



TULIP Project
GANIL



Producing short-lived isotopes by fusion-evaporation reactions in the TULIP TISS at GANIL

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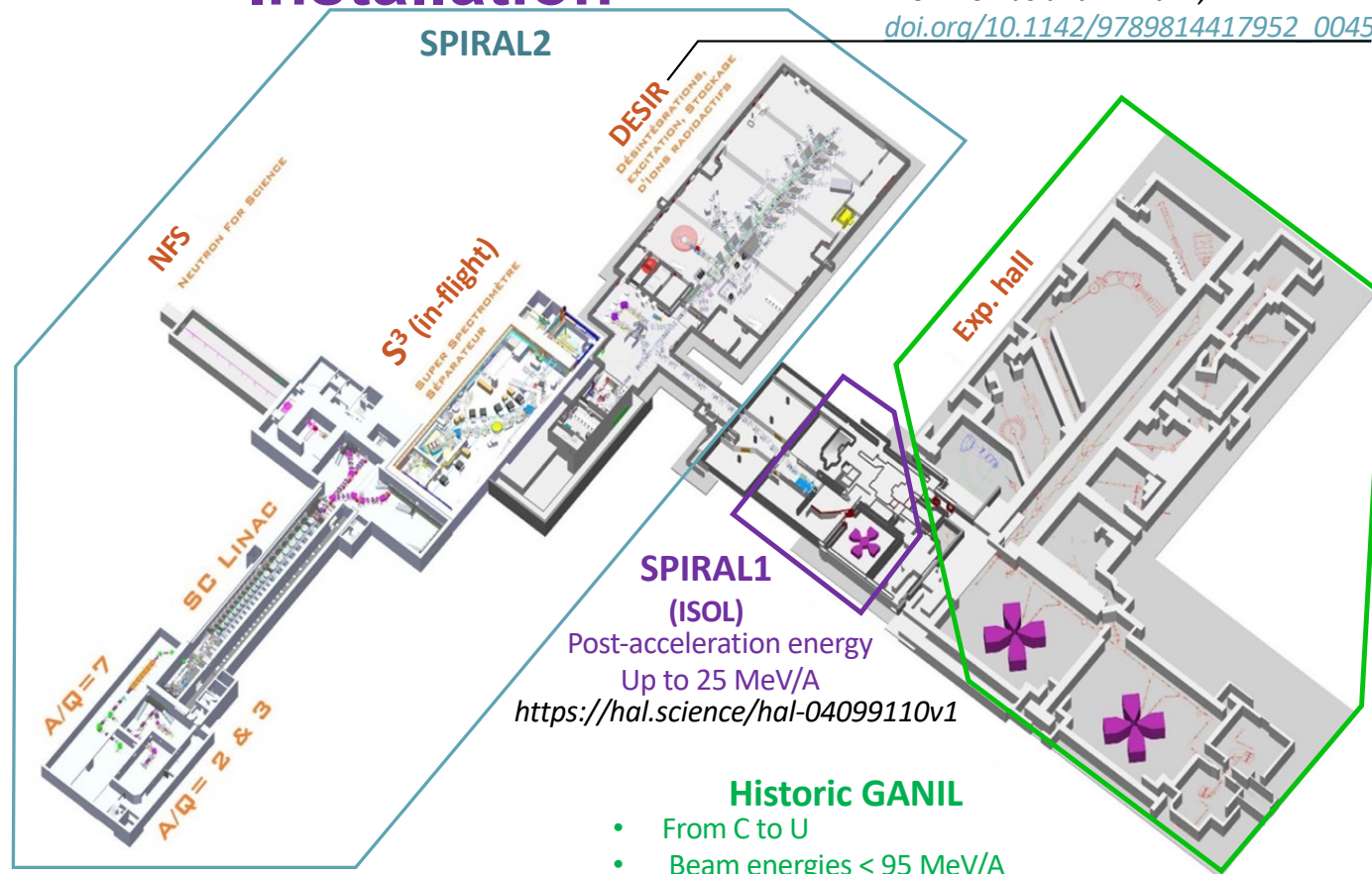
GANIL Installation

Energy ranges :

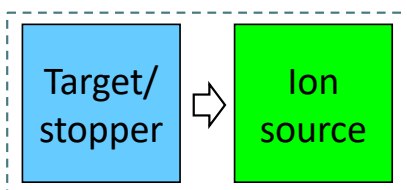
33 MeV for protons
40 MeV for deuterons
< 14.5 MeV/A for heavy ions

High intensities:

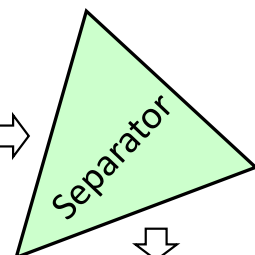
up to 5 mA for deuterons
< 1mA for heavier ions



The ISOL technique

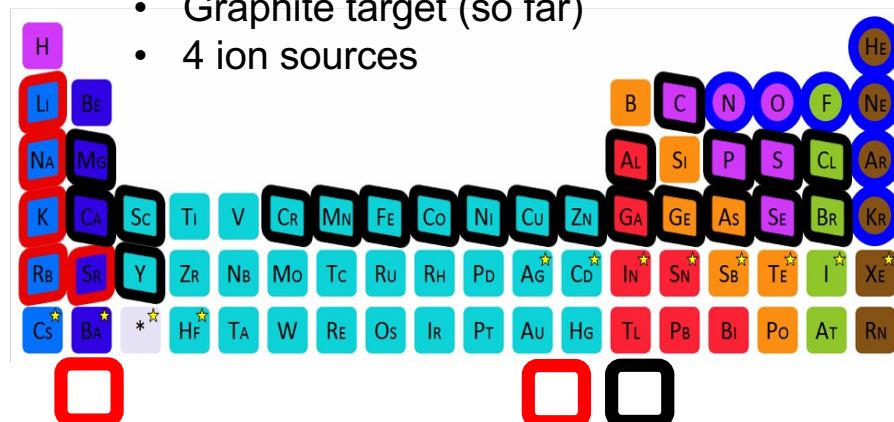


TISS

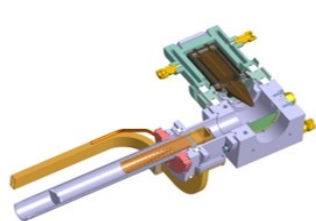


To user/
acceleration

- 58 primary beams from ^{12}C to ^{238}U
- Graphite target (so far)
- 4 ion sources



N+



Nanogan
For gas

1+



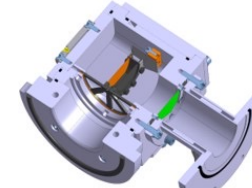
FEBIAD
For condensable

1+



MonoNaKe
For alkaline

1+



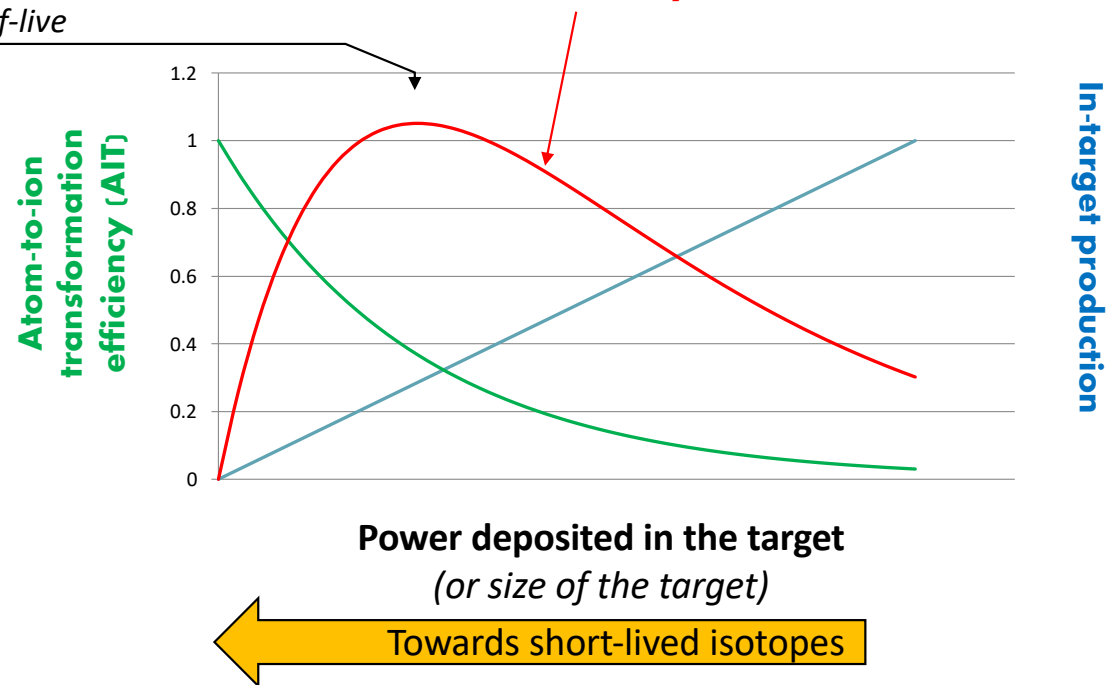
TULIP (SIS or FEBIAD)
For neutron deficient isotopes

For more details, see P. Chauveau, Proceedings of ICIS 2025 conference, Oxford

Origin of the TULIP project

*Position and magnitude
depend on the half-life*

Atom-to-ion transformation Eff x In-Target Production
= RIB intensity



TULIP project questions

Is it possible to produce competitive RIB intensities by favoring the Atom-to-Ion Transformation efficiency rather than the in-target production?

Why address this issue?

How to go about it?

Ideal case

Use of a code able to predict the optimum design
=> Requires all the right input parameters

Without code

Minimize the time of each process

- Diffusion
- Effusion
- Ionization

Chose the primary beam-target couple allowing this minimization

Case study: production of $^{74}\text{Rb}^+$. Why?

Regarding the TISS design

Possible to produce by fusion-evaporation reaction

- Low energy ($\sim$ Coulomb barrier)
 - Low recoil energy of the residues
 - Short range in material (few μm)
 - **>> fast diffusion**
 - Low power
 - small target volume
 - **>> fast effusion**
 -
 - Ionize by hot surface
 - High efficiency
 - Selective
 - And Selective in-target production

Regarding the physical interest

- Region of the nuclide chart little explored with ISOL systems
- Very exotic (11 neutrons from the ^{85}Rb)
- Short lived isotope (64,8 ms)
- N=Z nucleus, $0^+ \rightarrow 0^+$ superallowed emitter

Regarding the operation : low cost solution

- Reduced activation of the TISS and of its surrounding
- Limited production of artificial radioactivity
- Short delay before recycling



Technical principle for $^{74-78}\text{Rb}^+$ production

Production of radioactive isotopes by fusion-evaporation reactions

Production target
(Mo/ ^{nat}Ni /Mo)
(0,3 μm /4 μm /0,3 μm)

Window
Graphite, 2 μm thick

Graphite catcher
~1500°C
100 μm

Diffusion

Implantation Thickness (~10 μm)

Surface ionization

Electric field

Ions pushed towards the exit aperture

Accelerated ion beam

HV

TISS Exit aperture

High current for ohmic heating and E field creation

Collection of the residues in the cavity independent of

- The recoil energy spread
- The angular spread
- The charge state of residues

→ Target thickness can be maximized (up to ~6 μm)

Present design

Cavity: 12 mm long, 40 mm in diam.

Material: graphite

Heating power: up to ~3 kW + 70 W on catcher

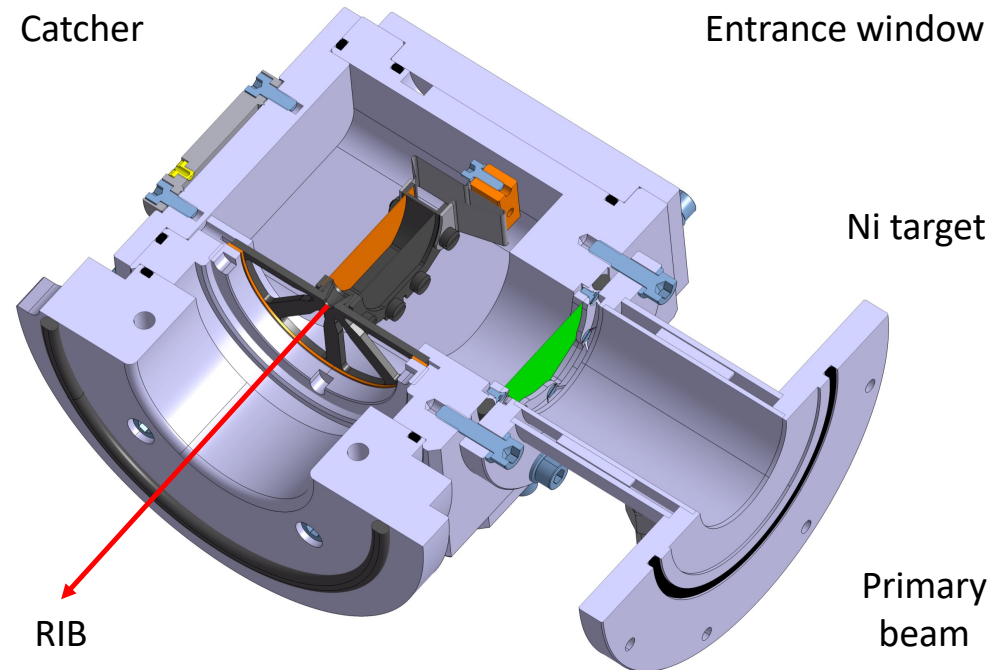
Temperature: 1500-1600°C

Target

0,3 μm Mo/4 μm $^{\text{nat}}\text{Ni}$ /0,3 μm Mo*

Beam power deposited: 30 W/100W of ^{20}Ne

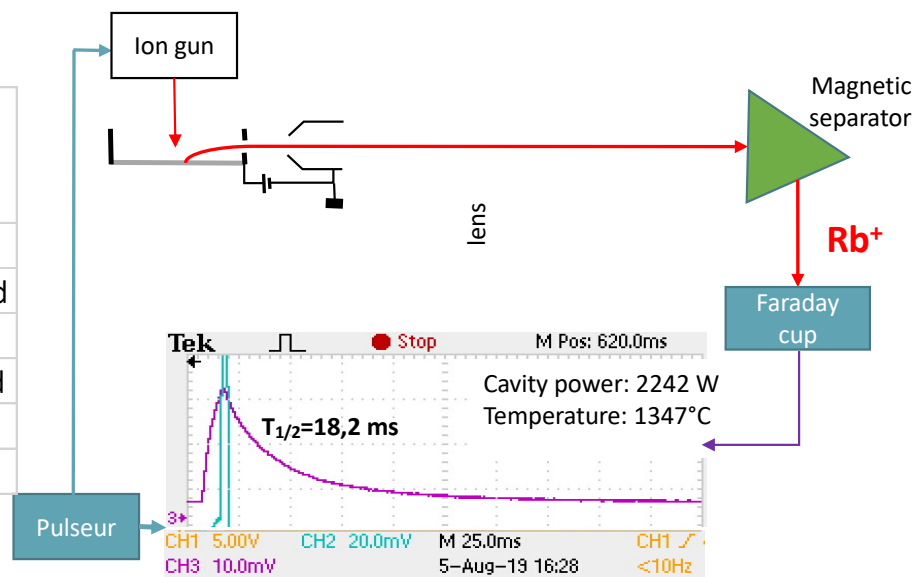
Life time observed at 3 kW > 3 days



* Thin layer deposition, <https://kerdry.com>

Effusion time expected (*Flight + sticking*)

Rb isotope	Temperature (°C)	Nbre of contacts	Sticking time/contact* (ms)	Time of flight (ms)	Total effusion time (ms)	
Without inner electric field, OFF-line						
85	1347	230	0,8	6,7	190,7	Calculated
With an inner electric field, OFF line						
85	1347	23	0,8	< 1	18,2	Measured
With an inner electric field, ON-line						
74	1500	23	0,2	< 1	4,6	Expected



*A. Pichard et al., Rev. Scient. Instrum. 81, 02A908 (2010)

Diffusion time expected

Production reaction: $^{20}\text{Ne@ 117 MeV}$ on Mo/Ni/Mo

Maximum range of ^{74}Rb residues in the carbon catcher: $5\text{ }\mu\text{m}$

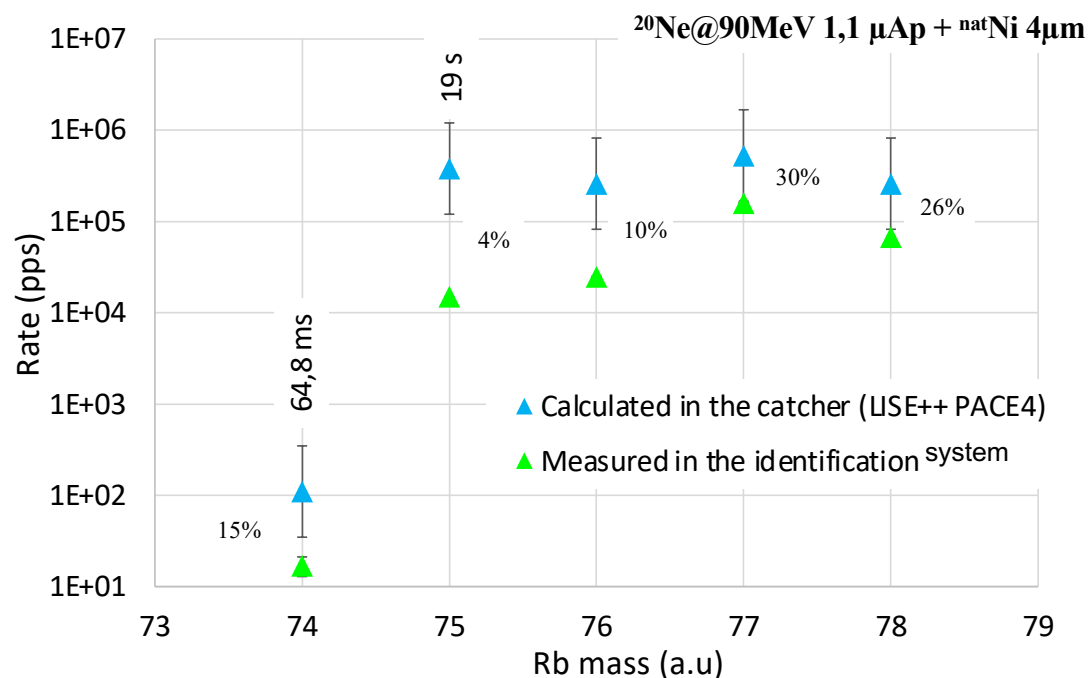
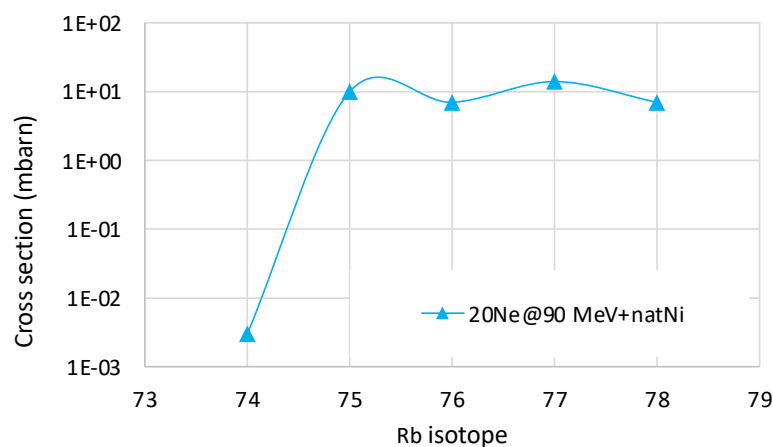
⇒ Release time out of the carbon catcher* at 1500°C : $\sim 2\text{ ms}$

Total atom-to-ion transformation time is expected to be of few milliseconds

* Extrapolated from dedicated measurements, J. Guillot et al., NIM B 526 (2022) 9-18

$^{74-78}\text{Rb}^+$ ion rates obtained in 2023

Reaction: ^{20}Ne @ 90 MeV on $_{\text{Mo}}/\text{natNi}/_{\text{Mo}}$
Temperature $\sim 1200^\circ\text{C}$ + $T_{\text{beam power}} (\sim 100^\circ\text{C})$

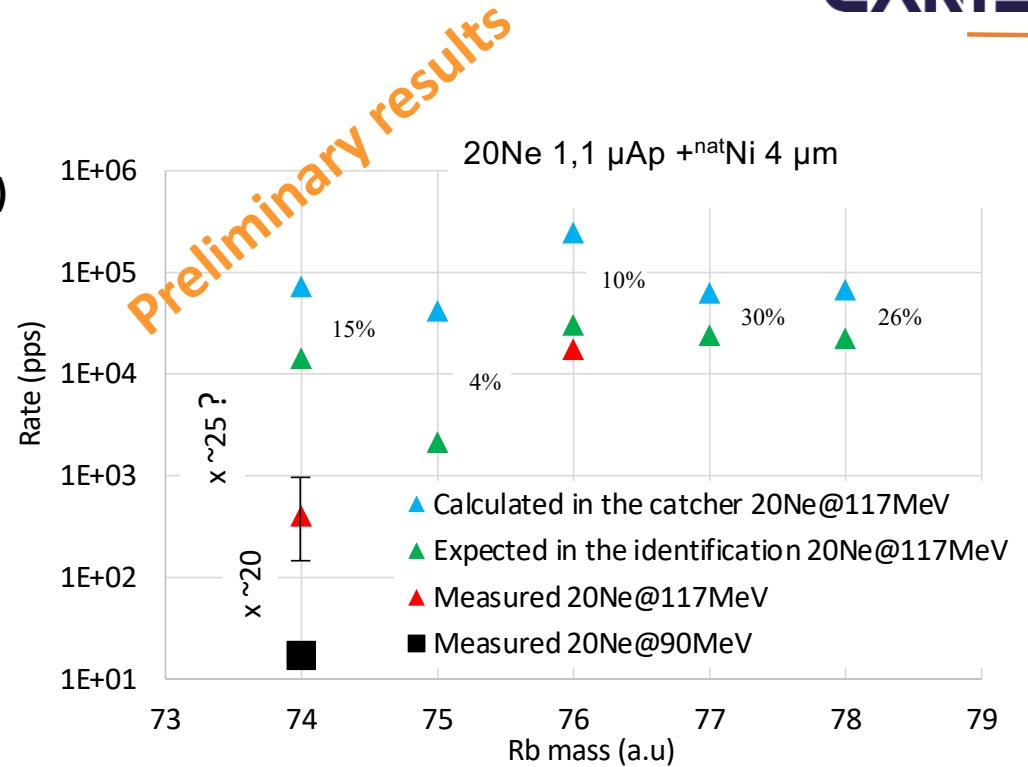
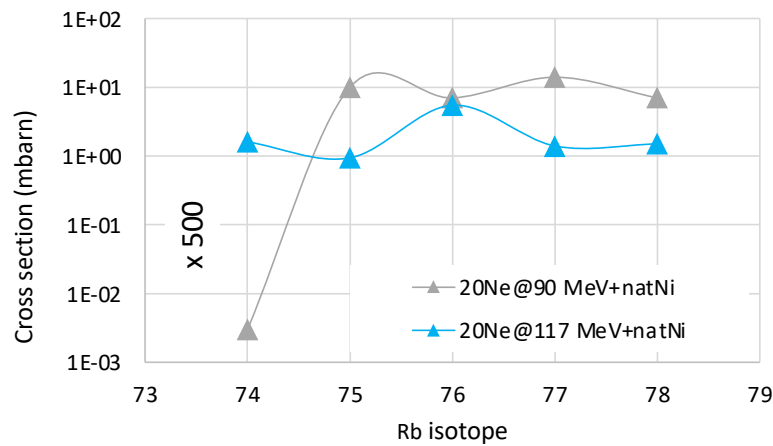


Consistent with the predictions

Implies an atom-to-ion transformation time shorter than the half-life of ^{74}Rb

2025: $^{74}\text{Rb}^+$ production

Optimized reaction: ^{20}Ne @ 117 MeV on $_{\text{Mo}}/\text{natNi}/_{\text{Mo}}$
Maximum heating ($T=1500^\circ\text{C}$) + beam power ($\sim 100\text{W}$)



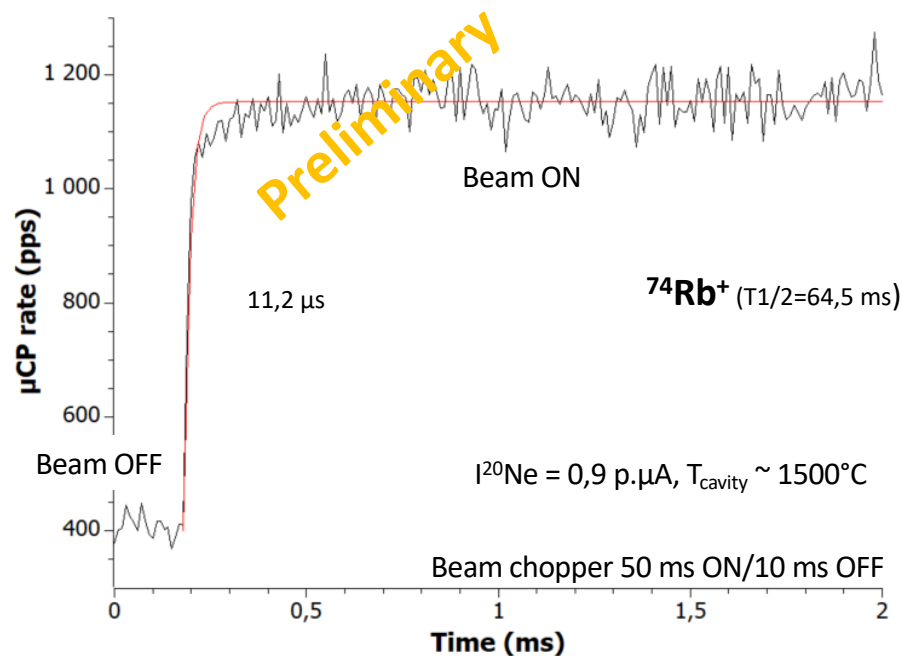
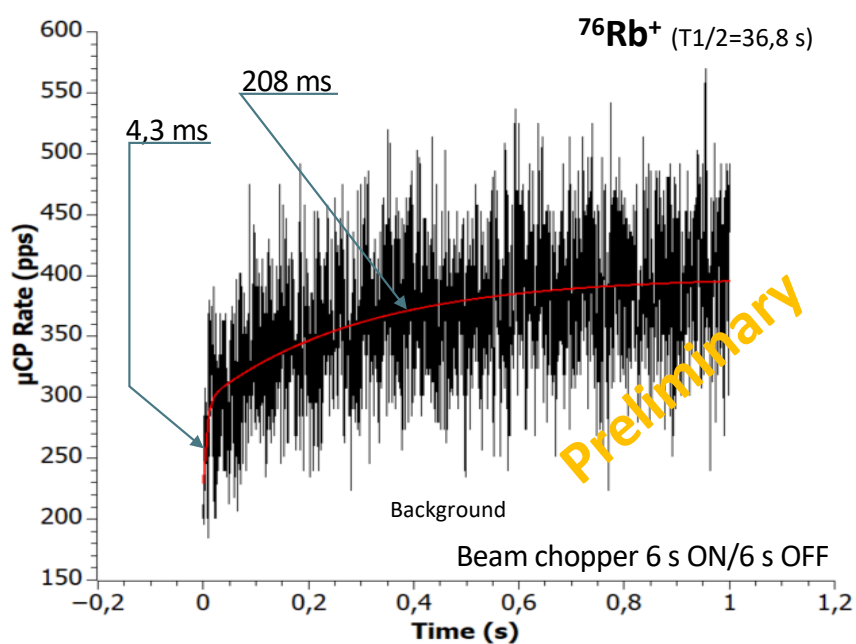
$^{74}\text{Rb}^+$ rate is not consistent with the prediction

Calculation: LISE++, PACE4

2025: $^{74}\text{Rb}^+$ atom-to-ion transformation time

20Ne @ 117 MeV on $\text{Mo}/^{nat}\text{Ni}/\text{Mo}$

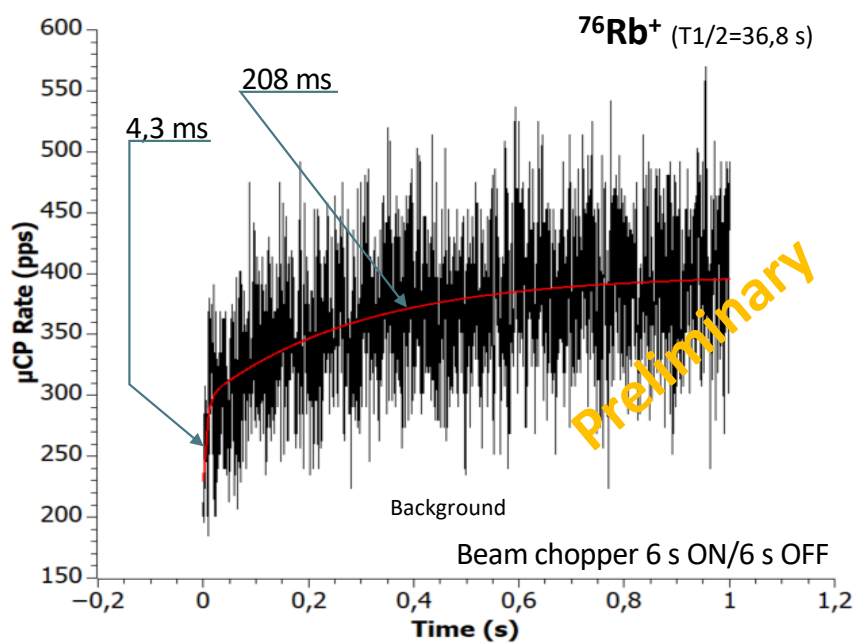
$I^{20}\text{Ne} = 0,2 \text{ p.}\mu\text{A}$, $T_{\text{cavity}} \sim 1300^\circ\text{C}$



2025: $^{74}\text{Rb}^+$ atom-to-ion transformation time

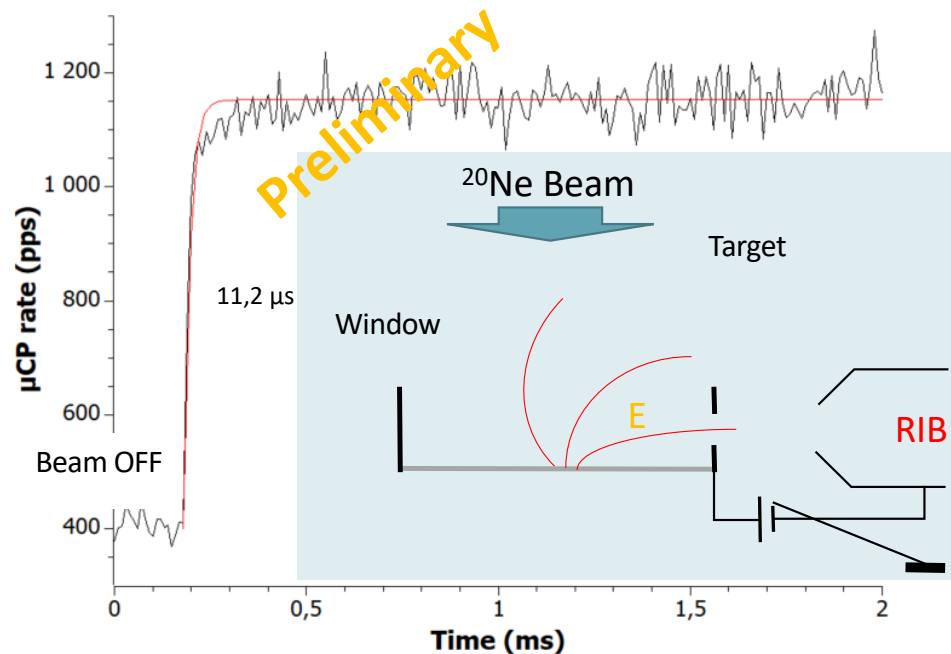
^{20}Ne @ 117 MeV on $\text{Mo}/^{nat}\text{Ni}/\text{Mo}$

$I^{20}\text{Ne} = 0,2 \text{ p.}\mu\text{A}$, $T_{\text{cavity}} \sim 1300^\circ\text{C}$



5 october 6:30

$I^{20}\text{Ne} = 0,9 \text{ p.}\mu\text{A}$, $T_{\text{cavity}} \sim 1500^\circ\text{C}$



7 october 2:00

Conclusion on $^{74-78}\text{Rb}^+$ production

The TISS works as expected (apart the window, which must be rendered more robust)

Long lived isotope ($^{75-78}\text{Rb}$) rates are convenient with the predictions

Few hundreds of $^{74}\text{Rb}^+$ produced within 11 μs

AIT time probably longer (~ 5 ms) with a no-broken window

$^{74}\text{Rb}^+$ probably higher

Towards the production of metallic elements around ^{100}Sn

Metallic elements $\Rightarrow$ first ionisation potential energy > 6 eV

$\rightarrow$ Association of the TULIP cavity (designed for alkali production) with a FEBIAD*

Thermo-mechanical tests at nominal currents

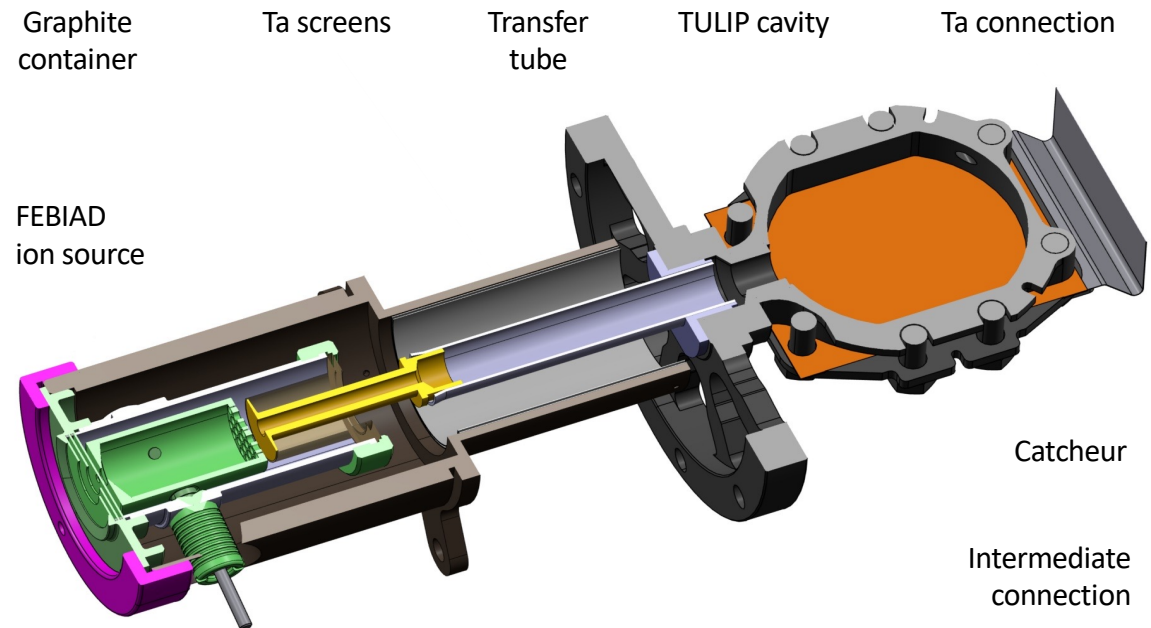
($I_{\text{cathode}} = 330$ A, $I_{\text{cavity}} = 500$ A, $T=1500^\circ\text{C}$)

No hint of aging after 4 days

Usual ^{40}Ar ionization efficiency: 20%

Expected for stable Sn, In, Cd, Ag: 10% to 40%

$\rightarrow$ Ready for online test



*V. Bosquet et al 2022 J. Phys.: Conf. Ser. 2244 012071

Expected production of ^{100}Sn

^{46}Ti at 203 MeV, 2.5×10^{12} pps on 4 μm thick $^{\text{nat}}\text{Ni}$ target

Production cross section of ^{100}Sn : 0.1 mbarn

Reduction factor of 10: 10 μbarn

In target production: ~ 600 pps

Max. range of ^{100}Sn in carbon catcher: 9 μm

Release efficiency of ^{120}Sn out of the catcher at 1500°C : $\sim 10\%$

Sn ionization efficiency assumed: few tens of %

PACE4

B. Blank et al., NIM B 416 (2018) 41-49

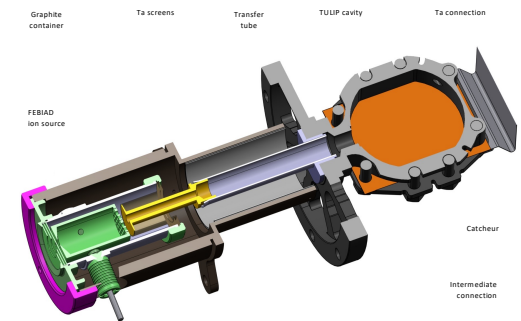
LISE++

A. Ribet et al. – Defect and Diffusion Forum – 439, 139-146

$\rightarrow ^{100}\text{Sn}^+$ rate at the exit of the TISS: few tens per second

Physics will decide

First test expected by Spring 2026

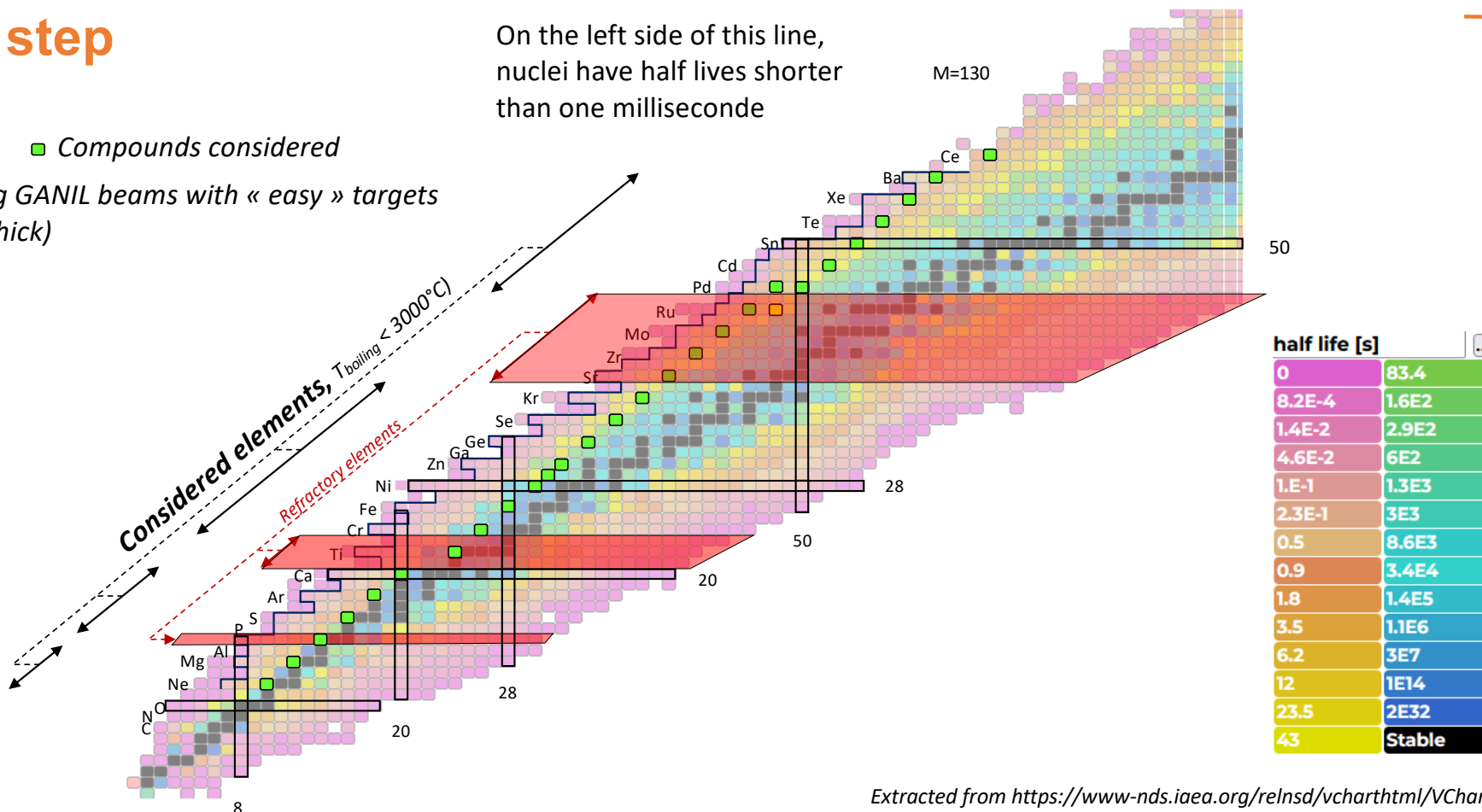


Next step

■ *Compounds considered*

Combining GANIL beams with « easy » targets
(few μm thick)

On the left side of this line,
nuclei have half lives shorter
than one millisecond



Extracted from <https://www-nds.iaea.org/relnsd/vcharthtml/VChartHTML.html>

**Thank you for
your attention**

For any question regarding the future beams, and for preparing the letters of intent, please contact the persons responsible for the development of the new beams via the following generic email address: chartbeams-spiral1@ganil.fr

