

Producción y uso de haces radioactivos

Master Interuniversitario de Física Nuclear

Berta Rubio IFIC (CSIC-Univ. Valencia)

B. Rubio. Master FN, Valencia 2024

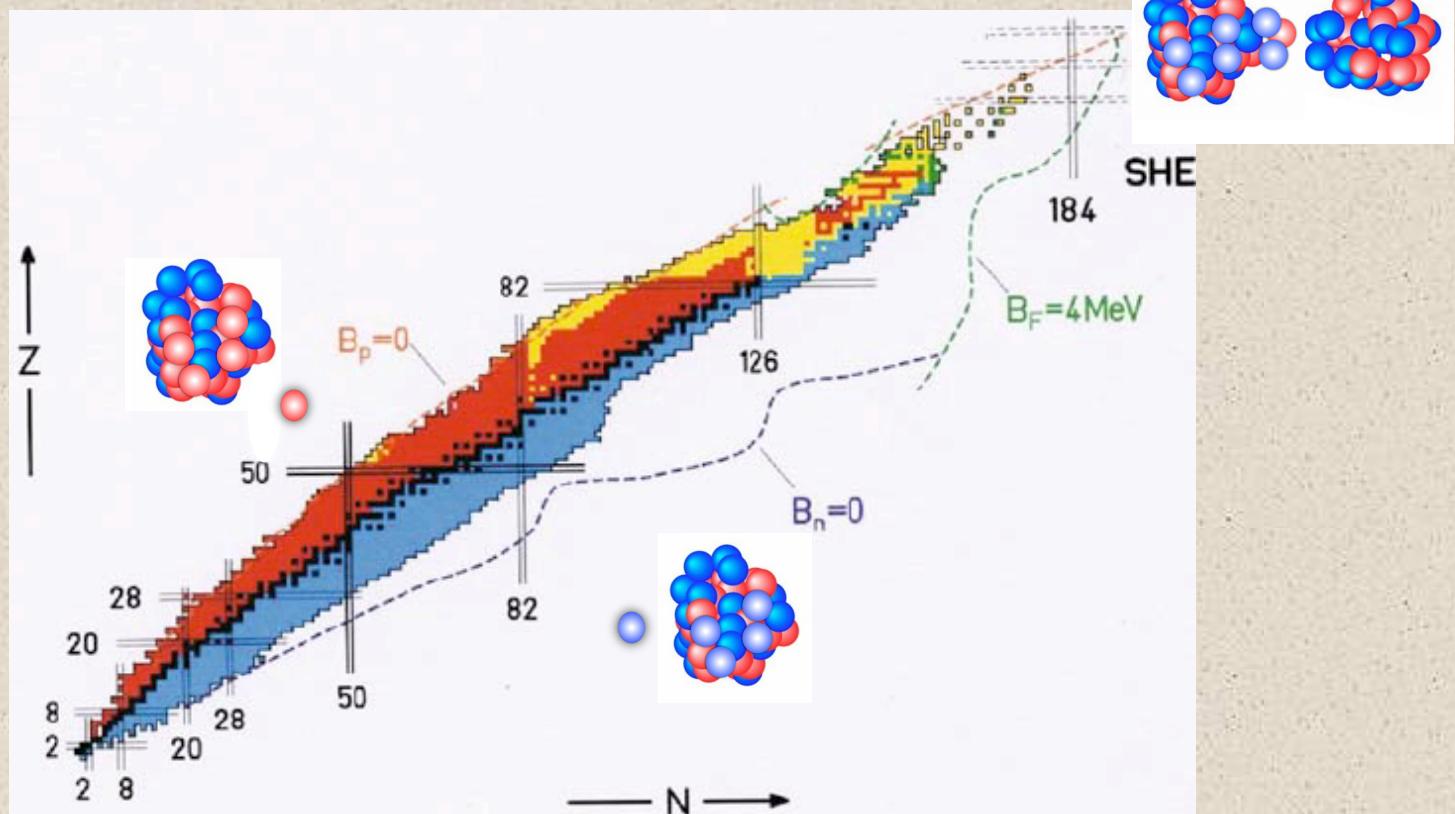
Porqué la producción de haces radioactivos ha marcado un antes y un después en la Física Nuclear

Reacciones nucleares y lo que nos enseñan de la estructura de los núcleos (M.J. García Borge, D. Cortina).

**Porqué necesitamos haces post acelerados
Porqué necesitamos alejarnos de la estabilidad
(J.L. Taín, A. Gadea, E. Nácher).**

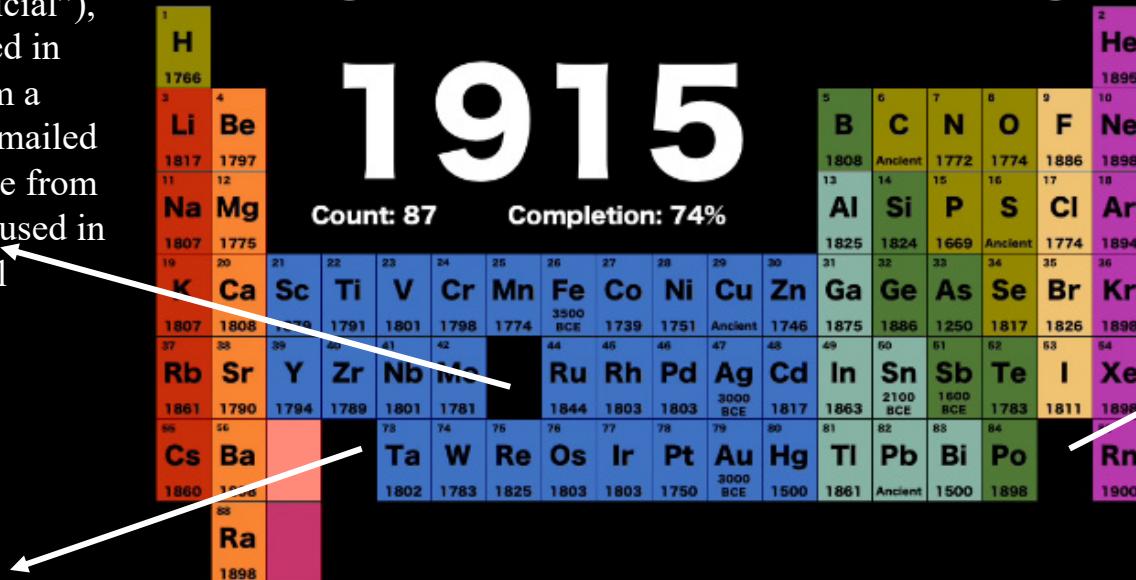
265 stable

About 3000 out
of 6000
synthesised in
our laboratories.



300 years of element discovery

Tachnetium (“artificial”), unstable, discovered in 1934 by Segre from a piece of cyclotron mailed by Ernest Lawrence from Berkely to Sicily. (used in millions of medical diagnosis)



Hafmium, relatively abundant in nature but very difficult to separate from Zn (X-ray spectrometry) (1923)

57	58	59	60
La	Ce	Pr	Nd
1839	1803	1885	1885

62	63	64	65	66	67	68	69	70	71
Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Rb
1879	1901	1880	1843	1886	1878	1843	1879	1878	1907

Alkali metals	Alkaline earth metals	Lanthanides	Actinides	Transition metals
Post-transition metals	Metalloids	Other non-metals	Halogens	Noble gases

Search for Super Heavy elements Z>100

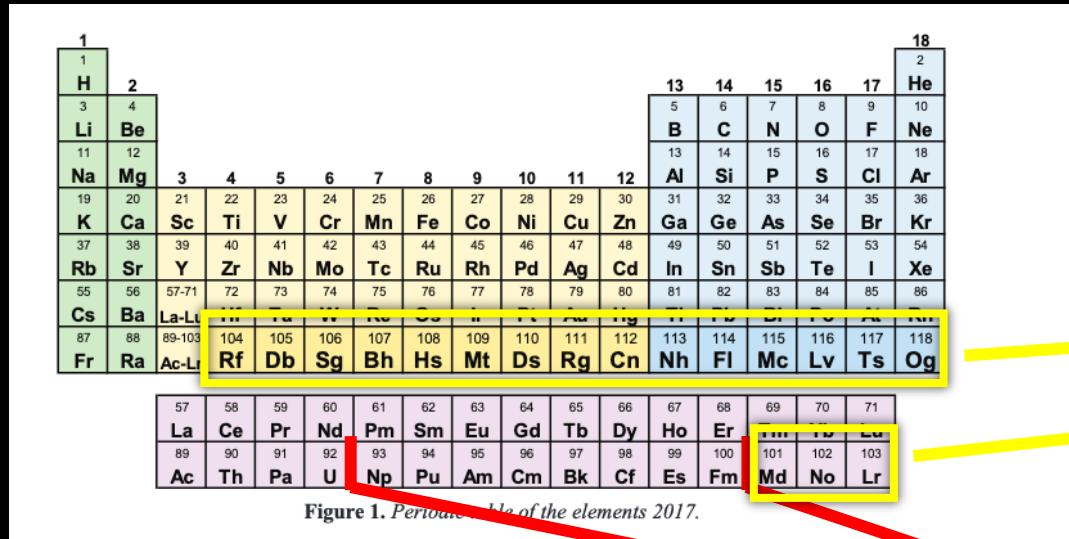


Figure 1. Periodic Table of the elements 2017.

Z	Name	Symbol	Year of Discovery	Discoverer
104	Rutherfordium	Rf	1969	Ghiorso et al. (LBNL Berkeley)
105	Dubnium	Db	1970	Ghiorso et al. (LBNL Berkeley)
106	Seaborgium	Sg	1974	Ghiorso et al. (LBNL Berkeley)
107	Bohrium	Bh	1981	Münzenberg et al. (GSI Darmstadt)
108	Hassium	Hs	1984	Münzenberg et al. (GSI Darmstadt)
109	Meitnerium	Mt	1982	Münzenberg et al. (GSI Darmstadt)
110	Darmstadium	Ds	1995	Hofmann et al. (GSI Darmstadt)
111	Roentgenium	Rg	1995	Hofmann et al. (GSI Darmstadt)
112	Copernicium	Cn	1996	Hofmann et al. (GSI Darmstadt)
113	Nihonium	Nh	2004	Morita et al. (RIKEN Wako-shi)
114	Flerovium	Fl	2004	Oganessian et al. (FLNR Dubna)
115	Moscovium	Mc	2010	Oganessian et al. (FLNR Dubna)
116	Livermorium	Lv	2004	Oganessian et al. (FLNR Dubna)
117	Tennessine	Ts	2010	Oganessian et al. (FLNR Dubna)
118	Oganesson	Og	2006	Oganessian et al. (FLNR Dubna)



Fm
discovered
in the 50ths



Meitnerium, the
only element
named after a
woman scientist

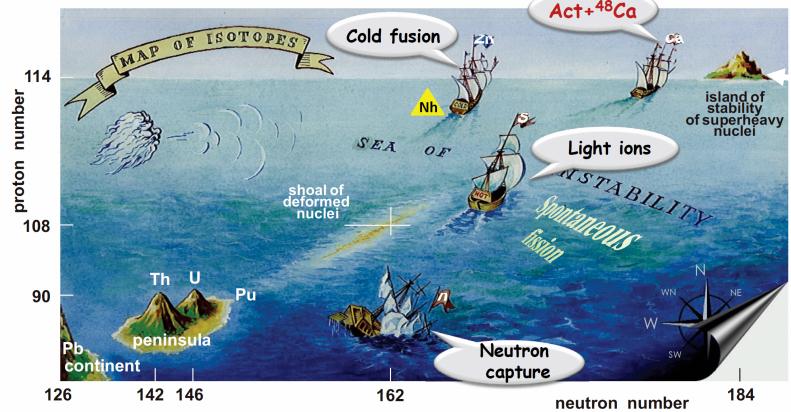


Figure 2. Voyage to the heaviest nuclei.

Will we prove the existence
of the island of stability?

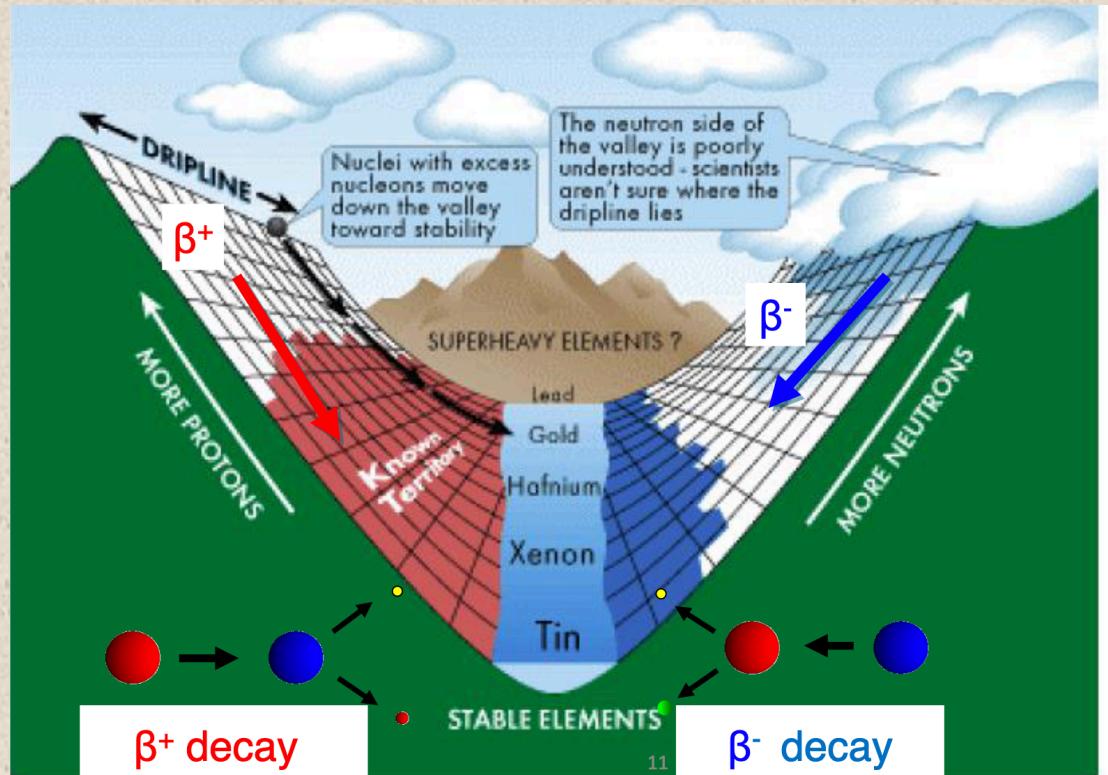
1	H	2																											
3	Li	4	Be																										
11	Mg	12																											
19	K	20	Ca																										
37	Rb	38	Sr																										
55	Cs	56	Ba																										
87	Fr	88	Ra																										
	Ac-Lr																												
57	La	58	Ce	59	Pr	60	Nd	61	Pm	62	Sm	63	Eu	64	Gd	65	Tb	66	Dy	67	Ho	68	Er	69	Tm	70	Yb	71	Lu
89	Ac	90	Th	91	Pa	92	U	93	Np	94	Pu	95	Am	96	Cm	97	Bk	98	Cf	99	Es	100	Fm	101	Md	102	No	103	Lr

Figure 1. Periodic table of the elements 2017.

Is Oganesson a nobel gas
or is neither nobel nor a gas

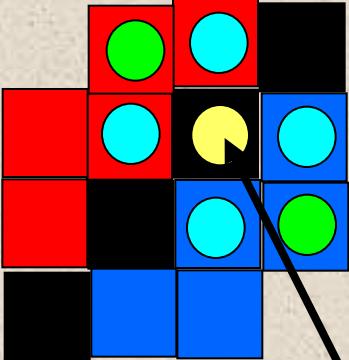
Does period number 8 exists?

Similarly to the chemical elements, the nuclei are better known and understood if they lie close to the stability

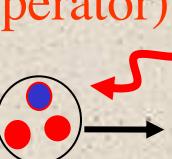


Let us see an example

What can we do if we have a stable target : we can perform nuclear reactions, for instance with light nuclei

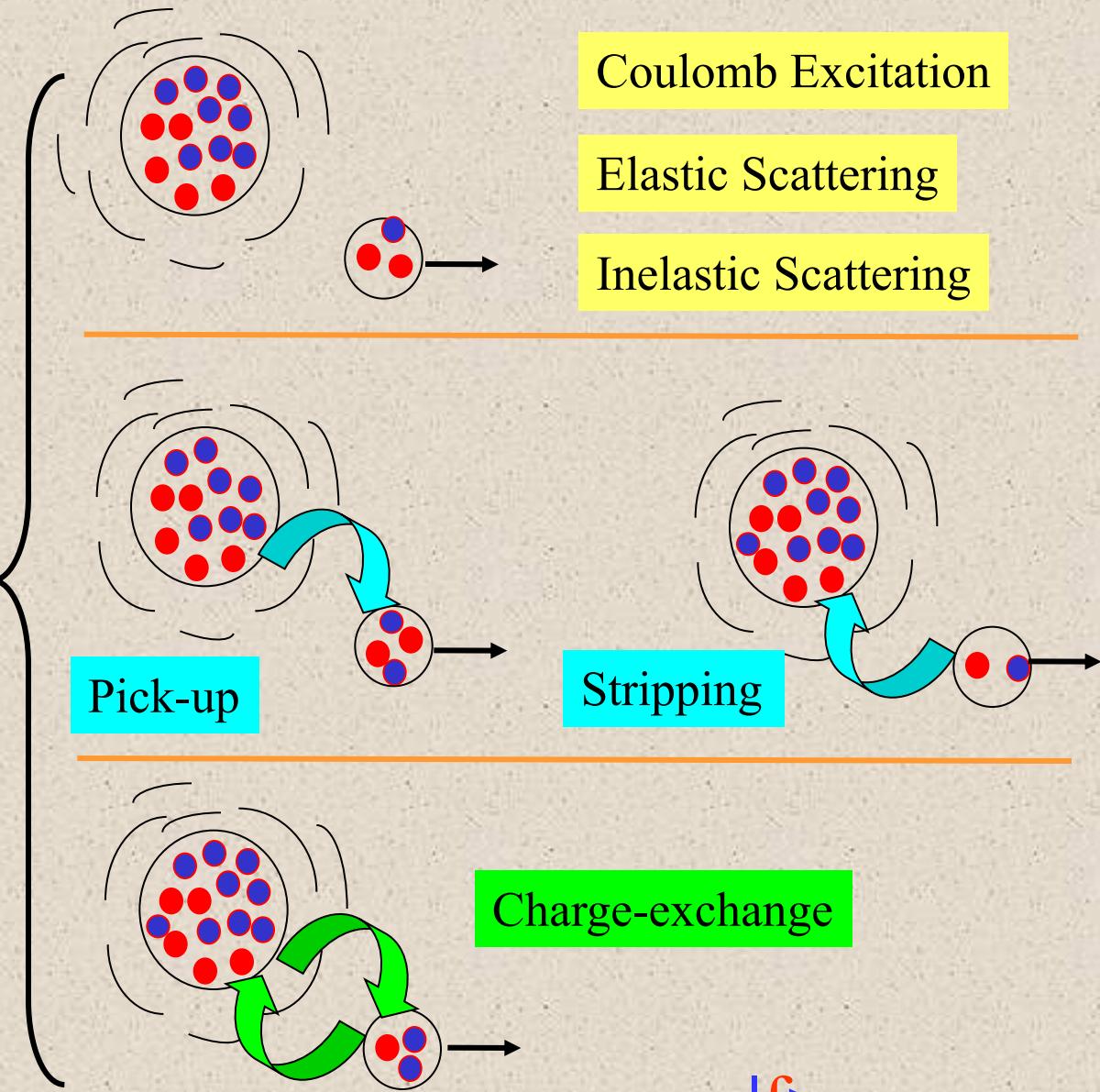


interaction (operator)

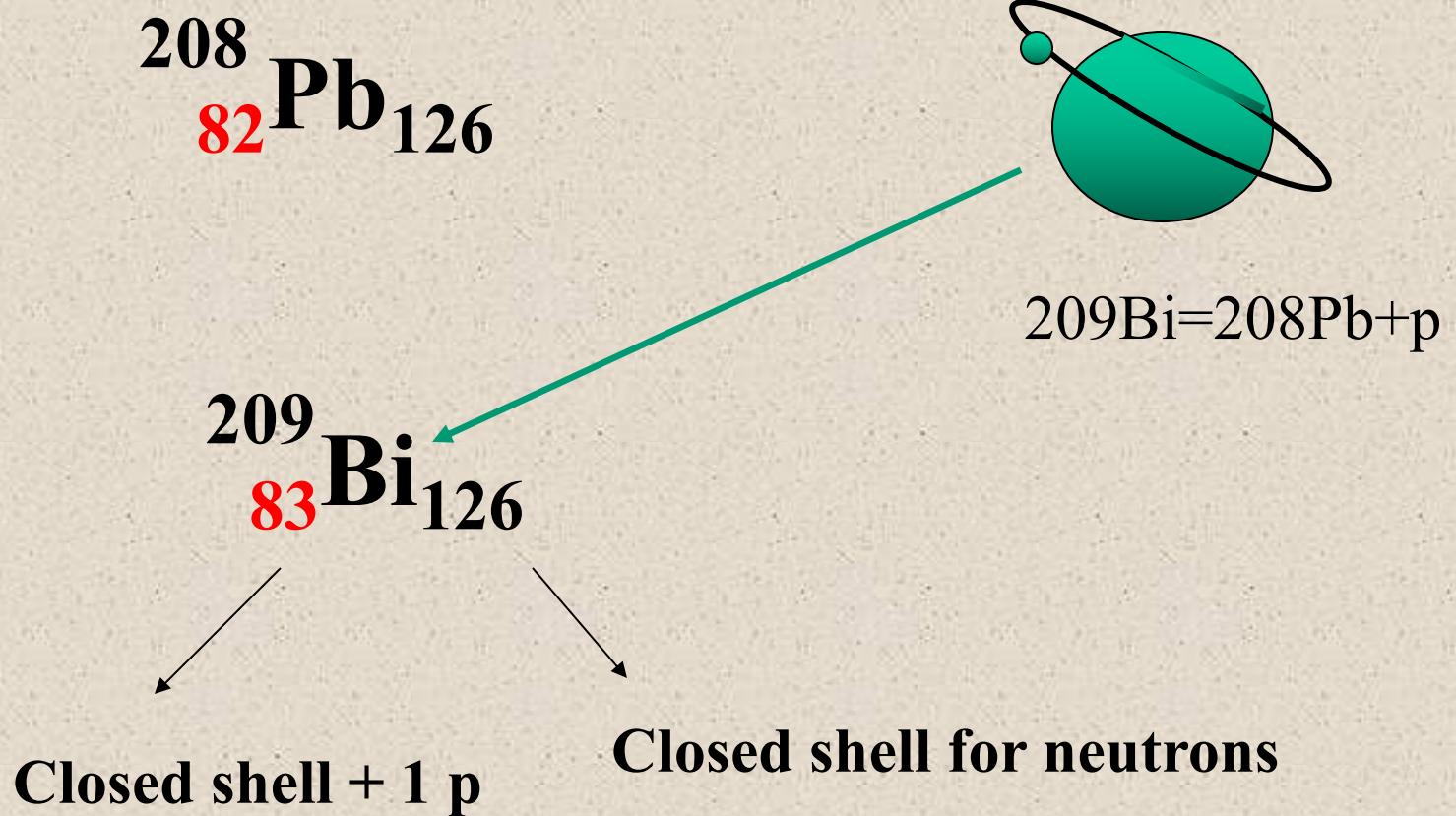


Target

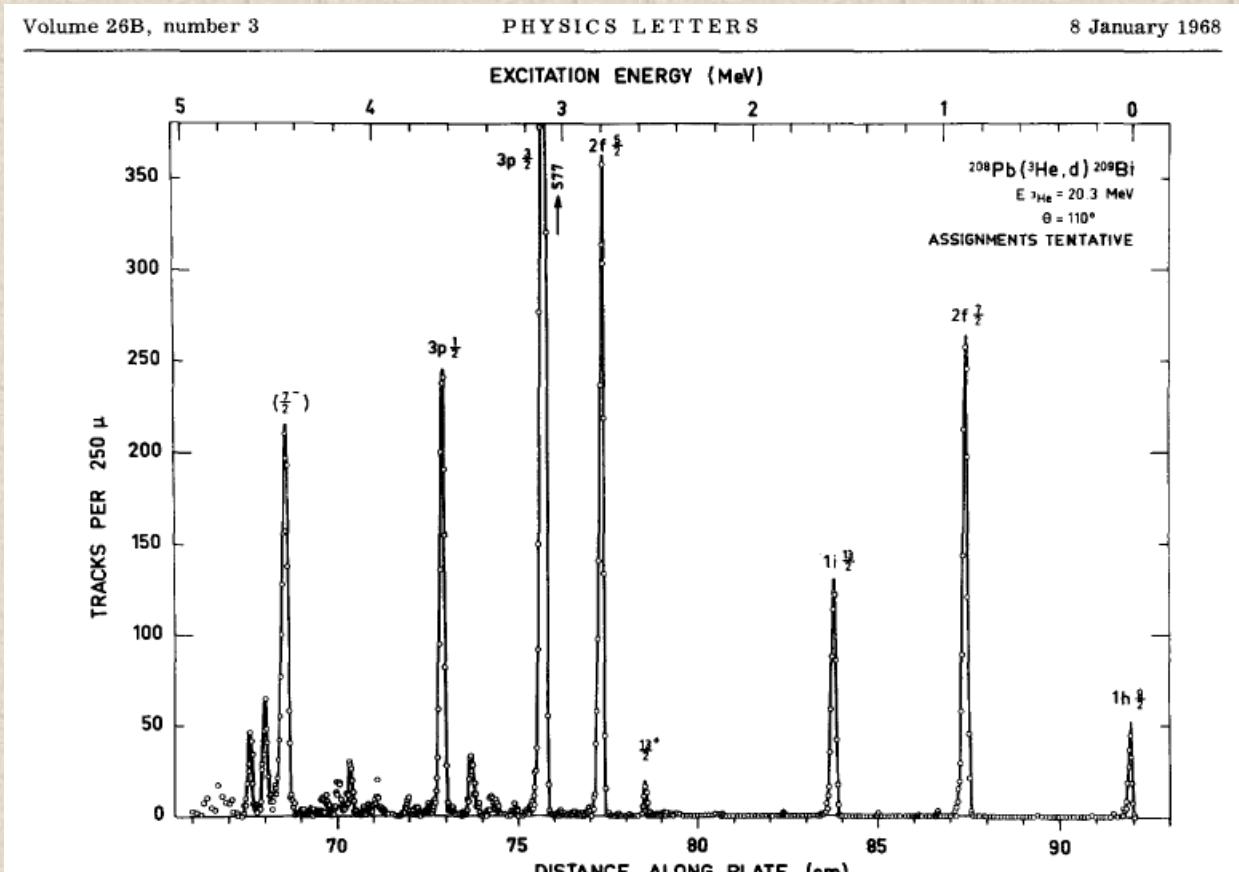
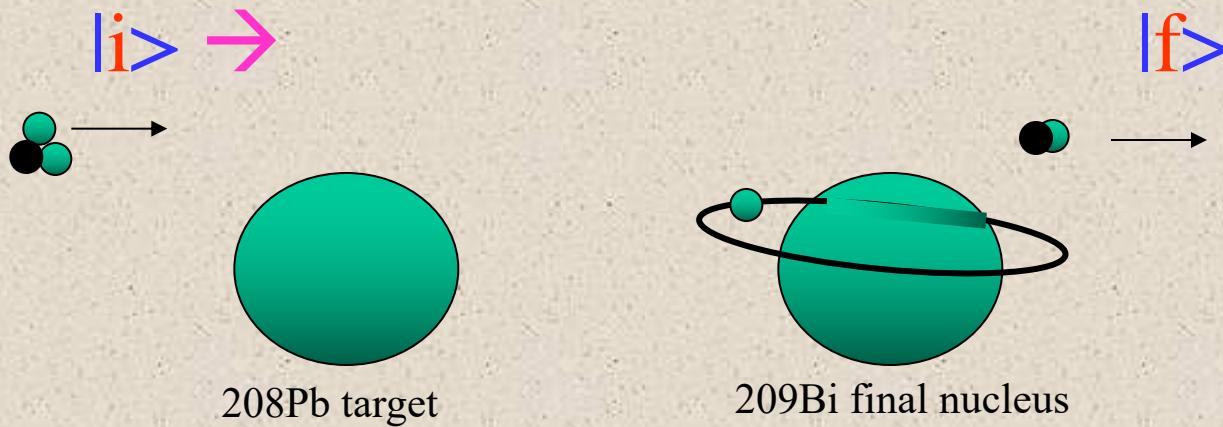
Projectile



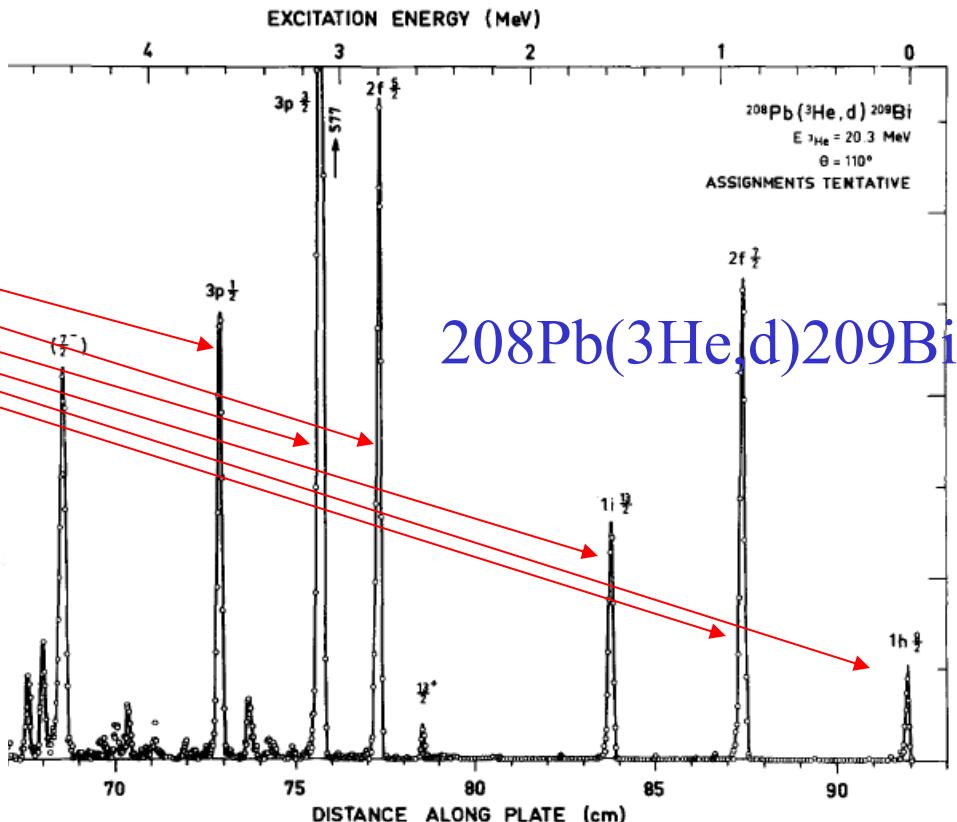
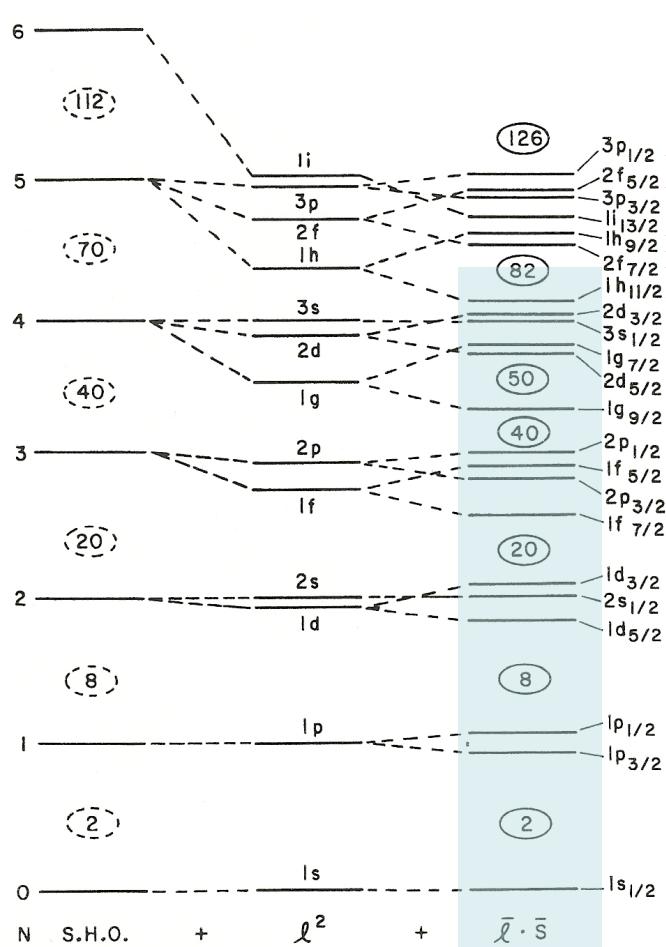
Let us use an illustrative example: ^{209}Bi



Reaction: $^{208}\text{Pb}(^3\text{He},\text{d})^{209}\text{Bi}$



Shell Model and Residual Interactions



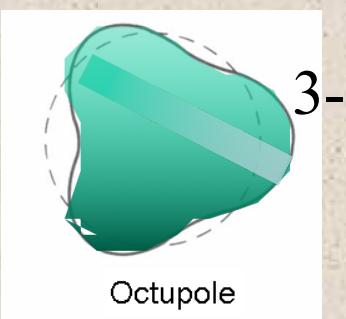
208Pb(3He,d)209Bi

Fig. 3.2. Single-particle energies for a simple harmonic oscillator (S.H.O.), a modified harmonic oscillator with l^2 term, and a realistic shell model potential with l^2 and spin orbit ($l \cdot s$) terms.

$209\text{Bi} = 208\text{Pb} + \text{p}$

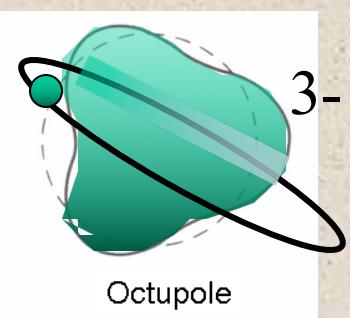
B. Rubio. Master FN, Valencia 2024

208Po



Octupole

$h_{9/2}$

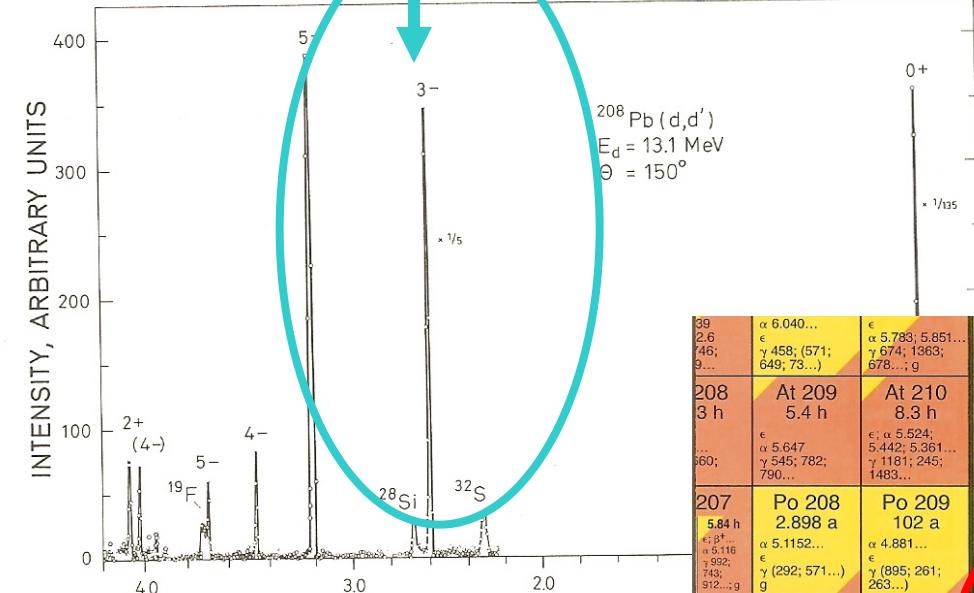


Octupole

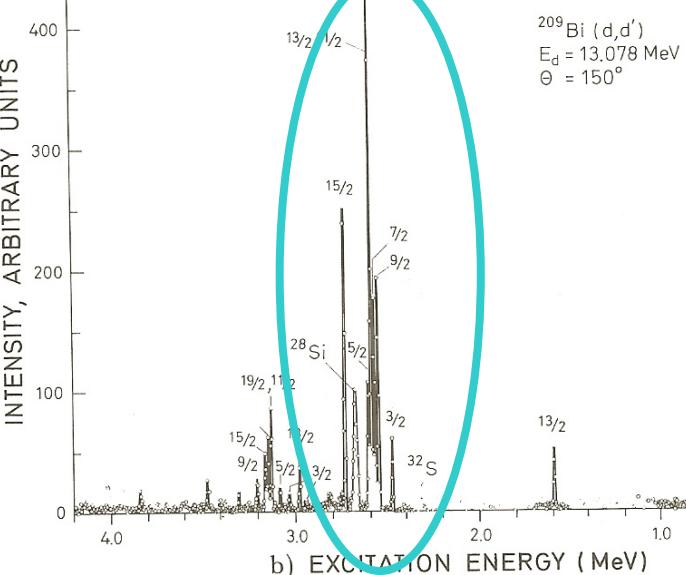
$209\text{Bi} = 208\text{Pb} + \text{p}$

$3^- \times h^{9/2}$

572 圖 VIBRATIONAL SPECTRA Ch. 6



$^{39}_{26}\text{Si}$	$\alpha 6.040...$	$\epsilon 5.783; 5.851...$	$\alpha 6.264...$	$\alpha 8.088; 7.252...$
$^{46}_{26}\text{Si}$	$\epsilon 4.458; (571; 649; 73...)$	$\gamma 674; 1363; 678...; g$	$\gamma 6.867...$	$\gamma 5.404...$
$^{9...}_{26}\text{Si}$			$\gamma (687...) g$	
$^{208}_{3}\text{h}$	$\text{At } 209$	$\text{At } 210$	$\text{At } 211$	$\text{At } 212$
	$5.4 h$	$8.3 h$	$7.22 h$	119 ms
	$\epsilon 5.647$	$\epsilon 5.524; 5.442; 5.361...$	$\epsilon 5.867...$	$\alpha 7.84; 7.90...$
	$\gamma 592; 740; 912...; g$	$\gamma 1181; 245; 1483...$	$\gamma (687...) g$	$\gamma 63...; \sigma$
207	$\text{Po } 208$	$\text{Po } 209$	$\text{Po } 210$	$\text{Po } 211$
	2.898 a	102 a	138.38 d	25.2 s
	$\epsilon 5.115...$	$\epsilon 4.881...$	$\alpha 5.0438...$	0.516 s
	$\beta^+...$	$\epsilon (292; 571...) g$	$\gamma (803); \sigma < 0.05$	$\alpha 11.6...$
	$\gamma 740; 912...; g$	$\gamma (895; 261; 263...) g$	$\gamma 570; 1064...$	$\gamma 26...$
206	$\text{Bi } 207$	$\text{Bi } 208$	$\text{Bi } 209$	$\text{Bi } 210$
4 d	31.55 a	$3.68 \times 10^5 \text{ a}$	100	0.10^6
	$\epsilon 5.116...$	$\epsilon 2615$	$\sigma 0.011 + 0.023$	5.013 d
	$\beta^+...$	$\epsilon 2615$	$\sigma_{\text{in}, \alpha} < 3E-7$	$\alpha 4.946...$
	$\gamma 570; 1064; 1770...$			$\gamma 4.649...$
205	$\text{Pb } 206$	$\text{Pb } 207$	$\text{Pb } 208$	$\text{Pb } 209$
10^7 a	24.1	22.1	52.4	3.253 h
	$\sigma 0.027$	$\sigma 0.61$	$\sigma 0.00023$	$\beta^- 0.6$
			$\sigma_{\text{in}, \alpha} < 8E-6$	$\alpha 0.00023$
204	$\text{Tl } 205$	$\text{Tl } 206$	$\text{Tl } 207$	$\text{Tl } 208$
8 a	70.48	3.7 m	4.20 m	3.053 m
	$\epsilon 0.11$	$\gamma 696; 453; 216; 256; 1021...$	$\beta^- 1.5...$	$\beta^- 1.8; 2.4...$
		$\gamma (803) g$	$\gamma 1000; 351$	$\gamma 2161; 583; 511; 860; 277...$
203	$\text{Hg } 204$	$\text{Hg } 205$	$\text{Hg } 206$	$\text{Hg } 207$
59 d	6.87	5.2 m	8.15 m	2.9 m
	$\sigma 0.4$	$\beta^- 1.5...$	$\beta^- 1.5...$	$\beta^- 1.8...$
		$\gamma 204...$	$\gamma 305; 650... g$	$\gamma 351; 997; 1637... m; g$
202	$\text{Au } 203$	$\text{Au } 204$	$\text{Au } 205$	$\text{Au } 205$
3 s	60 s	39.8 s	31 s	
	$\beta^- 2.0...$	$\beta^- 1.5...$	$\beta^- 1.5...$	
	$\gamma 218; 44; 51; 318; 369$	$\gamma 437; 1511; 692; 723; 1392...$	$\beta^- 379; 467; 946; 813...$	
201	$\text{Pt } 202$			
5 m	$\sim 43.6 \text{ h}$			
		β^-		



One could go to more sophisticated excitation modes such as An isovector dipole resonance (E1)

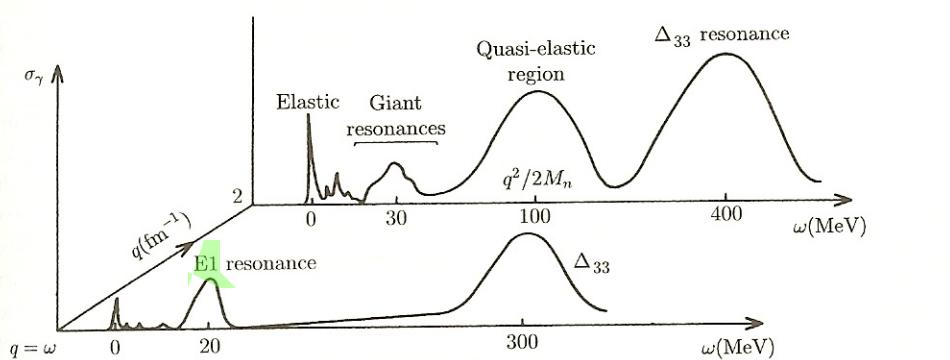
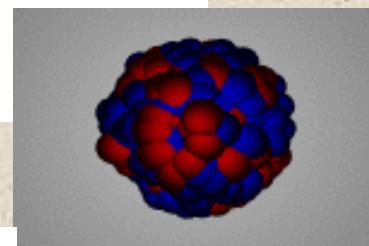
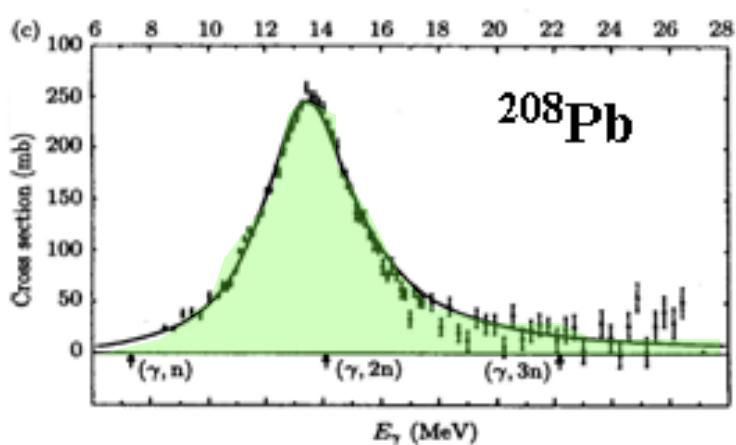
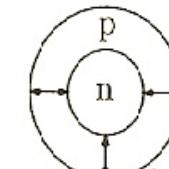


FIG. 1.1. Characteristic response (cross section σ) of an atomic nucleus as a function of the energy transfer ω and momentum transfer q . The lower curve is for photon absorption, the upper one for particle scattering with $q = 2 \text{ fm}^{-1} \neq \omega$.

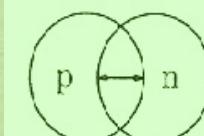


Isovector

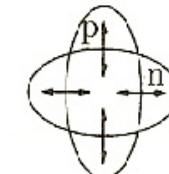
$\Delta T = 0$ 1



IVGMR



IVGDR



IVGQR

$\Delta T = 1$

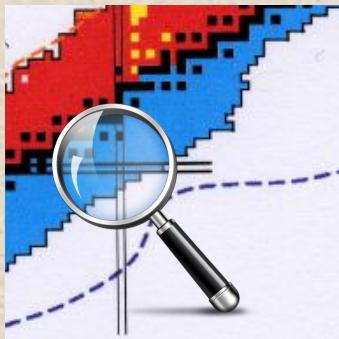
$\Delta S = 0$

E1

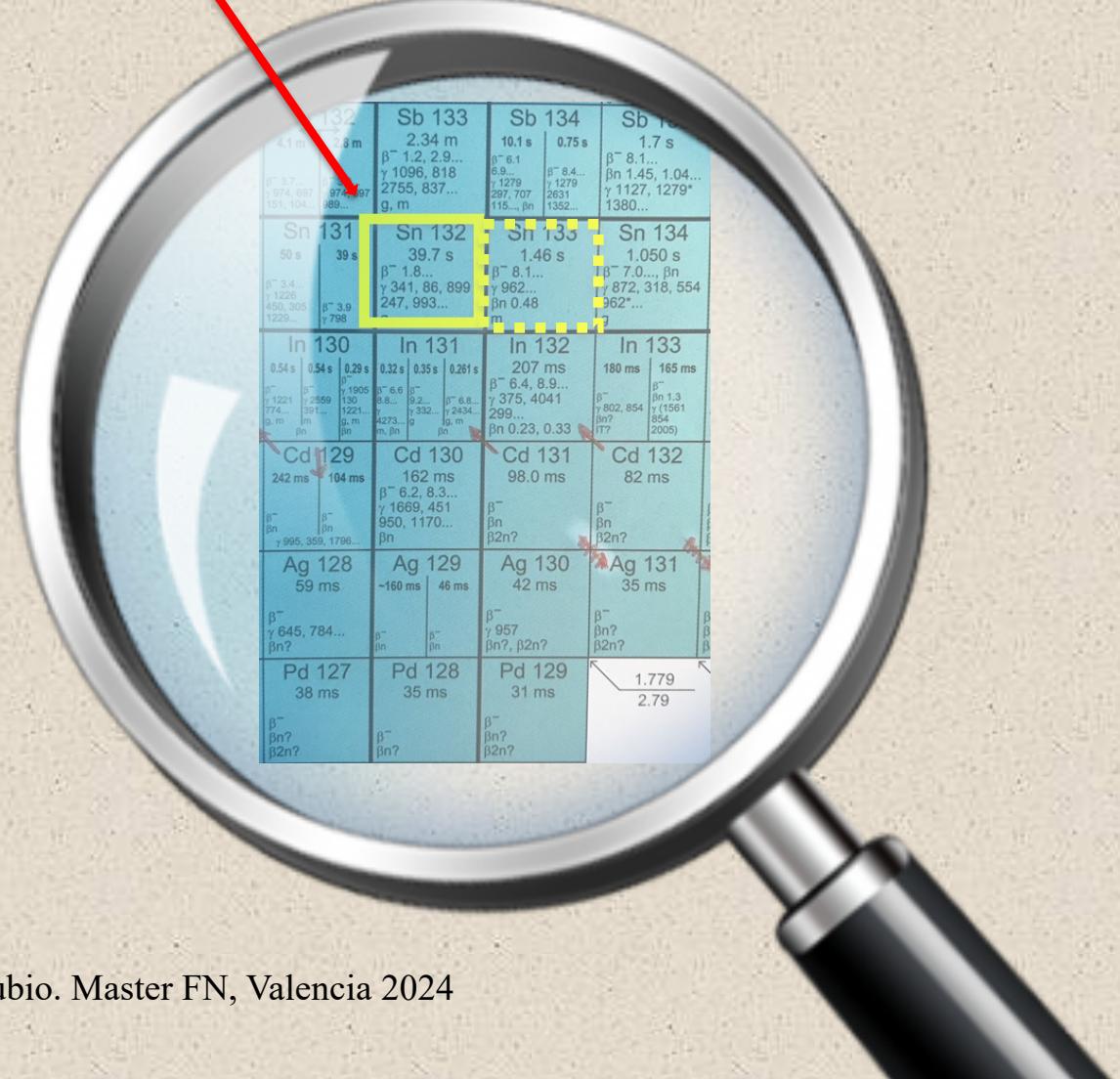
Qué ha sido necesario para poder hacer estos experimentos

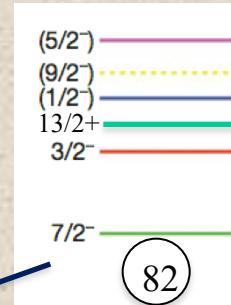
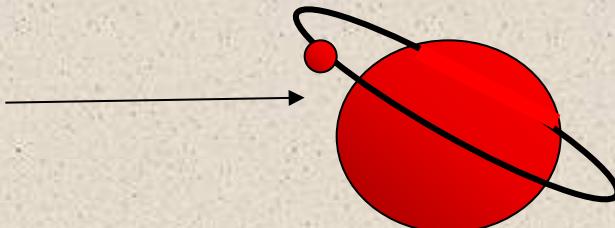
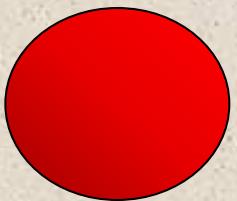
1. Definir la reacción nuclear de interés.
2. Blancos estables: ^{208}Pb y ^{209}Bi en este caso
2. (Un proyectil “sin estructura”)
3. (Un proyectil ligero (struggling))

Vamos a intentar aproximarnos a otro núcleo doblemente mágico:
 ^{132}Sn ($Z=50$, $N=82$)



^{132}Sn , unstable, $T_{1/2}=40\text{s}$





Sb 132	Sb 133	Sb 134	Sb 135
4.1 m β^- 3.7... γ 974, 697 151, 104...	2.34 m β^- 1.2, 2.9... γ 1096, 818 g, m	10.1 s β^- 6.1 8.5... 1.279 297, 707 115... (bn) 1352...	1.7 s β^- 8.1... βn 1.45, 1.04... γ 1127, 1279* 1380...
Sn 131	Sn 132	Sn 133	Sn 134
50 s β^- 3.4... 1226 450, 305 1229, 305 γ 798	39 s β^- 1.8... γ 341, 86, 899 247, 993...	1.46 s β^- 8.1... γ 962... βn 0.48	1.050 s β^- 7.0... (bn) 1872, 318, 554 362...
In 130	In 131	In 132	In 133
0.54 s β^- 1221... 774... g, m (bn)	0.54 s β^- 2559 391... g, m (bn)	0.29 s β^- 1905 0.35 s β^- 6.6... 0.261 s β^- 6.4, 8.9... 207 ms β^- 6.2, 8.3... γ 375, 4041 299... βn 0.23, 0.33	180 ms β^- 802, 854 165 ms β^- 8.1.3 γ (1561 177... 2005)
Cd 129	Cd 130	Cd 131	$^{133}\text{Sn}_{83-1}$
242 ms β^- 995, 359, 1796... βn	162 ms β^- 6.2, 8.3... γ 1669, 451 950, 1170... βn	98.0 ms β^- βn ? $\beta 2n$?	
Ag 128	Ag 129	Ag 130	
59 ms β^- 645, 784... βn ?	~160 ms β^- βn	46 ms β^- βn ? $\beta 2n$?	
Pd 127	Pd 128	Pd 129	
38 ms β^- βn ?	35 ms β^- βn ?	31 ms β^- βn ? $\beta 2n$?	

^{133}In β^- decay 1996Ho16, 2000Ho32

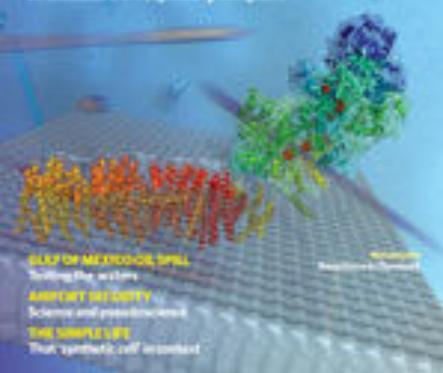
Type	Author	History	Citation
Full Evaluation	Yu. Khazov and A. Rodionov, F. G. Kondev		NDS 112, 855 (2011)

Parent: ^{133}In : E=0.0; $J^\pi=(9/2^+)$; $T_{1/2}=165$ ms; $Q(\beta^-)=12917$ SY; % β^- decay=100.0
 1995JoZZ, 1996Ho16, 2000Ho32: ^{133}In β^- [from ^{238}U (p,f), E=1 GeV]; measured $E\gamma$, $I\gamma$, $\gamma\gamma$, $\eta\gamma$, $\beta\gamma$ -c
 $(\beta n)\gamma(t)$. ISOLDE facility; plastic scin β detector, two liquid scin n detectors, two Ge γ -ray detectors

^{133}Sn Levels

$E(\text{level})^\dagger$	J^π^\ddagger
0.0	$7/2^-$
853.7 3	$3/2^-$
1560.9 5	$(9/2^-)$
2004.6 10	$(5/2^-)$

$1/2^-$ missing



nature

Vol 465 | 27 May 2010 | doi:10.1038/nature09048

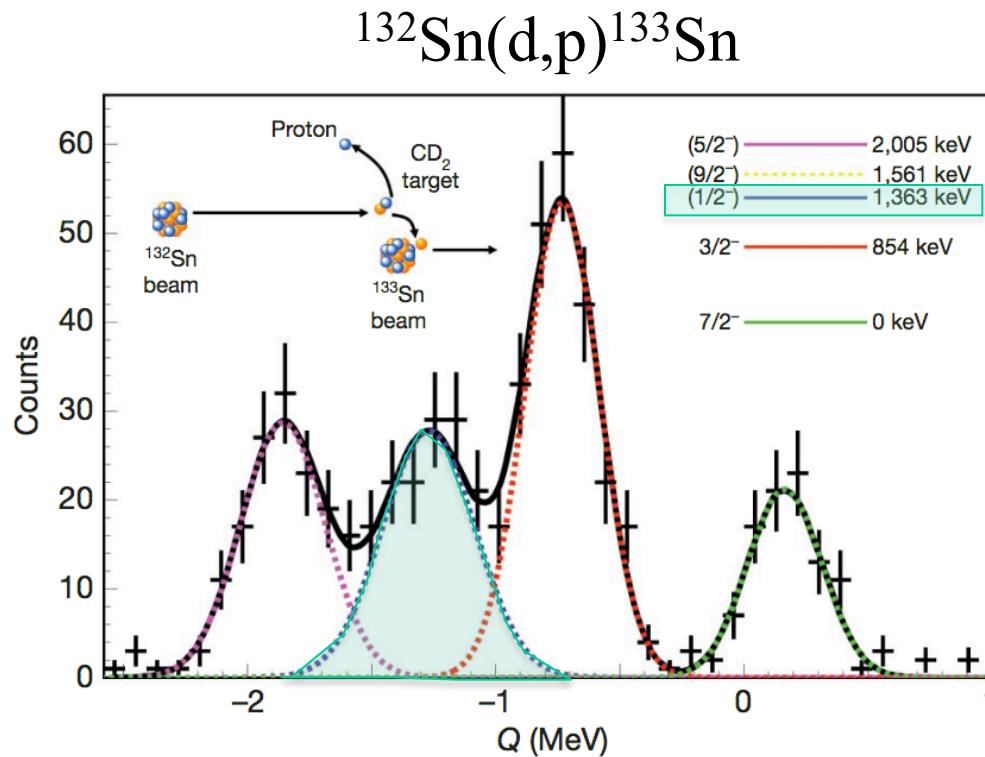
LETTERS

Year 2010

The magic nature of ^{132}Sn explored through the single-particle states of ^{133}Sn

K. L. Jones^{1,2}, A. S. Adekola³, D. W. Bardayan⁴, J. C. Blackmon⁴, K. Y. Chae¹, K. A. Chipps⁵, J. A. Cizewski², L. Erikson⁵, C. Harlin⁶, R. Hatarik², R. Kapler¹, R. L. Kozub², J. F. Liang⁴, R. Livesay², Z. Ma¹, B. H. Moazen¹, C. D. Nesaraja⁴, F. M. Nunes⁸, S. D. Pain², N. P. Patterson⁶, D. Shapira⁴, J. F. Shriner Jr⁷, M. S. Smith⁴, T. P. Swan^{2,6} & J. S. Thomas⁶

$(5/2^-)$	2,005 keV
$(9/2^-)$	1,561 keV
$(1/2^-)$	1,363 keV
$13/2^+$	
$3/2^-$	854 keV
$7/2^-$	0 keV



Conclusion of this part

It is important to have the possibility to perform reactions where either the target or the projectile are radioactive. But for most of the cases, a radioactive beam is more feasible than a radioactive target

We need to produce a beam of radioactive nuclei

The first thing to worry about is how to produce radioactive nuclei:
Tool: Reaction or Fission

The second thing to worry is to produce them in a “clean” way

For that it is important to separate them from other radioactive products

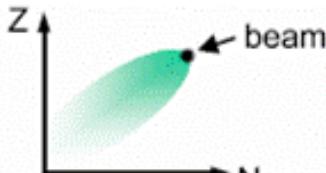
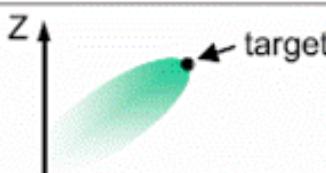
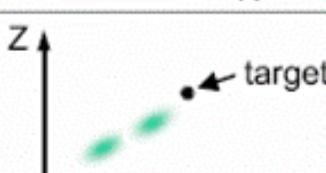
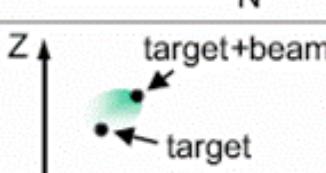
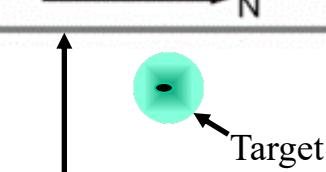
Postacceleration

There are two main ways: the Isol method and the fragmentation method

The first one is known since many years
(Isolde-CERN, since 1967)

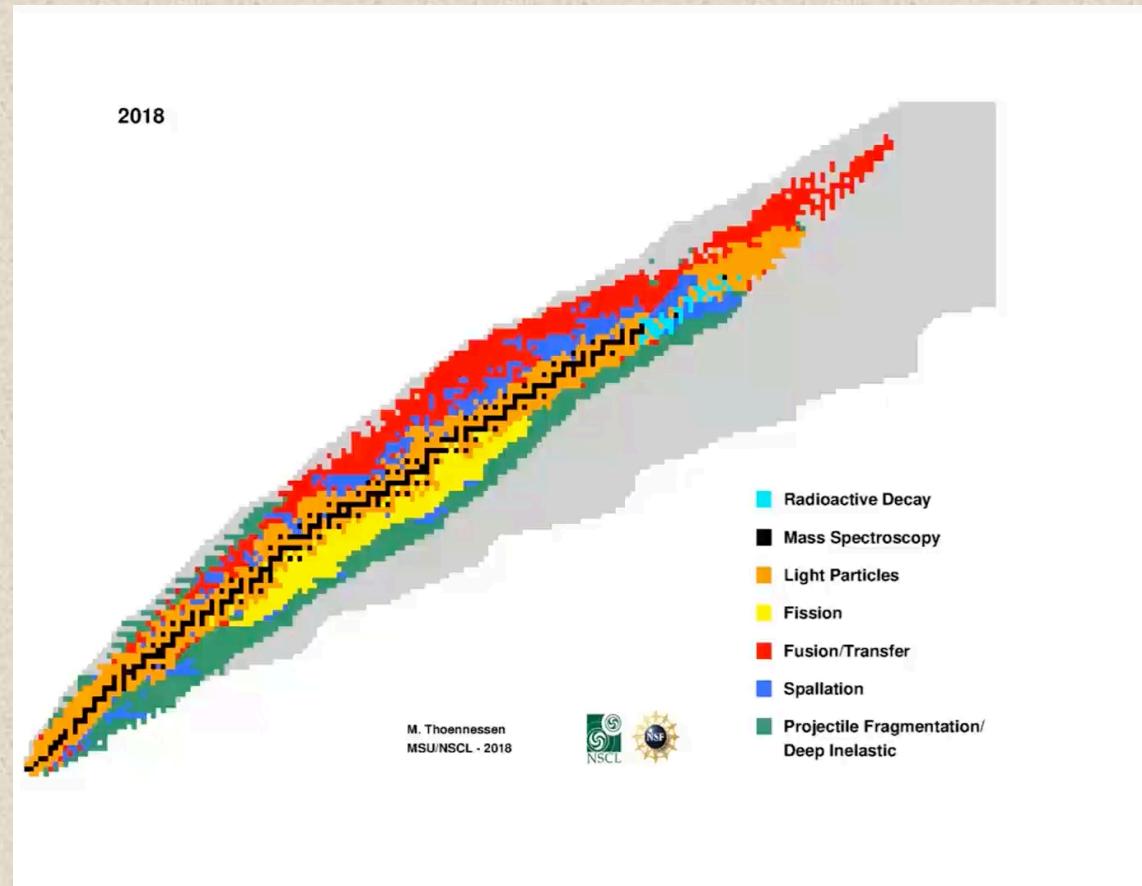
The first thing to worry about is how to produce radioactive nuclei.
Tool: Reaction or Fission

The reaction to produce Radioactive beams starting with stable beam and stable target

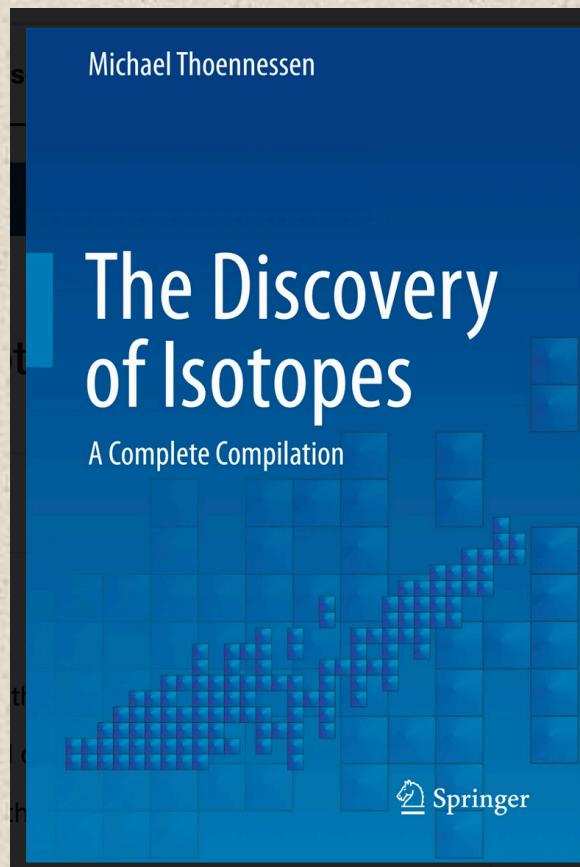
Beam + target \rightarrow products	Product energy	Number of Products	
GeV HI	“All nuclei” fragmentation RIKEN, GSI, MSU, FAIR, FRIB	$v_{\text{product}} = v_{\text{beam}}$ up to 1000	
GeV p	“All nuclei” spallation ISOLDE	few MeV/u up to 1000	
small p/n	“mainly neutron rich” fission Reactors, ISOLDE...	$\sim 1 \text{ MeV/u}$ few 100	
5 MeV/u HI	“mainly proton rich rich” fusion-evaporation “SPIRAL2”	$E_R = \frac{m_p}{m_p + m_t} E_p$ few (≤ 20)	
20 MeV/u	“close to the stability” Transfer reactions “Small facilities”	5-100 MeV/u Very few: 2-5	

Historical view to the discovery of new isotopes

<https://people.nscl.msu.edu/~thoennes/isotopes/2018-Isotope-Movie.mp4>

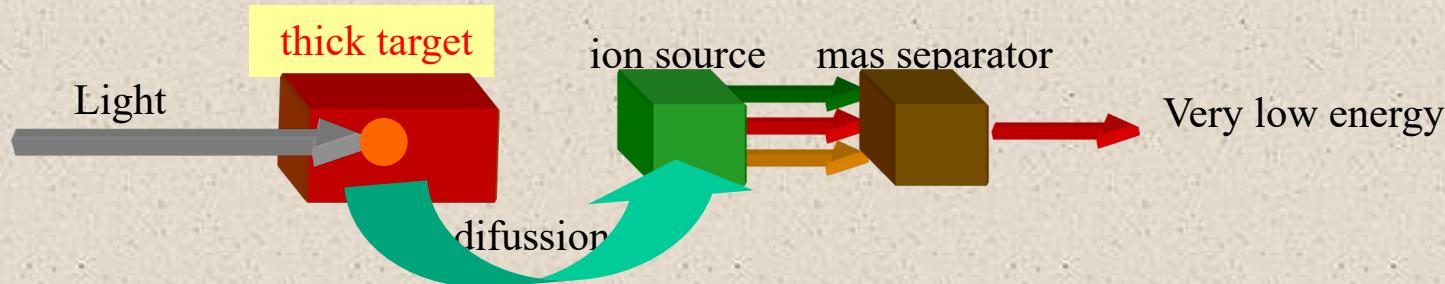


If you want to learn how all that happened, read this book

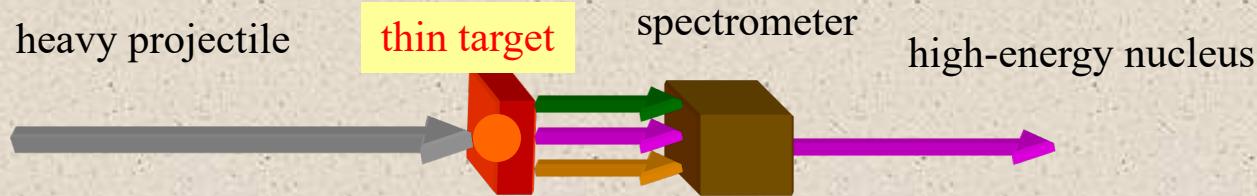


The second thing is to separate them from other radioactive products:
Radioactive nuclei production techniques

Isotopic Separation On-Line (ISOL)

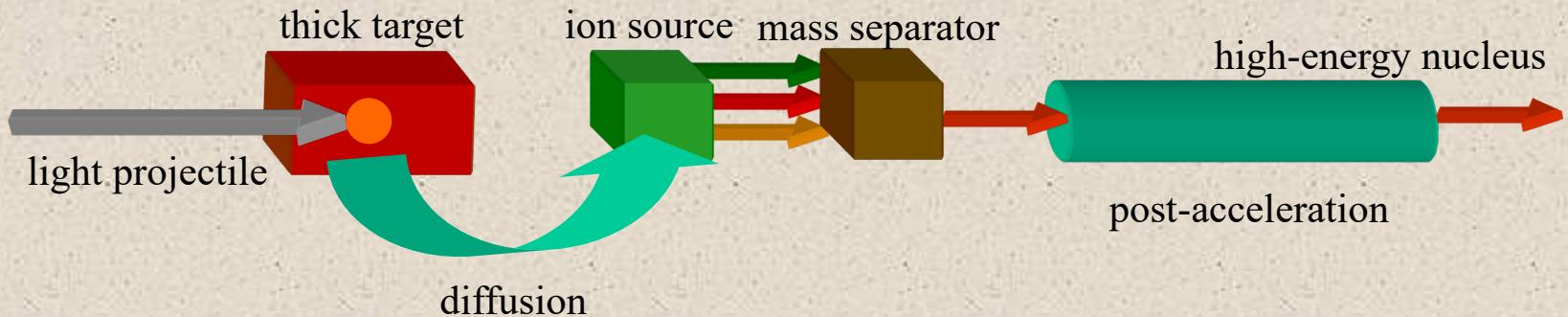


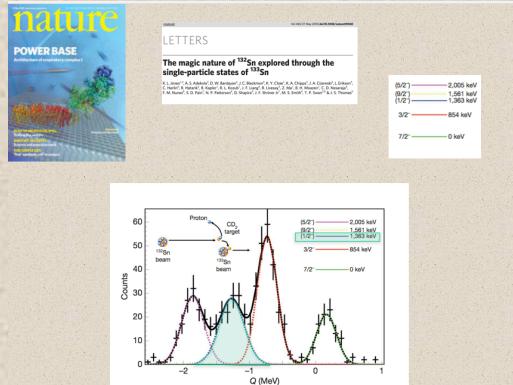
In-flight fragmentation



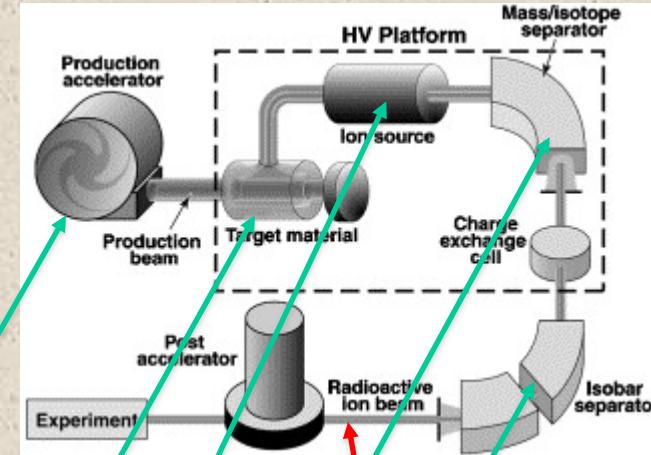
Production techniques Rad Beams: post acceleration

➤ Isotopic separation on-line (ISOL)





Radioactive beam
(40 sec):
inverse kinematics



1. Protons accelerated in the cyclotron
2. Uranium Carbide power target
3. About 100 species produced, some of them ionised
4. 132 Mass separated
5. 132Sn separated
6. 132Sn accelerated to 4.8 MeV x A
7. Experiment

On 16 October 1967, the first experiments were carried out at the Isotopic Separator On Line ISOLDE CERN.

CERN's longest serving experimental facility

Physics Reports 403–404 (2004) 459–469

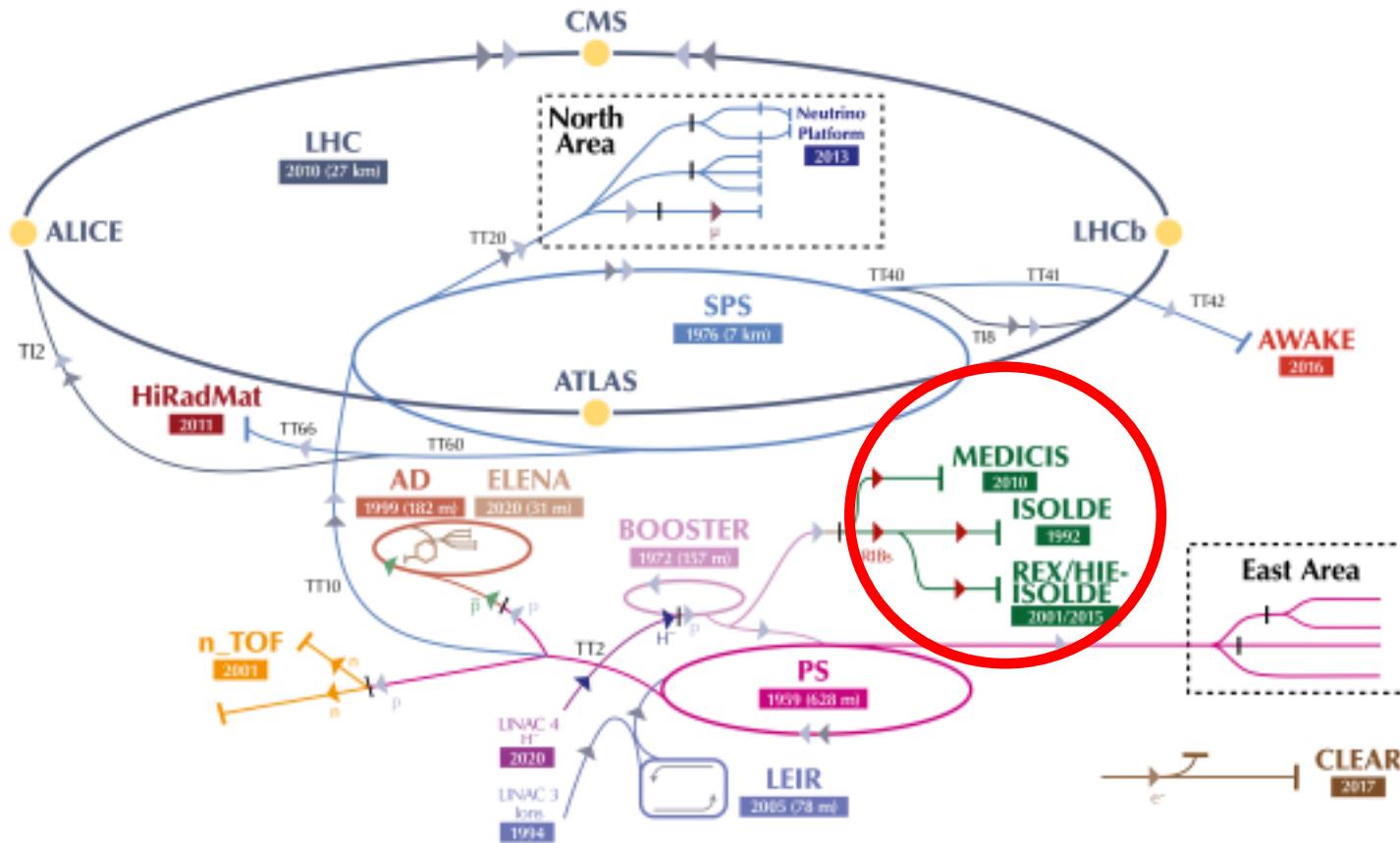
Juha Äystö

B. Rubio. Master FN



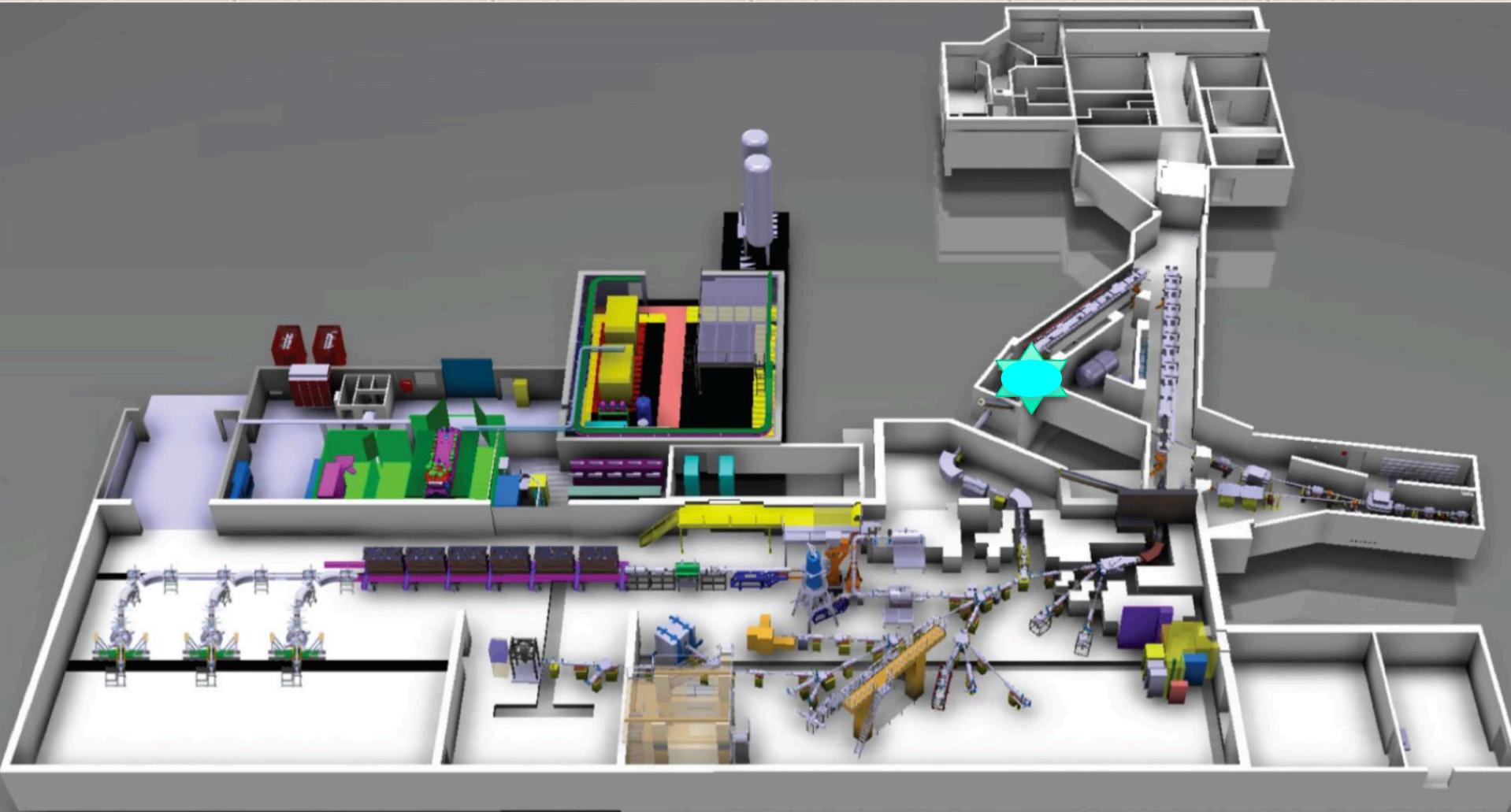
The CERN accelerator complex

Complexe des accélérateurs du CERN

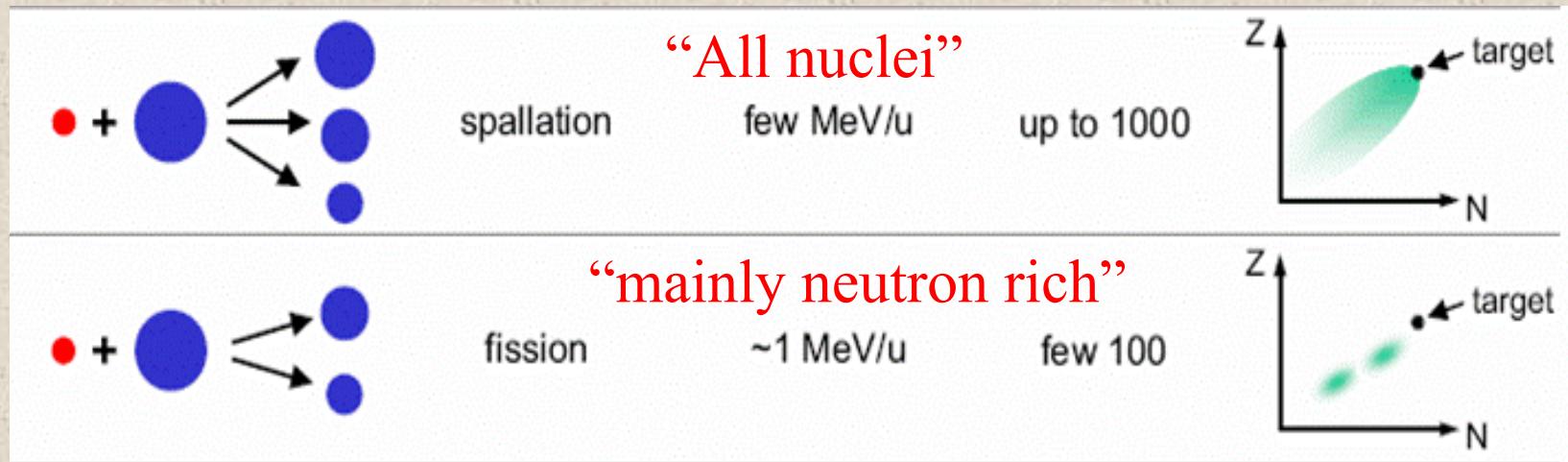


► H^- (hydrogen anions) ► p (protons) ► ions ► RIBs (Radioactive Ion Beams) ► n (neutrons) ► \bar{p} (antiprotons) ► e^- (electrons) ► μ (muons)

LHC - Large Hadron Collider // SPS - Super Proton Synchrotron // PS - Proton Synchrotron // AD - Antiproton Decelerator // CLEAR - CERN Linear Electron Accelerator for Research // AWAKE - Advanced WAKEfield Experiment // ISOLDE - Isotope Separator OnLine // REX/HIE-ISOLDE - Radioactive Experiment/High Intensity and Energy ISOLDE // MEDICIS // LEIR - Low Energy Ion Ring // LINAC - LINear ACcelerator // n_TOF - Neutrons Time Of Flight // HiRadMat - High-Radiation to Materials // Neutrino Platform



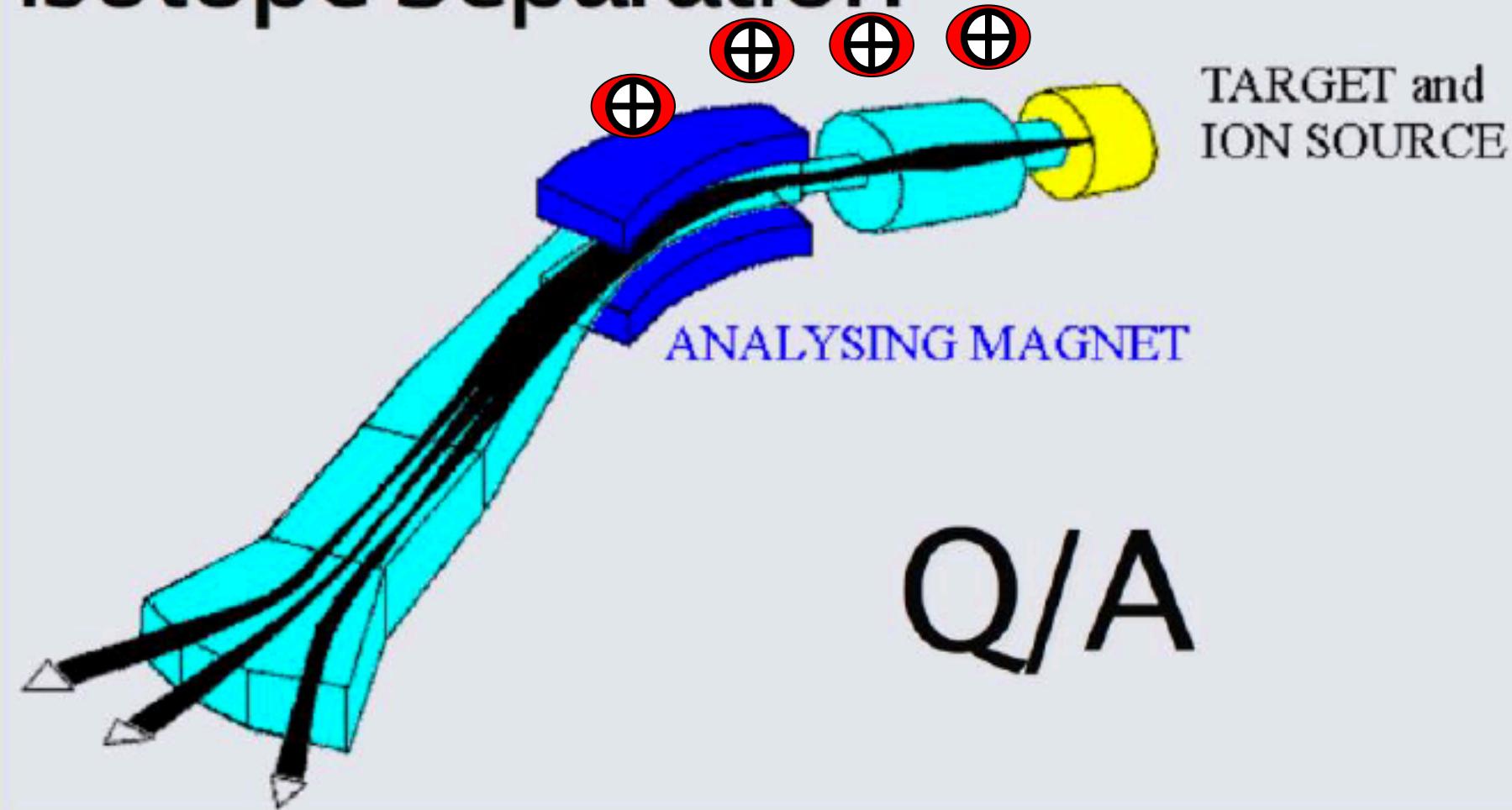
B. Rubio. Master FN, Valencia 2024



- 😊 High intensity p beam (upto 10^{16} s^{-1}) the same beam that serves LHC)
- 😊 thick target (100% energy range of p)
- 🙁 long extraction and ionization time (hundred's ms)
- 🙁 Chemistry dependent

The Separator: in general, only isobaric separation

Isotope Separation

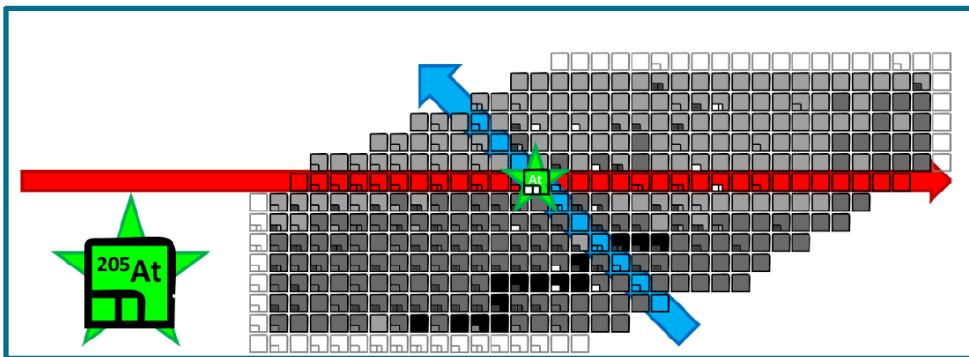
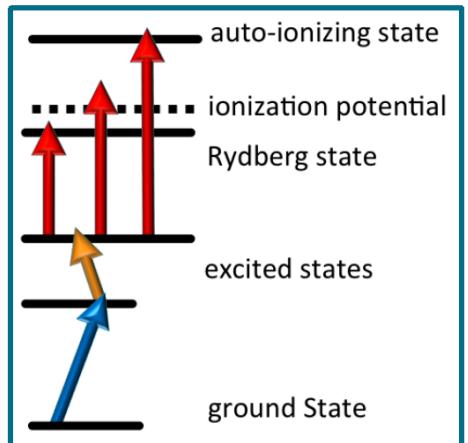
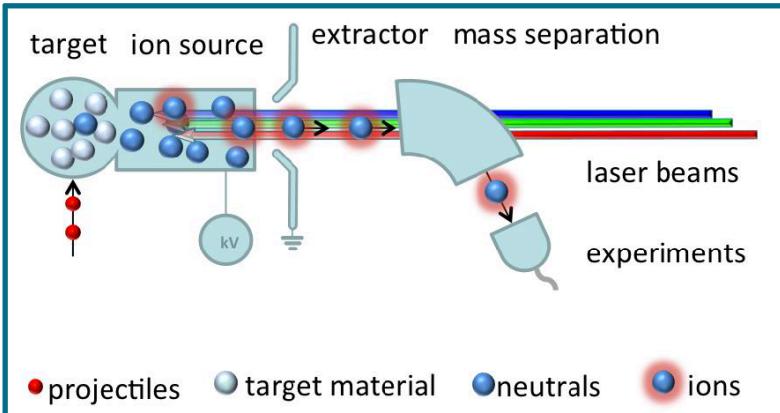




Ion sources

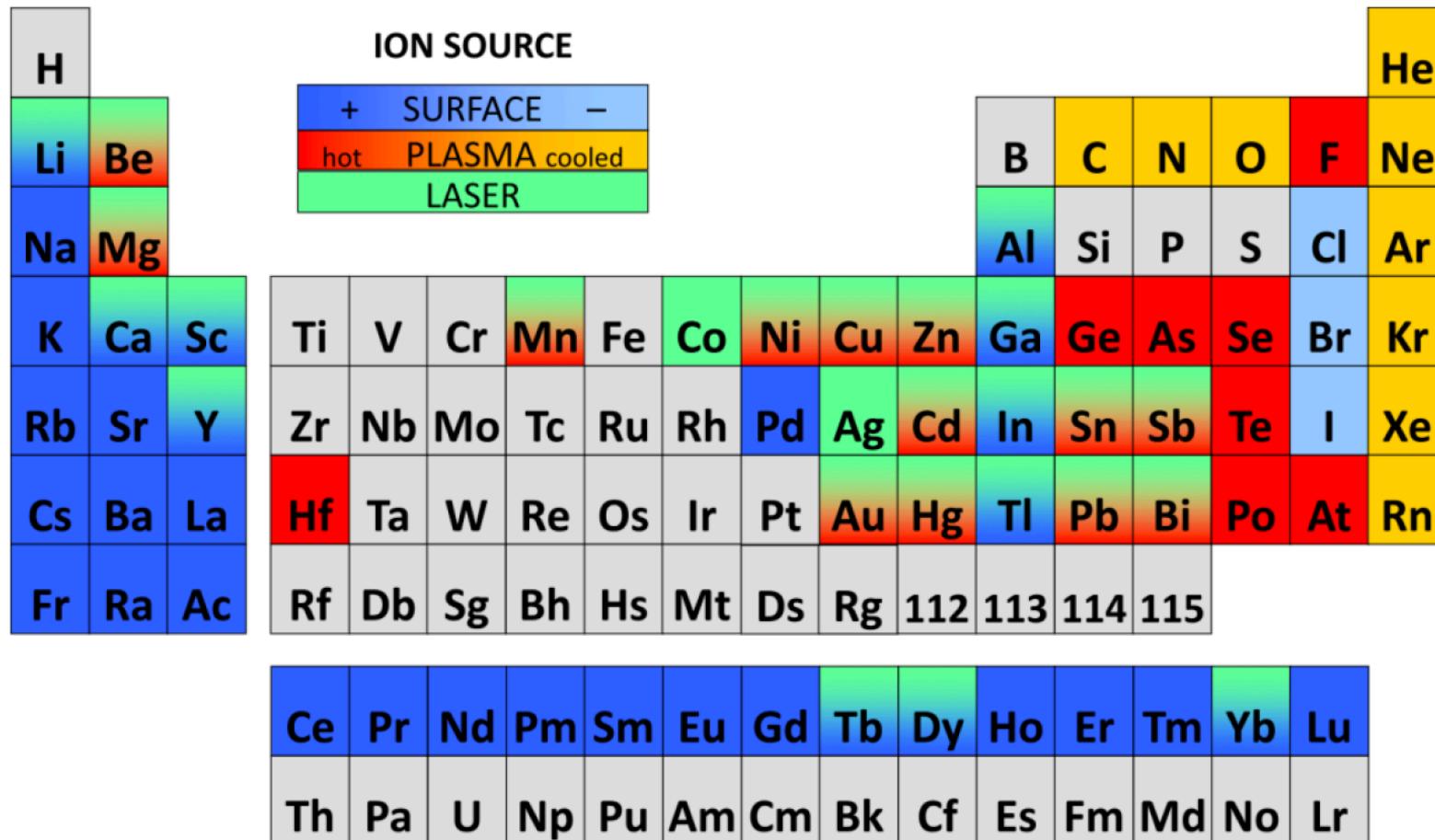
Laser ion source

- 2- or 3-step ionisation
- Isotope and isomer selection
- Universal (almost)



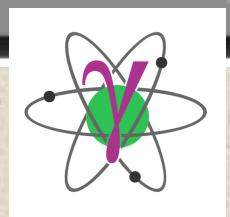
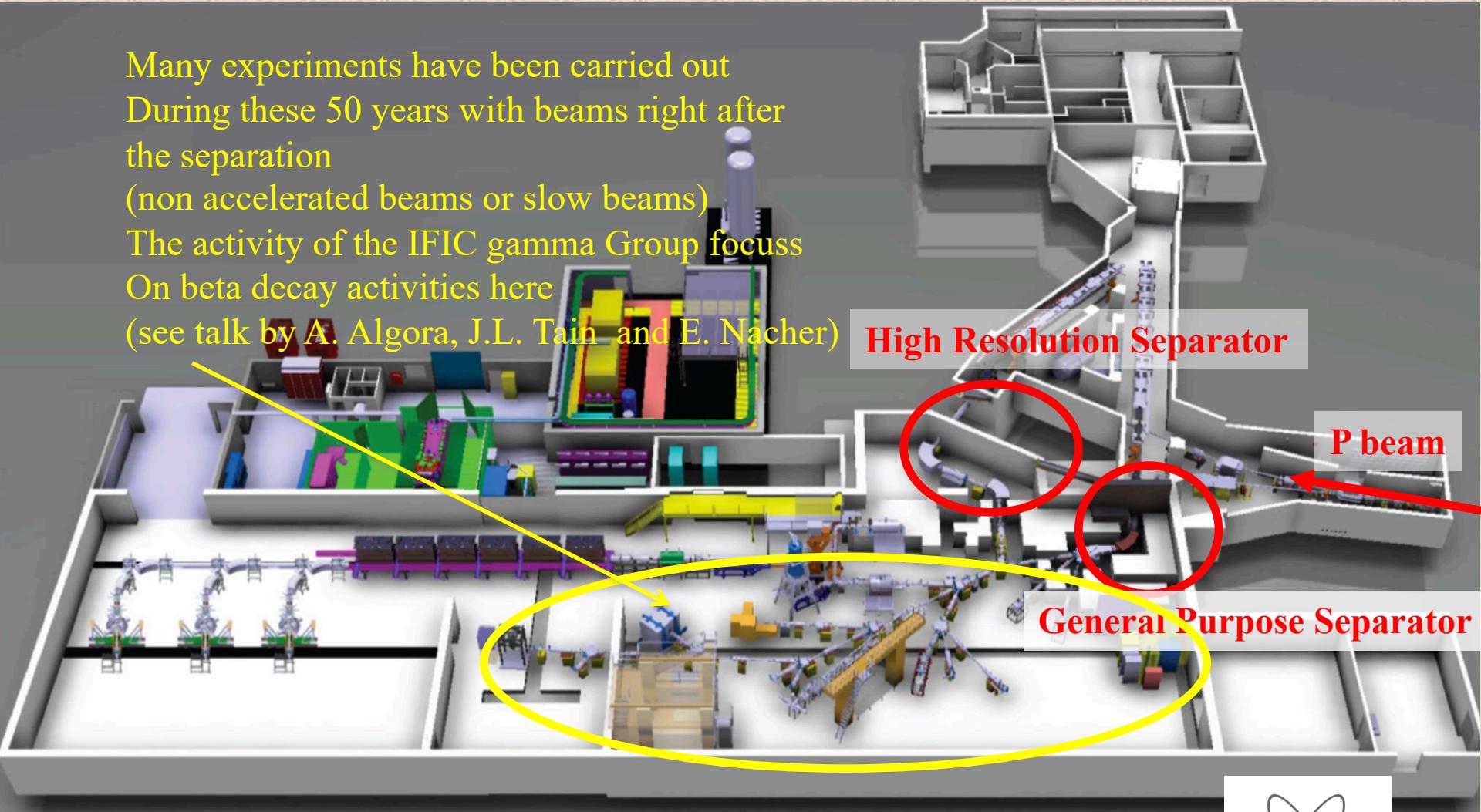
KU LEUVEN

Isotopes produced at ISOLDE



Many experiments have been carried out
During these 50 years with beams right after
the separation
(non accelerated beams or slow beams)

The activity of the IFIC gamma Group focuss
On beta decay activities here
(see talk by A. Algara, J.L. Tain and E. Nacher)



ISOLDE

The On-Line
beams

17 DECEMBER 1964
CERN approves the online
separator project

View 



ISOLDE

The On-Line Isotope Mass Separator ISOLDE is a facility dedicated to the production of a large variety of radioactive ion beams

26 JUNE 1992
First experiment at the ISOLDE Proton-Synchrotron Booster

View 

1951

2015

ISOLDE

The On-Line Isotope Mass Separator ISOLDE is a facility dedicated to the production of a large variety of radioactive ion beams

31 OCTOBER 2001
New accelerator, REX-ISOLDE, goes live

View 

1951

2015

ISOLDE

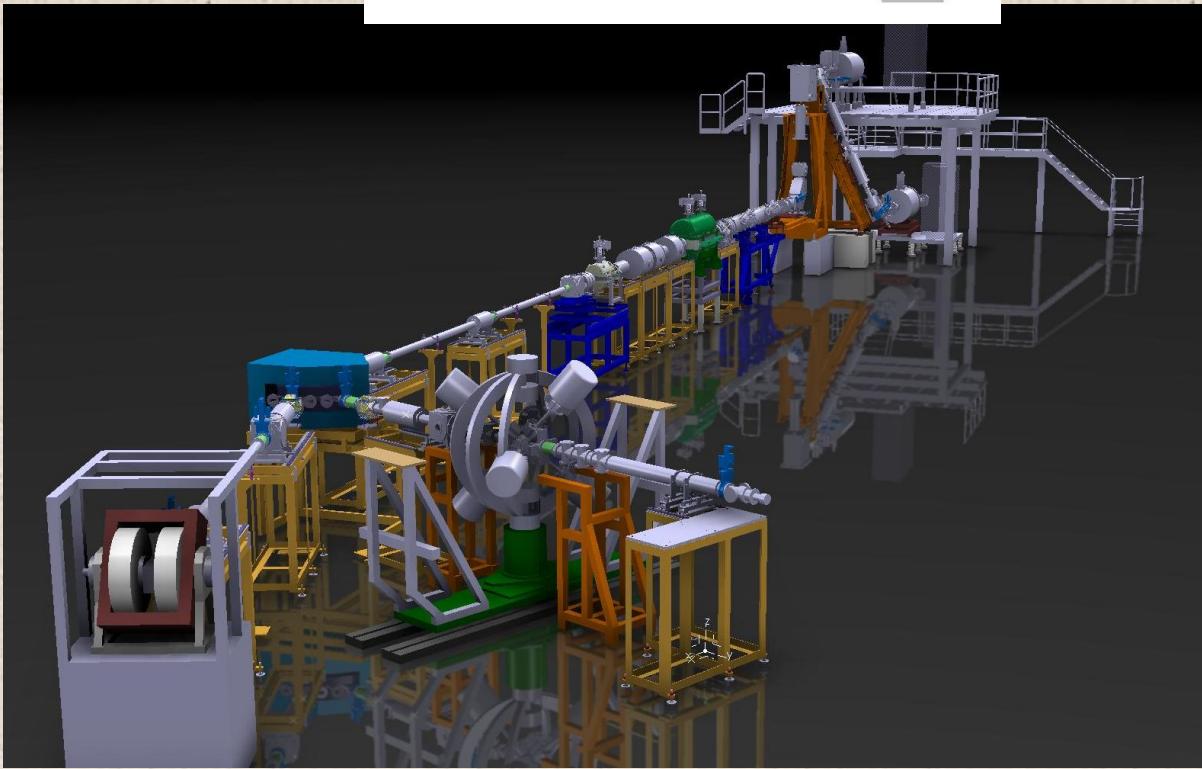
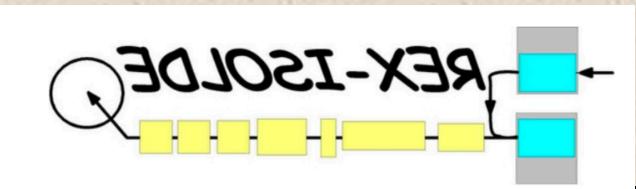
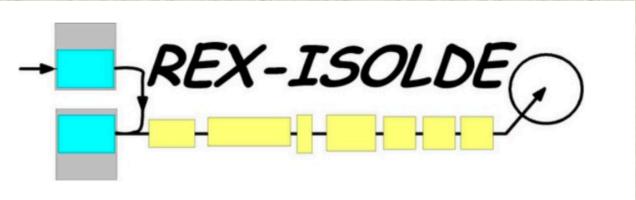
The On-Line Isotope Mass Separator ISOLDE is a facility dedicated to the production of a large variety of radioactive ion beams

22 OCTOBER 2015
First radioactive isotope beam accelerated in HIE ISOLDE

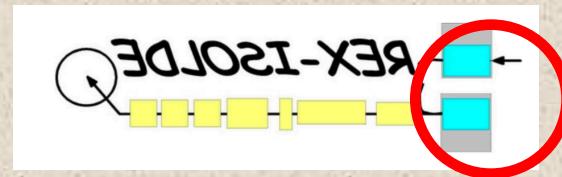
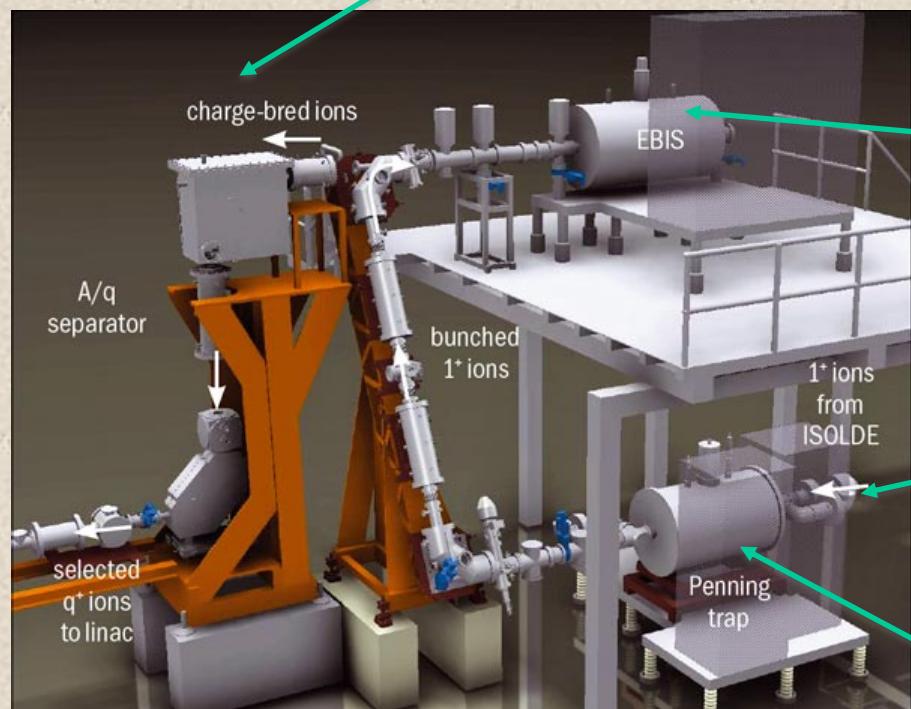
View 

1951

2015



EBIS Ion Source and beam handling: 5-15% efficiency

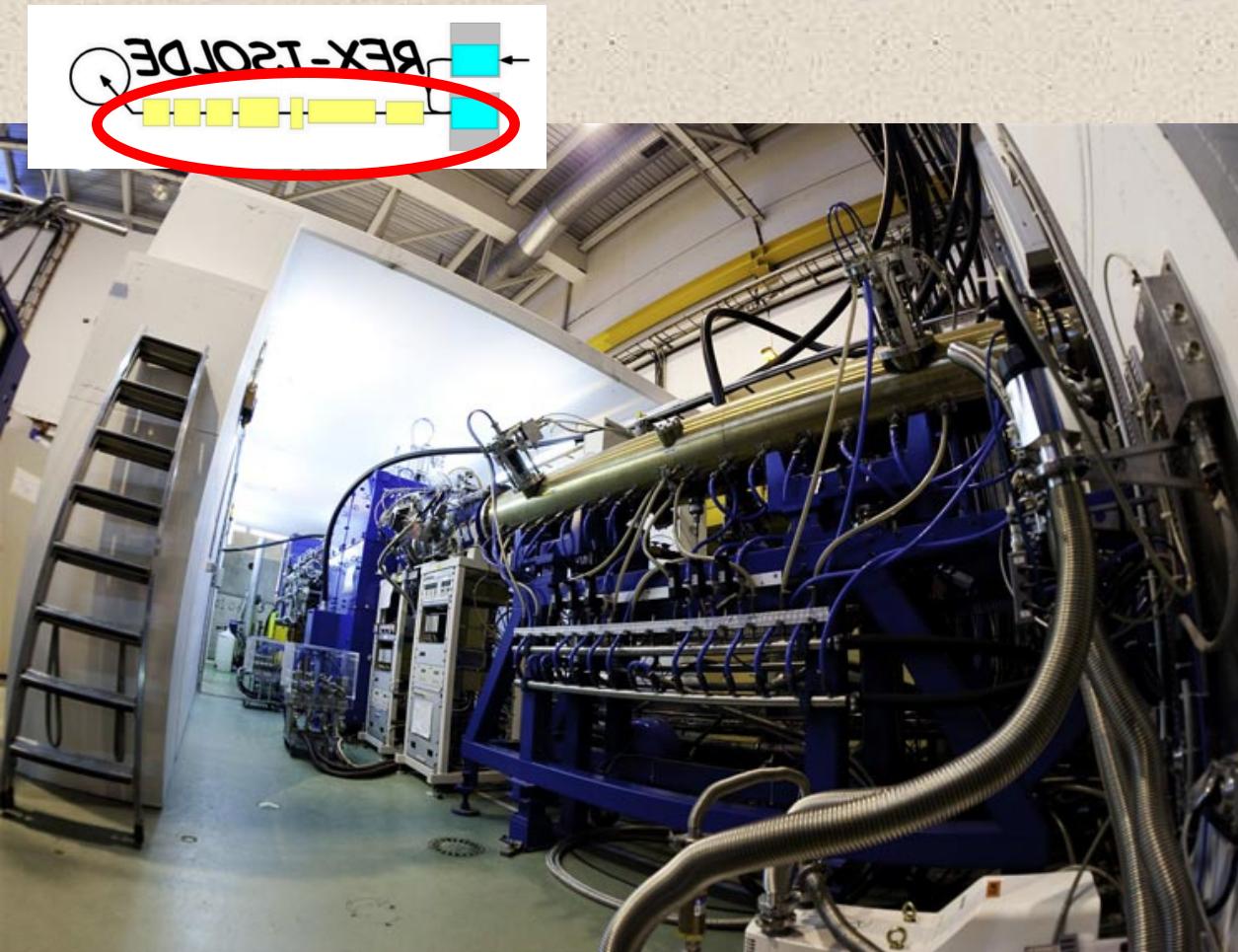


In the EBIS the 1^+ to n^+ ion conversion takes place

Beams from ISOLDE with
 1^+ charge state

Preparation
trap

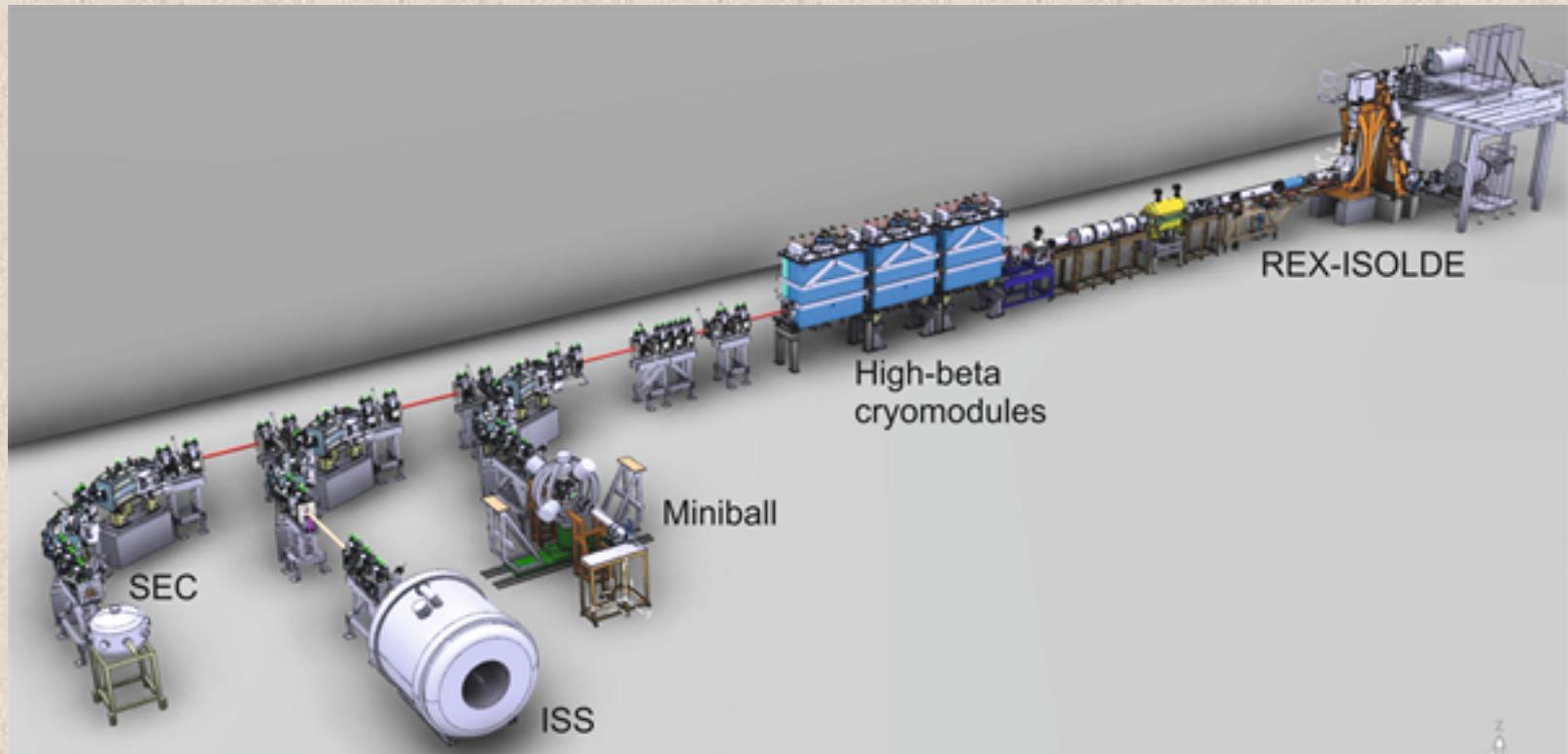
Post accelerator: Normal Conducting LINAC



However, energies only 0.8 to 3 MeV/u which are too low for Nuclear reactions

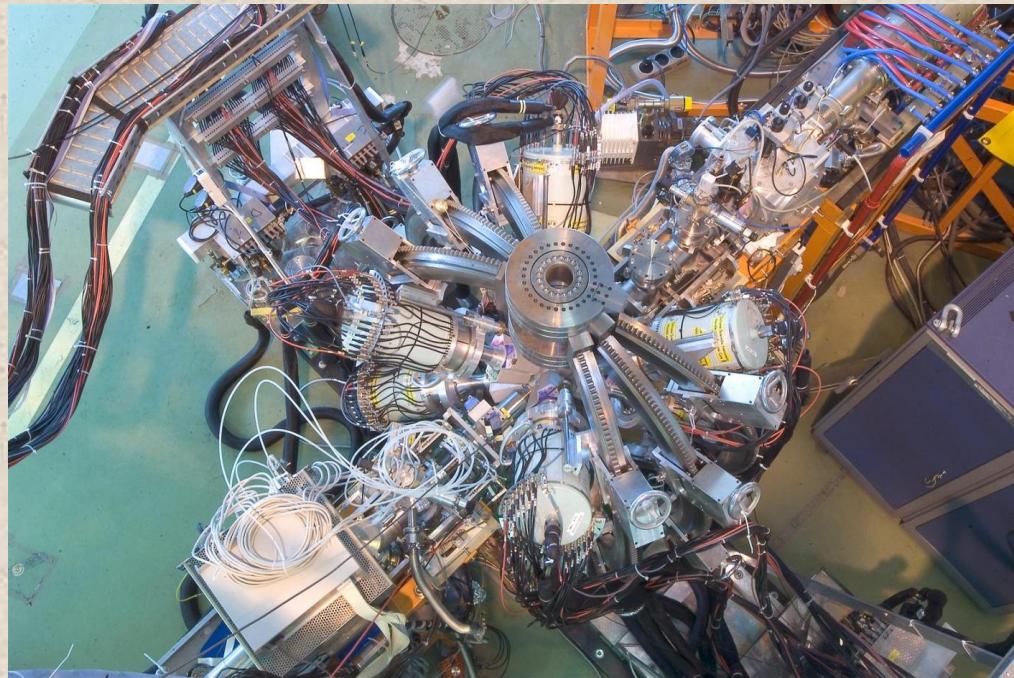
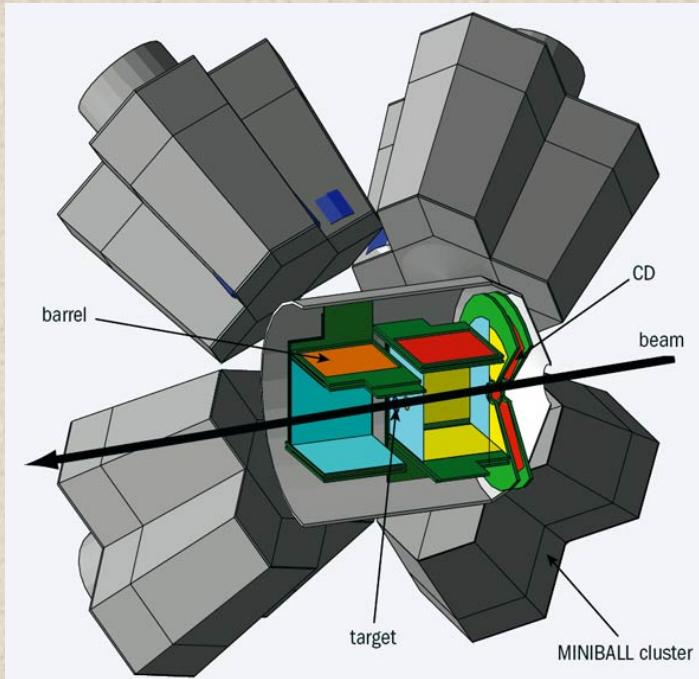
HIE ISOLDE (Superconducting) High Intensity and Energy ISOLDE (soon 10 MeV per nucleon)

4.3 MeV per nucleon



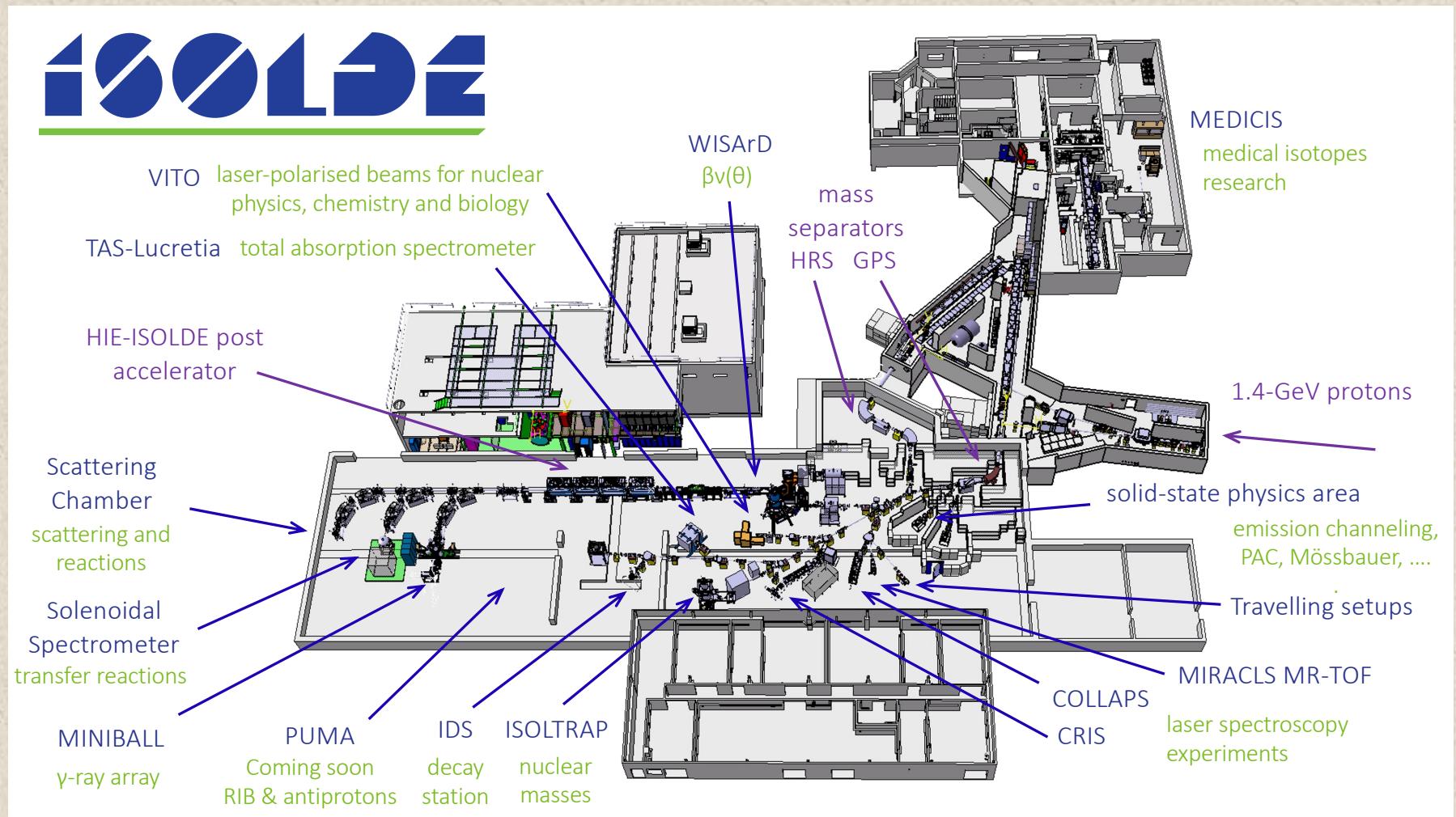
First experiments in 2015

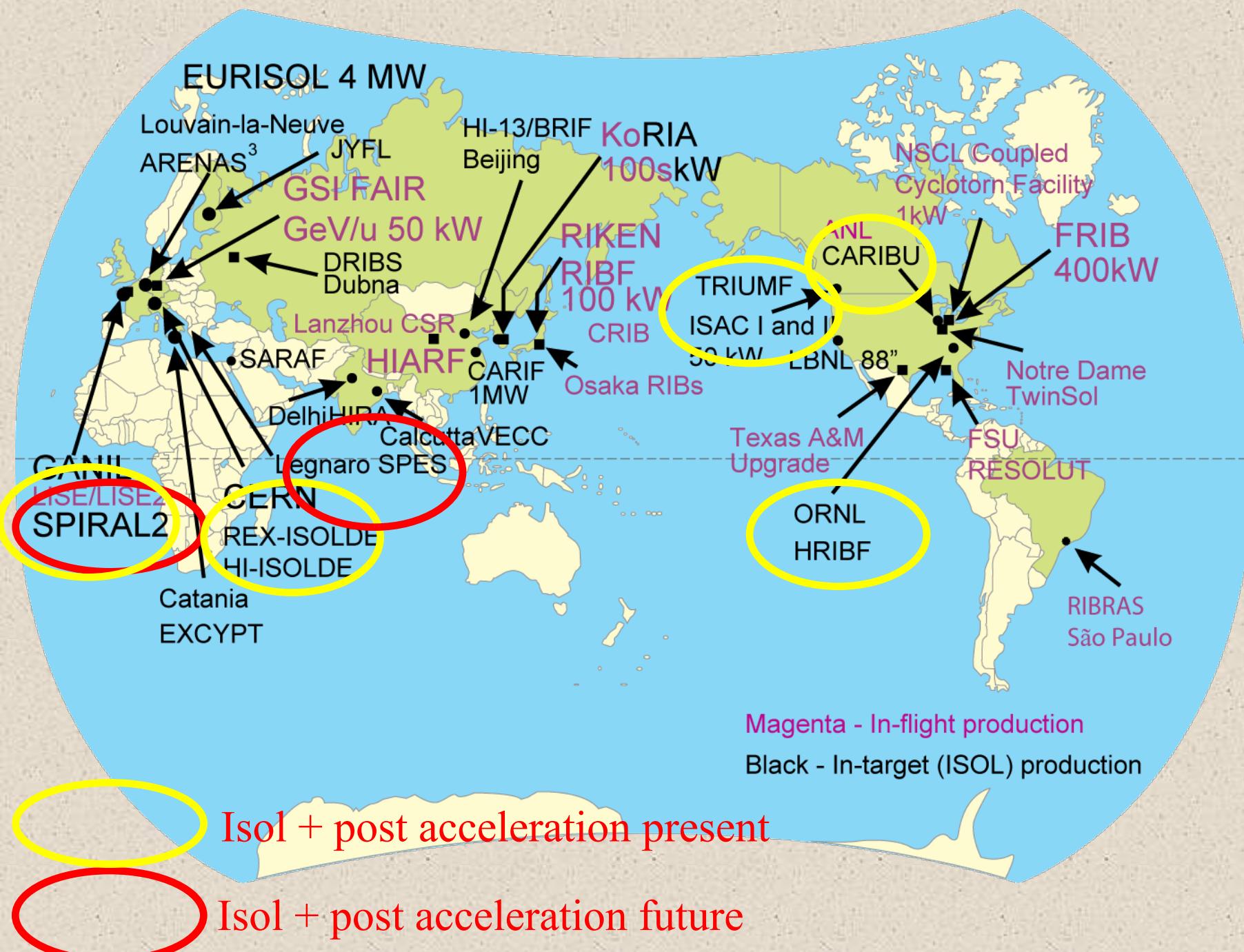
Post accelerated beams + sophisticated detection set-up: Miniball gamma array



24 six-fold segmented, tapered, encapsulated high-purity germanium crystals

ISOLDE lei out as will appear in the NUPECC long range plan 2024

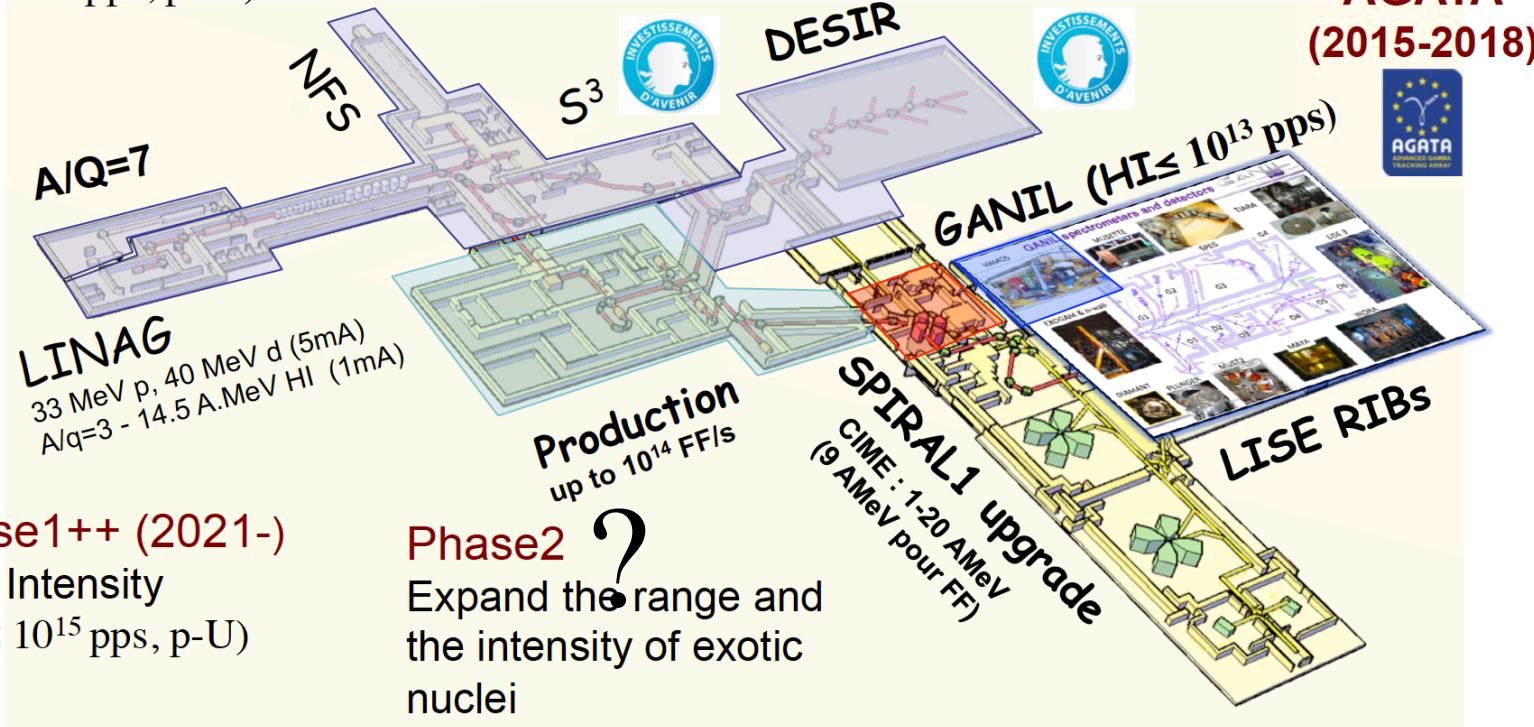




Future ISOL facilities

Phase1 (2016-)

Increase the intensity of stable beams
 High intense neutron source
 $(HI \leq 10^{15} \text{ pps, p-Ni})$



Phase1++ (2021-)

High Intensity
 $(HI \leq 10^{15} \text{ pps, p-U})$

Phase2 ?

Expand the range and
 the intensity of exotic
 nuclei

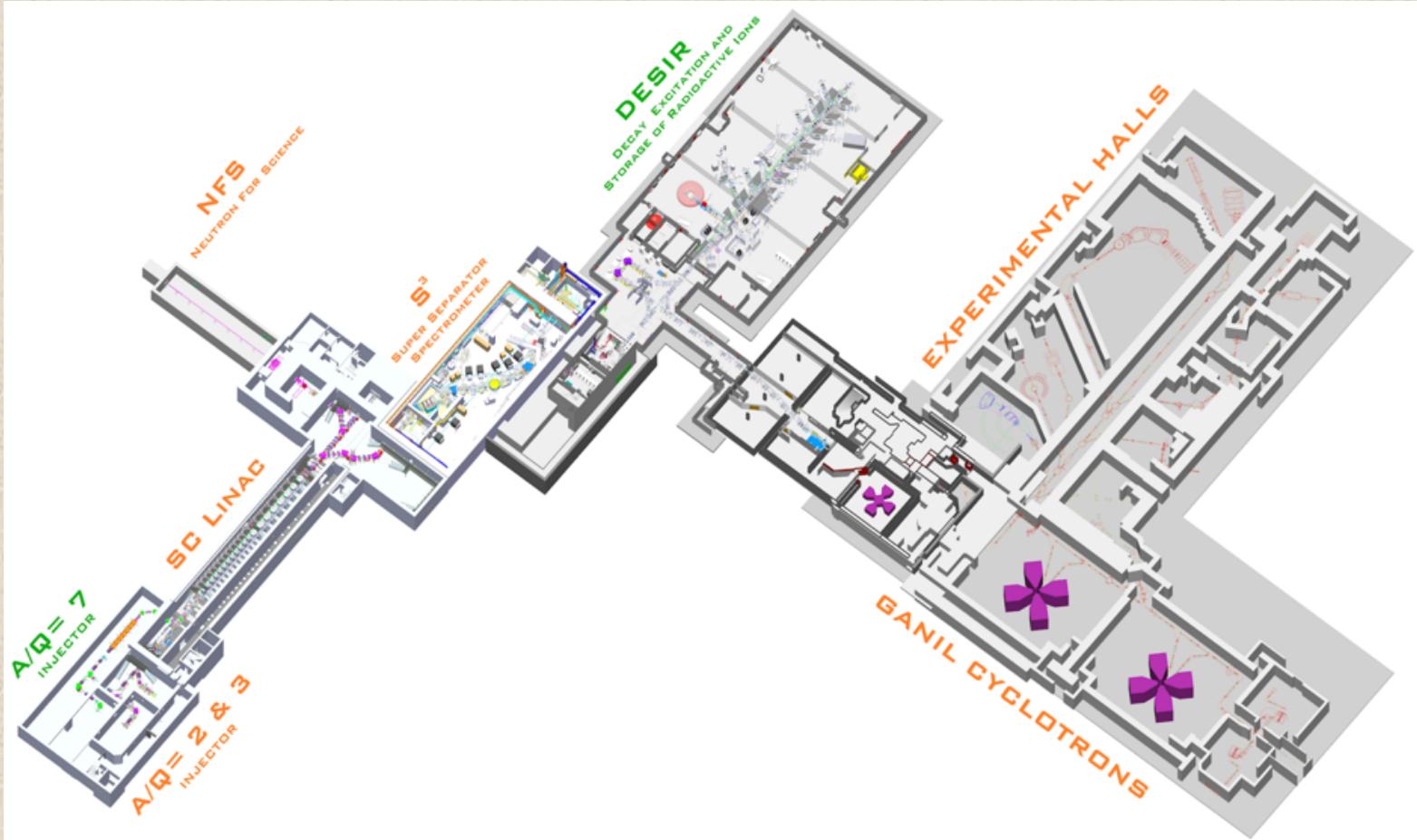
DESIR Phase1+ (2020-)

Low energy facility

AGATA
 (2015-2018)

SPIRAL1 Upgrade (2017-)
 New light RIBs from
 beam/target fragmentation

Unfortunately, the SPIRAL lei out, as it will appear in the Long Range plan 2024, does not include the ISOL part, but it is mentioned as future plan



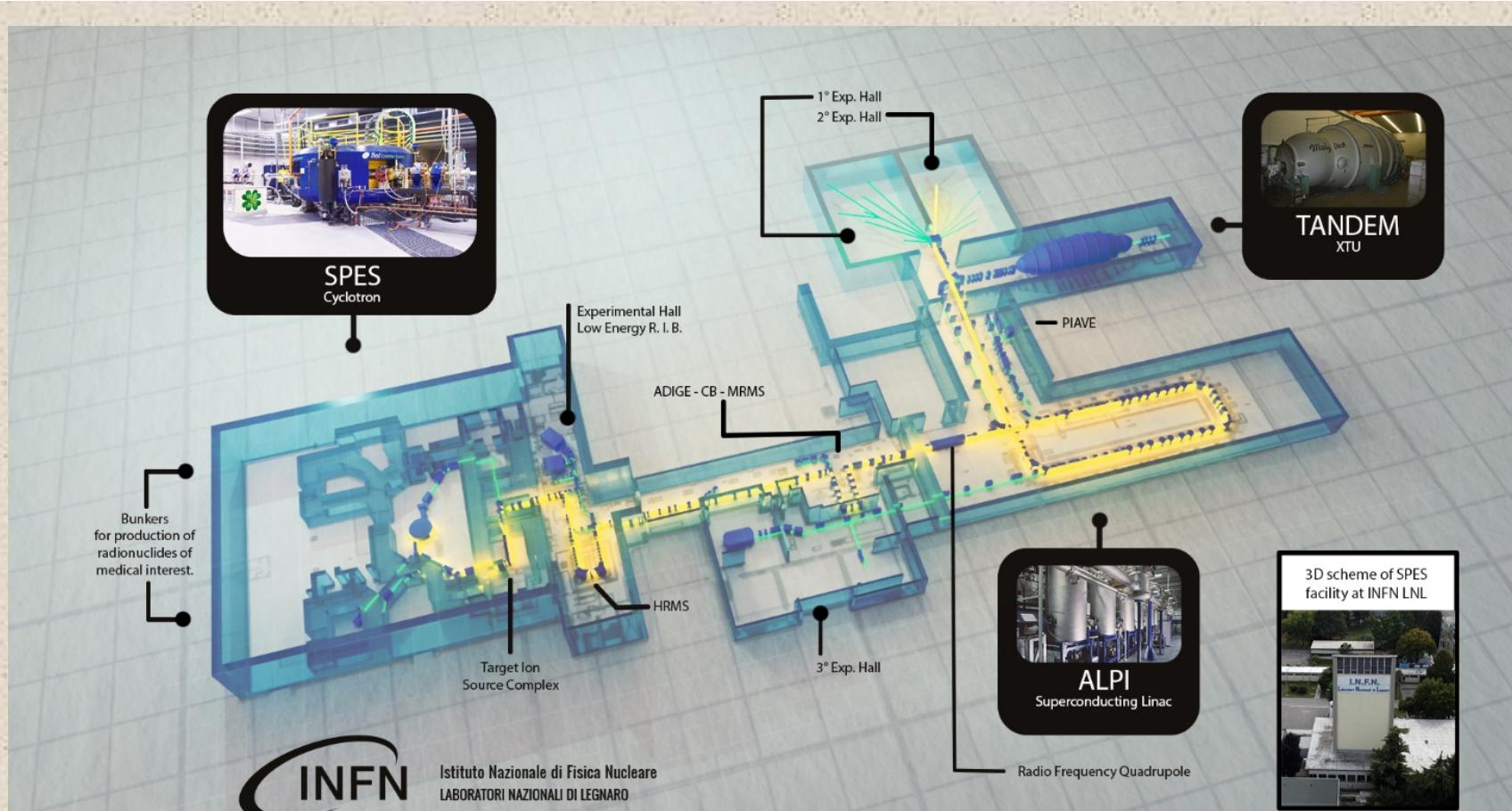
SPES

Selective Production of Exotic Species

Under construction

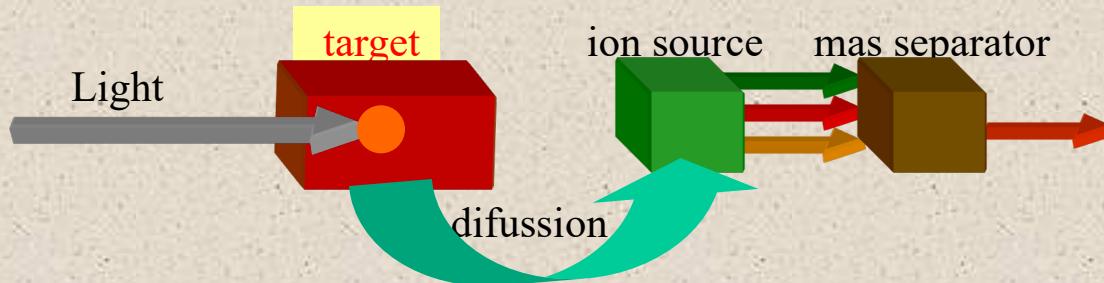


SPES Project



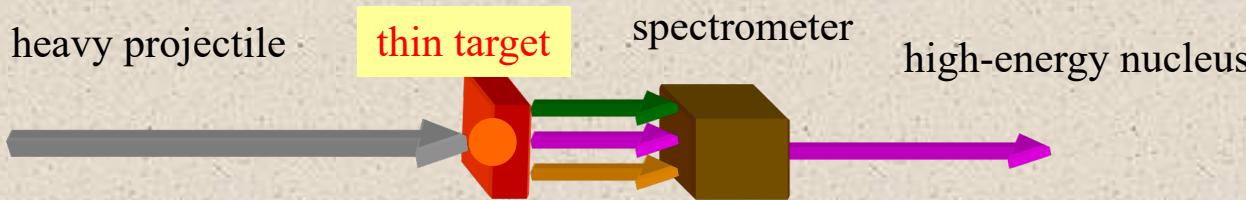
Radioactive nuclei production techniques

Isotopic separation on-line (ISOL)



- 😊😊 We can use thick production targets (100% of energy range)
→ and high injector beam current (upto 10^{16} s^{-1})
- 🙁 long extraction and ionization time (100 ms)
- 🙁 chemistry dependent

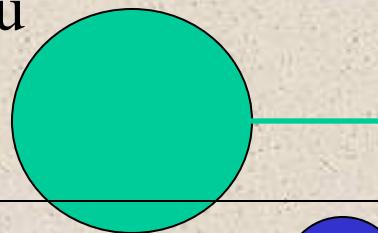
In-flight fragmentation



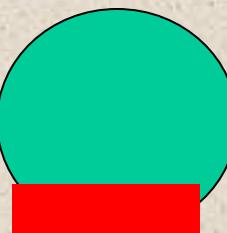
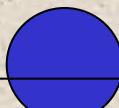
- 🙁 thinner targets (10% of range) => lower beam currents (upto 10^{12} s^{-1})
- 😊 short separation+identification time (100 ns)
- 😊 chemistry independent
- 😊 relativistic energies

Projectile Fragmentation Reactions

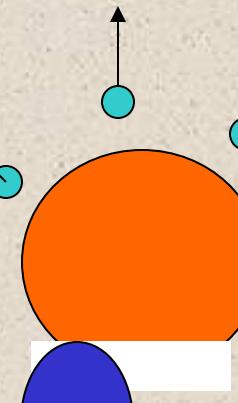
Projectile
1 GeV/u



target



hotspot



Excited
pre-fragment

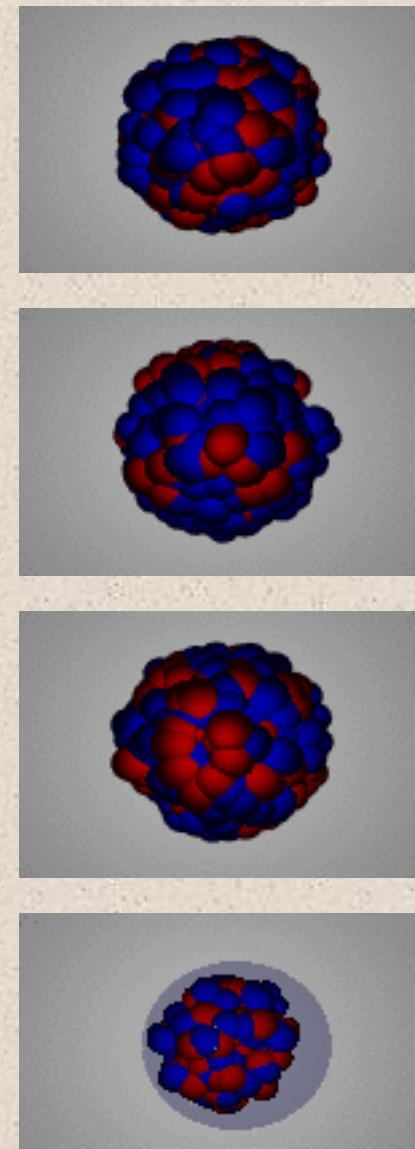
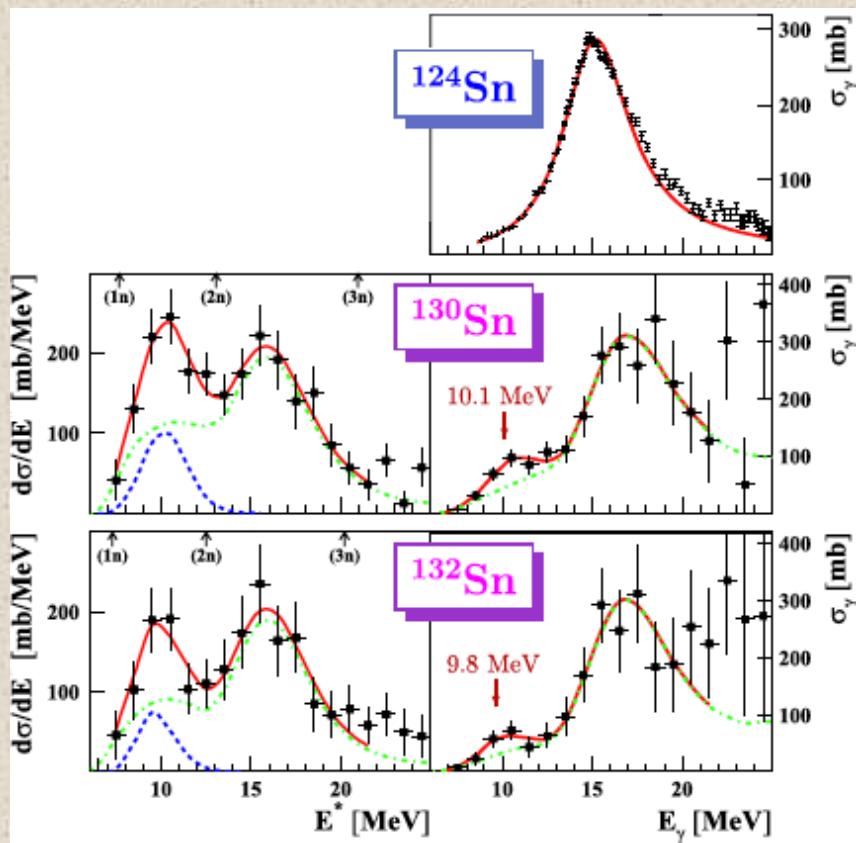
Final target like
fragment

Energy (velocity) of beam > Fermi velocity inside nucleus ~ 30 MeV/u

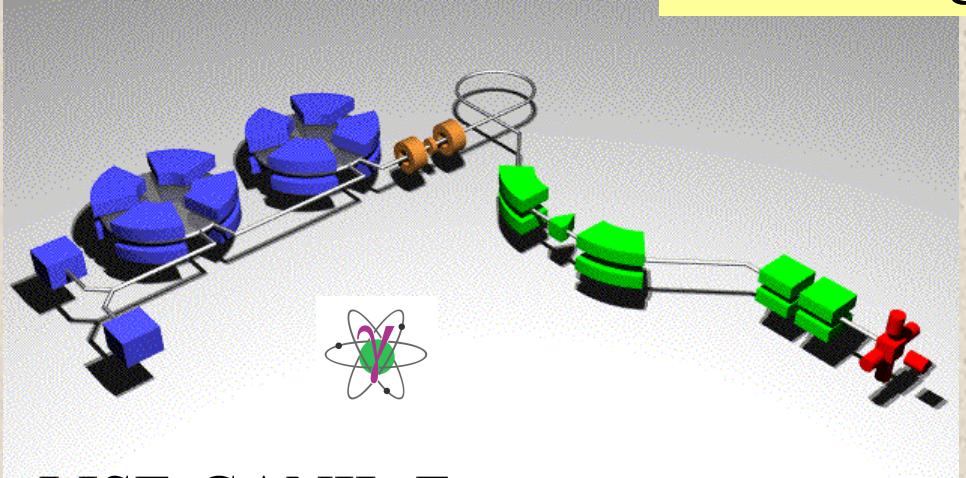
Can ‘shear off’ different combinations of protons and neutrons.

Large variety of exotic nuclear species created, all at forward angles with \sim beam velocity.

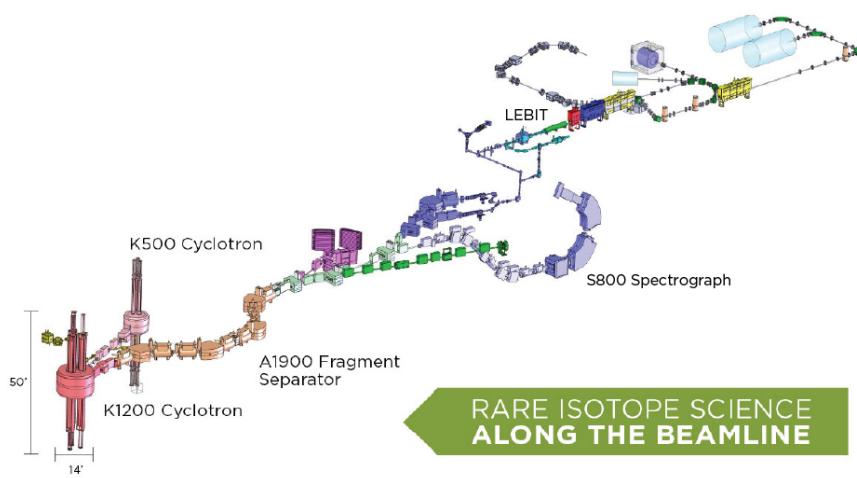
The energy issue: Some physics can only be reached with relativistic energies



Actual Fragment Separators

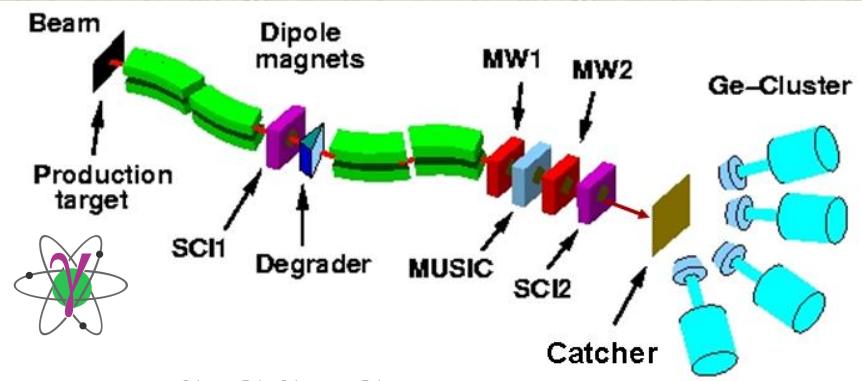


LISE-GANIL-France

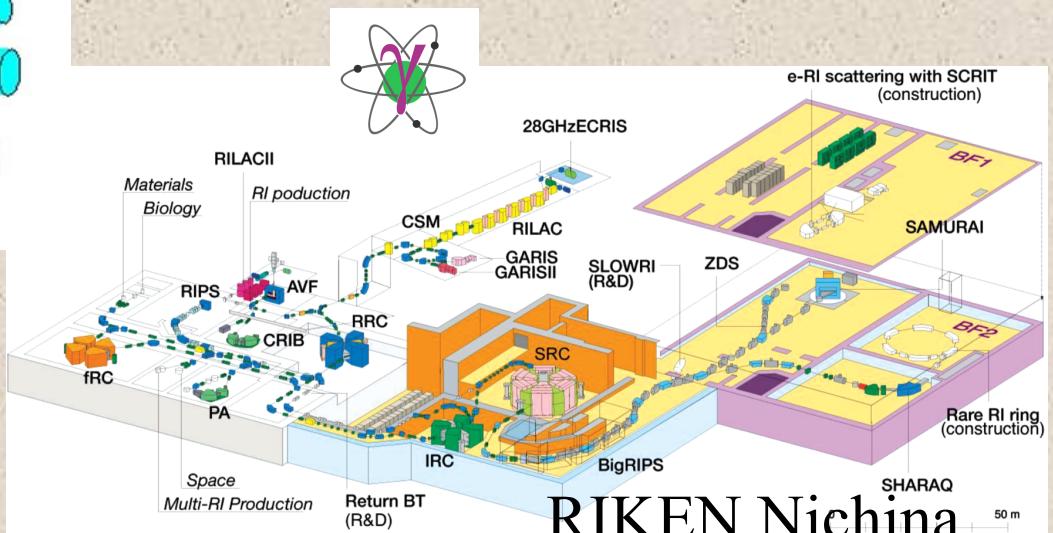


RARE ISOTOPE SCIENCE
ALONG THE BEAMLINE

NSCL-RIA→FRIB USA



FRS GSI Germany



RIKEN Nichina
Center, Japan

FRIB - Facility for Rare Isotope Beams

World-leading Next-generation Rare Isotope Beam Facility

- Rare isotope production via in-flight technique with primary beams up to 400 kW, 200 MeV/u uranium
- Fast, stopped, and re-accelerated beam capability
- Upgrade options
 - 400 MeV/u for uranium
 - ISOL production – multi-user capability

FRIB project start 6/2009

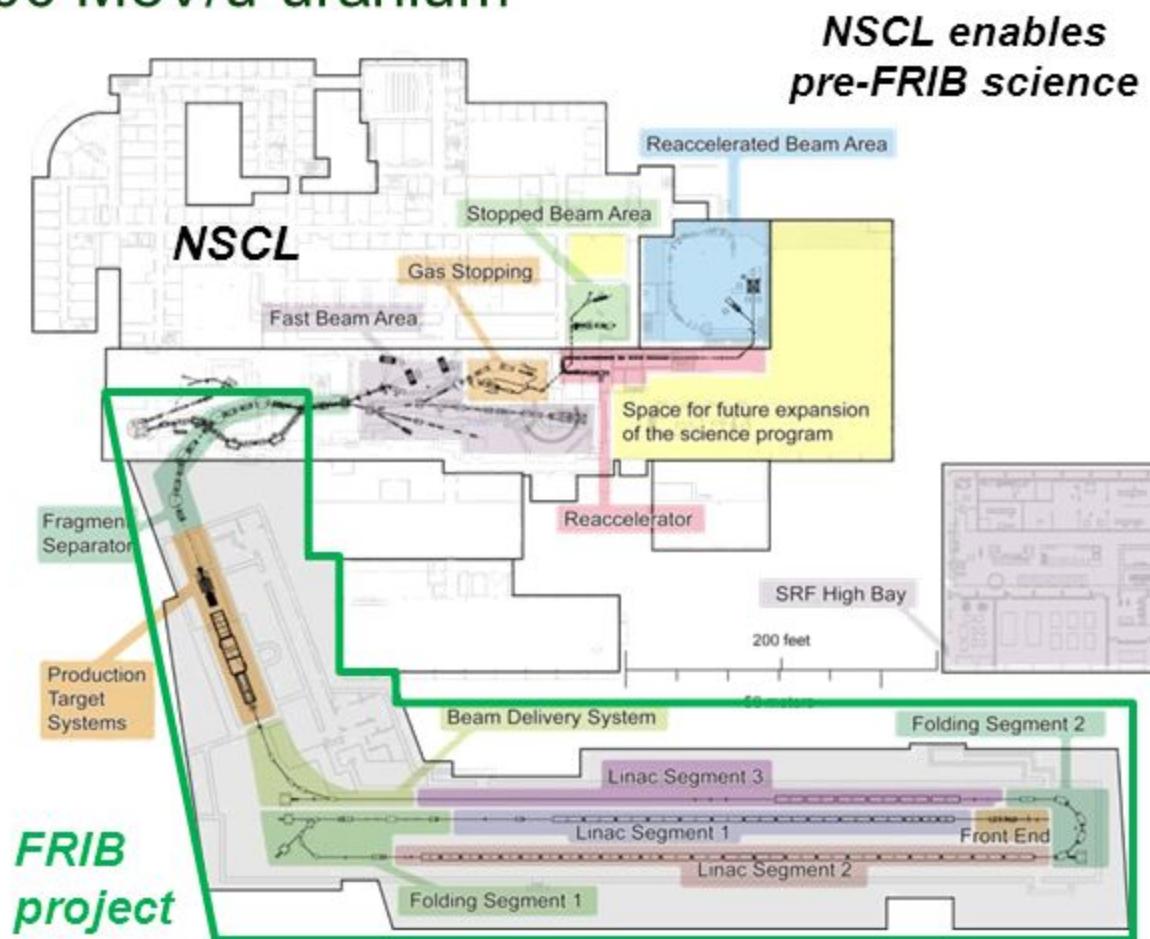
Civil construction started 3/2014

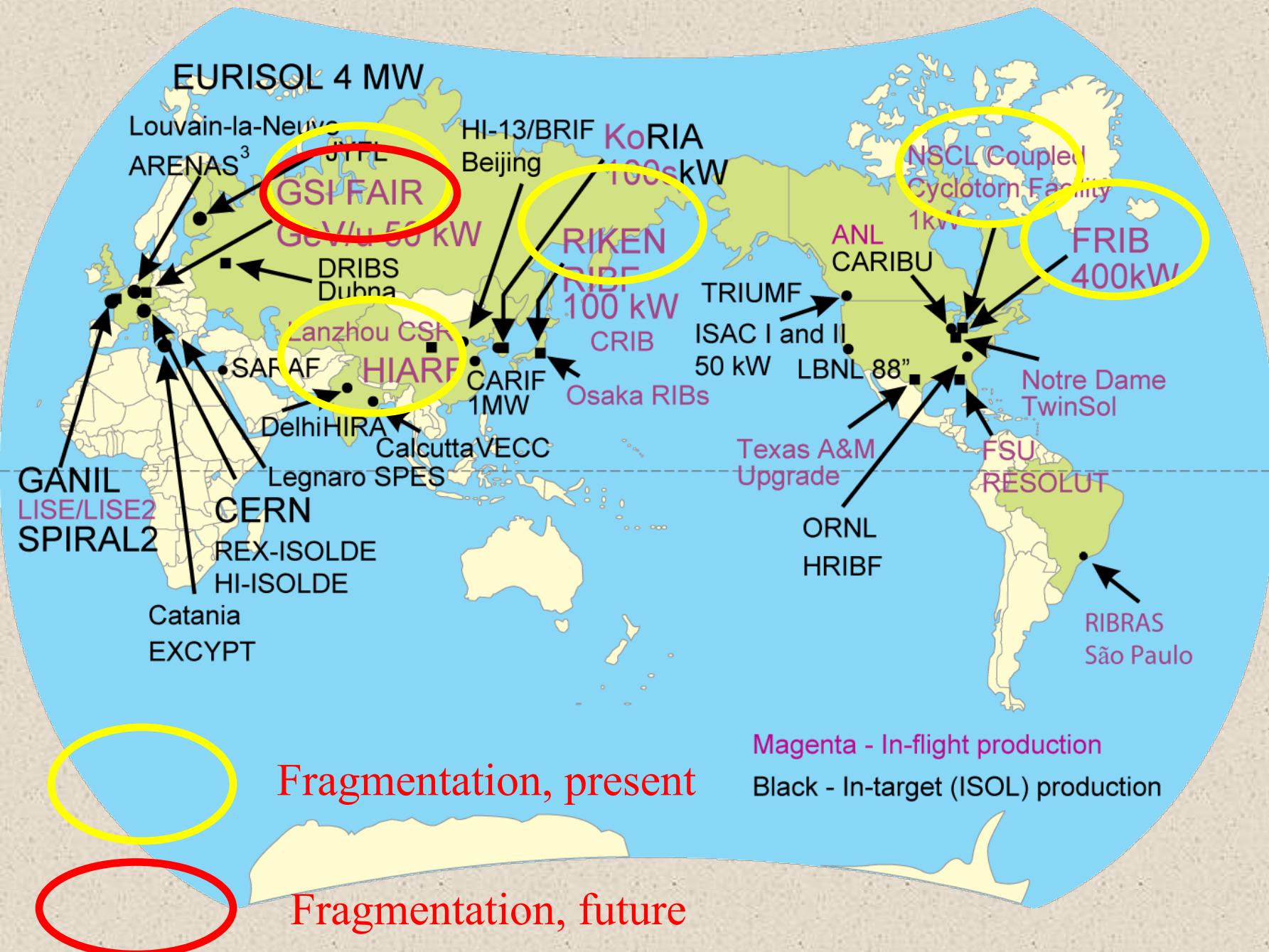
Technical construction started 10/2014

Managed to early completion 12/2020

CD-4 (project completion) 6/2022

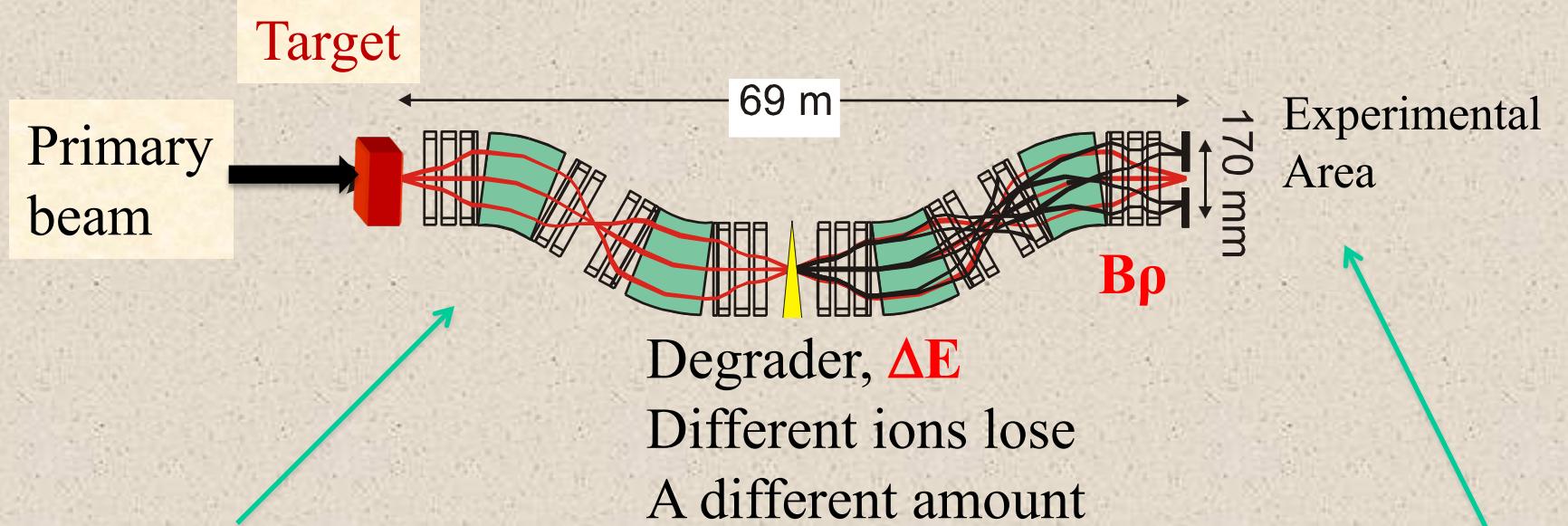
Total project cost \$730 million



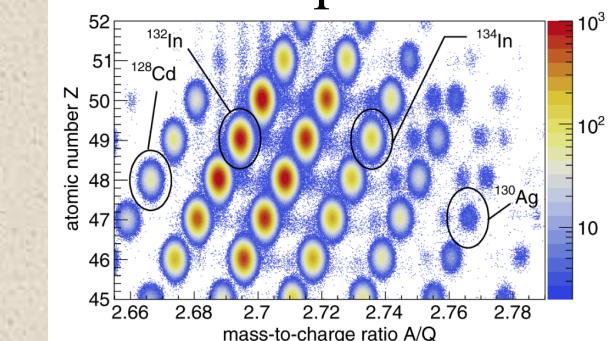


The FRS fragment separator at GSI

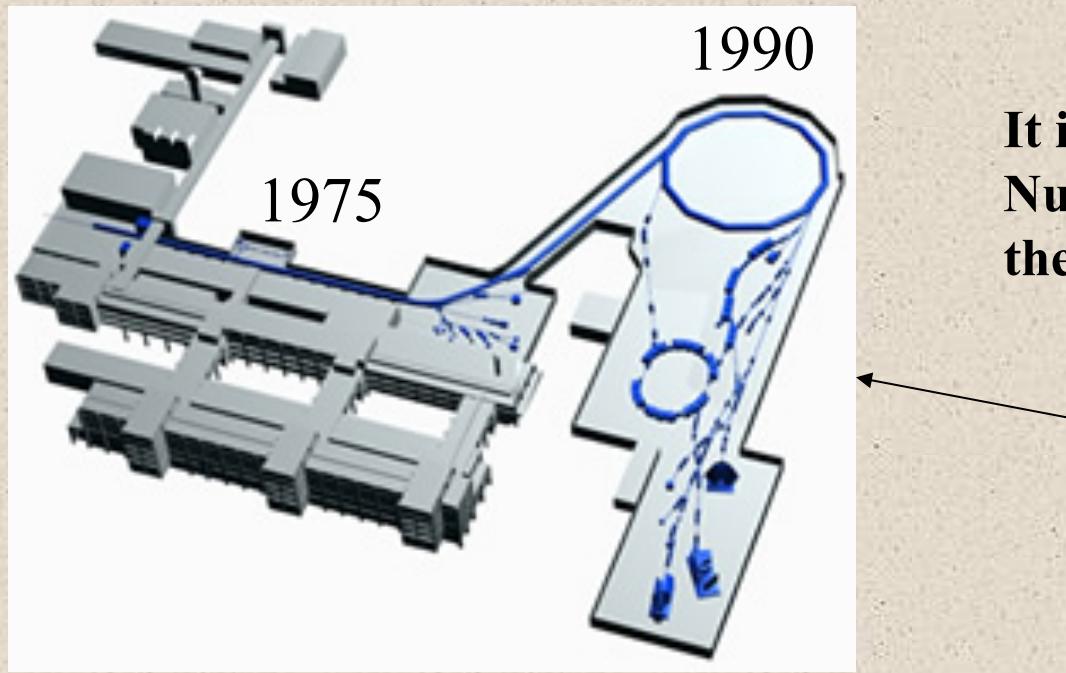
B ρ - Δ E-B ρ Separation Method



Ions are selected according to the momentum to charge ratio $Mv/q = Mv/Z = B\rho$. Primary beam is rejected.



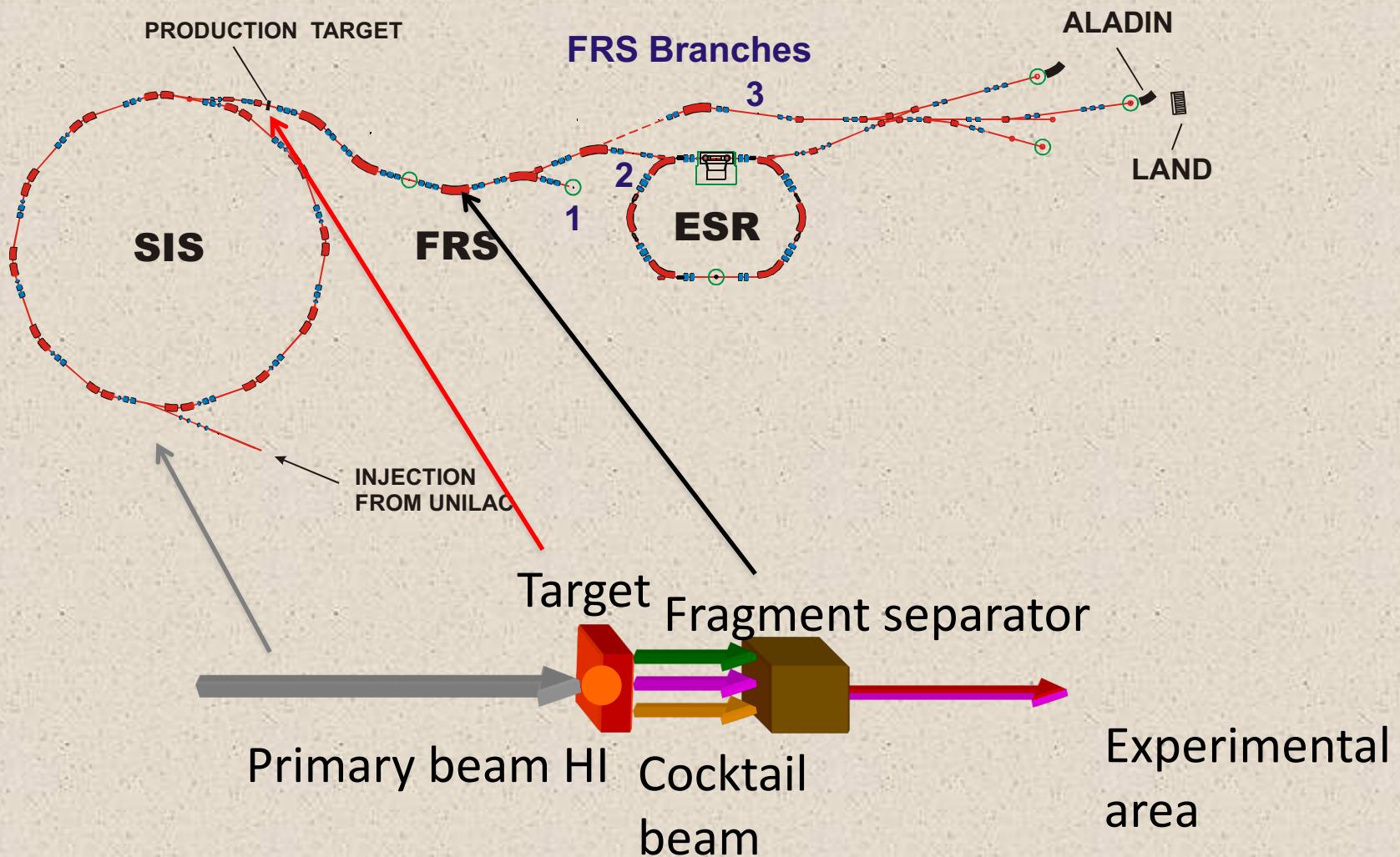
The GSI fragmentation facility



It is possible to accelerate stable Nuclei from H (Z=1) to U (Z=92), in the UNILAC + Synchrotron.

**Fragmentation of the primary ion beam:
“cocktail” of ions**

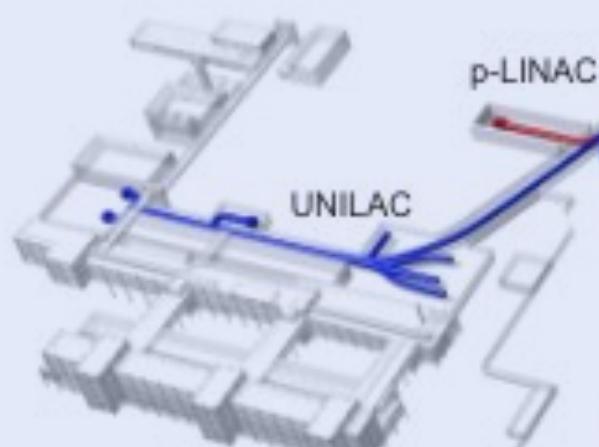




Near Future



Existing GSI facility



GSI

$Z = 1 - 92$

(from p to U)

Up to 2 GeV/nucleon

Beams at FAIR (future):

Intensity: factor 100 (prim. beams)

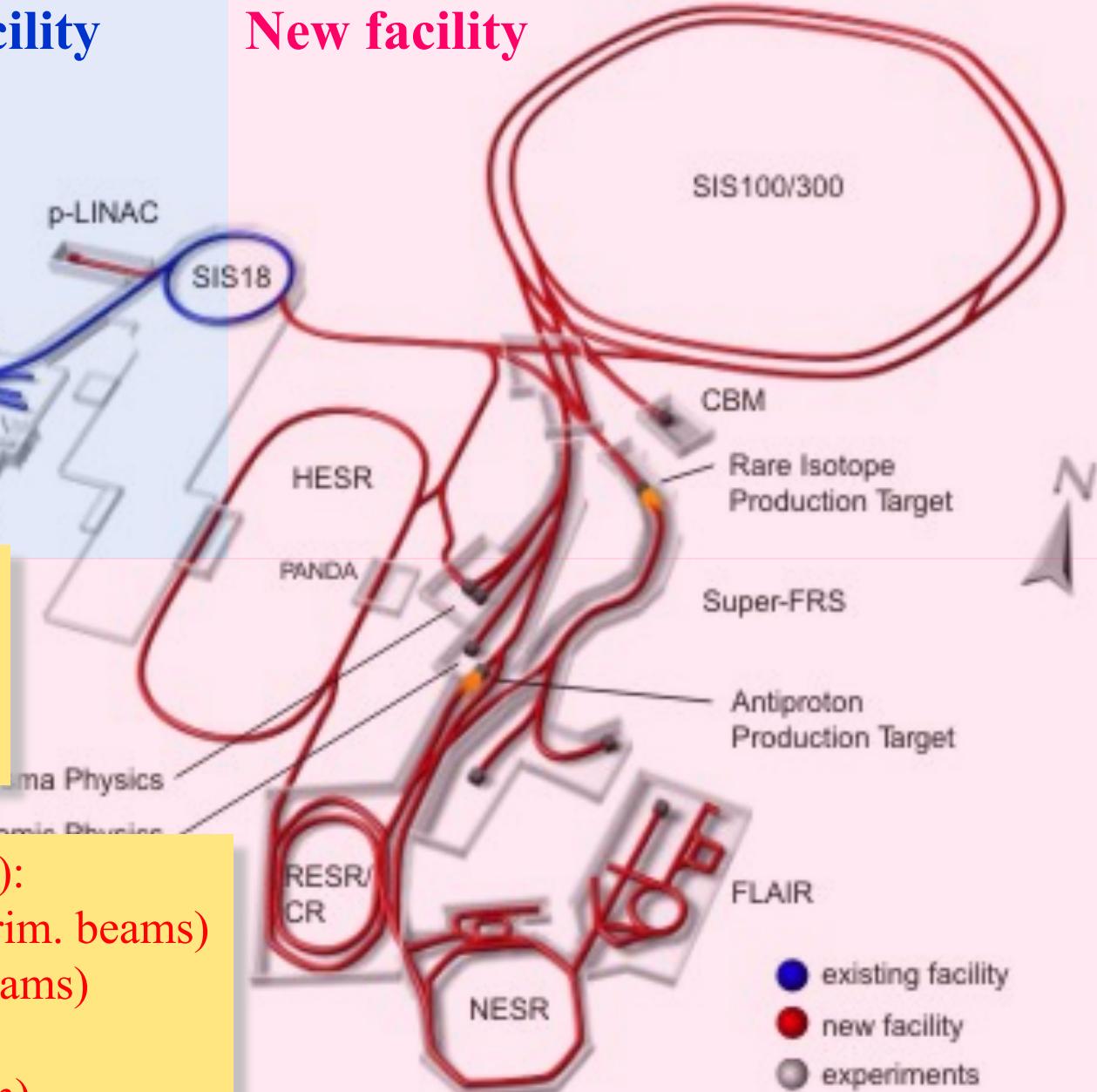
10 000 fold (second. beams)

$Z = -1 - 92$

(anti-protons to uranium)

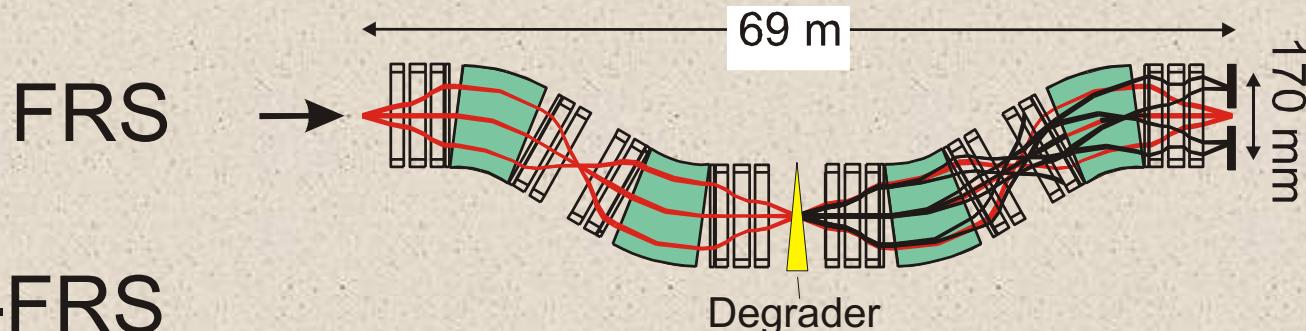
Up to 35 - 45 GeV/u

New facility

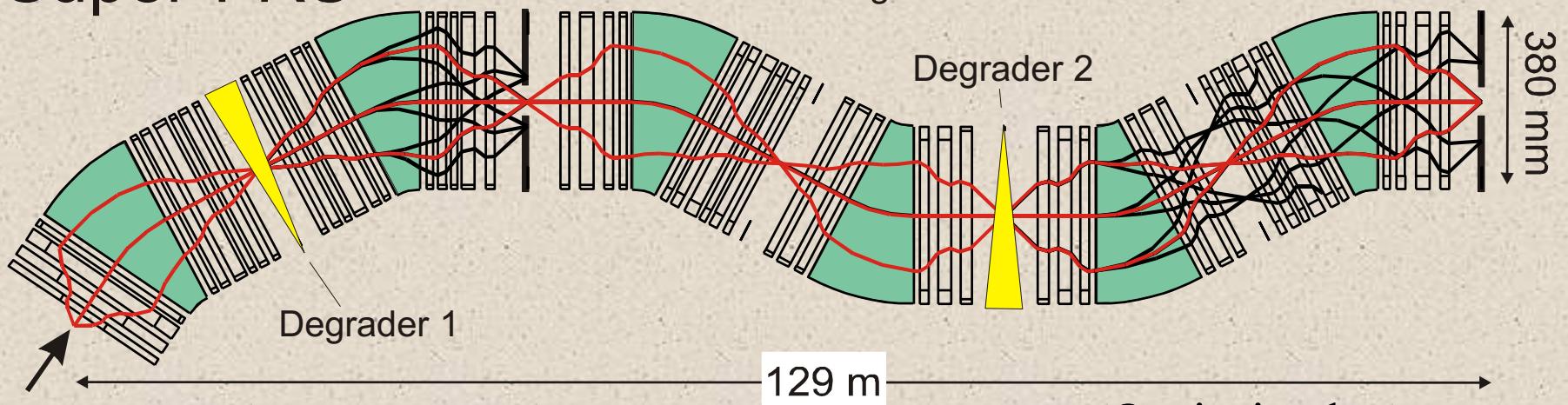


FRS to Super-FRS

B β - ΔE -B β Separation Method

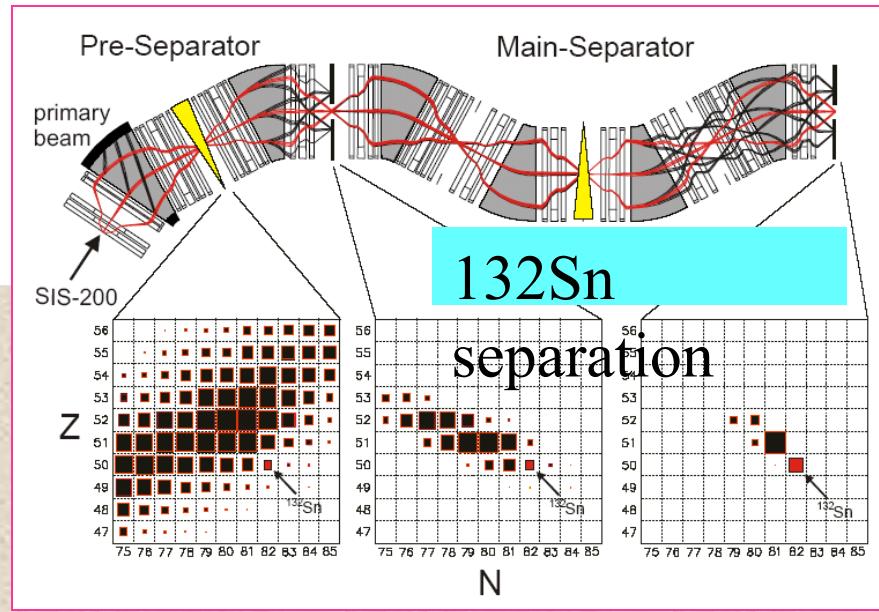
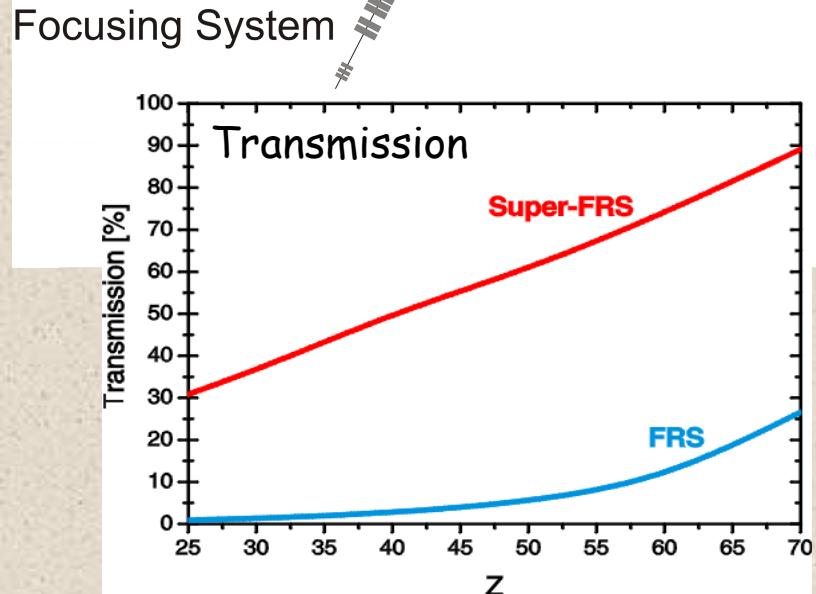
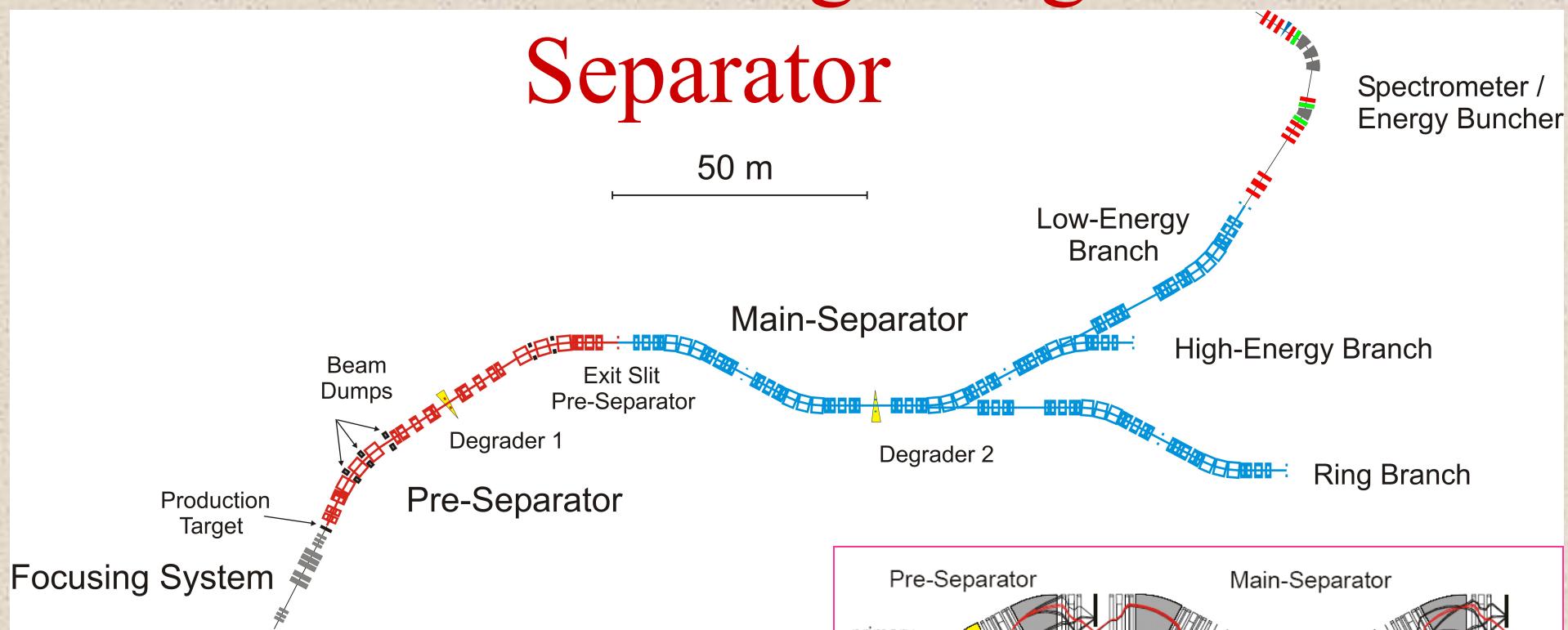


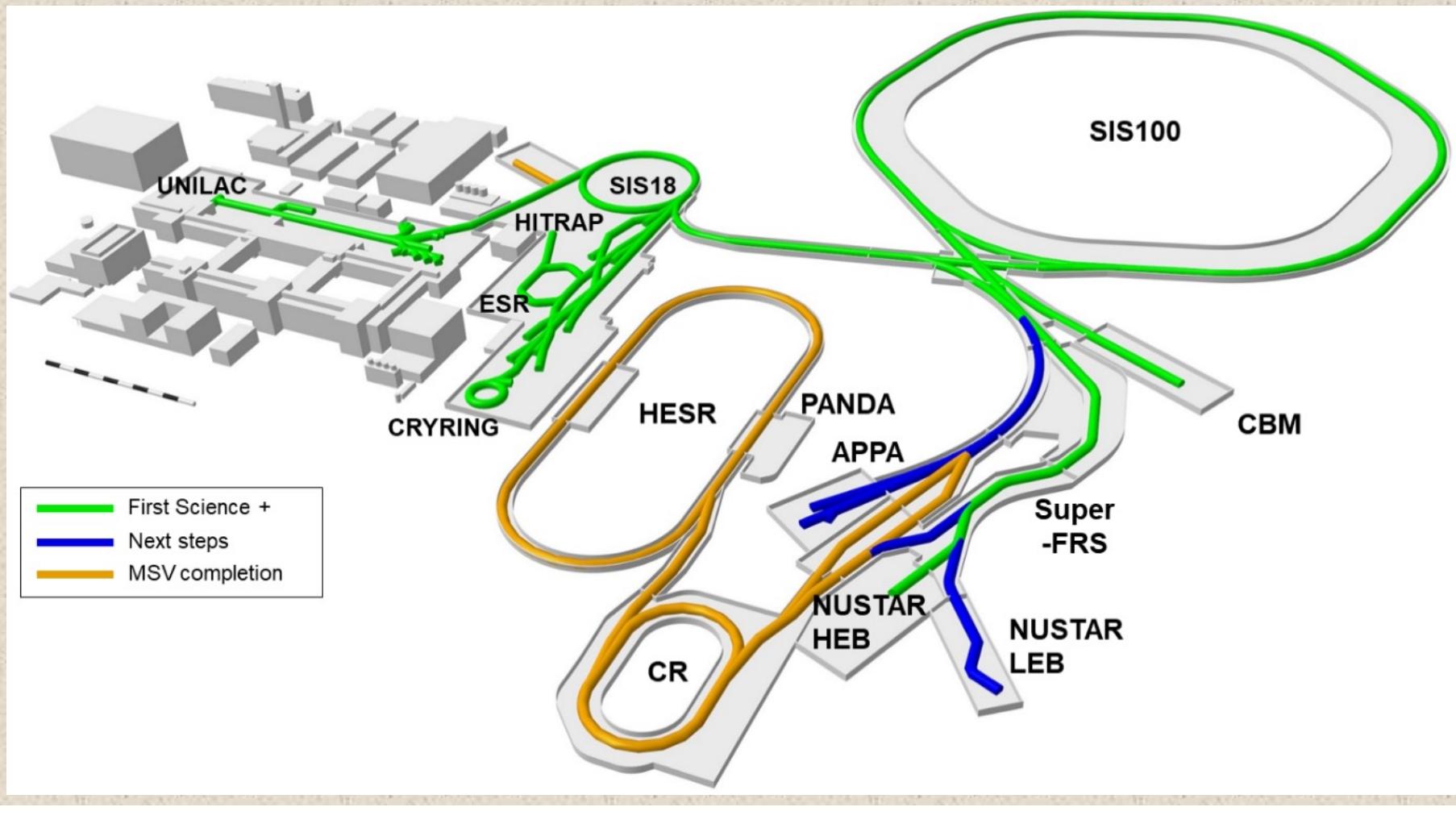
Super-FRS



Optimised to separate
very fast fragment or
fission products

SUPERconducting FRagment Separator





FAIR Construction Field





FAIR in
construction
as it was the last
Master Curse in
2022



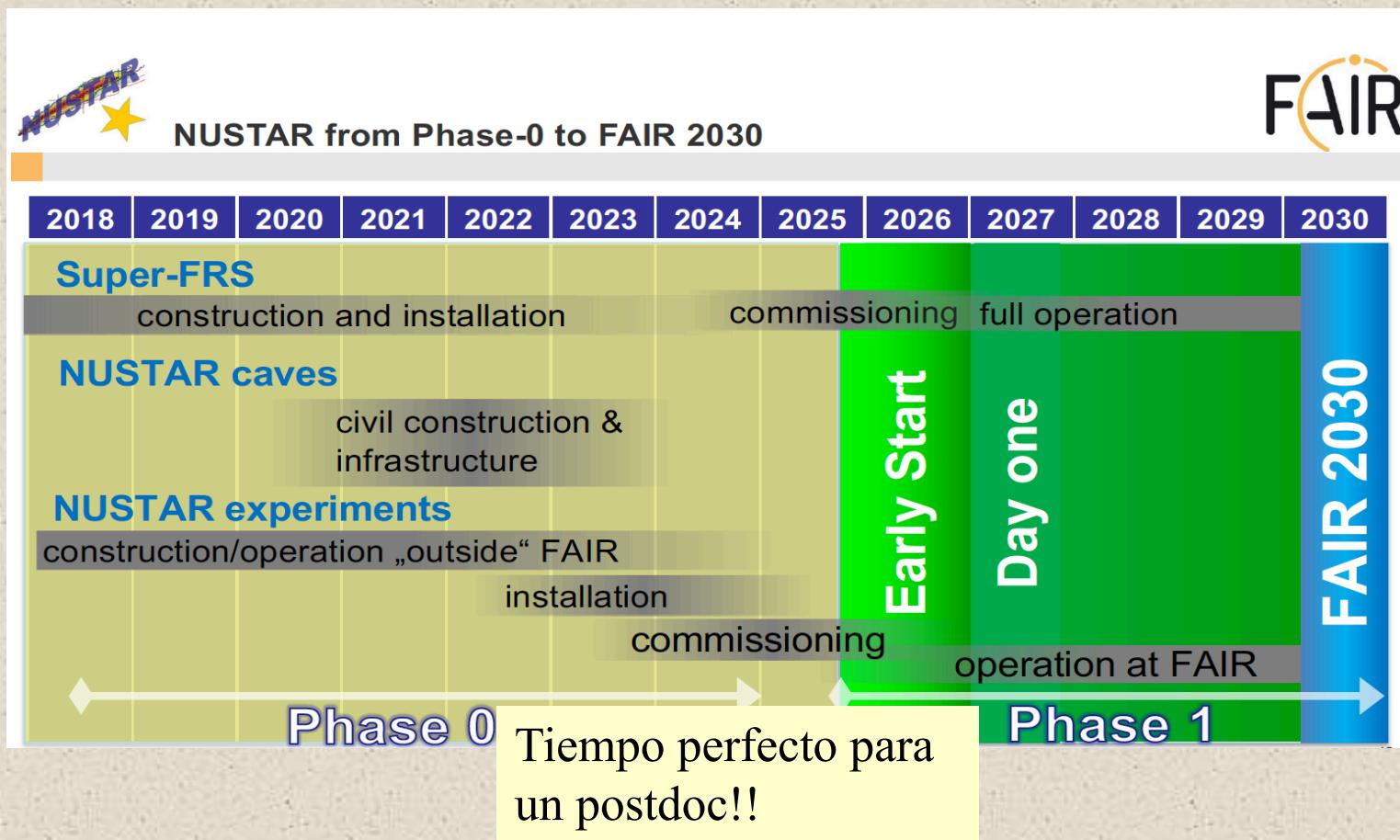
- FAIR GmbH
- GSI GmbH



The high Energy Cave 2023

(See talk by Dolores Cortina on
Reactions at Relativistic
Energies)

FAIR - NUSTAR schedule



ISOL and In-Flight facilities-Partners

It is probably true to say that if we worked at it virtually all experiments could be done with both types of facility but they are complementary.

In-Flight

- Relativistic beams
- Universal in Z
- Down to very short $T_{1/2}$
- More exotic beams

ISOL

- High intensity beams with ion optics comparable to stable beams
- Easy to manipulate beam energies from keV to 10s of MeV
- Cleaner

FIN