



Barium Tagging for EXO

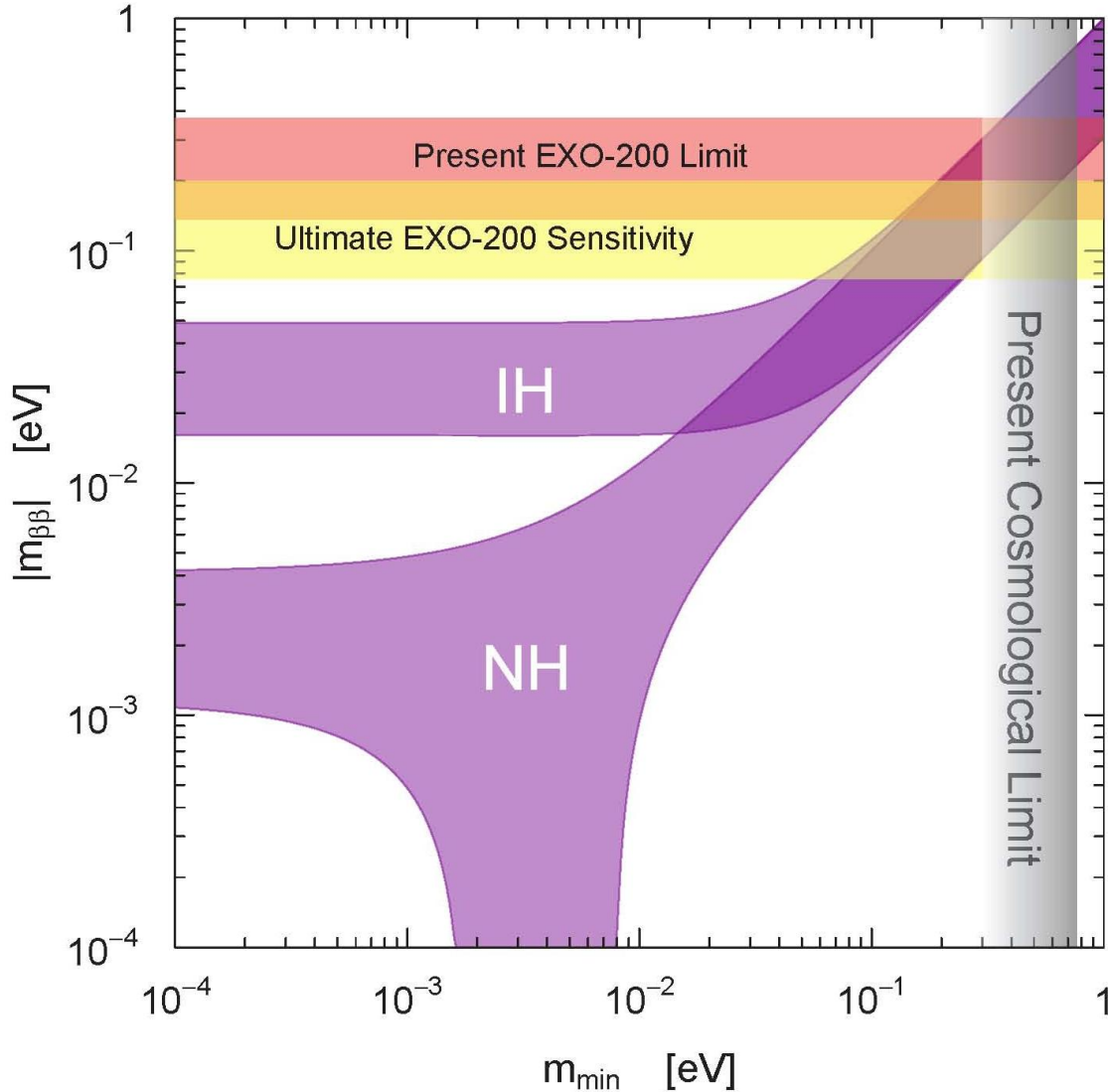
**Scott Kravitz
Stanford University**

APS DPF

UC Santa Cruz, 15 Aug. 2013



EXO Sensitivity Projections



Adapted from Bilenky & Giunti arXiv:1203.5250v2

Purple bands are 68%CL from oscillation experiments for inverted and normal mass hierarchy

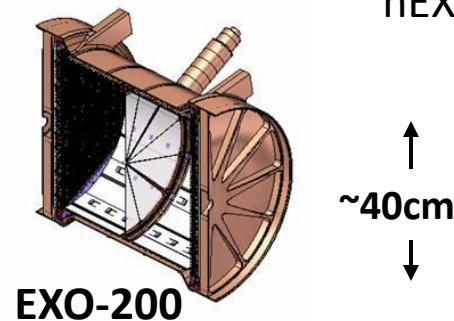
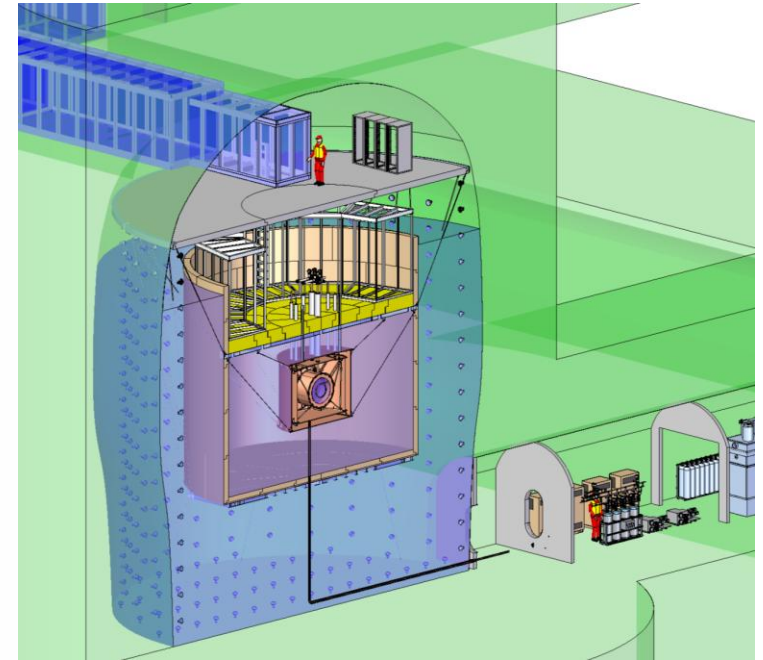
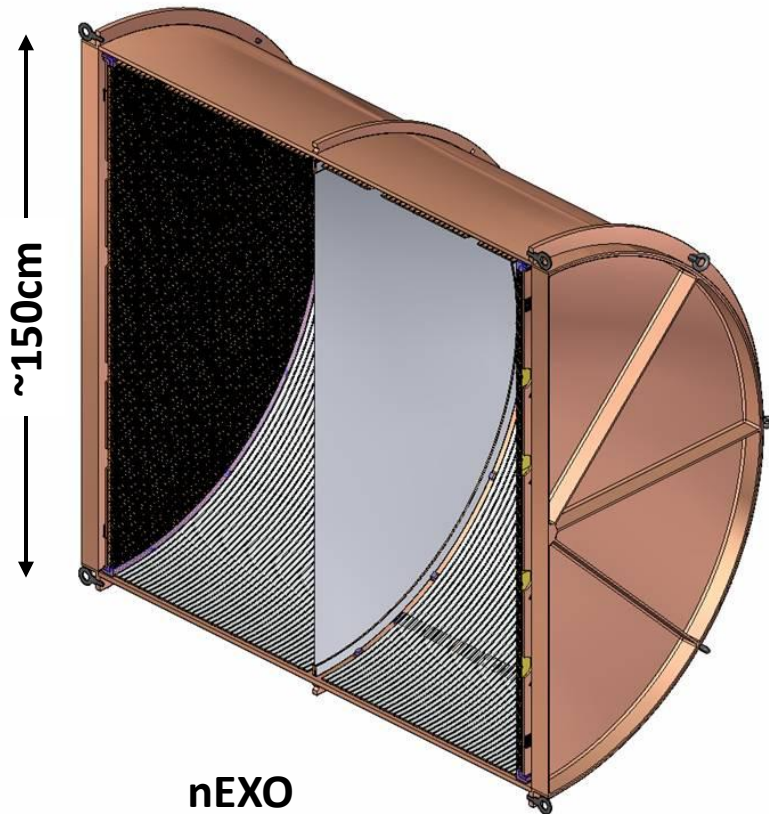
EXO-200 present limit is the 90%CL envelope of limits (for different NMEs) from PRL 109 (2012) 032505

EXO-200 ultimate sensitivity: 90%CL for no signal in 4 years livetime with new analysis and Rn removal



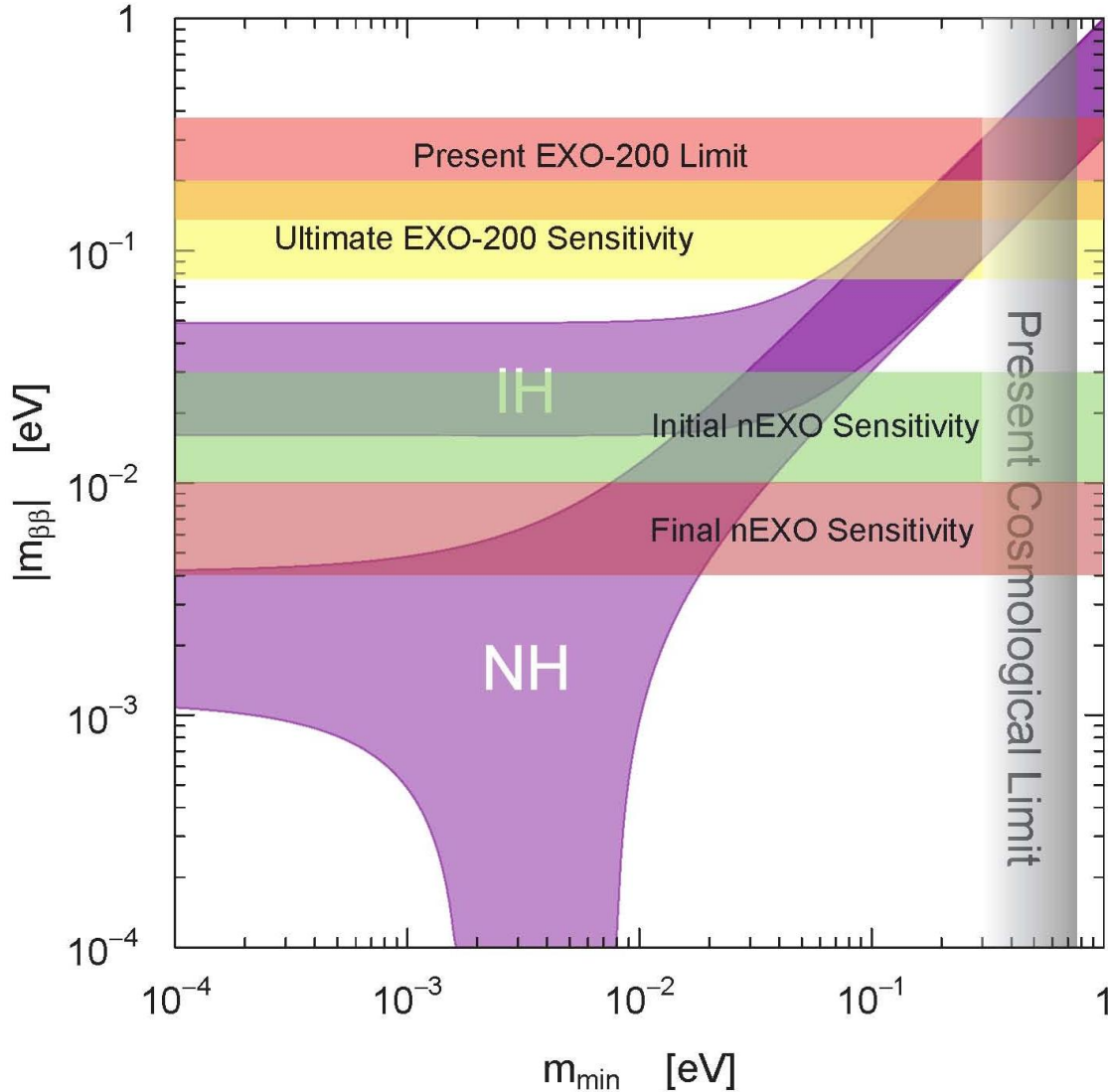
nEXO: EXO-200's Successor

- 5 tonne LXe time projection chamber (TPC) "as similar to EXO-200 as possible"
- Provide access ports for a possible later upgrade to barium tagging





EXO Sensitivity Projections



Adapted from Bilenky & Giunti arXiv:1203.5250v2

Purple bands are 68%CL from oscillation experiments for inverted and normal mass hierarchy

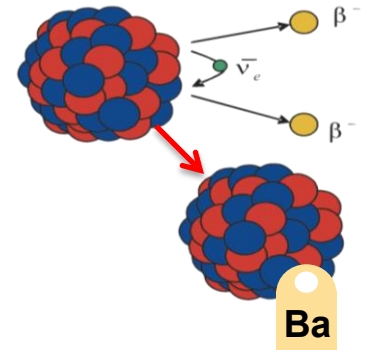
EXO-200 present limit is the 90%CL envelope of limits (for different NMEs) from PRL 109 (2012) 032505

EXO-200 ultimate sensitivity: 90%CL for no signal in 4 years livetime with new analysis and Rn removal

Initial nEXO band refers to a detector directly scaled from EXO-200, including its measured background and 10yr livetime.

Final nEXO band refers to the same detector and no background other than 2ν

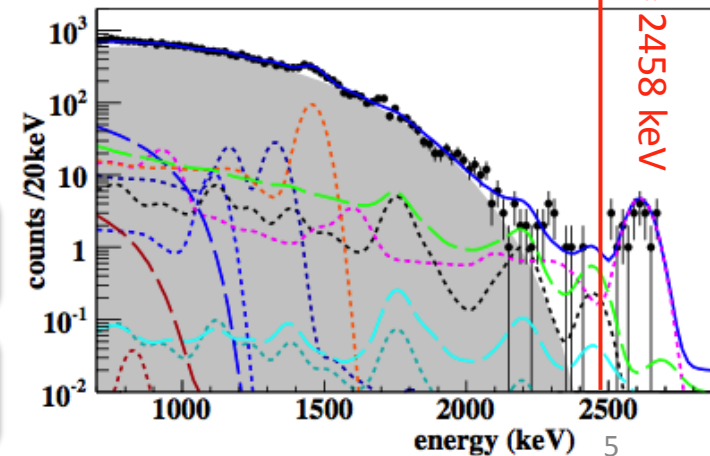
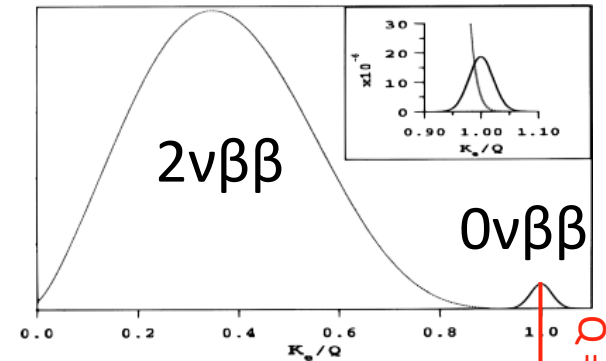
Ba Tagging Motivation



- **Goal: perform background free measurement**
 - Measure candidate decay event positions, energies
 - Recover and identify Ba ion
 - Tag event-recovery coincidences as $\beta\beta$
 - Reject untagged events
- **Achievement:**
 - A **background-free** detector
 - Better scaling of mass sensitivity
- **Strategy: multipronged approach**

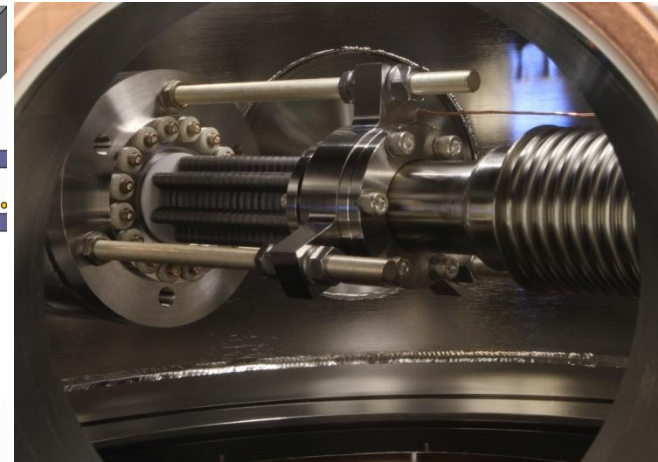
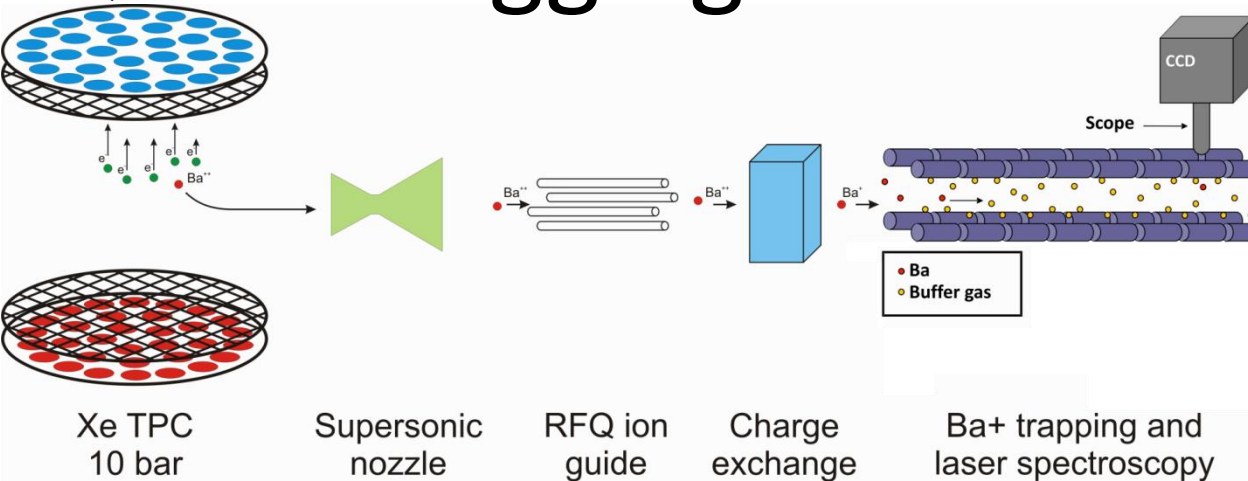
With background: $\langle m_\nu \rangle \propto 1/\sqrt{T_{1/2}^{0\nu\beta\beta}} \propto 1/(Nt)^{1/4}$

Without background: $\langle m_\nu \rangle \propto 1/\sqrt{T_{1/2}^{0\nu\beta\beta}} \propto 1/\sqrt{Nt}$





Tagging from Gas Xenon

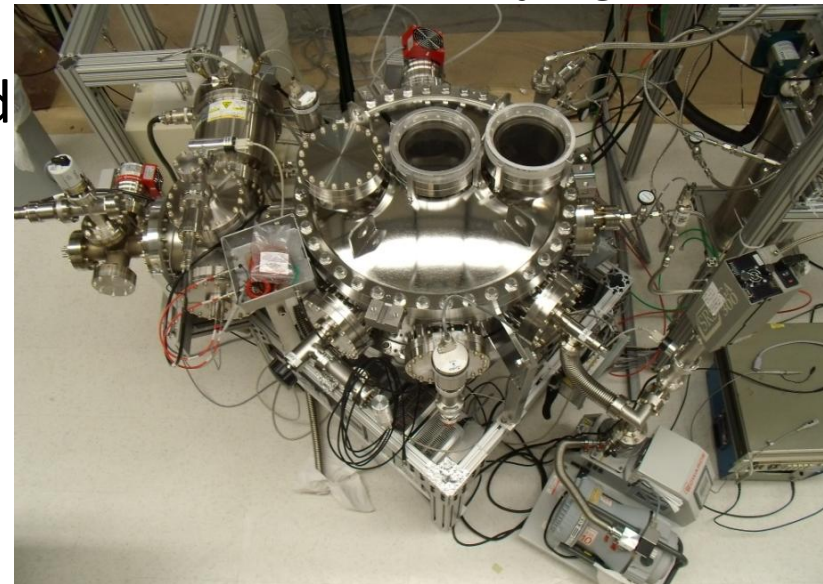


RF Funnel

How it works:

- Extract Ba⁺⁺ from TPC by shaping E-field
- Guide into vacuum
- Convert Ba⁺⁺ to Ba⁺ [1]
- Identify via laser spectroscopy [2]

Current work has demonstrated extraction of ions created by a test Ba source from 10 bar Xe into vacuum

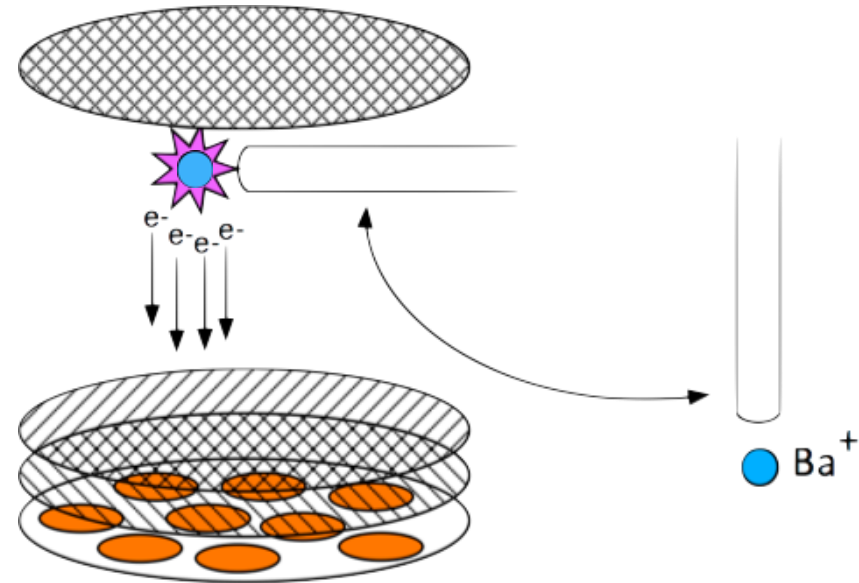


Current Stanford Setup

[1] J. of Phys.: Conf. Ser. 309(2011)12005 [2] Phys. Rev. A 76, 023404 (2007)

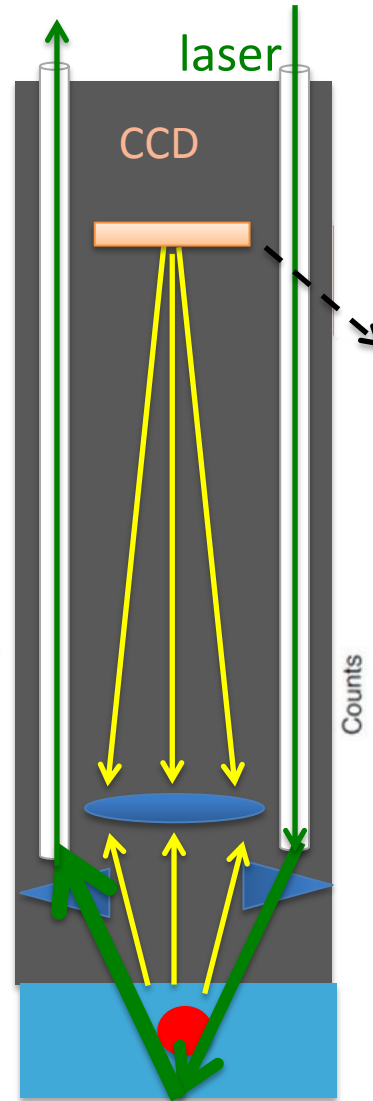
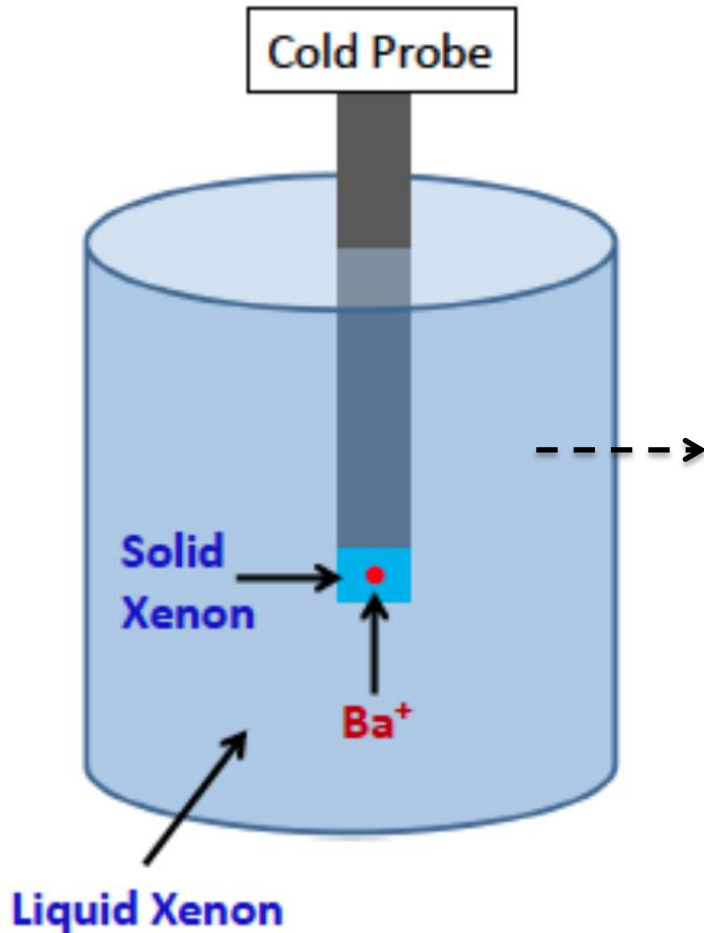
Tagging from Liquid Xenon

- Decay product in LXe is Ba^+
- Send probe into TPC, deposit Ba^+ onto probe tip (e.g. electrostatically)
- Remove probe to identification chamber
- Ba^+ moves slowly enough in LXe to remain in region of reconstructed decay event
- Investigating two methods of liquid tagging - CSU and Stanford



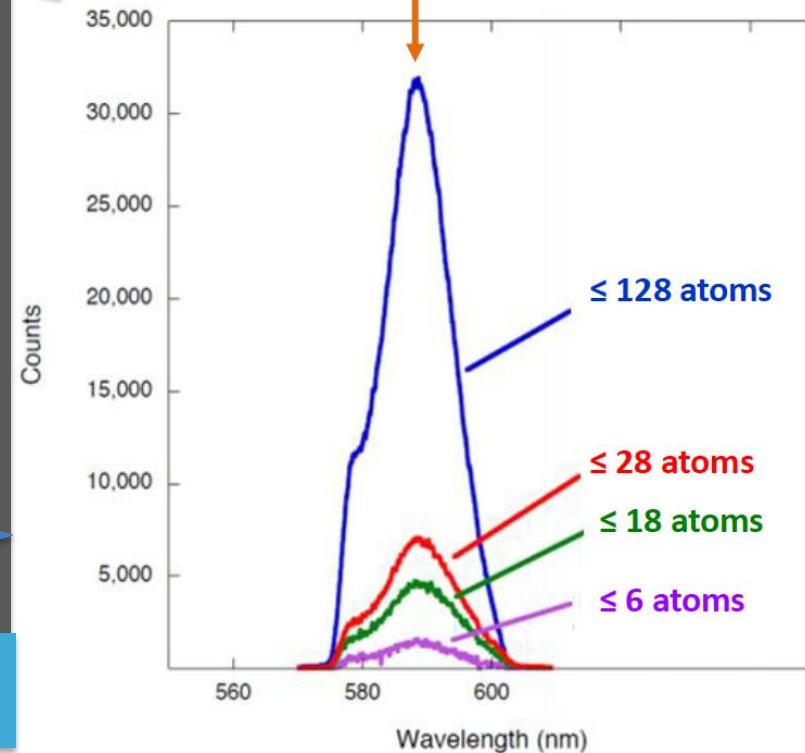
Tagging Spectroscopically in SXe (Colorado State University)

Trap barium daughter ion in solid xenon on a probe:



Detect single ion or atom on the probe with laser-induced fluorescence:

Fluorescence peak of Ba in SXe after excitation w/ 558 nm laser





Resonance Ionization Spectroscopy

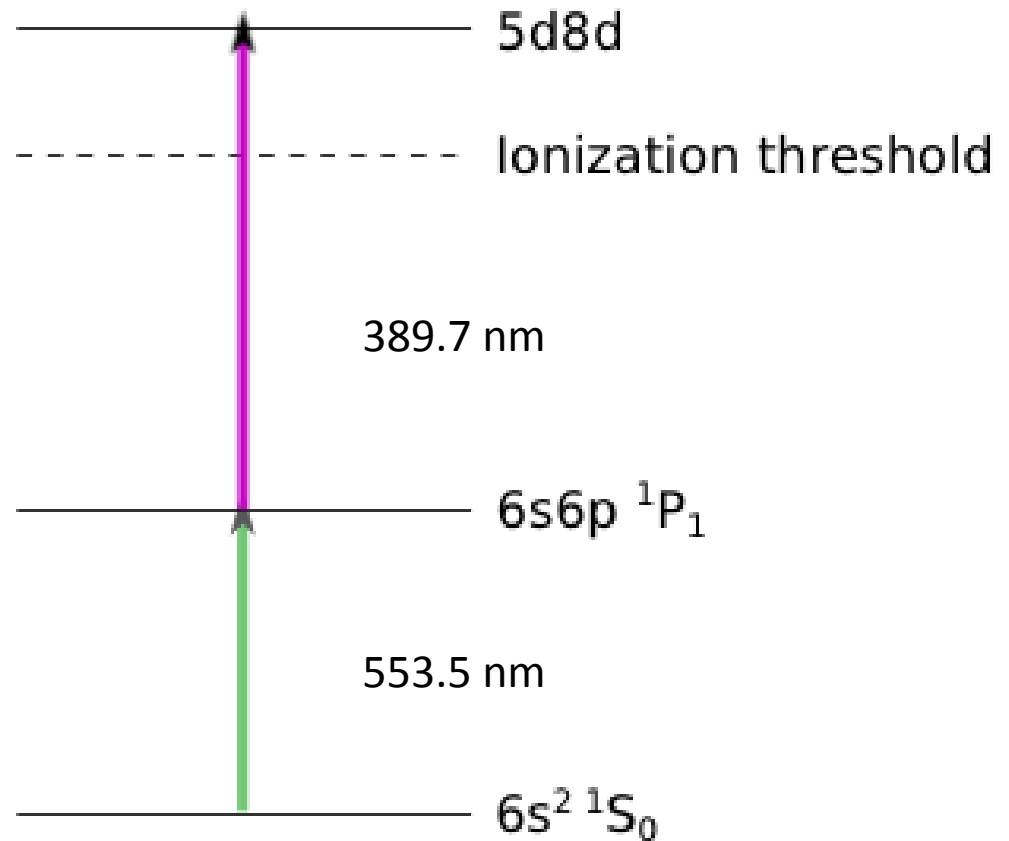
Challenge:

Recover and identify
single Ba, ignoring all else

Solution:

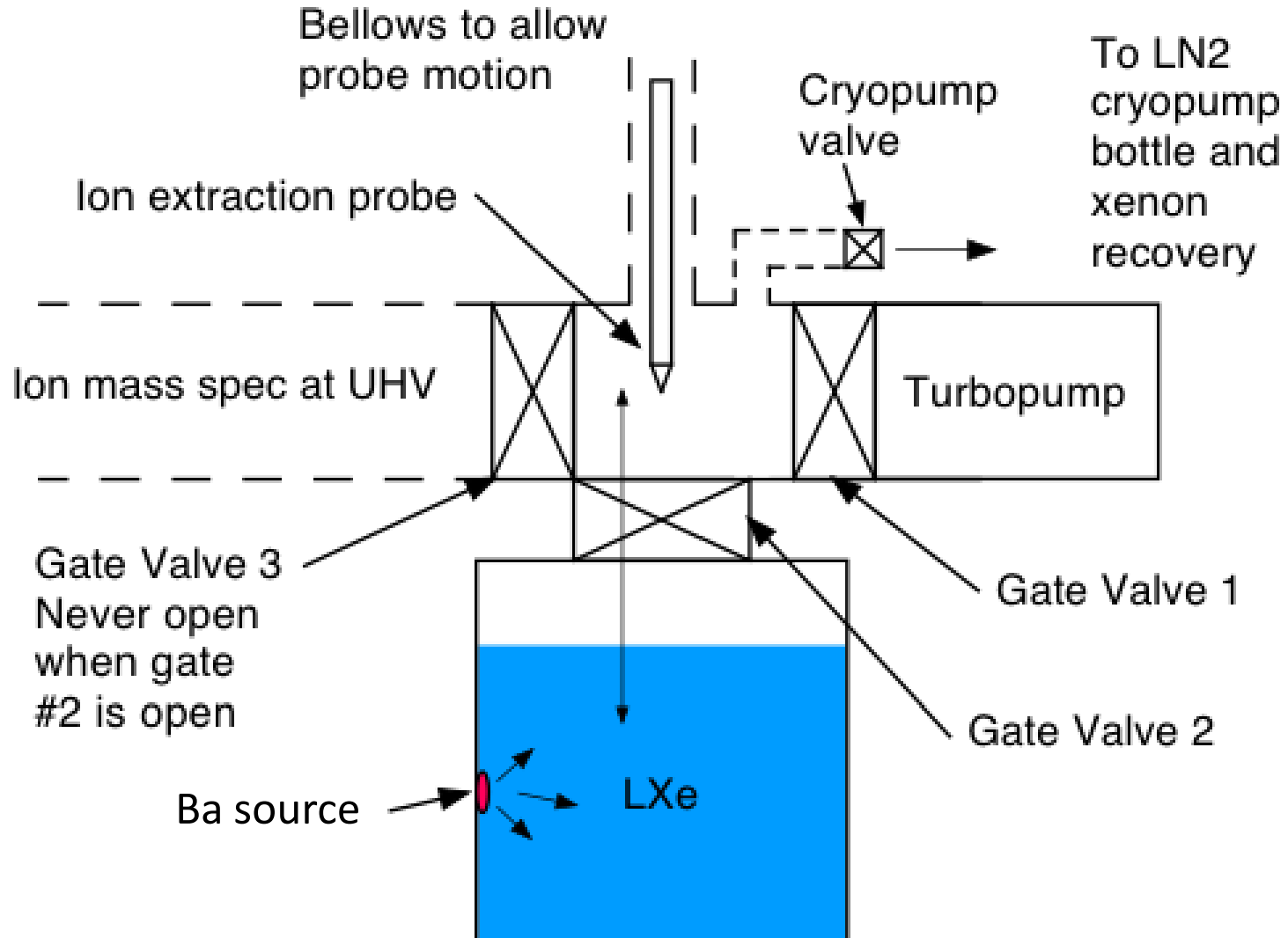
Efficiently and selectively
re-ionize barium
using its atomic spectroscopy

RIS is sensitive to single ions



Neutral barium spectroscopy

Liquid Tagging Setup Schematic



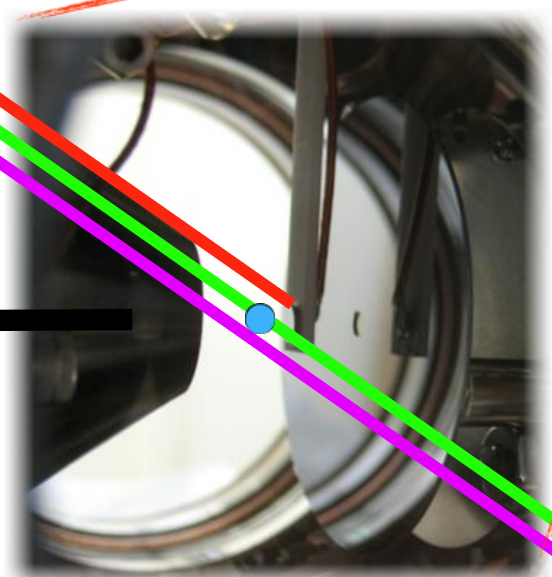
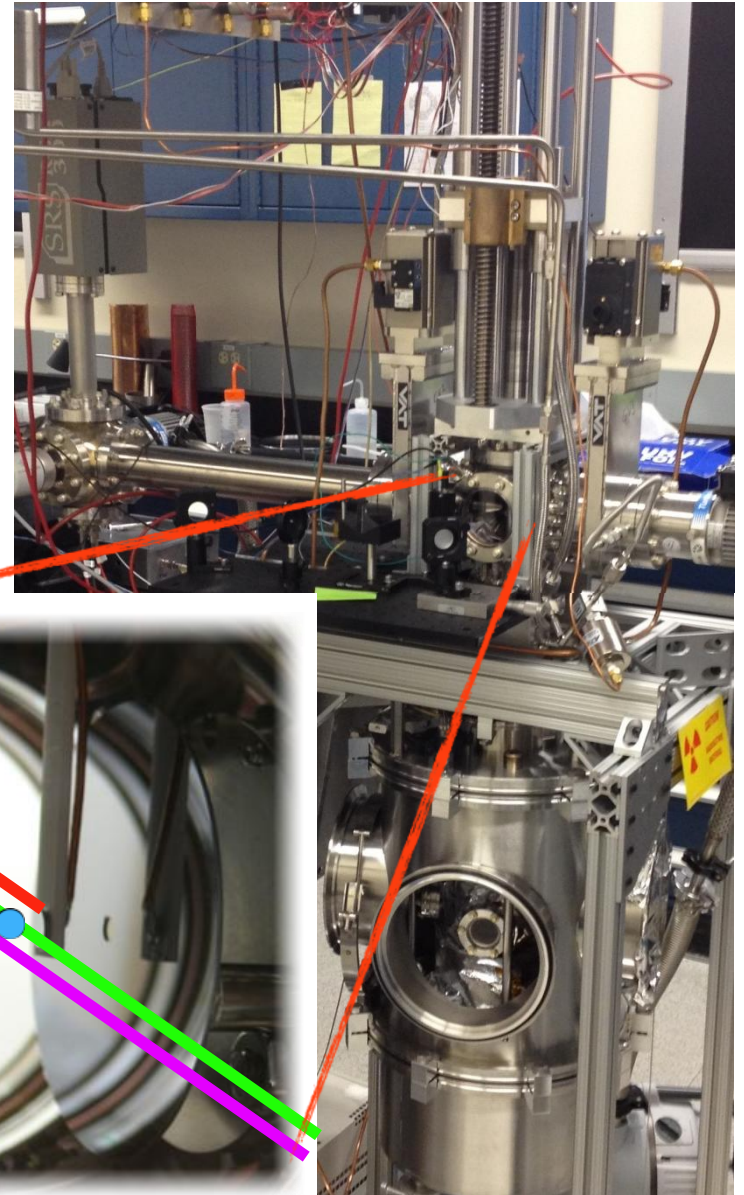
RIS at Stanford

- Thermally desorb with IR laser
- Ionize resonantly just Ba
- Detect and identify by mass using ToF spectrometer

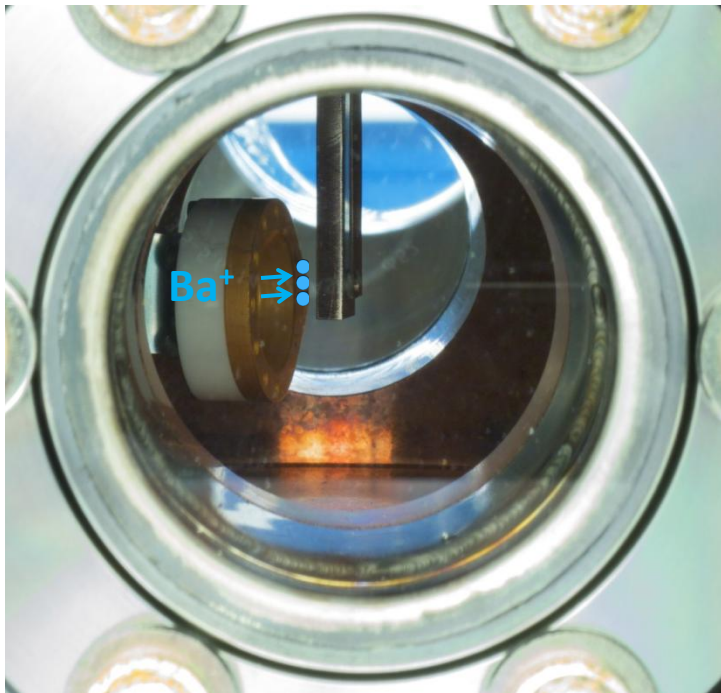
1064 nm
553.5 nm
389.7 nm

Ba⁺

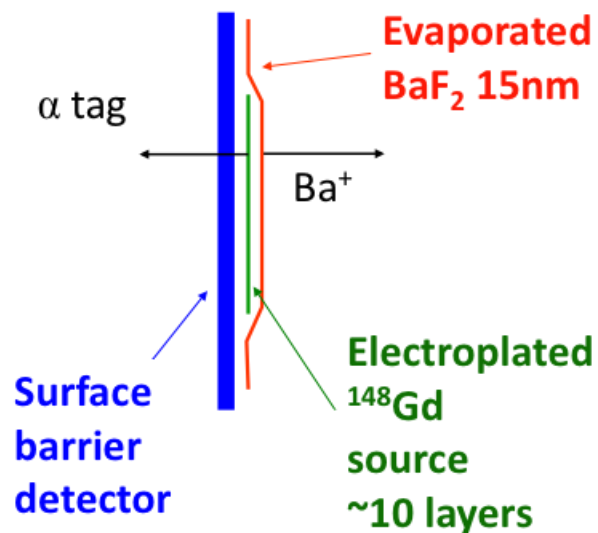
To Time of Flight
Spectrometer



Barium Loading in Vacuum

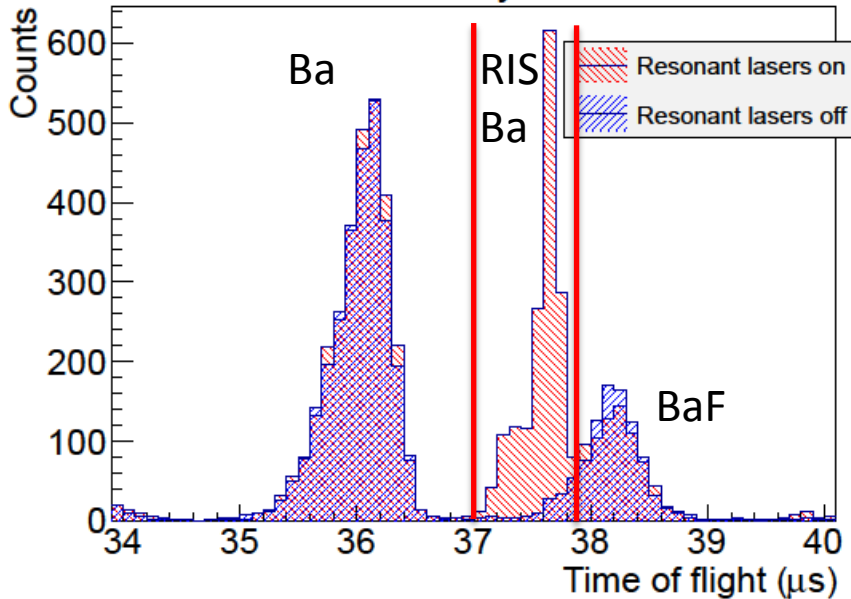


- Before liquid, test in vacuum
- ^{148}Gd -driven Ba ion source developed for testing
- Deposit ~ 10 - 100k Ba^+ initially

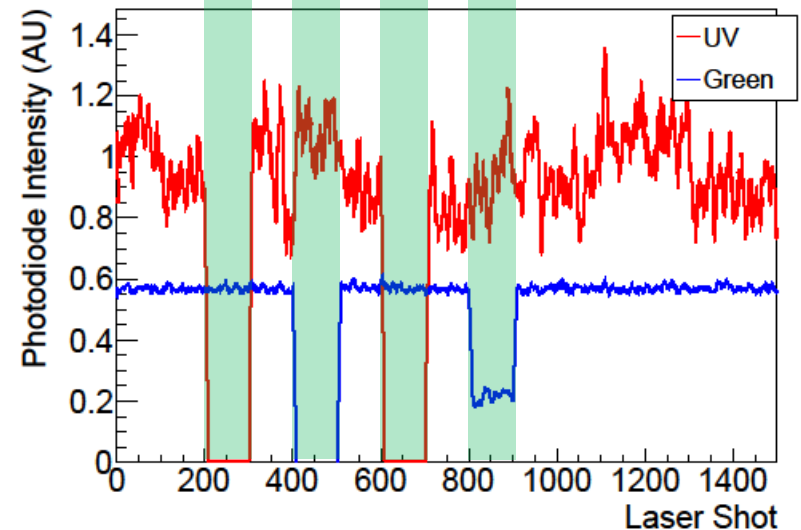
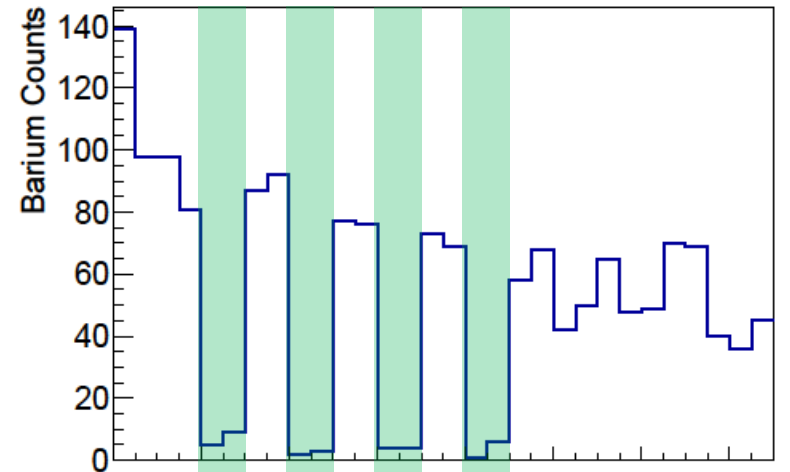


Full RIS Tests

Ablated and resonantly ionized barium



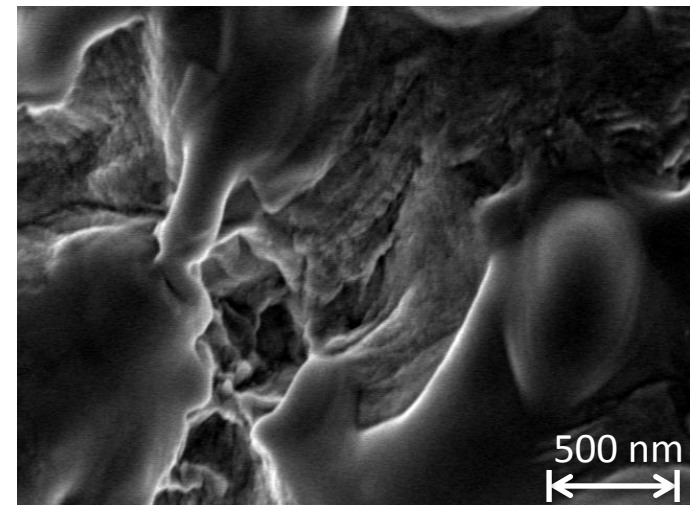
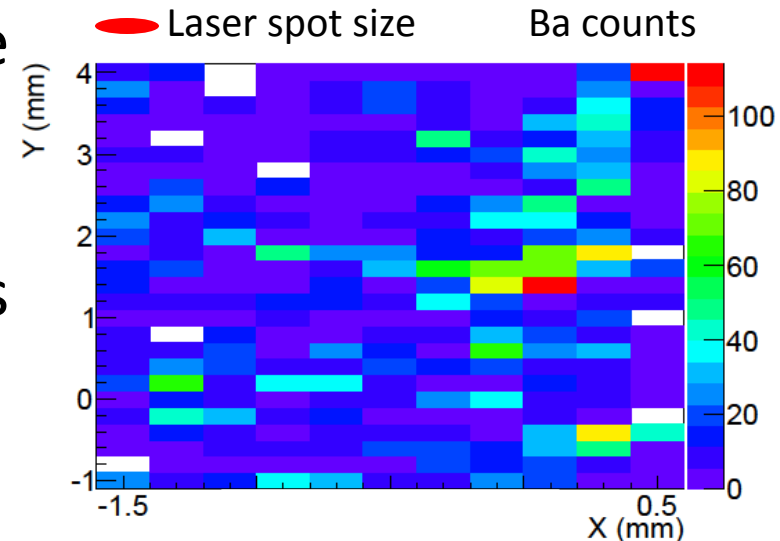
Resonantly ionized barium



- Many masses present
 - Alkalis dominate (low ionization E)
 - Clusters of substrate at high desorption power
 - Ba and BaF from desorption laser as well as RIS Ba
- Resonant behavior observed

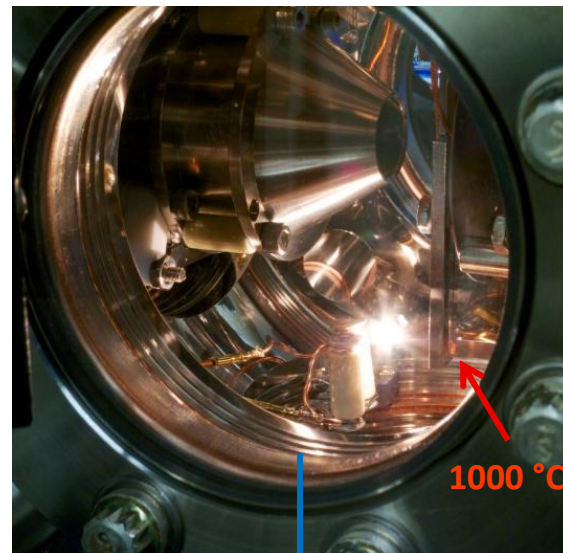
Desorption Challenges

- Usually desorption laser must be powerful enough to ablate several species to see RIS signal
- Ba signal from localized hotspots
- Surface contamination or structural defects could harm (help?) desorption
- Nature of bonding to surface is crucial; must choose the right material and surface prep

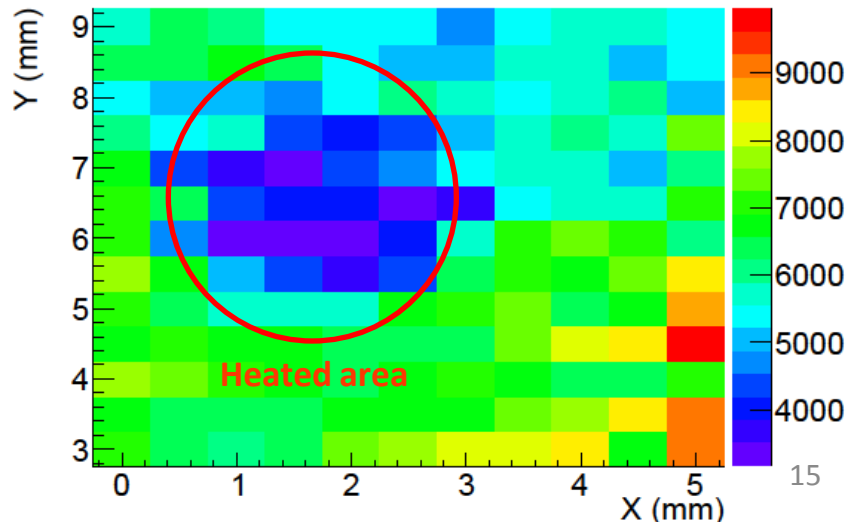


Surface Material and Prep

- Desired surface properties:
 - Weak Ba bond
 - Resistant to oxidation
 - Available at high purity
- Small/no RIS signal from Si, Re, Pt
- Notable RIS signal from W, Ni, Ta
- Surface cleaning:
 - Bare deposition substrate may facilitate Ba desorption
 - Reduce backgrounds
- Cleaning results:
 - Reduced backgrounds
 - No loss of signal

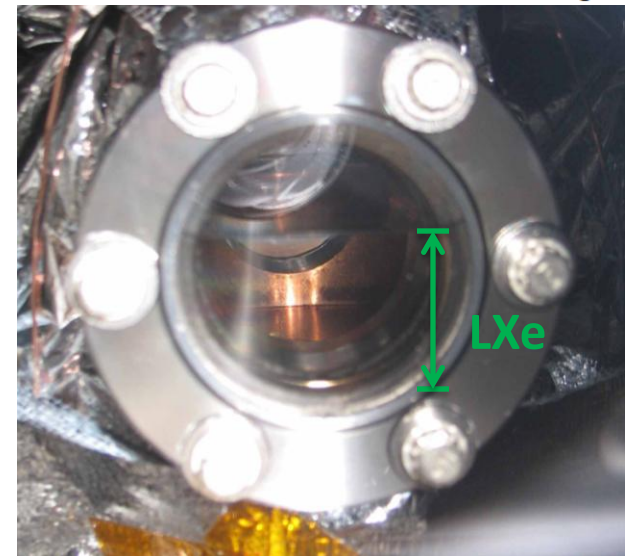
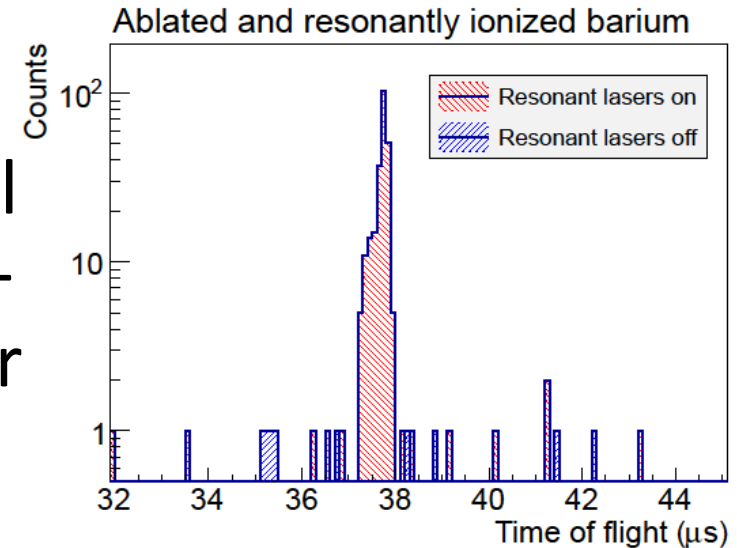


Total counts (all masses)



Results and Next Steps

- Proper choice of material and surface prep produces a signal with almost no backgrounds – not even Ba ionized by IR laser
- Ideal conditions + luck:
>5% efficiency from deposition to final detection
- **Next:**
 - Continued work on improving repeatability, efficiency
 - Testing in LXe



The EXO Collaboration



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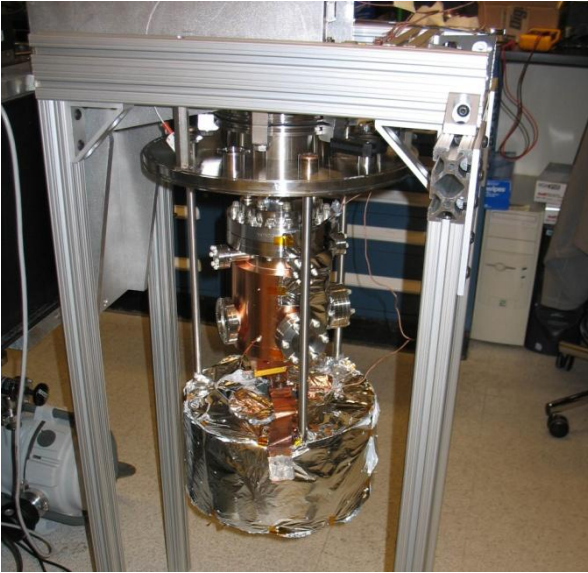
Stanford University, Stanford CA, USA - P.S. Barbeau, J. Bonatt, T. Brunner, J. Chaves, J. Davis, R. DeVoe, D. Fudenberg, G. Gratta, S. Kravitz, D. Moore, I. Ostrovskiy, A. Rivas, A. Schubert, D. Tosi, K. Twelker, L. Wen

Technical University of Munich, Garching, Germany - W. Feldmeier, P. Fierlinger, M. Marino



Backup Slides

LXe Cell and Ba Source

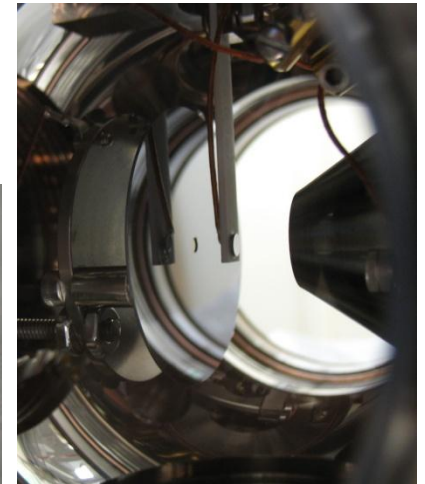
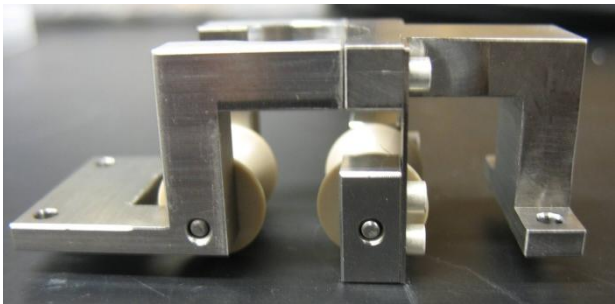


Loading $\approx 10^4$ Ba/hr

Effects of bias on Ba
deposition?

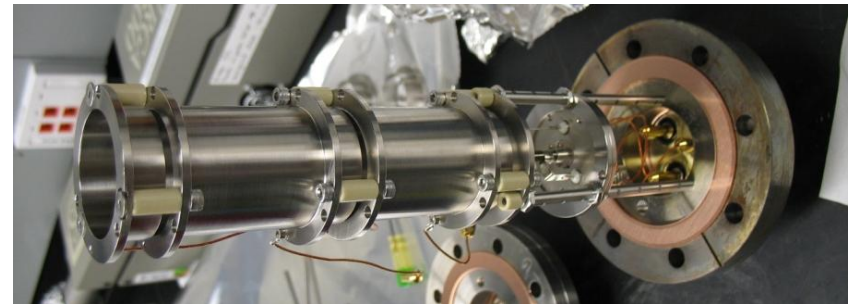
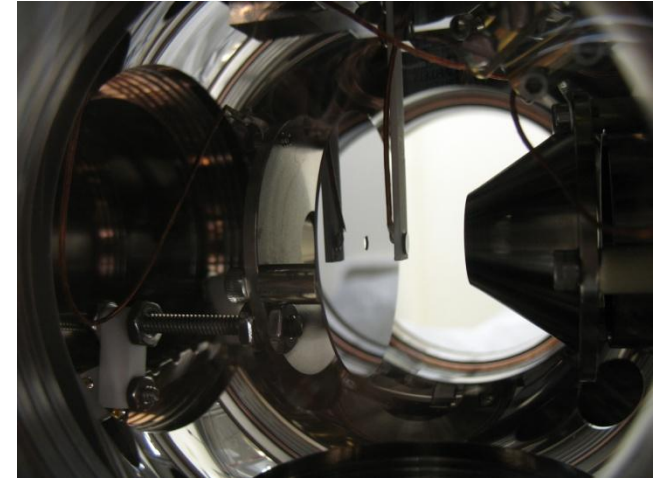
Probe

- Probe is a 0.5" diameter tube
- Rollers constrain lateral movement
- Target mounted on ceramic tab with vacuum-safe conductive epoxy
- Target charged via feedthrough at the top
- Second probe machined, allows for ohmic heating of the target

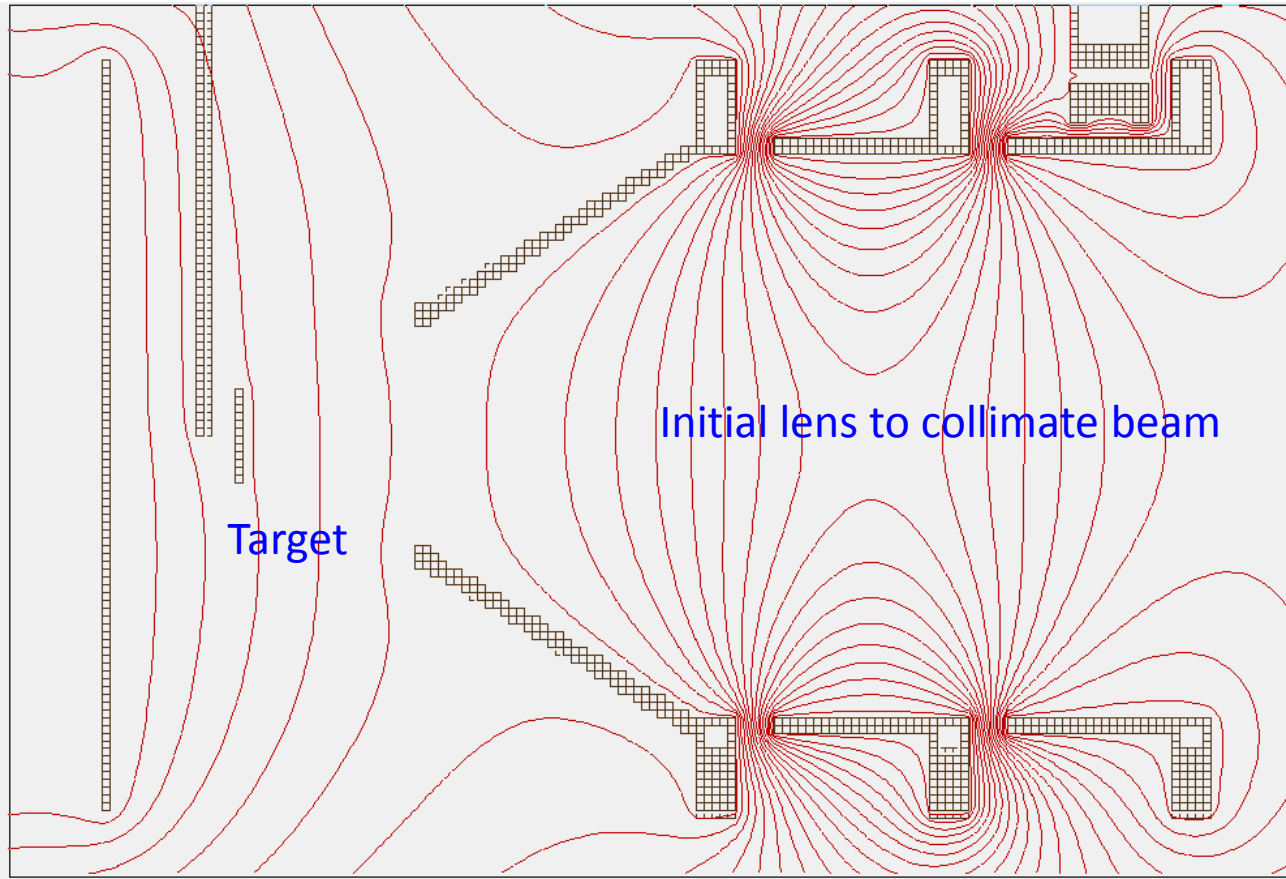


Spectrometer Design

- Ions are produced at target, guided by electrodes and detected by CEM
- Transit time determines ion mass
- Front electrodes capture, shape and accelerate ions
- Back electrodes refocus and detect ions

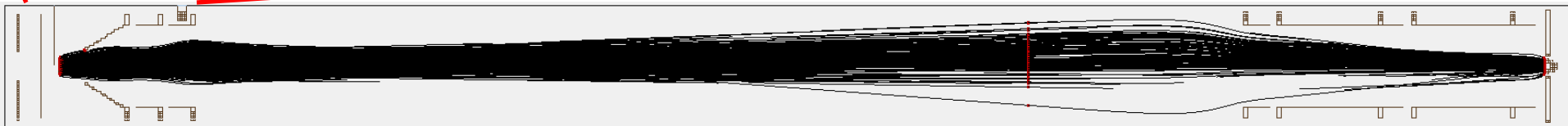


SIMION for time of flight



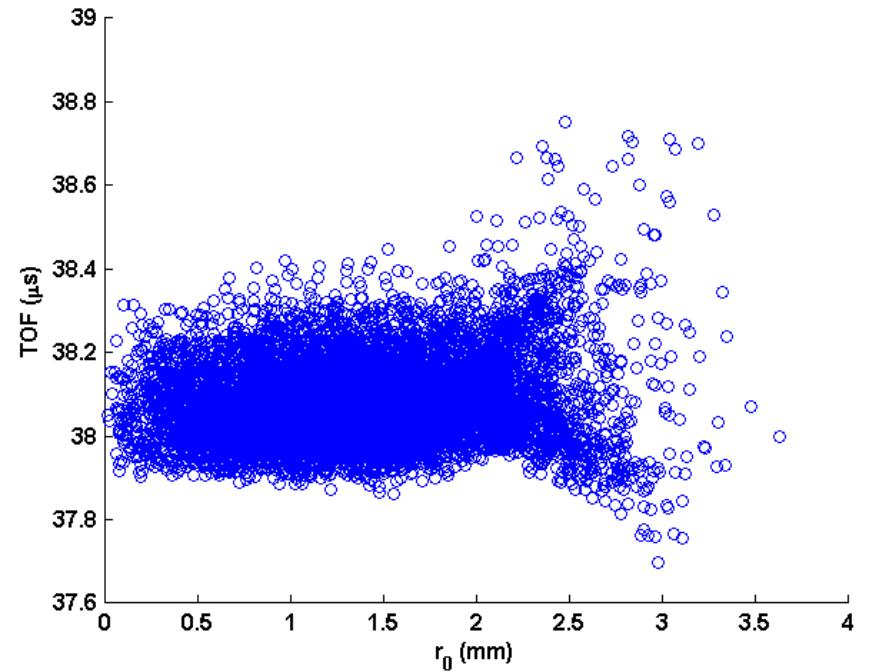
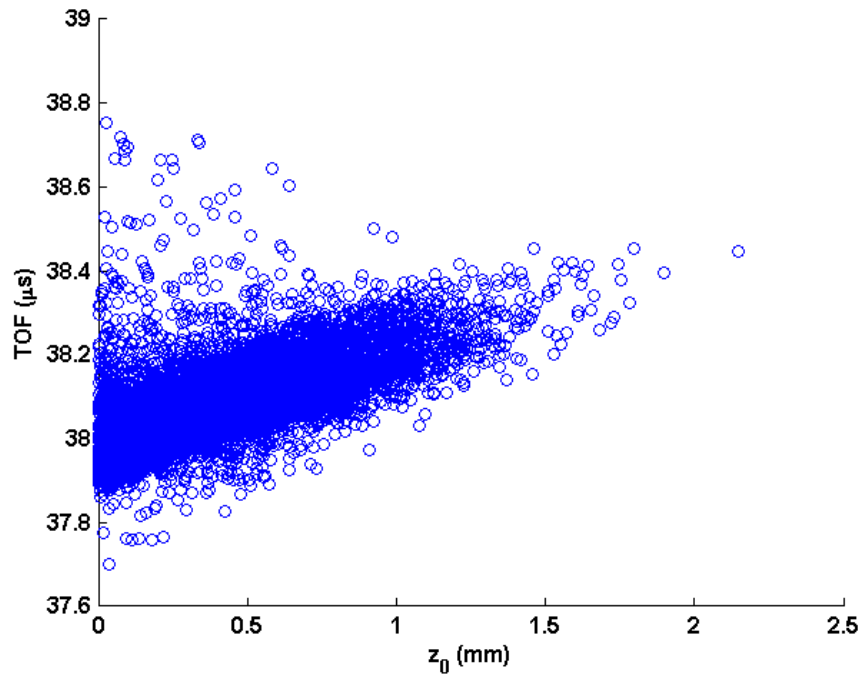
Predicted RIS time of flight:
38.6 us

This includes an initial energy
and spatial distribution for
barium.



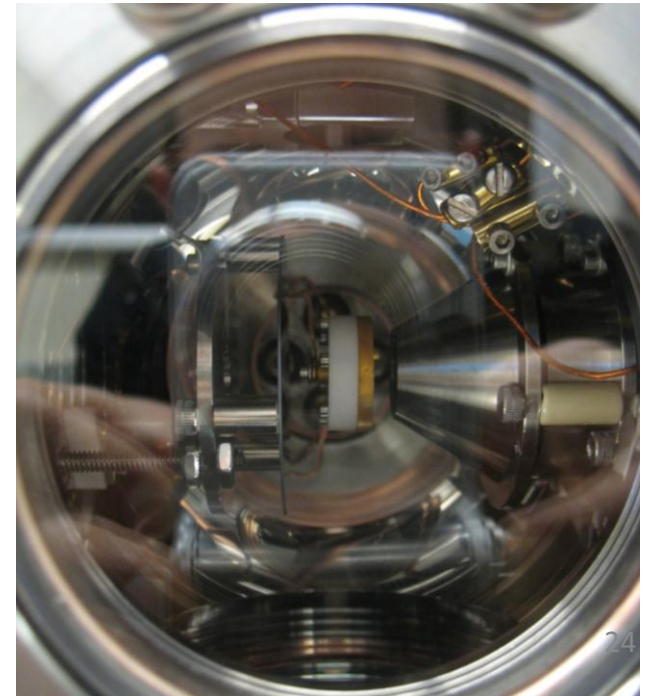
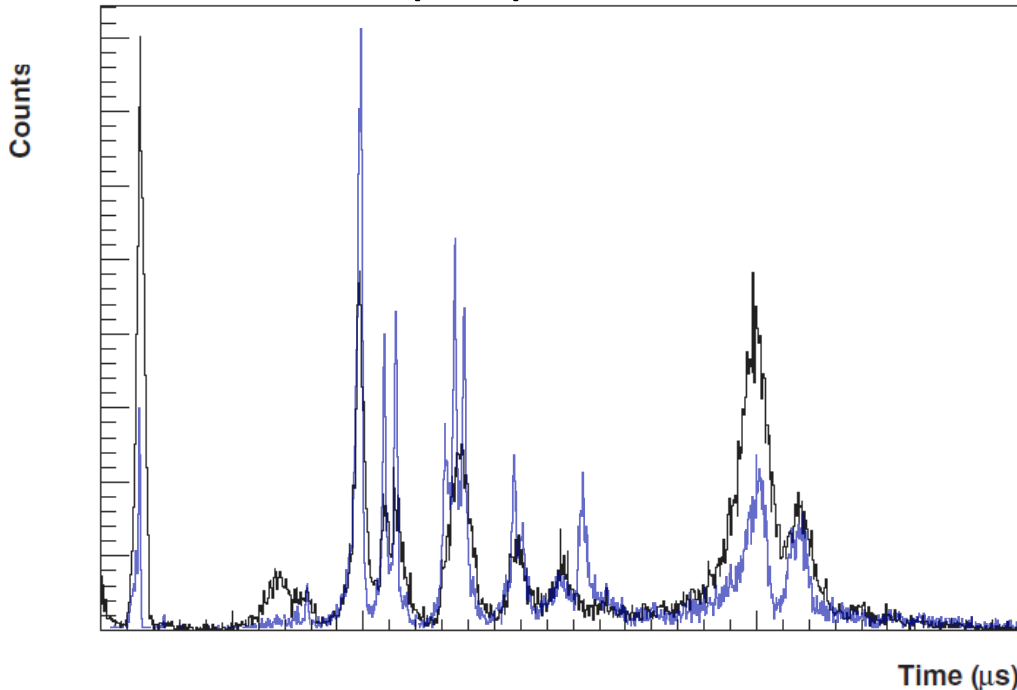


Target TOF Dependence

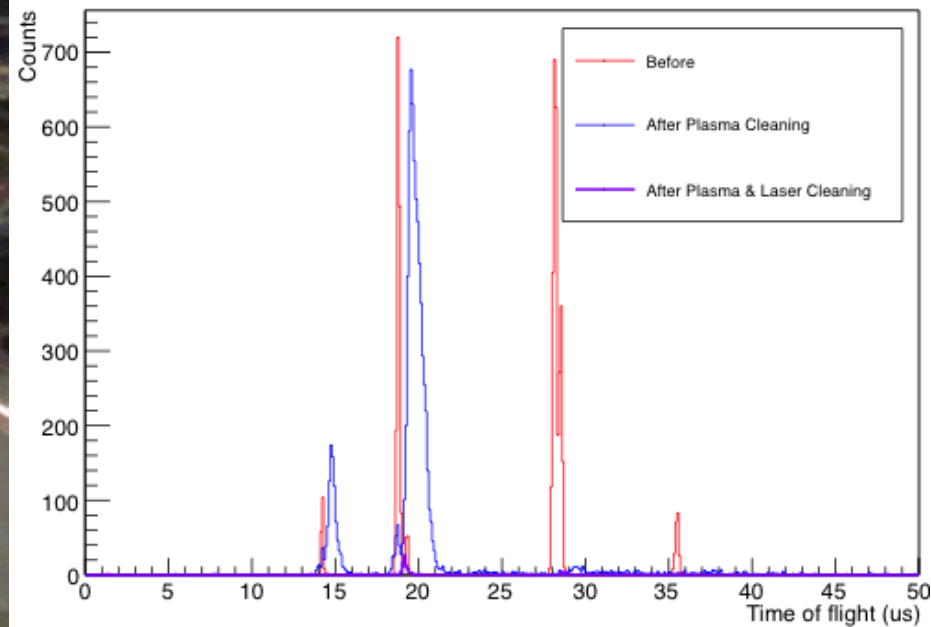
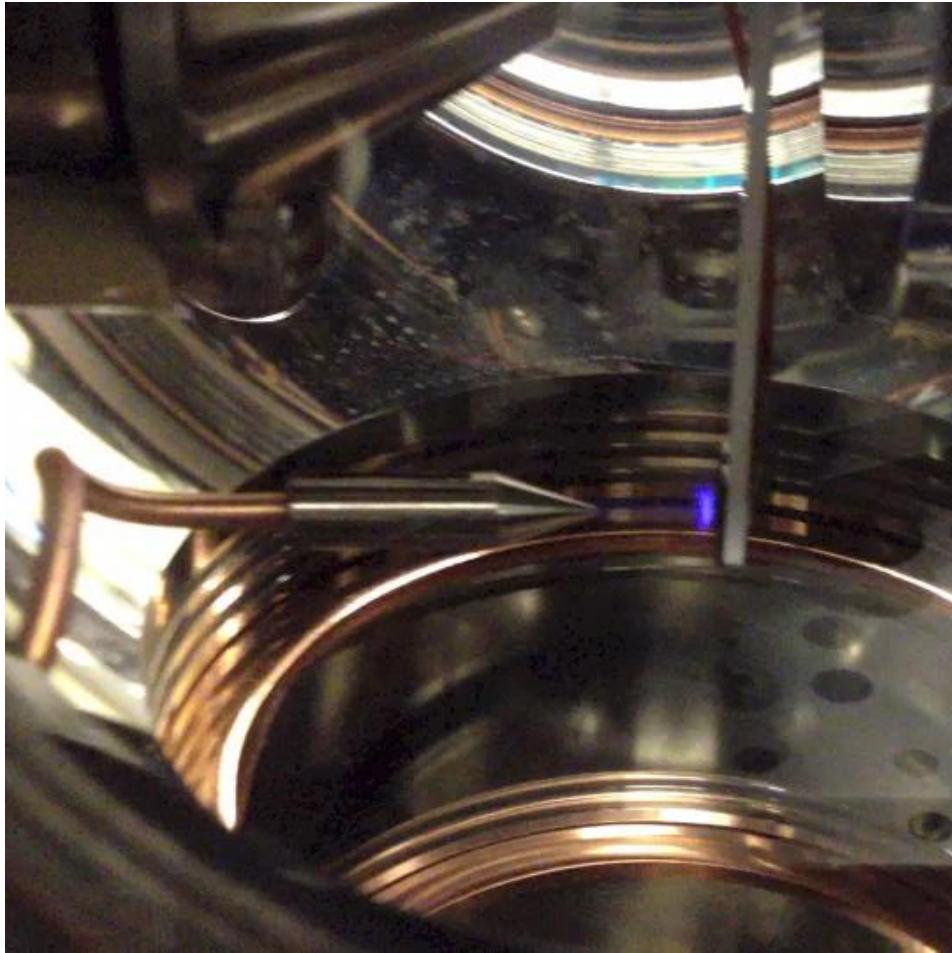


Direct Testing with Gd Source

- Spectrometer simulations: >99% efficiency, few amu resolution
- Want to test performance
- Move Ba source directly in front of spectrometer
- Previous spectrum is well-reproduced \longrightarrow success
- Peak ratios change (different source)
- Narrower/split peaks \longrightarrow better resolution

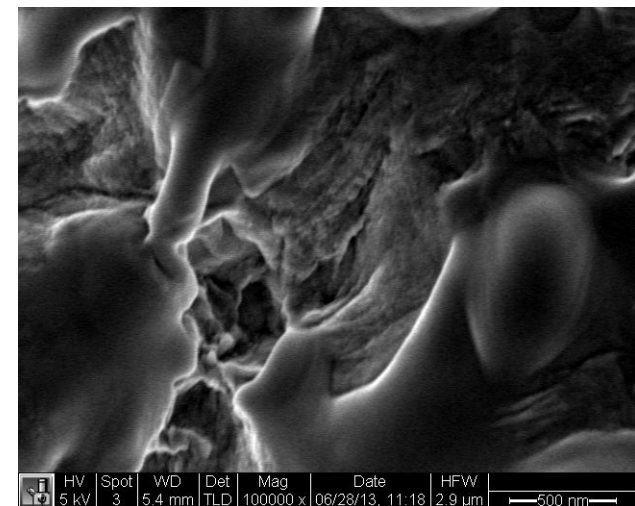
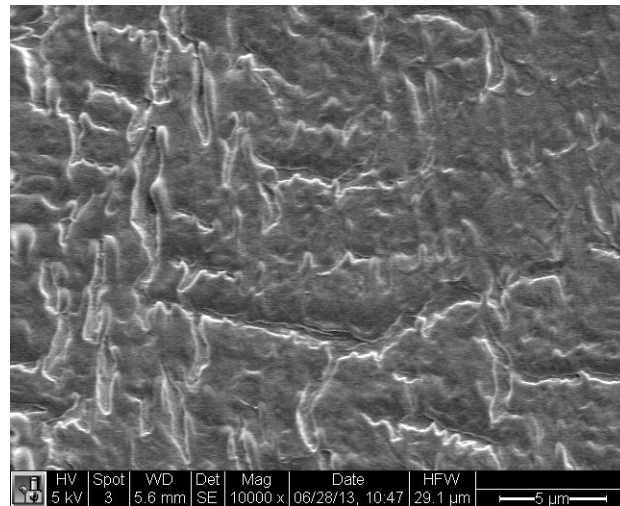
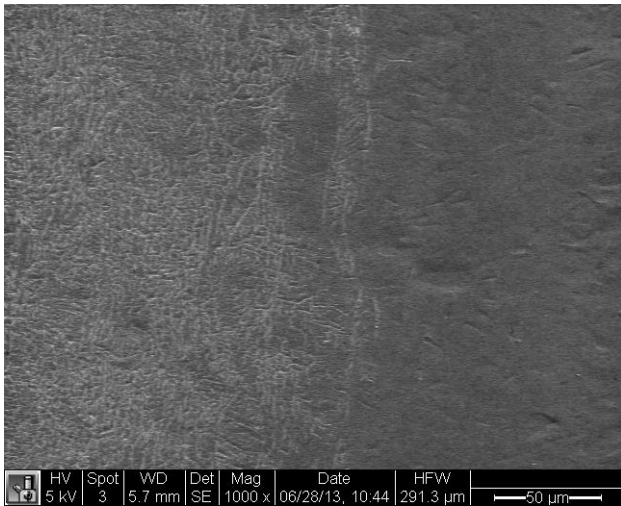


Xenon Sputtering

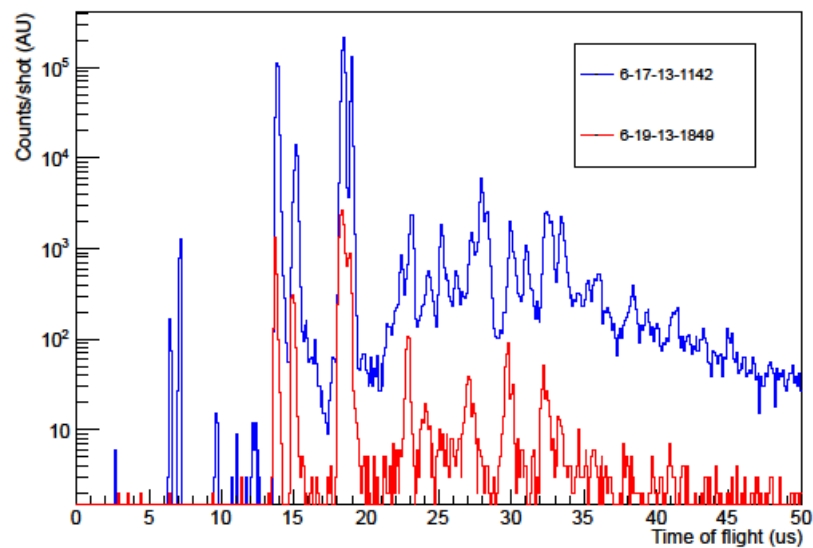


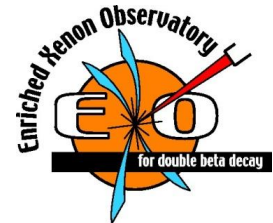
We can clean the surface by bombarding it with noble gas ions—xenon is handy!
The spike on the left is the anode, and the target (on the right) is the cathode.

Surface Damage



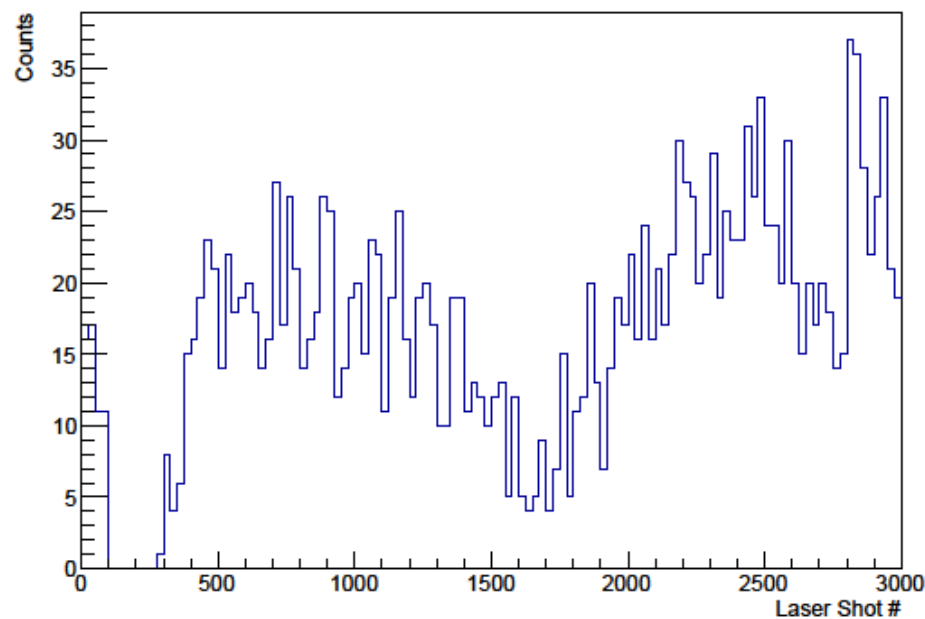
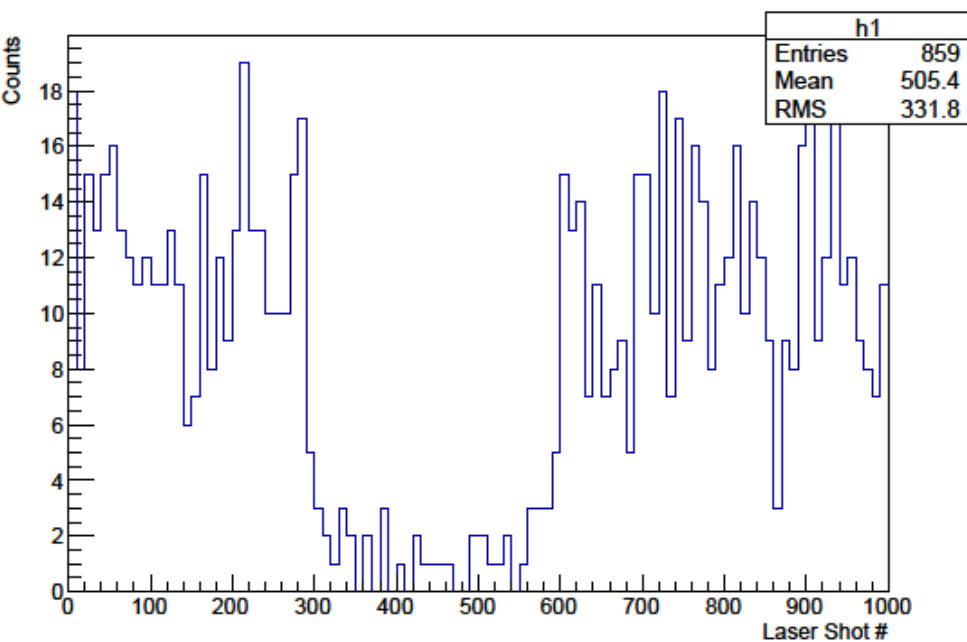
ToFs



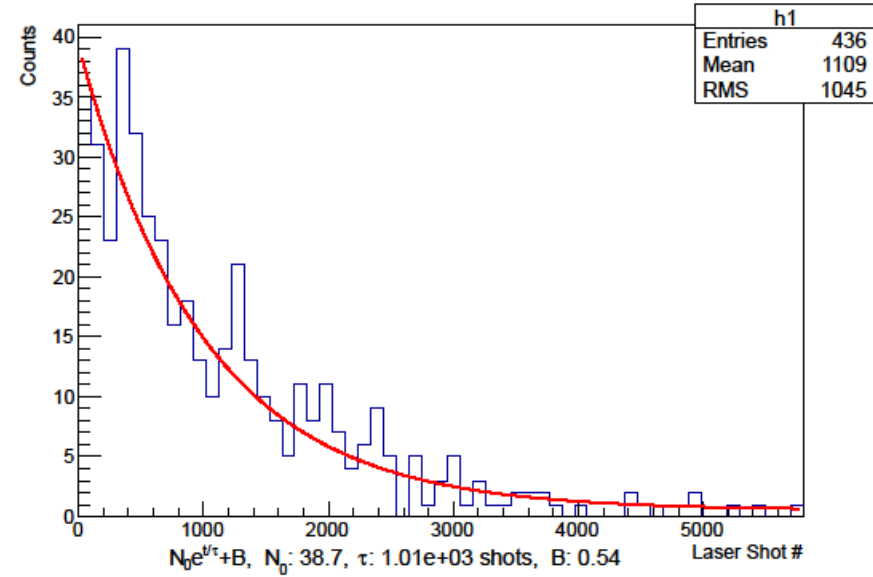
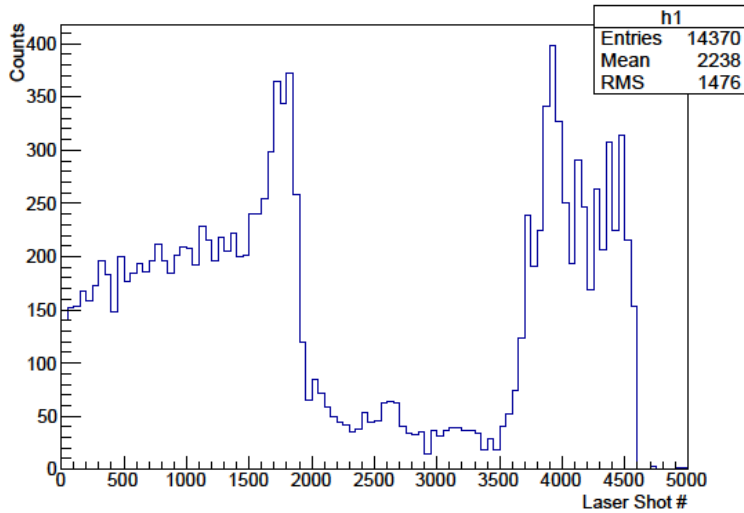


Detuning Tests

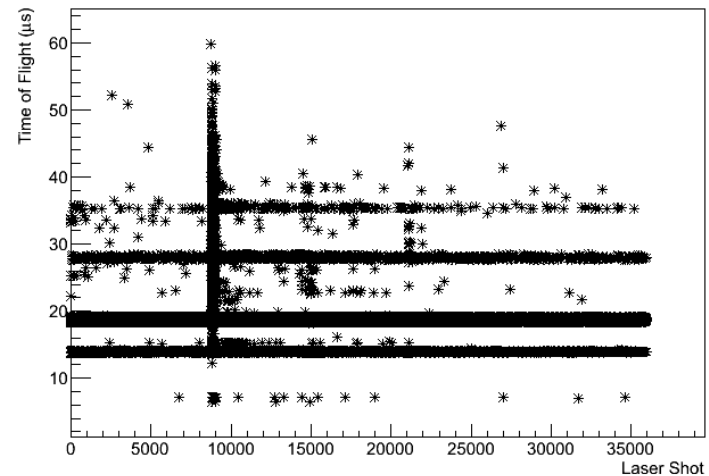
- Signal does depend on RIS laser wavelengths



Time Structure



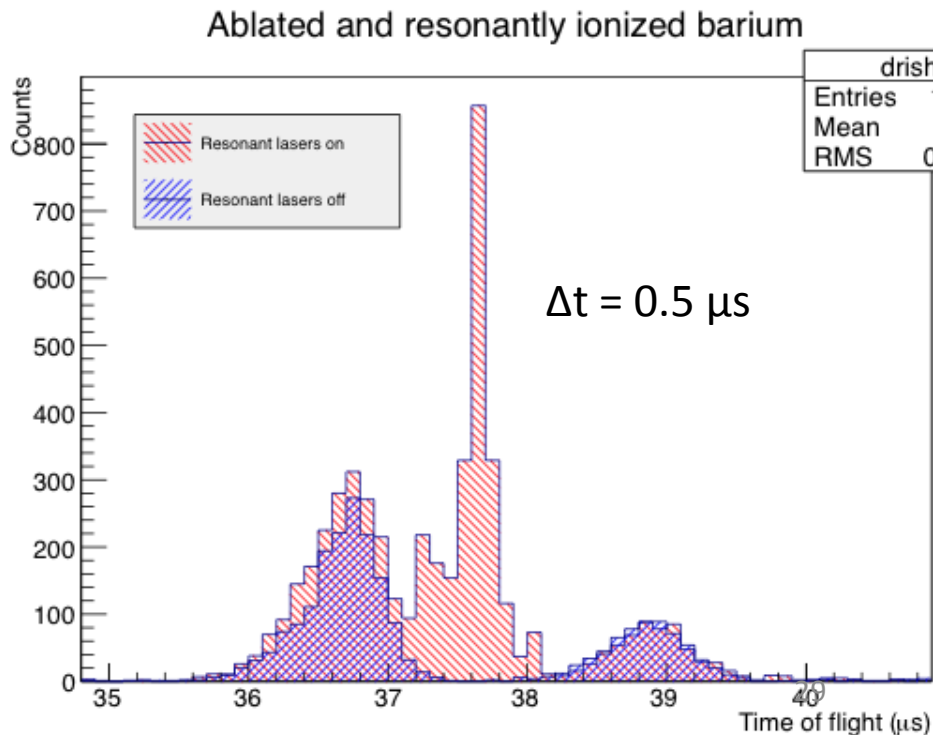
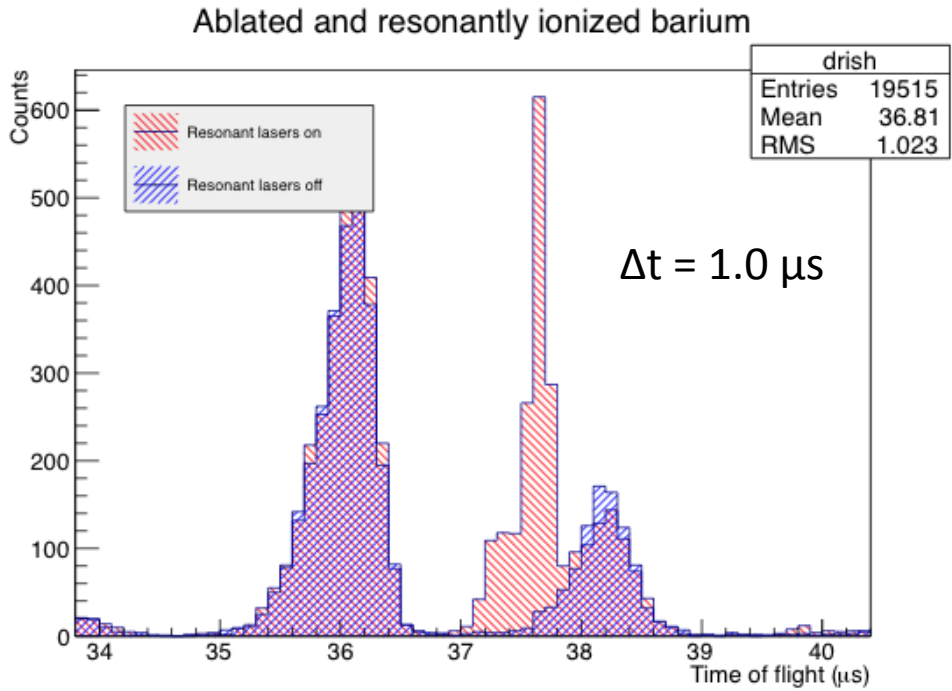
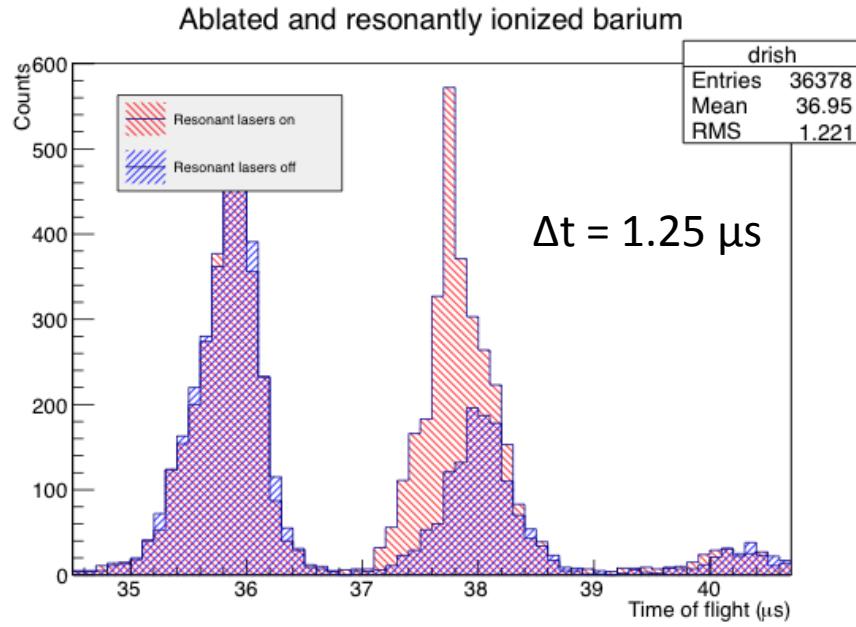
- Sometimes a nice exponential decay
- Others, have sudden arrival of signals





Laser Delay Shifts

By reducing the delay between the desorption laser and the RIS laser, we can shift the arrival time of the peak





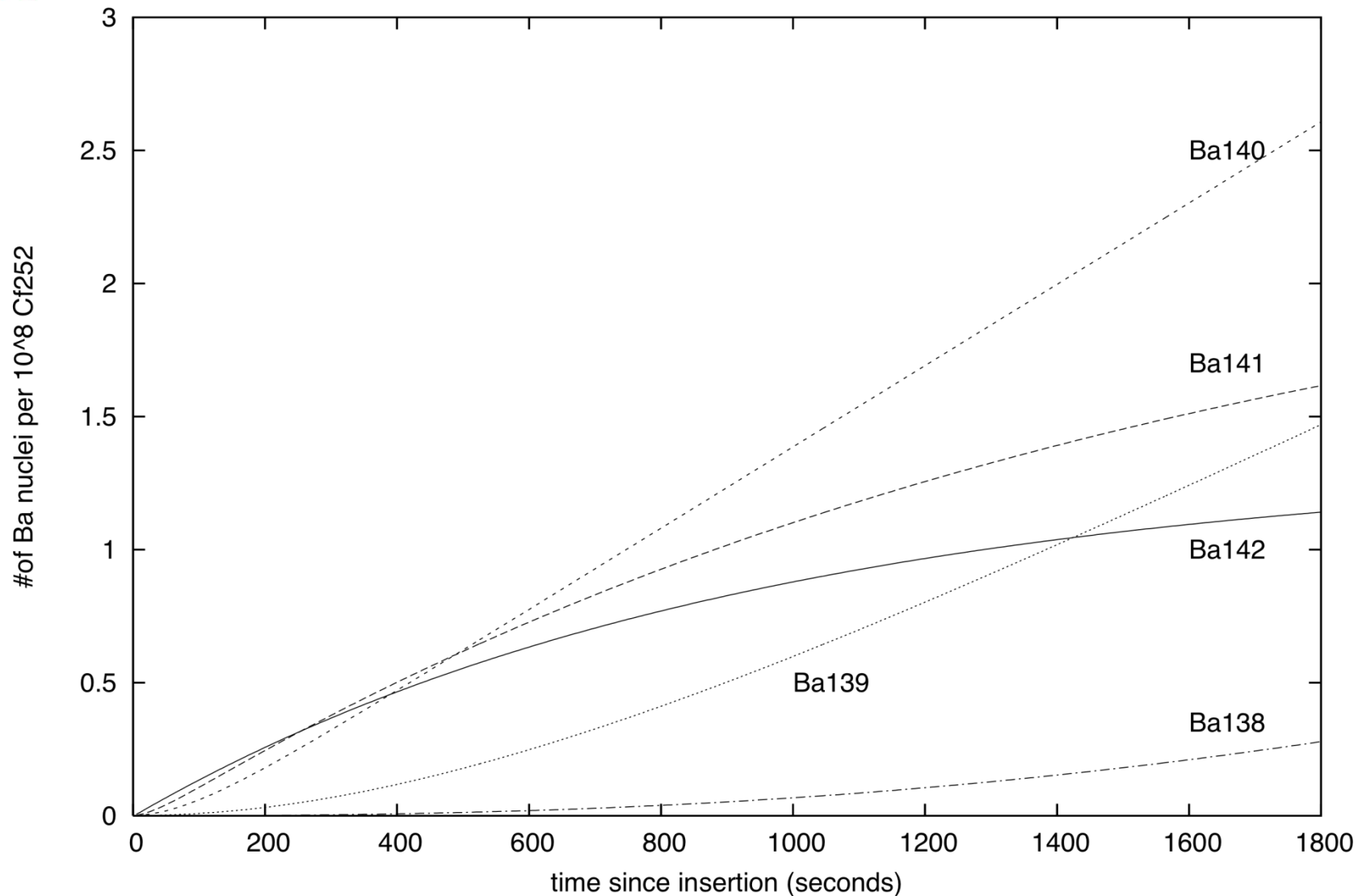
^{252}Cf Fission Source

- ^{252}Cf undergoes spontaneous fission
- Barium is $\sim 5\%$ of the fission fragments
- Most isotopes are radioactive, ^{138}Ba is not.
- The number of all Ba isotopes will constantly change.

Ba Isotope	Half-Life
138	Stable
139	83 min
140	13 days
141	18 min
142	11 min
143	15 sec
144	12 sec
145	4.3 sec
146	2.2 sec



Ba isotopes with large yields



10^8 ^{252}Cf is 0.83 Bq = 22 pCi

Plot courtesy of Petr Vogel ³¹