

Lifetime measurements of excited states in neutron-rich C and O isotopes

M. CIEMAŁA, IFJ PAN KRAKOW

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MILANO – IFJ PAN KRAKOW

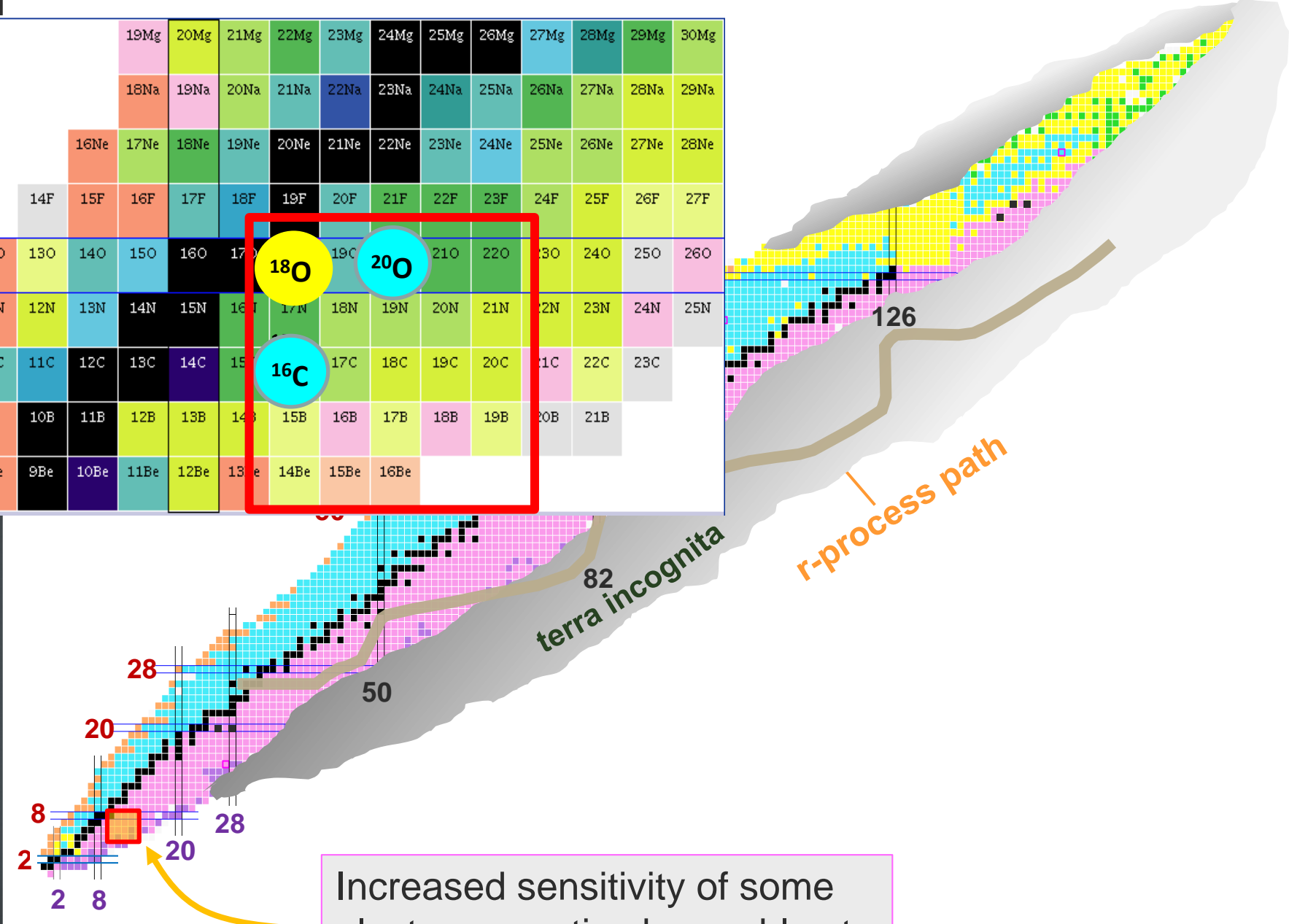
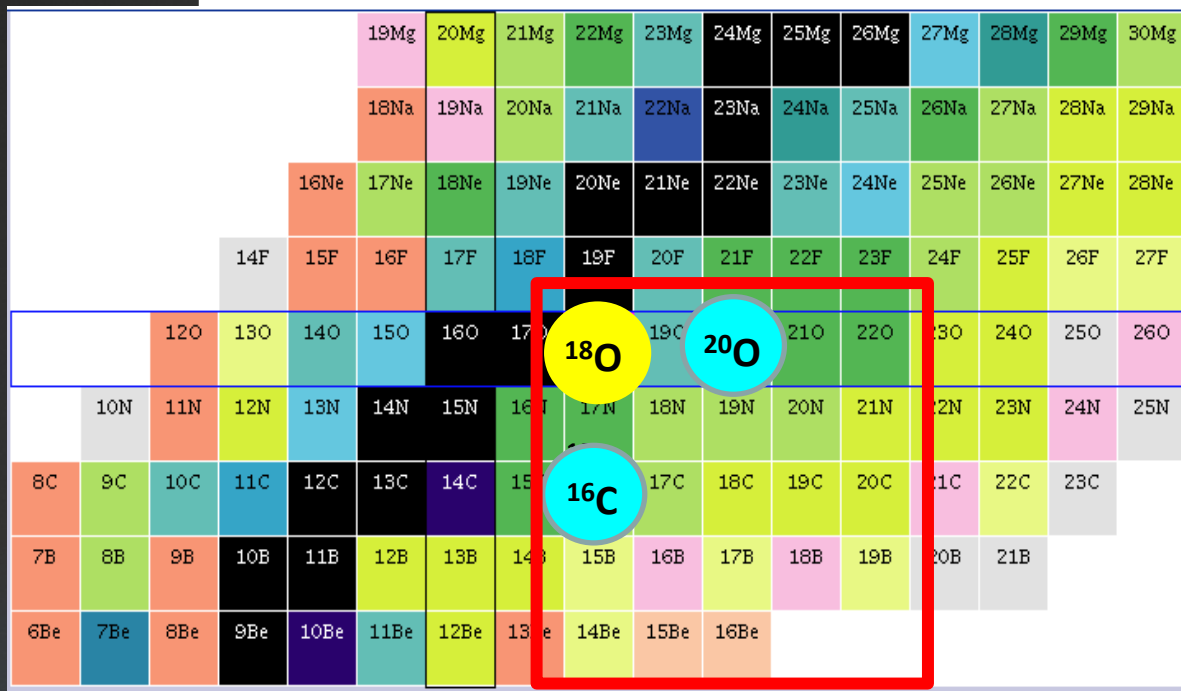
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VAMOS COLLABORATION, COORDINATOR: A. LEMASSON, GANIL, CAEN

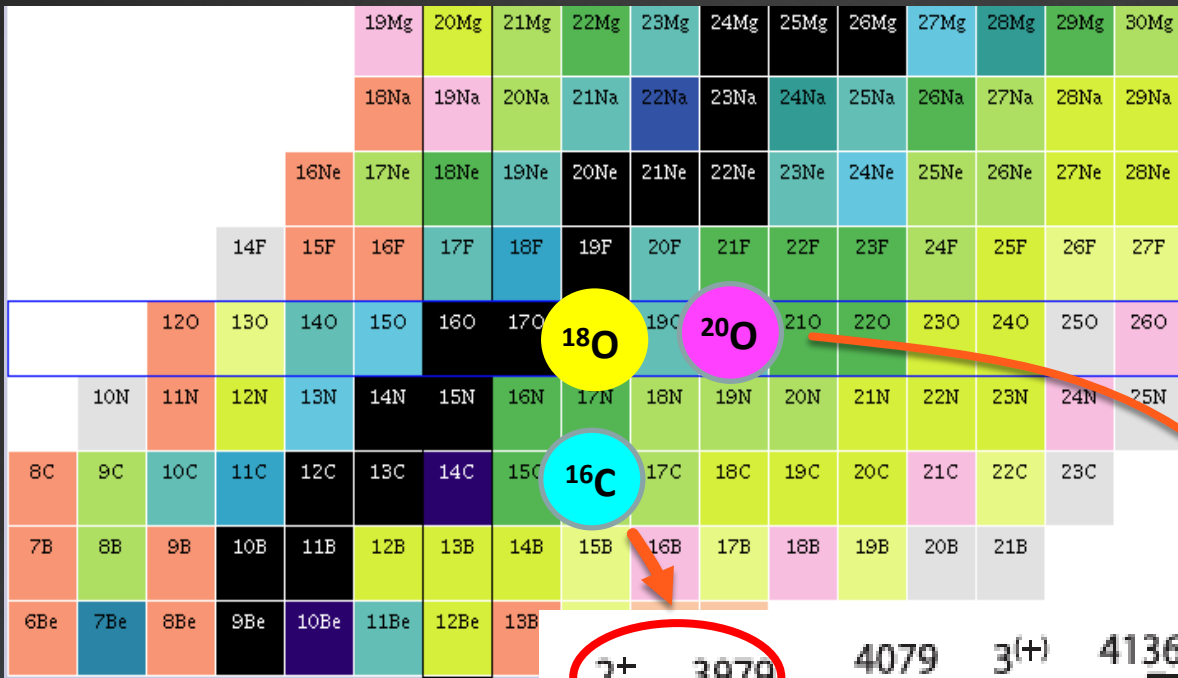
PARIS COLLABORATION, COORDINATOR A. MAJ, IFJ PAN, KRAKOW





Increased sensitivity of some electromagnetic observables to the details of n-n force

Z ↑

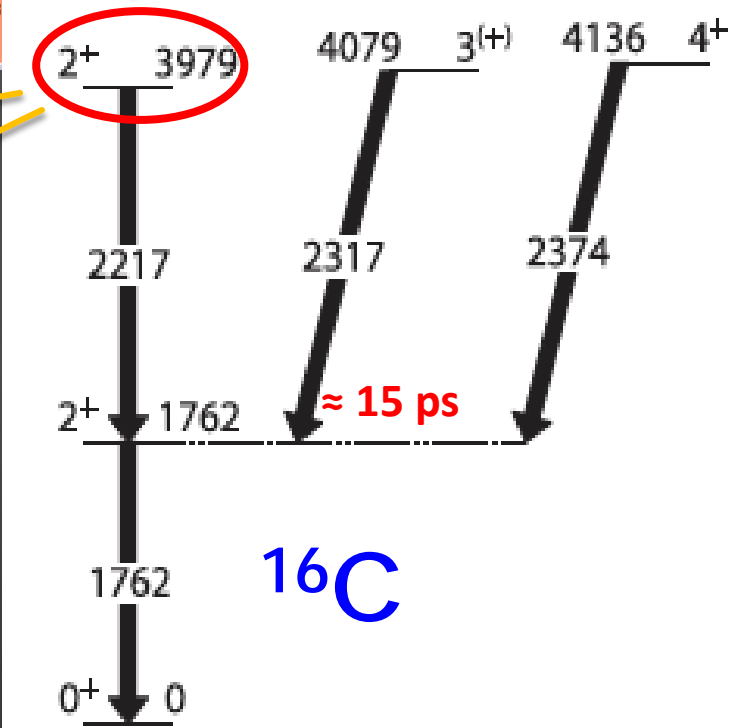


ab initio
Many-Body-Pert. Theory calculations
of the 2^+_2 lifetimes

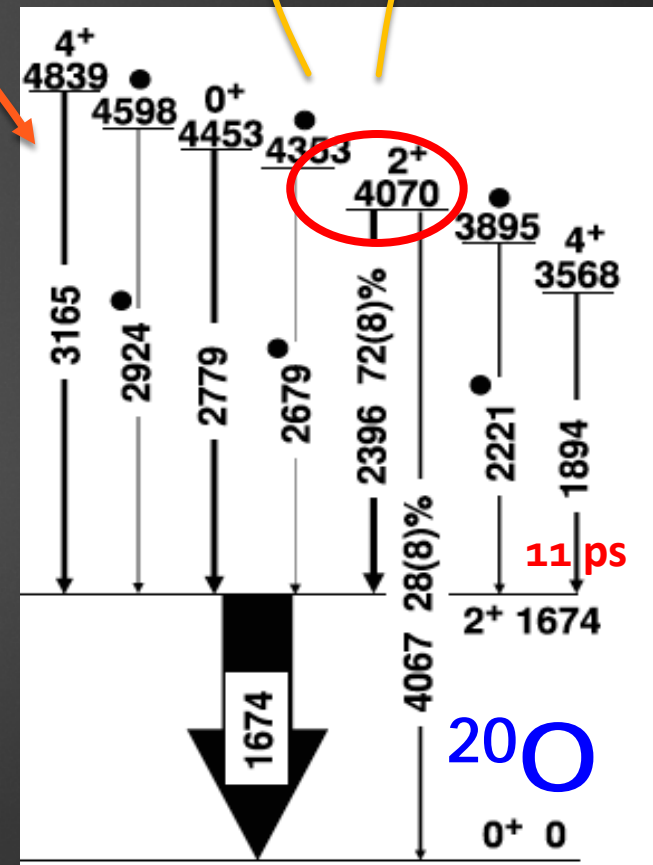
NN 0.32 ps
NNN 0.2 ps

NN 0.3 ps
NNN 0.08 ps

ab initio
No-Core-SHELL Model
calculations

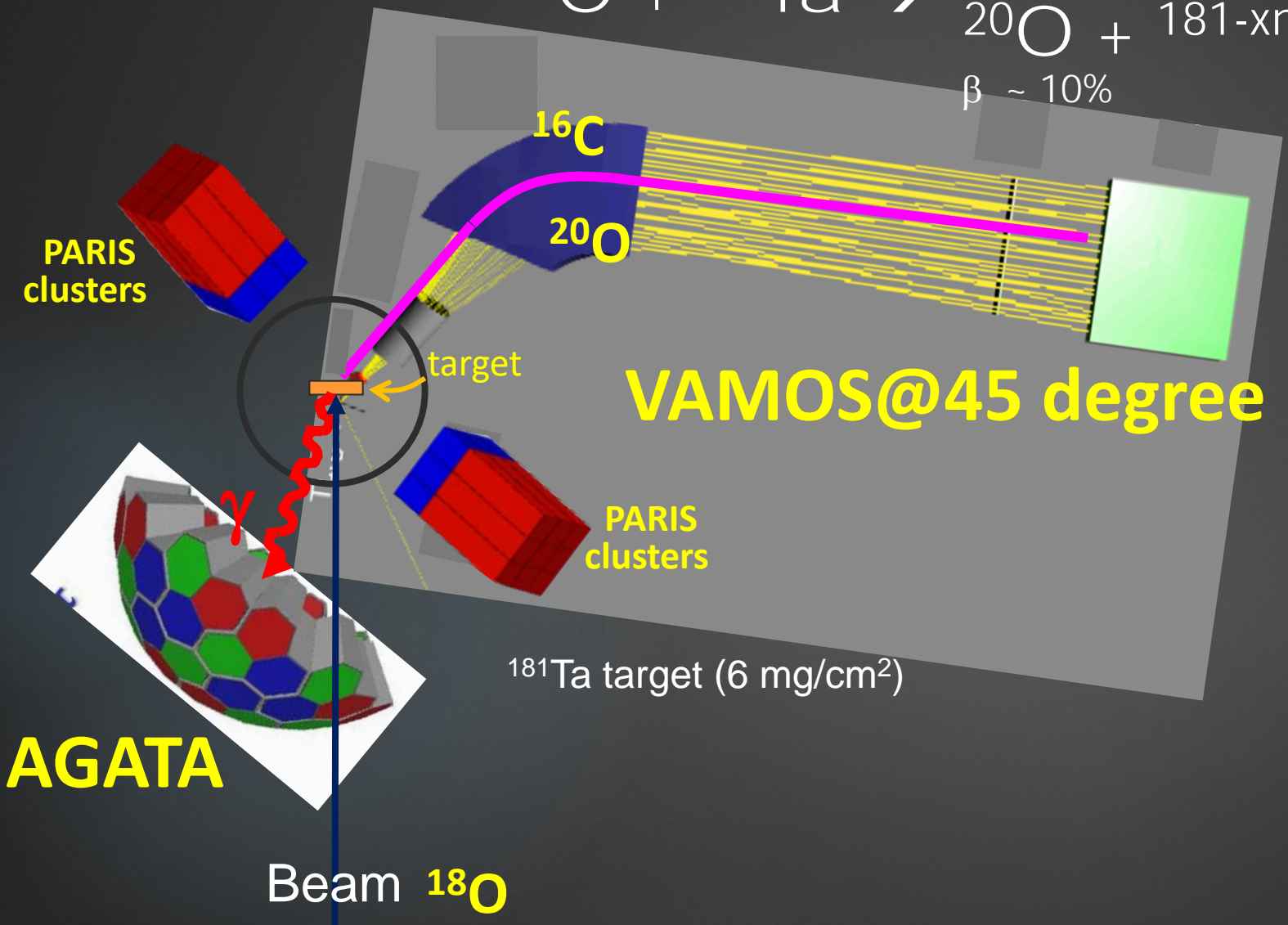
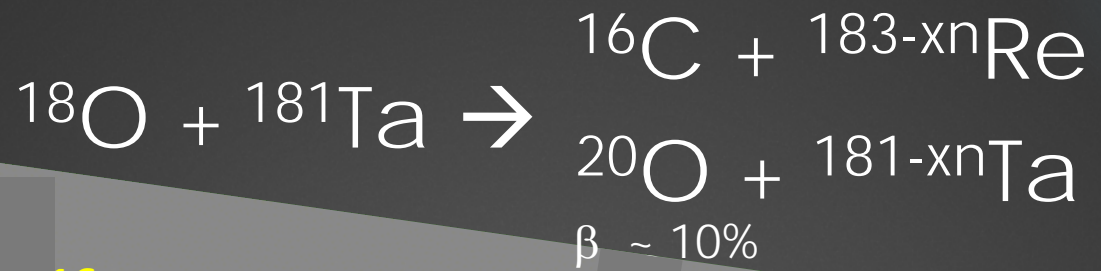


M. Petri et al., Phys. Rev. C 86, 044329 (2012).

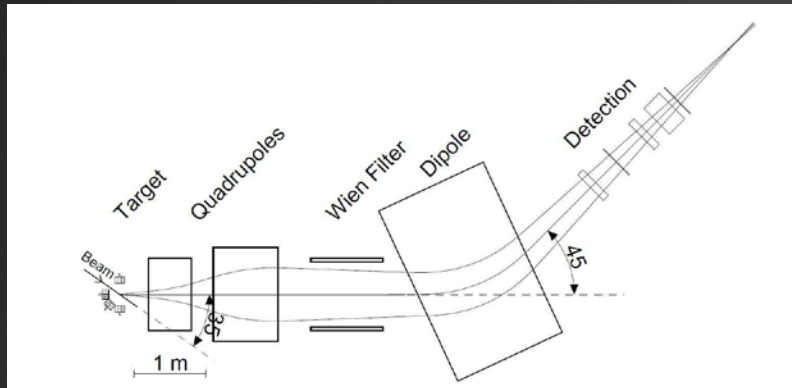


M. Wiedeking et al., Phys. Rev. Lett. 94, 132501 (2005).

Experimental setup - E676@GANIL



Experimental setup –VAMOS++ spectrometer



Scheme of VAMOS++

VAMOS++ at 45 degree

VAMOS entrance detector:

2 DC (for ions angle)

VAMOS focal plane:

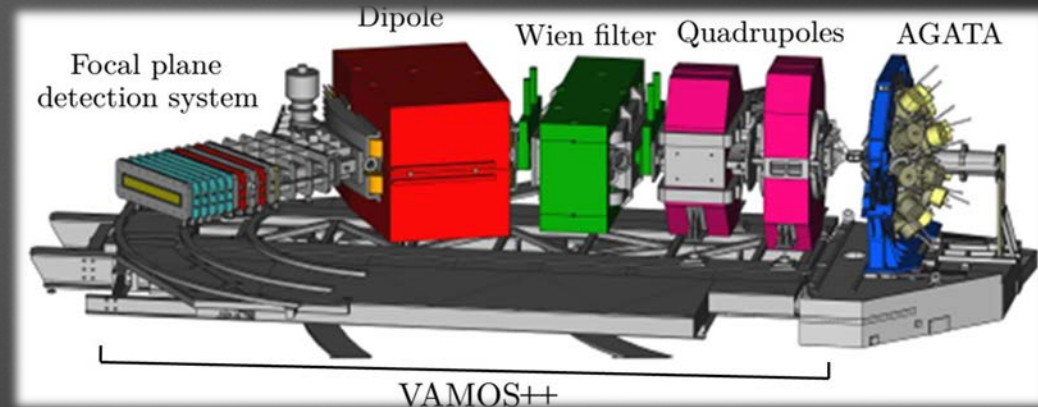
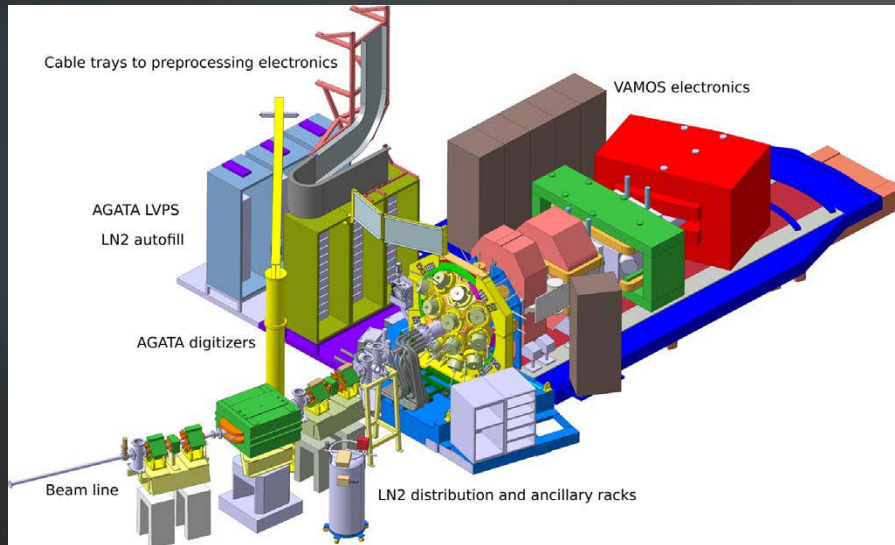
DC (for Brho reconstruction),

6 rows of IC (for ΔE)

Plastic (for trigger and ToF)

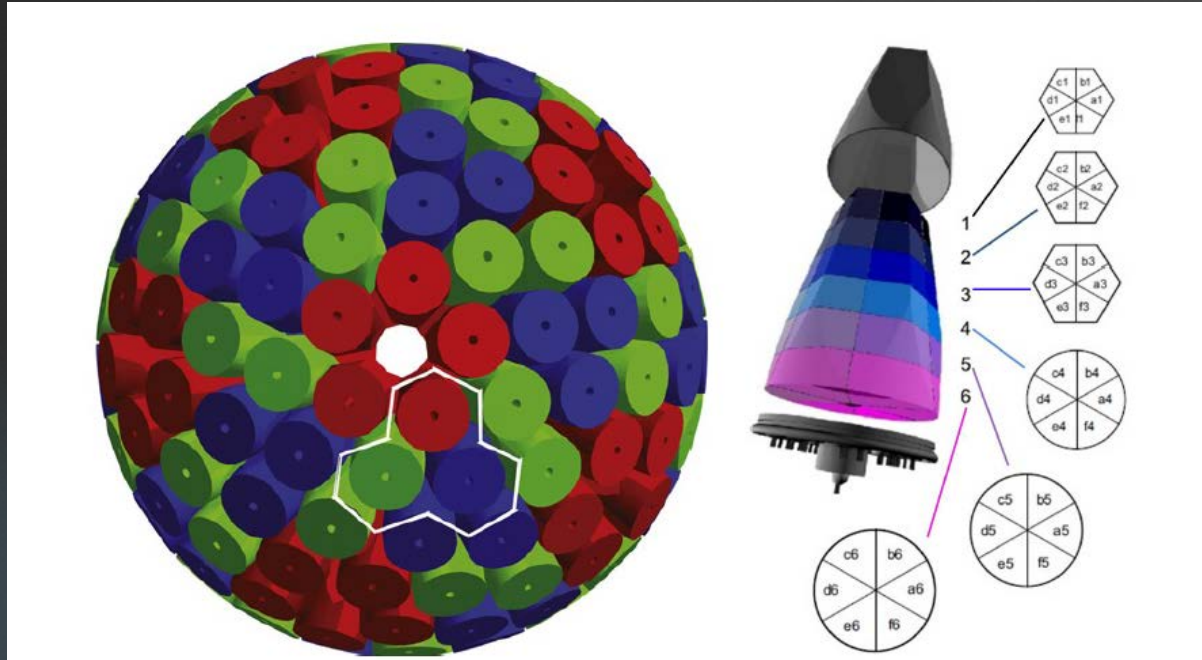
Angular acceptance:

+/- 6 degree in theta, +/- 15 degree in phi

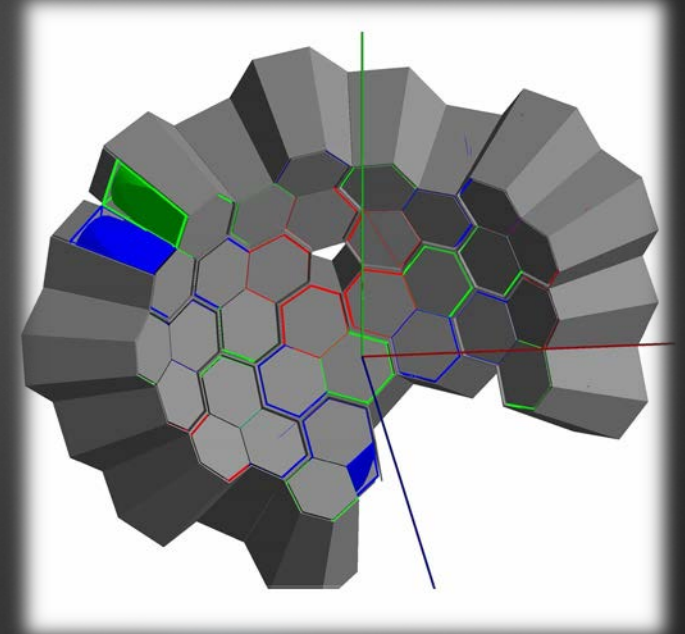


E. Clement et al. NIMA 885, 1-12 (2017)

Experimental setup -AGATA



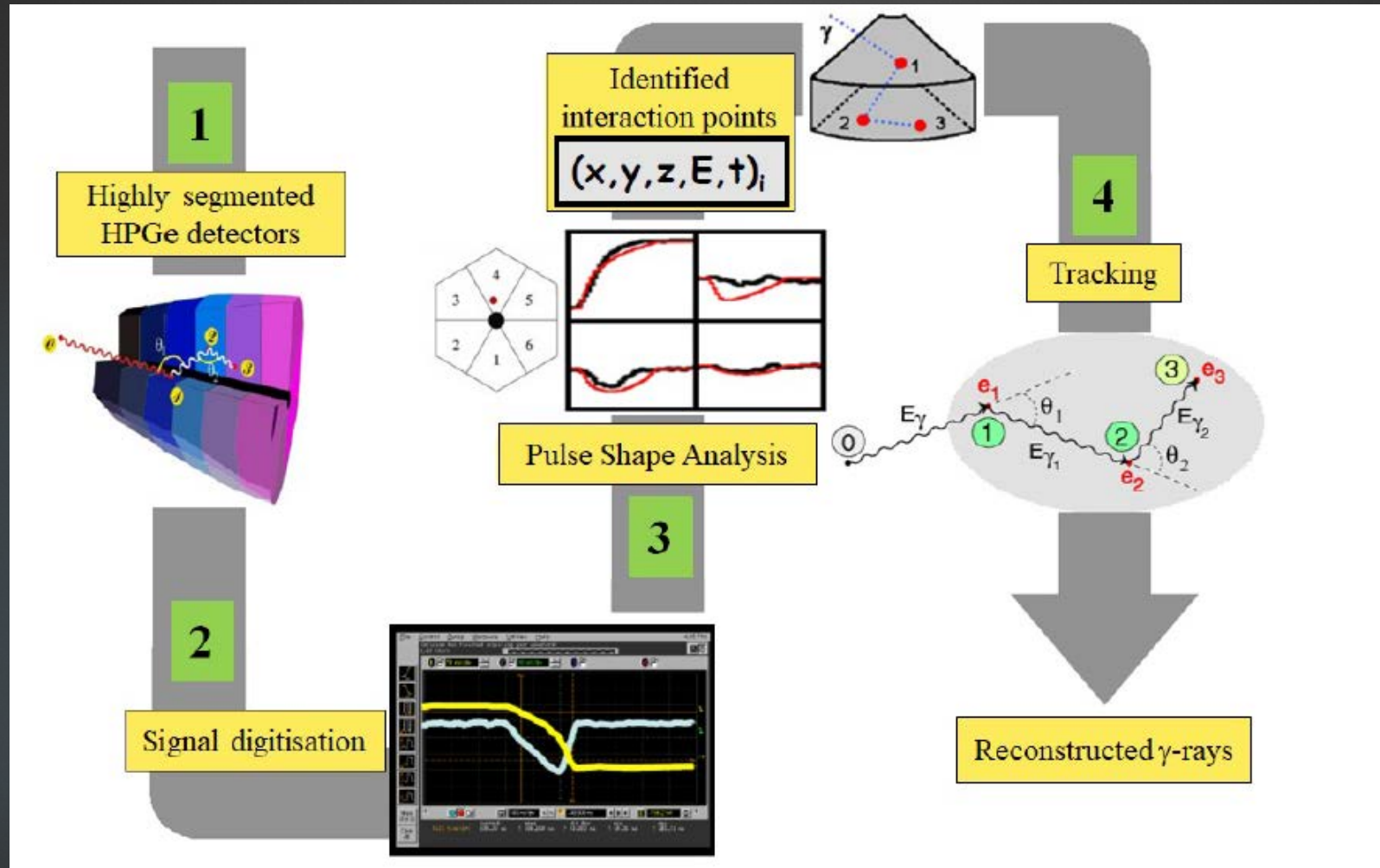
- **AGATA**
31 crystals (backward angles)
Working during our experiment
at GANIL



AGATA - *Advanced Gamma Tracking Array*.
36 segments per crystal (6 longitudinal rings and 6 transversal sectors)

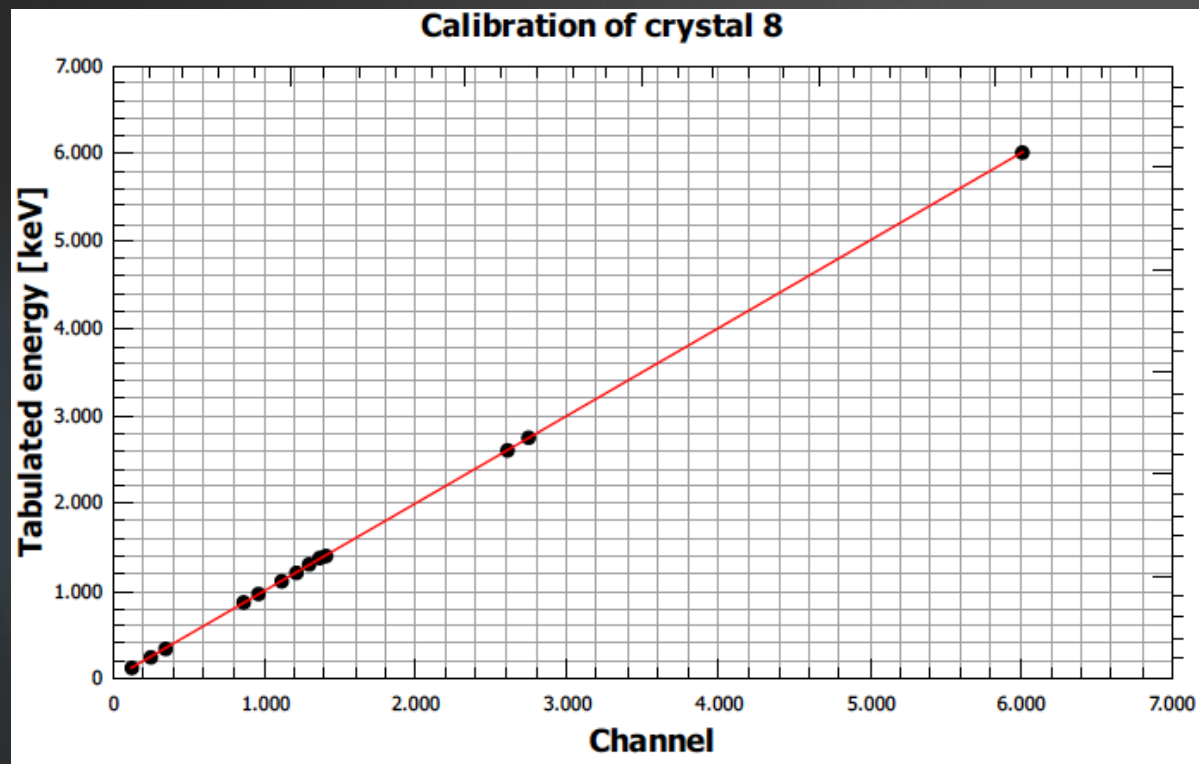
Tracking algorithms allows to reconstruct the path of the interacting γ -ray in the detection material, its energy and incoming direction.

Experimental setup –AGATA



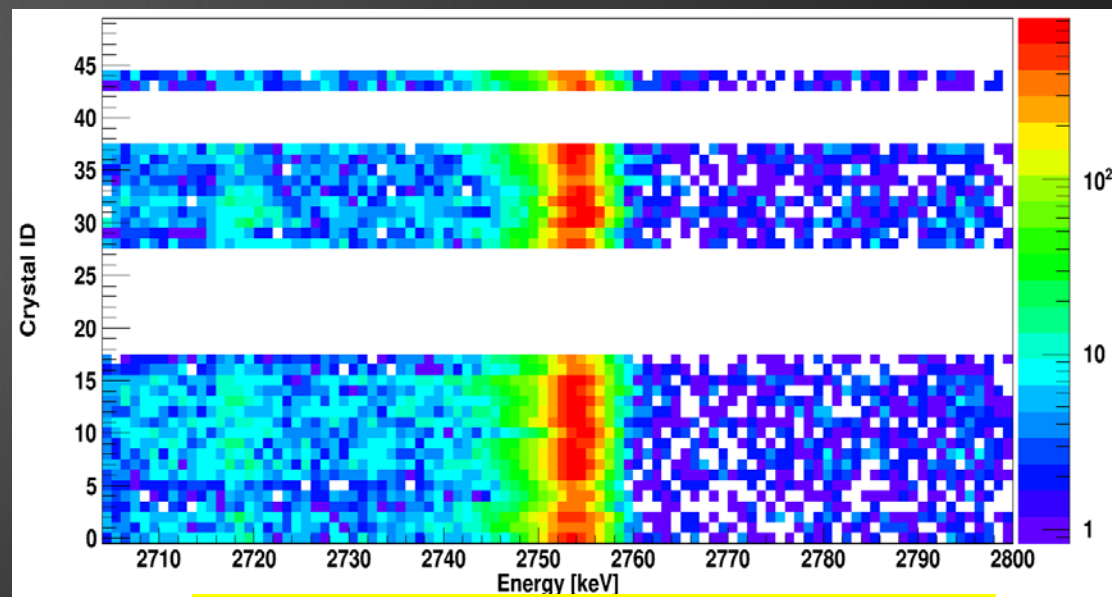
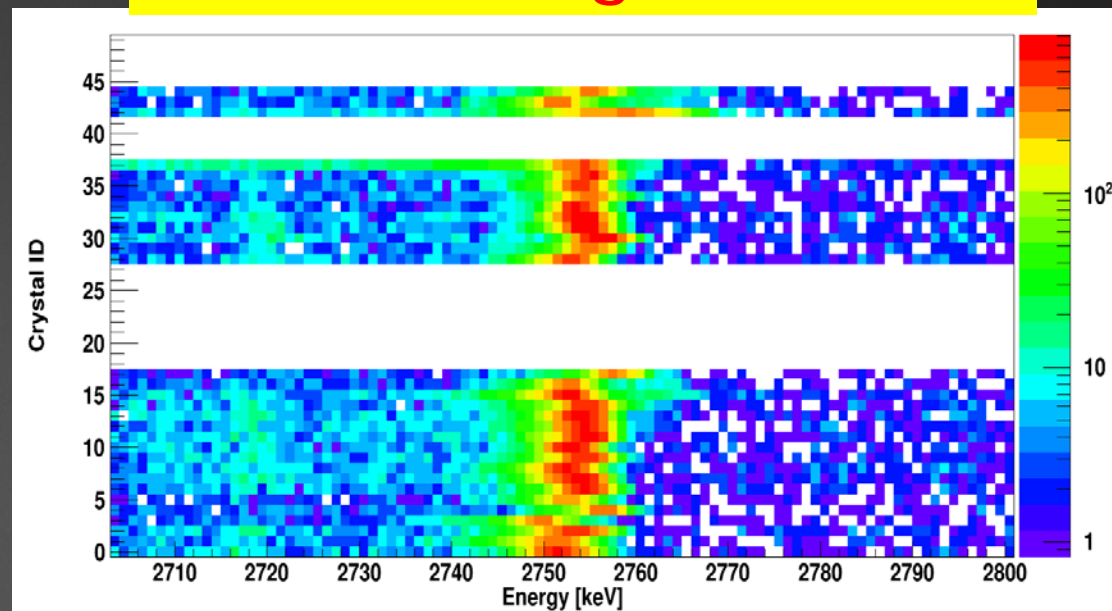
AGATA

Example of typical linear calibration curve



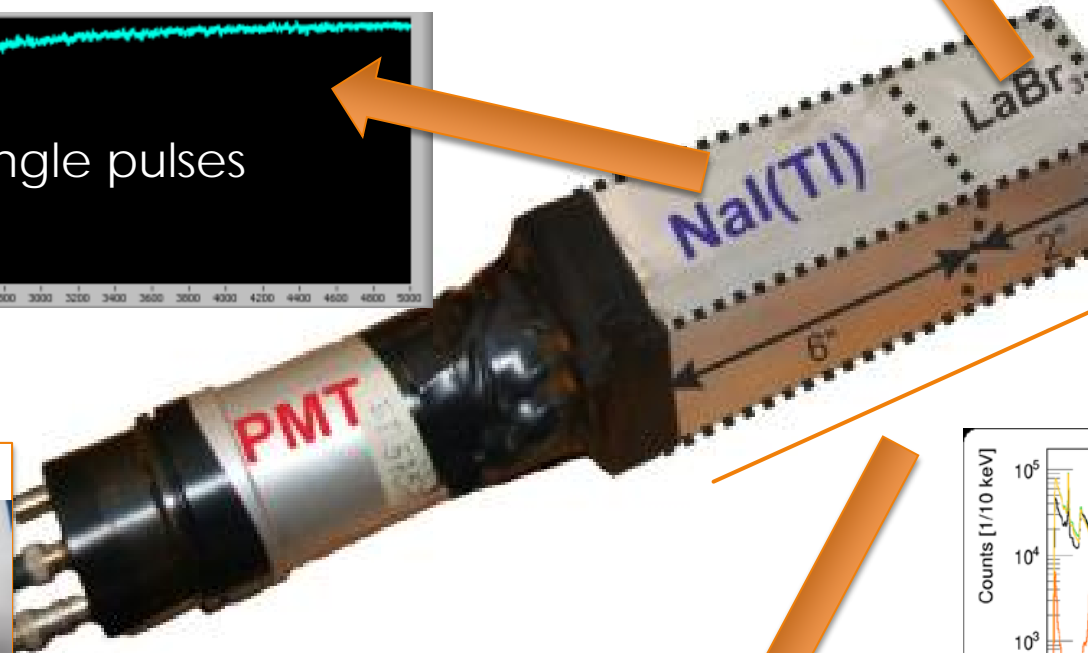
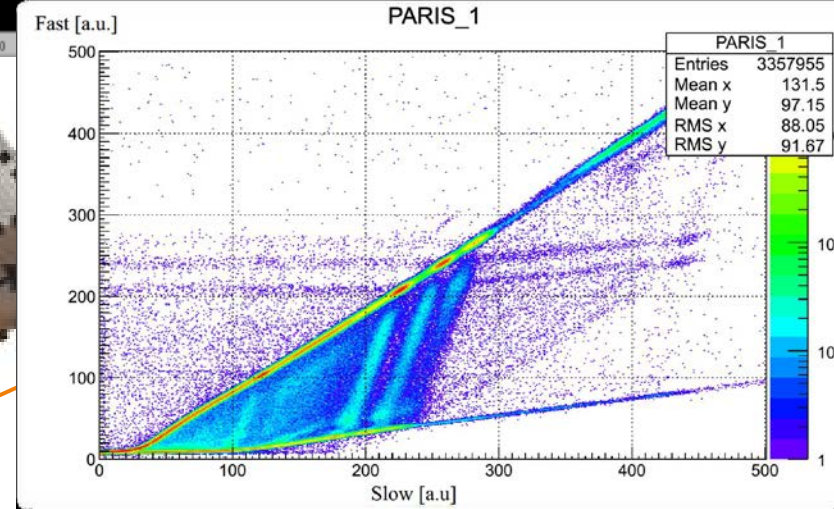
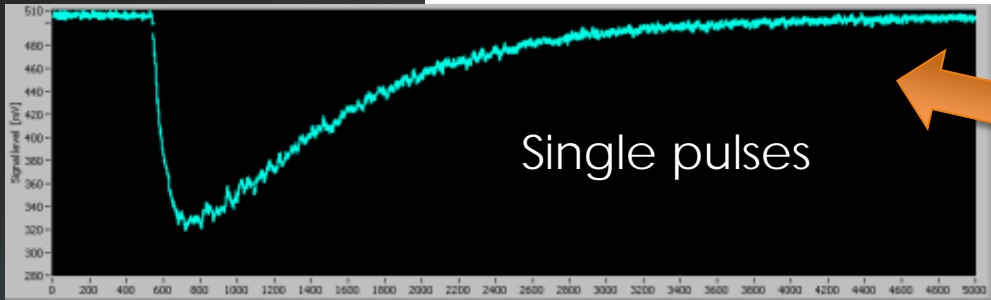
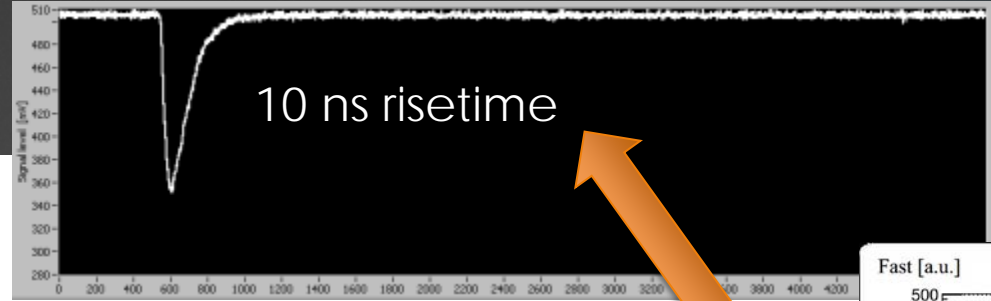
AGATA (re)calibration – performed by Milano group – S. Ziliani

Nearline sorting coefficients

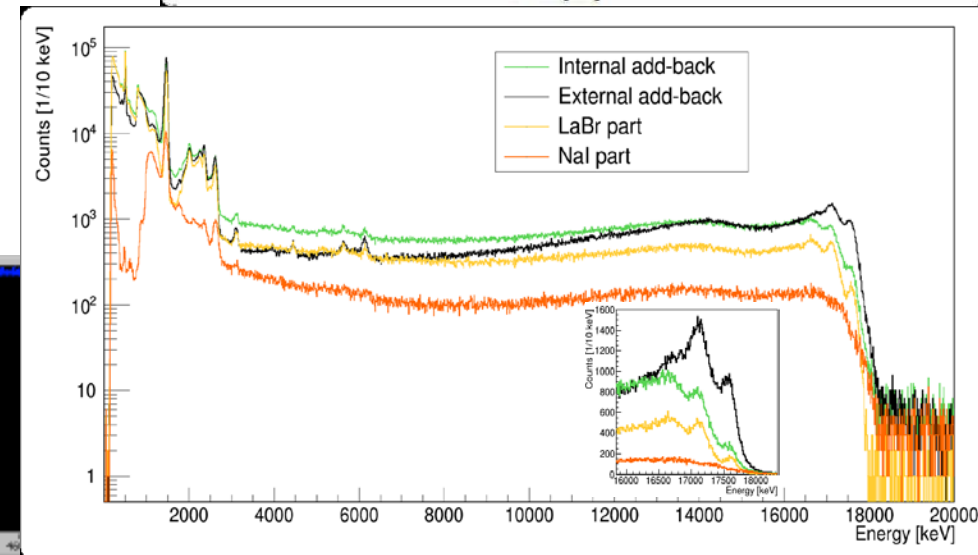
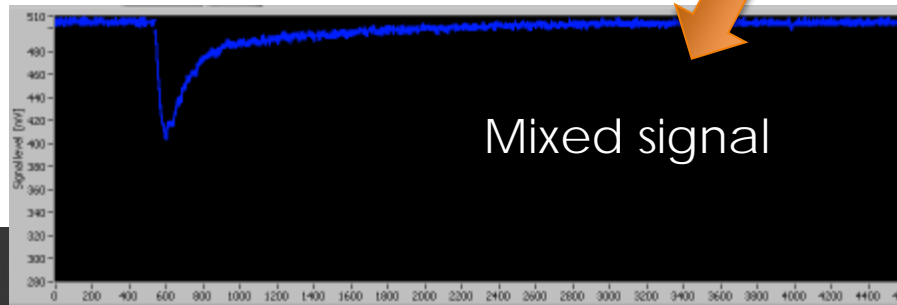


NEW sorting coefficients

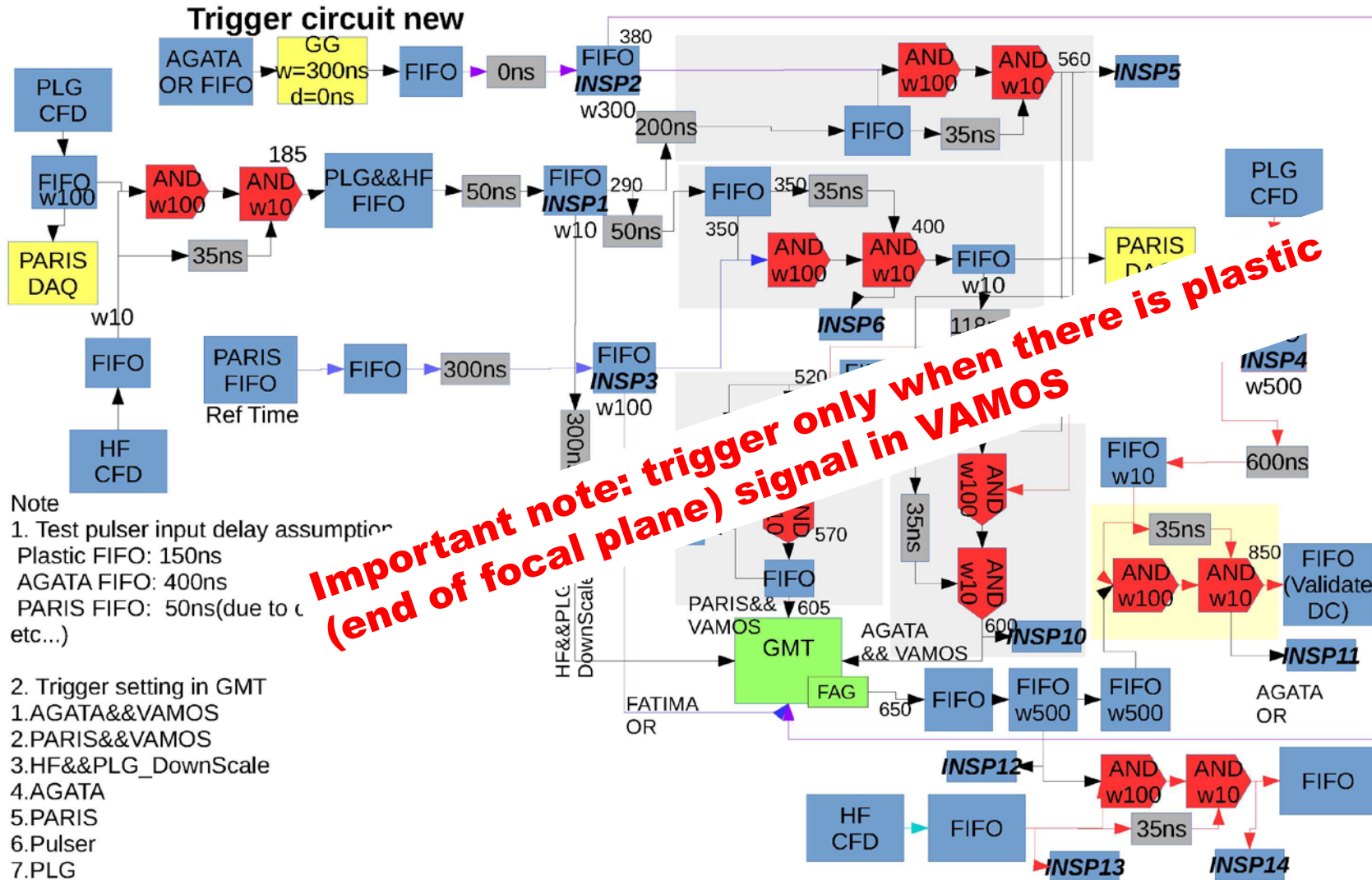
Experimental setup - PARIS



HAMAMATSU
Photomultiplier Tube 光電子増倍管
TYPE R7723-100
NO. ZK6699
浜松ホトニクス株式会社



Experimental setup – trigger conditions



VAMOS spectrometer, ion identification and velocity measurement

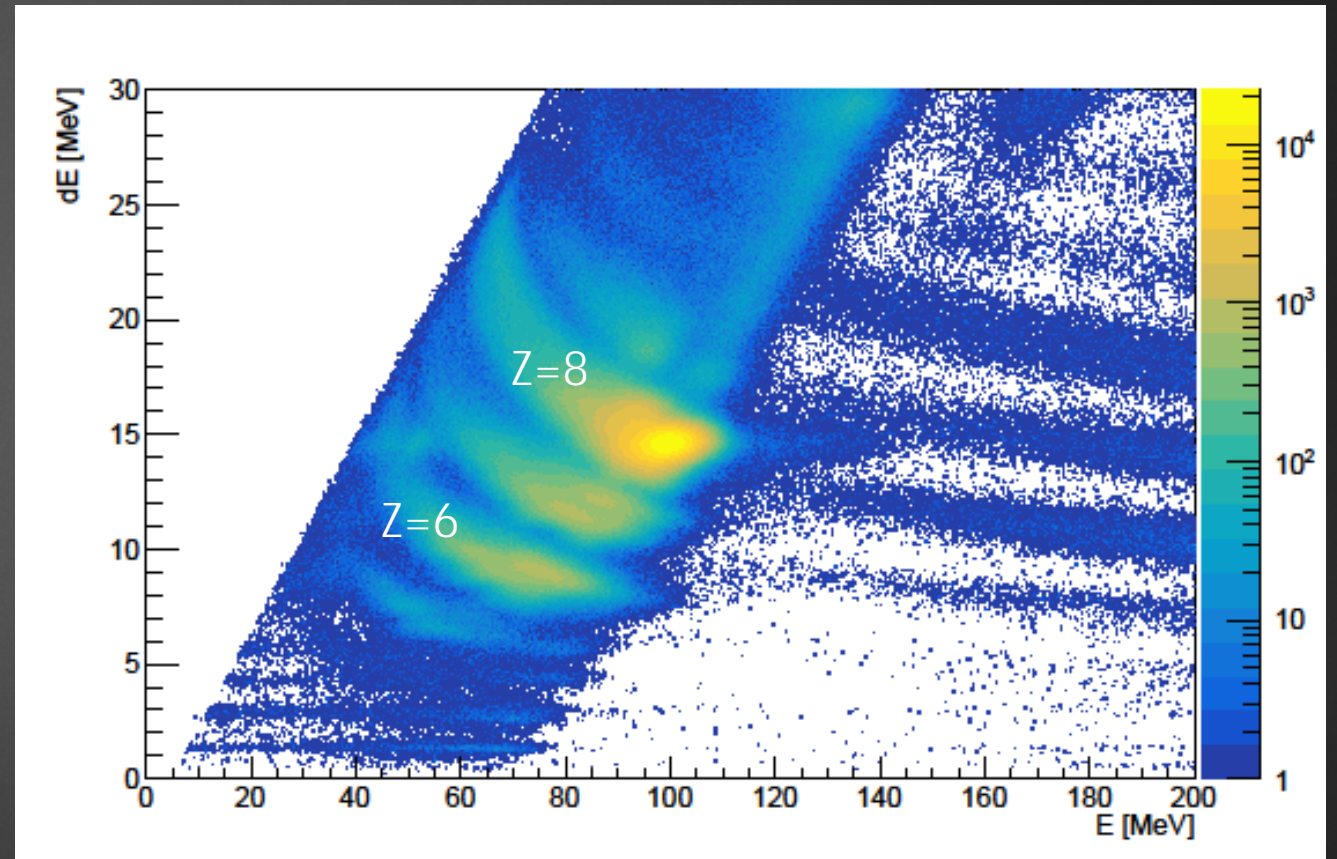
$$V = D/\text{TOF}$$

TOF time of flight calculated as a difference between the RF and the plastic detector time signals

D is ion path length

Z determination:

dE (ionization chambers) versus E (IC + plastic E deposit)



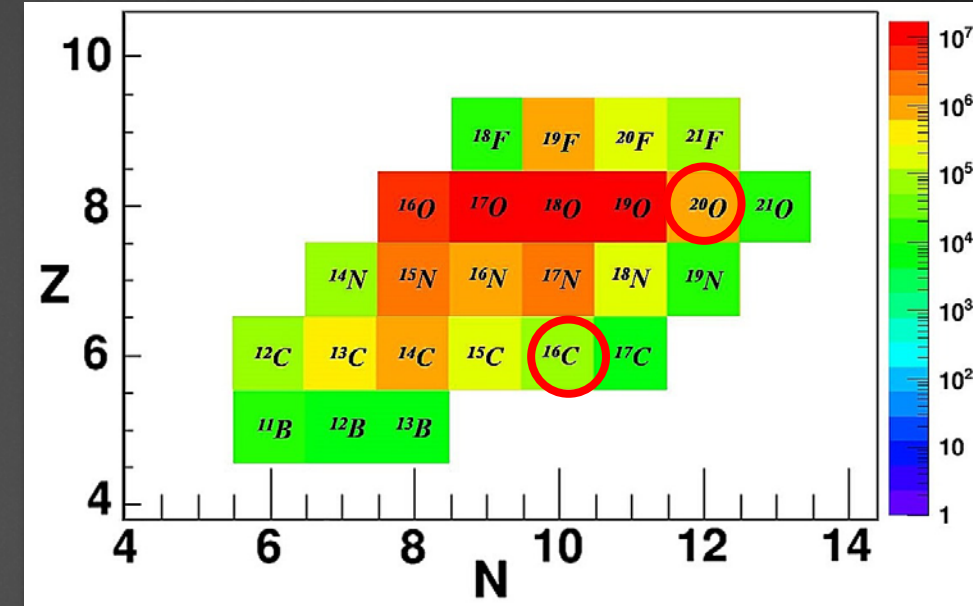
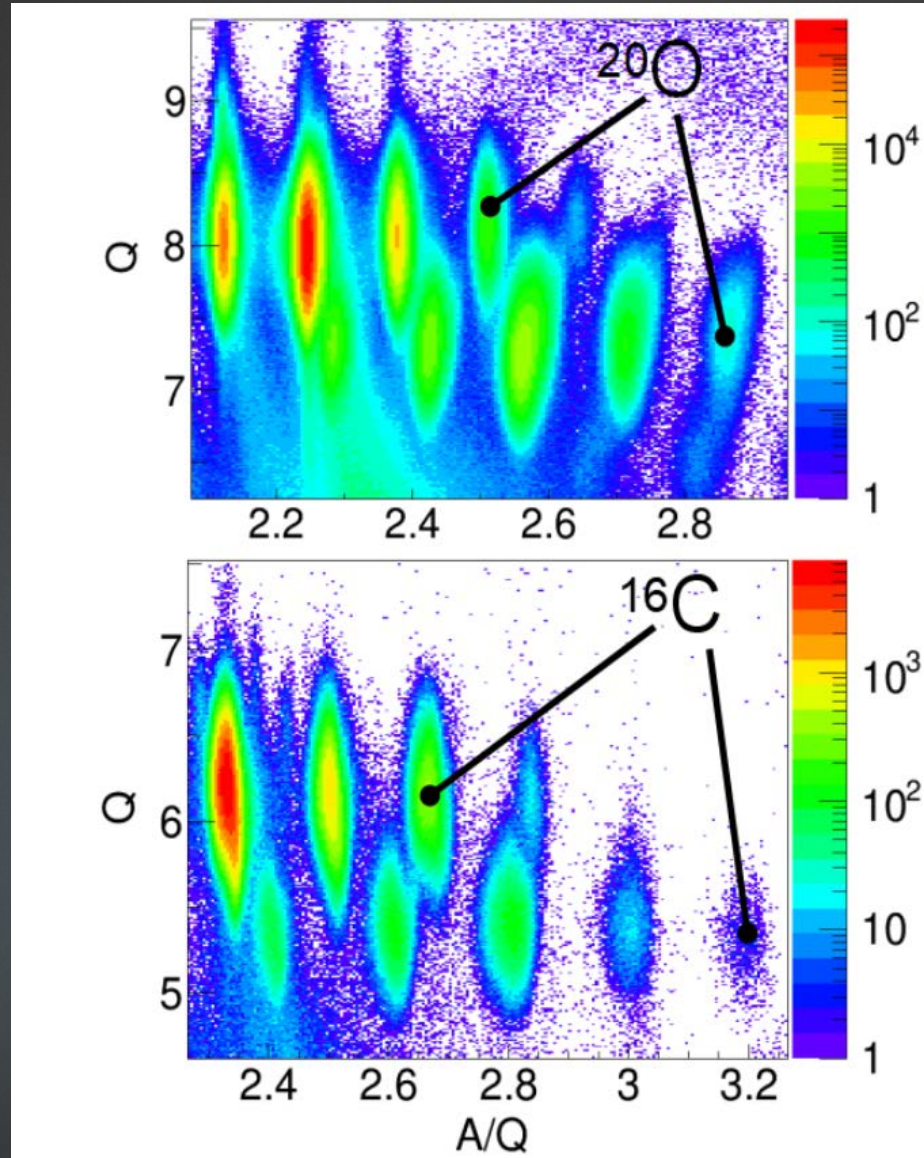
VAMOS spectrometer, ion selection

$$M/Q = B\rho/(3.105\gamma\beta)$$

$$M_0 = E/(931.5(\gamma - 1))$$

$$Q = M_0/(M/Q)$$

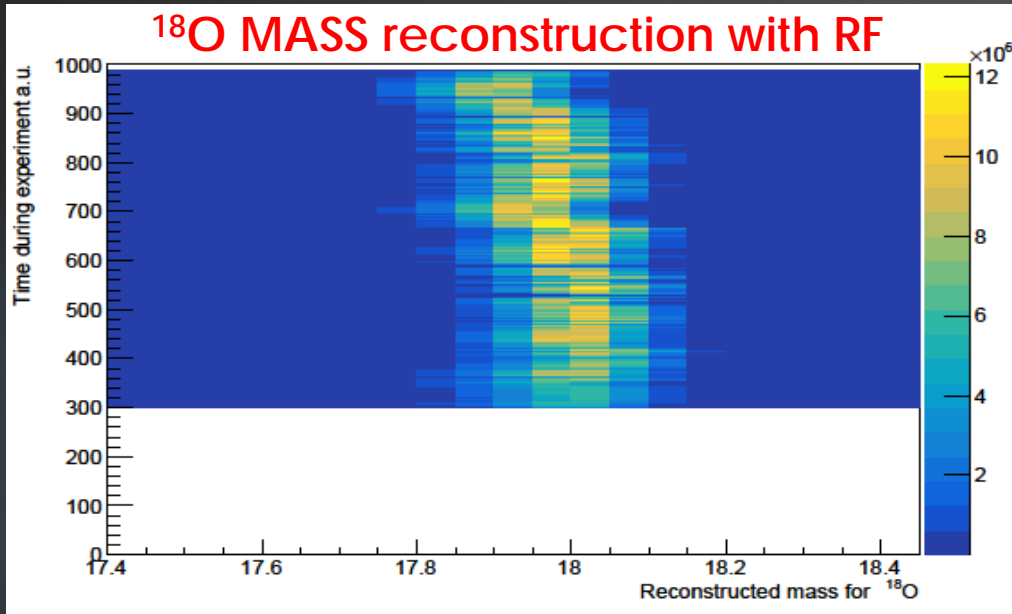
$$M_r = (M/Q)Q_{\text{int}}$$



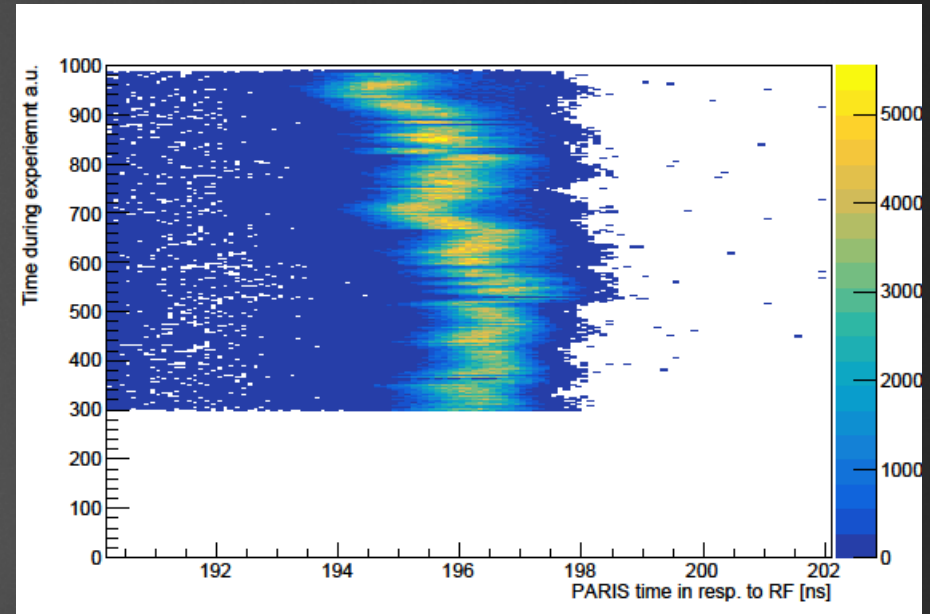
Identified isotopes from B to F.

PARIS timing – correction to velocity

We measure V by path in spectrometer and time between **RF** and **Plastic** at the end of focal plane.



Time of experiment ↑

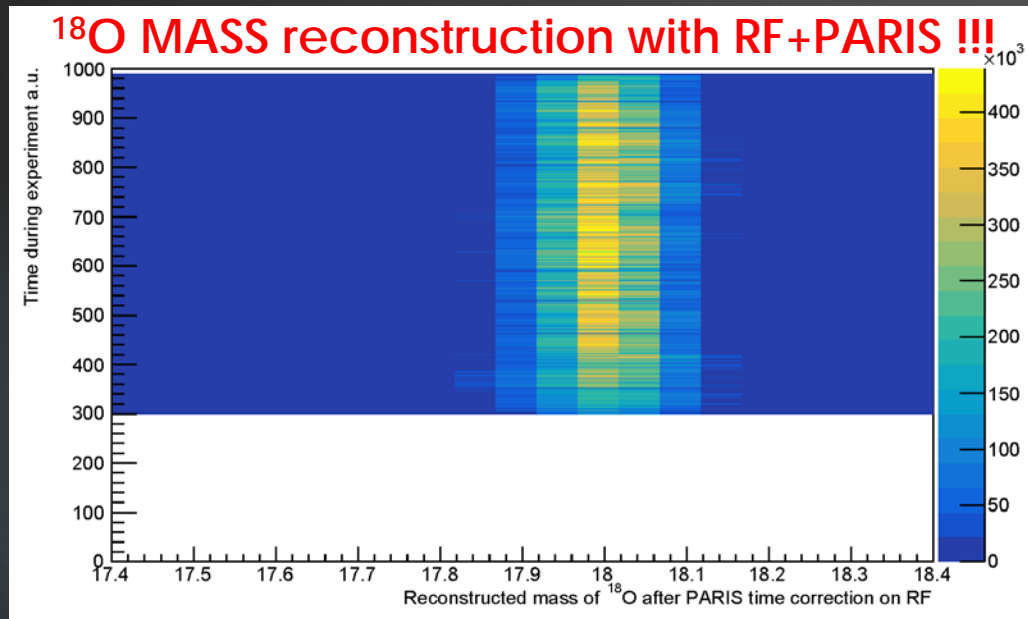


But **RF** signal is **NOT stable** in time in respect to beam on target – best observable is Mass (calculated from Brho and V)

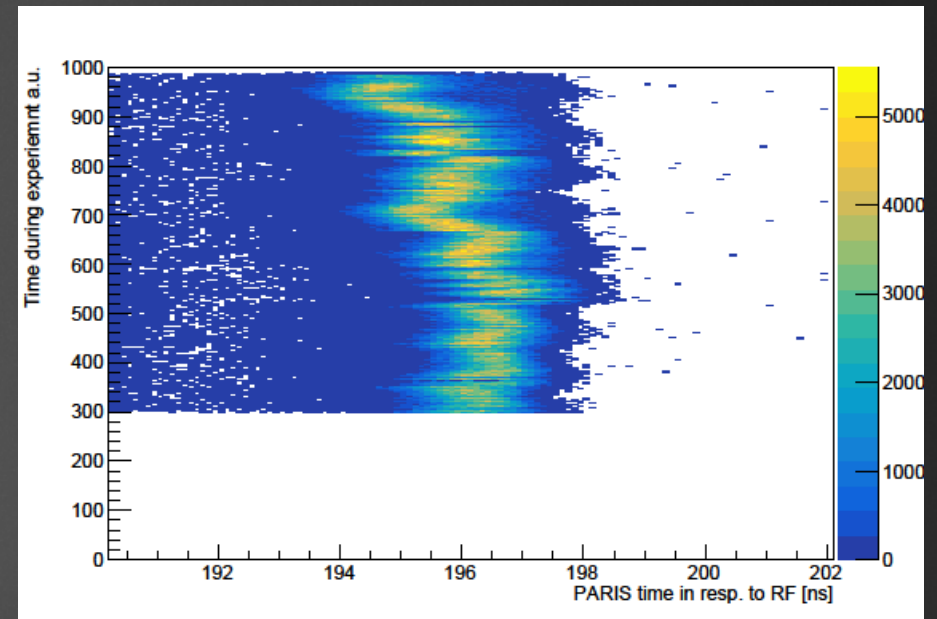
We are using PARIS (LaBr part) vs. RF timing to correct RF fluctuations (up to 2 ns, especially at the end of exp.)

PARIS timing – correction to velocity

We measure V by: measure path in spectrometer and time between **RF** and **Plastic** at the end of focal plane.



Time of experiment ↑



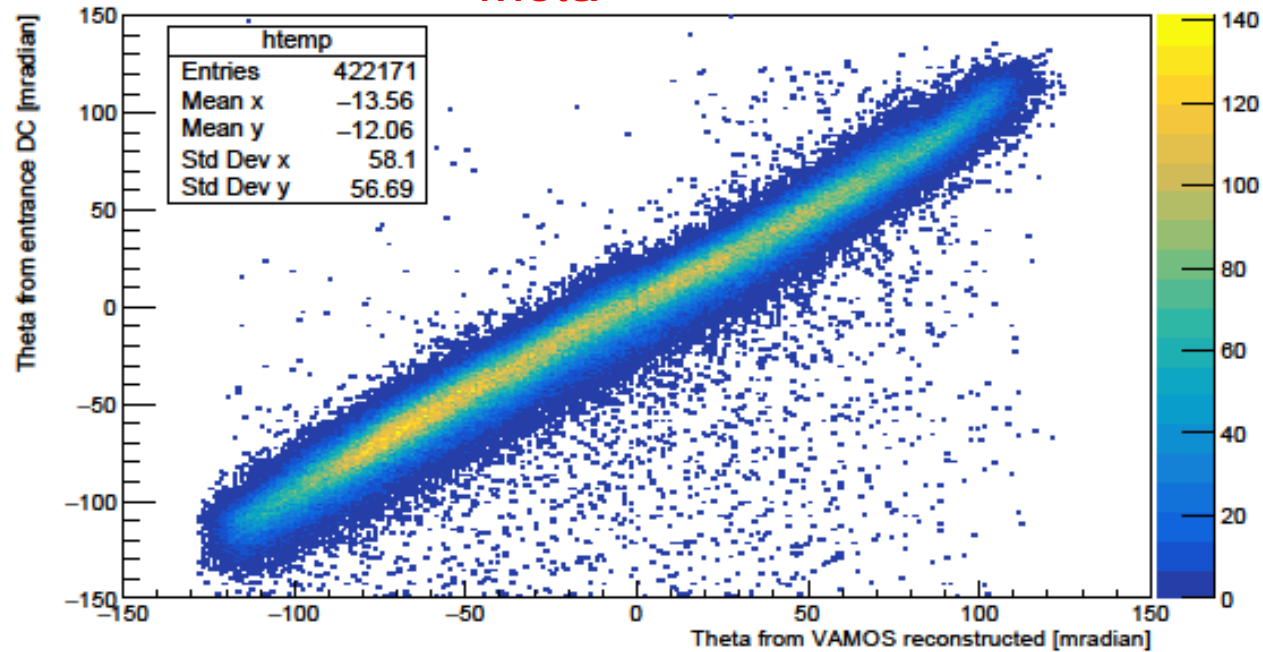
Thanks to **PARIS timing** we recovered **good A** reconstruction/stability (it means **also good V**)!

We are using (mean) PARIS vs. RF timing to correct RF fluctuations (up to 2 ns, especially at the end of exp.)

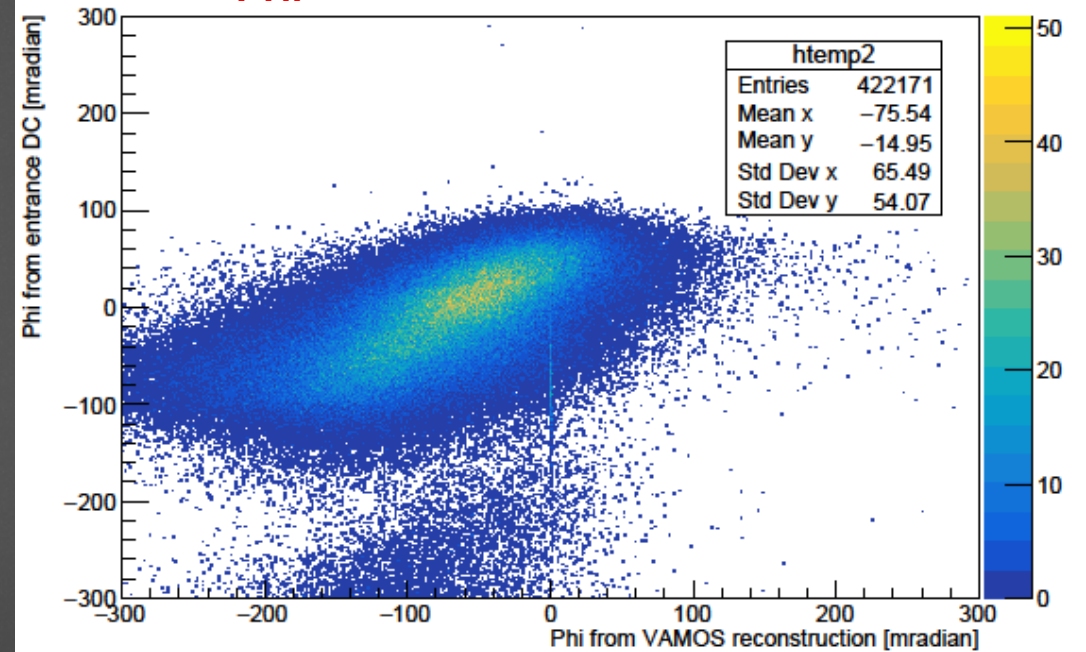
Doppler Correction (DC): ion angle determination



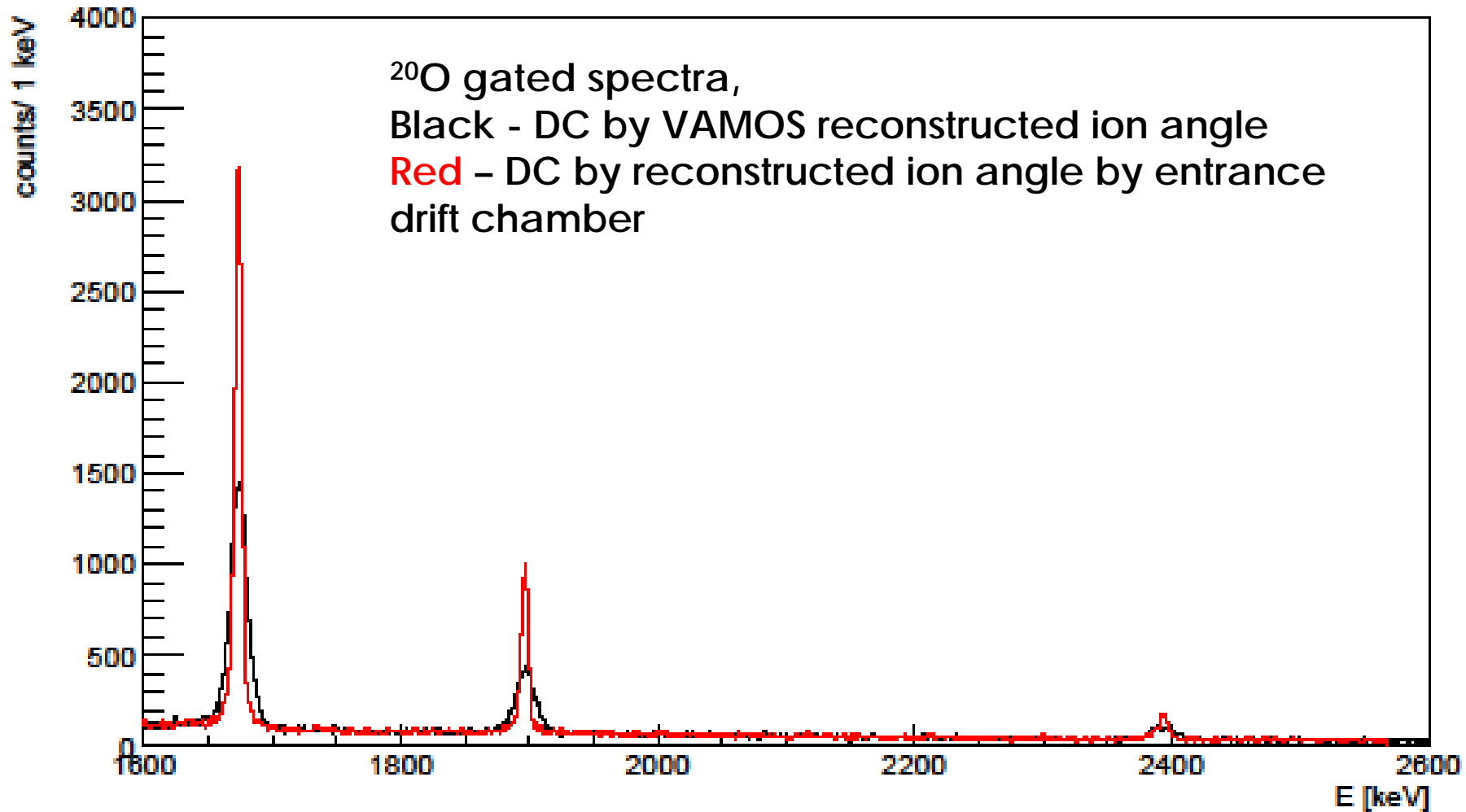
Theta



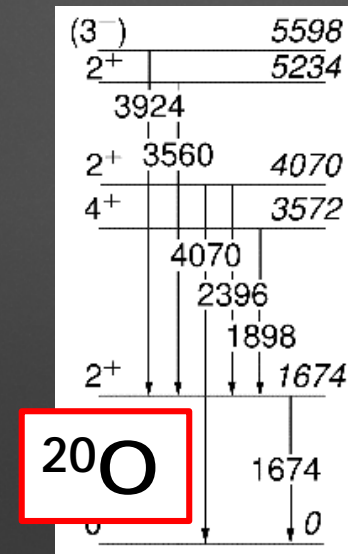
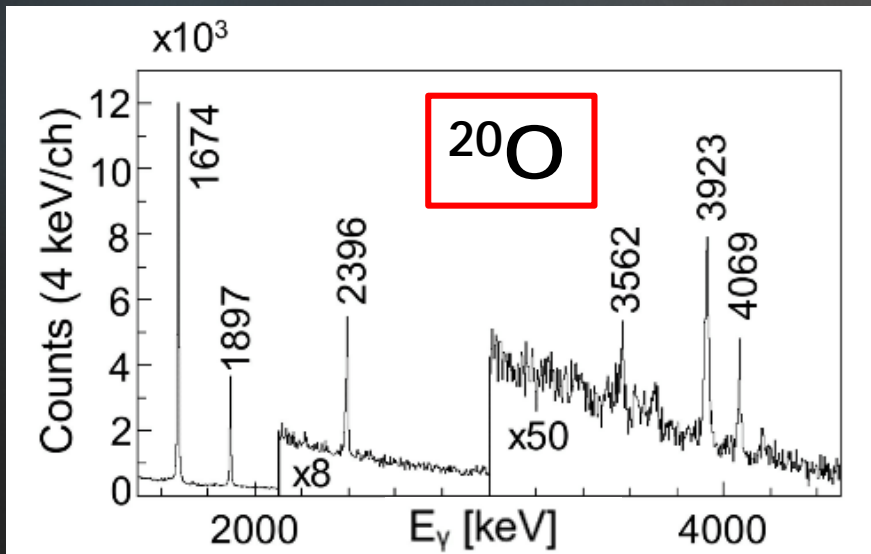
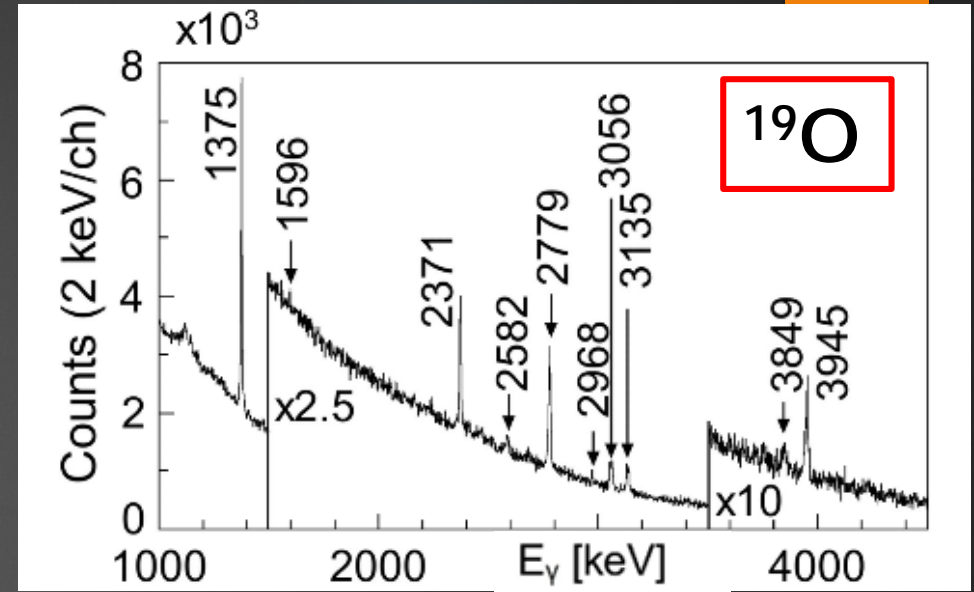
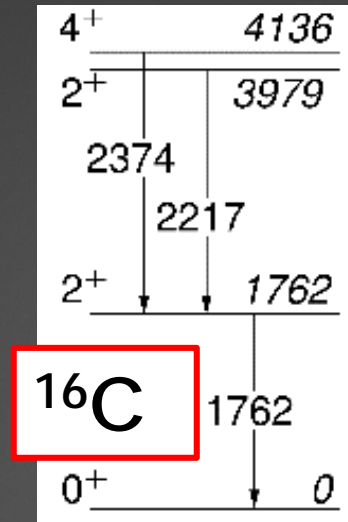
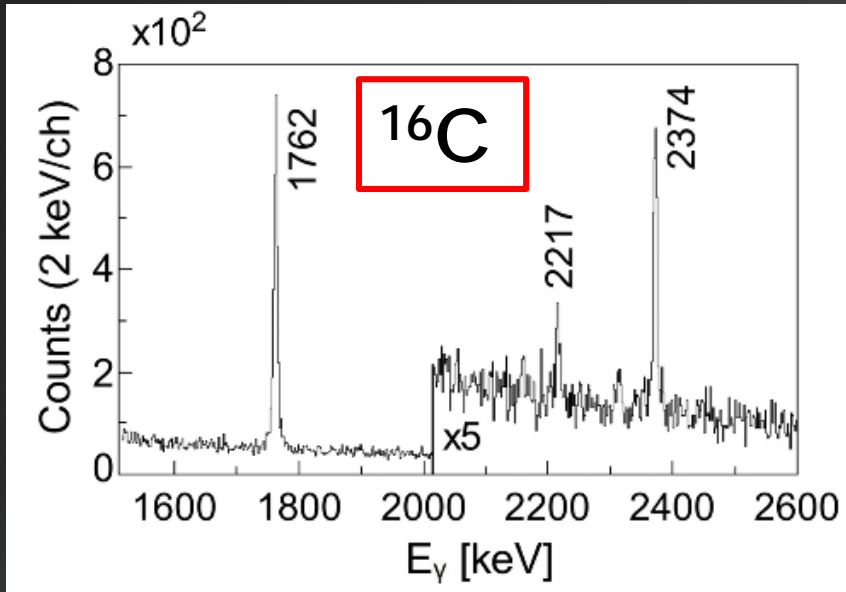
Phi



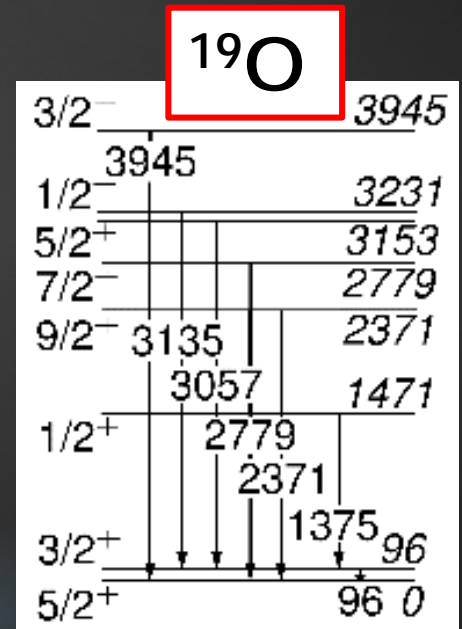
Doppler Correction (DC) – importance of entrance drift chambers



AGATA Spectra – Tracked and Doppler Corrected



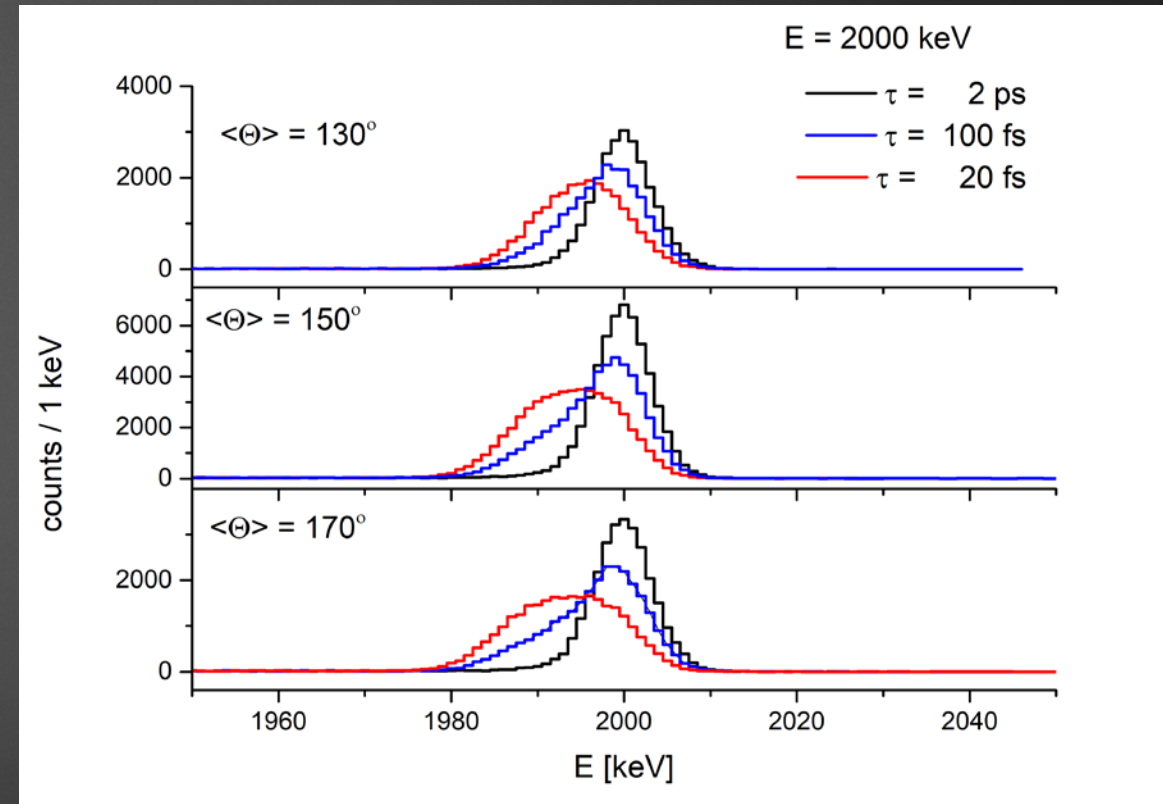
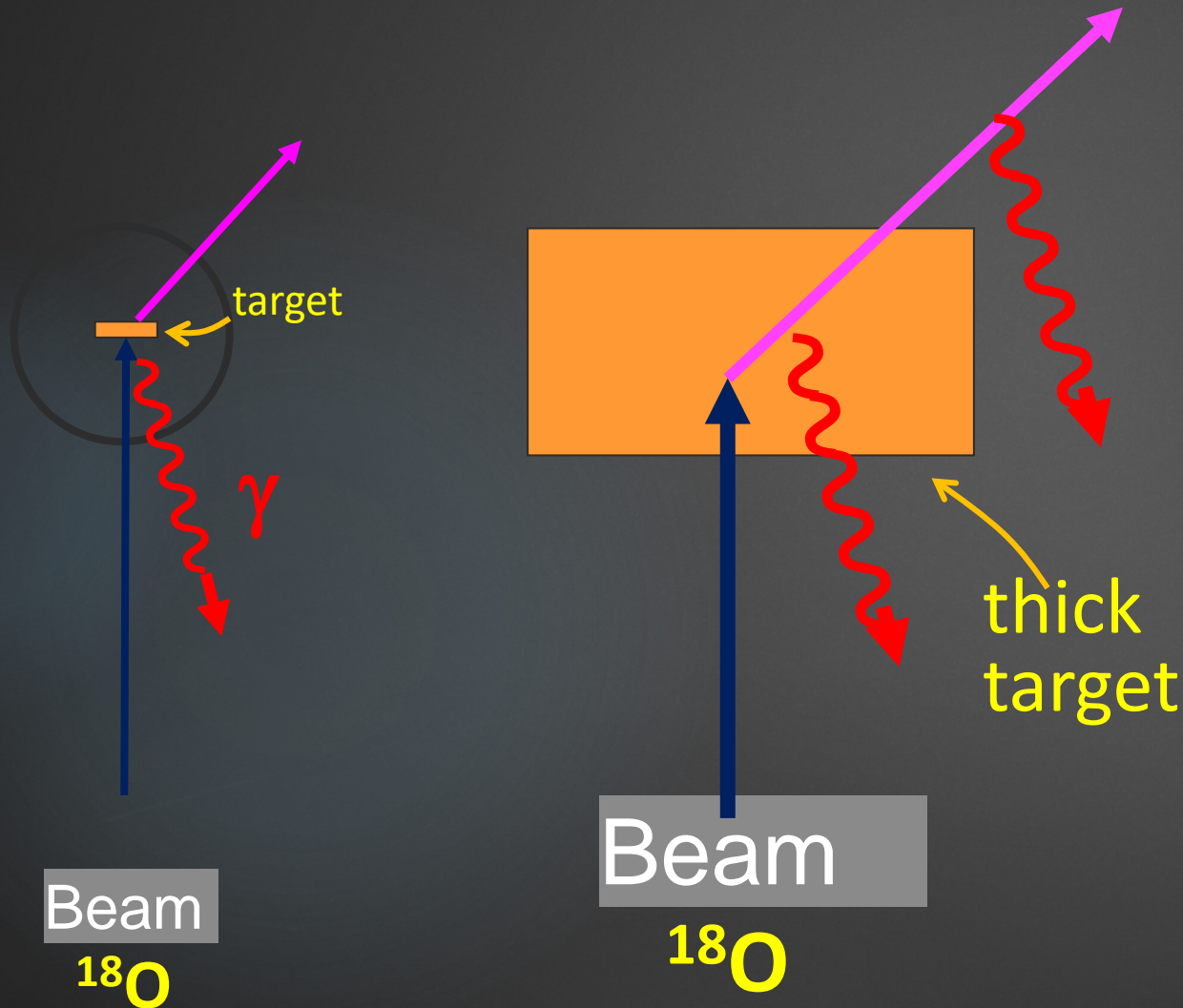
gated by
VAMOS
spectrometer



METHOD: Doppler shift dependence on the point of gamma emission

$$E = \frac{E_0}{\gamma(1 - \beta \cos\Theta)}$$

^{18}O (7 MeV/A) + ^{181}Ta target (6 mg/cm²)

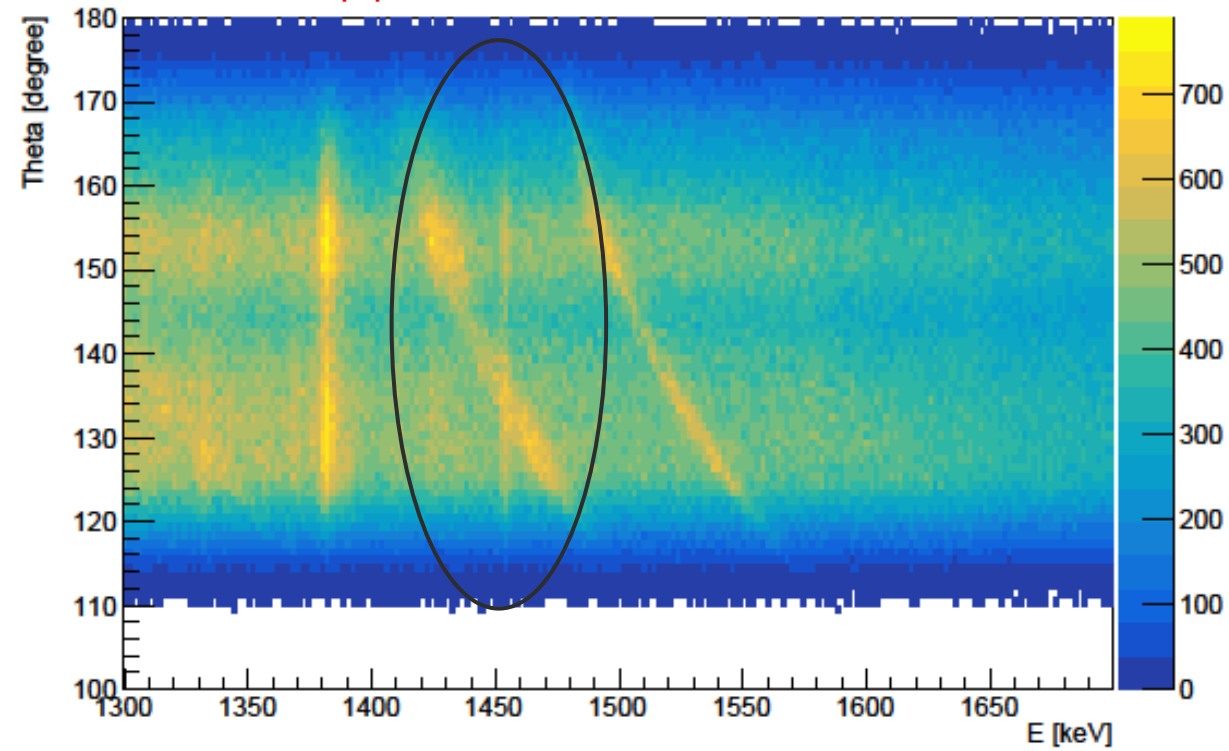


Possible problems - lines from partners

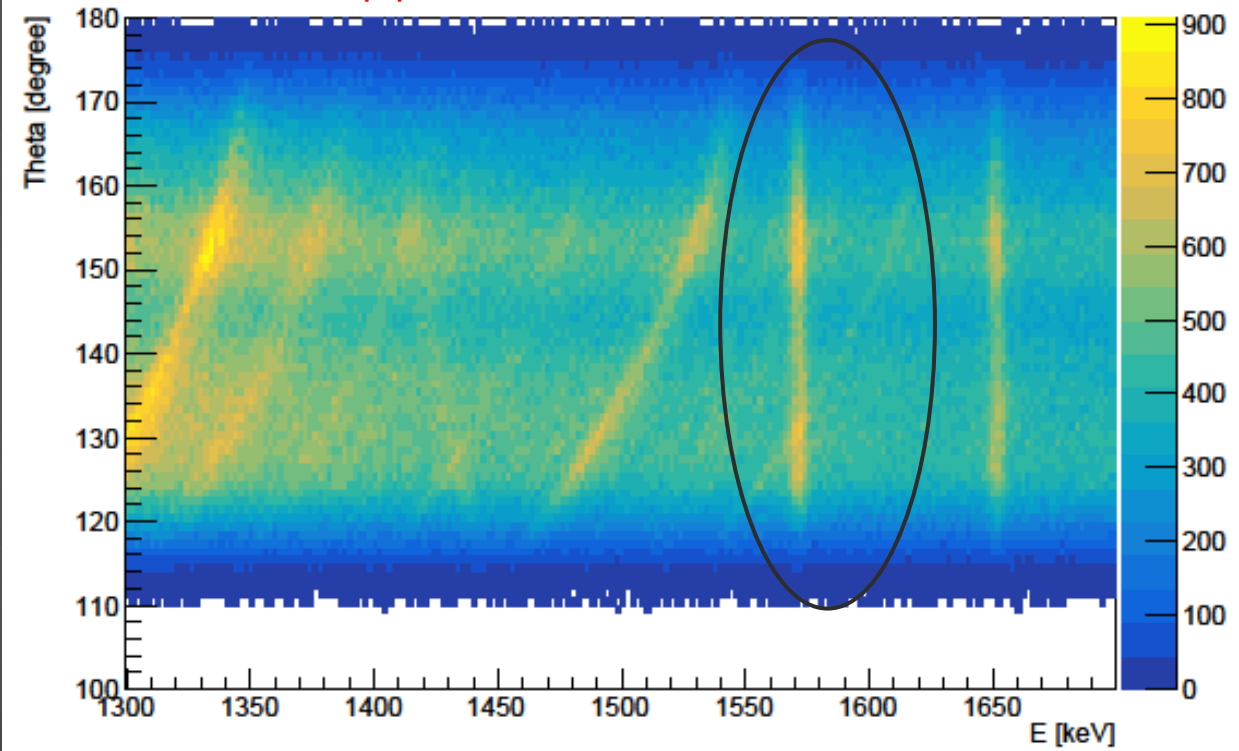
Example, ^{18}O



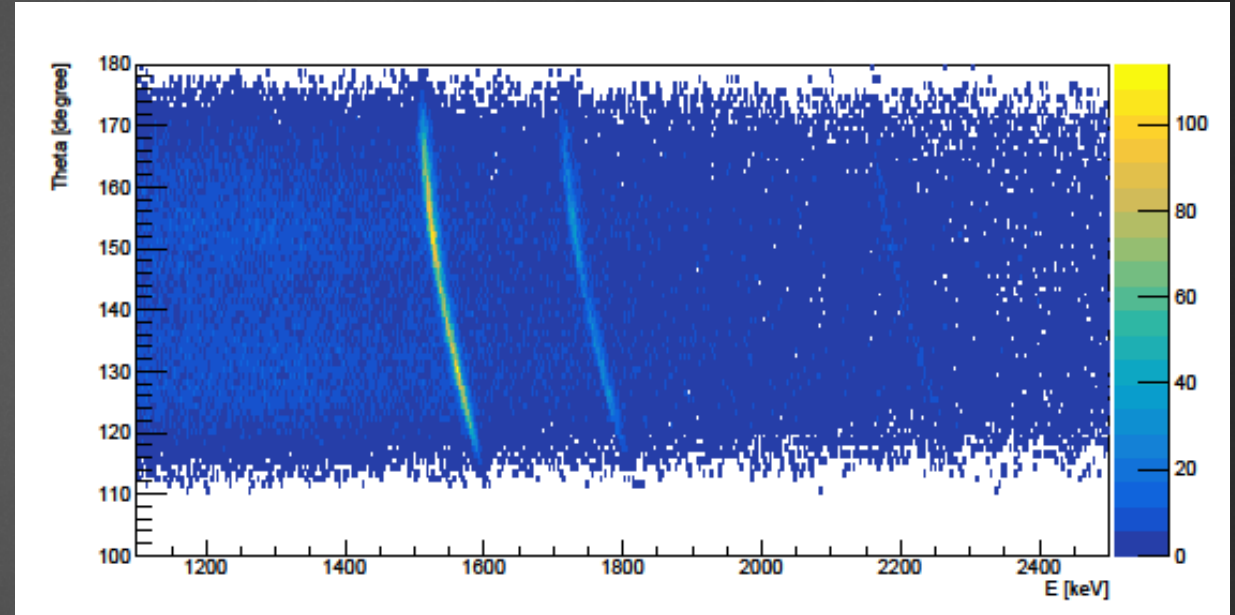
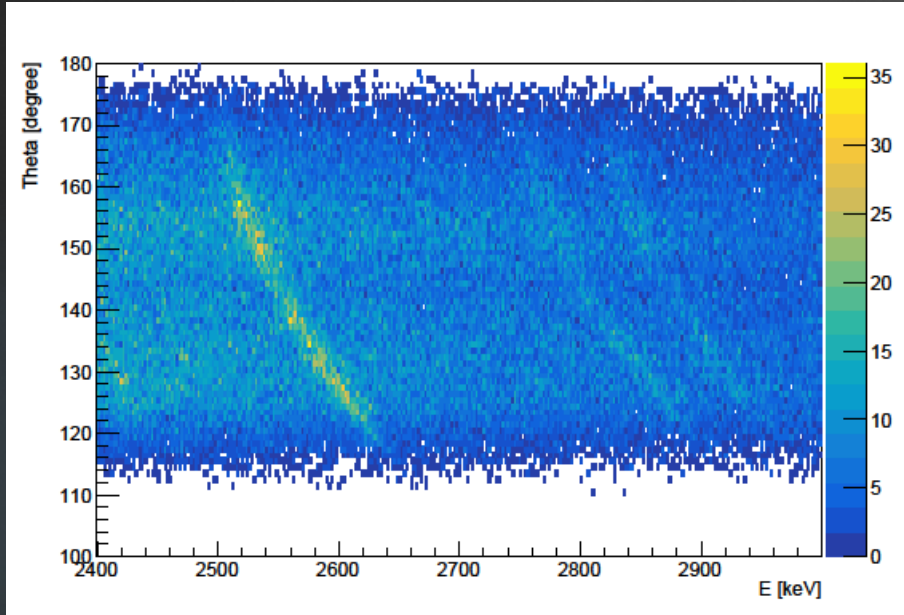
Not Doppler corrected



Doppler corrected



No contamination in all presented cases

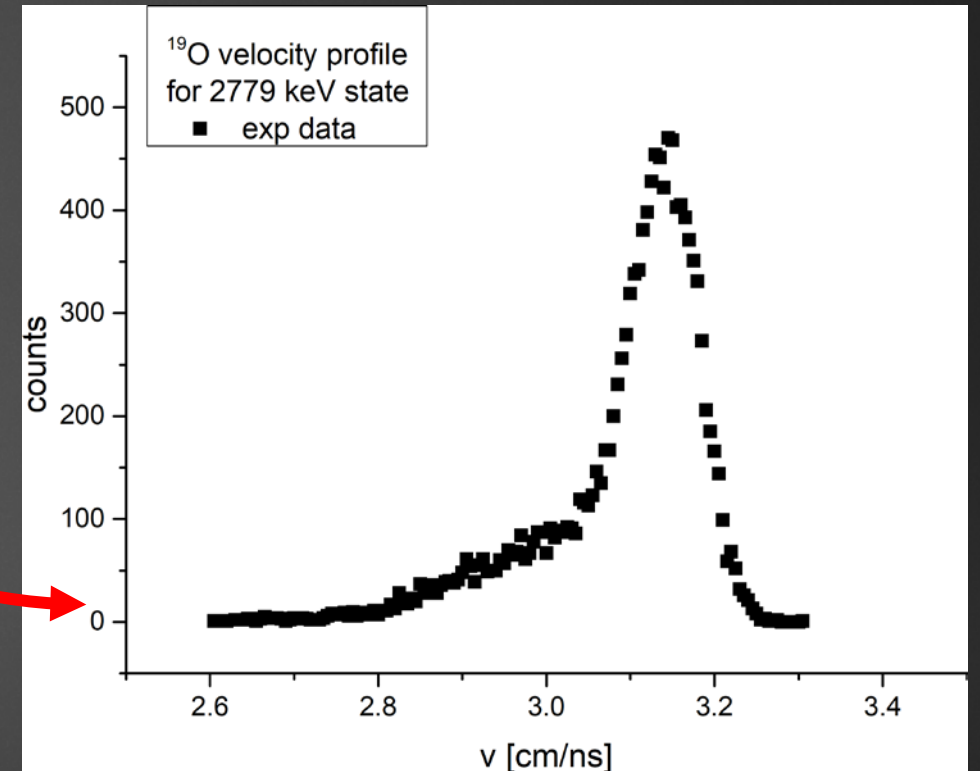
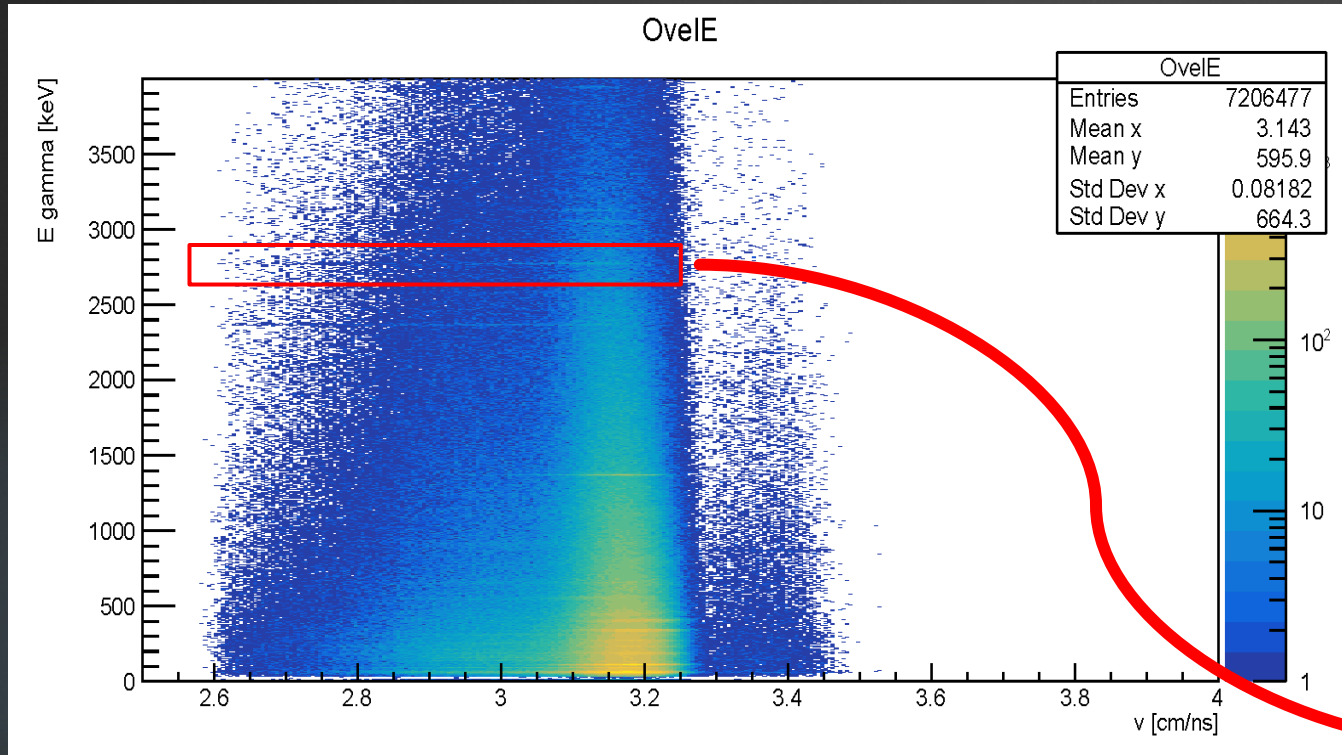


Not corrected gamma-ray spectra for ^{19}O and ^{20}O

Simulations are needed to extract lifetimes from γ lineshape

1. The beam is passing through the target decreasing its energy. Multi nucleon transfer reactions occur inside the target. An excited level is let to decay with fixed lifetime.
2. Simulation (with GEANT4 package) of AGATA response.
3. AGATA simulated data are tracked (similarly to experimental data) and Doppler corrected.
3. Experimental energy resolution of AGATA crystals and differences in counting rates are included in the simulation.
3. 2D χ^2 maps are used to determine optimum lifetime.

Simulation ingredients - velocity

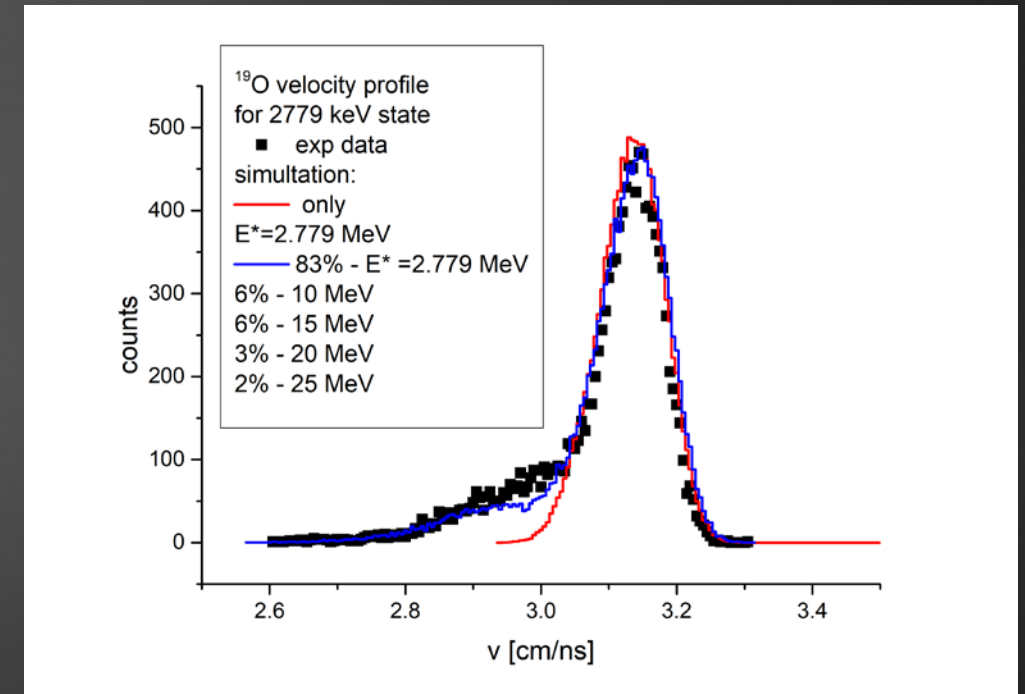
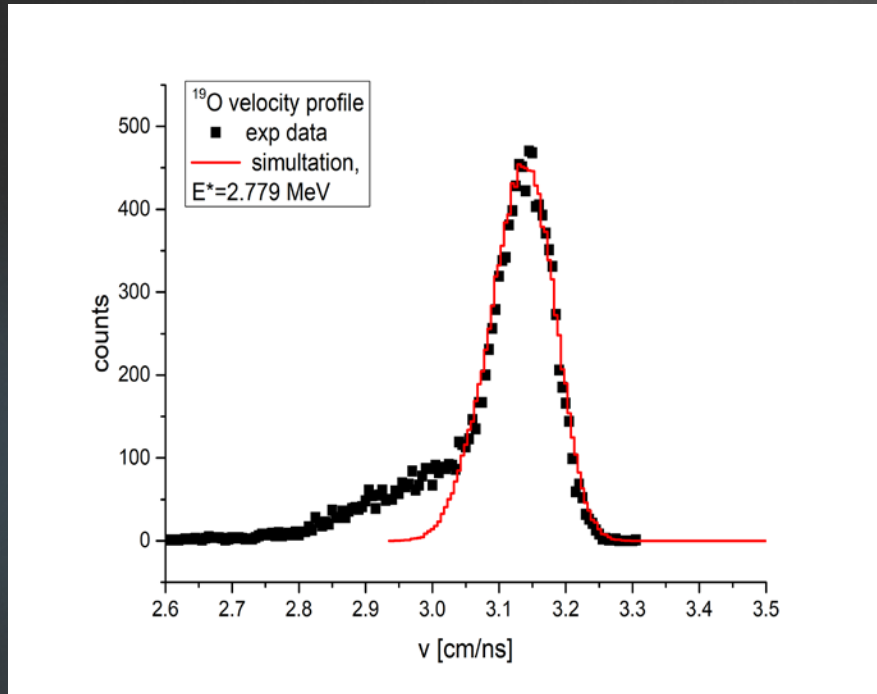


Example:
¹⁹O ion velocity and gamma-ray matrix

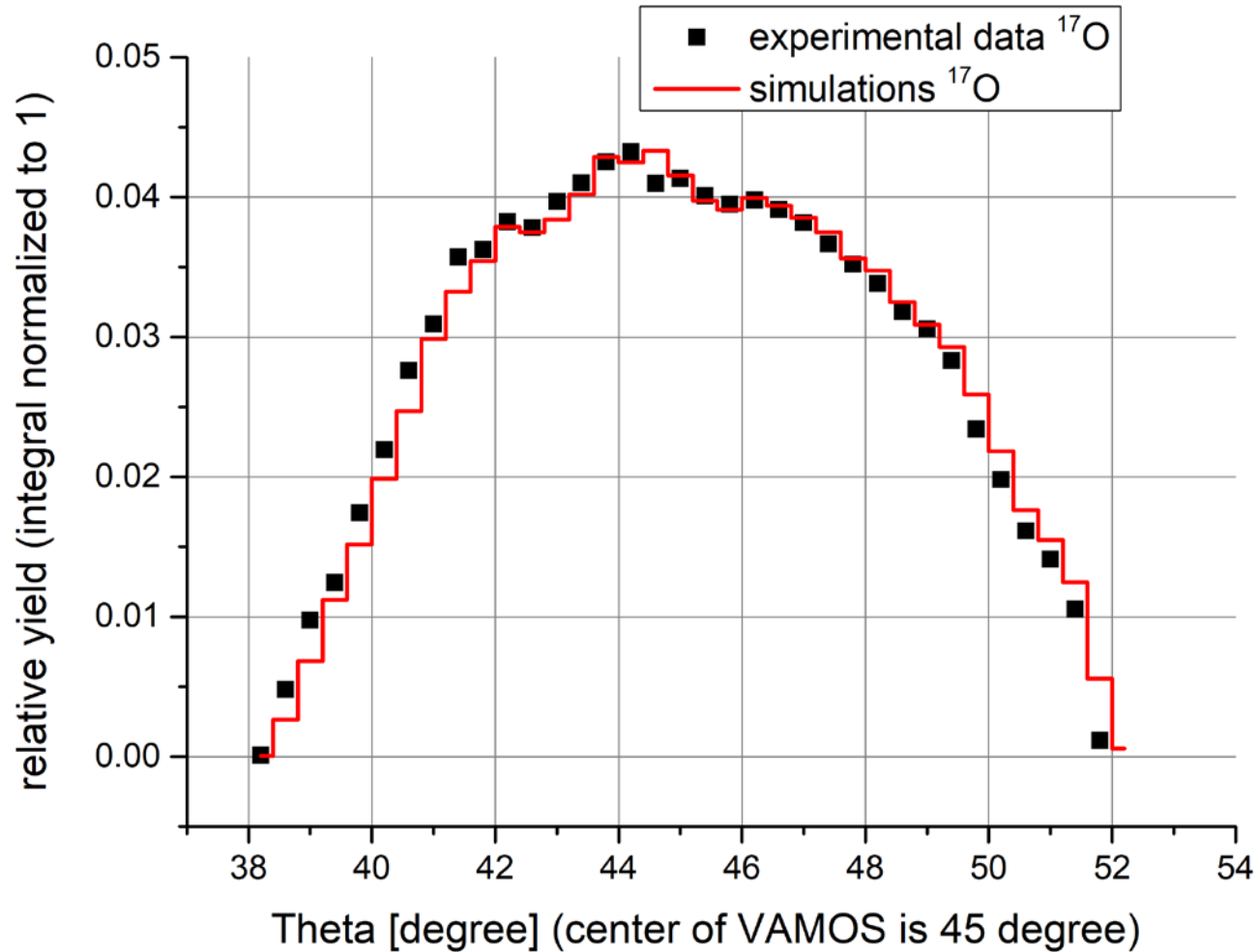
Simulation ingredients - velocity

Example:

Iterative method to reconstruct initial ion velocity from measured velocity

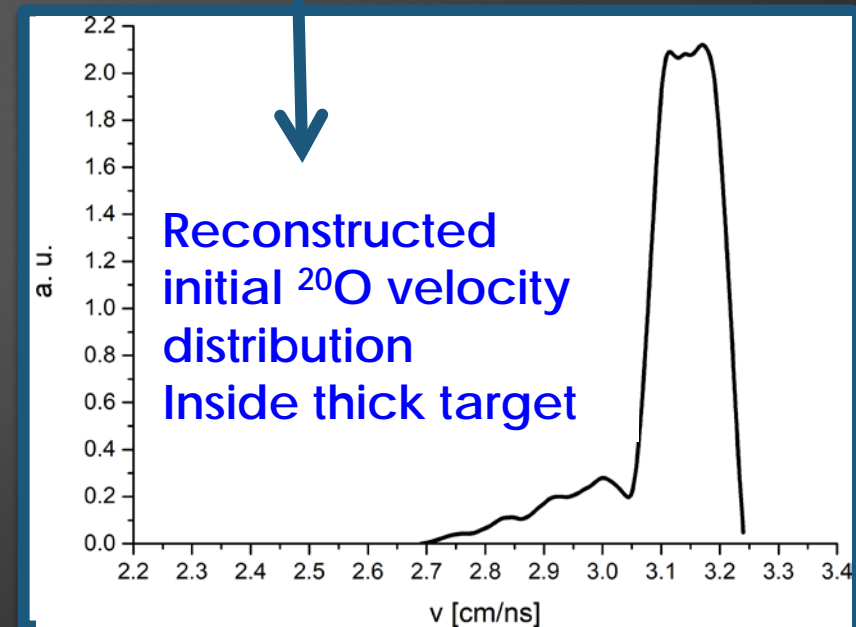
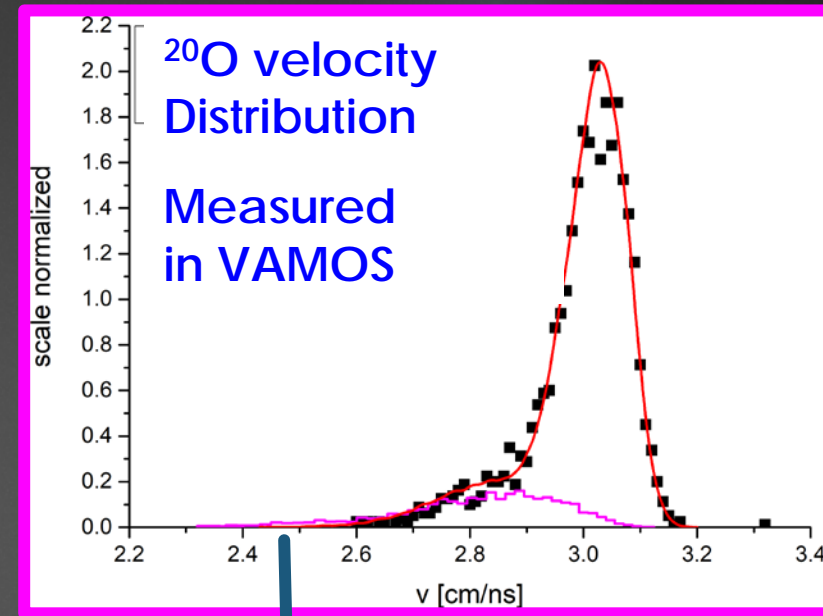
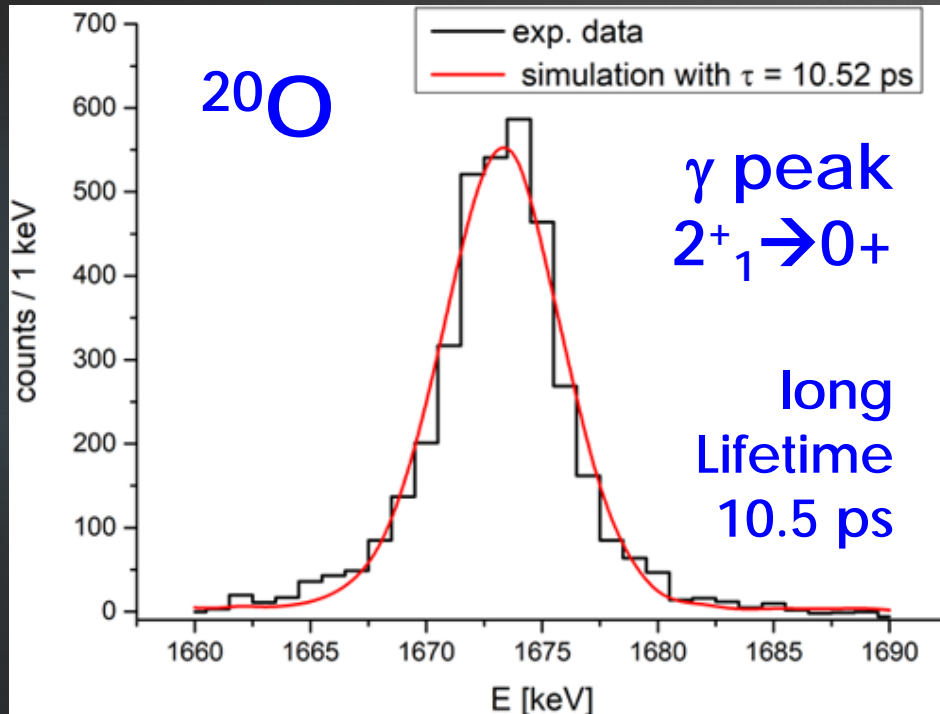


Simulation ingredients - theta

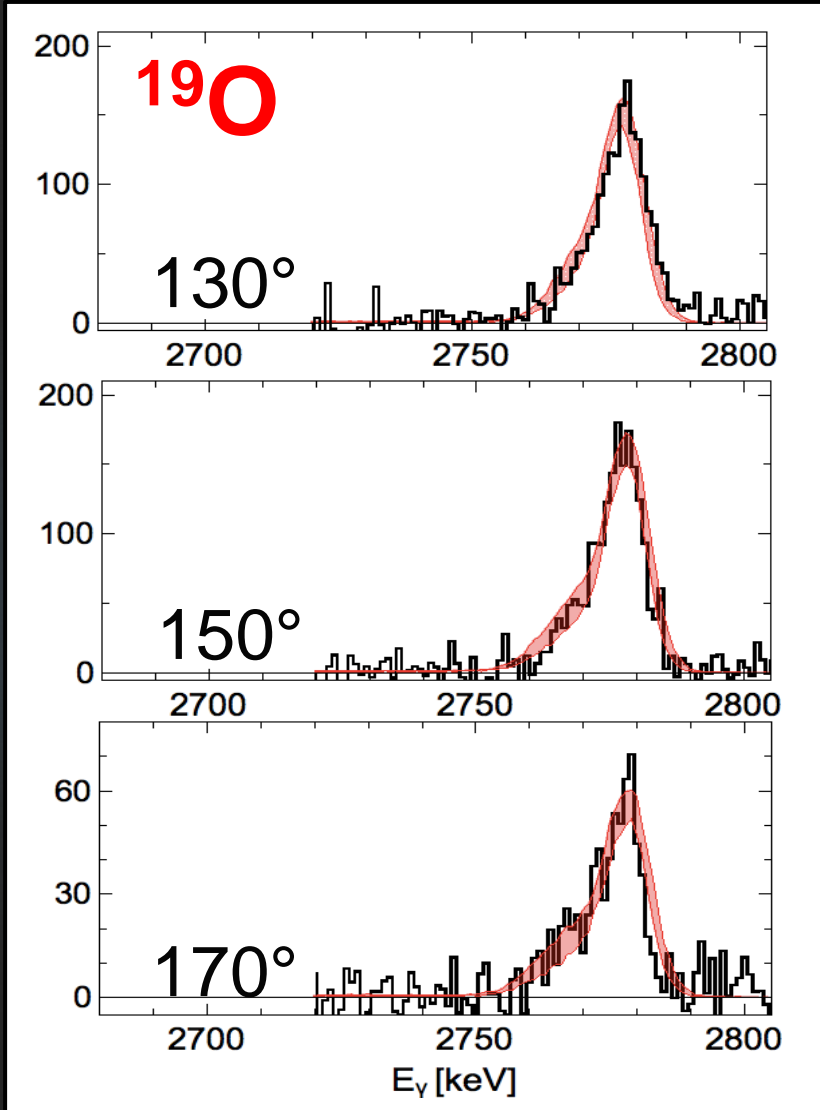


Simulation ingredients - v

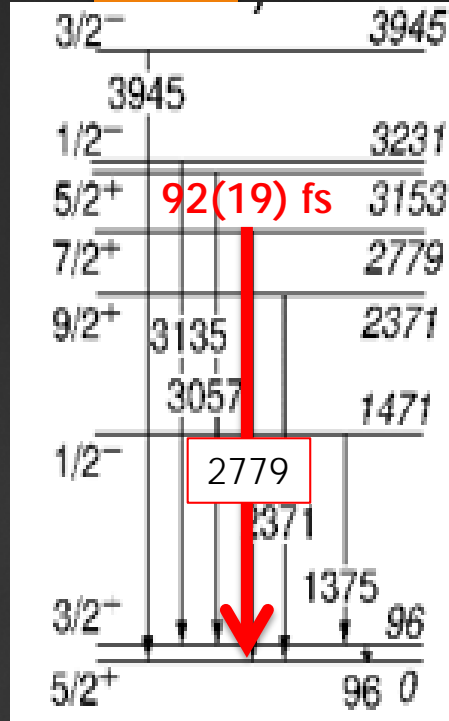
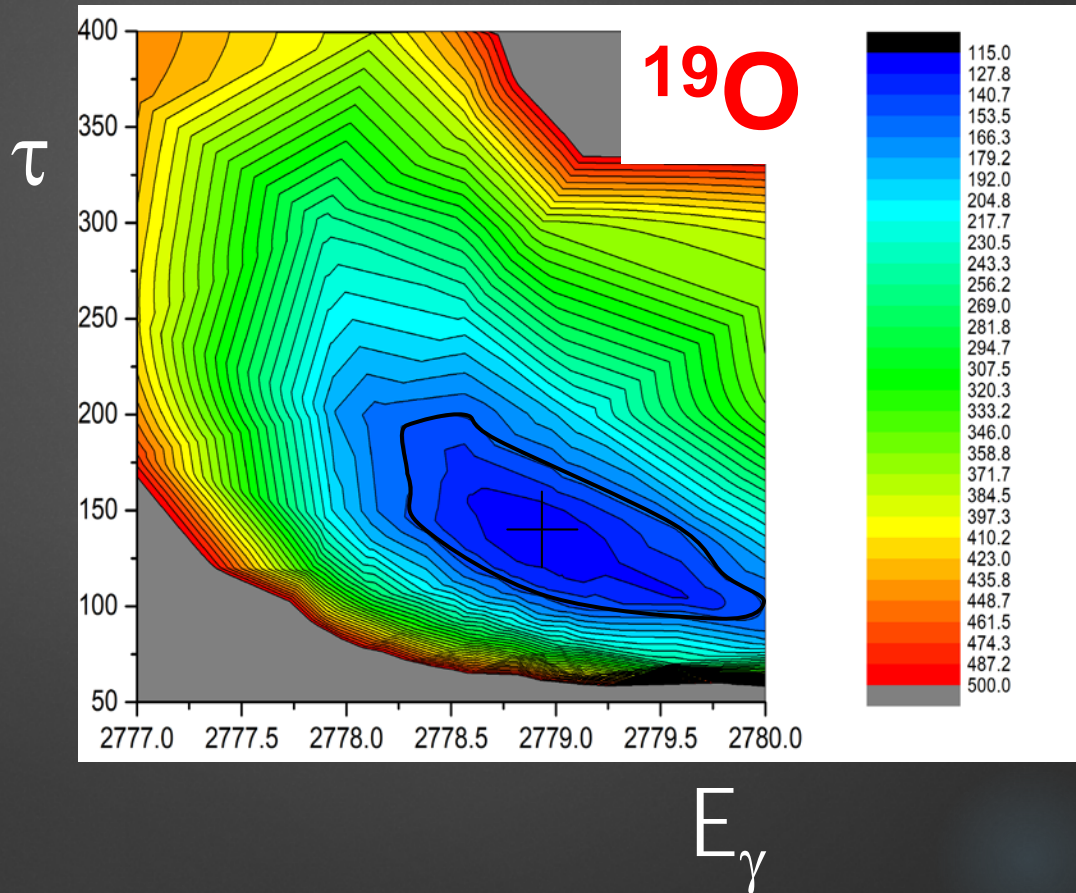
^{20}O – excited state of interest



TEST of KNOWN lifetimes in the 100s femtoseconds region



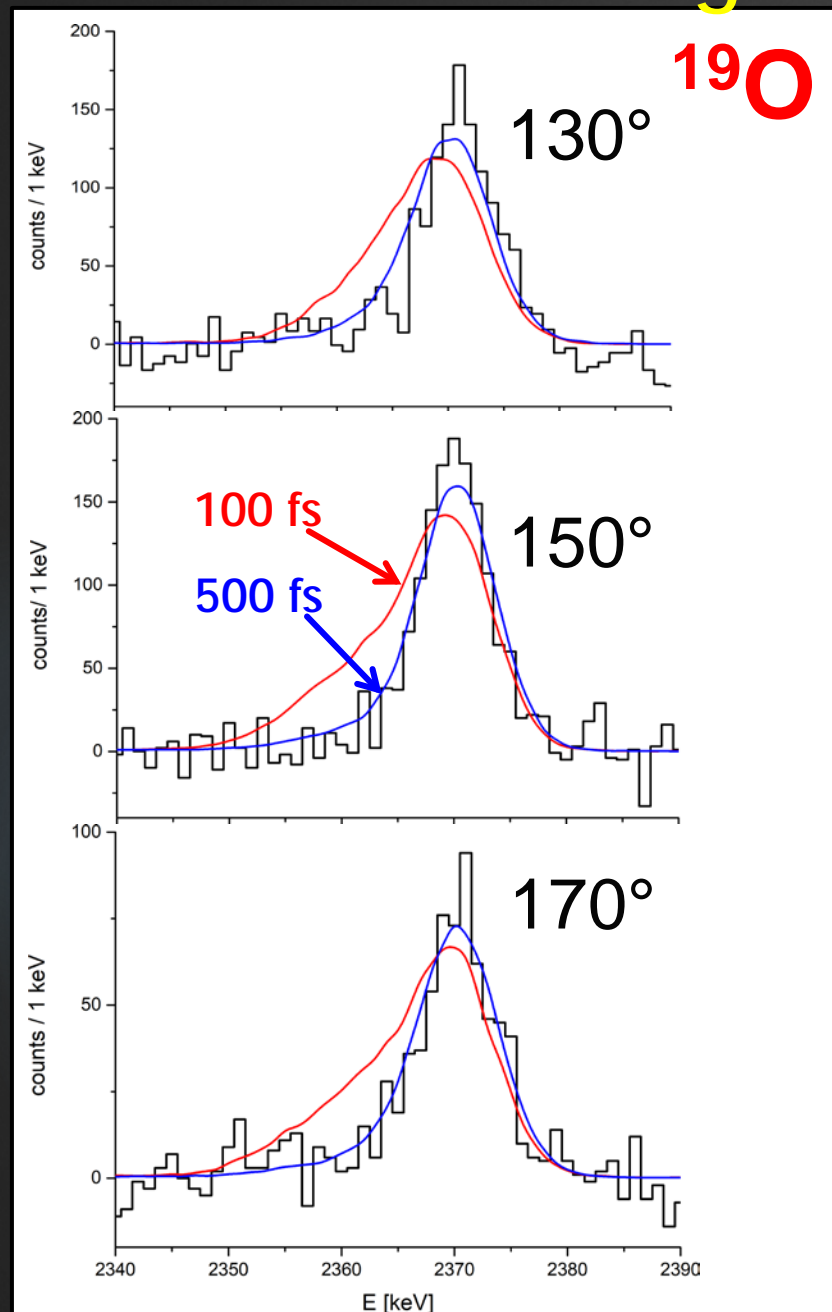
^{18}O (7 MeV/A) + ^{181}Ta target (6 mg/cm²)



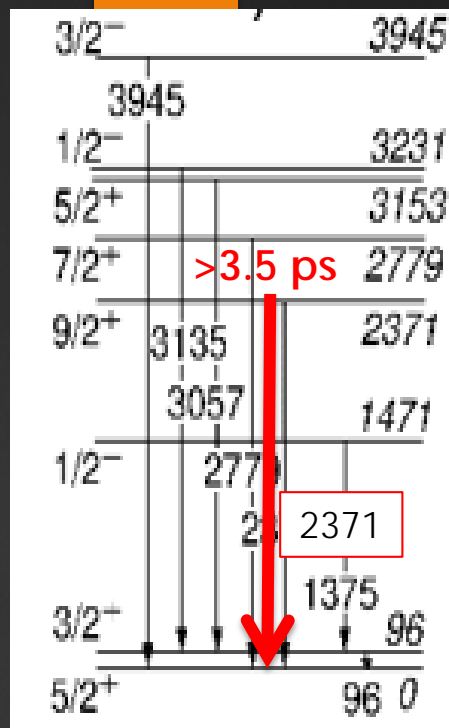
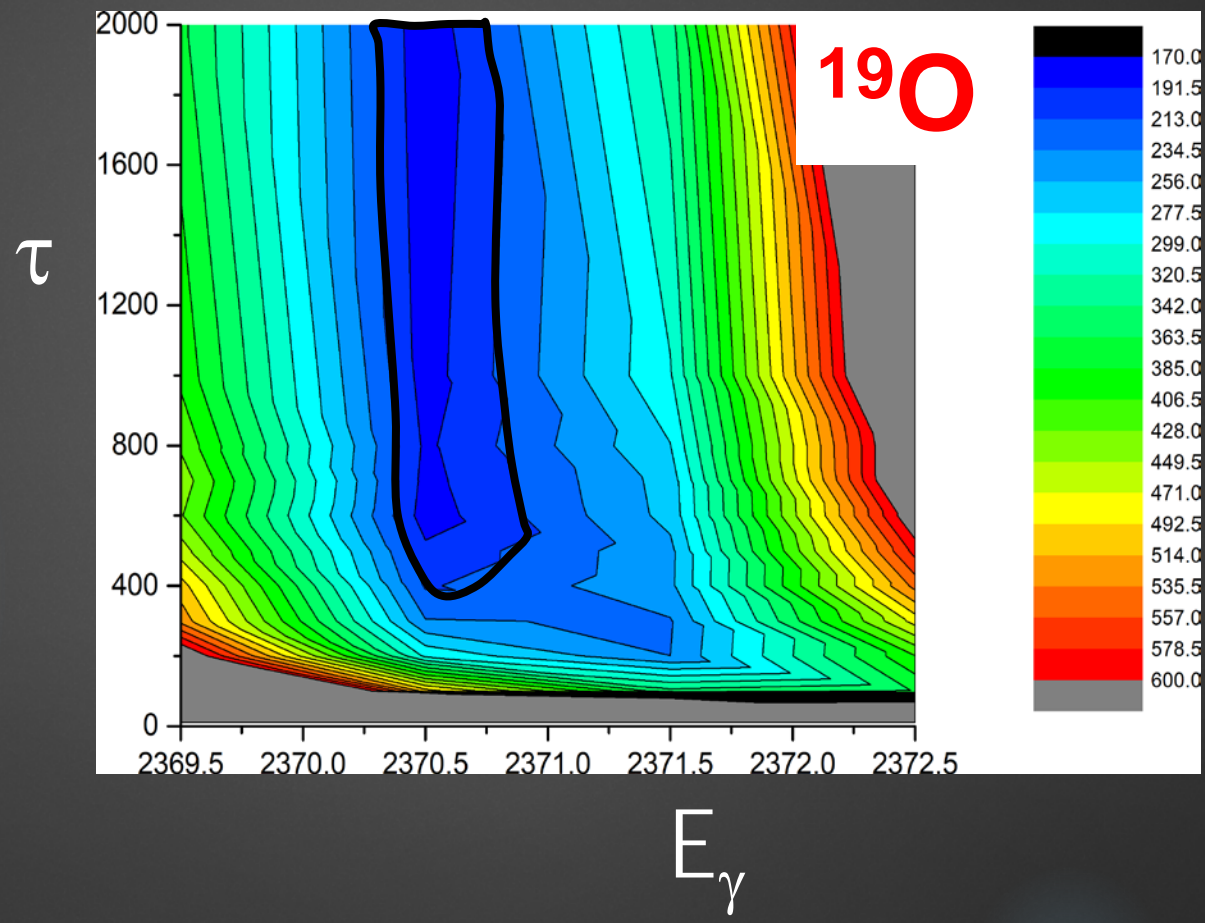
$$\tau = 140^{+50}_{-40} \text{ fs}$$

Very old literature values (1971)
 $\tau = 70(26) \text{ fs}$
 $\tau = 117(26) \text{ fs}$

TEST of KNOWN long lifetime in the region

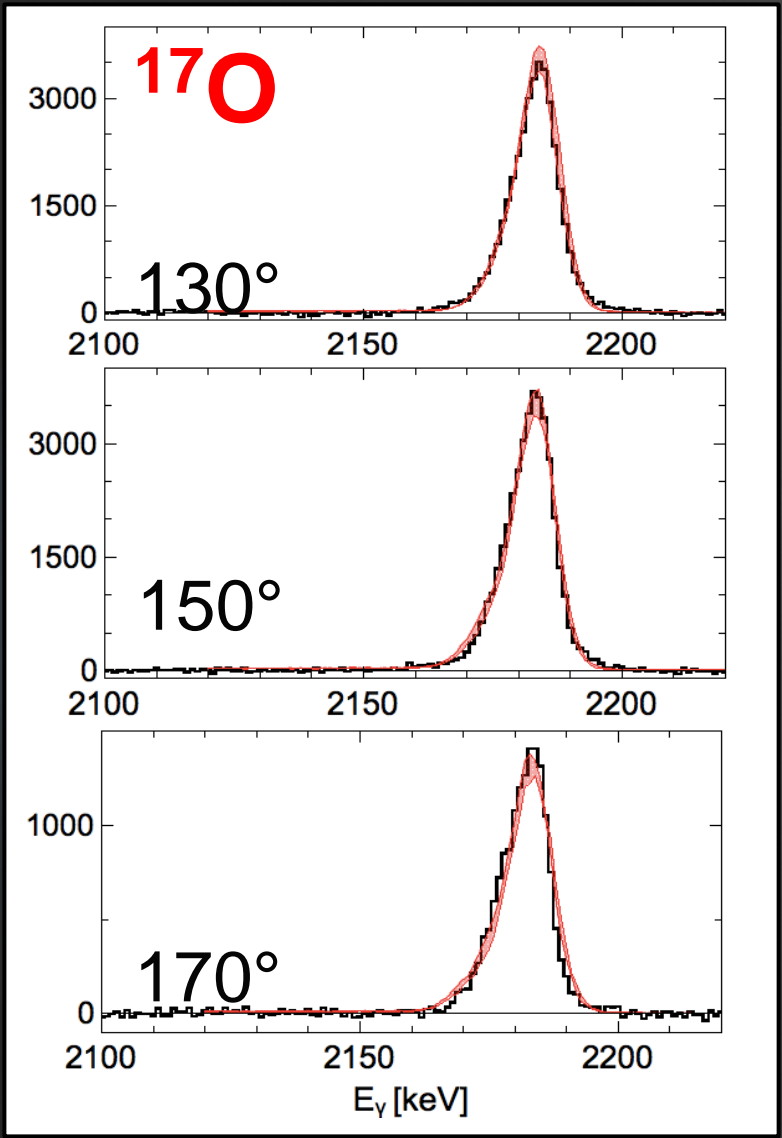


¹⁸O (7 MeV/A) + ¹⁸¹Ta target (6 mg/cm²)

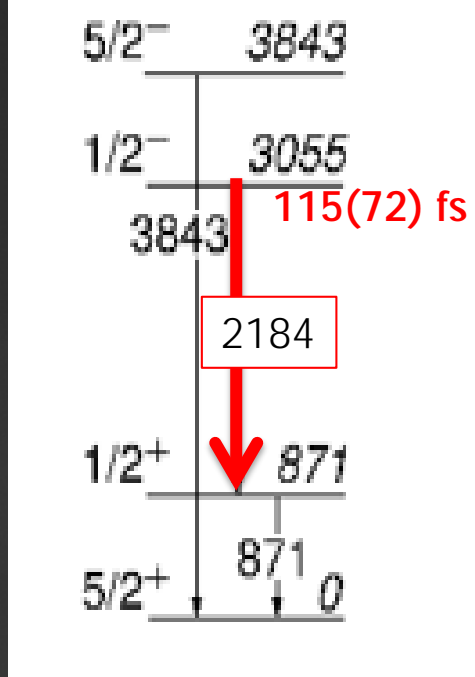
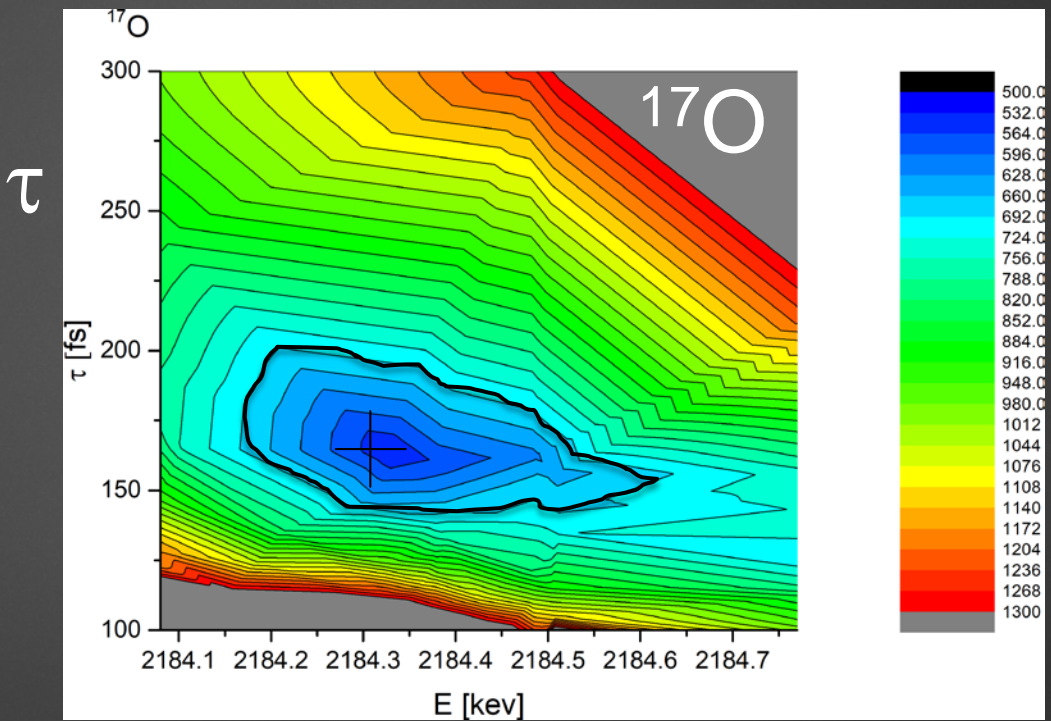


$\tau > 400$ fs

TEST of KNOWN Lifetimes in the 100s femtoseconds region



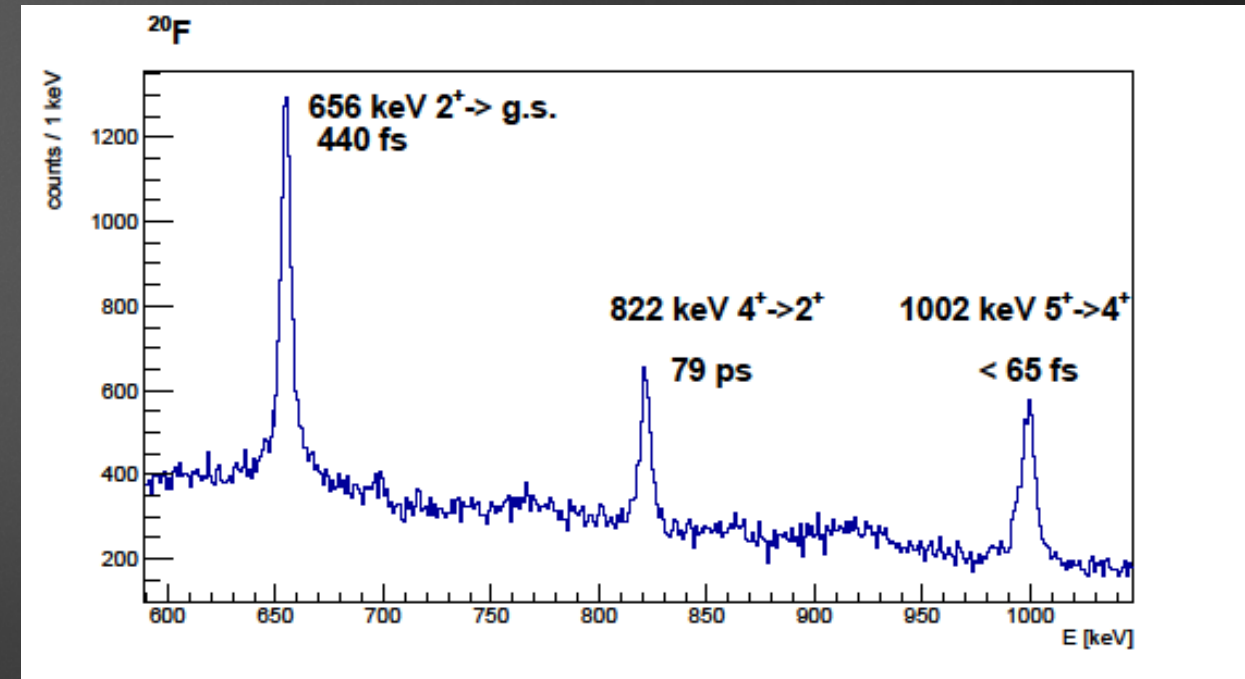
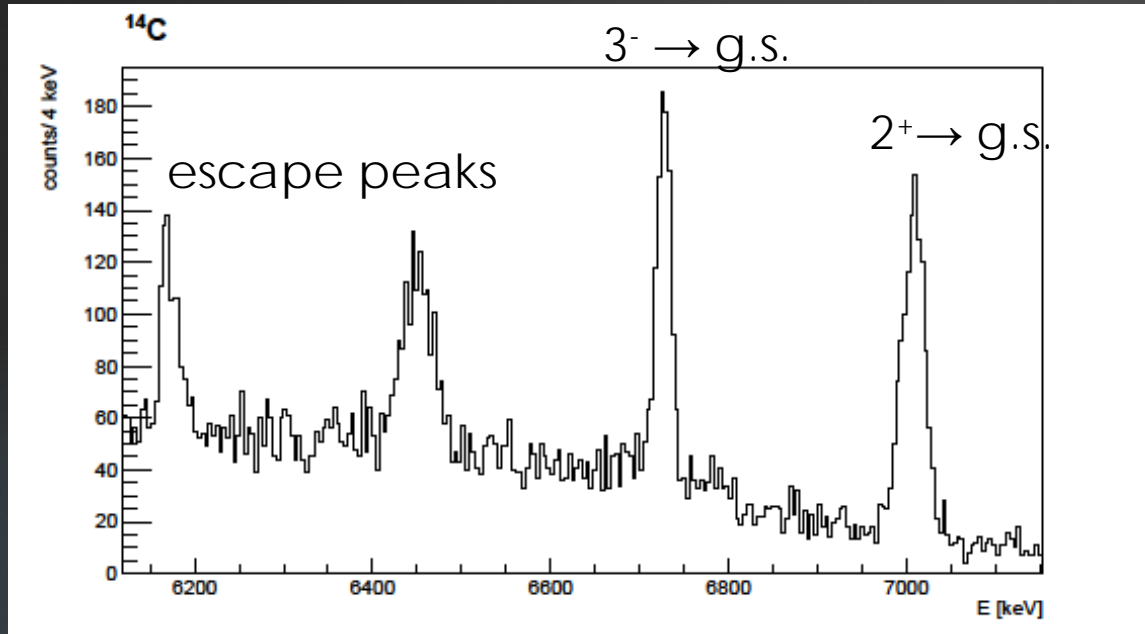
^{18}O (7 MeV/A) + ^{181}Ta target (6 mg/cm²)



$$\tau = 159^{+40}_{-20} \text{ fs}$$

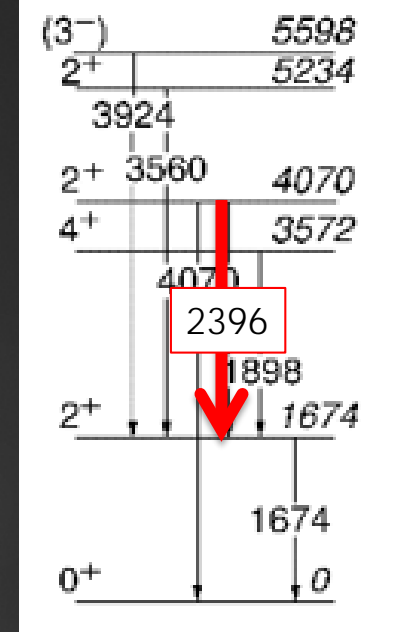
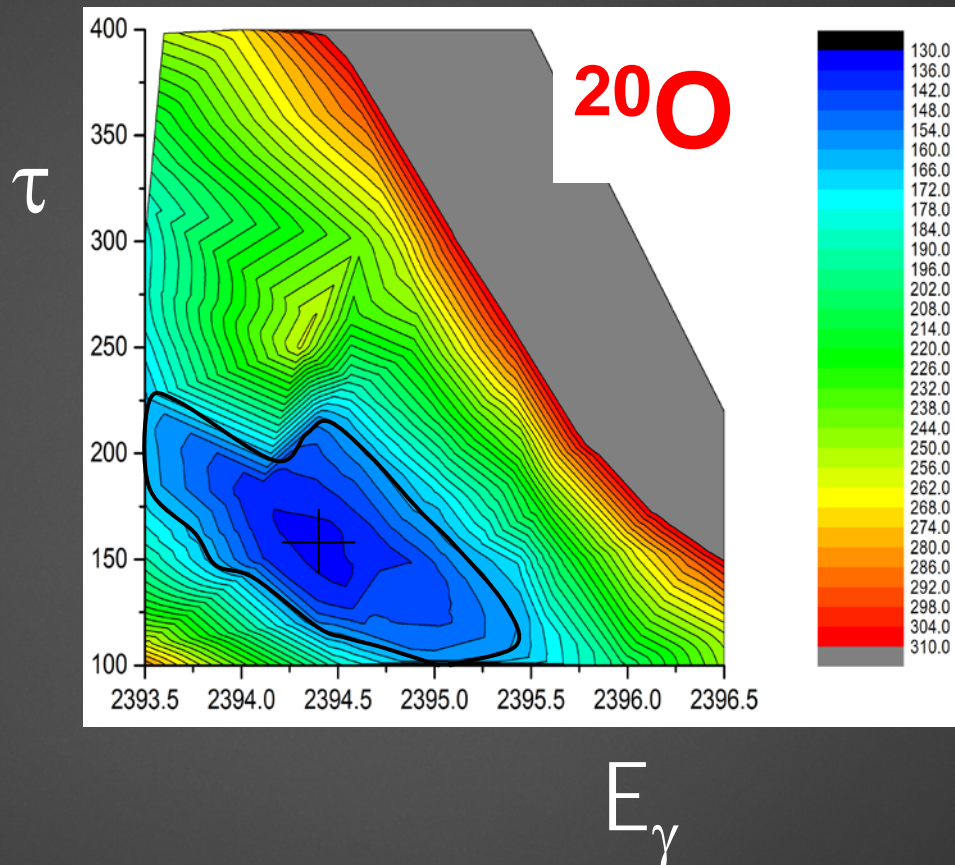
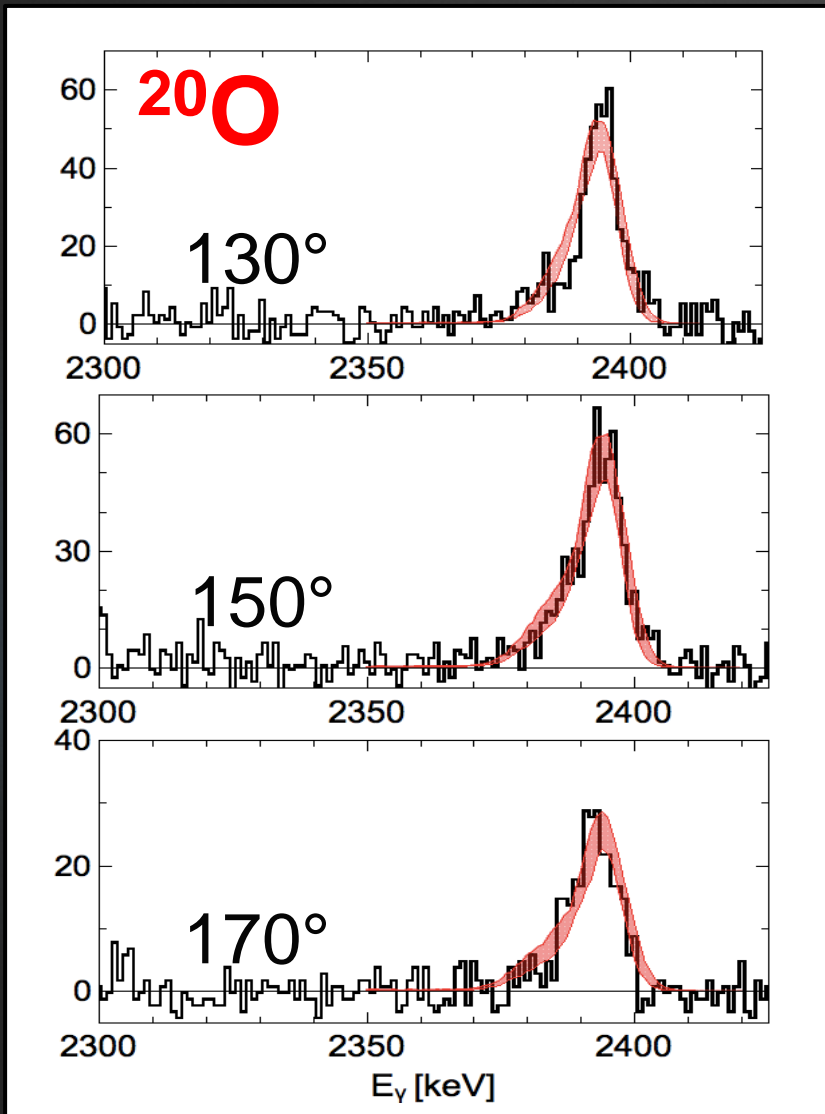
Very old literature values (1964)
 $\tau = 120(+80, -60) \text{ fs}$

Observed other known lifetimes – i.e. in ^{14}C , ^{20}F ...



OUR Case of interest – ^{20}O

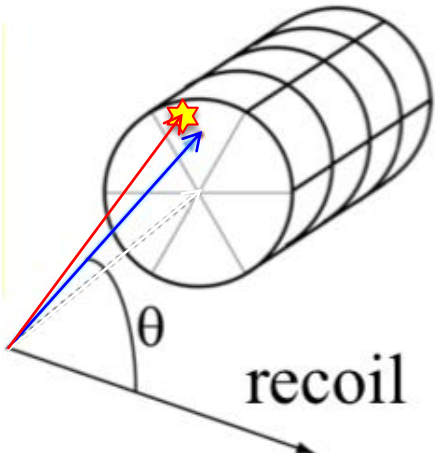
^{18}O (7 MeV/A) + ^{181}Ta target (6 mg/cm 2)



$$\tau = 150^{+80}_{-30} \text{ fs}$$

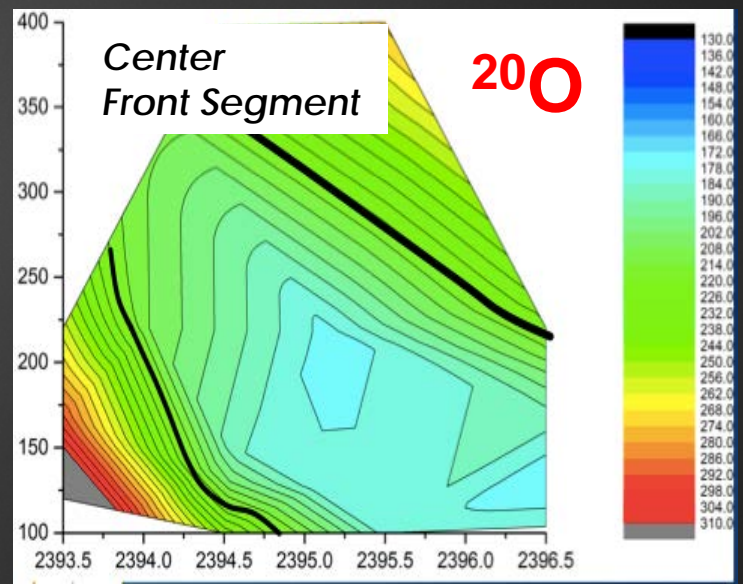
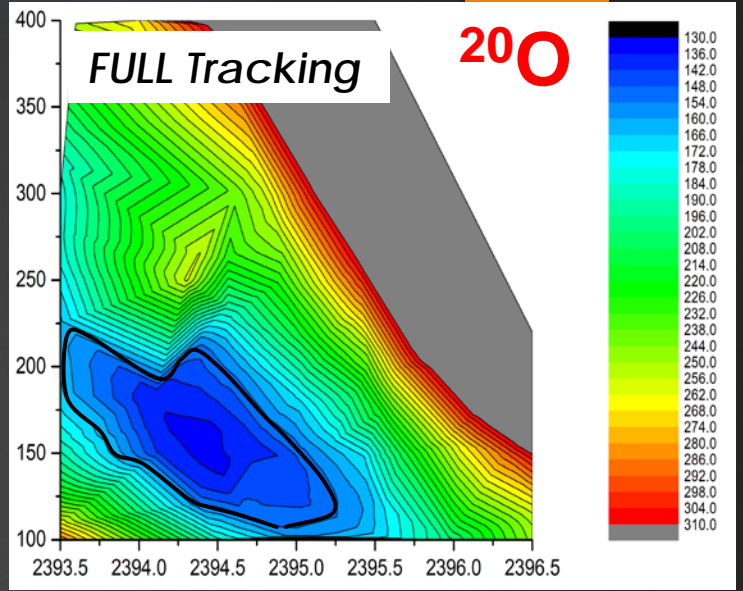
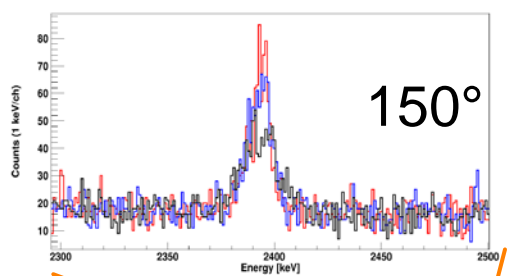
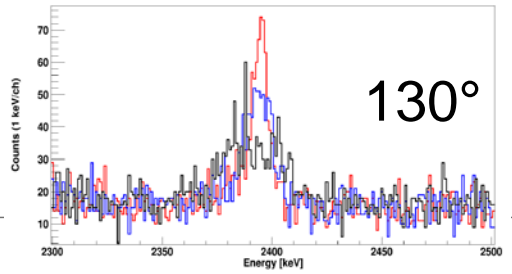
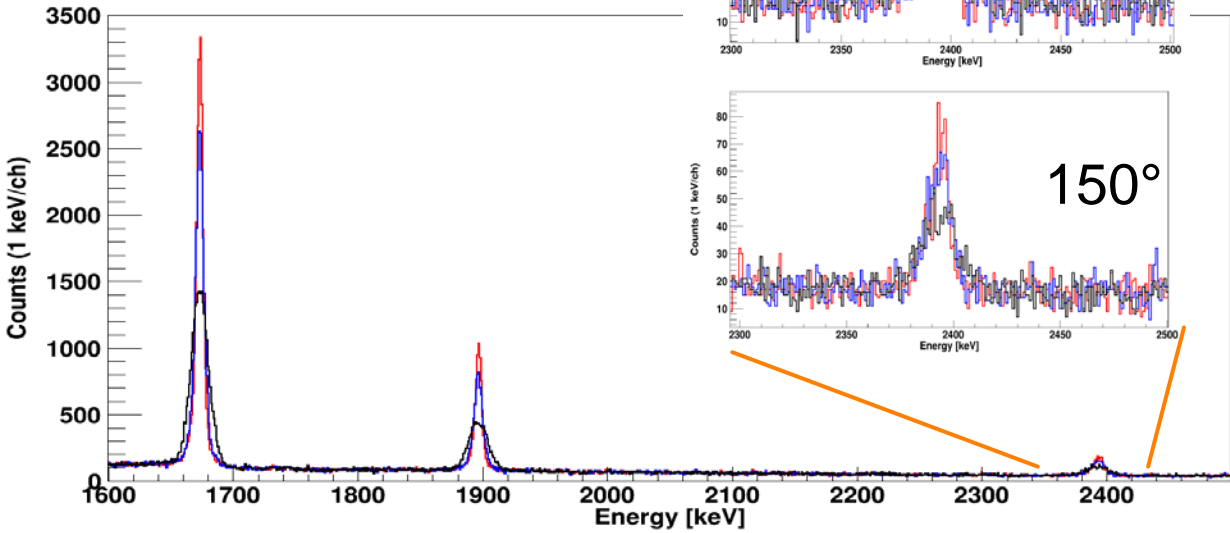
For $2_2 \rightarrow 2_1$ decay (79(5)% branching ratio),
 partial $\tau = 190^{+102}_{-39} \text{ fs}$

Large improvement with AGATA



$$\Delta E_{\gamma} = 2E_{\gamma 0} \frac{v}{c} \sin \theta_{\gamma} \sin \Delta \theta$$

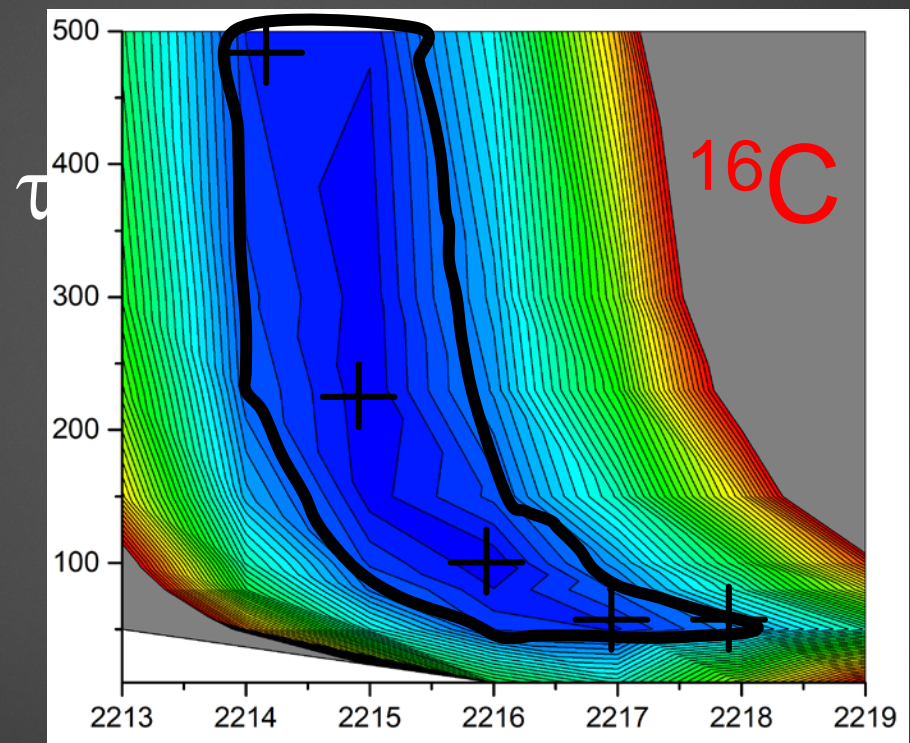
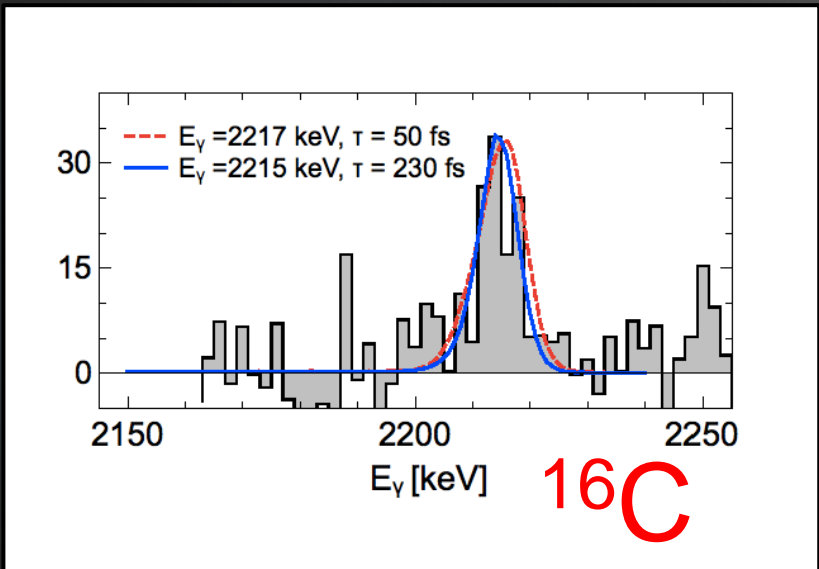
$2^+_2 \rightarrow 2^+_1$ (2396 keV)



OUR Case of interest – ^{16}C



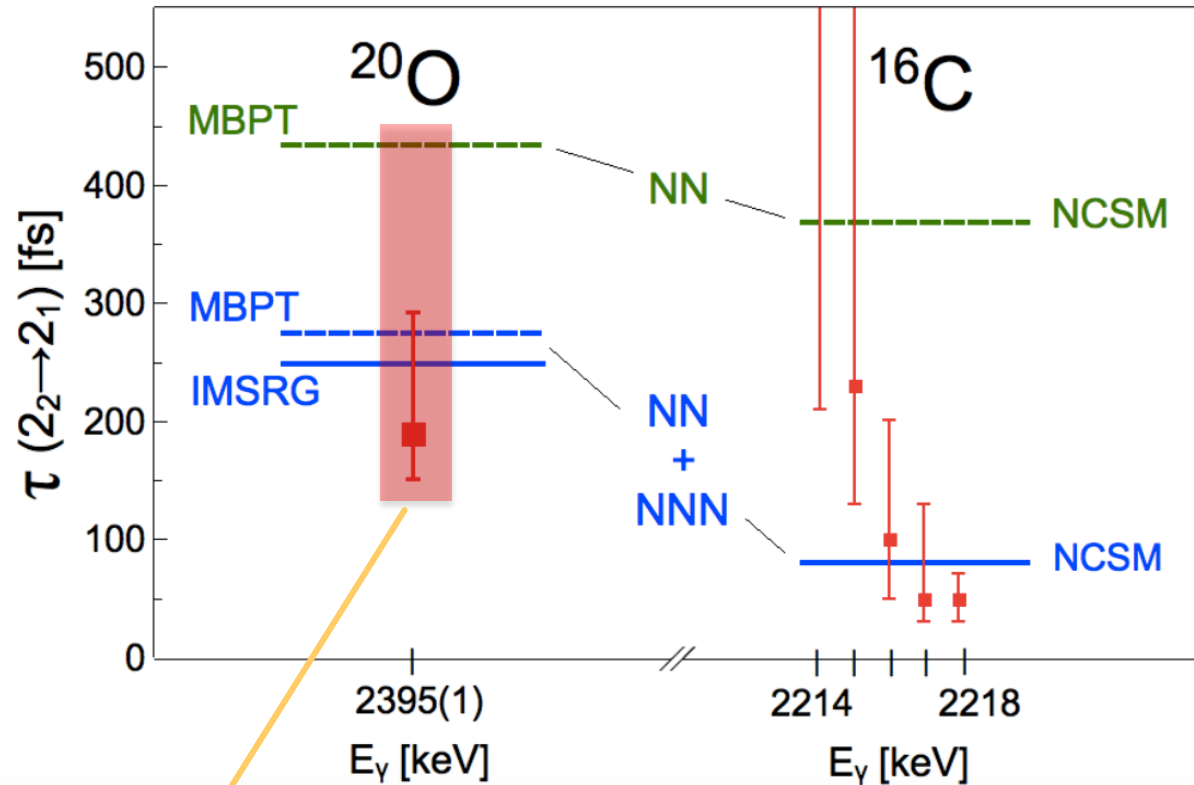
^{18}O (7 MeV/A) + ^{181}Ta target (6 mg/cm 2)



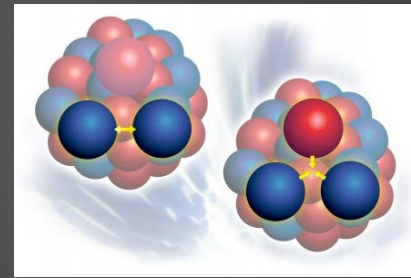
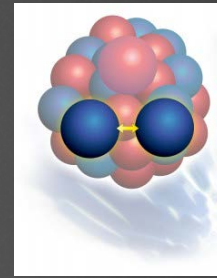
E_γ

ab initio predictions
3 body force $\tau = 80$ fs
2 body force $\tau = 360$ fs

Summary: Theory vs. exp. results comparison



**NO sensitivity would be obtained
with conventional HPGe detectors**



Give For ^{16}C most precise measurement
es $E = 2217(2)$ keV,
which do not allow
to determine exact lifetime value (for now).

Conclusions

- ▶ Italian-Polish-French collaboration allows to perform successful experiment at GANIL with combined **AGATA+VAMOS+PARIS** setup.
- ▶ (Re)-measurement of ^{19}O lifetimes – confirmation that DSAM method for AGATA and simulations works well.
- ▶ We measured lifetime of **second 2^+ in ^{20}O** : 150 fs ($2_2 \rightarrow 2_1$ decay partial $\tau = 190$ fs), which is consistent with *ab initio* calculations, including three body interactions.
AGATA tracking is crucial to obtain the needed sensitivity
- ▶ Extracted estimates of lifetime of **second 2^+ in ^{16}C** : it depends on non-shifted gamma-ray energy.
- ▶ Ongoing work on $^{18,19}\text{N}$ isotopes

The work **significantly broadens** the possibilities for nuclear structure high-precision measurements in hard-to-reach exotic systems

More comprehensive tests of *ab initio* theory approaches will be possible exploiting EM decays

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