

Producción y uso de haces radioactivos

Master Interuniversitario de Física Nuclear

Berta Rubio IFIC (CSIC-Univ. Valencia)

B. Rubio. Master FN, Valencia 2024

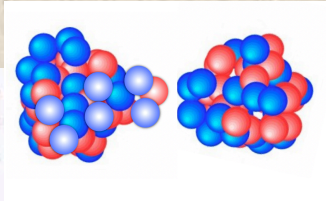
Porqué la producción de haces radioactivos ha marcado un antes y un después en la Física Nuclear

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.



Search for Super Heavy elements $Z > 100$

1																	18							
1																	2							
H																	He							
3	4															10	11	12	13	14	15	16	17	18
Li	Be															B	C	N	O	F	Ne			
11	12	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18							
Na	Mg	Al	Si	P	S	Cl	Ar																	
19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36							
K	Ca	Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr							
37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54							
Rb	Sr	Y	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Te	I	Xe							
55	56	57-71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86							
Cs	Ba	La-Lu	Hf	Ta	W	Re	Os	Ir	Pt	Au	Hg	Tl	Pb	Bi	Po	At	Rn							
87	88	89-103	104	105	106	107	108	109	110	111	112	113	114	115	116	117	118							
Fr	Ra	Ac-Lr	Rf	Db	Sg	Bh	Hs	Mt	Ds	Rg	Cn	Nh	Fl	Mc	Lv	Ts	Og							
57	58	59	60	61	62	63	64	65	66	67	68	69	70	71										
La	Ce	Pr	Nd	Pm	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu										
89	90	91	92	93	94	95	96	97	98	99	100	101	102	103										
Ac	Th	Pa	U	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No	Lr										

Figure 1. Periodic table of the elements 2017.

Super Heavy elements
Can only be produced at accelerators facilities

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	Darmstadtium	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)



Reactors

Fm discovered in the 50ths



Meitnerium, the only element named after a woman scientist

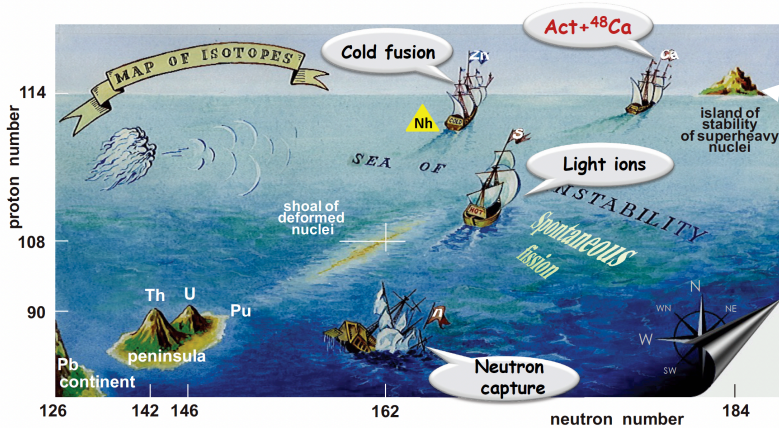


Figure 2. Voyage to the heaviest nuclei.

Will we prove the existence of the island of stability?

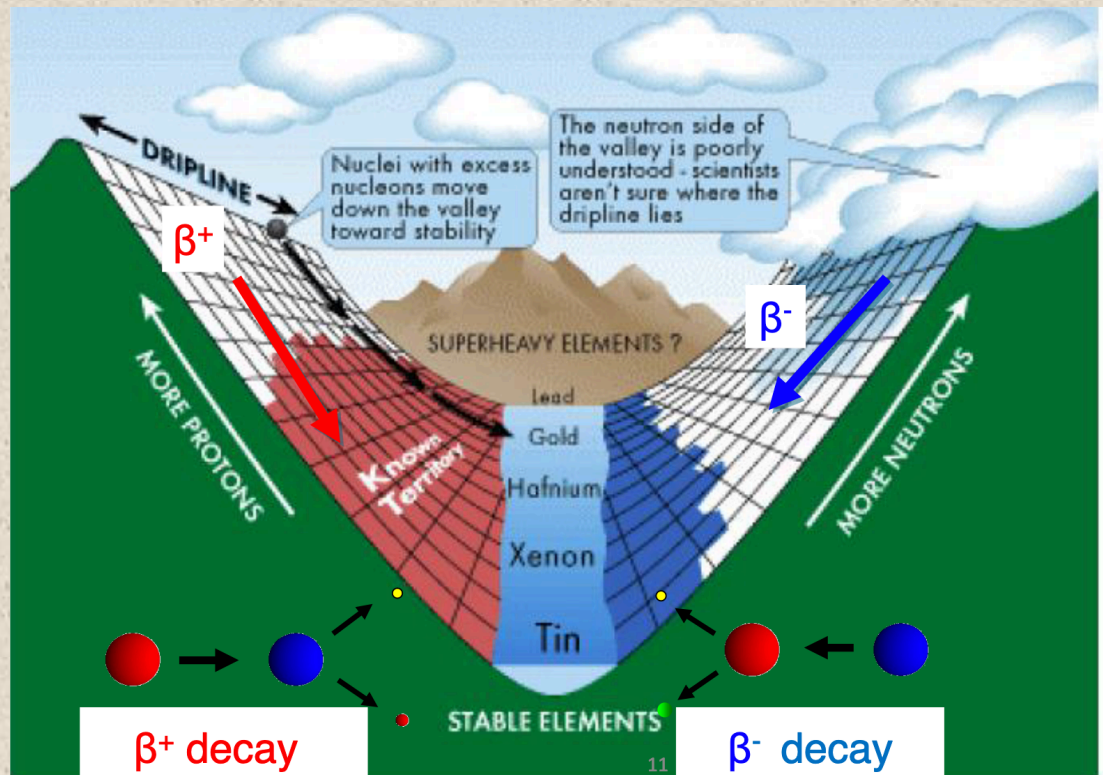
1 H	2 He																	18
3 Li	4 Be																	10 Ne
11 Na	12 Mg	13 Al	14 Si	15 P	16 S	17 Cl	18 Ar											36 Kr
19 K	20 Ca	21 Sc	22 Ti	23 V	24 Cr	25 Mn	26 Fe	27 Co	28 Ni	29 Cu	30 Zn	31 Ga	32 Ge	33 As	34 Se	35 Br	36 Kr	54 Xe
37 Rb	38 Sr	39 Y	40 Zr	41 Nb	42 Mo	43 Tc	44 Ru	45 Rh	46 Pd	47 Ag	48 Cd	49 In	50 Sn	51 Sb	52 Te	53 I	54 Xe	86 Rn
55 Cs	56 Ba	57-71 La-Lu	72 Hf	73 Ta	74 W	75 Re	76 Os	77 Ir	78 Pt	79 Au	80 Hg	81 Tl	82 Pb	83 Bi	84 Po	85 At	86 Rn	118 Og
87 Fr	88 Ra	89-103 Ac-Lr	104 Rf	105 Db	106 Sg	107 Bh	108 Hs	109 Mt	110 Ds	111 Rg	112 Cn	113 Nh	114 Fl	115 Mc	116 Lv	117 Ts	118 Og	
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

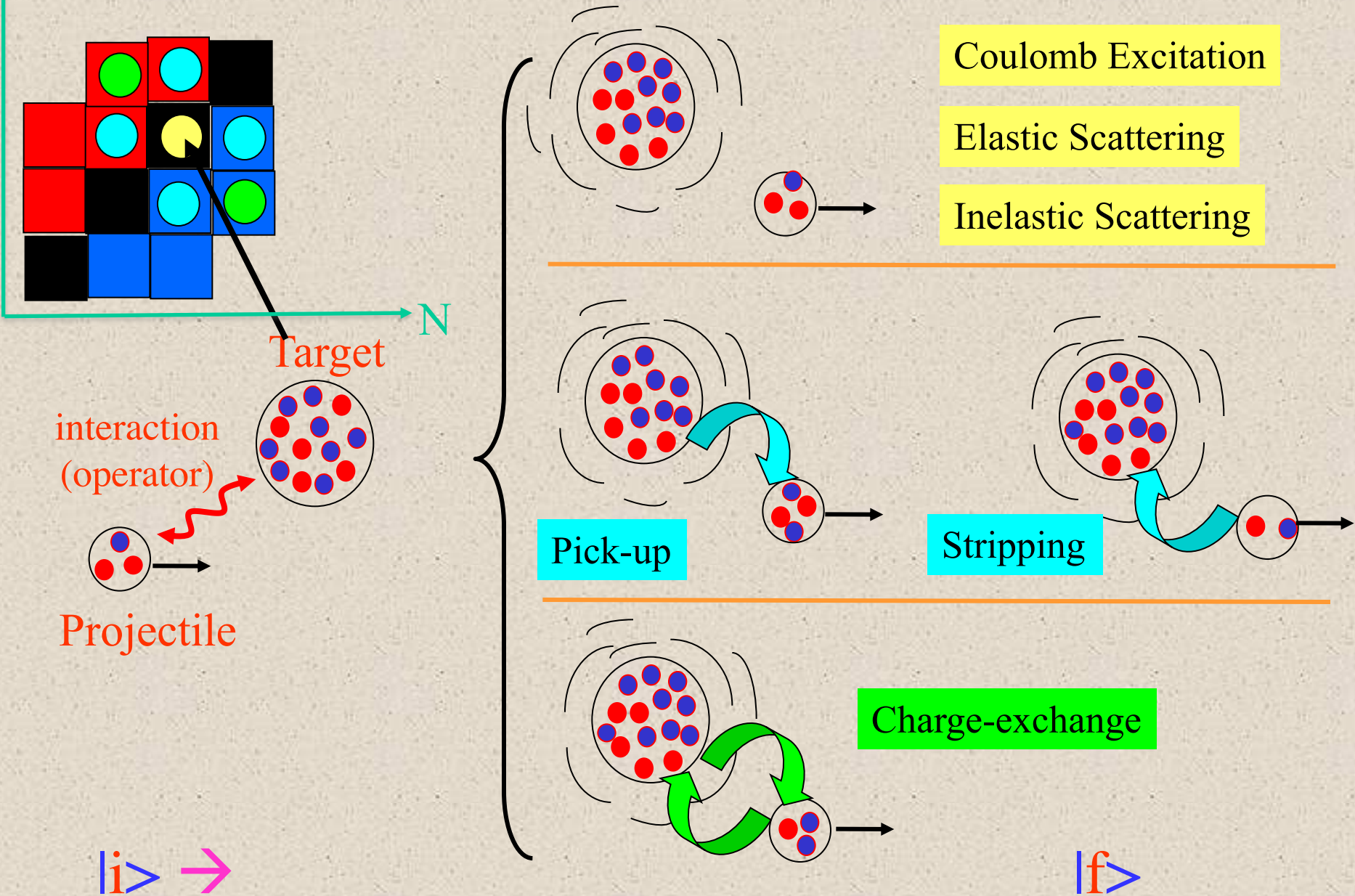
Does period number 8 exists?

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

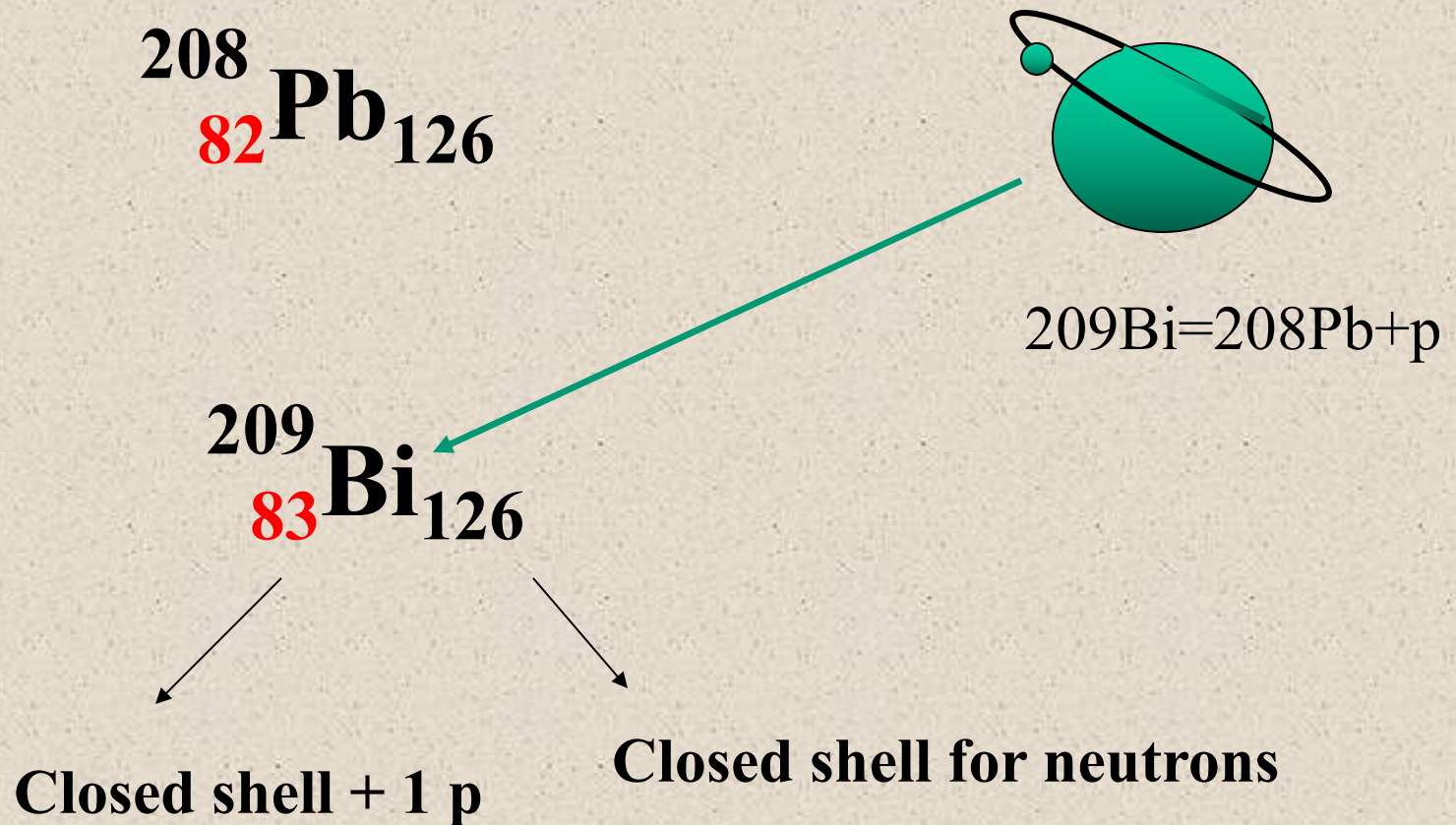


Let us see an example

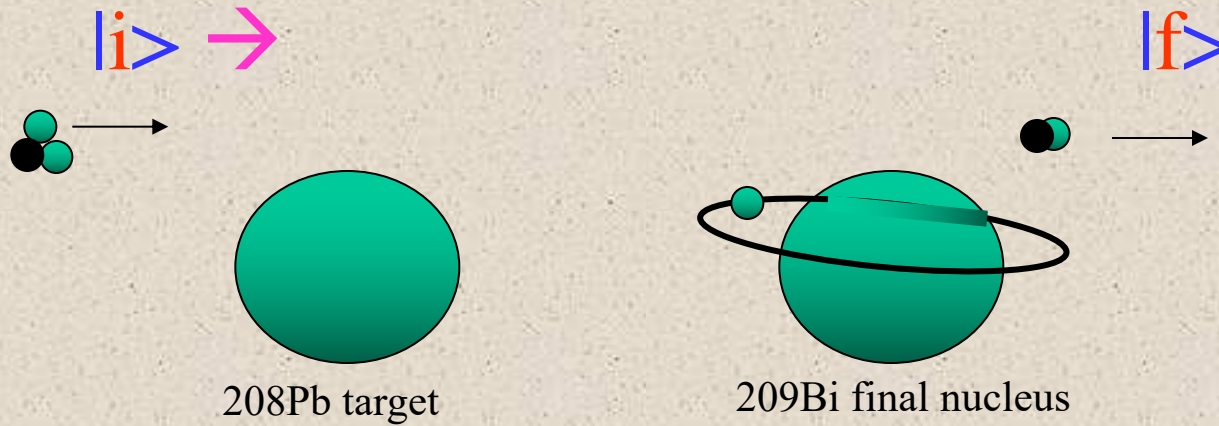
What can we do if we have a stable target : we can perform nuclear reactions, for instance with light nuclei



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



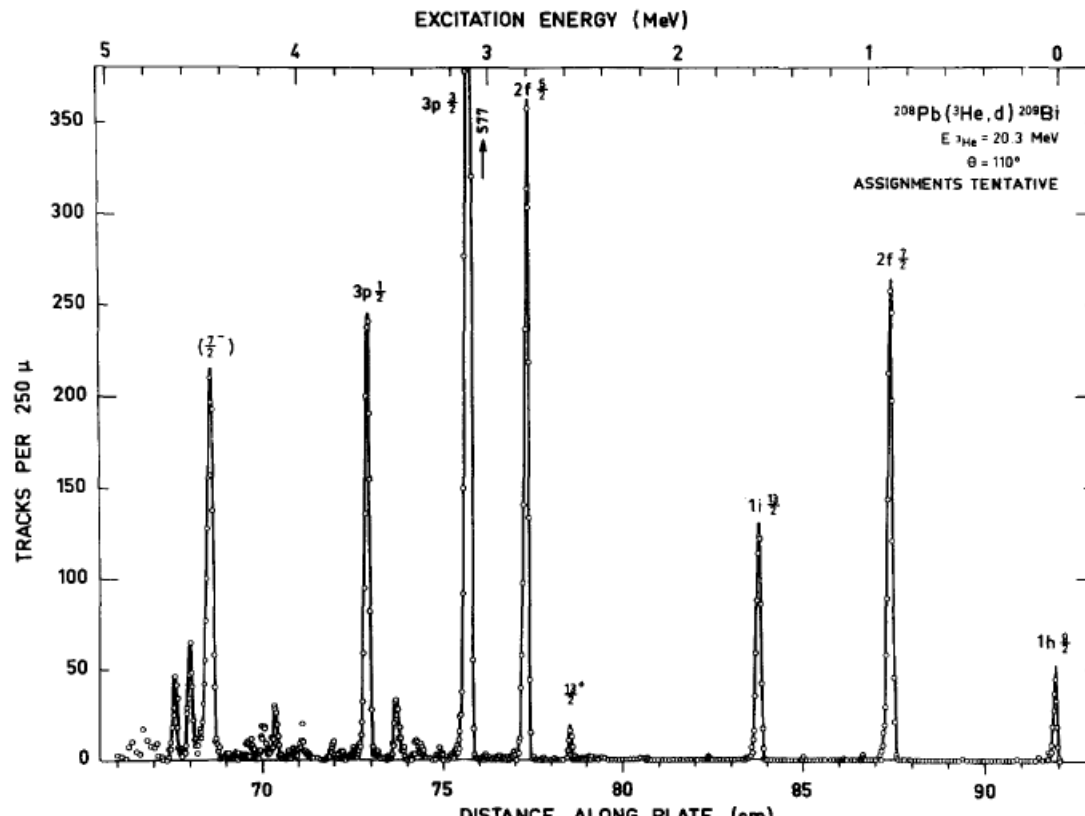
Reaction: $^{208}\text{Pb}(^3\text{He},d)^{209}\text{Bi}$



Volume 26B, number 3

PHYSICS LETTERS

8 January 1968



Shell Model and Residual Interactions

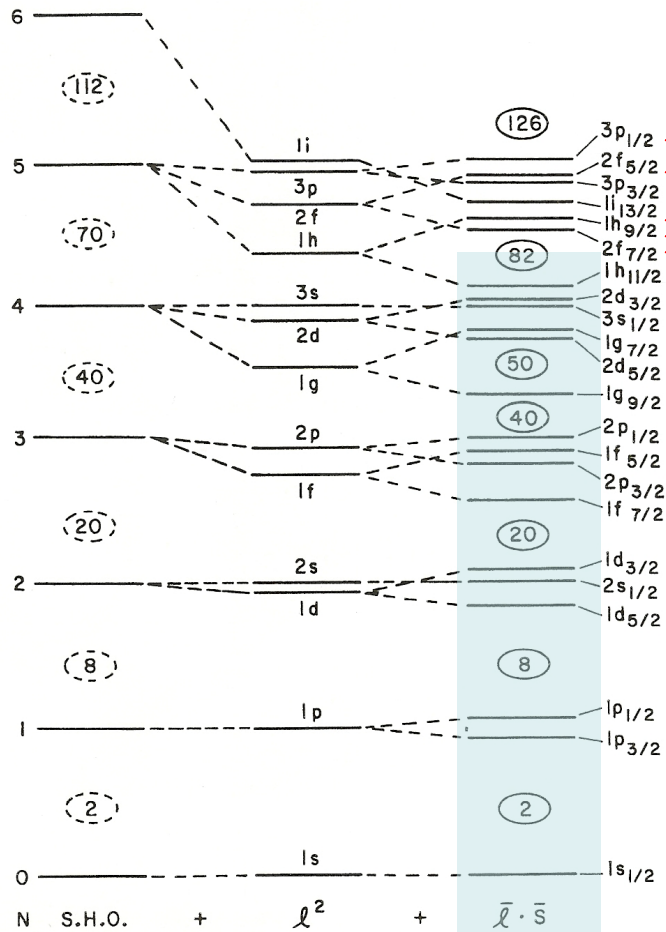
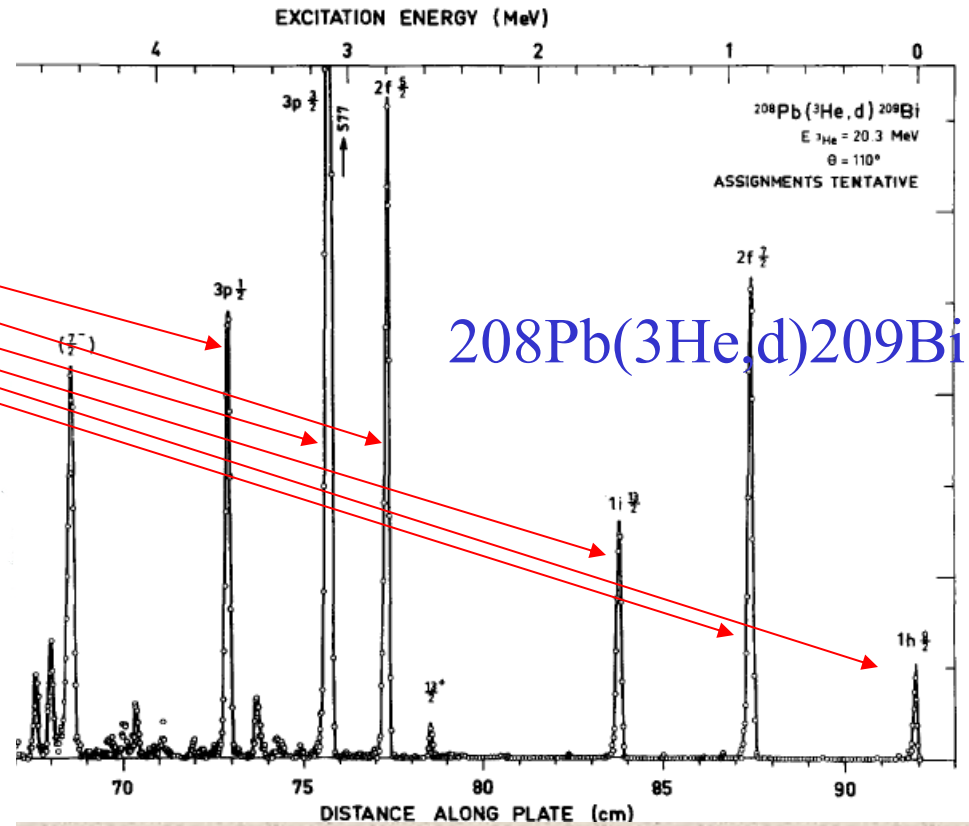
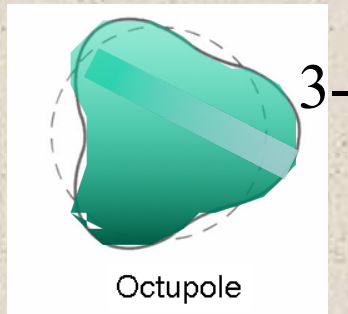


Fig. 3.2. Single-particle energies for a simple harmonic oscillator (S.H.O.), a modified harmonic oscillator with l^2 term, and a realistic shell model potential with l^2 and spin orbit ($l \cdot s$) terms.

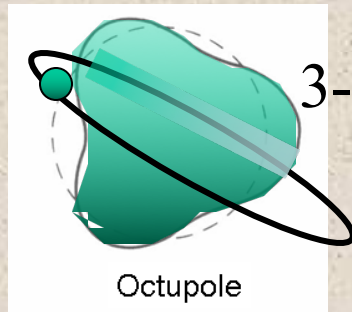


$$^{209}\text{Bi} = ^{208}\text{Pb} + p$$

208Po



$h_{9/2}$



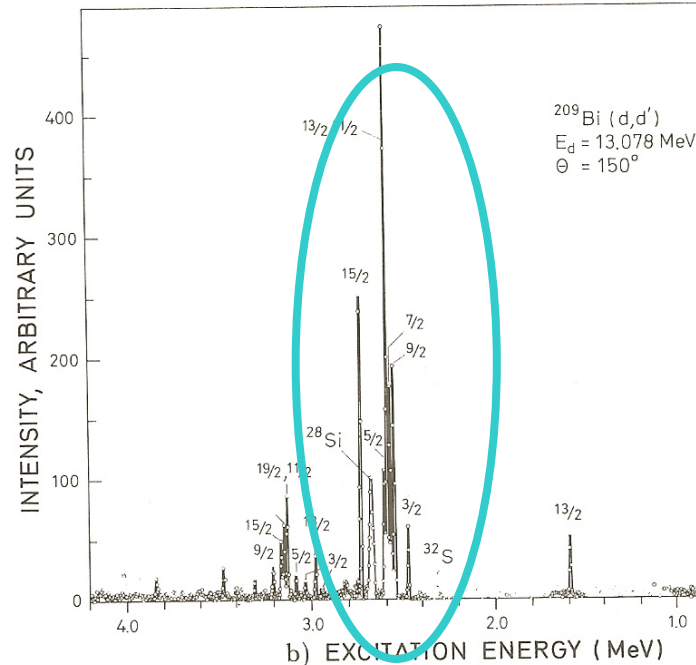
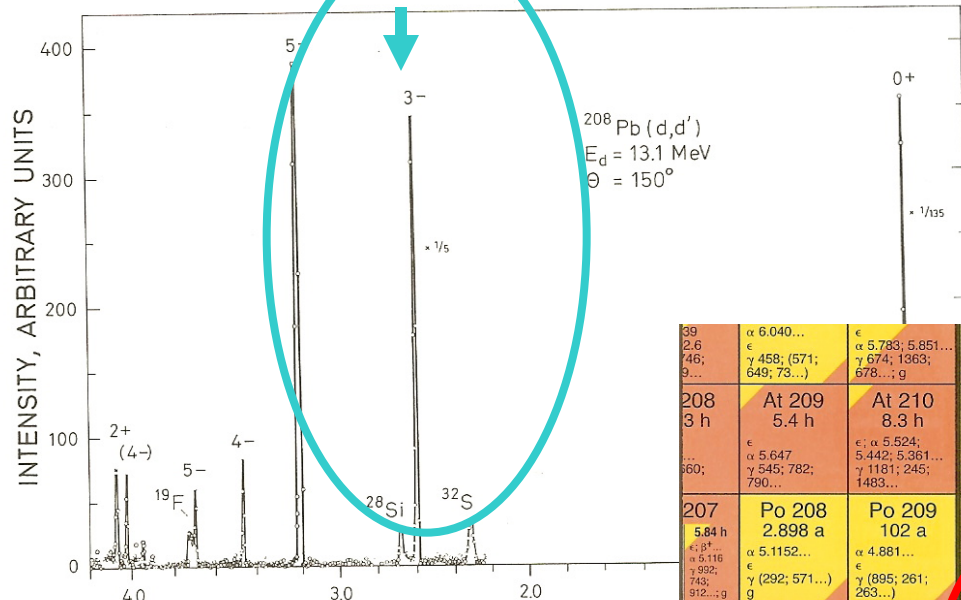
$209\text{Bi} = 208\text{Pb} + p$

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

572



VIBRATIONAL SPECTRA Ch. 6



39 2.6 46; 9...	α 6.040... ϵ 5.783; 5.851... γ 458; 571; 649; 73...	ϵ 5.783; 5.851... γ 474; 1363; 678...; g	α 6.264... γ	α 8.088; 7.252... γ 540...	ly 16... m 10...
208 3 h	At 209 5.4 h	At 210 8.3 h	At 211 7.22 h	At 212 119 ms 314 ms	α
ϵ 5.647 γ 545; 782; 790...	ϵ 5.524; 5.442; 5.361... α 5.867... γ (687...) g	ϵ 5.524; 5.442; 5.361... α 5.867... γ (687...) g	ϵ 5.524; 5.442; 5.361... α 5.867... γ (687...) g	α 7.84; 7.90... 7.62... γ 63... g	α
207 5.84 h	Po 208 2.898 a	Po 209 102 a	Po 210 138.38 d	Po 211 25.2 s 0.516 s	451... 11.6... 26... 383... ly
ϵ 5.116... α 5.1152... γ 992; 745; 912...; g	ϵ 4.881... α 4.881... γ (292; 571...) g	ϵ 4.881... α 4.881... γ (292; 571...) g	ϵ 4.881... α 4.881... γ (292; 571...) g	α 7.450... γ (896; 570...) ly	α
206 4 d	Bi 207 31.55 a	Bi 208 3.68 · 10 ⁵ a	Bi 209 100	Bi 210 5.013 d	α 11.6... β 11.6... γ 11.6... α 11.6... β 11.6... γ 11.6...
ϵ 5.116... α 5.1152... γ 992; 745; 912...; g	ϵ 4.881... α 4.881... γ (292; 571...) g	ϵ 4.881... α 4.881... γ (292; 571...) g	ϵ 4.881... α 4.881... γ (292; 571...) g	α 7.450... γ (896; 570...) ly	α
205 10 ⁷ a	Pb 206 24.1	Pb 207 22.1	Pb 208 52.4	Pb 209 3.253 h	β 1.8... γ 1.8... α 1.8... β 1.8... γ 1.8...
σ 0.027	σ 0.61	σ 0.00023 σ_n α < 3E-7	σ 0.00023 σ_n α < 3E-7	β 0.6 no γ	σ
204 8 a	Tl 205 70.48	Tl 206 3.7 m 4.20 m	Tl 207 1.3 s 4.7 m	Tl 208 3.053 m	β 1.8... γ 1.8... α 1.8... β 1.8... γ 1.8...
ϵ	σ 0.11	β 1.5... γ (803...)	β 1.4... γ (898...)	β 1.8... γ 1.8... α 1.8... β 1.8... γ 1.8...	β
203 59 d	Hg 204 6.87	Hg 205 5.2 m	Hg 206 8.15 m	Hg 207 2.9 m	β 1.8... γ 1.8... α 1.8... β 1.8... γ 1.8...
σ 0.4	β 1.5... γ 204...	β 1.5... γ 305; 650... g	β 1.5... γ 305; 650... g	β 1.8... γ 1.8... α 1.8... β 1.8... γ 1.8...	β
202 3 s	Au 203 60 s	Au 204 39.8 s	Au 205 31 s		
β 2.0... γ 218; 44; 51; 318; 369	β 2.0... γ 218; 44; 51; 318; 369	β 2.0... γ 218; 44; 51; 318; 369	β 2.0... γ 218; 44; 51; 318; 369		
201 5 m	Pt 202 ~43.6 h				
β	β				

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

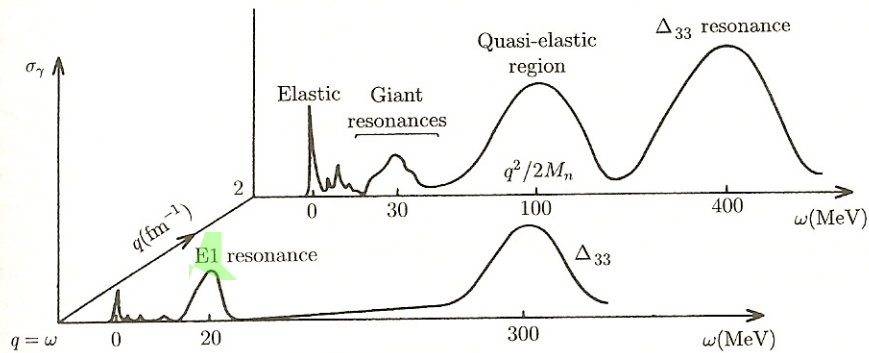
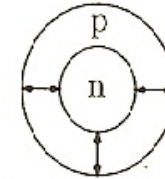


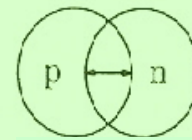
FIG. 1.1. Characteristic response (cross section σ) of an atomic nucleus as a function of the energy transfer ω and momentum transfer q . The lower curve is for photon absorption, the upper one for particle scattering with $q = 2 \text{ fm}^{-1} \neq \omega$.

Isovector

$\Delta T=0 \ 1$

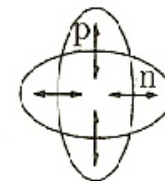


IVGMR



IVGDR

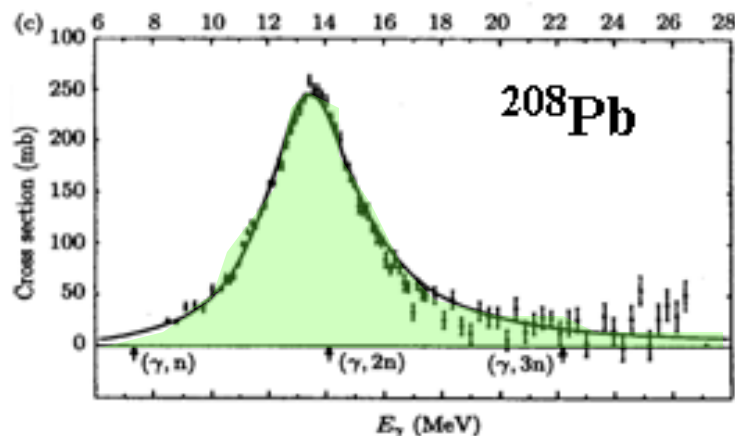
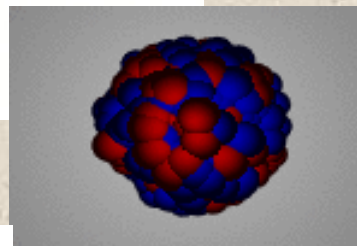
E1



IVGQR

$\Delta T = 1$

$\Delta S = 0$



Qué ha sido necesario para poder hacer estos experimentos

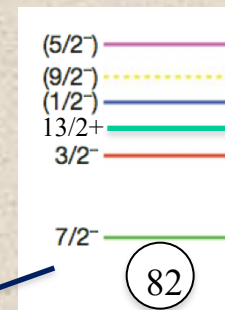
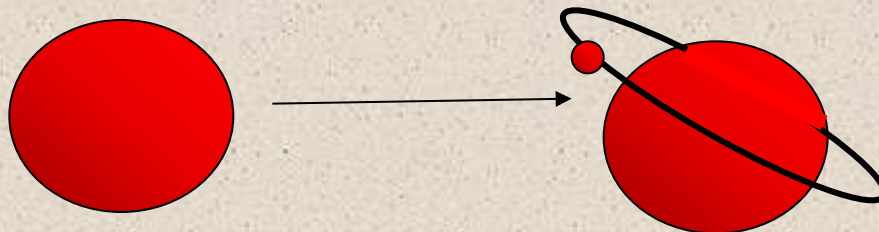
1. Definir la reacción nuclear de interés.
2. Blancos estables: ^{208}Pb y ^{209}Bi en este caso
2. (Un proyectil “sin estructura”)
3. (Un proyectil ligero (struggling))

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 4.1 m β^- 3.7... γ 974, 697 151, 104...	Sb 133 2.34 m β^- 1.2, 2.9... γ 1096, 818 2755, 837... g, m	Sb 134 10.1 s β^- 6.1... γ 1279 297, 707 115... 1352...	Sb 135 1.7 s β^- 8.1... βn 1.45, 1.04... γ 1127, 1279* 1380...
Sn 131 50 s β^- 3.4... γ 1226 480, 305 1229...	Sn 132 39.7 s β^- 1.8... γ 341, 86, 899 247, 993...	Sn 133 1.46 s β^- 8.1... γ 962... βn 0.48 m	Sn 134 1.050 s β^- 7.0... βn 0.72, 318, 554 962*...
In 130 0.54 s β^- 1221 174... g, m βn	In 131 0.32 s β^- 6.6... γ 1905 130 391... g, m βn	In 132 207 ms β^- 6.4, 8.9... γ 375, 4041 299... βn 0.23, 0.33	In 133 180 ms β^- 802, 854 IT? βn 1.3 βn 1561 854 2005)
Cd 129 242 ms β^- 995, 359, 1796... βn	Cd 130 162 ms β^- 6.2, 8.3... γ 1669, 451 950, 1170... βn	Cd 131 98.0 ms β^- 957 βn 7, βn 2	
Ag 128 59 ms β^- 645, 784... βn ?	Ag 129 ~160 ms β^- 957 βn ?	Ag 130 42 ms β^- 957 βn ?	
Pd 127 38 ms β^- 957 βn ?	Pd 128 35 ms β^- 957 βn ?	Pd 129 31 ms β^- 957 βn ?	

$^{133}_{50}\text{Sn}_{83}^{-1}$

From ENSDF - Evaluated October 2010

^{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 3; $Q(\beta^-)=12917$ SY; % β^- decay=100.0

1995JoZZ, 1996Ho16, 2000Ho32: ^{133}In β^- [from ^{238}U (p,f), E=1 GeV]; measured E γ , I γ , $\gamma\gamma$ -, $n\gamma$ -, $\beta\gamma$ -c (βn) γ (t). ISOLDE facility; plastic scin β detector, two liquid scin n detectors, two Ge γ -ray detectors

^{133}Sn Levels

E(level) [†]	J^π [‡]
0.0	$7/2^-$
853.7 3	$3/2^-$
1560.9 5	$(9/2^-)$
2004.6 10	$(5/2^-)$

$1/2^-$ - missing



nature

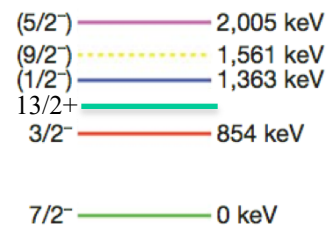
Vol 465 | 27 May 2010 | doi:10.1038/nature09048

LETTERS

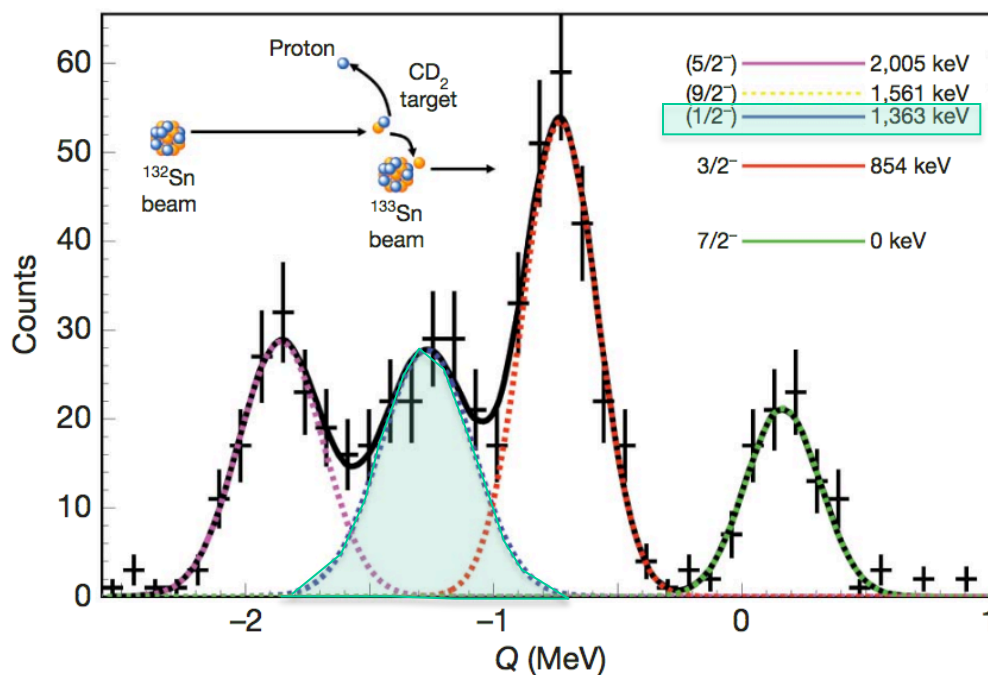
Year 2010

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. Livesay⁵, Z. Ma¹, B. H. Moazen¹, C. D. Nesaraja⁴, F. M. Nunes⁸, S. D. Pain², N. P. Patterson⁹, D. Shapira¹, J. F. Shriner Jr¹, M. S. Smith⁴, T. P. Swan^{2,6} & J. S. Thomas⁶



$^{132}\text{Sn}(d,p)^{133}\text{Sn}$



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

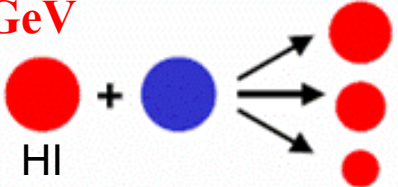
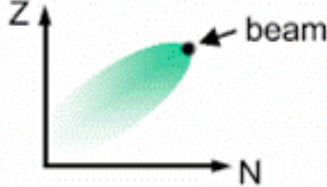
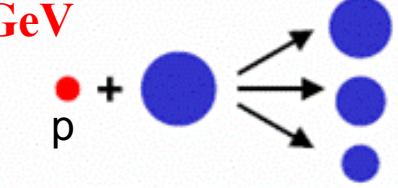
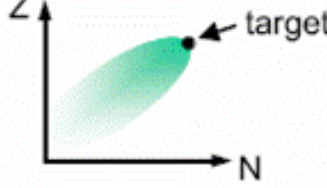
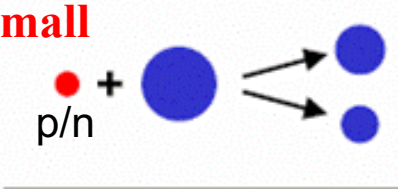
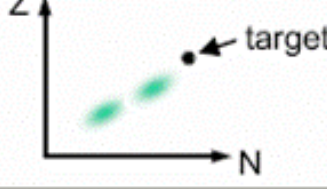
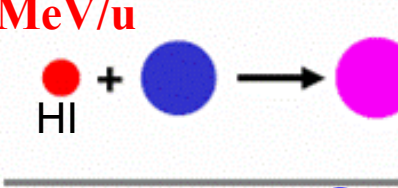
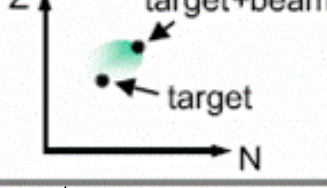
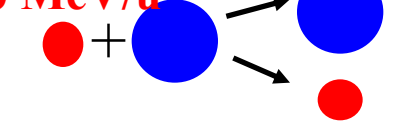
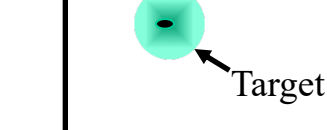
Postacceleration

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

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

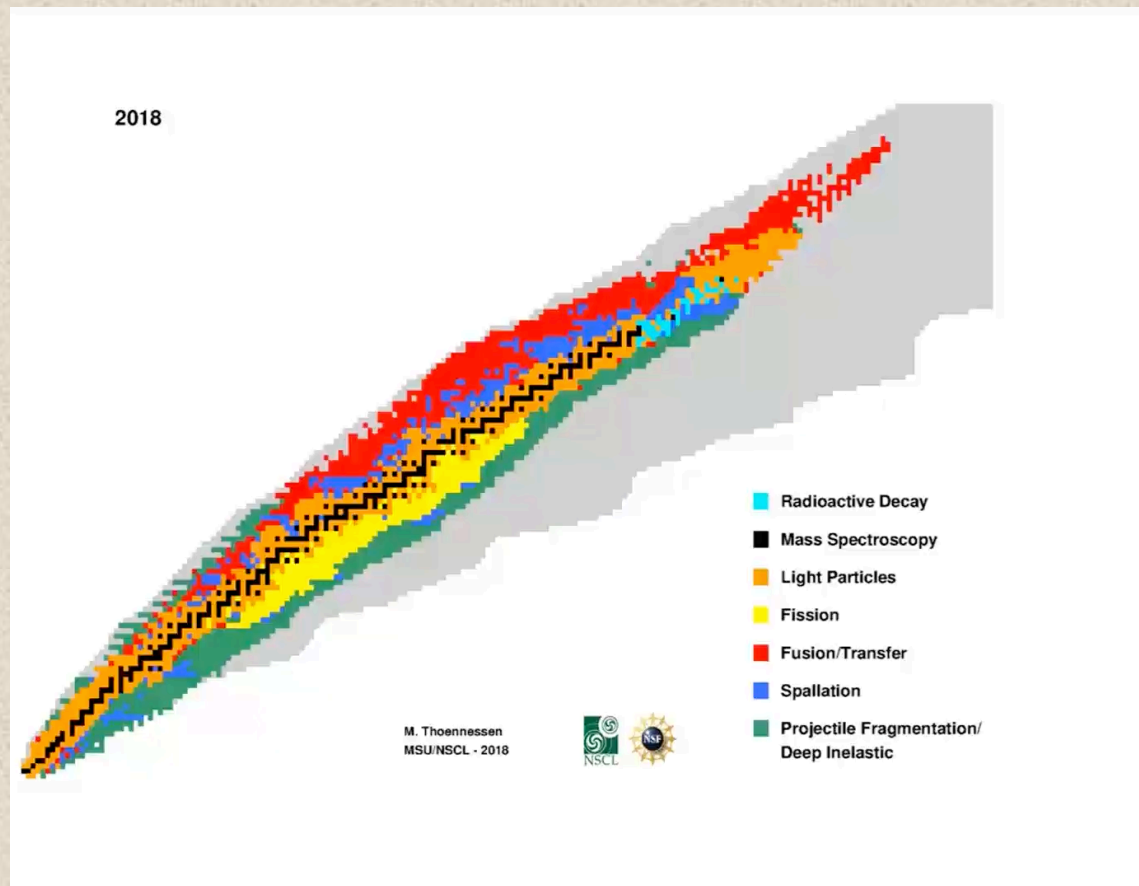
The first thing to worry about is how to produce radioactive nuclei.
Tool: Reaction or Fission

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

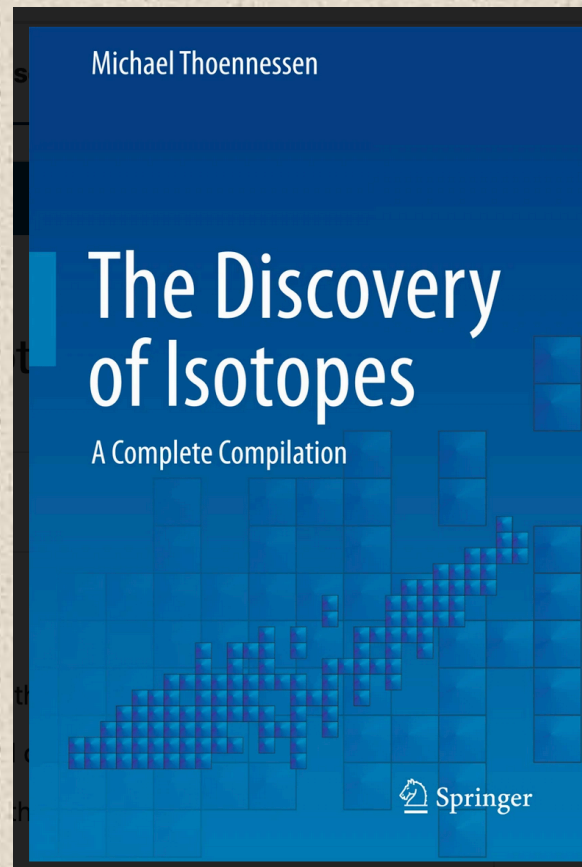
Beam + target → products	Product energy	Number of Products
GeV  HI	“All nuclei” fragmentation $v_{\text{product}} = v_{\text{beam}}$ RIKEN, GSI, MSU, FAIR, FRIB	up to 1000 
GeV  p	“All nuclei” spallation few MeV/u ISOLDE	up to 1000 
small  p/n	“mainly neutron rich” fission ~1 MeV/u Reactors, ISOLDE...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

Historical view to the discovery of new isotopes

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



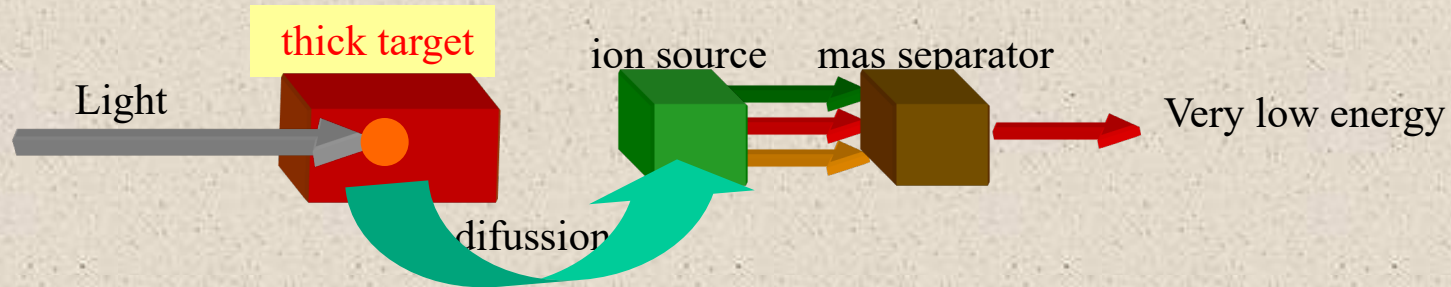
I you want to learn how all that happened, read this book



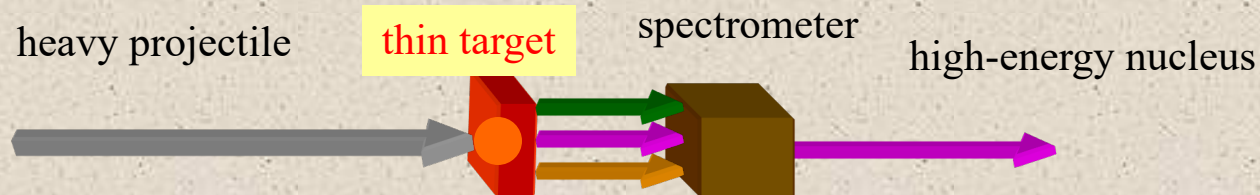
B. Rubio. Master FN, Valencia 2022

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

Isotopic Separation On-Line (ISOL)

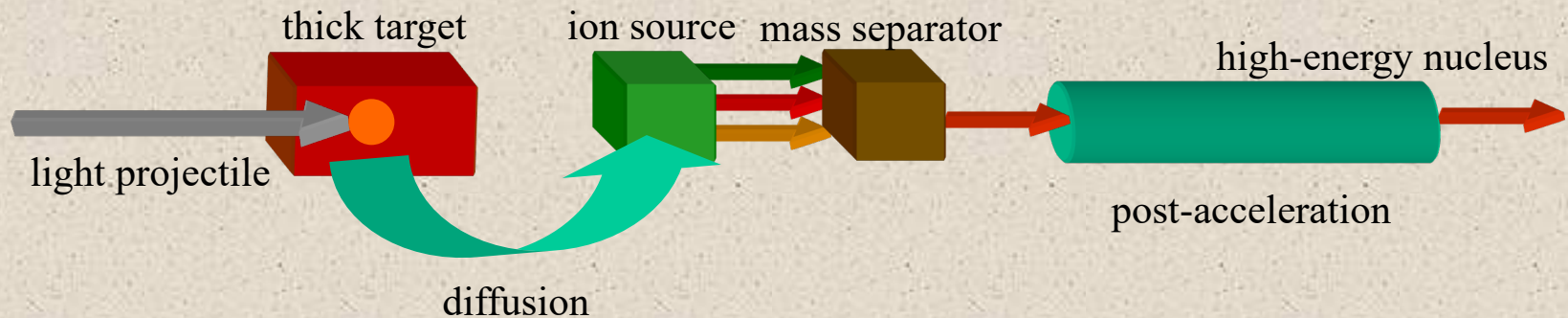


In-flight fragmentation



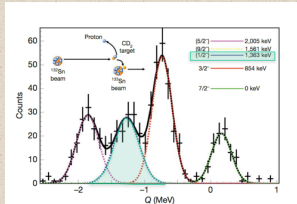
Production techniques Rad Beams: post acceleration

➤ Isotopic separation on-line (ISOL)

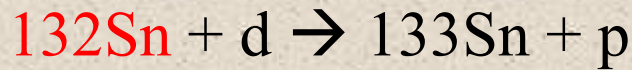




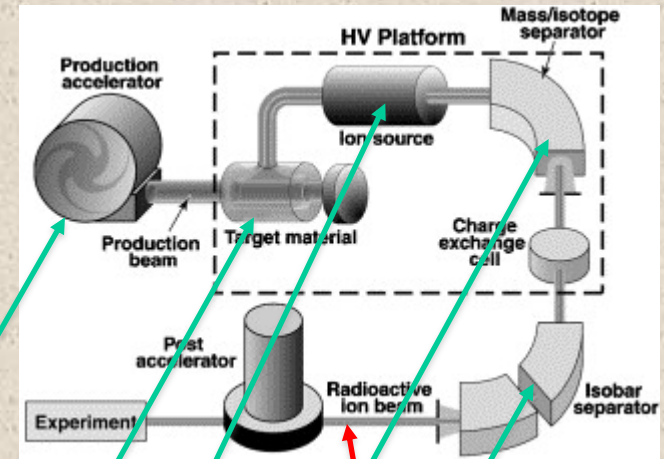
LETTERS
The magic nature of ^{132}Sn explored through the single-particle states of ^{132}Sn



5/2+ 2,000 keV
3/2+ 1,581 keV
1/2+ 1,363 keV
3/2+ 854 keV
7/2+ 0 keV



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}Sn Mass separated
5. ^{132}Sn separated
6. ^{132}Sn 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

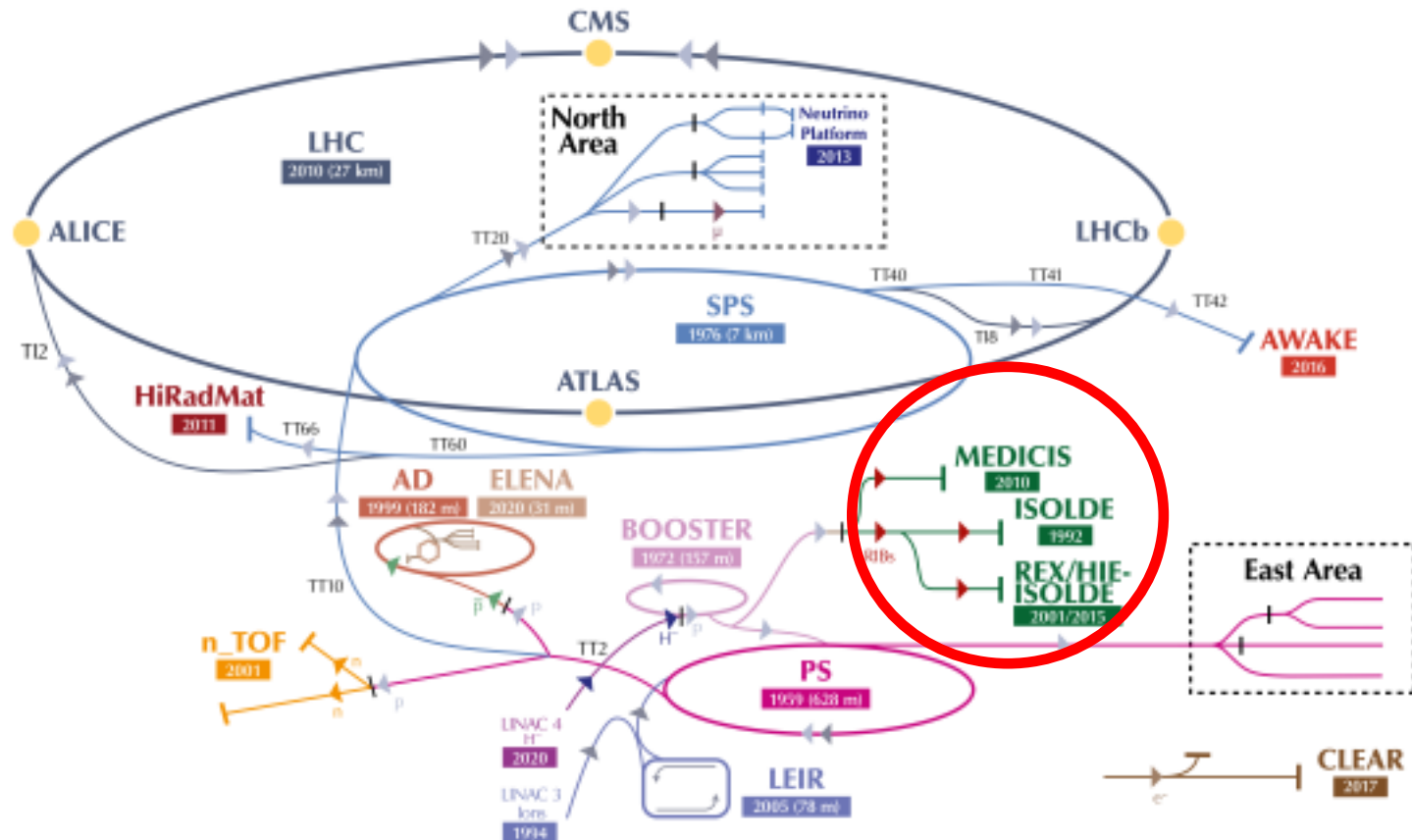
Juha Äystö



B. Rubio. Master FN

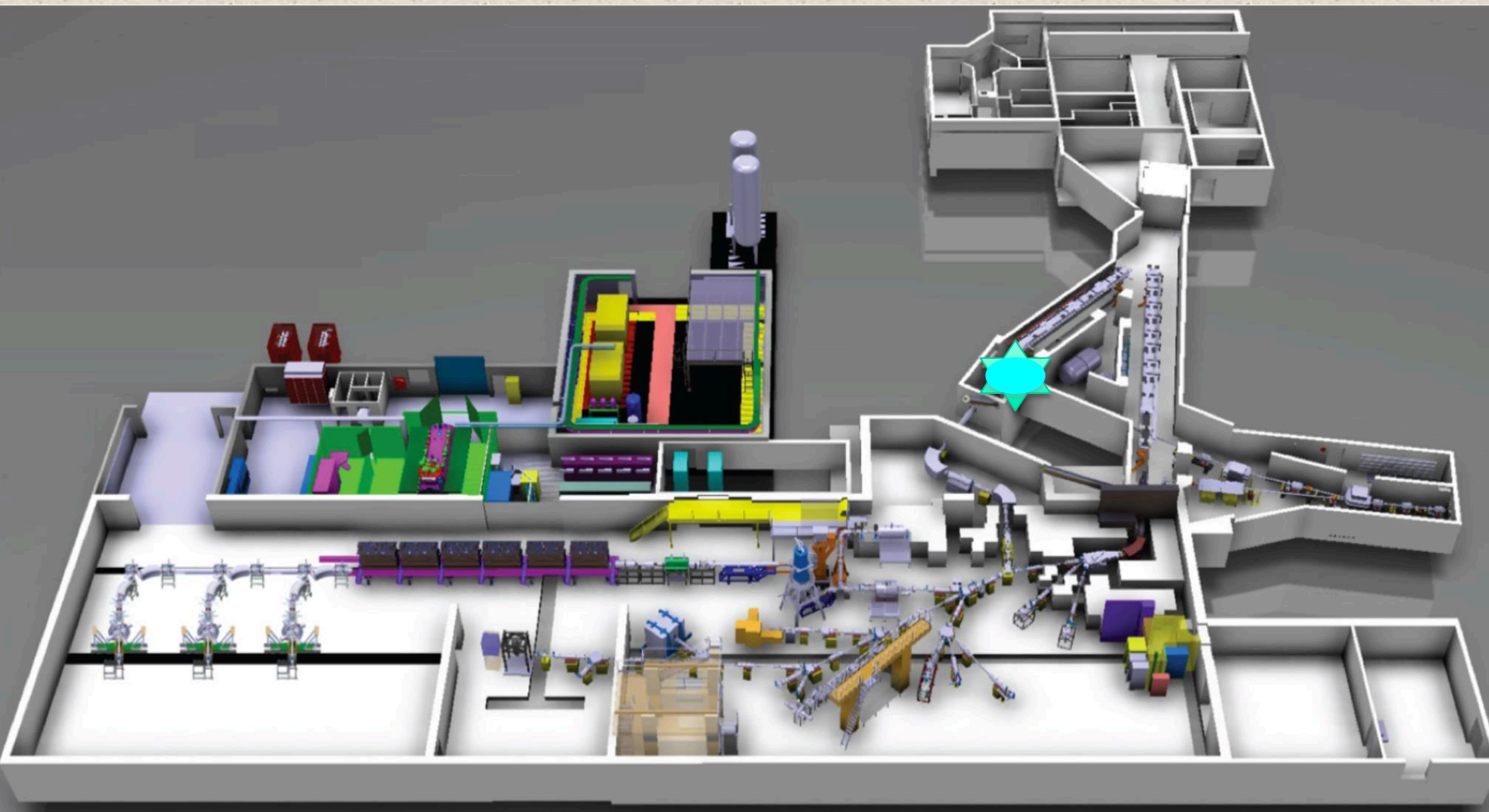
The CERN accelerator complex

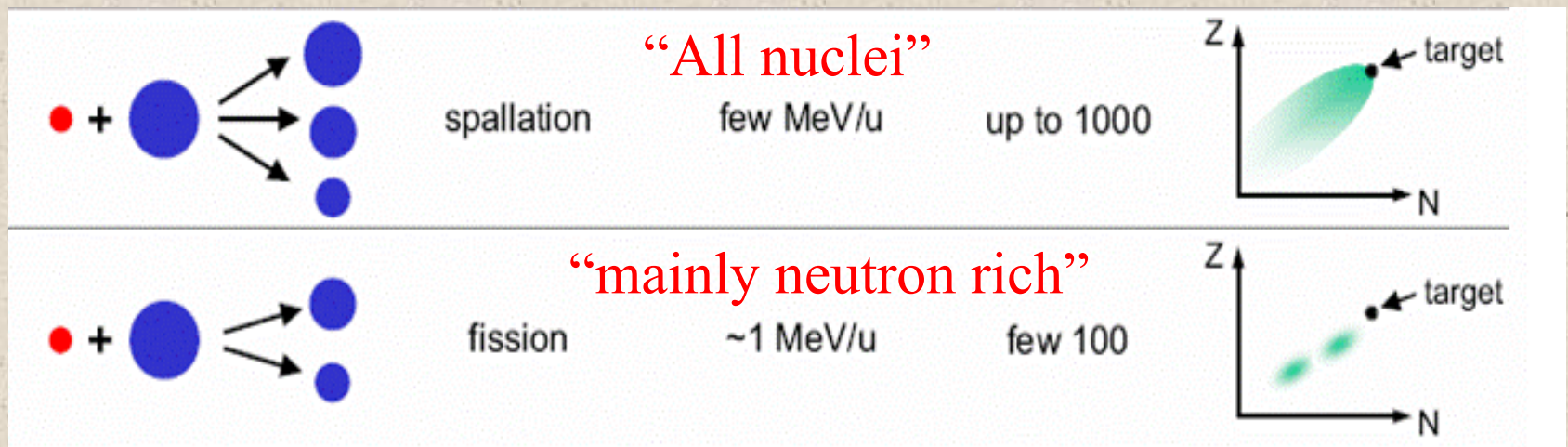
Complexe des accélérateurs du CERN



▶ 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

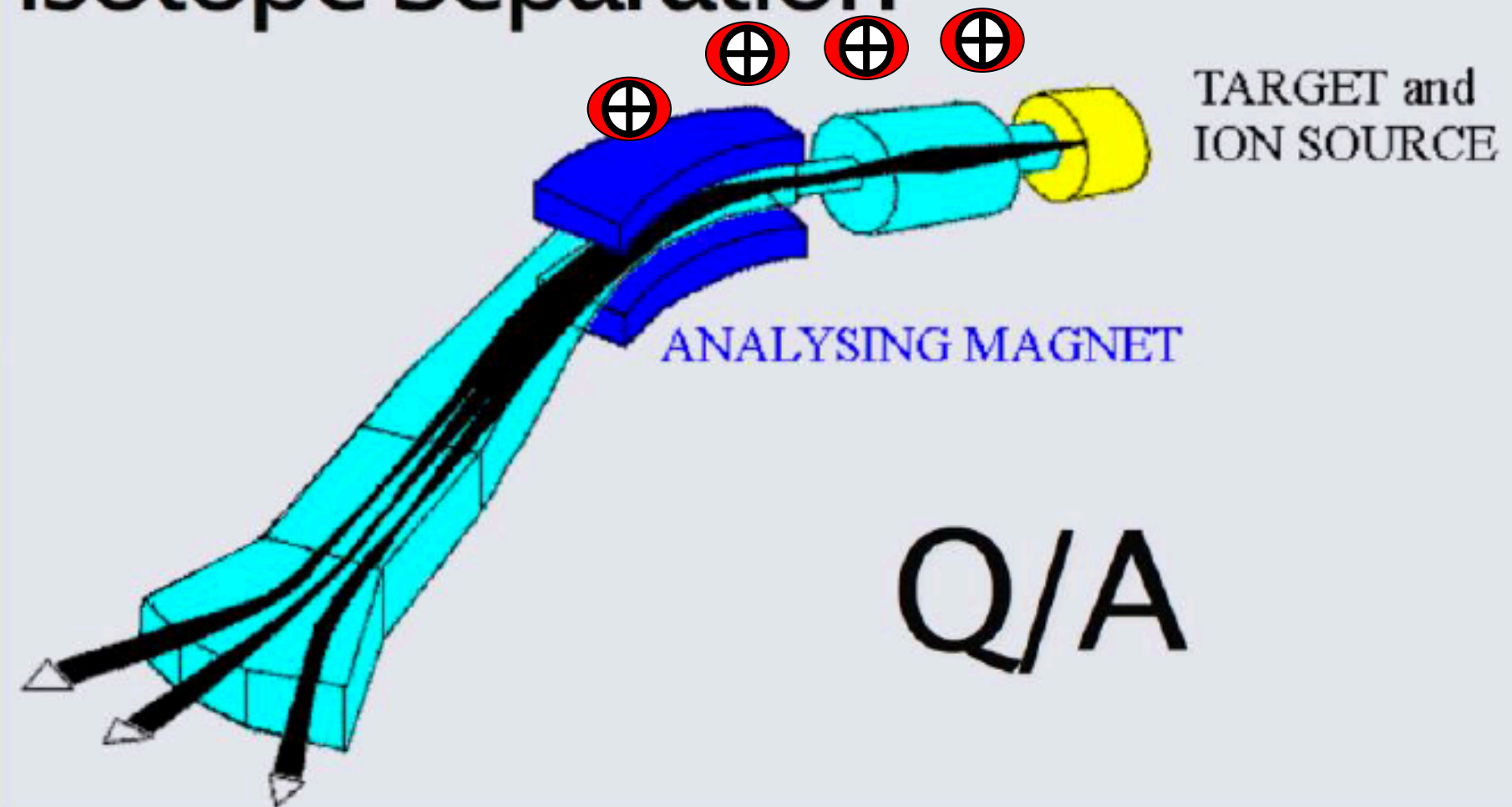




- 😊 High intensity p beam (upto 10^{16} s^{-1}) the same beam that serves LHC)
- 😊 thick target (100% energy range of p)
- 😞 long extraction and ionization time (hundred's ms)
- 😞 Chemistry dependent

The Separator: in general, only
isobaric separation

Isotope Separation

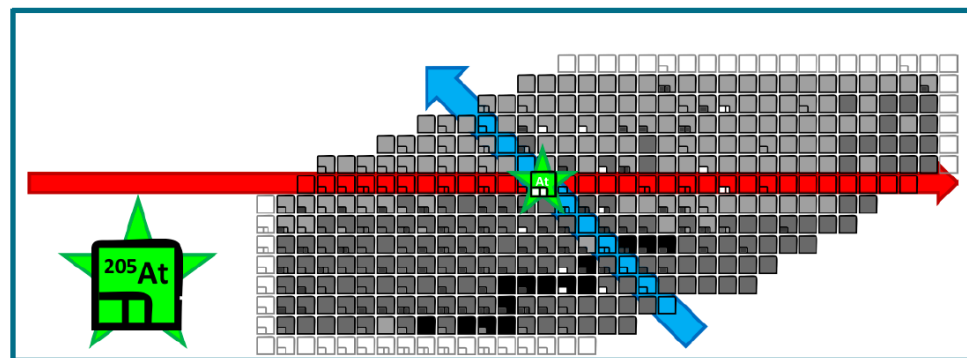
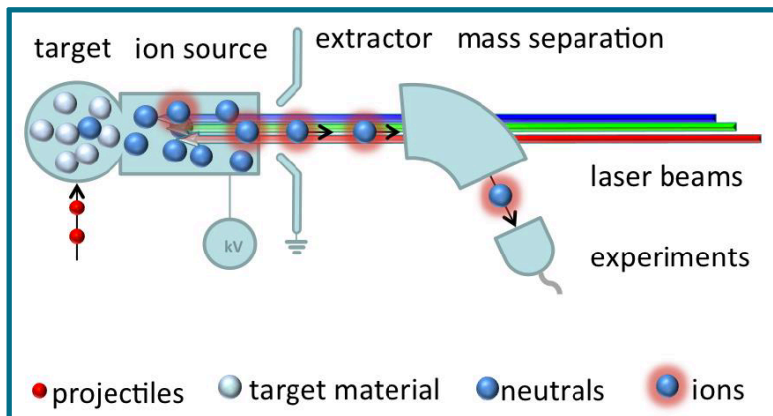
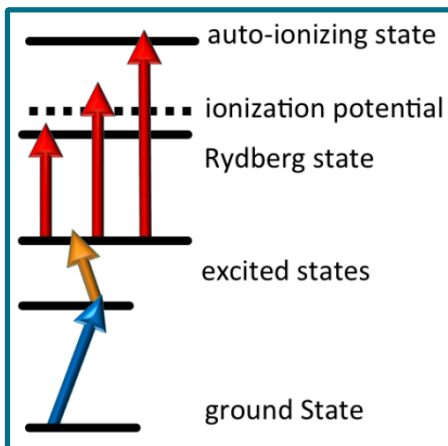




Ion sources

Laser ion source

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



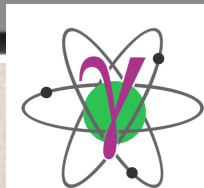
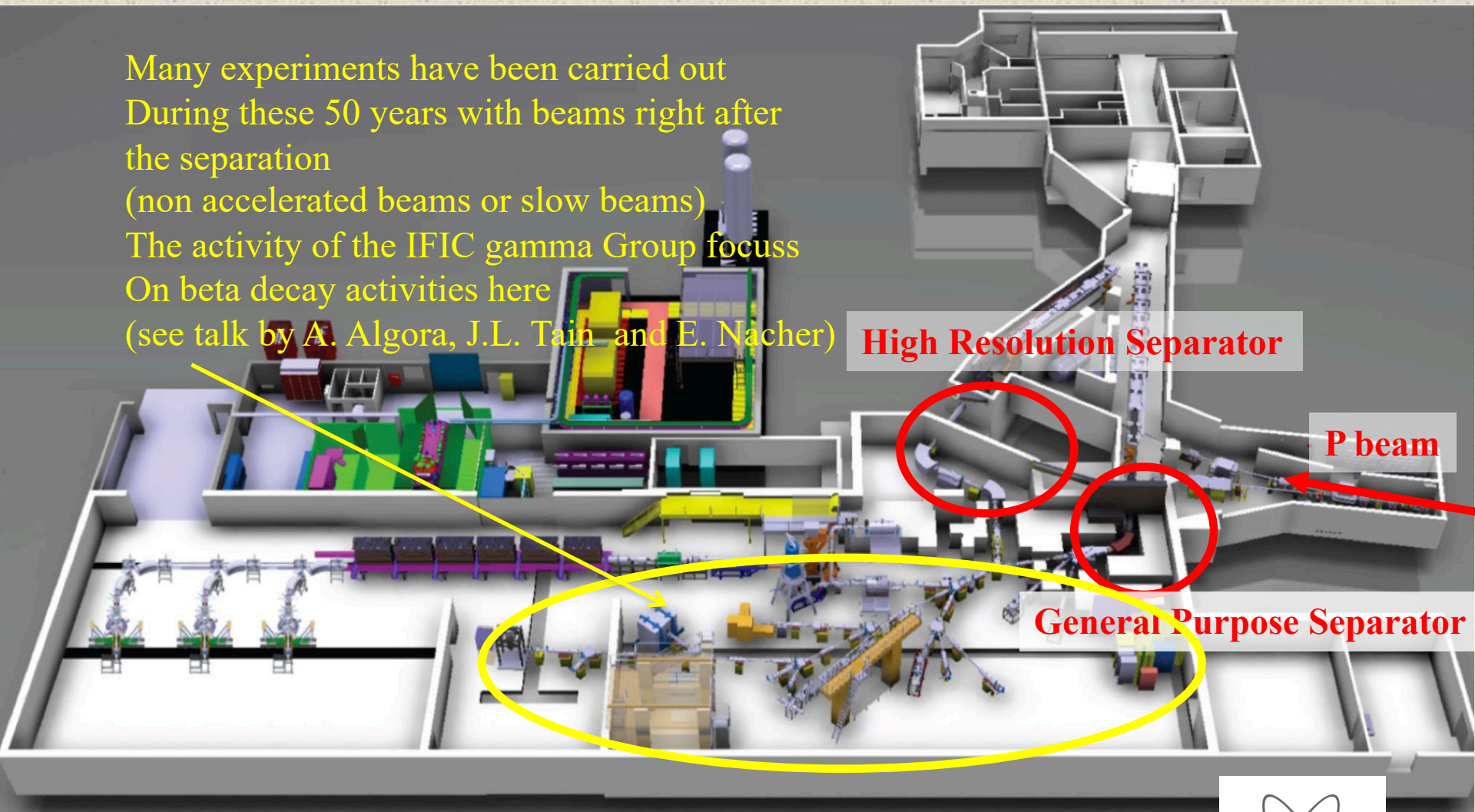
Isotopes produced at ISOLDE

H																	He						
Li	Be																	B	C	N	O	F	Ne
Na	Mg																	Al	Si	P	S	Cl	Ar
K	Ca	Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr						
Rb	Sr	Y	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Te	I	Xe						
Cs	Ba	La	Hf	Ta	W	Re	Os	Ir	Pt	Au	Hg	Tl	Pb	Bi	Po	At	Rn						
Fr	Ra	Ac	Rf	Db	Sg	Bh	Hs	Mt	Ds	Rg	112	113	114	115									
			Ce	Pr	Nd	Pm	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu							
			Th	Pa	U	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No	Lr							

ION SOURCE

+	SURFACE	−
hot	PLASMA	cooled
LASER		

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. Algora, J.L. Tain and E. Nacher)



ISOLDE

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



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 



ISOLDE

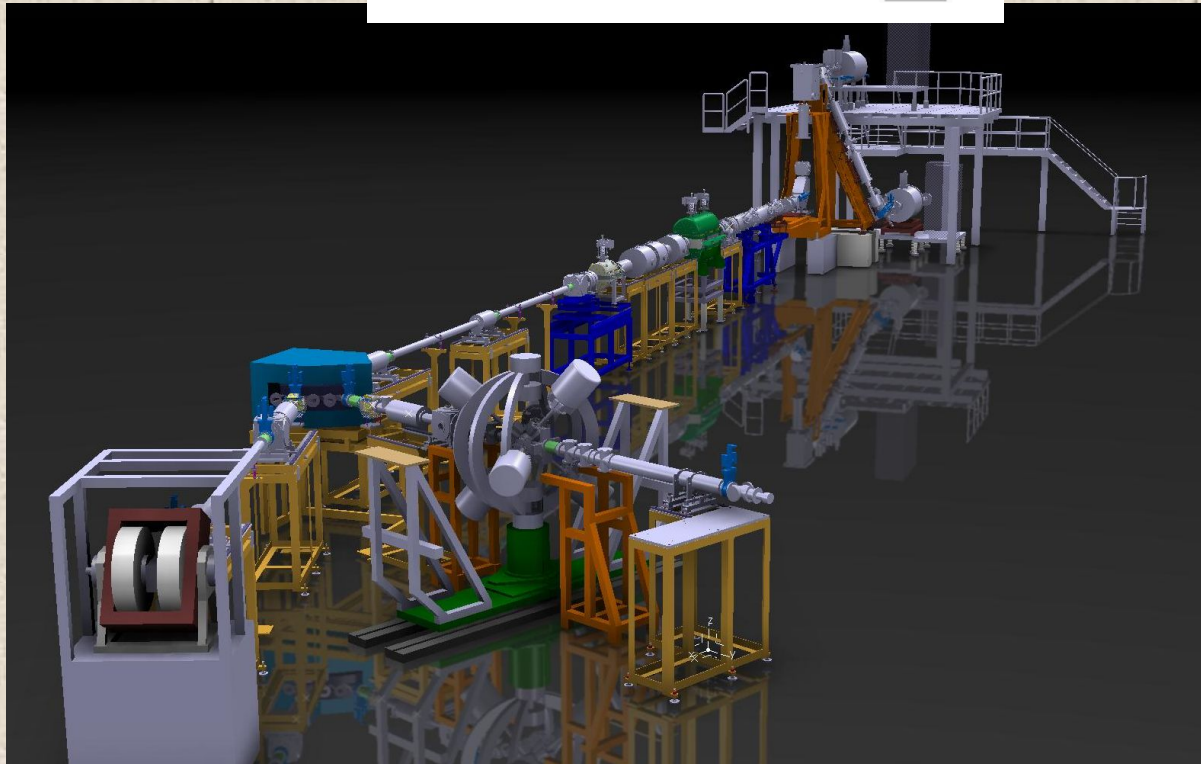
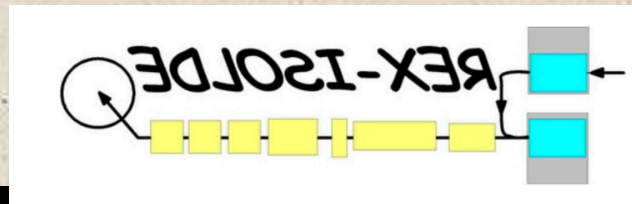
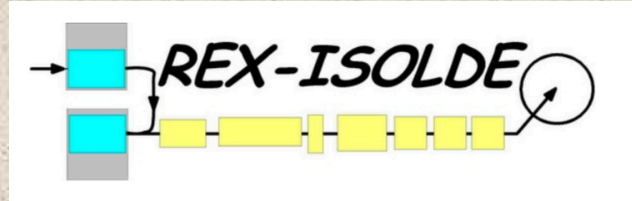
The On-Line Isotope Mass Separator ISOLDE is a facility dedicated to the production of a large variety of radioactive ion beams

22 OCTOBER 2015

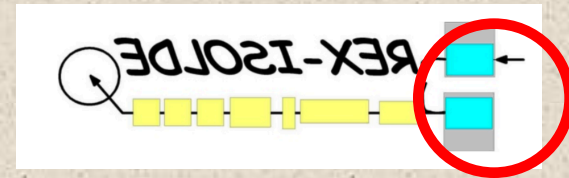
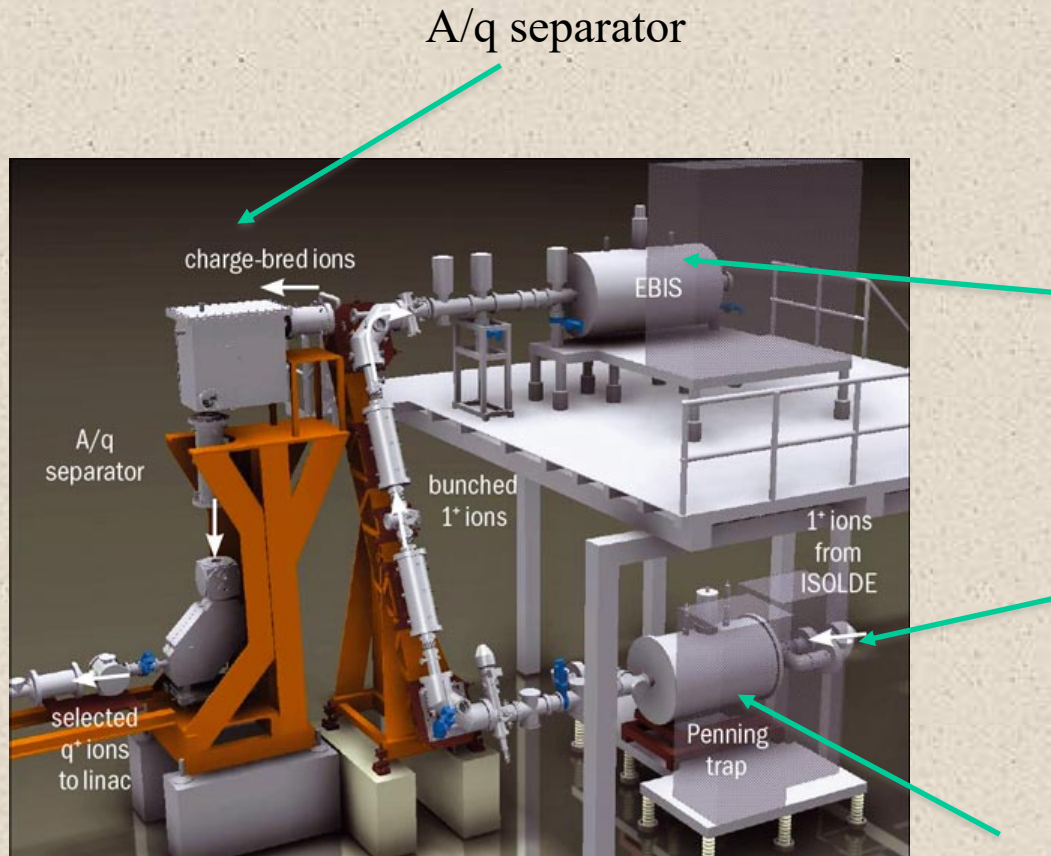
First radioactive isotope beam accelerated in HIE ISOLDE

View 





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

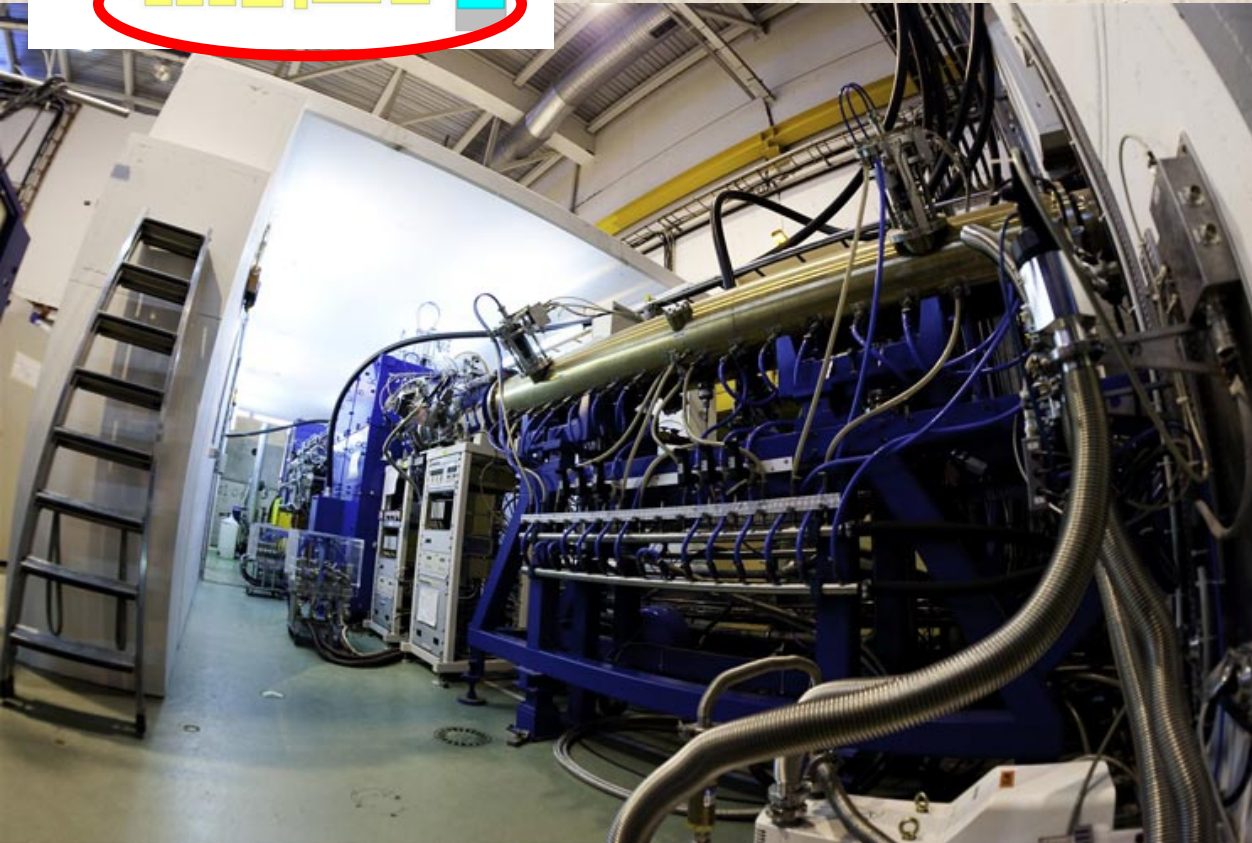
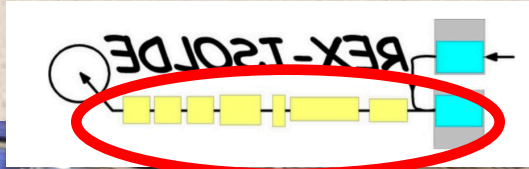


In the EBIS the 1⁺ to n⁺ ion conversion takes place

Beams from ISOLDE with 1⁺ charge state

Preparation trap

Post accelerator: Normal Conducting LINAC

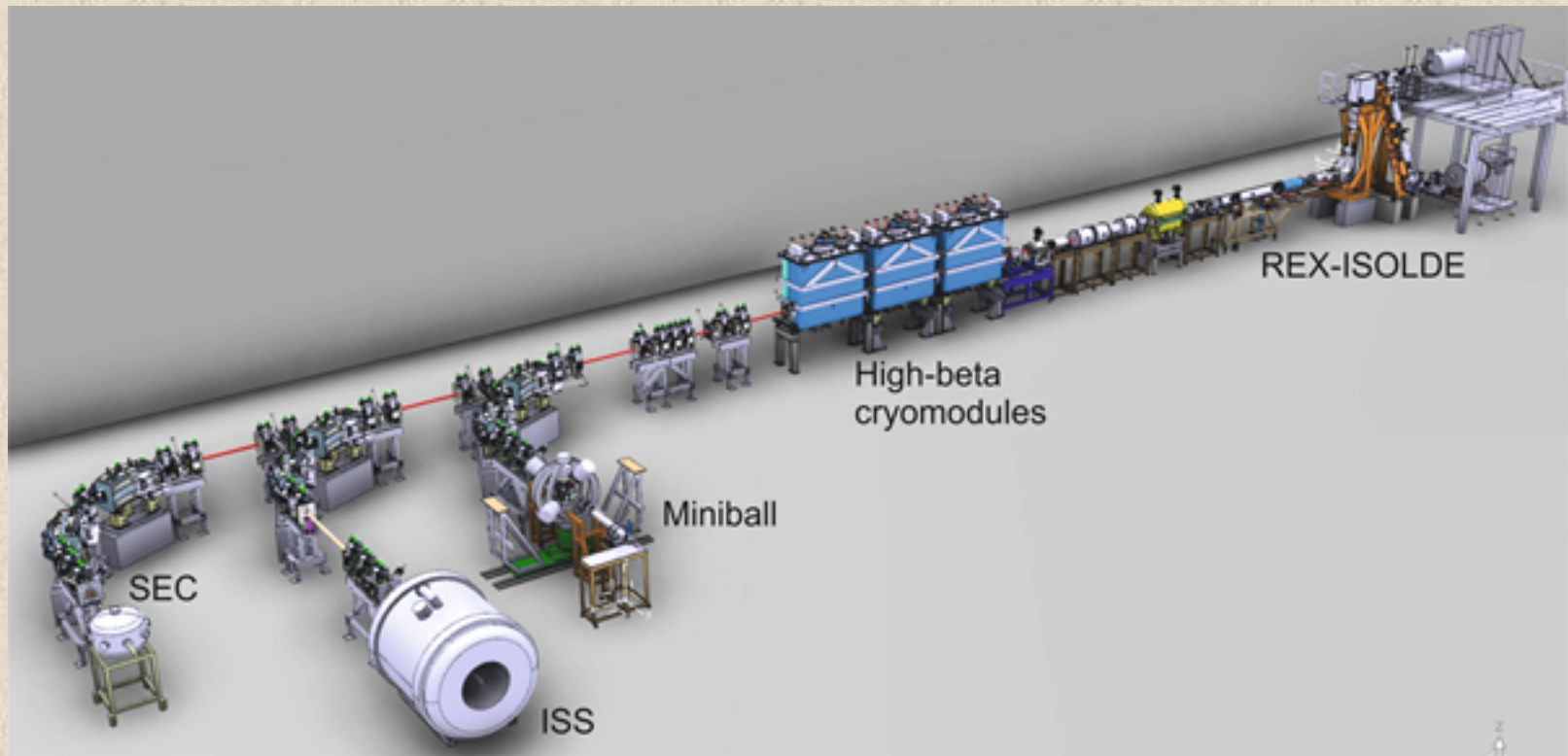


However, energies only 0.8 to 3 MeV/u which are too low for Nuclear reactions

HIE ISOLDE (Superconducting)

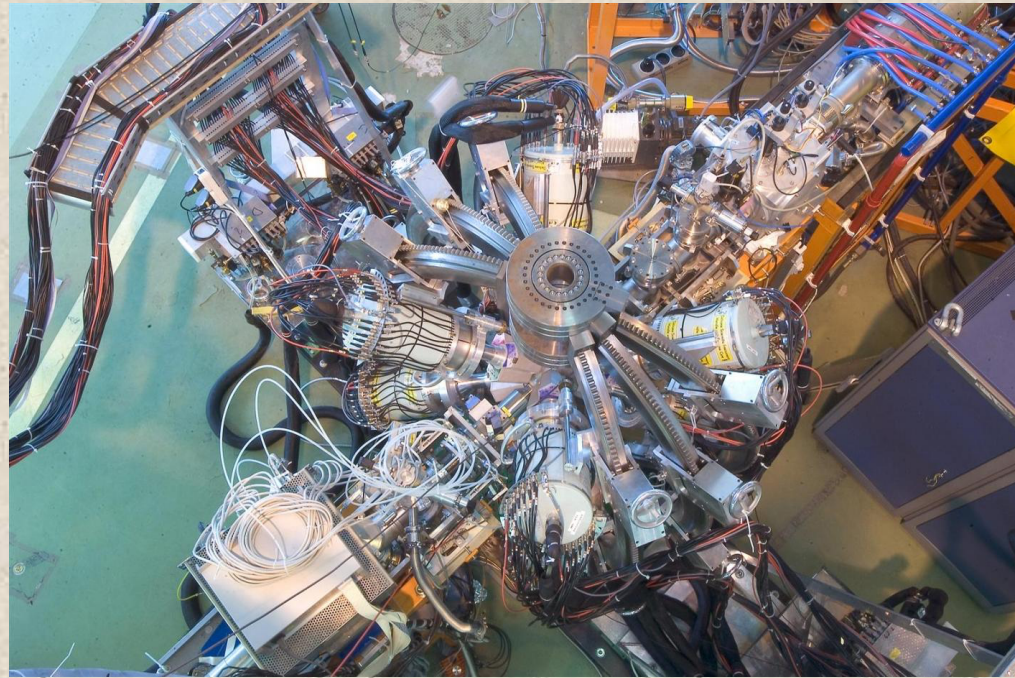
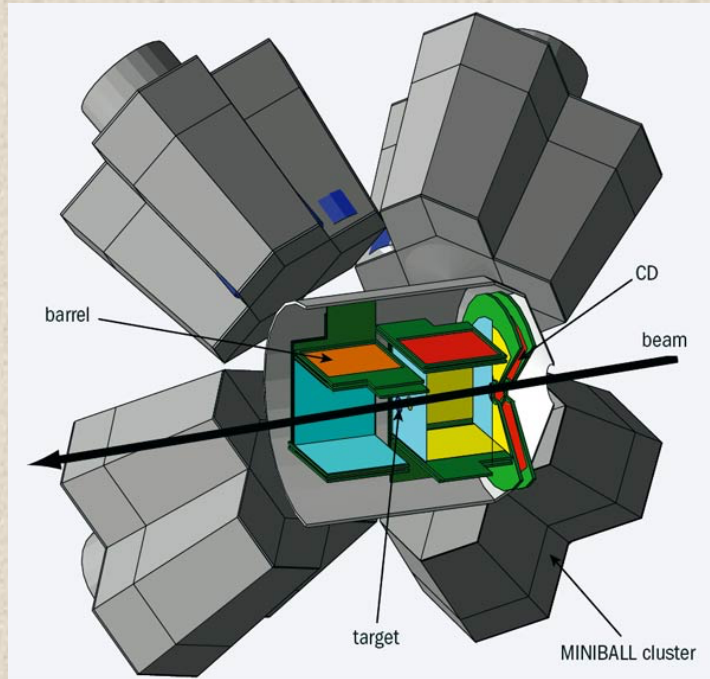
High Intensity and Energy ISOLDE (soon 10 MeV per nucleon)

4.3 MeV per nucleon



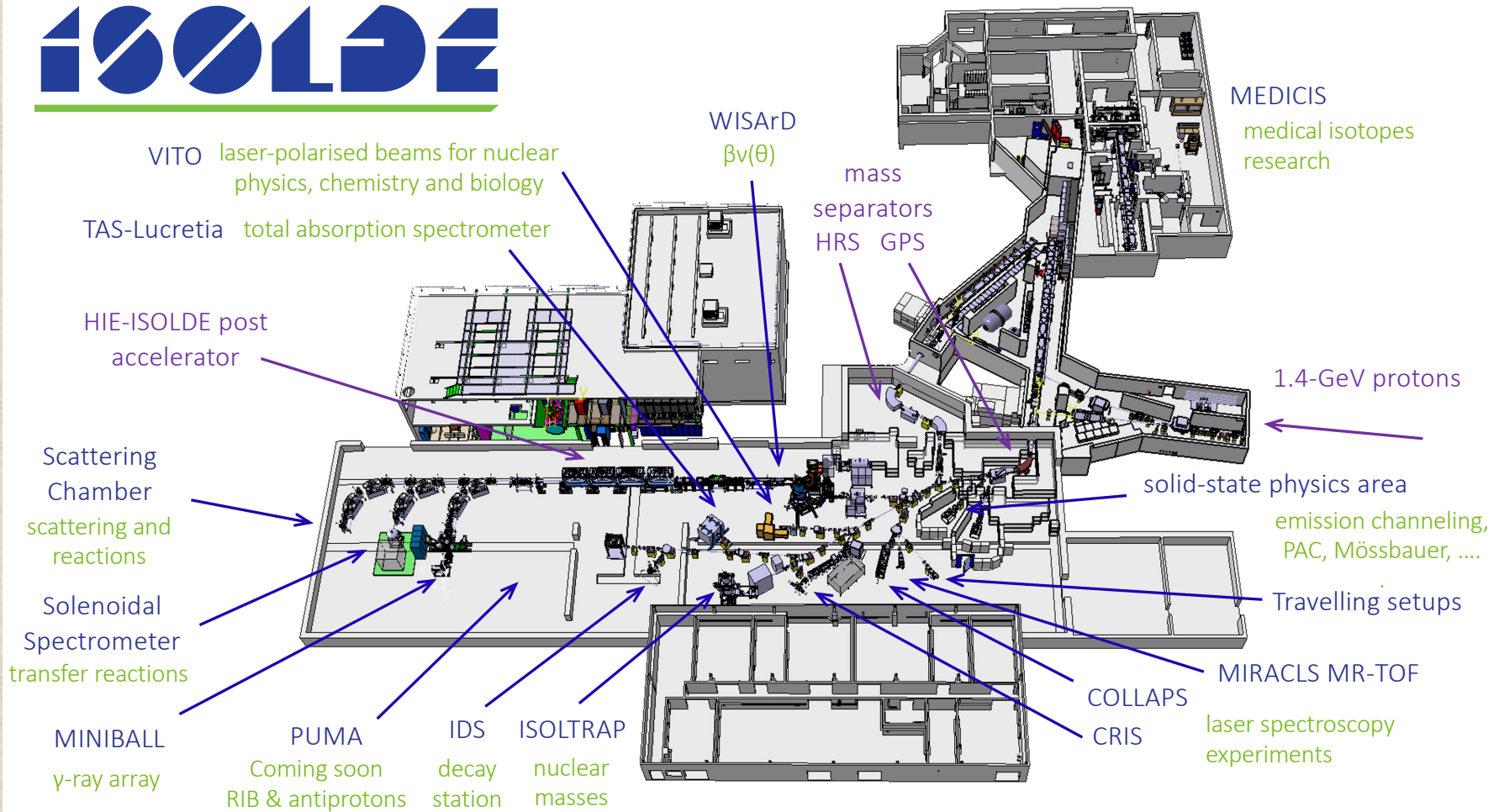
First experiments in 2015

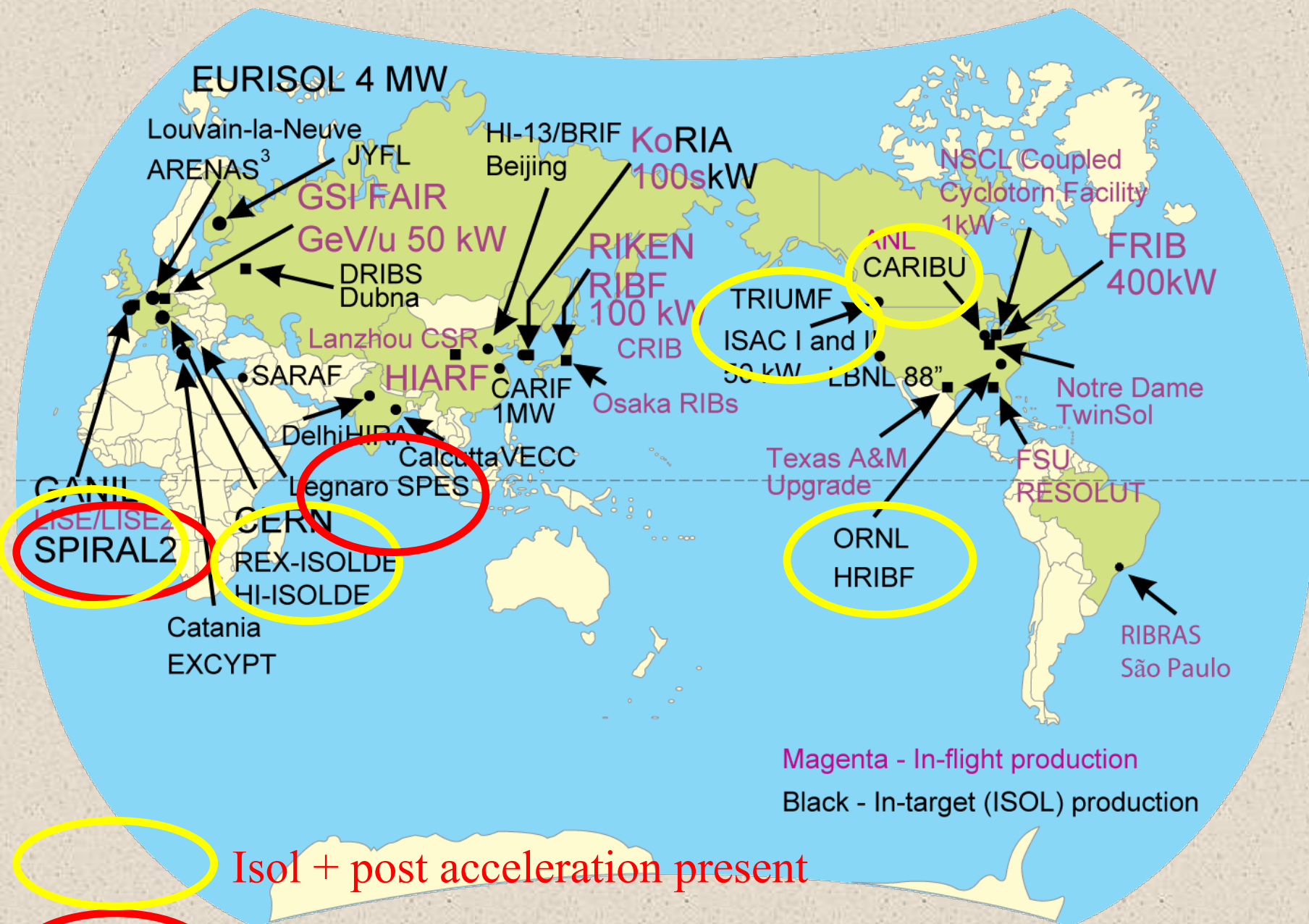
Post accelerated beams + sophisticated detection set-up: Miniball gamma array



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

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





Isol + post acceleration present

Isol + post acceleration future

Future ISOL facilities

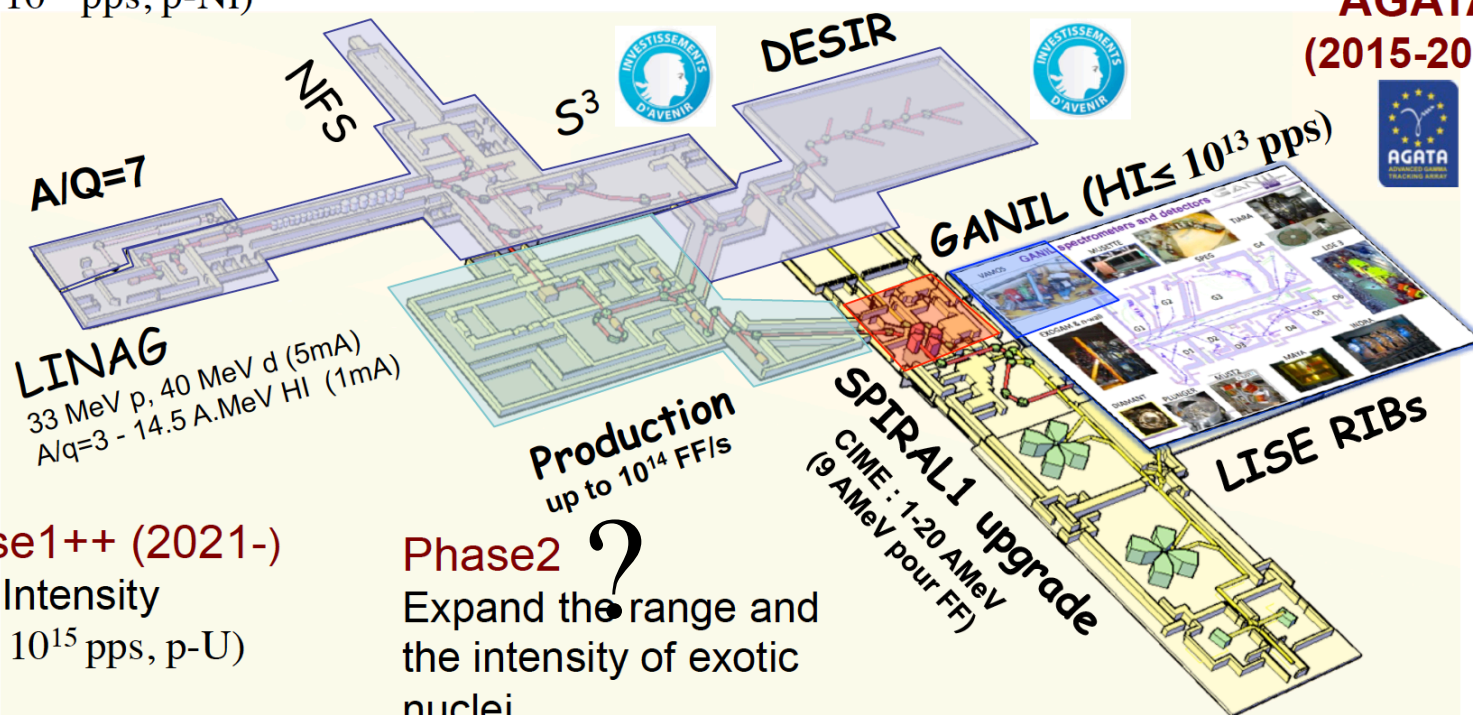
Phase1 (2016-)

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

DESIR Phase1+ (2020-)

Low energy facility

AGATA (2015-2018)



Phase1++ (2021-)

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

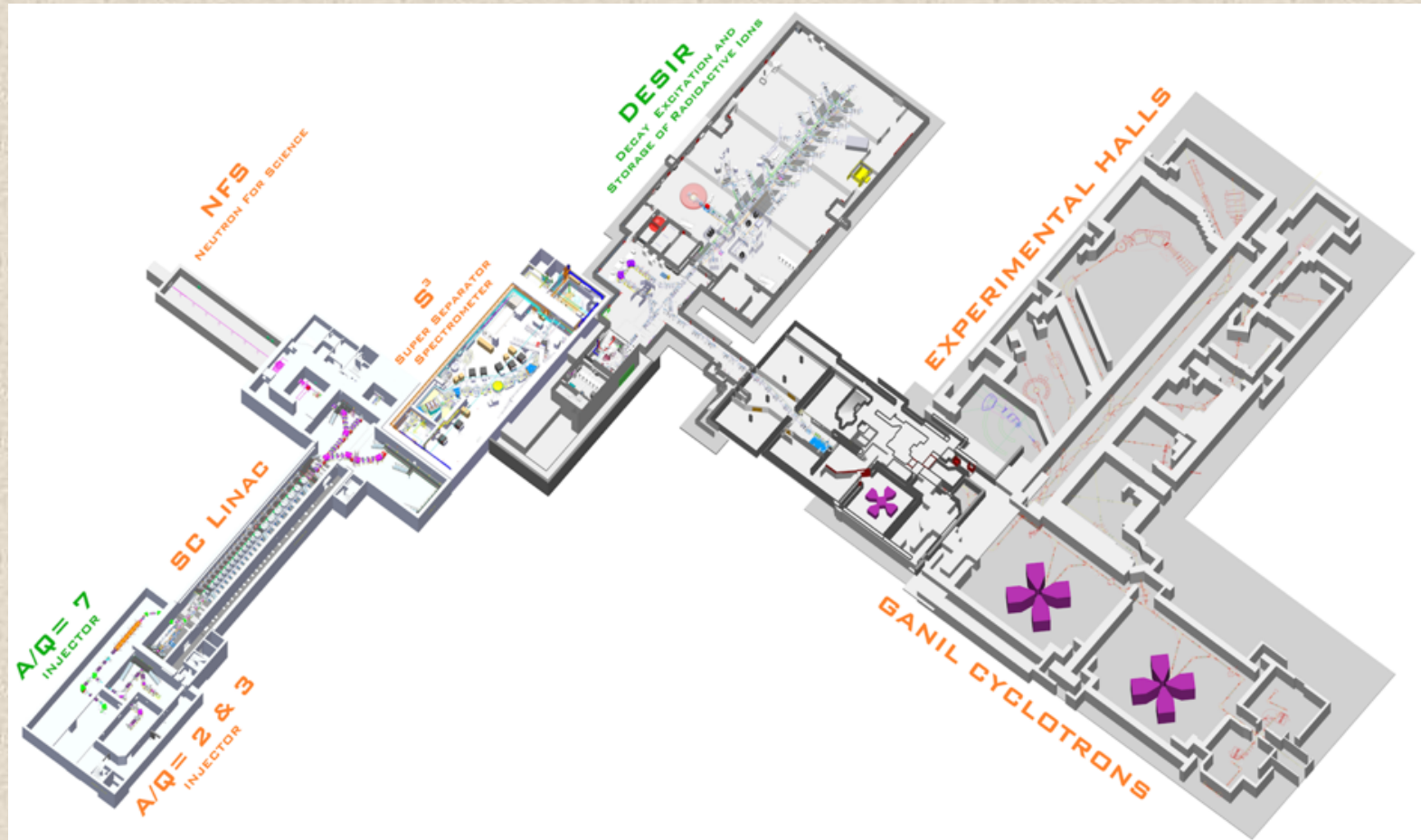
Phase2 ?

Expand the range and
the intensity of exotic
nuclei

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



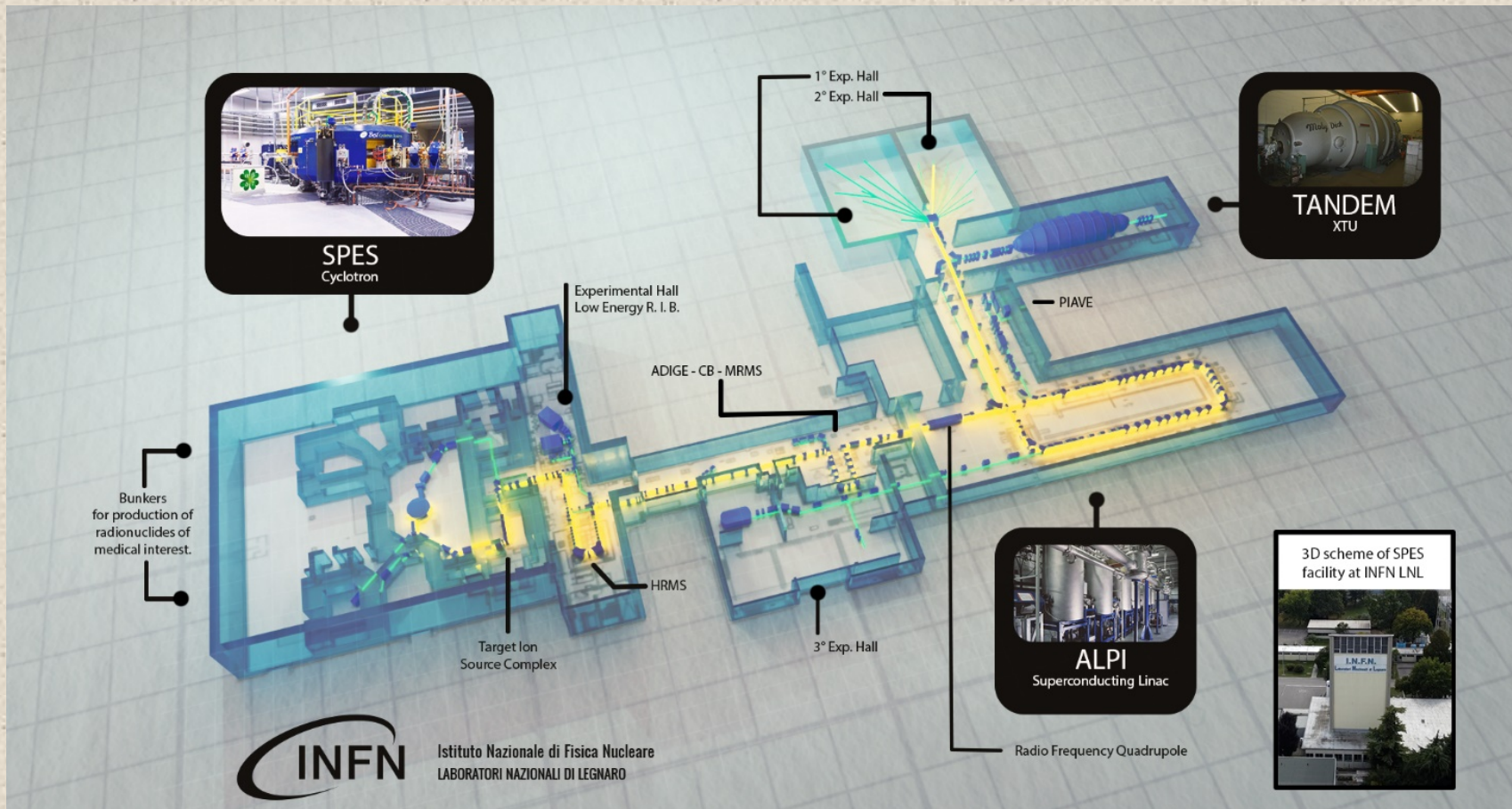
SPES

Selective Production of Exotic Species

Under construction

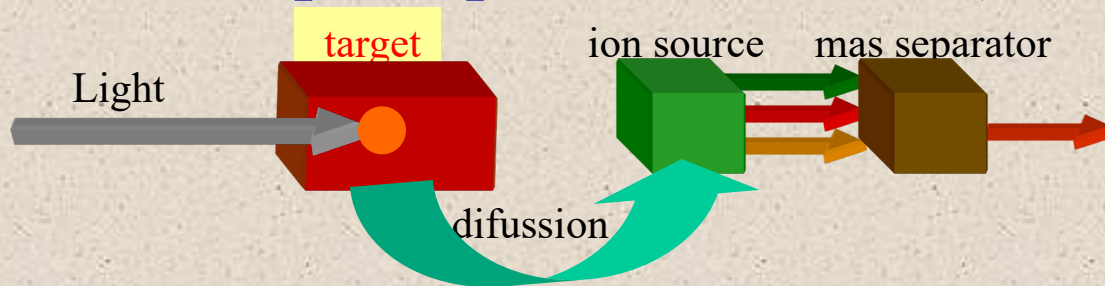


SPES Project



Radioactive nuclei production techniques

Isotopic separation on-line (ISOL)



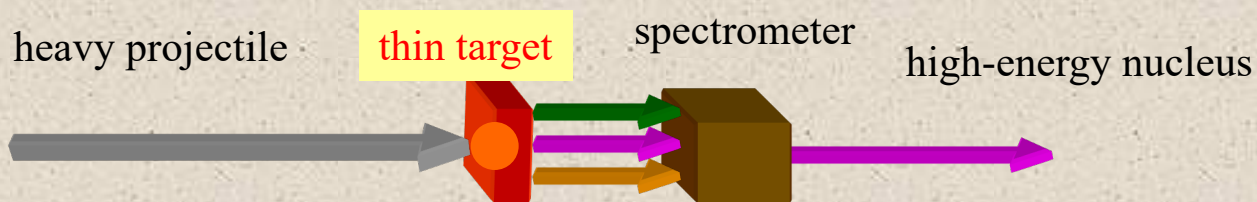
😊😊 We can use thick production targets (100% of energy range)

→ and high injector beam current (upto 10^{16} s^{-1})

😞 long extraction and ionization time (100 ms)

😞 chemistry dependent

In-flight fragmentation



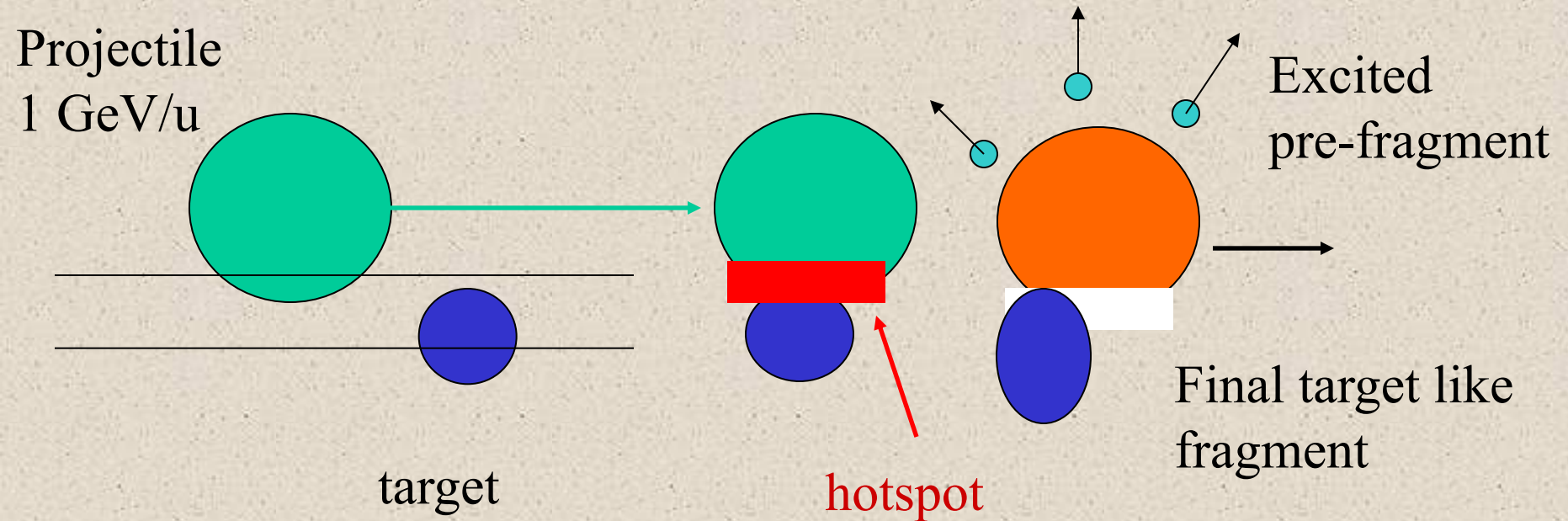
😞 thinner targets (10% of range) => lower beam currents (upto 10^{12} s^{-1})

😊 short separation+identification time (100 ns)

😊 chemistry independent

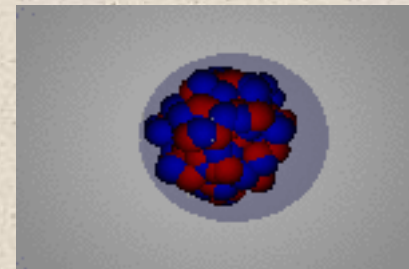
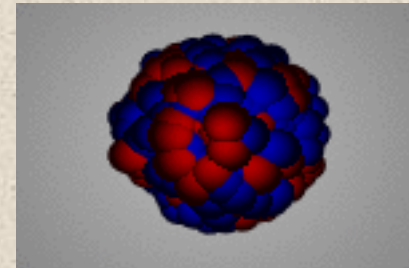
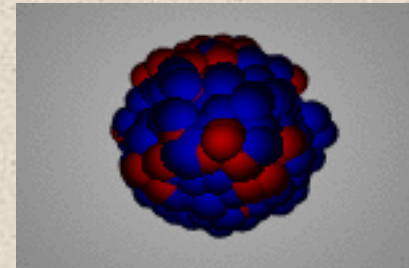
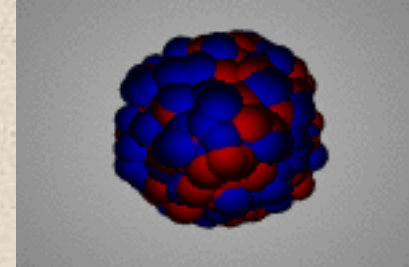
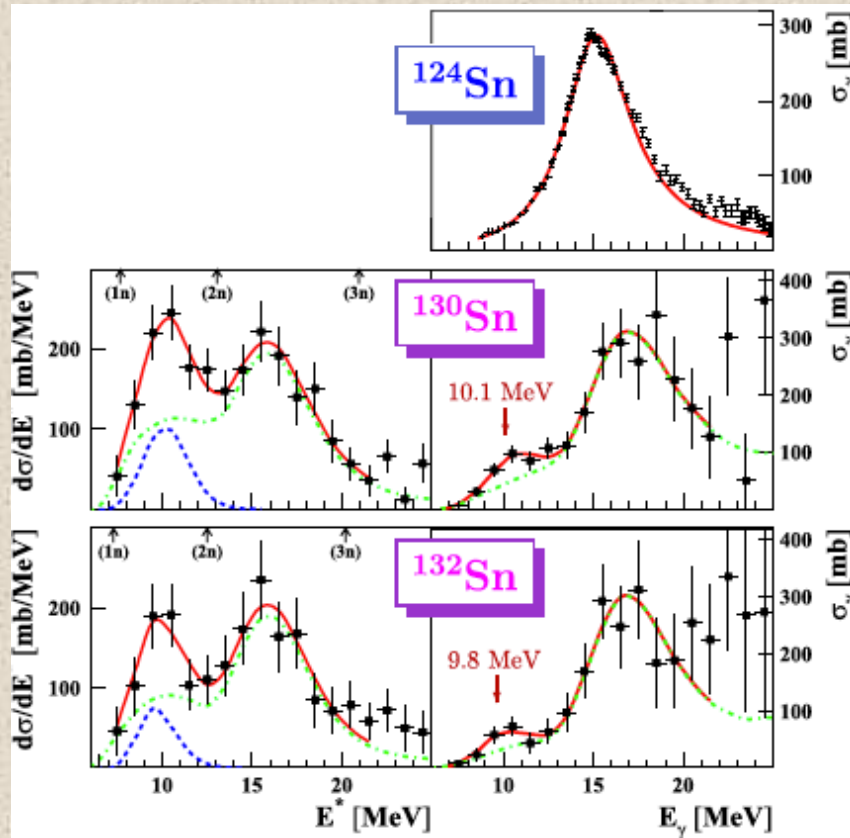
😊 relativistic energies

Projectile Fragmentation Reactions

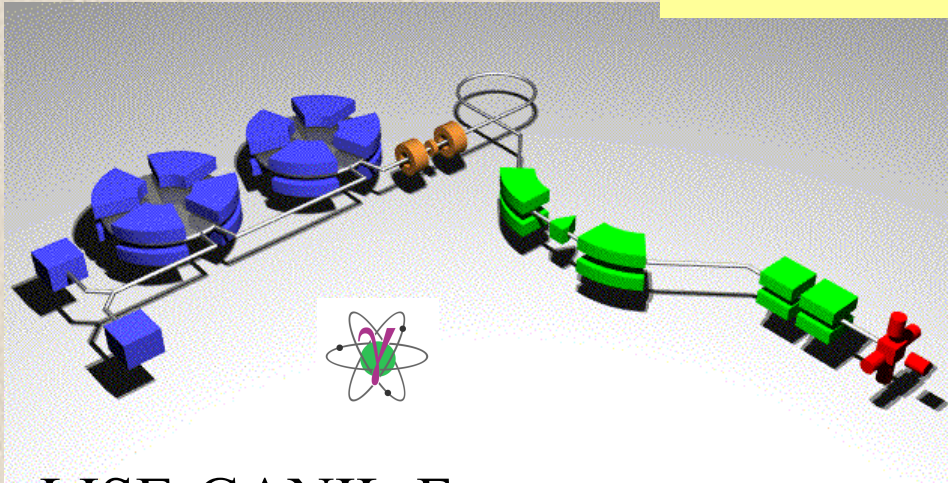


Energy (velocity) of beam > Fermi velocity inside nucleus ~ 30 MeV/u
Can 'shear off' different combinations of protons and neutrons.
Large variety of exotic nuclear species created, all at forward angles with \sim beam velocity.

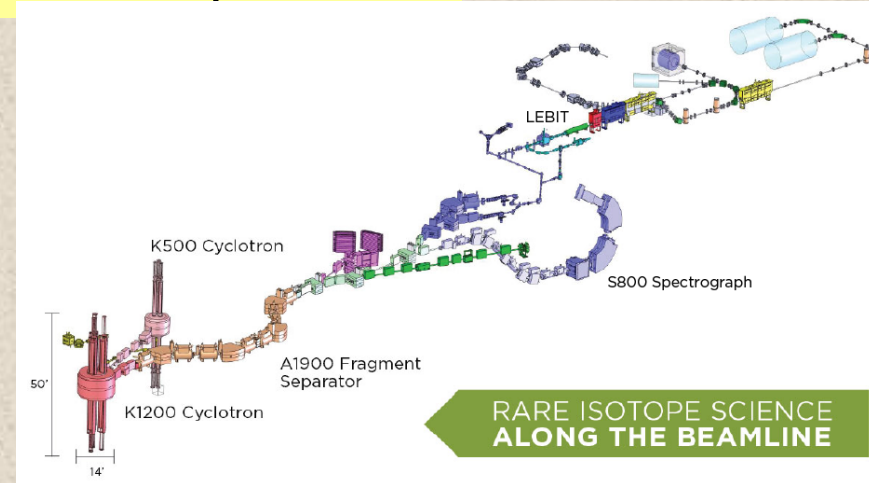
The energy issue: Some physics can only be reached with relativistic energies



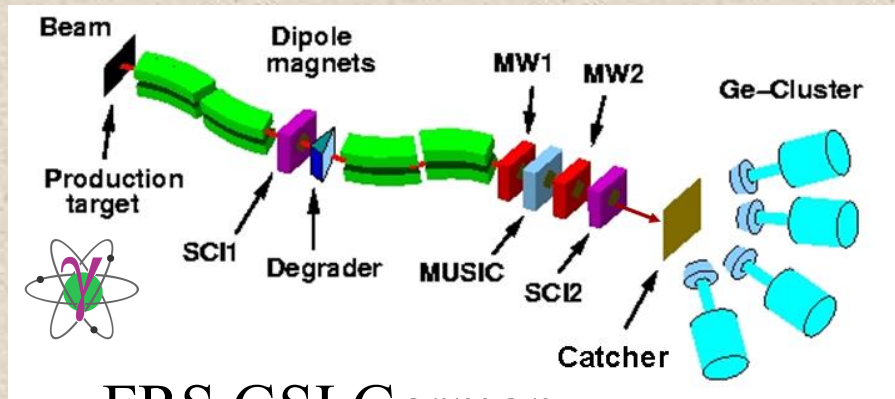
Actual Fragment Separators



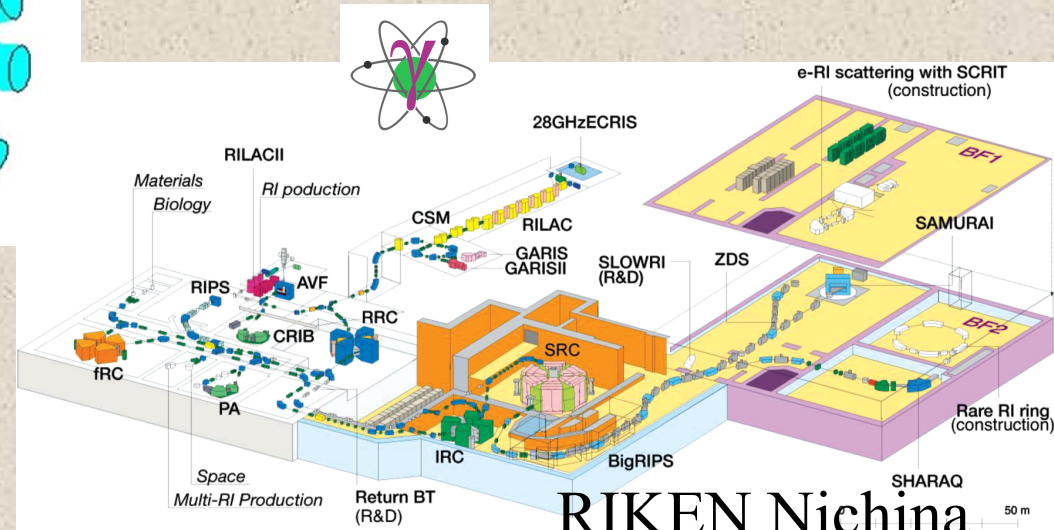
LISE-GANIL-France



NSCL-RIA-→FRIB USA



FRS GSI Germany



RIKEN Nishina
Center, Japan

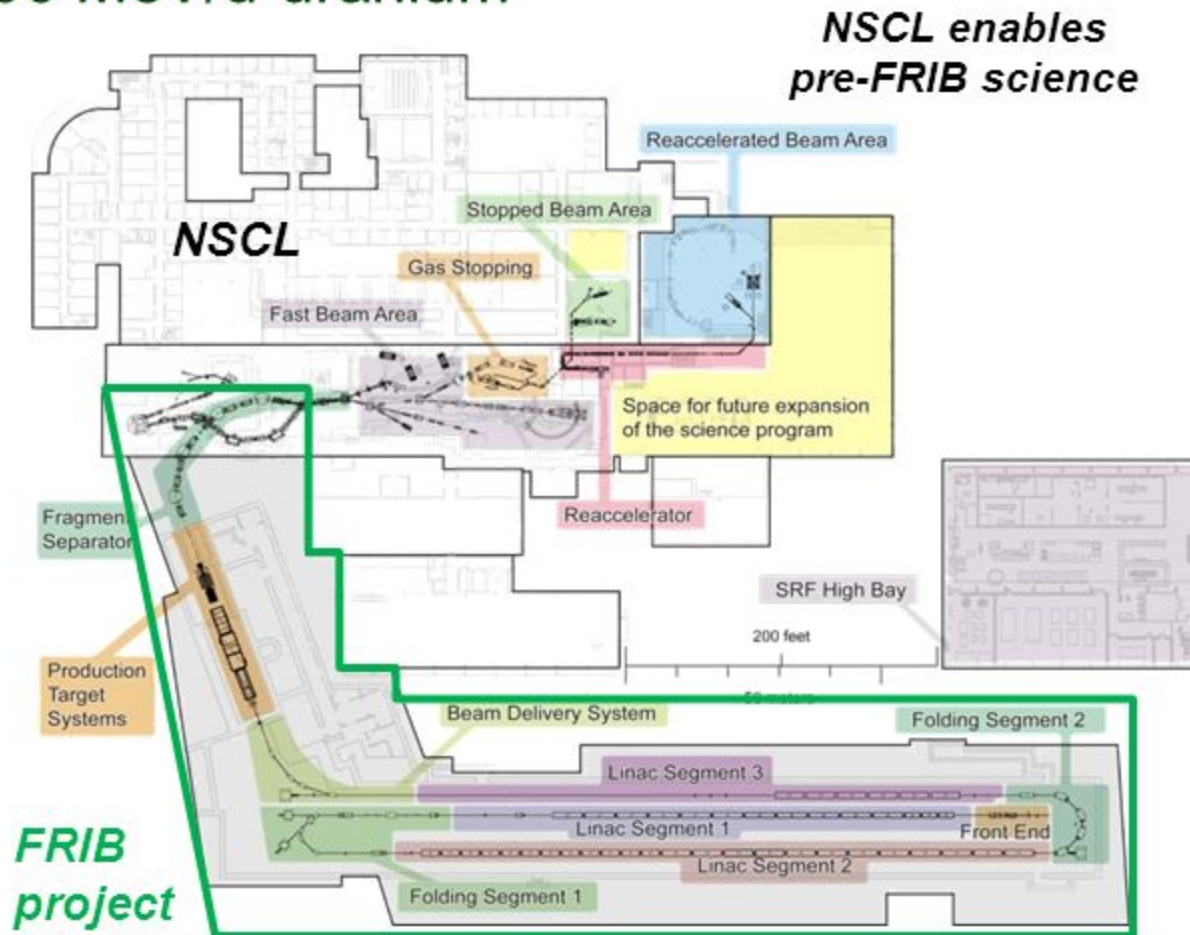
FRIB - Facility for Rare Isotope Beams

World-leading Next-generation Rare Isotope Beam Facility

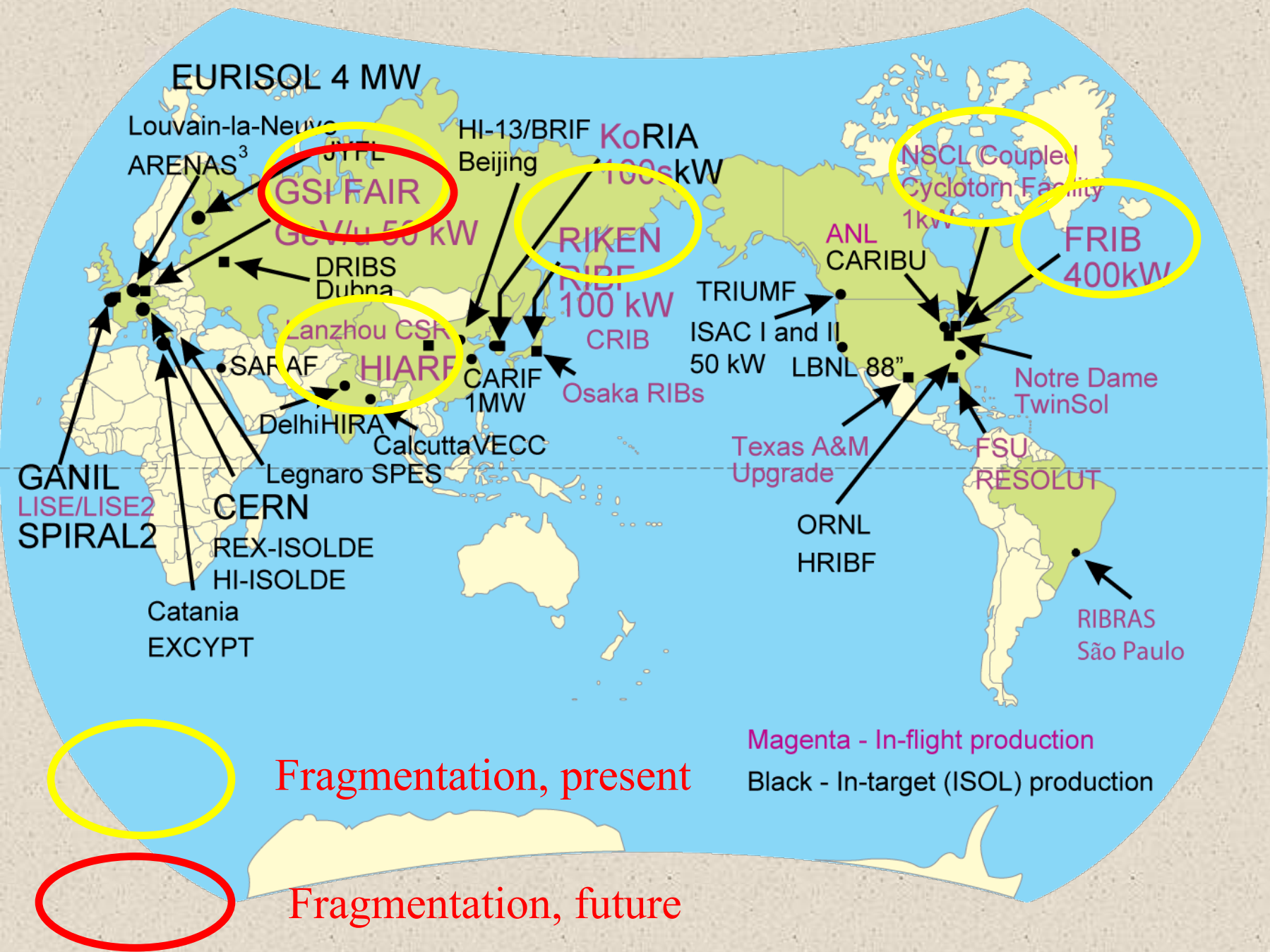
- Rare isotope production via in-flight technique with primary beams up to 400 kW, 200 MeV/u uranium
- Fast, stopped, and re-accelerated beam capability
- Upgrade options
 - 400 MeV/u for uranium
 - ISOL production – multi-user capability

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

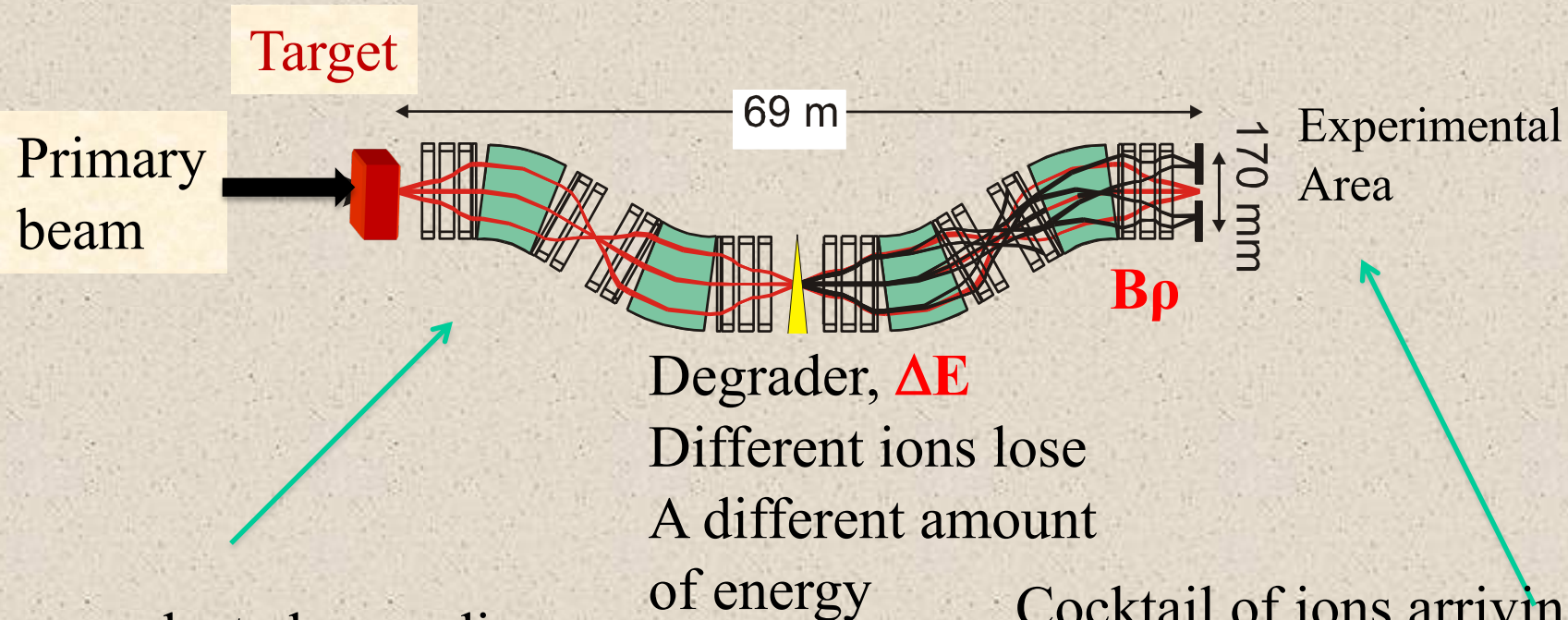


Facility for Rare Isotope Beams
U.S. Department of Energy Office of Science
Michigan State University



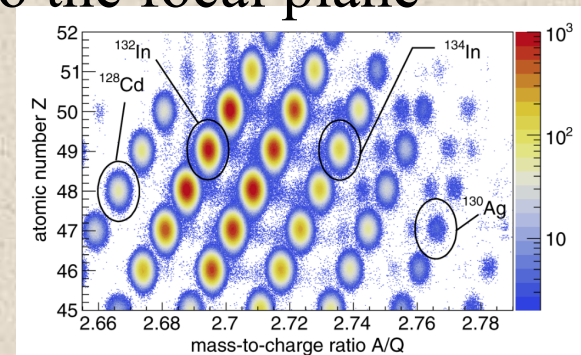
The FRS fragment separator at GSI

$B\rho$ - ΔE - $B\rho$ Separation Method

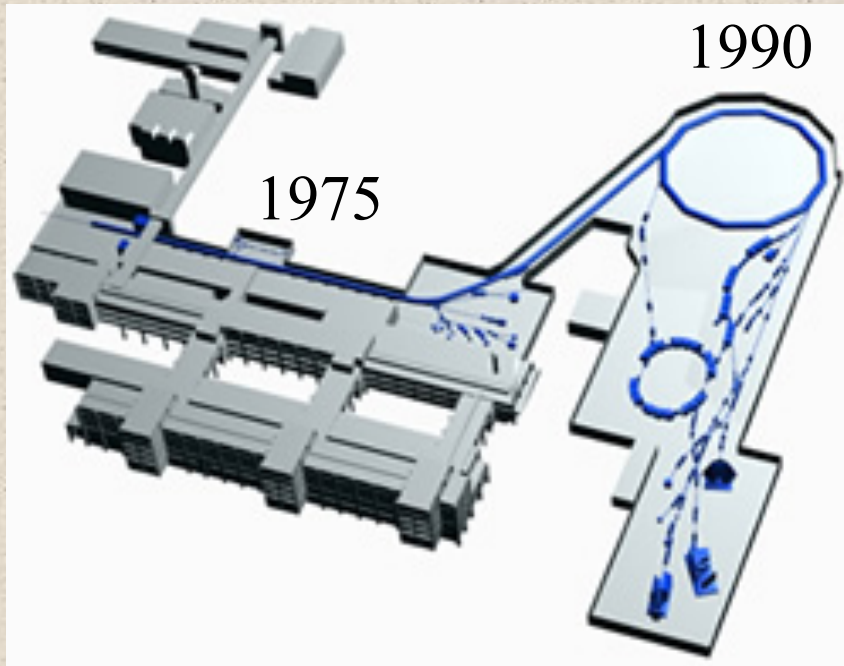


Ions are selected according
to the momentum to charge ratio
 $Mv/q = Mv/Z = B\rho$. Primary beam is rejected.

Cocktail of ions arriving
to the focal plane

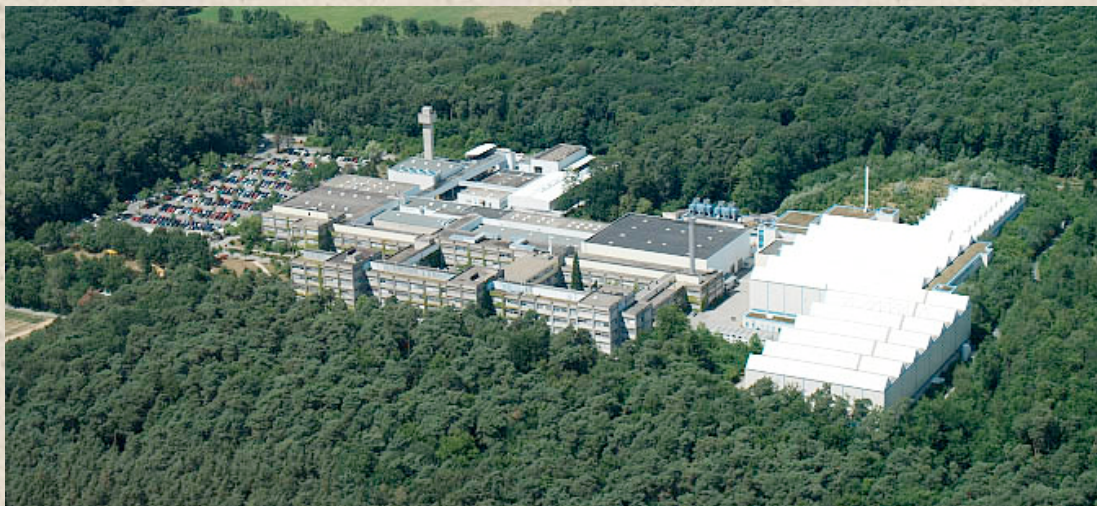


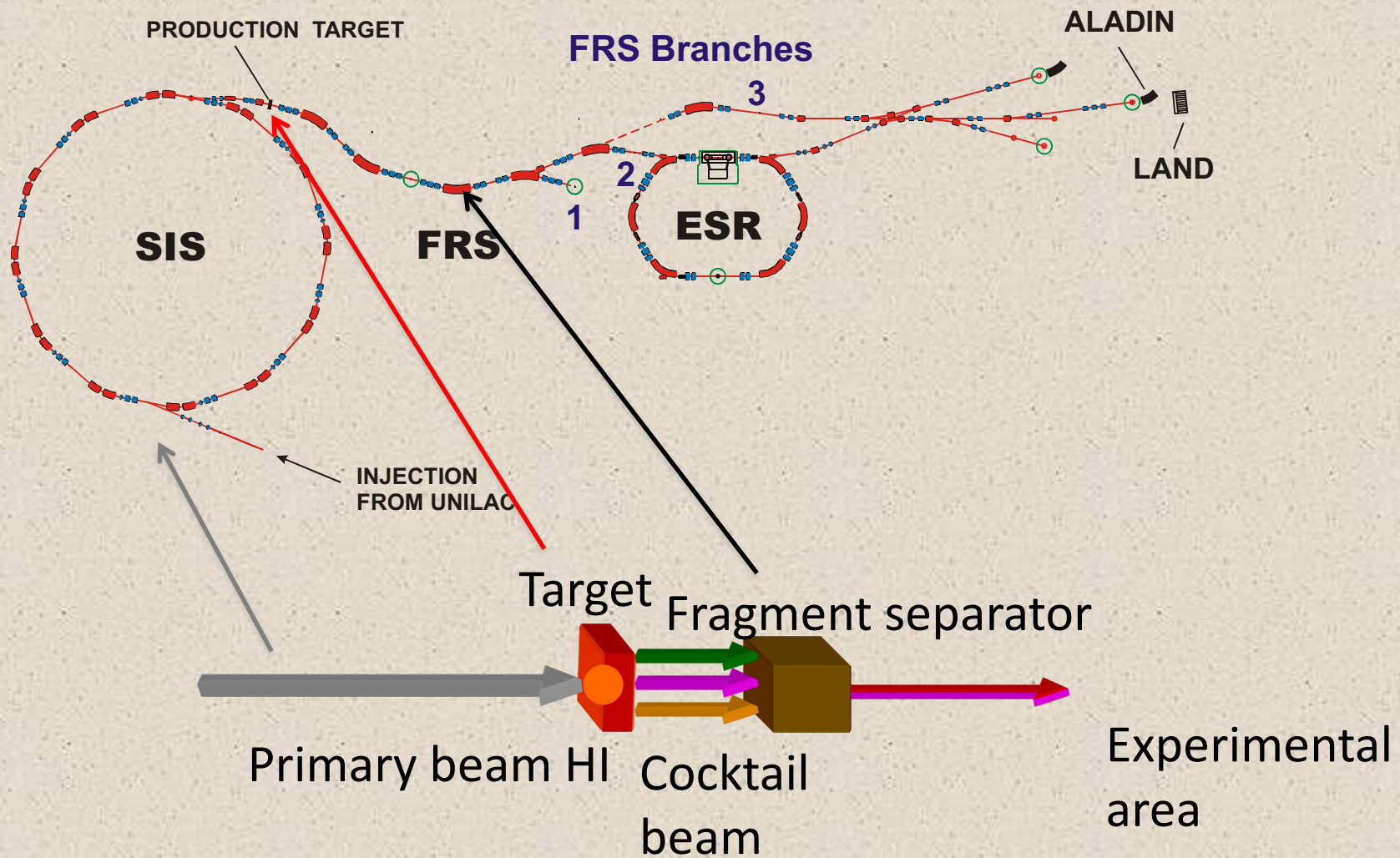
The GSI fragmentation facility



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





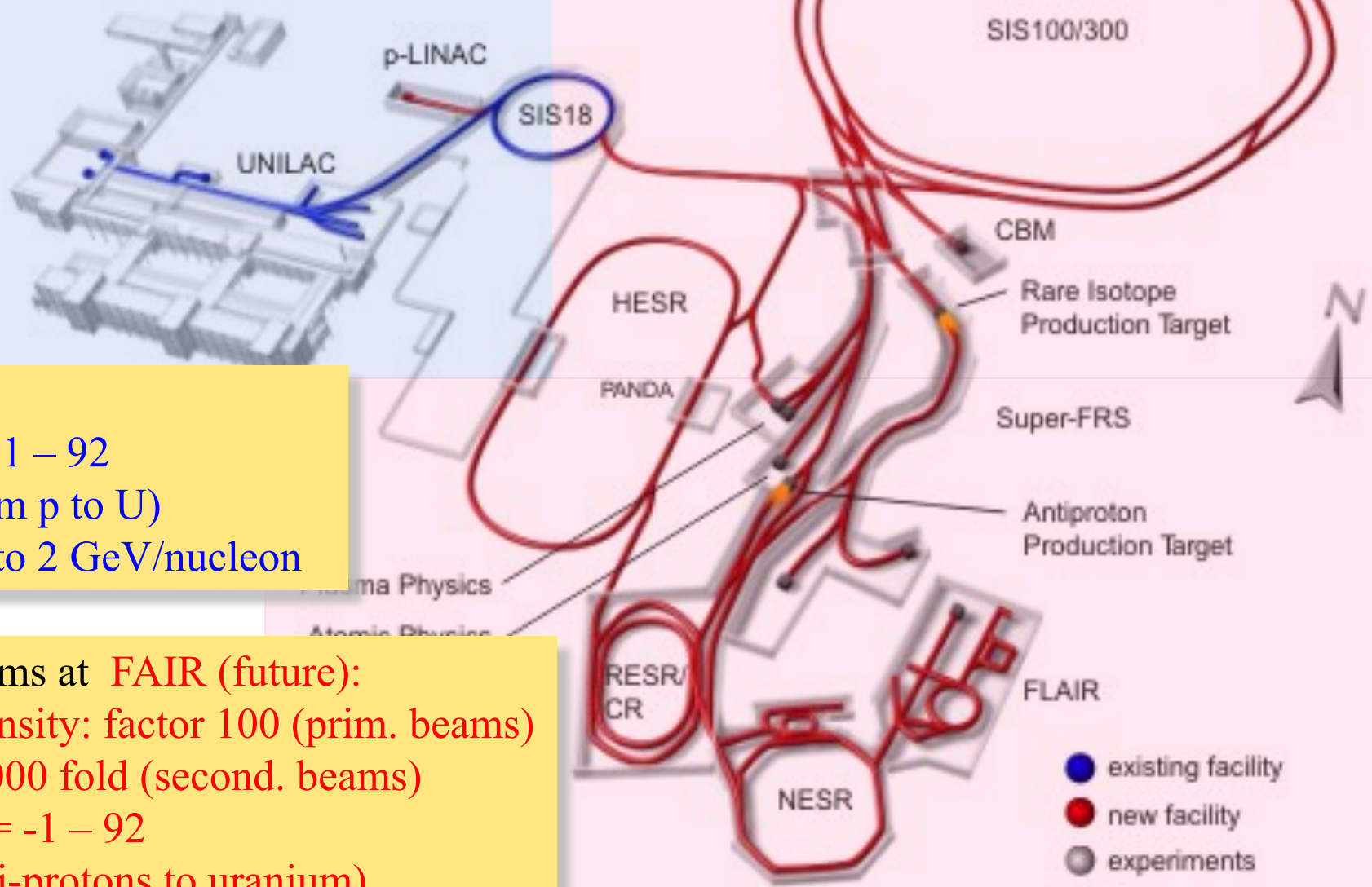
Near Future



B. Rubio. Master FN, Valencia 2024

Existing GSI facility

New 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

FRS to Super-FRS

B ρ - Δ E-B ρ Separation Method

FRS

69 m

170 mm

Degradar

Super-FRS

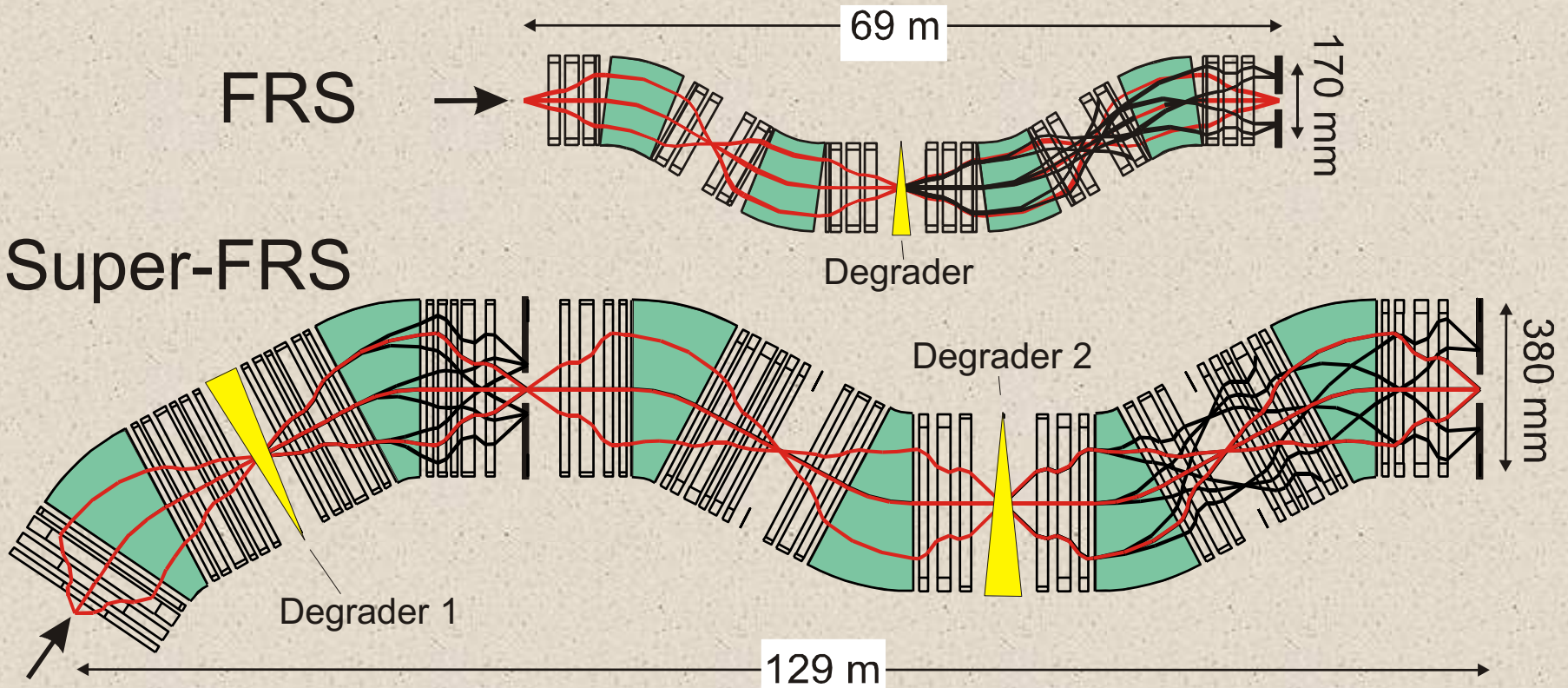
Degradar 2

380 mm

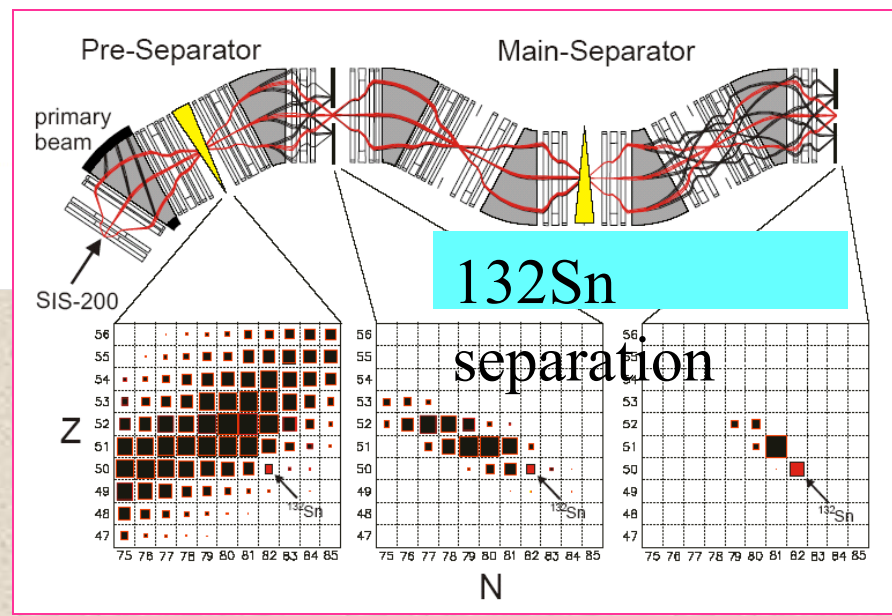
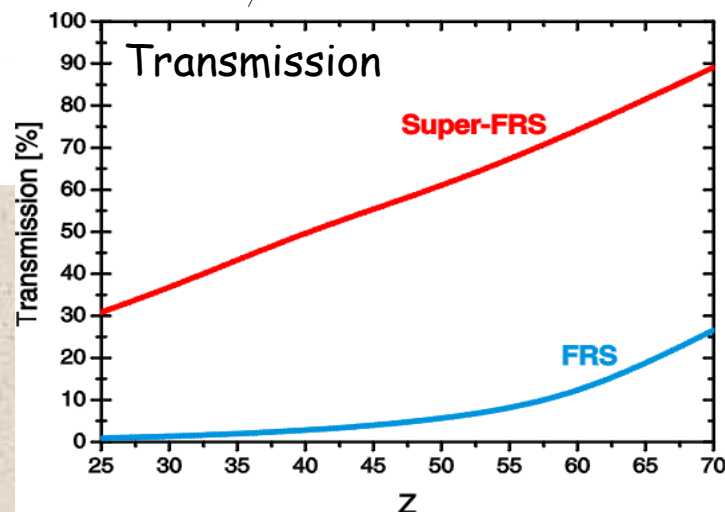
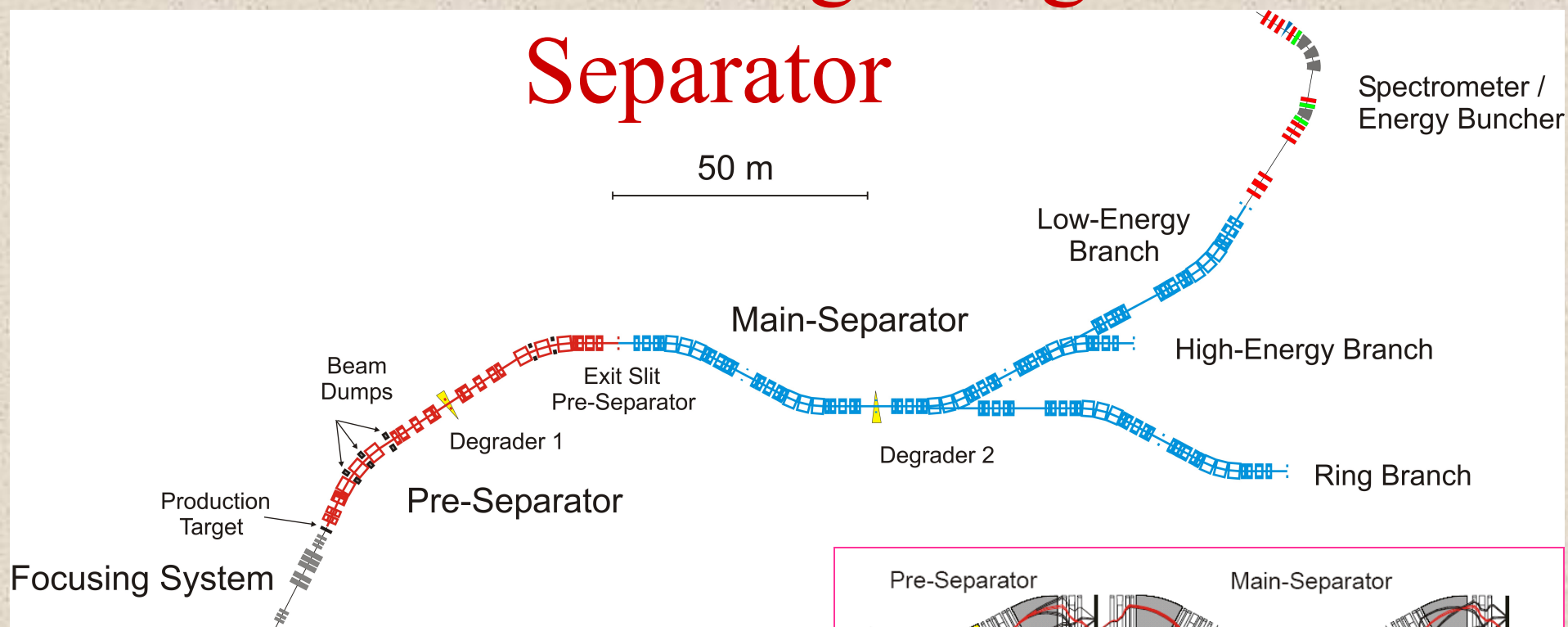
Degradar 1

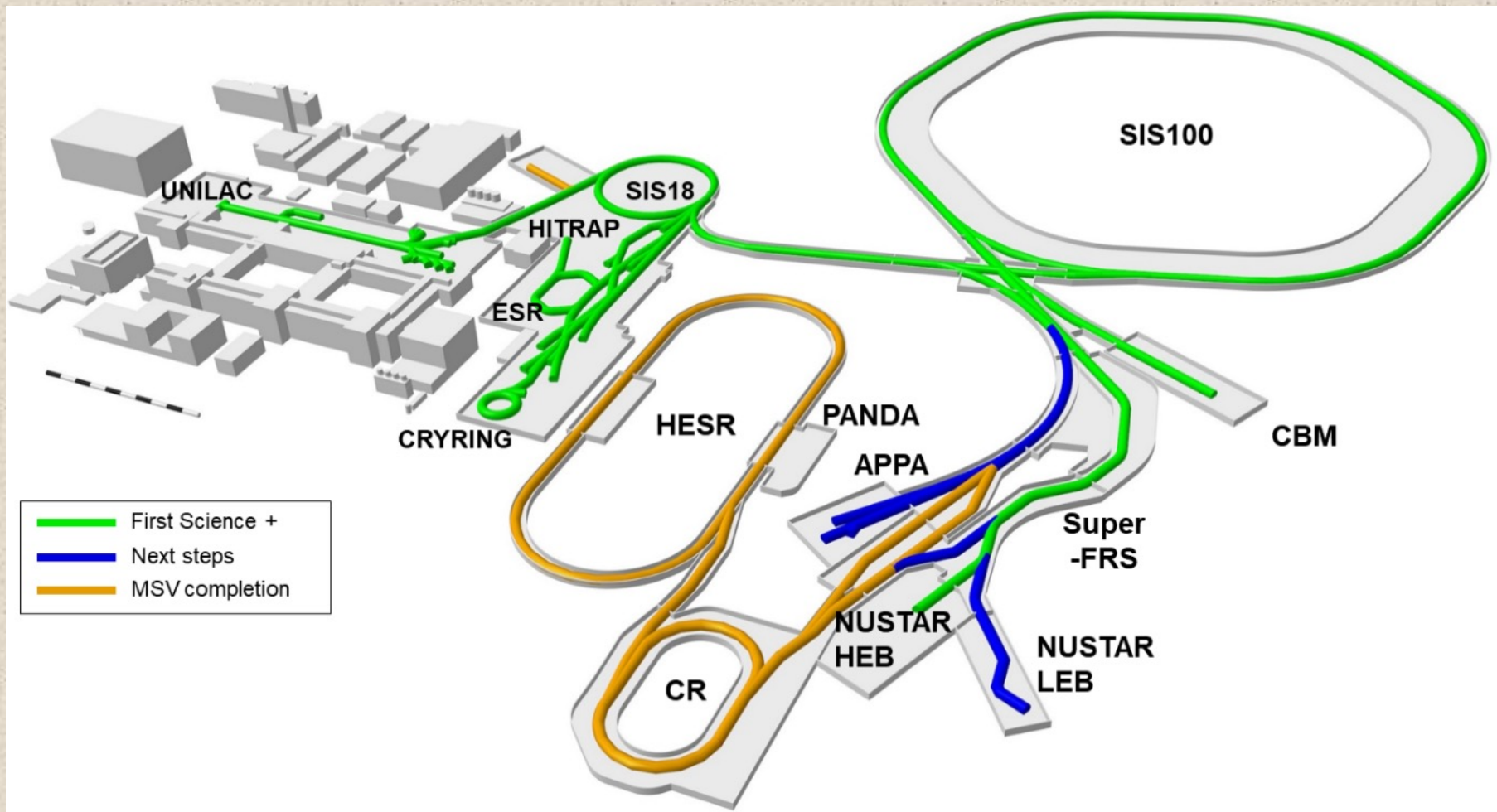
129 m

Optimised to separate
very fast fragment or
fission products



SUPERconducting FRagment Separator





FAIR Construction Field

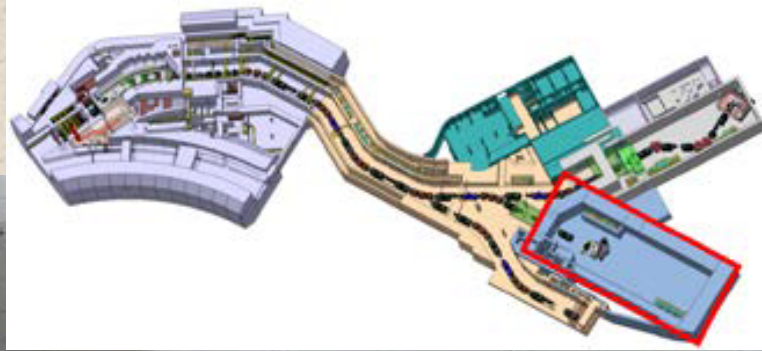




FAIR in construction 2023



- FAIR GmbH
- GSI GmbH

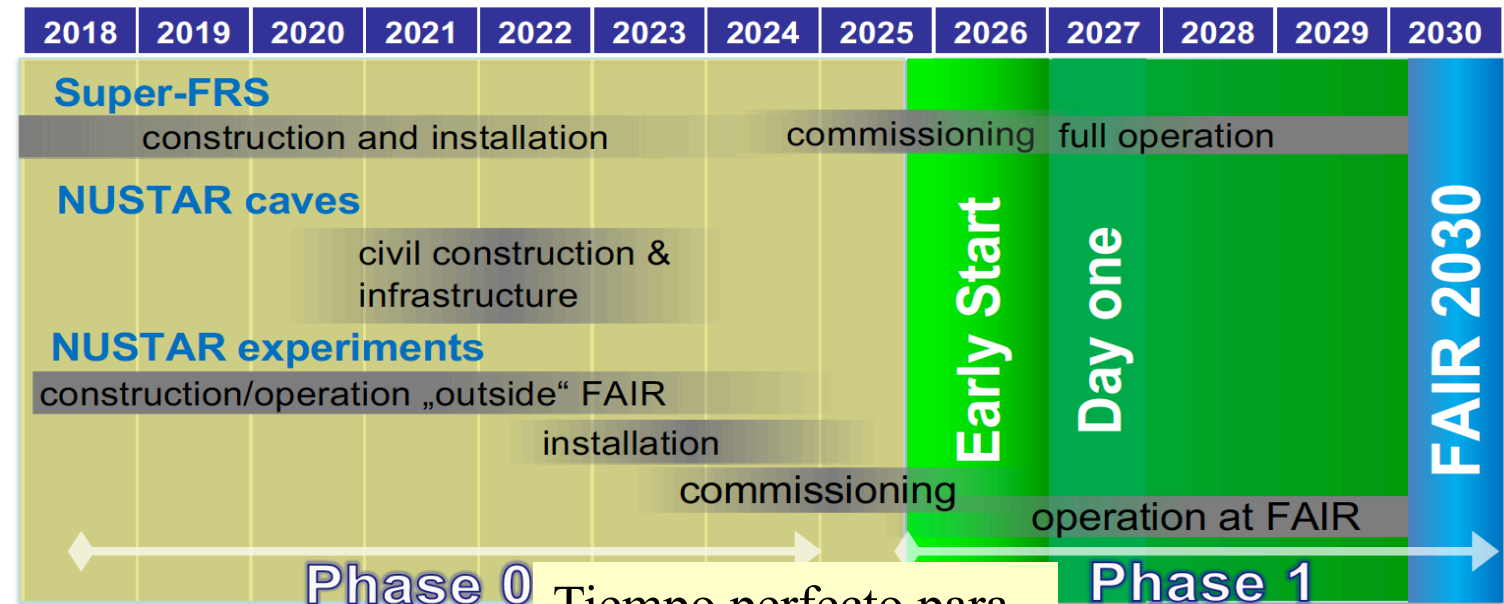


(See talk by Dolores Cortina on
Reactions at Relativistic
Energies)

FAIR - NUSTAR schedule



NUSTAR from Phase-0 to FAIR 2030



Tiempo perfecto para
un postdoc!!

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