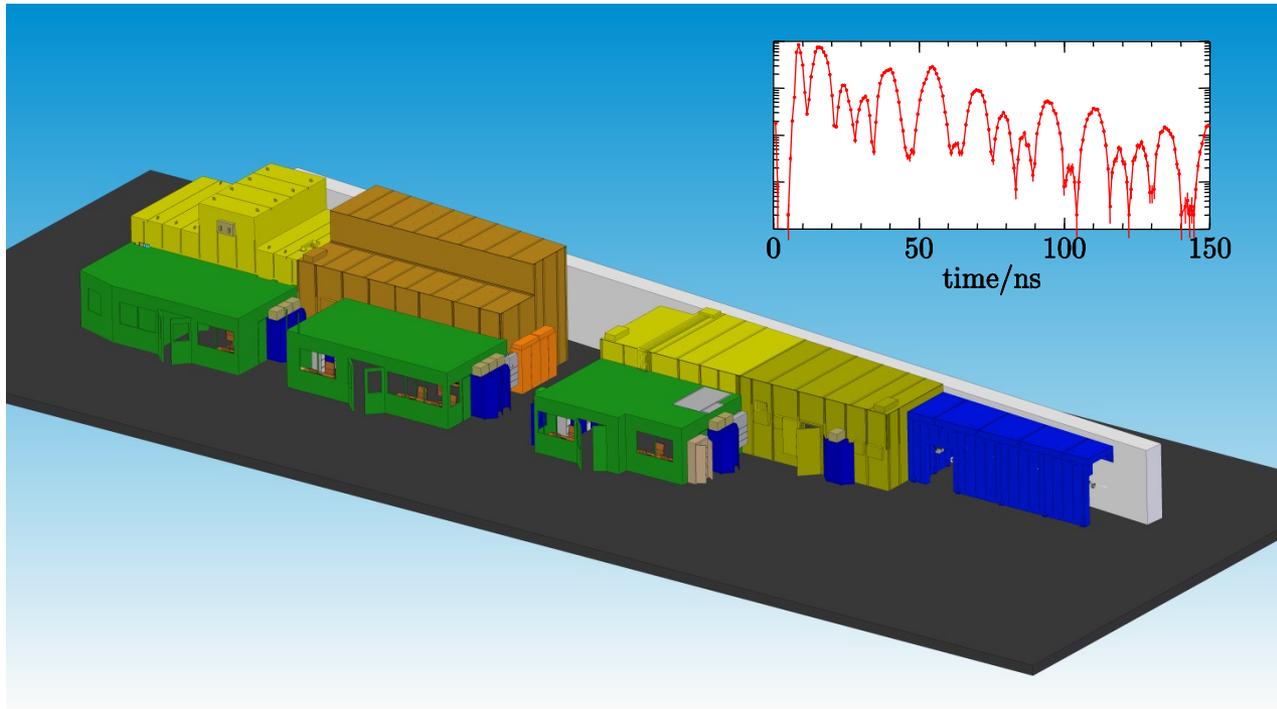


Timing experiments at the Dynamics beamline P01 of PETRA III

Why bunch purity is important for Nuclear Resonant Scattering



Hans-Christian Wille
Beschleuniger-Betriebsseminar

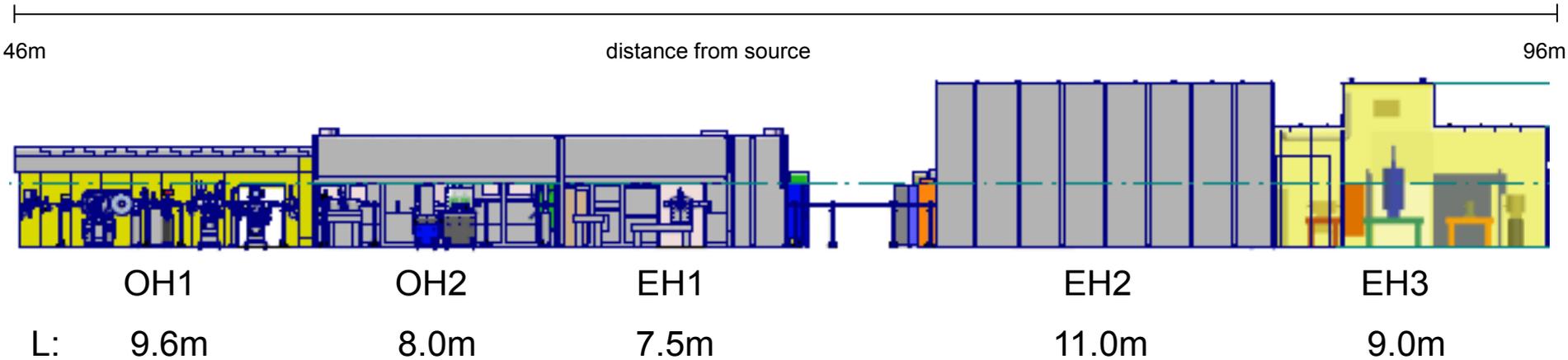
March 20 2013

Outline

- Beamline layout of P01
- Nuclear Resonant Scattering
- Some examples bunch purity / impurity
- Summary



P01 Beamline layout



OH1: High heat load optics (HFLM, mirrors, Be lenses)

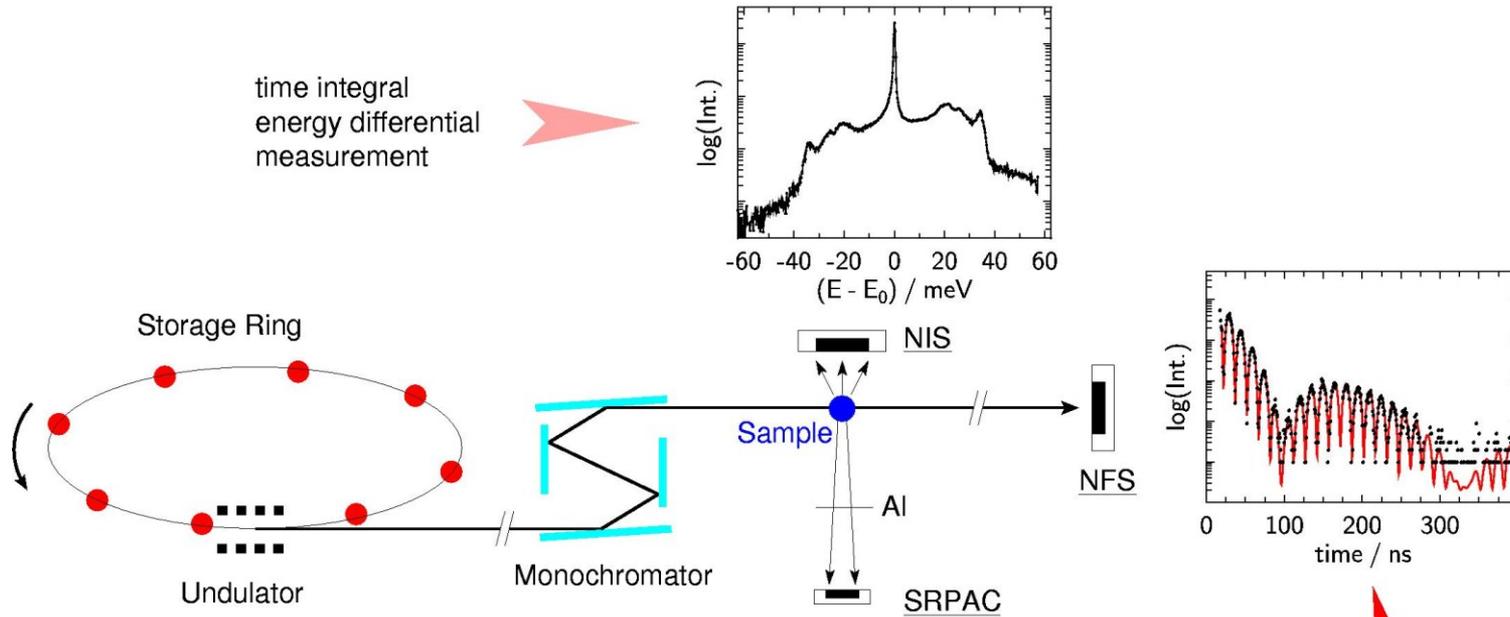
OH2: High resolution optics (HRMs)

EH1: Nuclear Resonant Scattering (raw focus, simple sample environment, θ - 2θ -setup)

EH2: IXS spectrometers (focused beam $1 \times 2 \mu\text{m}^2$, electronic excitations, RIXS ?)

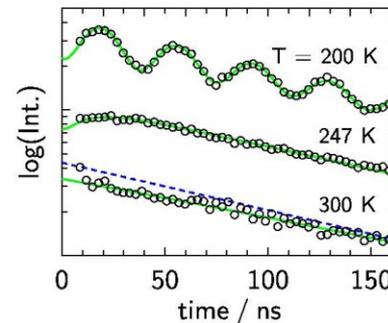
EH3: Nuclear Resonant Scattering (focused beam, UHV NFS / GISAXS in-situ)

Nuclear Resonant Scattering Techniques



Only nuclear resonant photons are counted. These are delayed in time with respect to the prompt electronic scattering due to the lifetime of the nuclear level

-> fast detectors / electronics, timing mode



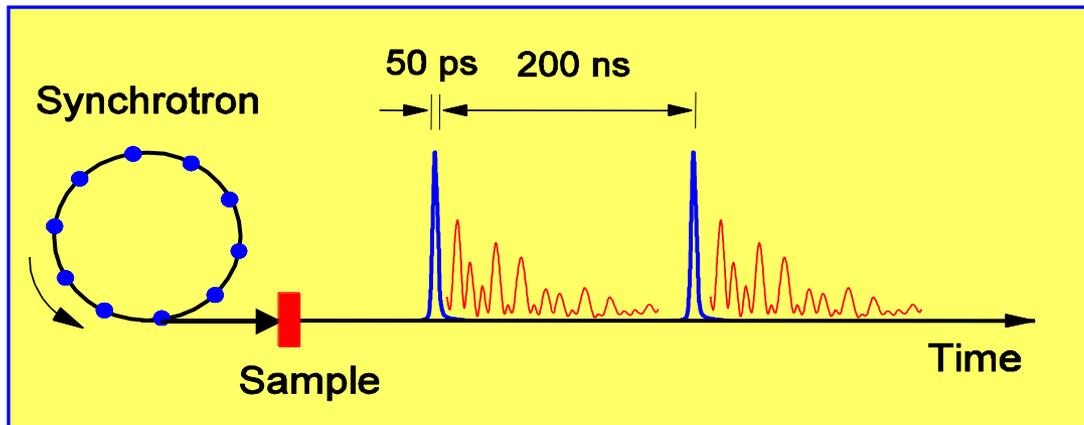
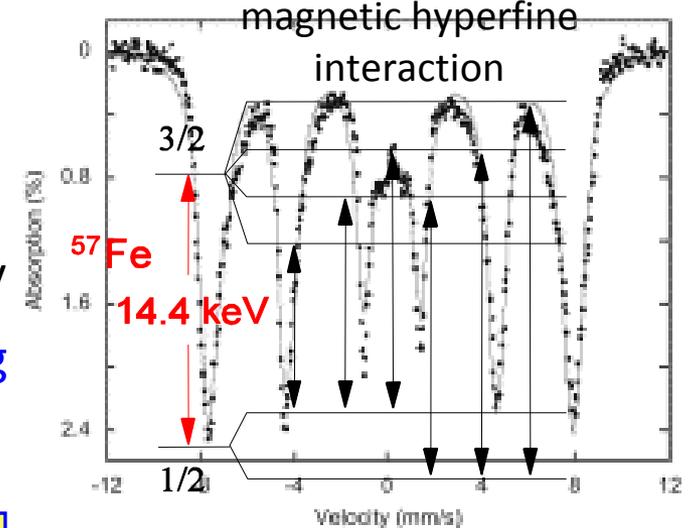
energy integral time differential measurements

Nuclear Forward Scattering (Mössbauer effect)

The Mössbauer effect:

Recoilless absorption and emission of photons

- In the energy domain: Mössbauer spectroscopy
- In the time domain : Nuclear forward scattering of synchrotron radiation



The 14.4 keV nuclear resonance of ^{57}Fe

$$\tau_0 = 141 \text{ ns}, \Gamma_0 = 4.7 \text{ neV}$$

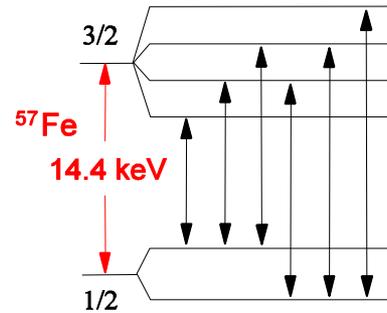
$$\Delta E/E = 3.3 \times 10^{-13}$$

Magnetic splitting: direction and value of \mathbf{B} at the nucleus, magnetic moments

Quadrupole splitting: electronic state, e.g. valence state

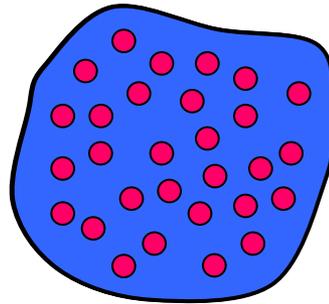
Nuclear Resonant Forward Scattering of Synchrotron Radiation

Pulsed broadband excitation of all hyperfine-split nuclear levels simultaneously



The 14.4 keV nuclear resonance of ^{57}Fe

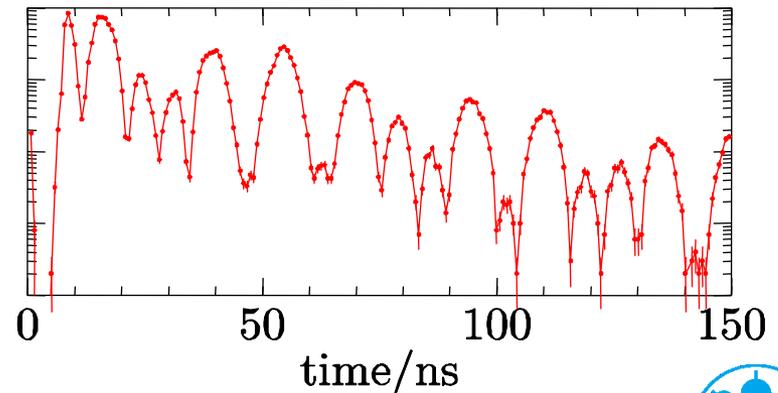
$$\tau_0 = 141 \text{ ns}, \Gamma_0 = 4.7 \text{ neV}$$



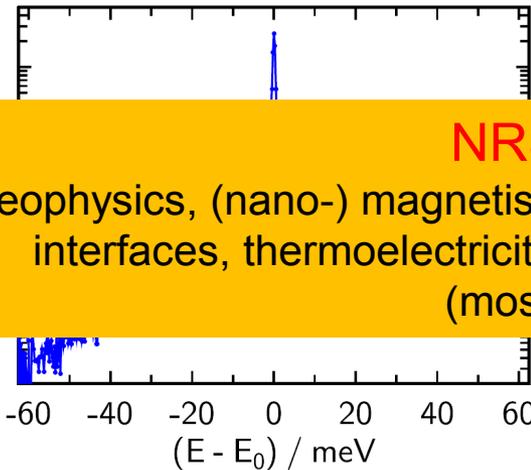
All levels are excited simultaneously and give rise to an interference beat pattern in time

The beat pattern is a fingerprint of the magnetic structure of the sample.

Temporal beats



Nuclear Inelastic Scattering (NIS)

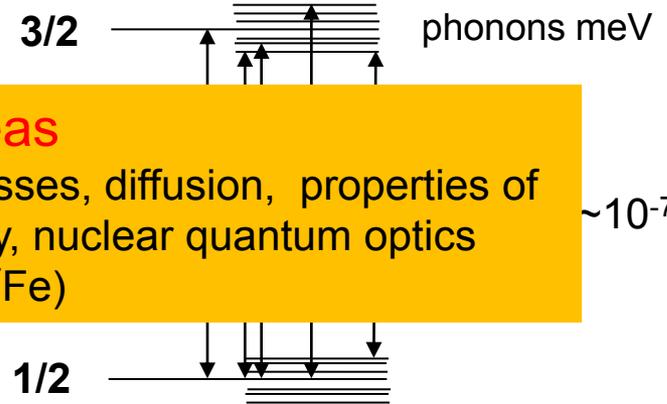


NIS spectrum of Fe at ambient T

NRS Research areas

geophysics, (nano-) magnetism, dynamics in glasses, diffusion, properties of interfaces, thermoelectricity, molecular biology, nuclear quantum optics
(most of it covered by ^{57}Fe)

$\sim 10^{-7}$

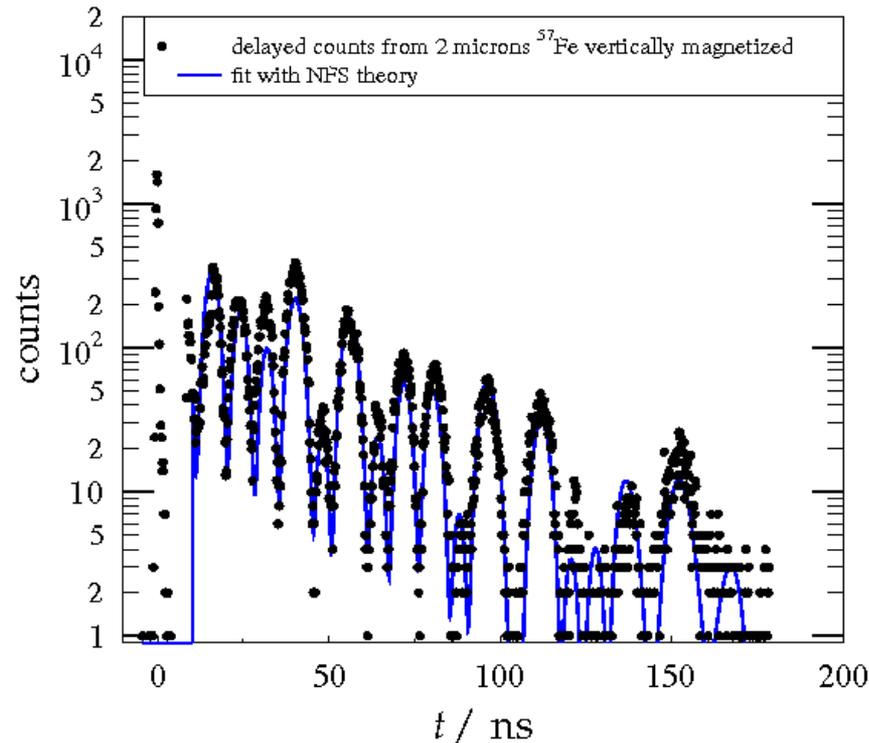


Nuclear inelastic scattering is a tool to study vibrational dynamics.

Accomplishes other techniques like inelastic neutron scattering and is unique:

- ❖ Direct extraction of element specific VDOS (no background, no models)
- ❖ Element specific dynamics in complex systems, probe atoms, probe layers...
- ❖ Small sample size, nano structured materials

Nuclear Forward Scattering of enriched iron ^{57}Fe

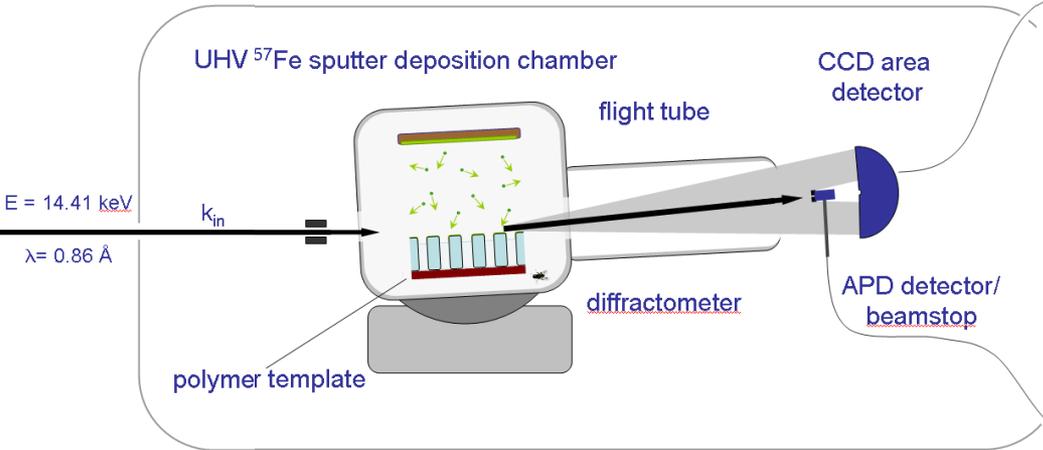


2 microns ^{57}Fe , 40 bunch mode 192 ns bunch spacing, 2m U23 undulator
Can be done in a few ms today...

In-situ Synchrotron Radiation 3D Microscopy

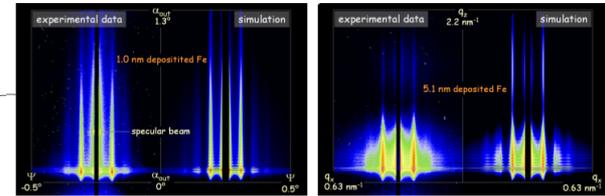
By courtesy of Kai Schlage:

CCD for diffuse scattered x-rays: GISAXS
structural correlations (vertical and lateral),
particle geometry

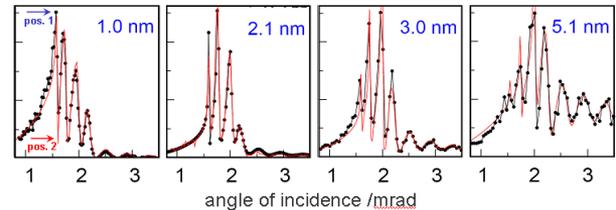


APD for specular reflection: NFS, NRXR
chemical and magnetic sensitivity to ^{57}Fe ,
magnetization dynamics up to/close to GHz regime,
high resolution iron depth-profile

Grazing incidence small angle scattering – surface morphology



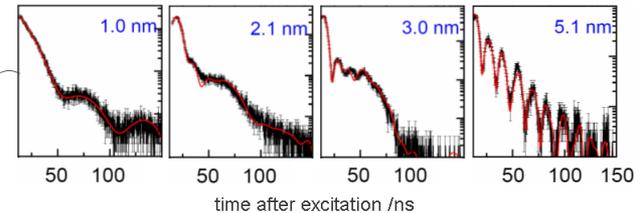
Nuclear resonant reflectometry – high resolution iron depth profile



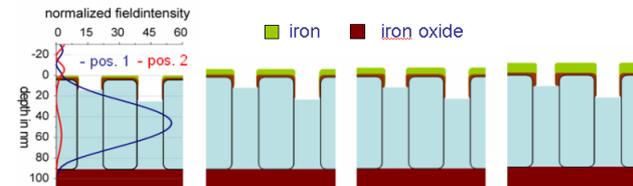
nuclear (delayed) intensity

stepwise iron sputter deposition

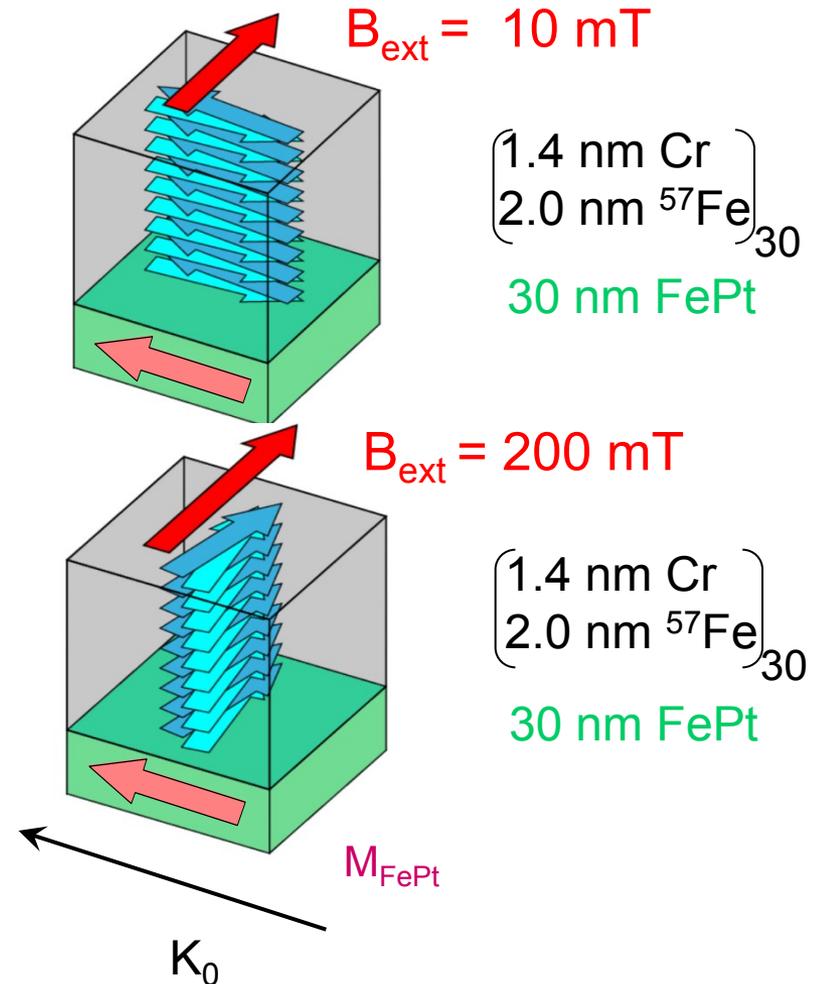
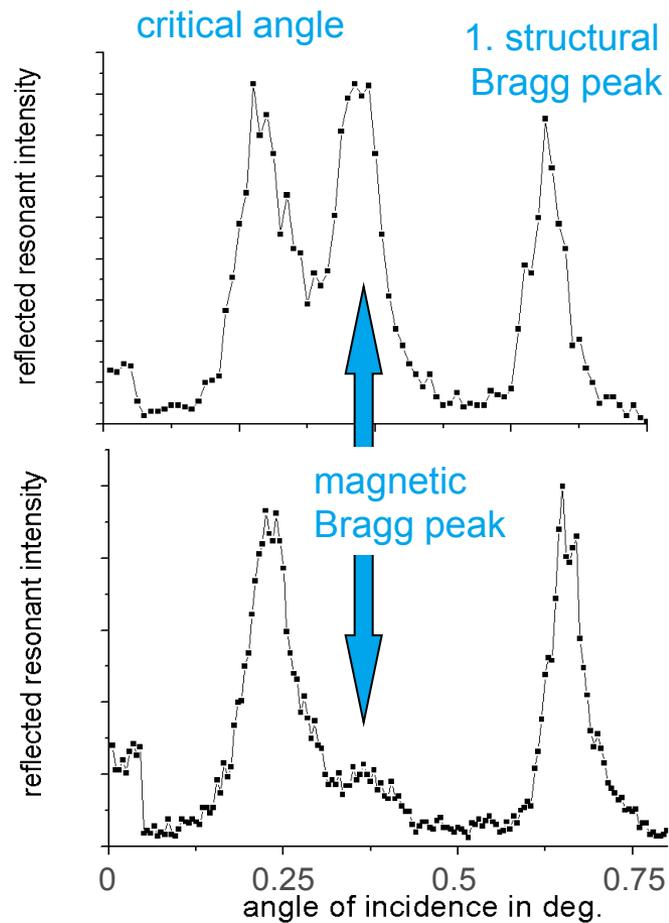
Nuclear time dependent scattering – chemical, magnetic state of iron



nuclear de-excitation (cts)



Resonant Reflectometry of Antiferromagnetic Multilayers



Intensity of the magnetic Bragg peak as tool for characterization of magnetic reorientation phenomena

T. Guryeva, K. Schlage

Raw estimate of count rates

Natural line width $5 \cdot 10^{-9}$ eV

We get 10^{10} photons per second in 1 meV

$$10^{13} \text{ photons / s / eV} \cdot 5 \cdot 10^{-9} \text{ eV}$$

$$= 5 \cdot 10^4 \text{ photons / s}$$

In the condition 1 photon per nucleus

*6 by the number of transitions and other factors < 1 like the Lamb Mössbauer factor, Natural abundance of ^{57}Fe (~2%)

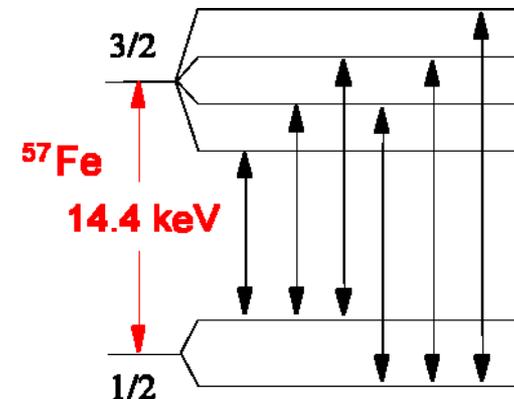
The most reducing factors is small cross section due to the sample size / thickness (nano science) and the sample environment (high P, low T, Fe embedded in optical thick material etc.)

Typical count rates are 0.1 to 100 Hz
One strength of the method: it is (usually) background free @ bunch purity of 10^{-10}

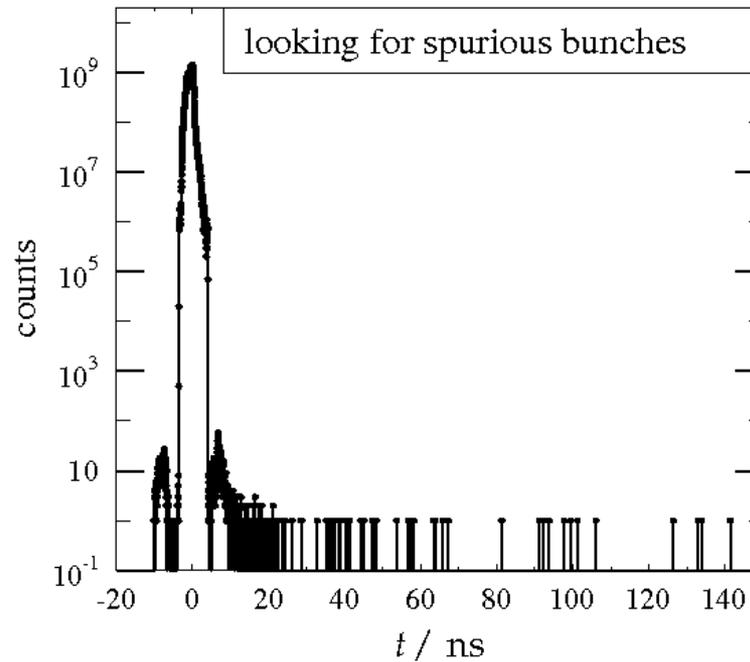
The 14.4 keV nuclear resonance of ^{57}Fe

$$\tau_0 = 141 \text{ ns}, \Gamma_0 = 4.7 \text{ neV}$$

$$\Delta E/E = 3.3 \cdot 10^{-13}$$



PETRA III bunch purity in 40 bunch mode (60 mA with top up 2011)

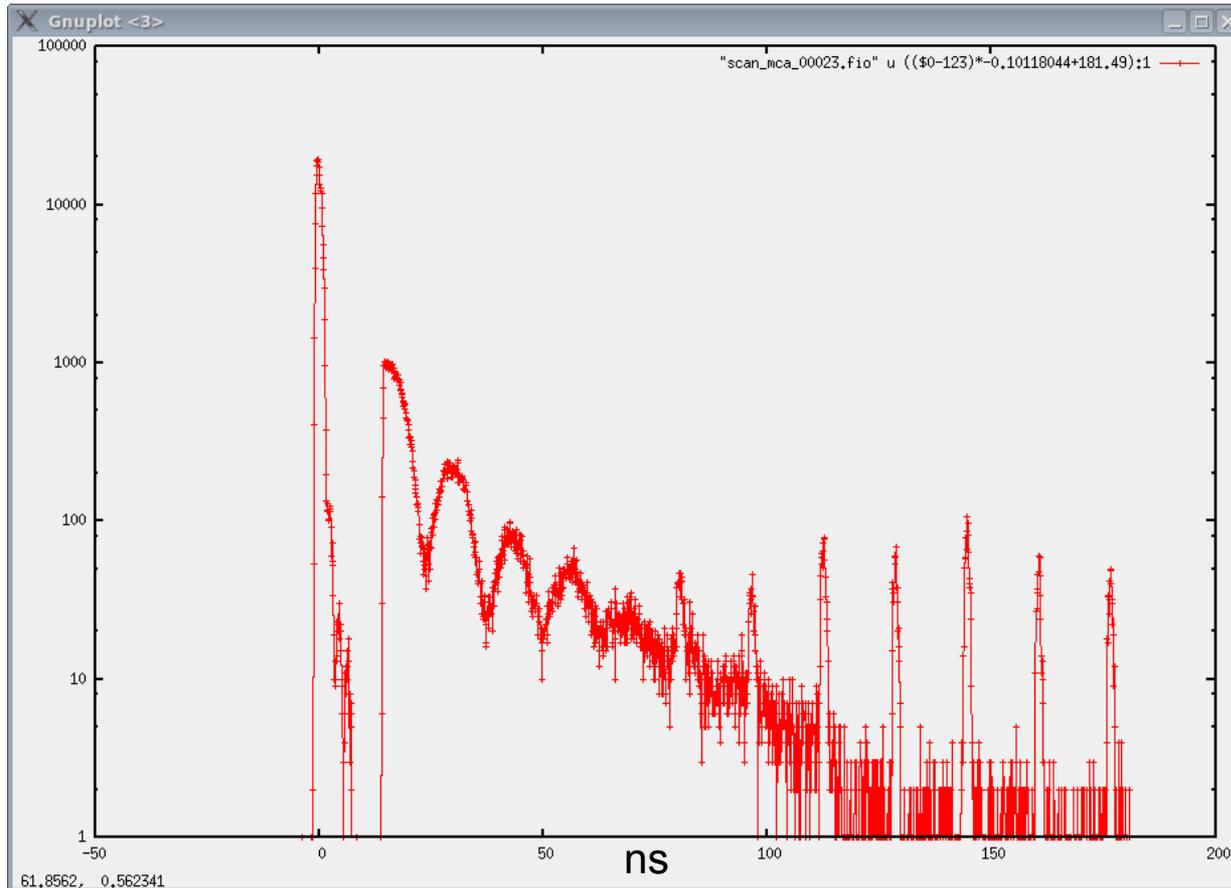


Spurious bunches at $\pm 8\text{ns}$ with 10^{-7} intensity

APD noise up to 20ns ?

Clean from 20 ns up

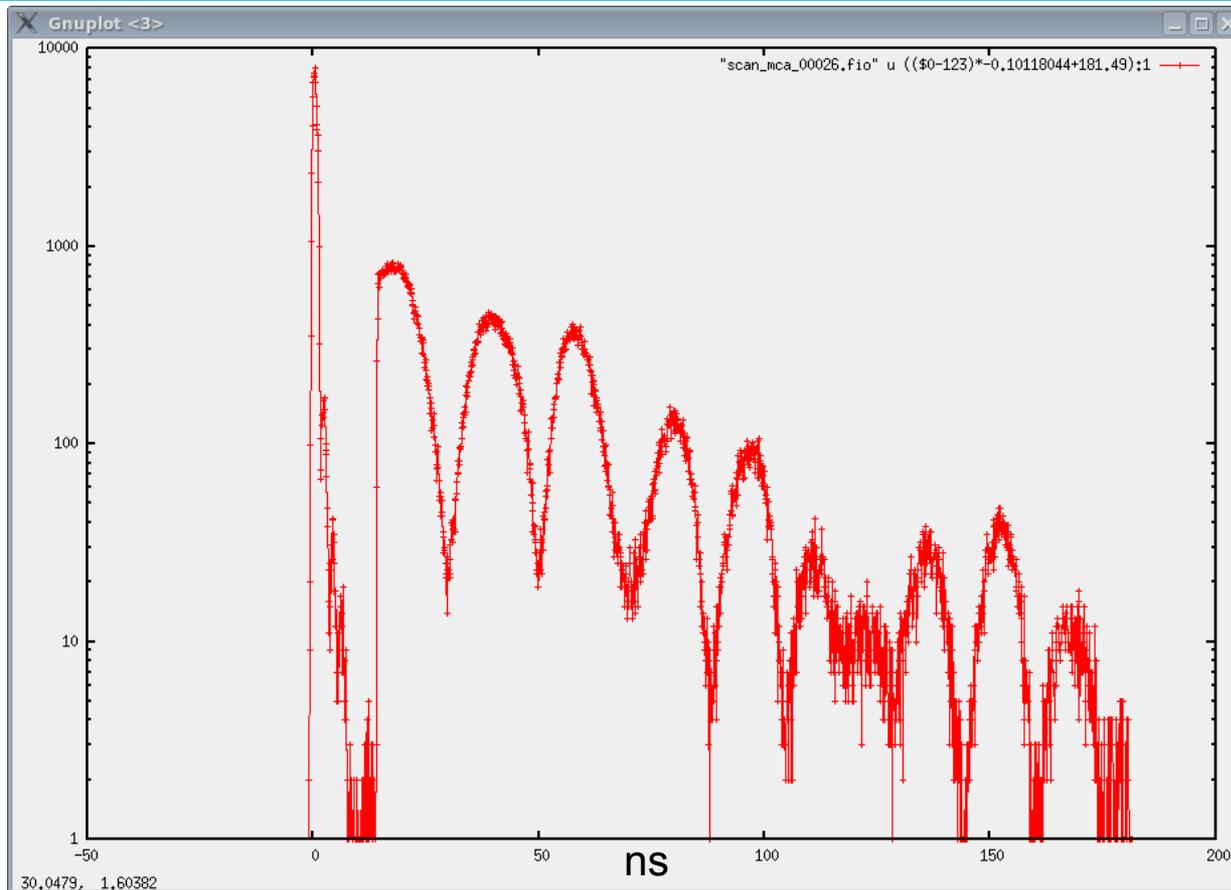
Spurious bunch problem 2013



2 h sample time, sp. bunch rate < 5 Hz, regular with 16ns distance
prompt at $t = 0$ ns measured only for fraction of a second
detector noise < 0.1 Hz



Spurious bunch problem

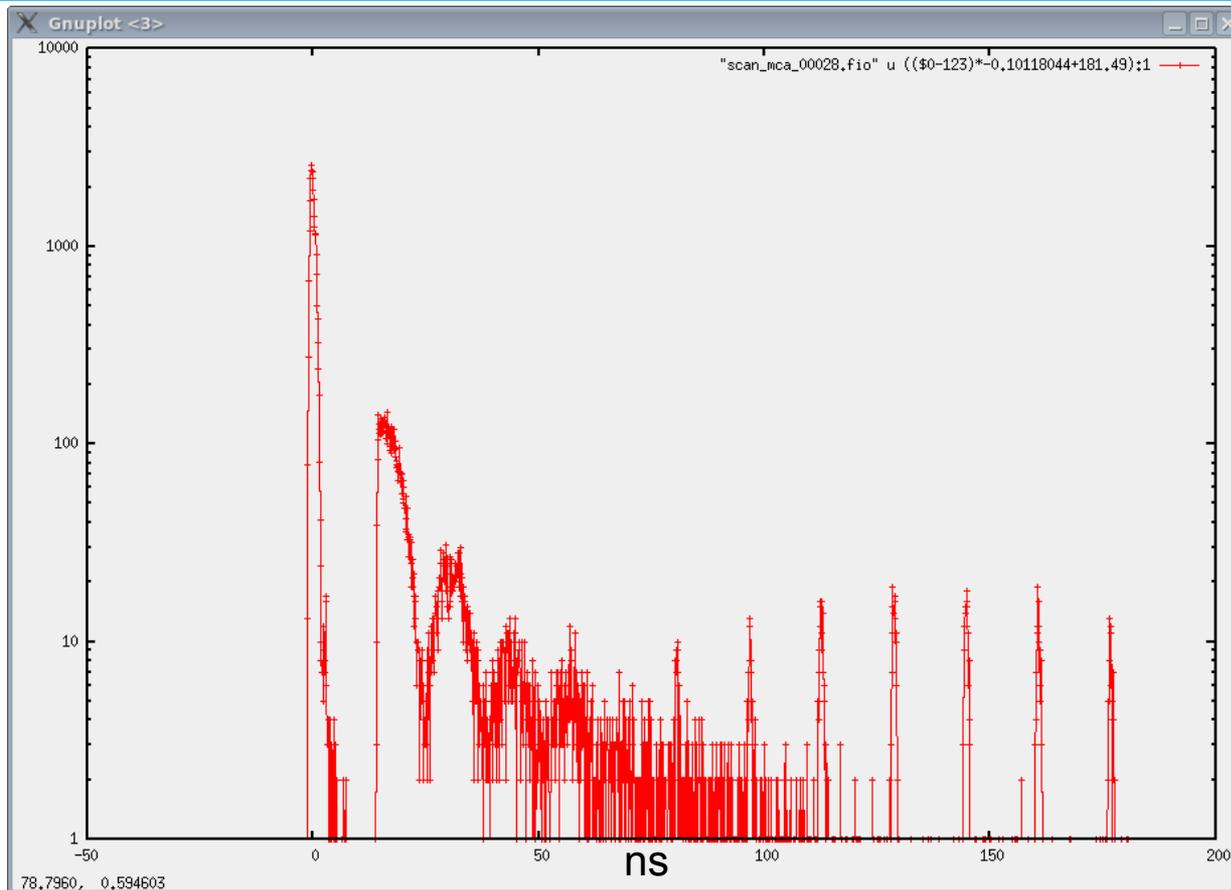


600 s sample time, sp. bunch rate < 5 Hz

detector noise < 0.1 Hz



Spurious bunch problem

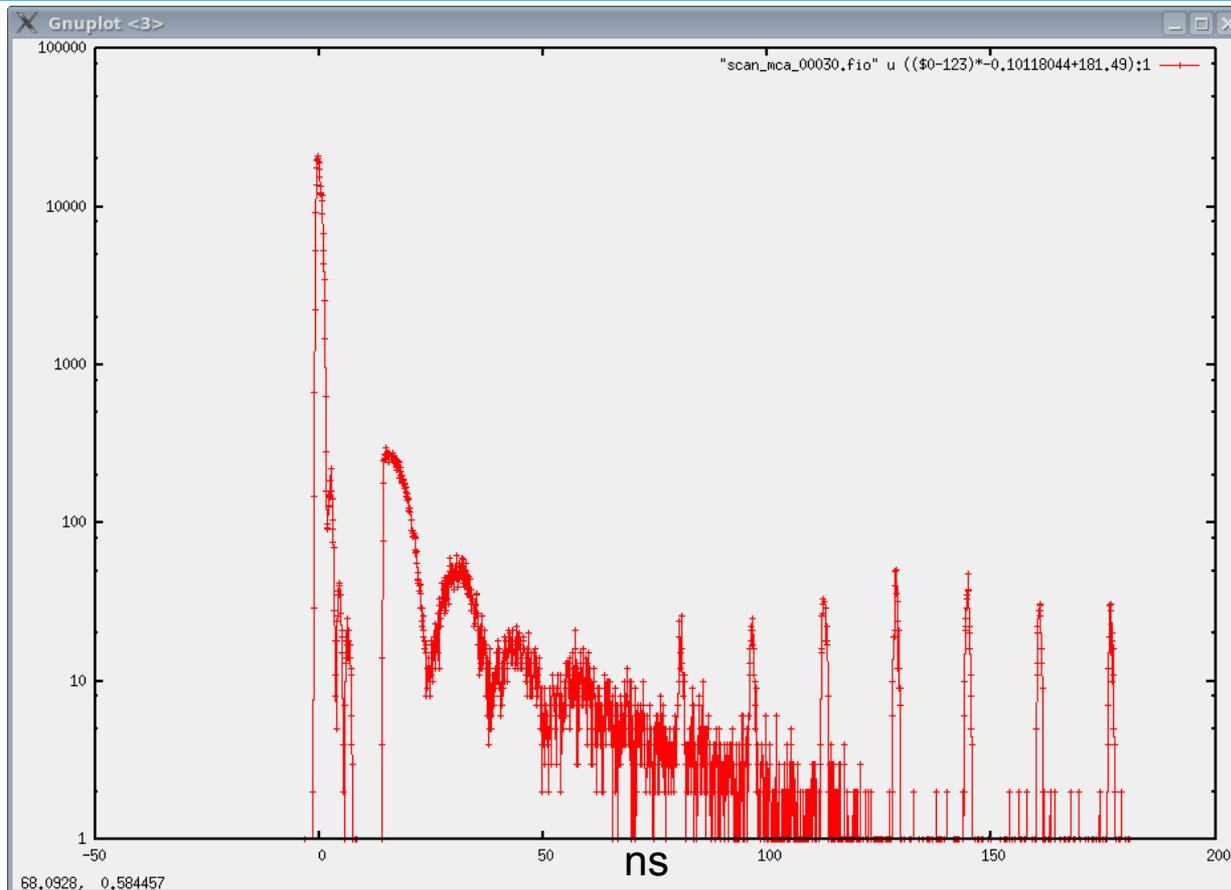


1000s sample time, sp. bunch rate < 5 Hz

detector noise < 0.1 Hz



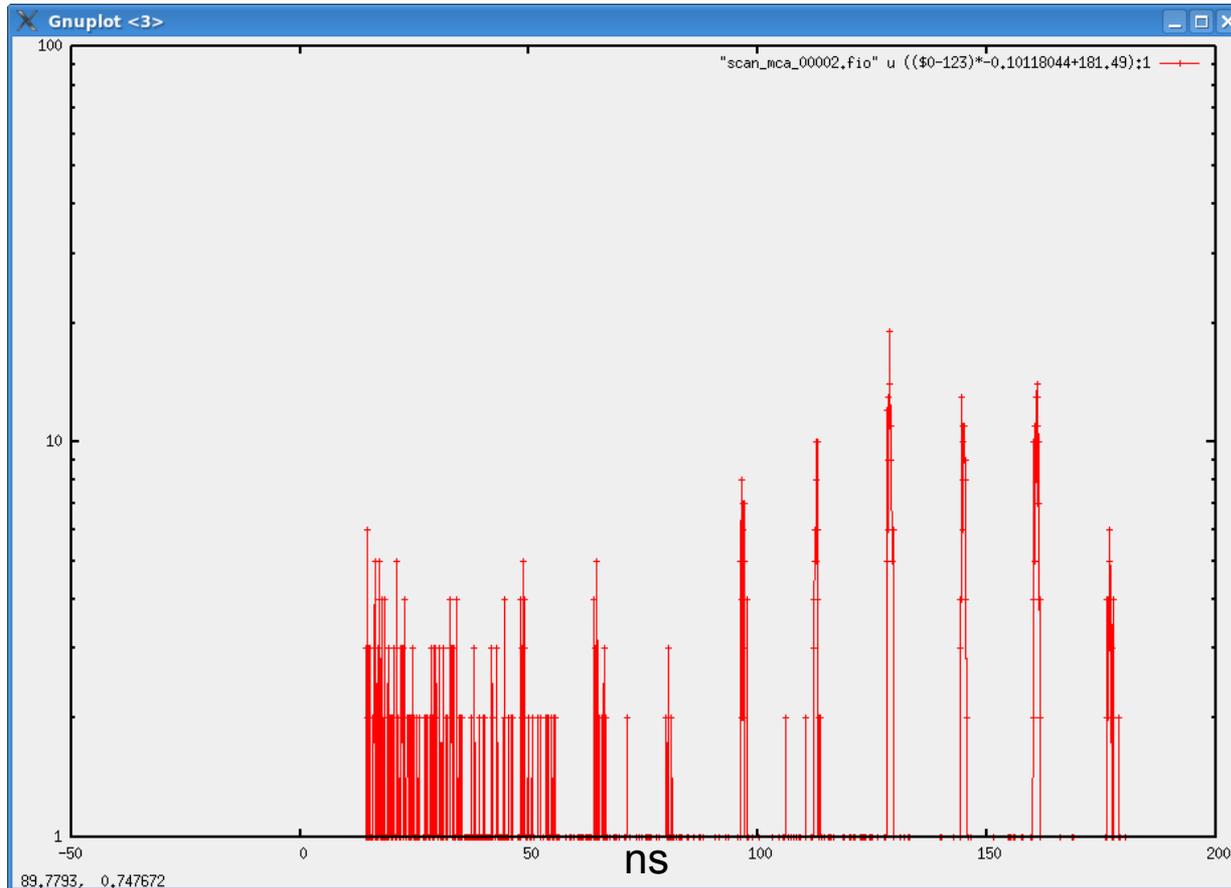
Spurious bunch problem



2700 s sample time, sp. bunch rate < 5 Hz

detector noise < 0.1 Hz

Spurious bunch problem



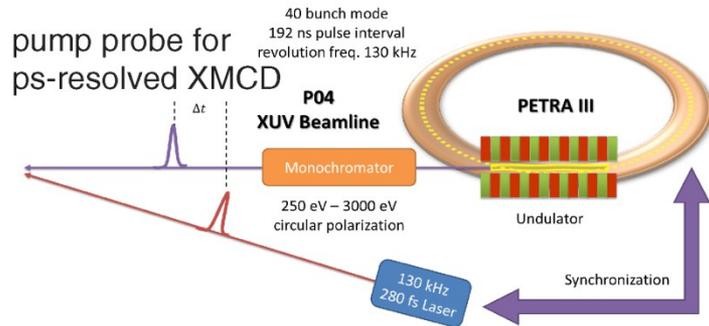
1000 s sample time, sp. bunch rate < 5 Hz

detector noise < 0.1 Hz

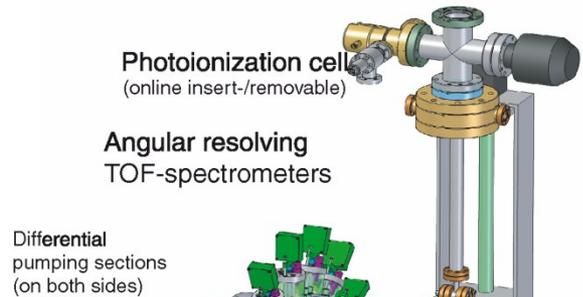


Timing Experiments at the XUV Beamline P04

> Time resolved XMCD-Imaging

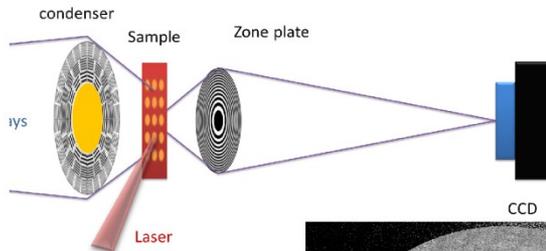


> P04 beamline diagnostics and in-house research

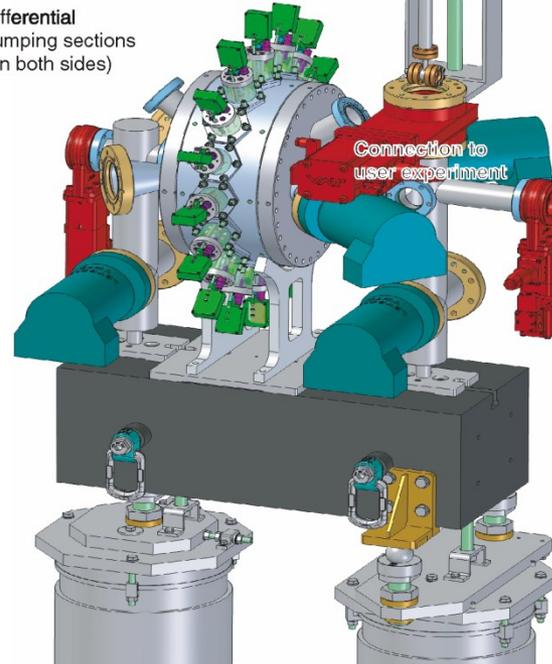
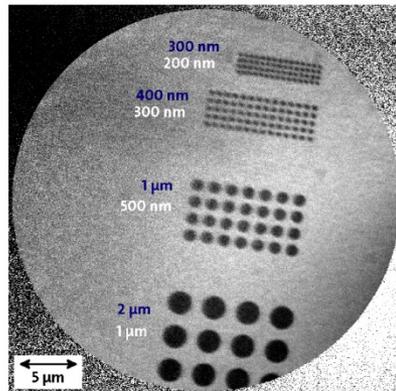


Electron time-of-flight system:

- Photon flux (accuracy $\leq 1\%$ abs.)
- Beam position (accuracy $\leq 1 \mu\text{m}$)
calibration possible with "BPMs"
- Photon energy ($\leq \Delta E$, $\leq 1/10,000$)
- Degree of polarization (accuracy $\leq 1\%$)
user warning if S_i below
a user-defined threshold.



Flatfield corrected
TXM image of
nanostructured
Co/Pt ML-system



> One third of P04 users
requests timing mode

> Beamlines using timing mode: P01, P02, P04, P06, P08, P10, P11, (P24)

Summary

About half of the PETRA III beamlines are performing timing experiments

For most of them a decent bunch purity of 10^{-5} is sufficient

NRS experiments rely on a purity of at least 10^{-9}

(This is not new and had been established until summer 2012... lets go for it again !!)



MANY THANKS TO...

Hasan Yavas

Alessandro Kropmanns

Hermann Franz

Tatyana Guryeva

Olof Gutowski

Hanns-Peter Liermann + P02 team

Michael Sprung + P10 team

FS-BT

FS-EC

FS-TI

Safety + Machine Group

...

Kai Schlage

Alexander Scholl

Ralf Röhlsberger

Denise Erb

Anita Ehnes

Frank-Uwe Dill

Yilmaz ‚Jimmy‘ Bican

Wolfgang Caliebe

Balaram Sahoo

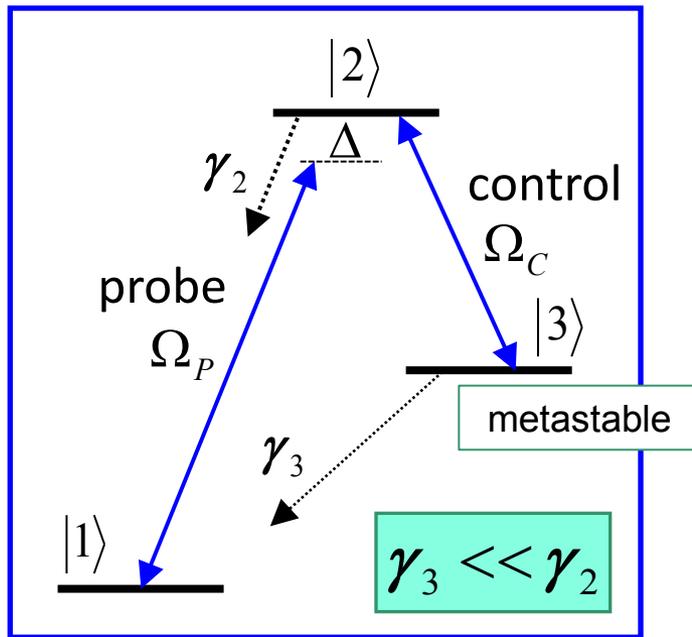
Daniel Schumacher



!! THANK YOU FOR YOUR ATTENTION !!



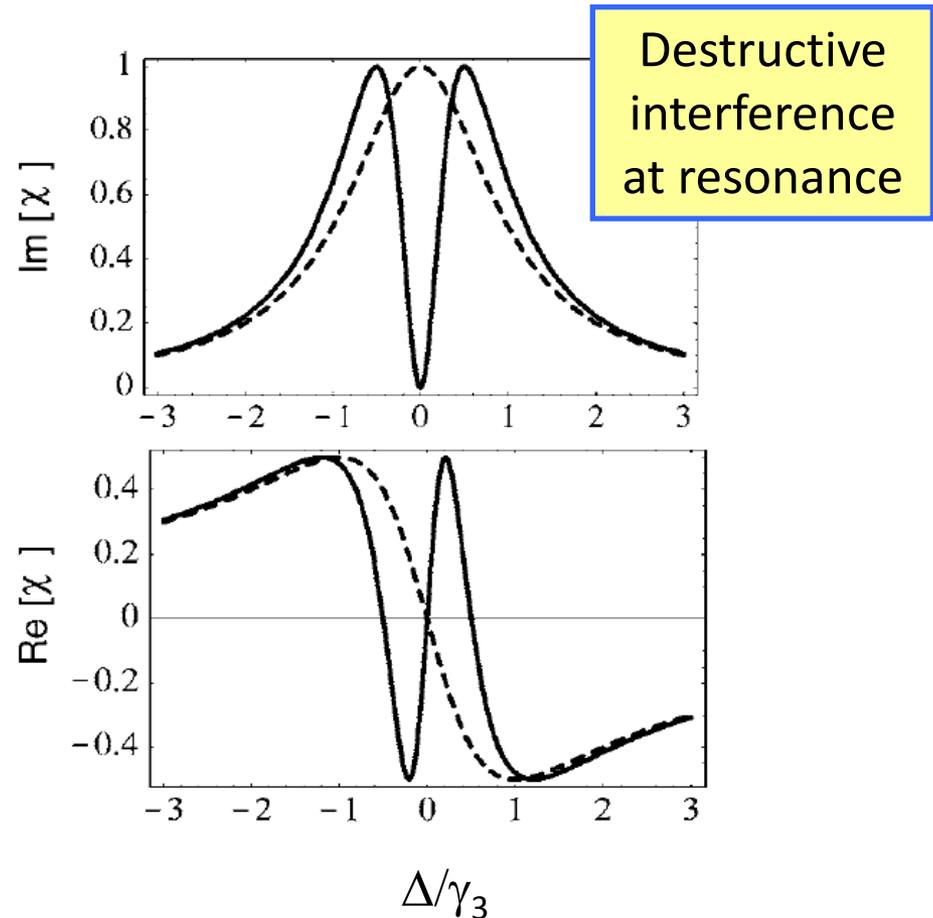
Electromagnetically Induced Transparency (EIT)



1st - order susceptibility

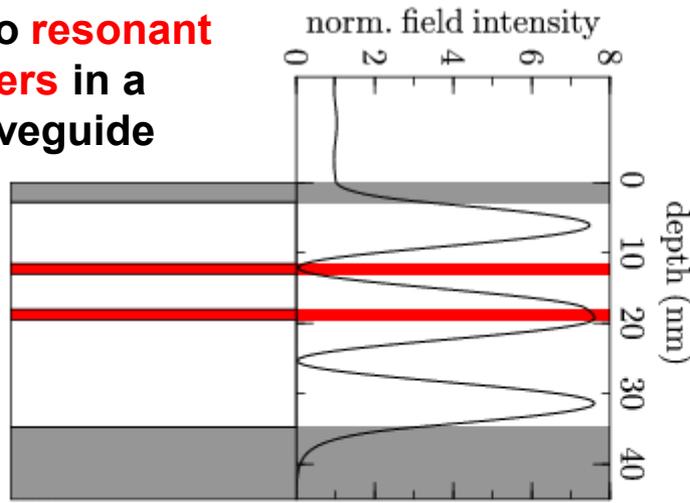
$$\chi(\Delta) = \frac{iP_{12}(i\Delta + \gamma_3)}{(i\Delta + \gamma_3)(i\Delta + \gamma_2) + |\Omega_C|^2}$$

M. Fleischhauer, A. Imamoglu, J. P. Marangos,
Rev. Mod. Phys. 77, 633 (2005)

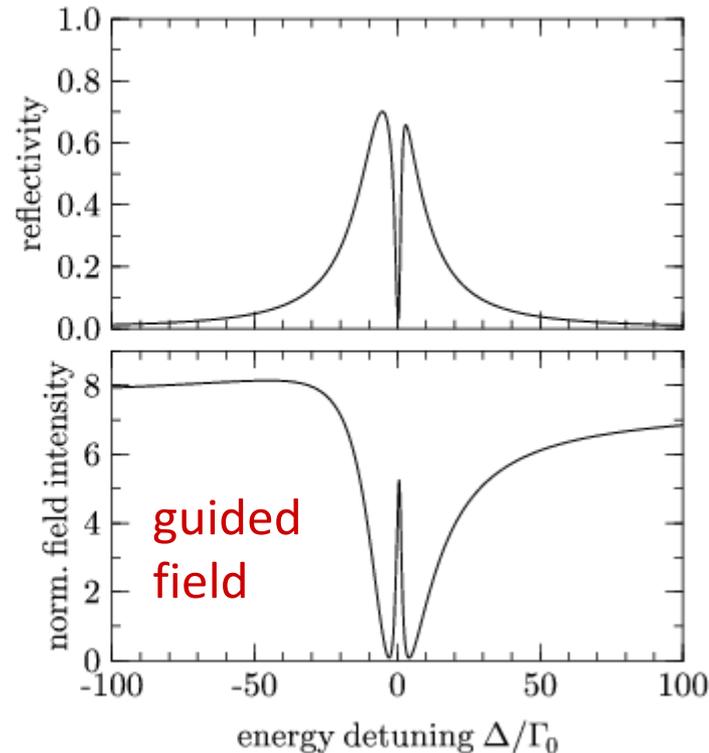


Electromagnetically Induced Transparency

Two resonant layers in a waveguide



reflectivity



The ^{57}Fe layers occupy node (natural decay, metastable) and antinode (superradiant, fast) of the field

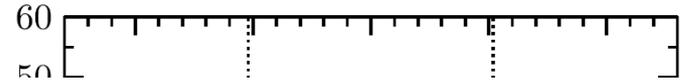
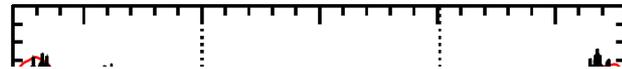
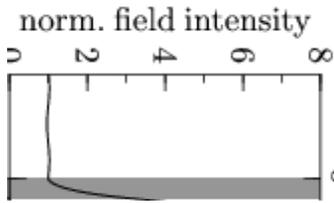
Electromagnetically Induced Transparency (EIT)

O. Kocharovskaya, Ya. I. Khanin, Sov. Phys. JETP 63, 945 (1986)

K. J. Boller, A. Imamoglu, S. E. Harris, Phys. Rev. Lett 66, 2593 (1991).

Nuclear Resonant EIT in a Cavity: Experiment

Experiment at P01 (R. Röhlsberger et al., May 2011)
(Four lines of nuclear resonance due to magnetic hyperfine splitting)

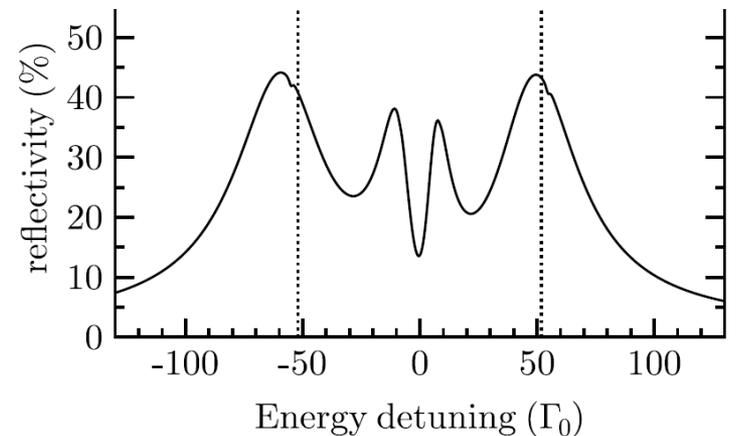
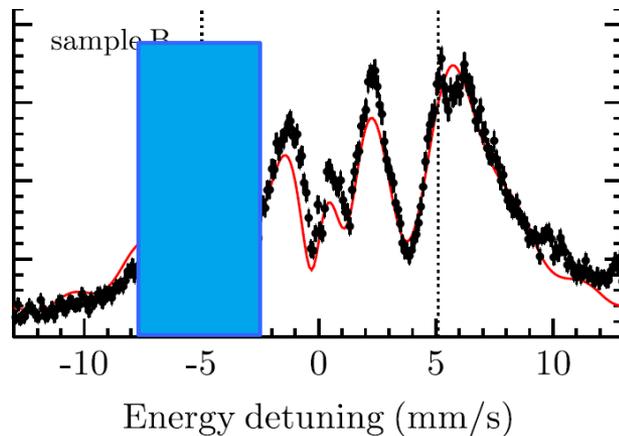
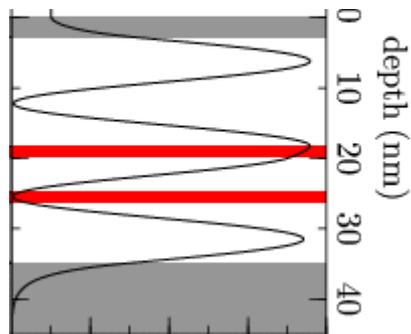


LETTER

doi:10.1038/nature10741

Electromagnetically induced transparency with resonant nuclei in a cavity

Ralf Röhlsberger¹, Hans-Christian Wille¹, Kai Schlage¹ & Balaram Sahoo¹



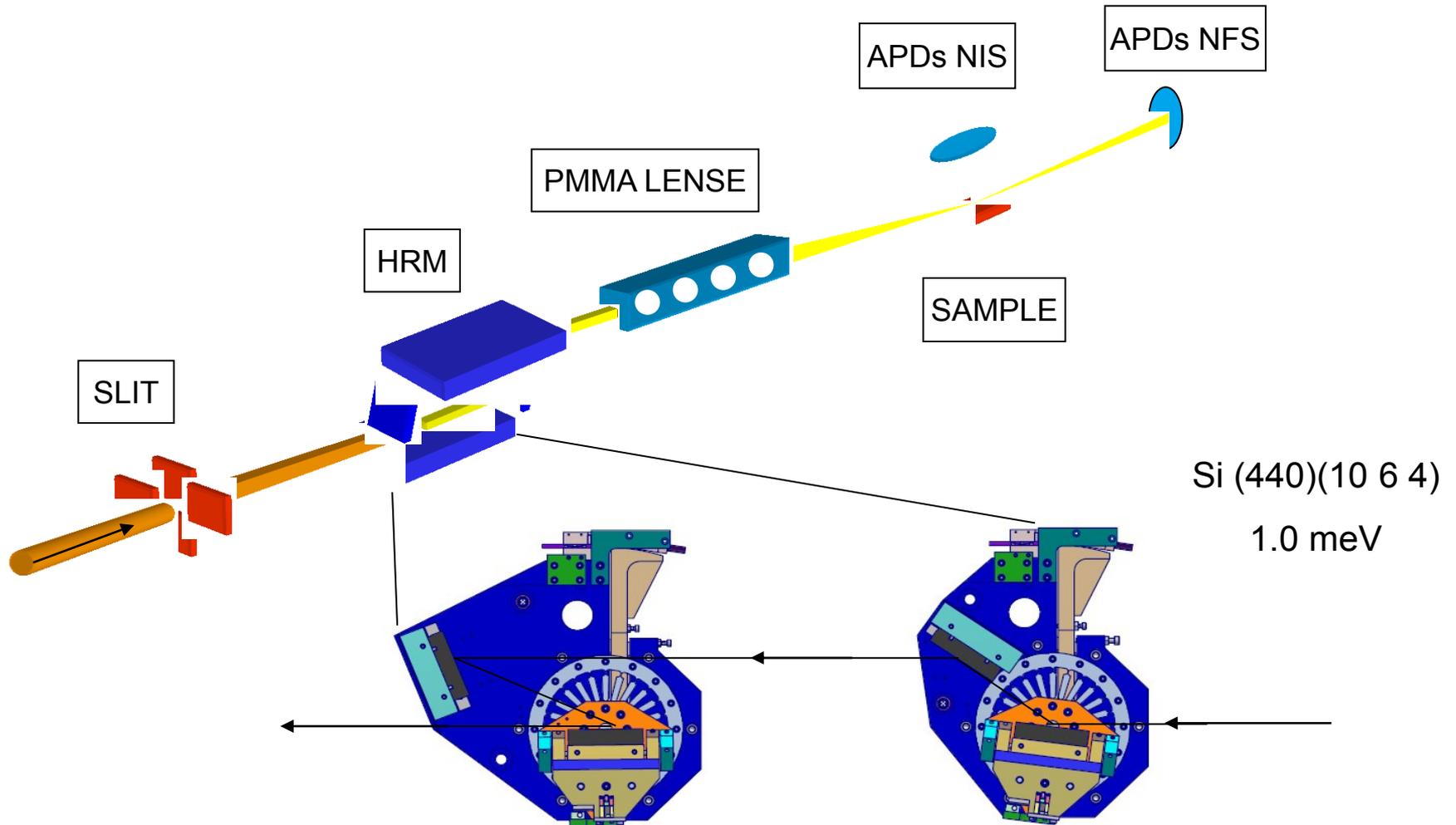
Time stamps...what happened so far @ P01

- 17.05.2010 First beam from 2m U23 (P02) in OH1
- 11.06.2010 First ^{57}Fe resonant counts
- 27.06.2010 First time spectrum from 2 microns thick ^{57}Fe foil (6 meV HR Monochromator)
- 15.07.2010 2.4meV HRM installed
- 25.07.2010 First Nuclear Inelastic Scattering (NIS) spectrum
Shutdown from 25.07.2010 until 20.08.2010
- 20.08.2010 3 weeks of timing mode
- 20.09. – no timing mode U23 back to P02
- 11.11.2010 Commissioning of the 5m U32
- 03.12.2010 3 weeks of timing mode

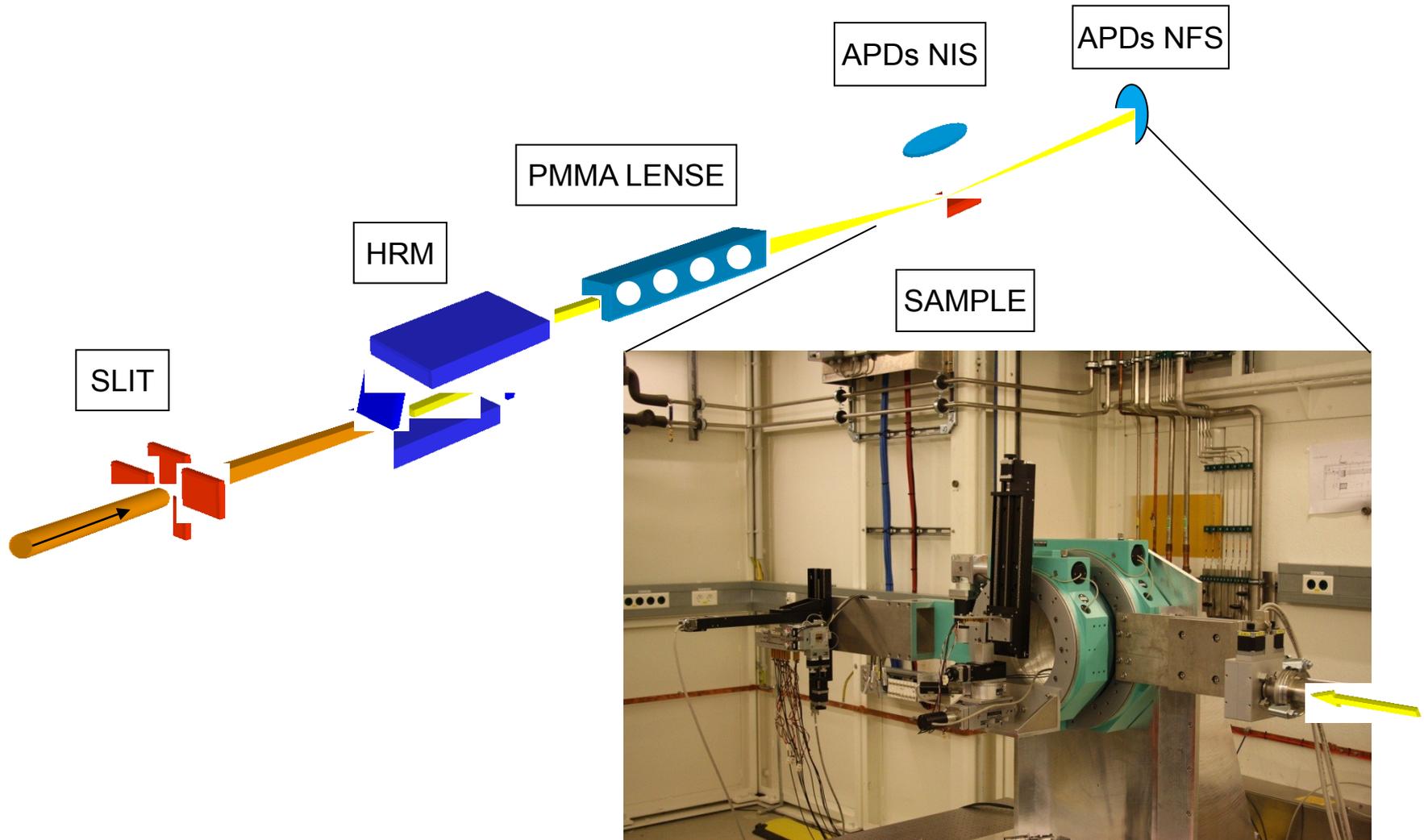
In total 6 weeks of timing mode beamtime since August 2010



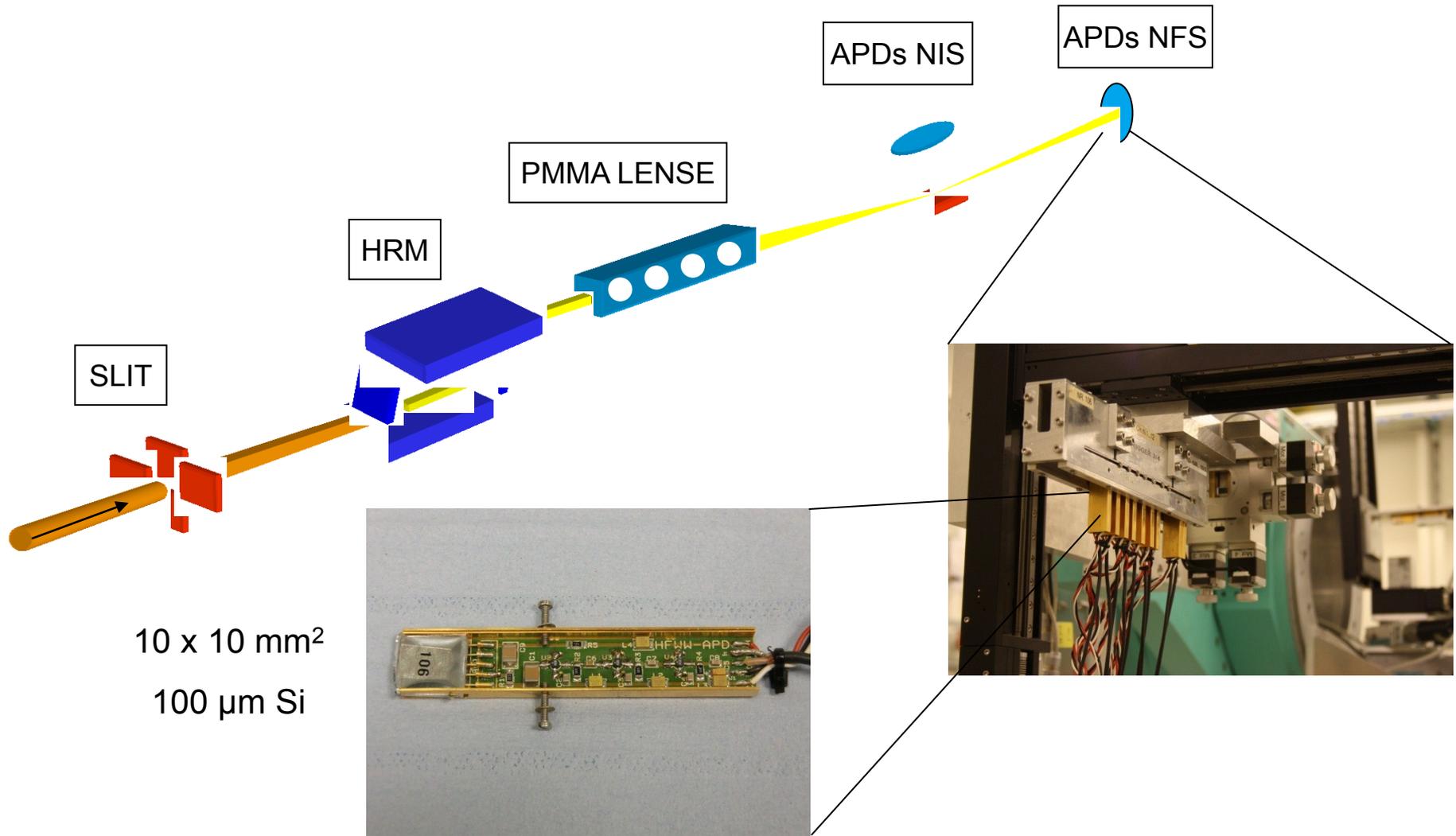
Setup θ - 2θ with focused beam (EH1)



Setup θ - 2θ with focused beam (EH1)



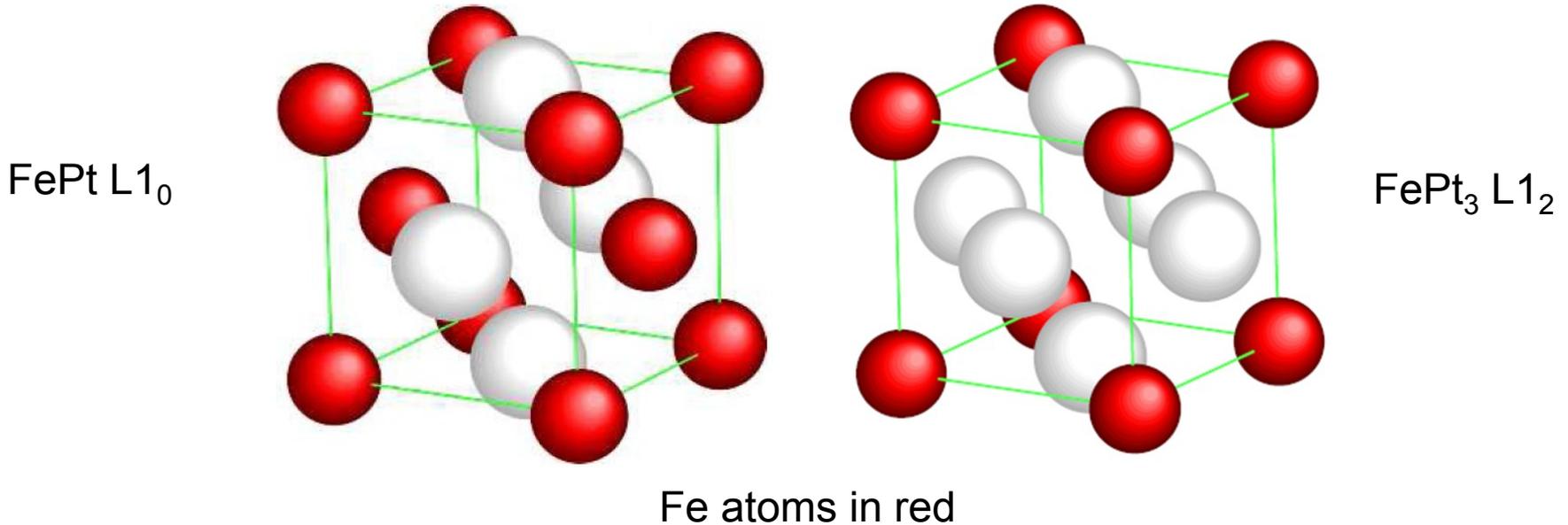
Setup θ - 2θ with focused beam (EH1)



Site specific phonon density of states in $L1_x$ FePt $_x$ alloys (S. Couet)

FePt is a magnetic material with a huge magneto-crystalline anisotropy (good hard magnet)

⇒ This originates from the crystal anisotropy

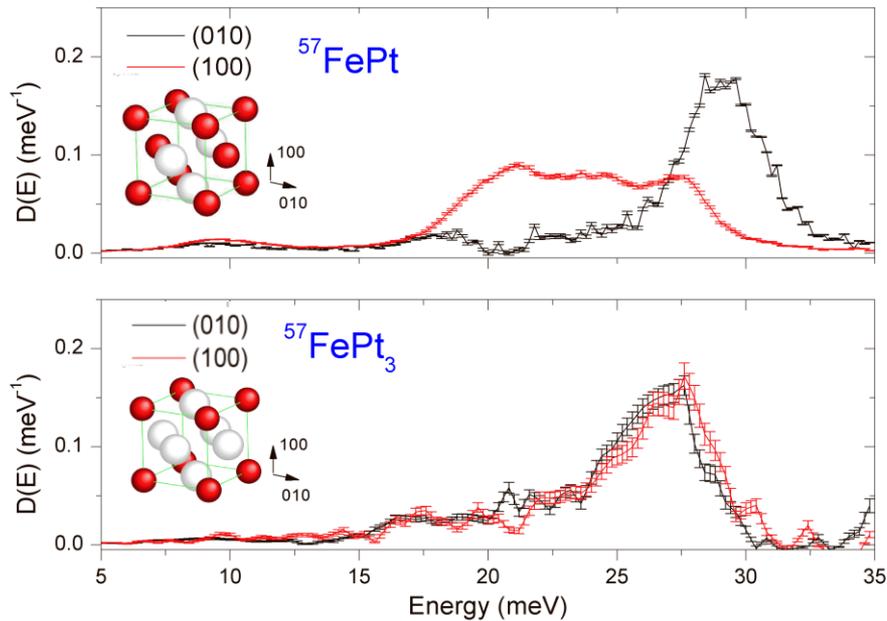


- How the anisotropy translate on the vibration spectrum?
 - Which atoms contribute to which part of the PDOS (non equivalent atom position)?
- ⇒ Compare the PDOS of FePt and FePt $_3$

Site specific phonon density of states in $L1_x FePt_x$ alloys

Measurement carried out on epitaxial $FePt$ and $FePt_3$ thin films (30 nm) grown on $MgO(100)$ and $MgO(110)$ substrates

Measurement on $^{57}FePt$ carried out at ID18, ESRF¹



- $FePt(010)$ PDOS close to $FePt_3$
- Peak shift induced by different strain in the two layers
- No major modes belonging to the “face-centered” Fe atom?

¹S. Couet et al., Phys. Rev. B 82, 094109 (2010)

Size dependent PDOS of ^{57}Fe islands on MgO (S. Couet)

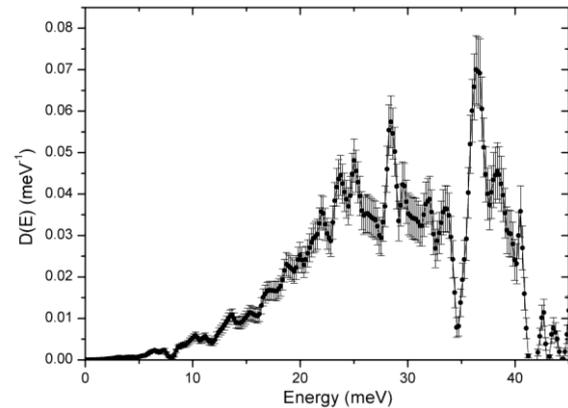
Tuning of the island size by control of the nominal thickness of Fe deposited at 700°C

5 nm

2 nm

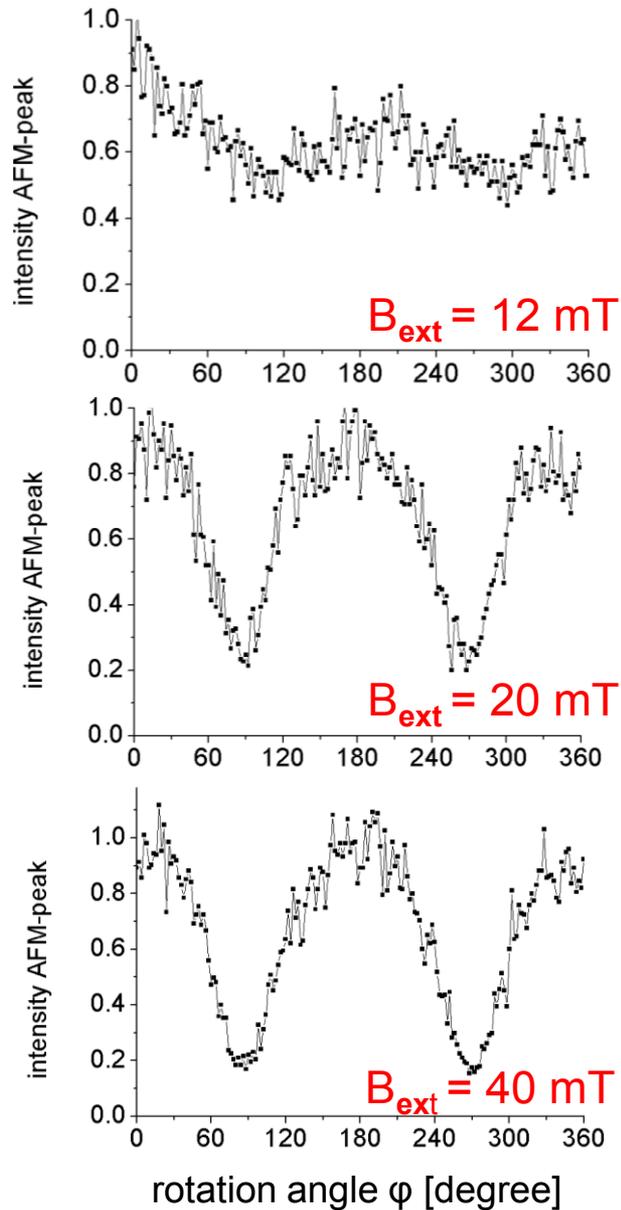
1 nm

⇒ Search for phonon confinement/quantum effects in nanostructures

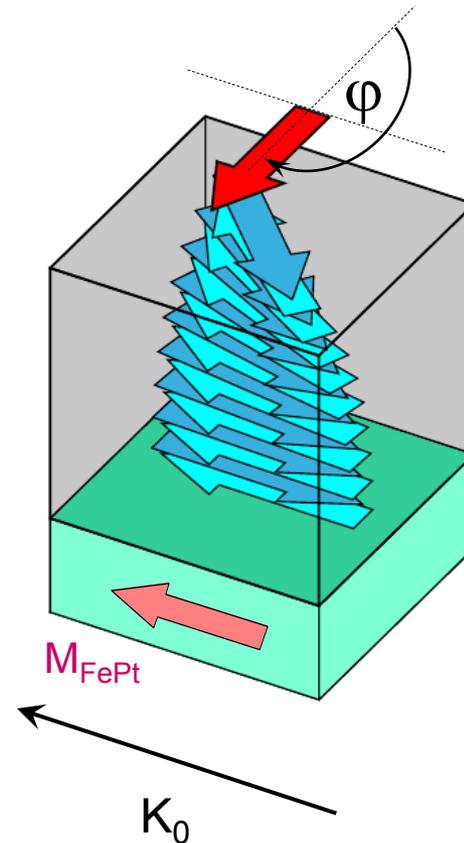


Still to be measured

Resonant Reflectometry on Magnetically Coupled Thin Films



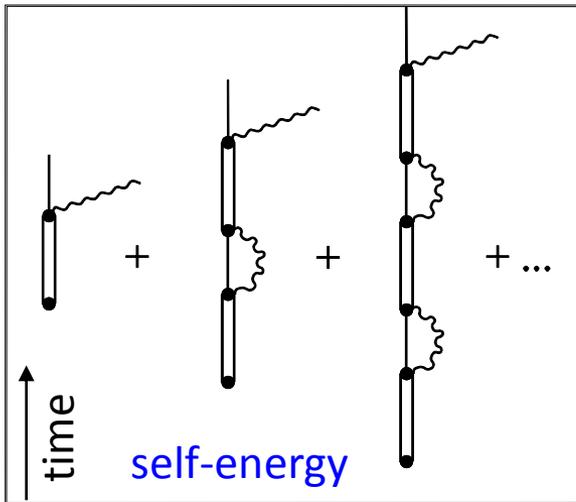
azimuthal rotation of the external field:



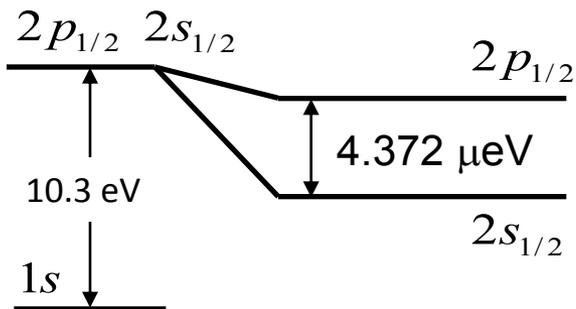
When Nanoscience meets Quantum Optics (R. Röhlsberger)

The Collective Lamb Shift

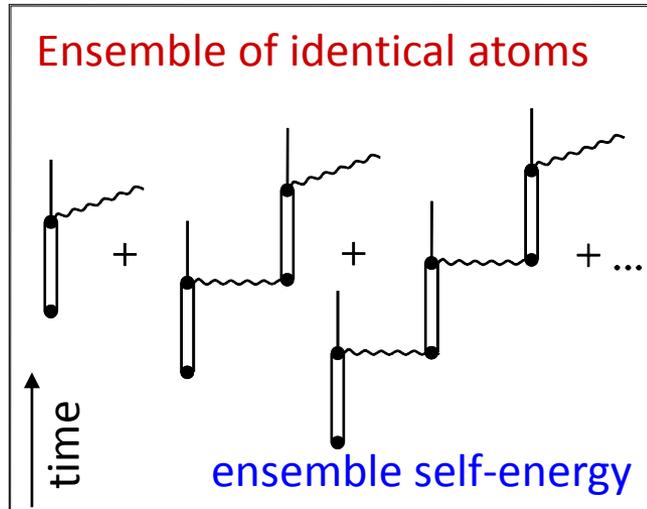
Single-atom Lamb shift



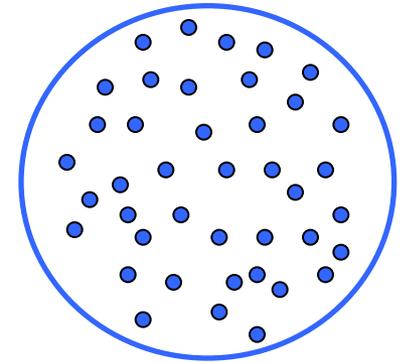
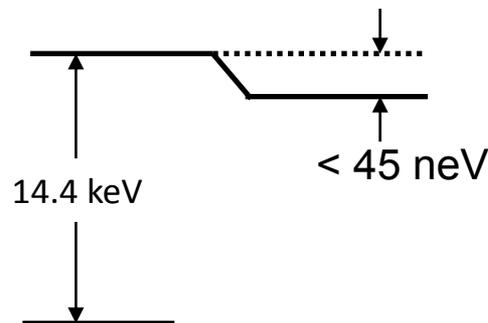
Hydrogen energy levels



Collective Lamb shift



^{57}Fe nuclear resonance

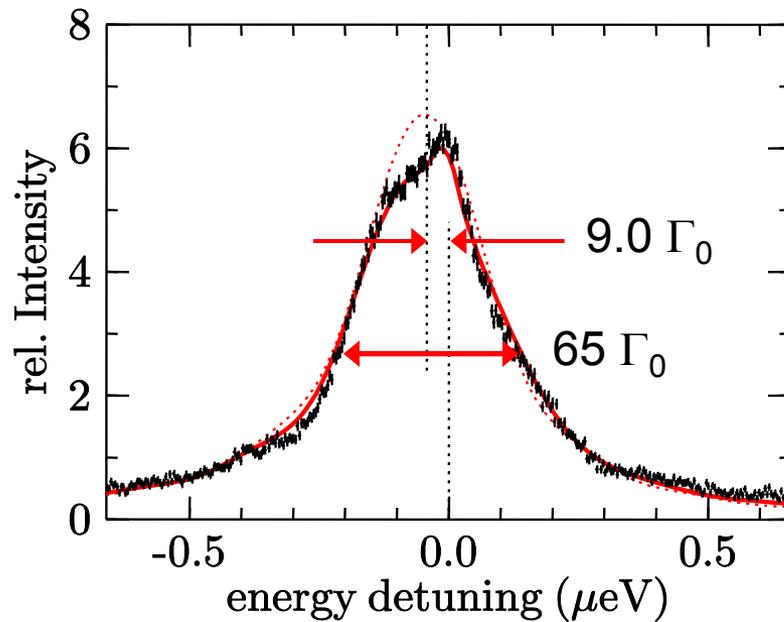
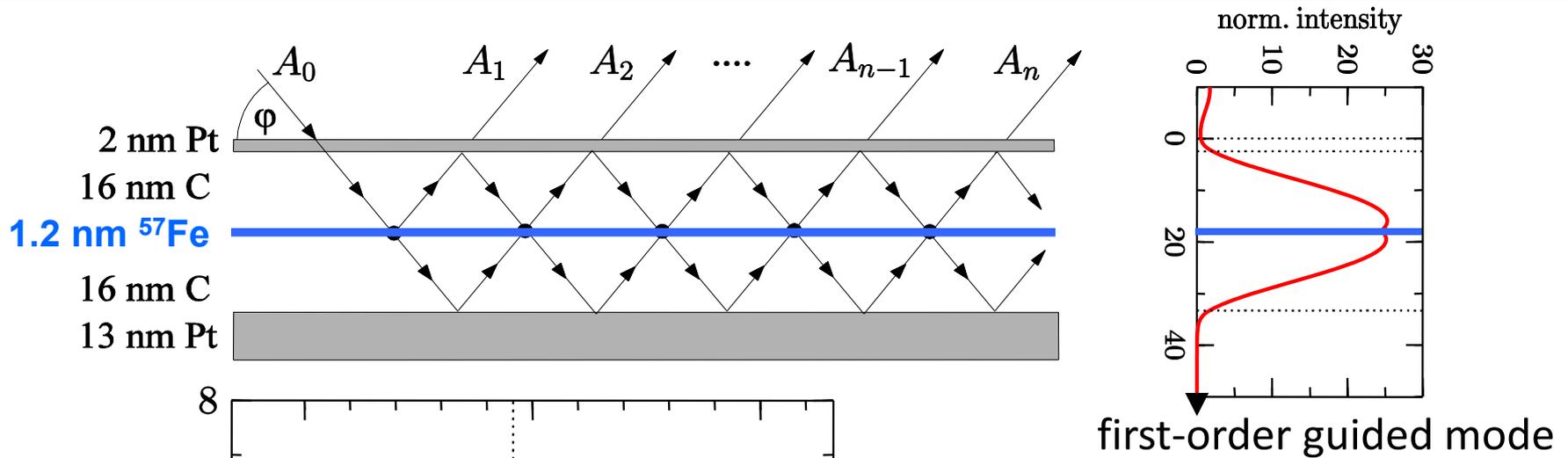


Cloud of N identical atoms acting as a giant atom

R. Friedberg et al.,
Phys. Rep. C 7, 101
(1973)
M. O. Scully, Phys. Rev.
Lett. 102, 143601 (2009)



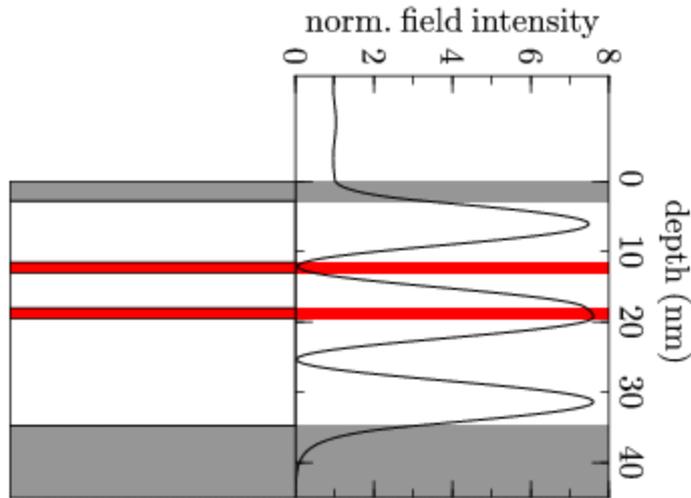
Experimental results (ID18 ESRF): The collective Lamb shift



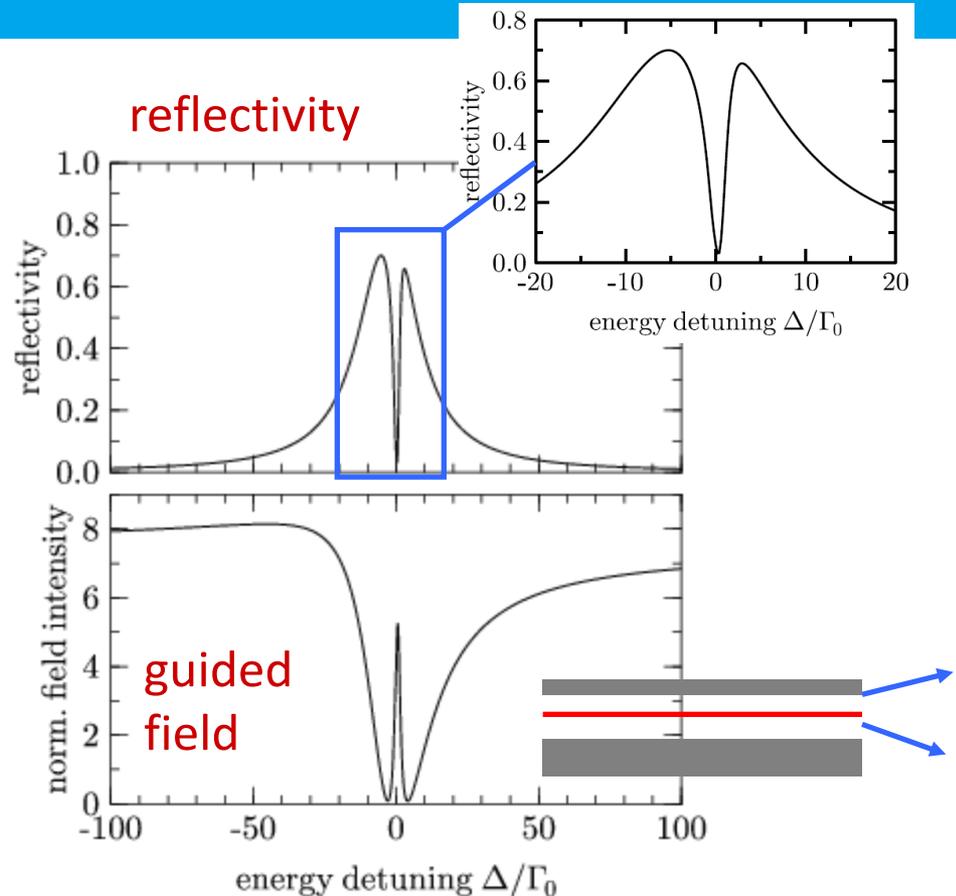
R. Röhlberger, K. Schlage, B. Sahoo, S. Couet, R. Rüffer,
Science 328, 1248 (2010)



Two resonant layers in a cavity



The layers occupy node and antinode of the cavity field



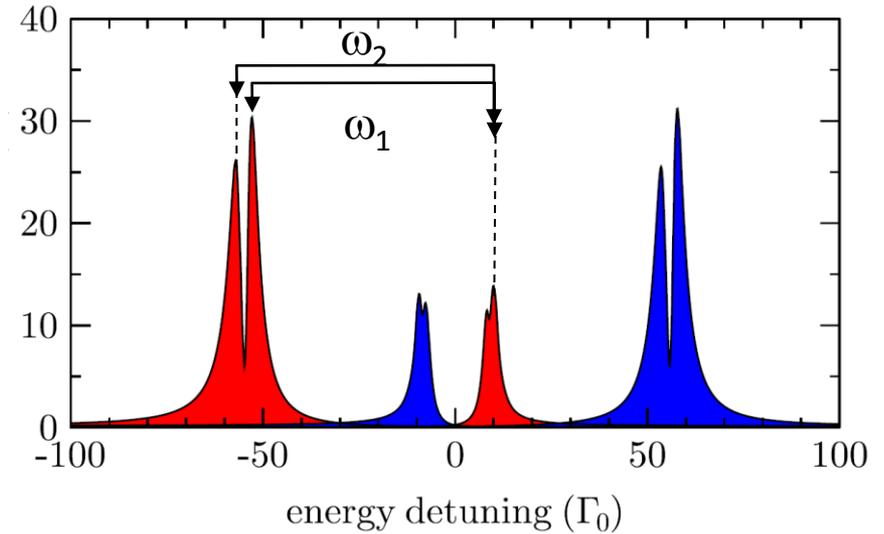
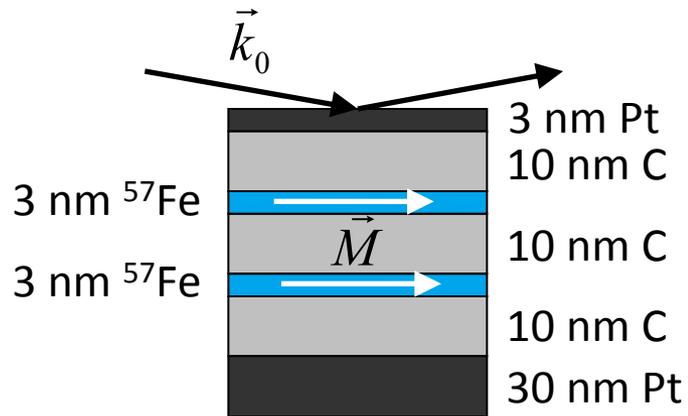
**Electromagnetically
induced transparency**

O. Kocharovskaya, Ya. I. Khanin, Sov. Phys. JETP 63, 945 (1986)

K. J. Boller, A. Imamoglu, S. E. Harris, Phys. Rev. Lett 66, 2593 (1991).

Electromagnetically induced transparency

Possible evidence seen at P01



DATA NOT YET PUBLISHED AND NOT SHOWN IN THIS VERSION

Flux compared to ESRF

	P01	ID18 (4.8m U23)
<u>Now (5m U32):</u>		
Flux after HHLM (in 2eV)	$2.6 \cdot 10^{13}$	$5.0 \cdot 10^{13}$
Flux after 2.4 meV HRM	$6.0 \cdot 10^8$	$1.2 \cdot 10^{10}$
Efficiency of HRM	2% (channel cut)	20%
<u>June 2011 (2x5m U32):</u>		
Flux after HHLM (in 2eV)	$5.2 \cdot 10^{13}$	$5.0 \cdot 10^{13}$
Flux after 1.0 meV HRM	$7.5 \cdot 10^9$	$5.7 \cdot 10^9$
Efficiency	30%	23%
<u>2012 / 2013 (2x5m U32 + 2x5m U21):</u>		
Flux after HHLM (in 2eV)	$1.5 \cdot 10^{14}$	$5.0 \cdot 10^{13}$
Flux after 1.0 (0.7) meV HRM	$2.2 \cdot 10^{10}$	$5.7 \cdot 10^9$

In the end a factor of 4 more flux at P01 All values for 100 mA ring current



Summary

First nuclear forward and nuclear inelastic measurements have been performed

Theoretical flux from 5m U32 undulator confirmed, new HRM setup and 10m optimized undulators strongly desired

Even with raw focussing and low flux enriched nano structured samples can be studied

First experiments on vibrational dynamics in nano systems and on magnetic nano layer systems have been performed successfully

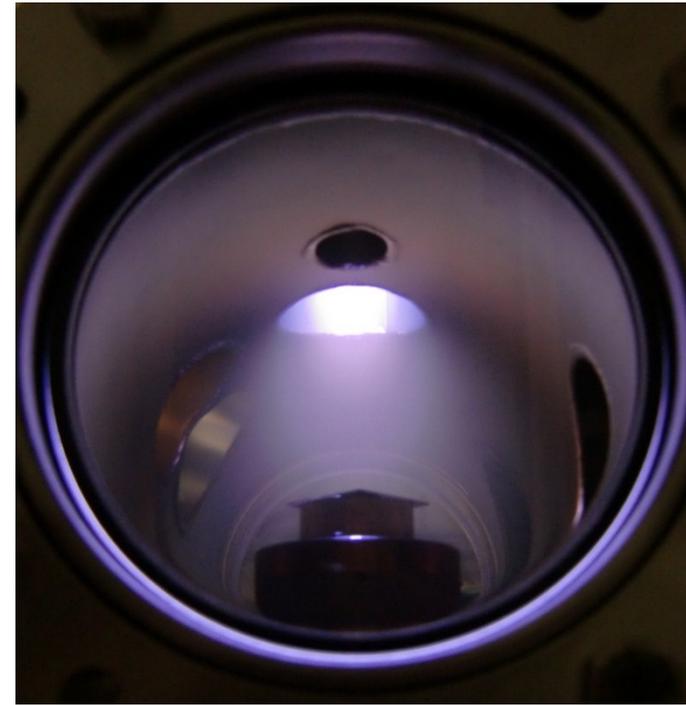
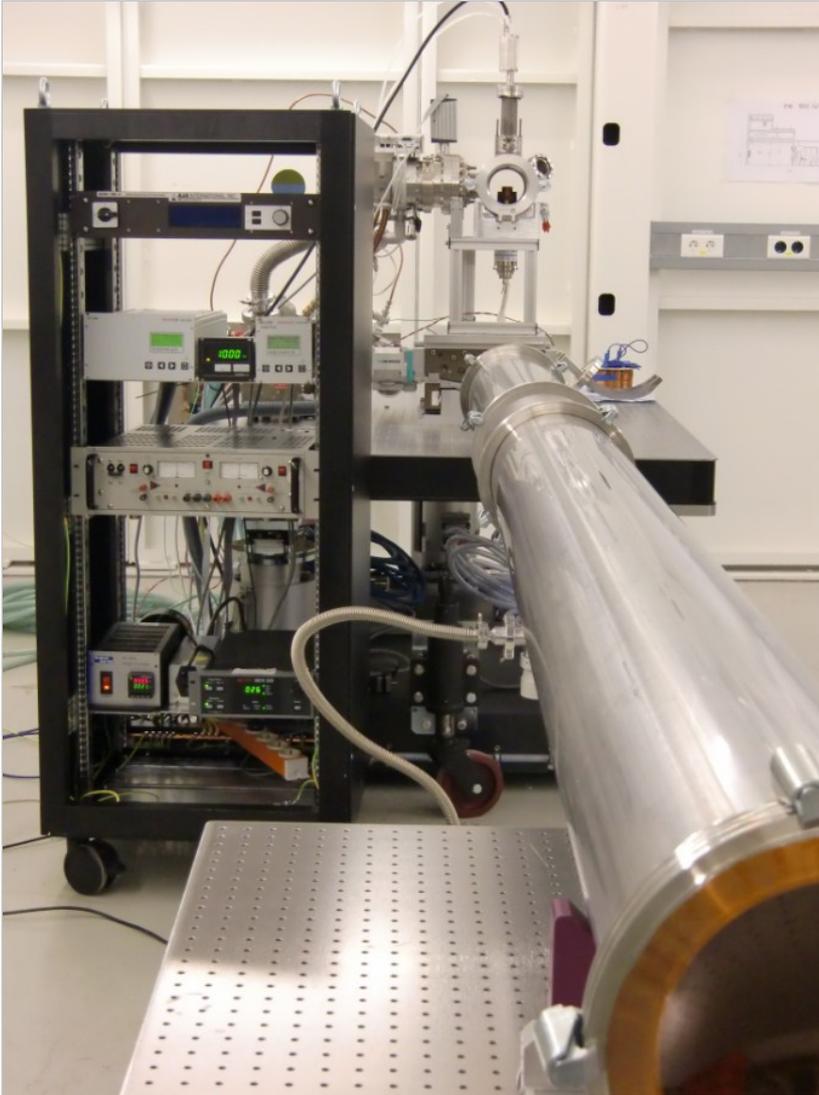
Due to its outstanding brilliance and focusing options P01 will offer unique possibilities for NRS and IXS in nano - and extreme condition science

First official user experiments in November 2011

Application deadline for first round is 1st of March 2011



Outlook: Combined In-Situ NFS-GISAX Setup



- sputter deposition chamber
- flexible flight tube setup (1 m to 6 m)
- guard slits + beam stops
- 2D-PMMA x-ray lenses
- x-ray area detector
- APD stage for nuclear line scans

What's coming next @ P01?

April	2011	TÜV check of EH2 and EH3
May	2011	Installation of the 2nd 5m U32 undulator
June	2011	Installation of the 1 meV HRM
October	2011	Installation of the first KB mirror 1 Tesla magnet, He flux (^3He) and He displacer
November	2011	First official user experiments on NRS
2012		Reflecting focusing mirrors for IXS installed in OH1 IXS diffractometers installed in EH2 Backscattering HRM 1meV for isotopes with $E_0 > 20$ keV in EH3
201x		Installation of 2 x 5m U21

!! Proposals are accepted now 1st deadline is March 01 2011 !!

