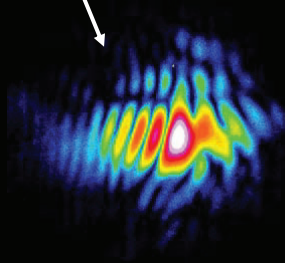
First demonstrated at SLS (40h)
Dierolf et al. *Nature* 467 (2010)

New 3D technique using coherence

Ptychography: raster scan with overlap, and rotation



Measurement of diffraction
Pattern at each point



First demonstrated at SLS (40h)
Dierolf et al. *Nature* 467 (2010)

Imaging techniques are called « lensless »

Long term stability (few hours)
Small intensity
Angular precision (coherence)

Conclusions

Increase of brilliance raises important stability issues:

Stability of monochromators (cooling systems)

To be scaled to the source size

Reproducibility (energy scans)

Temperature stability (thermal drifts)

Precision of angles (diffraction)

Precisions of all actuators for scanning (in the nm range)

Constraints on sample environments (low T, high P)

Thank you!