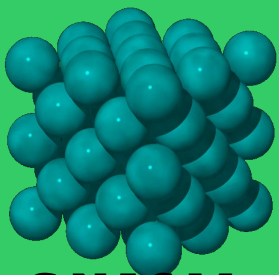




Francium trapping

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CNISM

FUNTRAP12, 4th of December 2012

The “traprad”/“francium”/“wade” collaboration



Ferrara University and INFN:

**R.Calabrese, S.N.Atutov, T.Ishikawa, G.Mazzocca,
(Z.Peshev), (G.Stancari), L.Tomassetti**



Legnaro National Laboratories:

L.Corradi, A.Dainelli



Pisa University:

P.Minguzzi, (S.Sanguinetti), M.L.Chiofalo

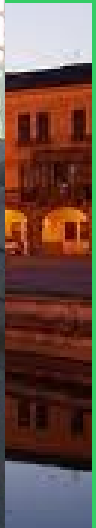


Siena University:

**L.Moi, (C.de Mauro), A.Khanbekyan, C.Marinelli,
L.Marmugi, E.Mariotti, (S.Veronesi)**

Ferruccio Renzoni (UCL), Leonardo Ricci (UniTrento)

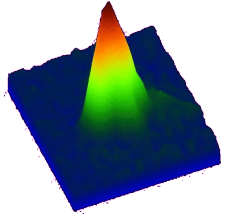
The “traprad”/“francium” location (?)



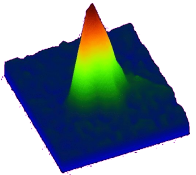
The “**traprad**”/“**francium**” (real) location



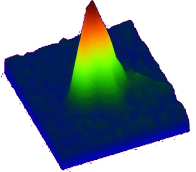
Outline



Introduction/Motivation (if any...)



The LNL apparatus

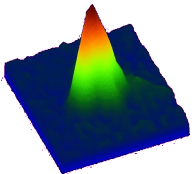


Precision frequency measurements

Laser spectroscopy diffusion coefficients meas.

Detection of lines by change in trapped atom number

Application of Light Induced Atom Desorption



Perspectives

(Bad) facts about francium

	Mass no. (A)	Half-life
Fr	202	0.34 s
	203	0.55 s
	205	3.85 s
	206	15.9 s
	207	14.8 s
	209	50 s
	211	3.1 min
	213	34.6 s
	220	27.4 s
	223	21.8 min
	224	3.3 min
Fr	225	4.0 min
	226	48 s
	227	2.47 min
	228	39 s
	230	19.1 s
	232	5 s

Fr has no stable isotopes

The longest lifetime is 22min

There is at most a tea spoon of francium in the whole Earth at any given time

⇒ **continuous production and trapping for further studies is necessary**



Facts about francium

-First spectroscopy measurements at CERN (ISOLDE):

S. Liberman *et al.*, C. R. Acad. Sci. Ser. B **286**, 253 (1978).
Francium is produced by spallation reactions in Th or U Carbide targets bombarded with protons: 10^9 Fr/s.

-Francium Magneto-Optical Trap (MOT):

J.E. Simsarian *et al.*, PRL **76**, 003522 (1996). (STONY-BROOK)
S.N. Atutov *et al.*, JOSA B **20**, 953 (2003). (LEGNARO)
Nuclear fusion-evaporation reactions in a Au target: 10^6 Fr/s.

Z.-T. Lu *et al.*, PRL **79**, 994 (1997). (Boulder/Berkeley)
Radioactive source: Francium produced in the decay chain
 $^{229}\text{Th} \rightarrow ^{225}\text{Ra} \rightarrow ^{225}\text{Ac} \rightarrow ^{221}\text{Fr} \Rightarrow 10^4$ Fr/s.

(Interesting) facts about francium

spectroscopically poorly known

simple electronic structure

several isotopes suitable for trapping

enhanced relativistic effects

enhanced P and T violations

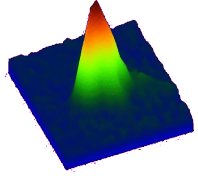


Parity Violation for the 7S - 8S transition:
test of neutral weak interactions

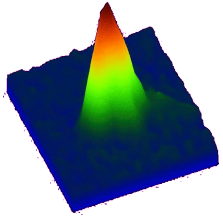
Parity Violation for the ground state hyperfine atomic
transition: measurement of Nuclear Anapole Moment
(TRIUMF/ISAC, production rate 10^{10} Fr/s, trap number 10^7 Fr)

Search for permanent Electric Dipole Moment:
test of Time reversal violation and SUSY
(RNCP @Sendai, trap number 10^8 Fr)

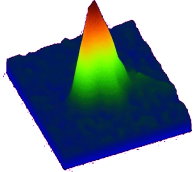
Outline



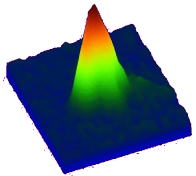
Introduction/Motivation (if any...)



The LNL apparatus

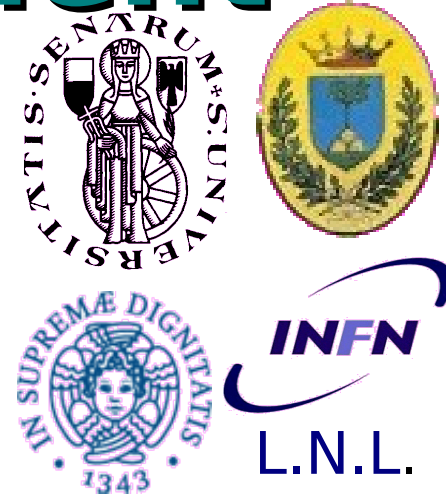
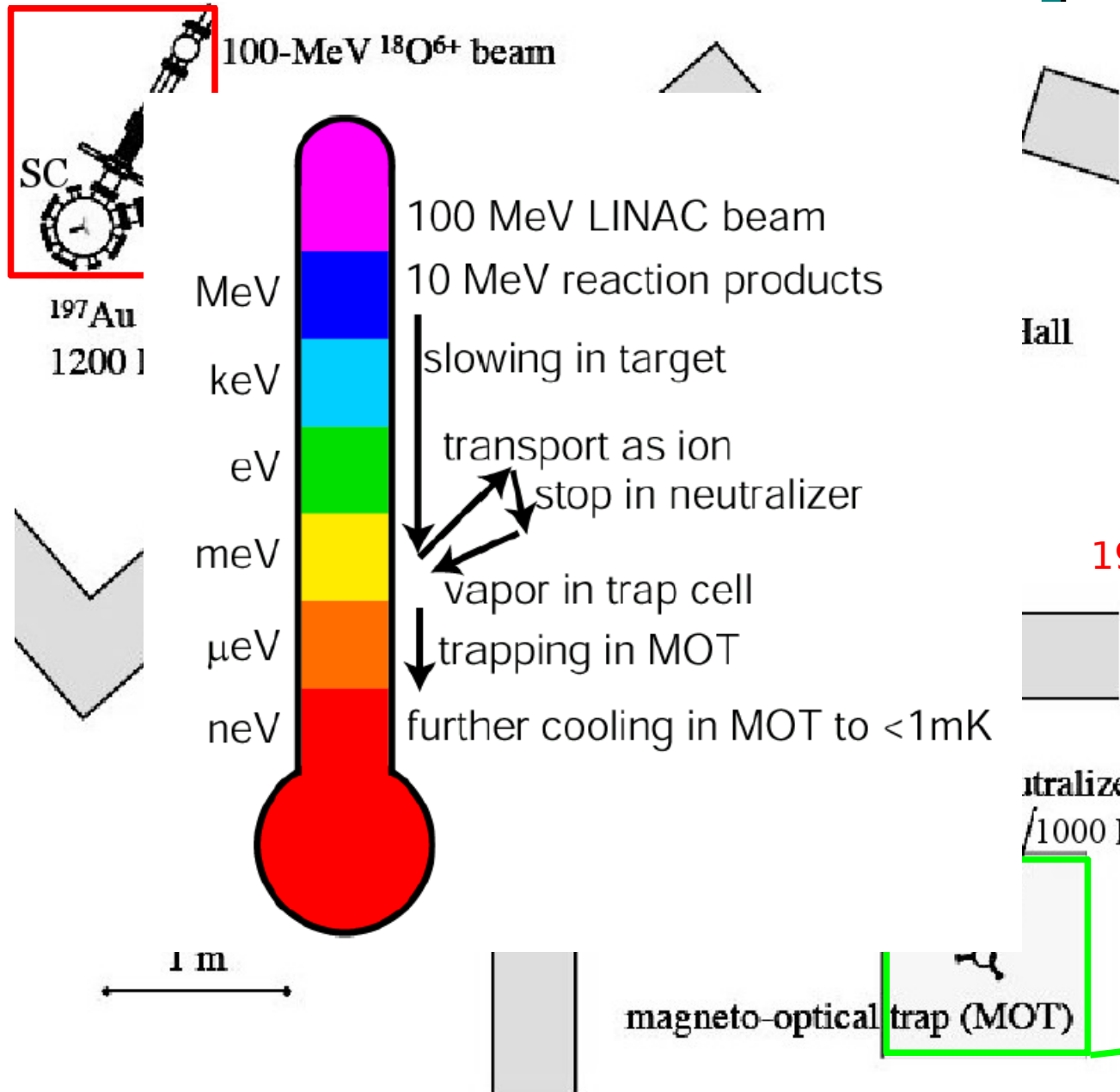


Precision frequency measurements
Laser spectroscopy diffusion coefficients meas.
Detection of lines by change in trapped atom number
Application of Light Induced Atom Desorption



Perspectives

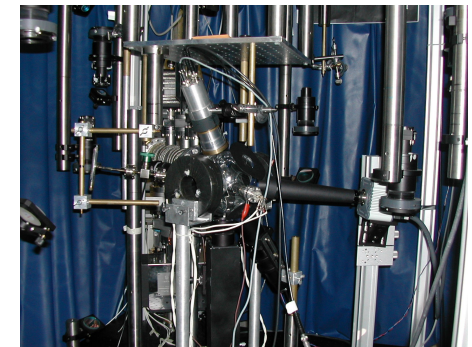
The TRAPRAD experiment



Fr production



Fr^+ (and Rb^+) ions
transport
at 3 keV



The laser setup

Ti:Sa laser pumped by Ar⁺ laser

covering of both Fr (718 nm) and Rb (780 nm) D2 line
for trapping transition

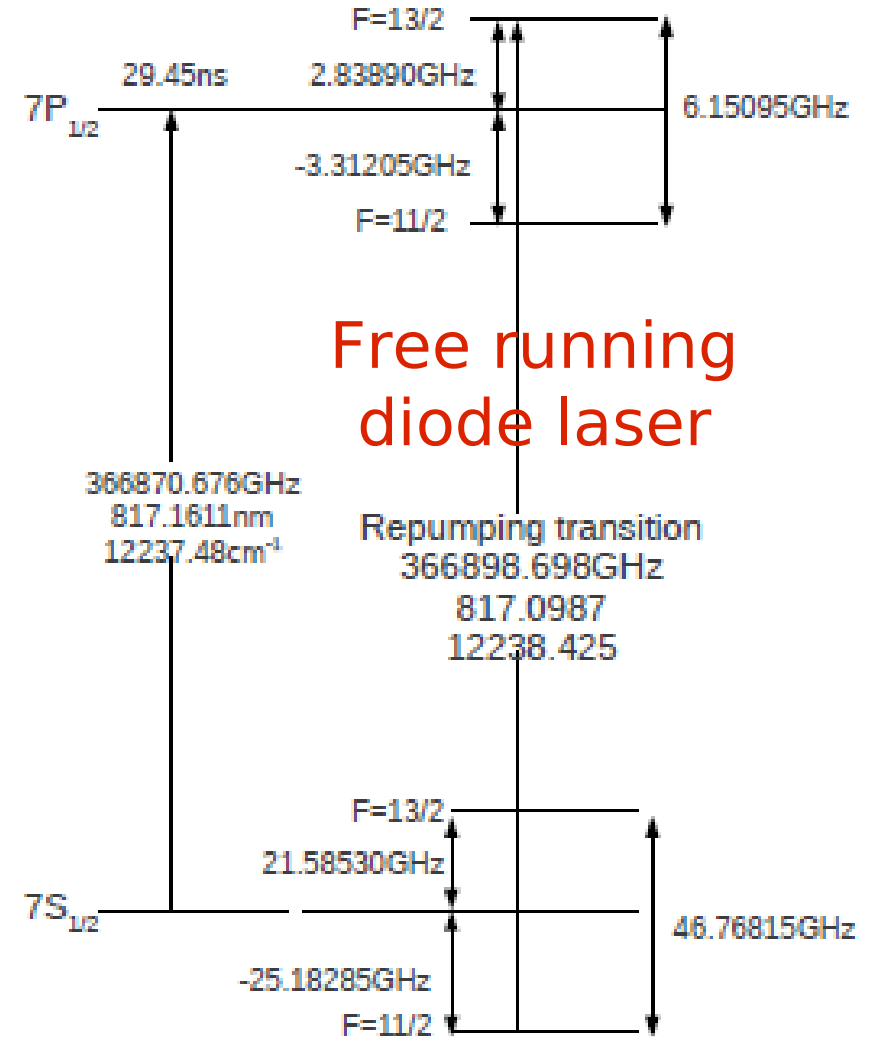
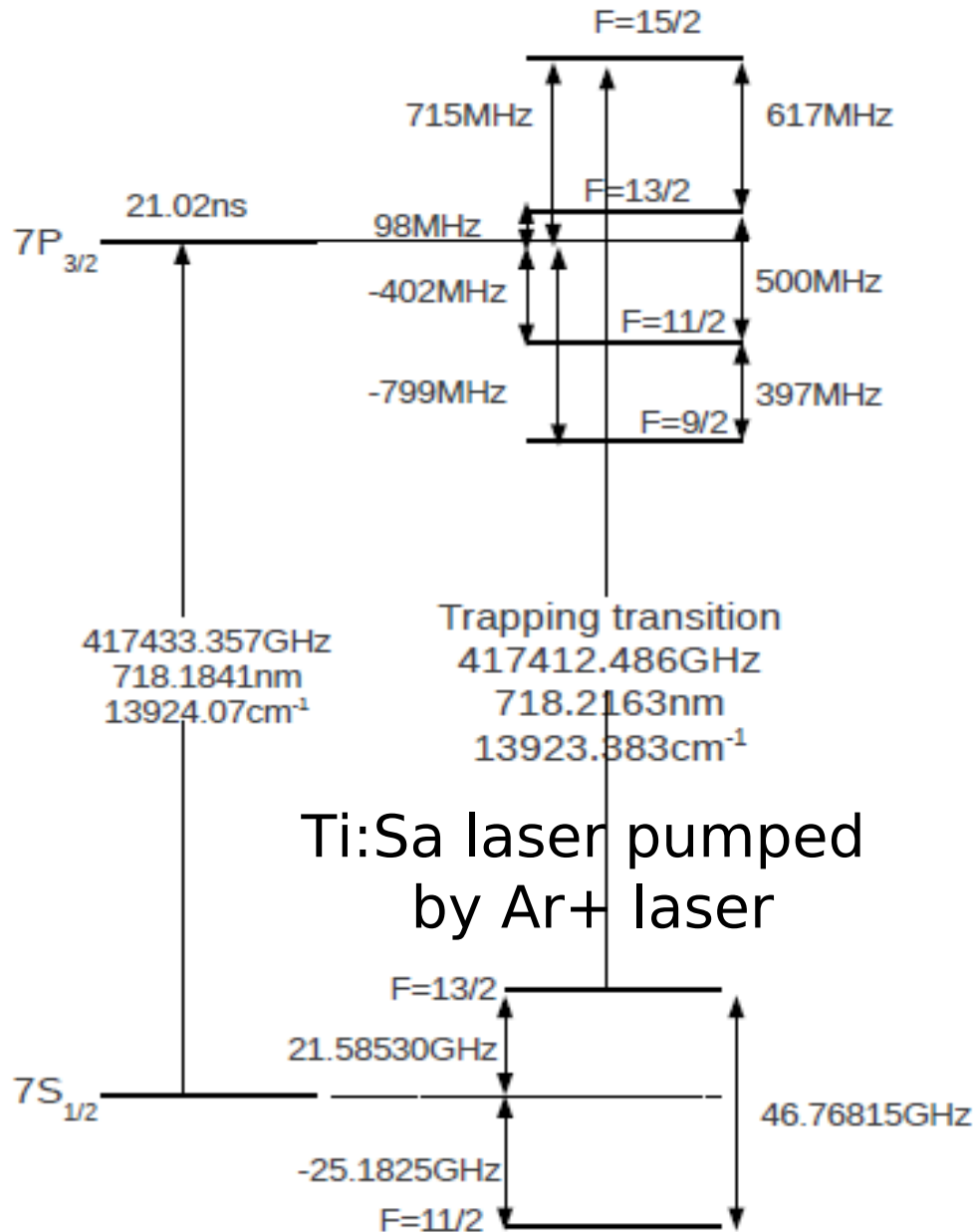
diode lasers for repumping

active stabilisation

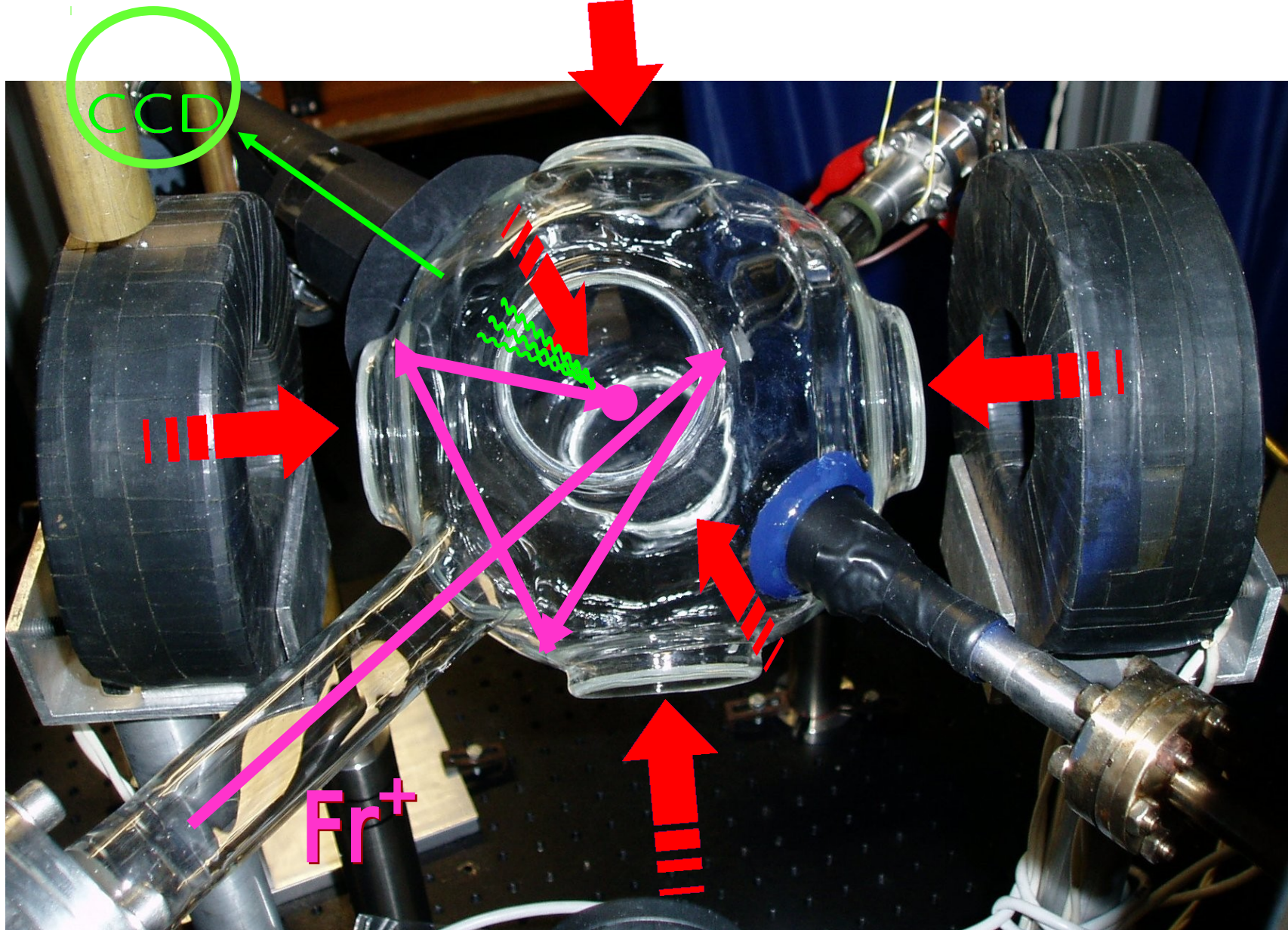
interferometric

wavelength measurement

The trapping and repumping transitions



The MOT cell



Dryfilm coated pyrex

The MOT cell

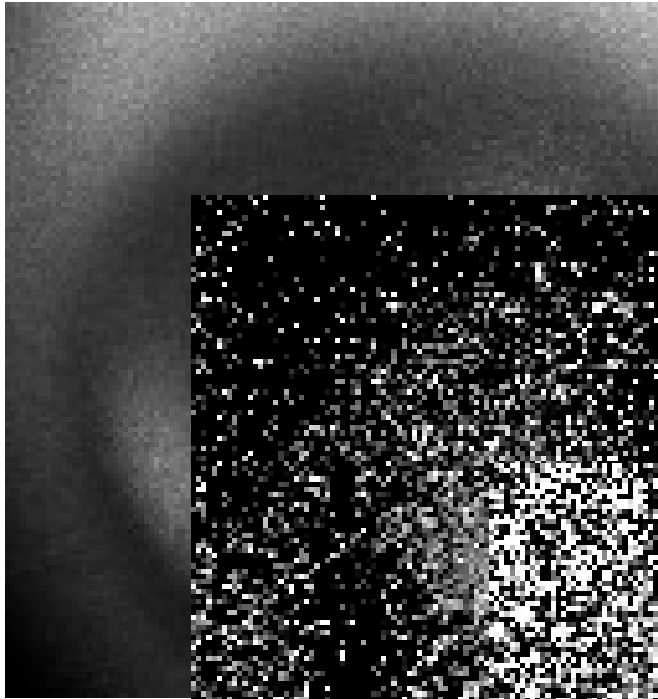


$$\frac{n_+}{n_a} = \frac{g_+}{g_a} \exp\left(\frac{\phi - I}{k_B T}\right)$$

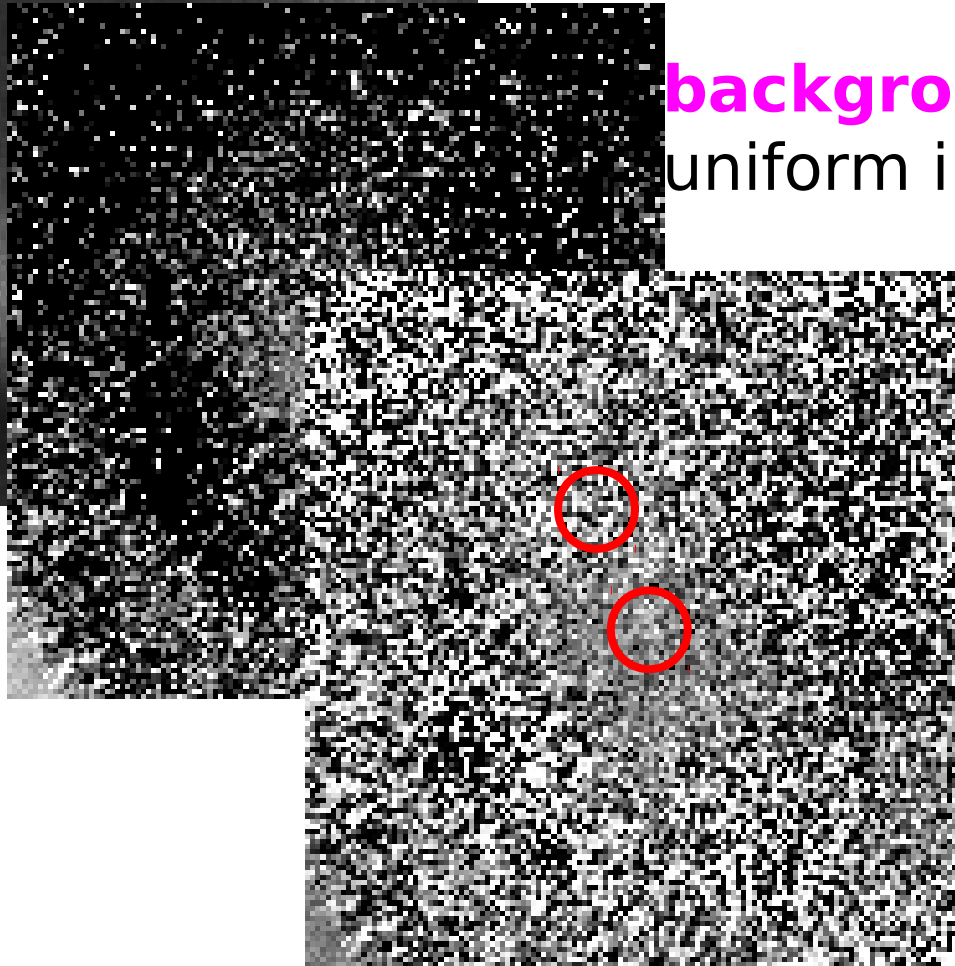
$$\begin{aligned}\phi_{\text{Au}} &= 5.1\text{eV} \\ \phi_{\text{Y}} &= 3.1\text{eV} \\ I &= 4.1\text{eV}\end{aligned}$$

CCD Detection of the MOT

we locate a dark region behind the MOT

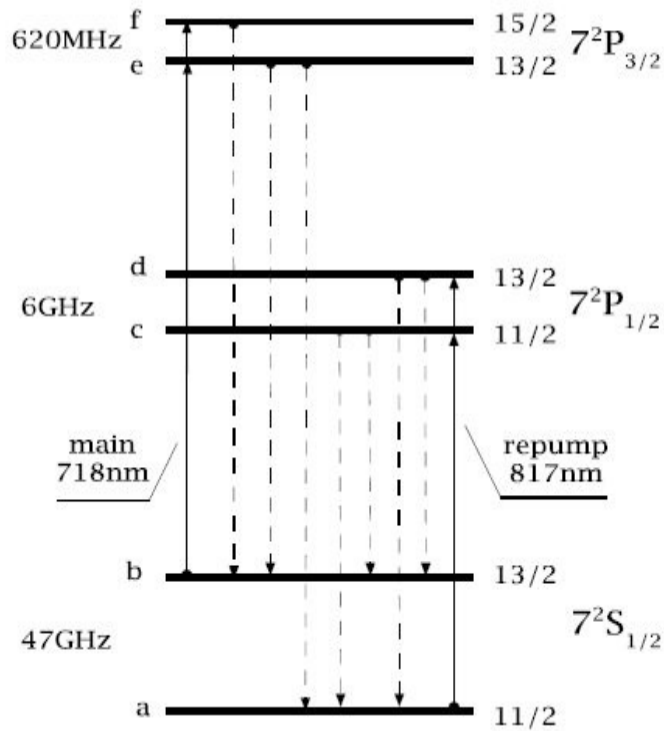


background subtraction:
uniform image



weighted background subtraction: compensation for laser intensity fluctuations

Calibration power-number of atoms



$$H = \hbar \begin{pmatrix} 0 & 0 & A & A & 0 & 0 \\ 0 & \omega_b & 0 & 0 & C & C \\ A^* & 0 & \omega_c & 0 & 0 & 0 \\ A^* & 0 & 0 & \omega_d & 0 & 0 \\ 0 & C^* & 0 & 0 & \omega_e & 0 \\ 0 & C^* & 0 & 0 & 0 & \omega_f \end{pmatrix}$$

hyp: trap laser detuning $-5\gamma_f$

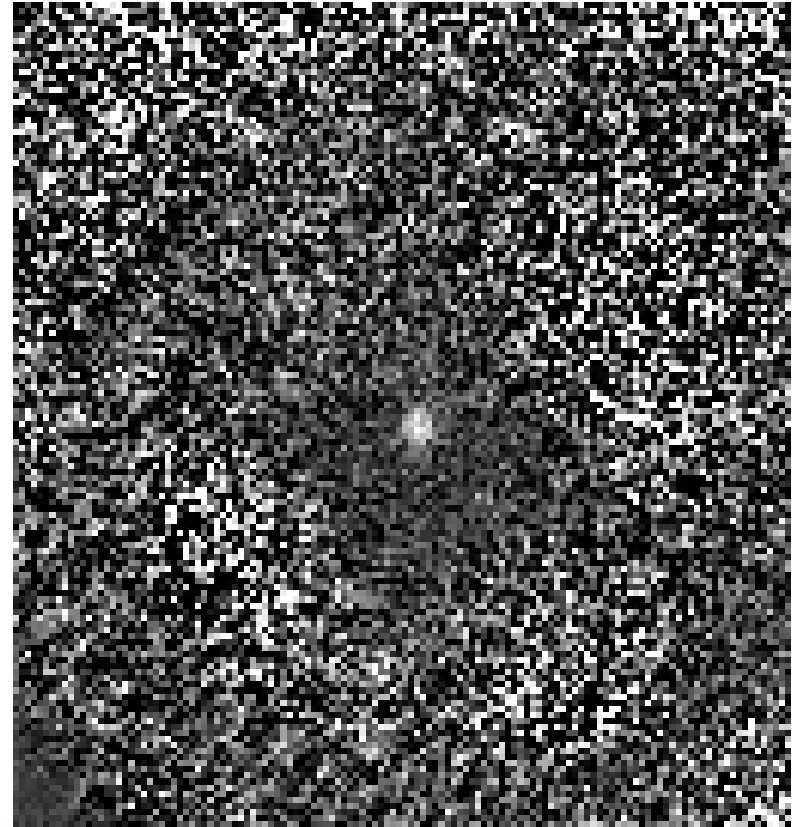
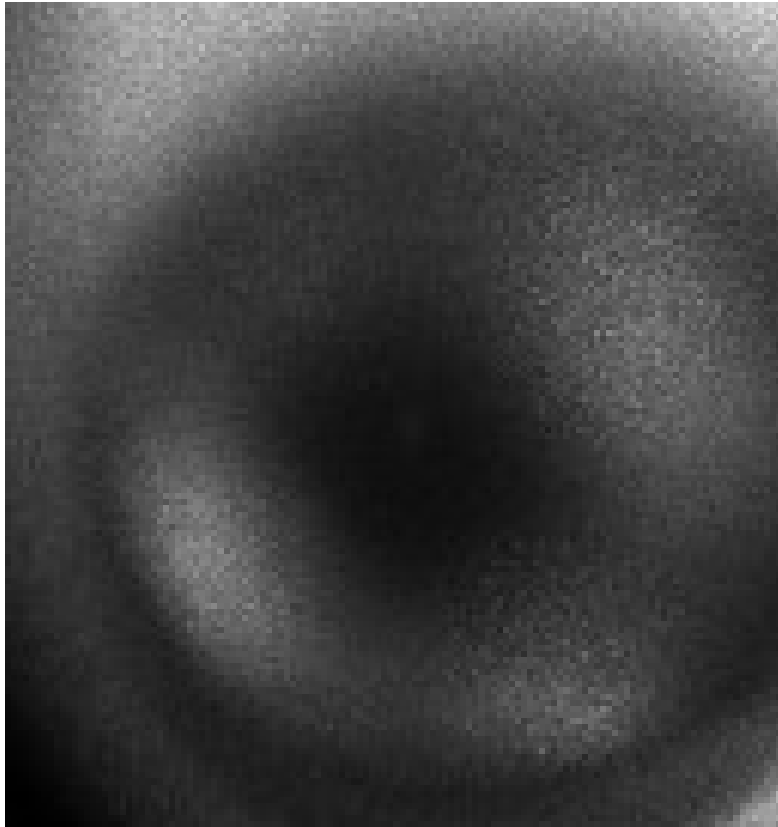
Rb: 1pW 1400 atoms

Fr: 1pW 1900 atoms

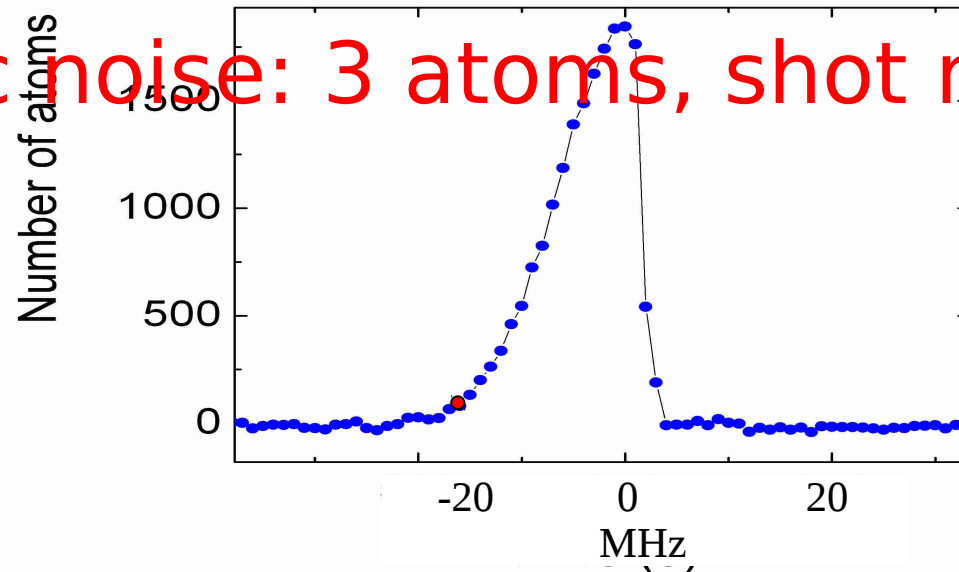
$$\Gamma = \begin{pmatrix} 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & \gamma_e & 0 & 0 & 0 \\ 0 & 0 & 0 & \gamma_e & 0 & 0 \\ 0 & 0 & 0 & 0 & \gamma_f & 0 \\ 0 & 0 & 0 & 0 & 0 & \gamma_f \end{pmatrix}$$

noise level less than 30 atoms

Tests on Rb MOT

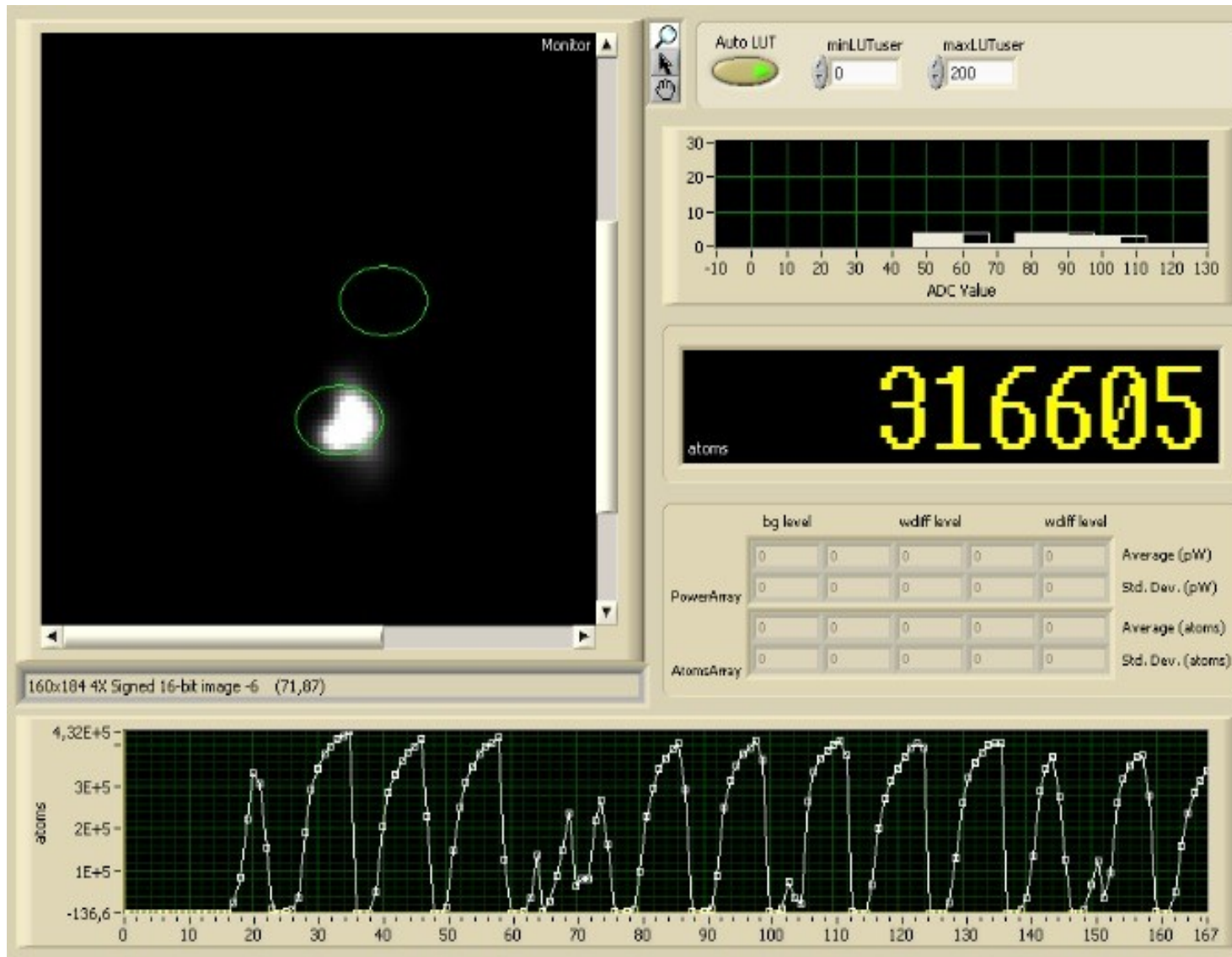


Electronic noise: 3 atoms, shot noise: 8 atoms



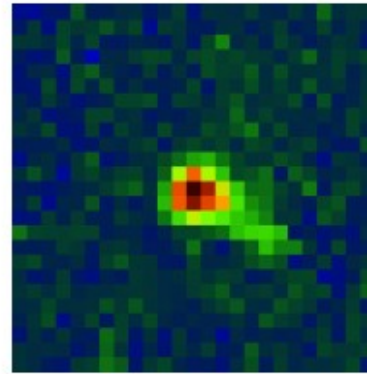
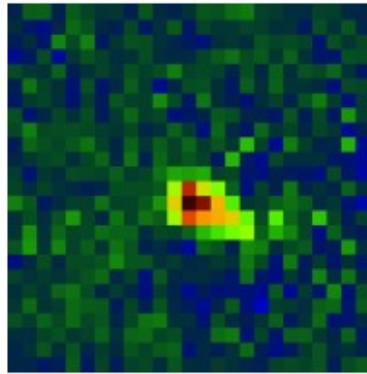
~ 80 atoms

Tests on Rb MOT



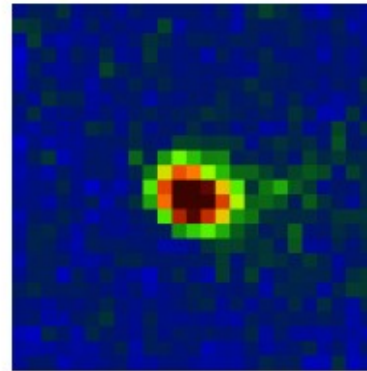
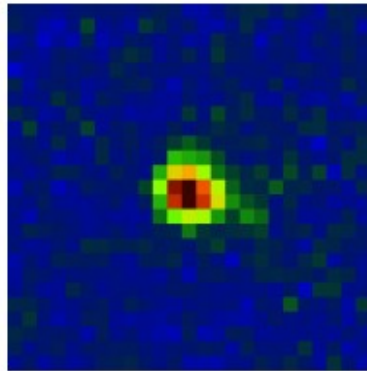
Francium trapping

220 atoms



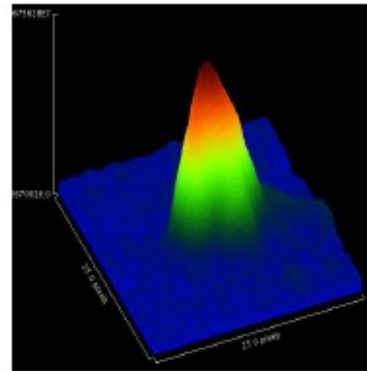
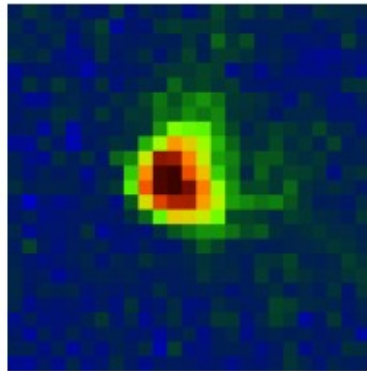
450 atoms

560 atoms



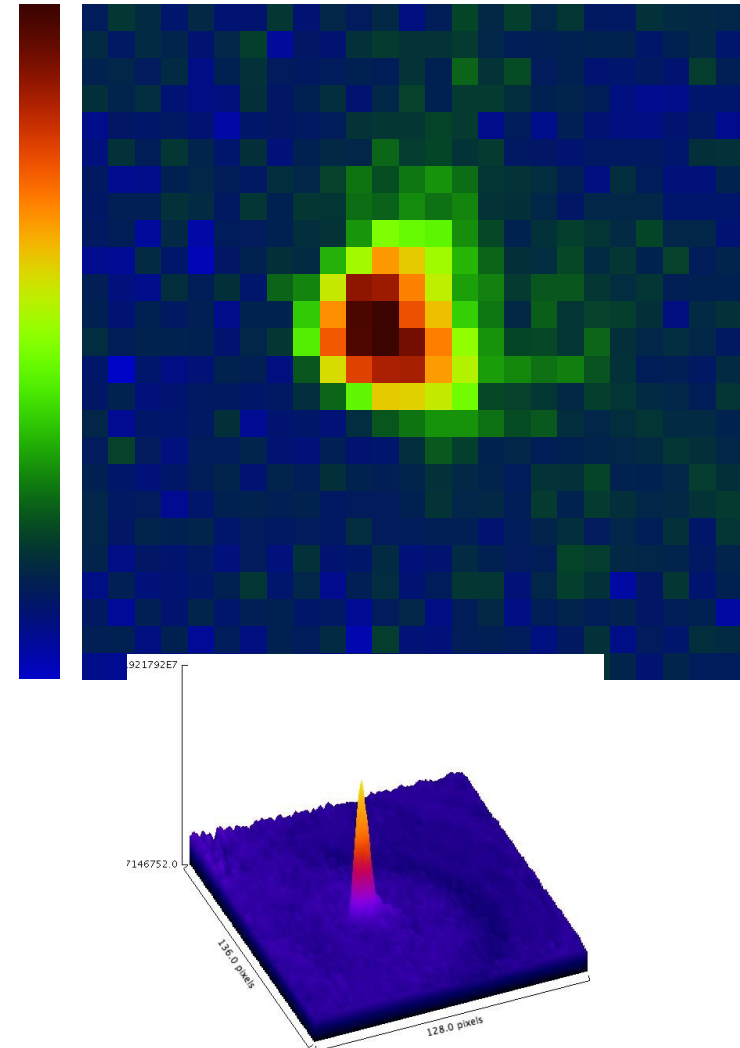
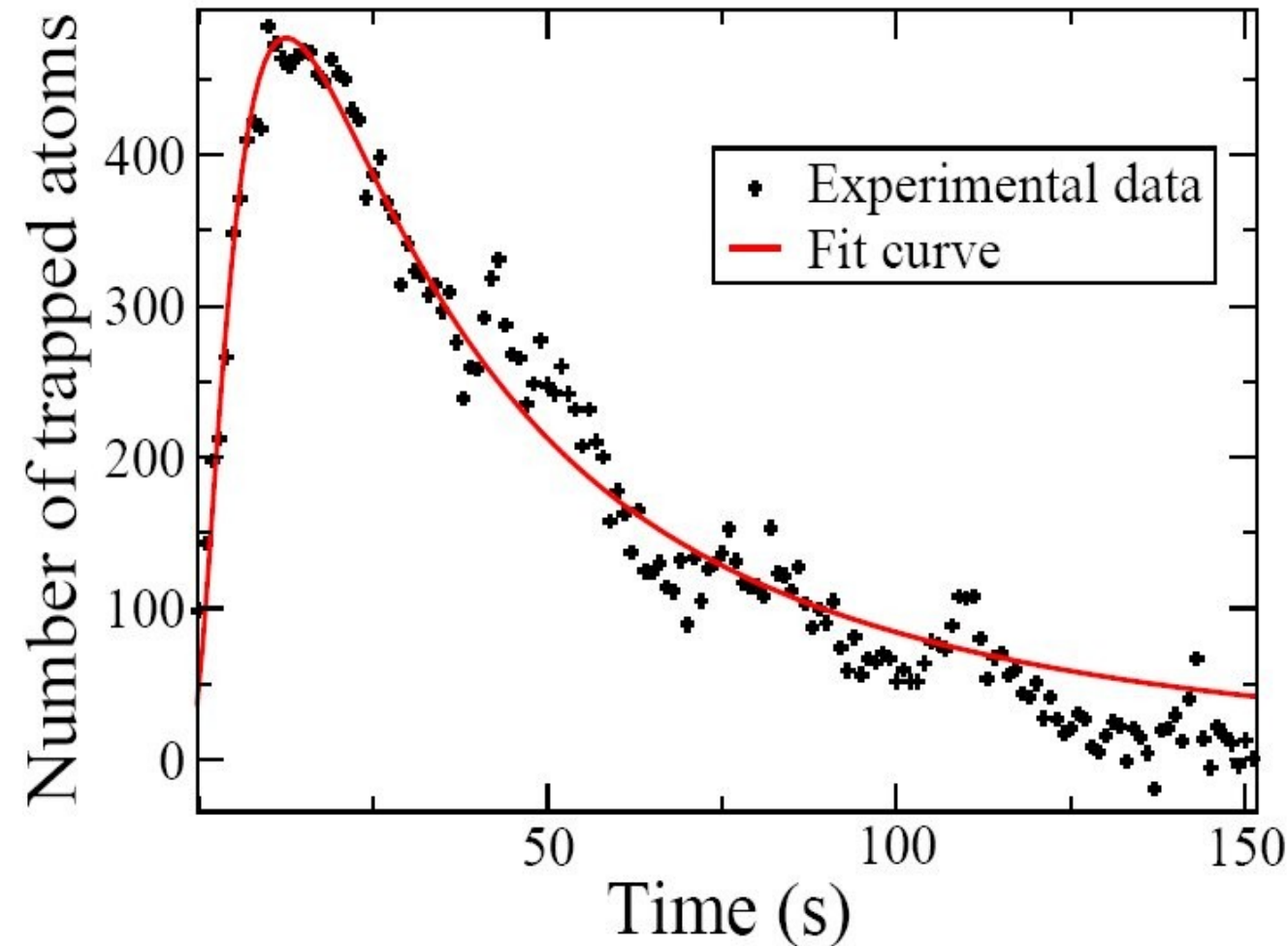
930 atoms

1100 atoms



...we wanted more...

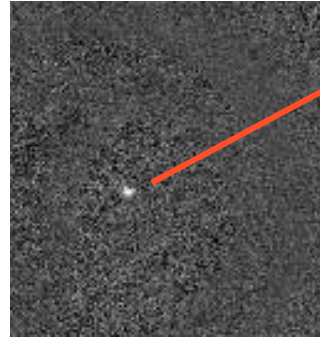
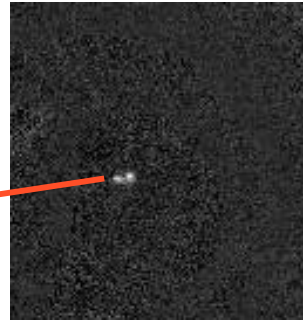
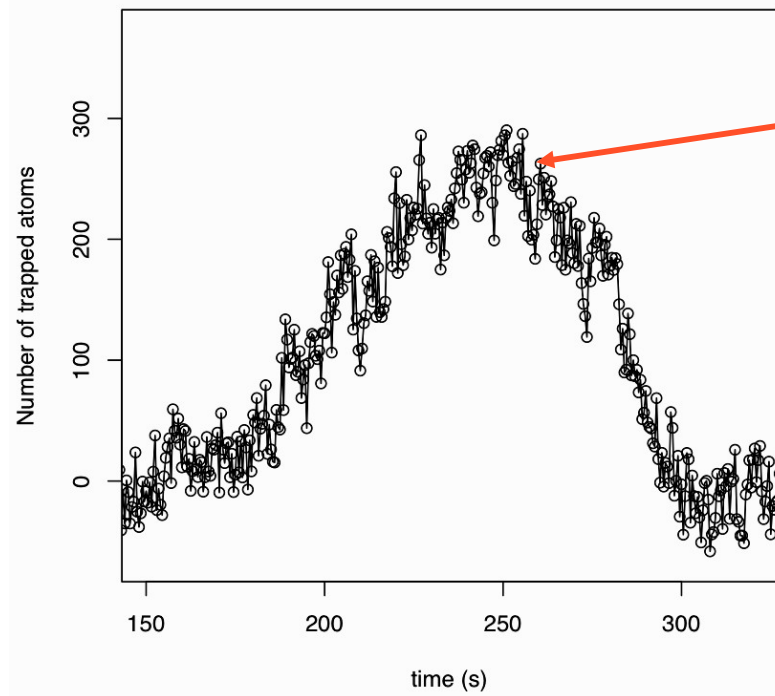
accumulation in the cold yttrium and fast release by suddenly switching on the heating of neutraliser



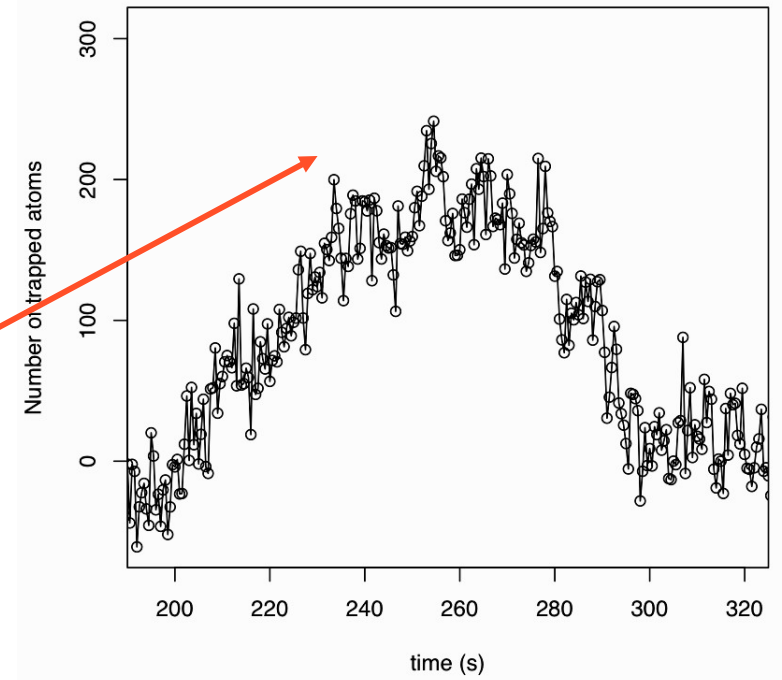
up to 10000 atoms !!

Fr 209, 211

Frequency scan of 209Fr trap



Frequency scan of 211Fr trap



Estimate of trapping efficiency

$$v_c \approx 3000 \text{ cm/s}, v_{th} \approx 1.7 \cdot 10^4 \text{ cm/s}, R_b = 1 \text{ cm}, l = 12 \text{ cm},$$

$$r = 1.25 \text{ cm}, C \approx 0.1 \text{ s}^{-1}, I \approx 10^5 \text{ atomi/s}, R = 7 \text{ cm}, \chi \approx 10000$$

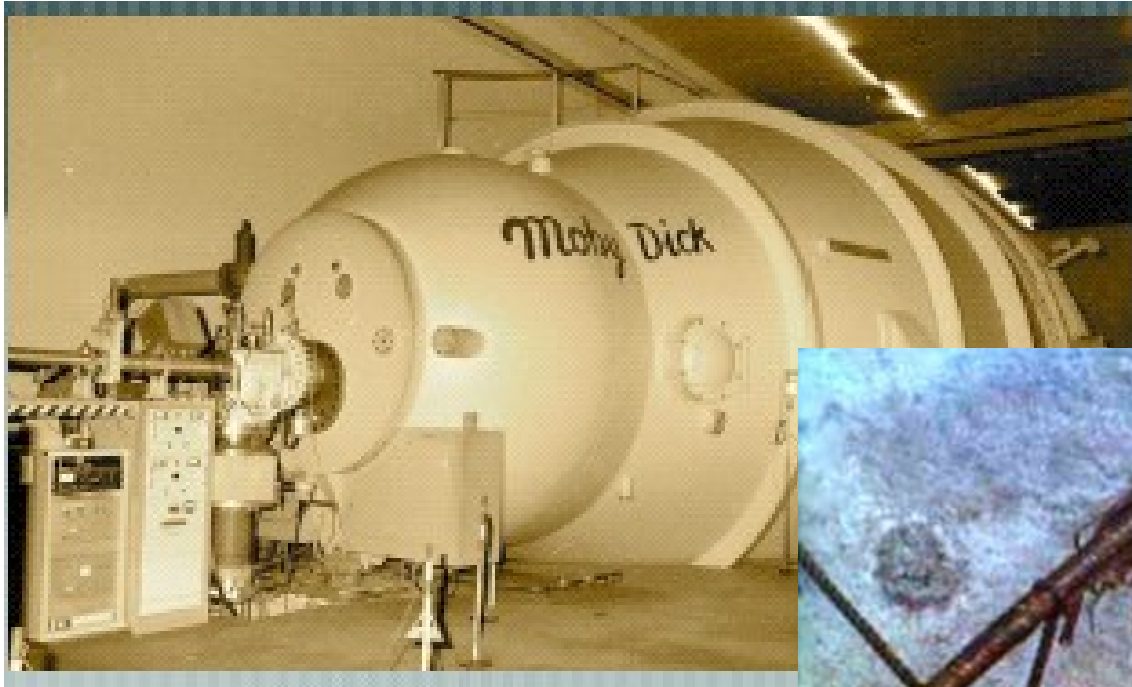
trapping
efficiency

$$\eta \equiv \frac{N_t}{I \tau_c}$$

$$\eta_{210} \simeq 2,3 \times 10^{-4}$$

$$\eta_{209} \simeq 3,2 \times 10^{-4}$$

Experimentalists' life



Francium collaborator

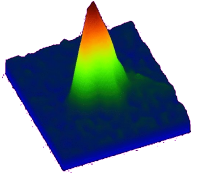
10 days of beam time per year maximum

Melting the target.....

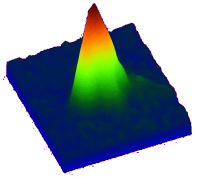


....means to start since the beginning!
(and fight with radioprotectors)

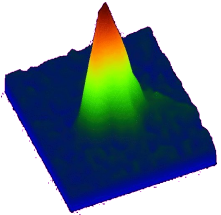
Outline



Introduction/Motivation (if any...)



The LNL apparatus

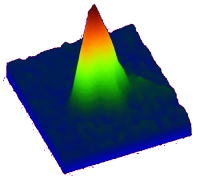


Precision frequency measurements

Laser spectroscopy diffusion coefficients meas.

Detection of lines by change in trapped atom number

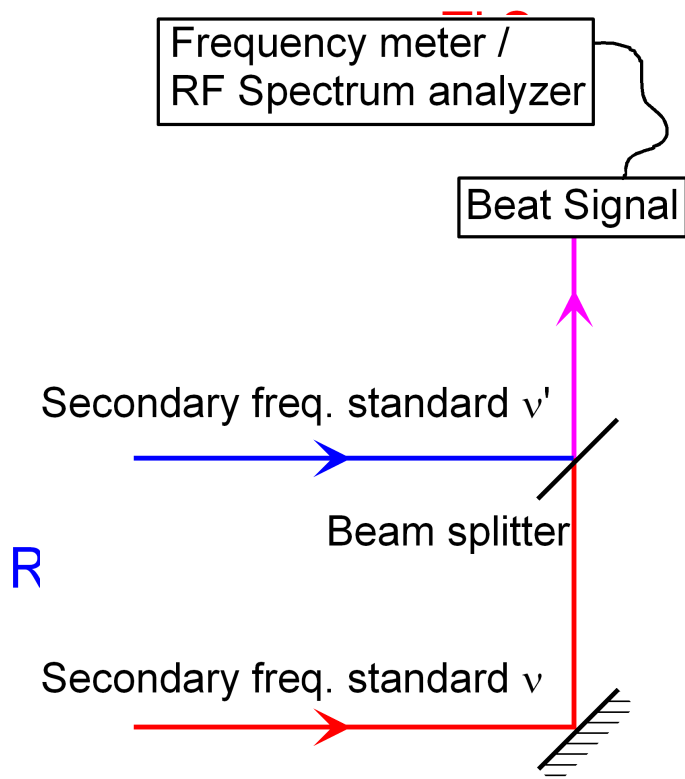
Application of Light Induced Atom Desorption



Perspectives

Precision measurements

We compare the frequency of 2 lasers transmitted by a confocal FP cavity (finesse 200, FSR 2 GHz)



$$\nu_i \cdot n(\nu_i) = \frac{c}{4d} \left(N_i - \frac{2\psi(\nu_i)}{\pi} \right), \quad i = 1, 2$$

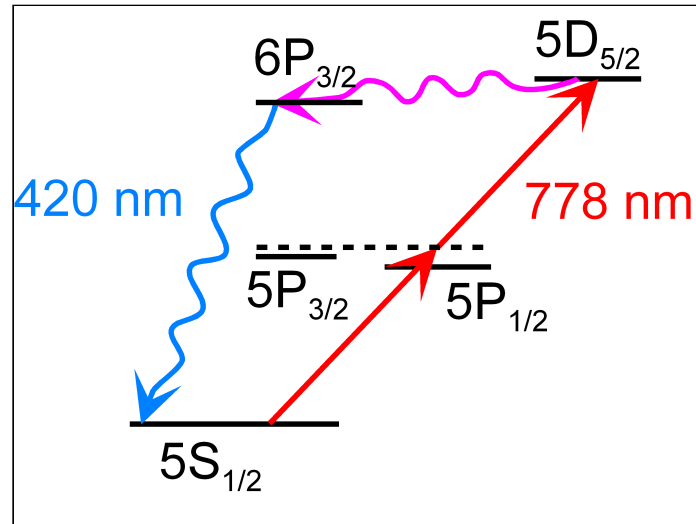
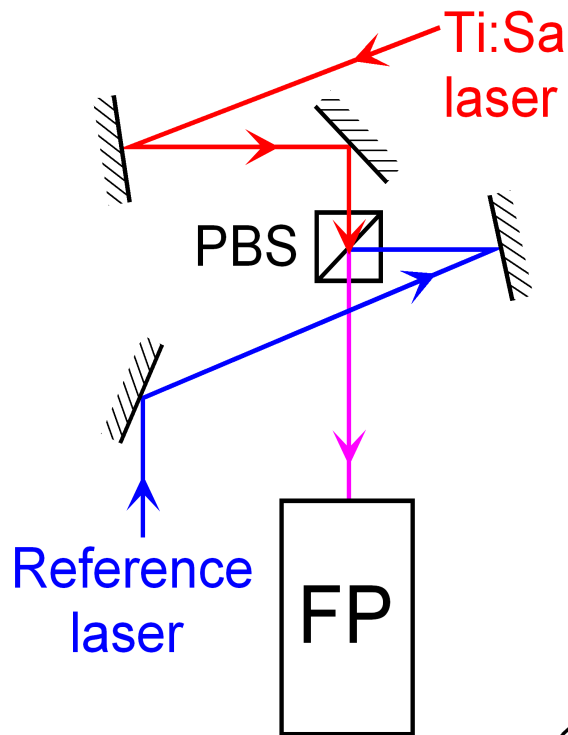
$$\Rightarrow \nu_2 n(\nu_2) - \nu_1 n(\nu_1) = \frac{c}{4d} (N_2 - N_1)$$

$$\nu_2 \frac{n(\nu_2)}{n(\nu_1)} = \nu_1 + \frac{c}{4d n(\nu_1)} \cdot (N_2 - N_1)$$

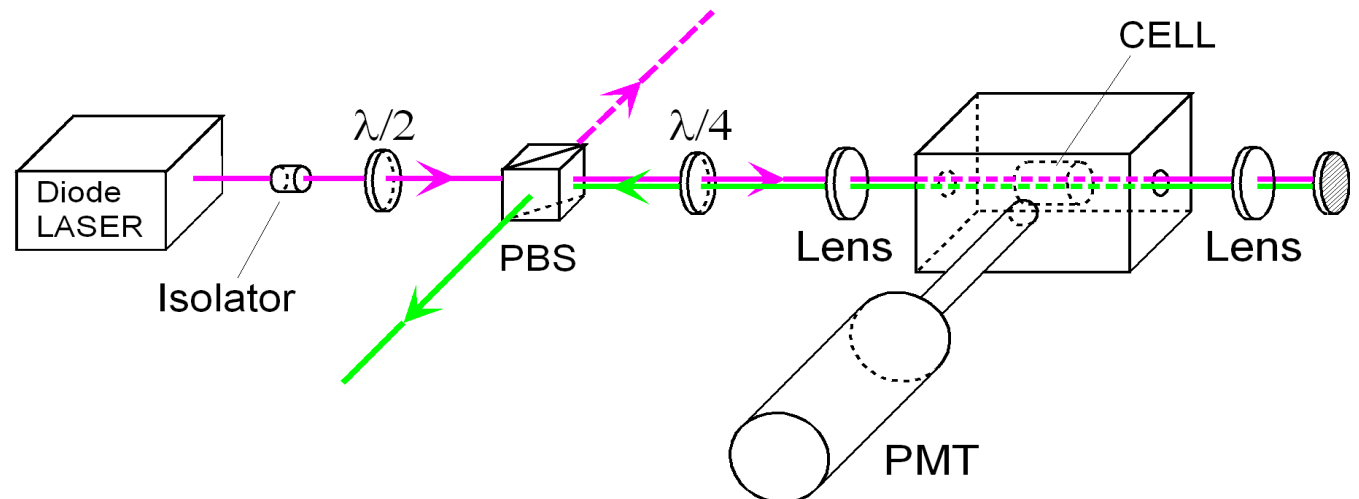
....Measuring the beat signal with a frequency meter (accuracy better than 300 kHz)

Precision measurements

Secondary frequency standard: Rb 5S - 5D_{5/2} 2 photon transition (@ 778 nm) measured with 8 kHz accuracy

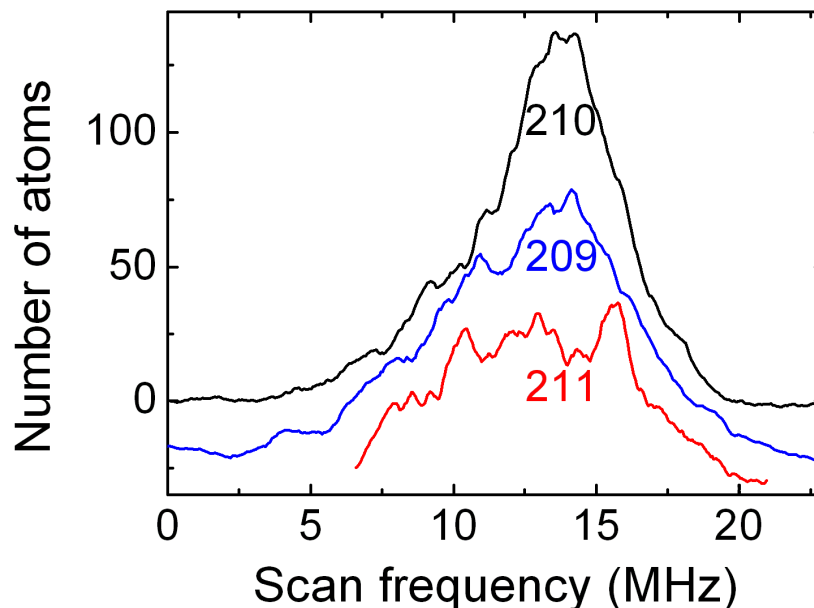
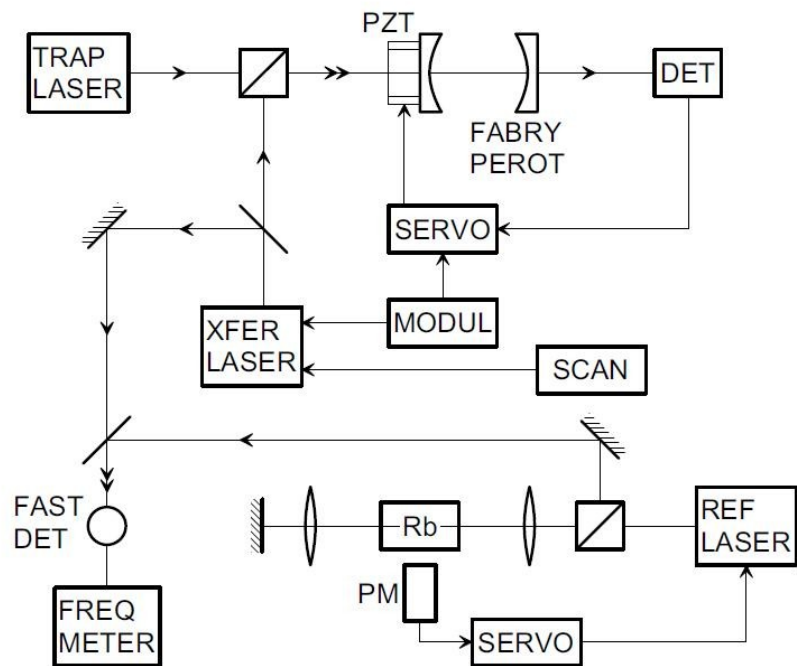


Lock-in detection + stabilization



Precision measurements

Secondary frequency standard: Rb 5S - 5D_{5/2} 2 photon transition (@ 778 nm) measured with 8 kHz accuracy



Preliminary measurements:



Isotope	209	210	211
Trapping freq. (GHz)	417415.0914(90)	417412.4493(90)	417412.6303(90)
Repumping freq. (GHz)	366897.43(5)	366898.70(5)	366895.57(5)

Accuracy:

Calibration ➡ 5 MHz

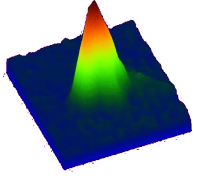
Fabry-Perot maxima ➡ 2 MHz

Refractive index of air ➡ 2 MHz

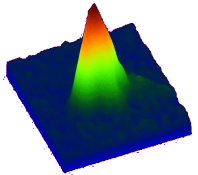
TOTAL ➡ 9 MHz

Published in
Opt. Lett. 34,
893,2009

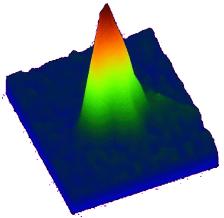
Outline



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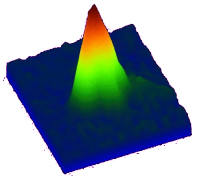


Precision frequency measurements

Laser spectroscopy diffusion coefficients meas.

Detection of lines by change in trapped atom number

Application of Light Induced Atom Desorption



Perspectives

MOT dynamics

$$\begin{cases} \dot{N}_t = LN_v - CN_t - BN_t^2 - AN_vN_t - \Gamma_r N_t \\ \dot{N}_v = -(L + W)N_v + CN_t + BN_t^2 + AN_vN_t - \Gamma_r N_v + f \end{cases}$$

$$\begin{cases} \dot{N}_t = LN_v - CN_t \\ \dot{N}_v = CN_t - LN_v - WN_v + I \end{cases} \quad L = \frac{1}{4\sqrt{\pi}} \frac{v_c^4}{v_T^3} \frac{3\pi w^2}{4\pi R^3}$$

collision rate *loading rate*

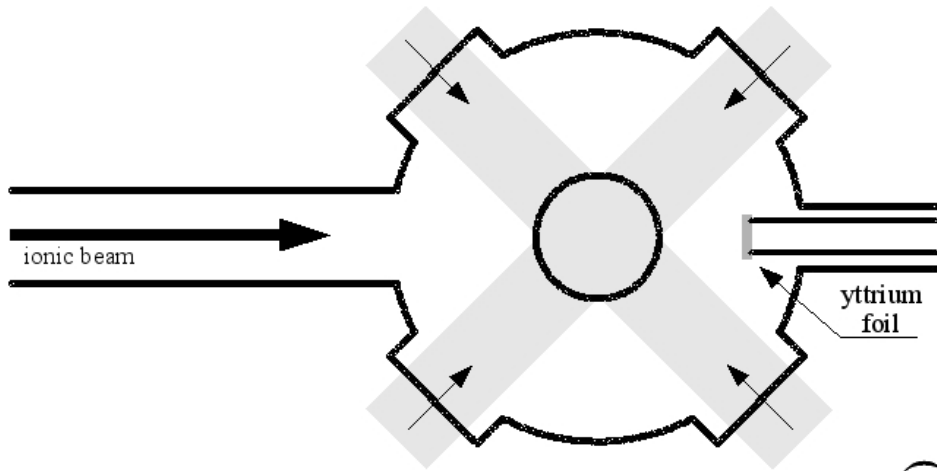
$$W \equiv \frac{1}{\tau_{loss}} \equiv \frac{1}{\tau_{esc}} + \frac{1}{\tau_{sto}} + \frac{1}{\tau_{dec}} \quad 275 \text{ s}$$

250 ms *1500 ms* *lossrate*

$$N_t = \frac{LI}{CW} \quad N_v = N_{eq} = \frac{I}{W}$$

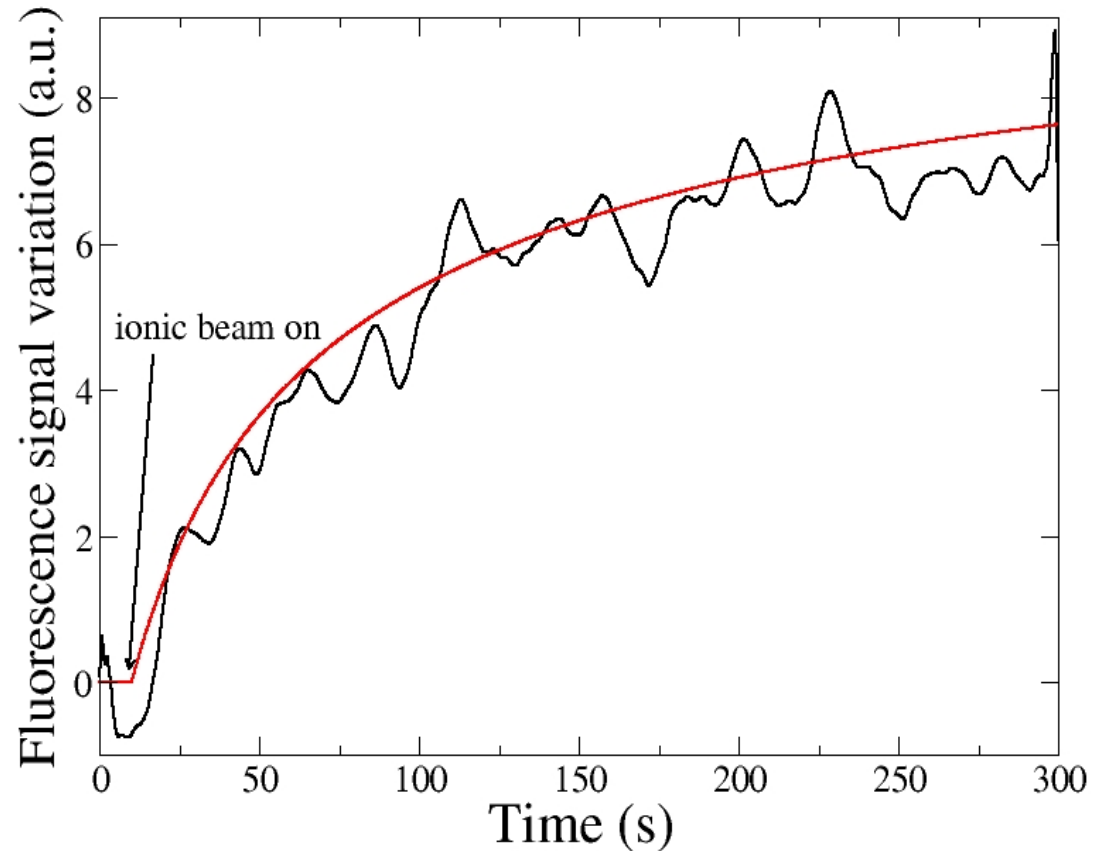
$$N_t = \frac{3}{8\sqrt{\pi}} \left(\frac{v_c}{v_{th}} \right)^4 \frac{R_b^2 l I}{C r^3} \frac{1}{1 + \frac{6lR^2}{\chi r^3}}$$

Tests on neutraliser



from calculated implantation depth of 49 Å diffusion coefficient is determined

$$\frac{\partial N(x,t)}{\partial t} = D \frac{\partial^2 N(x,t)}{\partial x^2} - \Gamma_p N(x,t) + \varphi(x,t)$$

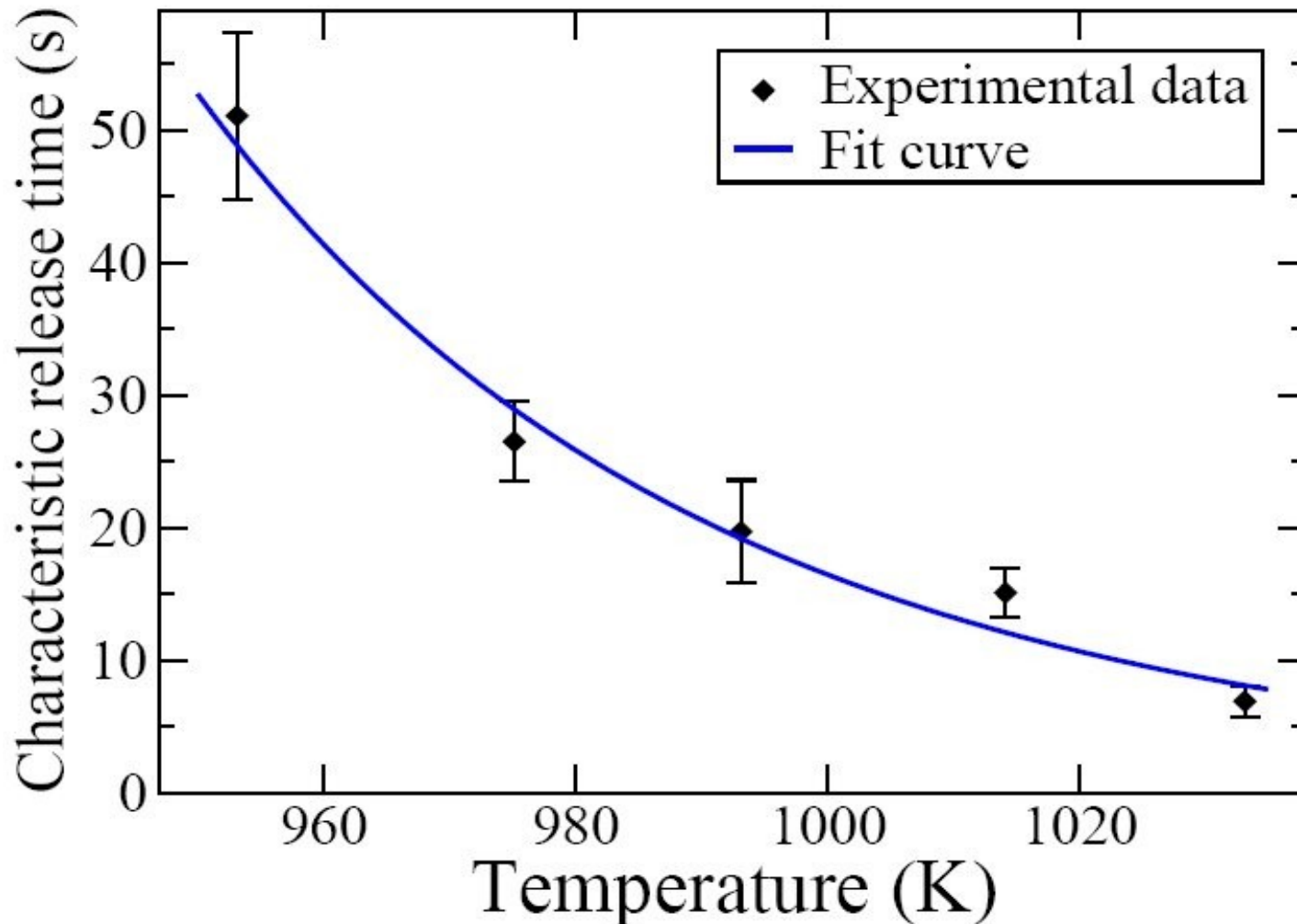


Diffusion time measurements

fit function comes from diffusion equation in the neutraliser

parameters of fit

$$\tau_d, A \propto N_0$$



from calculated implantation depth of 51 Å diffusion coefficient of Fr is determined for the first time

Published in
Phys.Rev.A **78**,
063415 2008

Diffusion time measurements

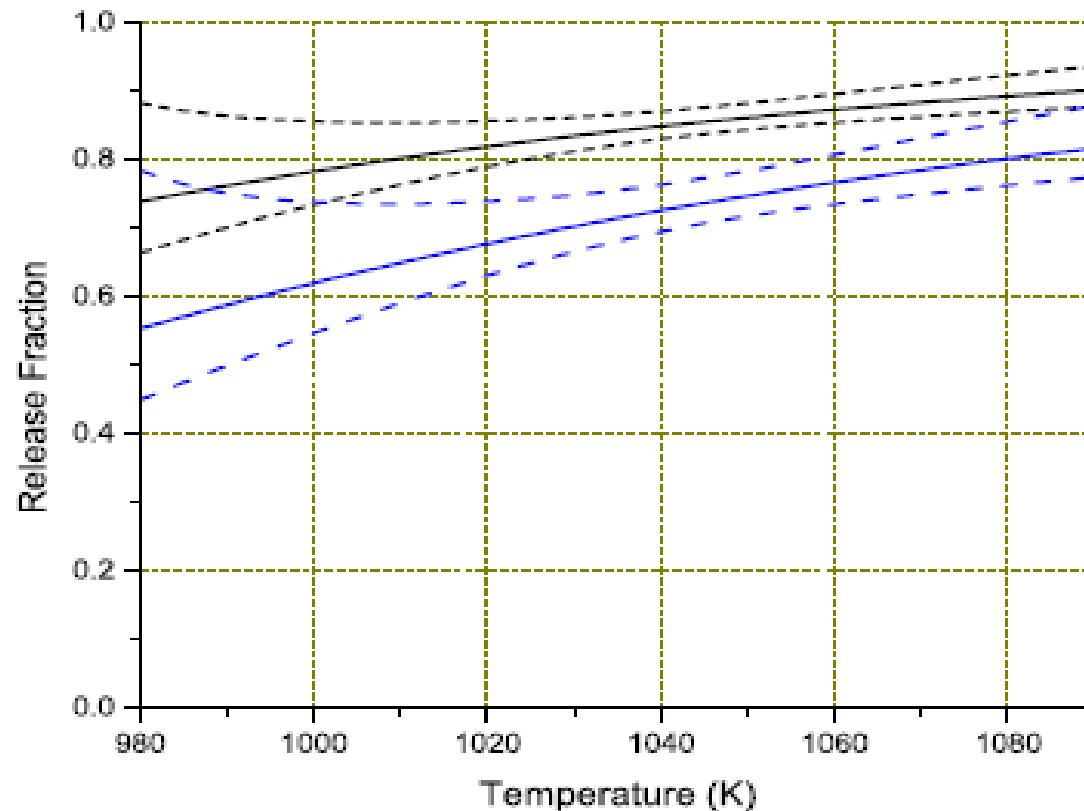
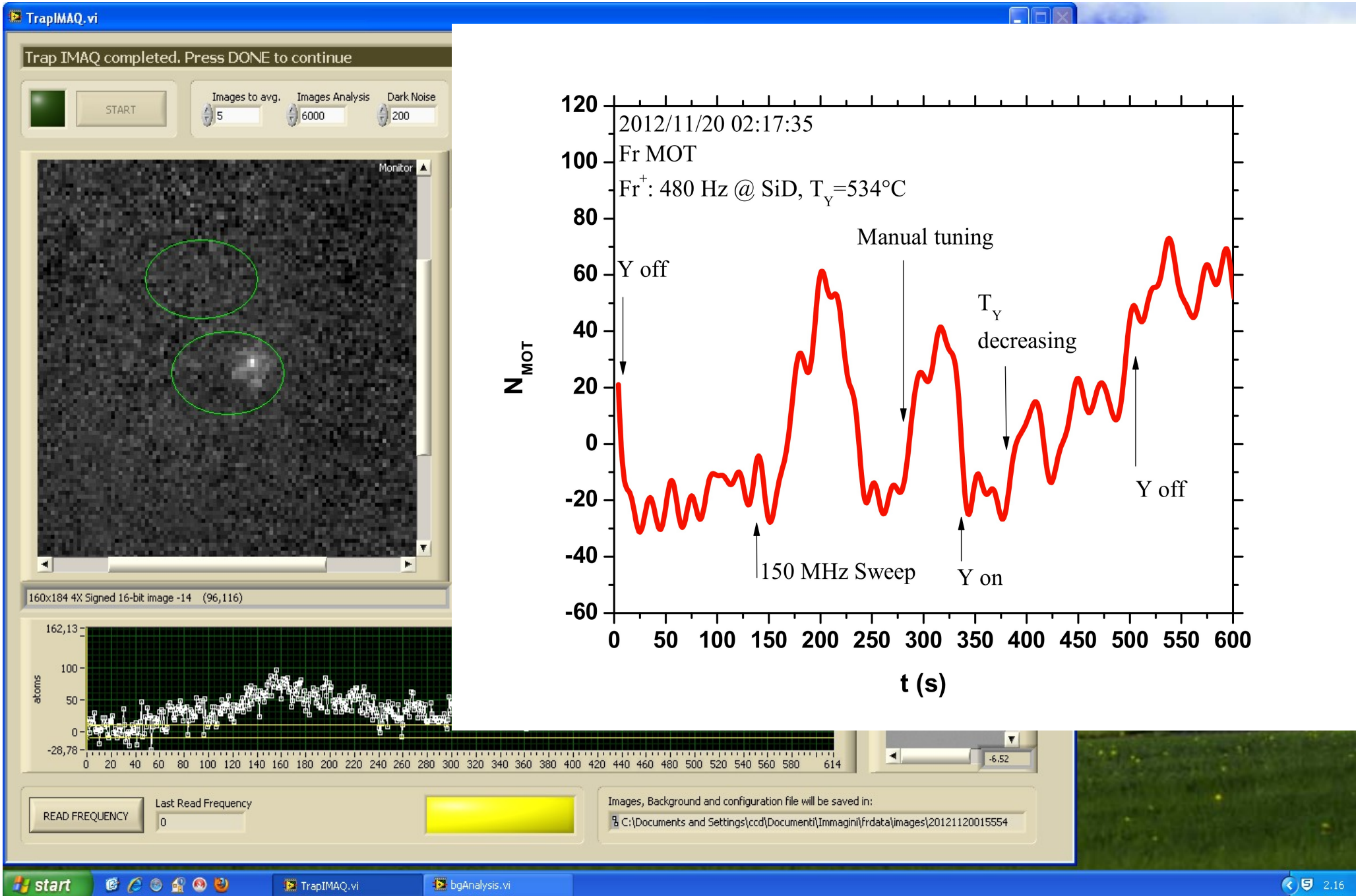


FIG. 8. (Color online) Calculated release fraction vs temperature for isotope 210 (black curve) and 209 (blue curve), according to the Arrhenius law and E_a and τ_{1000} given from our fit. Dashed curves give the uncertainty interval.

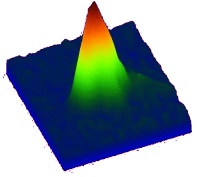
Neutralizer behaves pretty well!

Room temperature neutralizer trap!

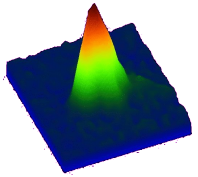


beam time of november

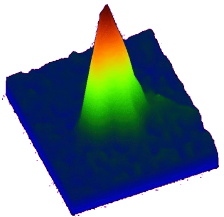
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The LNL apparatus

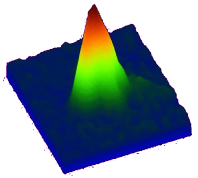


Precision frequency measurements

Laser spectroscopy diffusion coefficients meas.

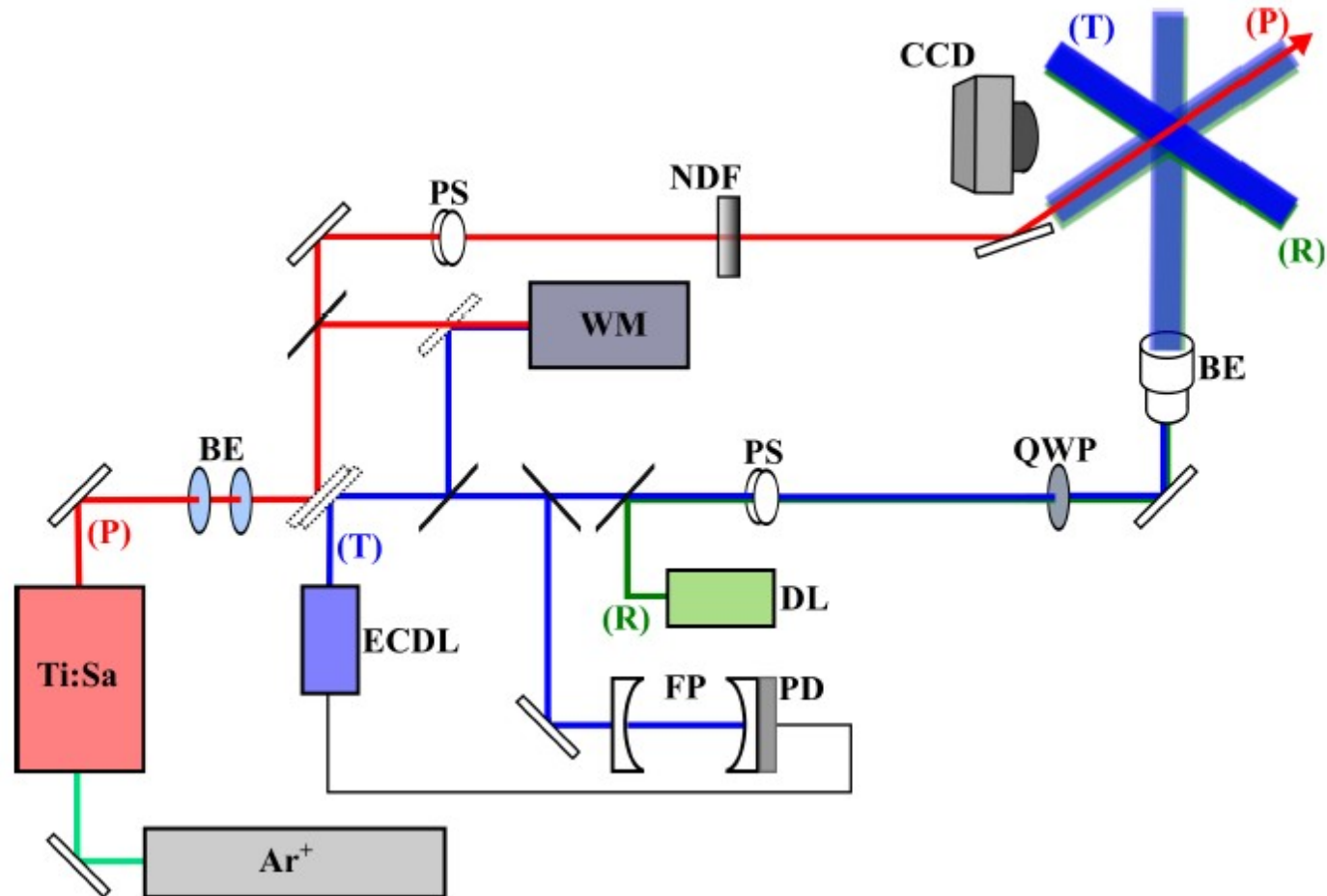
Detection of lines by change in trapped atom number

Application of Light Induced Atom Desorption

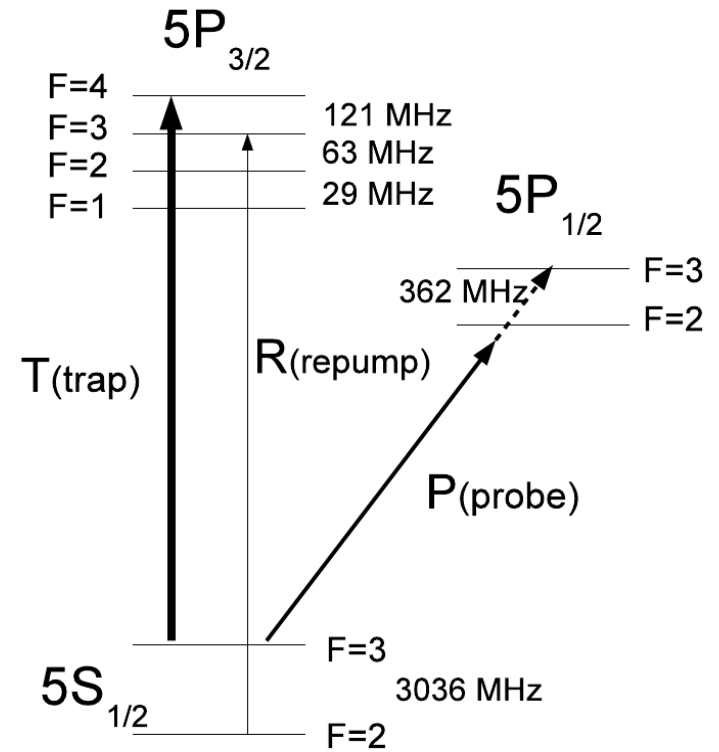
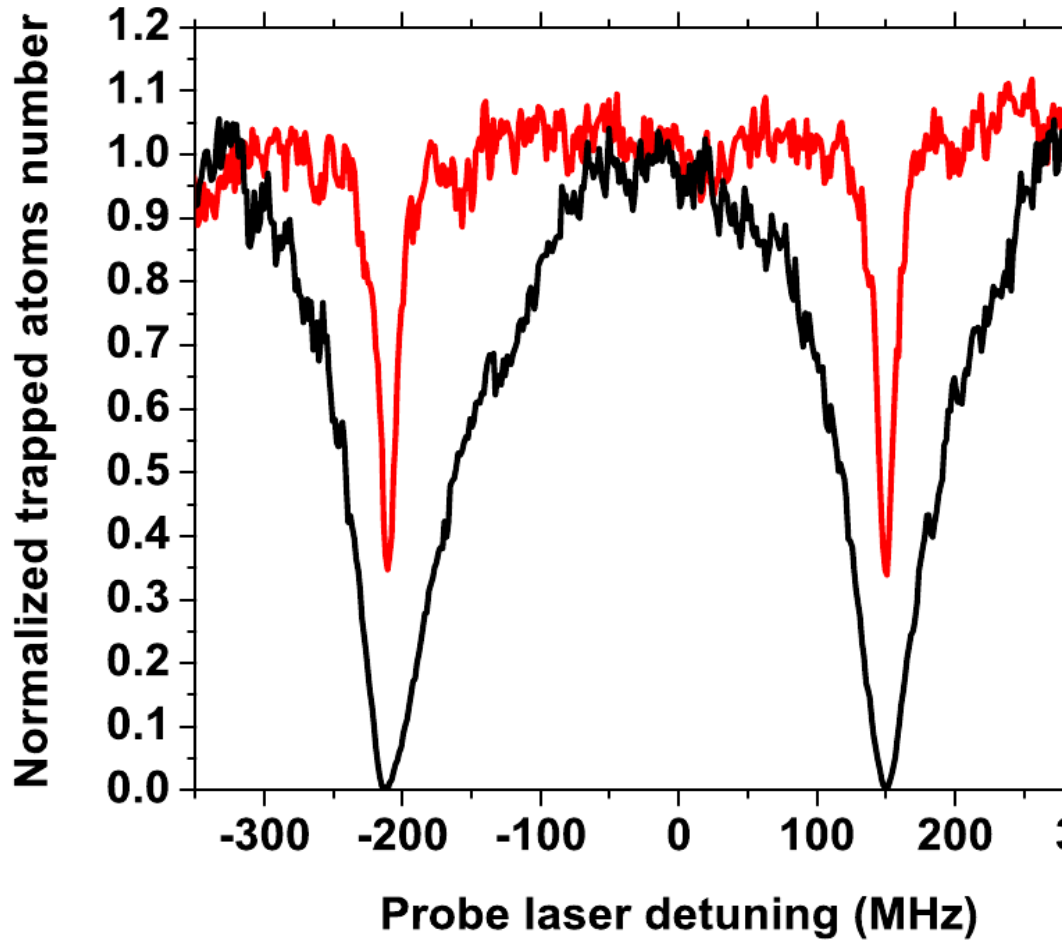


Perspectives

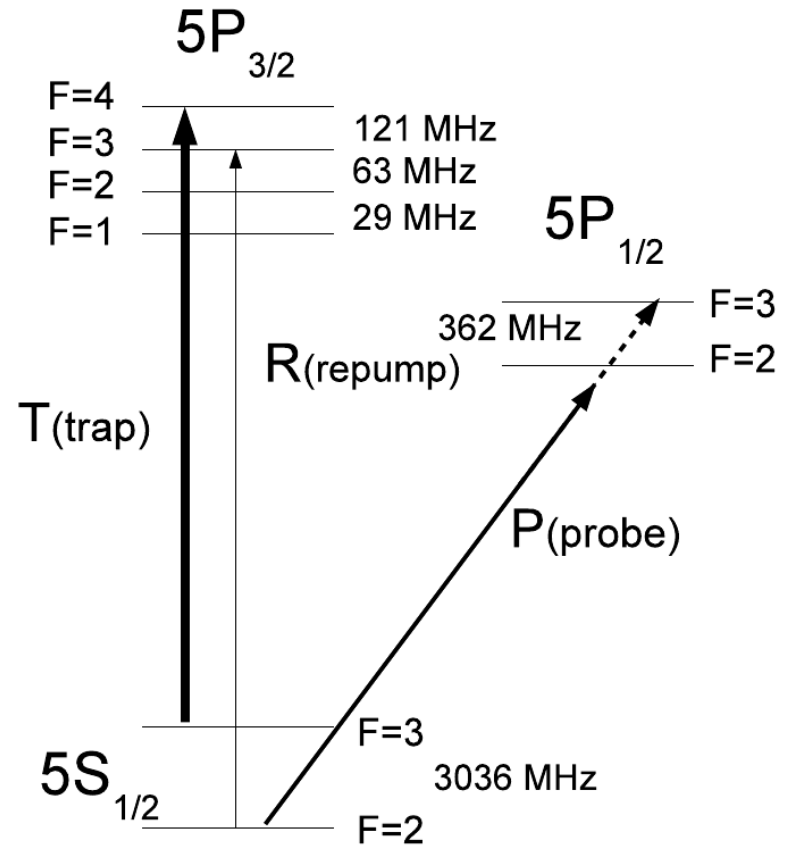
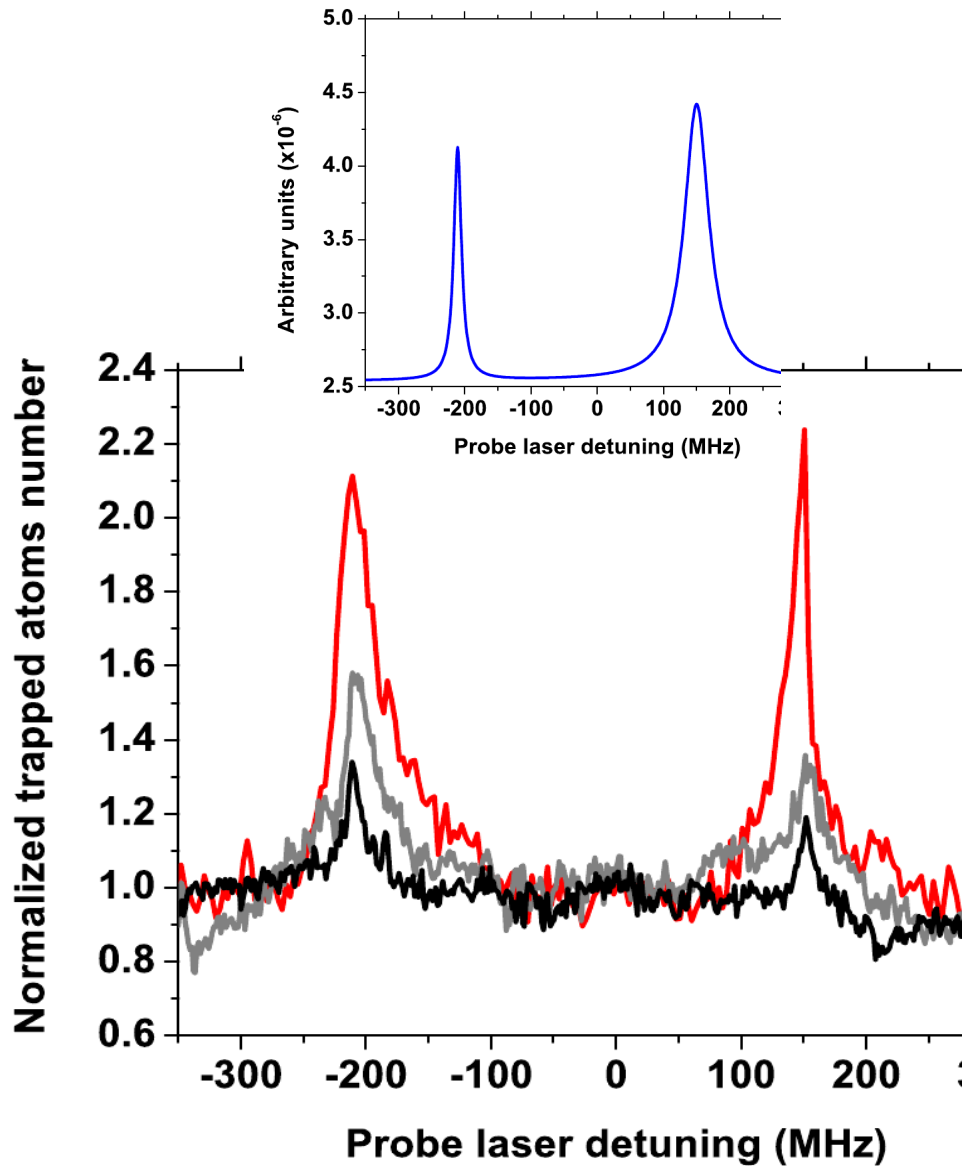
Detection setup (Rubidium)



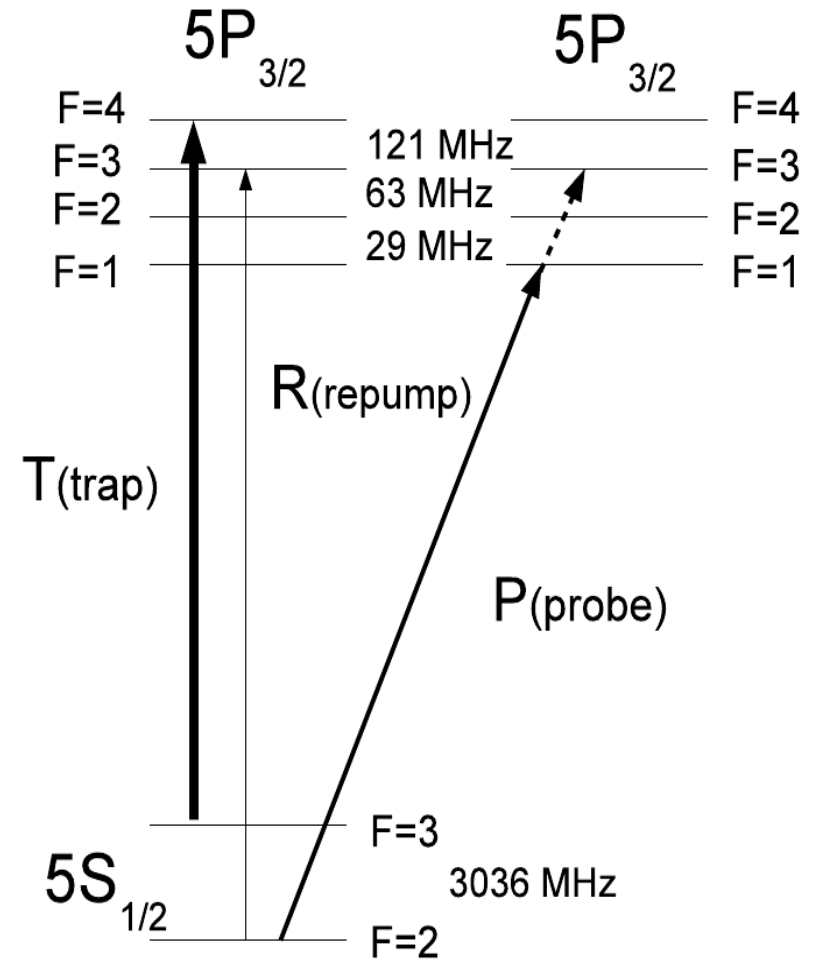
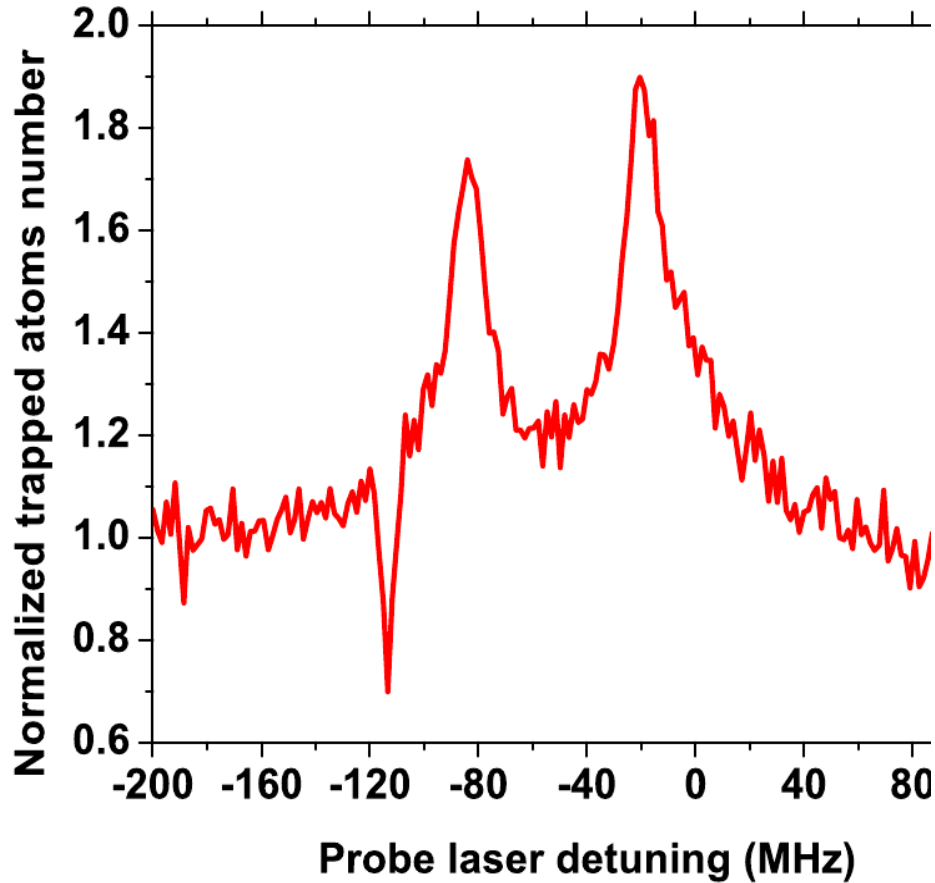
Detection results (Rb)



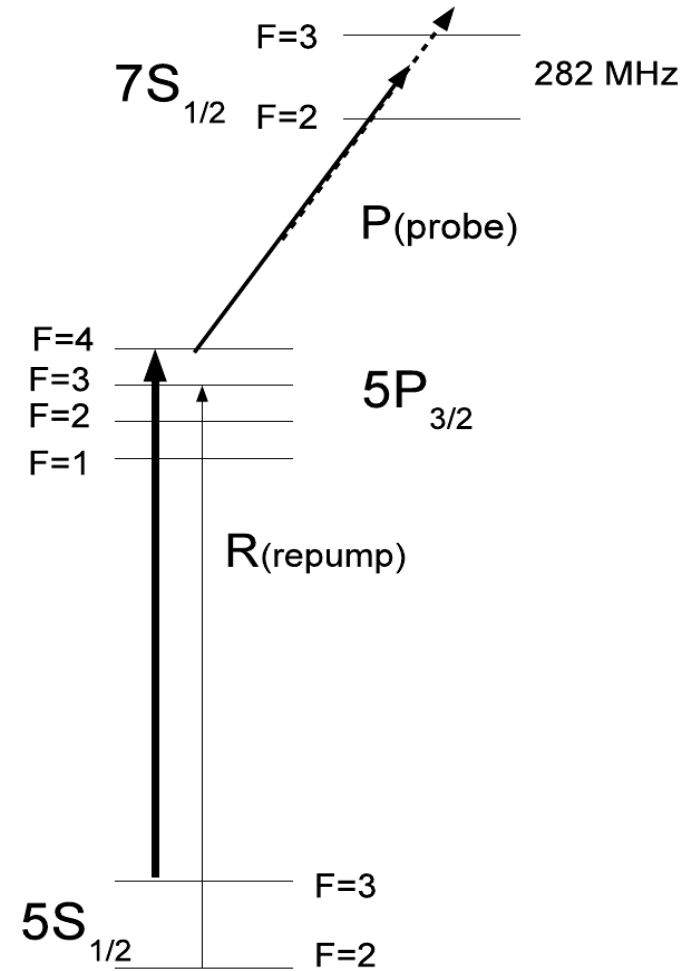
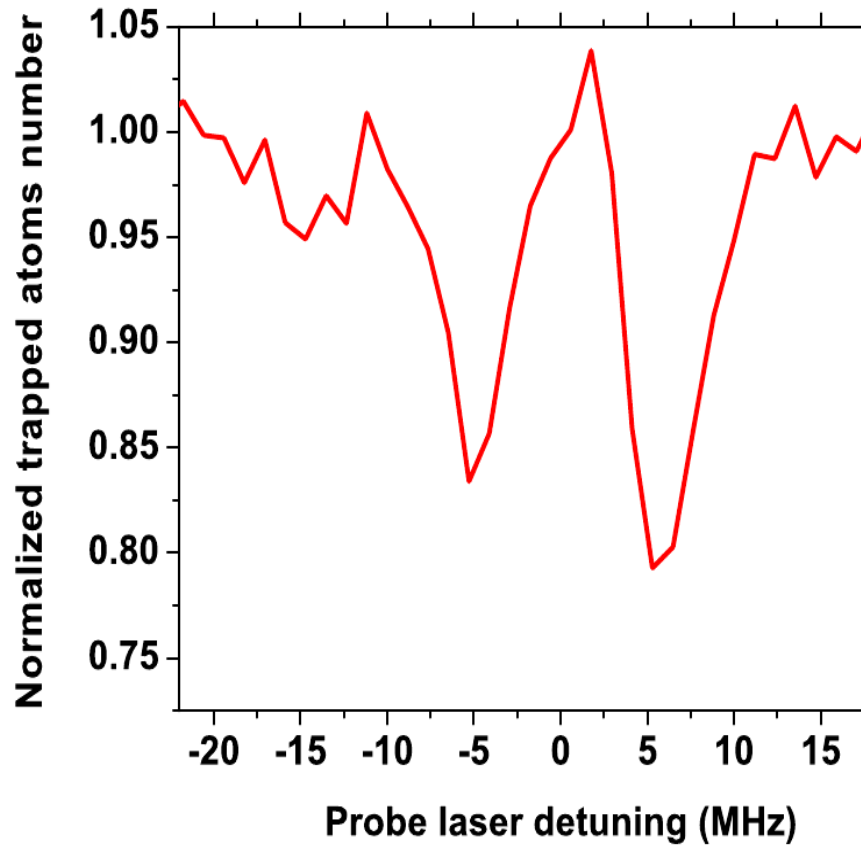
Detection results (Rb)



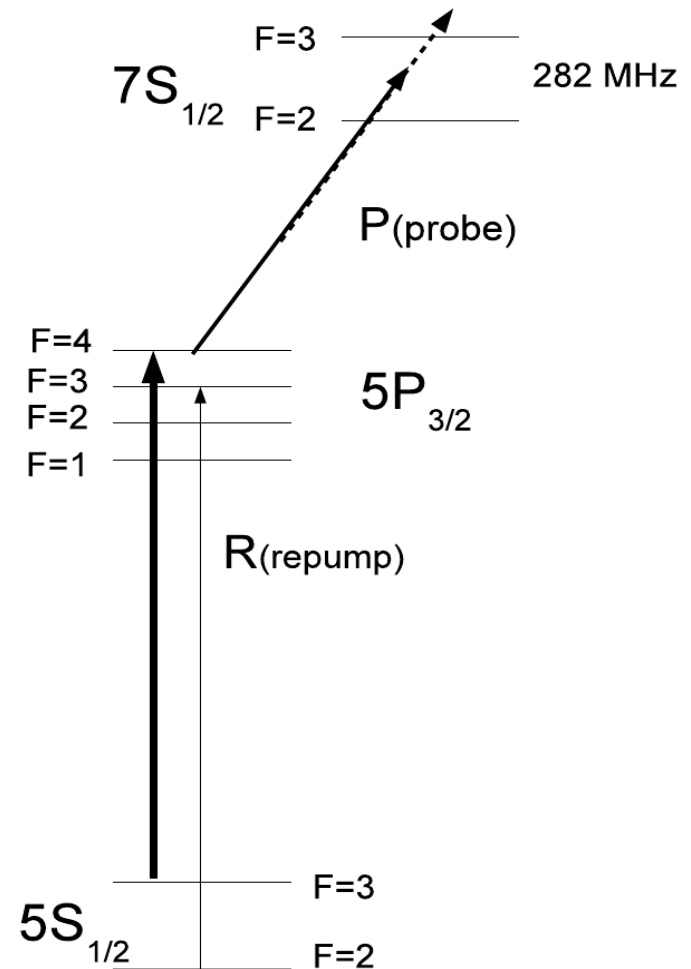
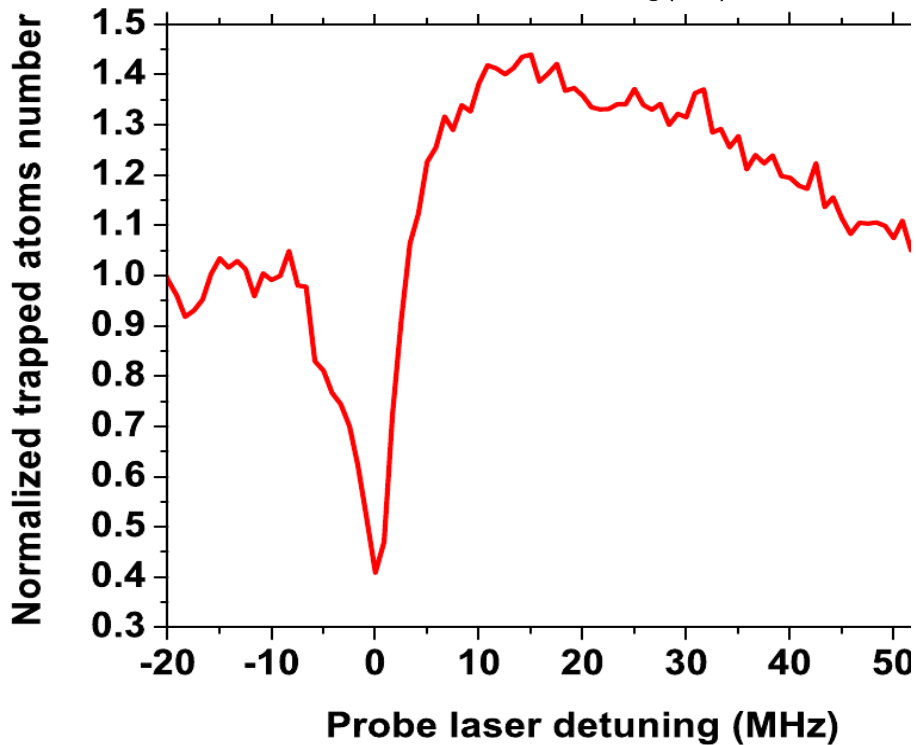
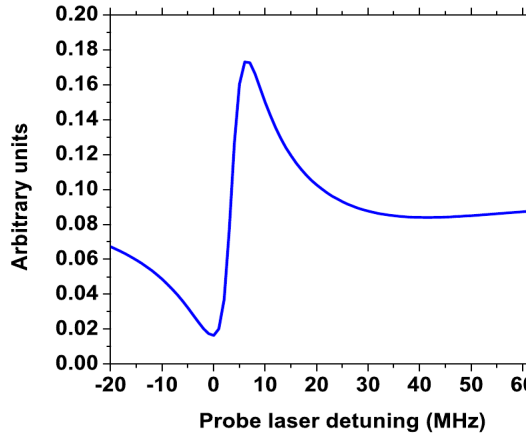
Detection results (Rb)



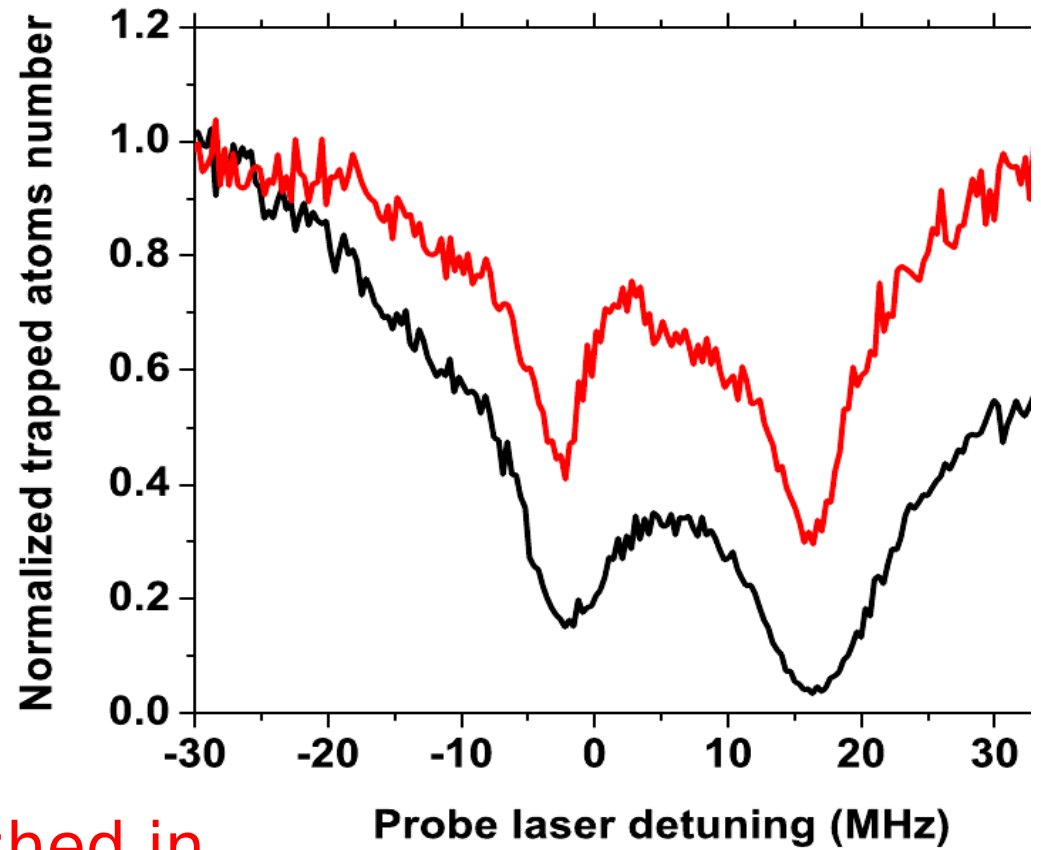
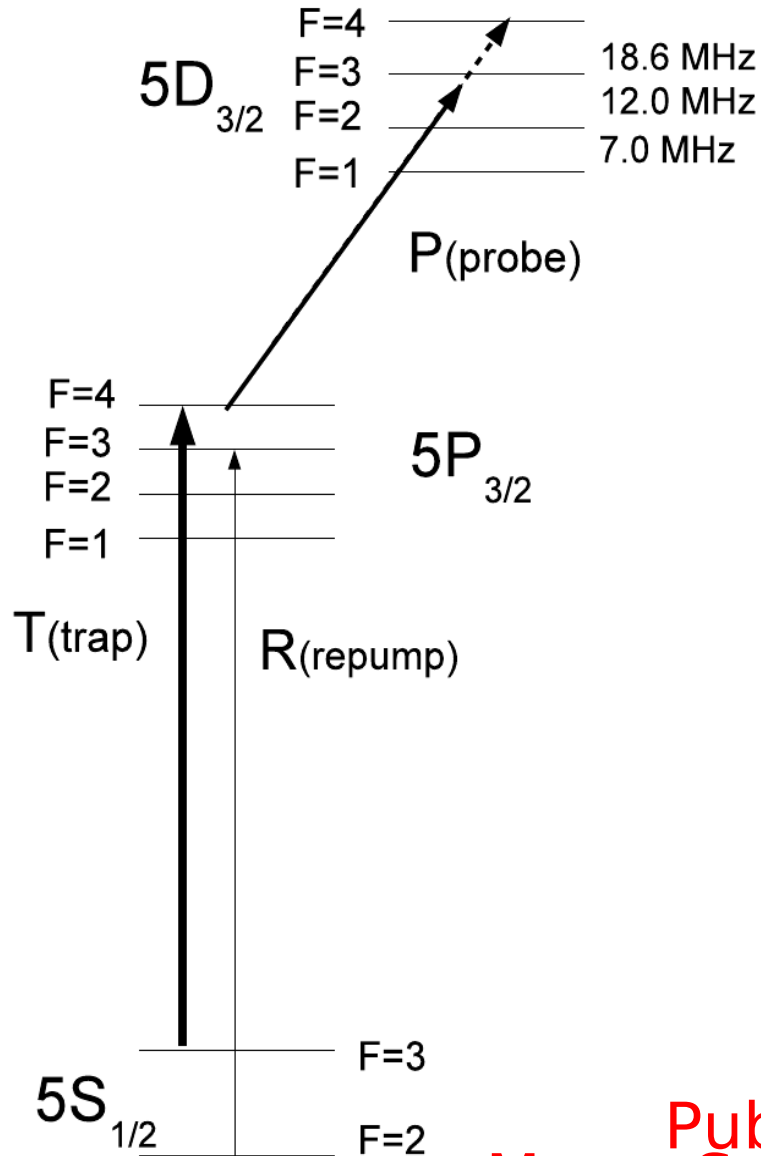
Detection results (Rb)



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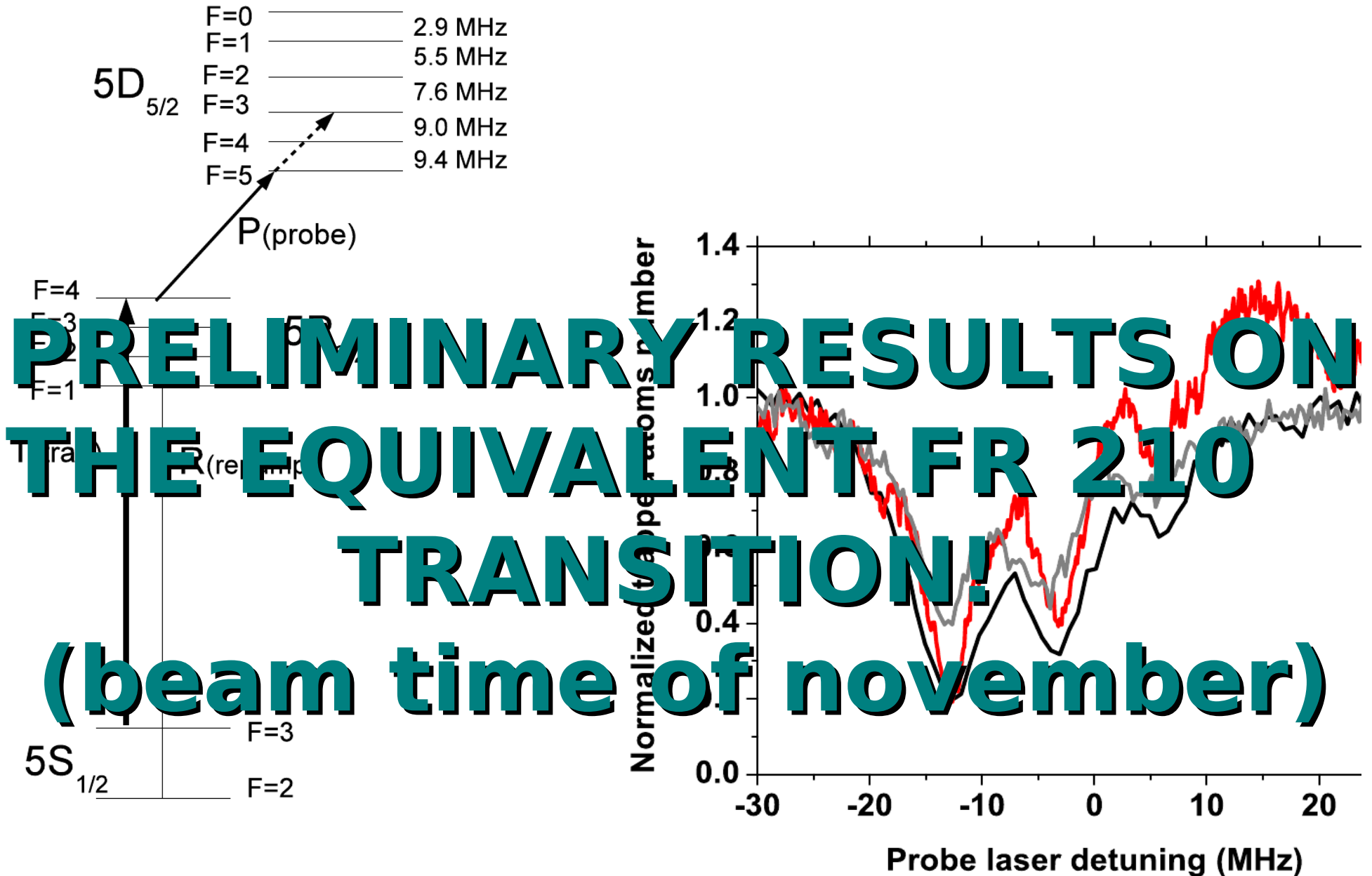


Detection results (Rb)

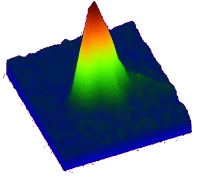


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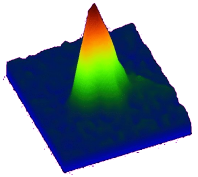
Detection results (Rb)



Outline



Introduction/Motivation (if any...)



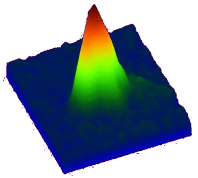
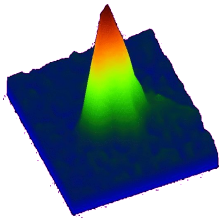
The LNL apparatus

Precision frequency measurements

Laser spectroscopy diffusion coefficients meas.

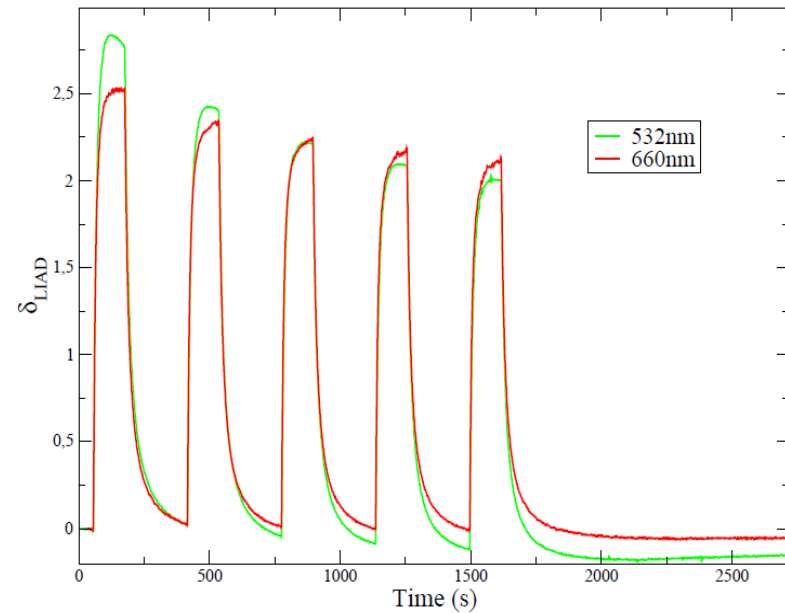
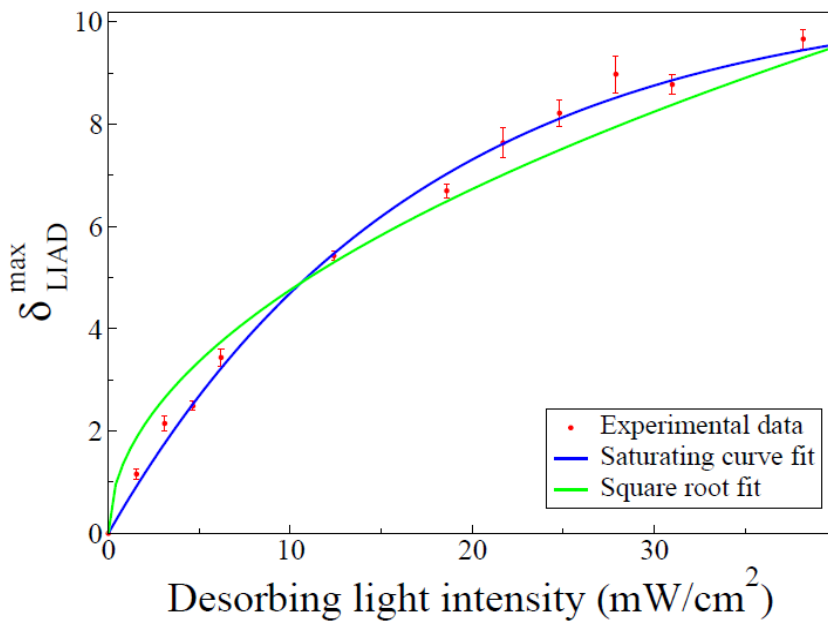
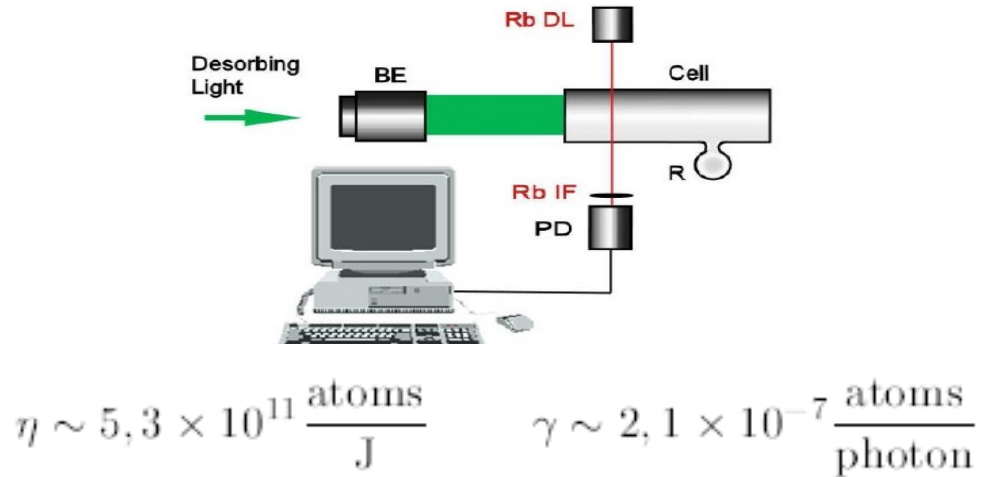
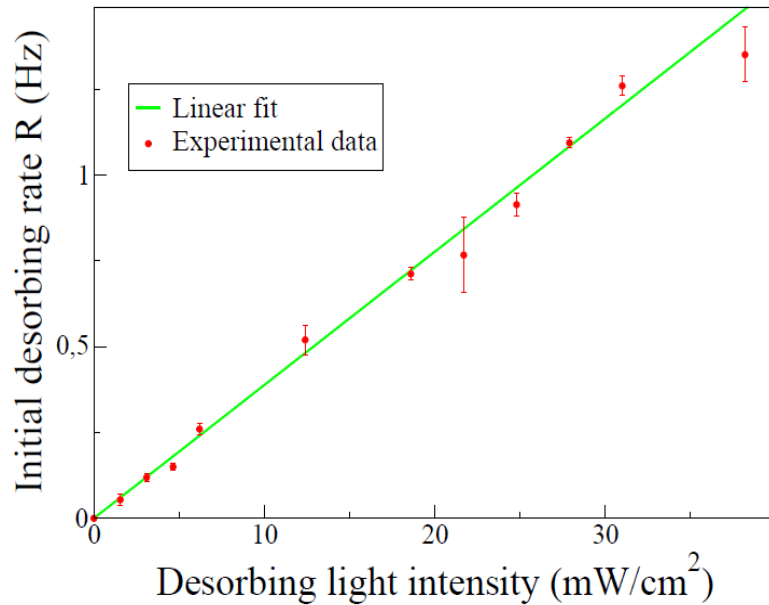
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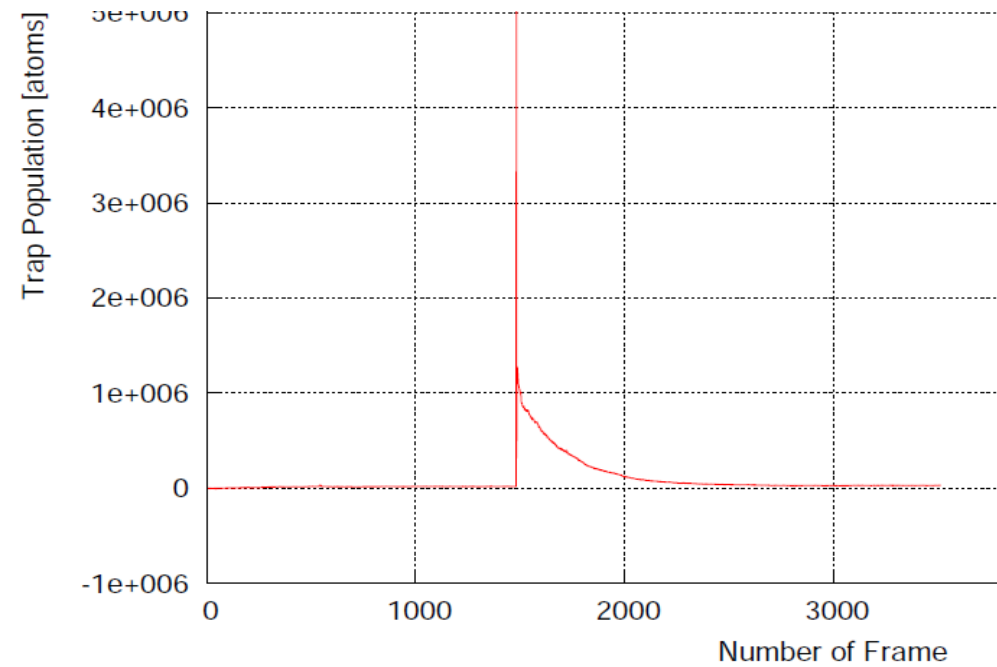
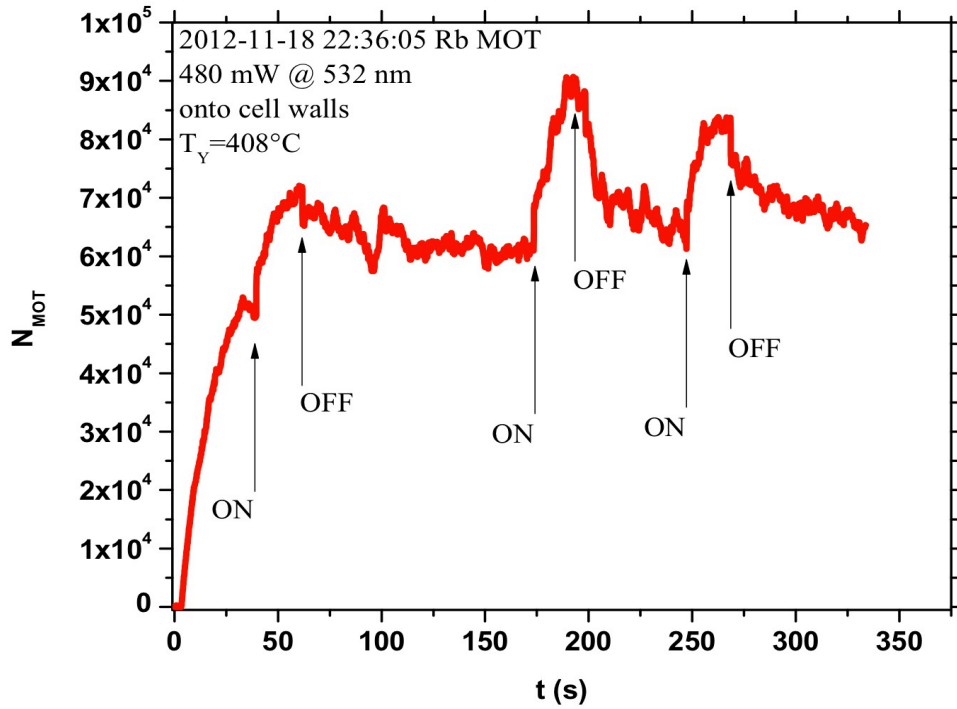


Perspectives

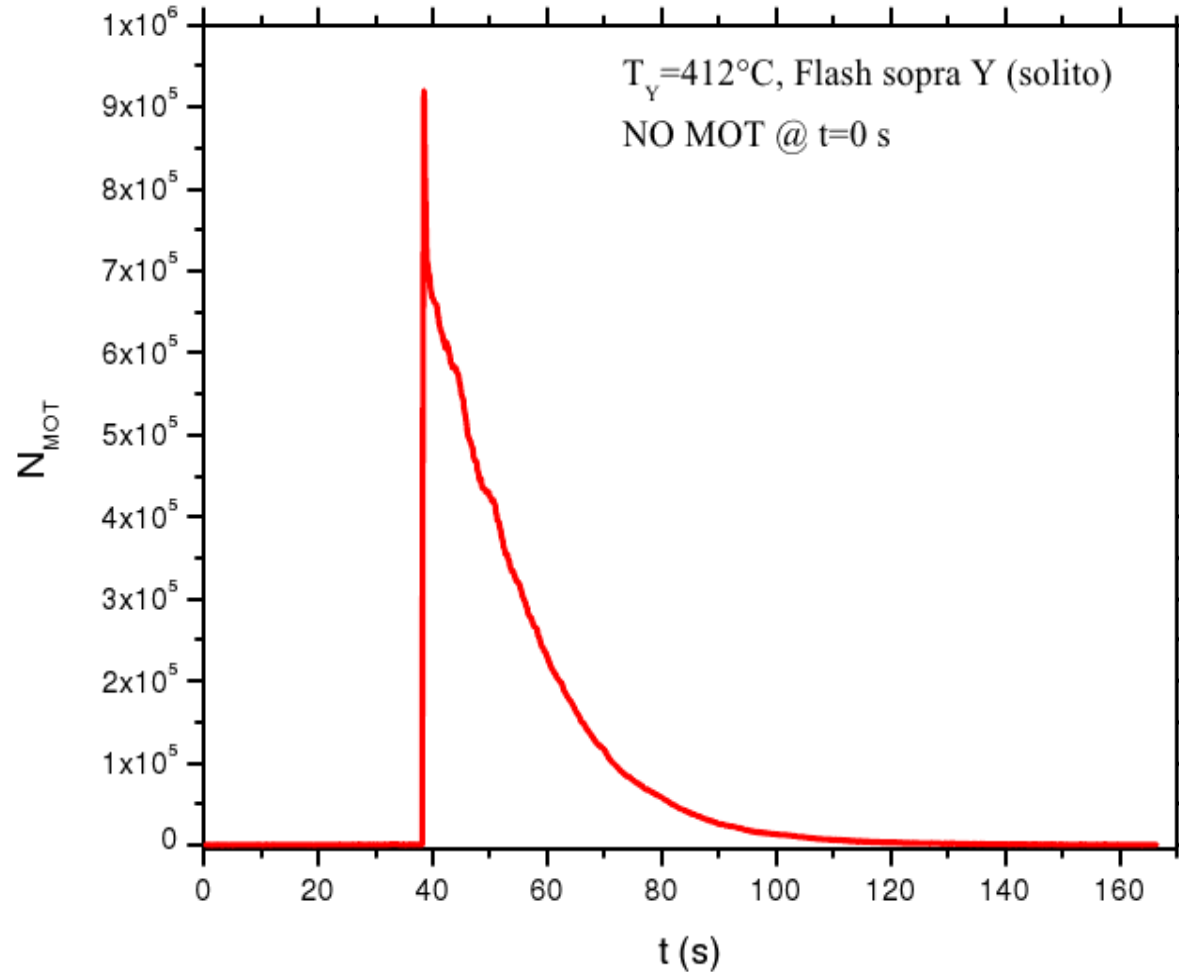
Rb desorption from OTS coated pyrex



Rb MOT loading from OTS

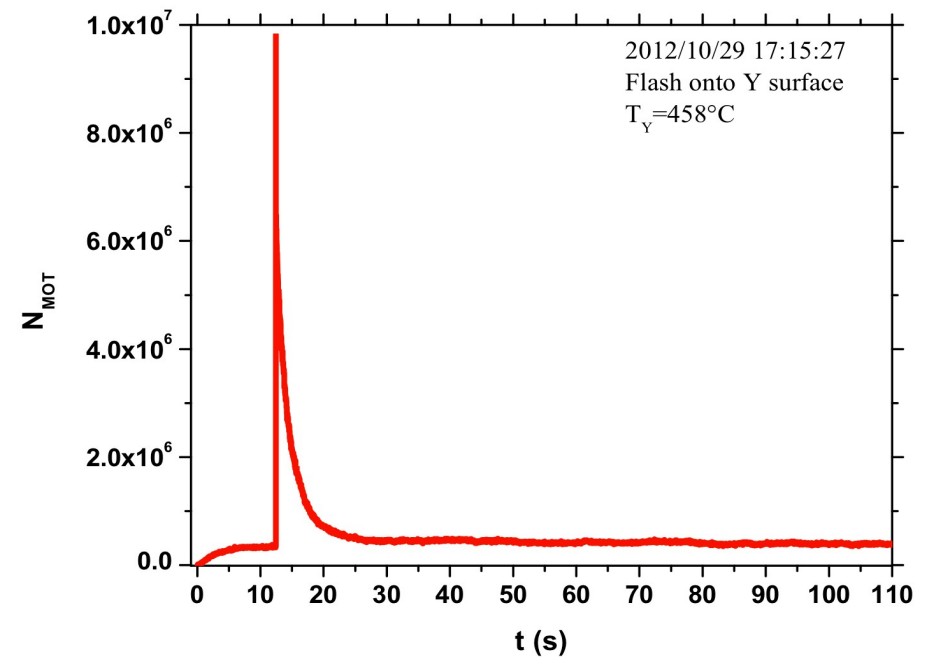
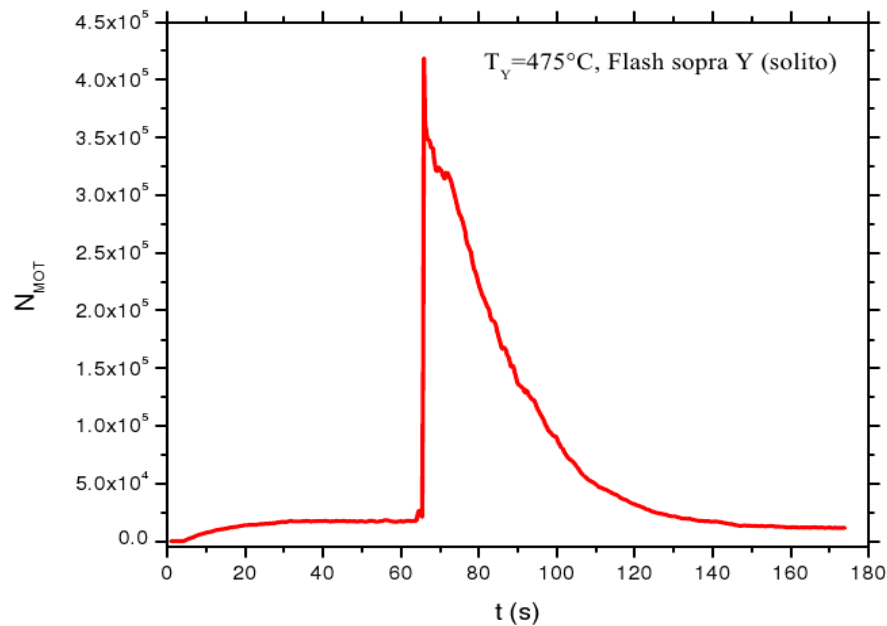


Rb MOT loading from Yttrium!



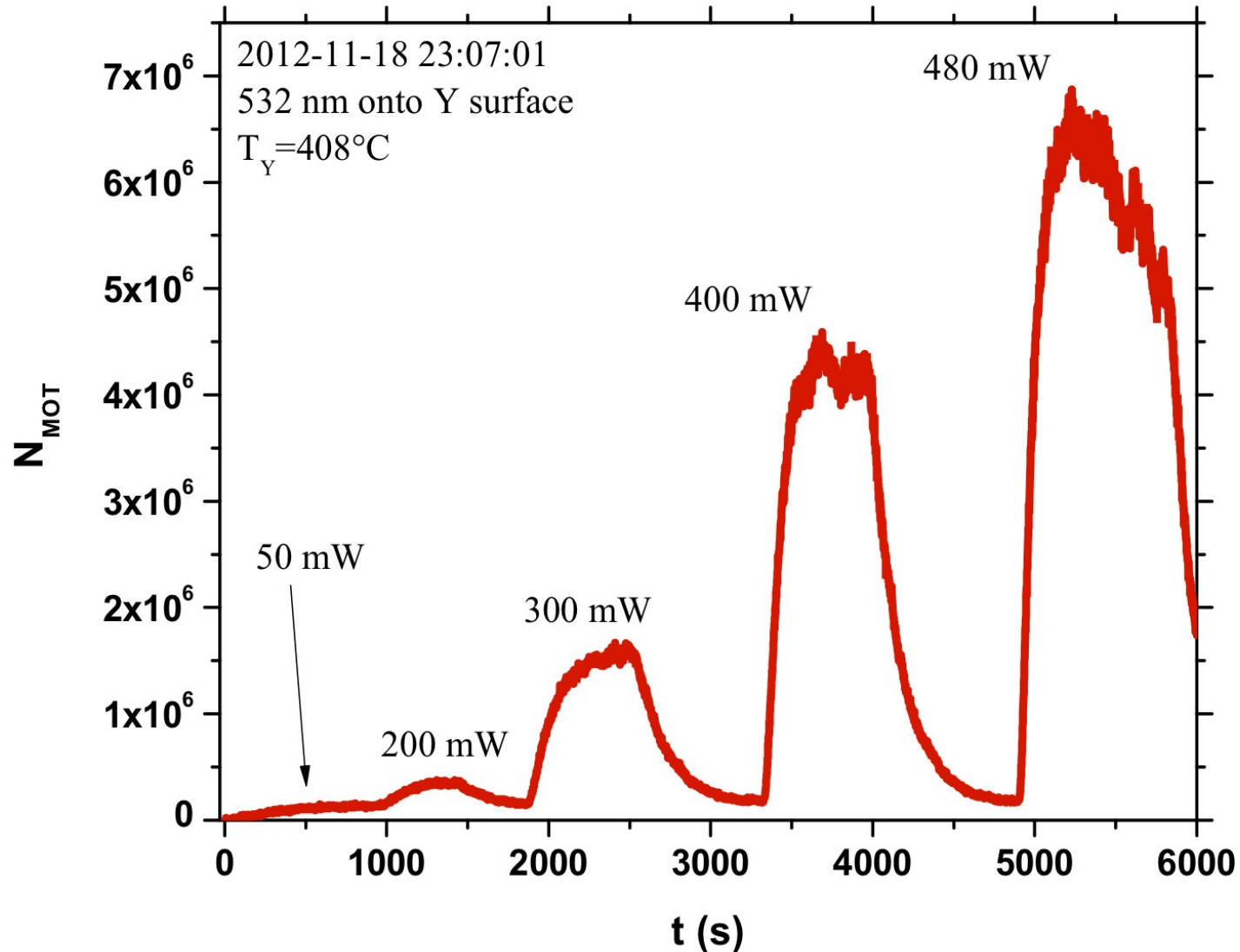
No MOT at the beginning
Very long restoring time

Rb MOT loading from Yttrium!



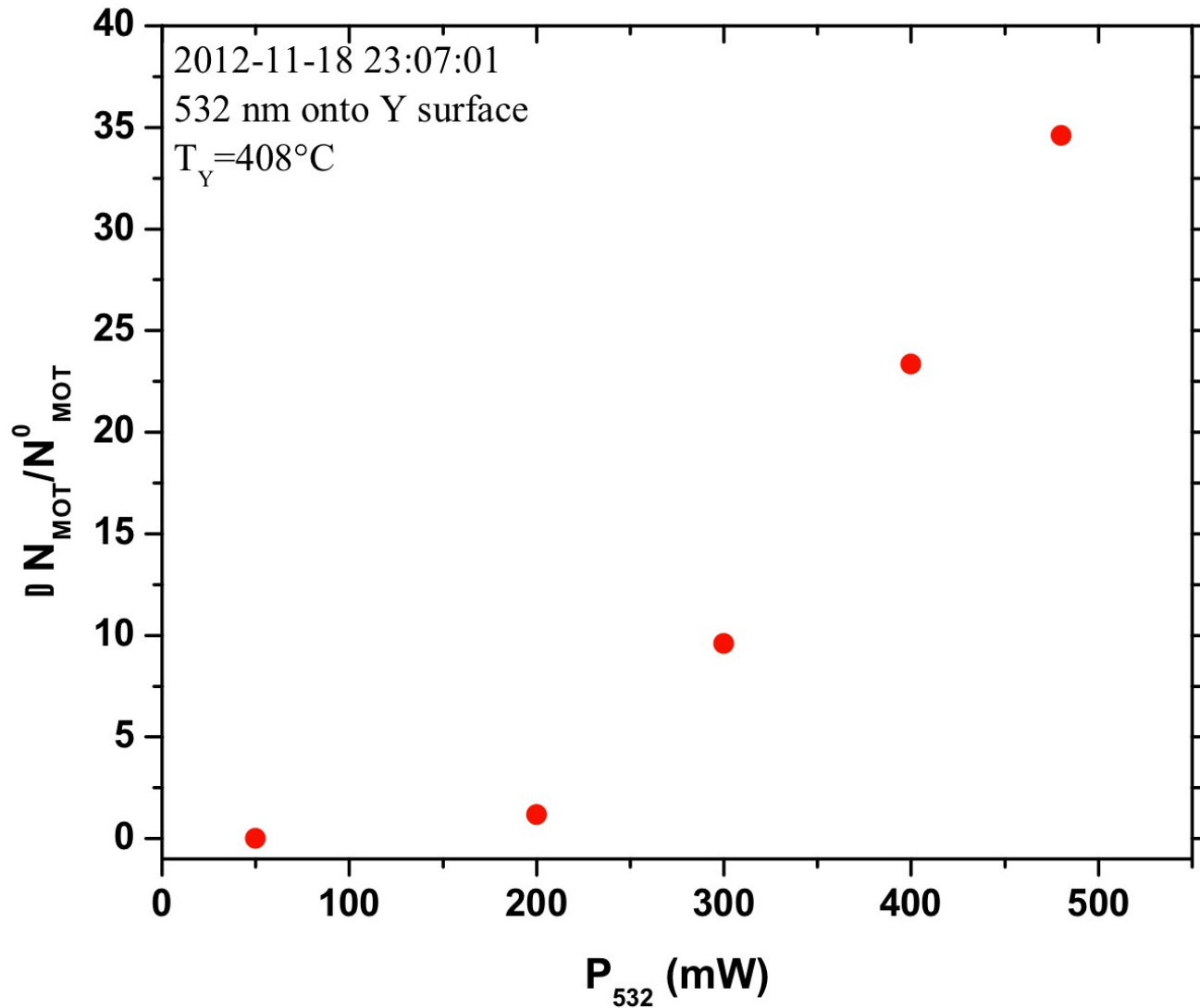
MOT at the beginning
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Rb MOT loading from Yttrium!



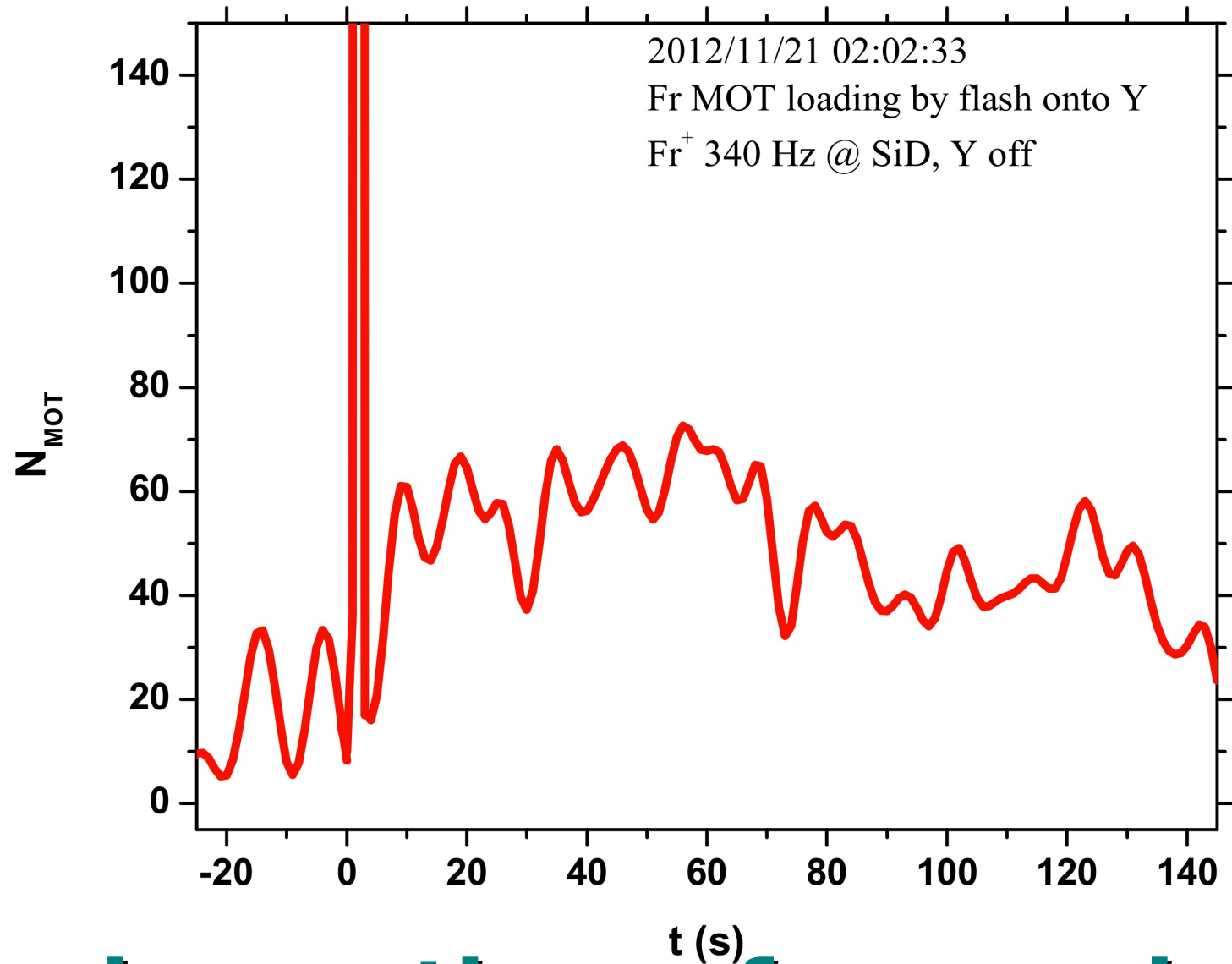
“Millennia” laser used as a desorption light

Rb MOT loading from Yttrium!



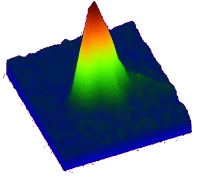
“Millennia” laser used as a desorption light

Fr LIAD MOT loading from Yttrium!!!

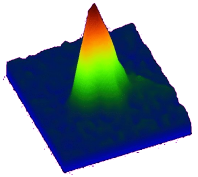


beam time of november

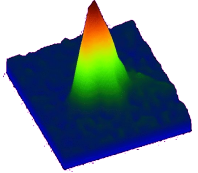
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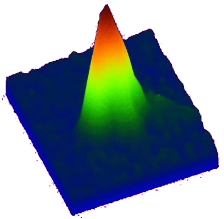


Precision frequency measurements

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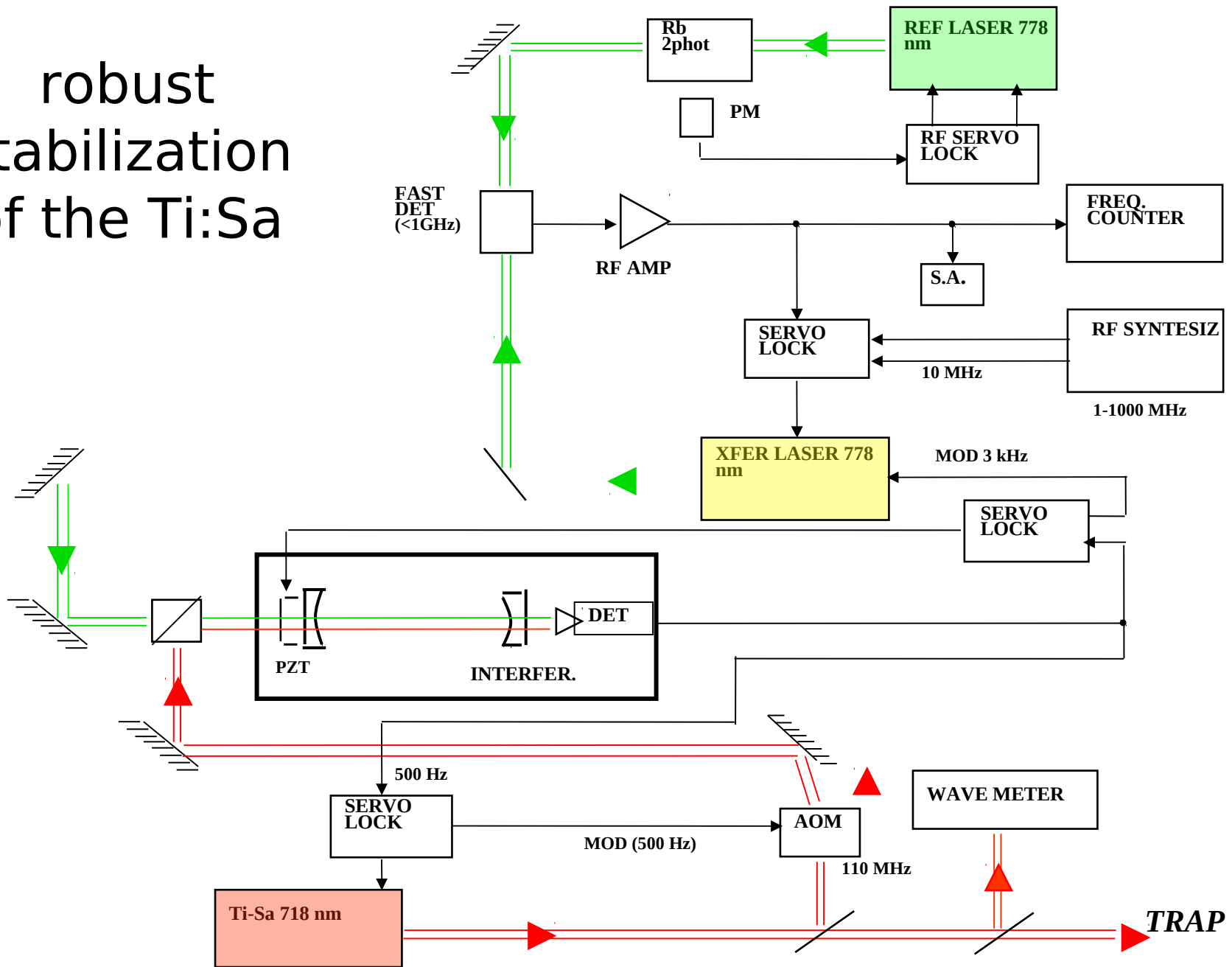
Perspectives

New spectroscopic measurements

- **ENERGY LEVEL DETERMINATION**
- **LIFETIMES MEASUREMENTS**
- **COLLISIONAL STUDIES**
- **DIMER FORMATION (??)**
- **RELATIVISTIC EFFECTS CONTROLS**

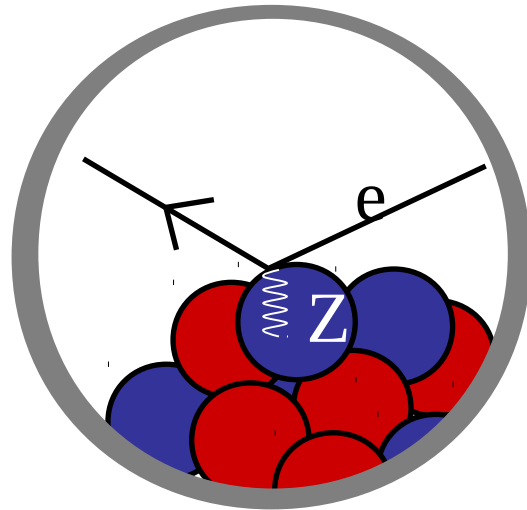
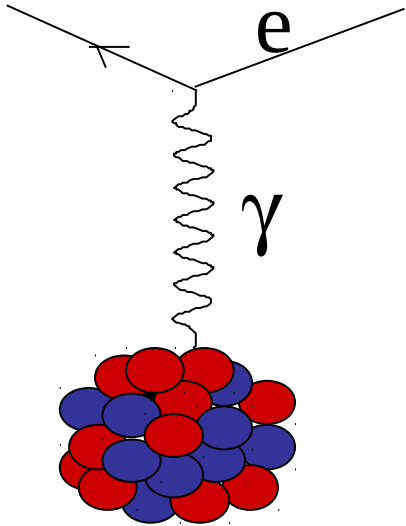
New spectroscopic measurements

robust
stabilization
of the Ti:Sa



Atomic parity violation

RELEVANT ELECTRON - HADRON PROCESSES



$$A_{\text{em}} \propto \frac{e^2}{p^2}$$
$$A_{\text{W}} \propto \frac{e^2}{p^2 + M_{Z_0}^2 c^2}$$

p is the momentum transfer
(inversely proportional to the Bohr radius)

$$p \sim m_e \alpha c$$

Atomic parity violation

Different transition probabilities for two mirror - image experiments

The amplitude A_W contains a part that is odd under space reflection and gives rise to a left - right asymmetry A_{LR} by interference with A_{em} .

$$P_{L/R} = |A_{em} \pm A_W^{odd}|^2$$

$$A_{LR} = \frac{P_L - P_R}{P_L + P_R} \simeq 2\text{Re} \frac{A_W^{odd}}{A_{em}}$$

$$\alpha^2 \left(\frac{m_{e^-}}{M_{Z_0}} \right)^2 10^{-15}$$

Atomic parity violation

Completely hopeless? No!

There are 2 factors of enhancement:

A. The so - called Z^3 law

- For valence electrons belonging to penetrating orbitals, the orbitals are deformed in the vicinity of the nucleus, where electrons “see” a Coulomb potential generated by a charge Ze . The orbital radius is given by a_0/Z , in such a way that p^2 is enhanced by Z^2 .
- The various nucleons add for their contributions coherently: the number of nucleons grows as Z

Atomic parity violation

B. The second source comes from the possibility of exciting highly forbidden transitions like $nS_{1/2} \rightarrow (n+1)S_{1/2}$ in alkalis. The electromagnetic selection rules strictly forbid the electric dipole transition; dipole magnetic transitions M_1 are allowed by the symmetry, not by the change in radial number.

$$M_1 \approx 4 \times 10^{-5} \frac{\mu_B}{c}$$

The weak interaction associated with the boson exchange breaks this rule and gives rise to a parity violating electric dipole amplitude $E^{(PV)}_1$:

$$E_1^{PV} \approx 10^{-11} ea_0$$

$$\text{Im } E_1^{PV} / M_1 \simeq 0.5 \times 10^{-4}$$

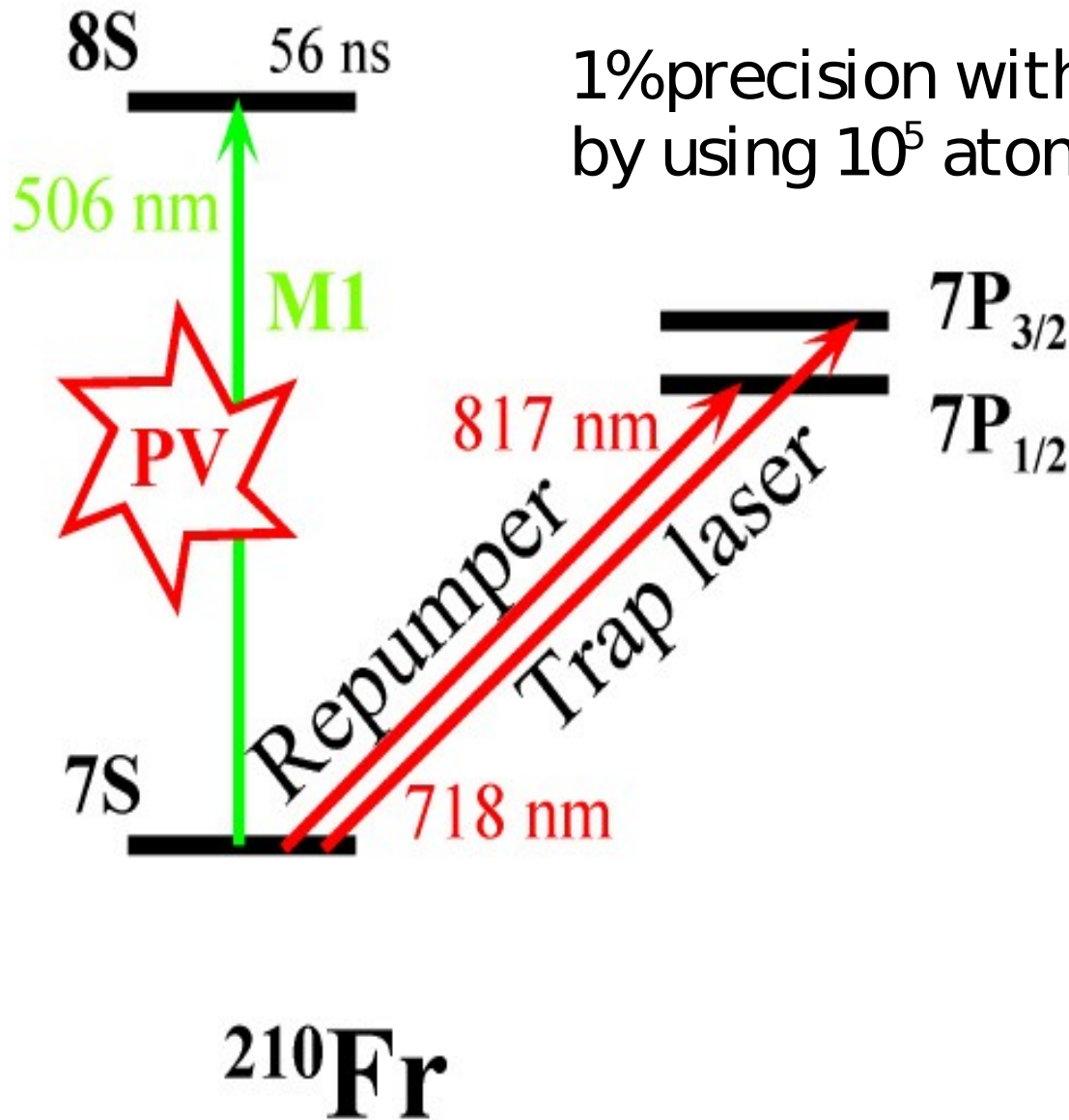
Atomic parity violation

A possible experimental approach:

1. Capture Fr atoms in a MOT
2. Accumulate and cool in the MOT
3. Transfer to a second trap (purely optical)
4. Establish a “coordinate system” by dc electric field, dc magnetic field, k vector of the exciting laser
5. Excite $7S$ to $8S$ using a build up cavity and detect using the $7S$ to $7P$ transition.
6. Reverse the coordinate axis.
7. Change isotope.

Towards APV measurement

$$\vec{d}_{F,F'}^{\text{eff}} = -\alpha\vec{E} - i\beta\vec{\sigma} \wedge \vec{E} + M_1'\vec{\sigma} \wedge \vec{k} - i\text{Im}(E_1^{PV})\vec{\sigma}$$



1% precision with 3 weeks measurements
by using 10^5 atoms

preliminary measure
of the ratios
 α/β , β/M_1 , $M_1/M_{1^{\text{hf}}}$:
in the MOT cloud
to “calibrate” APV

Expected signal to noise ratio

- ⇒ Fr production rate in Legnaro: up to 10^6 ions/s.
- ⇒ Trapping efficiency $\sim 10^{-2} \Rightarrow N = 10000$ atoms in 1 mm^3 (0.01 mm^3) (optical dipole trap).
- ⇒ **Laser intensity: 100 mW/mm^2** , enhanced by a factor $\zeta = 1000$ with a Fabry-Perot cavity (cf. Boulder) $\Rightarrow P/S = 10 \text{ kW/cm}^2$.
- ⇒ Fluorescence detection efficiency: $\eta \sim 10\%$.

$$\Rightarrow S/N = \frac{4\pi}{3\hbar c} \frac{1}{\hbar \Gamma} \frac{P}{S} \eta N \sqrt{t} = 0.009 \sqrt{t(\text{s})} \quad (1 \text{ for } t = 3 \text{ hours})$$

How can we improve S/N ?

- ⇒ Higher laser power, BUT: – heating due to photon scattering
– photoionization from $8S$ and $7P$.
- ⇒ Higher Fr^+ Rate: $\geq 4 \cdot 10^9$ ions/s at the ISOLDE facility.
 $\Rightarrow S/N = 0.55 \sqrt{t(\text{s})}$

⇒ **In 9 hours we can get $S/N = 100$**

Conclusions

