

**Исследование свойств ядер с помощью
лазерной резонансной фотоионизационной
спектроскопии в лазерном ионном источнике
на установке ISOLDE (CERN)**

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Ядерный семинар ОФВЭ

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Содержание (структура доклада?)

- Ядерный комплекс ISOLDE
- Лазерный ионный источник RILIS (ISOLDE)
- Фотоионизационная лазерная спектроскопия с помощью RILIS
 - Общее описание метода и экспериментальной установки
 - Развитие установки
- Применение фотоионизационной спектроскопии на установке ISOLDE
 - Поиск новых схем фотоионизации (Po, At) и измерение потенциала ионизации (At)
 - Разделение изомеров
 - Измерение изотопических сдвигов и сверхтонкой структуры атомных переходов (ядерные зарядовые радиусы и электромагнитные моменты)
- Перспективы

Exploring

exotic nuclei at ISOLDE

Production

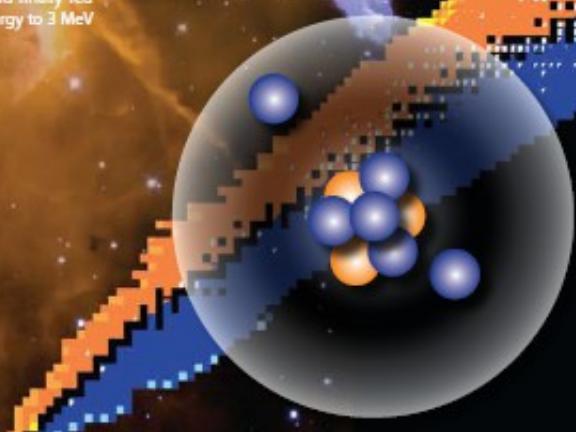
ISOLDE produces radioactive nuclei in reactions between protons at 1.4 GeV energy and nuclei in a variety of special targets. Several different types of reaction can take place, making a broad range of elements available. The targets are heated so that the new radioactive species diffuse out quickly before they decay. Scientists and engineers at ISOLDE have worked for decades to develop the best materials and designs for the targets.

Selection

To produce a beam of a chosen exotic nucleus requires not only the right choice of target material, but also methods to extract the nuclei as ions (with fewer electrons than atoms) and to separate them electromagnetically from other species. ISOLDE has pioneered a very selective ionization technique that uses several wavelengths of laser light simultaneously to pick out specific elements. ISOLDE can deliver more than 700 different beams of isotopes from 70 chemical elements.

Acceleration

To make the *fires* of the nuclei produced at ISOLDE, the REX ISOLDE system provides an acceleration stage. Here the nuclei are trapped, bunched, stripped of additional electrons, selected according to mass, and finally fed into a linear accelerator to boost their energy to 3 MeV per nucleon.



Nuclear mass surface

Ions that are confined almost at rest in devices called Penning traps can have their masses measured with very high precision. The large variety of nuclear species available at ISOLDE allows a comprehensive survey of the "nuclear mass surface" – in effect a map of the many nuclear masses. This gives important input for studies of fundamental symmetries, theoretical models of the atomic nucleus, and nuclear astrophysics.

Fundamental symmetries

The nuclei produced at ISOLDE, with proton-to-neutron numbers varying over a wide range, provide an interesting microscopic laboratory for low-energy tests of the Standard Model of elementary particle physics. The high quality of the beams allows high-precision measurements of beta decay, particle correlations and atomic masses.

Nuclear astrophysics

One of the most fundamental and challenging questions of the 21st century is how the elements from iron to uranium were created. Nuclear reactions occurring in explosive stellar environments, such as novae, supernovae and X-ray bursters, are believed to play an important role in the synthesis of these heavier elements. The pathways of the reactions leading to them involve short-lived radioactive exotic nuclei, which can be studied at ISOLDE and REX ISOLDE.

Sizes and shapes

Nuclei come in a variety of sizes and shapes, from spherical to deformed shapes, which can be "prolate" (cigar-shaped) or "oblate" (like a discus). Experiments at ISOLDE can investigate the transitions between extremes, for example, the development of a neutron-halo structure in lithium-11, which makes this nucleus with only 11 nucleons (neutrons and protons) as big as a lead nucleus with 208 nucleons.



Excited states

Nuclei are governed by the laws of quantum mechanics and exhibit "excited states" with well-defined energies and other properties predicted by theory. Radioactive decays and nuclear collisions can leave nuclei in excited states that decay to the ground state by emitting gamma rays. These can be detected by advanced germanium detectors cooled to liquid nitrogen temperature, as in the MINIBALL array. The properties of the gamma-rays (energy and angle) provide information on the excited states, which can be used to test theories.

ISOLDE – An Isotope Factory

Of the 3100 known isotopes, 256 are stable.

Only 83 radioactive isotopes are found in nature.

Up to now

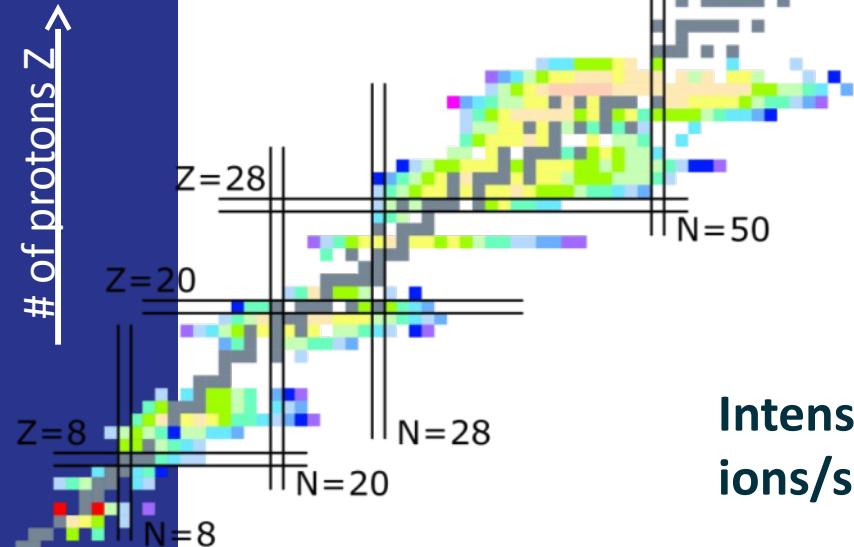
>800 isotopes of 70 elements have been produced at ISOLDE

Isotope range:

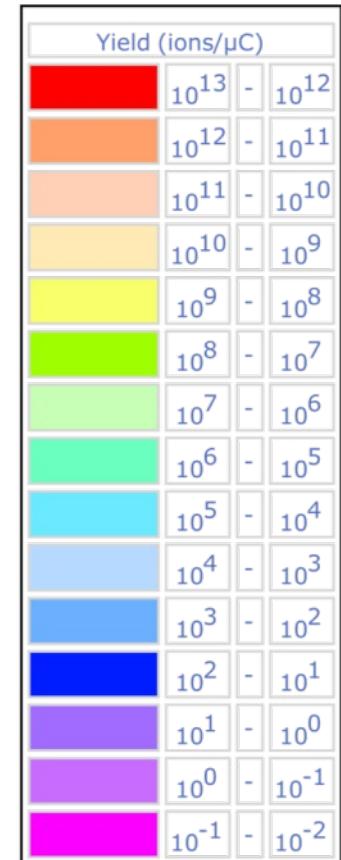
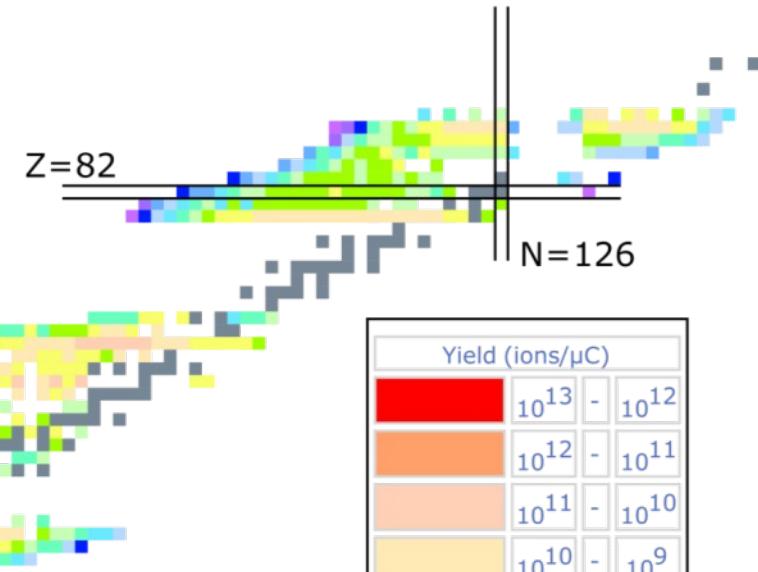
6He to 232Ra

Half lifes:

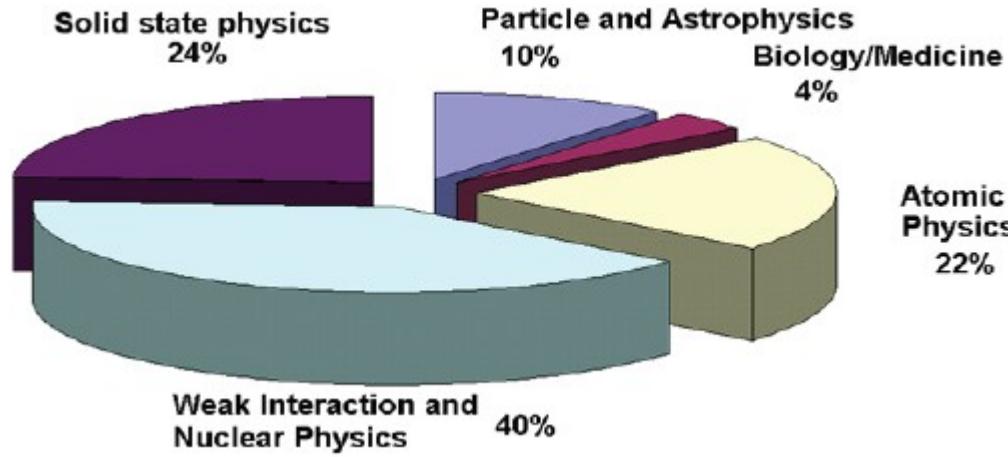
10 ms to stable



Intensity 10¹¹ down to 1 ions/s

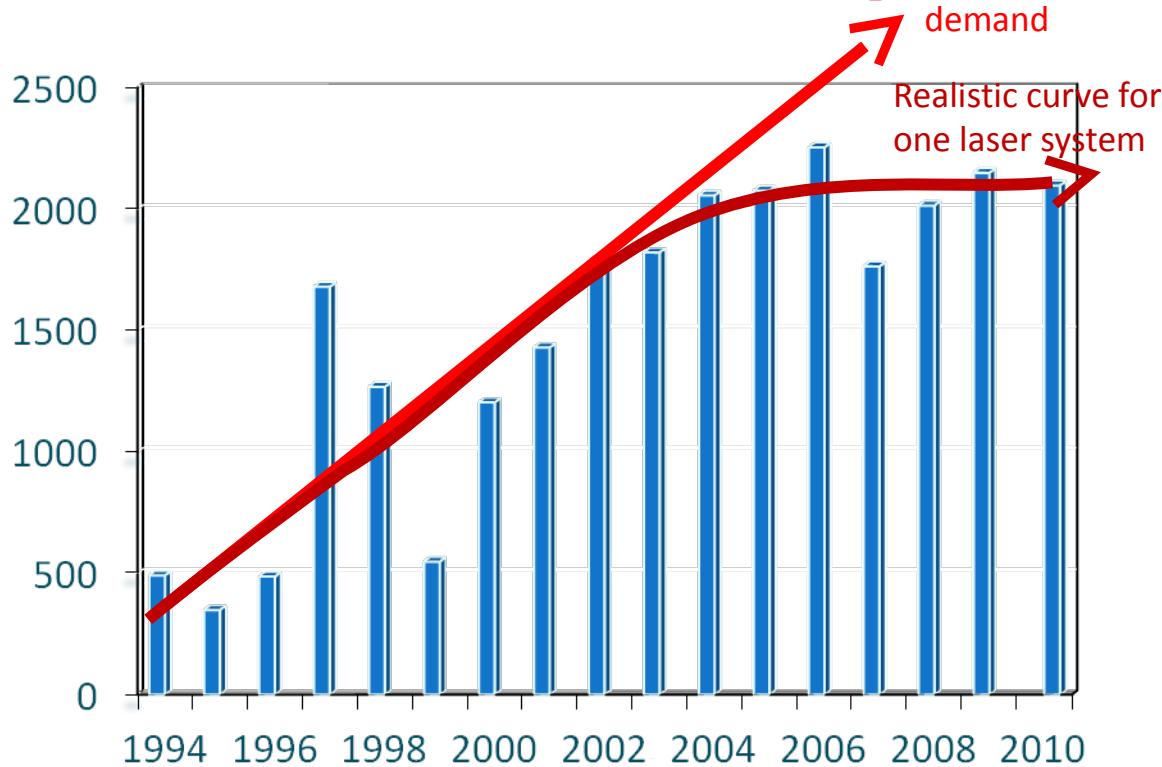


ISOLDE physics program



RILIS operation

Ion beams of 15 elements were produced



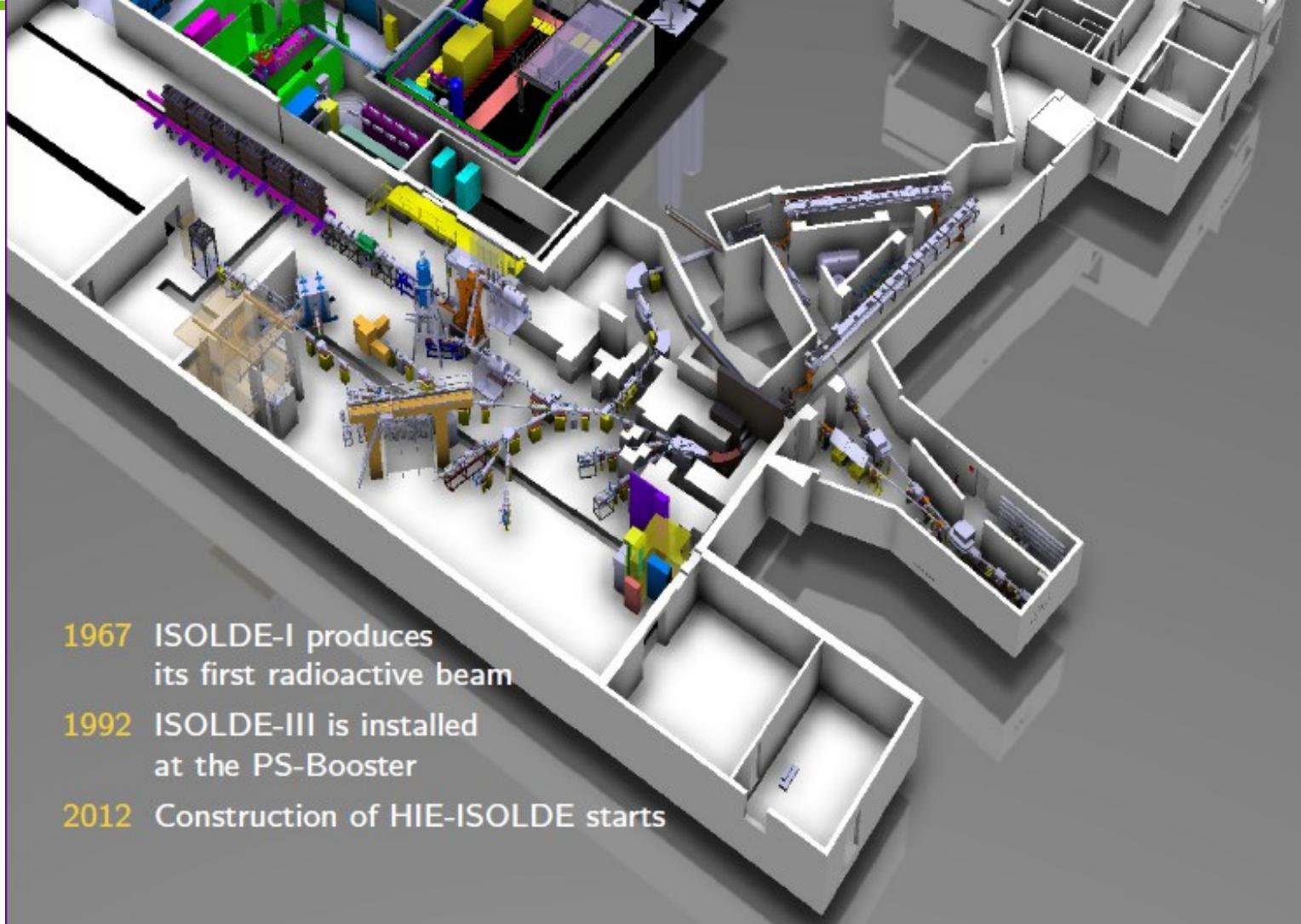
Laser ON time
in 2011:

2756 h – Total

2527 h – On-line

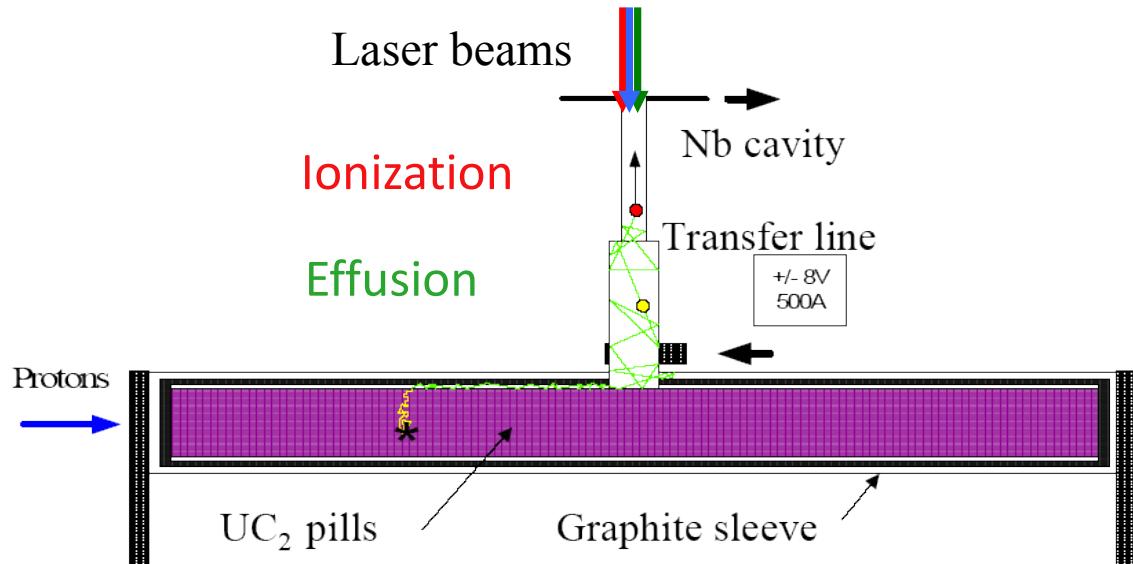
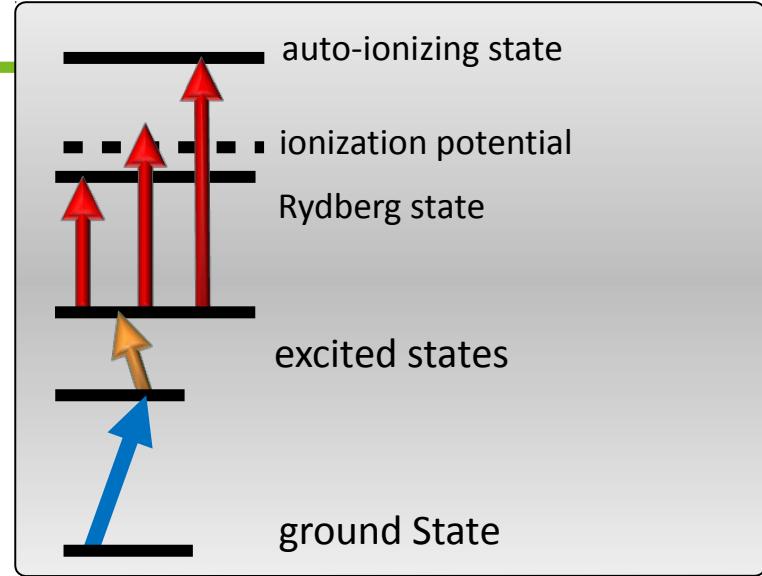
> 50 % of the
Total running time of
ISOLDE facility

Beam	Sm	Ga	Mg	At	Pb	Dy	Nd	Pr	Ag	Tl	Cd	Mn	Ni	Yb	Zn
Planned	80	104	256	160	184	64	136	-	112	208	288	120	272	-	376
Real	89	101	146	269	221	35	110	4	92	231	242	390	348	73	407

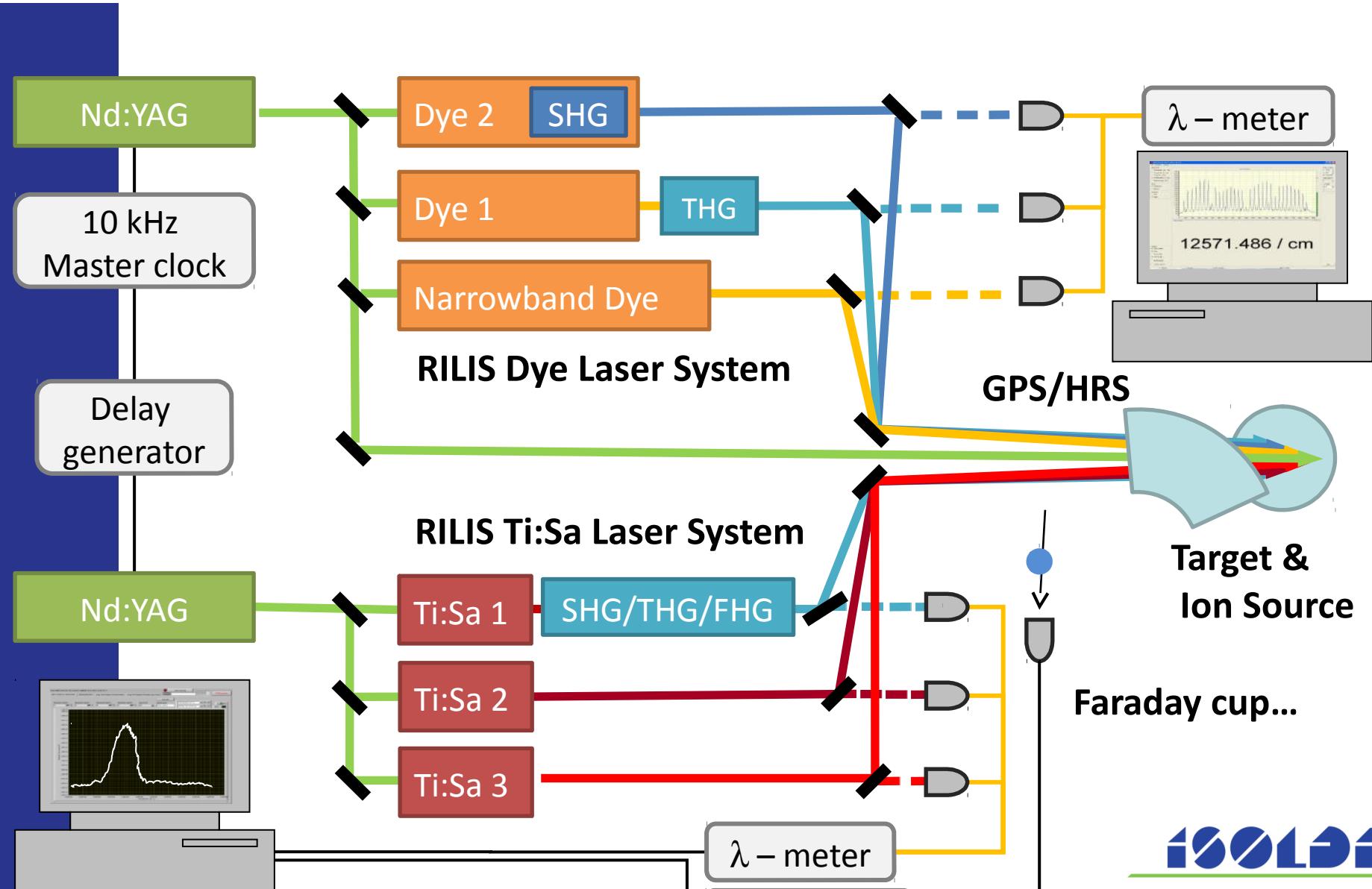


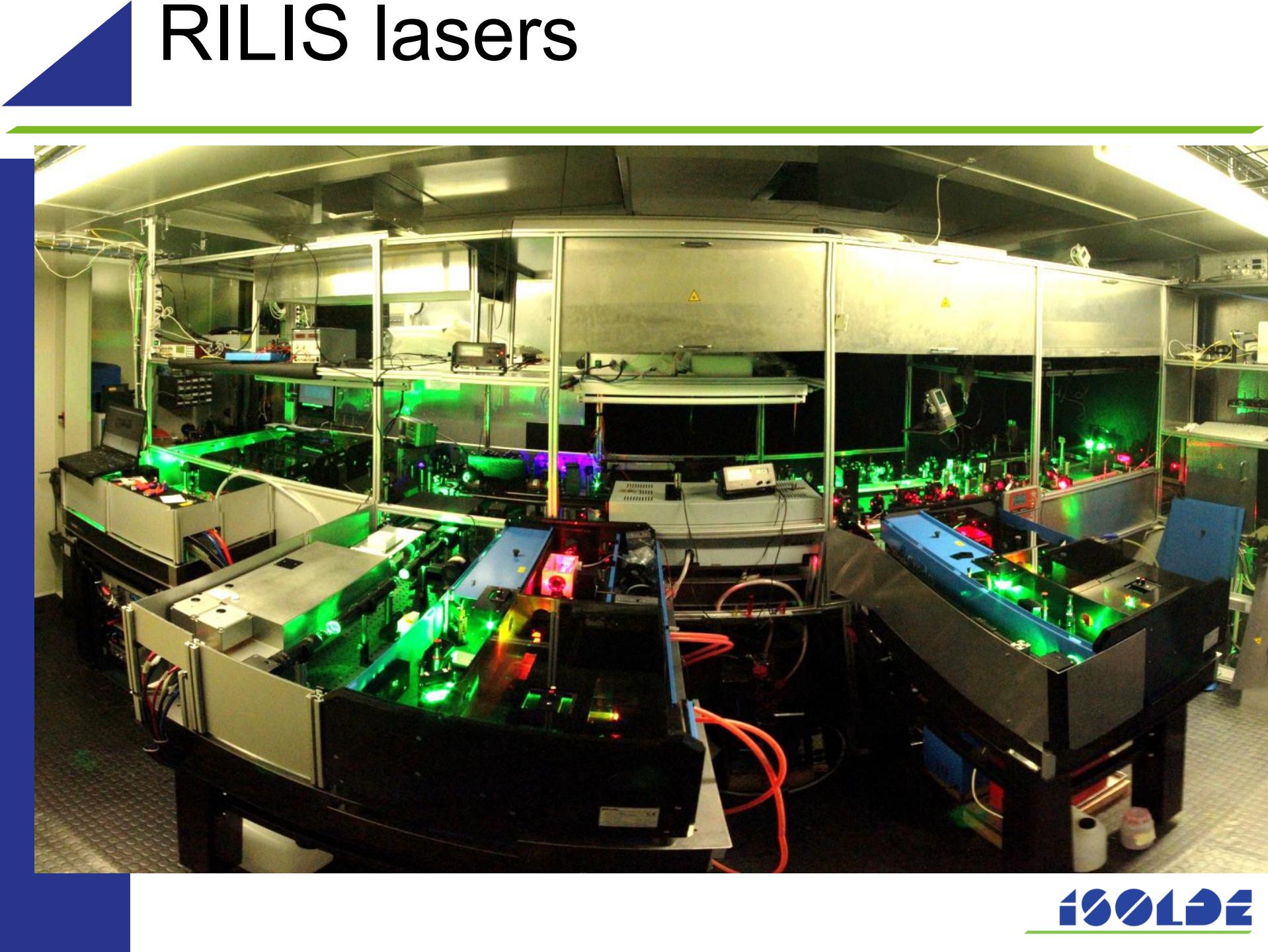
- 1967 ISOLDE-I produces its first radioactive beam
- 1992 ISOLDE-III is installed at the PS-Booster
- 2012 Construction of HIE-ISOLDE starts

Target –Ion Source unit



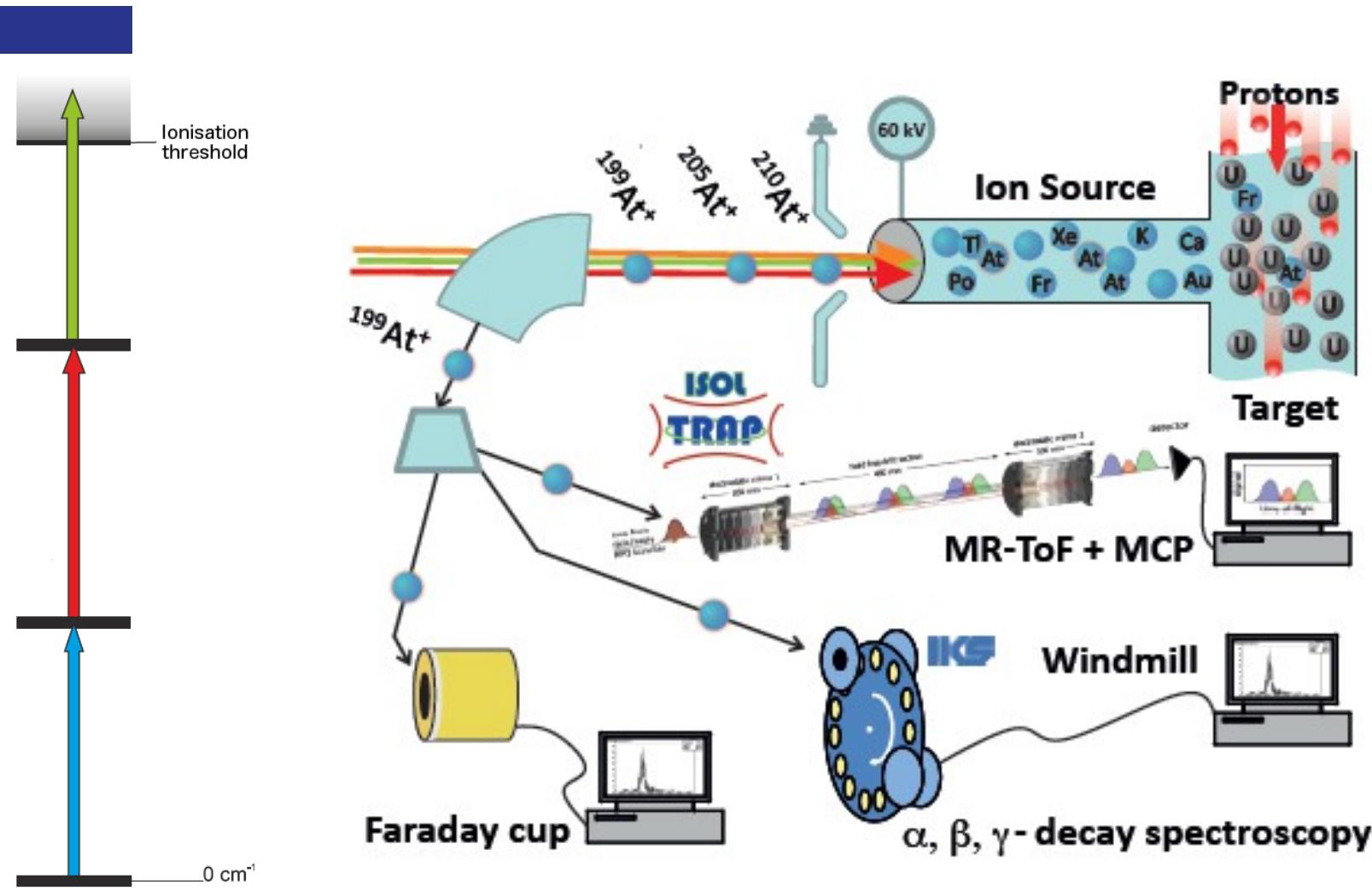
RILIS laser system

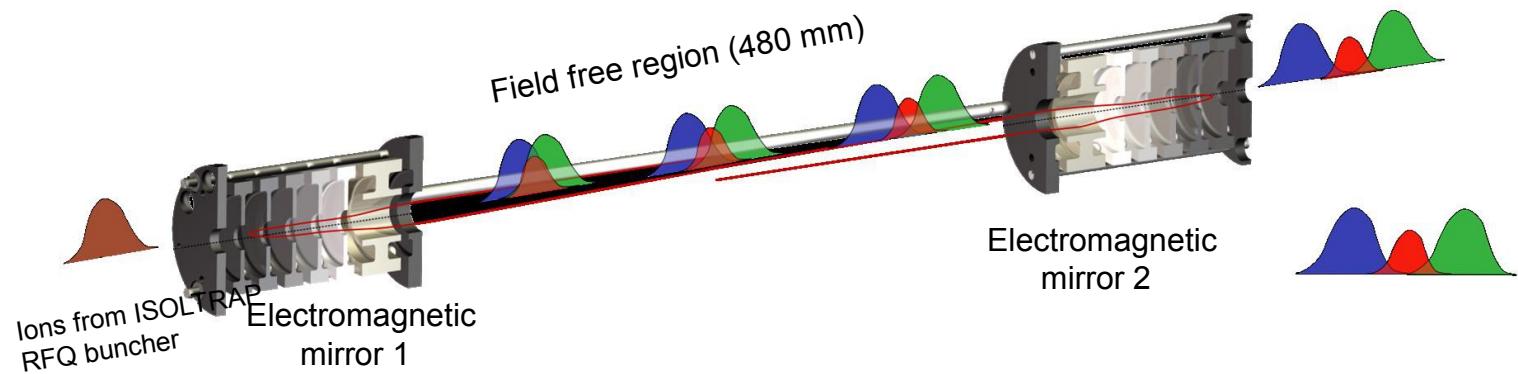
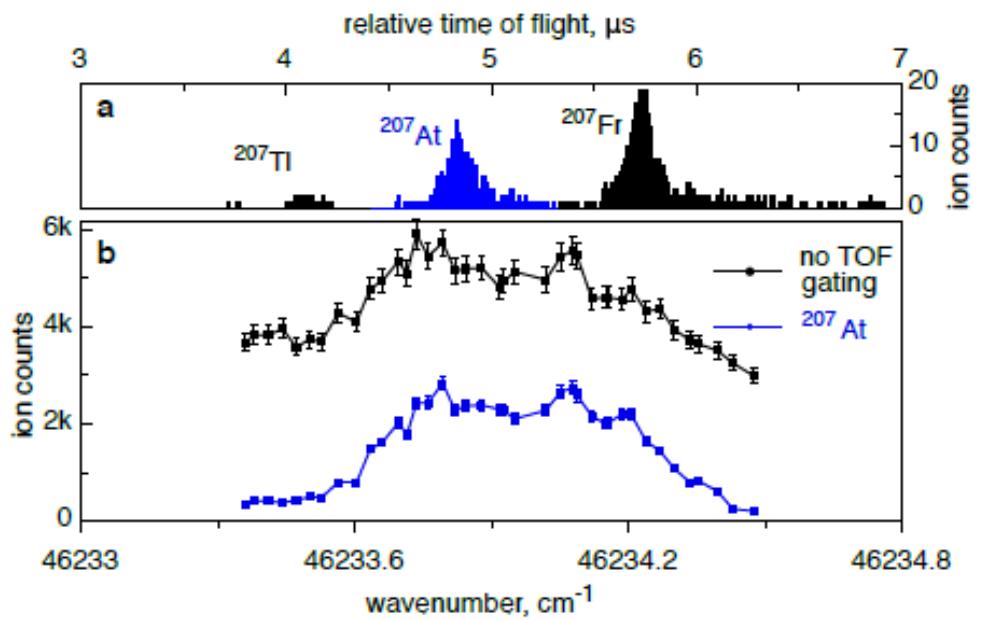


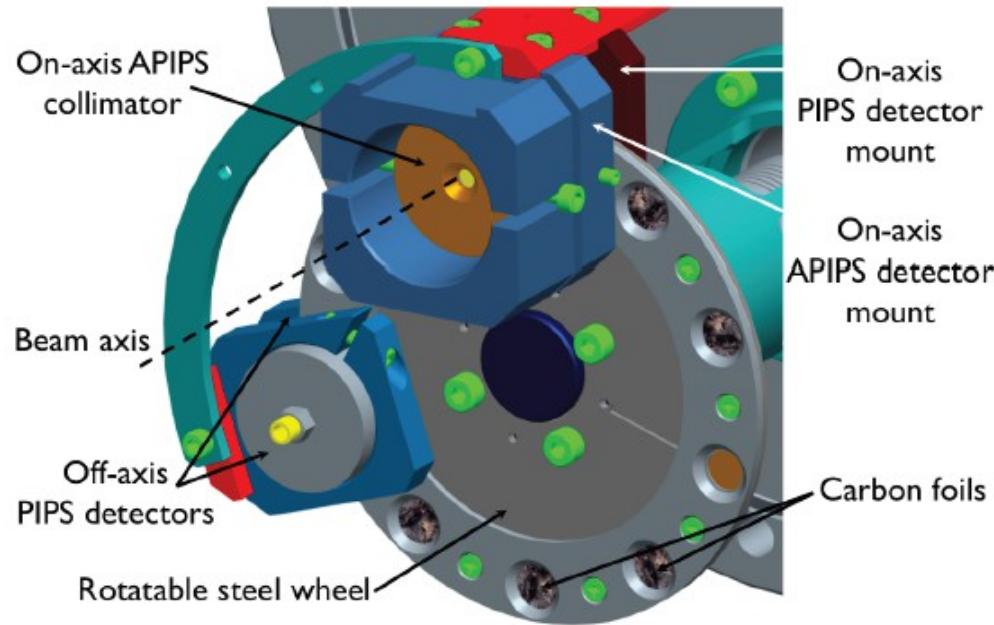


RILIS lasers



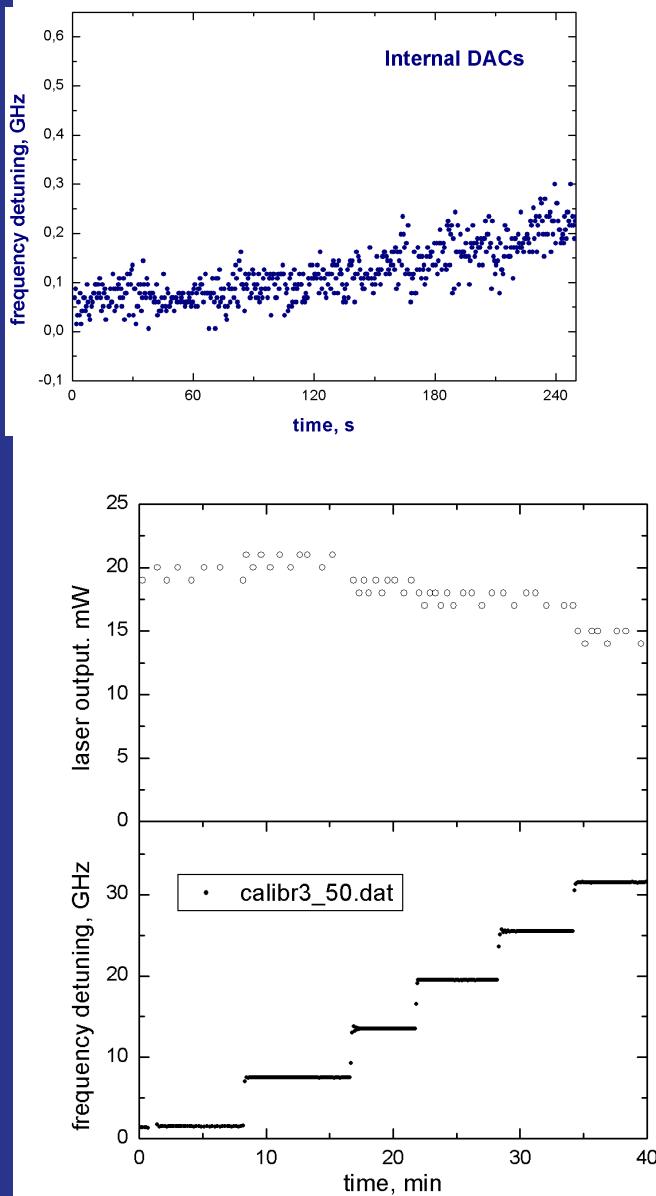




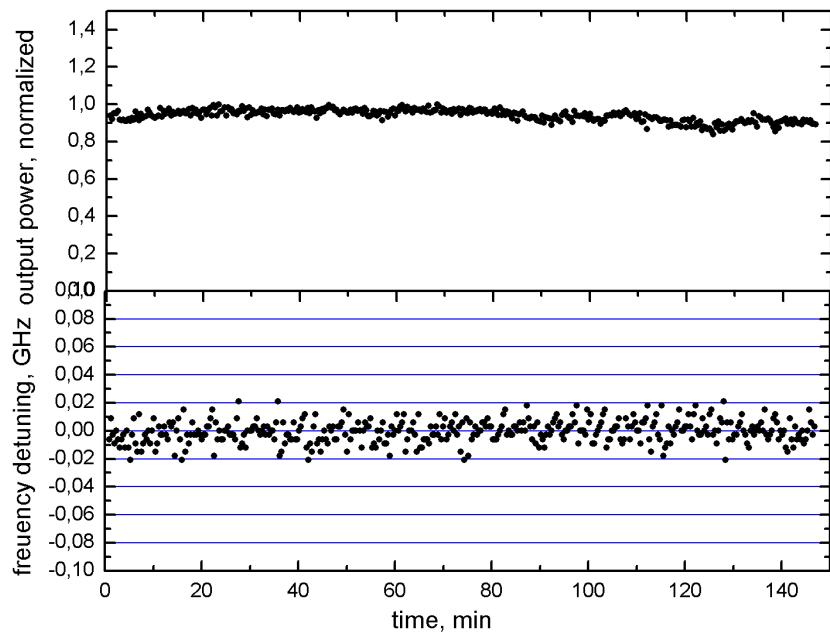


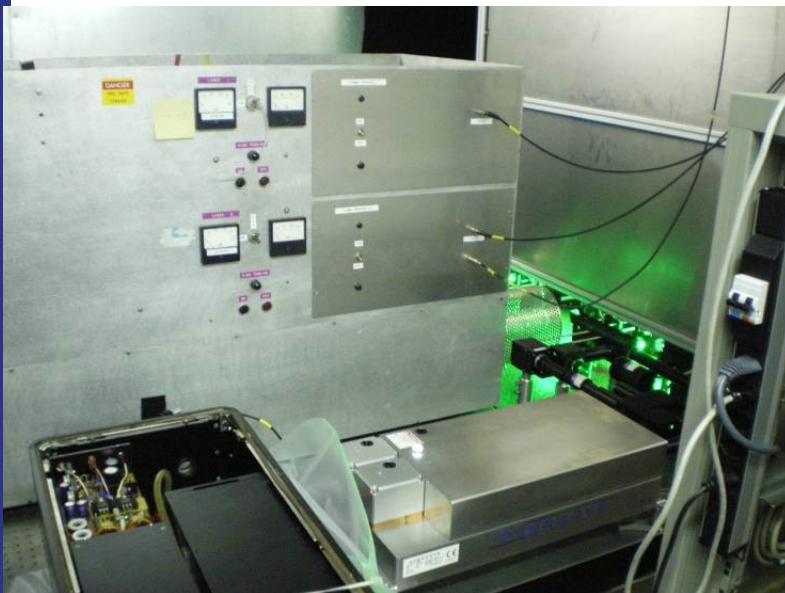


RILIS upgrade



20.02.2003 stability of the ISOLDE narrow-band laser
with Nd:YAG pumping (Mainz)





Advantages:

Better beam quality

Stability of operation

No desynchronization problems!

Complications:

New ionization schemes are needed (Mn, Au)

Service by manufacturer only

Shorter pulses

Copper Vapor Lasers are replaced by
Diode Pumped Solid State Nd:YAG Lasers
Total power > 100 W

Laser generates 3 beams at 10 kHz:

Main green beam

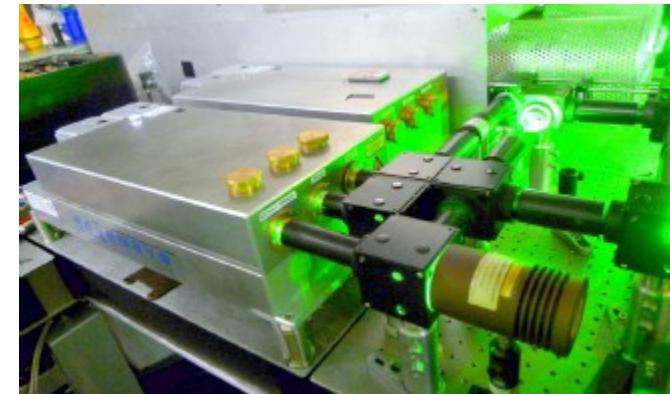
- 532nm, 70-80 W, 8 ns

Residual green beam

- 532 nm, 12-28 W, 9 ns

UV beam

- 355 nm, 18-20 W, 11 ns



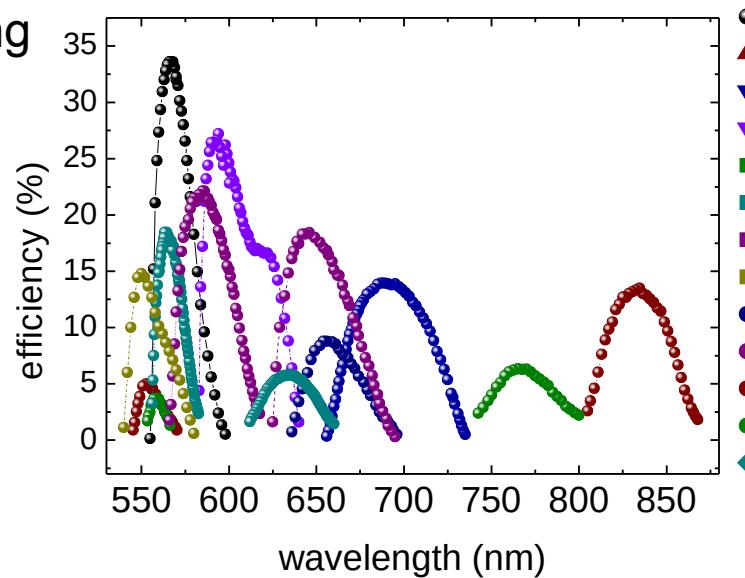
RILIS Dye Lasers

Optimized for Nd:YAG pumping
(UV+Visible)

Higher power

Tunable and **SCANNING**

Pump laser: Nd:YAG
(532 nm), Edgewave
Repetition rate: 10 kHz,
Pulse duration: 9 ns
Power: 100 W



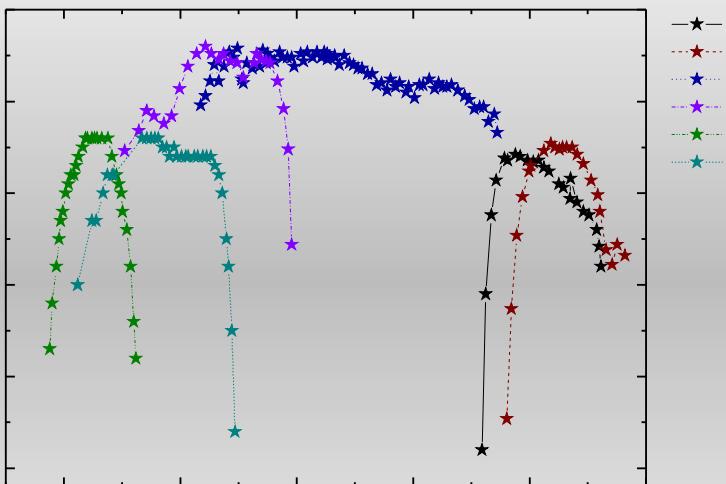
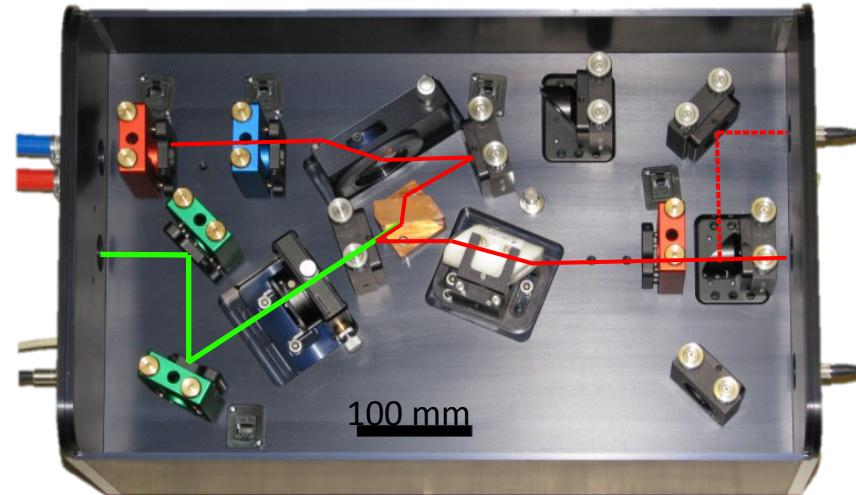
RILIS Ti:Sa lasers

Pump laser: Nd:YAG (532 nm), Photonics

Repetition rate: 10 kHz

Pulse length: 180 ns

Power: 60 W



Design & Construction:
S. Rothe (Uni Mainz)

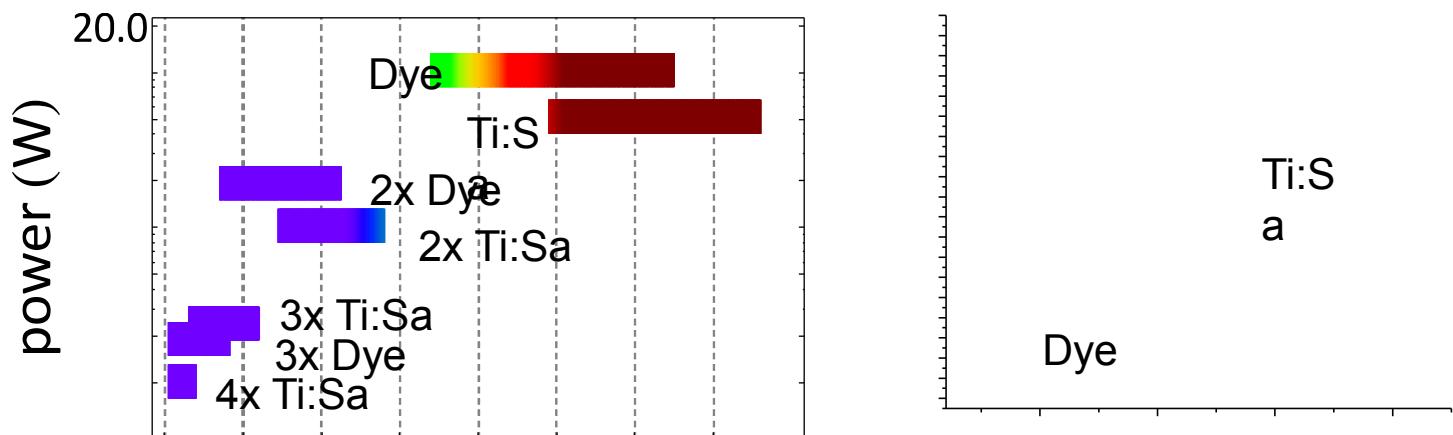
Wavelength tuning range:

- Fundamental (ω) **690** **940** nm (5 W)
- 2nd harmonic (2ω) **345** **470** nm (1 W)
- 3rd harmonic (3ω) **230** **310** nm (150 mW)
- 4th harmonic (4ω) **205** - **235** nm (50 mW)

6 resonator mirror sets cover the Ti:Sa range

Comparison dye vs. Ti:Sa system

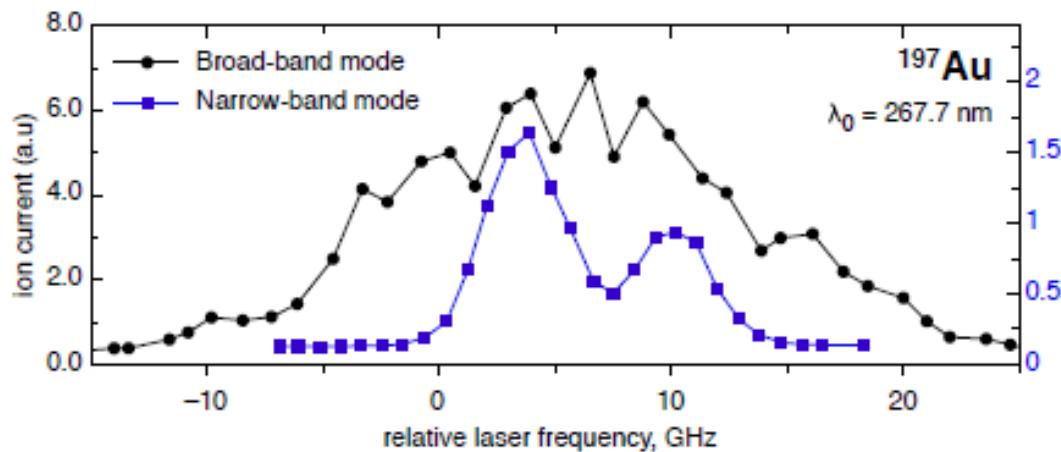
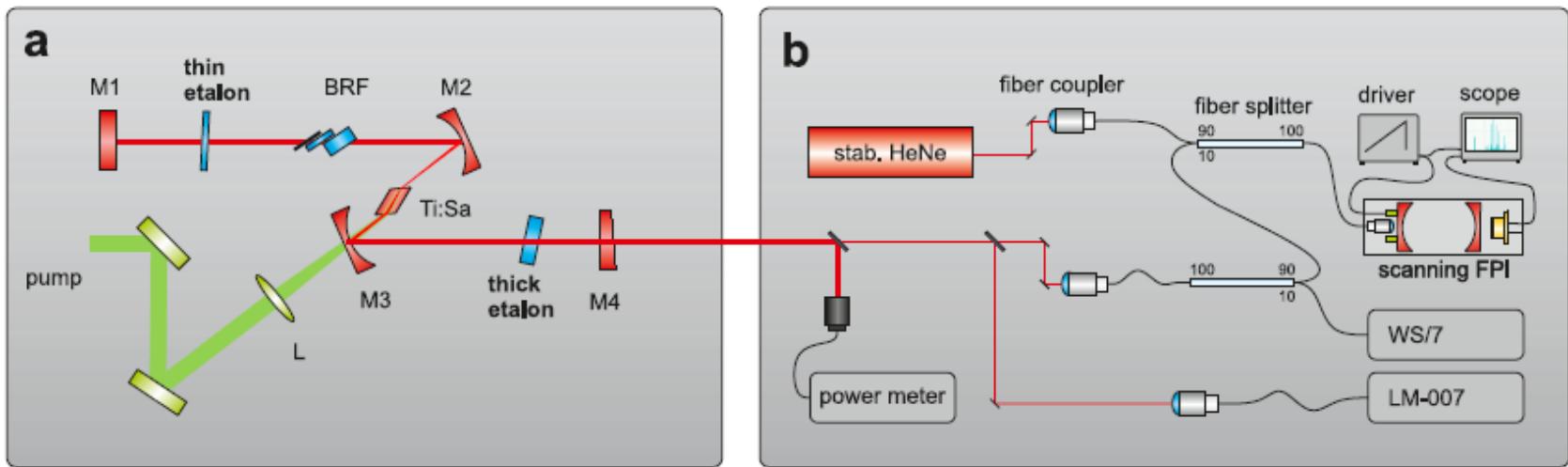
	Dye	Ti:Sa
Active Medium condition of aggregation	> 10 different dyes liquid (org. solvents)	=1 Ti:sapphire crystal solid-state
Tuning range	540 – 850 nm	680 – 980 nm
Power	< 12 W	< 5 W
Pulse duration	~8 ns	~50 ns
Synchronization	optical delay lines	q-switch, pump power
# of schemes developed	47	37
Maintenance	renew dye solutions	~ none



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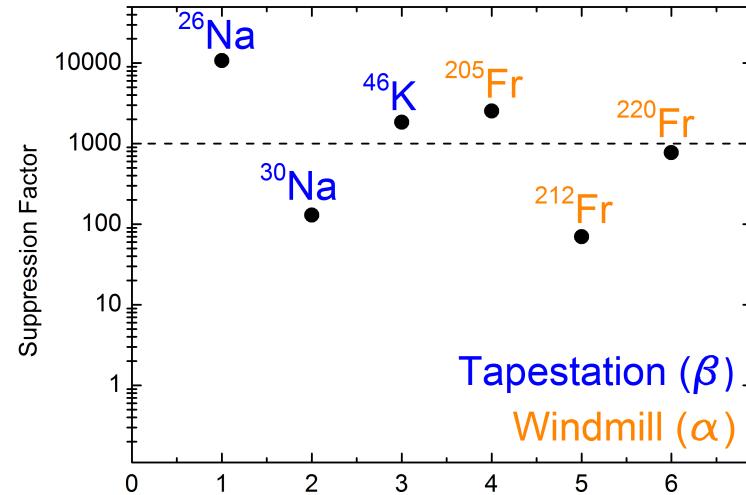
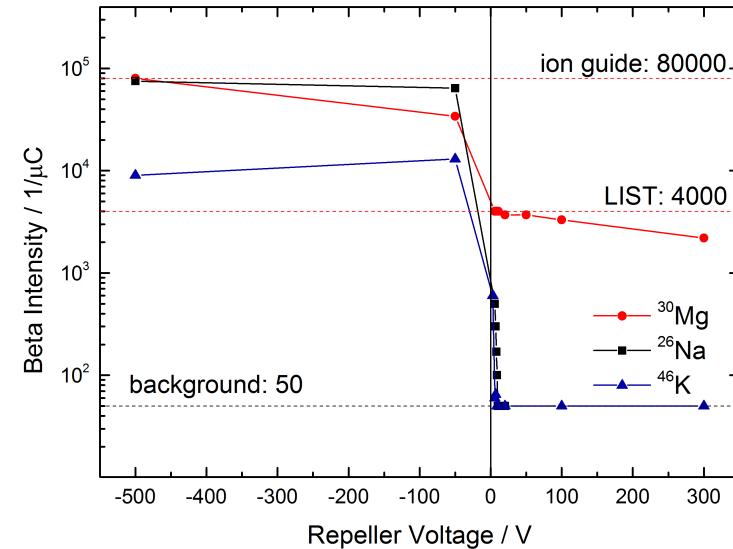
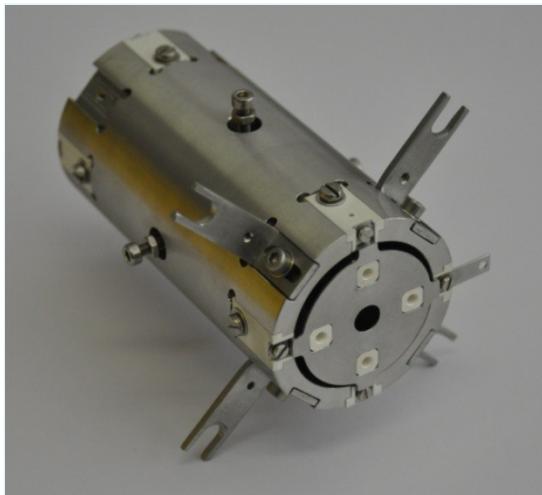
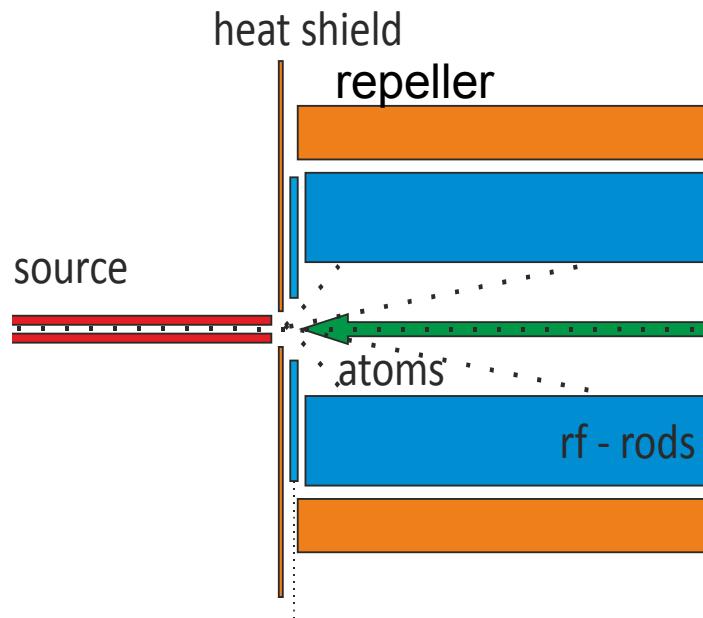
Ti:Sa system is complementary

Narrow-band scanning Ti:Sa laser

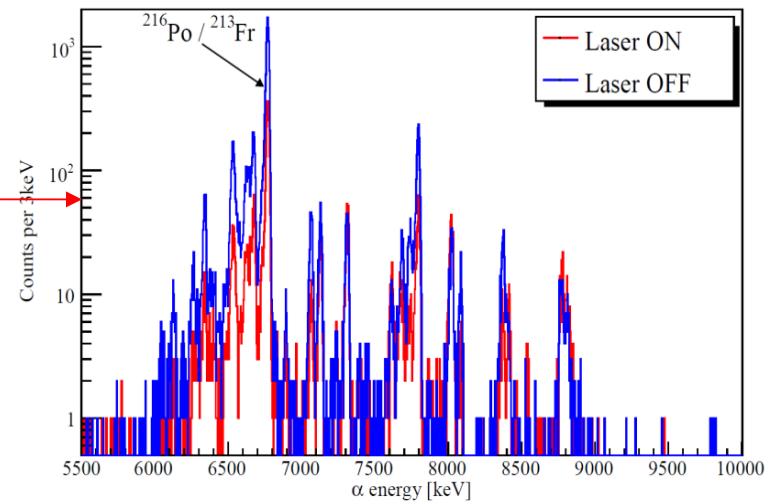
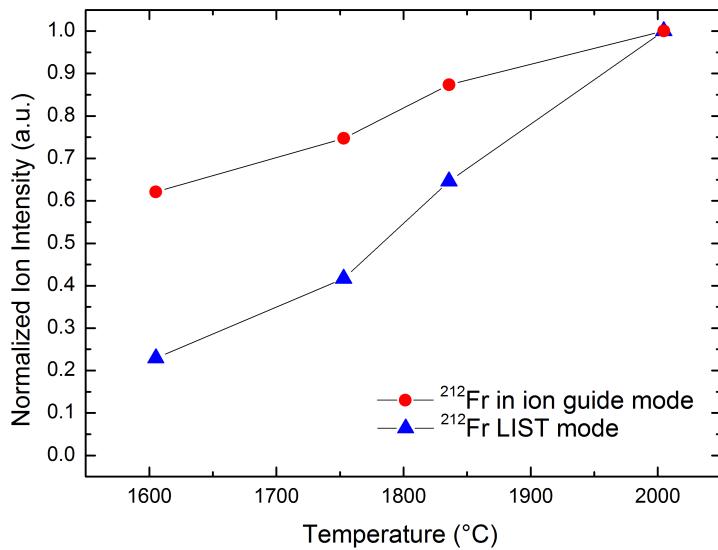


2012

LIST: Laser Ion Source Trap



LIST: Laser Spectroscopy of ^{217}Po



RILIS elements

Currently available RILIS elements are highlighted in red.

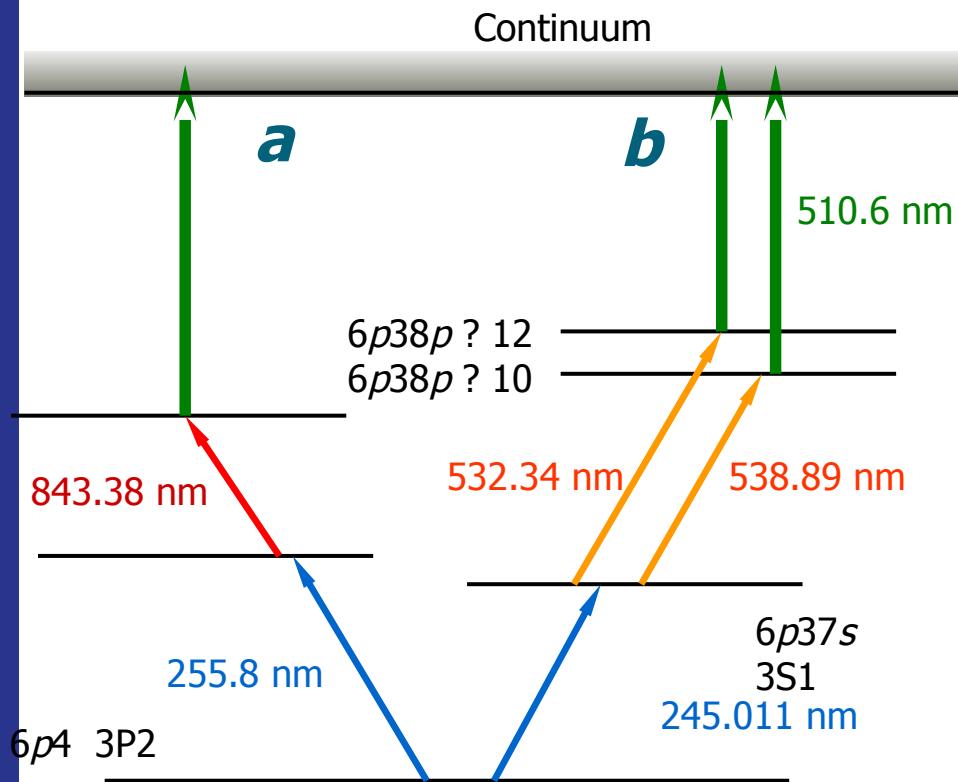
Elements for which ionization schemes have been tested to some extent but not yet applied are highlighted in green.

Elements for which ionization is feasible at RILIS but has not been tested are highlighted in yellow.

1 H																				2 He
3 Li	4 Be																			
11 Na	12 Mg																			
19 K	20 Ca	21 Sc	22 Ti	23 V	24 Cr	25 Mn	26 Fe	27 Co	28 Ni	29 Cu	30 Zn	31 Ga	32 Ge	33 As	34 Se	35 Br	36 Kr			
37 Rb	38 Sr	39 Y	40 Zr	41 Nb	42 Mo	43 Tc	44 Ru	45 Rh	46 Pd	47 Ag	48 Cd	49 In	50 Sn	51 Sb	52 Te	53 I	54 Xe			
55 Cs	56 Ba	57 La	72 Hf	73 Ta	74 W	75 Re	76 Os	77 Ir	78 Pt	79 Au	80 Hg	81 Tl	82 Pb	83 Bi	84 Po	85 At	86 Rn			
87 Fr	88 Ra	89 Ac	104 Rf	105 Db	106 Sg	107 Bh	108 Hs	109 Mt	110	111	112									

58 Ce	59 Pr	60 Nd	61 Pm	62 Sm	63 Eu	64 Gd	65 Tb	66 Dy	67 Ho	68 Er	69 Tm	70 Yb	71 Lu						
90 Th	91 Pa	92 U	93 Np	94 Pu	95 Am	96 Cm	97 Bk	98 Cf	99 Es	100 Fm	101 Md	102 No	103 Lr						

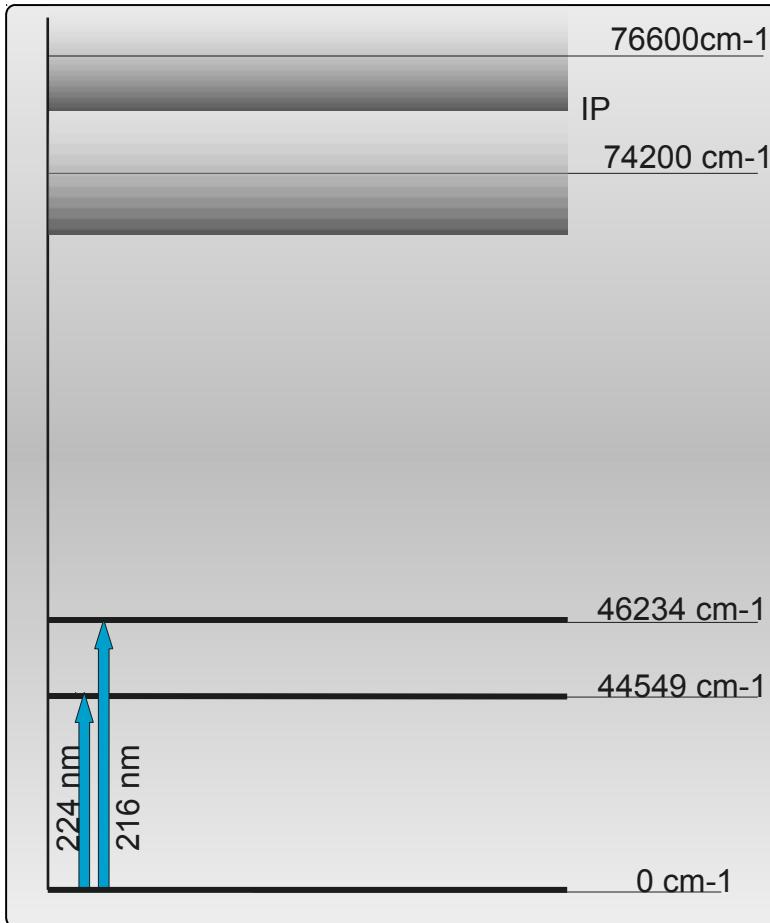
New photoionization schemes: Po



Po yields (scheme „a“)

Isotope	Half life, s	Yield, Atoms/ μ C
193gPo	0.42	7×10^1
193mPo	0.24	1×10^2
194Po	0.392	2.5×10^3
195gPo	4.64	2×10^4
195mPo	1.92	5×10^4
196Po	5.8	4.7×10^5
197gPo	53.6	2.5×10^5
197mPo	25.8	1.75×10^6
198Po	106.2	7×10^6

At: Photoionization scheme and Ionization potential



Recent interest in At and its IP:

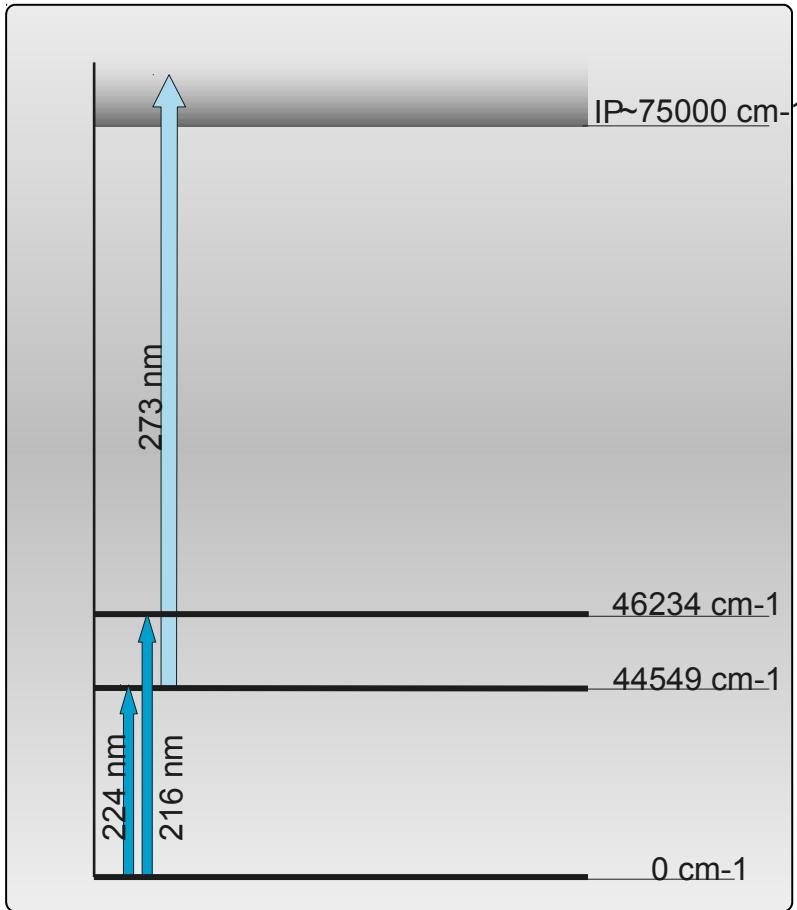
- targeted a therapy for cancer treatment
- Benchmark for theoretical chemistry of astatine
- Benchmark for calculations for IP(117Uus)
- At beam for ISOLDE users (β -delayed fission, laser spectroscopy)

Theoretical predictions of IP(At)

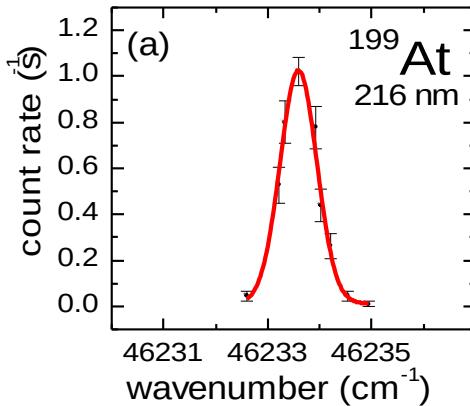
Reference	Year	IP (eV)	IP (cm ⁻¹)
[Fin55]	1955	9.2 ± 0.4	74 203 ± 6 500
[Kis60]	1960	9.5	76 623
[Kue91]	1991	9.4	75 816
[Mit06]	2006	9.24	74 526
[Cha10]	2010	9.35 ± 0.01	75 413 ± 160



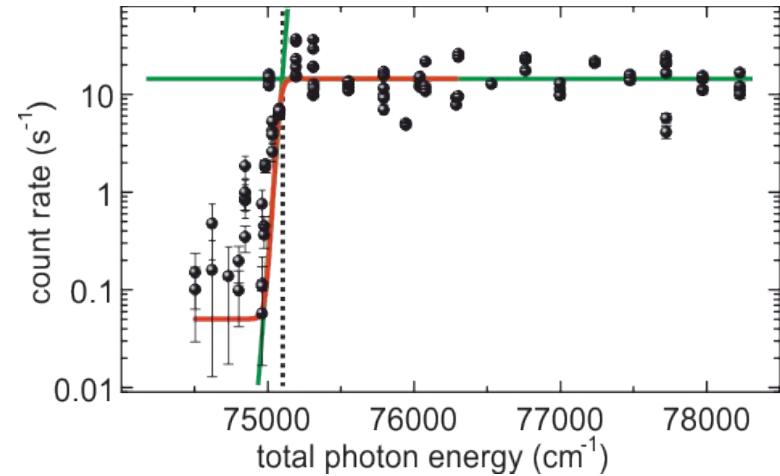
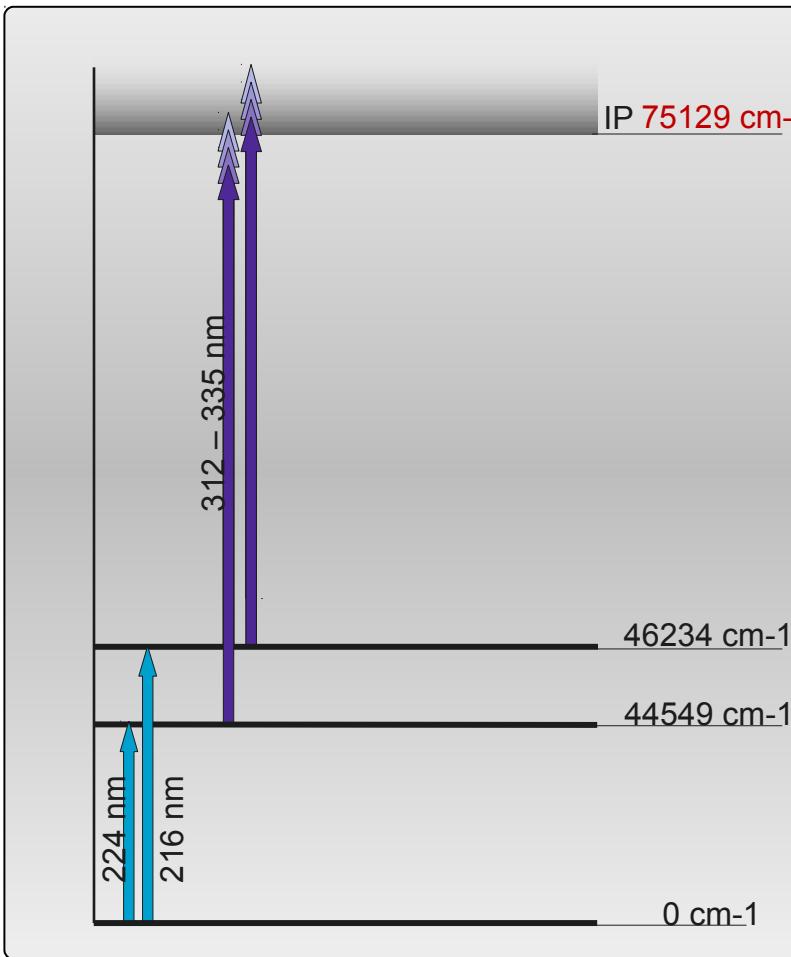
At: Photoionization scheme and Ionization potential



- ~2W @ 273 nm for non-resonant ionization
- Laser scans of 224 nm and 216 nm transitions
- Very low yields 1-10 s⁻¹
- ~5 min per wavelength step



At: Photoionization scheme and Ionization potential

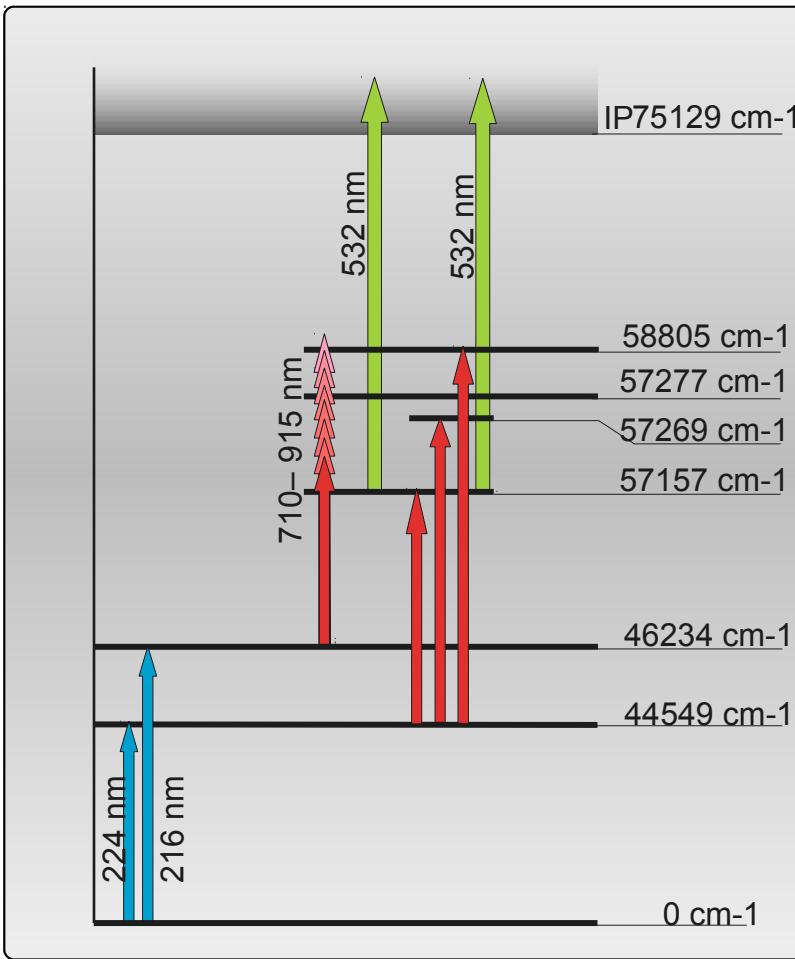


- Laser scan of second laser
- Low resolution
- Required ~6 h data taking

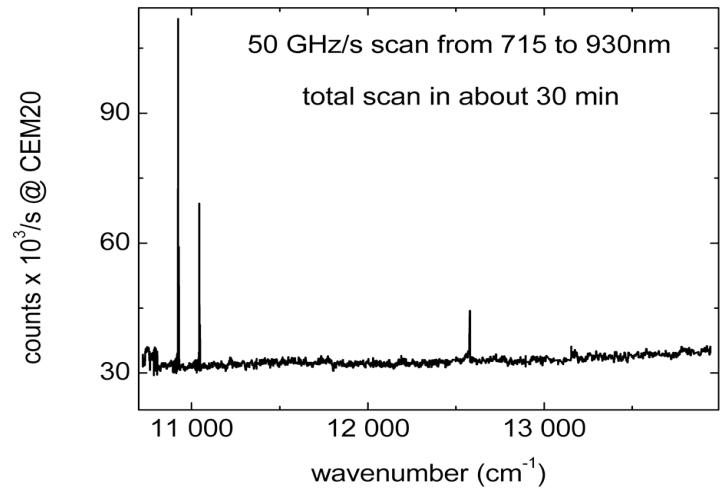
$$\text{IP}_{\text{threshold}}(\text{At}) = 75129(95) \text{ cm}^{-1}$$

- Higher resolution needed
- low yield due to low laser power in final step
- 3-color scheme allows use of 532 nm (50W)

At: Photoionization scheme and Ionization potential

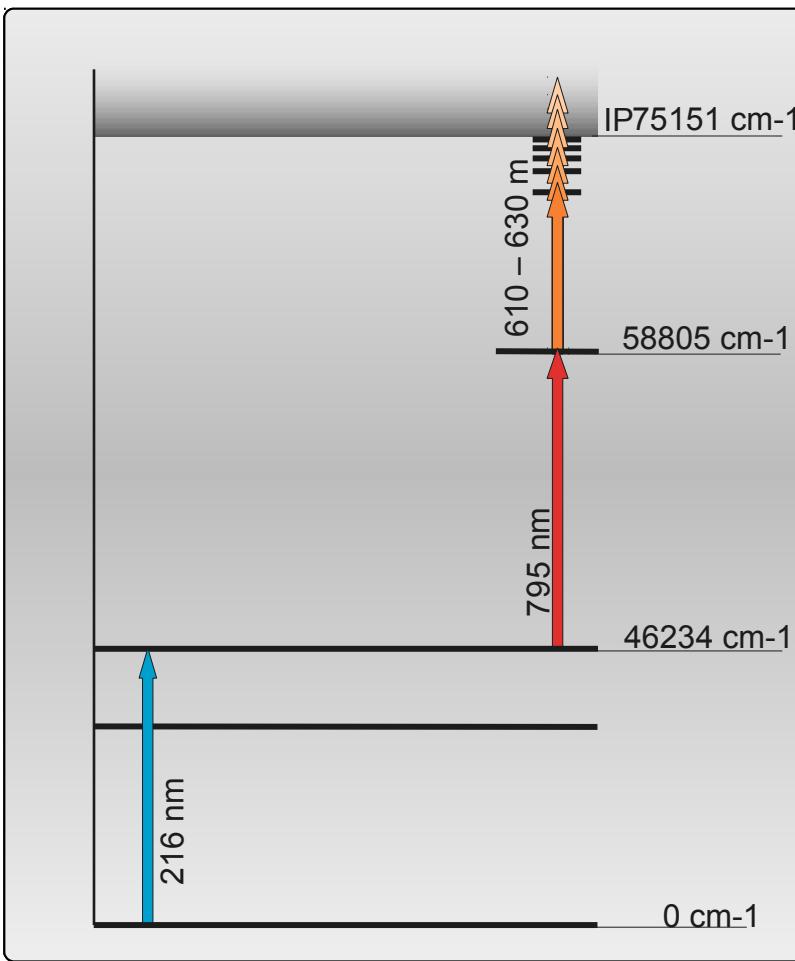


- Spectroscopy at ISAC/TRIUMF (199At)
- cw proton beam from cyclotron
- 200 nm scan: 3 new transitions
- Verified at ISOLDE/CERN (205At)

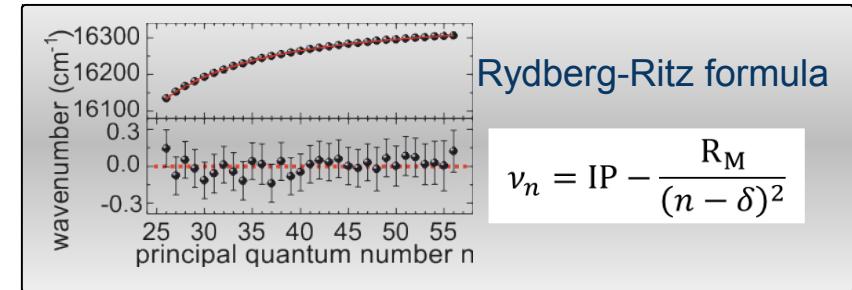
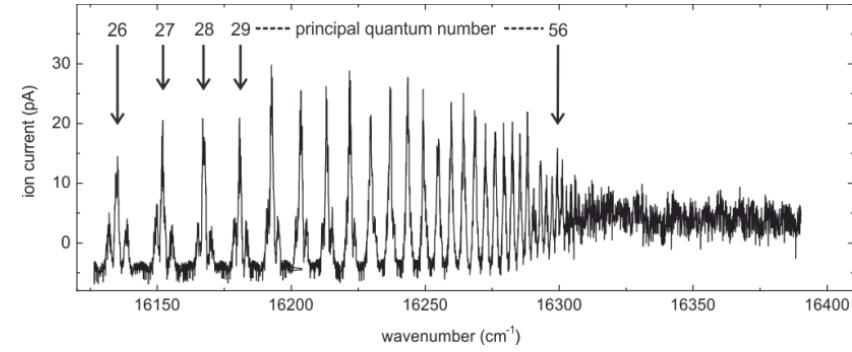


- 6 transitions, 4 new energy levels available
- Up to 150 pA of 205At
- Continuously measurable with Faraday cup

At: Photoionization scheme and Ionization potential



Spectroscopy of Rydberg levels



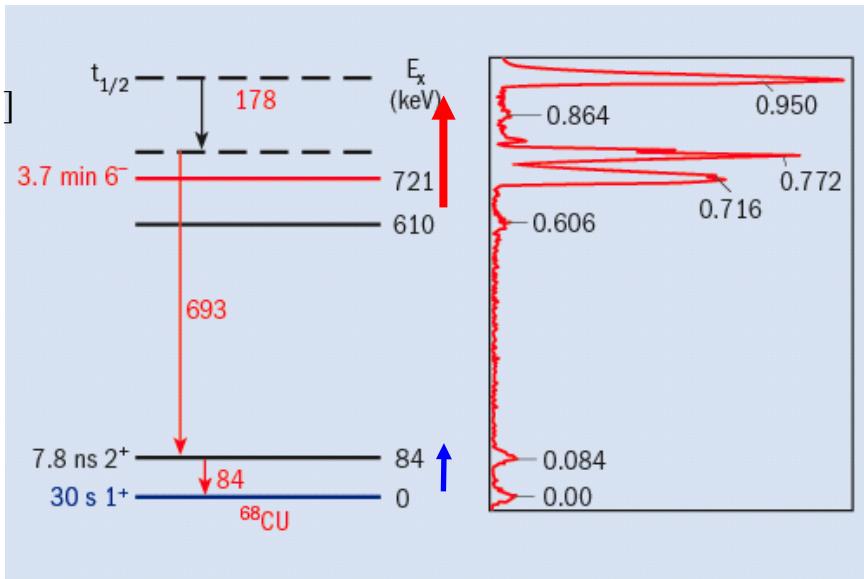
$$\text{IP}_{\text{Rydberg}}(\text{At}) = 75151(1) \text{ cm}^{-1}$$

$$\text{IP}_{\text{threshold}}(\text{At}) = 75129(95) \text{ cm}^{-1}$$

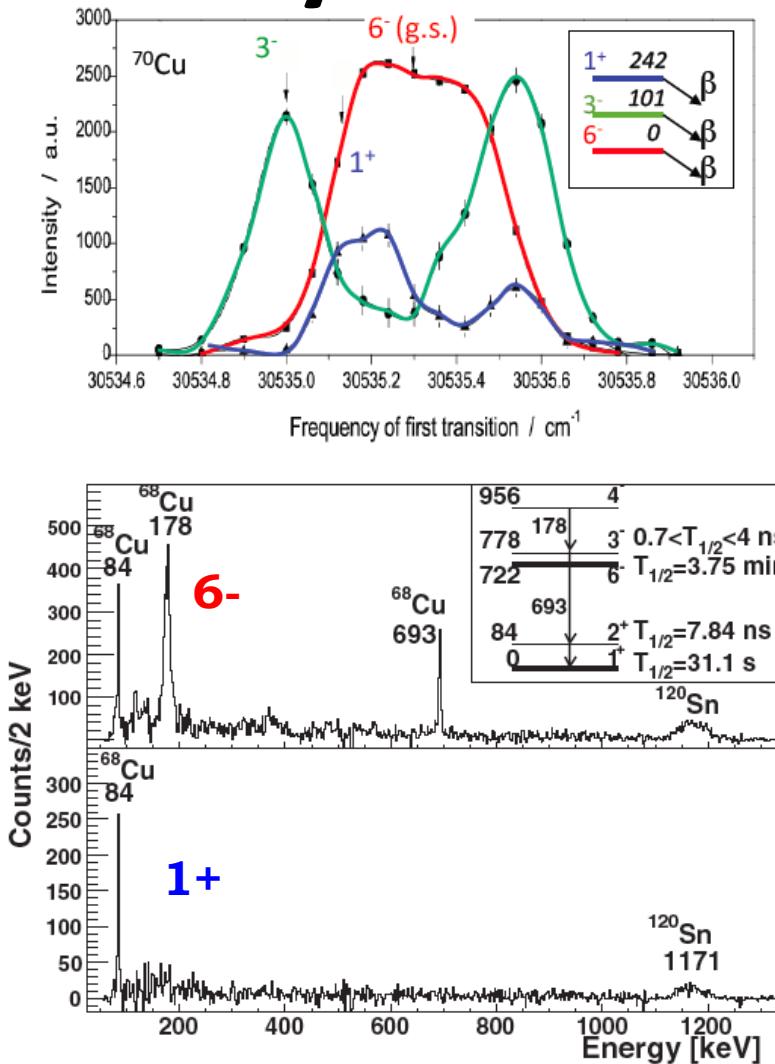
Isomer selectivity

Laser-> Maximize different isomers

$[\pi\ 1p_{3/2}\gamma\ 0g_{9/2}]$



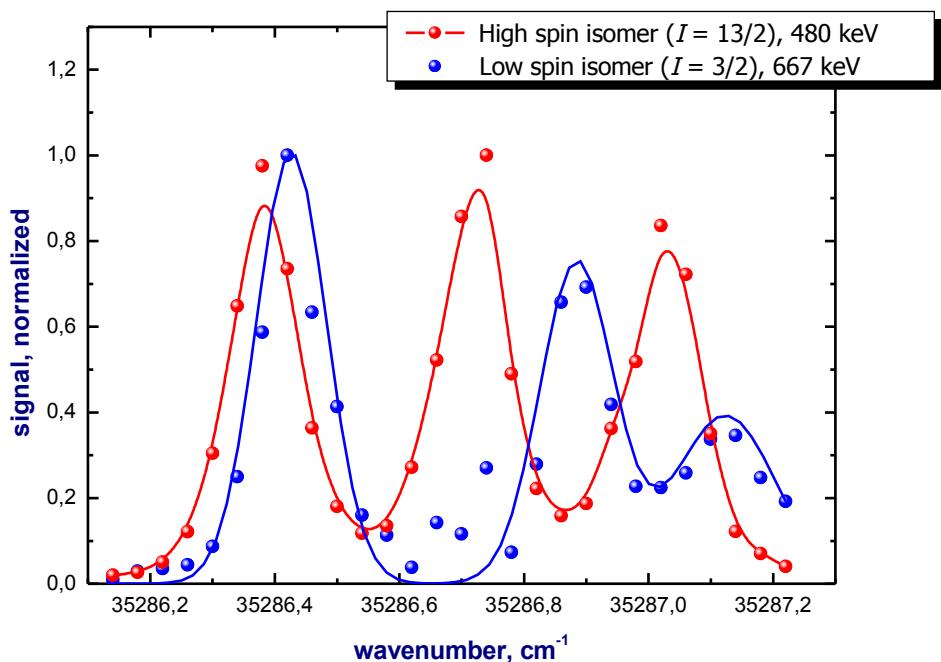
$[\pi\ 1p_{3/2}\gamma\ 1p_{1/2}]$



Isomer selectivity

Laser-> Maximize different isomers

189Pb

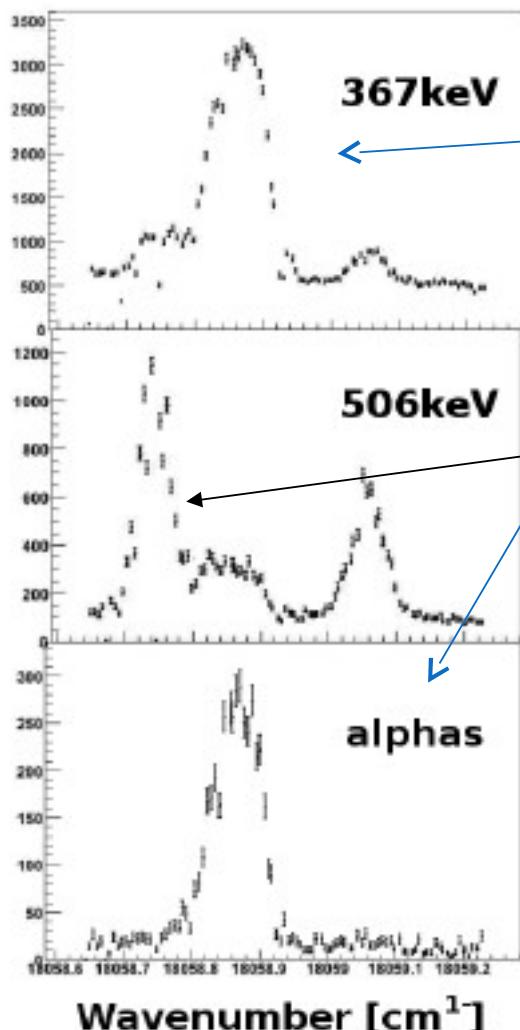


A level scheme of the ^{189}TI nucleus has been established from the β^+/EC decay study of the ^{189}Pb isomers using both nuclear spectroscopy and in-source laser spectroscopy experiments.

40 gamma lines belonging to the β/EC decay of ^{189}Pb have been identified:
386, 480, 700, 399....and 667keV are the main ones.

Isomer selectivity

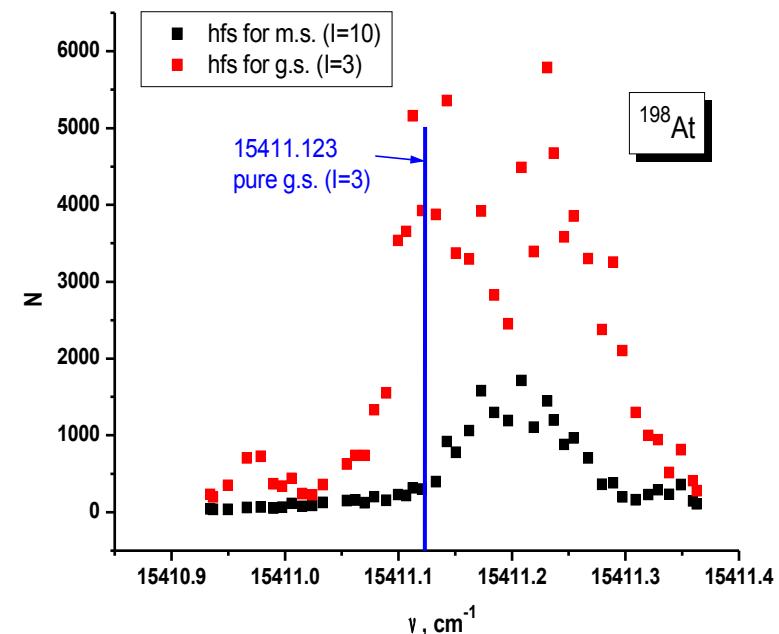
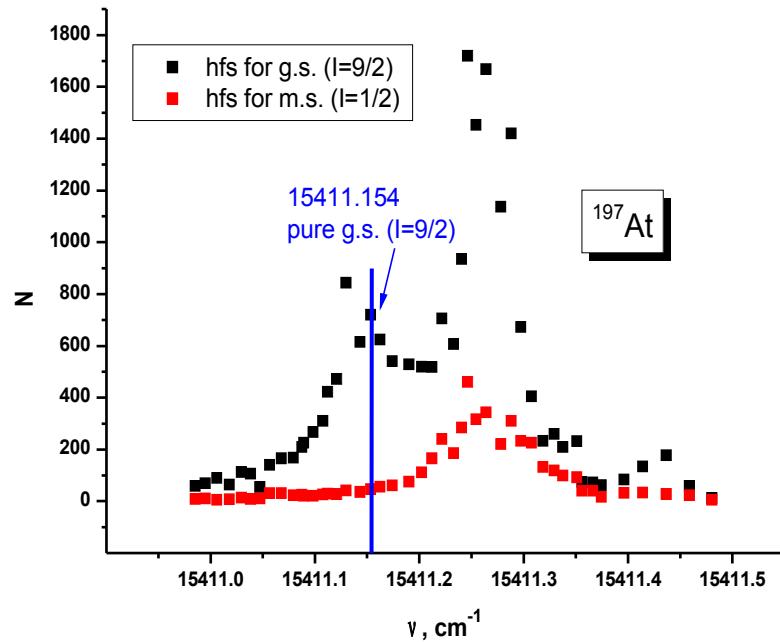
^{184}TI



Hyperfine structures observed for ^{184}TI with different detection modes

Isomer selectivity

197,198At



Isomer selectivity enable us to measure masses of 197g,198gAt and receive nuclear spectroscopic information for pure g.s.

Nuclear charge radii and electromagnetic moments

Isotope shift $\delta\nu_{A,A'}$

$$\delta\nu_{A,A'} = F \lambda_{A,A'} + MS$$

Rms charge radius

$$\lambda_{A,A'} = \delta r_2^{A,A'} + C_2 \delta r_4^{A,A'} + \dots = 0.93 \delta r_2^{A,A'}$$

Relative line position \rightarrow hyperfine constants A & $B \rightarrow ml, QS$

$$\nu_{F_i, F_f} = \nu_0 + \nu_{F_f} - \nu_{F_i}$$

$$\nu_F = A \cdot \frac{K}{2} + B \cdot \frac{0.75 \cdot K \cdot (K+1) - I \cdot (I+1) \cdot J \cdot (J+1)}{2 \cdot (2I-1) \cdot (2J-1) \cdot I \cdot J}$$

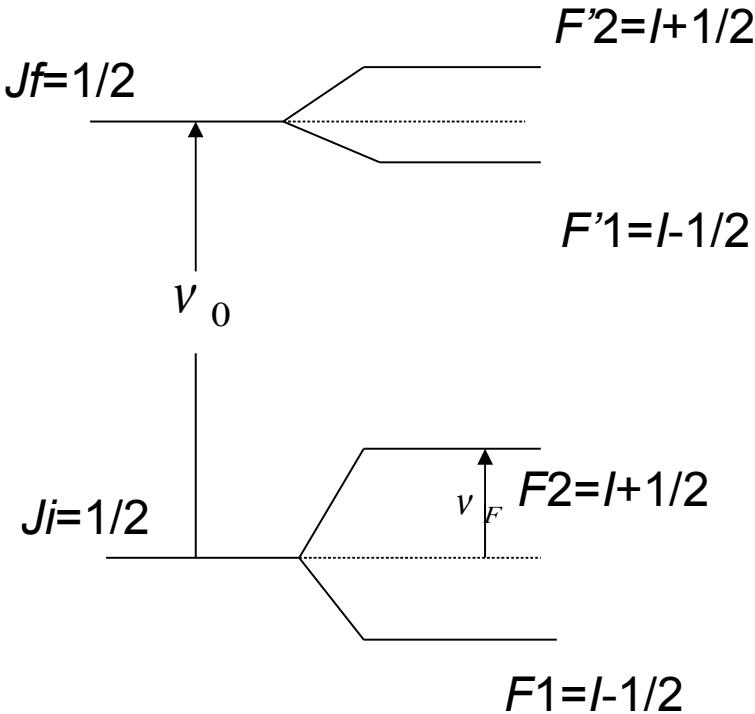
$$K = F \cdot (F+1) - I \cdot (I+1) - J \cdot (J+1)$$

$$\vec{F} = \vec{I} + \vec{J}, \quad F = |I - J|, |I - J| + 1, \dots, I + J$$

$$A \propto \mu, \quad B \propto Q$$

Amplitudes of the components:

$$S(F_i \rightarrow F_f) = \frac{(2F_f + 1) \times (2F_i + 1)}{2I + 1} \times \begin{Bmatrix} J_f & F_f & I \\ F_i & J_i & 1 \end{Bmatrix}^2$$



Nuclear deformation

Charge radii and deformation:

$$\langle r^2 \rangle_A \approx \langle r^2 \rangle_A^{sph} \left(1 + \frac{5}{4\pi} \langle \beta_2^2 \rangle_A \right)$$

$\langle r^2 \rangle_A^{sph}$ is the mean square radius of a spherical nucleus with the same volume. Usually evaluated using droplet model

Quadrupole moment and deformation:

$$Q_S = \frac{3K^2 - I(I+1)}{(I+1)(2I+3)} Q_0,$$

K is the projection of the nuclear spin on the symmetry axis of the nucleus.

$$Q_0 \approx \frac{3}{\sqrt{5\pi}} e Z R_0^2 \left(\beta_2 + \frac{2}{7} \sqrt{\frac{5}{\pi}} \beta_2^2 + \dots \right), \quad R_0 = 1.2A^{1/3} \text{ fm.}$$

Hyperfine components intensities: rate equations

$$N_i(v) = C_1 \int N_0^G(v') P_i(I^{L'}(v - v')) dv' + C_0$$

To take into account the saturation of transitions, pumping processes between hyperfine structure (hfs) components and a population redistribution of the hfs levels the number of photoions N_{ion} for each frequency step was calculated by solving the rate equations for the given photoionization scheme:

$$\begin{cases} \frac{dN_F}{dt} = \sum_k W_{F'_k F} N_{F'_k} - \sum_k W_{FF'_k} N_F - W_{F,ion} N_F \\ \vdots \\ \frac{dN_{ion}}{dt} = \sum_k W_{F'_k,ion} N_{F'_k} \end{cases}$$

$$W_{FF'} \sim S_{FF'}^* I(v + \Delta v^{FF'} - v'), \quad S_{FF'}^* = S_{FF'} / (2F + 1)$$

$$\text{At } t = 0: N_F^0 \sim 2F + 1$$

Hyperfine components intensities: rate equations

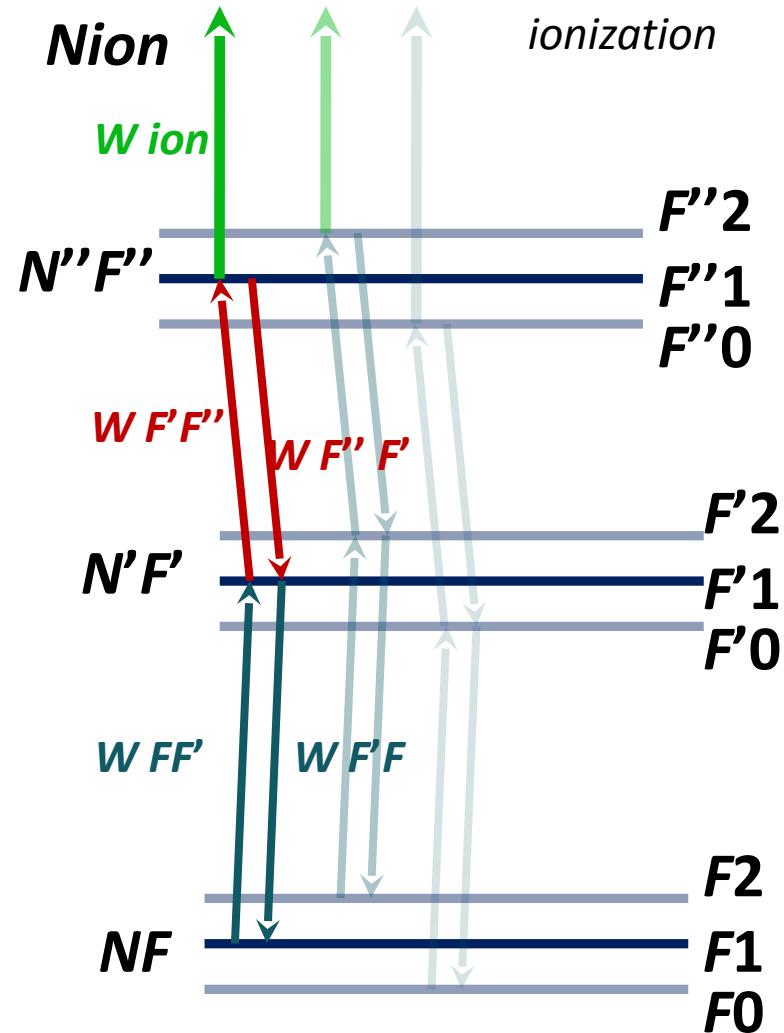
$$dN_{ion} = W_{ion} N''_{F''} dt + \dots$$

$$dN''_{F''} = - N''_{F''} (W_{ion} + W_{F''F'} + \dots) dt + \\ + N'_{F'} W_{F'F''} dt + \dots$$

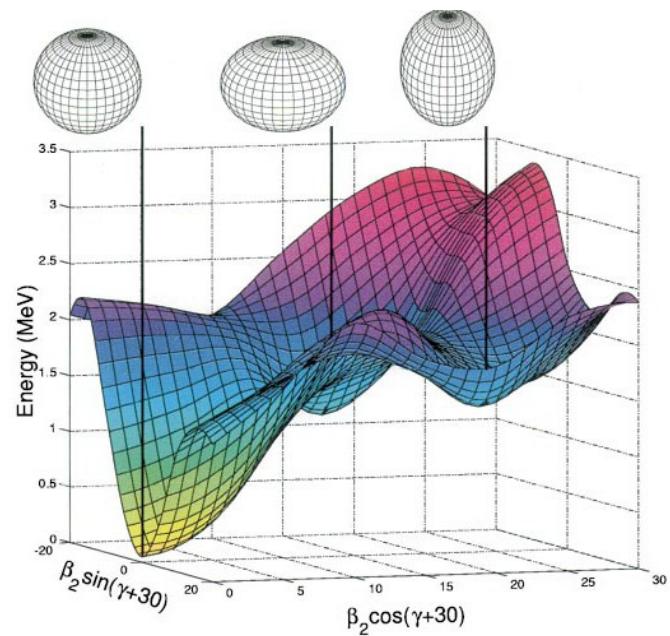
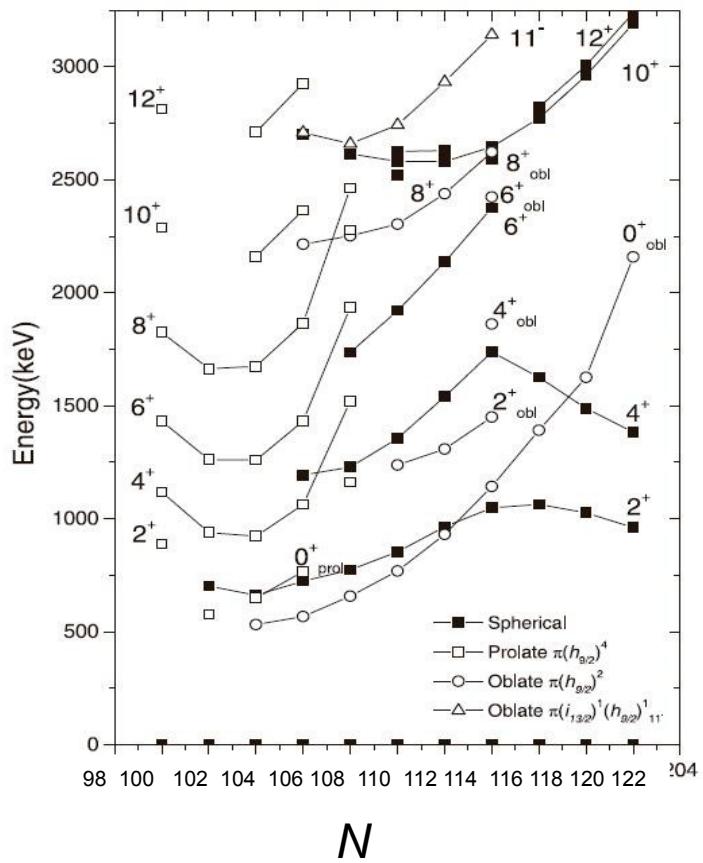
$$dN'_{F'} = - N'_{F'} (W_{F'F''} + W_{F'F} + \dots) dt + \\ + (N''_{F''} W_{F''F'} + N_F W_{FF'} + \dots) dt$$

$$dN_F = - N_F (W_{FF'} + \dots) dt + \\ + (N'_{F'} W_{F'F} + \dots) dt$$

$$\frac{W_{FF'}}{W_{F'F}} = \frac{2F'+1}{2F+1}$$



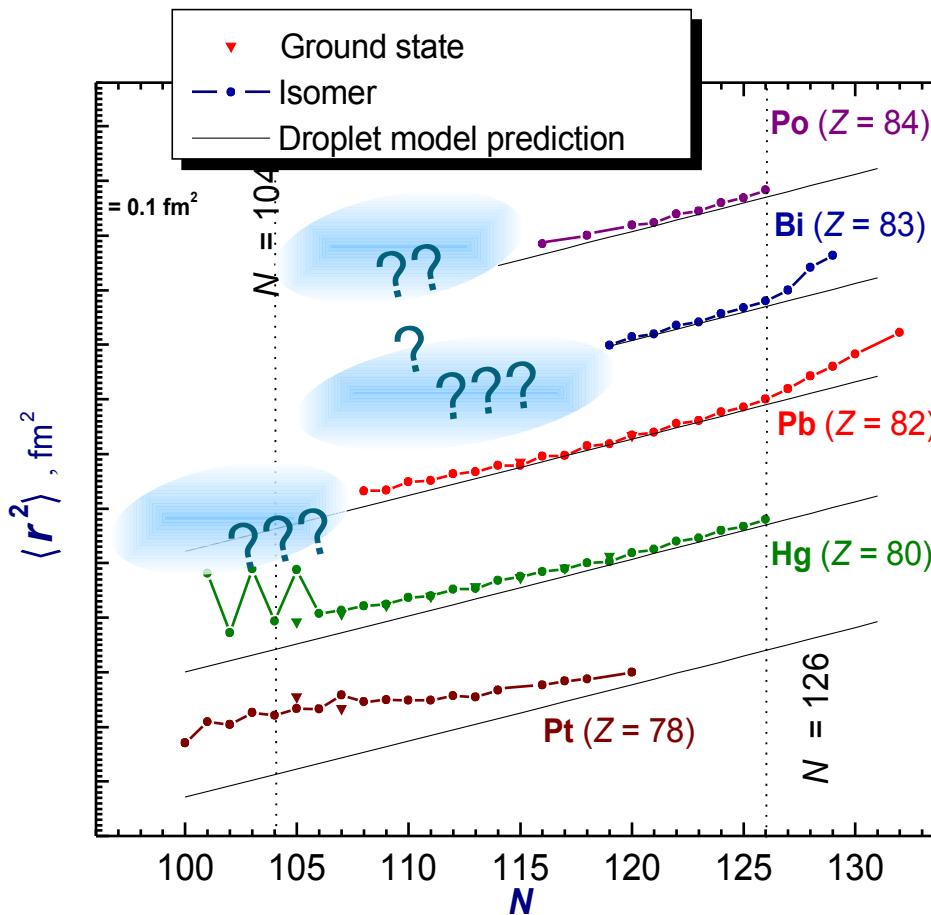
Shape coexistence in Pb region



Level systematics for the neutron-deficient lead isotopes.

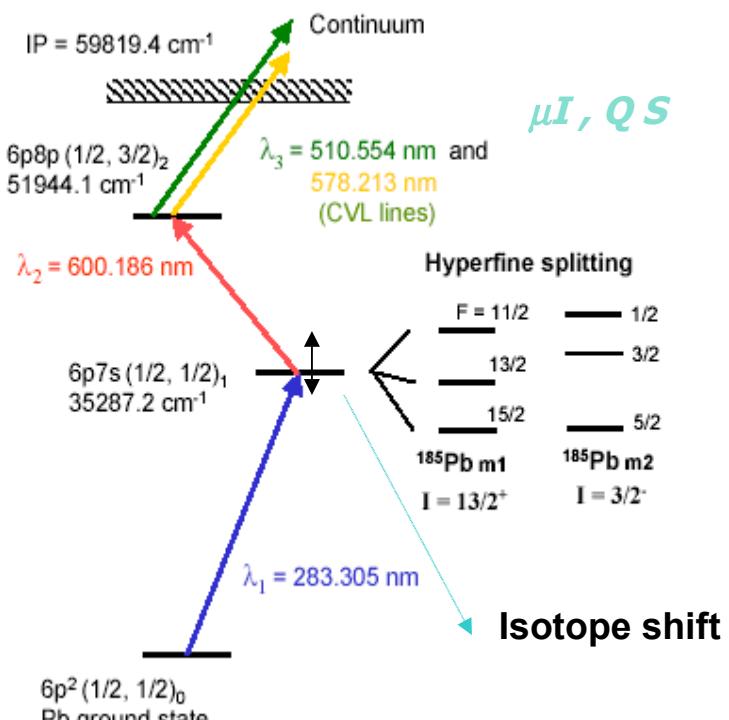
R. Julin et al., J. Phys. G: Nucl. Part. Phys. 27 (2001)

Nuclear charge radii around Z=82



nuclear ground and isomeric state properties : $\delta \langle r^2 \rangle$

Pb ($Z=82$)

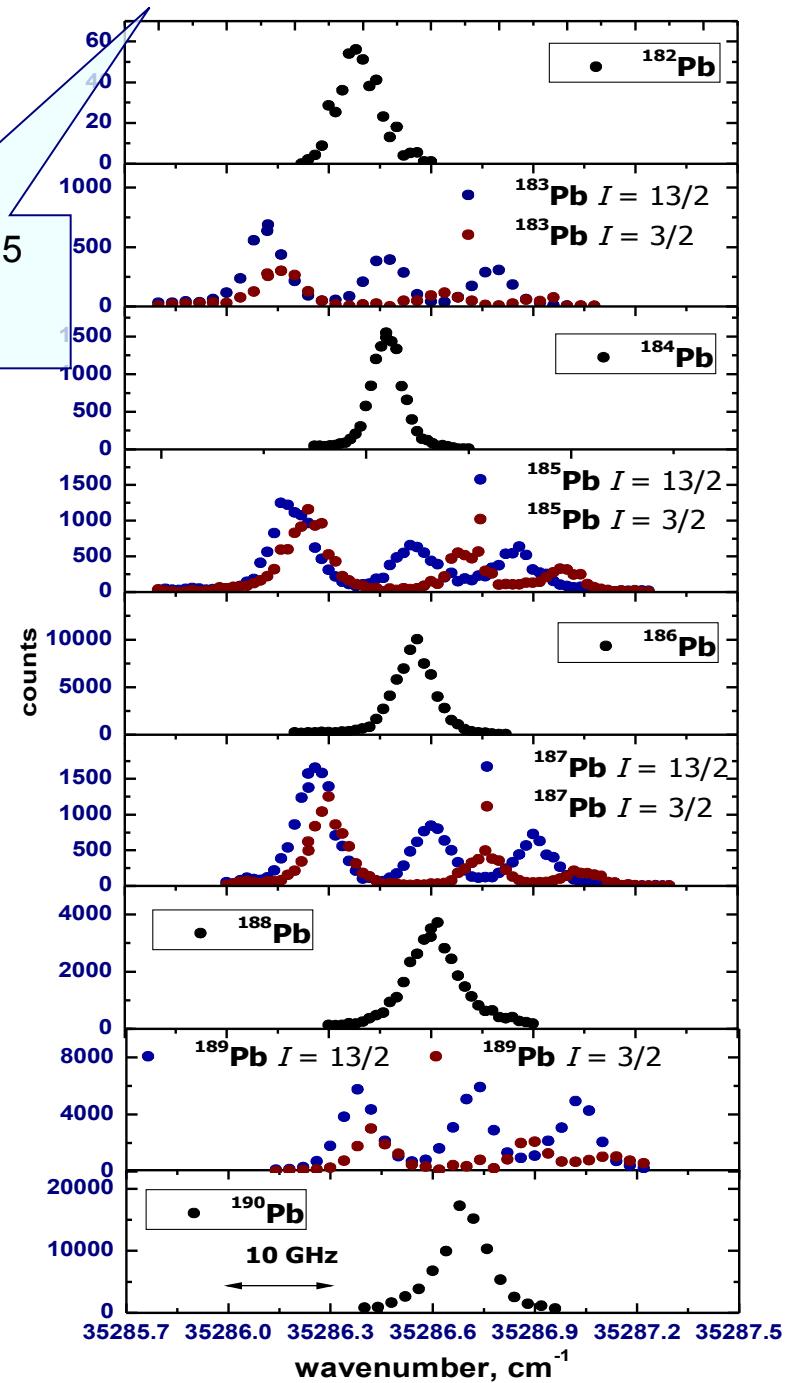


^{182}Pb : $T_{1/2} = 55$
ms
Yield: $\sim 1 \text{ s}^{-1}$

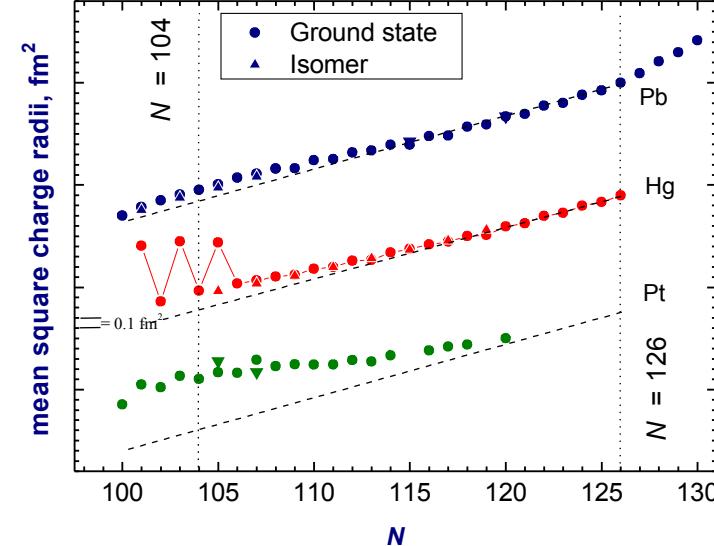
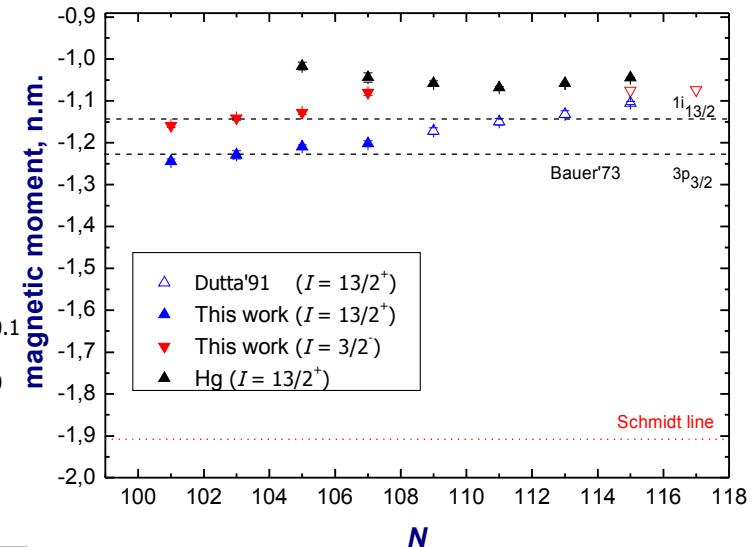
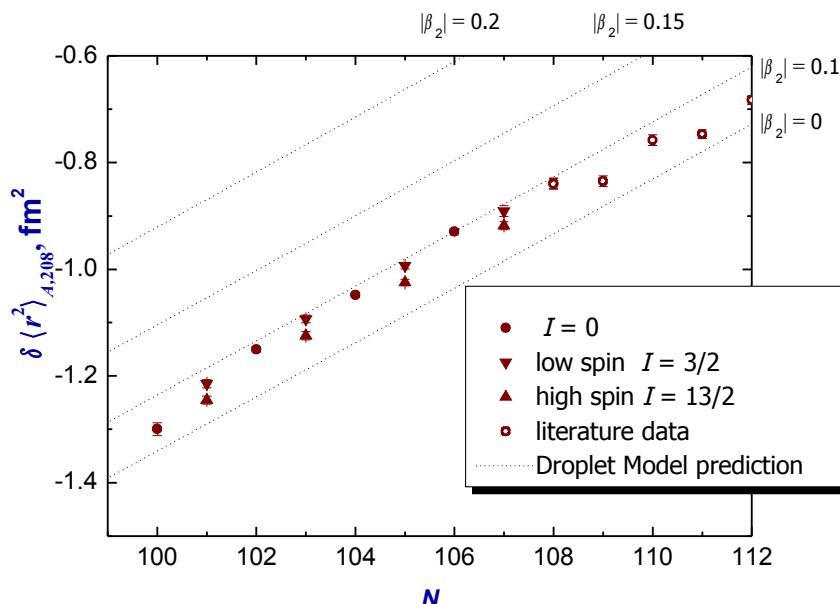
$\mu I, Q_S$

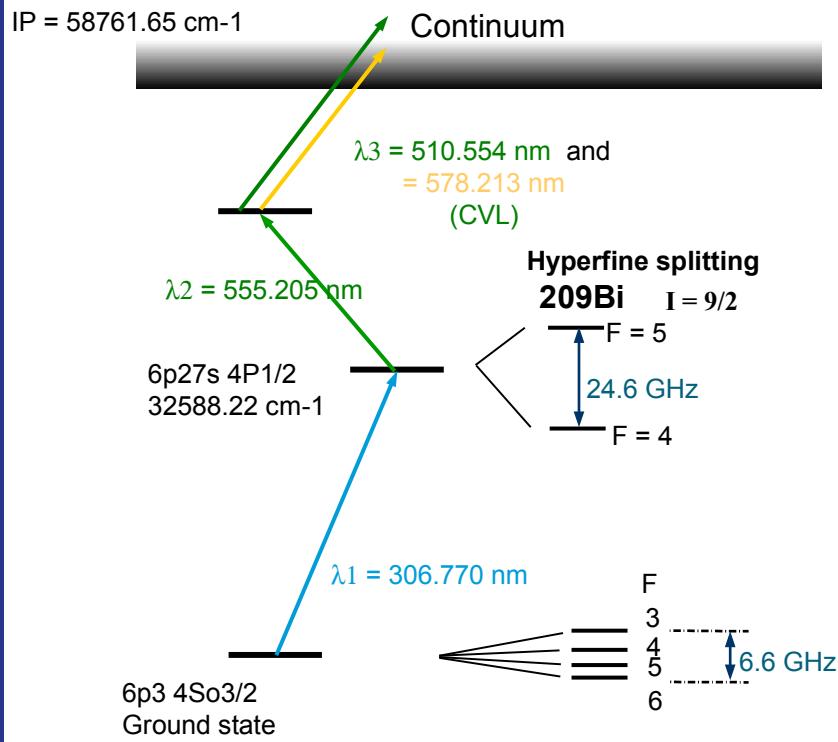
Isotope shift

$\delta \langle r^2 \rangle$

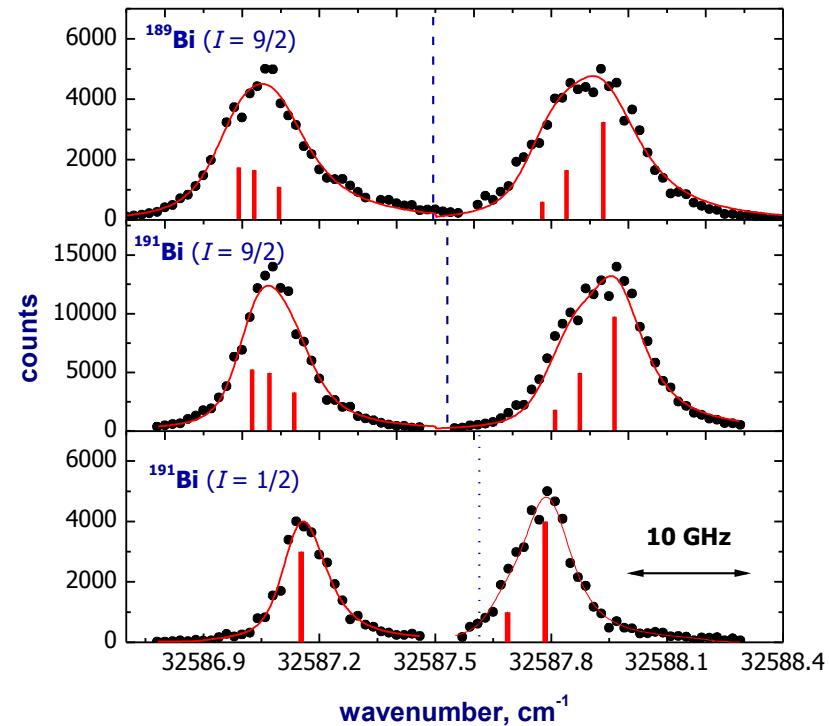


Pb: charge radii and magnetic moments





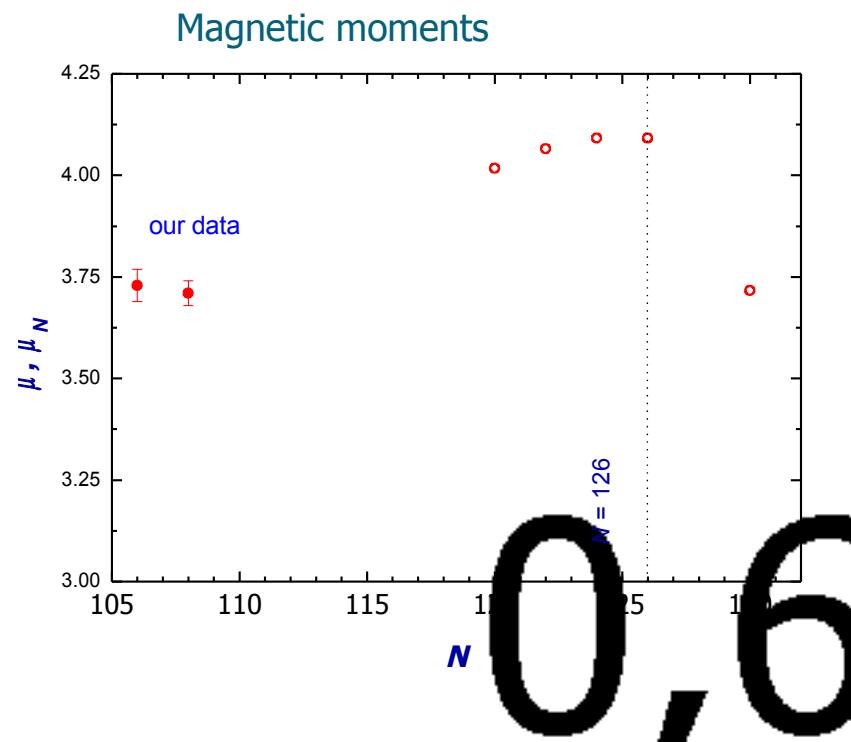
Bi



No reliable values for electronic factor
and specific mass shift constant :(

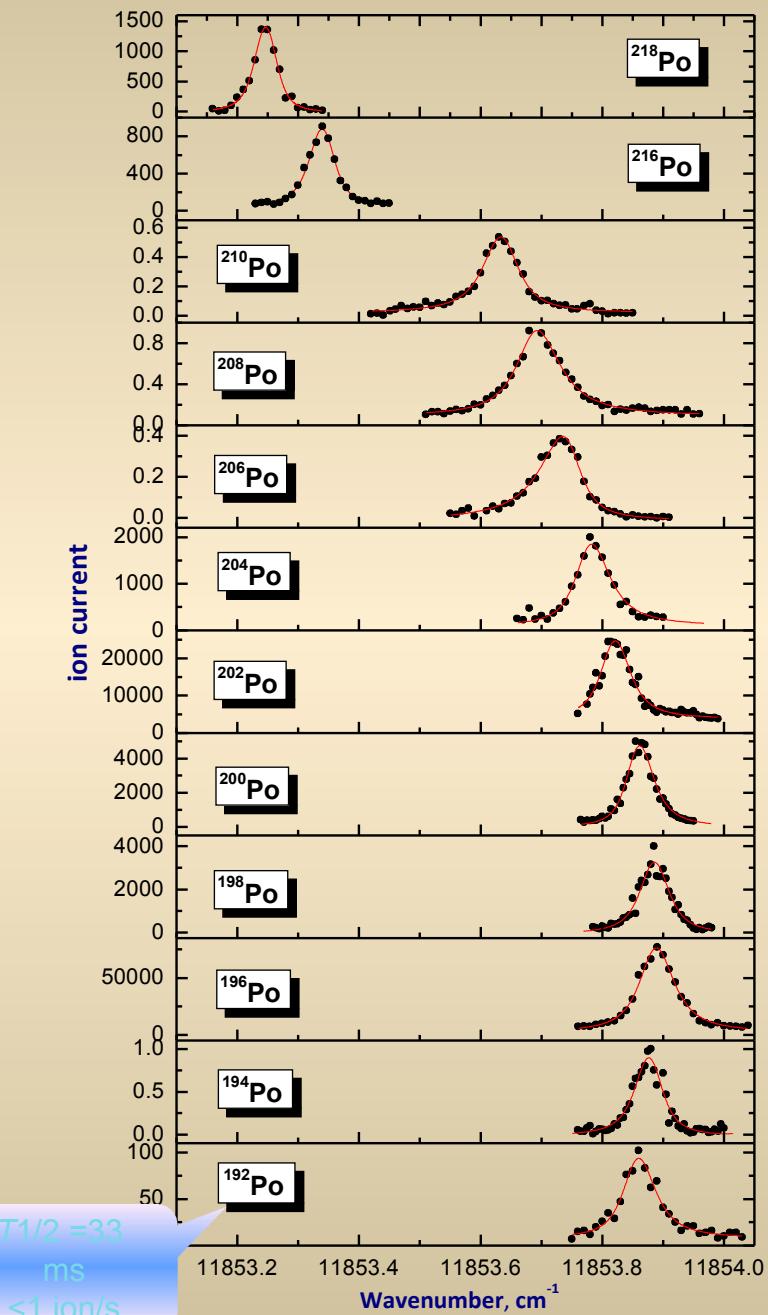
From the comparison of isotopes shifts
of Bi and Pb: **$F = 27(3)$ GHz/fm²**

*P. Campbell et al., Phys. Lett. B 346
(1995) 21*

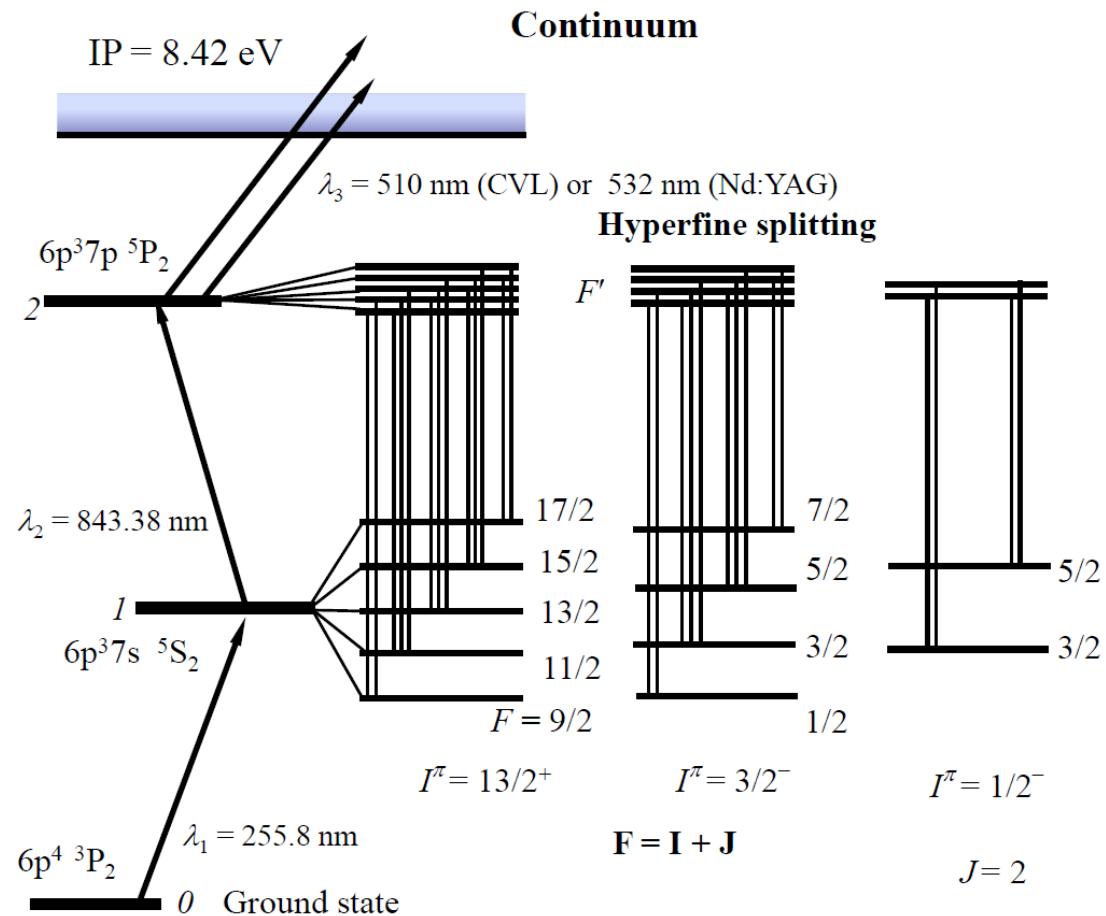


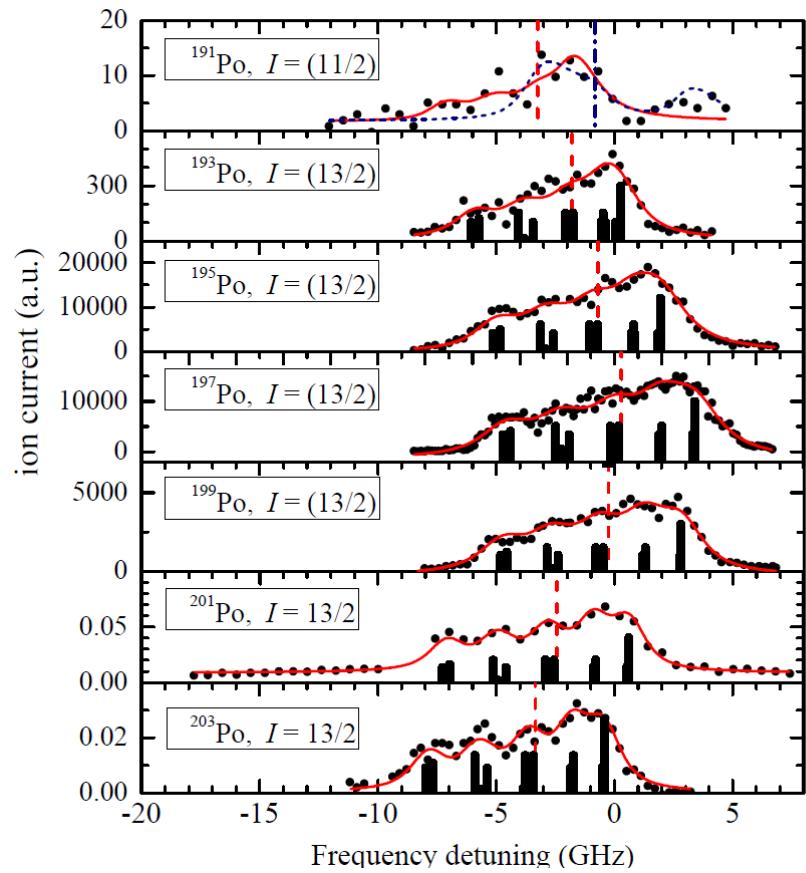
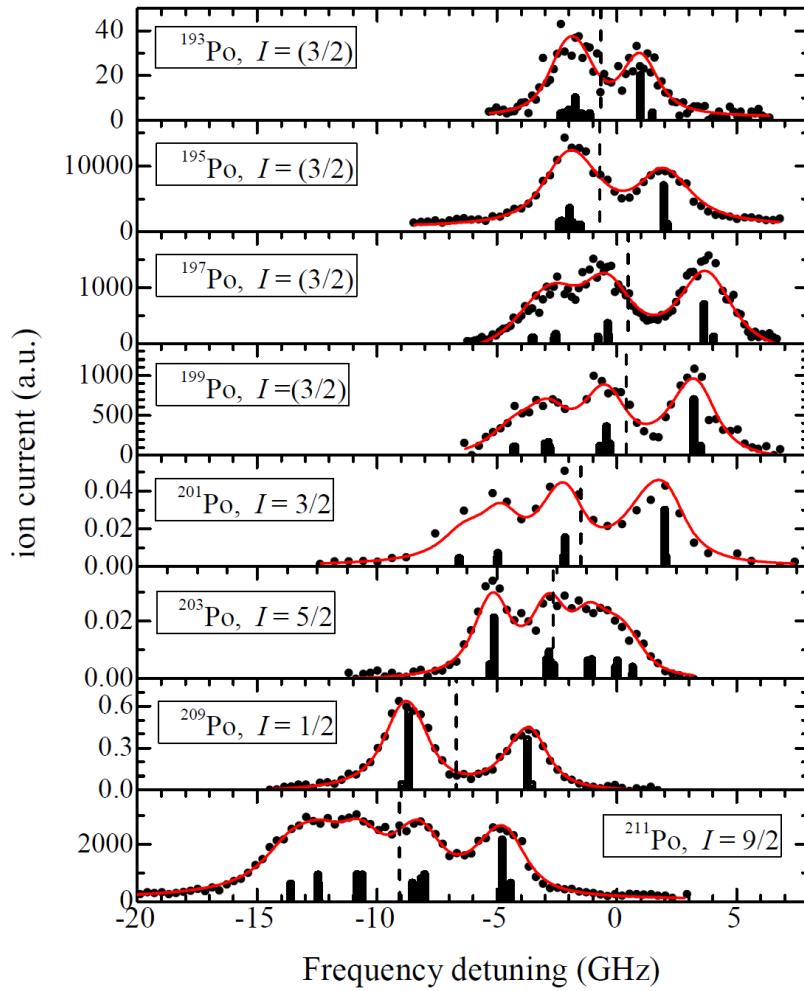
Po ($Z = 84$)

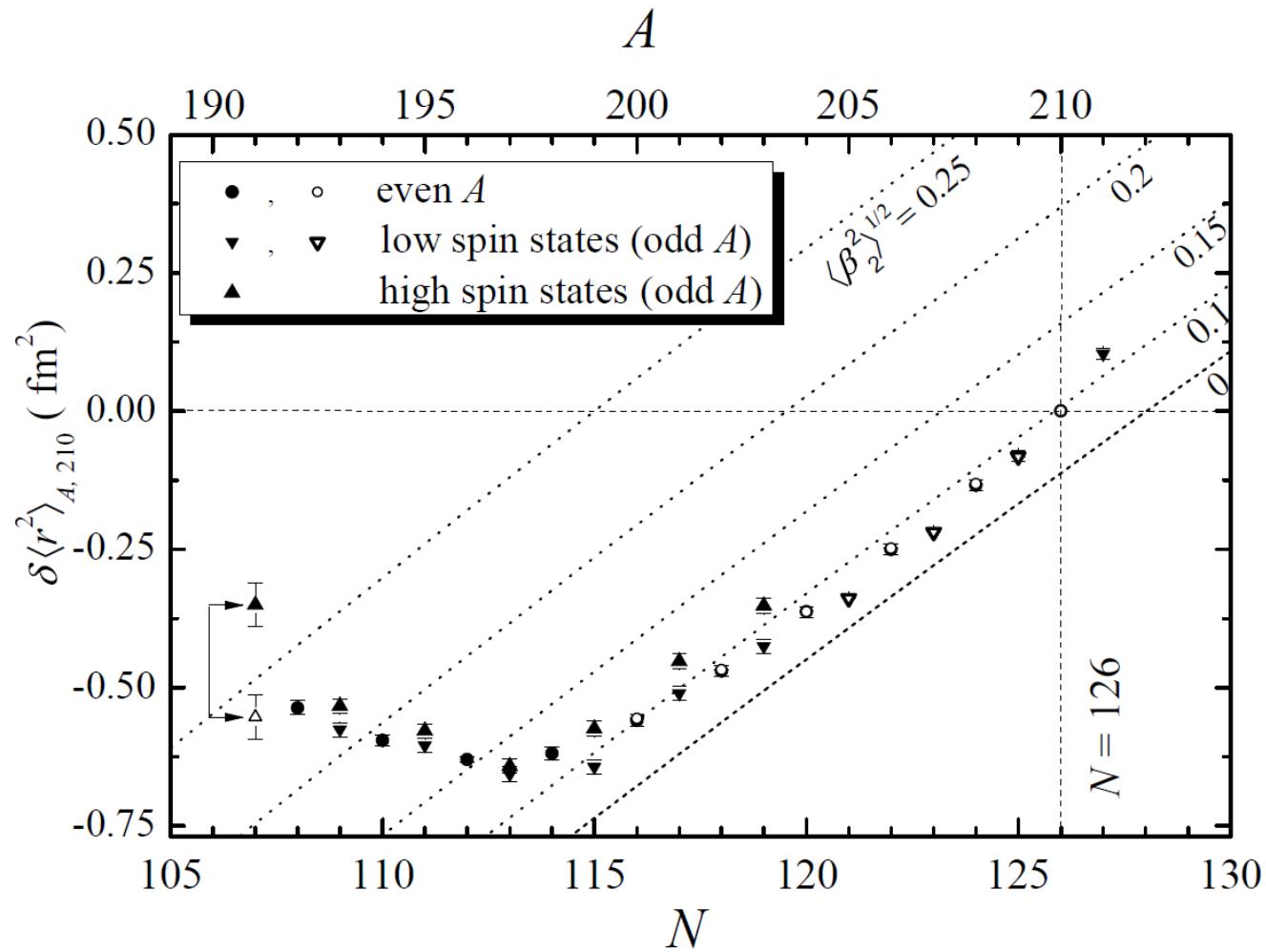
*This transition was already used for optical spectroscopy
(Kowalewska et al, Phys. Rev. A, 44 R1442, 1991.)



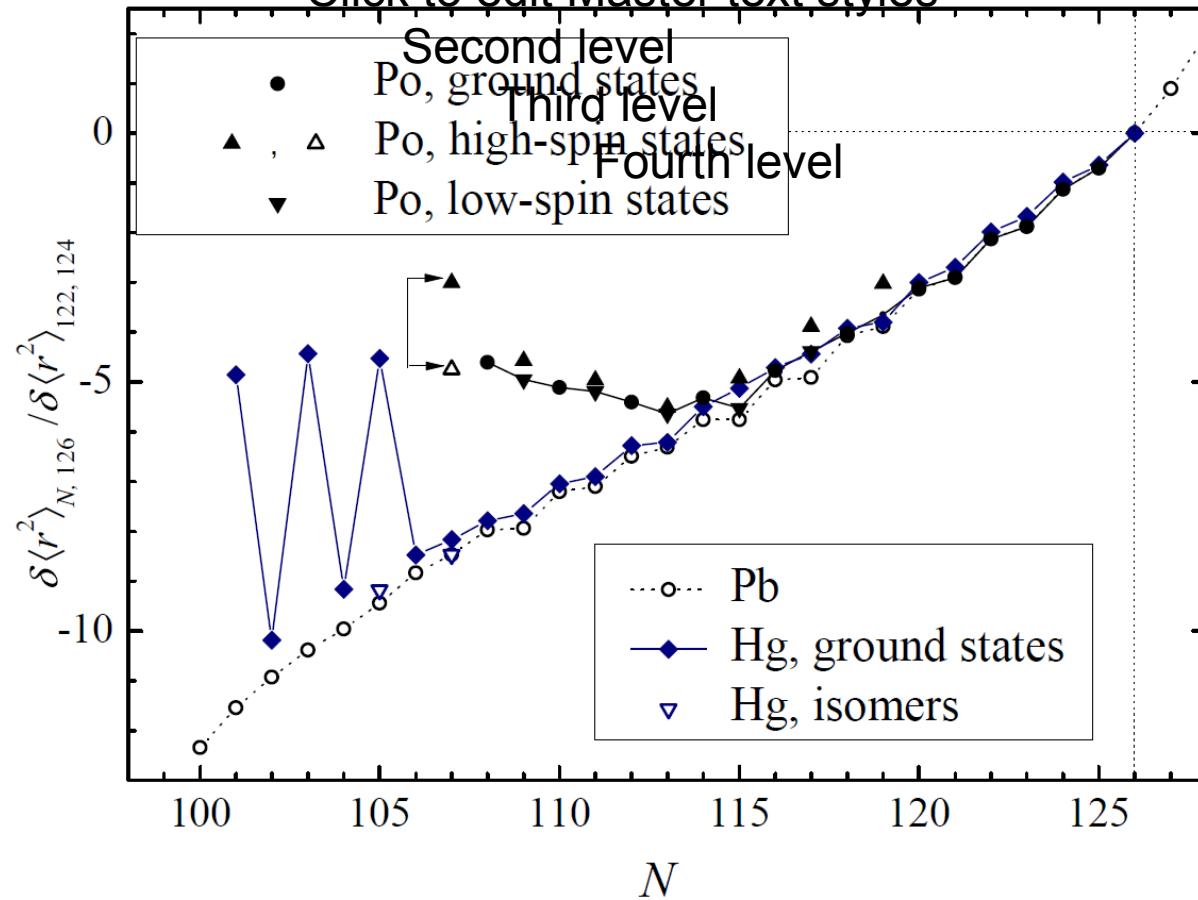
$T_{1/2} = 33$
ms
 $<1 \text{ ion/s}$



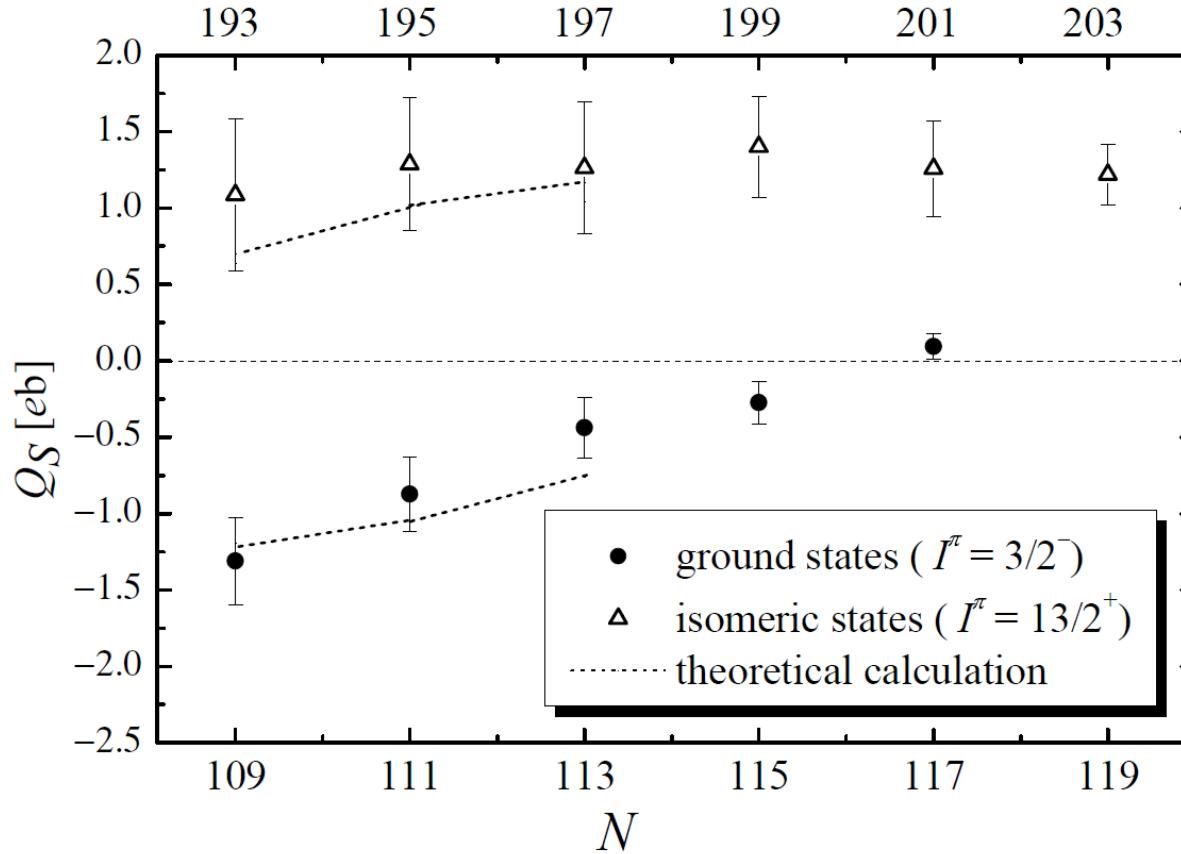


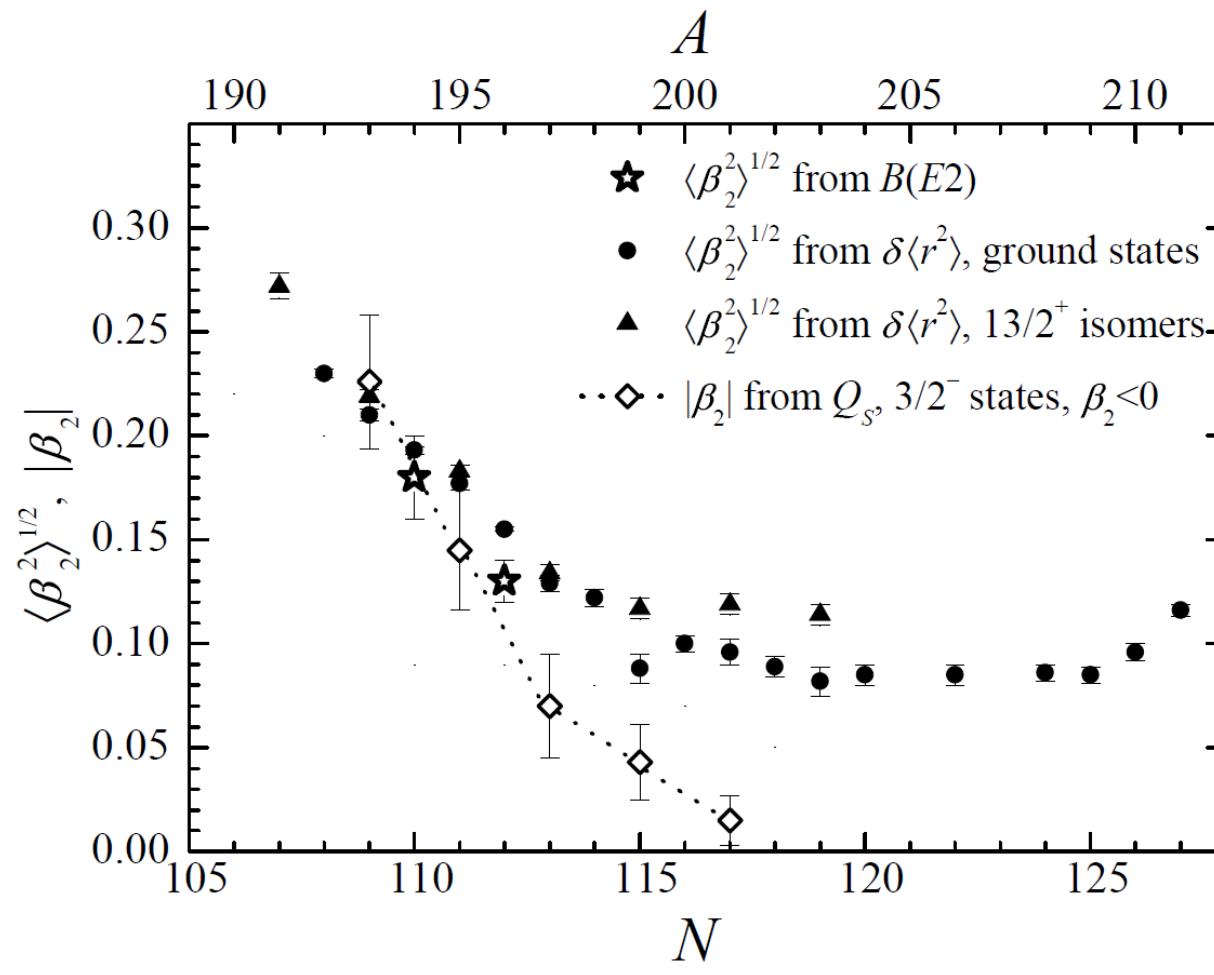


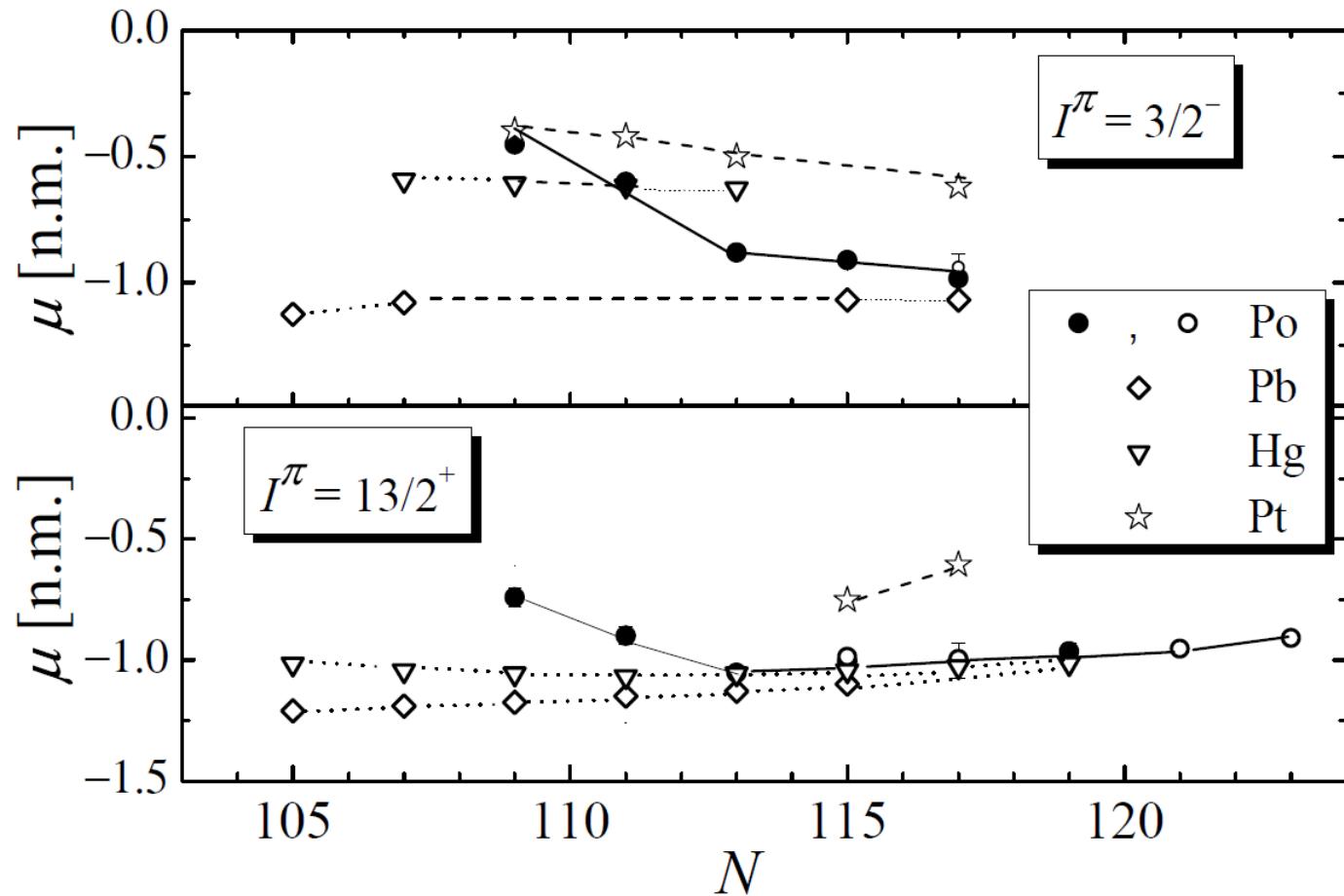
Click to edit Master text styles



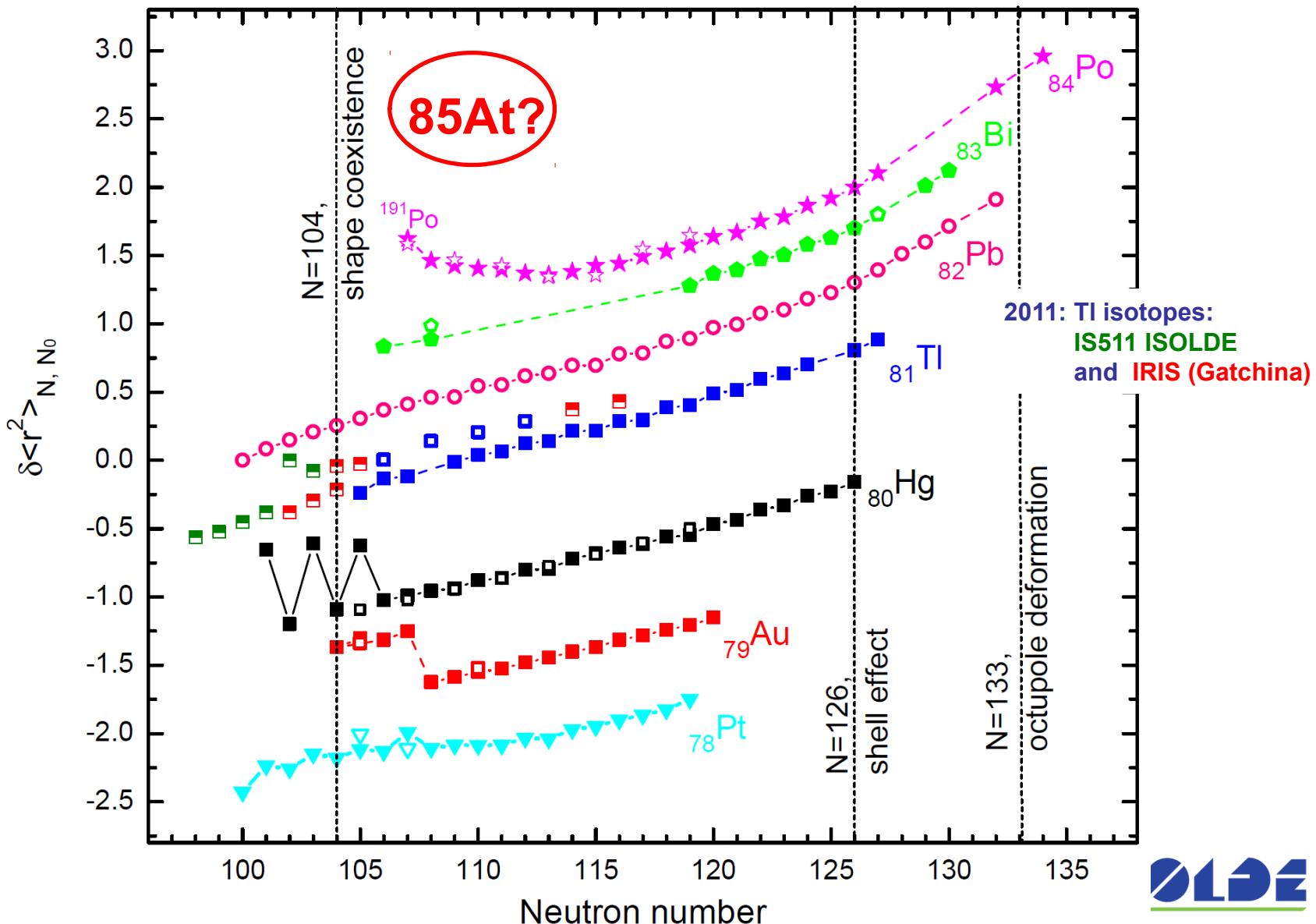
Click to edit Master text styles



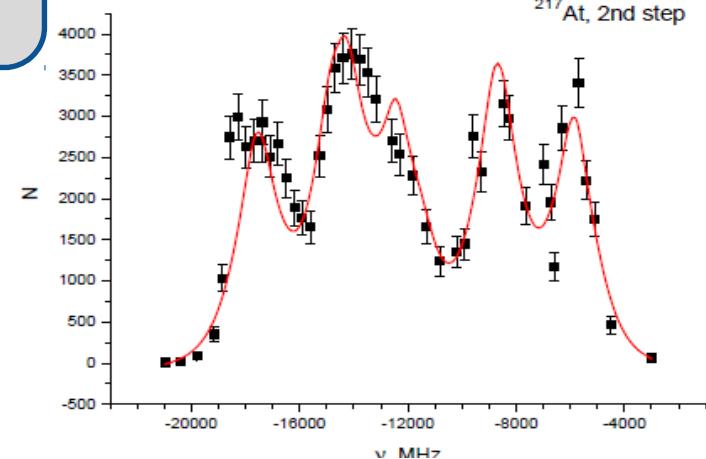
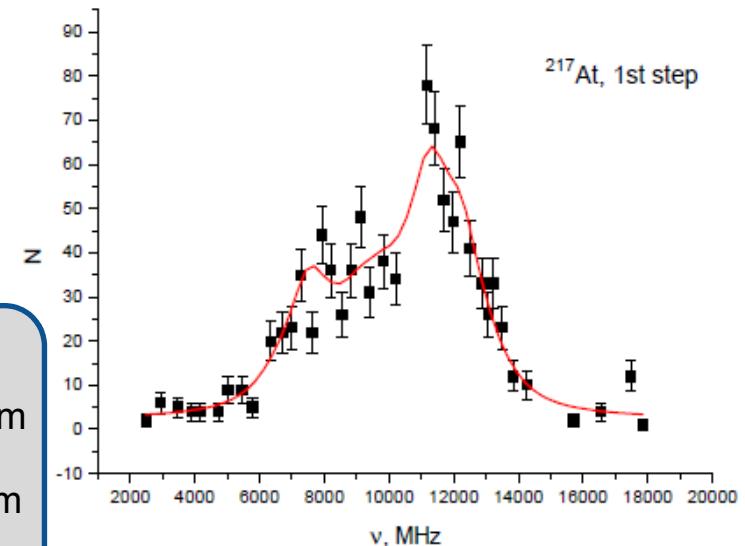
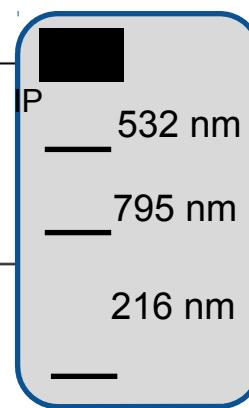
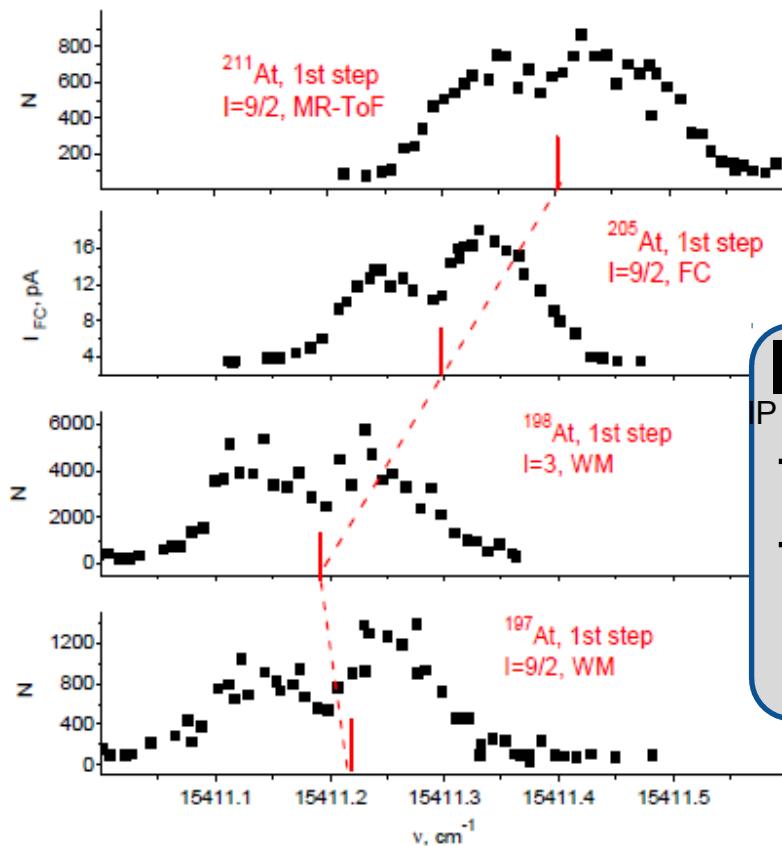




Next step: At

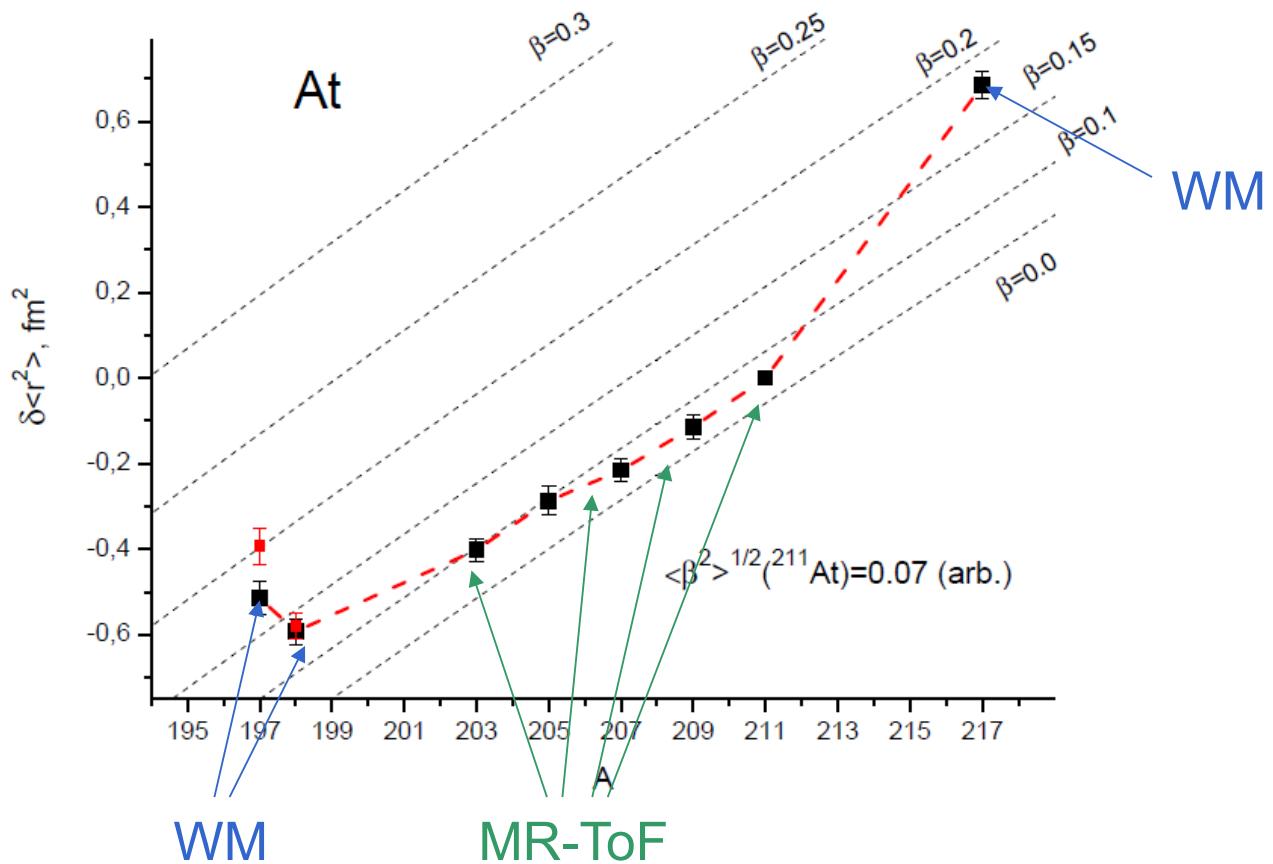


At (Z=85)

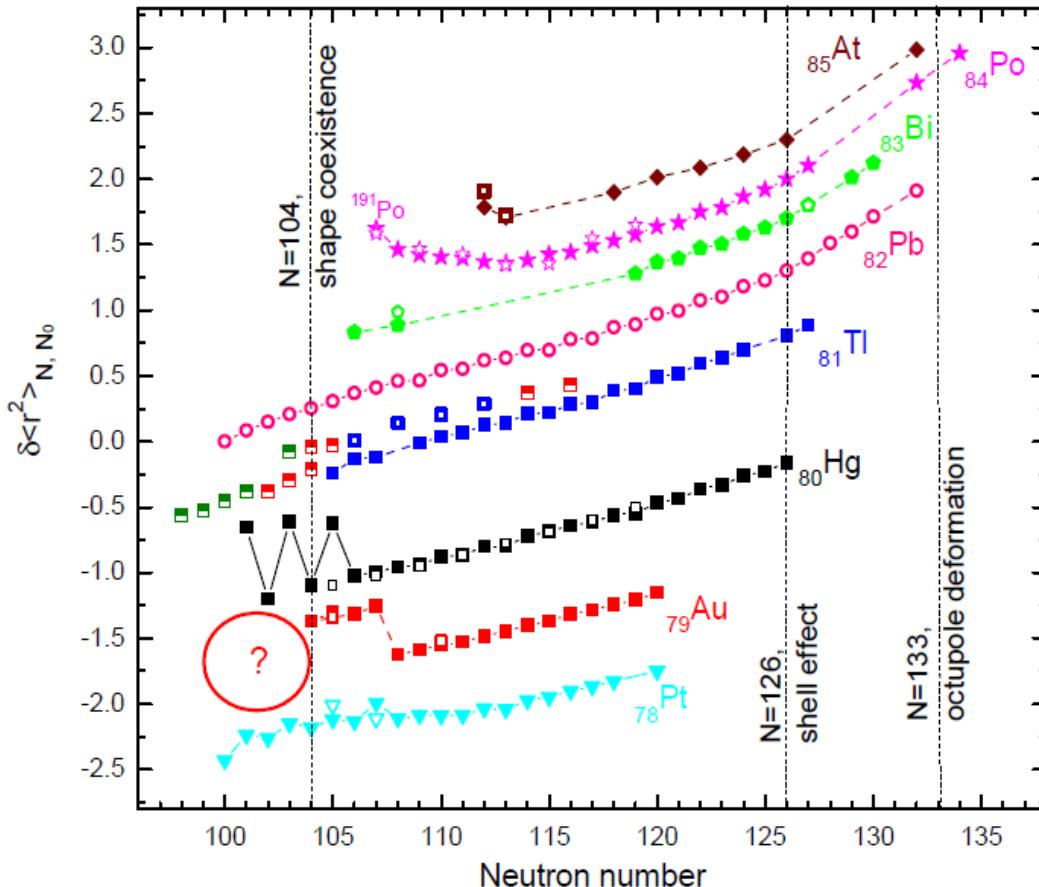


1st step scanning is better for $\Delta\langle r^2 \rangle$ extraction
2nd step scanning is better for hfs resolution
(Q and μ determination)

At: charge radii

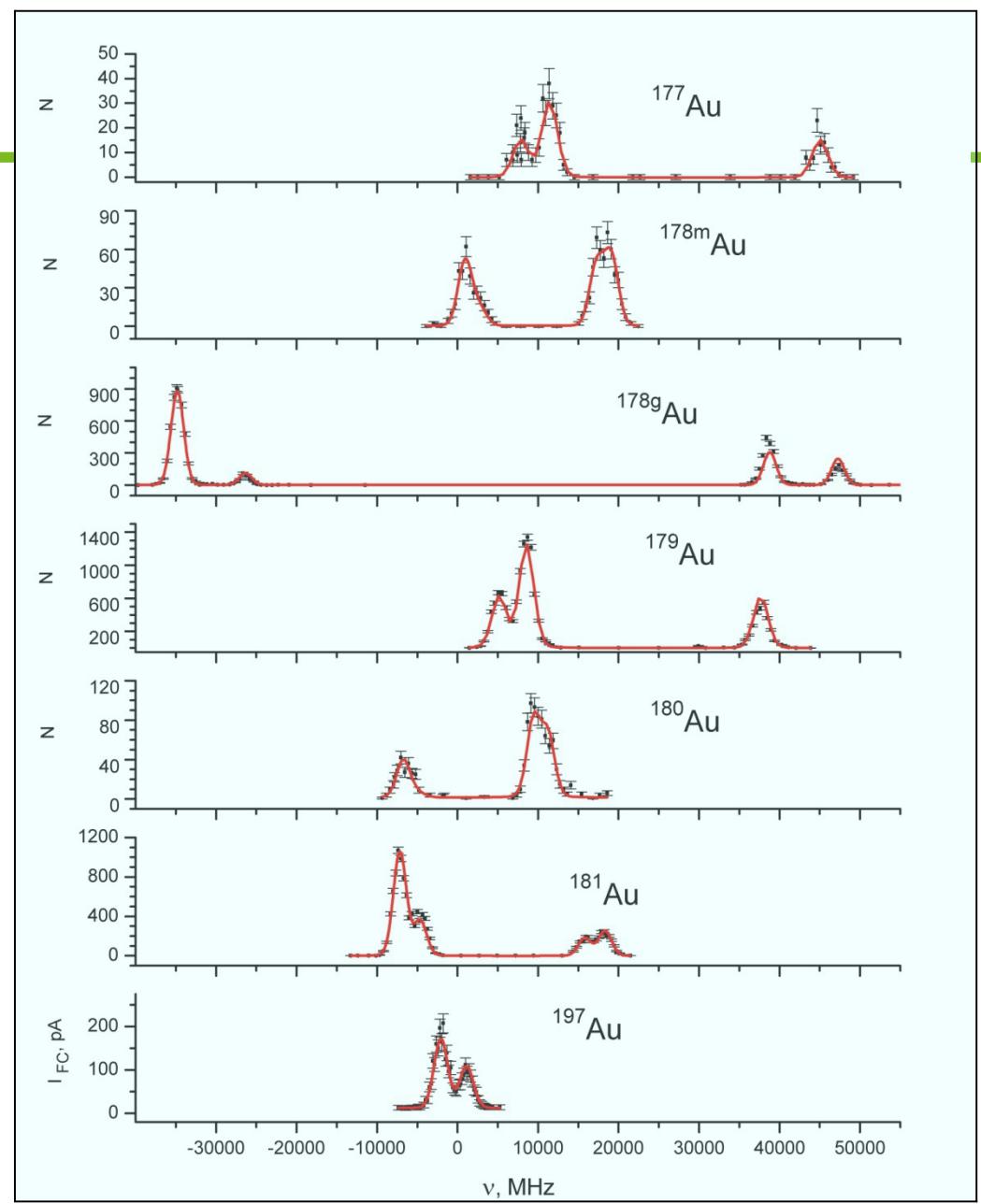
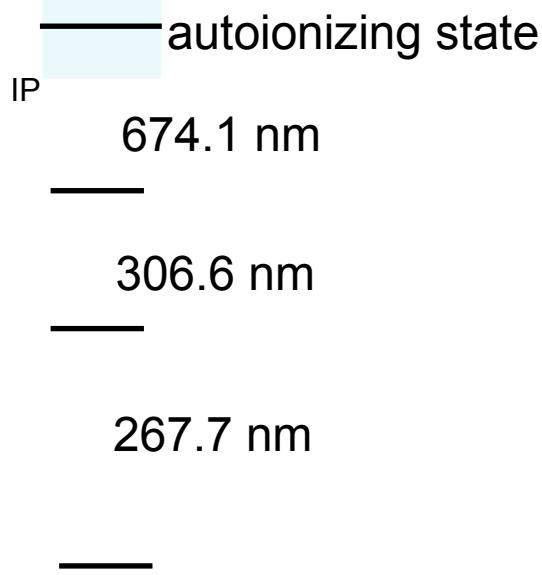


Next step: Au

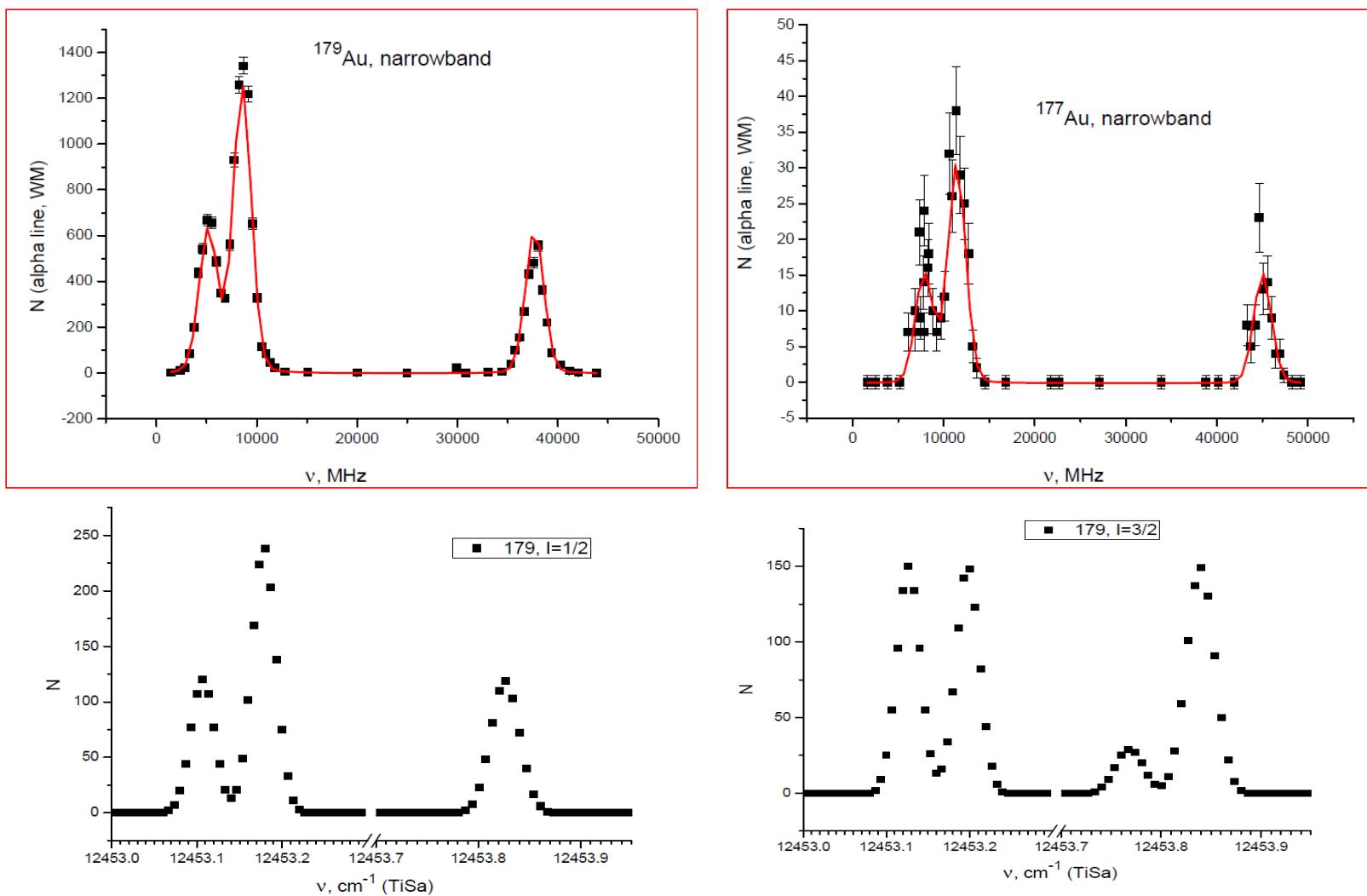


Au (Z=79)

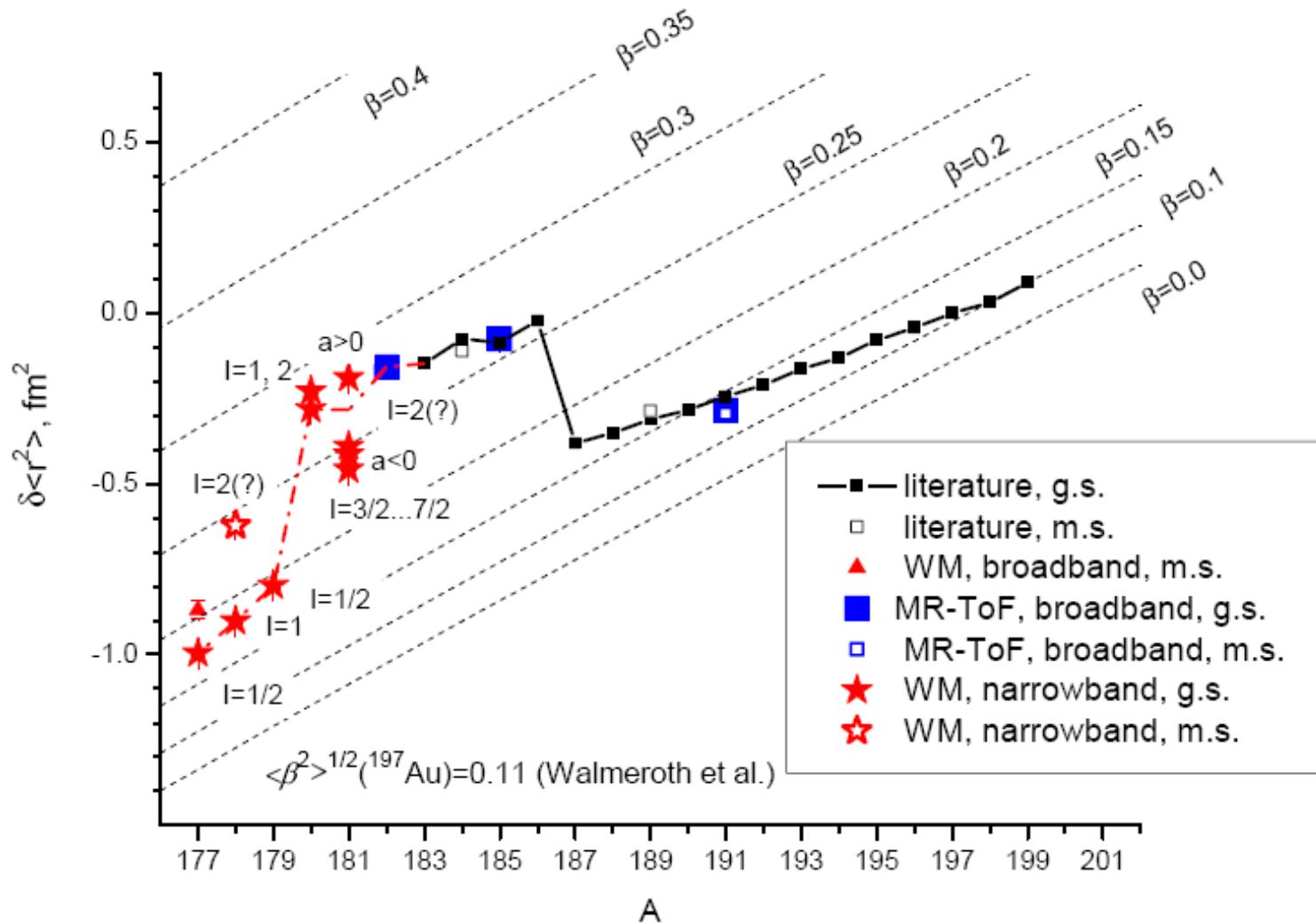
Au ionization scheme



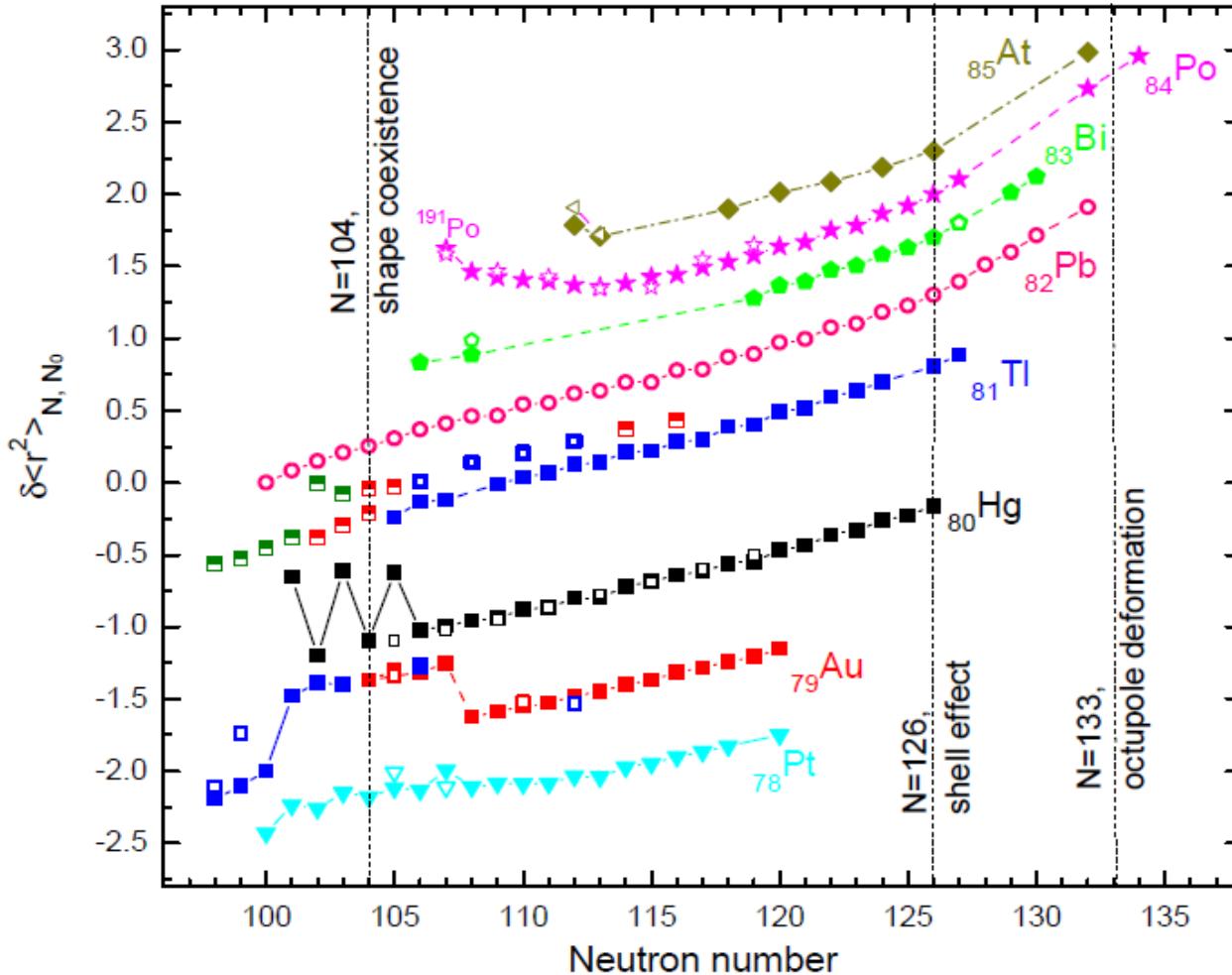
Spins of ^{177}g , ^{179}gAu



Au: charge radii



Charge radii in Pb region





Charge radii: summary

- **Pb:** 182, 183, 183m, 184, 185, 185m, 186, 187, 187m, 188, 189, 189m - *published*
- **Bi:** 189, 191, 191m *published (IS and electromagnetic moments)*
- **Po:** 191, 192, 193, 193m, 194, 195, 195m, 196, 197, 197m, 198, 199, 199m, 201, 201m, 203, 203m, 211, 216, 217, 218 *partly published*
- **Tl:** 179, 180, 181, 182, 183, 183m, 184, 184m
- **At:** 197, 197m, 198, 203, 205, 207, 209, 211, 217
- **Au:** 177, 178, 178m, 179, 180, 181

At and Au: proposal submitted

Hg: proposal in preparation



Collaboration

- ПИЯФ: А.Е. Барзах, Д.В. Фёдоров, П.Л. Молканов, Ю.М. Волков ...
- ISOLDE (CERN): B.N. Fedoseev, B. Marsh, S. Rothe, R.E. Rossel, D. Fink ...
- KU Leuven: P. van Duppen, M. Huyse, A. Andreyev, H. de Witte, T.E. Cocolios ...
- Mainz University

Рабочая группа по сотрудничеству с ЦЕРН

Расходование средств
на содержание российских специалистов на 20.09.2012
(в долларах США)

Эксперимент	Координатор	Распределение 2012 года	Истрачено по проектам	Процент истрачено	Новое распред.
ATLAS	А.М. Зайцев	817 000	538 685	65,9	915 000
CMS	В.А. Матвеев О.Ю. Лукина (FP)	817 000	562 553 + 20 177	71,3	915 000
ALICE	В.И. Манько	570 000	424 848	74,5	640 000
LHCb	А.И. Голутвин	466 000	362 970	77,9	520 000
MUCAP	А.А. Воробьев	32 000	18 924	59,1	36 000
LHC-MA	Ю.М. Иванов	70 000	31 846	45,5	80 000
LCG	В.А. Ильин	70 000	31 314	44,7	80 000
COMPASS	С.В. Донсков	92 000	56 939	61,9	105 000
DIRAC	Л.Л. Немёнов	32 000	31 888	99,7	42 000
NA61	А.Б. Курепин	32 000	18 143	56,7	36 000
NA62	В.Ф. Образцов	26 000	16 234	62,4	36 000
ICARUS	В.А. Матвеев	5 500	2 037	37,0	6 000
CAST	В.А. Матвеев	7 500	2 540	33,9	8 500
ISOLDE	Д.В. Фёдоров	6 500	5 044	77,6	7 000
RD50	А.Г. Залужный Е.М. Вербицкая	14 000	0	0	8 000 8 000
MEG	Ю.А. Тихонов	13 000	7 718	59,4	15 000
AEGIS	В.А. Матвеев	18 500	13 625	73,6	22 000
ADM+RES	В.И. Саврин	123 093	55 653	45,2	153 343
ИТОГО:		3 212 093	2 201 138	68,5	3 632 843