





Current status of MORA at IGISOL

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Contents

- CP Violation and D correlation, the search for baryogenesis
- MORA Setup, IGISOL
- Latest experimental developments
- Outlook and conclusions



CP Violation and D correlation, baryogenesis





In search of CP Violation

- Matter and antimatter should have been created in equal amounts
- Why do we only see baryonic matter?
- Sakharov conditions for baryogenesis
 - Baryon number violation
 - CPV
 - Process out of equilibrium

- CPV known in the SM
- Seen in decay of **K**, **B** and **D** mesons
- Not enough to explain baryogenesis
- Where to find CPV
 - In the SM: complex phase CKM matrix
 - Outside SM: D-correlation of beta decay





β decay D-Correlation

In a **GT-F** mixed β -decay the energy phase space can be written as *

$$\omega\left(\langle J\rangle|E_e,\ \Omega_e,\ \Omega_{\nu}\right)dE_ed\Omega_ed\Omega_{\nu} = \frac{F(\pm Z,E_e)}{(2\pi)^5}p_eE_e(E^0-E_e)^2dE_ed\Omega_ed\Omega_{\nu}$$

$$\times \xi \left\{ 1 + \underbrace{\mathbf{p}_e \cdot \mathbf{p}_\nu}_{E_e E_\nu} + \underbrace{\mathbf{p}_e \cdot \mathbf{p}_\nu}_{E_e} + c \left[\underbrace{\mathbf{p}_e \cdot \mathbf{p}_\nu}_{3E_e E_\nu} - \frac{(\mathbf{p}_e \cdot \mathbf{j})(\mathbf{p}_\nu \cdot \mathbf{j})}{E_e E_\nu} \right] \left[\frac{J(J+1) - 3\langle (\mathbf{J} \cdot j)^2 \rangle}{J(2J-1)} \right] \right\}$$

$$\frac{\langle \mathbf{J} \rangle}{J} \cdot \left[\underbrace{\mathbf{A}_{E_e}^{\mathbf{p}_e} + B \frac{\mathbf{p}_{\nu}}{E_{\nu}} + D \frac{\mathbf{p}_e \times \mathbf{p}_{\nu}}{E_e E_{\nu}}}_{} \right]$$

 $oldsymbol{eta}$ asymmetry

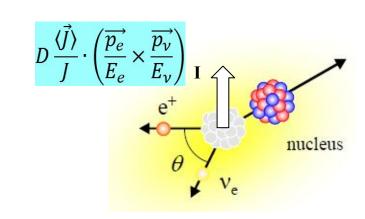
 β - ν correlation

Fierz interference

D correlation

D is **non-zero** for **T** reversal violation

* Jackson, J. D.; Treiman, S. B.; Wyld, H. W. (1957) Phy Rev 106(3), 517–521



A precise measurement of **D** would allow to:

- Give direct constraints on CP-violating Wilson coefficients in the nucleon-level EFT
- Probe specific BSM Models, like L-R symmetric model and Leptoquark model
- Measure Final State Interactions, predicted by the SM



Determination of the D-Correlation

As previously said: we need a **GT-F** mixed decay and **polarization of the ions**

polar higher $\frac{1}{2}$ Properties

We need to maximize F(X) and the polarization degree of ions to get the highest sensitivity for CPV

Proportional to mixture degree and axial vector-vector phase

Final state interactions (**FSI**), never measured using D-correlation

From neutron measurement (emiT): $\varphi_{AV}=180.013^{\circ}\pm0.028^{\circ}$ (68% CL)

T. E. Chupp, R. L. Cooper, K. P. Coulter, et al. Phys. Rev. C 86, 035505 (2012)

$$D_{FSI} \sim Z \alpha \frac{E_e}{M} \cdot A(\mu_f - \mu_i)$$
 Callan and Treiman, Phys. Rev. 162(1967)1494.



Selection of nuclei for D-Correlation

Maximal sensitivity to F(X)

Maximal spin **polarization**

Two cases to separate new physics from FSI

Well produced in GANIL, easy to polarize, short half-life, GT-F decay Shorter half-life, opposite sign for FSI, good candidate for IGISOL

	neutron	¹⁹ Ne	²³ Mg	³⁵ Ar	(39Ca)
Sensitivity F(X)	0,43	-0,52	-0,65	0,41	0,71
D ₁ (x10 ⁻⁴)	0,11	2,31	2,64	0,43	-0,47
D ₂ (x10 ⁻⁴)	0,02	0,17	0,16	0,01	-0,02

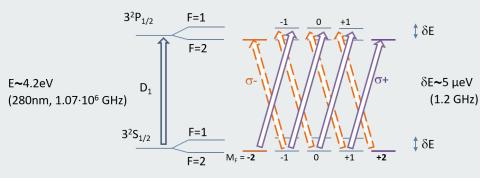
$$D_{neutron} = (-0.94 \pm 1.89 \pm 0.97) \cdot 10^{-4}$$
 $D_{19Ne} = (1 \pm 6) \cdot 10^{-4}$

$$D_{FSI}(p_e) = \left(D_1 \cdot \frac{p_e}{p_{emax}} + D_2 \cdot \frac{p_{emax}}{p_e}\right) \times 10^{-4}$$

Callan and Treiman, Phys. Rev. 162(1967)1494. Chen, Phys. Rev. 185(1969)2003.



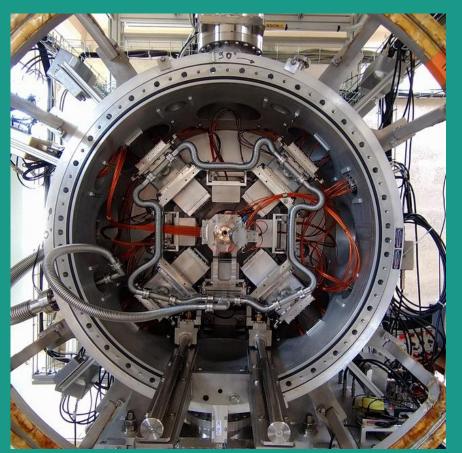
Laser polarization of ²³Mg



- Pump the ions to $M_F = \pm 2$ for spin J polarization
- We use a dipole excitation with a circular polarized laser to increase(decrease) the projection number M_F
- ³⁹Ca has a more complex structure to polarize than ²³Mg (2 lasers vs 1 laser)
- 23Mg is the best candidate for the first experiments at IGISOL
 - First candidate for DESIR
 - Easy to produce and polarize
 - High sensitivity to FSI and new physics



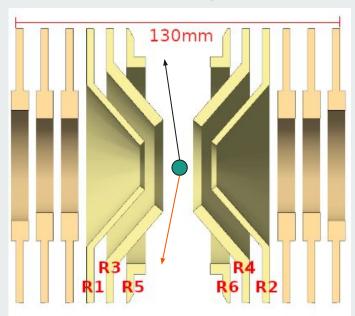
MORA Setup, IGISOL





MORA Trap

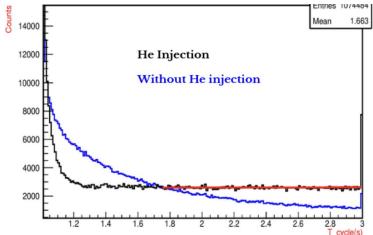
Open Paul Trap developed in LPC Caen



Paul Trap consisting on 3 pairs of **electrodes** (R1-R6) and 2 **Einzel lenses**

The electrodes (R1-R6) trap the ions

Einzel lenses focus the beam

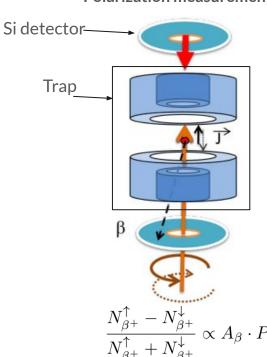


He gas is used to cool the ion cloud and optimize trapping efficiency



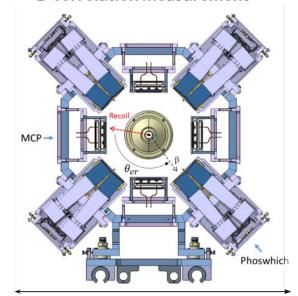
MORA Setup

Polarization measurement



- **Paul Trap** in the middle of the detection system
- 2 annular **Si detectors** in the line axis
 - For continuous polarization measurement by betas
- 4 MCP detectors, each 90° apart from each other
 - For recoil ion detection
- 4 phoswich detectors, between the MCP detectors
 - For beta detection
- Laser setup for polarization

D correlation measurement

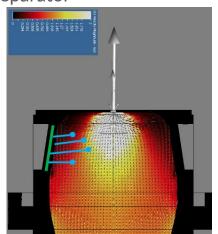


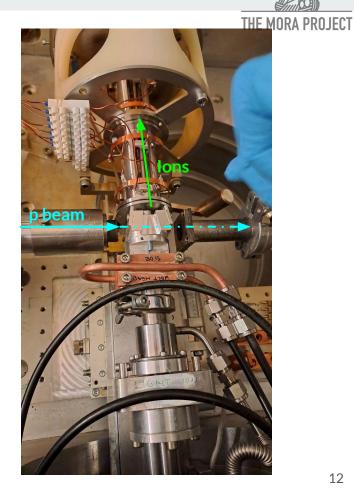
$$\frac{N_{coinc}^{+45^{\circ}} + N_{coinc}^{+135^{\circ}} - N_{coinc}^{-45^{\circ}} - N_{coinc}^{-135^{\circ}}}{N_{coinc}^{+45^{\circ}} + N_{coinc}^{+135^{\circ}} + N_{coinc}^{-45^{\circ}} + N_{coinc}^{-135^{\circ}}} = \delta \cdot D \cdot P$$

Ion Production at IGISOL

- Radioactive ions produced by the **IGISOL** method. Collision of a primary beam (proton, 3He...) with a target in a gas cell.
- Transfer reactions, ²⁴Mg(p,d) or ²⁷Al(³He, ⁴He)
- lons transported within **purified He gas** into the Sextupole Ion Guide (SPIG), later accelerated at 30keV towards the **Dipole Magnet** mass separator

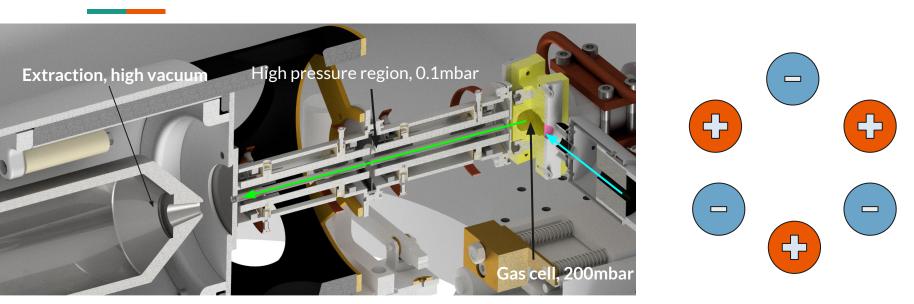








Sextupole Ion Guide



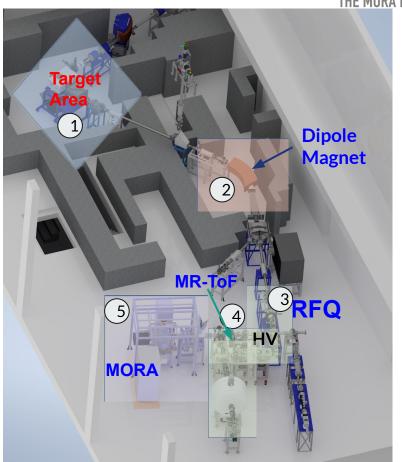
RF sextupole ion guide, 3-5MHz, $\sim 1kVV_{pp}$

Transports ions from **high pressure** (~0.1 mbar) section into **high vacuum**



IGISOL Beamline

- 1. Ion production
- 2. Isobar (A/q) separation
- 3. Ions enter the HV cage, at 30kV
 - a. Ions get cooled and bunched in the RFQ Cooler Buncher
 - b. When exiting, they get accelerated to 2keV
- 4. Ions travel through the MRToF, where mass separation can be done
- 5. Arrival to MORA, at ground potential
 - a. Pulsed Drift Tube (PDT) to slow down and bring to ground potential the ions
 - b. Second PDT to further slow down the ions
 - c. MORA Trap and detection setup
 - d. The ions get ejected from the trap





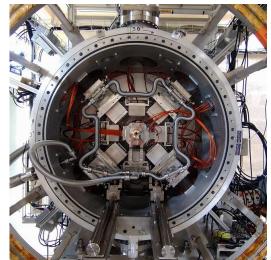
Latest developments

Laser polarization proof-of-principle Beam purification efforts

MORA Experiment at IGISOL

- Proton beam @30MeV hits the target, ²⁴Mg(p,d)²³Mg reaction.
- lons cool down and charge exchange to 1+, travel through SPIG and go to dipole magnet
- Choose Isobar (A/q=23) and send beam to RFQ
 Cooler buncher
- Beam gets cooled and gets bunched for ~2.7ms.
 Sending one bunch to MORA every 11s.
- Bunched beam arrives to MORA PDT, where it gets pulsed down to -1kV to send to ground, ions at ~1keV
- Bunched ions arrive to second PDT, where they get further deaccelerated to ~100eV
- lons get trapped for 6s

- After trap, the ions get ejected
- **5s of "off-trap" measurement** for active background
- Measurement repeated for two different circular polarizations, changing every hour



First conditions of MORA, Nov 2022

Typical conditions of IGISOL

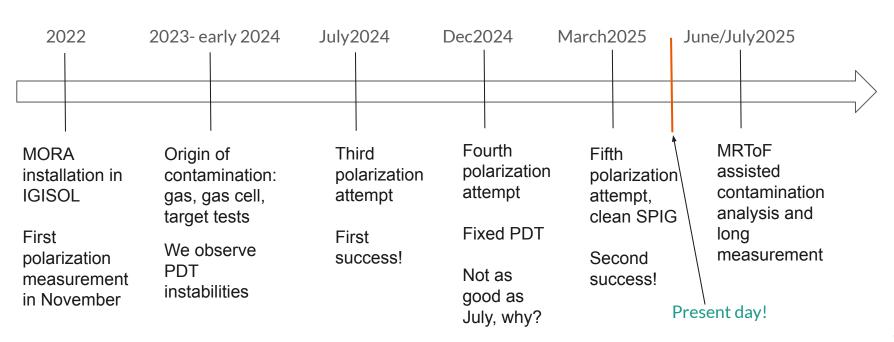
- He gas purified by zeolite cold trap and getter purified
- ~200mbar of He gas inside of gas cell
- Ti windows on the gas cell, nat Mg target facing the beam
- 5MHz SPIG RF, usual extraction voltages
- Usual Al ion guide

- Too high contamination, getting only ~10
 ²³Mg ions per trap cycle, even with high production (>10k ions/s)
- Impossible to do polarization proof-of-principle
- The culprit: ²³Na contamination
- The RFQ Cooler Buncher has a limited charge space, that gets saturated by too much ²³Na⁺
- Since the problem is in the Cooler, we cannot make use of the MRToF for cleaning the beam
- Need to tackle contamination on the production



MORA Timeline

Biggest challenge: Na contamination for trap efficiency





In search of contamination's origin

Possible origin: Target

- Traces of Na
- Nuclear reaction
- Surface ionization
- Sputtering

Various **target tests** done, **coatings** to prevent ionization and different **target material** (AI)

No improvement

Possible origin: Gas cell

 Surface ionization / sputtering from the gas cell

Changed usual Al to graphite gas cell baked at high temperature (1300°C) for long period of time to evaporate traces

No improvement

Possible origin: Gas

- Stopping He purification showed improvement
- Traces of Na coming from Zeolite
- Forming molecules with Na that could improve yield

Various gas tests done, different traces of gases over purified He and different getter and cold trap configurations

No improvement

In search of contamination's origin

Second half 2024

- **Progress** made for **polarization measurement**
 - Managed to get a first measurement of polarization inversion with 1.53σ precision
- Fixed PDT instabilities
- Better understanding of the source of contamination
- Only the SPIG can explain what we see
 - Sputtering inside of the SPIG, affected by gas purity
 - Voltages greatly affect the contamination
 - Na only appears with beam on

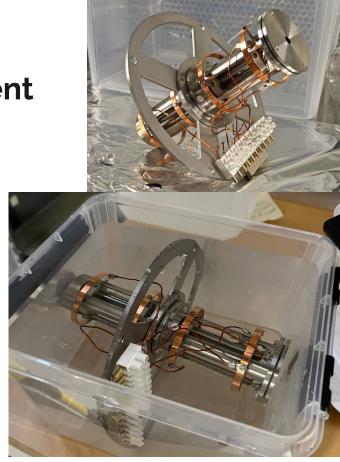


March 2025, Clean SPIG measurement

- Next step was removal of Na from all SPIG surfaces
- Used ACCLAB cooling loop water, conductivity of ~0.23uS/cm
- SPIG disassembled, cleaned and reassembled carefully to not spread contamination
- Minimal ratio of Na:Mg yet
- Best results to this date!

The **rods** are the most **critical part**, cleaned with special care

New Nb rods
 commissioned
 baked at high
 temperature for
 evaporating Na

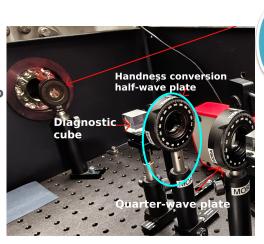


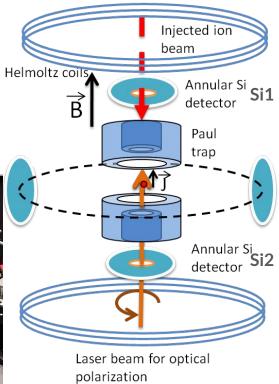
THE MORA PROJECT

Polarization proof-of-principle

- Using detections ($N_{\beta+}$) on the Si detectors (Si1 and Si2) in each polarization (+ or -, \uparrow or \downarrow) to measure the asymmetry (A)
- $A = \alpha \cdot A_{\beta} \cdot P$, where α is a sensitivity factor estimated from simulations, A_{β} is the beta asymmetry and P the polarization degree
- Needed ~10² ions per trap cycle to measure P
- We need at least ~10⁴ ions per trap cycle to attempt to measure D

$$A = \frac{N_{\beta^+}^{\uparrow} - N_{\beta^+}^{\downarrow}}{N_{\beta^+}^{\uparrow} + N_{\beta^+}^{\downarrow}} \propto A_{\beta} \cdot P$$







March 2025, Results

Asymmetry Polarization σ - : 0.72±0.25

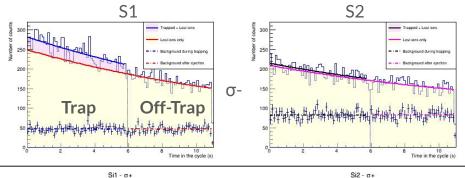
Asymmetry Polarization σ +:-0.42±0.16

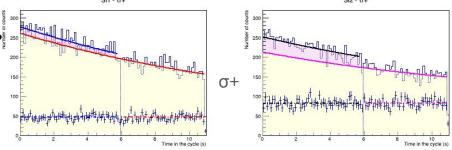
Asymmetry with full polarization of the cloud (from simulations): =0.51±0.01

 $A = 0.51 \pm 0.14$

55% < P < 100% at 90% C.L.

$A = \frac{N_{\beta^+}^{\uparrow} - N_{\beta^+}^{\downarrow}}{N_{\beta^+}^{\uparrow} + N_{\beta^+}^{\downarrow}} \propto A_{\beta} \cdot I$





Trapped ions/cycle

90±9 from Si detectors

145±55 from coincidences





Conclusions

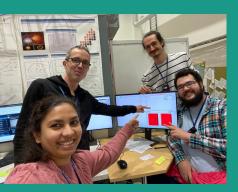
Recent developments on MORA permitted to **fix old problems** of **trapping stability** and reduce the **beam contamination** issue, allowing us to have a **first measurement of the polarization** proof-of-principle

Na contamination got low enough that we can use the MRToF to precisely measure the yield and contaminants, and explore how to improve them.

In the upcoming beamtime, we will use the new SPIG rods (Nb baked at high temperature), the contamination analysis and everything we have learned along the way, we will measure the polarization degree P with higher precision, and start the measurement of the D correlation

Thank you for your attention!







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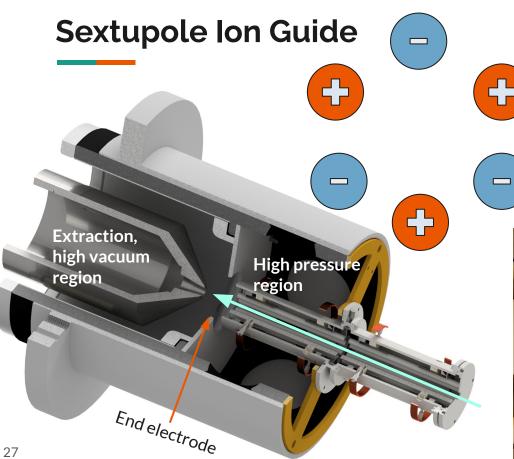






Extra slides



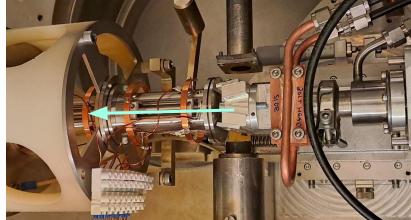


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RF sextupole ion guide, 3-5MHz, $\sim 1kVV_{DD}$

Transports ions from **high pressure** (~0.1 mbar) section into high vacuum





Source: taxget? gas cell? gas?

Data Acquisition: June2024

Installation of **baked C gas cell** trying to evaporate all Na \rightarrow No immediate improvement.

Second gas injection attempt, trying to find the gas that makes Na disappear

Gases planned to be used: CO2, N2, He+CF4, SF6, He+Kr, He+Ne

Gases used: CO2, He+CF4, He+Kr

Mass flow controller allows for precise measurement of gas flow we are introducing into the chamber





Source: taxget? gas ell? gas?

Data Acquisition: July2024

- **First polarization evidence** (1.53σ) measured
- The improvement was achieved mainly by having a full dewar of LN and fine tuning the SPIG voltages
- If the Na was produced at a lower energy/other mechanism, it is safe to assume that the extraction would be much different
- Further gas injection test did not prove useful, but it might be coming from the zeolite in the purification system
- PDT stability issues and how to fix them



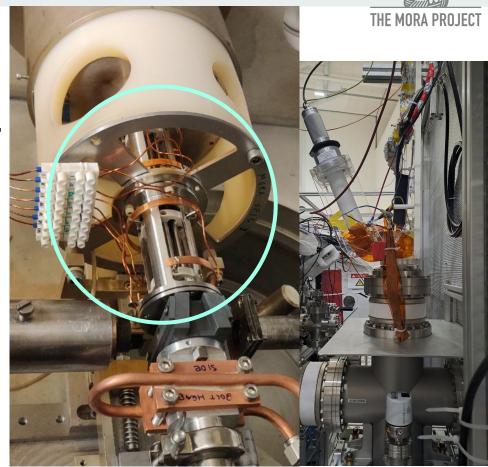
Ringing from the PDT after pulsing down



Source: taket? gas ell? gas?

Data Acquisition: Dec2024

- We used charcoal and SiO2 based cold trap to avoid any zeolite, and new filters inside of LN to simulate the events in July (full dewar)
- Immediately we saw **no improvement** in the yield Na:Mg
- The only suspect left was the SPIG
- The SPIG is coherent with all other observations:
 - Effects of gas purity/pressure
 - The Na+ can be trapped in the SPIG
 - Na+ only appears with beam on
- Fixed PDT!



In search of contamination's origin

July 2024 beamtime

- Observed improvement on ratio
 Na:Mg by completely filling the cold
 trap, getting the filter submerged in

 LN
- Instead of maximizing ²³Mg yield, we tried to minimize the contamination from ²³Na with the SPIG
- Trace gas injection did not provide useful for the Mg beam
- With this, we managed to get a first measurement of polarization inversion with 1.53σ precision

December 2024 beamtime

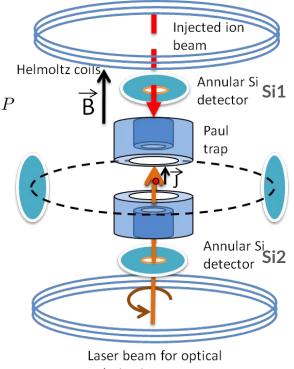
- Tried to replicate July condition, by using new cold traps with activated charcoal and SiO₂
- No improvement on contamination, but the PDT was finally fixed
- After discarding other suspects, the only possibility left was the SPIG as a source of Na.



Polarization proof-of-principle

- Using detections ($N_{\beta+}$) on the Si detectors (Si1 and Si2) in each polarization (+ or -, \uparrow or \downarrow) to measure the asymmetry (A)
- $A = \alpha \cdot A_{\beta} \cdot P$, where α is a sensitivity factor estimated from simulations, A_{β} is the beta asymmetry and P the polarization degree
- Needed ~10² ions per trap cycle to measure P
- We need at least ~10⁴ ions per trap cycle to attempt to measure D

- Using coincidences between Si and MCP to improve **signal to** noise ratio
- Then we need to use B₁, the neutrino asymmetry





Data Analysis: Calibration

The calibration fit is done with a PENELOPE simulation

Bi²⁰⁷ source for the Si detectors and 3 alpha sources for the RIDE detectors

Si calibration using 3 parameters and resolution

