

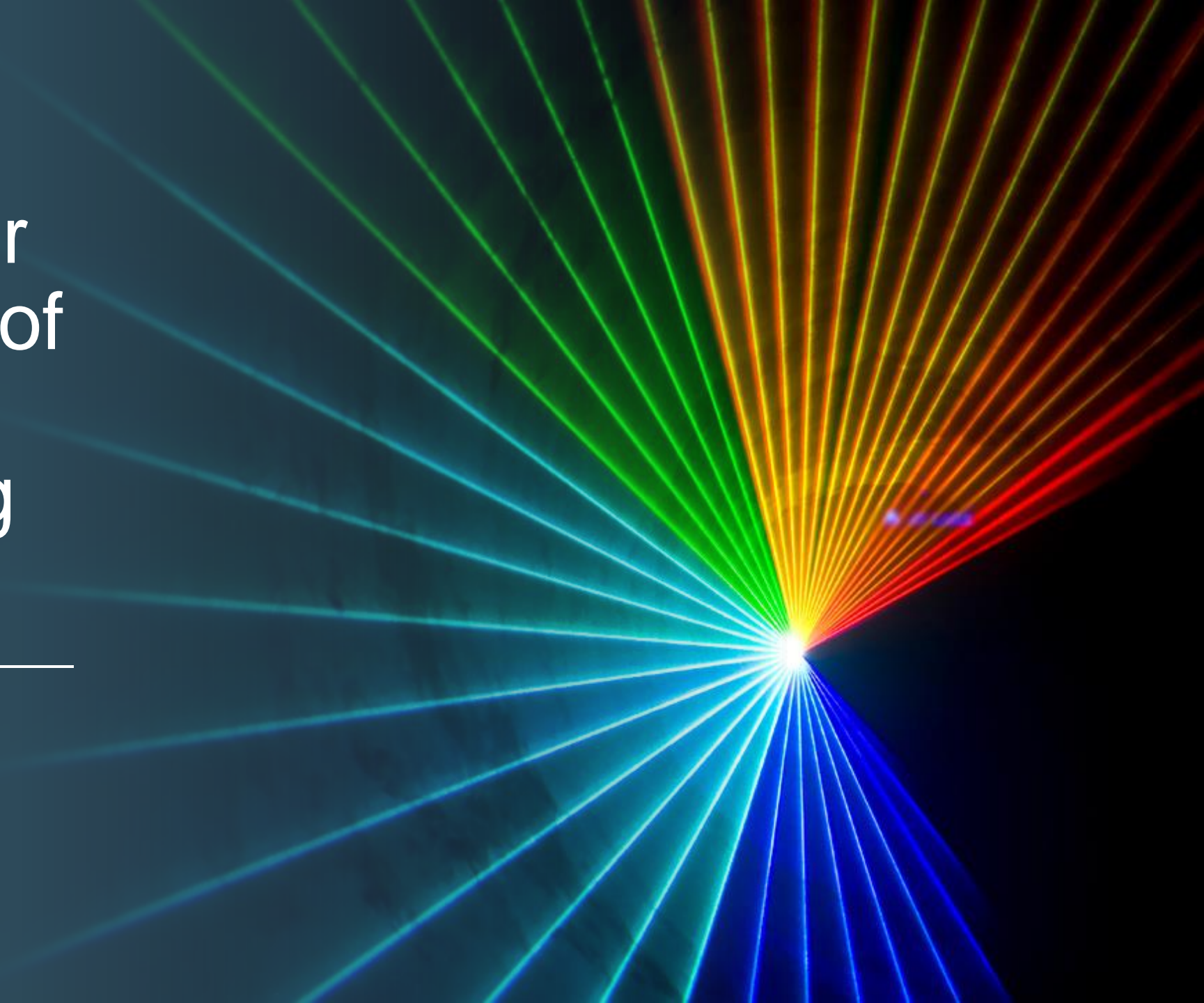
# Precision laser spectroscopy of radioactive isotopes using traps

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Ruben de Groote

EMIS XIX – 03.10. 22

Daejon, Korea



# Outline

- Laser spectroscopy of exotic isotopes
- Study of ultra-rare isotopes with particle-based detection schemes
  - Penning-trap assisted study of  $^{95,96}\text{Ag}$
  - Collinear resonance ionization spectroscopy of  $^{52}\text{K}$
- Atomic state manipulation with trapped ions
  - Optical pumping of trapped radioactive cobalt isotopes
- Development of precision spectroscopy of trapped ions
  - Future plans towards magnetic octupole moment measurements of  $^{87}\text{Sr}$

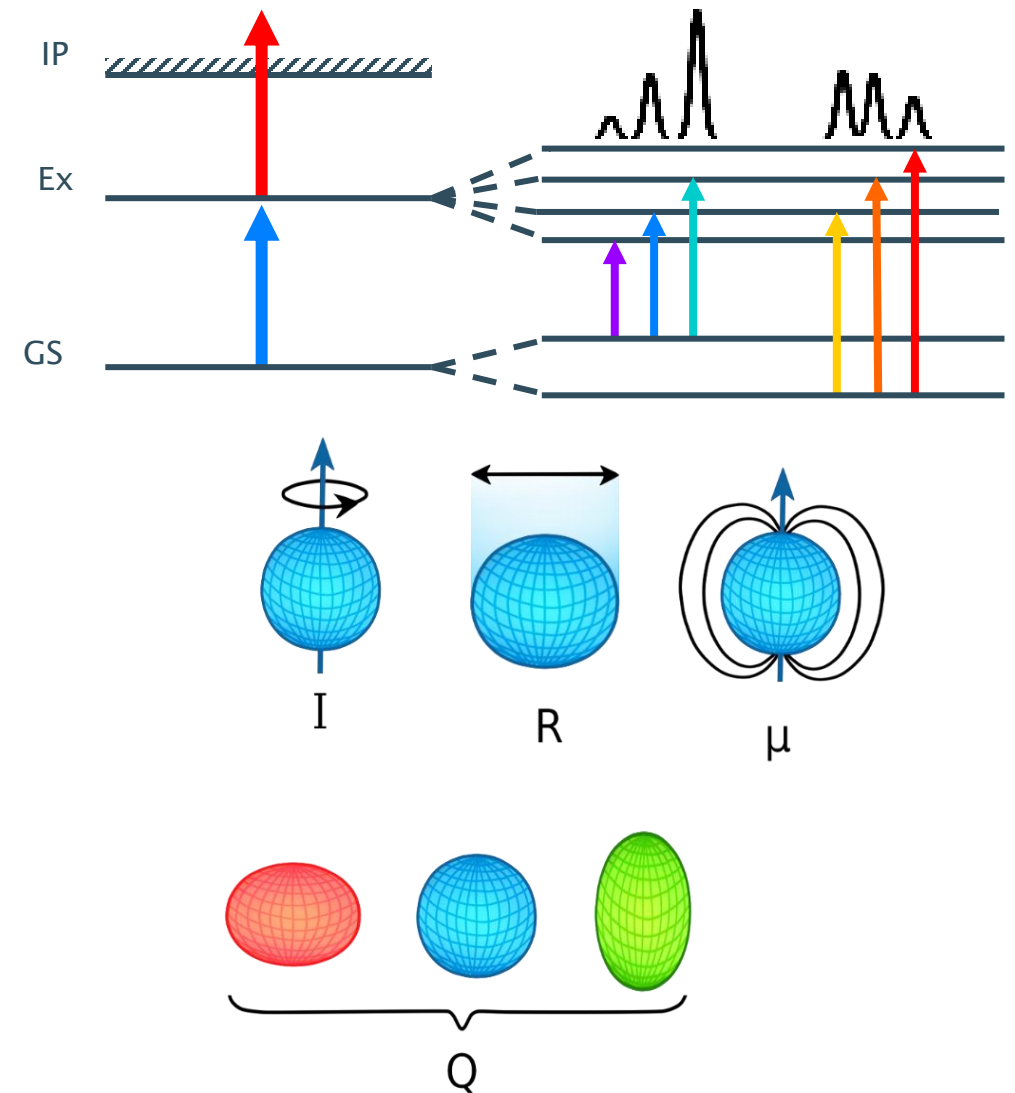
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# Some fundamentals of laser spectroscopy

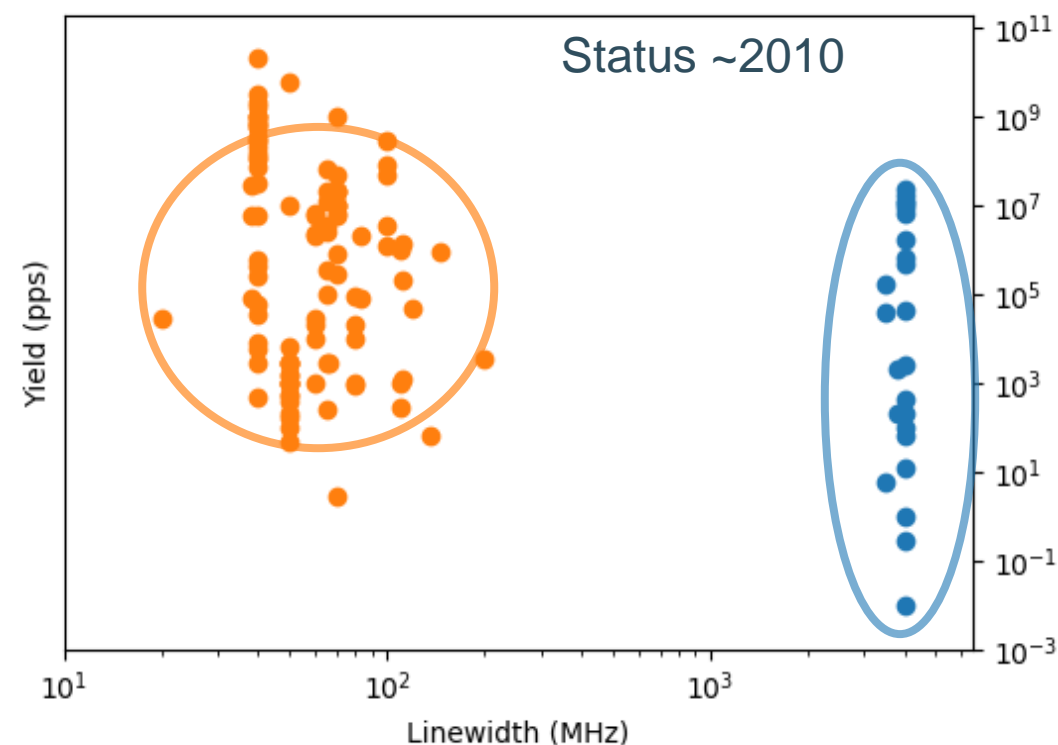
# Laser spectroscopy: study of atom and nucleus

- Electromagnetic interaction between orbiting electrons and the nucleus
  - Isotopes have a **finite size** leading to isotope shifts  
→ nuclear size
- Orbiting charges and intrinsic nucleon moments lead to **nuclear magnetic moments**  
→ valence configuration, wf purity
- Charge can assume non-spherical charge distribution, and thus **electric moments**  
→ deformation, shape, collectivity



# Laser spectroscopy approaches

- In-source laser spectroscopy
  - **Hot cavity**
  - Gas cell
- Collinear laser spectroscopy
  - **Fluorescence detection**
  - **Resonance ionization**
  - Beta-NMR/collisional ionization/state-selective charge exchange/...
- Methods tailed to specific cases
  - **Spectroscopy on trapped ion/atoms** (faaaar to the left on this diagram)



# Study of ultra-rare isotopes with particle- based detection schemes

# Spectroscopy very far from stability

- Many scientific questions require the study of nuclei far from stability
- This means we are forced to deal with a double-whammy of pain:
  - Production rates of ions of interest will be low;
  - Production rates of isobaric contamination will be high.
- The challenge thus lies in devising a spectroscopic scheme that is **efficient** and **immune to the influence of contaminants**.



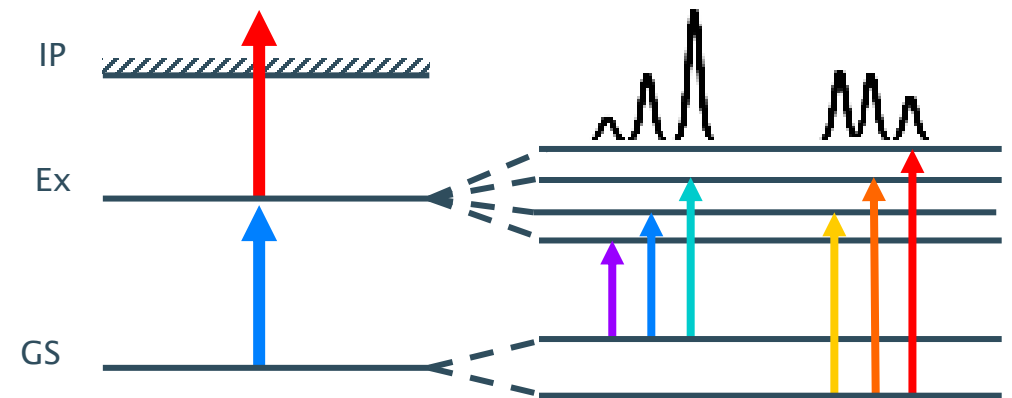
# Penning-trap mass spectrometry as an isobar remover

## Laser ionization spectroscopy

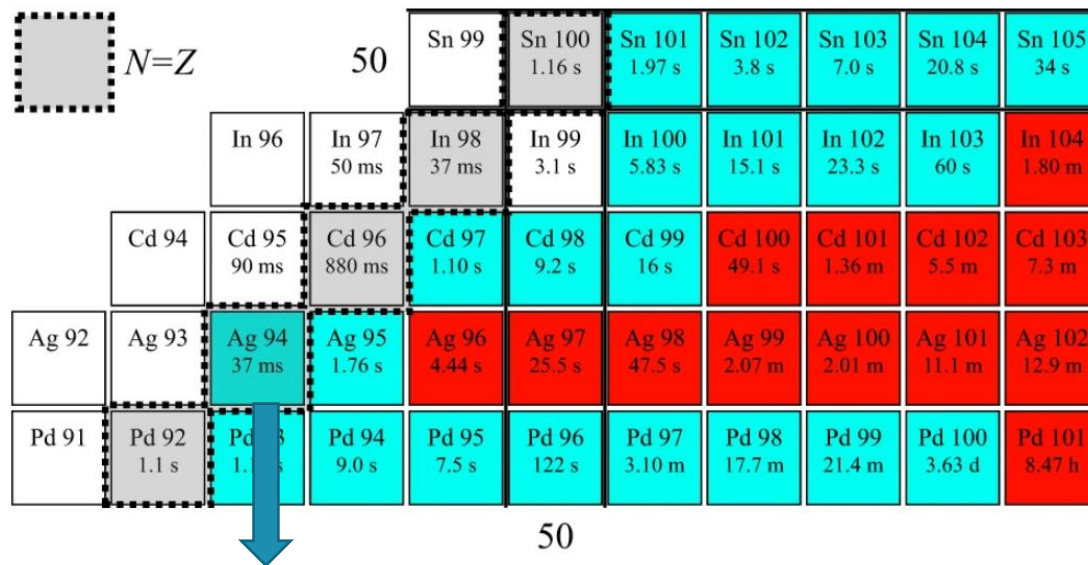
- Multi-photon, multi-color process to stepwise excite and ionize an atom
- This process is very **efficient**, and **selective**.
- The photon resonance condition is translated into the **creation of an ion**

This ion can then be further manipulated!

- By injecting these laser-induced ions into a Penning trap mass spectrometer, influence of isobars can be **completely removed**.



# Penning-trap assisted RIS of silver isotopes



One of the most fascinating isotopes on the chart.

- N=Z
- High-spin isomerism
- Double proton decay?

- Production mechanism?

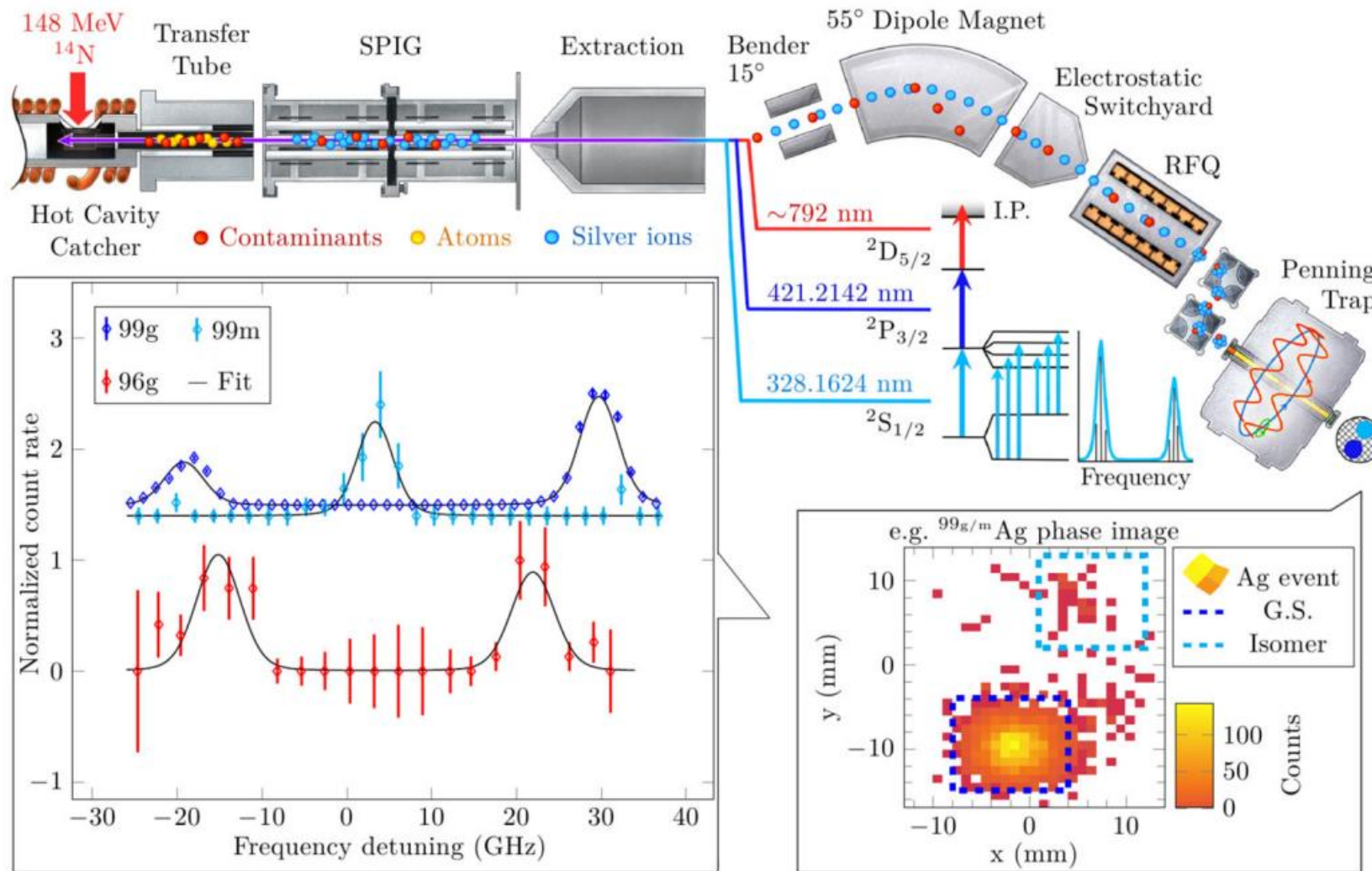


- Silver isotopes: dip well below pps
- Other isotopes: much greater quantities.
- $^{\text{nat}}\text{Mo}$  get knocked out of target => more contamination

=> Easily thousands of ions per second, of which  $\ll 1$  pps of interest.

=> High-resolution separator required!

# Penning-trap assisted RIS of silver isotopes

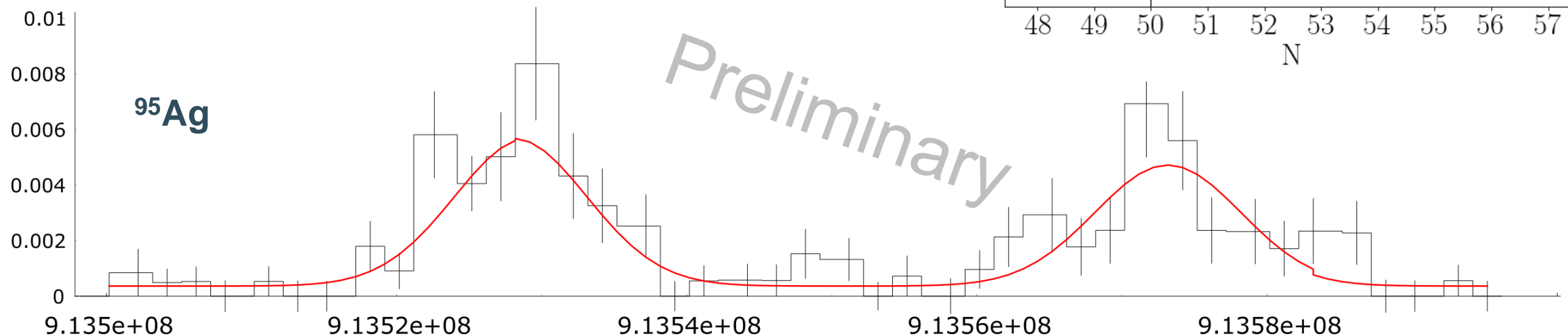
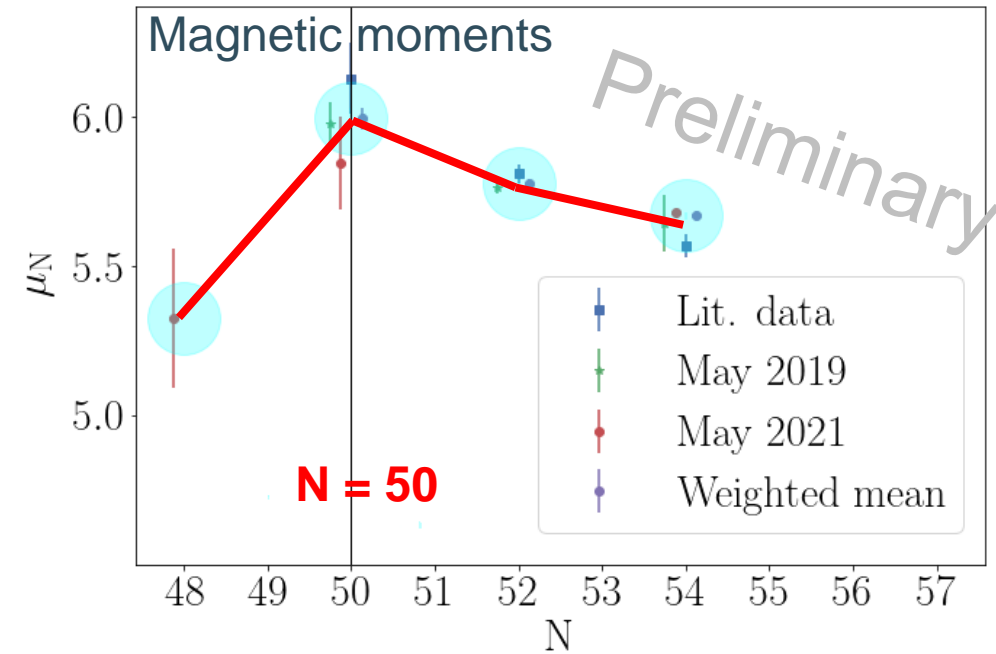


PI-ICR:  
Poster Isaac Yandow

# Penning-trap assisted RIS of silver isotopes

- $^{104-96}\text{Ag}$  using  $^{14}\text{N}(^{92}\text{Mo}, 2\text{pxn}) \text{Ag}$   
and  
 $^{96,95}\text{Ag}$  using  $^{40}\text{Ca}(^{58/60}\text{Ni}, \text{pxn}) \text{Ag}$
- *Totally* background-free measurements  
Production rate: **< 1 per minute**  
Detection rate: 1 per 20 minutes...

See poster  
M. Reponen!



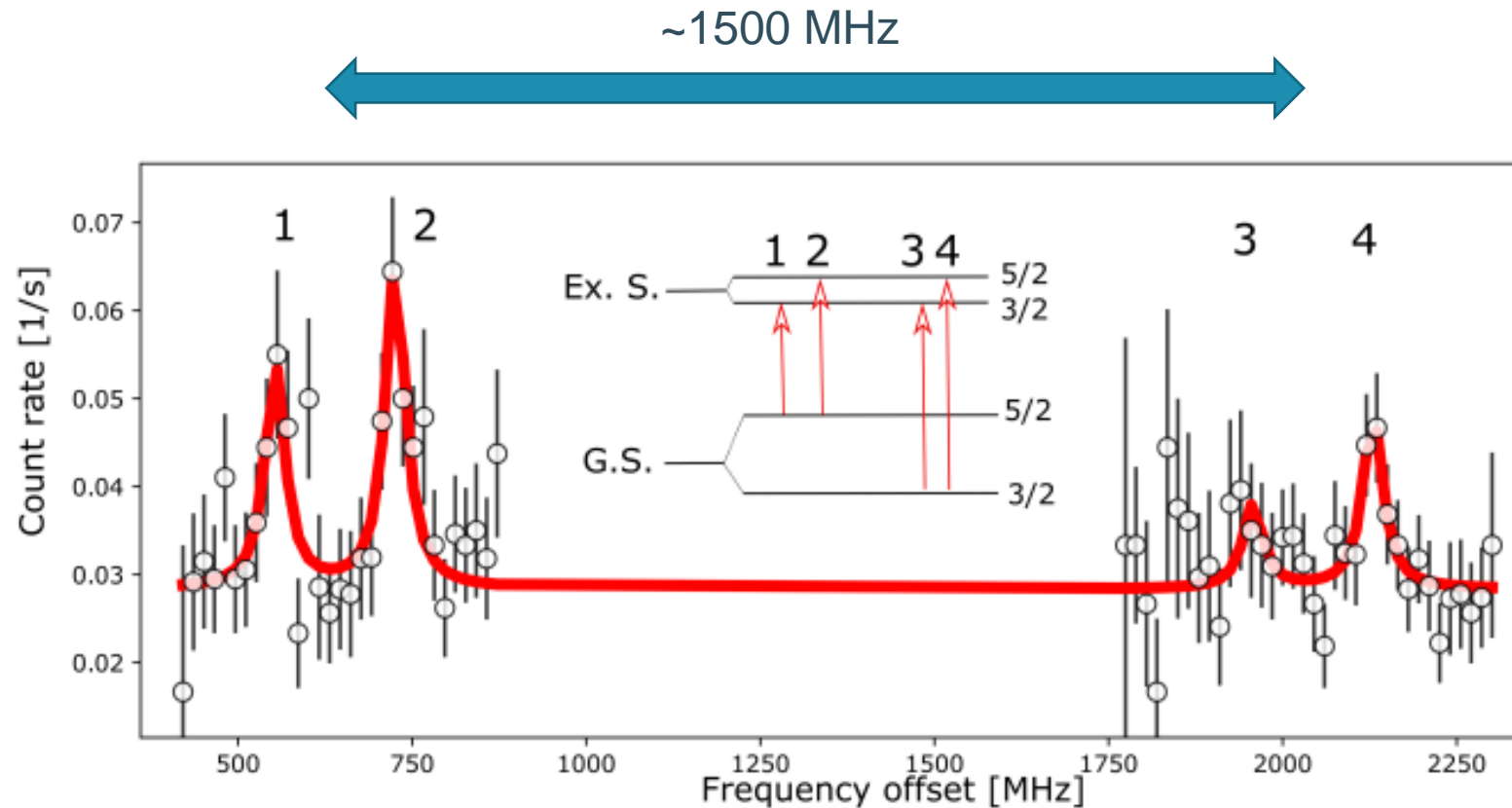
# Collinear resonance ionization spectroscopy of potassium

- Study of potassium isotopes beyond the proposed shell closure of  $N=32$ 
  - Issue: production rates below 1000/s
  - Issue: large isobaric chromium content
  - Issue: small hyperfine structure, small isotope shift
  - **Solution: collinear geometry, efficient RIS, use of decay detection**
- Laser ionization spectroscopy can be implemented on fast ion beams as well in a collinear geometry:  
**Collinear Resonance Ionization Spectroscopy**

*See talks by*

- *M. Athanasakis-Kaklamanakis – CRIS at ISOLDE (this session)*
- *S. Kujanpää – CRIS at IGISOL, Finland (next session)*
- Advantage: **removal of broadening effects** enhances the resolution of the measurement
  - In-source spectroscopy (without modern adaptation): cannot study all elements and extract all information... ☹

# Results

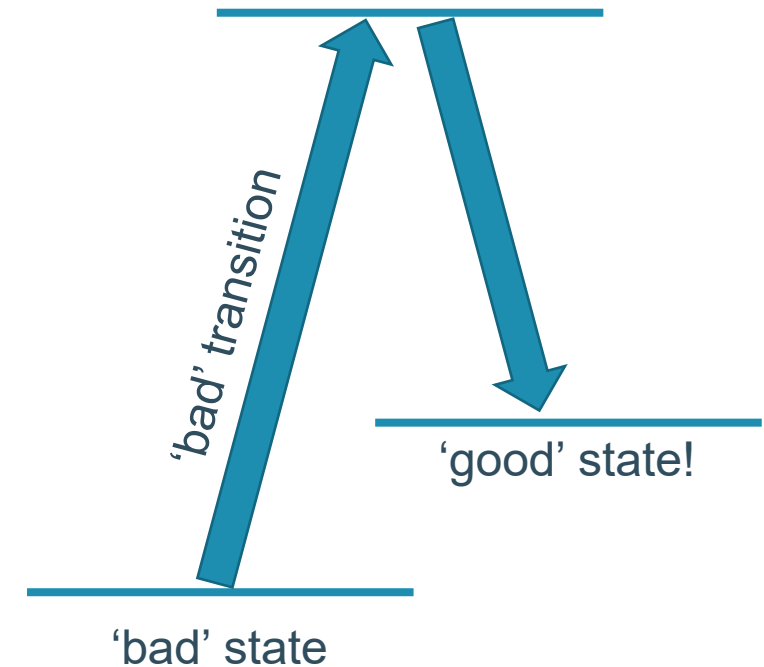


- 300 pps  $^{52}\text{K}$
- **>10<sup>6</sup> pps stable contamination**
- Beta-detection: remove stable contaminants
- < 1 day of data!!
- (background in part due to faulty electronics - 'user error'....!)

# Atomic state manipulation with trapped ions

# Atomic state manipulation

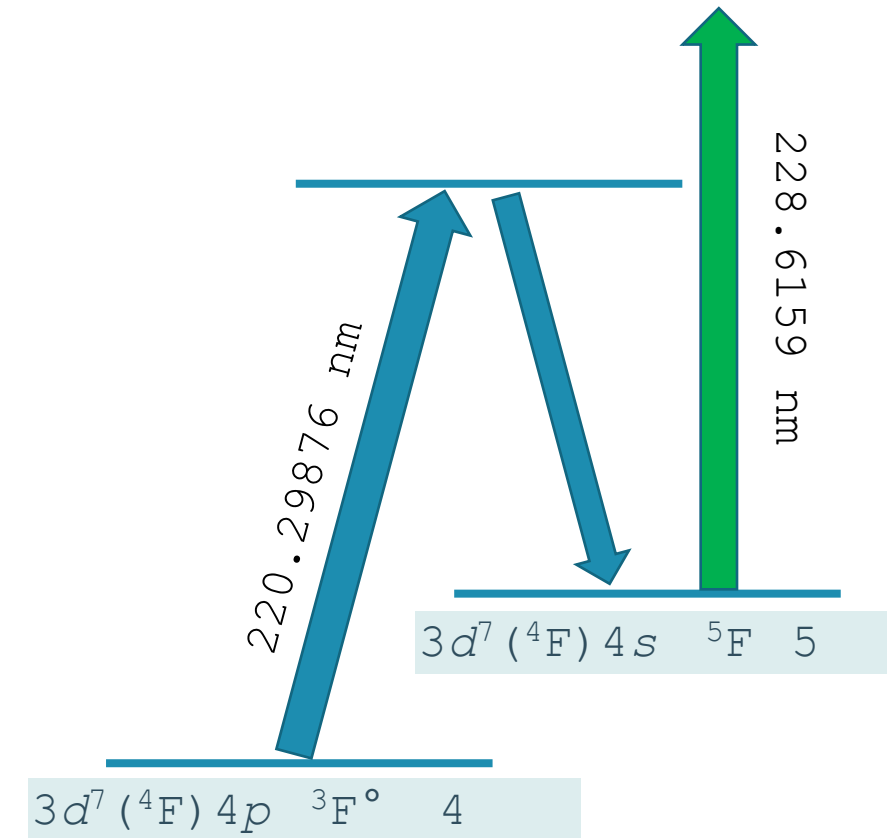
- Sometimes the state the atom is produced in is not suitable for spectroscopy
  - Atomic spin may be too low to access moments;
  - There may not be transitions accessible with current laser technology;
  - Transitions may not be sufficiently strong, may not have sensitivity to the nuclear size.
- Through a careful choice of optical process, we can change the state of the atom
  - Optical pumping: through many cycles of excitation and spontaneous decay, the electron eventually ends up in a more favourable atomic state.
  - This process takes *time*. Thus, it is applied to ions inside an ion trap.
  - First developed and used in IGISOL laboratory  
*B. Cheal et al, PRL **102**, 222501, 2009*





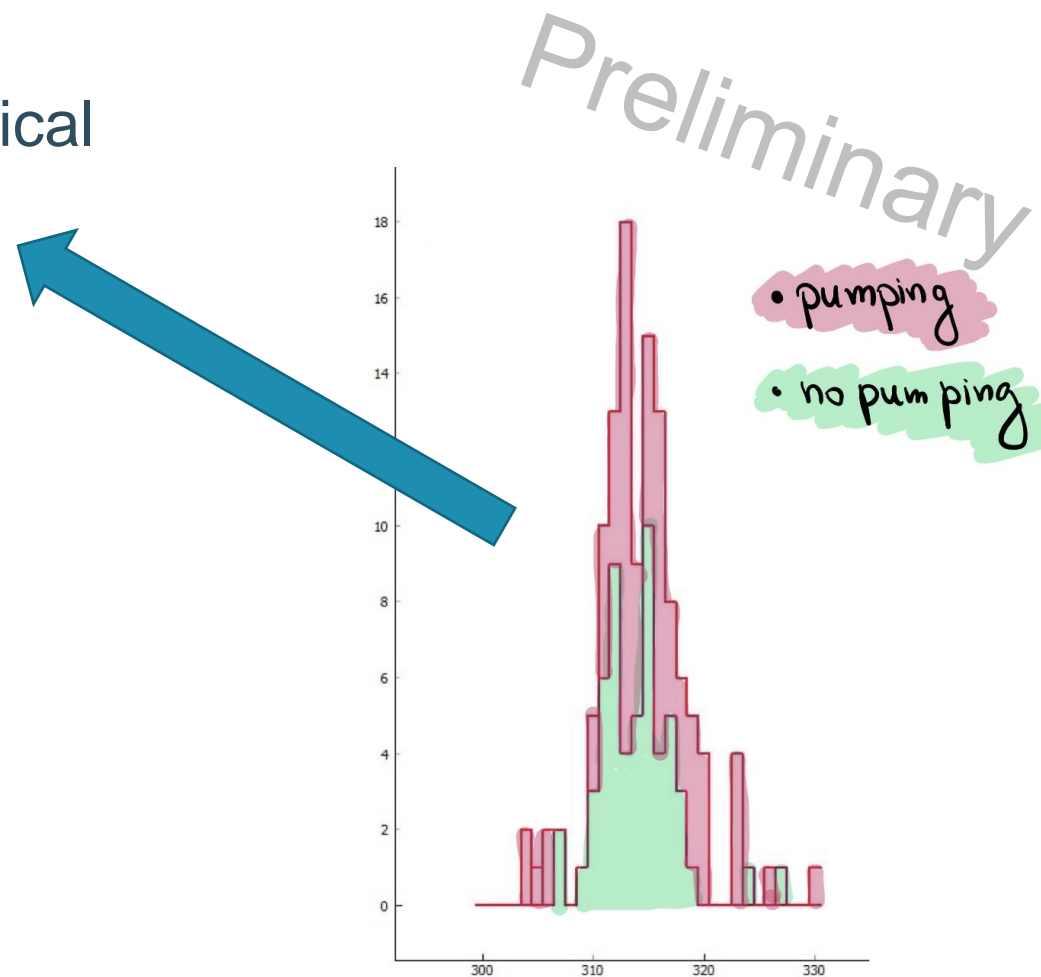
# Laser spectroscopy of optically pumped cobalt

- With 27 protons, cobalt is an ideal probe of the magic nickel isotopes
  - Radii, moments, ... would provide lots of insight!
- But, Co chain never studied with laser spectroscopy
  - Refractory: not available in ISOLDE
  - Complex atomic/ionic system:
    - Lots of metastable states
    - deep-UV lasers required
    - Ionic ground-state not very useful...
- Optical pumping at 221 nm with 10 kHz pulsed-TiSa laser
- Spectroscopy at 229 nm



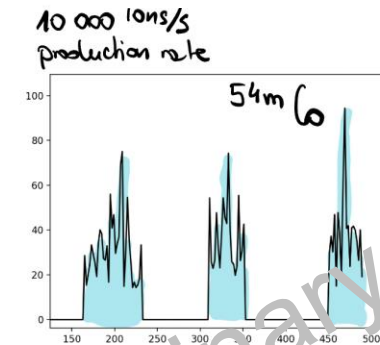
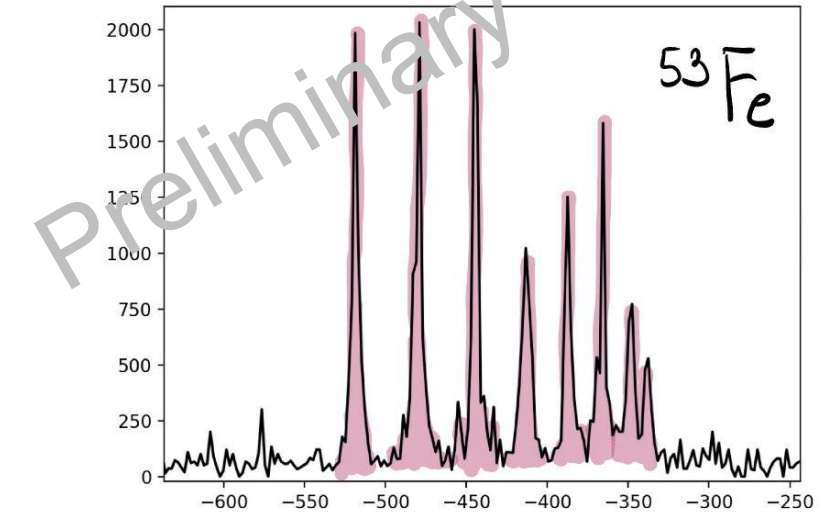
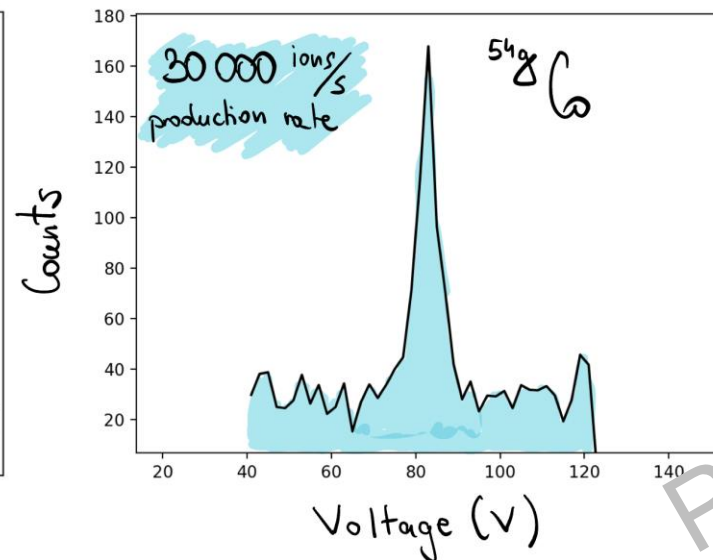
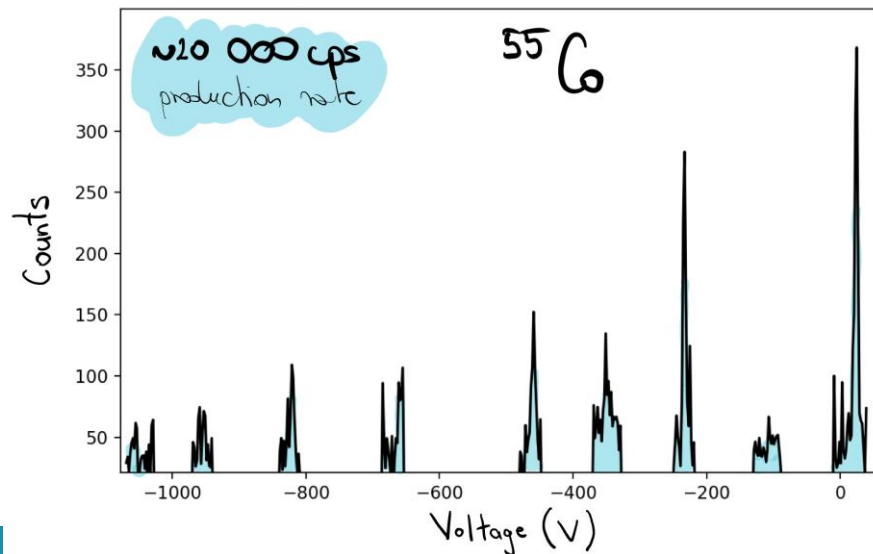
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- Factor 3 enhancement of signal due to optical pumping
- Enabled study of  $^{54,55,57-59}\text{Co}$ !



# Laser spectroscopy of optically pumped cobalt

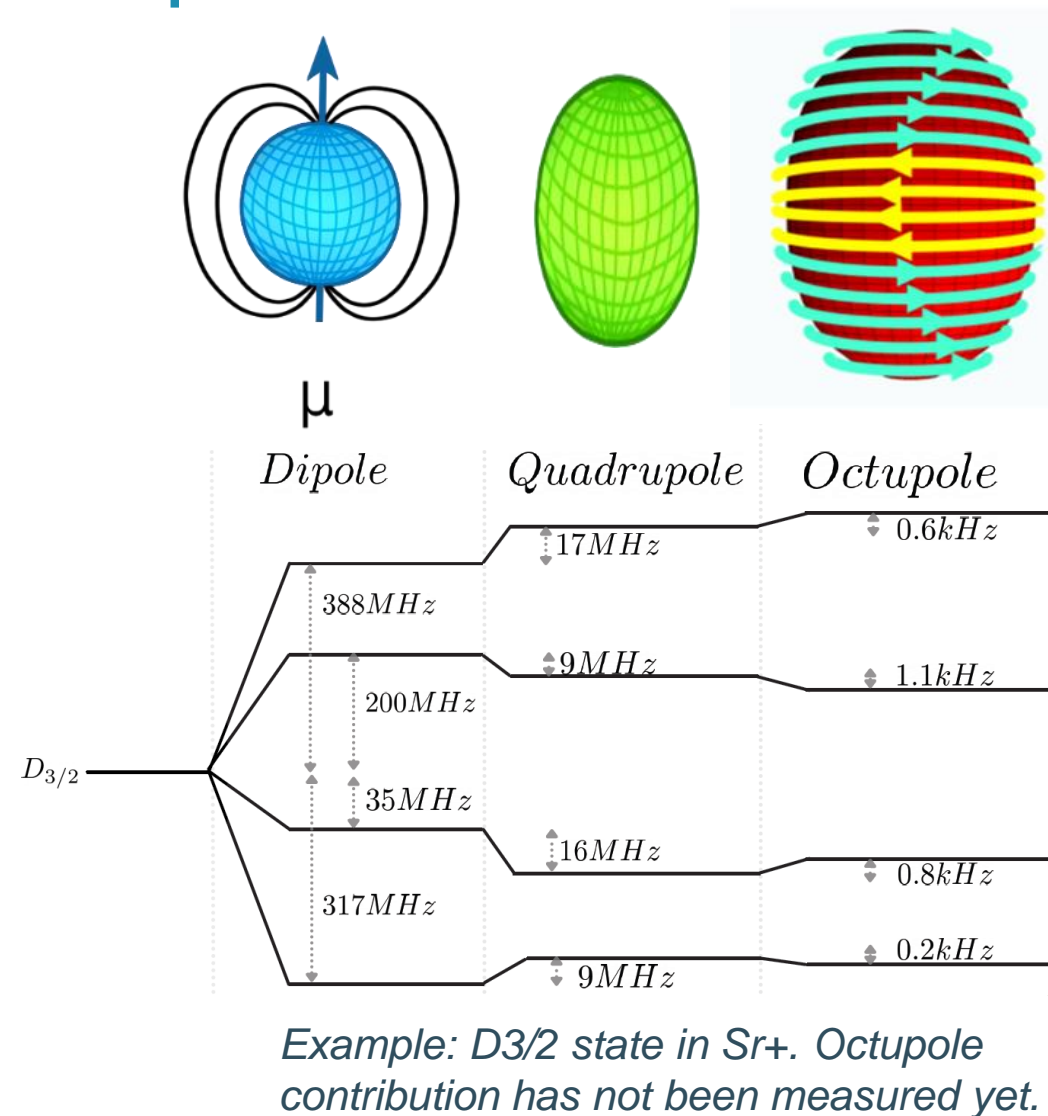
- Factor 3 enhancement of signal due to optical pumping
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- Same experiment:  $^{53-58}\text{Fe}$ !



# Development of precision spectroscopy of trapped ions

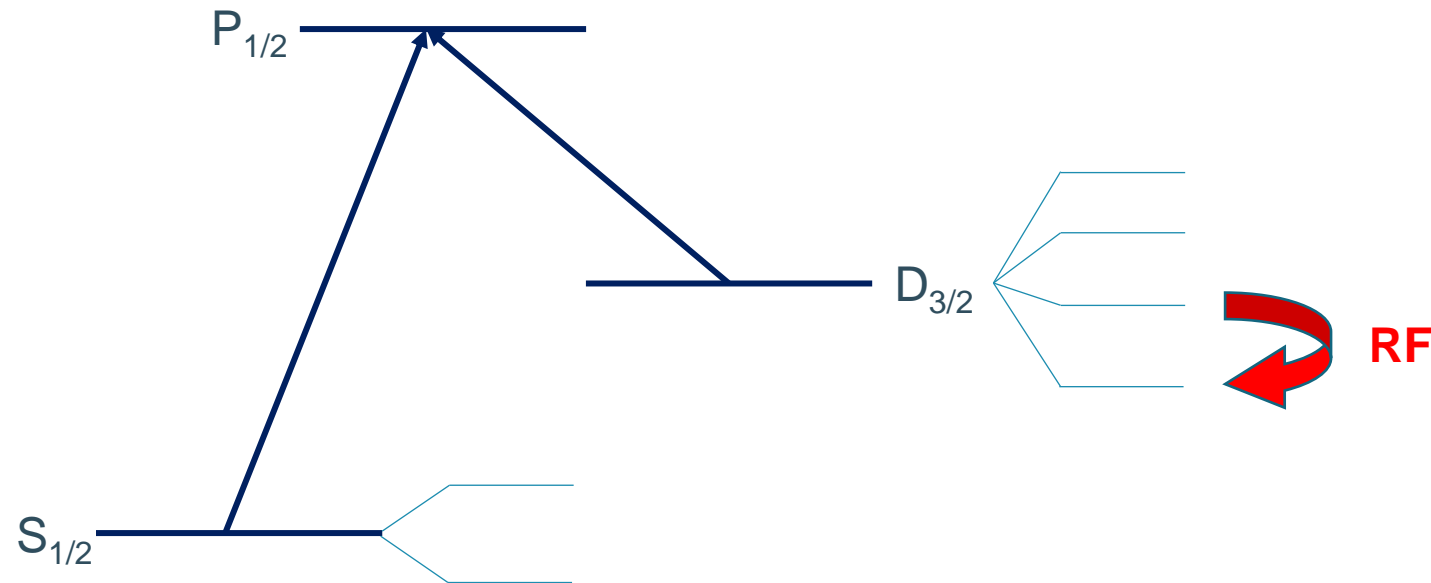
# New observables through higher precision spectroscopy

- Nuclear moments lead to shift and splitting of fine structure levels (*hyperfine structure*)
  - Multipole expansion *does not stop at the quadrupole!*
    - Magnetic octupole moment: 20 known values [1]
  - Contribution of higher-order magnetic octupole moment is three or more orders of magnitude smaller
- ⇒ Three or more orders of magnitude precision boost has to be found.
- **There are other observables we could hope to access!**



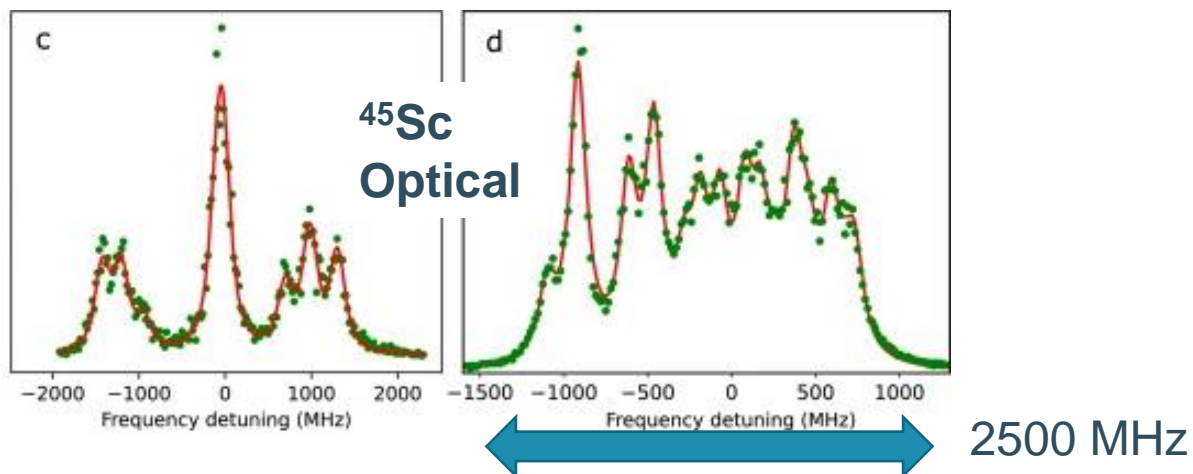
# New observables through higher precision spectroscopy

- Laser-radiofrequency double-resonance spectroscopy
  - Lasers are used for state preparation and readout;
  - Spectroscopy is performed using a radiofrequency excitation;
  - Linewidth:  $\Delta\nu \propto \frac{1}{\tau}$



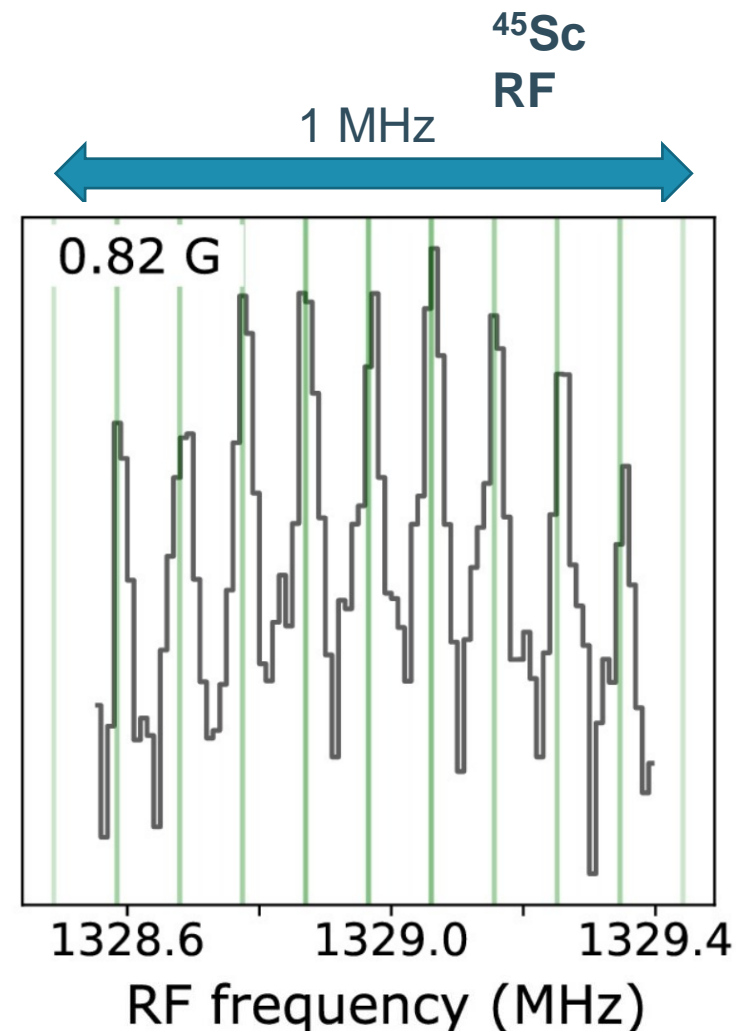
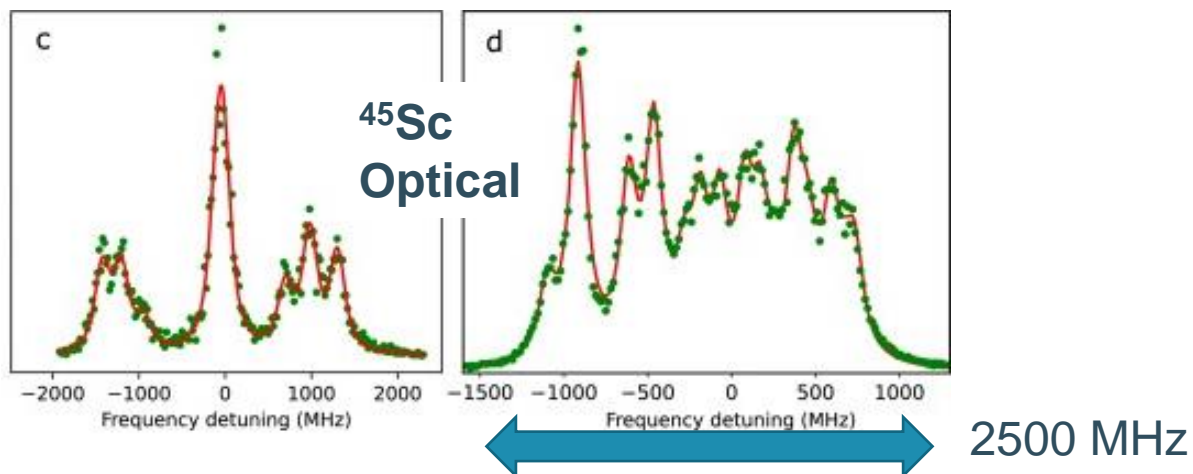
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- On a thermal beam?



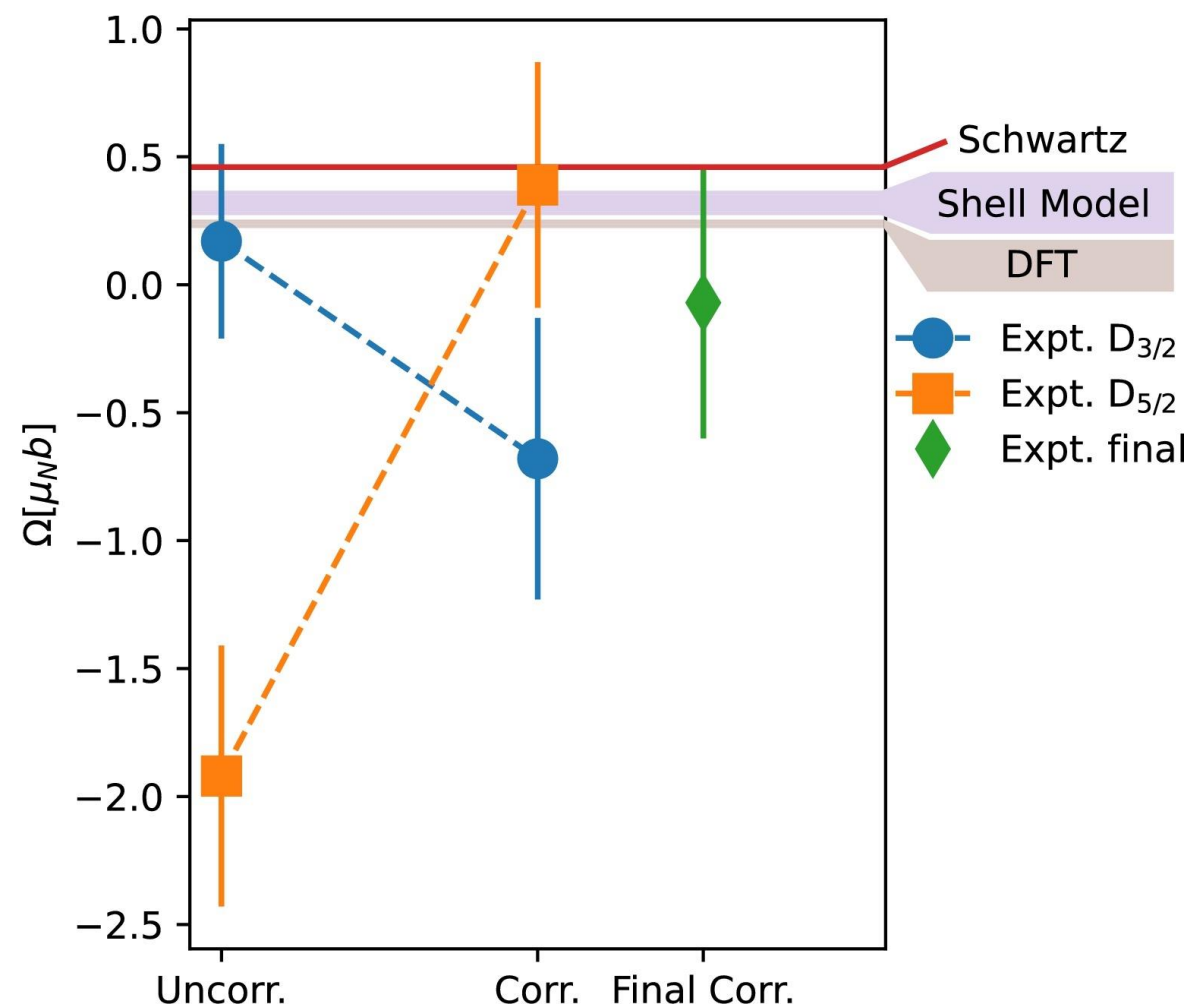
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- On a thermal beam?
  - Rf interaction zone of a few cm  $\Rightarrow \Delta\nu = 10$  kHz





		Theory this work	Expt. Ref. [17]		Expt. this work	
			Uncorrected	Corrected	Uncorrected	Corrected
$D_{3/2}$	A [MHz]	271(3)	269.556(1)	269.558(1)	269.55817(5)	269.55844(7)[3]
	B [MHz]	-27.5(5)	-26.346(4)	-26.360(8)	-26.3531(9)	-26.3596(5)[5]
	C [kHz]	—	—	—	-0.010(22)	0.039(28)[2]
	$\Omega$ [ $\mu_N b$ ]	—	—	—	0.17(38)	-0.68(49)[6]
$D_{5/2}$	A [MHz]	108(2)	109.032(1)	109.033(1)	109.03275(7)	109.03297(5)[3]
	B [MHz]	-38.5(5)	-37.387(12)	-37.373(15)	-37.3954(12)	-37.3745(8)[15]
	C [kHz]	—	1.7(10)	1.5(12)	0.31(8)	-0.062(59)[17]
	$\Omega$ [ $\mu_N b$ ]	—	-10.7(63)	-9.4(75)	-1.92(51)	0.39(37)[11]



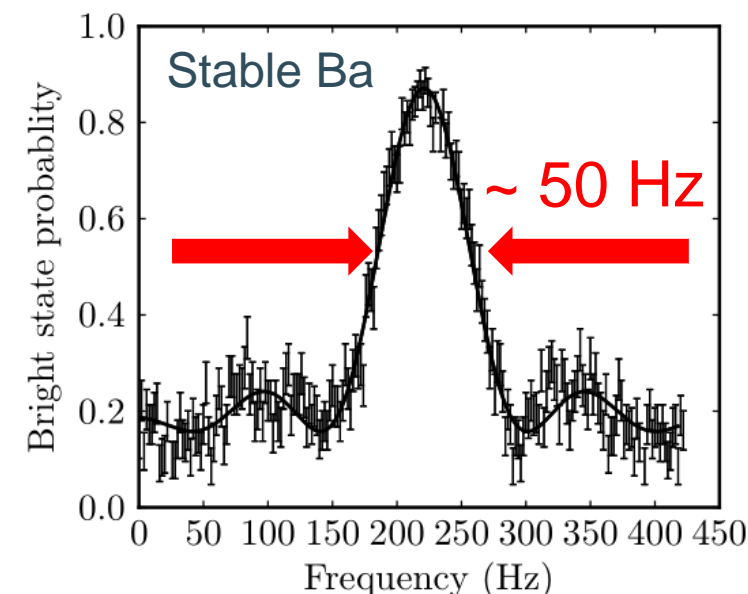
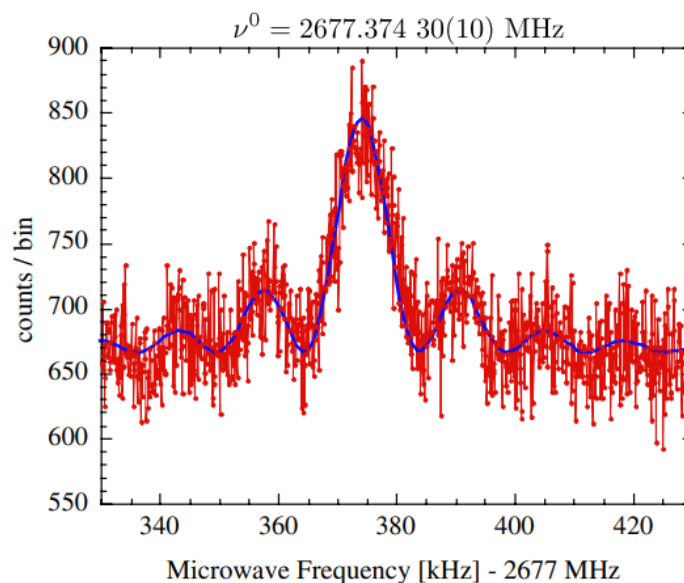
# New observables through higher precision spectroscopy

- On a thermal beam?
  - Rf interaction zone of a few cm =>  $\Delta\nu = 10 \text{ kHz}$
- In a collinear geometry?
  - Interaction time can be a few microseconds  $\Delta\nu \geq 100 \text{ kHz}$
  - Implementation in RAPTOR experiment at IGISOL planned

# New observables through higher precision spectroscopy

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- In an atom or ion trap...?
  - Offline: **Hz-level!**
  - Only on-line so far:  ${}^7, {}^{11}\text{Be}$  in RIKEN

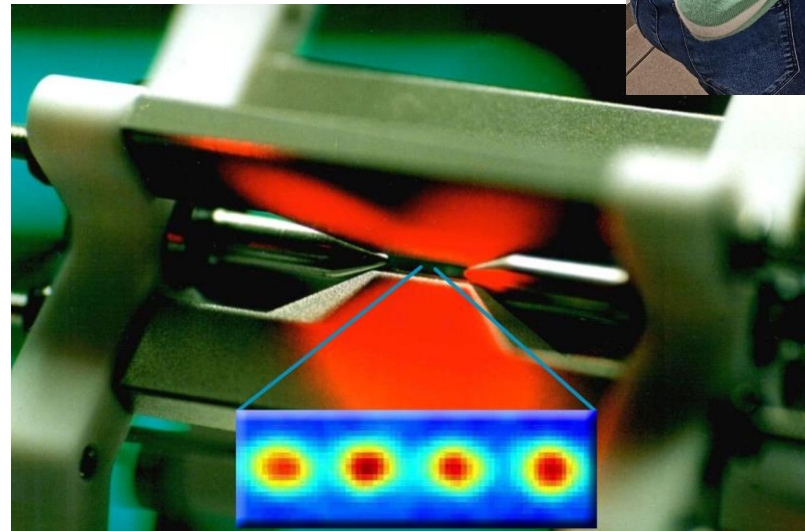
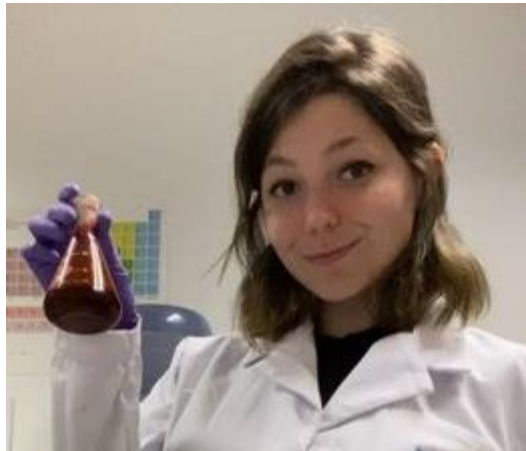
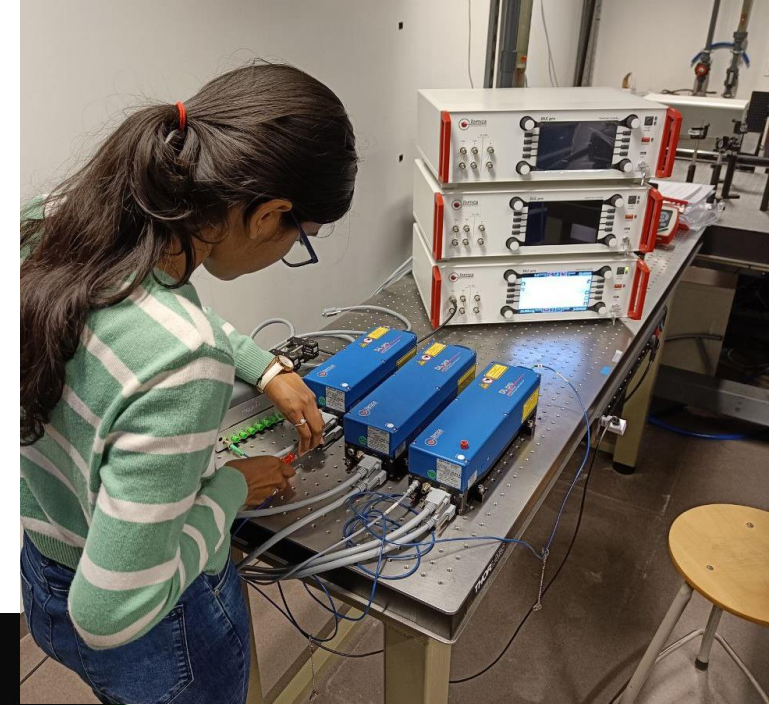


PhD thesis N. C. Lewty, U of Singapore

A. Takamine et al., PRL **112**, 162502 (2014)

Isotope	$A$ [MHz]
${}^7\text{Be}$	$-742.772\ 28(43)^c$
${}^9\text{Be}$	$-625.008\ 837\ 048(10)^d$
${}^{11}\text{Be}$	$-2677.302\ 988(72)^b$

# Our aim: magnetic octupole moment of strontium



PhD project Anaïs Dorne

5μm

Picture: Univ of Innsbruck. Group R. Blatt



UNIVERSITY OF JYVÄSKYLÄ

**KU LEUVEN**

# Thank you all for your attention!

Thanks to many people for their hard work and enthusiasm (and slides)

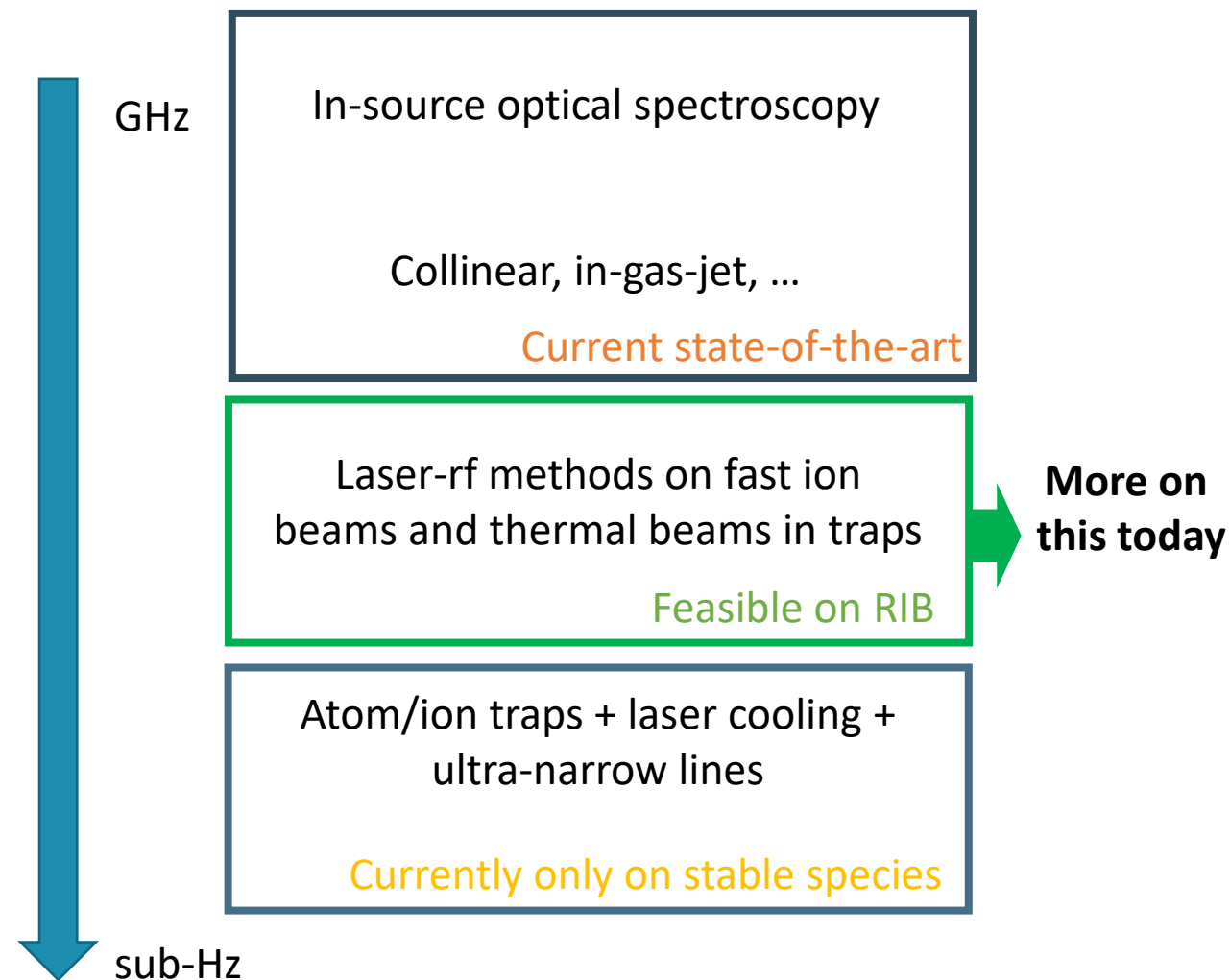


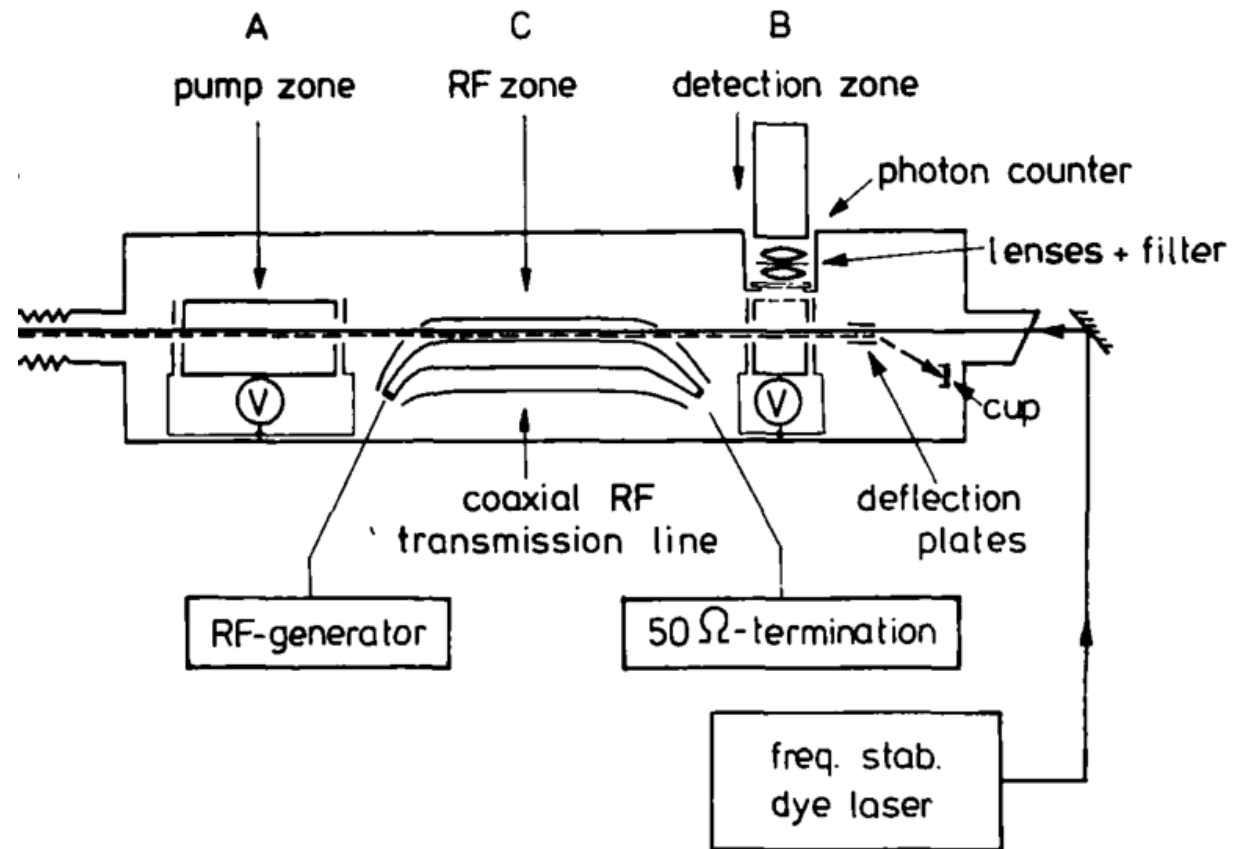
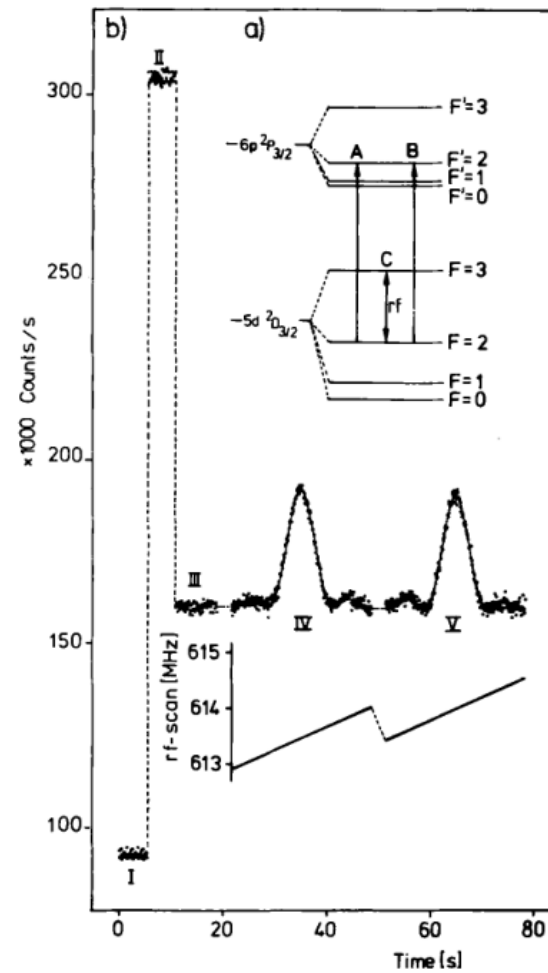




# Wider context: higher-precision spectroscopy of radioactive ions

- Magnetic dipole moments
- Electrical quadrupole moments and charge radii
- Hyperfine anomaly
  - Relates to the distribution of magnetization inside nuclear volume
- Higher-order moments
  - Magnetic octupole, electric hexadecupole, ...
- Higher-order moments of the charge radii
  - E.g.  $\langle r^4 \rangle$  relates to surface thickness of nuclear density
- Beyond-standard model physics from Hz-level isotope shift spectroscopy

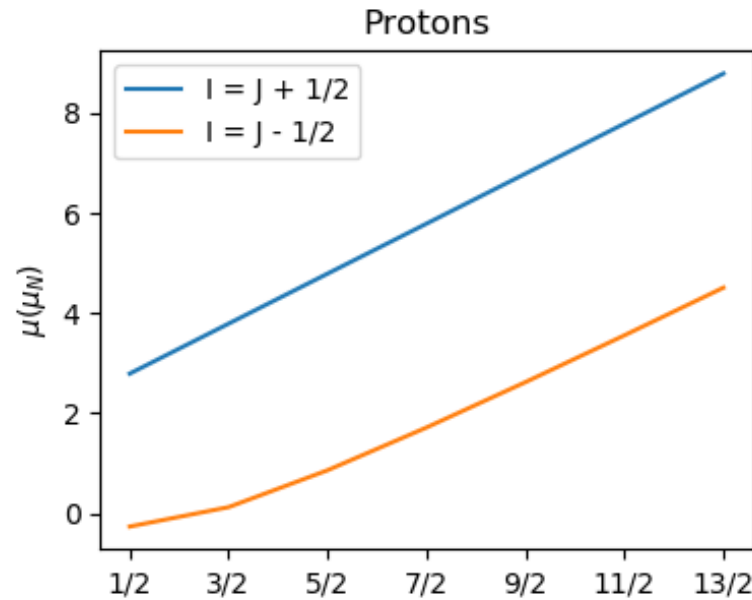






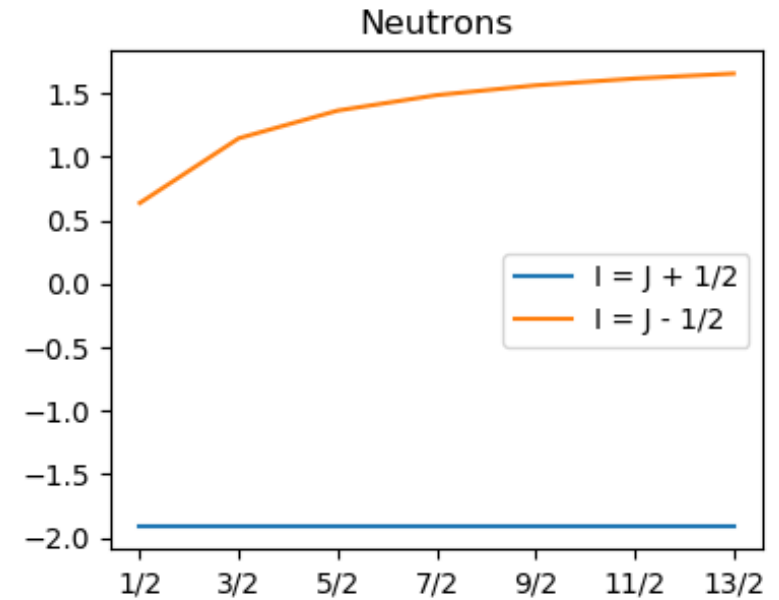
# Magnetic dipole moments

- Most basic model: assume only the valence nucleon contributes; all other particles paired to  $I=0$   
=> Schmidt lines
- Provides a very powerful tool to establish valence nucleon configurations!



$$\mu = j - \frac{1}{2} + \frac{g_s^\pi}{2} \quad \text{for } j = l + \frac{1}{2}$$

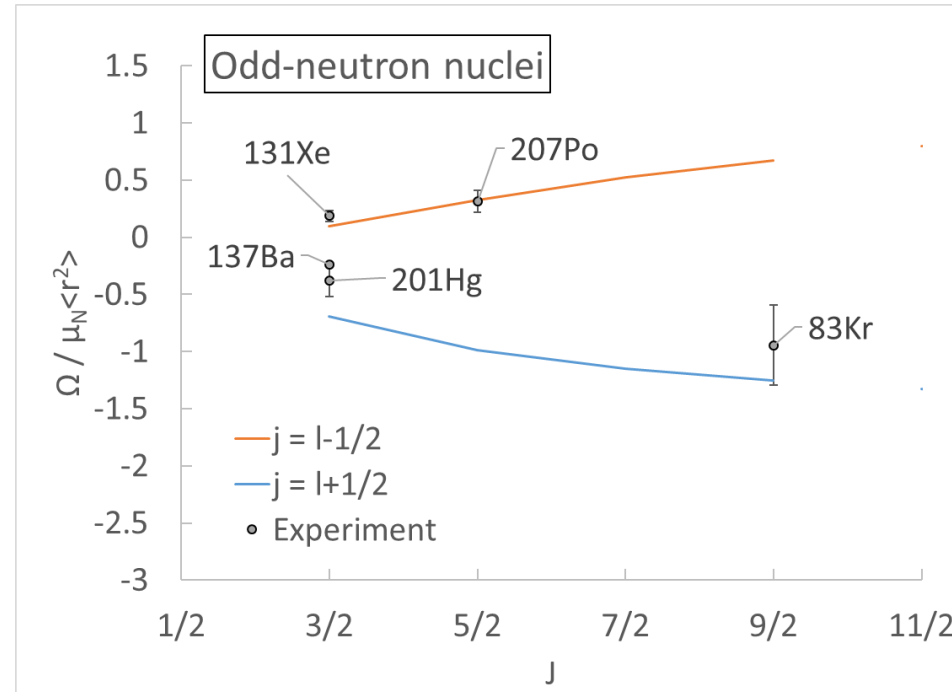
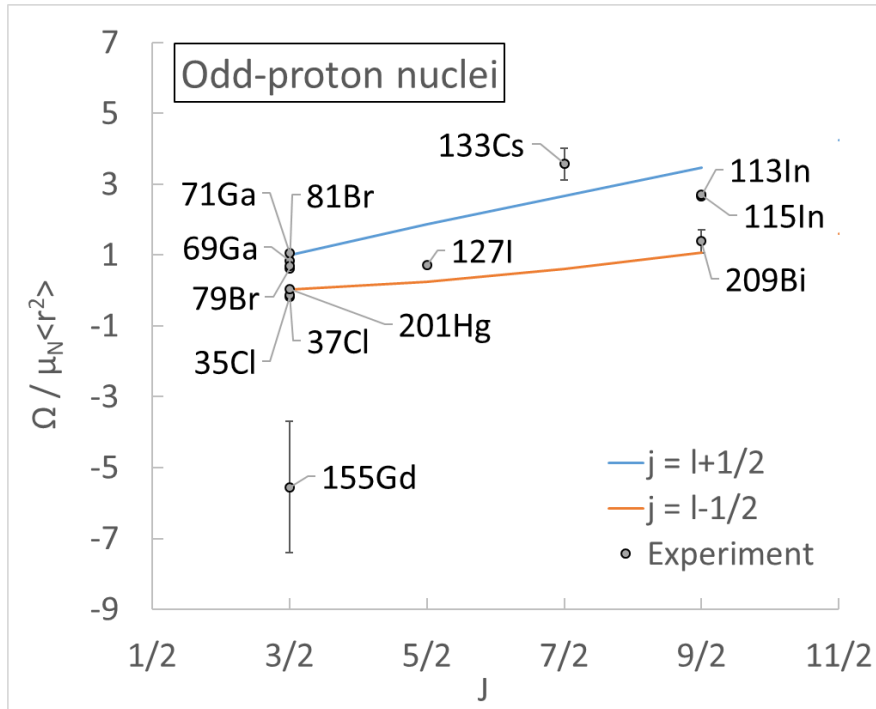
$$\mu = \frac{j}{j+1} \left( j + \frac{3}{2} - \frac{g_s^\pi}{2} \right) \quad \text{for } j = l - \frac{1}{2}$$



$$\mu = \frac{g_s^v}{2} = -1.913 \quad \text{for } j = l + \frac{1}{2}$$

$$\mu = -\frac{j}{j+1} \frac{g_s^v}{2} \quad \text{for } j = l - \frac{1}{2}$$

# Nuclear magnetic octupole moments



$$\frac{\Omega}{\mu_N \langle r^2 \rangle} = \frac{3}{2} \frac{2I - 1}{(2I + 4)(2I + 2)} \times \begin{cases} (I + 2)[(I - \frac{3}{2})g_l + g_s], & I = l + \frac{1}{2} \\ (I - 1)[(I + \frac{5}{2})g_l - g_s], & I = l - \frac{1}{2} \end{cases}$$