

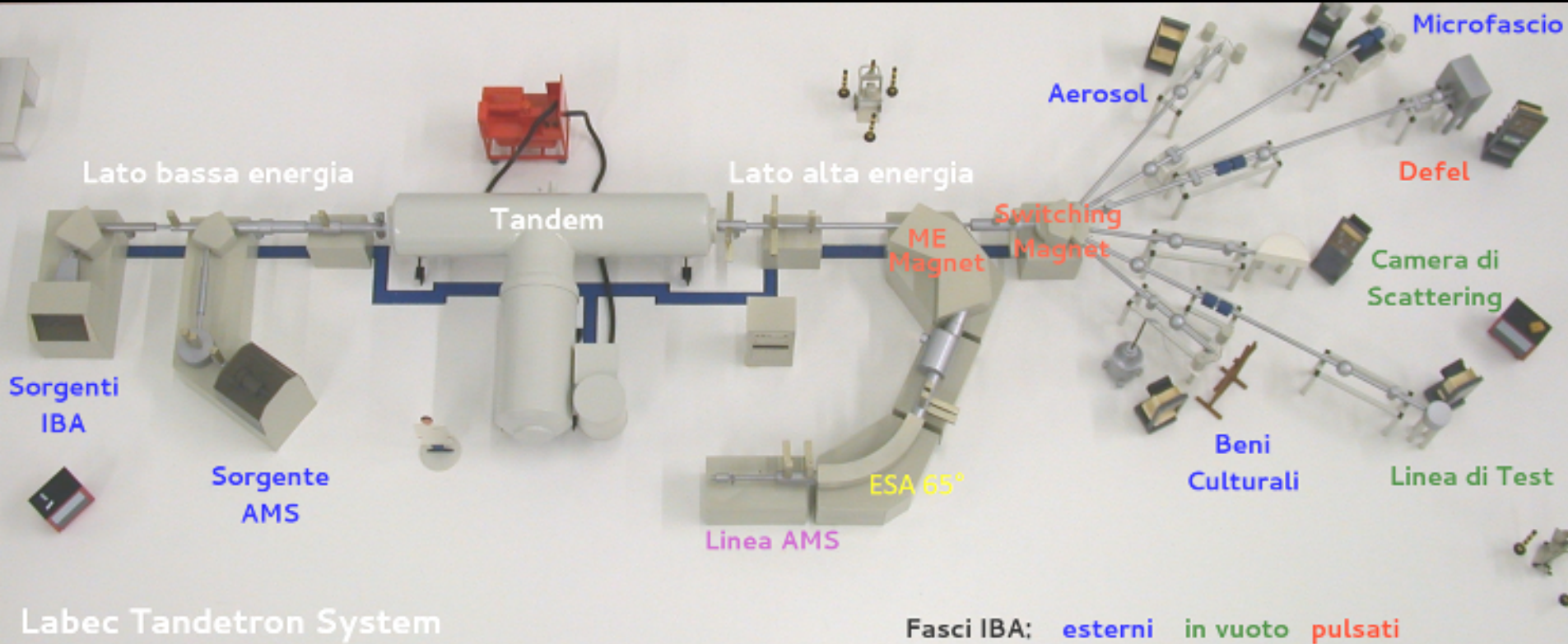


# ***Tuning of ion beams and measurements at the LABEC TANDEM***



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***per il gruppo LABEC Università&Sezione***

# Measurements at LABEC



Labec Tandetron System

Fasci IBA: **esterni** **in vuoto** **pulsati**



# Outline

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- ✓ Why measuring by Accelerator Mass Spectrometry (AMS)
  - The example of radiocarbon
- ✓ How to perform an AMS measurement by a Tandem accelerator facility
  - The example of LABEC accelerator
- ✓ **Laboratory session**
  - Beam transport
  - Data analysis



# Accelerator Mass Spectrometry (AMS)

- ✓ AMS technique has been introduced to measure the relative abundance of **rare isotopes**
  - Radioactive isotopes of medium-long half life
  - Isotopes characterized by a very low abundance

**$^{14}\text{C}$**   
( $t_{1/2} = 5730$  yrs)



*Archaeometry  
Geology  
Palaeoclimate*

**$^{129}\text{I}$**   
( $t_{1/2} = 1.6 \cdot 10^7$  yrs)



*Environmental  
science  
Nuclear  
safeguards*

**$^{10}\text{Be}$**   
( $t_{1/2} = 1.5 \cdot 10^6$  yrs)



*Environmental  
science  
Geology*

**$^{36}\text{Cl}$**   
( $t_{1/2} = 3 \cdot 10^5$  yrs)

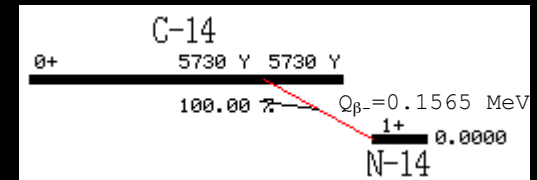


*Environmental  
science  
Geology*



# The example of $^{14}\text{C}$

- ✓ Radiocarbon is the only natural radioactive carbon isotope
- ✓ It is produced in the atmosphere and enters in all the biosphere

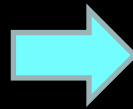


→ **equilibrium condition:** 
$$^{14}R = \frac{^{14}\text{C}}{C_{\text{totale}}} \approx \frac{^{14}\text{C}}{^{12}\text{C}} \approx 1.2 \cdot 10^{-12}$$

- ✓ After the death of an organism, its  $^{14}\text{C}$  content starts to decrease due to the radioactive

**Dating of organic findings**

$$^{14}R(t) = ^{14}R_0 e^{-t/\tau}$$

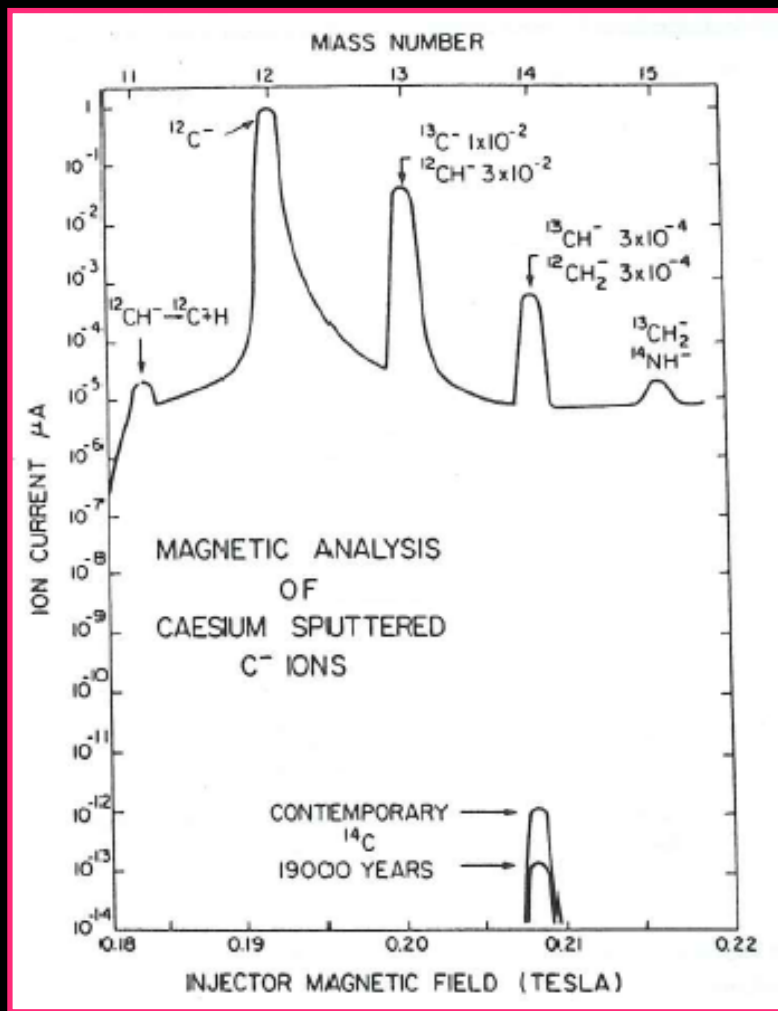


$$t = \tau \cdot \ln \left( \frac{^{14}R_0}{^{14}R(t)} \right)$$





# ***<sup>14</sup>C: a needle in a haystack***



✓ The typical sensitivity of traditional mass spectrometry is not sufficient



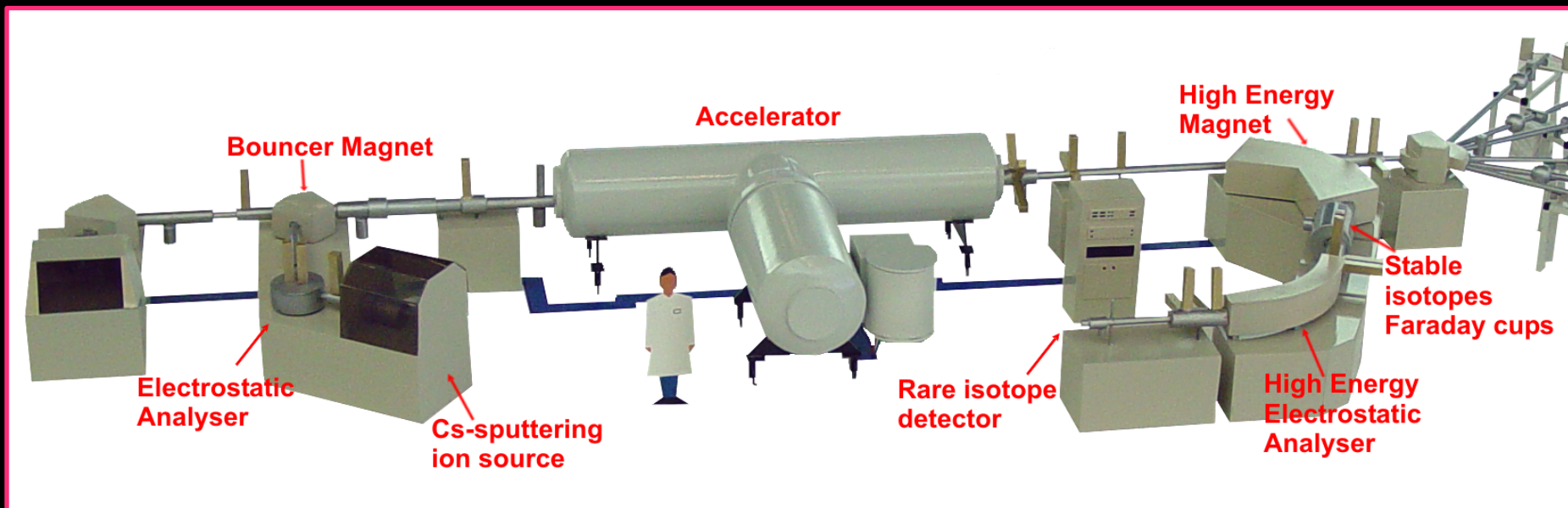
**In AMS, electrostatic and magnetic "filters" are coupled to a Tandem electrostatic accelerator**

***The accelerator itself is used as a sort of high sensitivity detector***





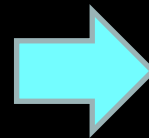
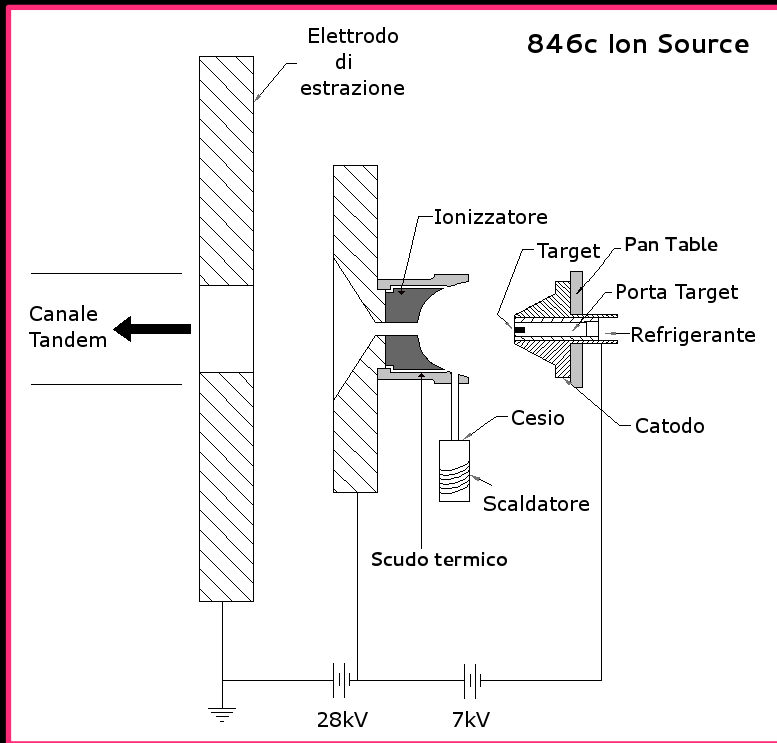
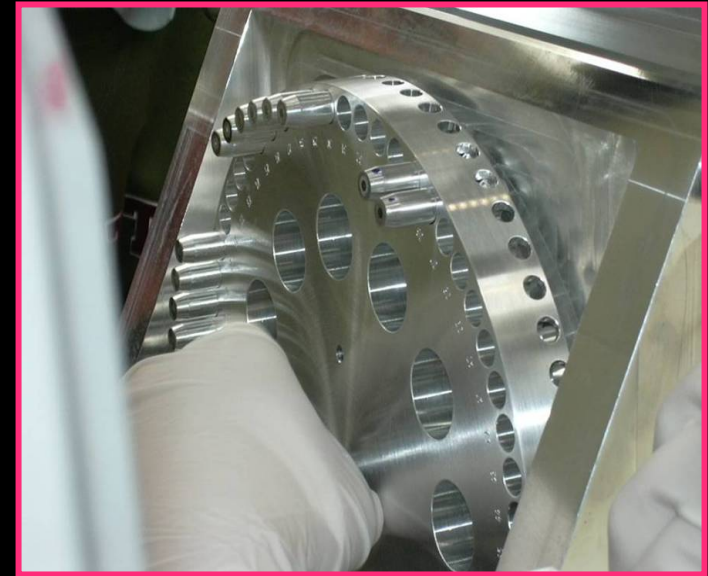
# The AMS beam line at LABEC





# From the sample to the ions: the source

- ✓ Samples are inserted in the Cs-sputtering ion source as graphite pellets ( $m < 1 \text{ mg}$ )



**Negative ions are extracted from the source  
→ the isobar  $^{14}\text{N}$  is suppressed**

# Suppression of $^{14}\text{N}$

Ionization Potentials and Electron Affinities of the Elements

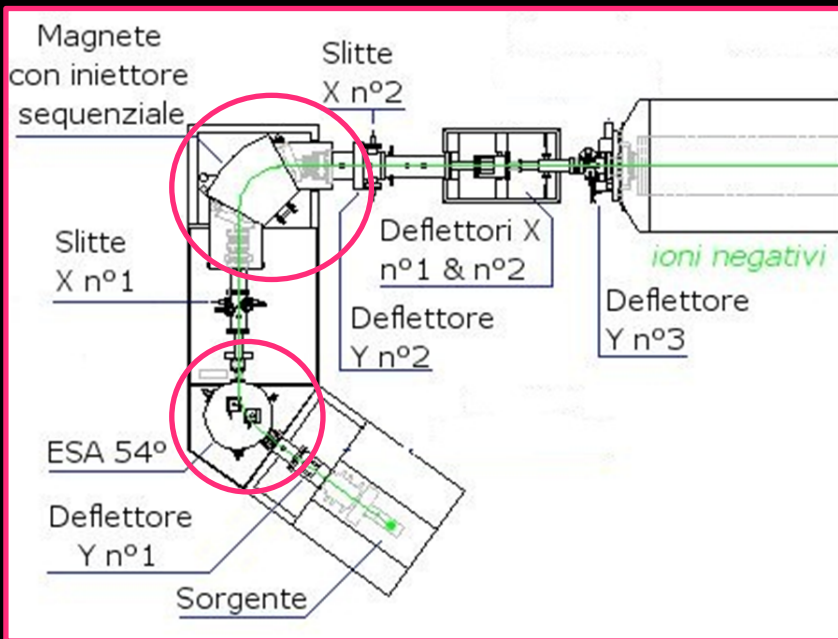
IA	IIA	IIIA	IVA	VA	VIA	VIIA	VIIIA
$^1\text{H}$ 13.59 0.754	Ionization Potential Electron Affinity						$^2\text{He}$ 24.48 0.078*
$^3\text{Li}$ 5.39 0.618	$^4\text{Be}$ 9.32 0.195*	$^5\text{B}$ 8.30 0.277	$^6\text{C}$ 11.26 1.263	$^7\text{N}$ 14.53 -0.07	$^8\text{O}$ 13.61 1.461	$^9\text{F}$ 17.42 3.399	$^{10}\text{Ne}$ 21.56 < 0
$^{11}\text{Na}$ 5.14 0.548	$^{12}\text{Mg}$ 7.64 < 0	$^{13}\text{Al}$ 5.98 0.441	$^{14}\text{Si}$ 8.15 1.385	$^{15}\text{P}$ 10.48 0.747	$^{16}\text{S}$ 10.36 2.077	$^{17}\text{Cl}$ 13.01 3.617	$^{18}\text{Ar}$ 15.76 < 0
$^{19}\text{K}$ 4.34 0.501	$^{20}\text{Ca}$ 6.11 0.043	$^{31}\text{Ga}$ 6.00 0.30	$^{32}\text{Ge}$ 7.90 1.2	$^{33}\text{As}$ 9.81 0.81	$^{34}\text{Se}$ 9.75 2.021	$^{35}\text{Br}$ 11.81 3.365	$^{36}\text{Kr}$ 14.00 < 0
$^{37}\text{Rb}$ 4.18 0.486	$^{39}\text{Sr}$ 5.70 < 0	$^{49}\text{In}$ 5.79 0.3	$^{50}\text{Sn}$ 7.34 1.2	$^{51}\text{Sb}$ 8.64 1.07	$^{52}\text{Te}$ 9.01 1.971	$^{53}\text{I}$ 10.45 3.059	$^{54}\text{Xe}$ 12.13 < 0
$^{55}\text{Cs}$ 3.89 0.472	$^{56}\text{Ba}$ 5.21 < 0	$^{81}\text{Tl}$ 6.11 0.2	$^{82}\text{Pb}$ 7.42 0.364	$^{83}\text{Bi}$ 7.29 0.946	$^{84}\text{Po}$ 8.42 1.9	$^{85}\text{At}$ 9.5 2.8	$^{86}\text{Rn}$ 10.75 < 0

\*Metastable

*R. Middleton, A negative-ion cookbook*

- ✓  $^{14}\text{N}$  does not form negative ions in the stable state
- ✓ Lifetime of negative excited states are too short with respect to the time needed to travel to the injection into the accelerator
- ✓ Possible fragments from molecules (e.g.  $\text{NH}$ ) are discriminated thanks to the following charge-energy-mass analysis

# The analysis on the low energy side



- ✓ First selection according to **energy/charge**

$$\epsilon r = 2 \frac{E}{q}$$

$$q = -1$$
$$E = 35 \text{ keV}$$

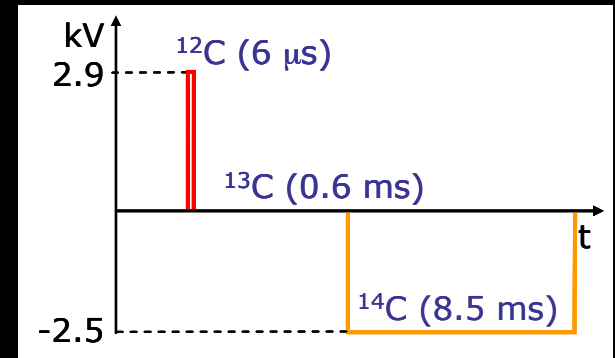
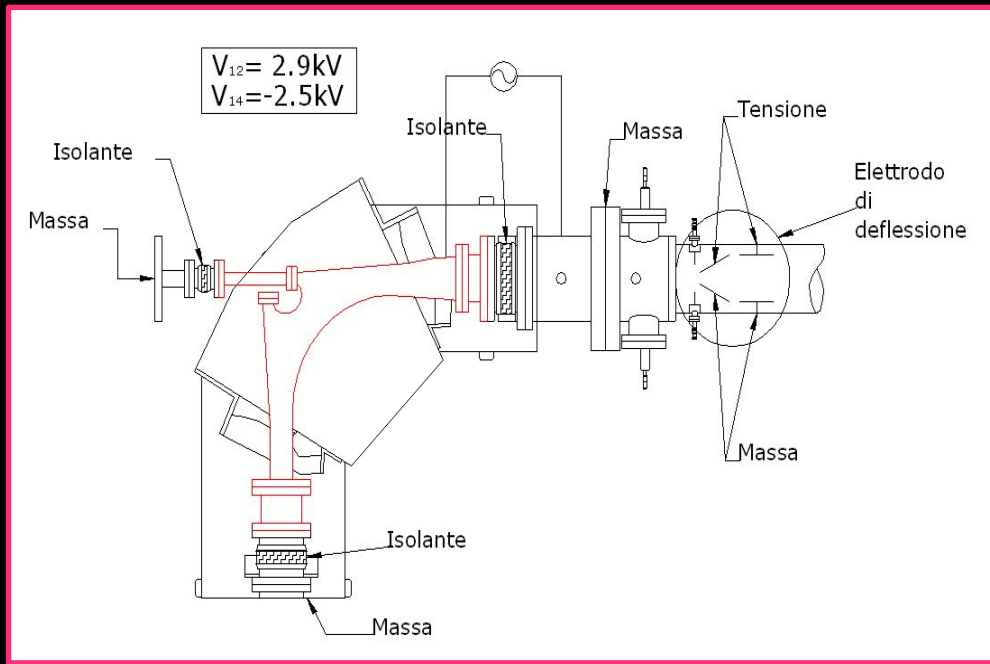
- ✓ Second selection according to **impulse/charge**

$$Br = \frac{\sqrt{2mE}}{q}$$

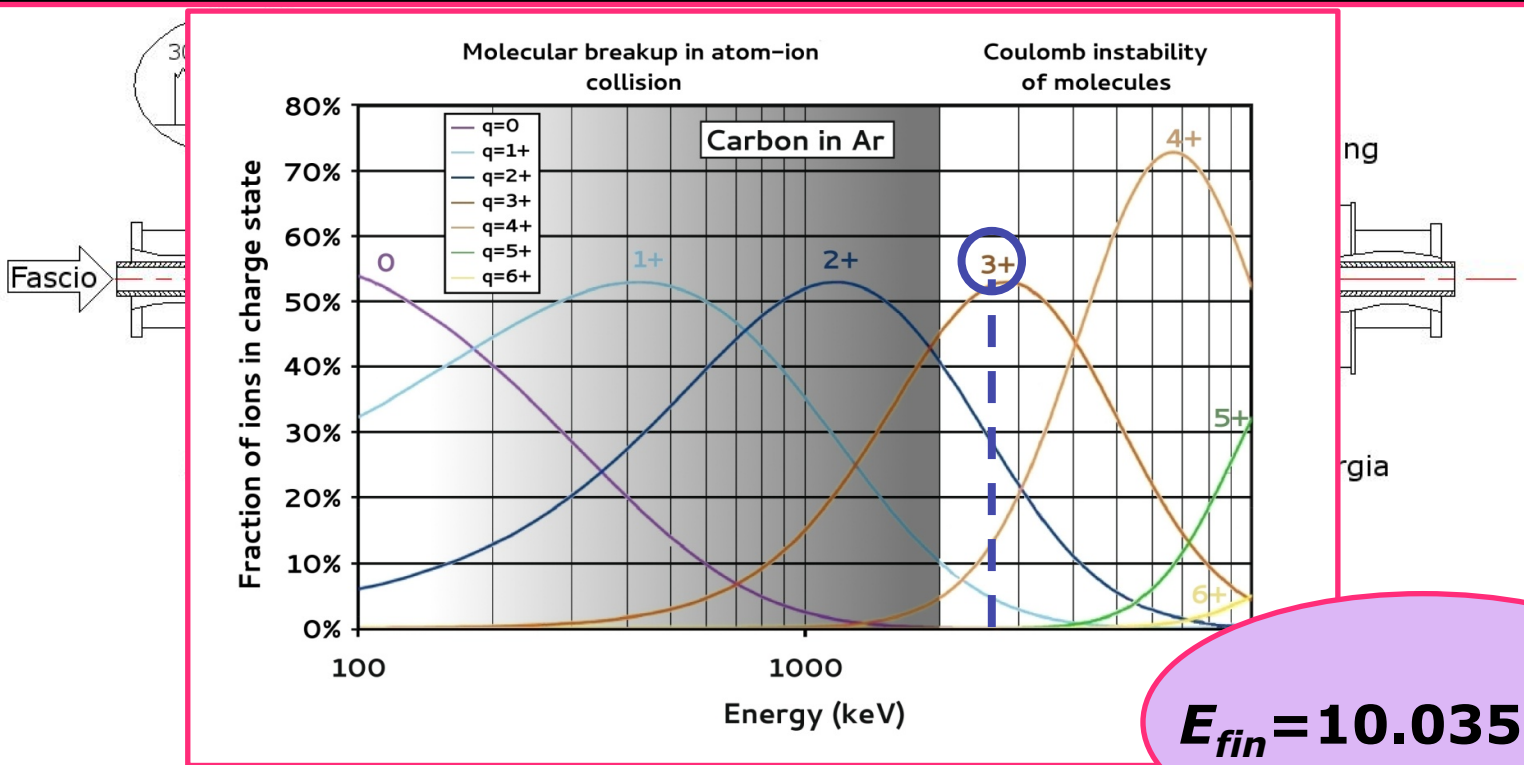
→ to select the three masses 14, 12 and 13

# Sequential injection...

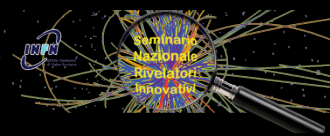
- ✓ The magnetic field is kept fixed to optimize the transmission of a mass ( $m=13$ )
- ✓ The other masses are transmitted by changing their energy inside the magnet chamber



# ...into the accelerator

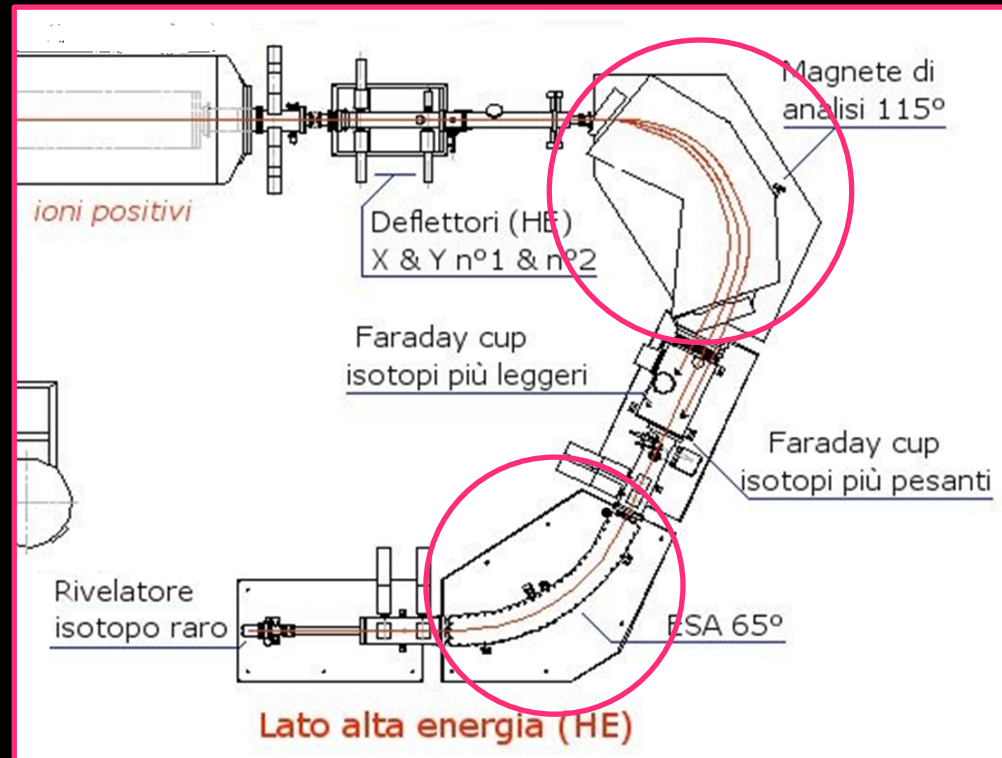


$$E_{fin} = (E_{inj} + e \cdot V_t) \cdot \frac{m_i}{m_{tot}} + q \cdot e \cdot V_t$$



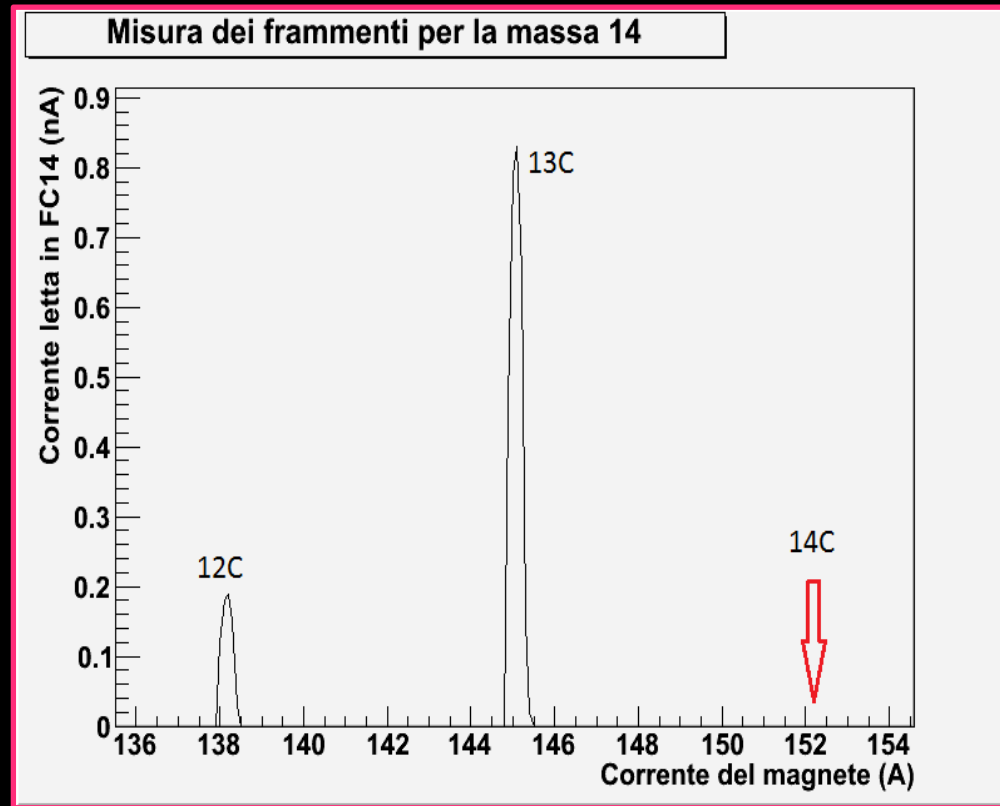
# The analysis on the high energy side

- ✓ First selection according to impulse/charge  
→ **discrimination of  $^{14}\text{C}^{3+}$  ions**
- ✓ Second selection according to energy/charge  
→ rejection of possible residual interferences



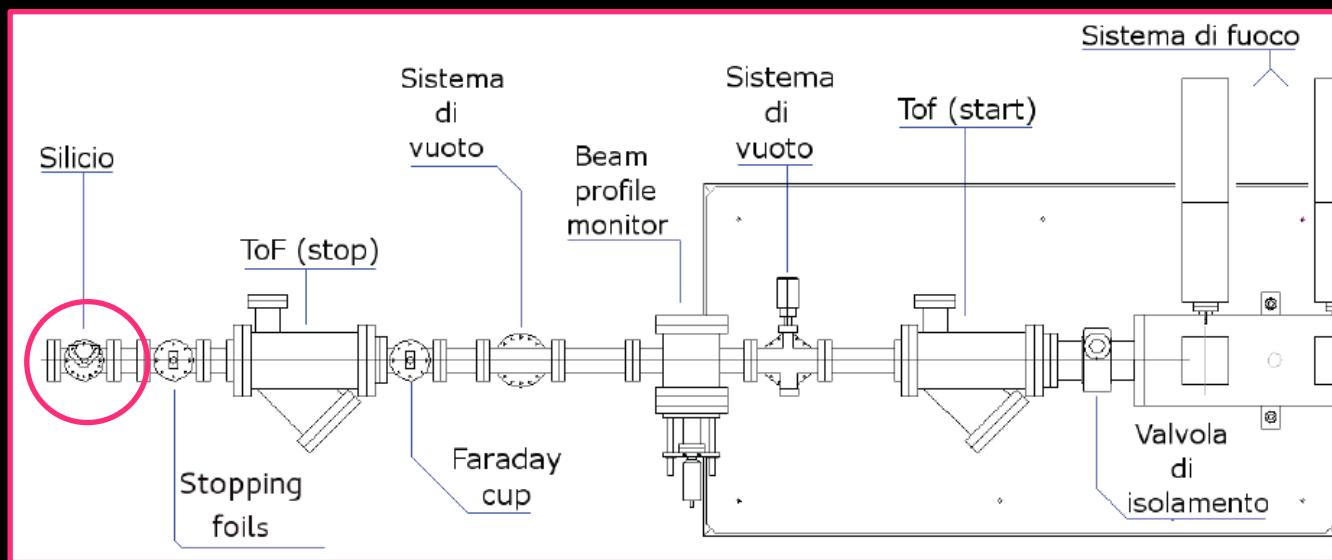
# What happens to molecular isobars?

- ✓ As a consequence of the stripping process, molecules can be split into their atomic constituents





# Counting the radiocarbon ions



Hamamatsu  
Si-  
photodiode

CAEN A422A  
Charge-  
sensitive  
preamplifier

ORTEC 570  
Amplifier

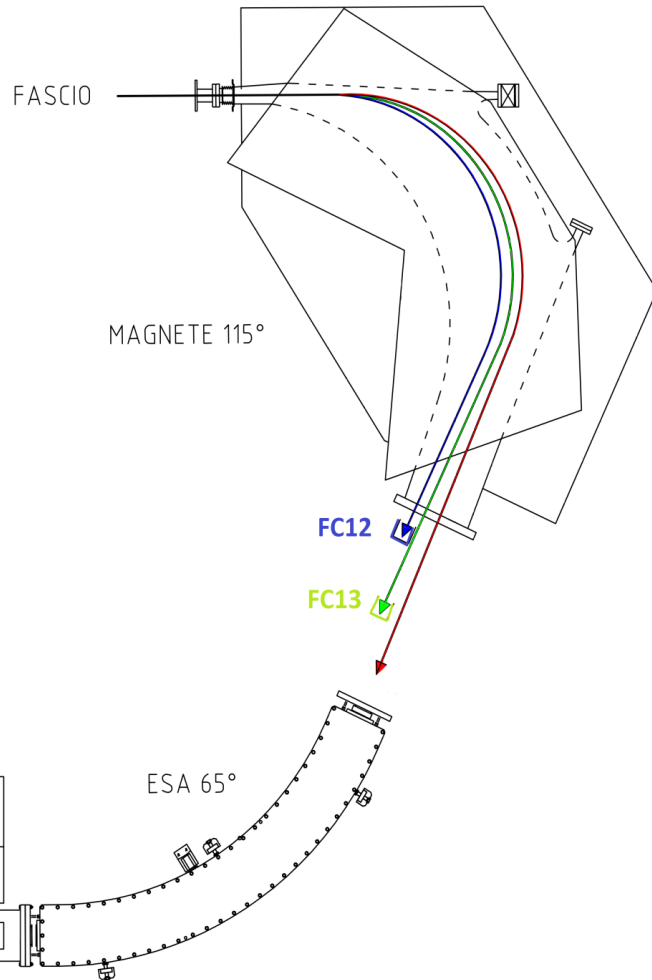
ORTEC 550A  
SCA

ORTEC 661  
Ratemeter

ADC - PC

# What about the stable isotopes?

- ✓  $^{12}\text{C}^{3+}$  and  $^{13}\text{C}^{3+}$  are measured in offset Faraday cups at the exit of the high-energy magnet



Measurement of  $^{12}\text{C}$  →  
measurement of the **isotopic**

$$\text{ratio } \frac{^{12}\text{C}}{^{14}\text{C}}$$

Measurement of  $^{13}\text{C}$  →  
measurement of **isotopic fractionation correction**

# ***The importance of beam transport***

- ✓ The expected  $^{14}\text{C}$  counting rate is low (few tens of Hz at maximum) → the transmission has to be optimized
- ✓ A good analysis of the beam depends on the stability of machine parameters (e.g. energy of the beam, i.e. terminal voltage) → we can exploit feedback systems

***Set focusing elements (Einzel lens and quad doublets) and steerers to optimize the transmission***

***Set electrostatic analysers and magnets to efficiently discriminate the components of the beam***

***Check Faraday cups, Beam Profile Monitors (BPM) and  $^{14}\text{C}$  counting rate to monitor the beam transport***

# Working with the beam

**F4 846B Source panel - BIAMS**

Panel Filing Edit Panel View Diagnostics Machine AMS Math Help

846 Cathode Voltage	7.00	7.0	846 Cathode Current	0.15	
846 Ionizer Current	17.00	17.0	HV2 846	1.3 e-5	mBar
846 Cesium Reservoir	73.00	71	HV3 BI	4.4 e-7	mBar
BI 54° ESA	3941.97	3956	Current Converter	9.69	%
846 Extraction Voltage	28.00	28.0	846 Extraction Current	0.06	mA
846 Einzellens 1 Neg.	20.70	20.7	846 Steerer Y1	-124.00	123

**Current Converter**

BI FC2 100 µA

Current: 9.7 µA

Auto Range

Bypass Dose Control

**Beam Profile Monitor**

2 9.4

**Vacuum Status Overview**

Complete Layout

7.1 e-6 4.4 e-7 1.5 e-6 1.1 e-6 7.8 e-9

860 358

1.3 e-5

Carrousel 846

Argon

Active 9.7 µA

Start F4 846B Source pane... 2:27 PM

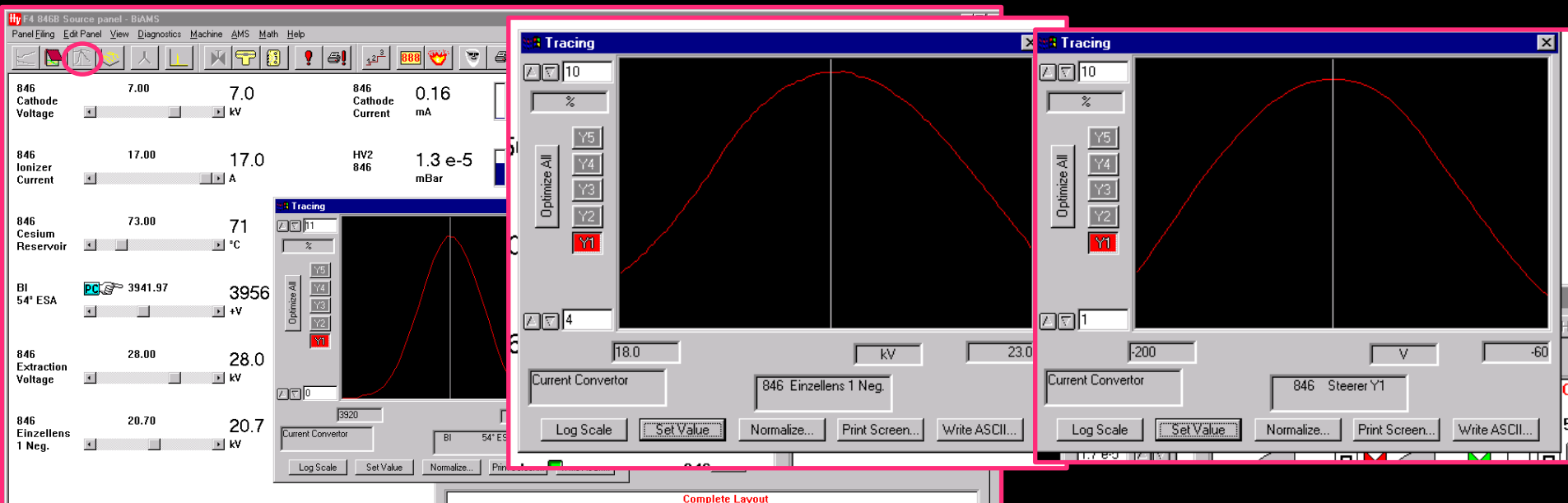
# Beam in FC2

✓ Three elements to set

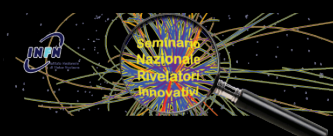
**Electrostatic  
Analyser (ESA)**

**Einzel Lens**

**Steerer Y**

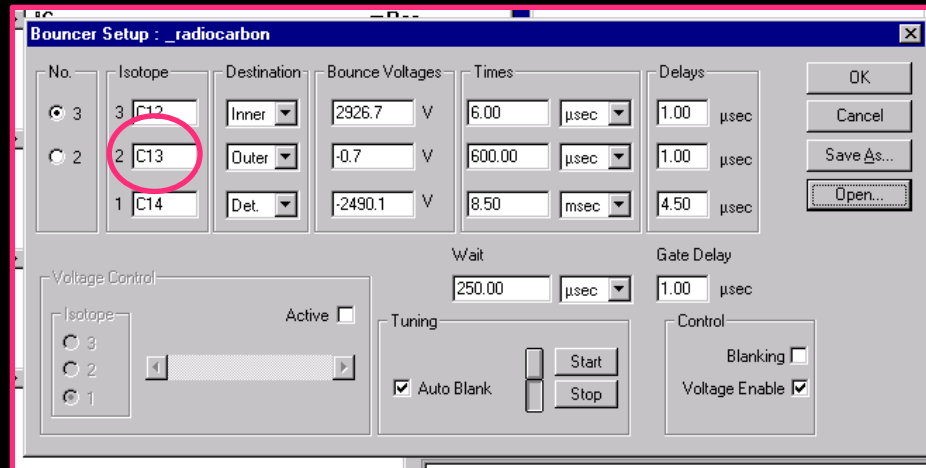


Complete Layout



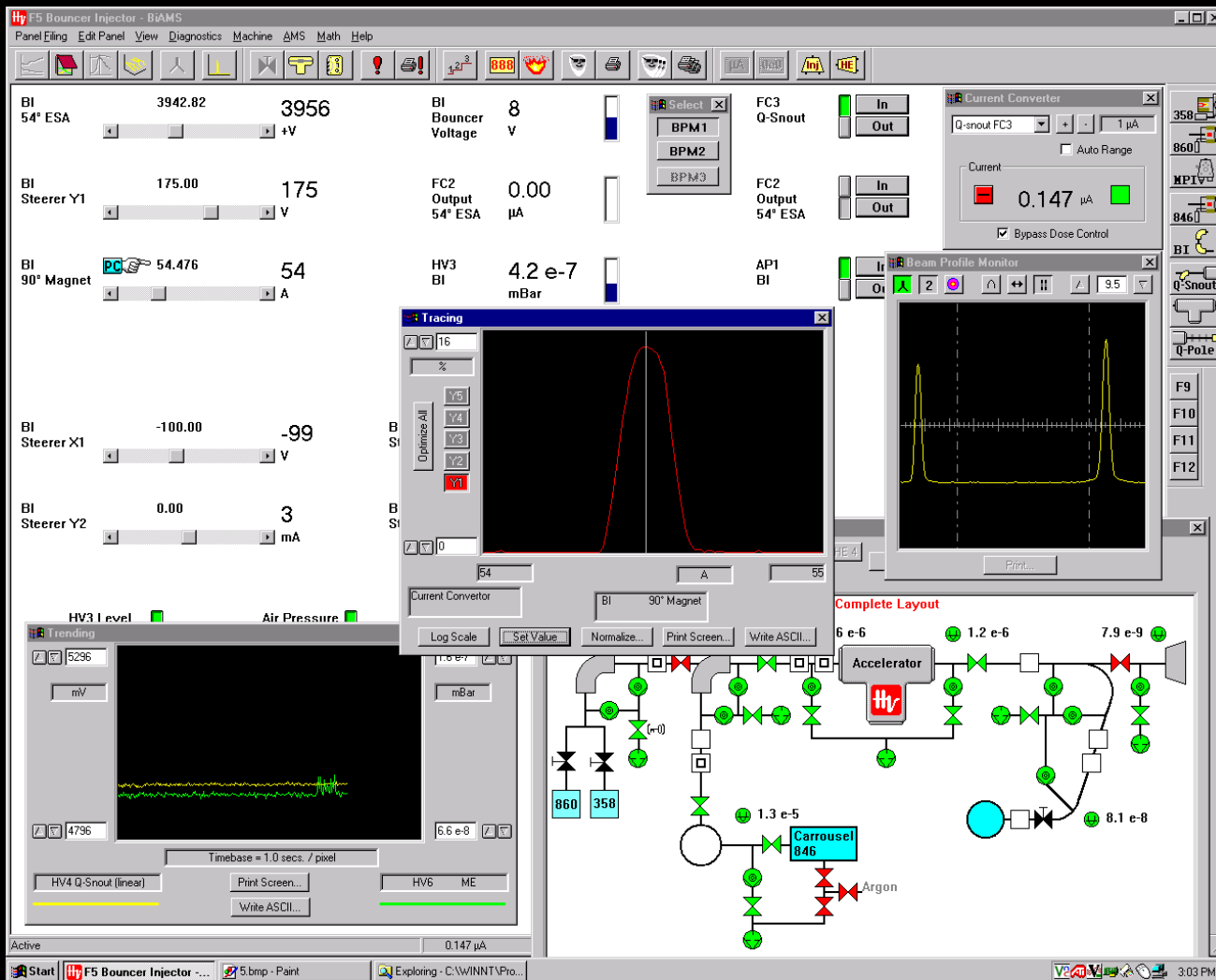
# Beam in FC3

✓ Set the magnet → which mass?



**Optimize the current in the Faraday cup and monitor beam spatial distribution in the BPM**

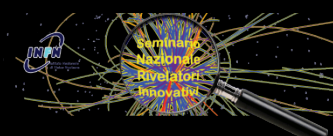
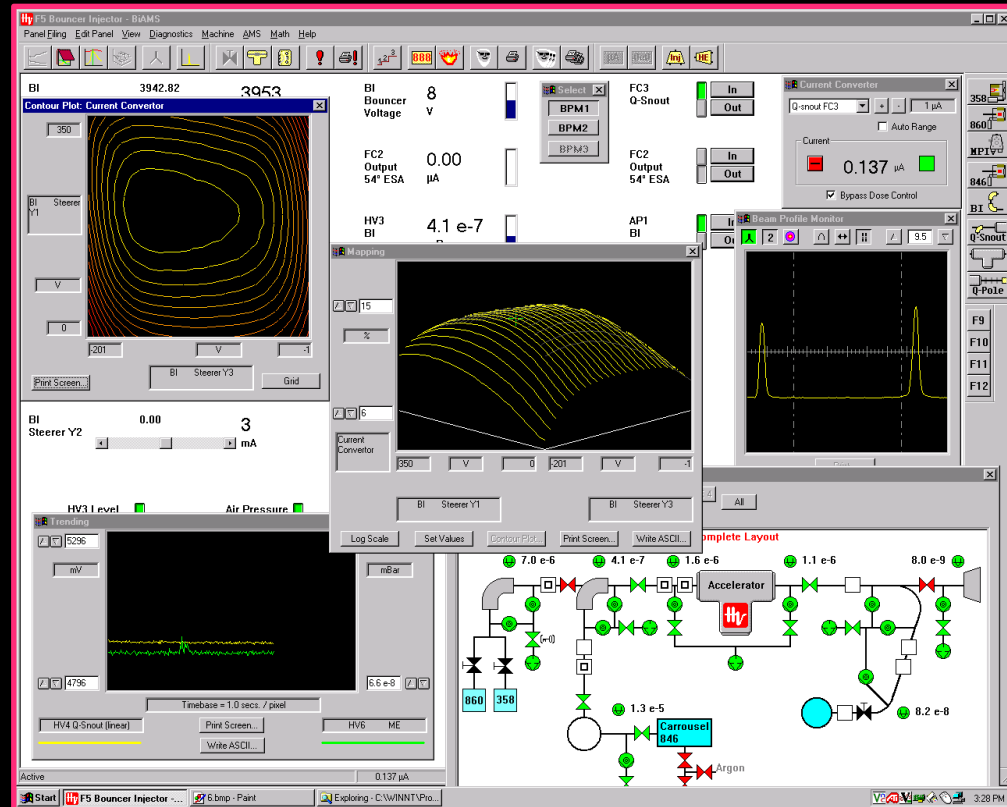
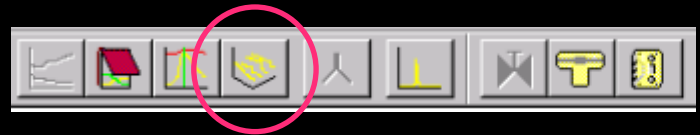
# Beam in FC3





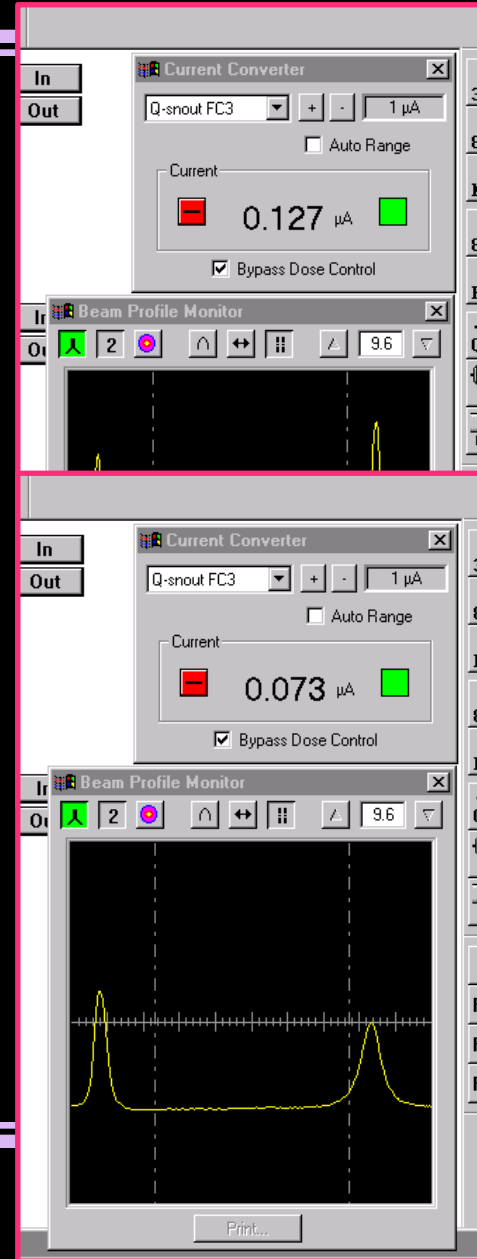
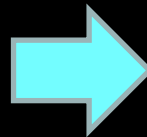
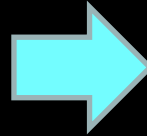
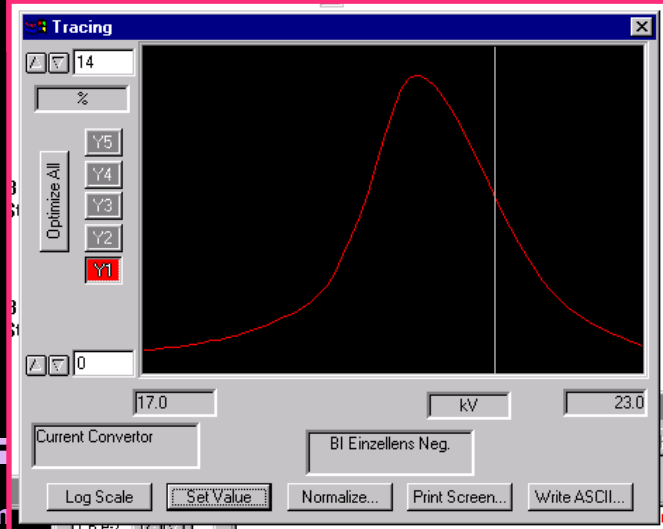
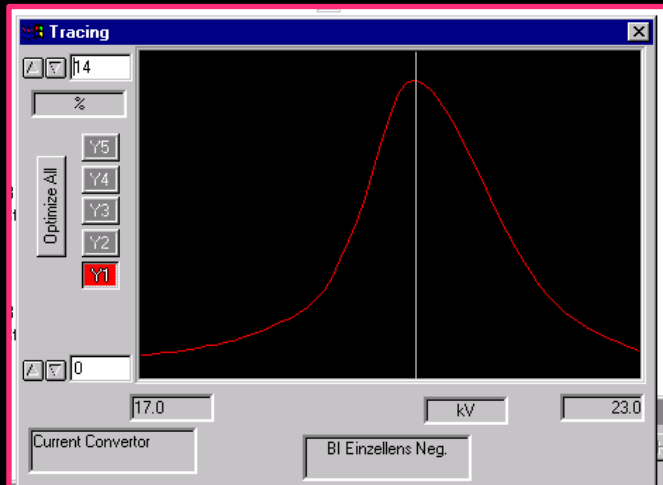
# Beam in FC3

- ✓ A couple of Y steerers
- ✓ A couple of X steerers



# Beam in FC3

- ✓ Einzel lens to focus



# At the exit of the accelerator

- ✓ Roughly set the magnet on the high energy side → monitor  $^{13}\text{C}^{3+}$  current in the dedicated offset Faraday cup

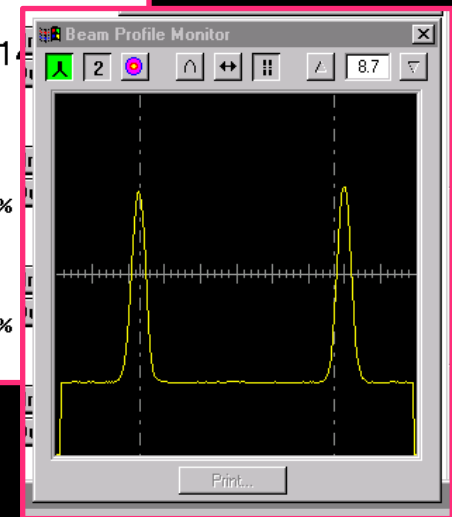
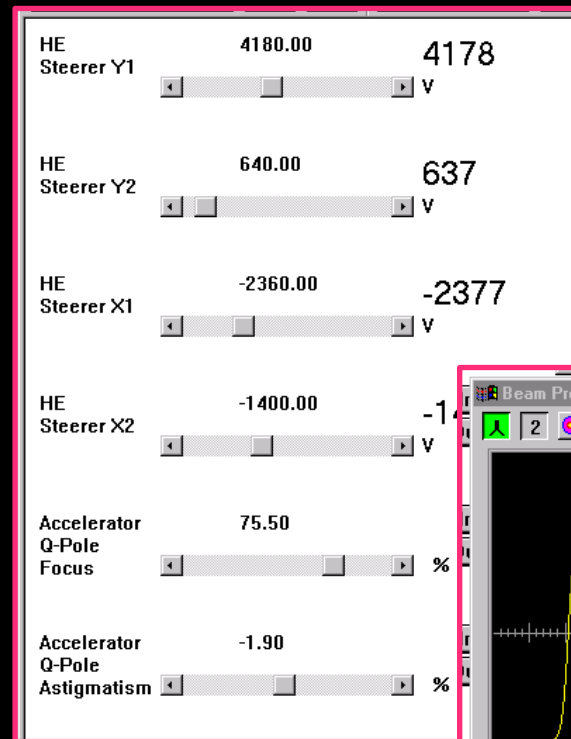
The screenshot displays the control interface for the F9 Multi Element Separation - BiAMS. The interface includes a menu bar (Panel, Filing, Edit Panel, View, Diagnostics, Machine, AMS, Math, Help) and a toolbar with various icons. The main control area features several panels:

- Magnet Control:** ME 115° Magnet slider set to 153 A (circled in red). Other magnet controls include ME 65° ESA (80.00 to 80.0 +kV), ME Q-Pole 1 Focus (0.00 %), ME Q-Pole 1 Astigmatism (0.00 %), Accelerator Q-Pole Focus (74.80 %), and Accelerator Q-Pole Astigmatism (-2.00 %).
- HV Control:** HV6 ME reading of 8.6 e-8 mBar (circled in red).
- Error Monitoring:** ME Slit Error of -0.74 % (circled in red).
- Keithley Readouts:** A window showing two current readings: 7.512e-011 A (Keithley 1) and 5.129e-008 A (Keithley 2) (circled in red).
- Current Converter:** A window showing a current reading of 0.037  $\mu\text{A}$ .
- Beam Profile Monitor:** A window showing a graph with two peaks.

Other visible elements include a 'Select' menu with BPM1, BPM2, and BPM3 options, and a 'Vacuum Status Overview' window at the bottom.

# At the exit of the accelerator

- ✓ Set Q-snout
- ✓ Set quadrupoles (focus and astigmatism) and high energy steerers



# ***Downstream the high-energy magnet***

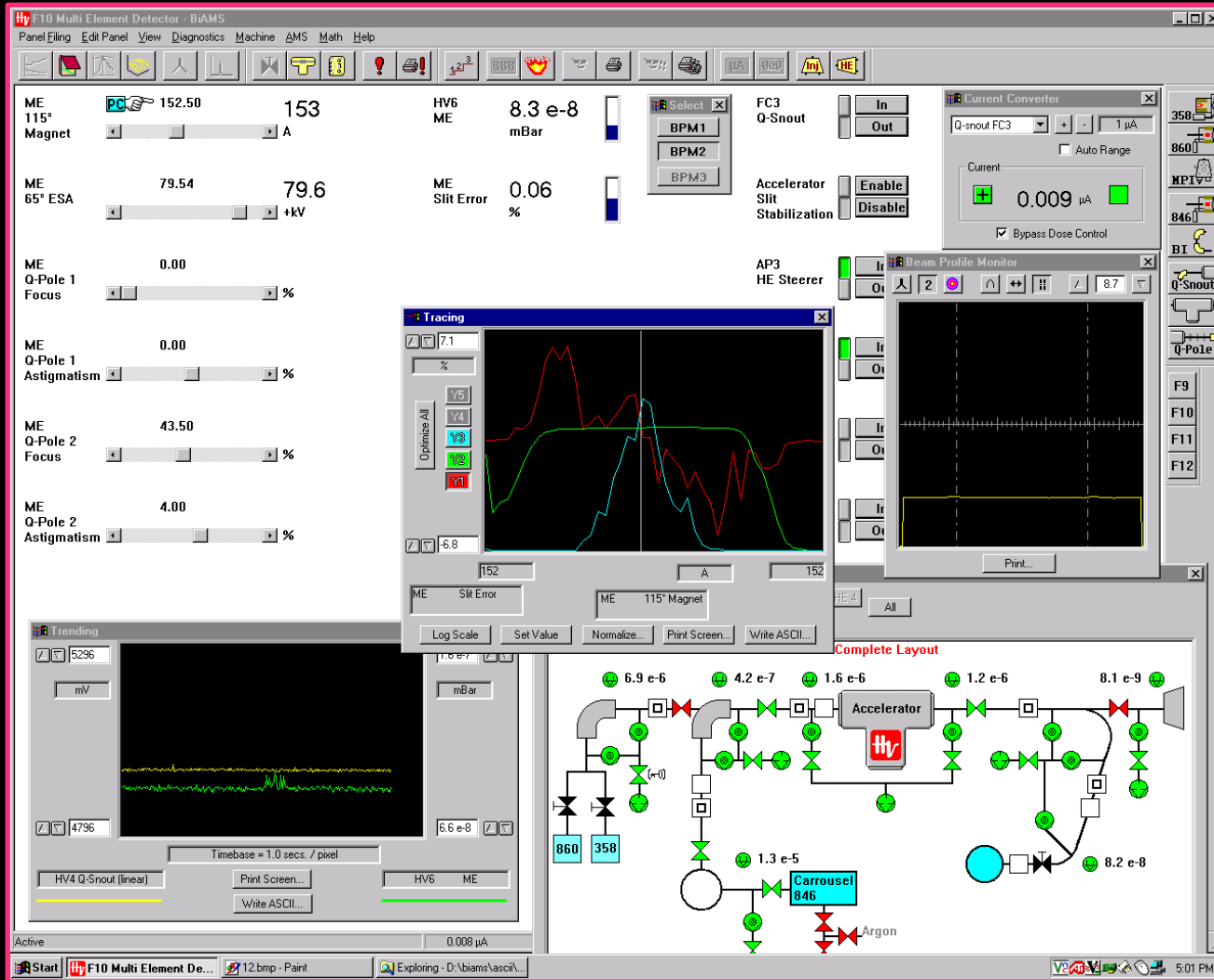
- ✓ Typical beam currents are too low to use Faraday cup or BPM

***Check the counting rate in the  $^{14}\text{C}^{3+}$  silicon detector (roughly, looking at the output signal of SCA)***

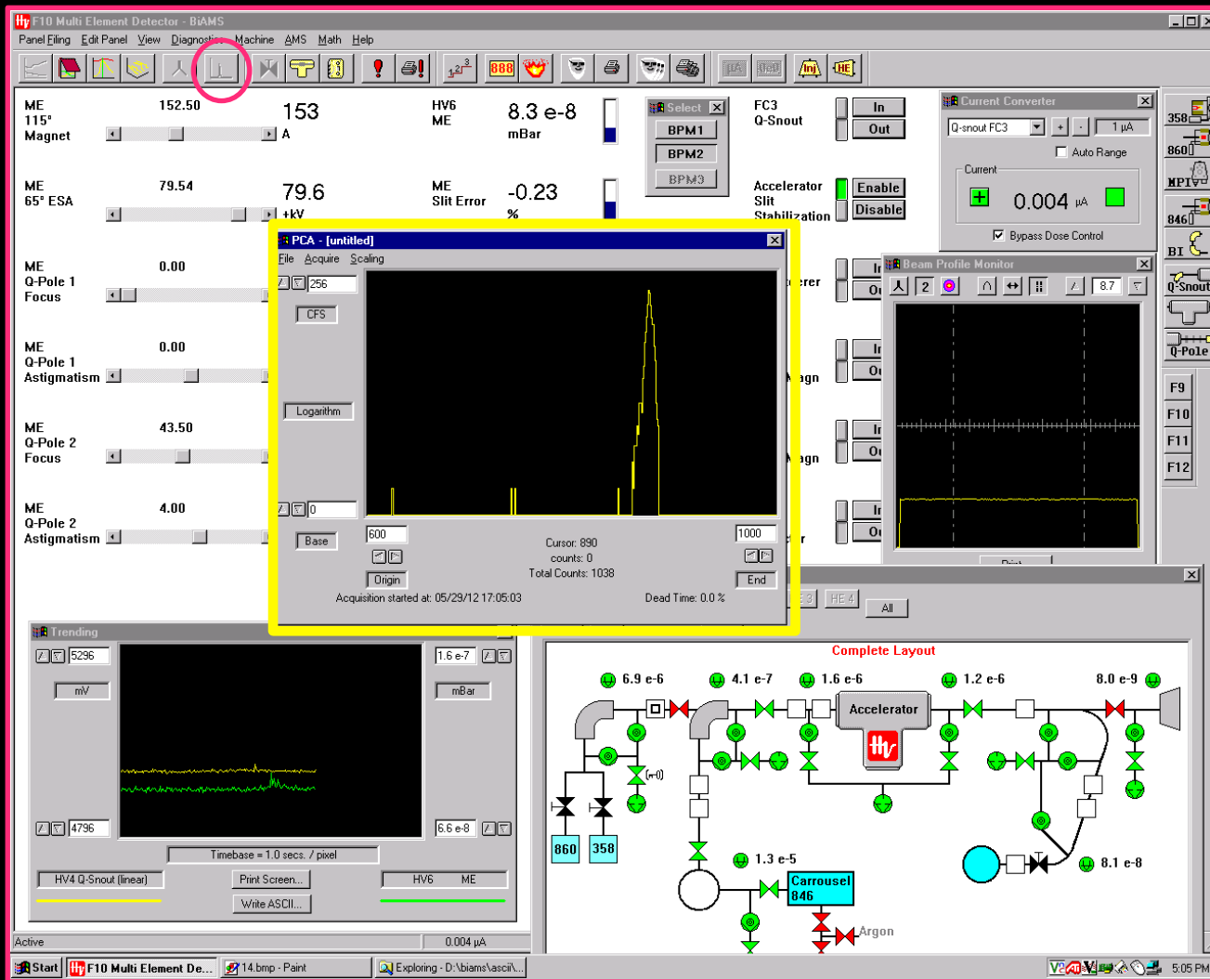
- ✓ Set high-energy electrostatic analyser and quadrupoles



# Final check of the three isotopes



# Our goal





# The AMS measurement

- ✓ Your set of samples to be measured:
  - **Standard** samples → to normalize measured isotopic ratio
  - **Blank** samples → to subtract background counts (due to machine and sample preparation)
  - **Unknown samples**
  
- ✓ What is measured:
  - **$^{14}\text{C}$  counts**
  - **$^{12}\text{C}$  and  $^{13}\text{C}$  currents**



# Data Analysis

$${}^{14}R_{norm,sample} (pMC) = {}^{14}R_{cert,std} (pMC) \cdot \left( \frac{{}^{13}R_{meas,std}}{{}^{13}R_{meas,sample}} \right)^2 \cdot \left( \frac{{}^{14}R_{meas,sample} - {}^{14}R_{meas,blk}}{{}^{14}R_{meas,std} - {}^{14}R_{meas,blk}} \right)$$

**Certified  
concentration  
of standard**

**Isotopic  
fractionation  
correction**

**Normalization  
to standard and  
subtraction of  
background**

$$t_{RC} (yrs BP) = 8033 yrs \cdot \ln \left( \frac{100 pMC}{{}^{14}R_{norm,sample}} \right)$$

**Calibration**

**Most probable  
calendar age**

# Lab practice

<b>14:30 – 15:40</b>	<i>Beam transport and optimization of measurement parameters</i>
<b>15:40 – 15:50</b>	<i>Start of AMS measurement</i>
<b>15:50 – 16:10</b>	<i>Sample preparation laboratory</i>
<b>16:10 – 17:00</b>	<i>Improvements on detection of molecular isobars</i>
<b>17:00 – 18:00</b>	<i>Data analysis</i>

***Have a good job!***

