

Studies of the Reaction $^{48}\text{Ca} + ^{238}\text{U}$ @ BGS

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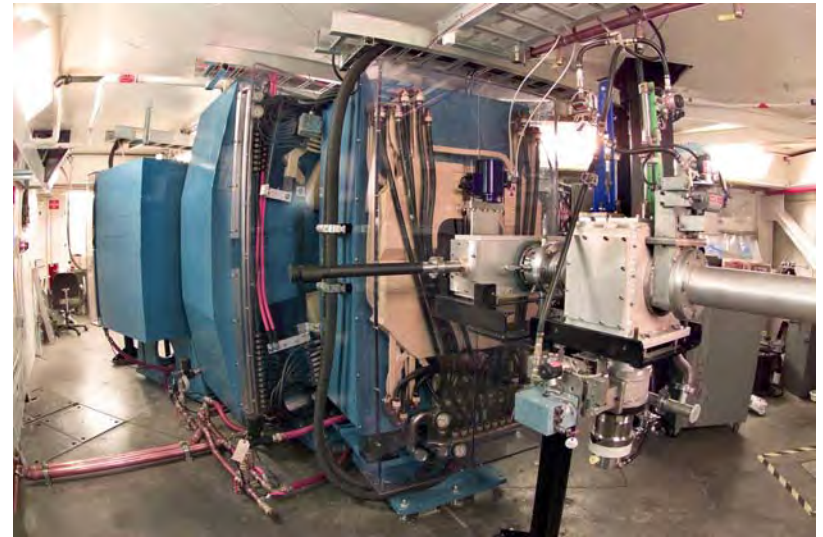
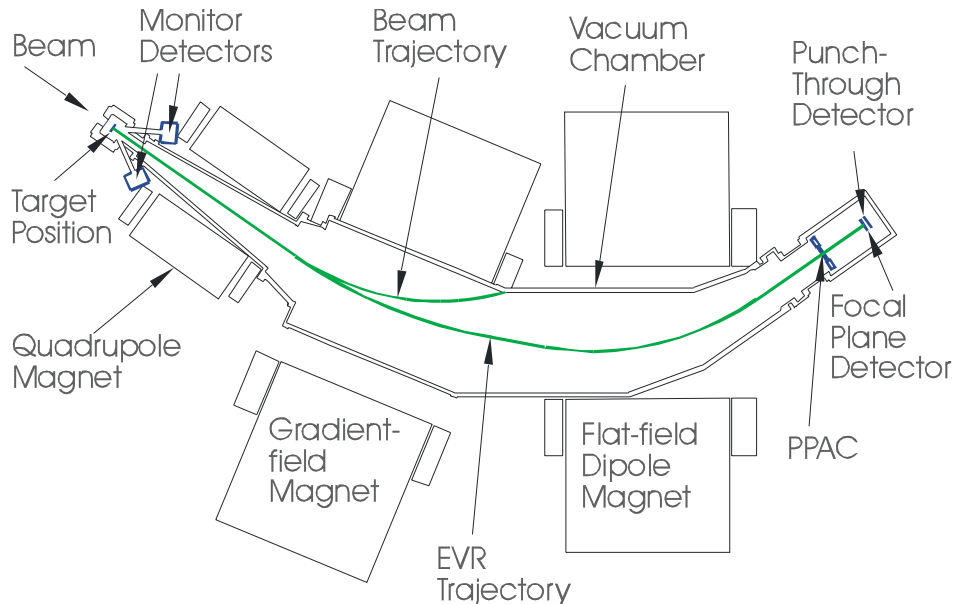
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(There has been NO INDEPENDENT CONFIRMATION of *ANY* isotope with $A > 277$)

Goal: **Independent confirmation** of *ONE* of the DUBNA SHE results

Berkeley Gas-filled Separator (BGS)

- Construction “completed” fall 1999
- Recycled Bevalac magnets
- Innovative design gives $\Omega=45\text{msr}$
- 70° bend gives superior separation
- ~ 1 mBar He fill gives full momentum and charge acceptance



- Beam rejection up to 10^{15}
- Transit time $\sim \mu\text{s}$
- Rotating target allows beam intensities up to μA range
- **Beam intensity, target thickness, and efficiency give 1 event/(picobarn*week)**

$^{283}_{112}$ history $^{238}\text{U}(^{48}\text{Ca},3\text{n})^{283}_{112}$

1999: **Vassilissa** 2 SF observed **5.6 pb** @ **231 ± 3 MeV** **half-life = 81 sec**
No SF observed **< 4.0 pb** @ **238 ± 3 MeV**

1999: **Vassilissa** SF observed after 10.29-MeV α -decay of $^{287}_{114}$ **new $^{283}_{112}$ half-life = 3 min**

2001: **BGS** No SF observed **< 1.6 pb** @ **228-234 MeV** $B_p(\text{in He}) = 2.19\text{-}2.31 \text{ Tm}$

2001: **Dubna** Chemistry **~2.0 pb** fissions could be long-lived $^{283}_{112}$ with Rn-like Chemistry

2002: **BGS** No SF observed **< 0.8 pb** @ **228-234 MeV** $B_p(\text{in He}) = 2.19\text{-}2.31 \text{ Tm}$

2003: **PSI@GSI** Hg-Rn chemistry gave inconclusive result (sensitive only to long-lived SF activity)

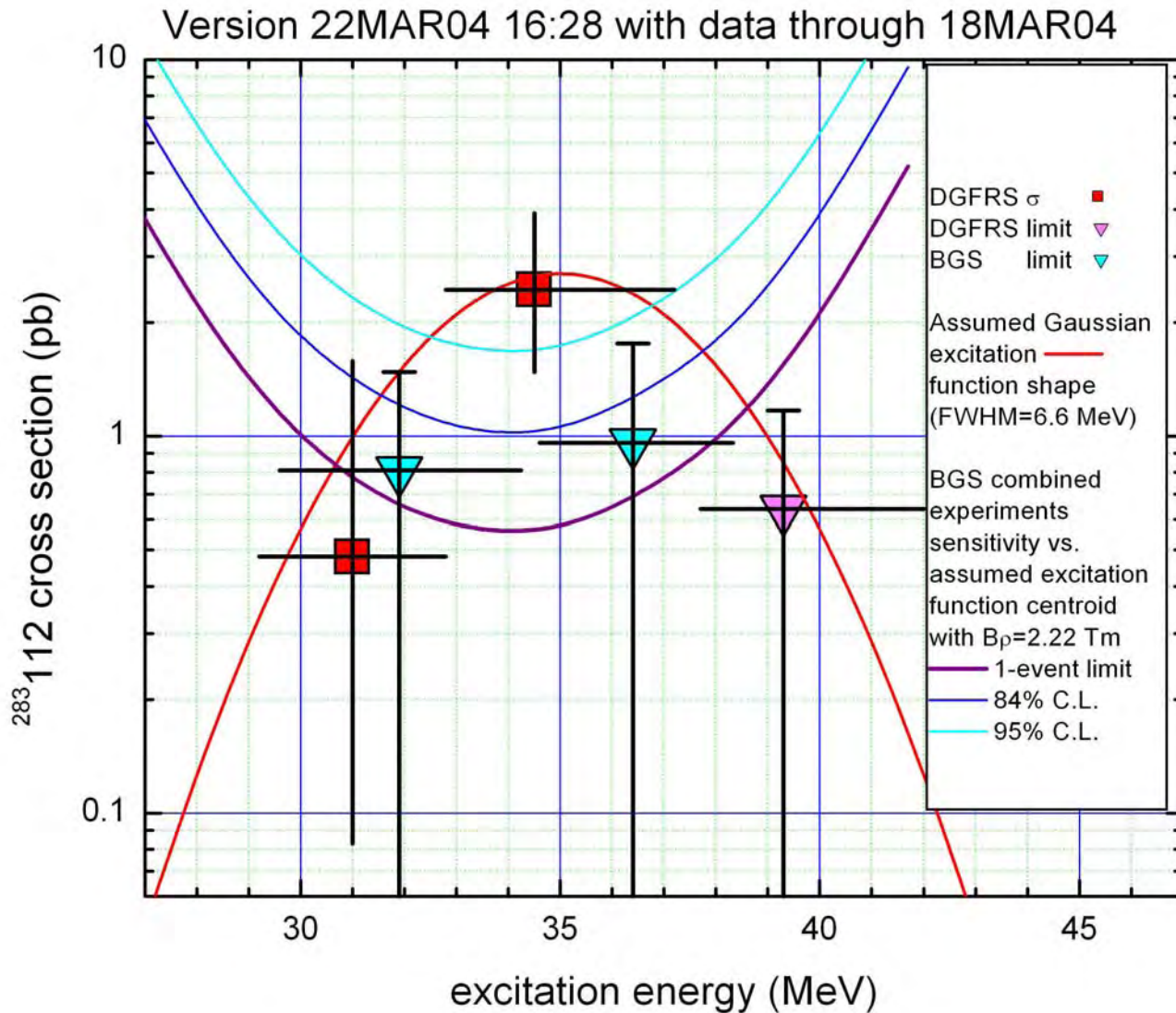
2003: **DGFRS** 9.5-MeV α after 10.0-MeV α -decay of $^{287}_{114}$ **half-life ~ 5s**
seen in both $^{244}\text{Pu}(^{48}\text{Ca},5\text{n})^{287}_{114}$ and $^{242}\text{Pu}(^{48}\text{Ca},3\text{n})$ reactions

2003: **Vassilissa** No SF observed **< 1.2 pb** @ **231±3 MeV**
2 SF observed **~ 4.0 pb** @ **234± 3 MeV** **new $^{283}_{112}$ half-life = 5.1 min**

2004: **DGFRS** 9.5 MeV α **~1.3 pb** @ **231±3 MeV** **half-life ~4s**
9.5 MeV α **~2.5 pb** @ **234±3 MeV** **half-life ~ 4s**
None observed **<1.3 pb** @ **240±3 MeV**

2004: **BGS** No SF observed **< 0.96 pb** @ **233-238 MeV** $B_p(\text{in He}) = 2.19\text{-}2.31 \text{ Tm}$

$^{283}\text{112}$ Cross Sections and Upper Limits



Why Don't We see 4-s 9.54-MeV $^{283}\text{112}$?

Is there a problem with the BGS targets?

Is there a systematic difference in beam energies from the accelerators at LBNL and Dubna?

Do we have a different beam energy in our uranium targets due to errors in energy loss calculations?

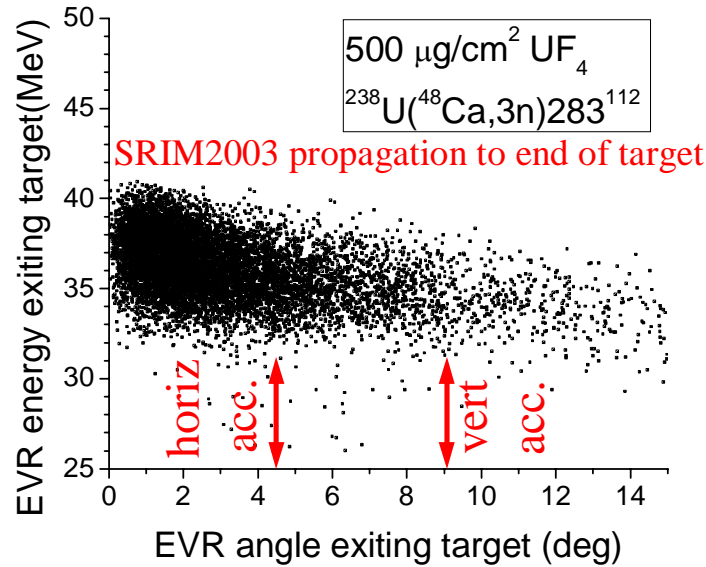
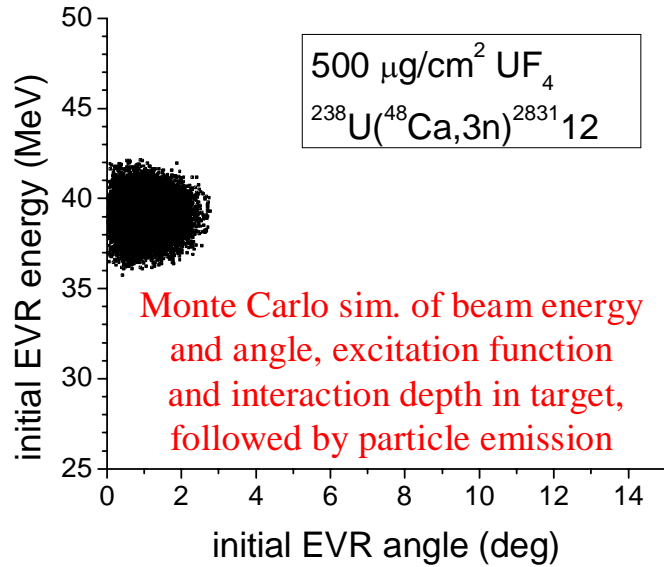
Did we run the BGS at the wrong magnet settings?

Are the UF₄ Targets Any Good?

Targets are 475-611 μg/cm² UF₄ evaporated onto 2-μm Al foils. This thickness is good for the BGS.

Are the UF₄ Targets Any Good?

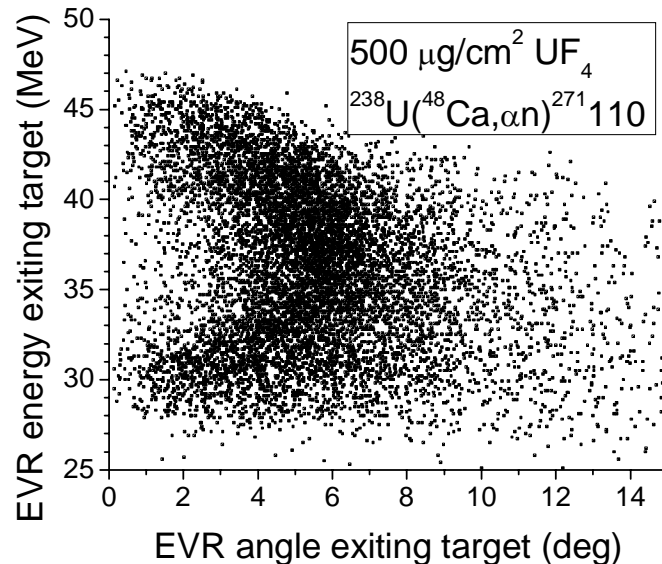
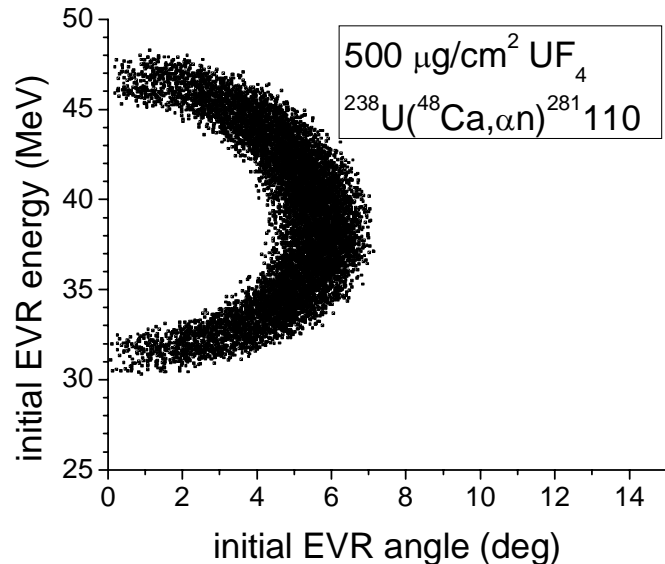
Are the 475-611 μg/cm² targets too thick?



Monte Carlo sim. of particle trajectories in gas-filled magnets

BGS efficiency for $^{238}\text{U}(^{48}\text{Ca},3n)^{283}\text{112}$

59%



BGS efficiency for $^{238}\text{U}(^{48}\text{Ca},\alpha n)^{281}\text{Ds}$

42%

Are the UF₄ Targets Any Good?

Targets are 475-611 μg/cm² UF₄ evaporated onto 2-μm Al foils. This thickness is good for the BGS.

On two sets of targets, some of the UF₄ flaked off the Al foils during the first minutes of irradiation (with low-intensity ⁴⁸Ca beams).

Luminosity is determined by # of Rutherford-scattered ⁴⁸Ca ions . . .
The cross section limits given are correct.

α-spectroscopy of the ²³⁸U shows no large change in the thickness of the UF₄ layer.

α-particle energy loss measurements indicate that there is no large change in of UF₄ thickness or Al thickness during the experiments.

Atomic Force Microscopy shows a change in the UF₄ structure.
Thickness variations are within acceptable limits.

Conclusion: Targets are good (although not perfect)

Why Don't We see 5-s 9.5-MeV $^{283}112$?

Is there a problem with the BGS targets?

Yes, but luminosity is determined with Rutherford-scattered beam.

Is there a systematic difference in beam energies from the accelerators at LBNL and Dubna?

No, $^{48}\text{Ca} + ^{208}\text{Pb}$ excitation functions match to within 2 MeV.

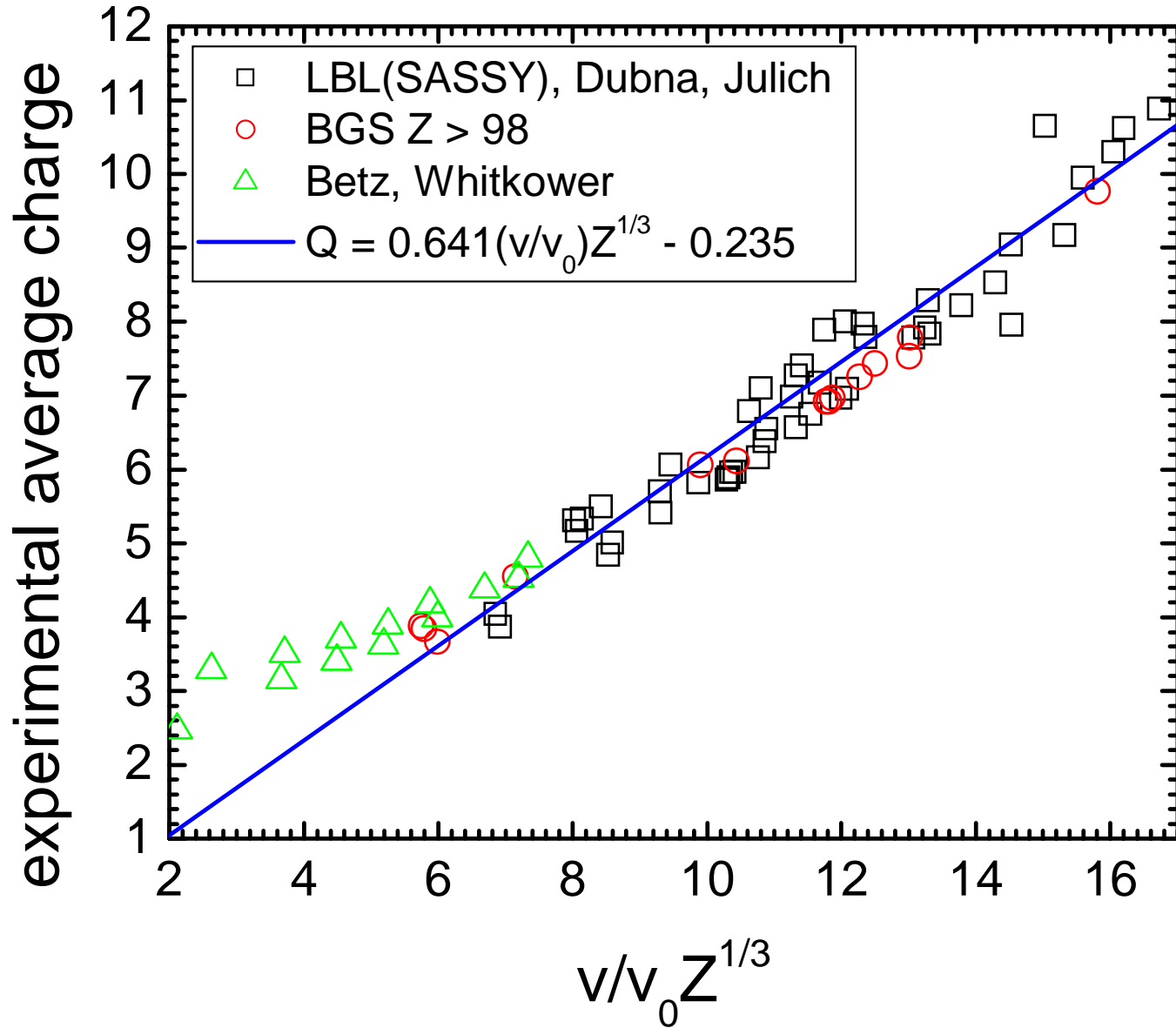
Do we have a different beam energy in our uranium targets due to errors in energy loss calculations?

No, pulse-height of Rutherford-scattered ^{48}Ca is within ~ 1 MeV.

Did we run the BGS at the wrong magnet settings?

What is the $^{283}112$ magnetic rigidity?

Back to basics ...



Back in 1948, Neils Bohr suggested a

$q = vZ^{1/3}$ dependence

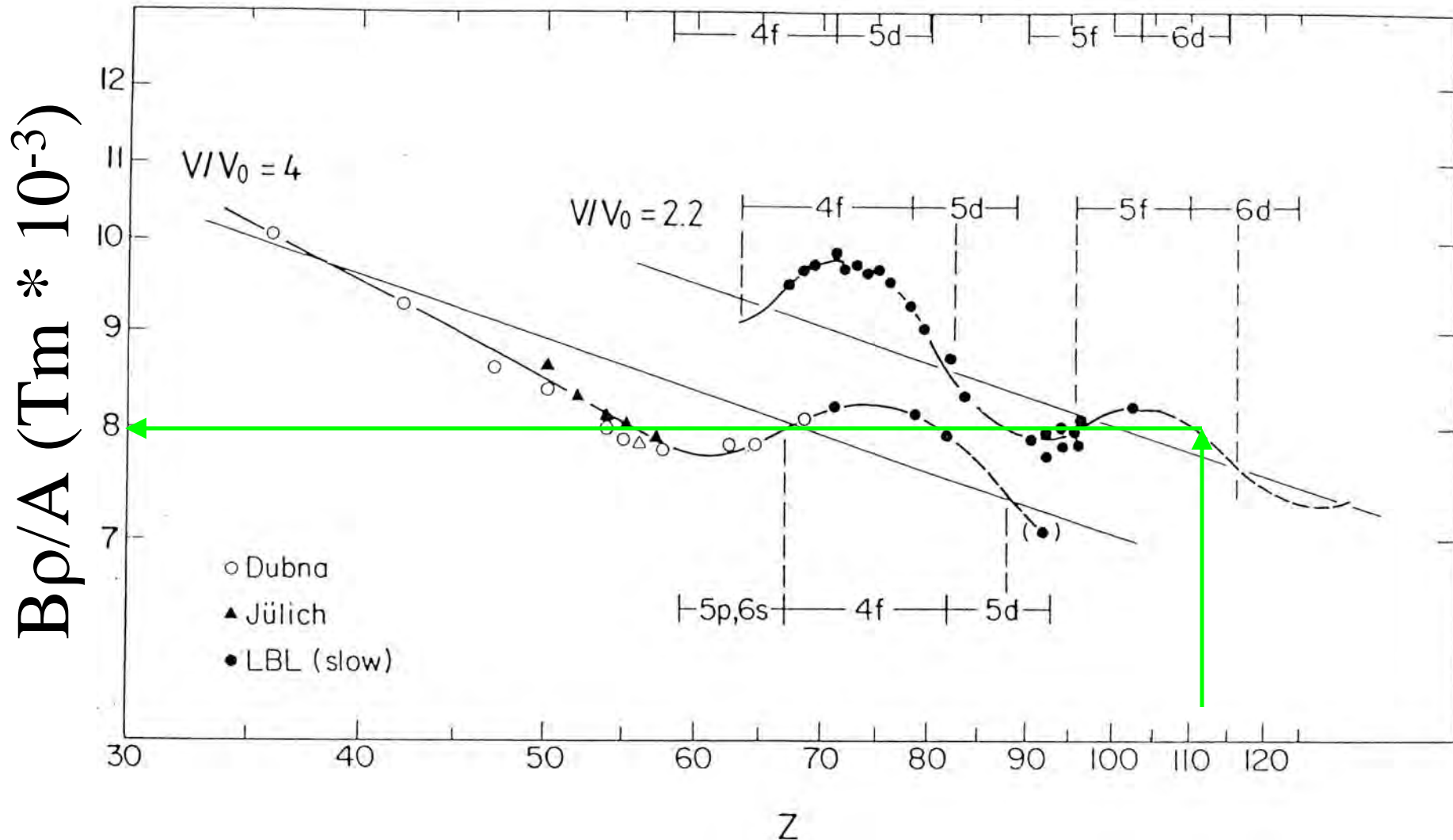
This fit shows much scatter. **Deviations are +/- 10%**. Can this be understood in terms of the electronic shell structure of the stripped ions?

What is the $^{283}112$ magnetic rigidity?

According to Ghiorso and Armbruster . . .

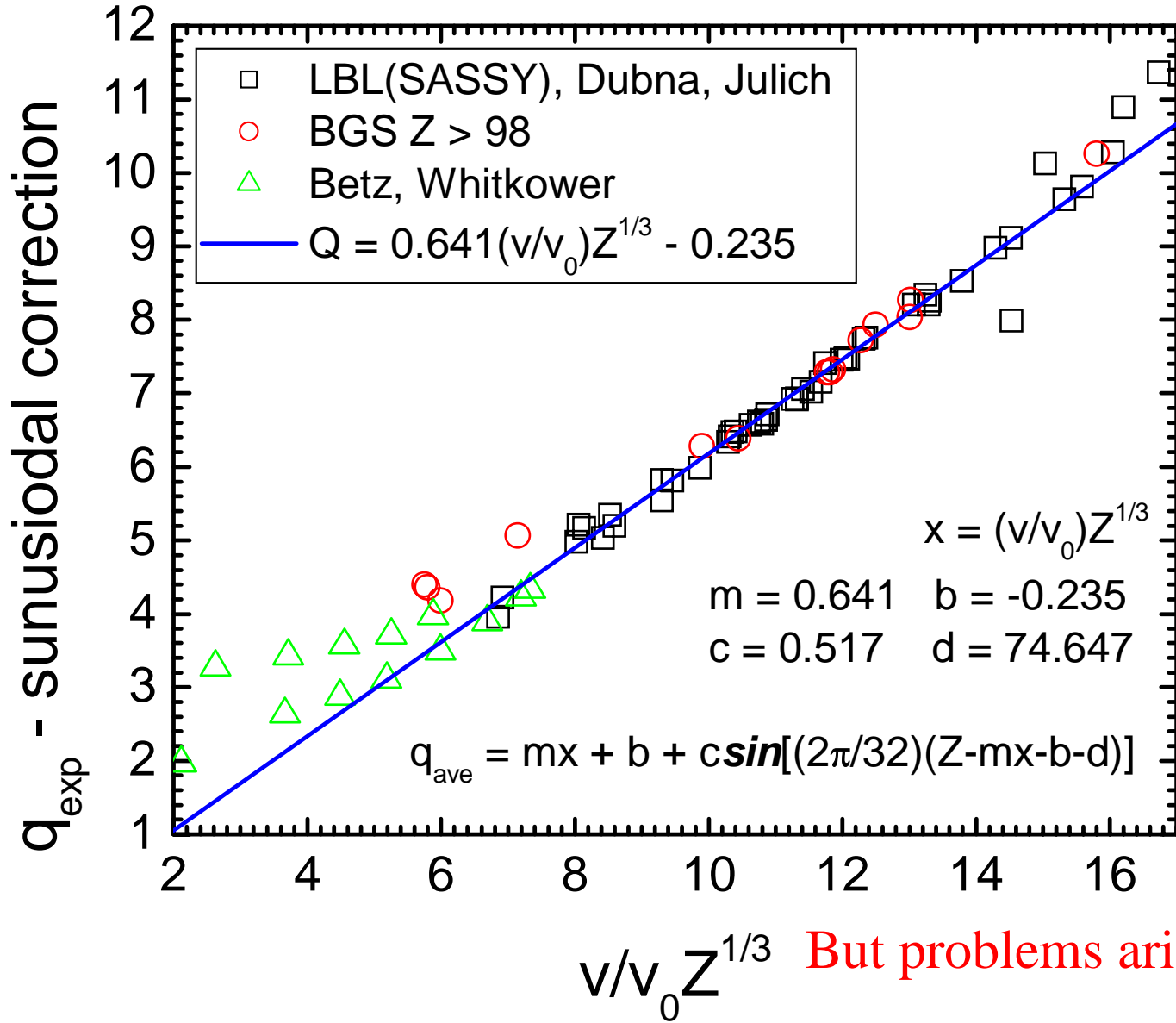
$$B\rho/A = 7.96 \quad B\rho = 2.25 \text{ Tm}$$

Clearly, the electronic shell structure of the stripped ion is important



What is the $^{283}112$ magnetic rigidity?

The Armbruster/Ghiorso plot suggests a sinusoidal correction . . .



Semi-empirical understanding of why this works:

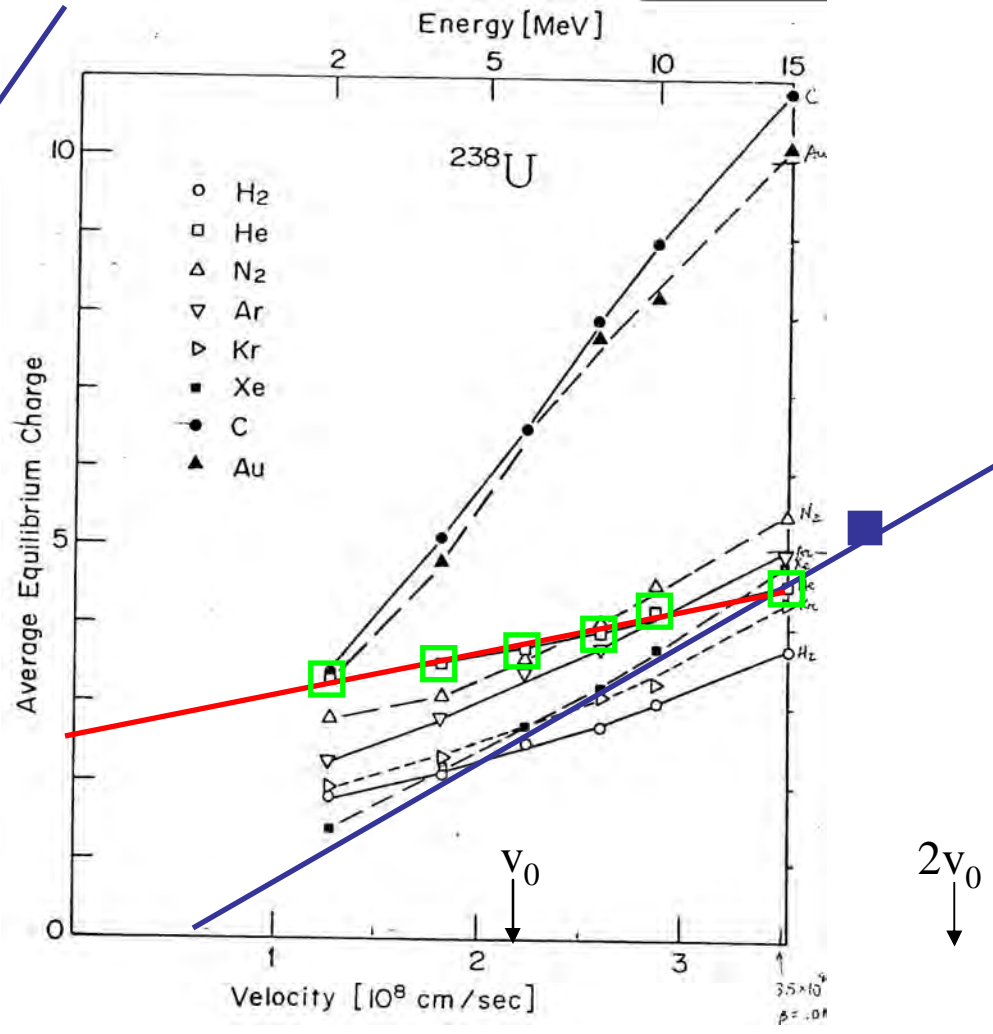
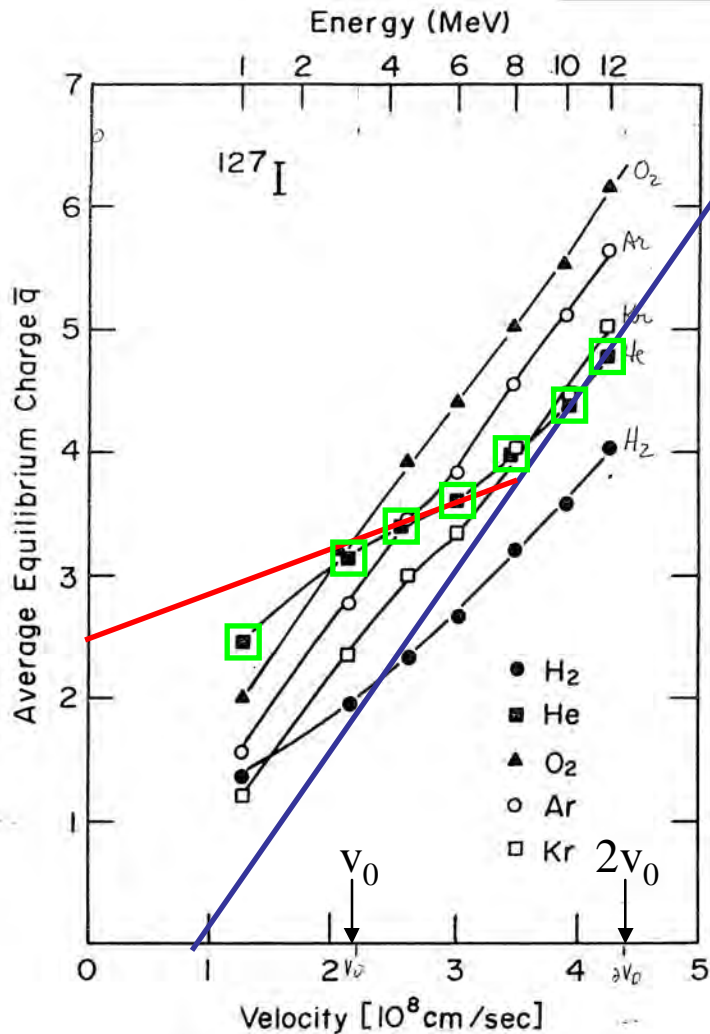
If the stripped ion is in an f-orbital, the most loosely bound electrons are inner electrons, and are less available for stripping by the gas, giving a lower q .

If the stripped ion is in a p-orbital, the most loosely bound electrons are outer electrons, and are readily available for stripping by the gas, giving a higher q .

But problems arise at low velocities!

What is the $^{283}112$ magnetic rigidity?

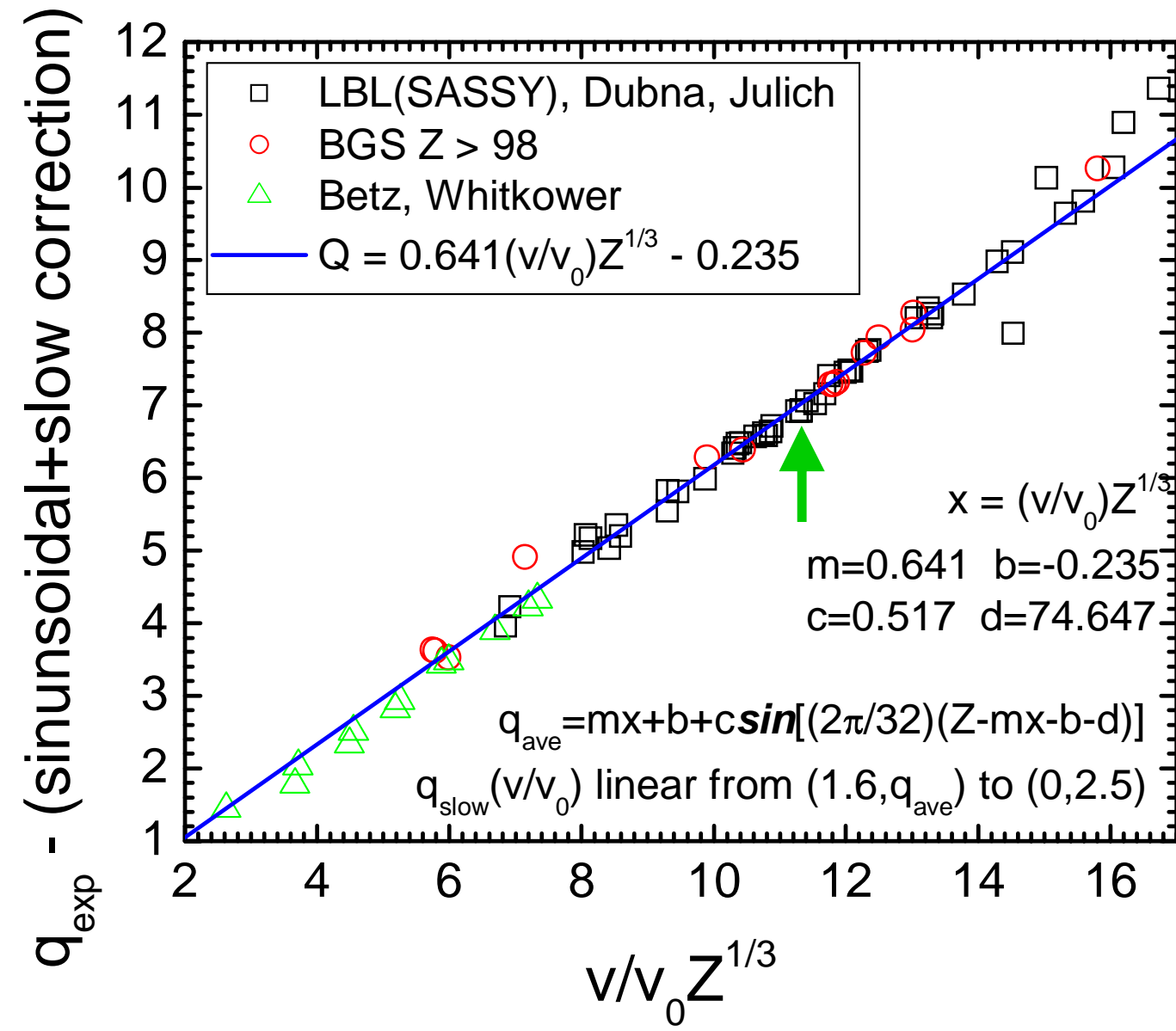
Iodine and uranium data show a break at $v = 1.6v_0$



The red lines trend toward $q = 2.5$ at $v = 0$ because the first of ionization potential of He is 25 eV. This is usually between the second and third ionization potentials of heavy elements.

What is the $^{283}112$ magnetic rigidity?

Putting it all together . . .



$v/v_0 = 2.35$ ($\gg 1.6$)
no slow EVR correction

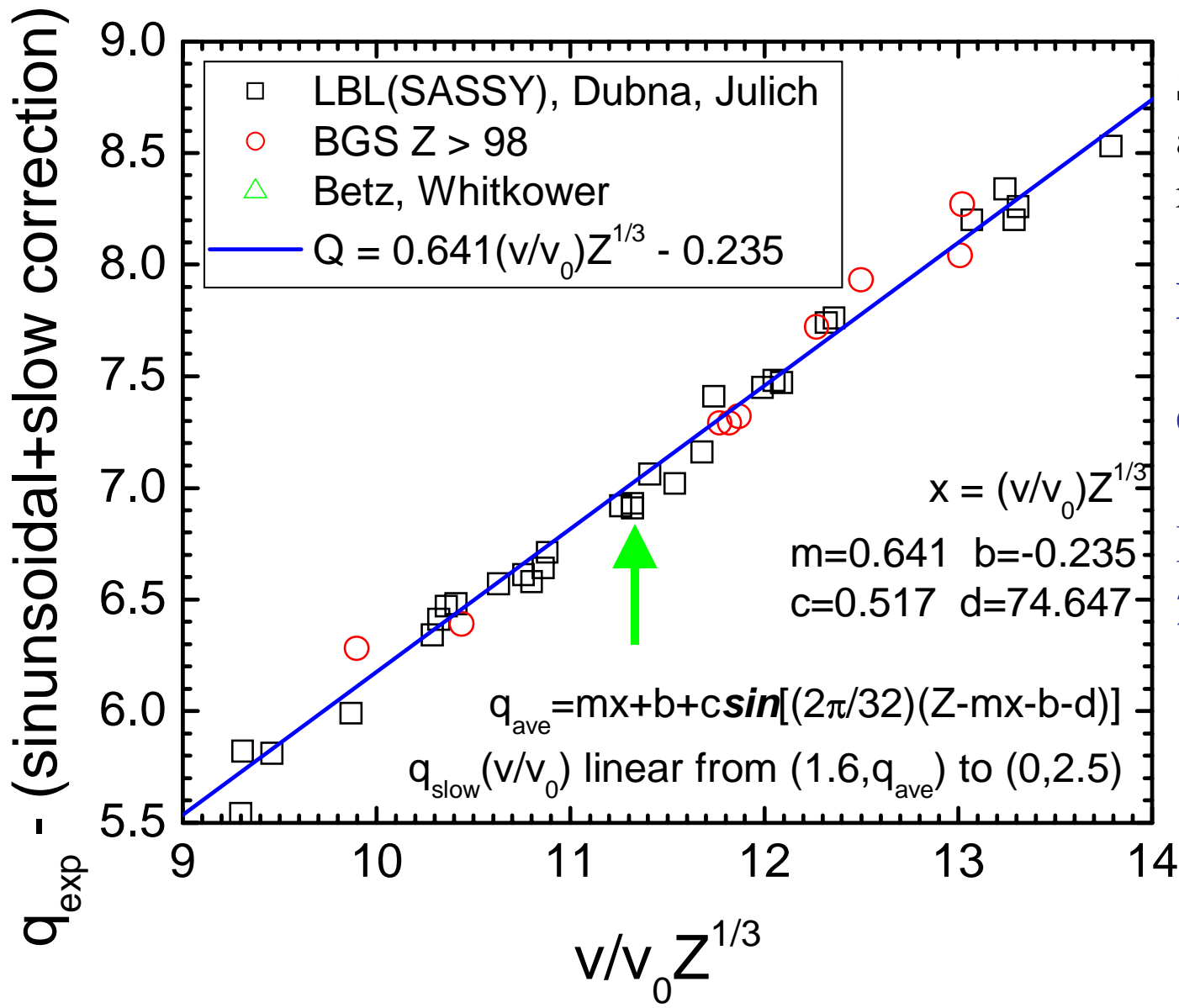
$(v/v_0)Z^{1/3} = 11.33$
gives $q = 6.86$
in the region of best fit
to q vs. $(v/v_0)Z^{1/3}$ data

$B\rho = 2.20$ Tm

compares well with
RIKEN and SASSY
predictions of
2.17 Tm and 2.25 Tm

What is the $^{283}112$ magnetic rigidity?

Estimating the uncertainty in the $^{283}112$ $B\rho$ prediction



Standard deviation about the fit in this region is 0.103

For our $^{283}112$:

$$q = 6.86 \pm 0.103$$

$$B\rho = 2.20 \pm 0.032 \text{ Tm}$$

What is the $^{283}112$ magnetic rigidity?

Effect of the uncertainty in the $^{283}112$ $B\rho$ prediction

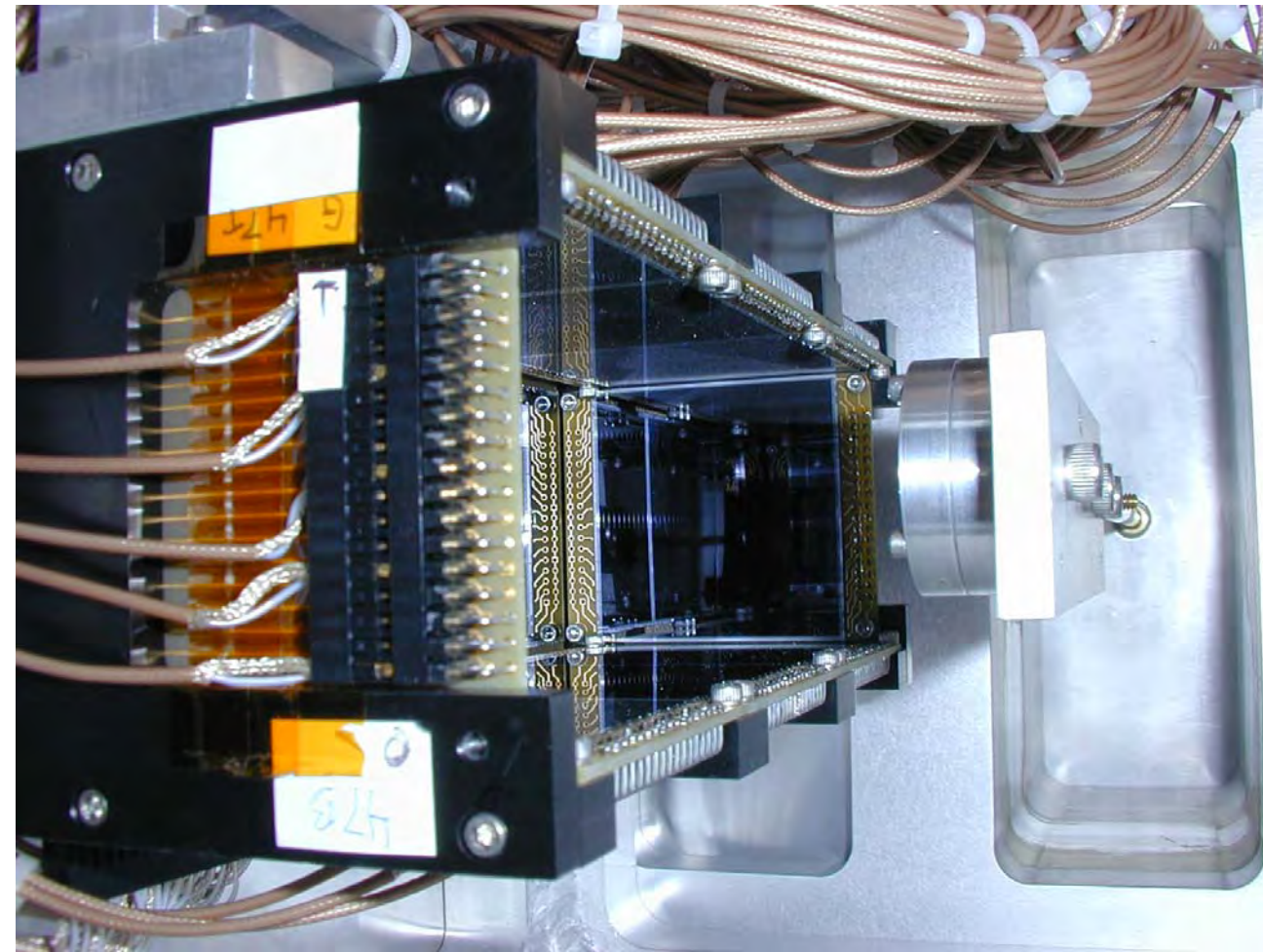
$$B\rho = 2.20 \pm 0.032$$

$$B\rho = 2.20 \pm 1.5\%$$

$$\text{BGS } \delta x' / \delta B\rho = 1.8 \text{ cm}/\%$$

16-cm wide MWAC and
18-cm wide Si-strip array:
detector covers **9%** in $B\rho$
($\pm 4.5\%$)

a **3σ** $B\rho$ error results in
half of the EVR distribution
missing the detector, and
would double the cross
section upper limits.



Why Don't We see 5-s 9.5-MeV $^{283}112$?

Is there a problem with the BGS targets?

Yes, but luminosity is determined with Rutherford-scattered beam.

Is there a systematic difference in beam energies from the accelerators at LBNL and Dubna?

No, $^{48}\text{Ca} + ^{208}\text{Pb}$ excitation functions match to within 2 MeV.

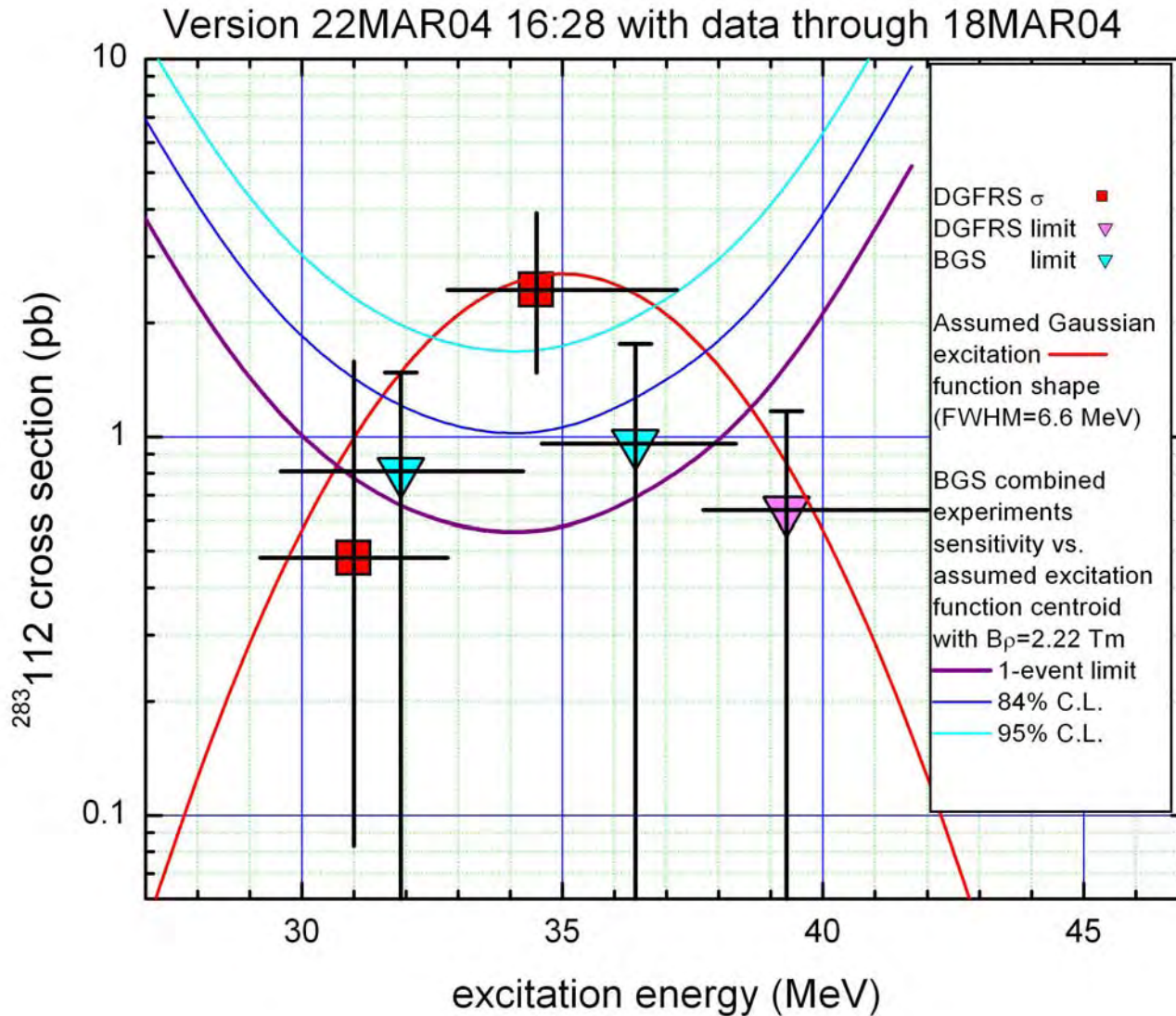
Do we have a different beam energy in our uranium targets due to errors in energy loss calculations?

No, pulse-height of Rutherford-scattered ^{48}Ca is within ~ 1 MeV.

Did we run the BGS at the wrong magnet settings?

No, this would require a 3σ deviation from systematics.

$^{283}\text{112}$ Cross Sections and Upper Limits



What's next?

Development of ^{244}Pu target capability

$^{244}\text{Pu}(^{48}\text{Ca},\text{xn})^{292}114$ in 2005

Radiochemistry with BGS-RTC using reactions such as

Rf:	$^{244}\text{Pu}(^{22}\text{Ne},5\text{n})^{261}\text{Rf}$	$t_{1/2} = 78 \text{ s}$	$^{208}\text{Pb}(^{50}\text{Ti},\text{n})^{257}\text{Rf}$	$t_{1/2} = 4 \text{ s}$
Db:	$^{244}\text{Pu}(^{23}\text{Na},5\text{n})^{262}\text{Db}$	$t_{1/2} = 34 \text{ s}$	$^{209}\text{Bi}(^{50}\text{Ti},\text{n})^{258}\text{Db}$	$t_{1/2} = 4 \text{ s}$
Sg:	$^{244}\text{Pu}(^{26}\text{Mg},5\text{n})^{265}\text{Sg}$	$t_{1/2} = 17 \text{ s}$		
Bh:	$^{244}\text{Pu}(^{27}\text{Al},4\text{n})^{267}\text{Bh}$	$t_{1/2} = 17 \text{ s}$		
Hs:	$^{244}\text{Pu}(^{30}\text{Si},5\text{n})^{269}\text{Hs}$	$t_{1/2} = 19 \text{ s}$		
112:	$^{244}\text{Pu}(^4\text{Ca},3\text{n})^{287}114 \rightarrow ^{283}112$	$t_{1/2} = 4 \text{ s}$		

...

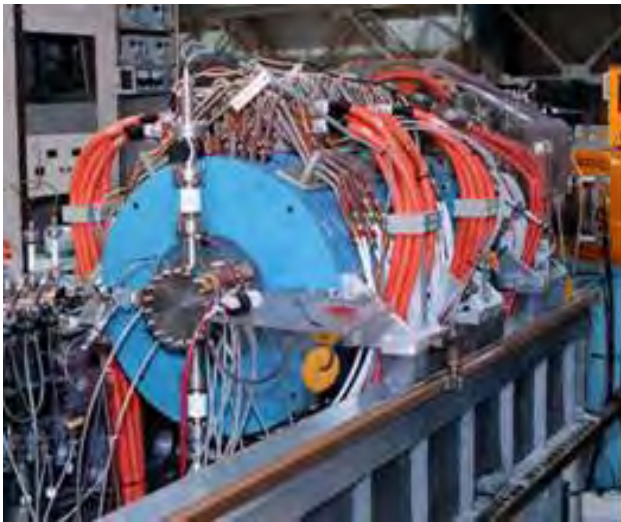
The LBNL 88-Inch Cyclotron



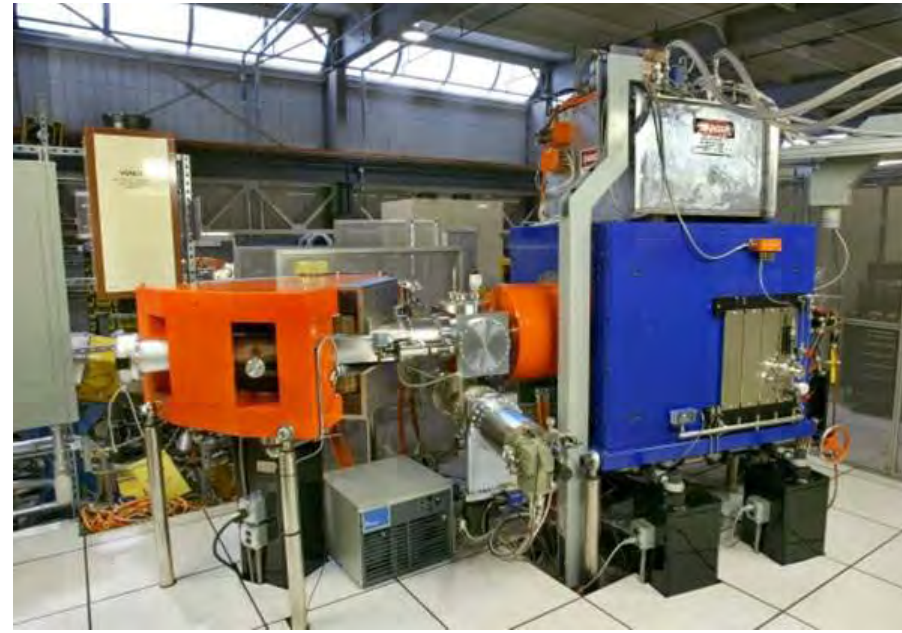
K130 Sector focused cyclotron
 $A/q \leq 5$ for Coulomb Barrier



First Operation in 1961

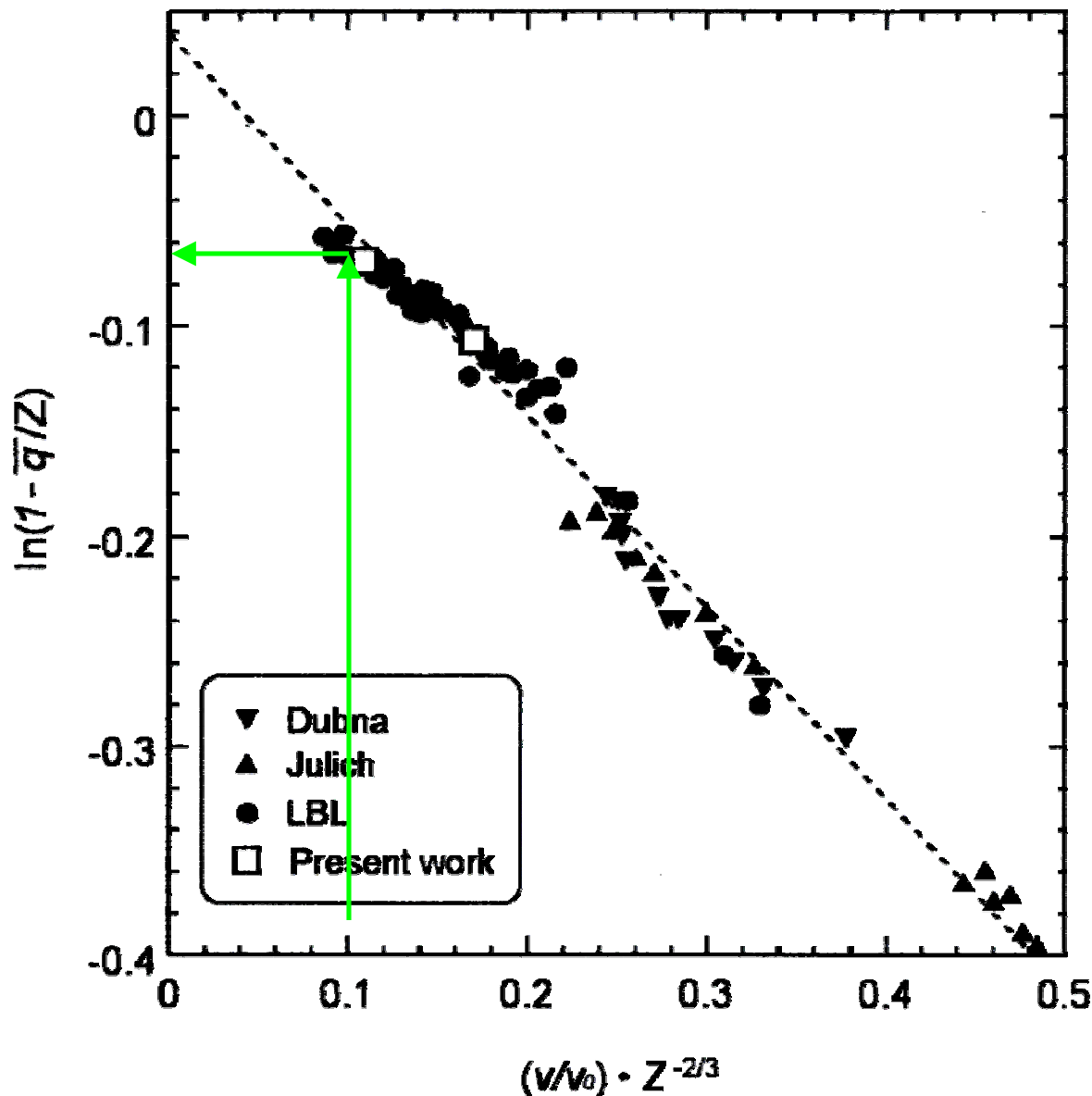


AECRU
(present)
and
VENUS
(Spring '05)



What is the $^{283}112$ magnetic rigidity?

According to RIKEN GARIS systematics . . .



$$v/v_0 = 11.33$$

$$(v/v_0)Z^{-2/3} = 0.1012$$

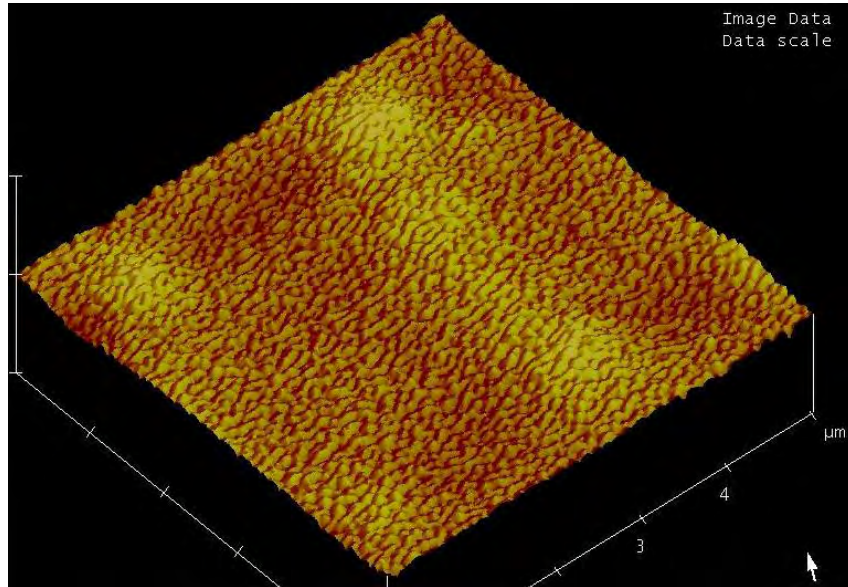
$$\ln(1 - q/Z) = -0.064$$

$$q = 6.94$$

$$B\rho = 2.17 \text{ Tm}$$

for small values of $\ln(1 - q/z)$
small errors in $\ln(1 - q/Z)$
lead to large errors in q

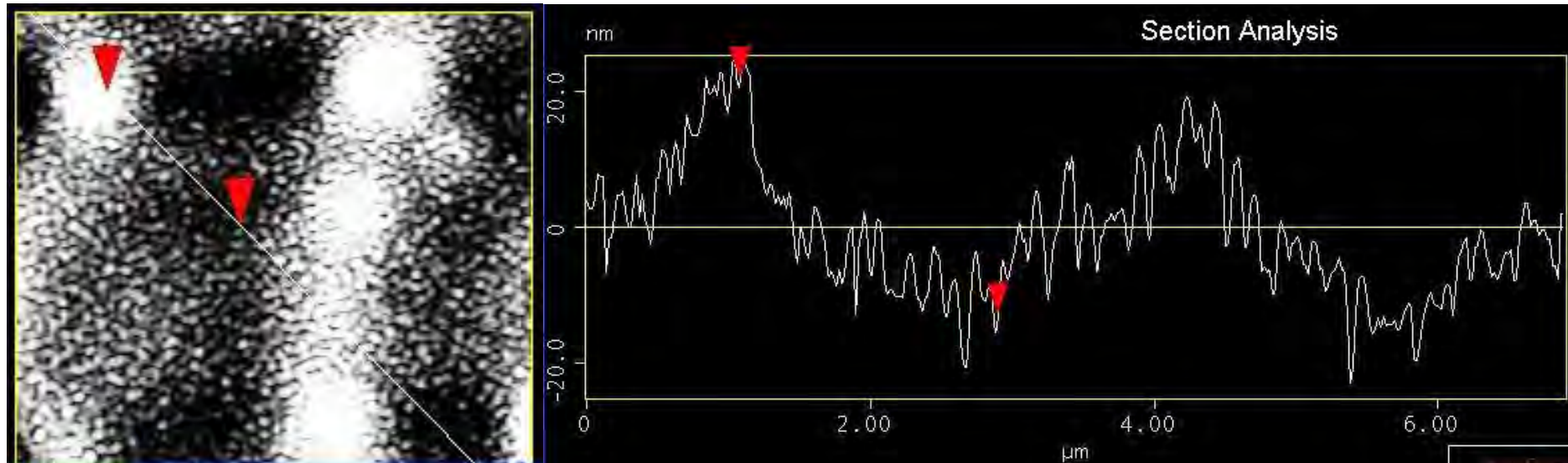
AFM of the edge of the UF₄ layer (outside the visible beam stripe)



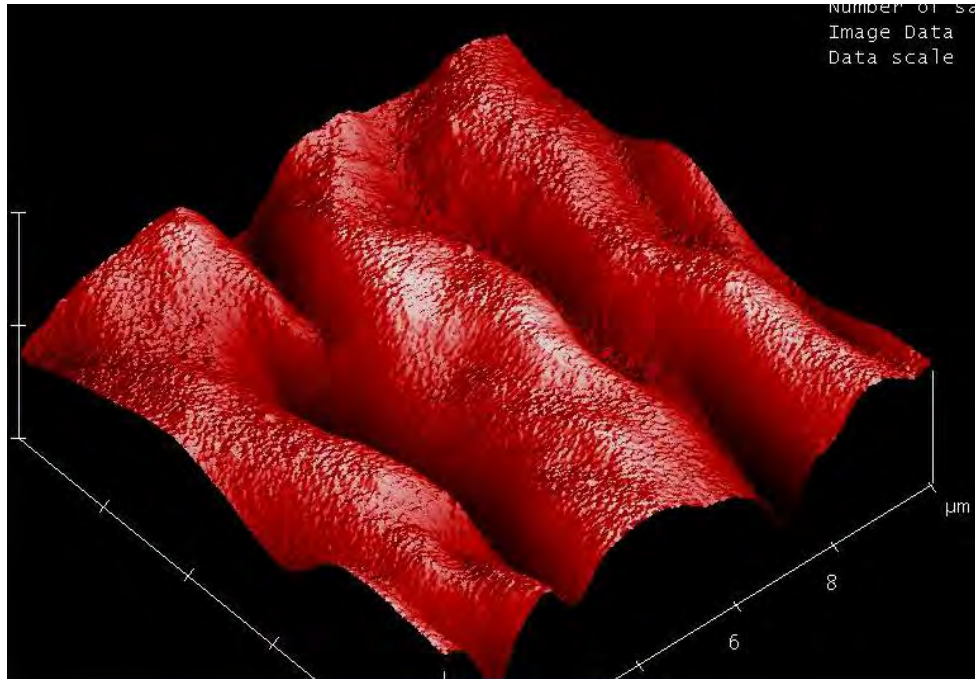
Overall UF₄ thickness is 900 nm

Crystalline structure

Thickness variations up to +/- 2%



AFM of the center of the UF4 layer (inside the visible beam stripe)

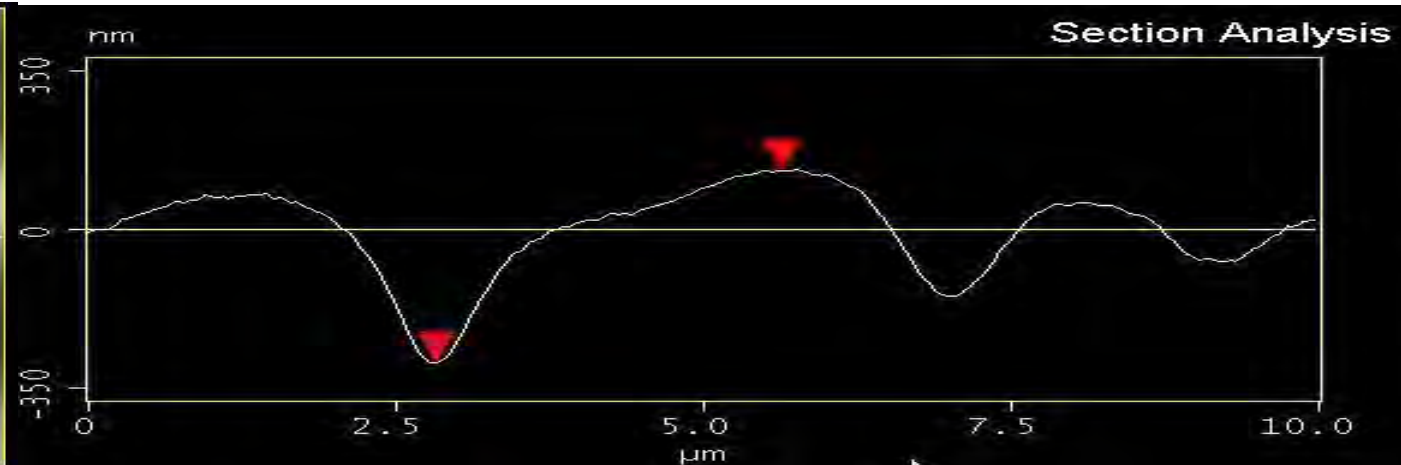
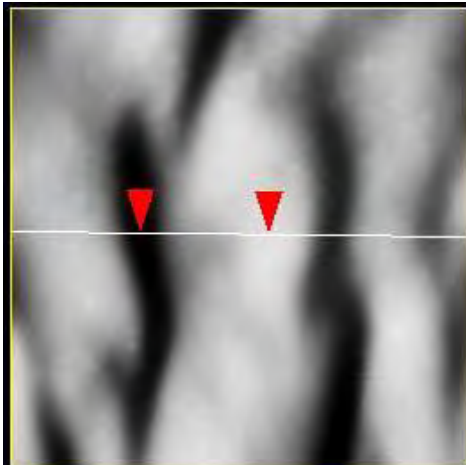


Overall UF4 thickness 900 nm

large-scale melting of UF₄

Variations up to +/- 20%

RMS thickness variations are
much less than 10%



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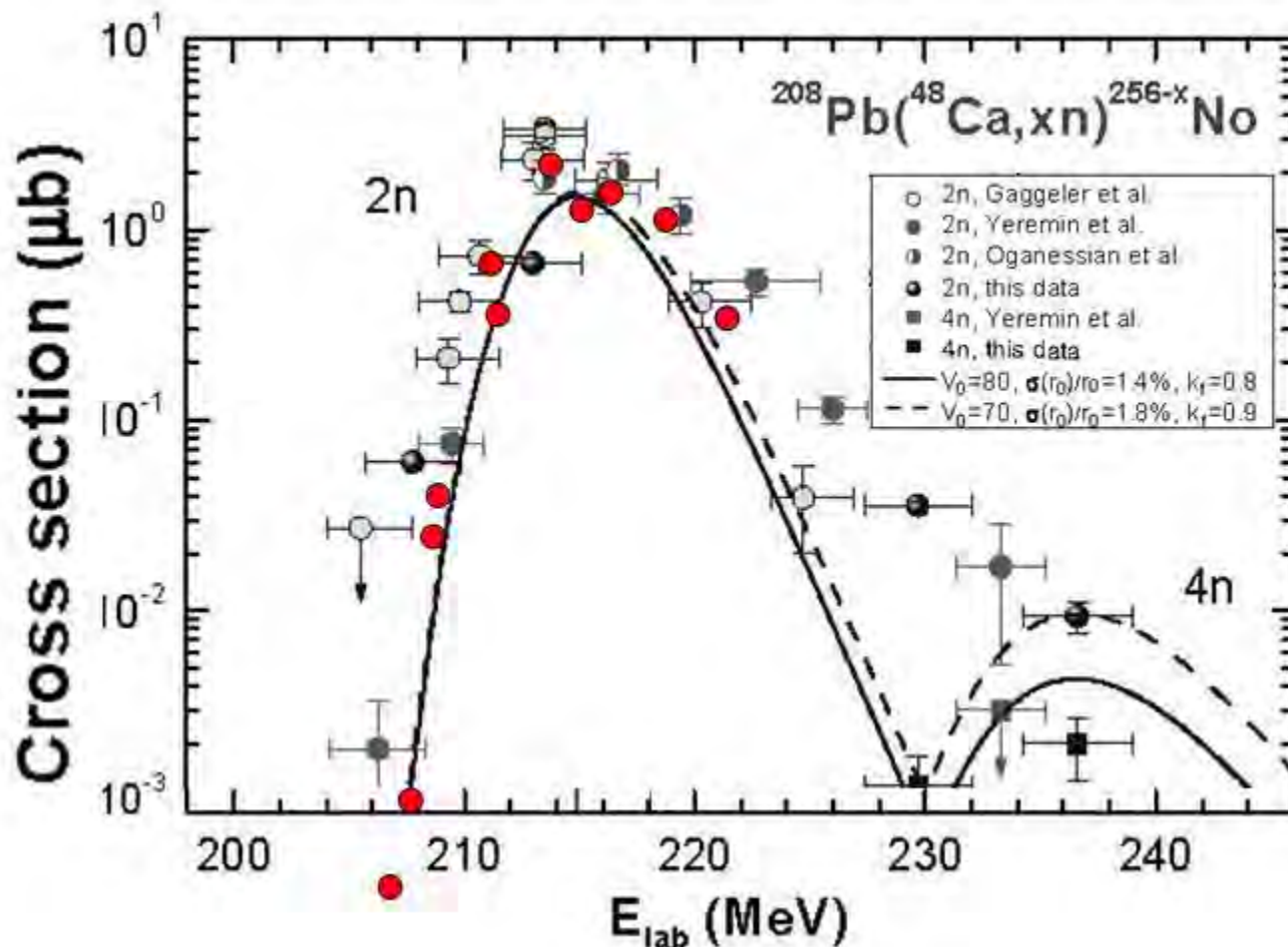
Did we run the BGS at the wrong magnet settings?

Is there a systematic difference in beam energies from the two accelerators?

Measurement of $^{208}\text{Pb}(^{48}\text{Ca}, 2n)^{254}\text{No}$ shows that . . .

Absolute beam energy is accurate to within 1% (from comparison of excitation functions at right)

Beam energy reproducibility is accurate to within 0.5% FWHM (from a plot of pulse-height in a PIN diode vs. the square of the cyclotron frequency)



Are there errors in the energy loss calculation?

Measurement of pulse-heights for Rutherford-scattered beam particles, and a comparison with those for the Ca + Pb reactions show that the energies in our $^{48}\text{Ca} + ^{238}\text{U}$ experiments were accurate to within 0.5% (1.2 MeV)

Conclusion: There is no doubt that our **TWO BEAM ENERGIES** cover the peak of the Dubna excitation function.

Run052	RE Ca + ^{206}Pb	RW Ca + ^{206}Pb	RE Ca + ^{238}U	RW Ca + ^{238}U
48Ca E at end of tgt	211.7	211.7	227.6	227.6
^{48}Ca E after scatter	202.7	202.4	218.0	217.6
E-P.H.D.	197.7	197.4	212.8	212.4
peak channel	1743.7	2348.5	1884.5	2527.5
E by Ruth. Ratio			213.7	212.4

Run070	RE Ca + ^{207}Pb	RW Ca + ^{207}Pb	RE Ca + ^{238}U	RW Ca + ^{238}U
48Ca E at end of tgt	211.8	211.8	233.5	233.5
^{48}Ca E after scatter	202.8	202.5	223.6	223.3
E-P.H.D.	197.8	197.5	218.3	218.0
peak channel	1556.7	1585.8	1724.2	1763.5
E by Ruth. Ratio			219.1	219.6