

Probing the absolute neutrino mass scale with ^{163}Ho : the **HOLMES** project

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on behalf of the HOLMES collaboration

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^{163}Ho electron capture

$$^{163}\text{Ho} + e^- \rightarrow \text{Dy}^* + \nu_e \quad \frac{d\lambda_{EC}}{dE_c} = \frac{G_\beta^2}{4\pi^2} (Q - E_c) \sqrt{(Q - E_c)^2 - m_\nu^2} \times \sum n_i C_i \beta_i^2 B_i \frac{\Gamma_i}{2\pi} \frac{1}{(E_c - E_i)^2 + \Gamma_i^2/4}$$

$Q \sim 2.8 \text{ keV}$, capture only from shell $\geq M1$

De Rujula & Lusignoli, Phys. Lett. B 118 (1982) 429

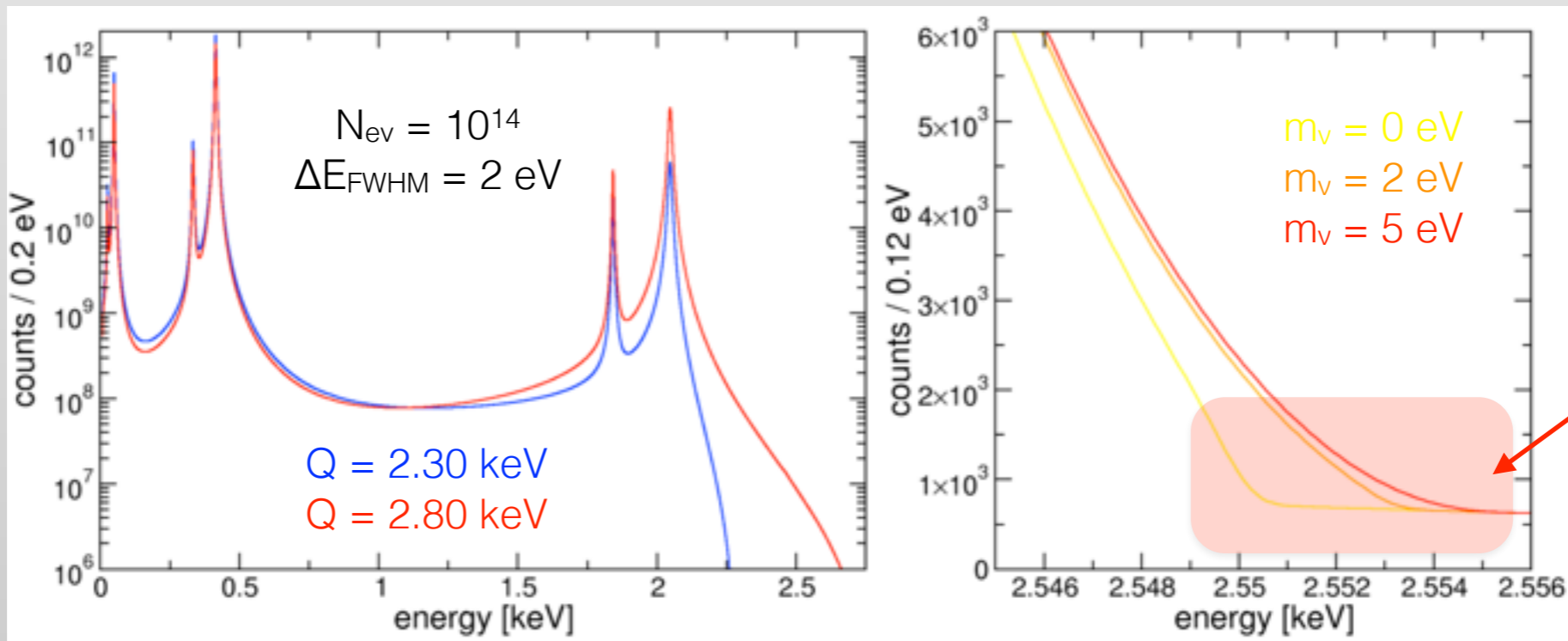
same factor as β decay

(total de-excitation energy E_c instead of E_e)

Breit-Wigner shapes

- calorimetric measurement of Dy^* de-excitation
- “good” event rate and ν mass sensitivity depends on Q-value and capture peak position (roughly $\sim 1/(Q - E_{M1})^3$)
- $\tau_{1/2} \sim 4570$ years \rightarrow few active nuclei needed

simulated spectra with different Q-value



^{163}Ho electron capture

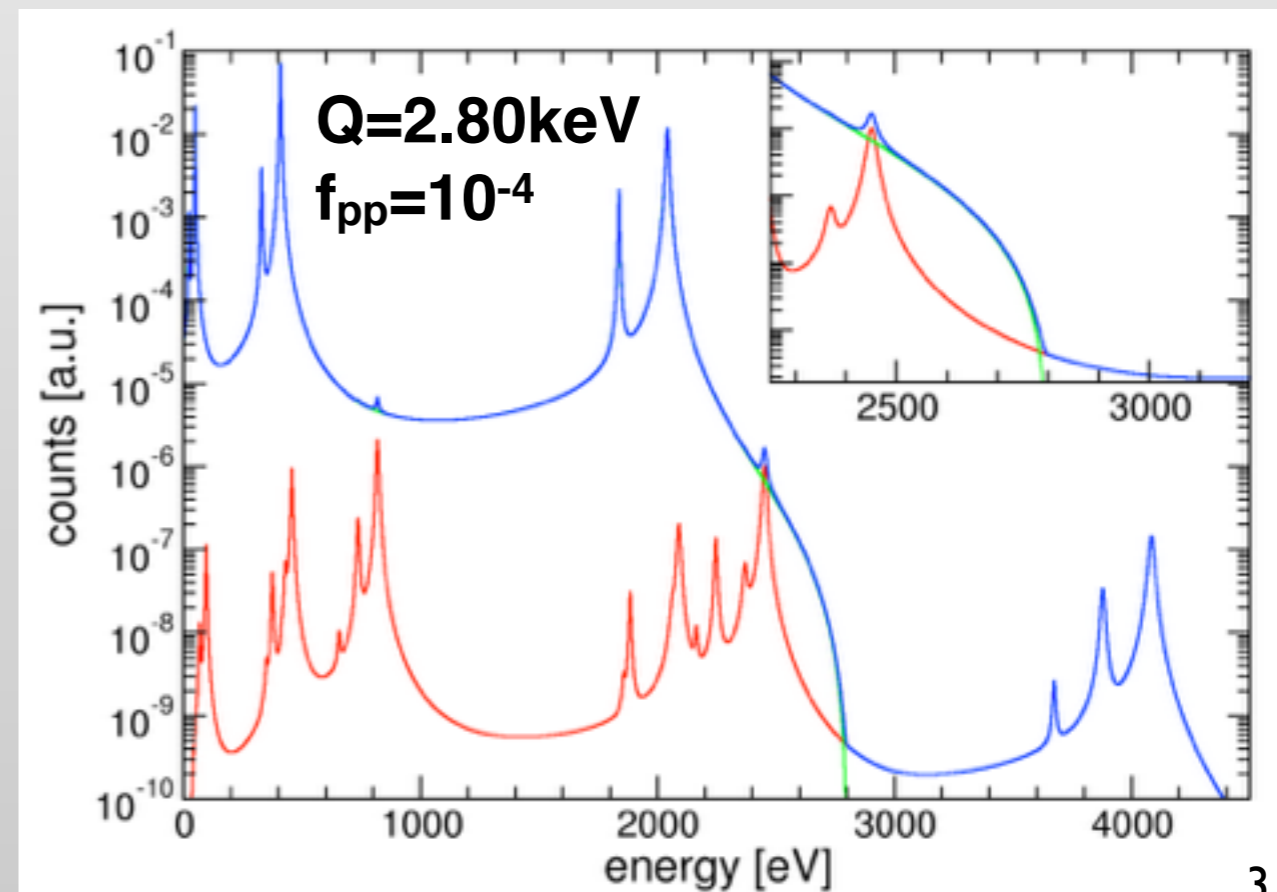
Complex pile-up spectrum: end-point is dominated by $((Q-E_c)^2 - m_\nu^2)^{1/2}$ but expected distortions due to pile-up:

$$N_{pp}(E) = f_{pp} N_{EC}(E) \otimes N_{EC}(E)$$

Pile-up occurs when multiple events arrive within the resolving time of the detector. In a first approximation, fraction of unresolved pile up is given by **$f_{pp} = \tau \times A_{EC}$** .

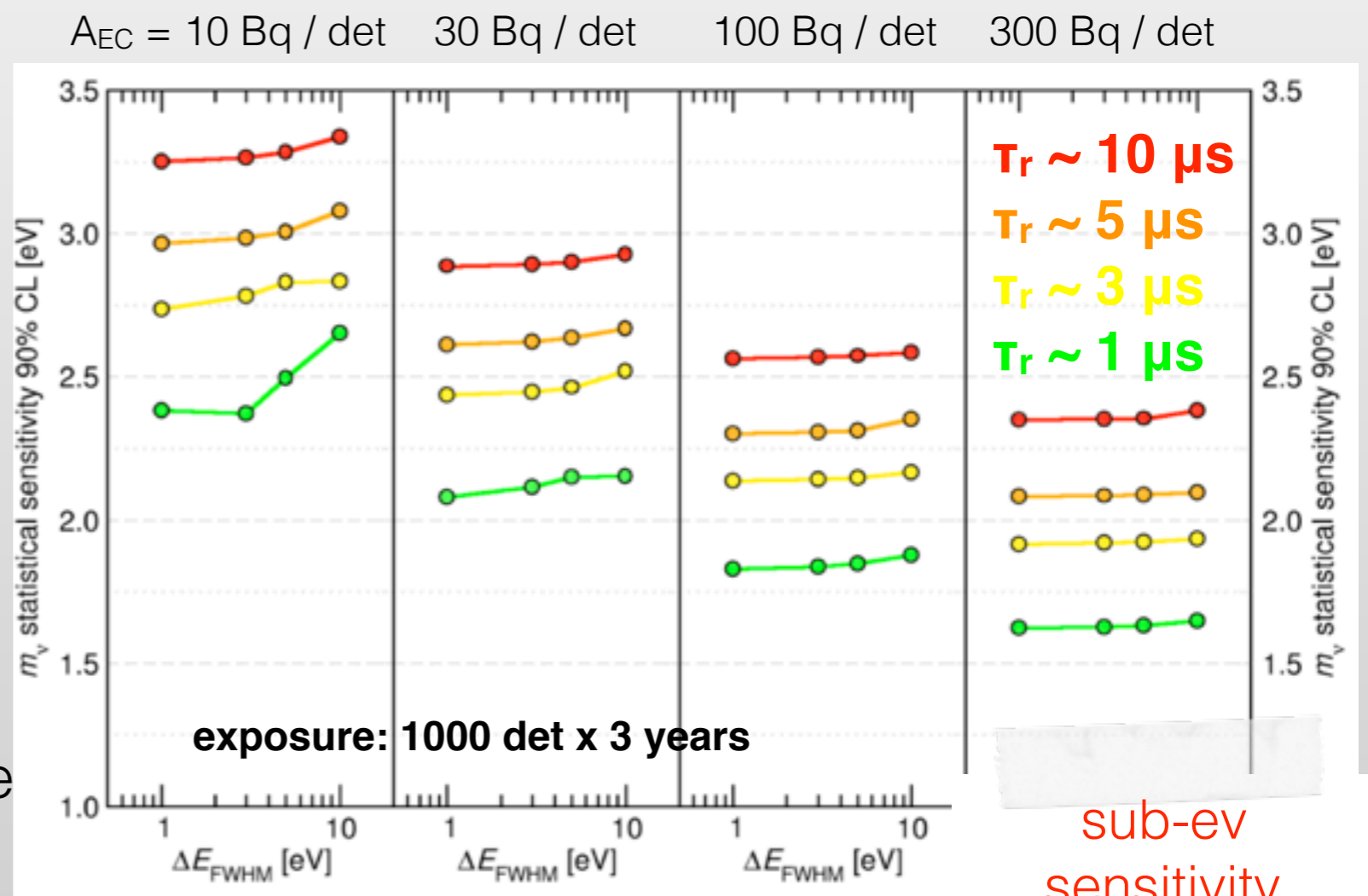
In order to reduce pile-up:

- **trade-off** between **activity** and **statistic**;
- detector with **fast signal rise time** τ_r ;
- pile-up **resolving algorithm**.



The **H_VLMES** project in a nutshell

- **Direct neutrino mass measurement** with statistical sensitivity around 1 eV
- **Usage of Transition Edge Sensor (TES) based micro-calorimeters** with ^{163}Ho implanted Au absorber:
 - 6.5×10^{13} nuclei / det
 - $A_{\text{EC}} \sim 300$ Bq / det
 - $\Delta E \sim 1\text{eV}$, $\tau \sim 1\mu\text{s}$
- **1000 channels array:**
 - 6.5×10^{16} total nuclei
 - $O(10^{13})$ events / year
- **Should prove the technique potential and scalability** by:
 - assessing EC spectral shape
 - assessing systematic errors



^{163}Ho production

^{163}Ho does not exist in nature: it is produced from ^{162}Er neutron activation at nuclear reactor:

- $^{162}\text{Er} (n,\gamma) ^{163}\text{Er}$, $\sigma_{\text{therm}} \sim 20 \text{ b}$
- $^{163}\text{Er} + e^- \rightarrow ^{163}\text{Ho} + \nu_e$ ($\tau_{1/2} \sim 75 \text{ m}$)
- high yield
 - $\sim 3 \times 10^{12}$ ^{163}Ho nuclei/mg ^{162}Er /h
- requires ^{162}Er enrichment and oxide chemical form (Er_2O_3)

Tm 163 1.81 h	Tm 164 5.1 m / 2.0 m	Tm 165 30.06 h	Tm 166 7.70 h	Tm 167 9.25 d	Tm 168 93.1 d
Er 162 0.139	Er 163 75 m	Er 164 1.601	Er 165 10.3 h	Er 166 33.503	Er 167 2.3 s / 22.869
Ho 161 6.7 s / 2.5 h	Ho 162 68 m / 15 m	Ho 163 1.1 / 4570 a	Ho 164 37 m / 29 m	Ho 165 100	Ho 166 1200 a / 26.80 h
Dy 160 2.329	Dy 161 18.889	Dy 162 25.475	Dy 163 24.896	Dy 164 28.260	Dy 165 1.3 m / 2.35 h
Tb 159	Tb 160	Tb 161	Tb 162	Tb 163	Tb 164



Collaboration with
ILL (Grenoble)
ITN (Lisboa)
Paul Scherrer Institute

But contaminations from other isotopic species. Main one:

- $^{165}\text{Ho} (n,\gamma) ^{166\text{m}}\text{Ho} (\beta, \tau_{1/2} \sim 1200 \text{ years})$
- from Ho contamination or $^{164}\text{Er} (n,\gamma)$
- **need high purification of sample:**
 - radiochemical separation
 - mass separation with magnetic dipole

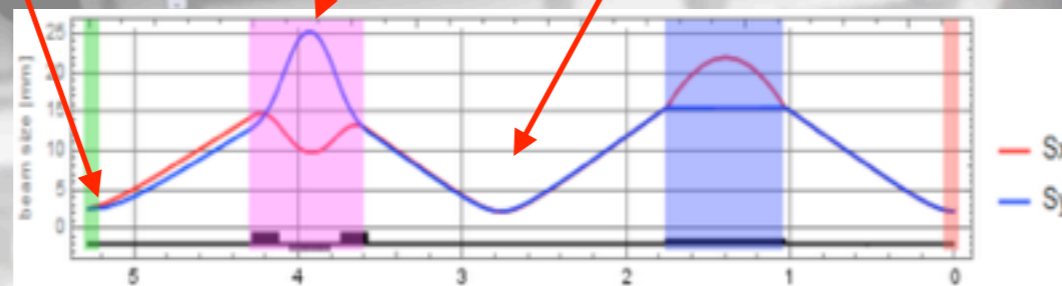
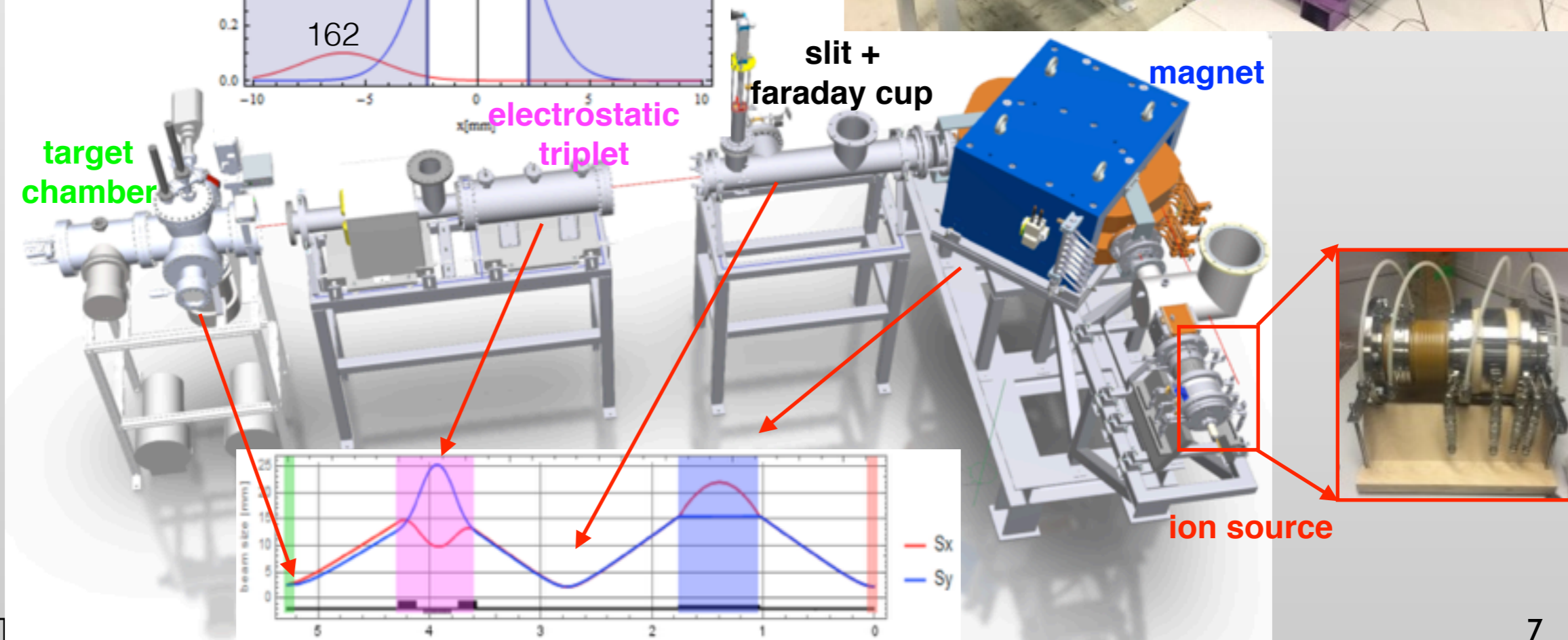
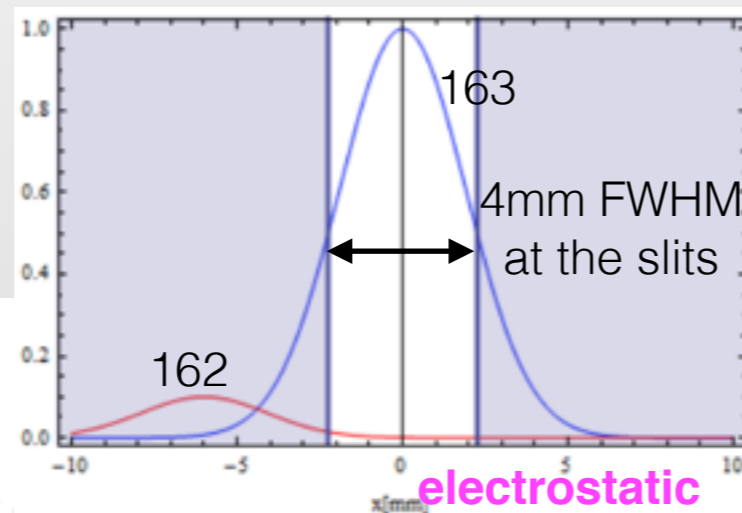
^{163}Ho purification

- Enriched Er_2O_3 samples irradiated at ILL (Grenoble) and post-processed at PSI
 - 25mg, 55 days irradiation $\rightarrow A(^{163}\text{Ho}) \sim 5 \text{ MBq}$
 - 150mg, 50 days irradiation $\rightarrow A(^{163}\text{Ho}) \sim 38 \text{ MBq}$
- **Ho radiochemical separation is performed via ion-exchange resins in hot-cell at PSI**
 - efficiency $> 80\%$ (provisional estimation)
- 540mg irradiated for 50 days at ILL in early 2017 are ready for purification
 - expected overall activity: $\sim 130 \text{ MBq}$ (enough for R&D and half pixels)



^{163}Ho mass separation and implantation

- **Implanter machine with 30/50 kV acceleration (10-50nm implantation depth) and magnetic dipole:**
 - $^{163}\text{Ho}/^{166\text{m}}\text{Ho}$ separation better than 10^5
 - first components delivered in Genova at beg. 2017, now under test (magnet, source, vacuum OK - full system test coming soon)
 - upgrade with focusing triplet and magnetic xy scanning expected for late 2017 / beg. 2018

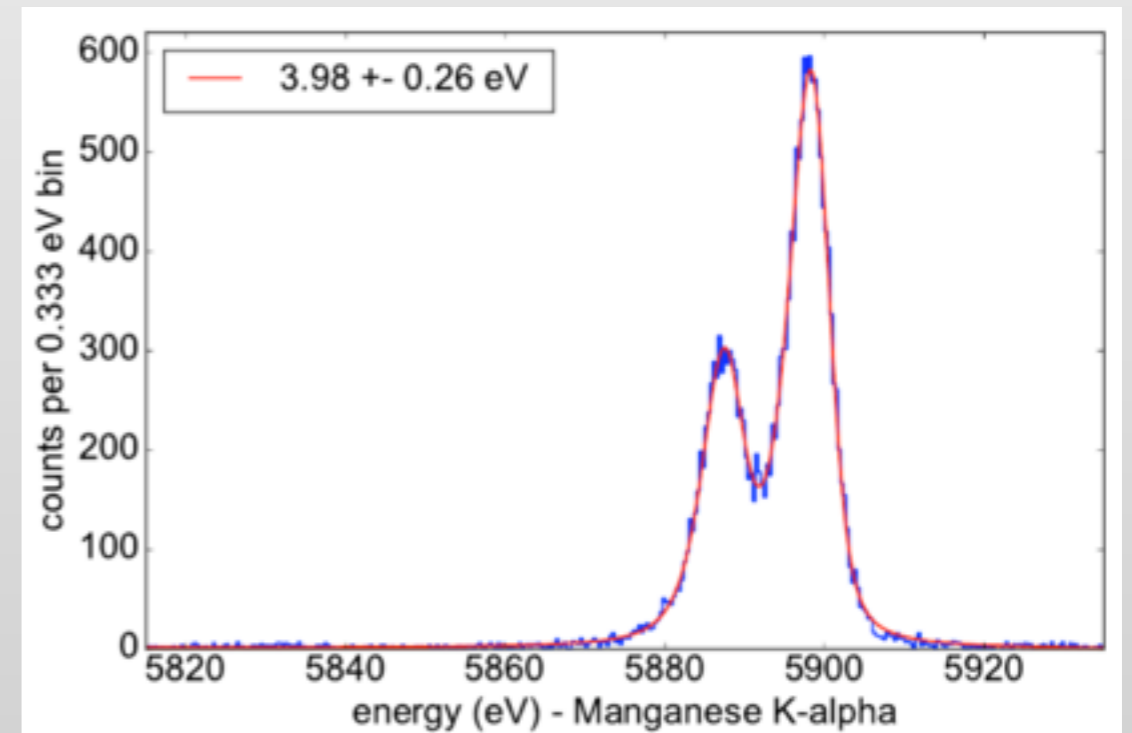
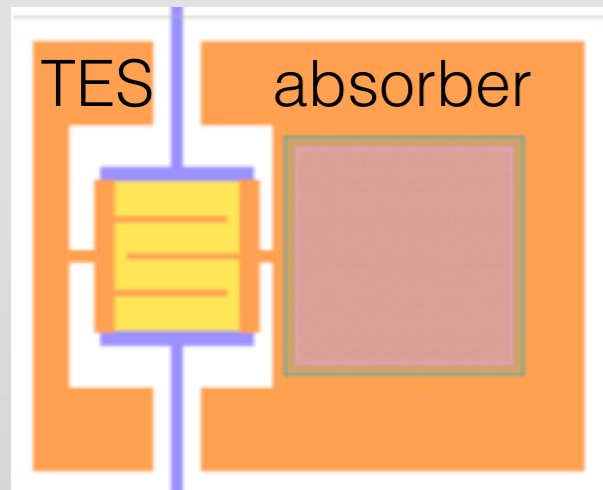


Detector design and test

- TES design, production and preliminary test is done @NIST
- **2 μm gold thickness for full absorption** of electrons and photons
- **“side car” configuration** to avoid TES proximization and allow G engineering for τ control
- Design optimized to obtain **best compromise between resolution and time response.**

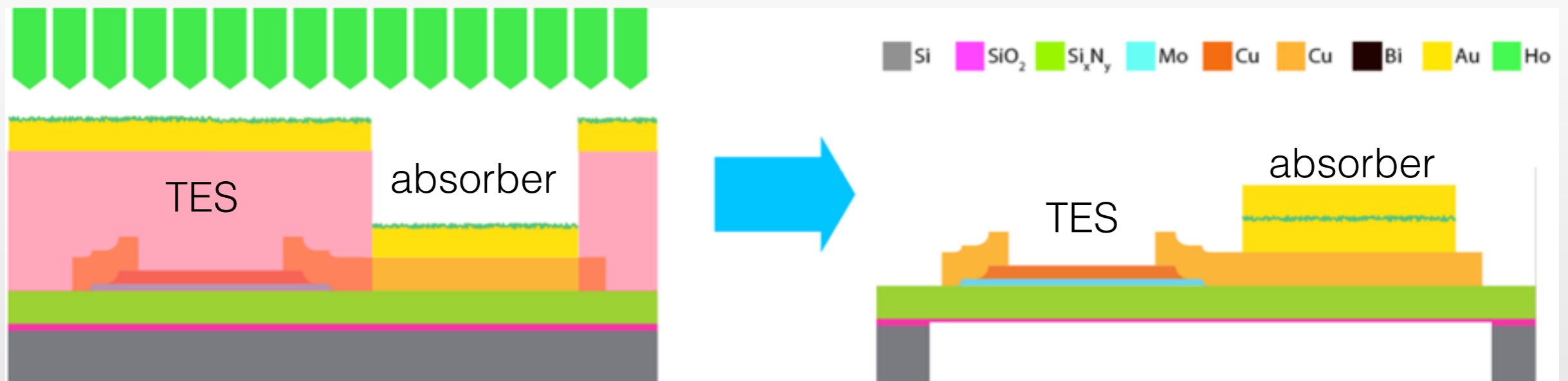
Target (@3keV):

- $\Delta E_{\text{FWHM}} \sim 1\text{eV}$
- $\tau_{\text{rise}} \sim 1\ \mu\text{s}$
- $\tau_{\text{decay}} \sim 100\ \mu\text{s}$
- RF-SQUID readout with microwave MUX



- $\Delta E_{\text{FWHM}} \sim 4\text{eV @}6\text{keV}$ ($\sim 3\text{eV @ } Q_{\text{EC}}$)
- $\tau_{\text{rise}} \sim 3\ \mu\text{s}$
- $\tau_{\text{decay}} \sim 130\ \mu\text{s}$

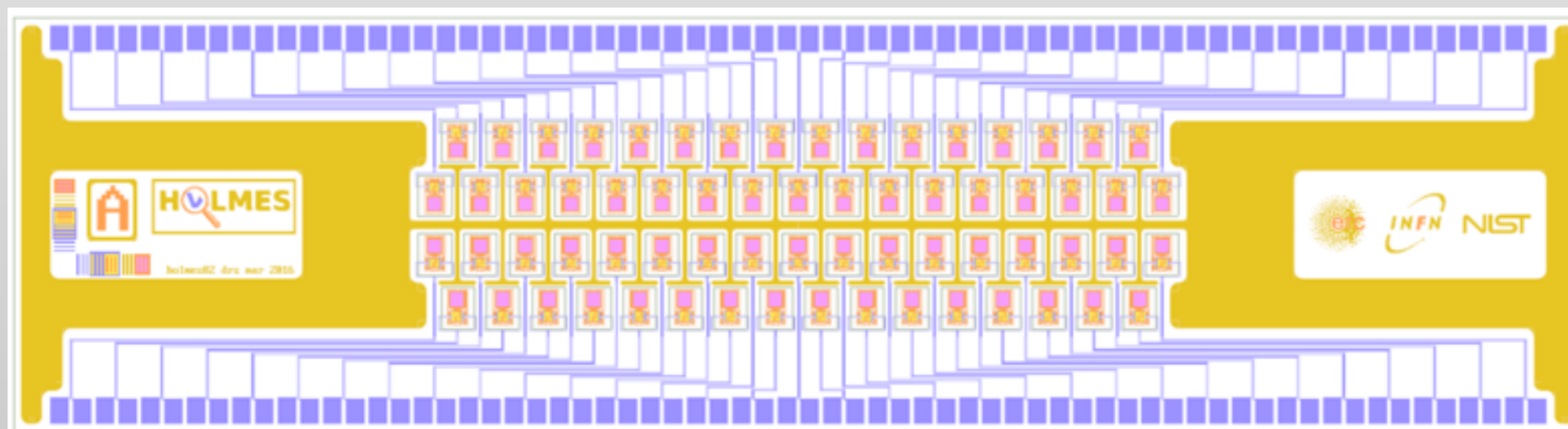
Detector fabrication



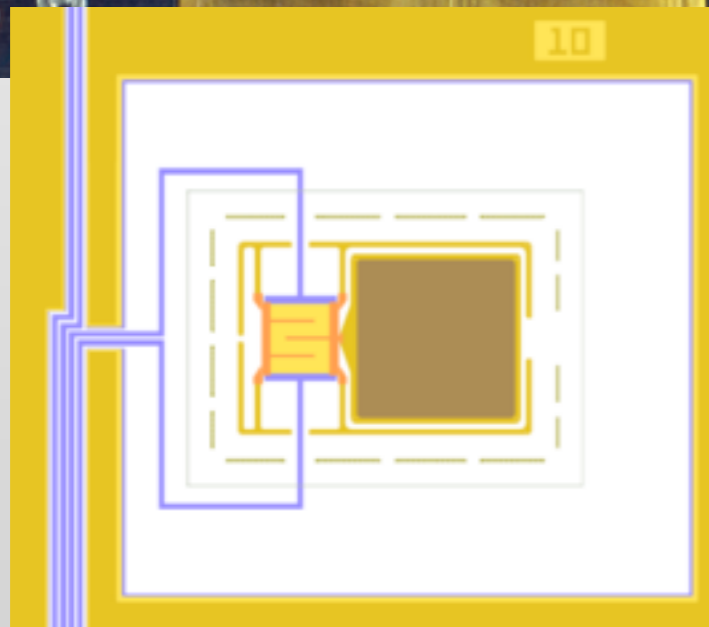
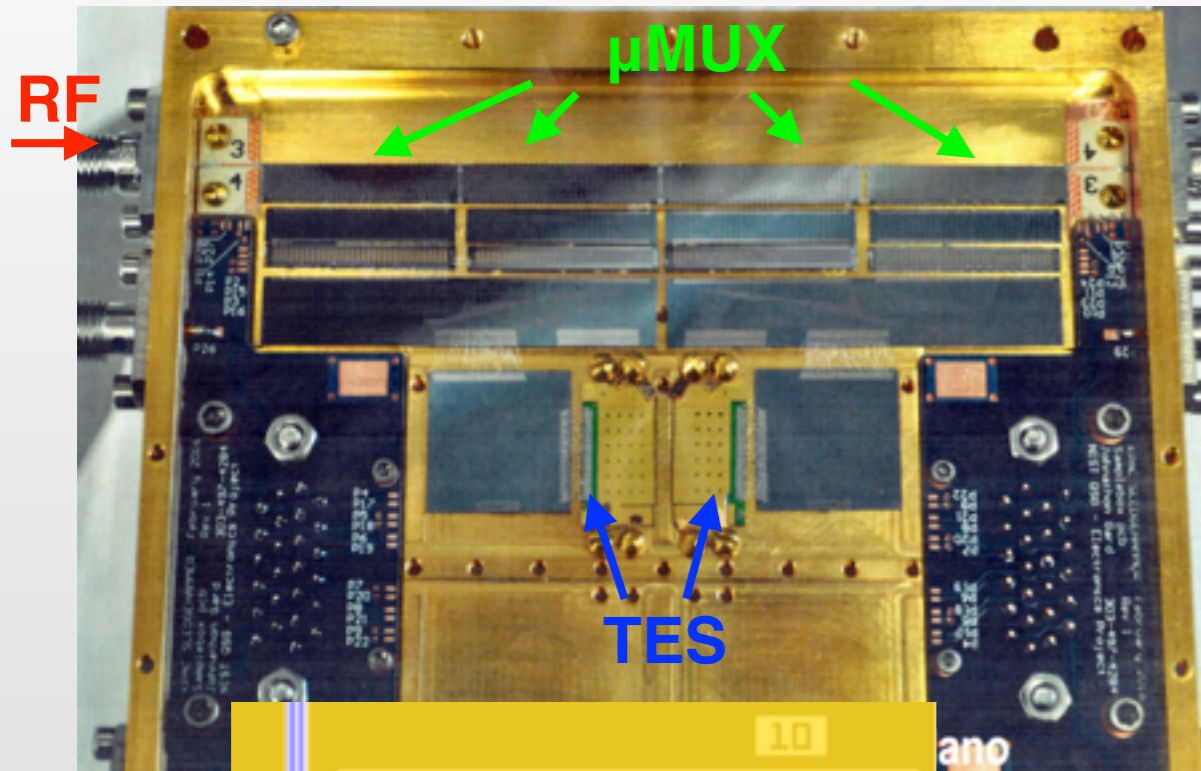
Detector fabrication is done with a **multi-step procedure**:

- 1) TES array is **produced @NIST**
- 2) ¹⁶³Ho is **implanted @Genova**
- 3) 1 μm **Au final layer is deposited over Ho implantation** (“complete” the absorber)
- 4) final fabrication processes definition is on-going

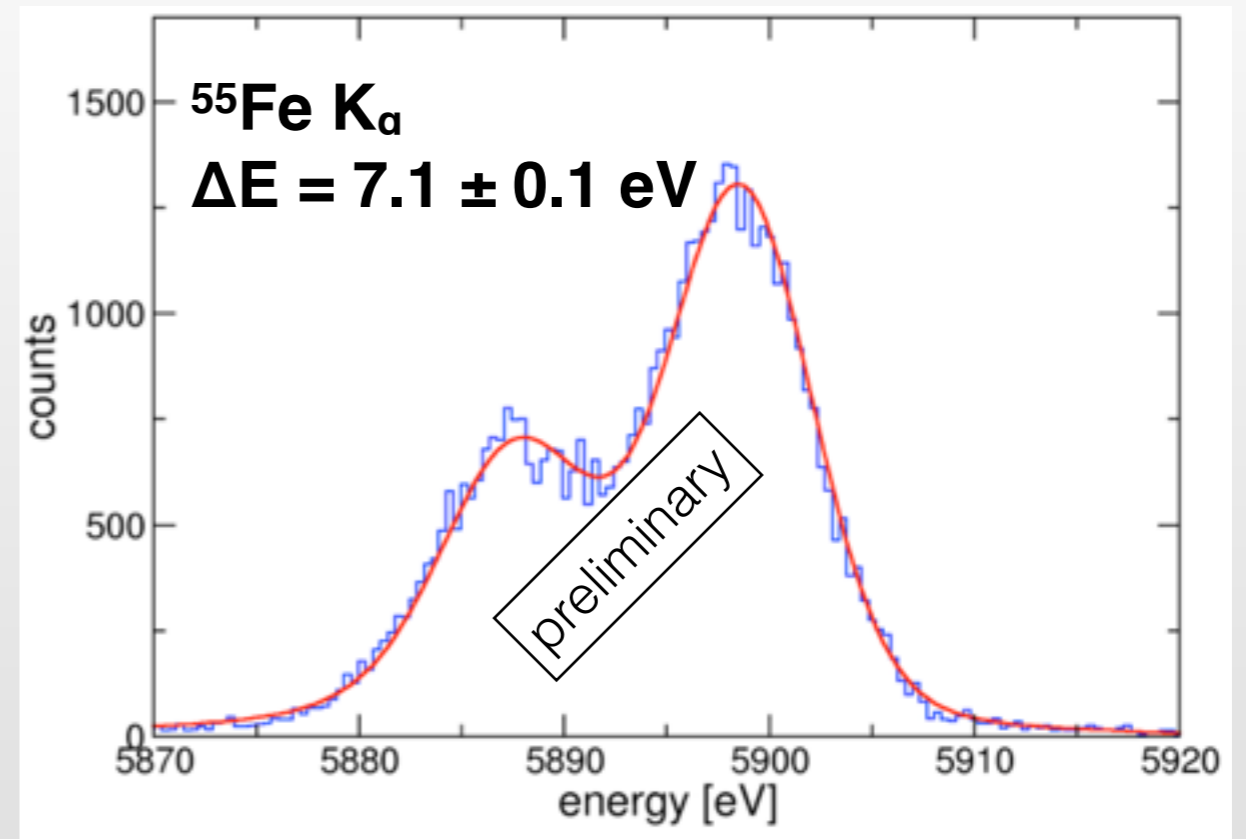
4 x 16 linear array for implantation optimisation



Pixel testing with HOLMES DDAQ



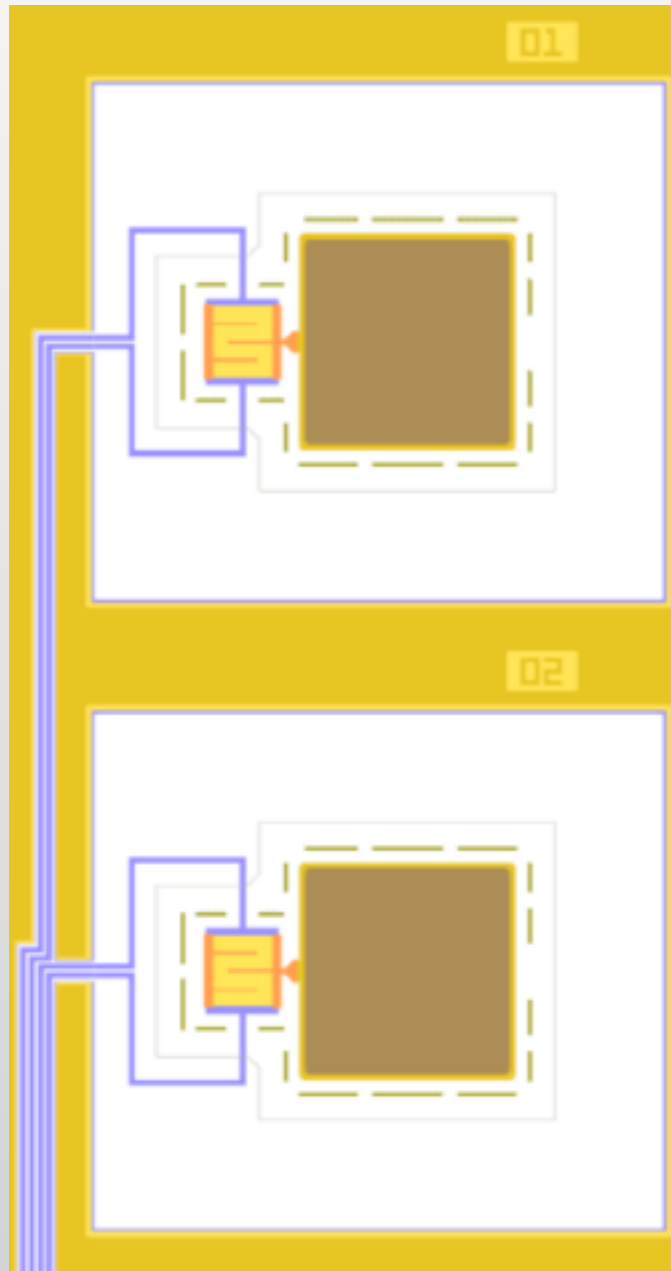
280 x 280 μm^2 absorber
C = 0.75 pJ/K
G = 330 pW/K



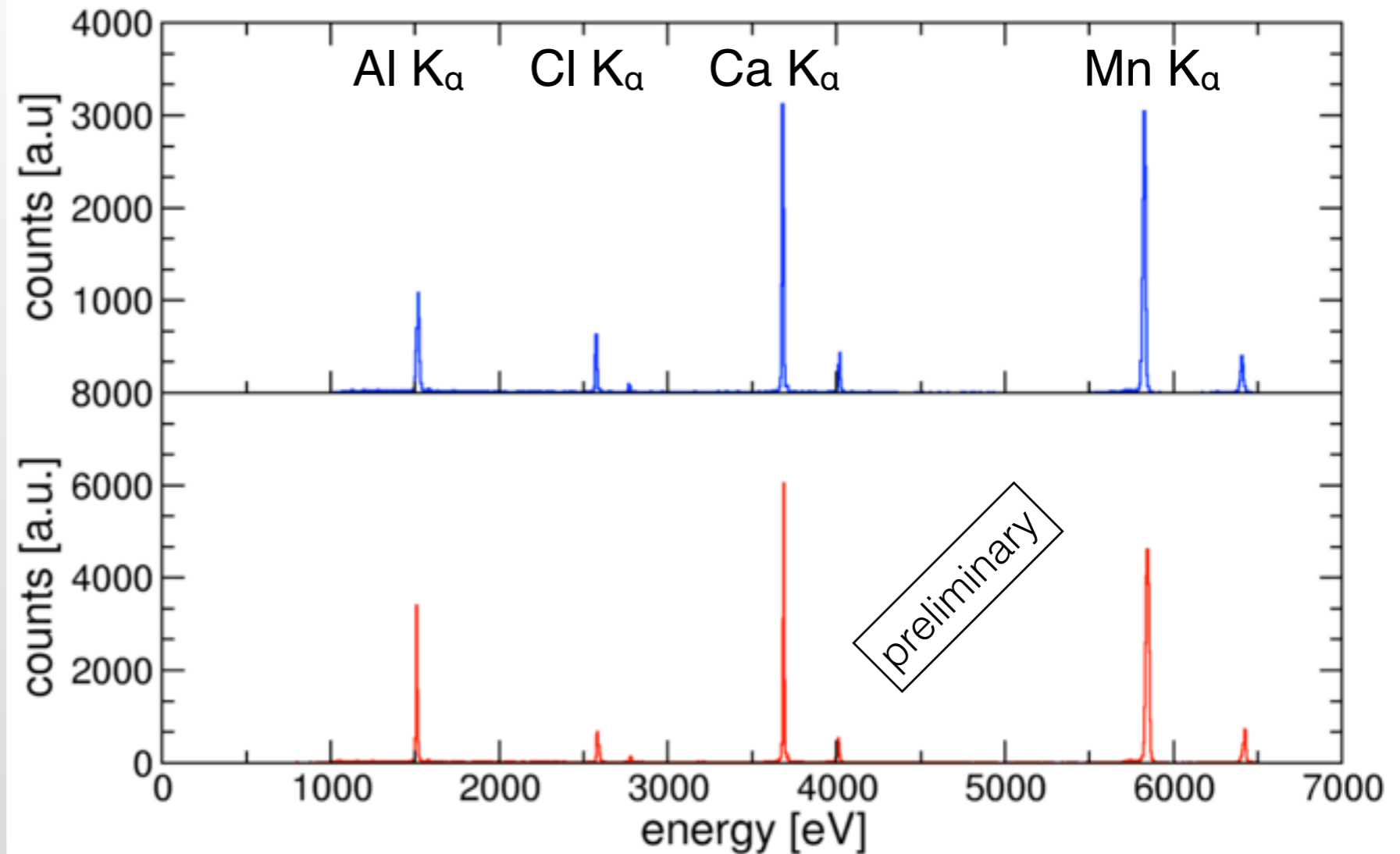
$f_{\text{samp}} = 500 \text{ kS/s}$
 $T_{\text{rise}} = 6.5 \mu\text{s}$
 $T_{\text{decay}} = 67 \mu\text{s}$

Pixel testing with HOLMES DDAQ

HOLMES-like pixels
without collimator

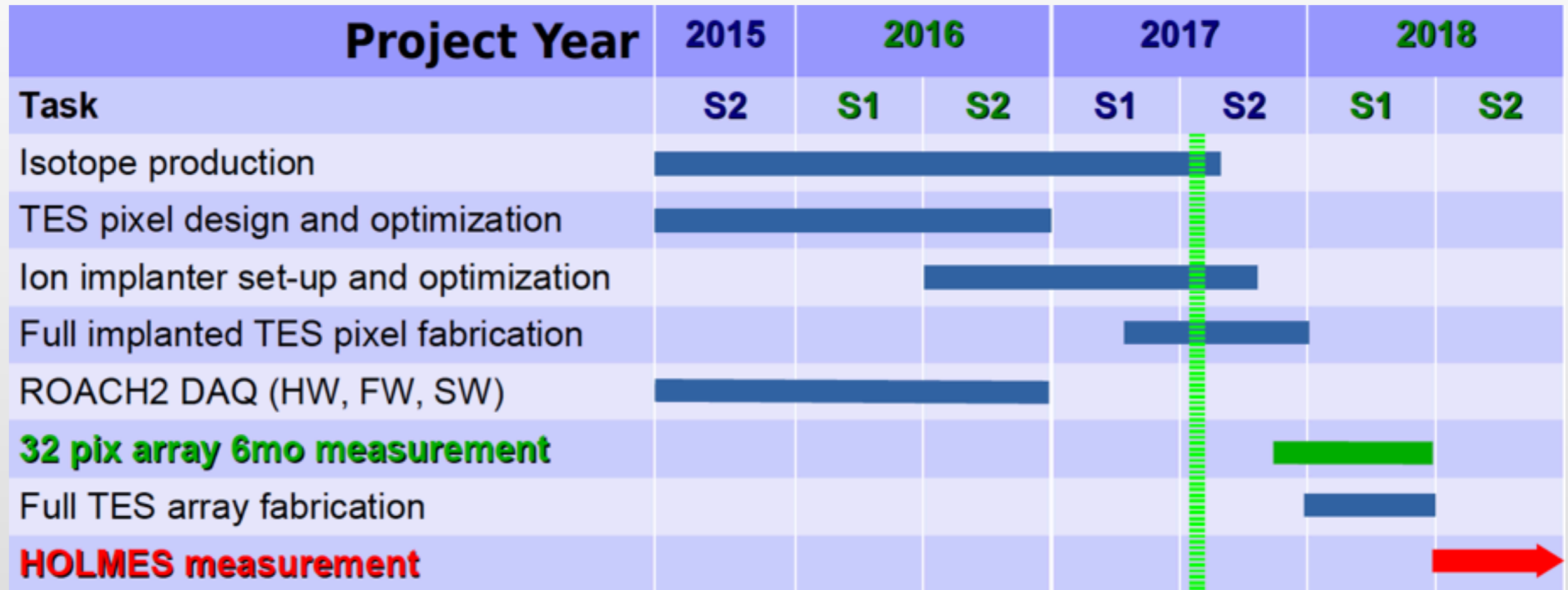


380 x 380 μm^2 absorber
C = 0.9 pJ/K
G = 480 pW/K



$\Delta E \sim 9 \text{ eV @ } 2.6 \text{ keV}$
 $T_{\text{rise}} = 20 \mu\text{s}$
 $T_{\text{decay}} = 140 \mu\text{s}$

Current status and schedule



HOLMES project status:

- TES array and DAQ ready
- Ion implanter setting up is in progress
- First ^{163}Ho implantation coming shortly
- Spectrum measurements will begin late in 2017
- **32 pixels for 1 month $\rightarrow m_\nu$ sensitivity ~ 10 eV**

Back up slides

The **HOLMES** collaboration



ERC Advanced Grant 2013
Research proposal [Part B1]

Principal Investigator (PI): *Prof. Stefano Ragazzi*
PI's Host Institution for the project: *Istituto Nazionale di Fisica Nucleare*



**The Electron Capture Decay of ^{163}Ho to Measure the
Electron Neutrino Mass with sub-eV sensitivity**

HOLMES

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A.Nucciotti
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G.Pessina
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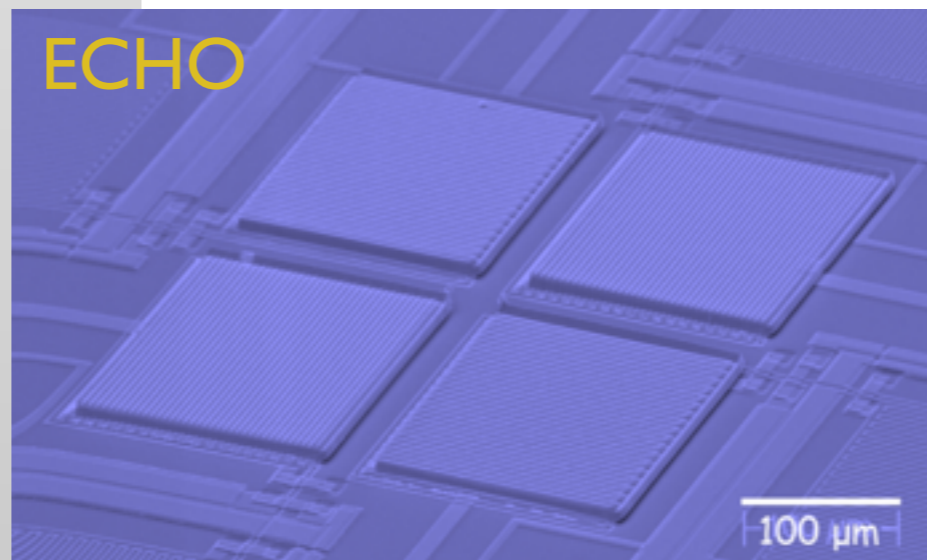
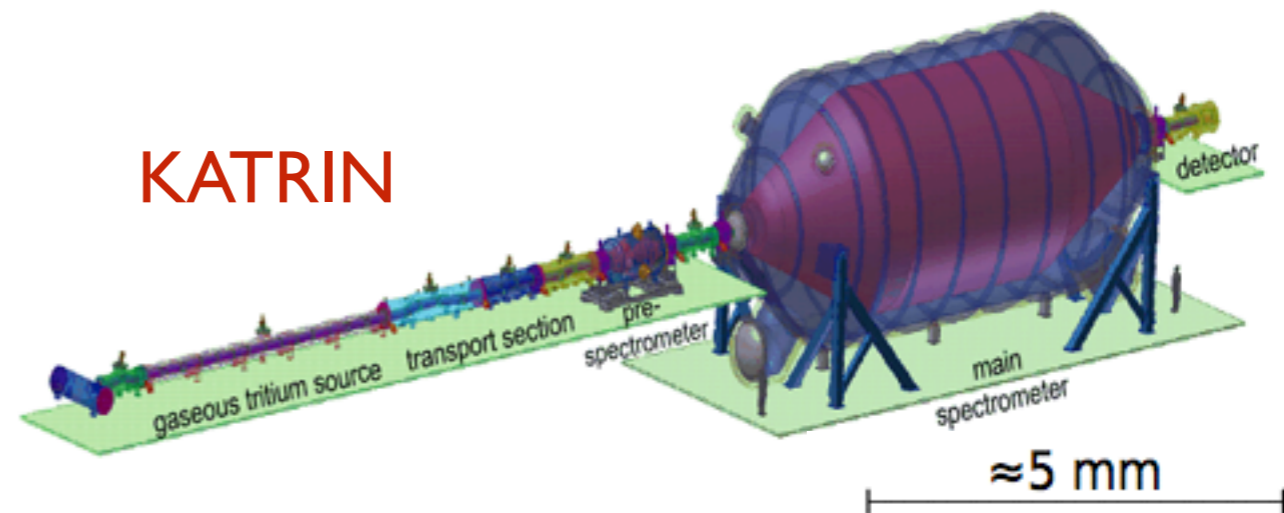
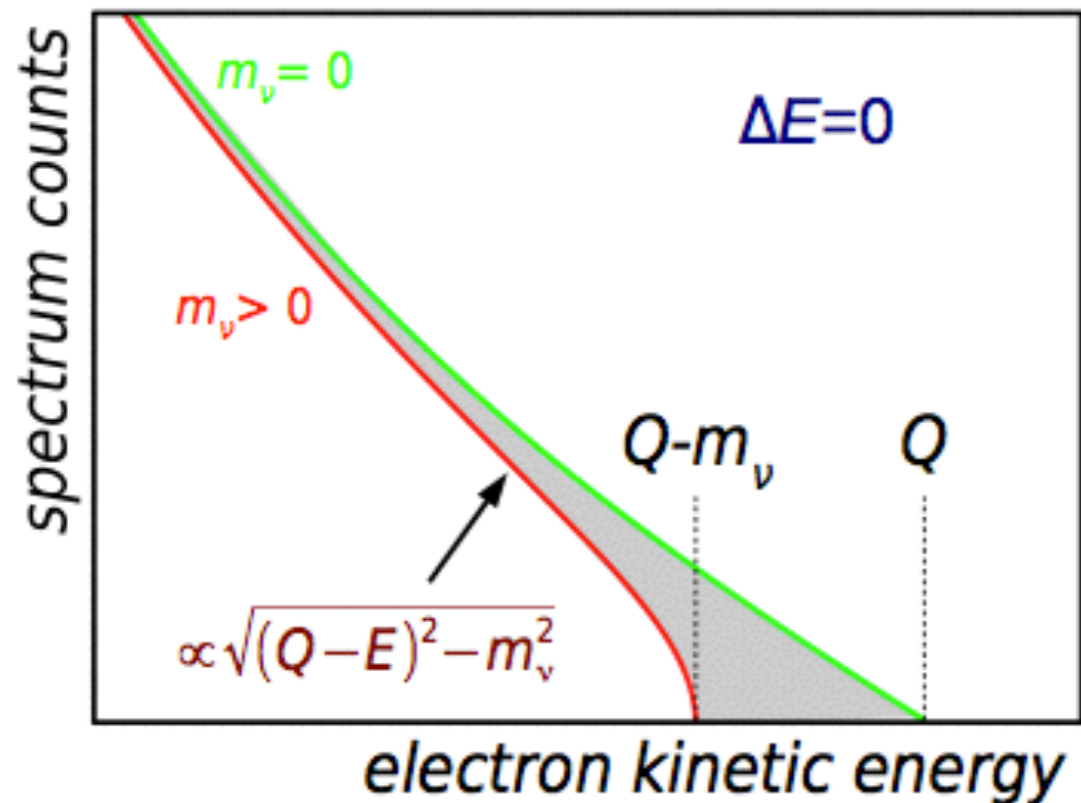
Direct ν mass measurement

Kinematics of weak decay with ν emission:

- low Q nuclear β decays (${}^3\text{H}$, ${}^{187}\text{Re}$, ${}^{163}\text{Ho}$...)
- model independent: only E, p conservation
- **ν mass appears as a distortion in the Kurie plot**

2 different approaches:

- **spectrometry**: source placed outside the detector (**KATRIN** approach)
- **calorimetry**: source embedded inside the detector (**ECHO**, **MARE**, HOLMES approach) \Rightarrow low T μ -calorimeters



Spectrometry vs calorimetry

General requirements for a ν mass experiment:

- High statistics near the end point
 - low Q-value (stat $\sim 1/Q^3$)
 - high activity/efficiency of the source
- Energy reso order $\sim eV$ or below (comparable with m_ν)
- S/N ratio
- small systematic effects

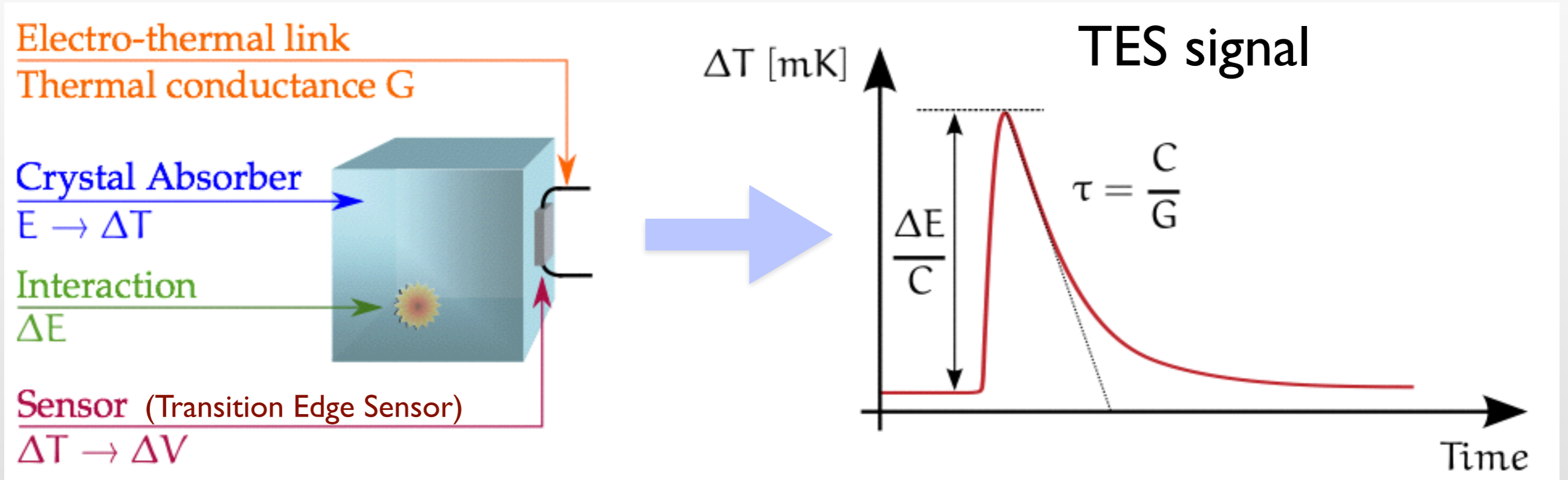
Spectroscopy: source $\not\subset$ detector

- high statistics
- high energy resolution (below eV)
- systematics due to the source (energy loss)
- systematics due to decay to excited states
- background

Calorimetry: source \subset detector

- no backscattering
- no energy loss in source
- no solid state excitation
- no atomic/molecular final state effects
- good energy resolution ($\sim eV$)
- limited statistics
- systematics due to pile-up
- background

Low T calorimetry in a nutshell



- Complete energy thermalization (ionization, excitation \rightarrow heat \rightarrow calorimetry)
- $\Delta T_{\max} = E/C$, C is the total thermal capacity
 - absorber with low thermal capacity
 - for superconductors below T_C and dielectric: $C \sim (T/\theta_D)^3$ (Debye law)
 - very low T is needed (10 ÷ 100 mK)
- $\Delta E_{\text{rms}} = (k_b T^2 C)^{1/2}$ due to statistical fluctuations of internal energy
- $\Delta T(t) = E/C e^{-t/\tau}$, $\tau = C/G$ and G is the thermal conductance

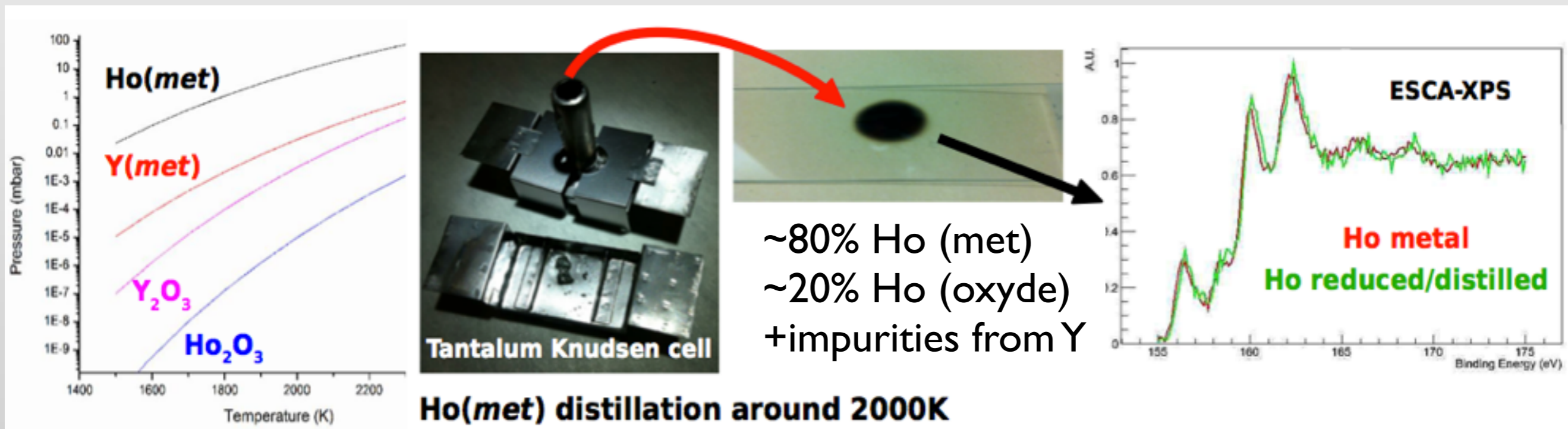
Ho production and purification

^{163}Ho separation from Dy, Er and others...

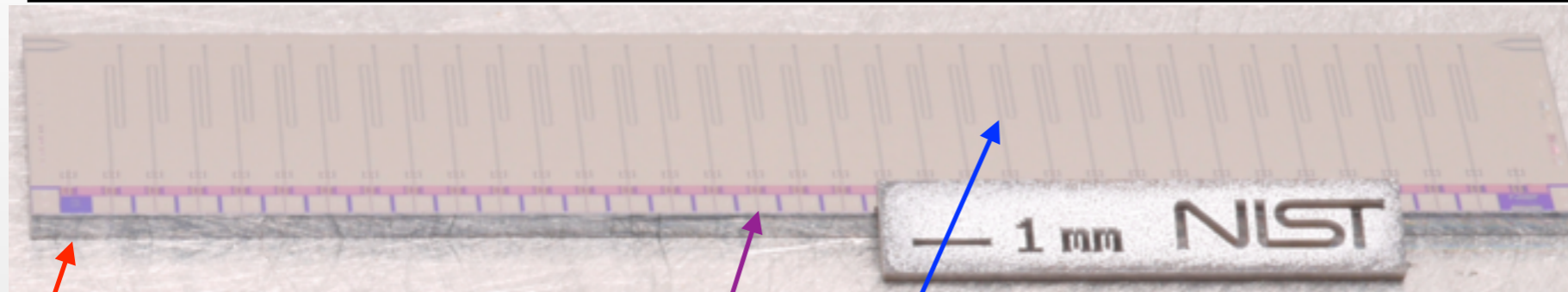
- radiochemistry (before/after activation process)
- magnetic mass separation

Ho_2O_3 thermoreduction in Knudsen cell provides a metallic sample for the implantation:

- $\text{Ho}_2\text{O}_3 + \text{Y}(\text{met}) \rightarrow \text{Ho}(\text{met}) + \text{Y}_2\text{O}_3 @2000\text{K}$
- First test already performed in Genova

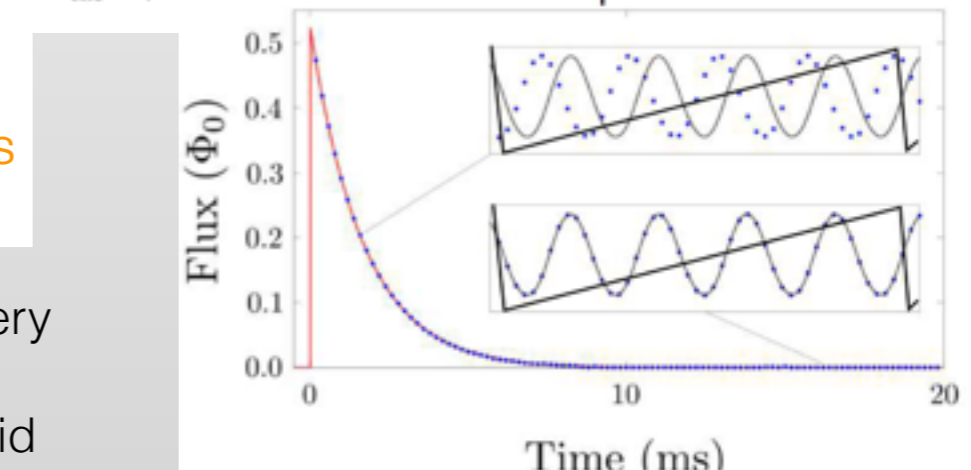
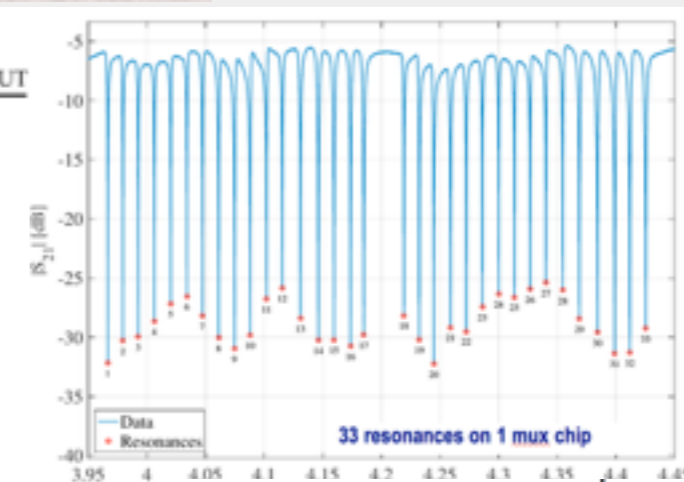
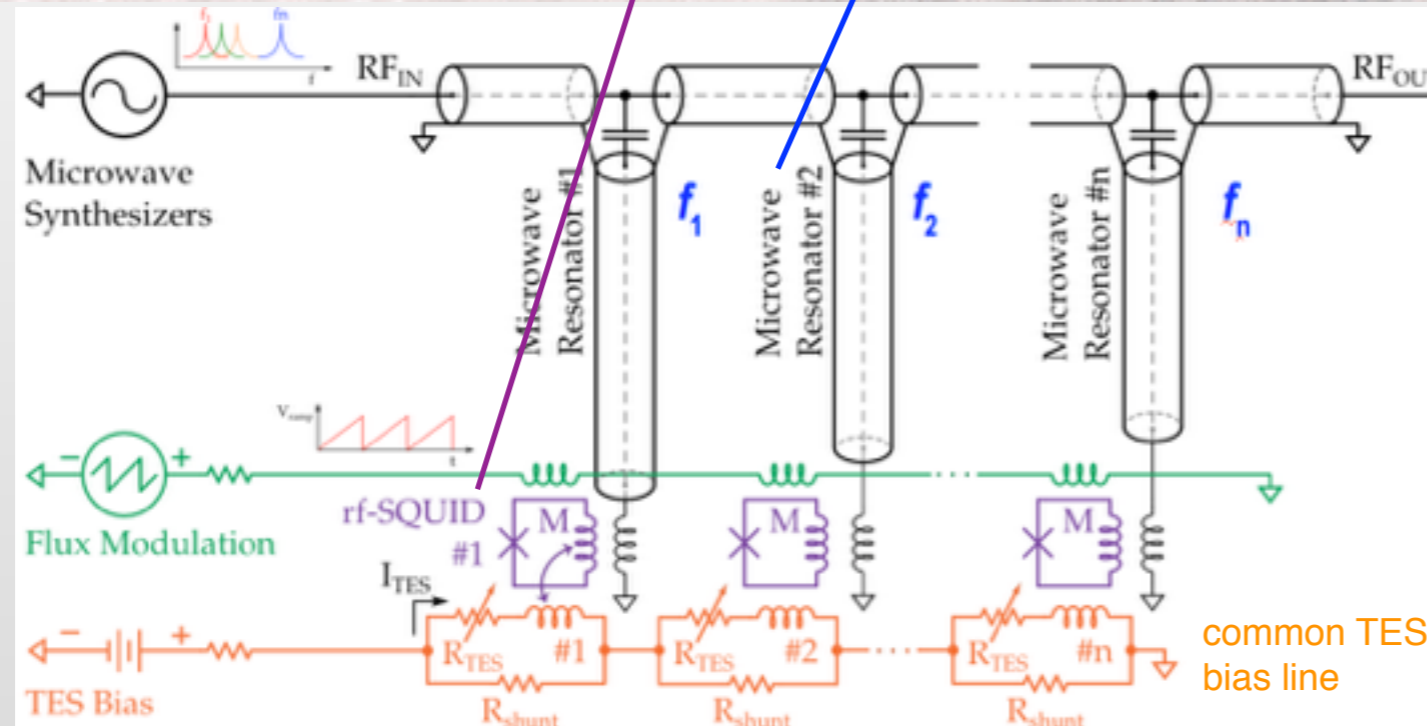


Array readout: rf-SQUID μ wave mix



$\lambda/4$ resonators coupled to common feedline tuned at different frequencies for multiplexing

common SQUID modulation line



To linearize the SQUID response, a voltage ramp is constantly applied to every SQUID trough a common line.

The signal is reconstructed from the phase shift of the SQUID oscillation (solid line), with respect to a reference sine function (dotted line).

The ramp frequency is the effective pulse sampling

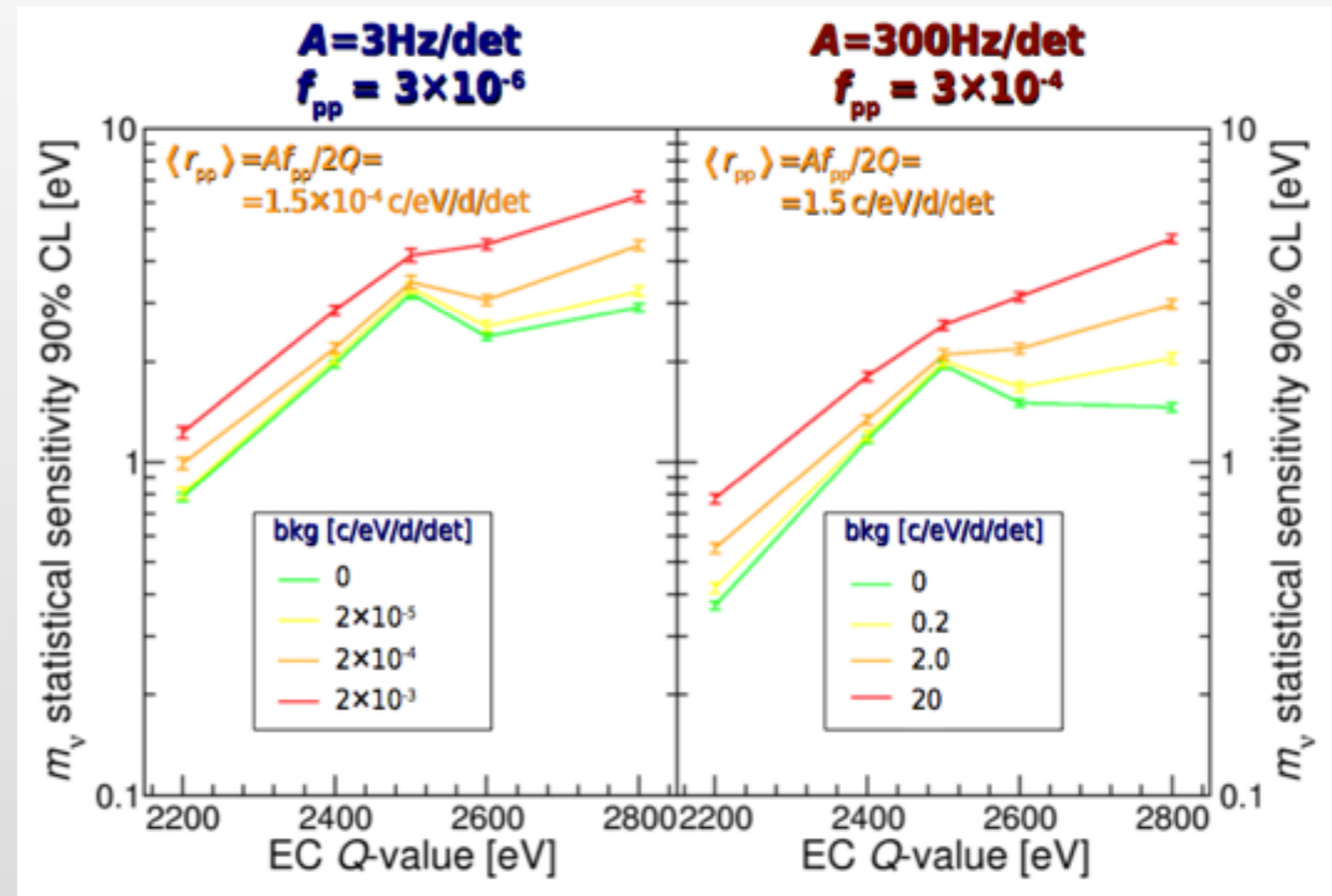
Each rf-SQUID is coupled to a GHz range resonator

- resonance bandwidth has to match the SQUID oscillation frequency i.e. 2 MHz

- resonance spacing has to be tuned to maximise multiplexing factor avoiding crosstalk i.e. 14 MHz

Source of background

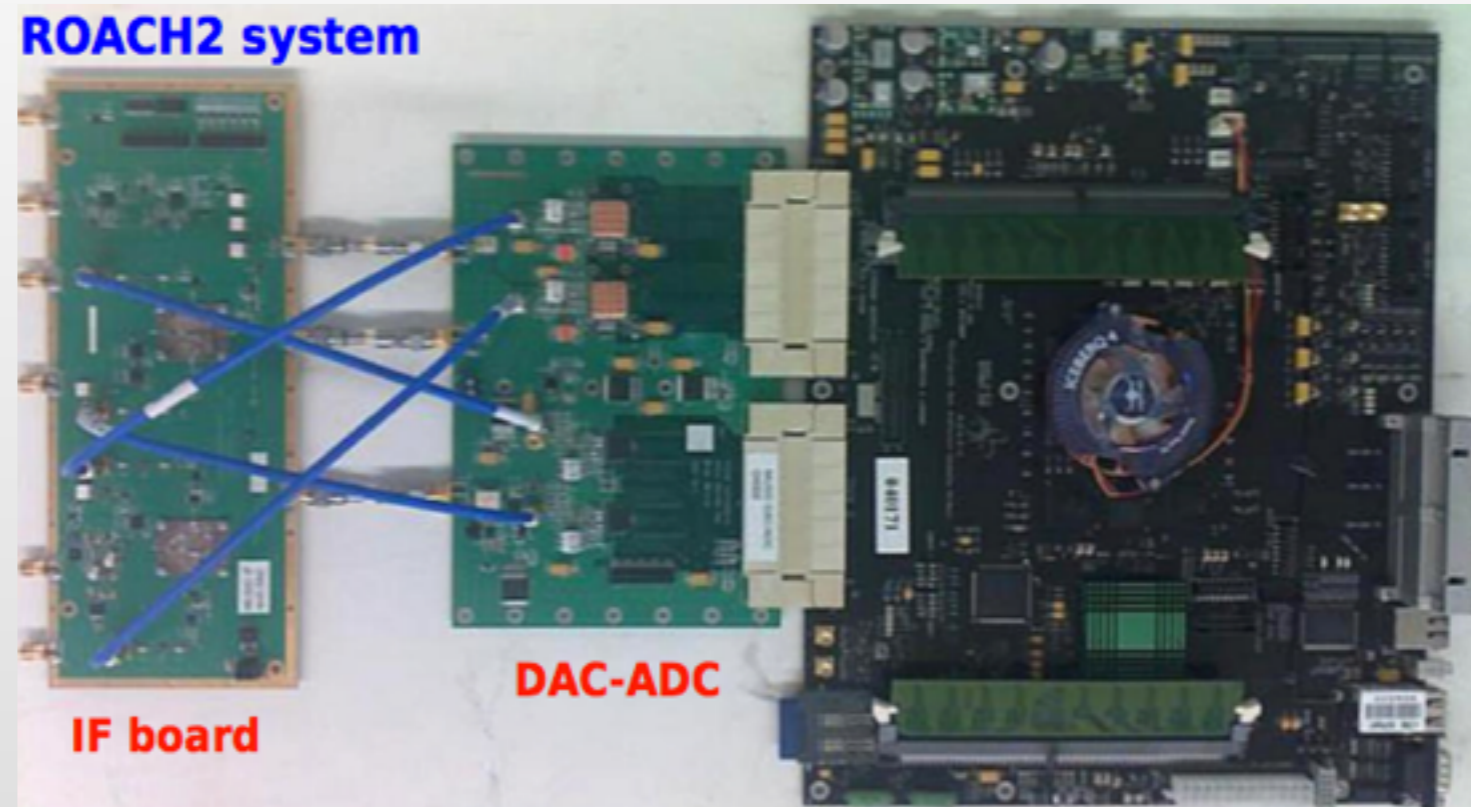
- Environmental γ radiation
 - Compton interactions, photoelectric interactions with p.e. escape
 - Fluorescent X-rays and X-ray escape line
- Cosmic rays
 - GEANT4 simulation for CR at sea level (only muons)
 - Au pixel $200 \times 200 \times 2 \mu\text{m}^3 \rightarrow$ bkg $\sim 5 \times 10^{-5}$ c/eV/day/det (0 - 4 keV)



- Internal radionuclides
 - $^{166\text{m}}\text{Ho}$ (β^- , $\tau_{1/2} = 1200$ y, produced along with ^{163}Ho)
 - Au pixel $200 \times 200 \times 2 \mu\text{m}^3 \rightarrow$ bkg ~ 0.5 c/eV/day/det/Bq($^{166\text{m}}\text{Ho}$)
 - $A(^{163}\text{Ho}) = 300\text{Bq/det}$ ($\sim 6.5 \times 10^{13}$ nuclei/det)
 - if $\text{bkg}(^{166\text{m}}\text{Ho}) < 0.1$ c/eV/day/det
 - $\rightarrow A(^{163}\text{Ho})/A(^{166\text{m}}\text{Ho}) > 1500$
 - $\rightarrow N(^{163}\text{Ho})/N(^{166\text{m}}\text{Ho}) > 6000$

ROACH2-based multiplexing

- Reconfigurable Open Architecture Computing Hardware (ROACH) designed by the Collaboration For Astronomy Signal Processing and Electronics Research (CASPER);
- Xilinx Virtex FPGA based digital data processing;
- Frequency comb generation (≈ 60 tones in the $0 \div 550$ MHz range);
- Quadrature frequency upmixing (500 MHz \rightarrow 5 GHz) and down-mixing (5 GHz \rightarrow 500 MHz);
- Signal channelizing and rf-SQUID signal de-modulation
- Real time signal processing;
- Strongly tested for MKIDs read-out (ARCONS, 2048 pixels)



Holmes design:

- $4 \times 256 = 1024$ pixels;
- Target: 64 resonances per ROACH-module;
- Complete system composed by 16 modules.