

Photo emission electron microscopy

Outline

Introduction

Electron optics:
Resolution
Transmission

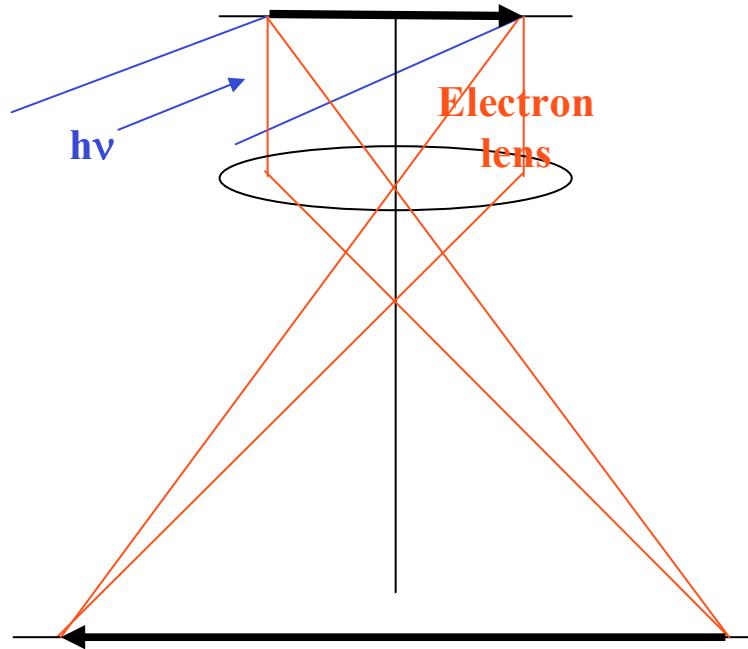
Instruments:
PEEM
PEEM + LEEM
SPELEEM

Methodic

Applications:
Magnetic imaging

Photo Emission Electron Microscopy (PEEM)

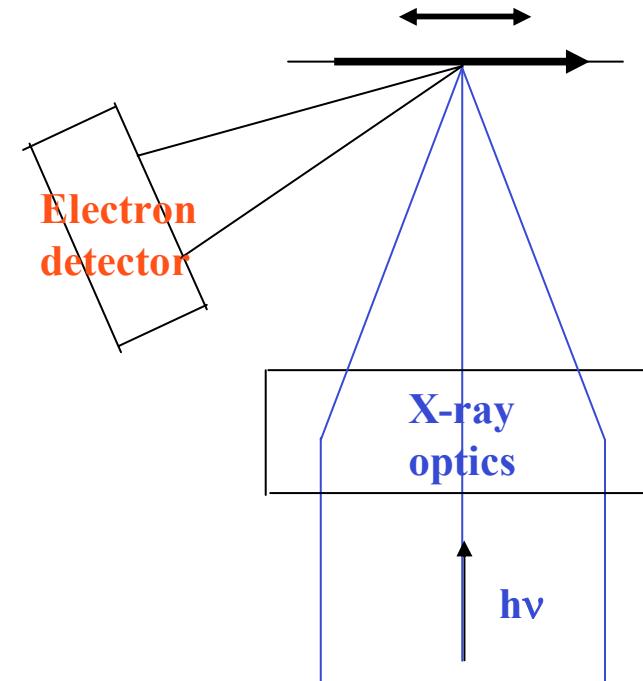
2 types



broad illumination

Full field
sample fixed

Bauer, Locatelli



focused illumination

PEEM

Scanning
sample scanned

Kiskinova

3 imaging modes

1 XPEEM

Photo electrons PE

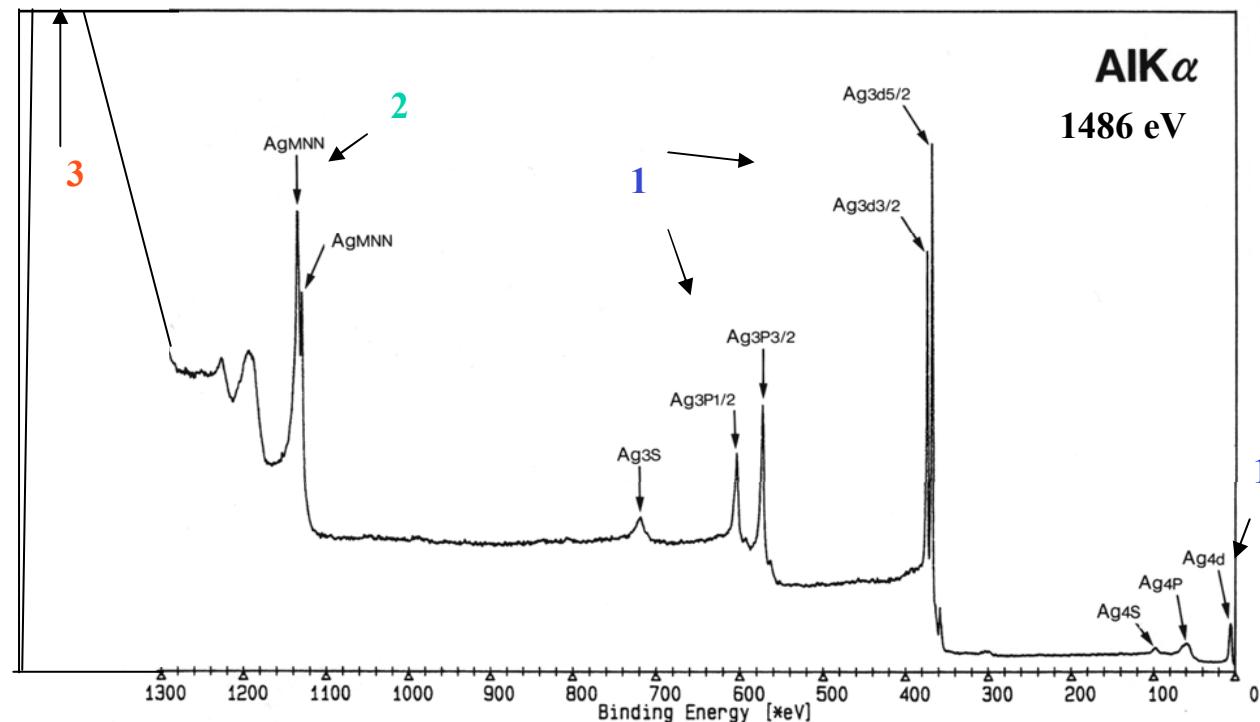
} with energy filter

2 XAEEM

Auger electrons AE

3 XSEEM

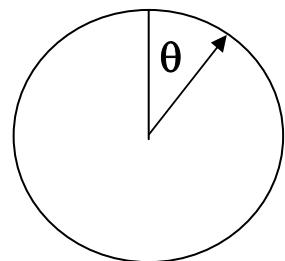
Secondary electrons SE



Angular distribution

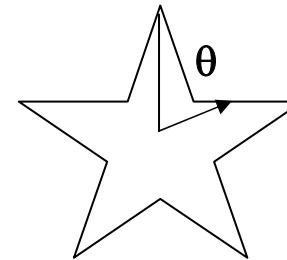
Internal

Amorphous, polycrystalline, SE



$$I_i(\theta) = \text{const.}$$

single crystalline, PE, AE

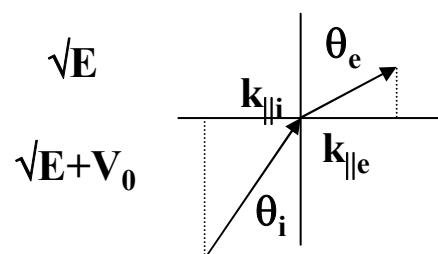


$$I_i(\theta) \text{ due to diffraction}$$

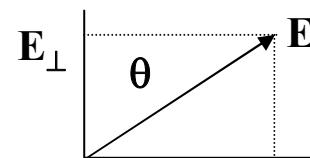
Internal (i) → External (e)

$n \sim$

Refraction



k_{\parallel} conservation $k_{\parallel e} = k_{\parallel i}$

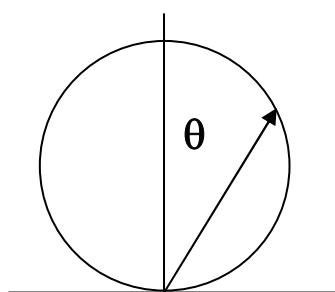


$$E_{\perp} = E \cos \theta$$

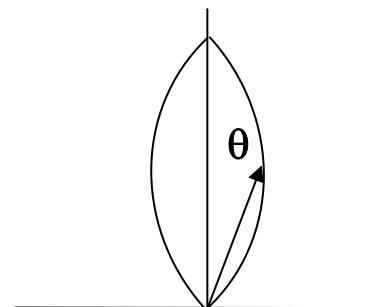
For escape

necessary:
 $E_{\perp} > \Phi$ (work function)
 I (ionization energy)
 U (HOMO)

$$I_e(\theta) = \cos \theta$$



External



Electron optics

The cathode lens

In emission microscopy $\theta \equiv \alpha_0$ is large

Electron lenses can accept only small $\theta \equiv \alpha_0$ because of
large chromatic and spherical aberrations

Solution of problem: accelerate electrons to high energy before lens



Immersion objective lens = cathode lens

$$n \sin \theta = \text{const}$$

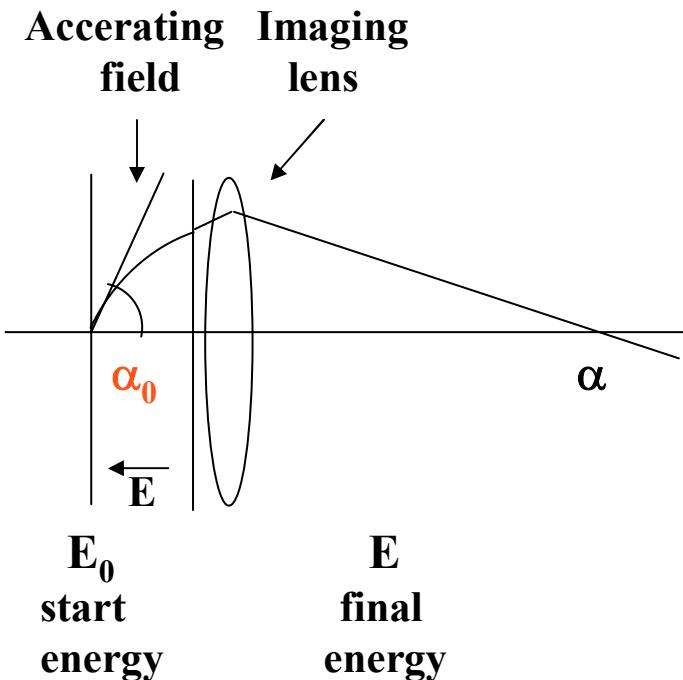
$$n \sim v \sim \sqrt{E}$$

$$\theta \rightarrow \alpha$$

$$\sin \alpha / \sin \alpha_0 = \sqrt{E_0/E}$$

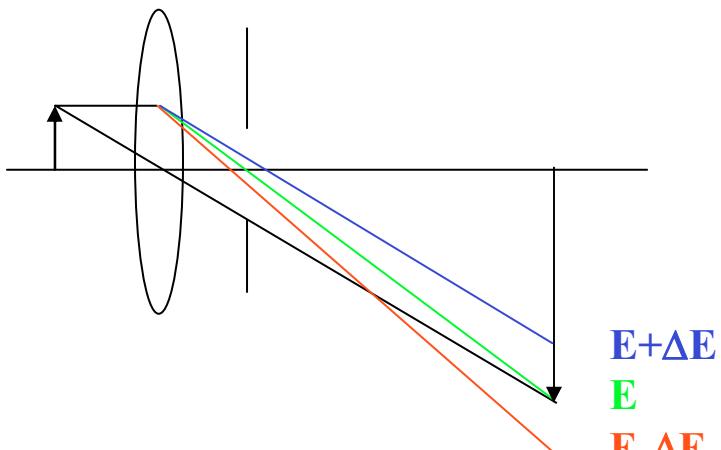
Example for $E = 20000$ eV:

E_0	2 eV	200 eV
α for $\alpha_0 = 45^\circ$	0.4°	4.5°



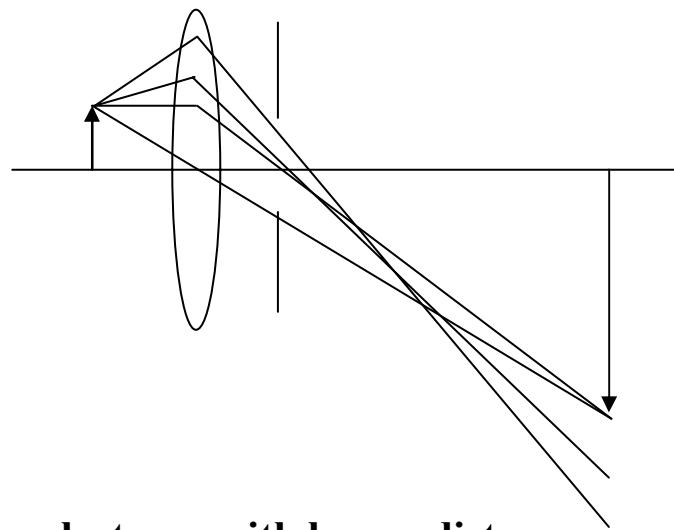
Aberrations

chromatic



slower (faster) electrons
are more (less) deflected

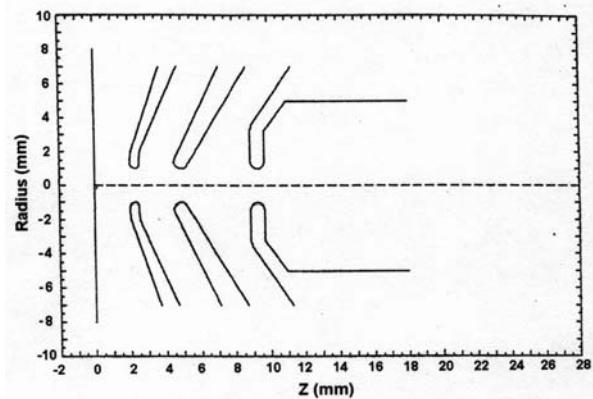
spherical



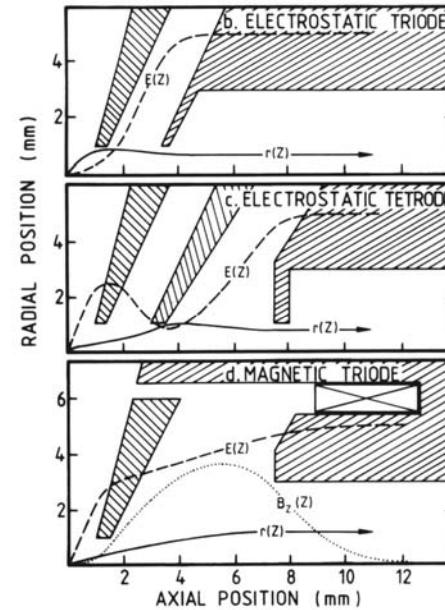
electrons with larger distance
from axis are more deflected
(stronger field!)

Cathode lens types

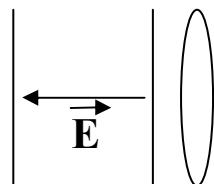
Electrostatic tetrode



Lens comparison



Estimation of aberrations:
Separate lens into acceleration and imaging regions

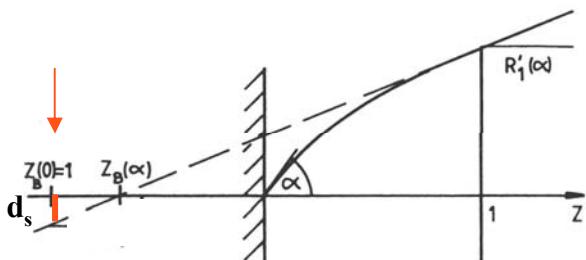


At low energies the aberrations of the accelerating region dominate

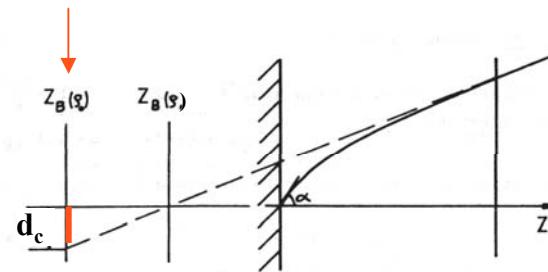
Aberrations of homogeneous acceleration field

$$\rho_0 = E_0/E \quad \varepsilon = \Delta E_0/E \quad \rho = \rho_0 + \varepsilon$$

Spherical aberration d_s



Chromatic aberration d_c



Analytical solution

Approximation: ρ_0 and $\varepsilon \ll 1/\cos \alpha^2 > 1$

Example: $E_0 = 100 \text{ eV}$, $\Delta E_0 = 1 \text{ eV}$, $E = 20000 \text{ eV}$

$$\varepsilon = \rho_0 / 100, \quad \rho_0 = 1/200$$

$$d_s \approx 2 \rho \sin \alpha (1 - \cos \alpha)$$

$\approx \rho \alpha^3$ for small α

$$d_c \approx 2 \rho \sin \alpha (\sqrt{\rho_0 / \rho} - 1)$$

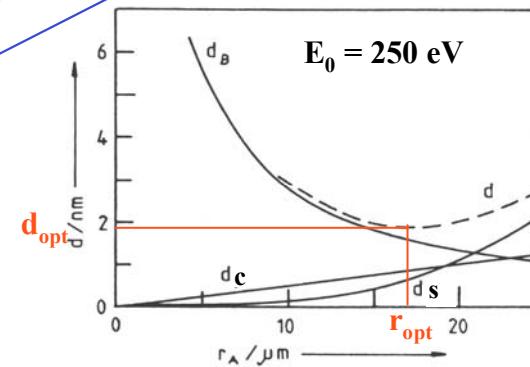
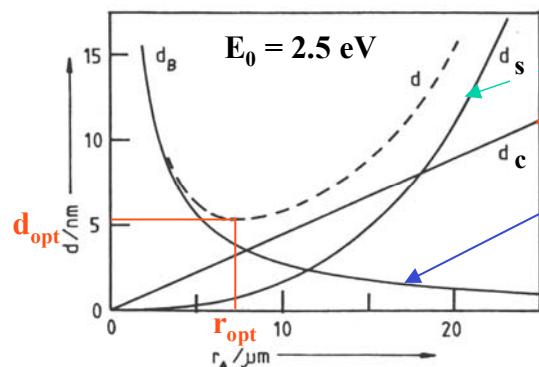
$\approx \varepsilon \sin \alpha$ for $\varepsilon \ll \rho_0 \approx \rho$
 $\approx \varepsilon \alpha$ for small α

α -dependent aberrations require α -limitation by angle-limiting aperture (“contrast aperture”) with radius r_A



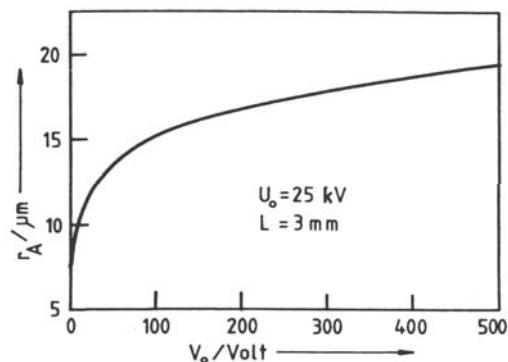
Diffraction by aperture: diffraction disc of confusion $d_B = 0.6 \lambda / r_A$

$$\text{Approximate resolution } d = \sqrt{d_s^2 + d_c^2 + d_B^2}$$

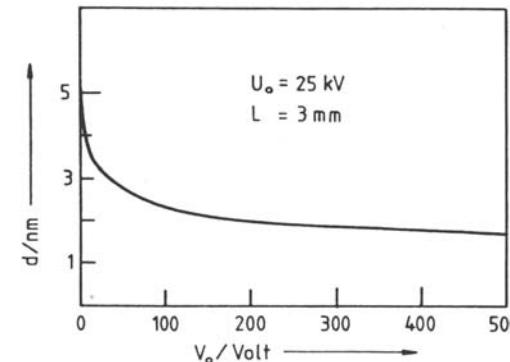


$$L = 3 \text{ mm} \quad E = 25000 \text{ eV} \quad \Delta E_0 = 0.25 \text{ eV}$$

Optimum aperture radius



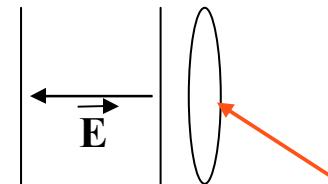
Optimum resolution



Note: small angle approximation $\sin \alpha \approx \alpha \sim r$

Complete lens

Combine acceleration and imaging regions

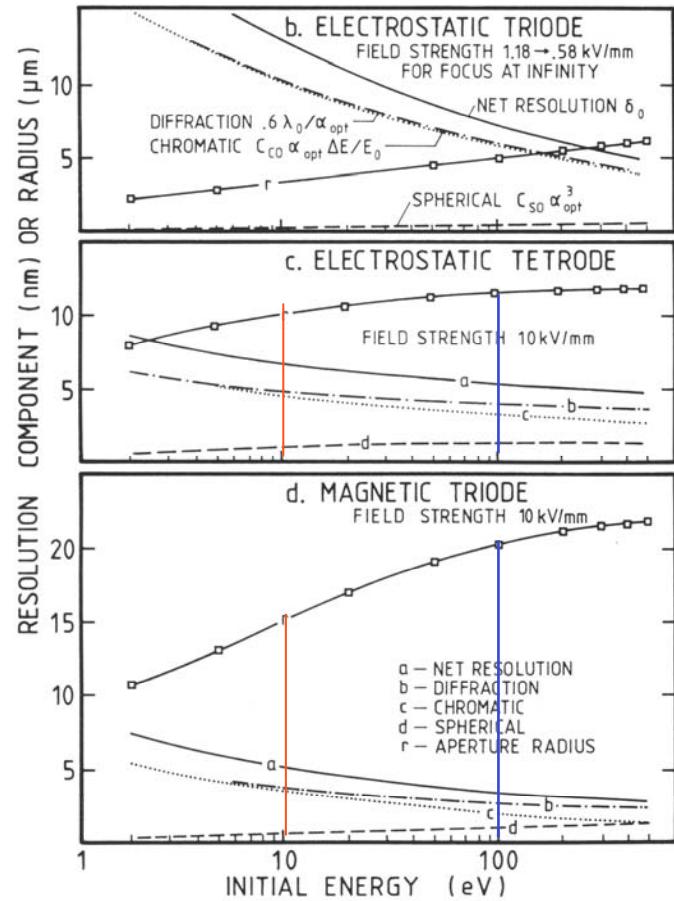


At low energies aberrations of accelerating region dominate
but

at high energies the spherical aberration of second part of lens becomes important

Resolution and optimum aperture of real lenses

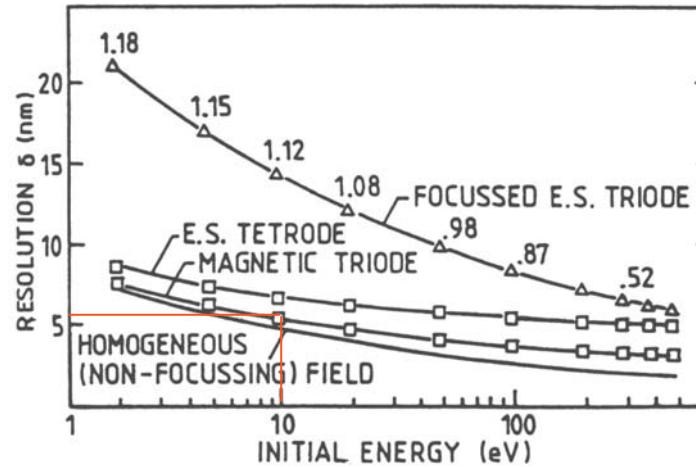
Optimum aperture r and resolution-limiting contributions



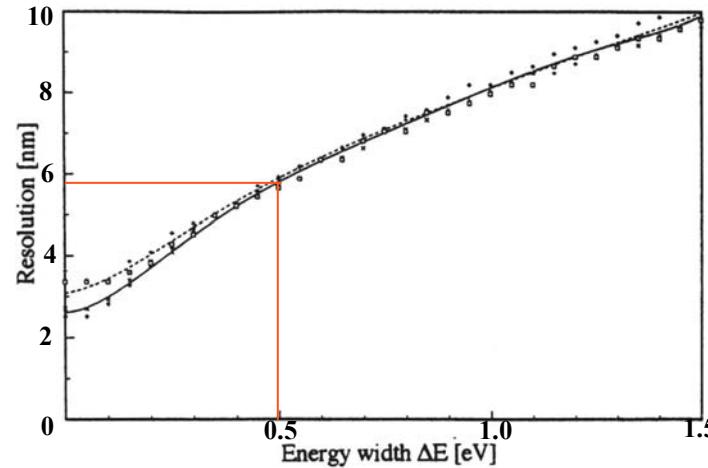
J. Chmelik et al, Optik 83 (1989)155

T. Müller, M.S. thesis, TU Clausthal 1995

Resolution with optimum aperture E -dependence at fixed $\Delta E = 0.5 \text{ eV}$, $U_0 = 20 \text{ keV}$



ΔE -dependence at fixed $E = 10 \text{ eV}$, $U_0 = 18 \text{ kV}$
magnetic triode



Transmission

limited by angle accepted by contrast aperture (r_A)

Axial distance (in back focal plane) of electron starting at angle α

$$r \approx f \sin \alpha \sqrt{E_0/E} \quad (f \text{ focal length})$$

$$\sin \alpha \approx (r/f) \sqrt{E_0/E}$$

Examples for $f = 10 \text{ mm}$, $E = 20000 \text{ eV}$, $r_A = 10 \mu\text{m}$

E_0	2 eV	200 eV
$\sin \alpha$	0.2	0.02
α	11.5°	1.15°

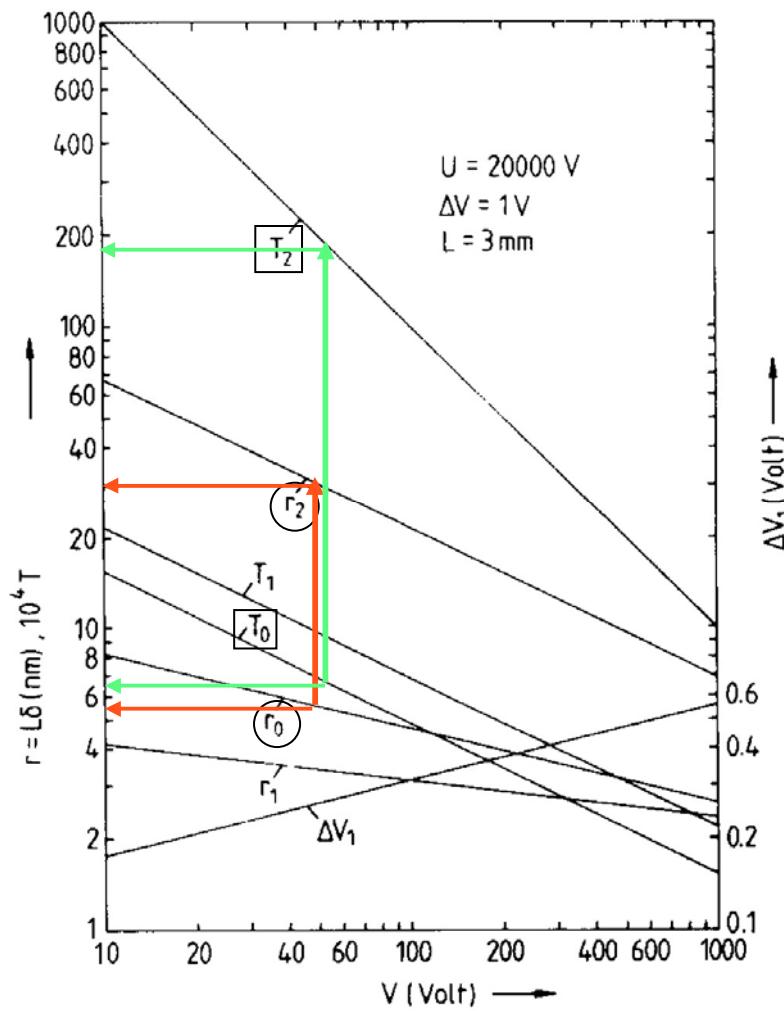
In emission microscopy (wide α range) optimum resolution condition reduces transmission T , therefore

optimize T^n/d^2 instead of $1/d^2$

For $\cos \alpha$ distribution $T = \pi \sin^2 \alpha$

$$T^n/d^2 = \pi \sin^2 n/d^2$$

Transmission T_n , resolution r_n of homogeneous field

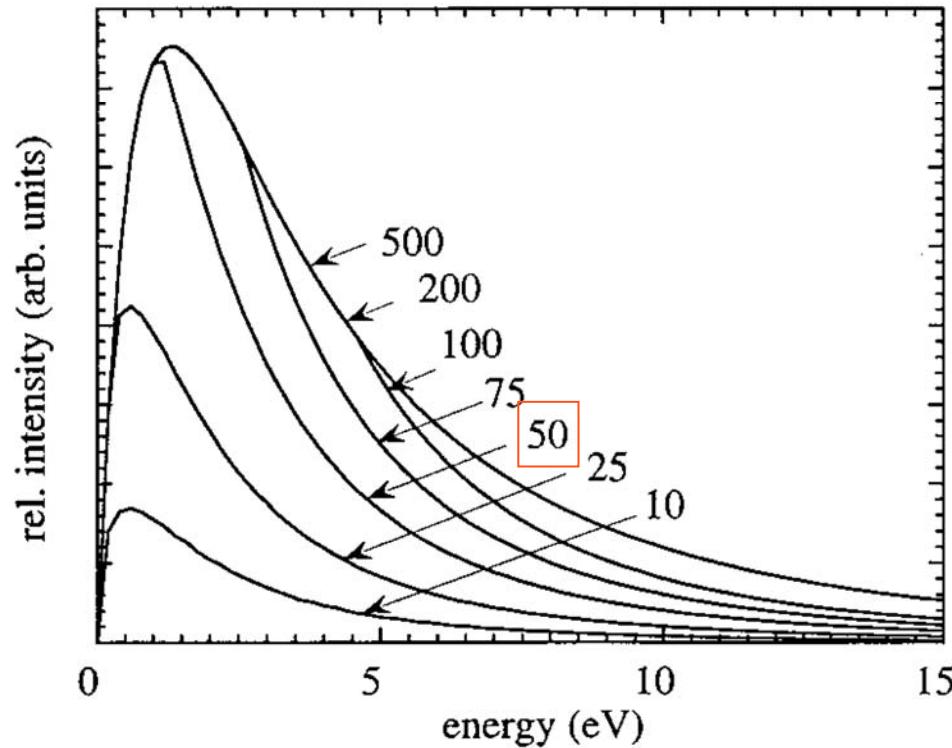


T_2

**T and d
equally
weighted:**

50 eV:
 $\Delta T \approx 30$
 $\Delta d \approx 5$

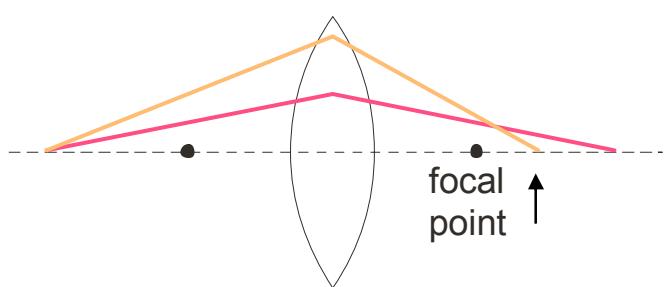
Influence of angle-limiting aperture on the energy distribution of secondary electrons



Work function $\Phi = 4$ eV, accelerating voltage $V = 20$ kV
Parameter: aperture diameter in μm , ALS PEEM

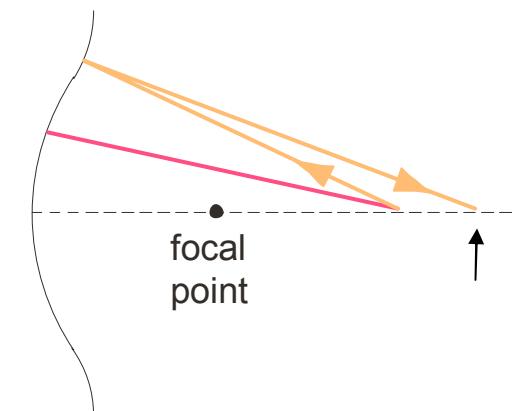
Aberration correction in electron optics

Round **convex** lenses



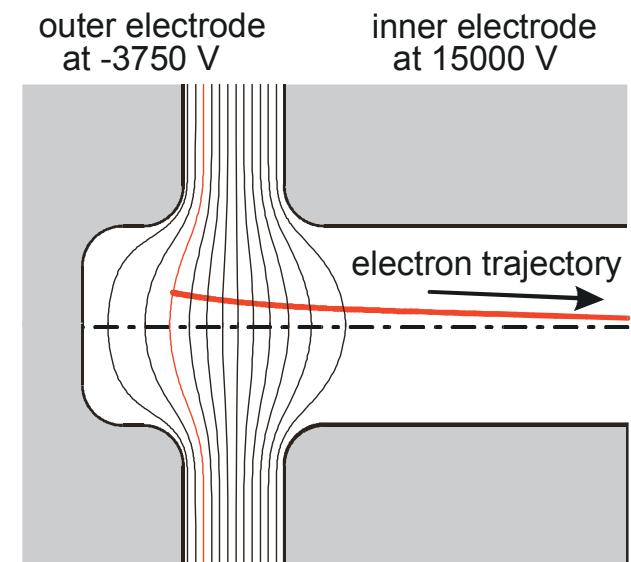
Spherical aberration

electrostatic mirror



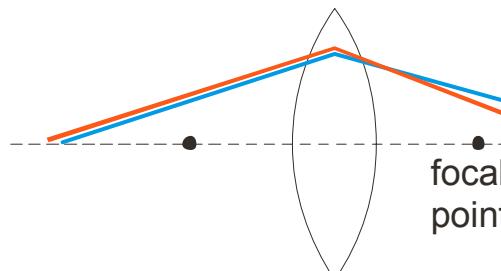
focal point

Equipotential surfaces
in a diode mirror



outer electrode
at -3750 V

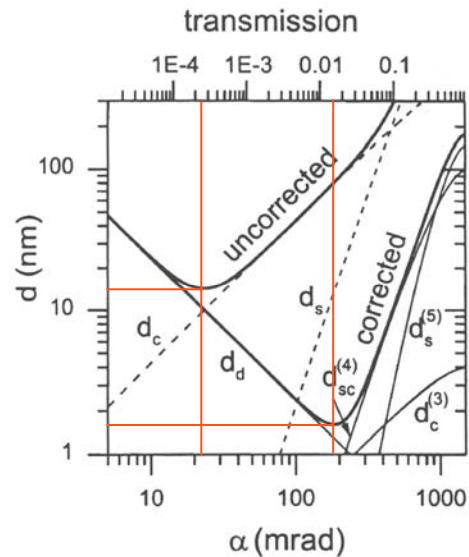
inner electrode
at 15000 V



Chromatic aberration

Resolution and transmission improvement with aberration correction

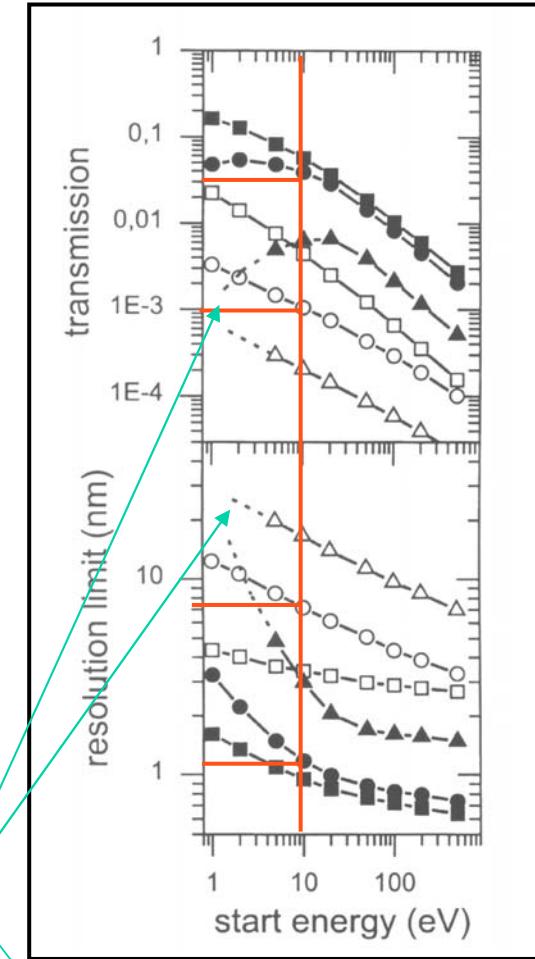
Example: SMART



$$E_0 = 10 \text{ eV}, \Delta E = 2 \text{ eV}, F = 5 \text{ kV/mm}$$

Calculations: D. Preikszas
From Th. Schmidt et al,
Surf. Rev. Lett. 9 (2002) 223

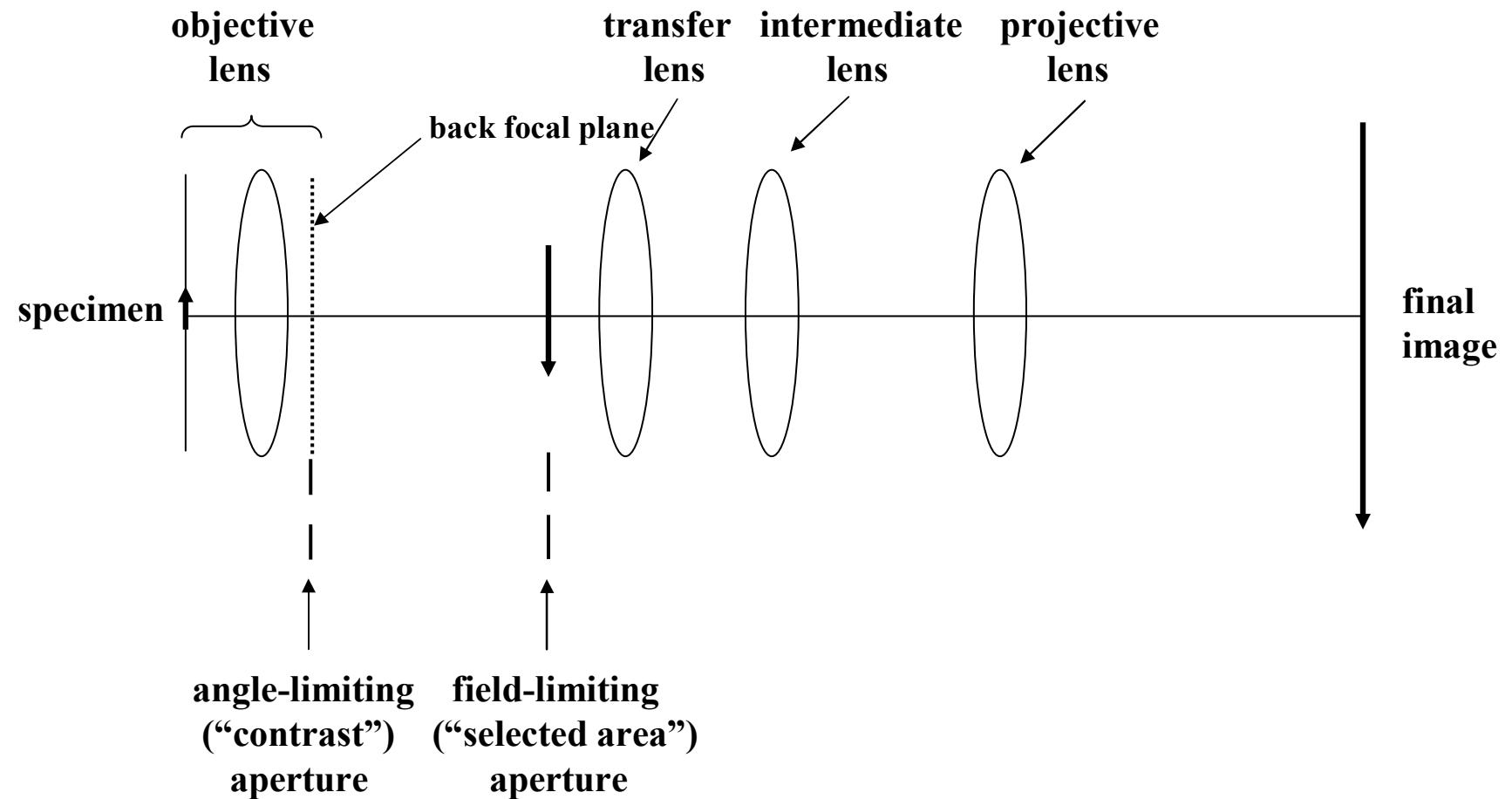
ΔE (eV)
 0.1 □ ■
 1.0 ○ ●
 5.0 △ ▲
 without correction with correction



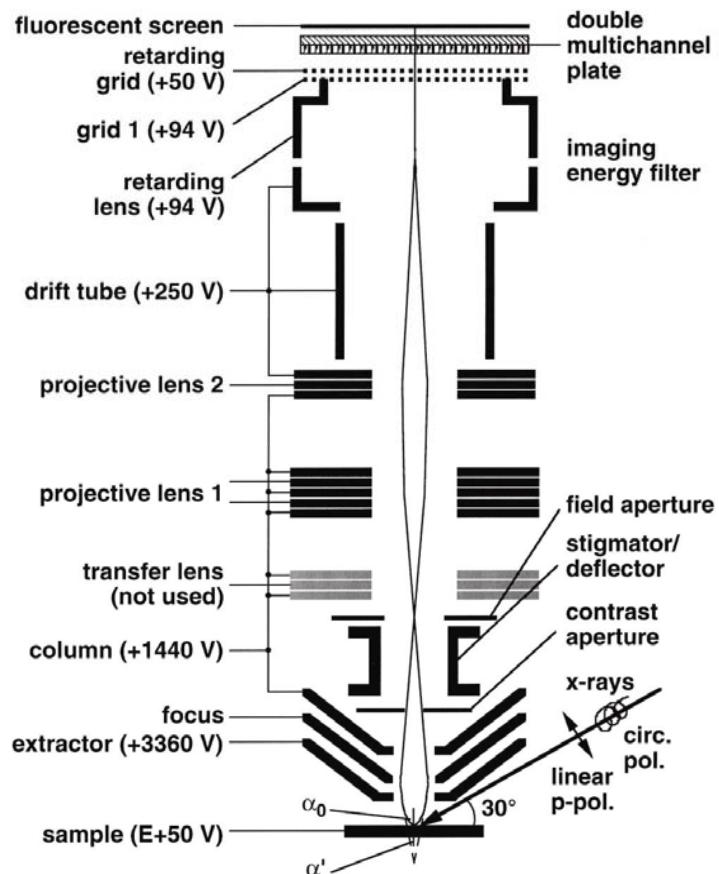
Energy filter needed for
secondary electrons

Instruments

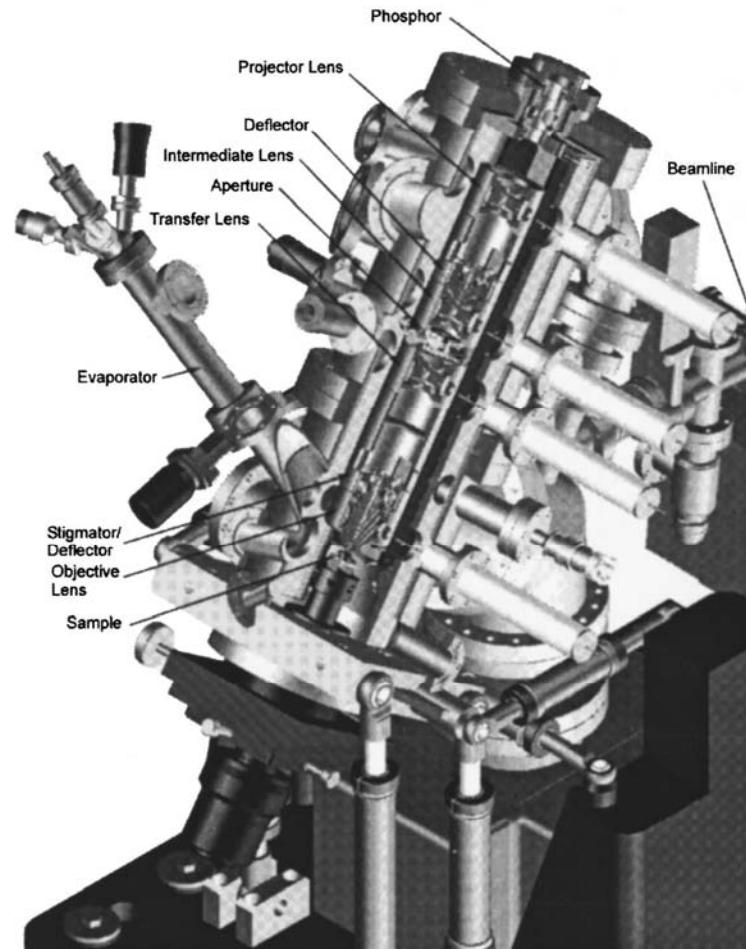
Basic PEEM schematic



Electrostatic PEEM examples

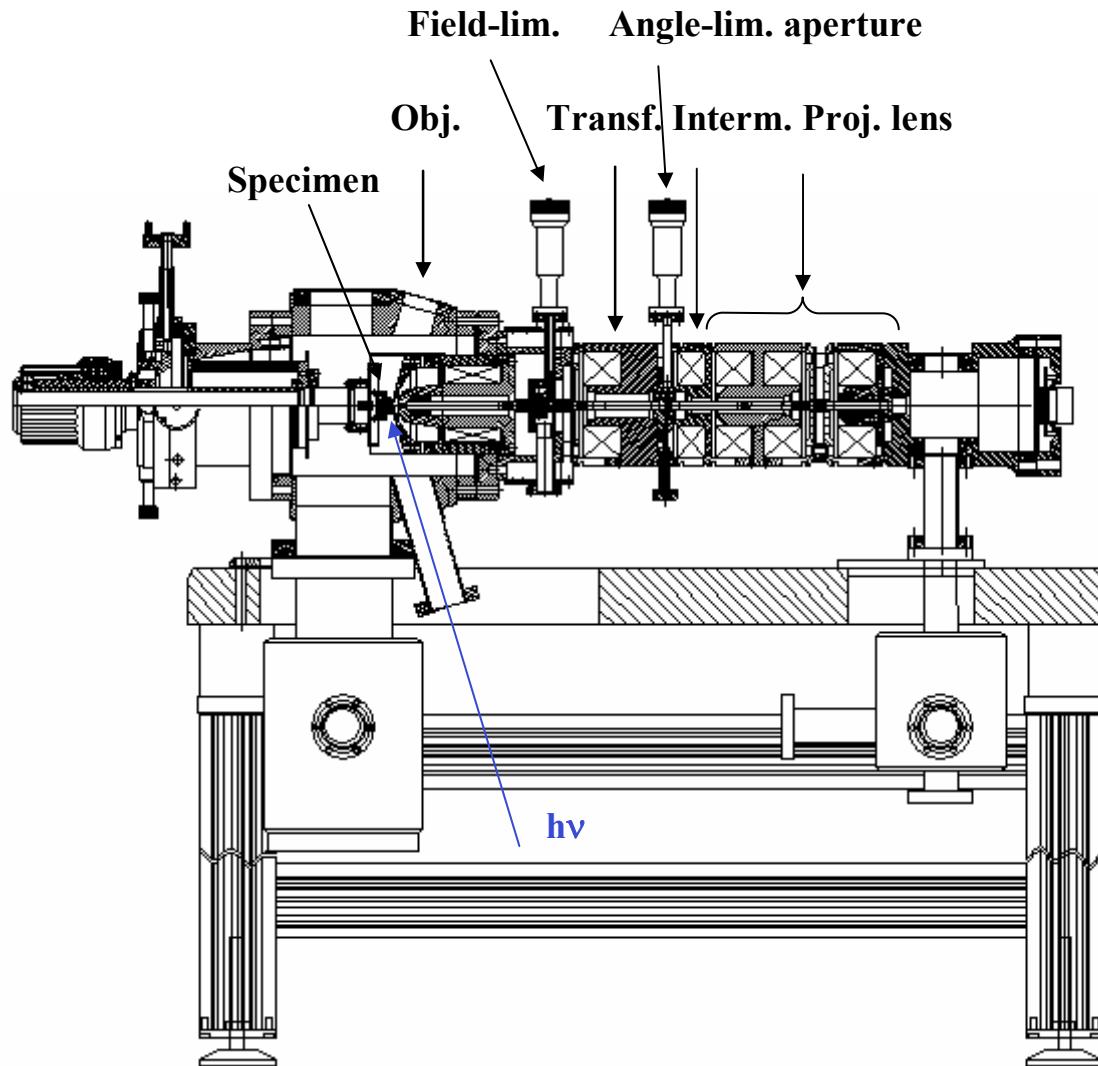


**Focus PEEM
with high pass filter**



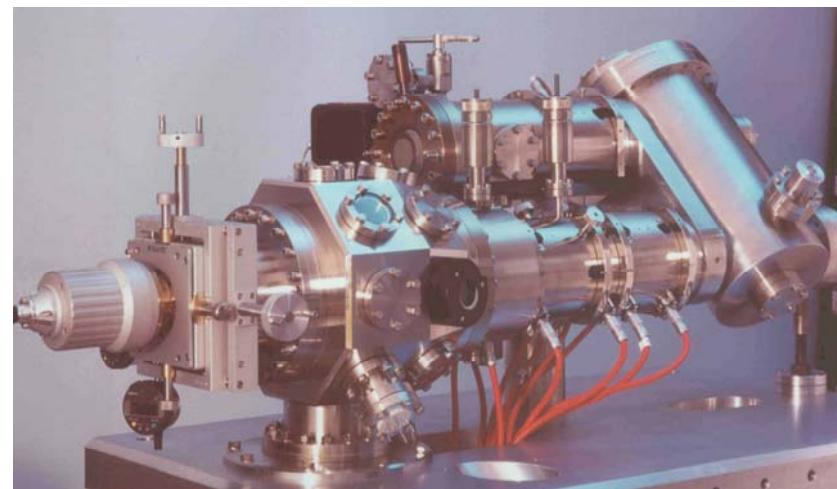
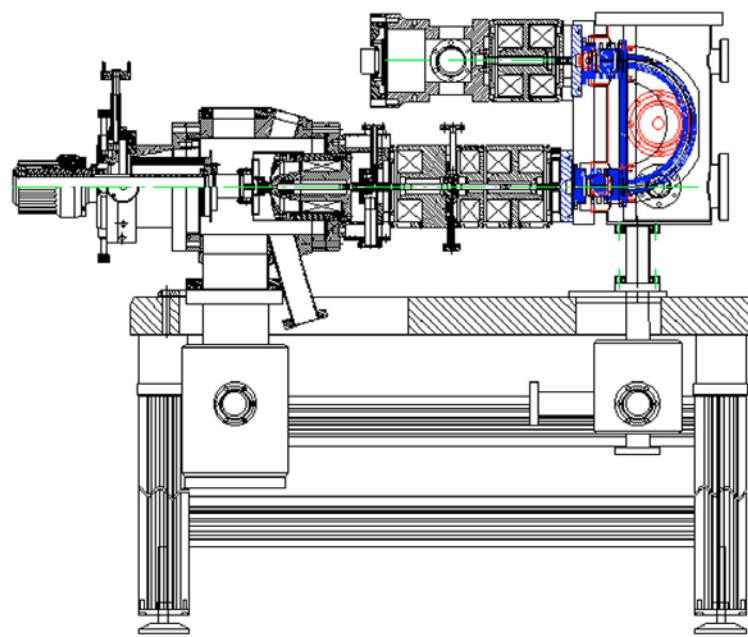
ALS PEEM II

Magnetic PEEM (ELMITEC)



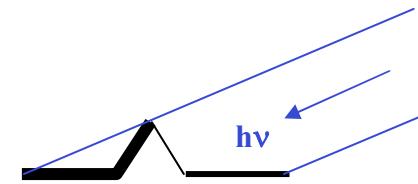
Spectroscopic PEEM with band pass filter

ELMITEC



Contrast mechanisms

- 1 Topographic contrast due to oblique illumination and field distortion
- 2 Work function contrast at low E_0 (escape probability!)
- 3 Chemical contrast due to inner shell ionization
- 4 Magnetic contrast via XMCD and XMLD

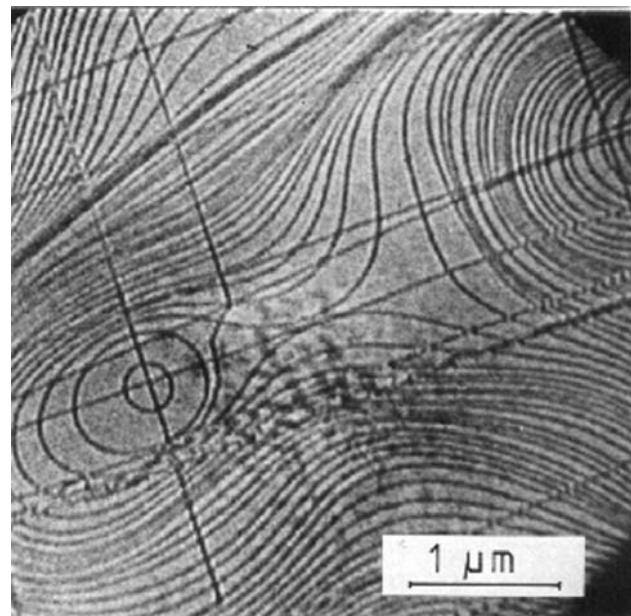


No structural contrast, therefore combination with
Low Energy Electron Microscopy
(LEEM)

The usefulness of LEEM

Properties not visible with PEEM, but with LEEM

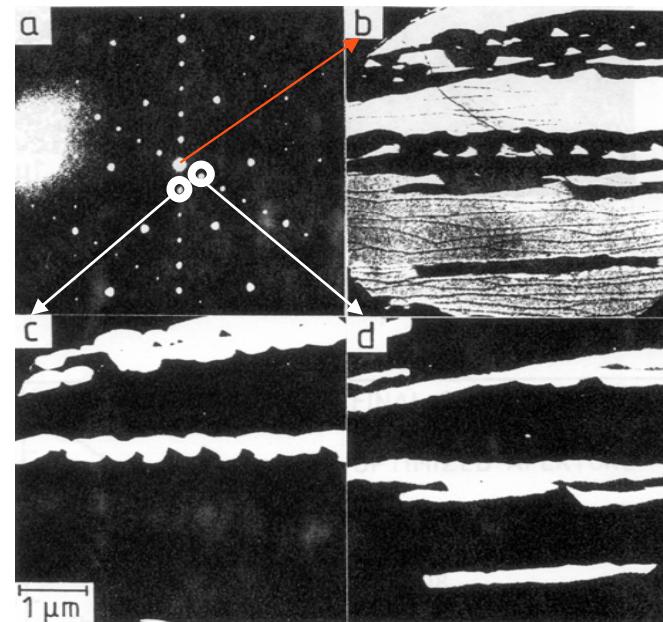
atomic steps



Mo(110)

Interference contrast

domain orientations



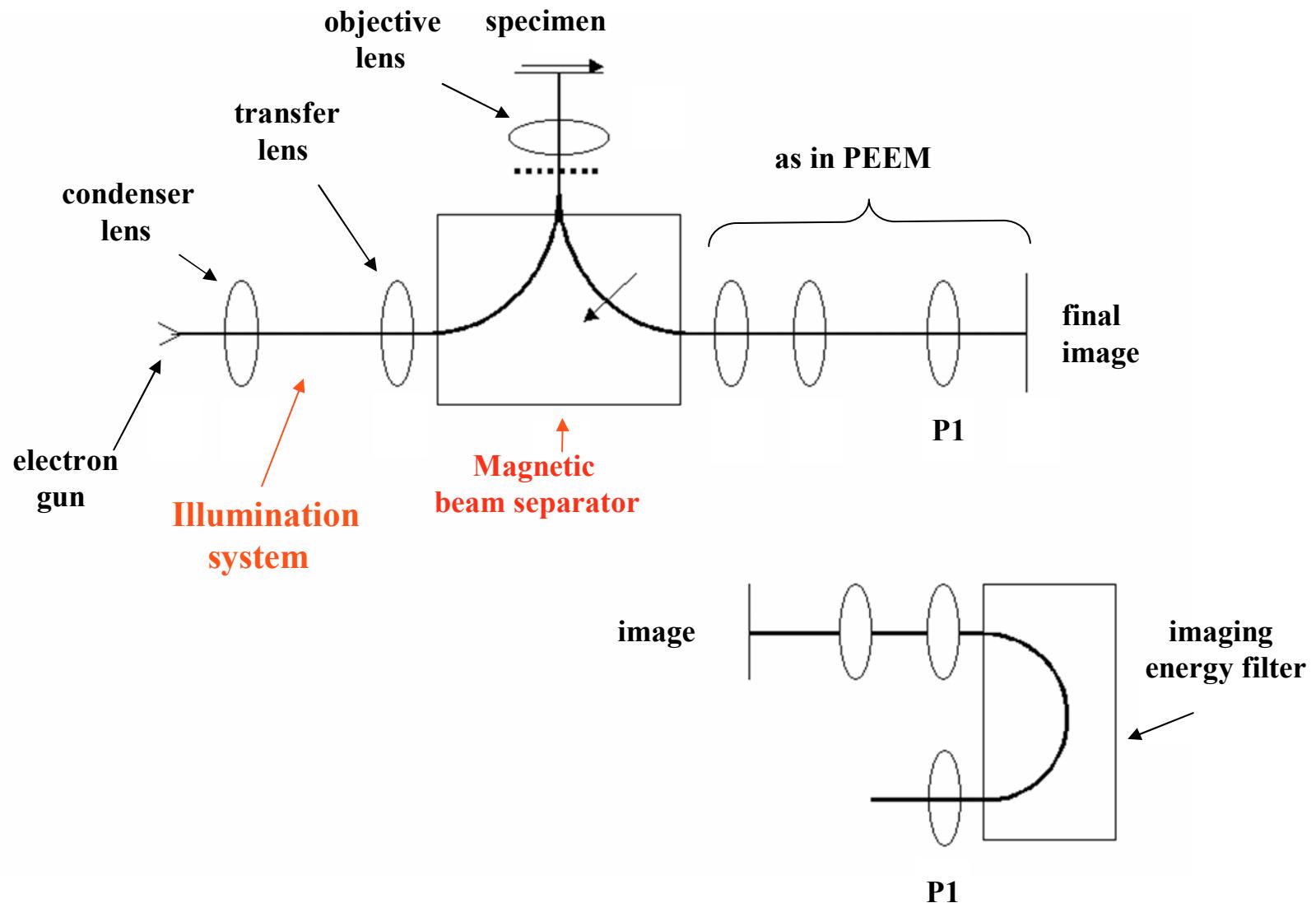
Au($\sqrt{3} \times \sqrt{3}$)-R30° + Au(5 × 2) on Si(111)
b
c,d

Diffraction contrast

LEEM also much brighter and better resolution ⇒ use for focusing in XPEEM

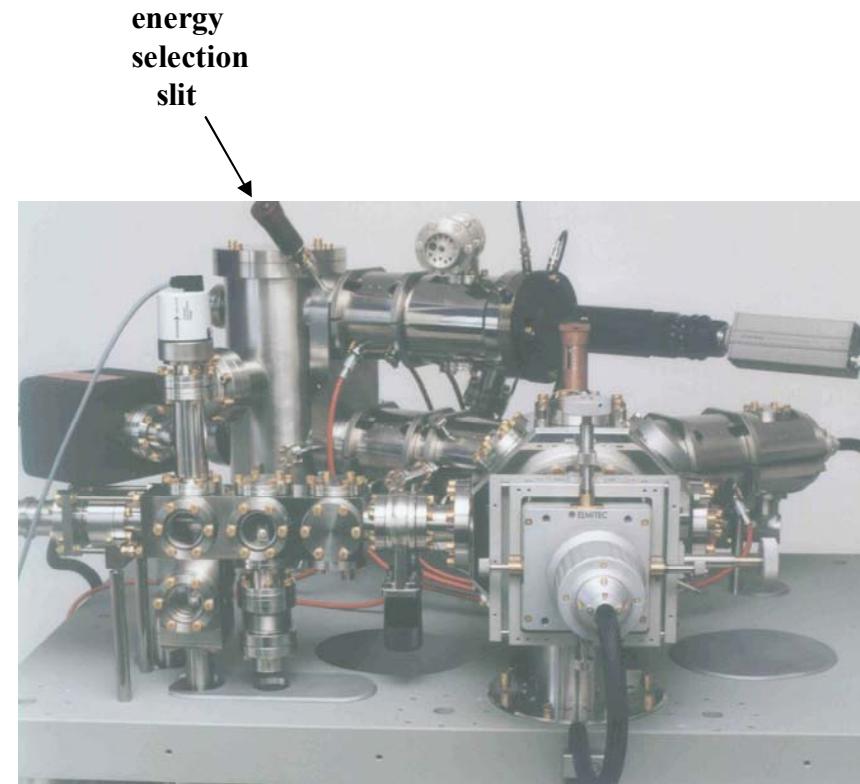
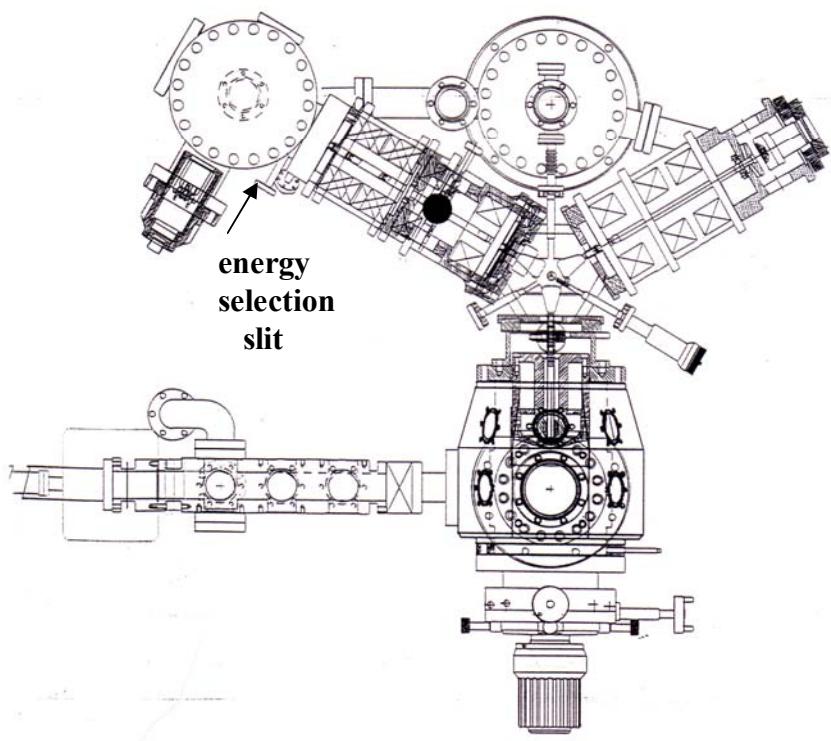
LEED much easier to interpret than PED ⇒ use for structure analysis

Basic LEEM schematic

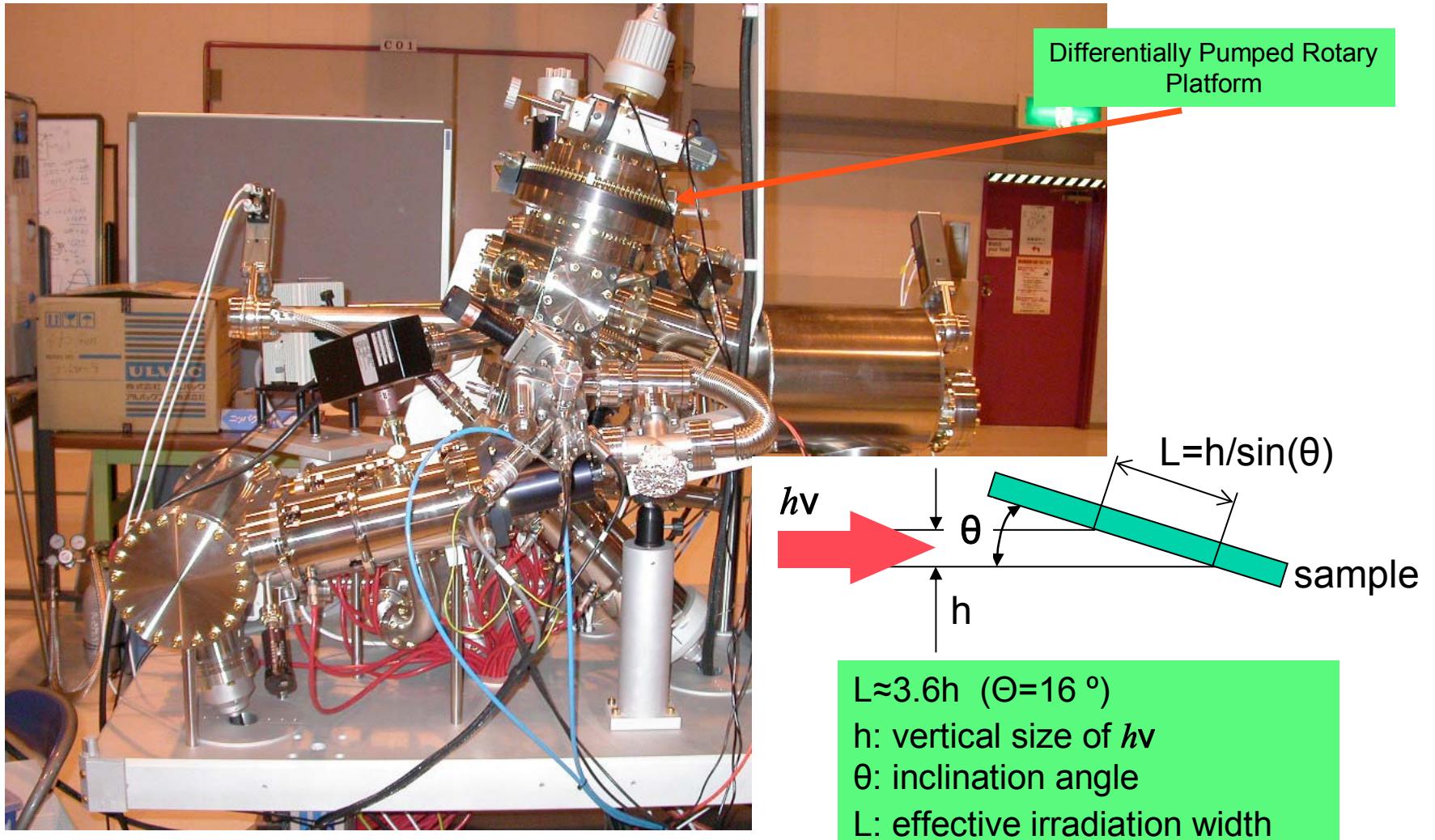


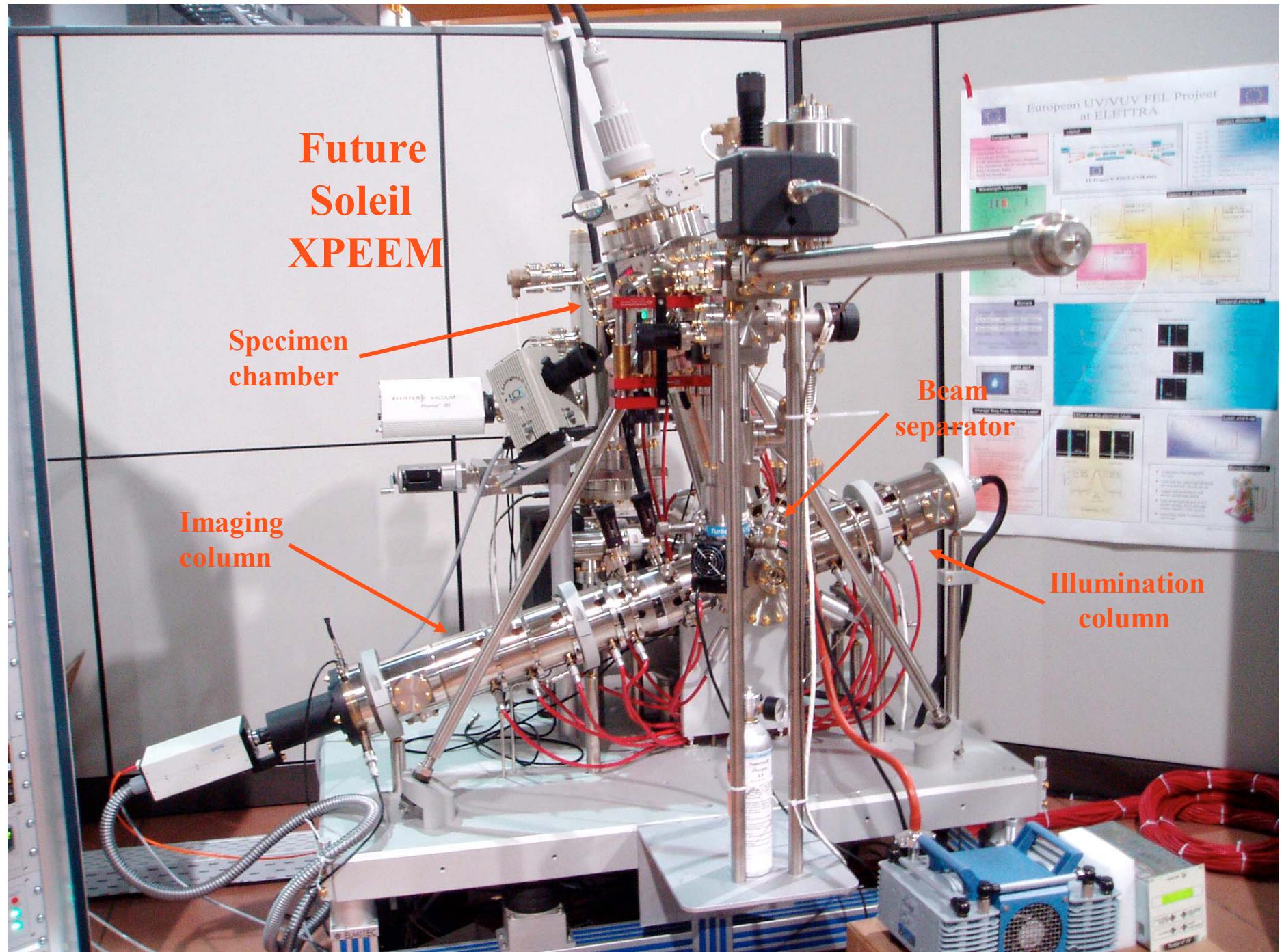
Spectroscopic Photo Emission and Low Energy Electron Microscope

SPELEEM ELMITEC



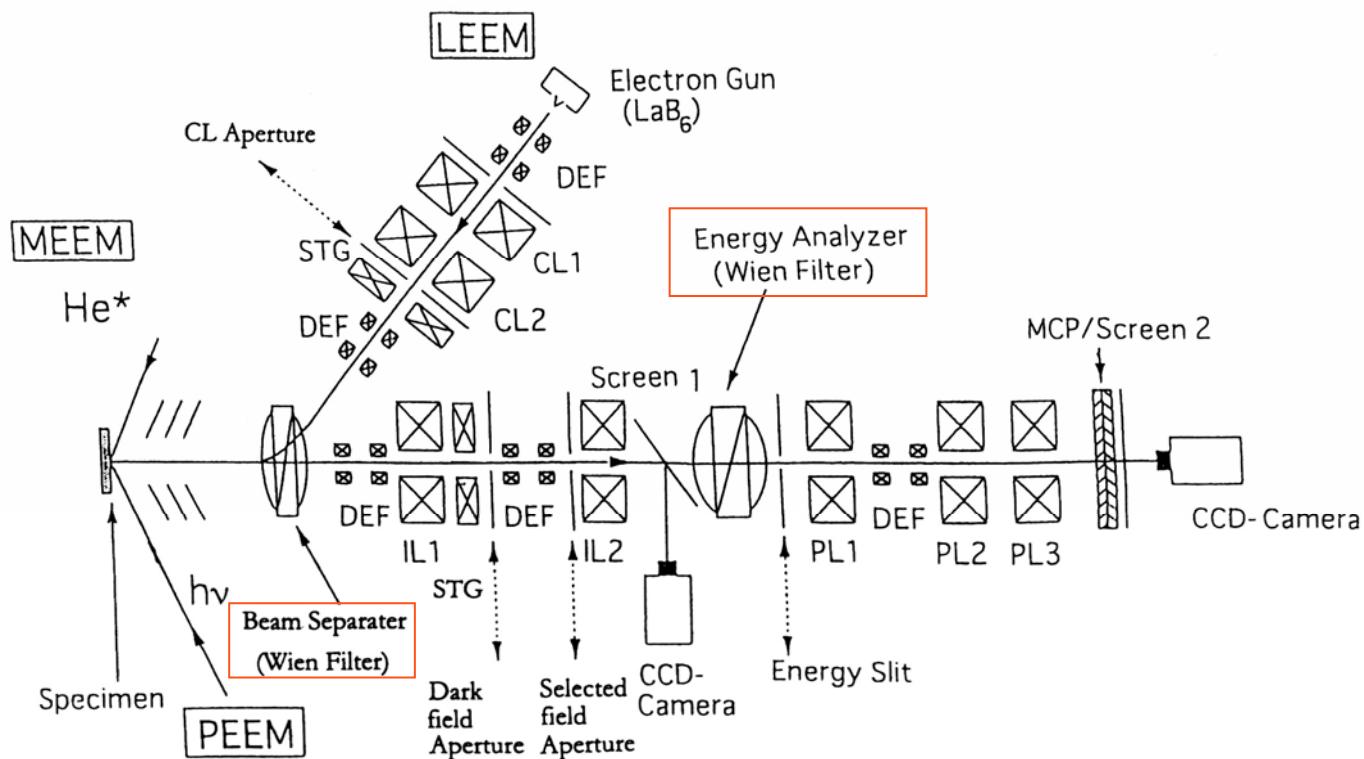
SPELEEM side view





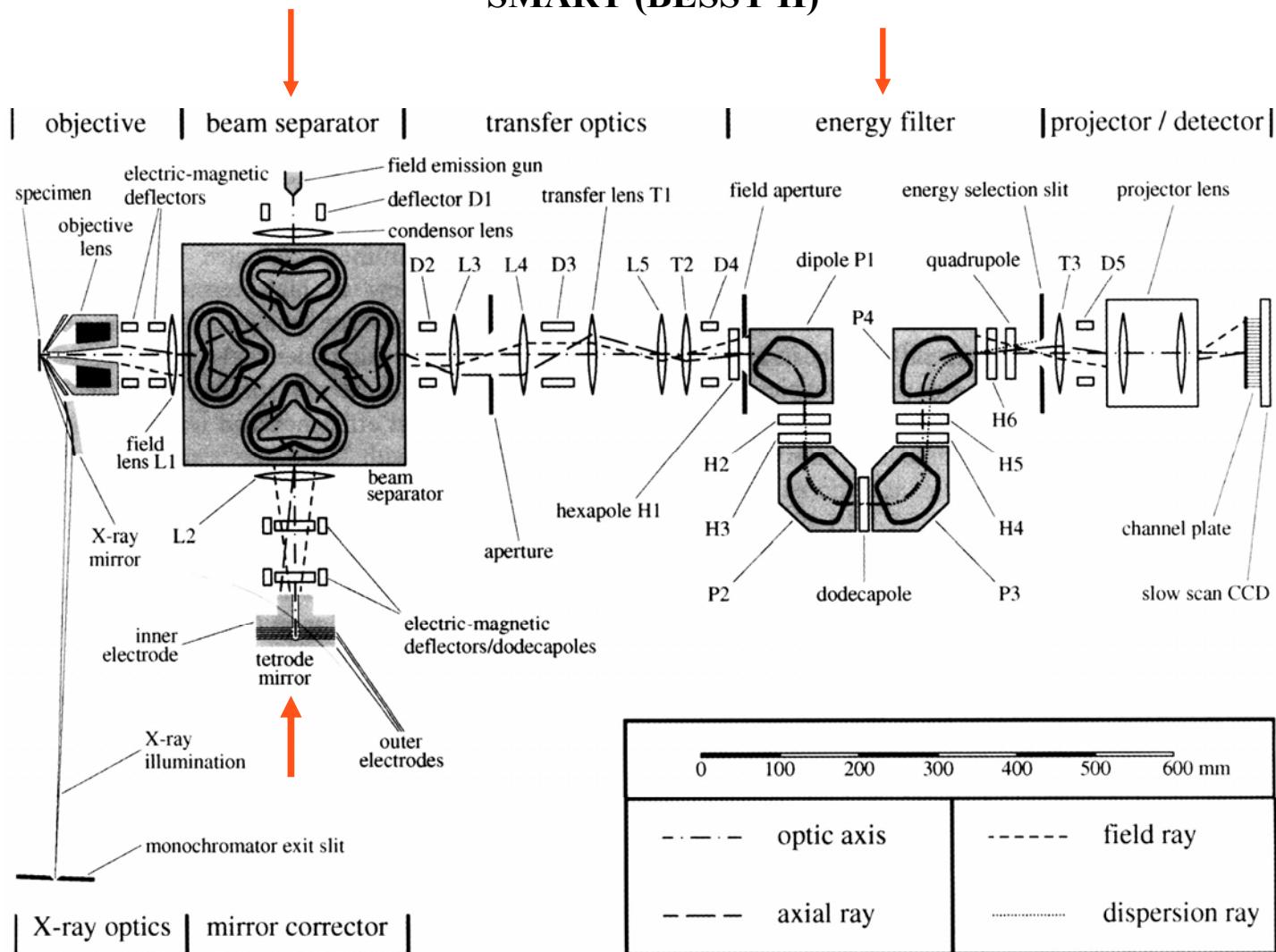
LEEM with energy filter

JEOL



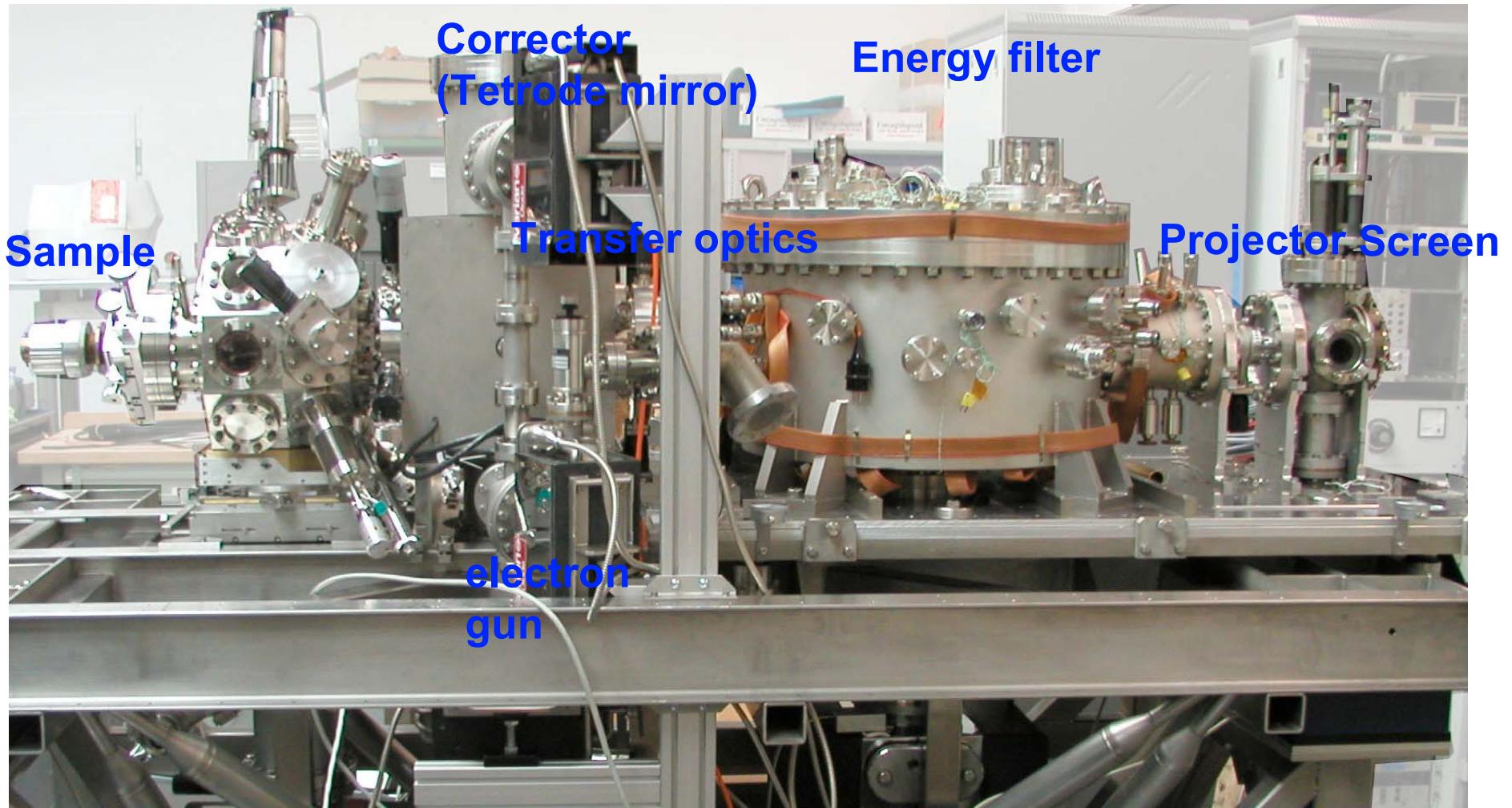
Aberration-corrected SPELEEM

SMART (BESSY II)



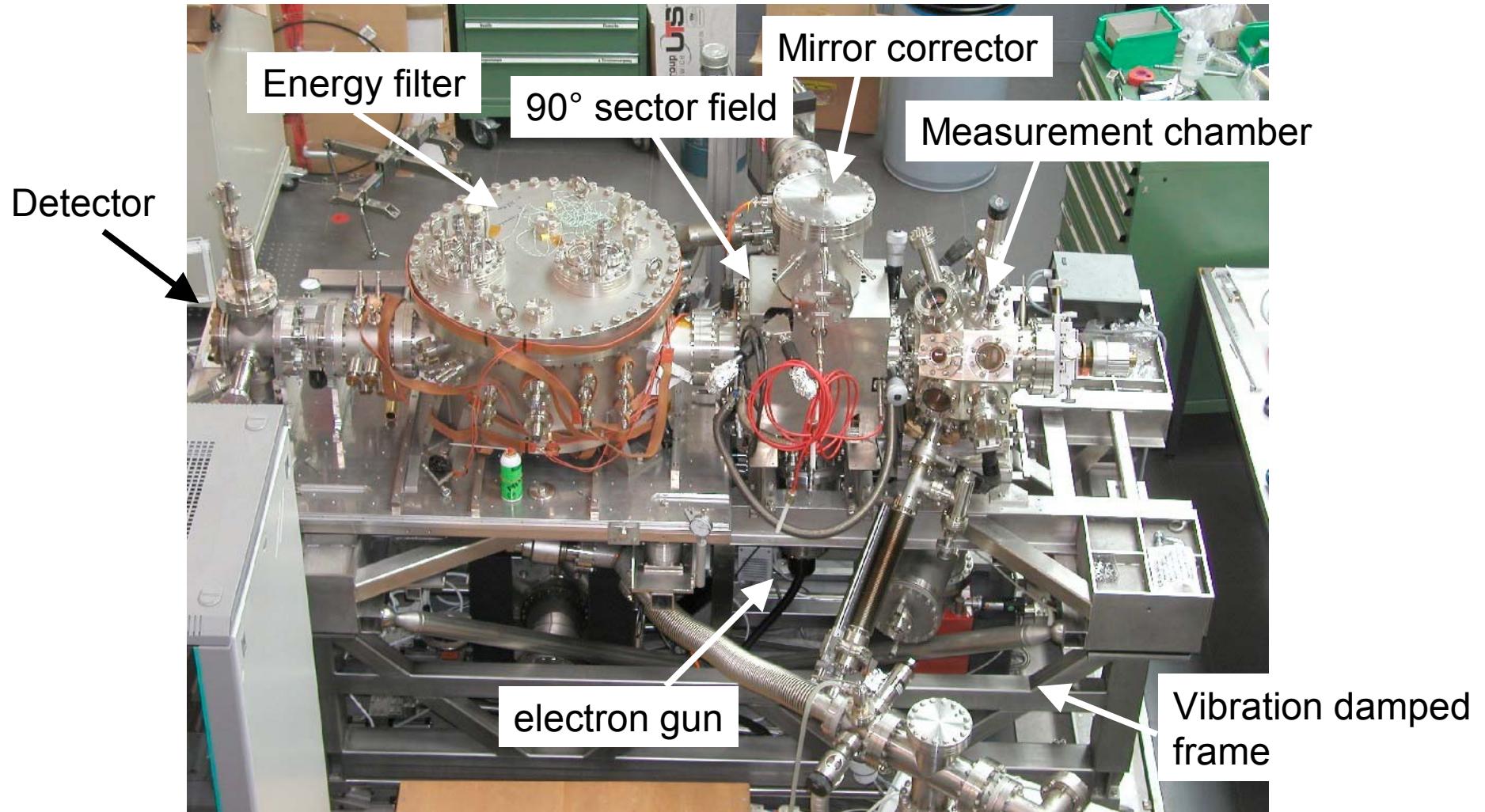
SMART side view

Aberration corrected PEEM/LEEM with energy filtering



Th. Schmidt April 2004

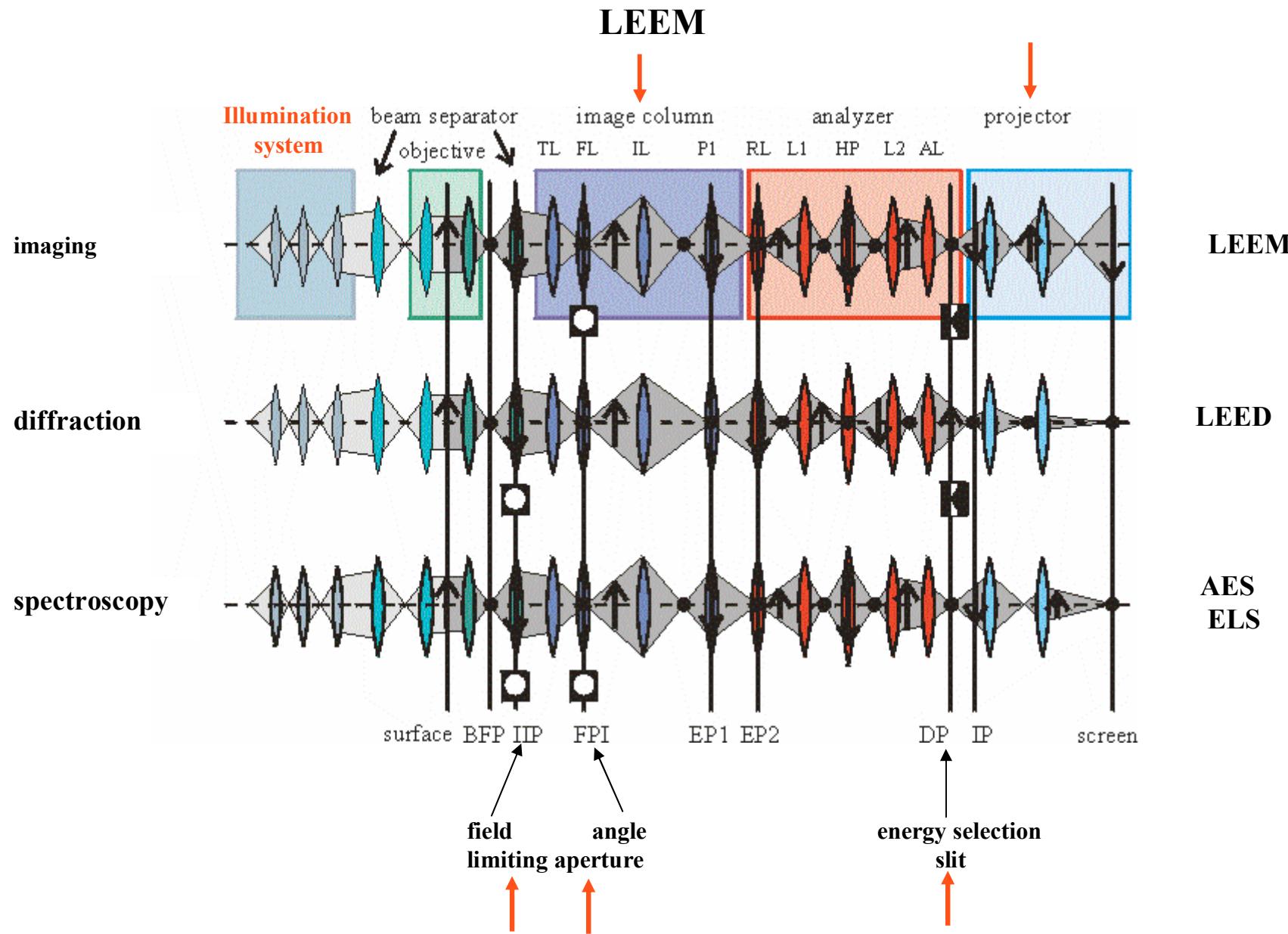
SMART top view



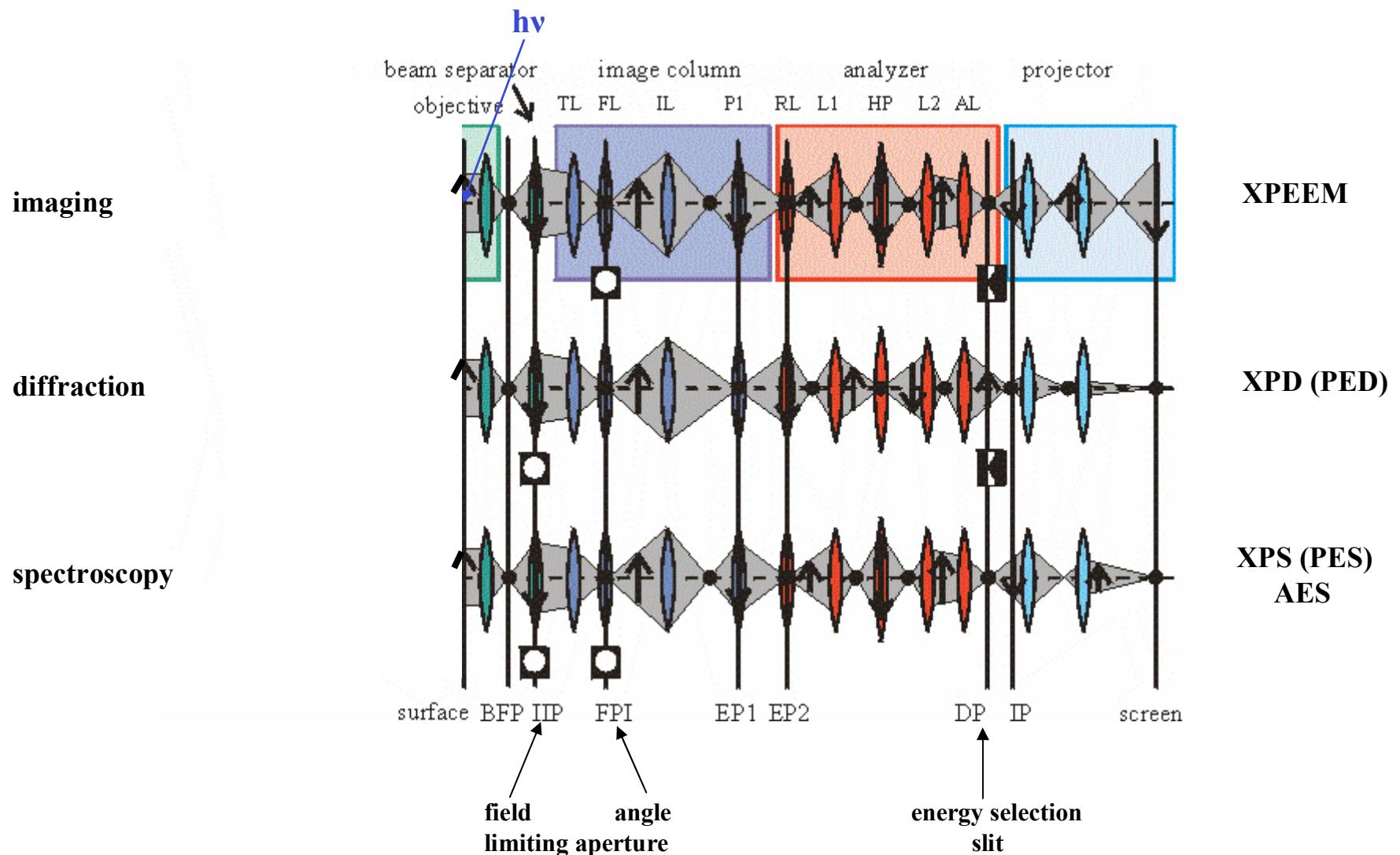
Th. Schmidt April 2004

Methodic

Operation modes of a SPELEEM



Operation modes of a SPELEEM with photons



PEEM practice

**Ultrahigh vacuum (low 10^{-10} torr range)
but experiments up to 10^{-5} torr range possible**

Surface cleaning: heating, sputtering or chemical reactions, e.g with oxygen for carbon removal

Choice of optimum photon energy:

**Secondary electron imaging: $h\nu \approx E_i$ in:
XANES, NEXAFS, XMCD, XMLD**

Photo electron imaging: $E_i + 50 \text{ eV} < h\nu < E_i + 200 \text{ eV}$

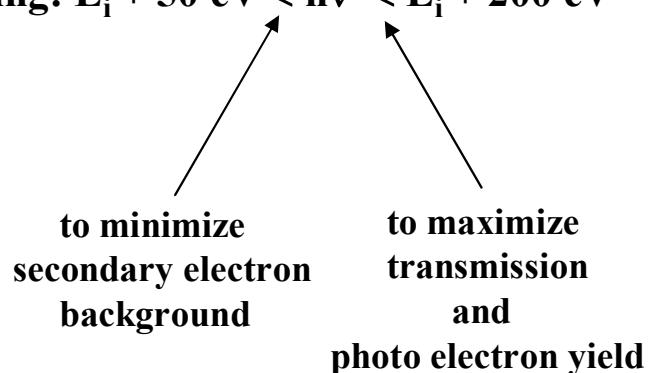
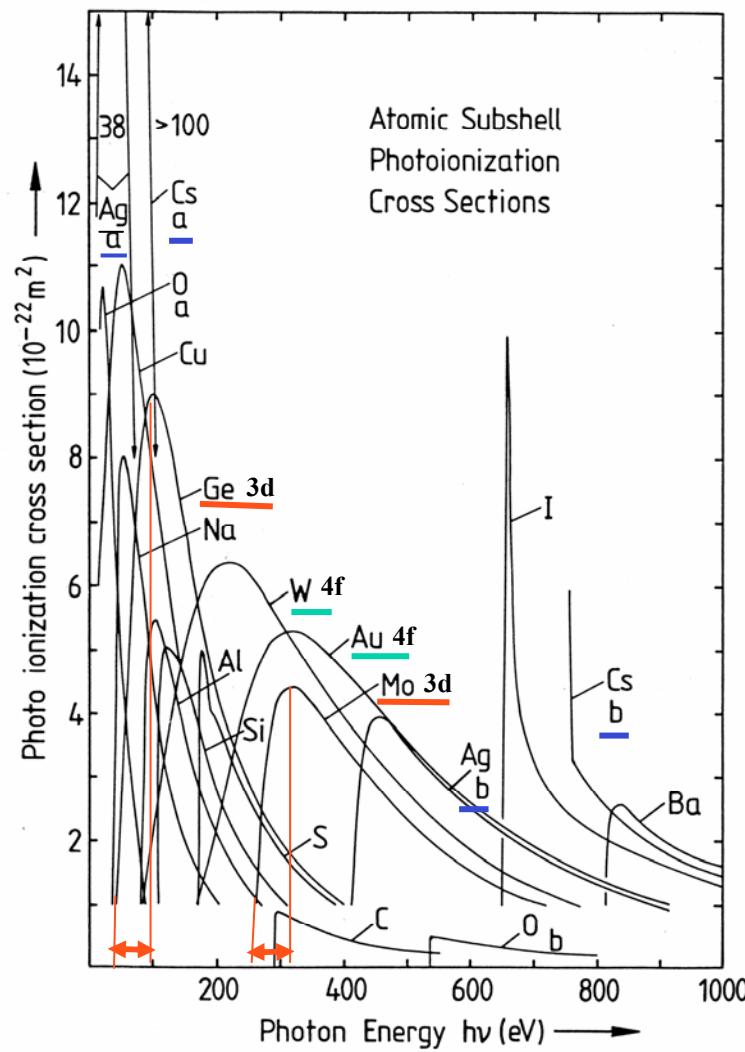


Photo ionization cross sections

Photon energy selection



Binding energies (eV)

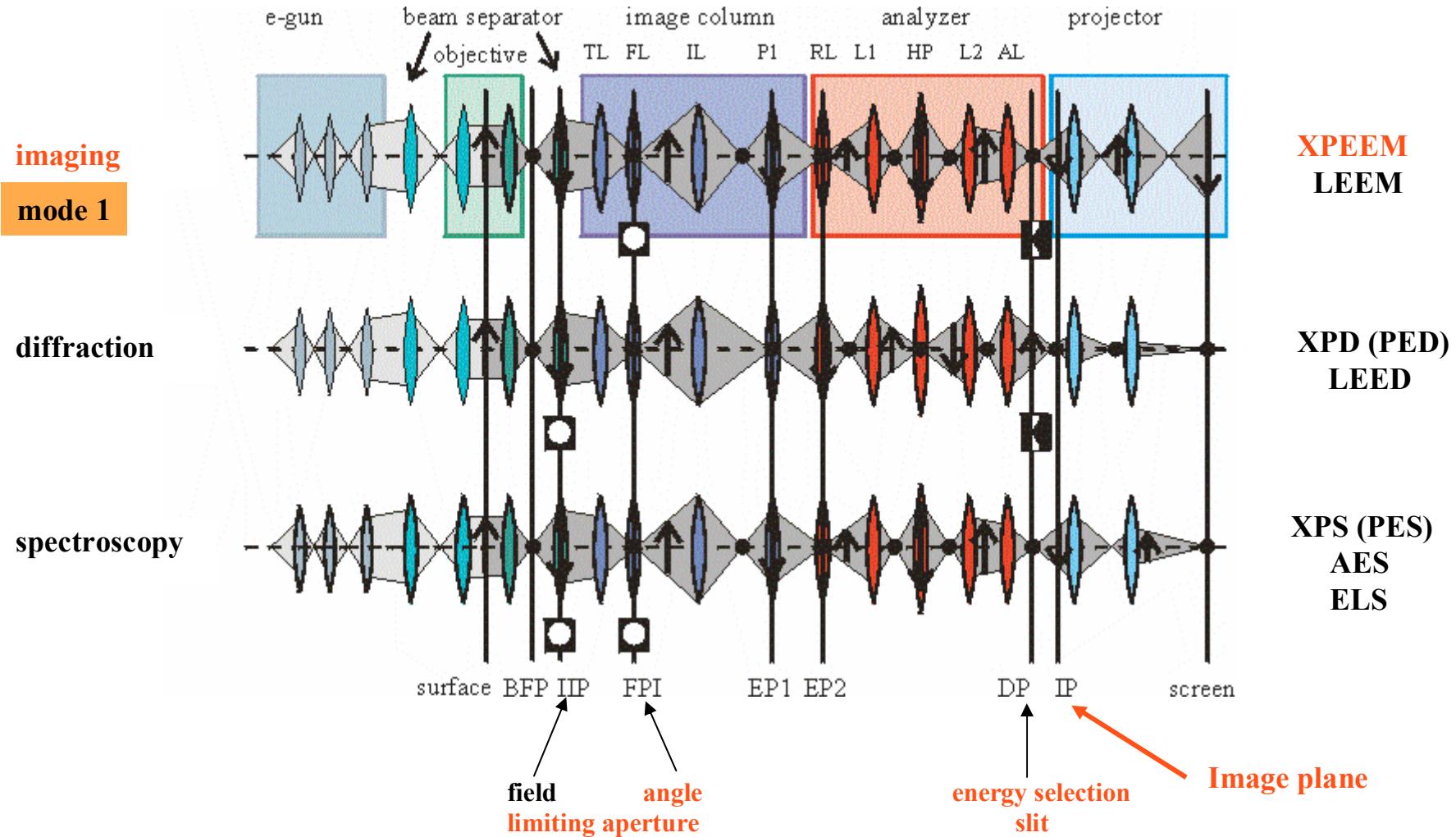
Ge 3d 29.8, 29.2
Mo 3d 231.1, 227.9

W 4f 33.6, 31.4
Au 4f 87.6, 84.0

Ag
a 4d ≈ 5
b 3d 374.0, 368.3
Cs
a 4d 79.8, 77.5
b 3d 740.5, 726.6

J.J. Yeh and I. Lindau,
Atomic Data 1985

Operation modes of a SPELEEM



Chemical imaging (mode 1)

secondary electrons

spatial resolution

$$\sigma_{\text{Ag}4d} \approx 5 \sigma_{\text{W}5d}$$

$$hv = 65 \text{ eV} \gg E_i < 10 \text{ eV}$$

$$\Delta E_F \leq 1 \text{ eV}$$

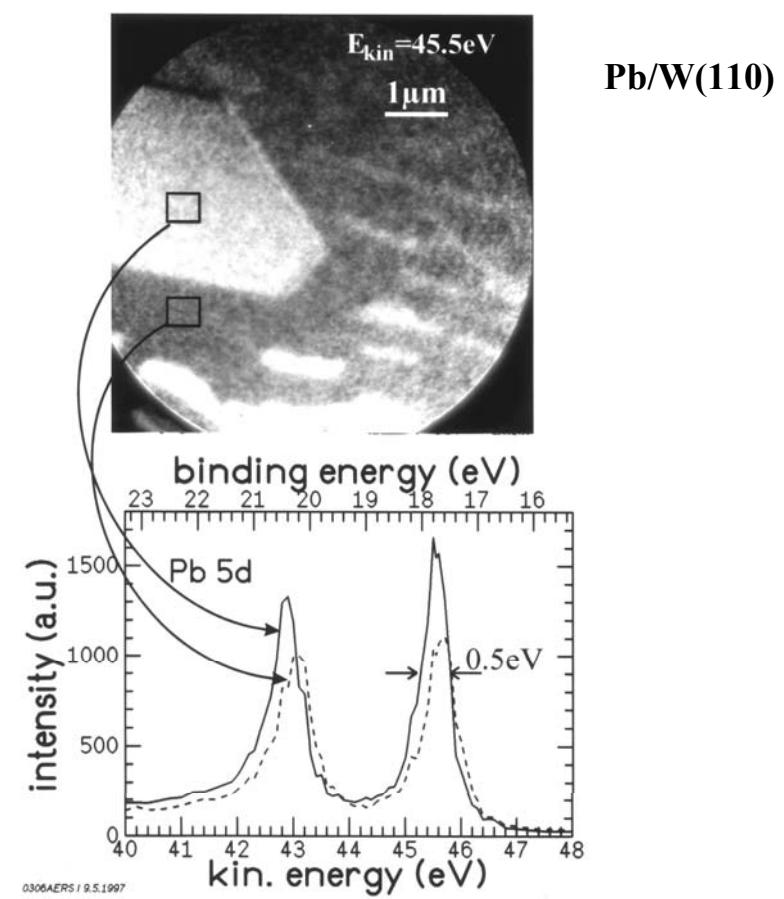
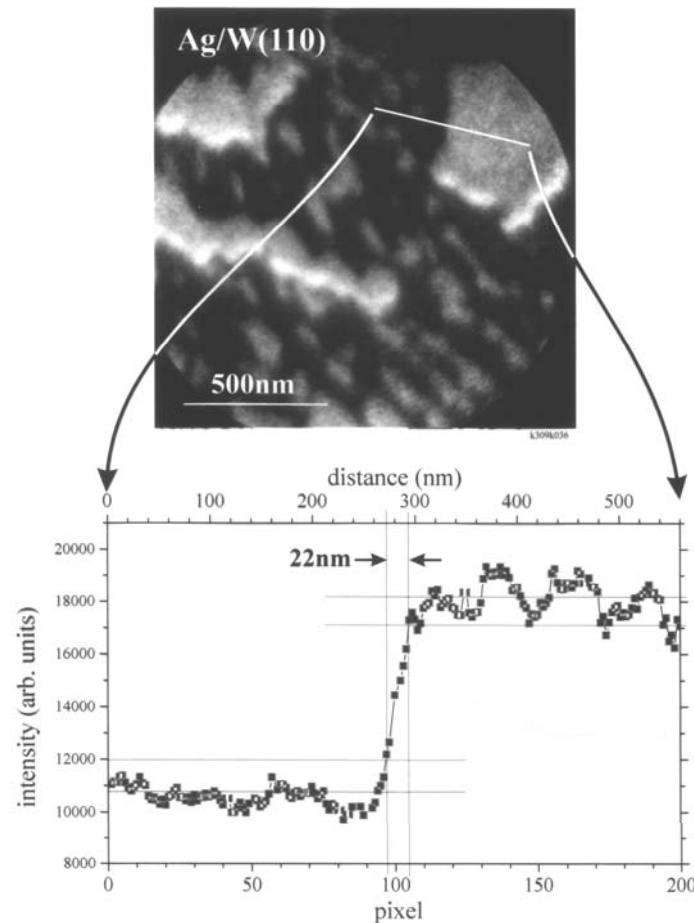
photo electrons

energy resolution

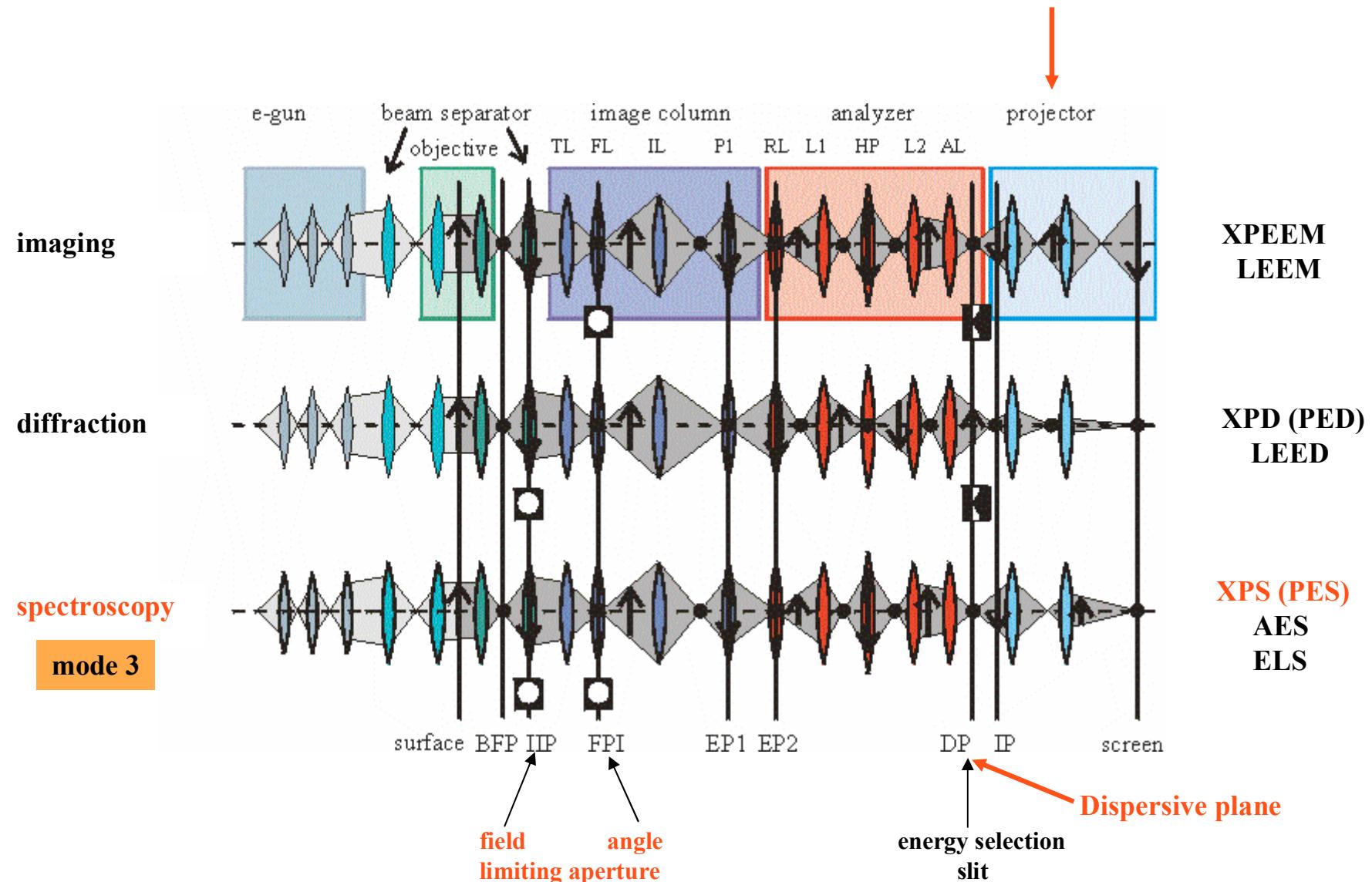
$$hv = 65 \text{ eV, images in } 0.2 \text{ eV steps}$$

10-60 sec/image $0.25 \mu\text{m}^2$ areas

$$\Delta E_F \leq 0.5 \text{ eV}, \Delta E_{\text{chem}} \approx 0.15 \text{ eV}$$

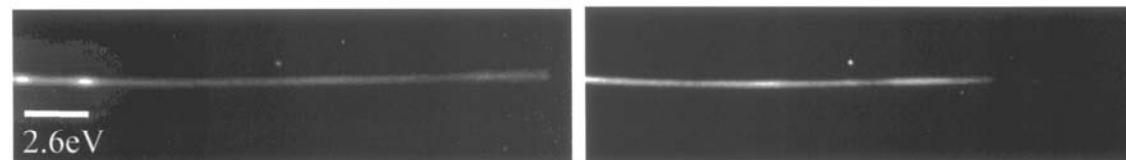


Operation modes of a SPELEEM



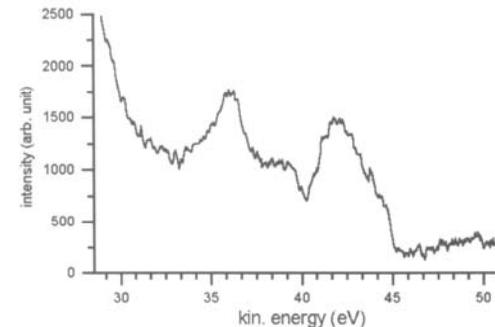
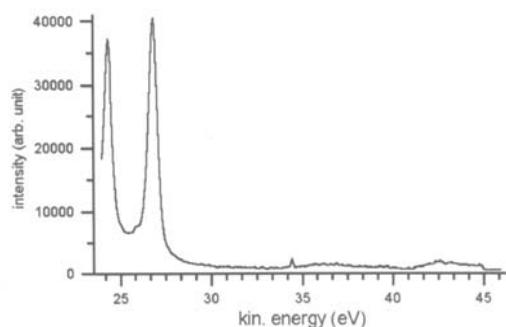
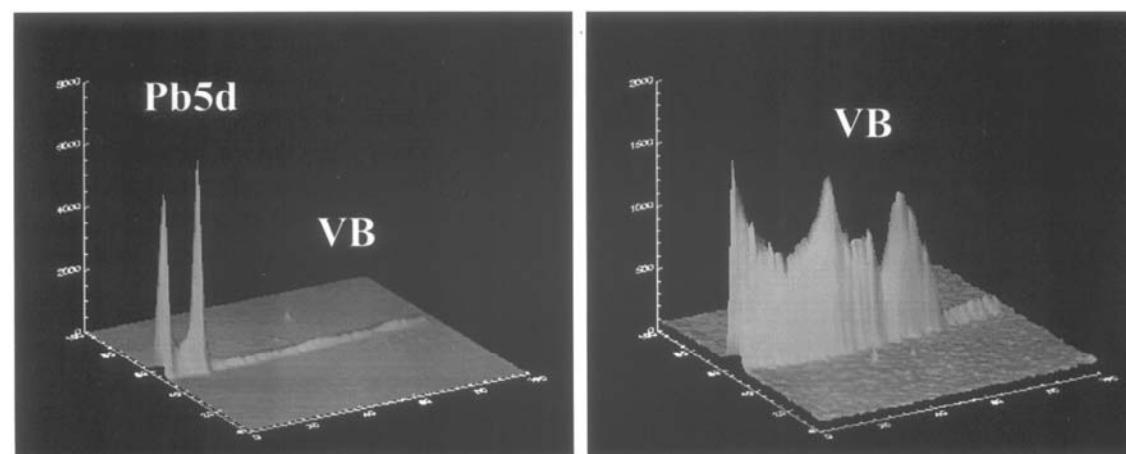
Fast local spectroscopy by imaging the dispersive plane (mode 3)

$\alpha = 8^\circ$ (contrast aperture), $0.8\mu\text{m}^2$ area (selected field aperture)
20 eV full dispersion, 60 sec
 $h\nu = 48$ eV



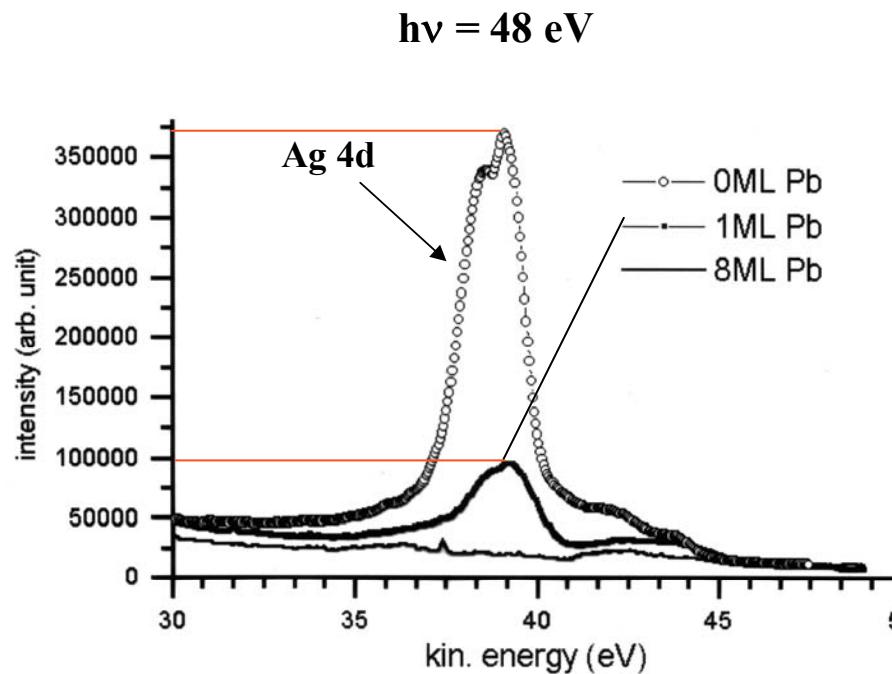
Dispersive plane

8 monolayers
Pb on Si(111)-
 $\sqrt{3}\times\sqrt{3}$ -Ag



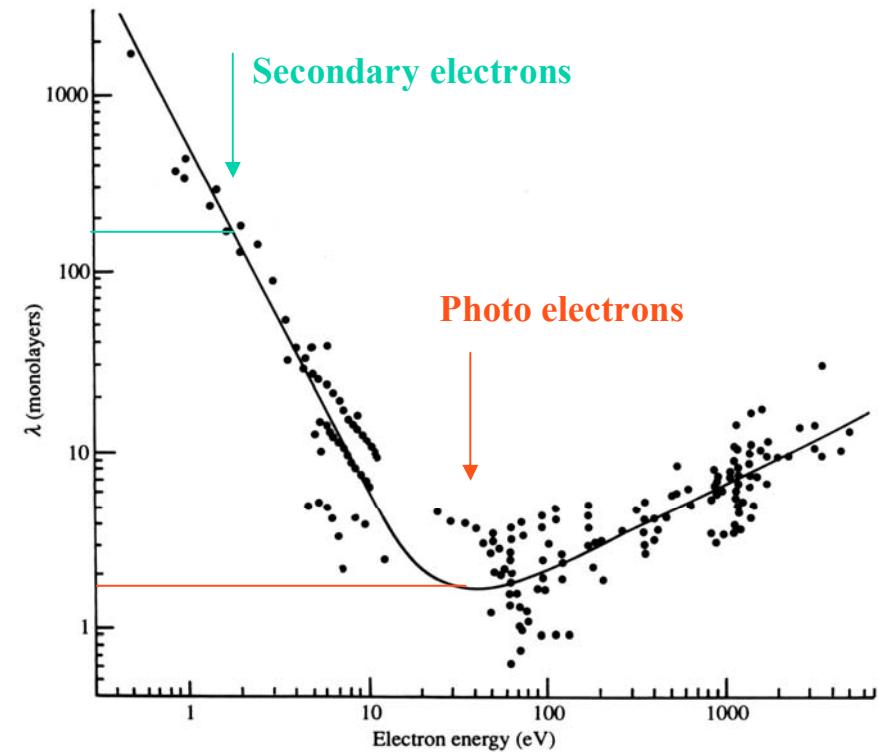
Surface sensitivity of photo electrons versus secondary electrons

valence band region



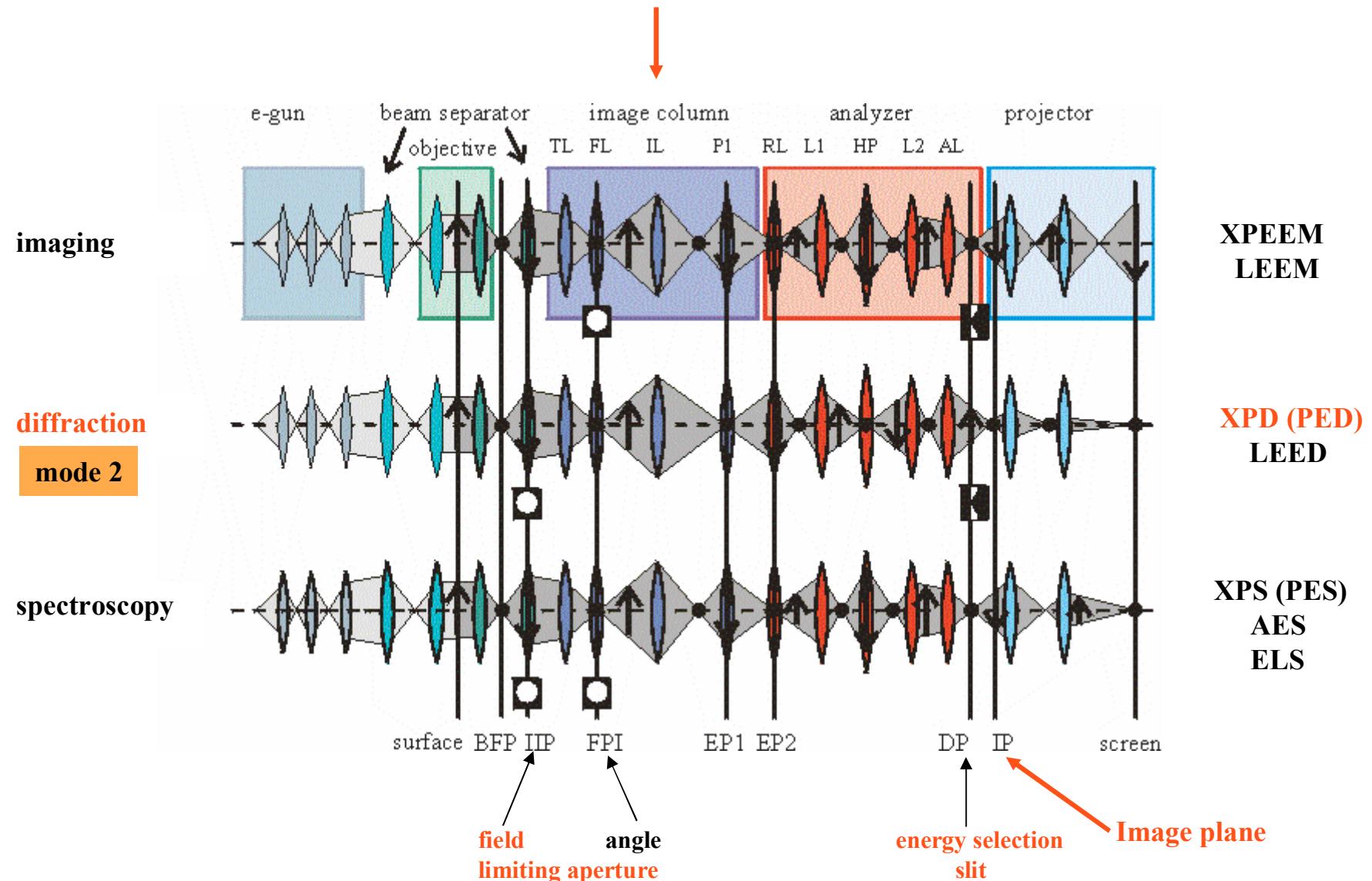
Pb on Si(111) – Ag ($\sqrt{3} \times \sqrt{3}$) – R30°
(1 monolayer Ag)

Th. Schmidt et al, 1998



Inelastic mean free path
("universal curve")
determines sampling depth

Operation modes of a SPELEEM



Local photo electron diffraction (mode 2)

Pb 5d photo electrons

from $0.8\mu\text{m}^2$ area (selected field aperture)

E_{kin}

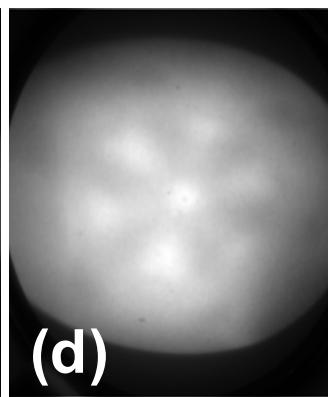
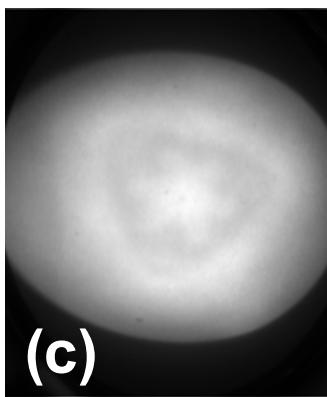
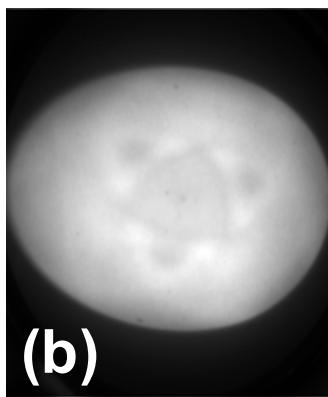
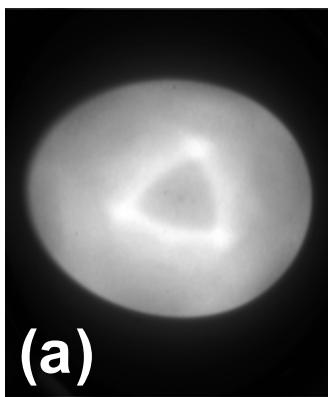
23.0 eV

28.0 eV

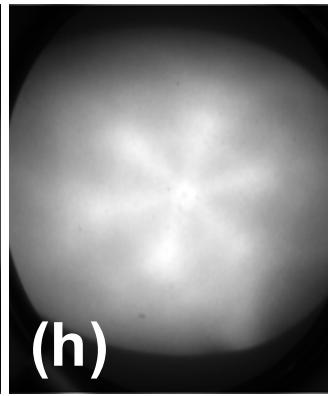
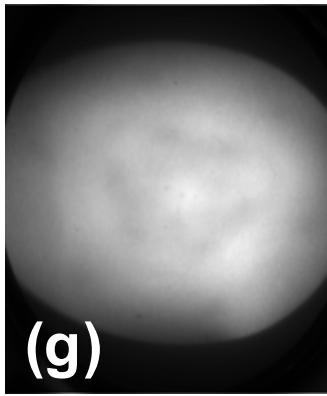
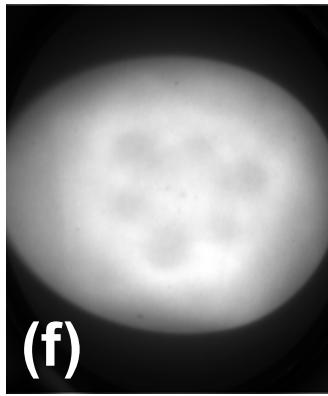
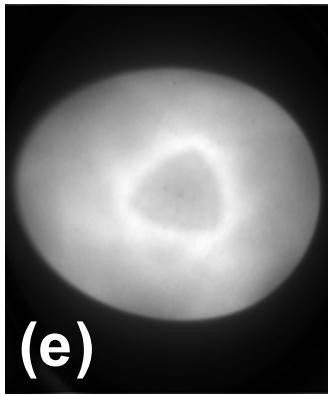
33.0 eV

38.0 eV

$5d_{3/2}$



$5d_{5/2}$



E_{kin}

25.6 eV

30.6 eV

35.6 eV

40.6 eV

$\hbar\nu$

43.5 eV

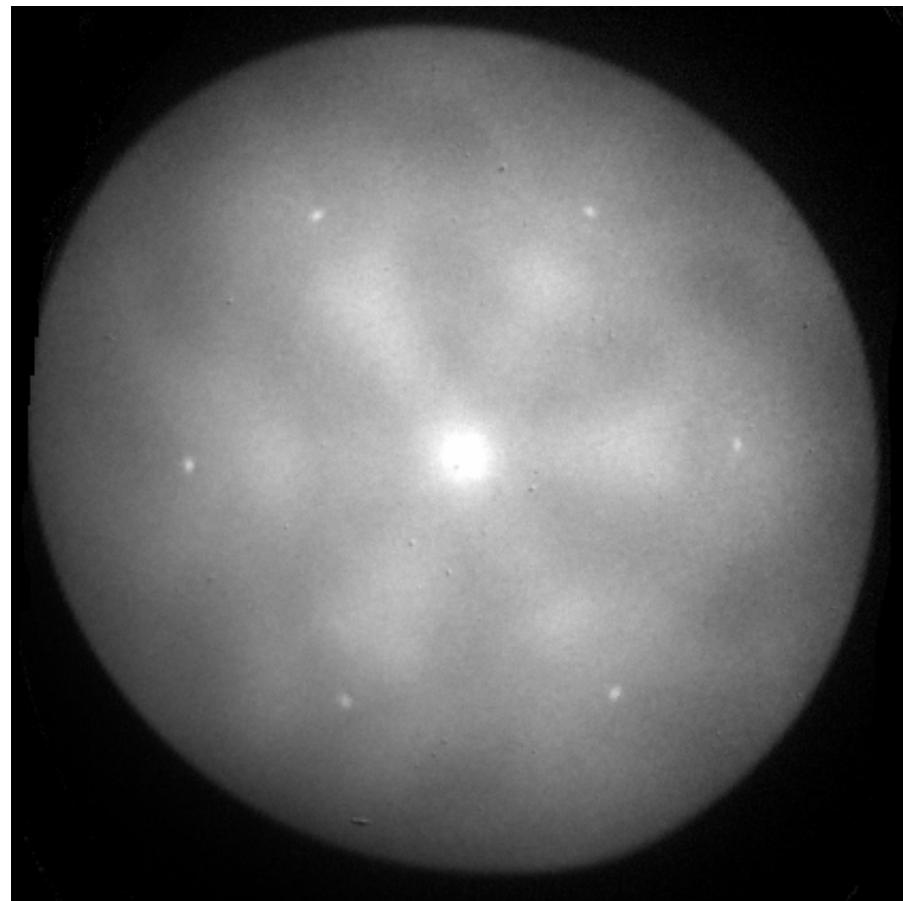
48.5 eV

53.5 eV

58.5 eV

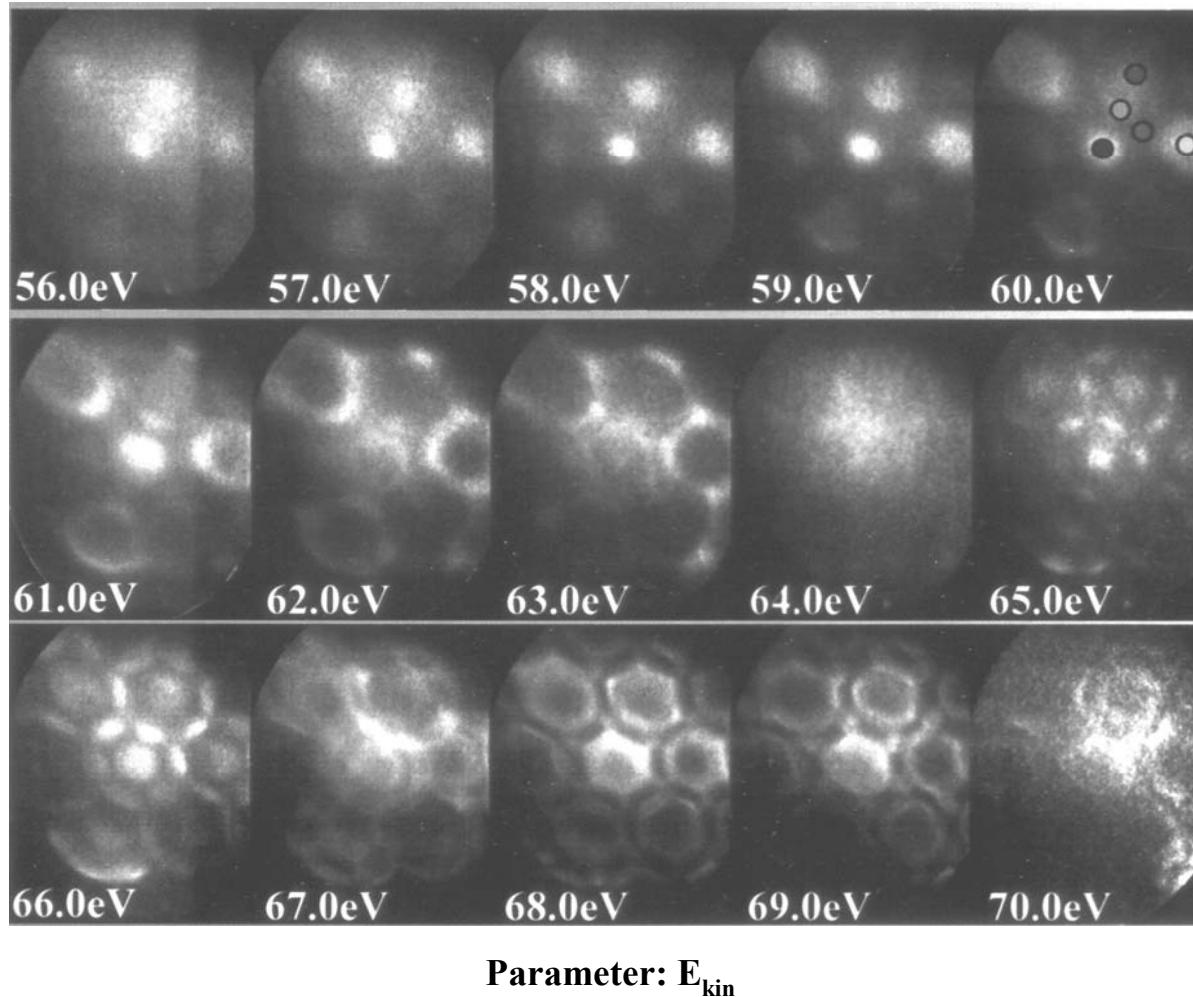
Simultaneously acquired PED and LEED pattern

Pb 5d 38 eV



Local band structure analysis (mode 2)

Conduction band of Pb(111)
5 Pb monolayers on Si(111) – Au $\sqrt{3} \times \sqrt{3}$ – R30°
 $h\nu = 73$ eV, 0.8 μm^2 area (selected field aperture)

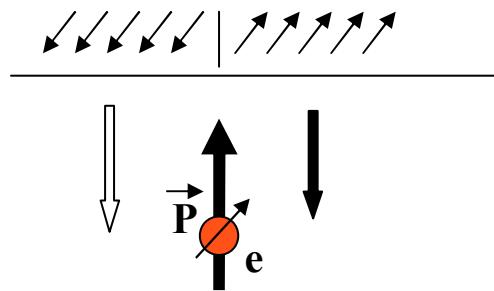


Magnetic imaging

XMCD, XMLD

Methods

SPLEEM

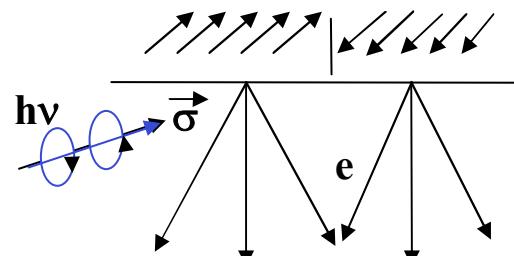


$$E_i = E_r = 0 - 20 \text{ eV}$$

$$I = I_{0r} + c \vec{P} \bullet \vec{M}$$

2 images with opposite \vec{P} or $\vec{\sigma}$: I_+ , I_-

XMCDPEEM



$$E_e = 0 - 20 \text{ eV (SE)}$$

$$I = I_{0e} + c \vec{\sigma} \bullet \vec{M}$$

$$I_+ - I_- = \text{magnetic image}$$

$$2 I_{0r}$$

structural image

$|\vec{P}|$ typical 20%

$$2 I_{0e}$$

chemical topographic image

$|\vec{\sigma}|$ typical 80%

$$(I_+ - I_-) / (I_+ + I_-) \text{ asymmetry image}$$

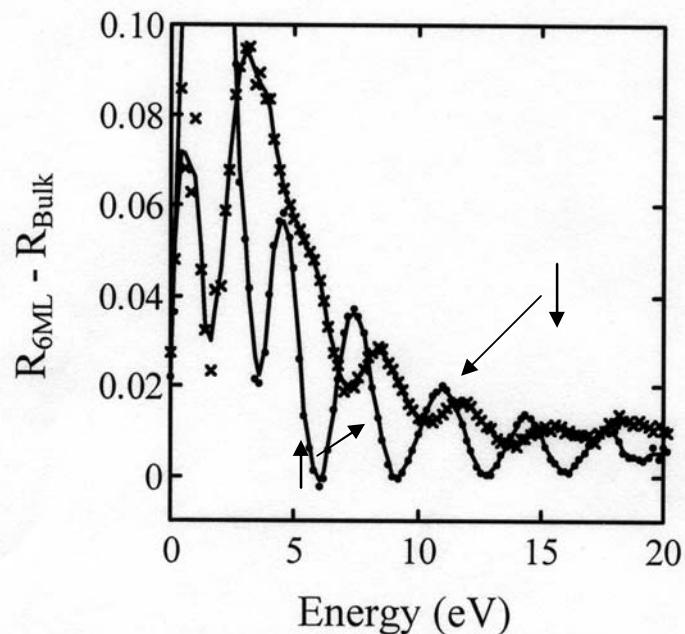
Basic mechanisms

SPLEEM

Spin-dependent scattering cross-section
due to exchange interaction

↓
Spin-dependent reflectivity

6 ML Fe on W(110)

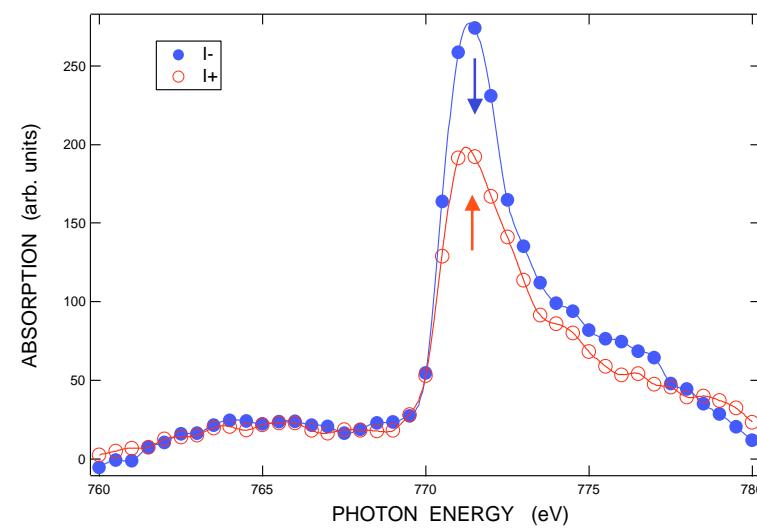


XMCDPEEM

Helicity-dependent transition probability
from 2p to unoccupied 3d ($\uparrow\downarrow$) states

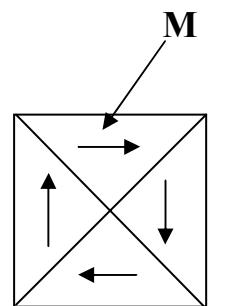
↓
Spin-dependent secondary electron emission

Secondary electron yield around Co 2p_{3/2} edge

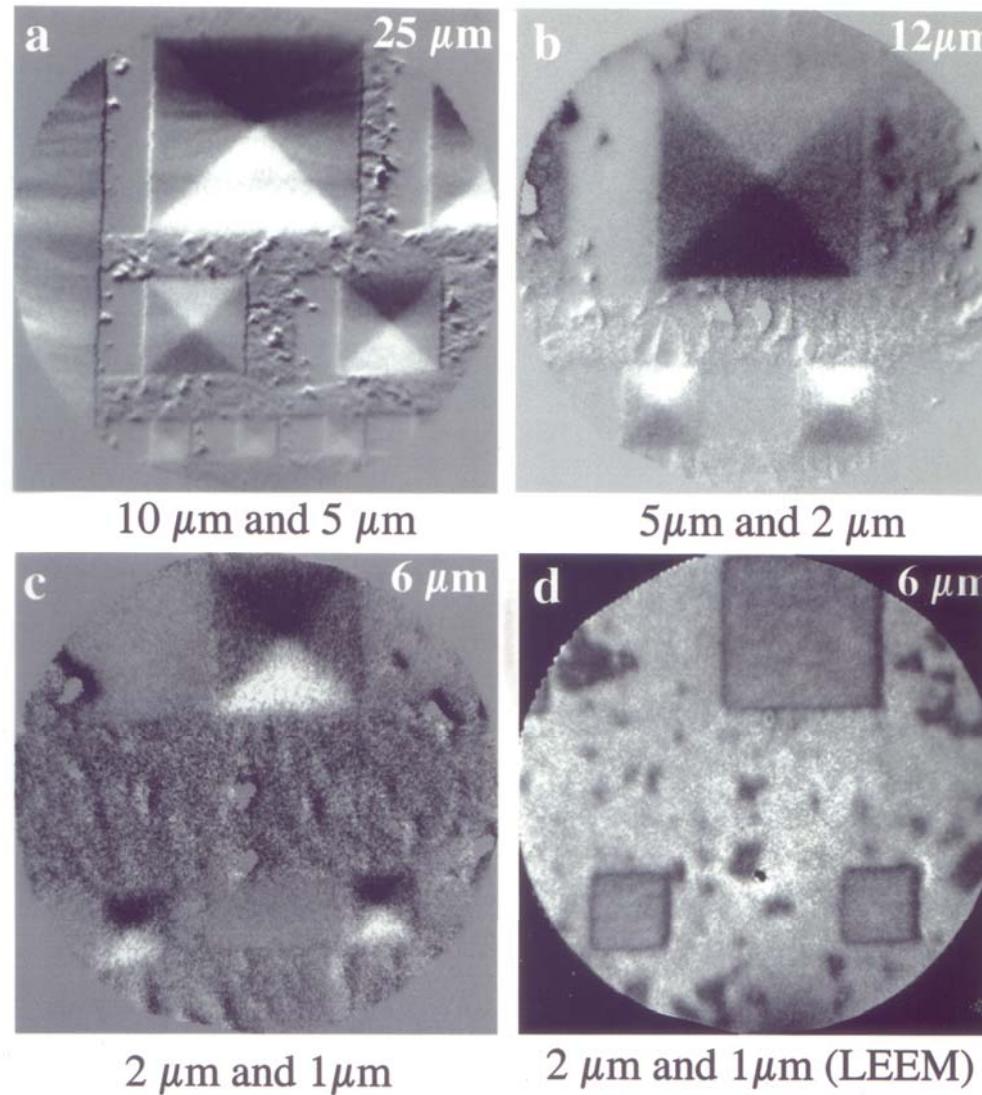
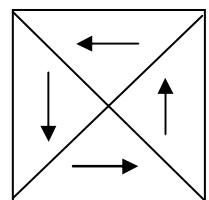


Closure domains in permalloy squares

Ion beam milled from permalloy film



$h\nu$
σ
Fe L₃ edge



Conclusion

Full field XPEEM is one of the most important applications of the high brilliance of third generation synchrotron light, in particular when combined with a **band pass energy filter** and with **LEEM** because it allows a complete characterization of surfaces and thin films, presently on the 10 nm lateral **resolution** scale, with aberration correction in the future on the 1 nm scale (hopefully).

The main benefit of **aberration correction**, however, will be the strong increase of the **transmission** of the system which will reduce image acquisition time considerably. This will allow dynamical studies, which are presently limited to LEEM, also with XPEEM.

General references

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