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Development of New Apparatus of Ion Trap with ICP-MS For Trace Isotope Analysis

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Introduction

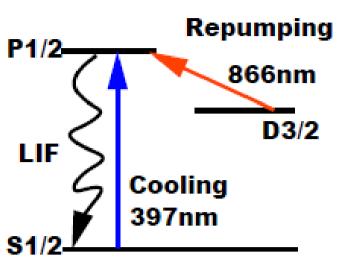
- ⁴¹Ca (lifetime 1.04×10⁵ years, γ decay 3.3keV) [1]
- . Geochronometry: date determination
- **.** Biomedicine: tracer, bone metabolism
- . Geoscience: tracer
- Planetary science: meteorite

Accelerator Mass Spectrometry [2] requires a large facility and laser based methods[3] such as Resonance Ionization Mass Spectrometry[4], Atom Trap Trace Analysis[5] have been developed.

The lon trap - Laser cooling technique has the advantages to easily manipulate atomic and molecular ions and detect single isotope ions kept in a trap.

Ca II isotope shift

Mass number (abundance)	Isotope shifts [MHz]		Neclear
	S-P: 397 nm	D-P: 866 nm	spin
40 (96.9%)	0	0	0
41 (~ 10 ⁻¹⁴)	unknown	unknown	7/2
42 (0.647%)	425(6)	-2350(4)	0
43 (0.135%)	c.g.:688(17)	c.g.:-3465(5)	7/2
44 (2.09%)	842(3)	-4495(4)	0
46 (0.004%)	1287(4)	-6478(8)	0
48 (0.187%)	1696(6)	-8288(7)	0
(1) (1)			-



V_{PZT}

IR1 On

80

120

Long focus ion lens system

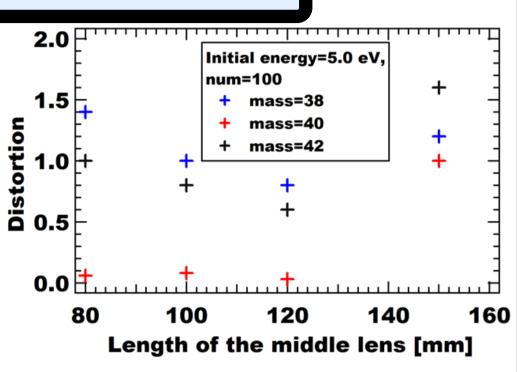
To obtain high vacuum environment at ion trap region, low conductance ion guide system from ICP-MS to Ion trap is required. A long focus ion lens system is suitable to this requirement. Simulation of the mass dependency at long focus lens system was performed with **SIMION**;

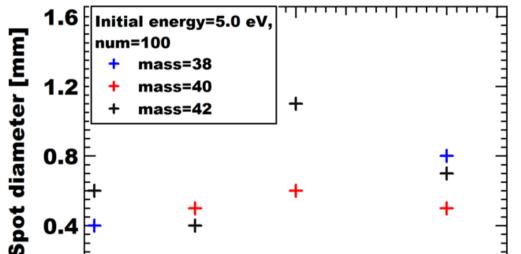
- Three ring electrodes : Inner diameter 10mm,
- Length of the first and last lens 10mm
- Spot diameter : 0.2~1.2 mm

The distortion of the ion beam is calculated as a function of the length of the middle lens.

The dependence of the spot size (at 300mm behind the lens) on the mass is also simulated.

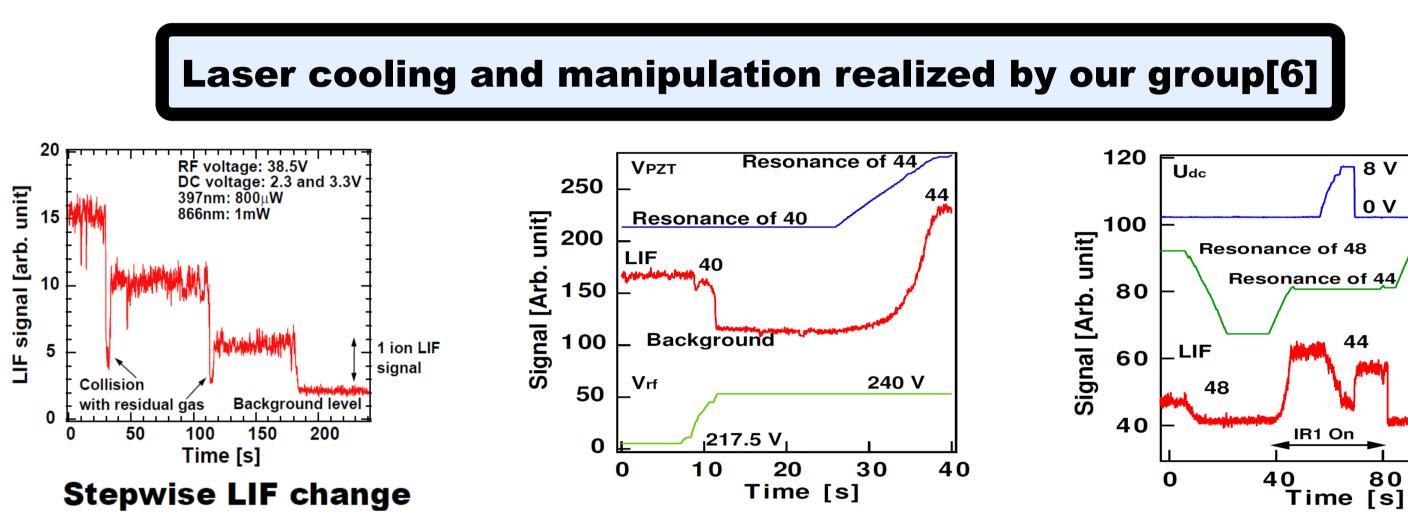
Based on these calculations, the length of the middle lens is determined as 100 mm. A following ion guide





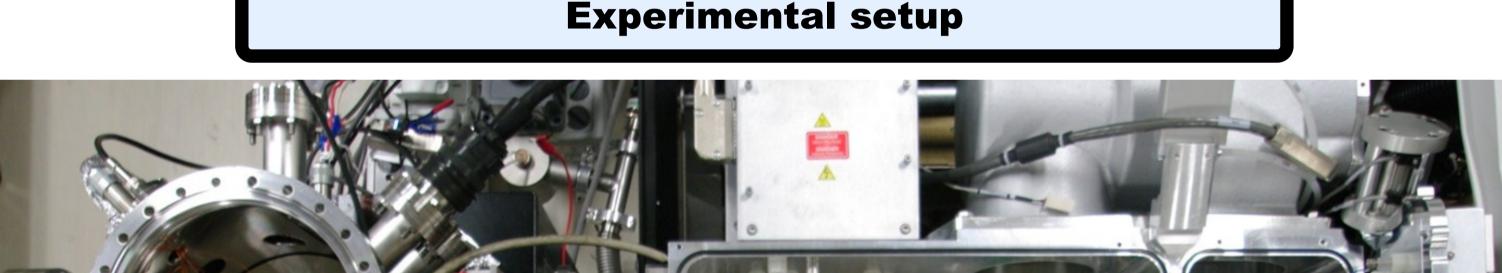
Direct injection of a liquid sample to the apparatus is preferable in analytical chemistry, which can be realized by ICP-MS.

We have combined two techniques and developed a new apparatus for trace isotope analysis.



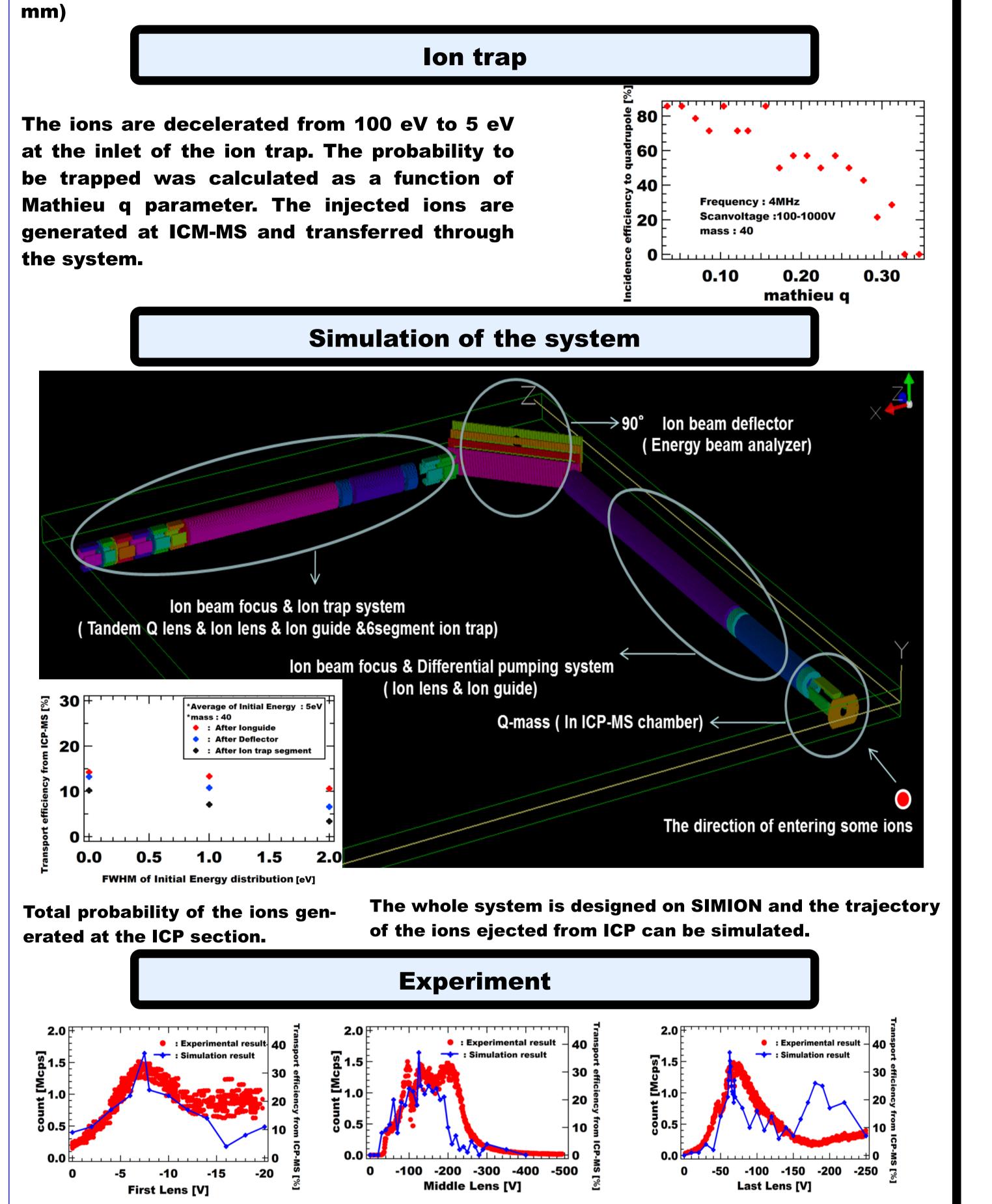
Single ions detection: stepwise difference of LIF signal corresponds to individual single ions.

Manipulation of each isotope: low and high pass filters by nonlinear resonance of Mathieu parameters



with static electric field limits the conductance, the diameter of which is 4 mm. (Length = 300mm and 150

160 Length of the middle lens [mm]



parallel plate type **ICP-MS** chamber static deflector Ion guide Einzel lens sample Q-mass filter Q-len DC shutter by static plate **Design Requirements of the system 1. Differential pumping** lon guide => long focus ion lens + static ion guide Trap segment **2. Deflection of ion's trajectory** z [mm] channeltron => laser and direct detection **3.** Ion trap => laser cooling

⁴⁰Ar⁺ ions are extracted from ICP-MS to the detector placed behind the deflector to investigate the performance of the ion les system. The experimental results are in good agreement with the numerical ones. The main reason of the discrepancy may derive from the space charge effect.



- We have designed and constructed a novel apparatus of ICP-MS-TRAP.
- SIMION calculations were performed to obtain the parameters of the ion optics to maximize the transport efficiency of ions.
- The extraction of the ions from ICP-MS was performed and the ion beams

Parallel deflector

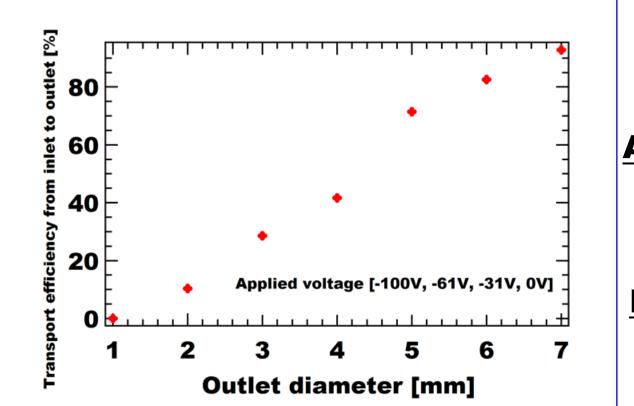
Laser

A simple deflector which consists of four parallel plates is used for the deflection region. The advantages of the deflector are;

- Simple configuration
- Suitable for beams with low kinetic energy
- Beam focusing at outlet plane

The inlet diameter of the deflector is 5.6 (4 sqrt{2}) mm. According to the simulation, larger outlet diameter gives higher efficiency as expected.

The outlet diameter of the deflector is determined as 6 mm because the diameter of the following lens is 6 mm.



were measured at the detector behind the deflector.

Future works

- Guidance of the ion beams to the trap
- Laser cooling of ions with this apparatus
- Realization of trapping ions from ICP-MS
- Optimization of the experimental system for detecting trace isotopes

Acknowledgments

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References

[1] Science <u>236</u>, 725(1987). Nucl. Instrum. Methods Phys. Res., Sect. B <u>52</u>, 531(1990). Nucl. Instrum. Methods Phys. Res., Sect. B <u>52</u>, 572(1990). [2] Nucl. Instrum. Methods Phys. Res., Sect. B <u>172</u>, 399 (2000). [3] Rev. Sci. Instrum. 74, 1169(2003). [4] Nucl. Instrum. Methods Phys. Res., Sect. B 229, 519 (2005). [5] Phys. Rev. Lett. <u>92</u>, 153002(2004). [6] J. Nucl. Sci. Technol. <u>43</u>, 300(2006). Spectrochimica Acta. B 63, 645(2008). Int. J. Mass Spectrom. 279, 163(2009).