

The HOLMES experiment

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

ERC-Advanced Grant 2013

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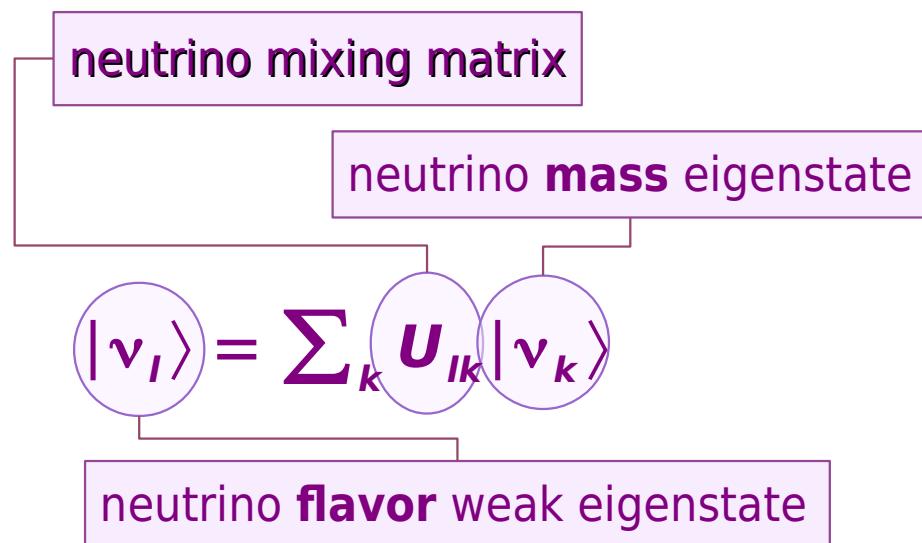


- absolute neutrino mass
- neutrino mass measurements
- low temperature detectors
- ^{163}Ho EC decay for direct neutrino mass measurements
- **HOLMES** experiment
 - sensitivity MC simulations
 - experiment design
 - task development status
- conclusions

Neutrino properties



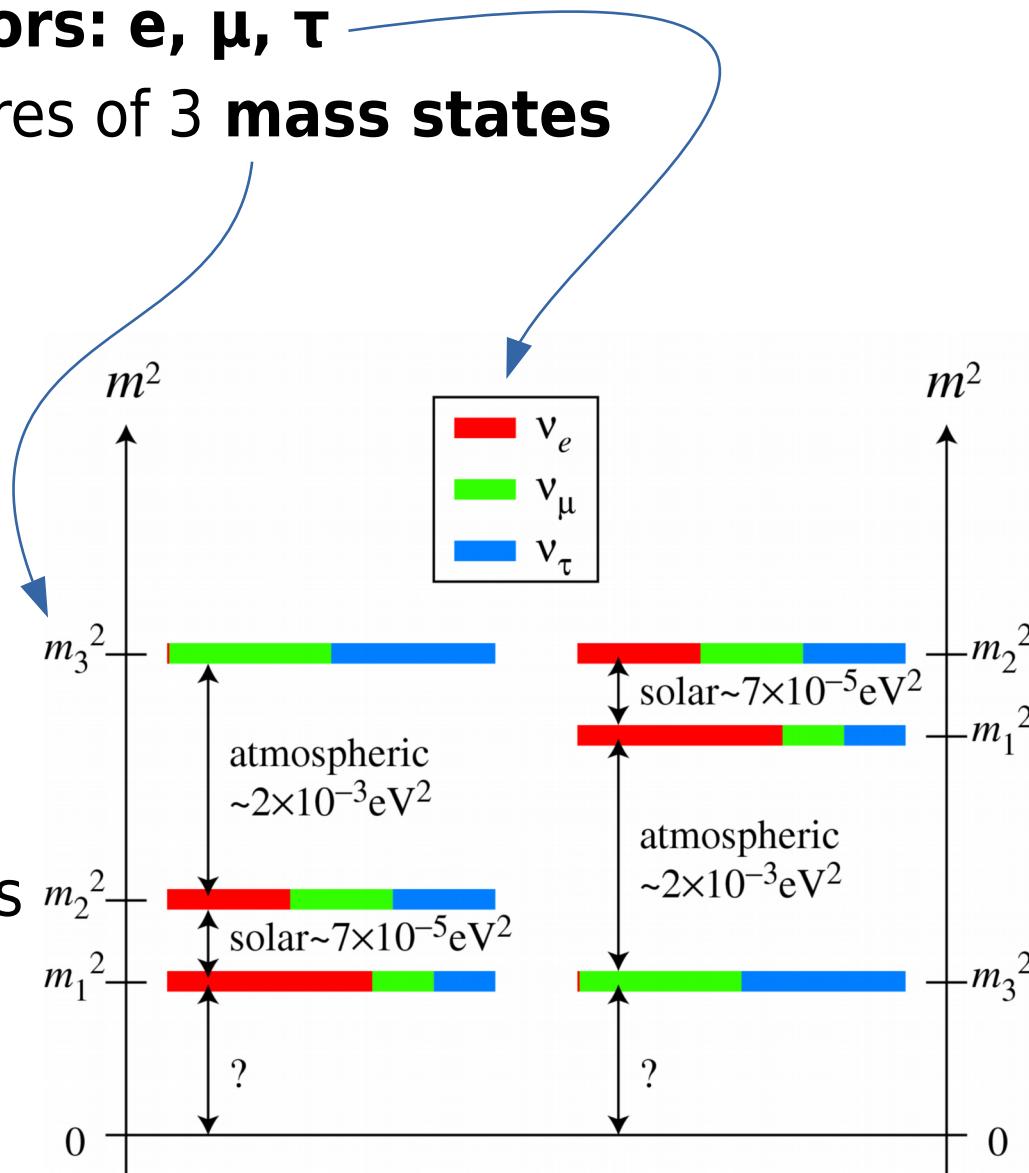
- neutrinos are massive fermions
 - there are 3 active neutrino **flavors: e, μ , τ**
 - neutrino flavor states are mixtures of 3 **mass states**



from neutrino oscillation experiments

$$\Delta m_{ik}^2 = |m_i^2 - m_k^2|$$

$$\sin^2 2\vartheta_k = f(|U_k|^2)$$

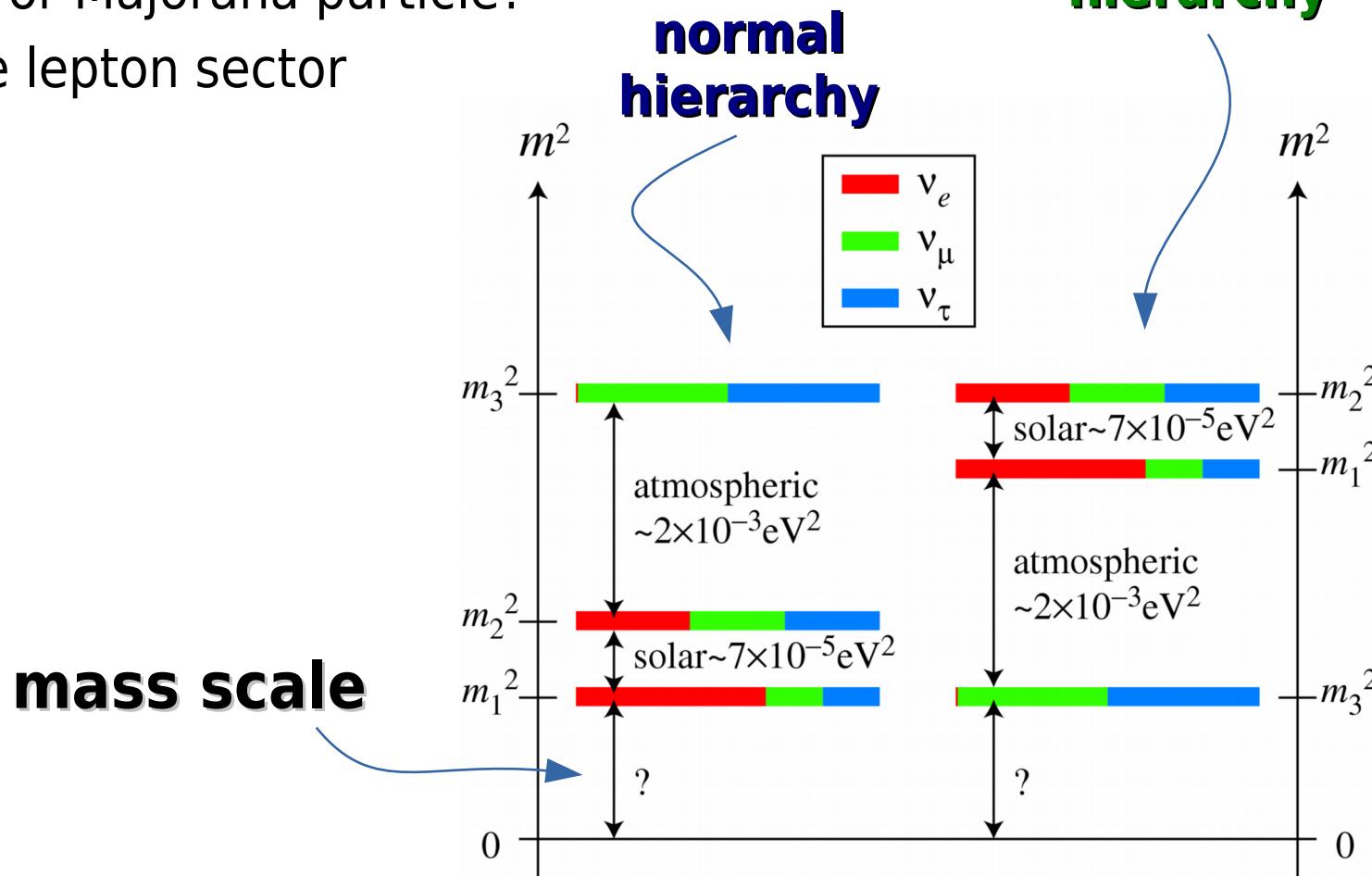


Neutrino open questions



- mass scale: i.e. mass of the lightest ν
- degenerate ($m_1 \approx m_2 \approx m_3$) or hierarchical masses
 - mass hierarchy: $m_1 < m_2 \ll m_3$ or $m_3 \ll m_1 \approx m_2$?
- $\nu = \bar{\nu}$? i.e. Dirac or Majorana particle?
- CP violation in the lepton sector

inverted
hierarchy



Direct neutrino mass measurements



■ kinematics of weak decays with ν emission

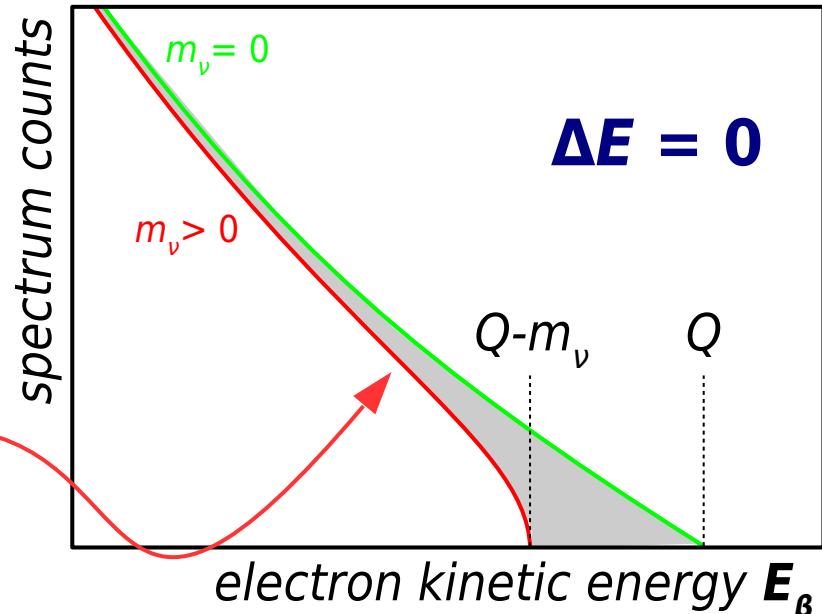
- ▶ low Q nuclear beta decays (${}^3\text{H}$, ${}^{187}\text{Re}$...)
- ▶ only energy and momentum conservation
- ▶ no further assumptions



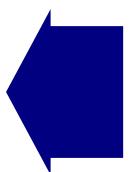
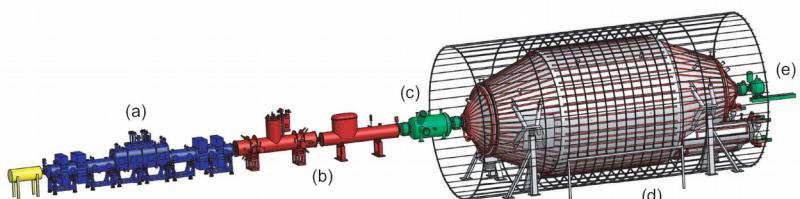
$$N(E_\beta) \propto p_\beta E_\beta (Q - E_\beta) \sqrt{((Q - E_\beta) - m_\nu^2)} F(z, E_\beta) S(E_\beta)$$

■ 2 approaches with different systematics:

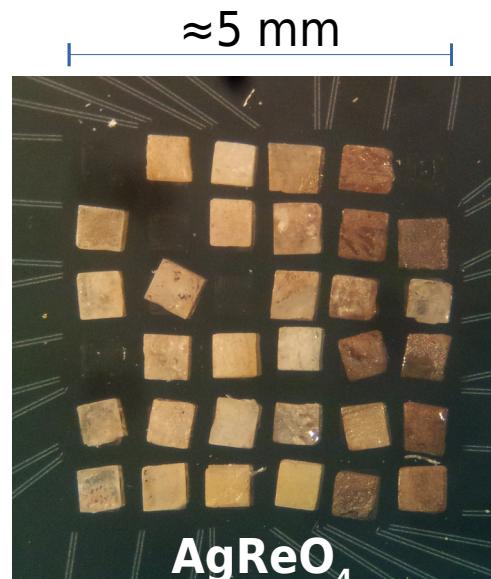
- ▶ **spectrometry** with the β source outside
- ▶ **calorimetry** with the β source inside



$$\Delta E = 0$$



KATRIN
large MAC-E filter
spectrometer with ${}^3\text{H}$



MARE/ECHO/HOLMES
array of low temperature
microcalorimeters
with ${}^{187}\text{Re}$ or ${}^{163}\text{Ho}$

Direct neutrino mass measurements / 2



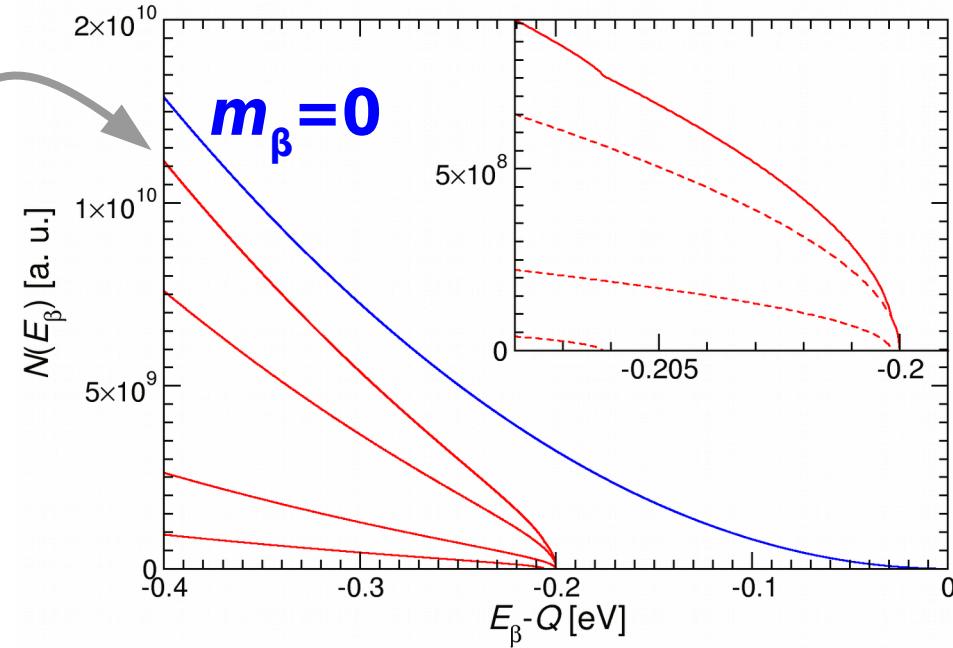
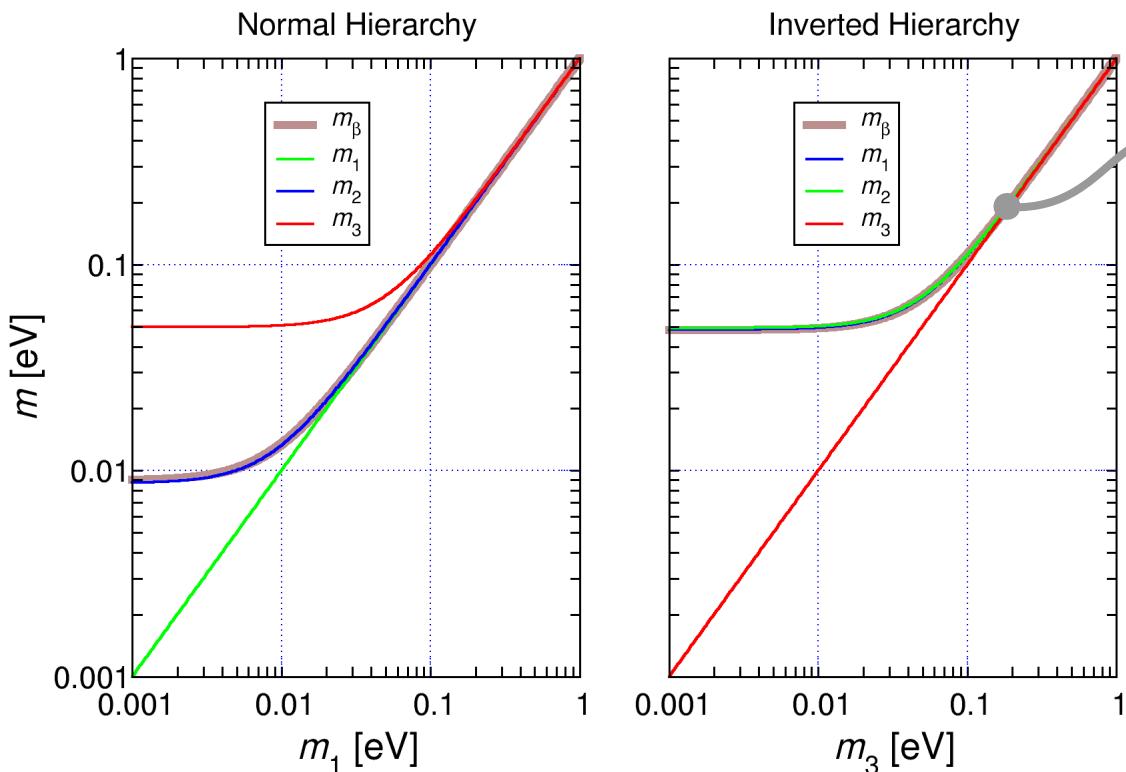
■ kinematics of weak decays

- observable in nuclear beta decays with $\bar{\nu}_e$ (ν_e) emission

$$|\nu_e\rangle = \sum_k \mathbf{U}_{ek} |\nu_k\rangle \quad N(E) \propto p_\beta E_\beta (Q - E_\beta) \sum_k |\mathbf{U}_{ek}|^2 \sqrt{(Q - E_\beta)^2 - \mathbf{m}_{\nu_k}^2} F(Z, E_\beta) S(E_\beta)$$

$$\mathbf{m}_\beta = \sqrt{\sum_k \mathbf{m}_{\nu_k}^2 |\mathbf{U}_{ek}|^2}$$

$$\rightarrow N(E) \approx p_\beta E_\beta (Q - E_\beta) \sqrt{(Q - E_\beta)^2 - \mathbf{m}_\beta^2} F(Z, E_\beta) S(E_\beta)$$



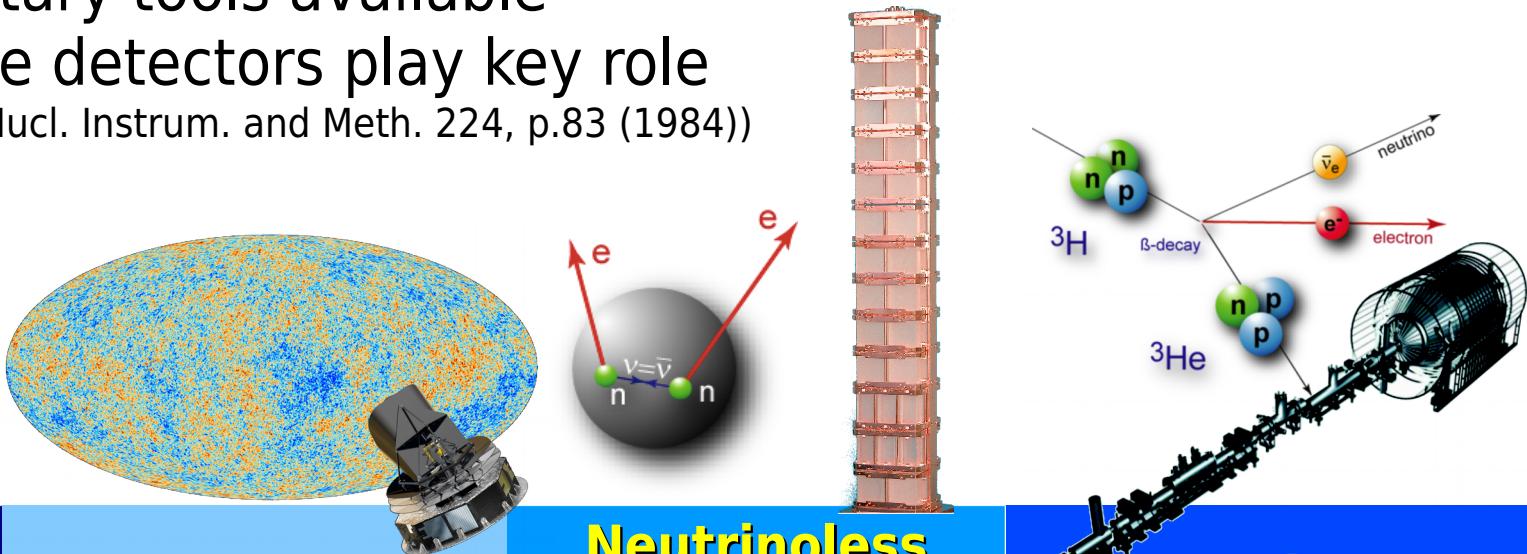
Mass scale: experimental tools / 1



three complementary tools available

→ low temperature detectors play key role

(E. Fiorini and T. Niinikoski, Nucl. Instrum. and Meth. 224, p.83 (1984))



tool	Cosmology CMB+LSS+...	Neutrinoless Double Beta decay	Beta decay end-point
observable	$m_{\Sigma} = \sum_k m_{\nu_k}$	$m_{\beta\beta} = \sum_k m_{\nu_k} U_{ek}^2 $	$m_{\beta} = (\sum_k m_{\nu_k}^2 U_{ek} ^2)^{1/2}$
present sensitivity	≈ 0.1 eV	≈ 0.1 eV	2 eV
future sensitivity	0.05 eV	0.05 eV	0.2 eV
model dependency	yes ☹	yes ☹	no ☺
systematics	large ☹	yes ☹	large ☹

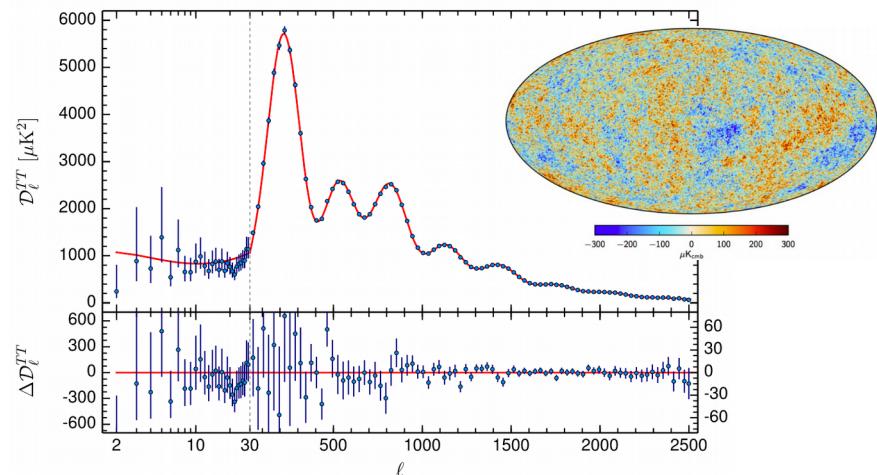
Experimental status for neutrino mass / 1



Cosmological measurements:

- Planck TT, TE, EE+lowP+BAO:
 $m_{\Sigma} < 0.17 \text{ eV}$ (95%)

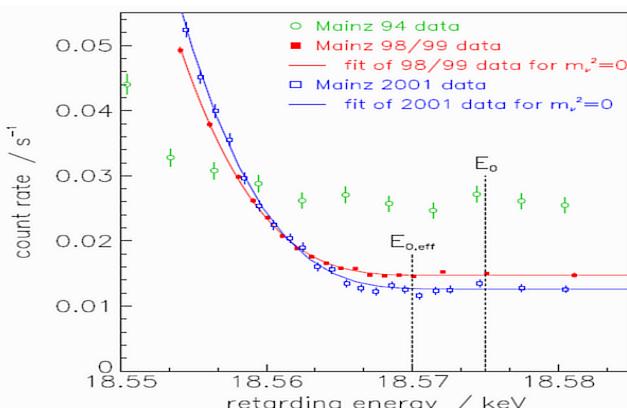
P. A. R. Ade et al., A&A 594, A13 (2016)



Tritium beta decay end-point measurements:

- Troitsk + Mainz experiments: $m_{\beta} < 2.2 \text{ eV}$ (95%)

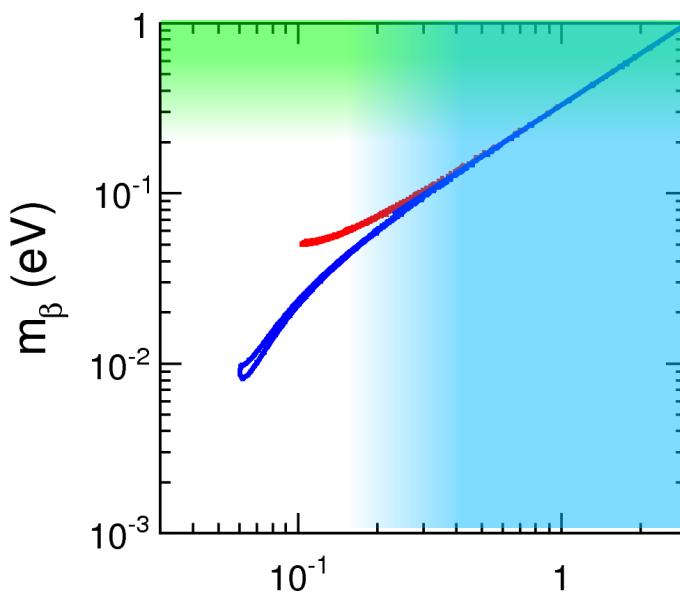
Ch. Kraus et al., Eur. Phys. J. C 73, (2013) 2323;
V. N. Aseev, Phys. Rev. D 84, (2011) 112003.



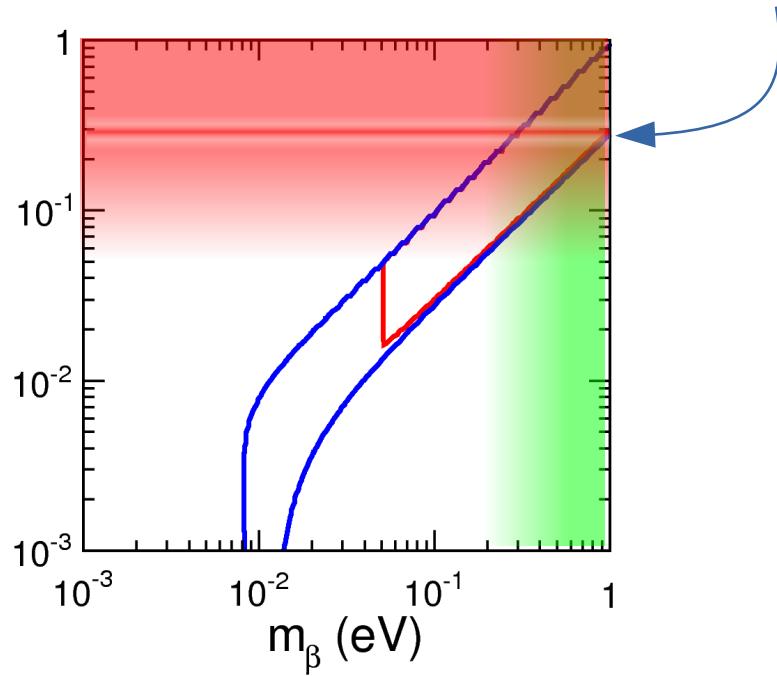
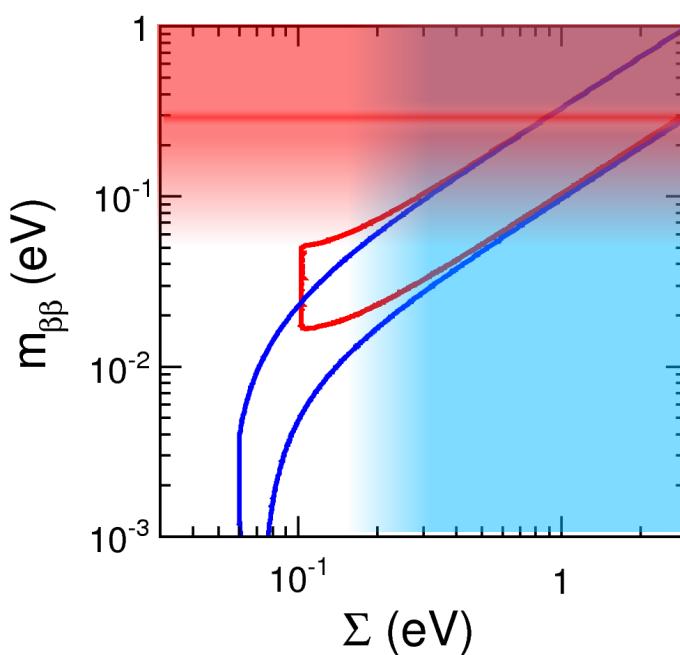
Neutrinoless double-beta decay searches:

- GERDA (^{76}Ge): $m_{\beta\beta} < 0.15 \div 0.33 \text{ eV}$ (90%) Nature 544 (2017) 7648
- KamLAND-Zen (^{136}Xe): $m_{\beta\beta} < 0.06 \div 0.16 \text{ eV}$ (90%) Phys. Rev. Lett. 117 (2016) 082503
- CUORE-0 (^{130}Te): $m_{\beta\beta} < 0.27 \div 0.76 \text{ eV}$ (90%) Phys. Rev. Lett. 115 (2015) 102502

Direct ν mass measurements: the status



situation about now



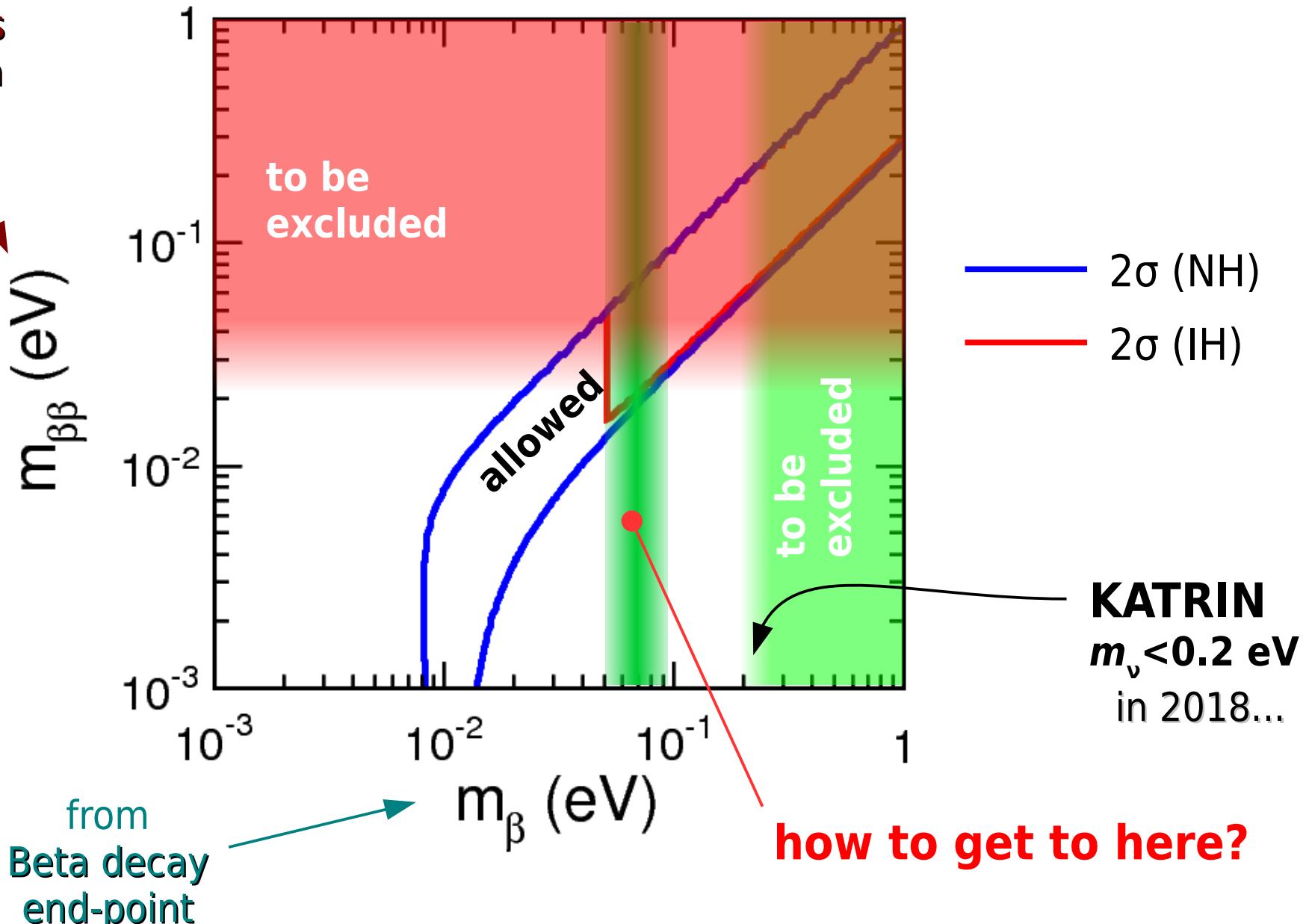
claim for $\beta\beta$ -0 ν observation in ^{76}Ge
H.V. Klapdor-Kleingrothaus et al.
Mod. Phys. Lett. A, 21 (2006) 1547

Direct ν mass measurements: the challenge



expected for the next few years
how to explore the Inverse Hierarchy?

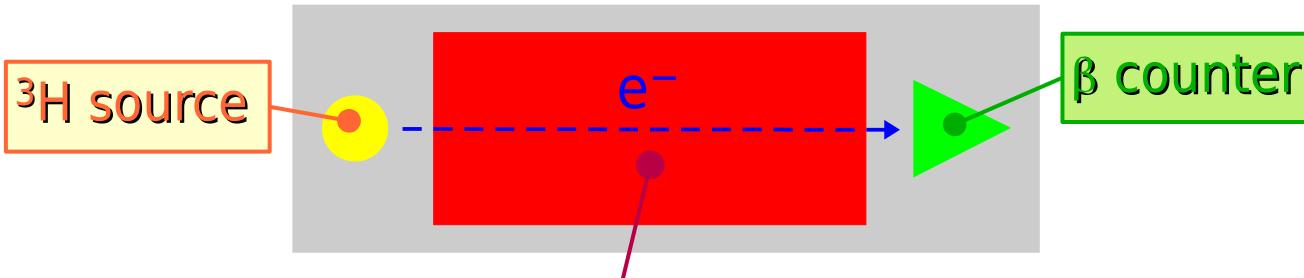
from
Neutrinoless
Double Beta
decay



Direct ν mass measurements: experimental



Spectrometers: source \neq detector

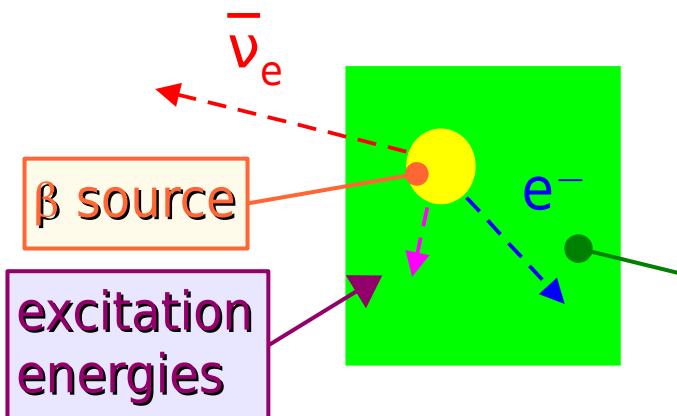


β differential or integral spectrometer

β s from the ${}^3\text{H}$ spectrum in δE are magnetically and/or electrostatically selected and transported to the counter

- ▲ high statistics
- ▲ high energy resolution
- ▼ large systematics
 - ▶ source effects
 - ▶ decays to excited states
- ▼ background

Calorimeters: source \subseteq detector



β calorimeter

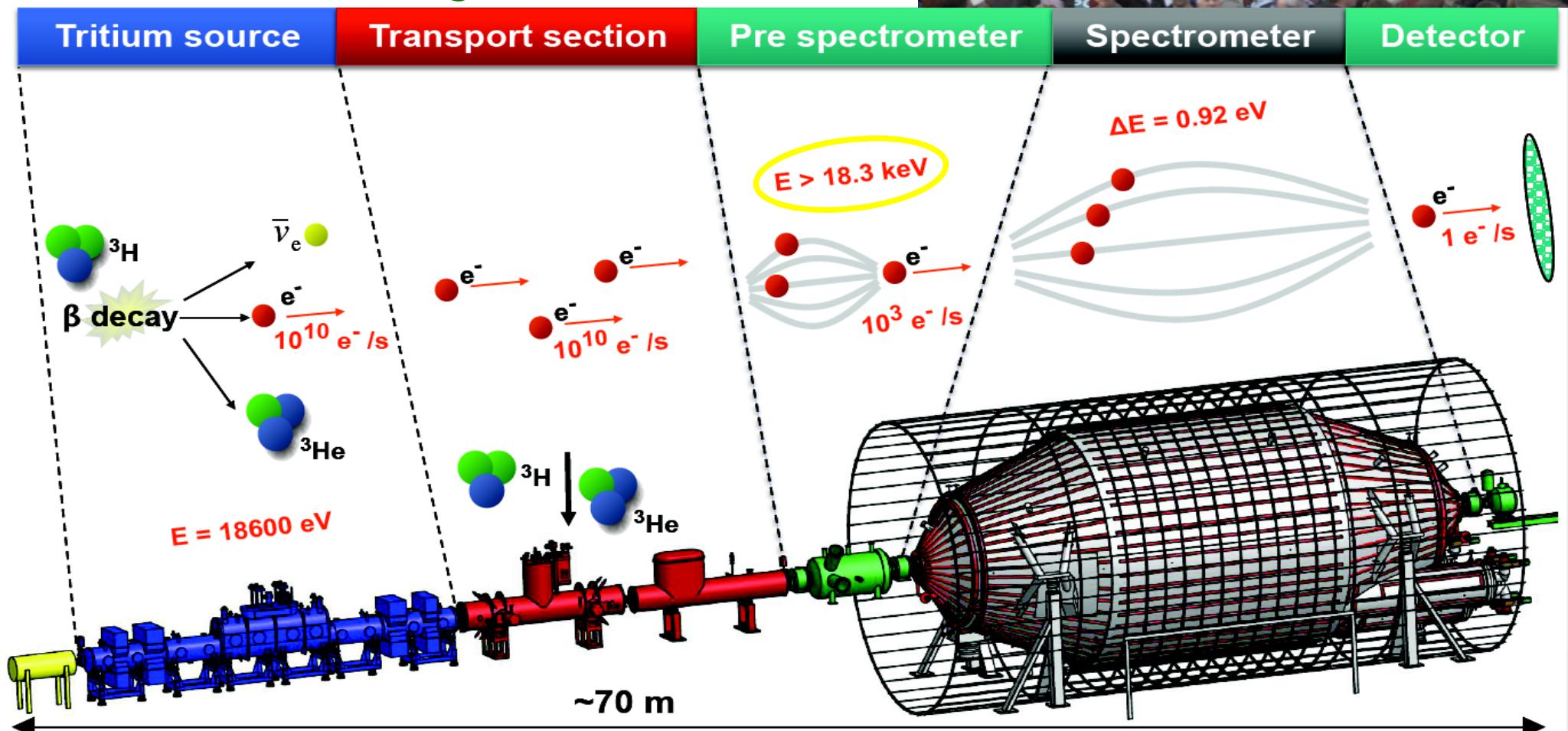
ideally measures all the energy E released in the decay except for the $\bar{\nu}_e$ energy: $E = Q - E_{\bar{\nu}}$

- ▲ no backscattering
- ▲ no energy losses in source
- ▲ no decay final state effects
- ▲ no solid state excitation
- ▼ limited statistics
- ▼ pile-up background
- ▼ spectrum related systematics

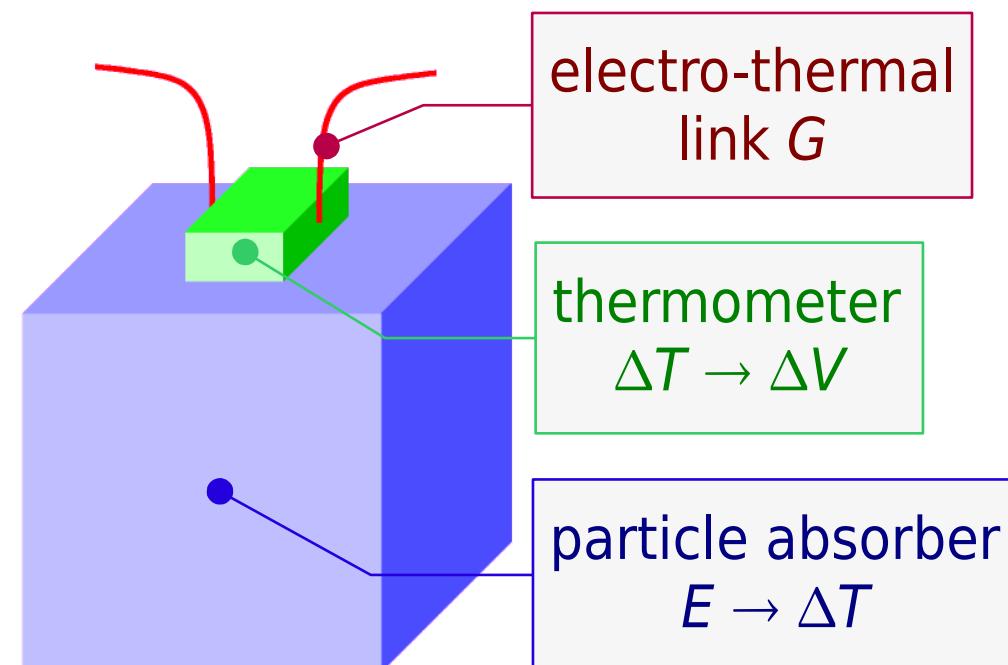
Spectrometers: KATRIN

largest electrostatic spectrometer
with gaseous ${}^3\text{H}$ source ($E_0 = 18.6\text{keV}$)

- expected statistical sensitivity
 $m_{\nu_e} < 0.2 \text{ eV } 90\% \text{ CL}$
- start data taking in 2017



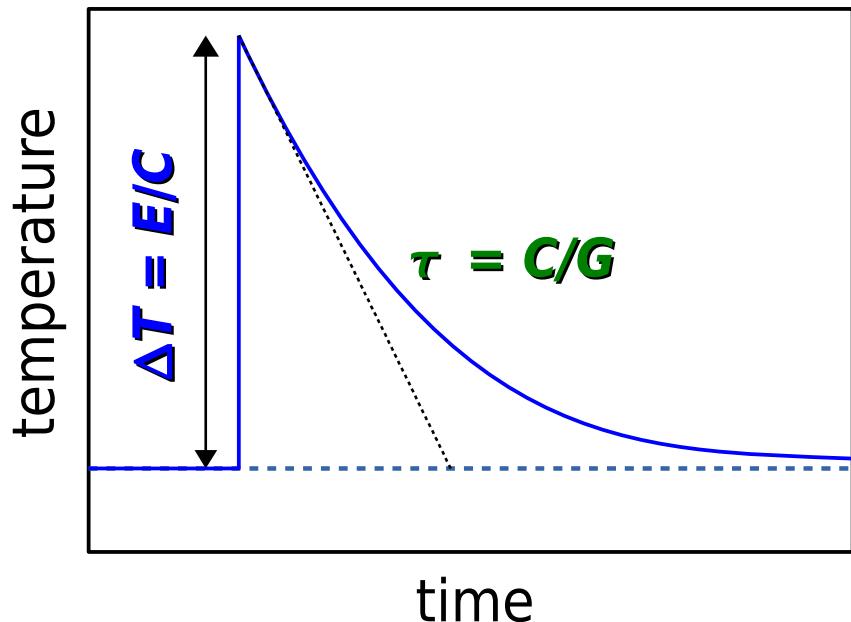
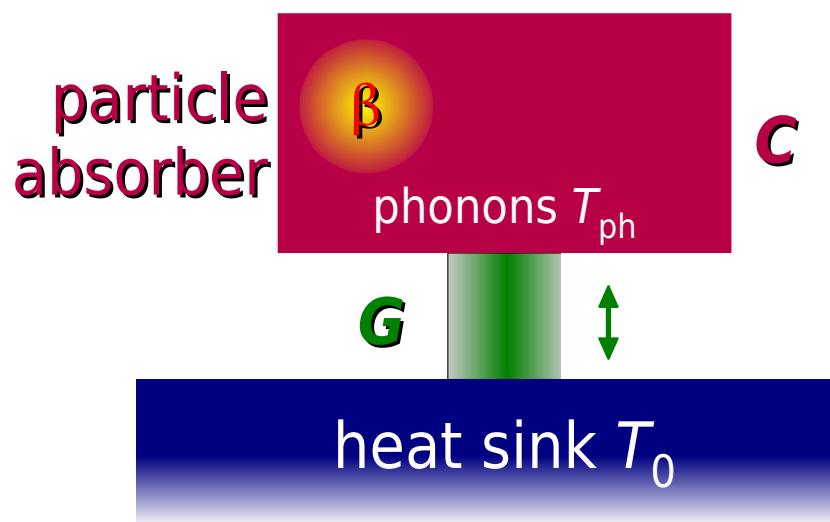
Low temperature detectors as calorimeters



- (quasi-)equilibrium thermal detector
- complete energy *thermalization*
 - ▶ **calorimetry**
- $\Delta T = E/C$ (C thermal capacity)
 - ▶ low C
 - ▷ low T (i.e. $T \ll 1\text{K}$)
 - ▷ dielectrics, superconductors
- Pros and cons
 - ▲ high energy resolution
 - ▲ large choice of absorber materials
 - ▲ true calorimeters
 - ▼ only energy and time informations
 - ▼ slow time response



Low temperature detector principles



$$C(T_{ph}) \frac{dT_{ph}}{dt} + G(T_{ph}, T_0) = P(t)$$

$$P(t) = \Delta E \delta(t) \rightarrow T_{ph}(t) = T_0 + \frac{\Delta E}{C} e^{-t/\tau}$$

for $t > 0$ and with $\tau = C/G$

- resolution limit: random energy flow through G
- statistical fluctuations of internal energy U

$$U = \langle U \rangle \pm \Delta U_{rms}$$

$$N_{ph} = \frac{\langle U \rangle}{\langle E_{ph} \rangle} = \frac{CT}{k_B T}$$

$$\Delta U_{rms} = \sqrt{N_{ph}} (k_B T) = \sqrt{k_B T^2 C}$$

1 mg of Re @ 100 mK

$$C \sim T^3 \text{ (Debye)} \Rightarrow C \sim 10^{-13} \text{ J/K}$$

$$6 \text{ keV X-ray} \Rightarrow \Delta T \sim 10 \text{ mK}$$

$$G \sim 10^{-11} \text{ W/K} \Rightarrow \tau = C/G \sim 10 \text{ ms}$$

$$\Delta U_{rms} \sim 1 \text{ eV}$$

CUORE experiment



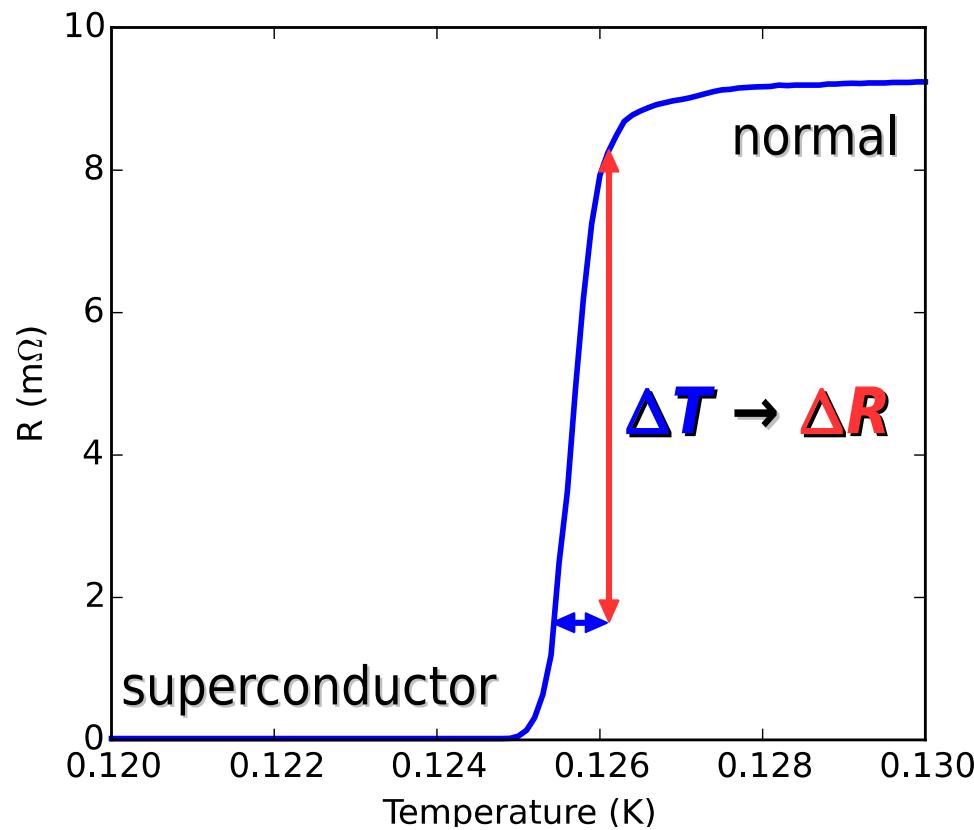
CUORE: Cryogenic Underground Observatory for Rare Events

- ▶ searches for ^{130}Te neutrinoless double-beta decay
- ▶ 988 natural TeO_2 750 g crystals as low temperature detectors
- ▶ total mass **740 kg TeO_2** \Rightarrow 206 kg of ^{130}Te
- ▶ **now cold at about 10 mK**

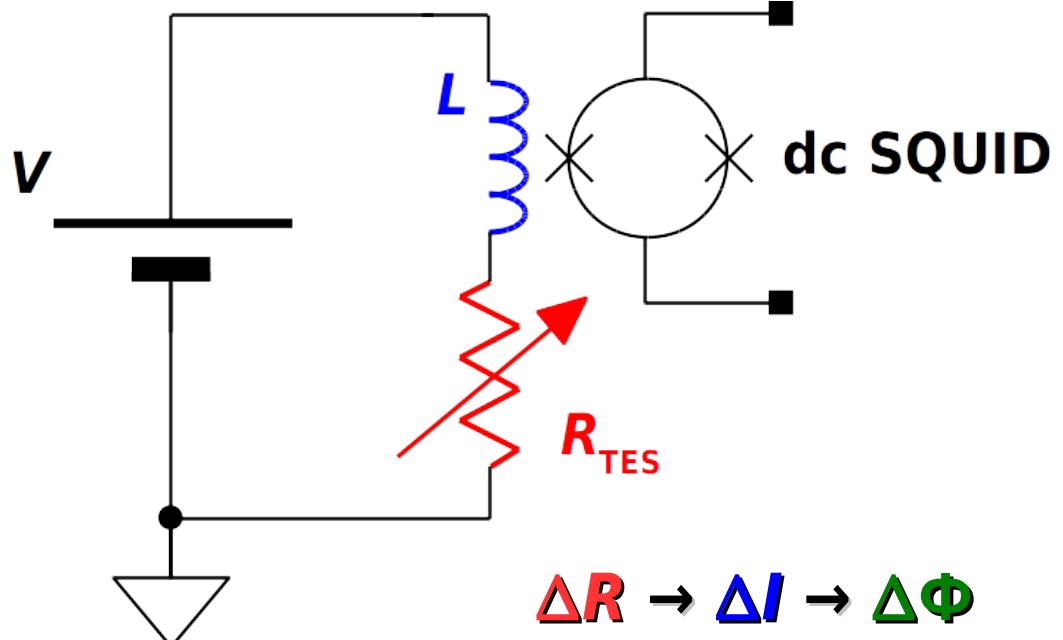
Superconducting transition edge sensors (TES)



- superconductor thin films operated inside the phase transition at T_c
 - ▶ elemental superconductors: Ir ($T_c = 112$ mK), W ($T_c = 15$ mK), ...
 - ▶ metal-superconductor bilayers \Rightarrow tunable T_c (20÷200 mK) : Mo/Cu, Ti/Au, Ir/Au, ...
- high sensitivity $TdR/(RdT) \approx 100$ \Rightarrow high energy resolution
 - ▶ as **thermal sensors** $\rightarrow \sigma_E^2 \approx \xi^2 k_B T^2 C$
- strong electron-phonon coupling \Rightarrow high intrinsic speed
- low impedance \Rightarrow SQUID read-out \Rightarrow multiplexing for large arrays



TES read-out: constant voltage bias



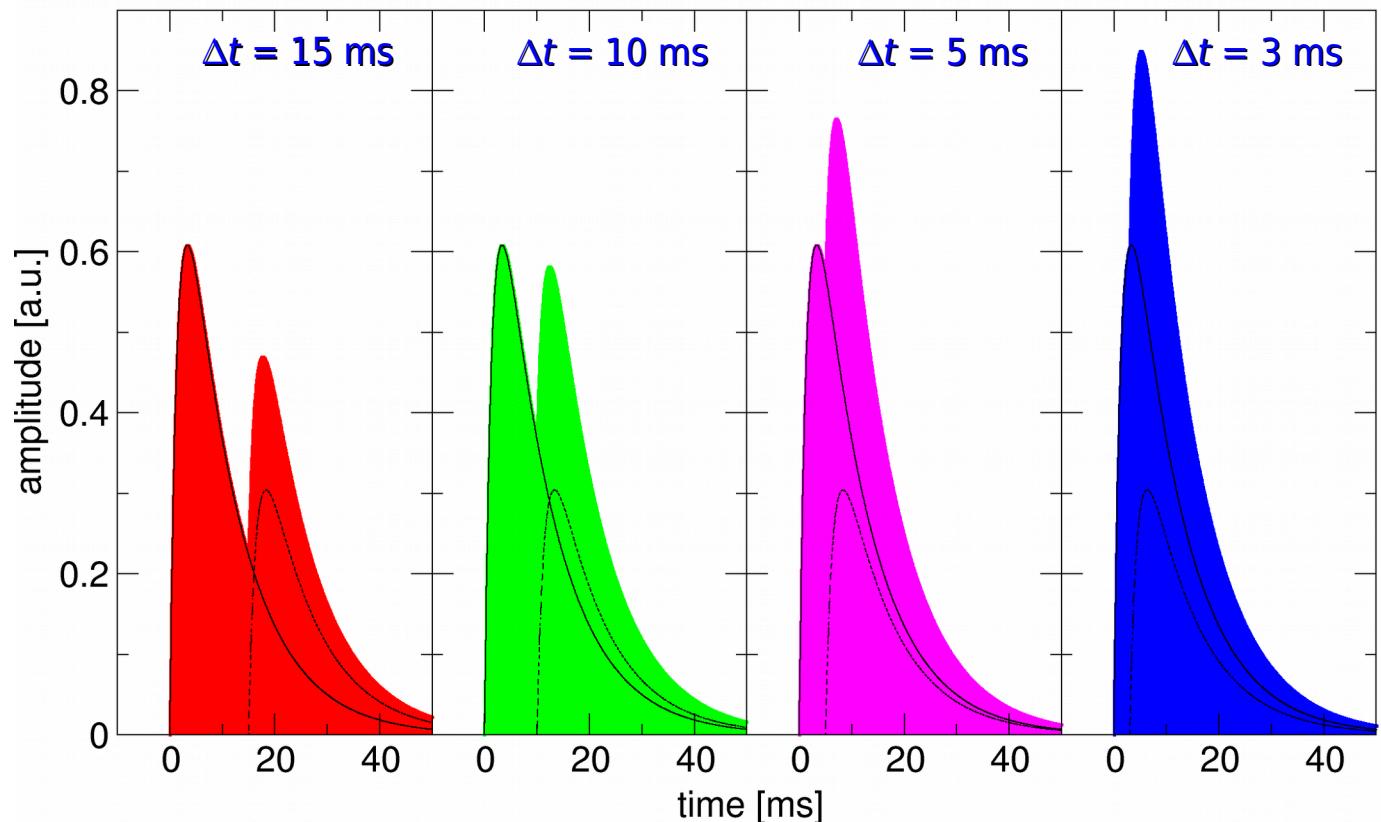


simple pulse model

$$A(t) = A_0(e^{-t/\tau_{decay}} - e^{-t/\tau_{rise}})$$

2 pulses with:

- $\tau_{rise} = 1.5 \text{ ms}$
- $\tau_{decay} = 10 \text{ ms}$
- $A_2/A_1 = 0.5$



resolving time $\tau_R \approx$ pulse rise time τ_{rise}

Calorimetry of beta sources



- calorimeters measure the **entire spectrum** at once
 - ▶ low $E_0 \beta$ decaying isotopes for more statistics near the end-point
 - ▶ best choice ^{187}Re : $Q = 2.5 \text{ keV}$, $\tau_{1/2} \approx 4 \times 10^{10} \text{ y} \Rightarrow F(\Delta E = 10 \text{ eV}) \approx (\Delta E/Q)^3 = 7 \times 10^{-8}$
 - ▶ other option ^{163}Ho electron capture: $Q \approx 2.6 \text{ keV}$, $\tau_{1/2} \approx 4600 \text{ y}$

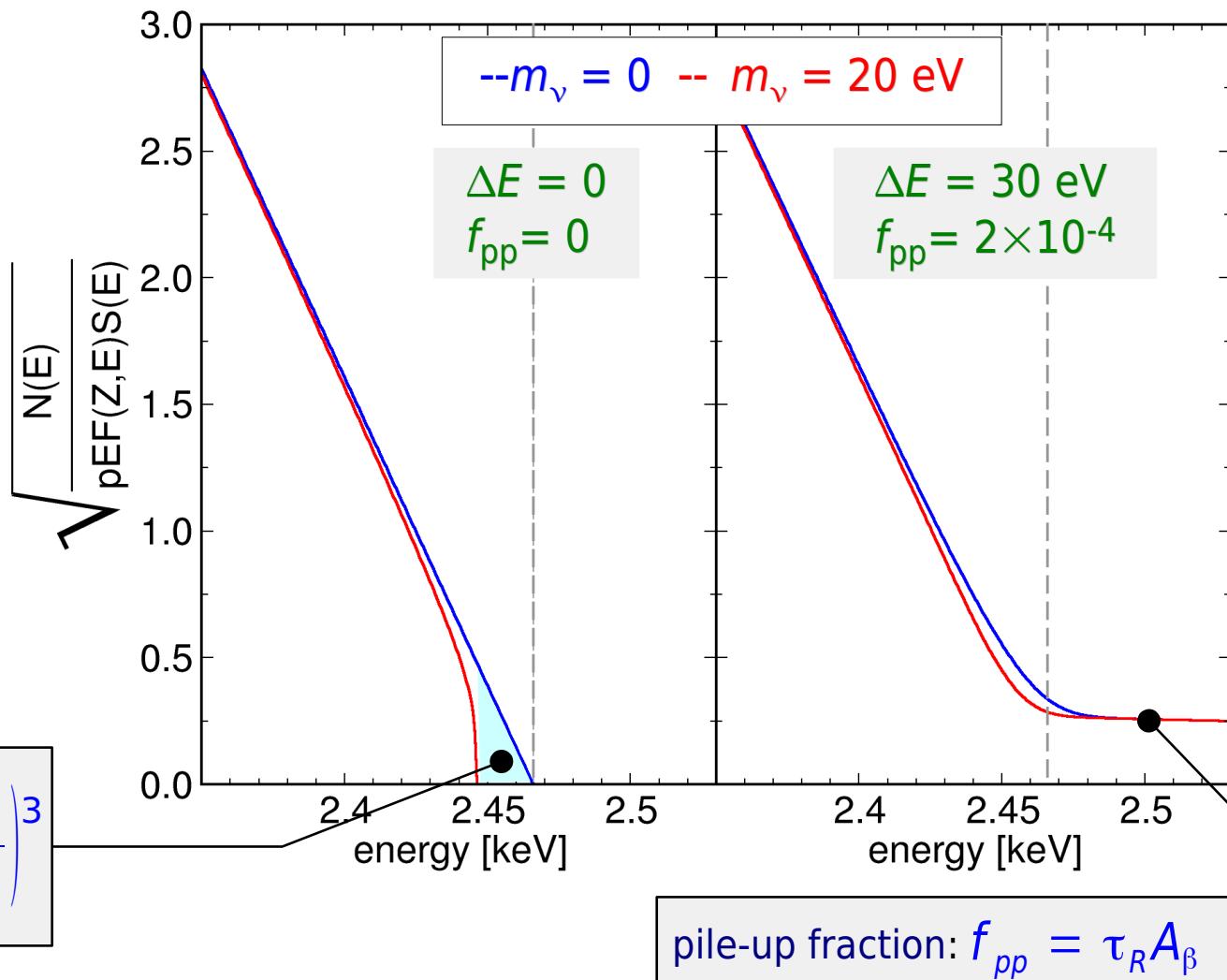
advantages

- ▲ no back-scattering
- ▲ no energy losses in the source
- ▲ no atomic/molecular final state effects
- ▲ no solid state excitation

drawbacks

- ▼ limited statistics
- ▼ pile-up background
- ▼ spectrum related systematics

$$F(\Delta E) \approx \left(\frac{\Delta E}{Q} \right)^3$$



β decay calorimetry statistical sensitivity



resolving time τ_R

analysis interval ΔE

source activity A_β

number of detectors N_{det}

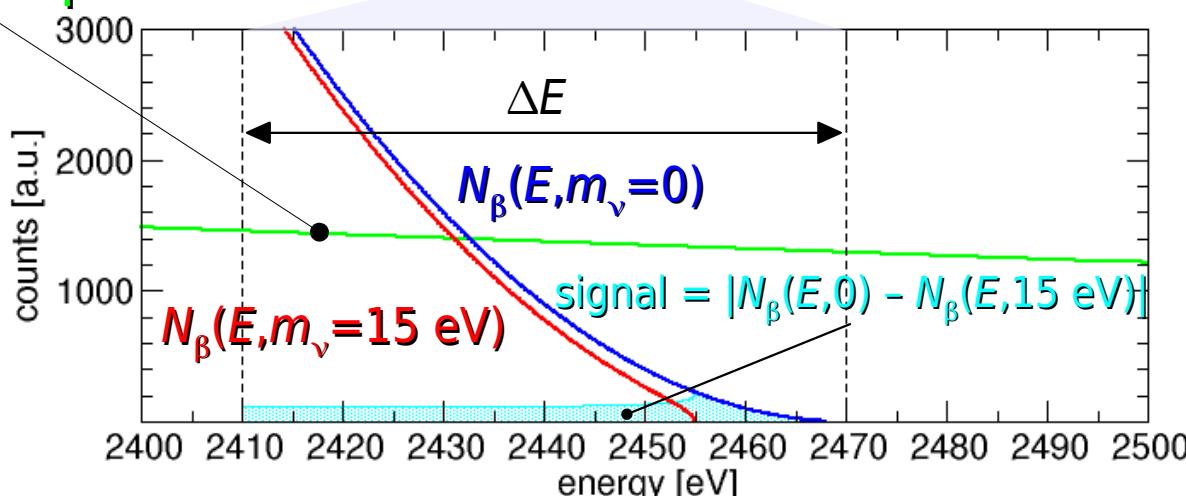
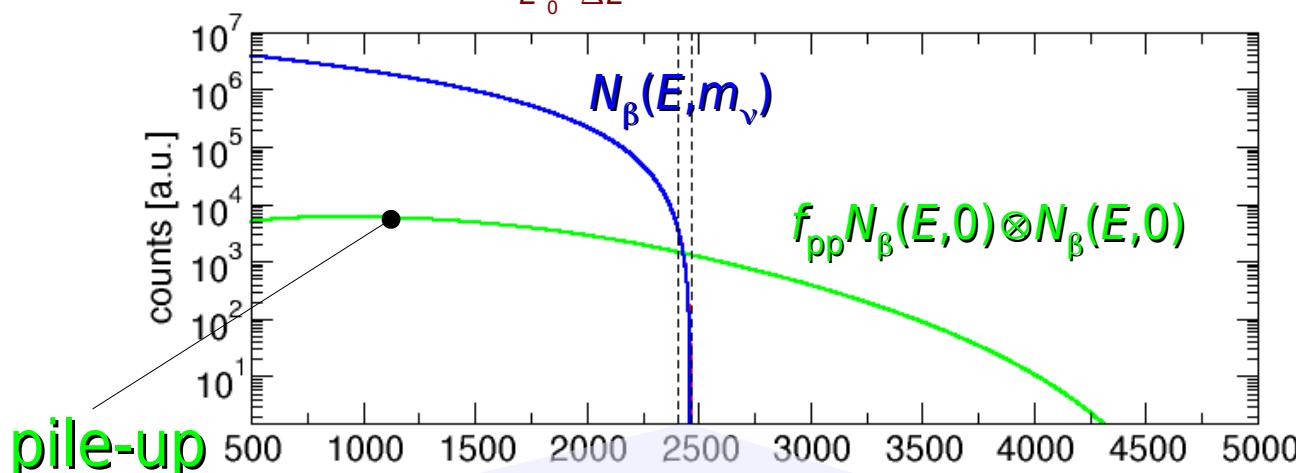
pile-up fraction $f_{\text{pp}} = \tau_R A_\beta$

experimental exposure $t_M = T \times N_{\text{det}}$

$$F_{\Delta E}(m_\nu) = A_\beta N_{\text{det}} \int_{E_0 - \Delta E}^{E_0} N_\beta(E, m_\nu) dE$$

$$N_\beta(E, m_\nu) \approx \frac{3}{Q^3} (Q - E)^2 \sqrt{1 - \frac{m_\nu^2}{(Q - E)^2}}$$

$$F_{\Delta E}(0) \approx A_\beta N_{\text{det}} \frac{\Delta E^3}{Q^3} \rightarrow {}^{187}\text{Re } Q = 2.5 \text{ keV}$$



$$f_{\text{pp}} = \tau_R A_\beta \ll \frac{\Delta E^2}{E_0^2} \quad \text{negligible pile-up}$$

$$\sum_{90} (m_\nu) \approx 0.89 \sqrt[4]{\frac{Q^3 \Delta E}{A_\beta t_M}}$$

- experimental challenges
- ▶ energy resolution ΔE
 - ▶ time resolution τ_R
 - ▶ exposure $t_M = N_{\text{det}} \times T$
 - ▶ detector activity A_β



^{187}Re β decay



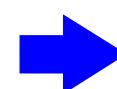
- $5/2^+ \rightarrow 1/2^-$ unique first forbidden transition $\Rightarrow S(E)$
- end point $Q = 2.47$ keV
- half-life time $\tau_{1/2} = 43.2$ Gy
- natural abundance a.i. = 63%
 - ▶ 1 mg metallic Rhenium $\Rightarrow \approx 1.0$ decay/s

■ **metallic rhenium** single crystals

▶ superconductor with $T_c = 1.6$ K

▶ NTD thermistors

▶ **MANU experiment (Genova)**

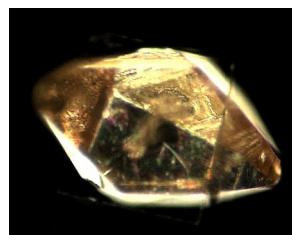


$m_\nu < \approx 15$ eV

■ **dielectric rhenium compound** (AgReO_4) crystals

▶ Silicon implanted thermistors

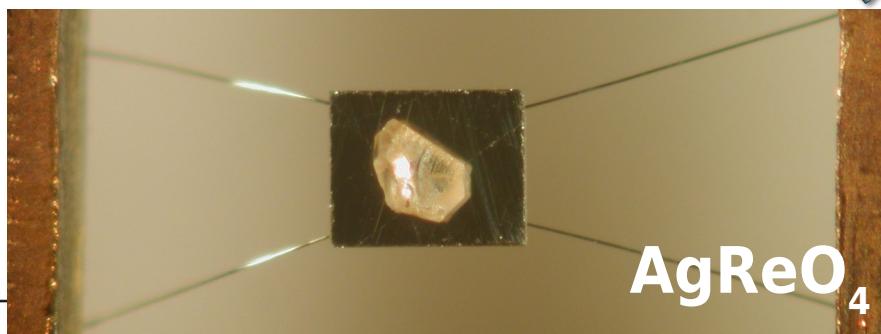
▶ **MIBETA experiment (Milano)**



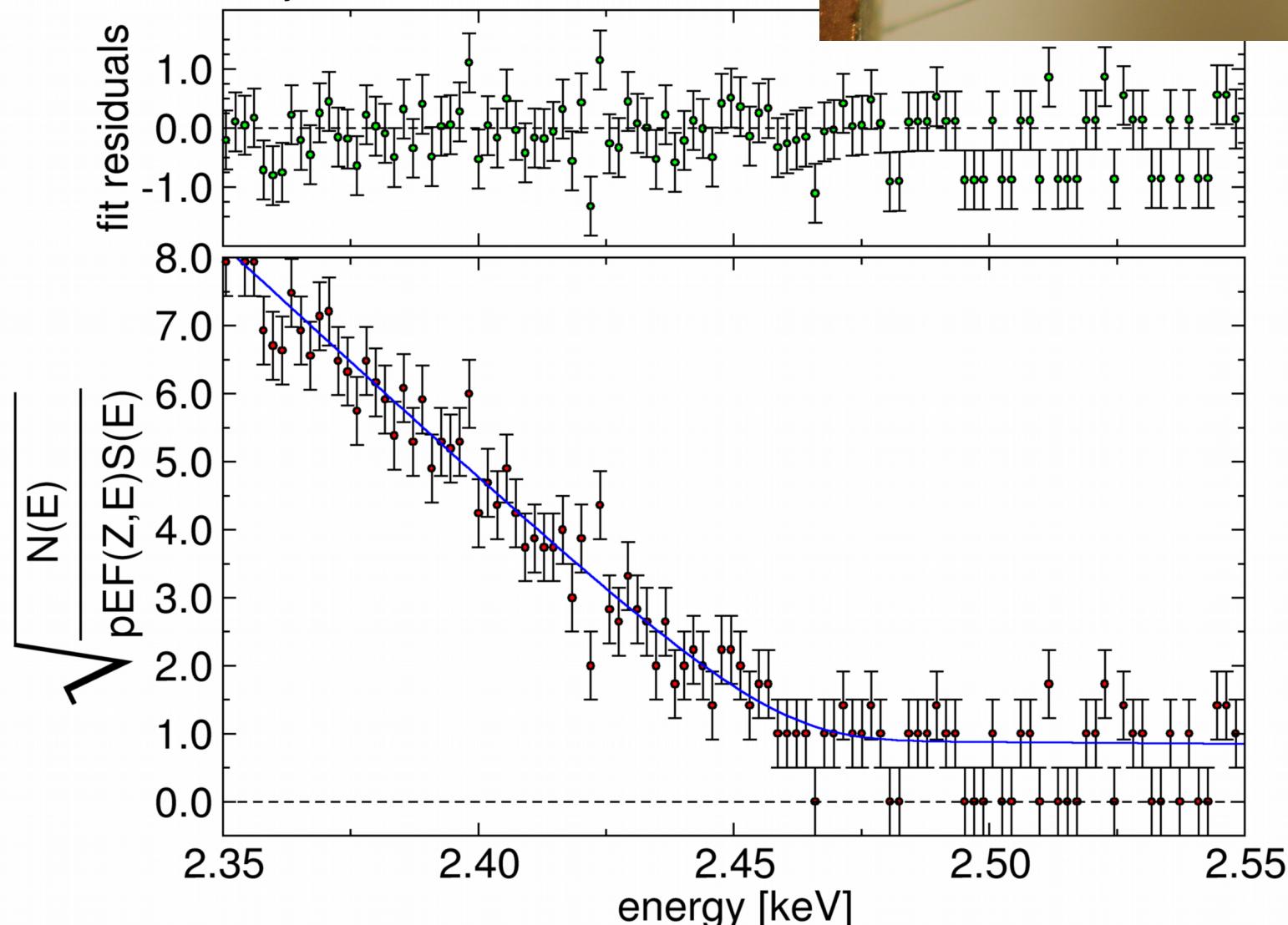
MIBETA experiment results



- 0.6 years live time (0.45 years only β)
- 6.2×10^6 ^{187}Re decays above 700 eV
- $m_\nu^2 = -96 \pm 189_{\text{stat}} \pm 63_{\text{sys}} \text{ eV}^2$
- $m_\nu < 15.2 \pm 2.0_{\text{sys}} \text{ eV (90 \% C.L.)}$



AgReO_4



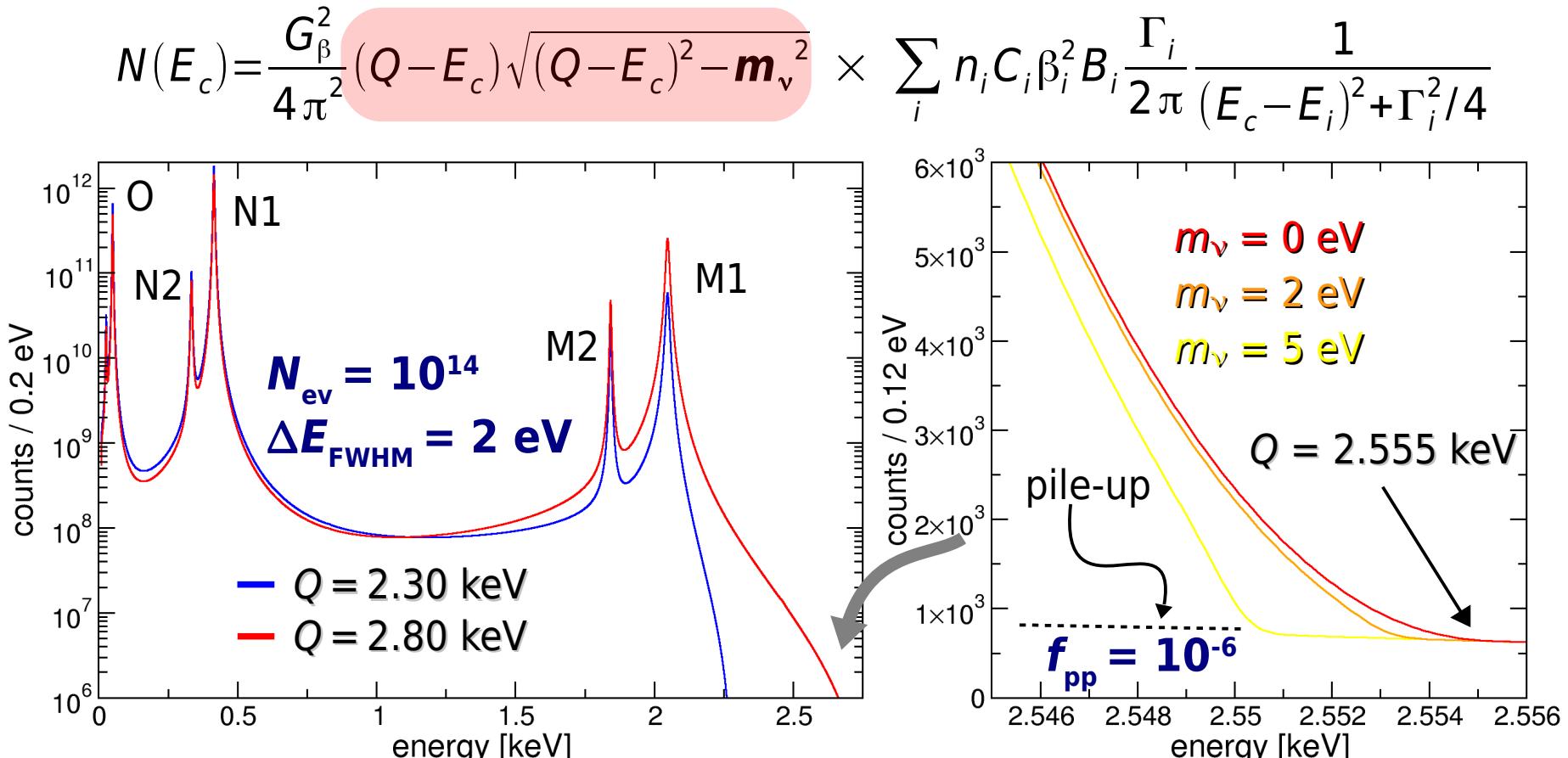
Electron capture calorimetric experiments



electron capture from shell $\geq M1$

A. De Rújula and M. Lusignoli, Phys. Lett. B 118 (1982) 429

- calorimetric measurement of Dy atomic de-excitations (mostly non-radiative)
- **$Q = 2.8 \text{ keV}$** (recently measured with Penning trap)
 - ▶ end-point rate and ν mass sensitivity depend on **$Q - E_{M1}$**
- **$\tau_{\nu} \approx 4570 \text{ years}$** → few active nuclei are needed ($2 \times 10^{11} \text{ }^{163}\text{Ho}$ nuclei $\leftrightarrow 1 \text{ Bq}$)



From ^{187}Re to ^{163}Ho calorimetric experiments



■ scaling up ^{187}Re experiments for sub-eV sensitivity

→ **MARE** (Microcalorimeter Array for a Rhenium Experiment)

- ▶ no clear understanding Re absorber physics in spite of 20 years of R&D
- ▶ low ^{187}Re specific activity → “large” masses → fabrication issues
- ▶ possible large systematics → Beta Environmental Fine Structure (BEFS)

■ **^{163}Ho seems to be better than ^{187}Re**

- ▶ higher specific activity → *Holmium detector* not needed
- ▶ *self calibrating* → better control of systematics
- ▶ **but**
 - higher Q → maybe less sensitive
 - pile-up spectrum
 - chemical effects on Q

■ two active projects presently

- ▶ **ECHo** (Heidelberg)
- ▶ **MARE** (→ now **HOLMES**)
- ▶ Los Alamos National Lab., Standford University ?, ...

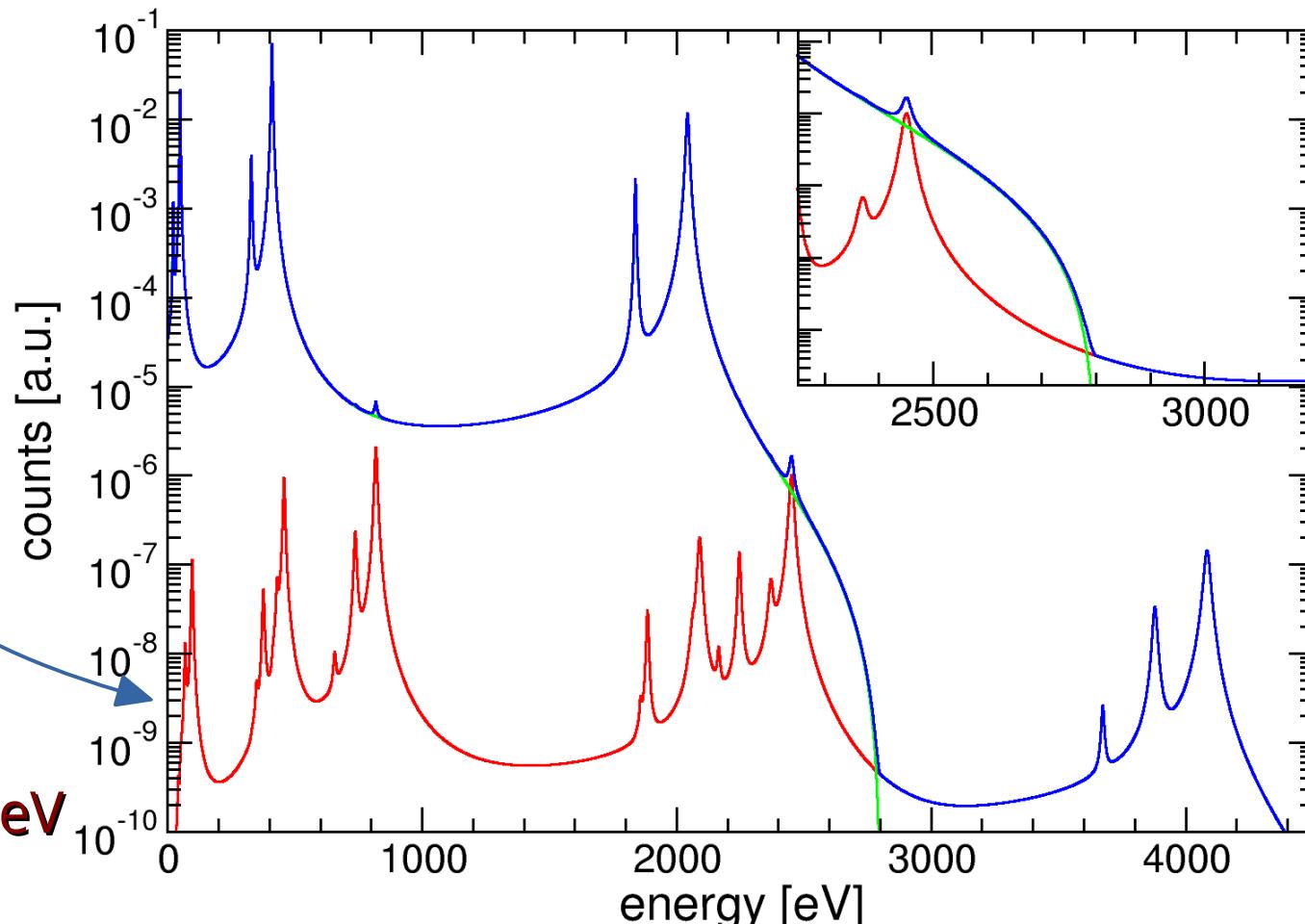
Electron capture end-point experiment



- no direct calorimetric measurement of Q (end-point) so far
- complex pile-up spectrum

► $N_{pp}(E) = f_{pp} N_{EC}(E) \otimes N_{EC}(E)$ with $f_{pp} \approx A_{EC} \tau_R$

A_{EC} EC activity per detector
 τ_R time resolution (\approx rise time)



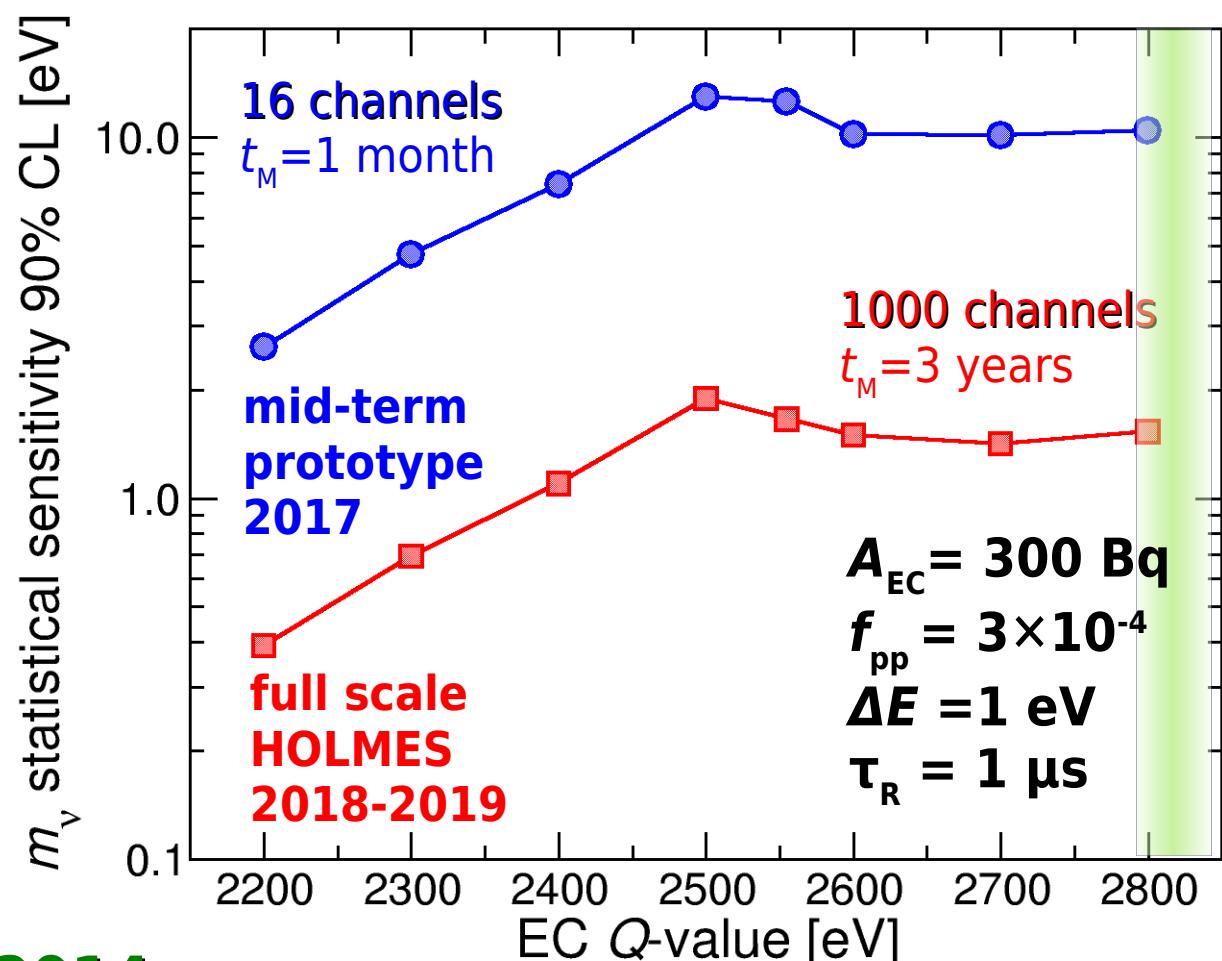
goal

- neutrino mass measurement: m_ν statistical sensitivity as low as 0.4 eV
- prove technique potential and scalability:
 - ▶ assess EC spectral shape
 - ▶ assess systematic errors

baseline

- **TES microcalorimeters** with **implanted ^{163}Ho**
 - ▶ 6.5×10^{13} nuclei per pixel
 - $A_{\text{EC}} = 300 \text{ dec/sec}$
 - ▶ $\Delta E \approx 1 \text{ eV}$ and $\tau_R \approx 1 \mu\text{s}$
- **1000 channel array**
 - ▶ $6.5 \times 10^{16} {}^{163}\text{Ho}$ nuclei
 $\rightarrow \approx 18 \mu\text{g}$
 - ▶ 3×10^{13} events in **3 years**

→ started on February 1st 2014





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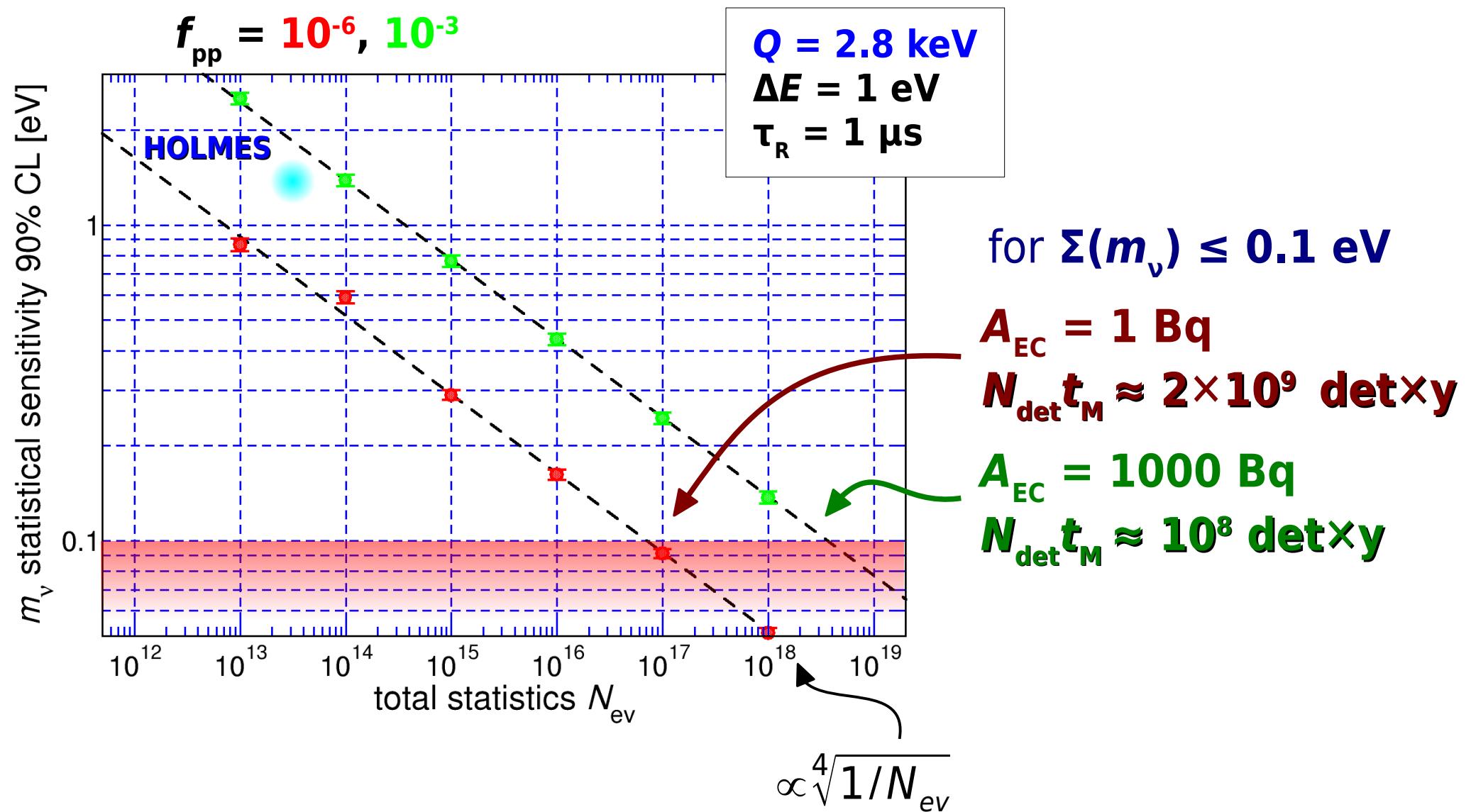
R.Dressler
S.Heinitz
D.Schumann

CENTRA-IST M.Ribeiro-Gomes

ILL
U.Koester



Potential sensitivity: Montecarlo simulations



More on EC end-point experiments / 1



- shake-up/shake-off → double hole excitations

- ▶ n -hole excitations possible but less probable

- ▶ authors do not fully agree on energies and probabilities

- even more complex pile-up spectrum

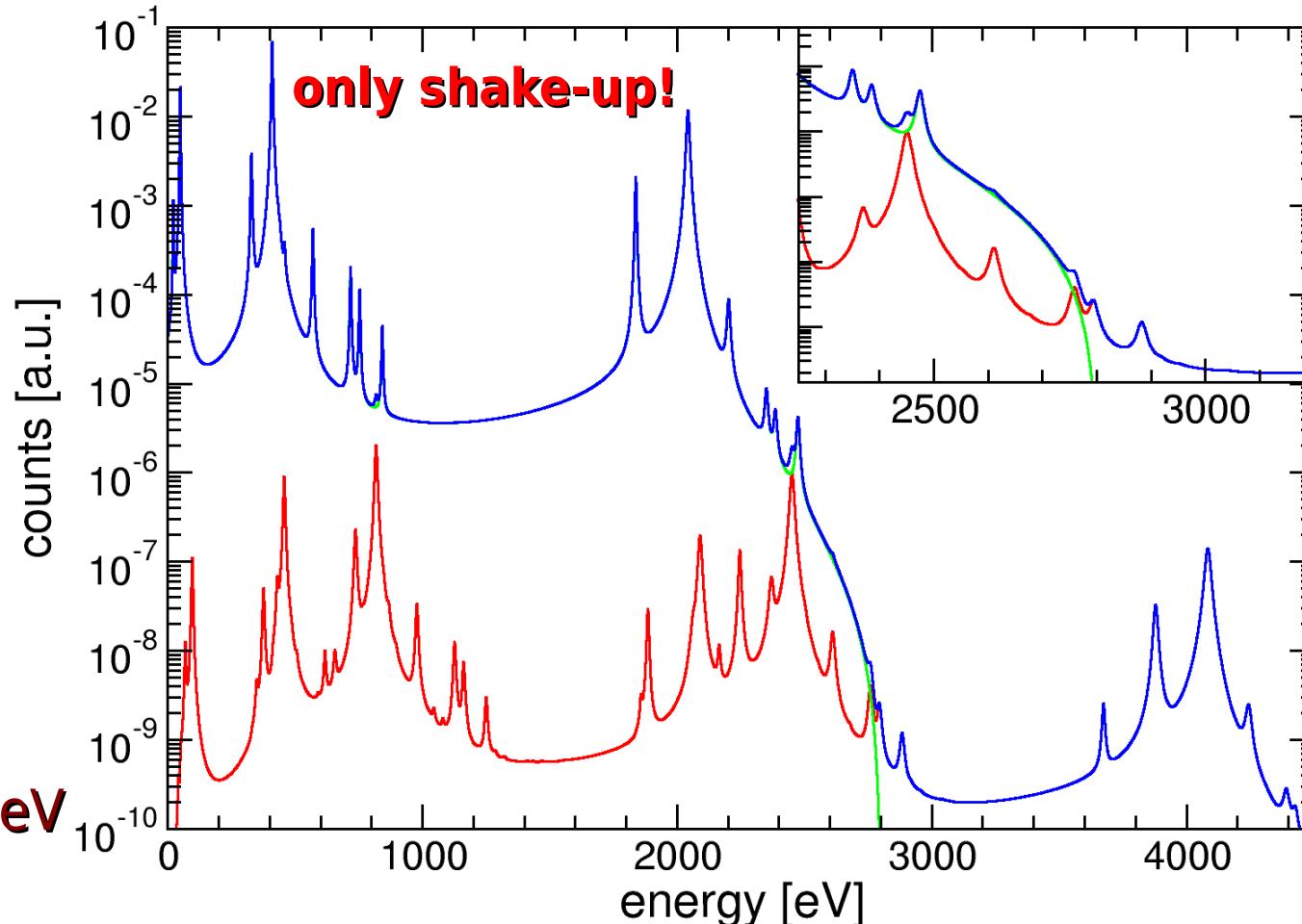
- ▶ it may be worth keeping f_{pp} smaller than 10^{-4}



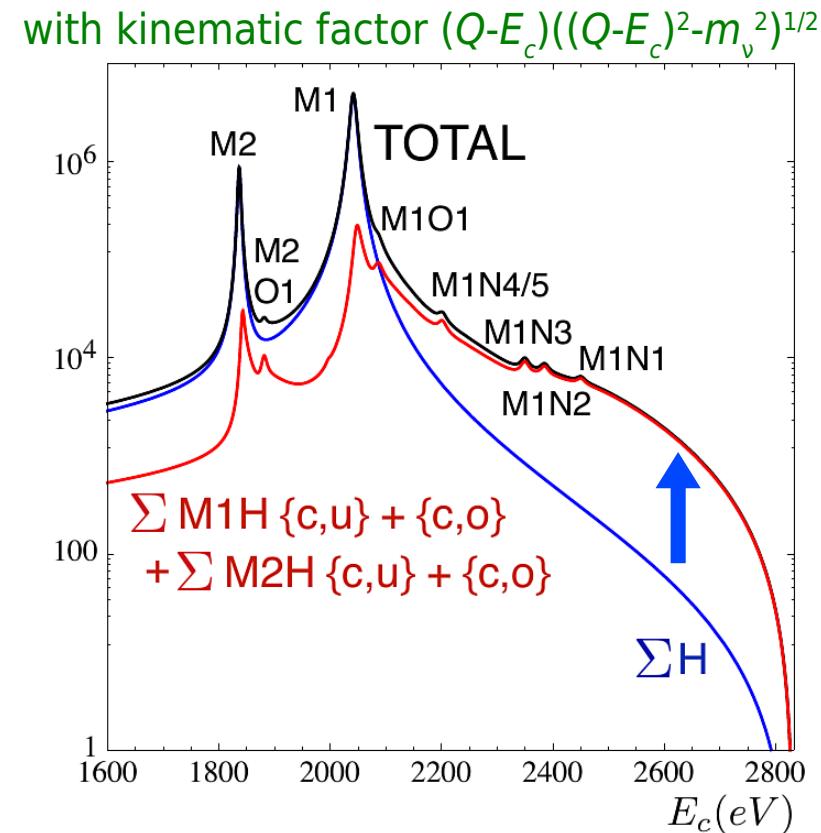
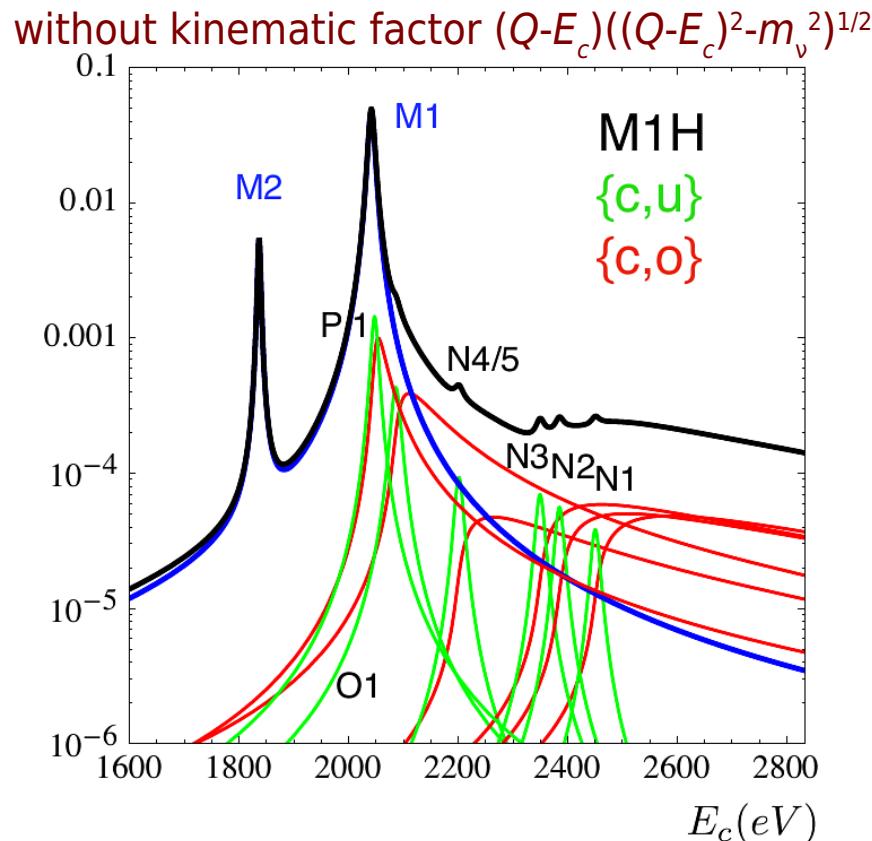
A.De Rújula, arXiv:1305.4857

R.G.H.Robertson, arXiv:1411.2906

A.Faessler et al., PRC 91 (2015) 45505



More on EC end-point experiments / 2



A.De Rújula & M. Lusignoli, J. High Energ. Phys. (2016) 2016: 15

- including **2-hole shake-off processes**



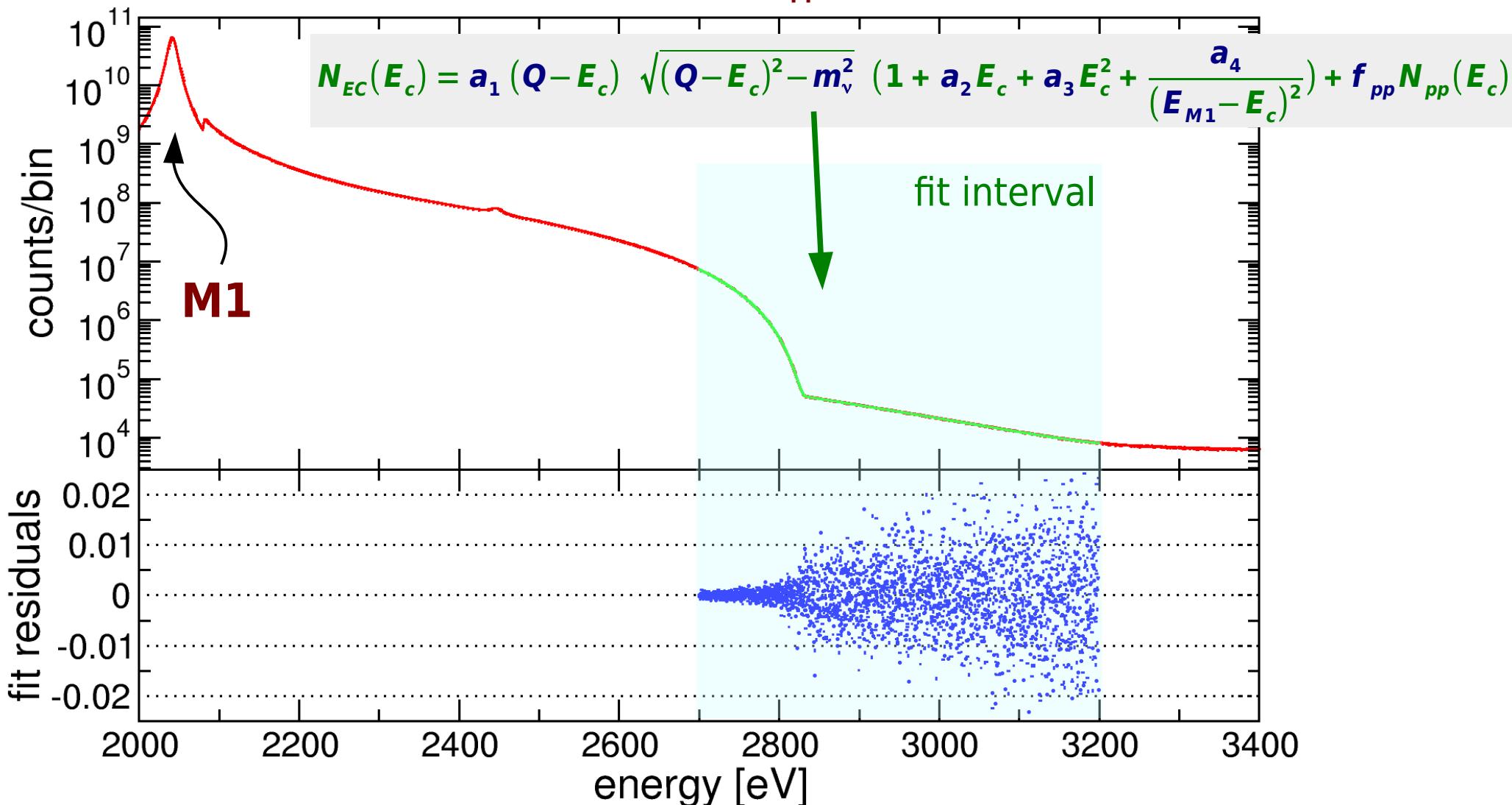
- dominate rate at end-point
 - **optimistic:** factor ~40 increase
 - no analytic description of spectral shape at end-point
- make pile-up less important

Statistical sensitivity: shake-off processes



HOLMES simulation with the *optimistic* spectrum from A.De Rújula & M. Lusignoli

$$Q = 2833 \text{ eV}, N_{\text{ev}} = 3 \times 10^{13}, f_{\text{pp}} = 3.0 \times 10^{-4}, \Delta E = 1.0 \text{ eV}$$



$$\text{statistical sensitivity } \Sigma(m_\nu) \approx 0.64 \pm 0.03 \text{ eV}$$

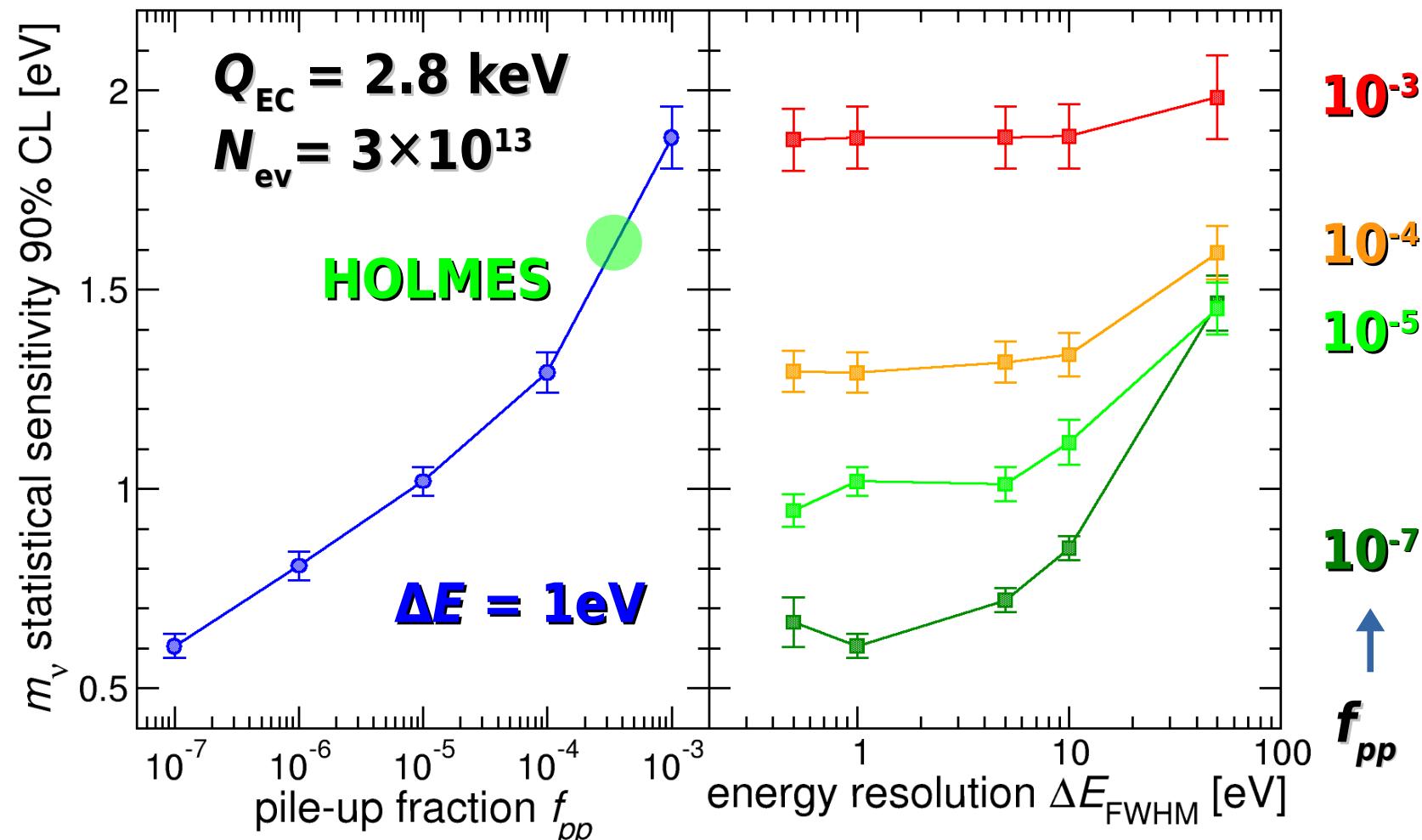
HOLMES design: more MC simulations...



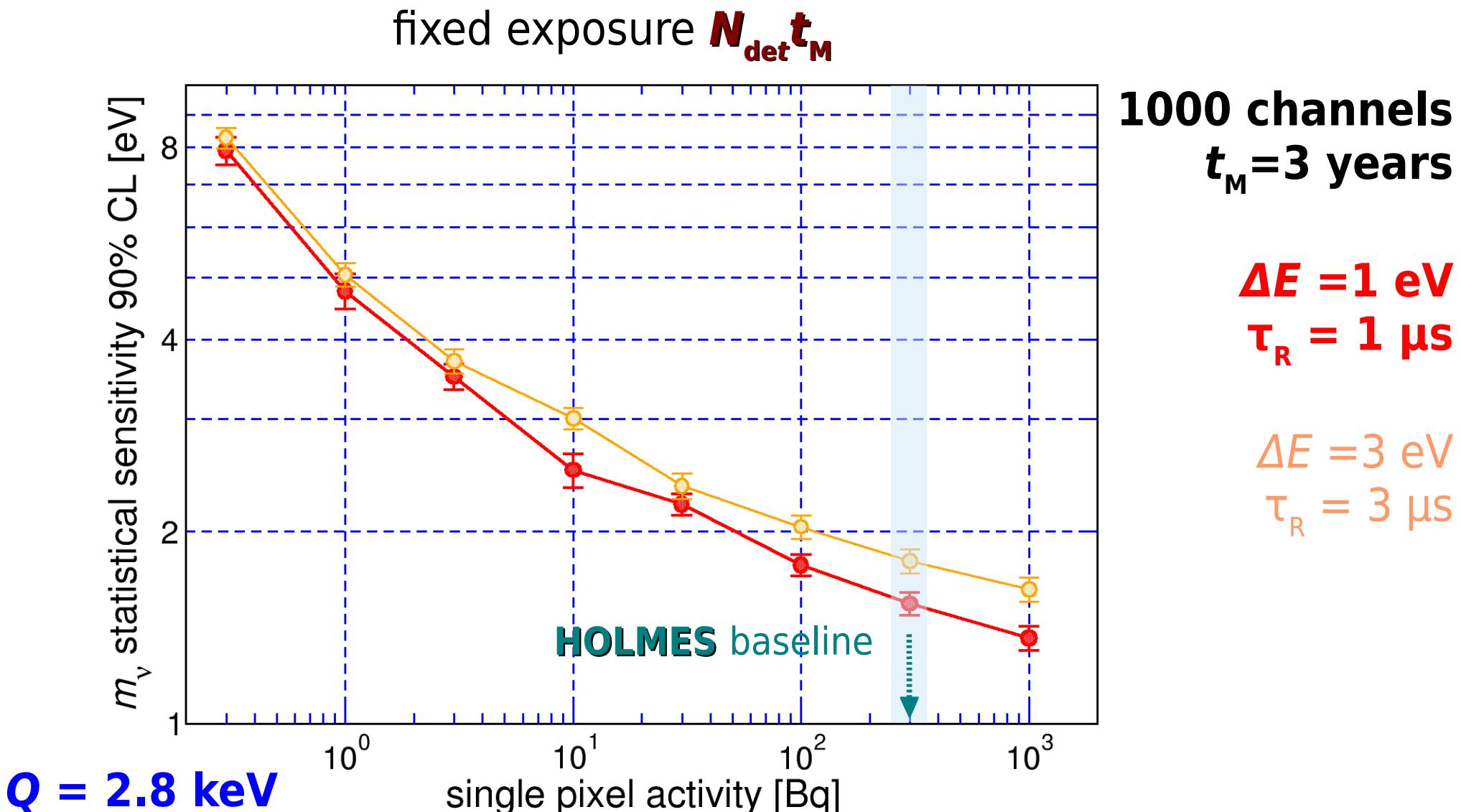
Statistical sensitivity $\Sigma(m_\nu)$ dependencies from MC simulations

- strong on statistics $N_{\text{ev}} = A_{\text{EC}} N_{\text{det}} t_M$: $\Sigma(m_\nu) \propto N_{\text{ev}}^{-1/4}$
- strong on rise time pile-up (probability $f_{pp} \approx A_{\text{EC}} \tau_R$)
- weak on energy resolution ΔE

t_M measuring time
 N_{det} number of detectors
 A_{EC} EC activity per detector
 τ_R time resolution (\approx rise time)



Statistical sensitivity and single pixel activity



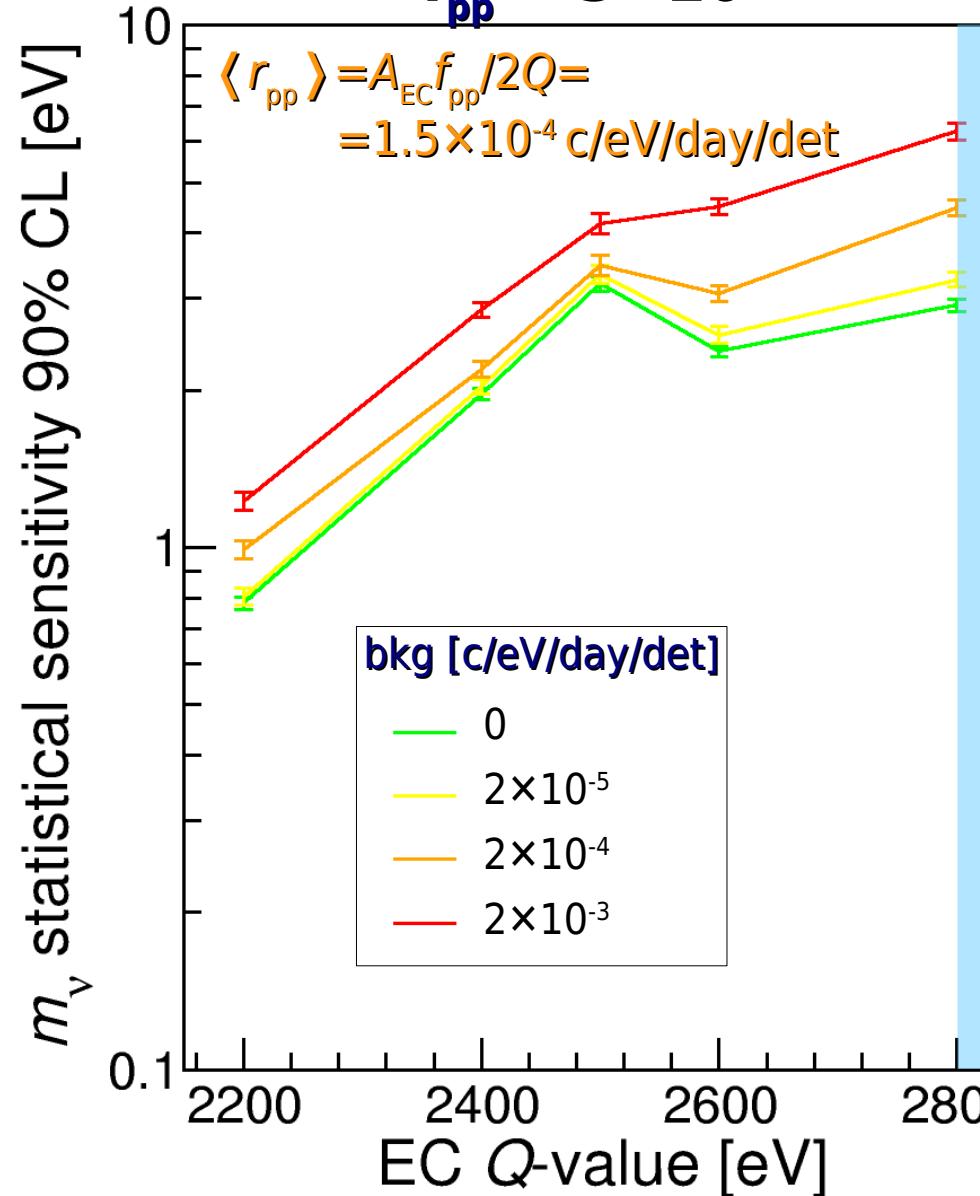
high activity \rightarrow robustness against (flat) background
 $A_{\text{EC}} = 300 \text{ Bq} \rightarrow bkg < \approx 0.1 \text{ counts/eV/day/det}$

Effect of background on sensitivity



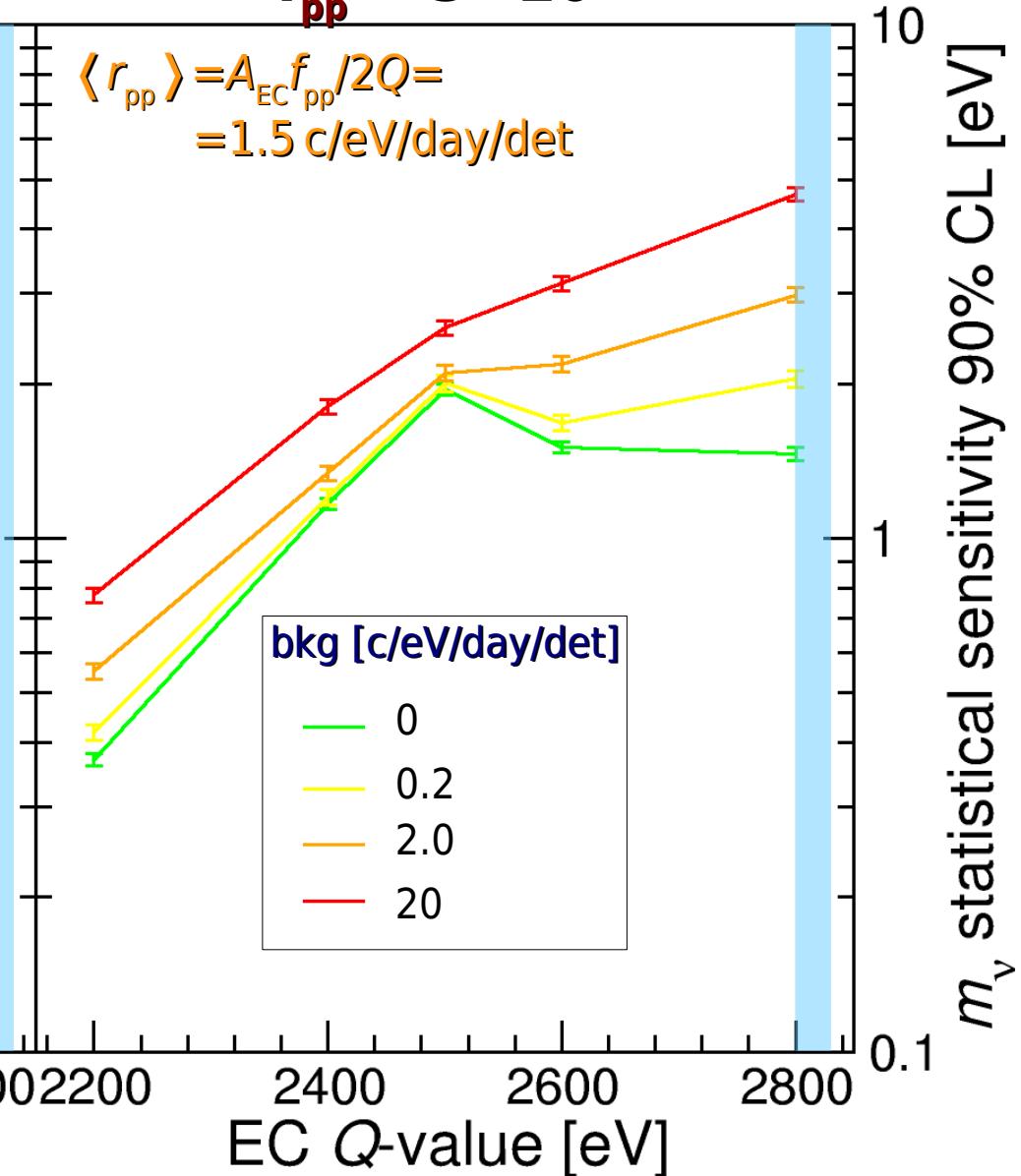
$A_{EC} = 3 \text{ Bq/det}$

$f_{pp} = 3 \times 10^{-6}$



$A_{EC} = 300 \text{ Bq/det}$

$f_{pp} = 3 \times 10^{-4}$



Low energy background sources



- environmental γ radiation
- γ , X and β from close surroundings
- **cosmic rays**
 - ▷ GEANT4 simulation for CR at sea level (only **muons**)
 - ▷ **Au pixel $200 \times 200 \times 2 \text{ } \mu\text{m}^3 \rightarrow bkg \approx 5 \times 10^{-5} \text{ c/eV/day/det (0 - 4 keV)}$**
- **internal radionuclides**
 - ▷ **^{166m}Ho (β^- , $\tau_{1/2} = 1200 \text{ y}$, produced along with ^{163}Ho)**
 - ▷ **Au pixel $200 \times 200 \times 2 \text{ } \mu\text{m}^3$**
 - GEANT4 simulation $\rightarrow bkg \approx 0.5 \text{ c/eV/day/det/Bq}({}^{166m}\text{Ho})$
 - ▷ **$A({}^{163}\text{Ho}) = 300 \text{ Bq/det}$** ($\leftrightarrow \approx 6.5 \times 10^{13} \text{ nuclei/det}$)
 - $bkg({}^{166m}\text{Ho}) < 0.1 \text{ c/eV/day/det}$** $\rightarrow A({}^{163}\text{Ho})/A({}^{166m}\text{Ho}) > 1500$
 - $\rightarrow N({}^{163}\text{Ho})/N({}^{166m}\text{Ho}) > 6000$

MIBETA experiment with $300 \times 300 \times 150 \text{ } \mu\text{m}^3 \text{ AgReO}_4$ crystals
 $bkg(2..5\text{keV}) \approx 1.5 \times 10^{-4} \text{ c/eV/day/det}$



- **^{163}Ho isotope production**
- **^{163}Ho embedding system**
- **TES pixel R&D**
- **TES array design and fabrication**
- **TES array multiplexed read-out**
- **Data Acquisition System**

^{163}Ho production by neutron activation



$^{162}\text{Er} (\text{n},\gamma) ^{163}\text{Er}$ $\sigma_{\text{thermal}} \approx 20\text{b}$
 $^{163}\text{Er} \rightarrow ^{163}\text{Ho} + \nu_e$ $\tau_{1/2}^{\text{EC}} \approx 75\text{min}$

Tm 163 1.81 h	Tm 164 5.1 m	Tm 165 30.06 h	Tm 166 7.70 h	Tm 167 9.25 d	Tm 168 93.1 d
$\epsilon; \beta^+ ...$ $\gamma 104; 69; 241; 1434; 1397 ...$	$\epsilon; \beta^+ 2.9 ...$ $\gamma 91; 1155; 769 ...$	$\epsilon; \beta^+ ...$ $\gamma 243; 47; 297; 807 ...$	$\epsilon; \beta^+ 1.9 ...$ $\gamma 779; 2052; 184; 1274 ...$	$\epsilon; \beta^+ ...$ $\gamma 532 ...$	$\epsilon; \beta^+ ...$ $\beta^- ...$ $\gamma 198; 816; 447 ...$
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
$\sigma_{n,\alpha} < 0.011$	$\epsilon; \beta^+ ...$ $\gamma (1114 ...)$	$\sigma_{n,\alpha} < 0.0012$	$\epsilon; no \gamma$	$\sigma_{n,\alpha} < 7E-5$	$\epsilon; 208$ $\sigma_{n,\alpha} 3E-6$
Ho 161 6.7 s	Ho 162 2.5 h	Ho 163 15 m	Ho 164 37 m	Ho 165 100	Ho 166 1200 a
$\epsilon; \gamma 26; 78 ...$ $\beta^-; e^-$ $\gamma 211$	$\epsilon; \beta^+ 1.1 ...$ $\gamma 185; 1220; 283; 937 ...$ e^-	$\epsilon; no \gamma$	$\epsilon; \beta^- 1.0 ...$ $\gamma 91; 73 ...$ e^-	$\sigma_{n,\alpha} < 2E-5$	β^- $\gamma 184; 810; 712$ $\sigma_{n,\alpha} 3100$
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
$\sigma_{n,\alpha} < 0.0003$	$\sigma_{n,\alpha} < 1E-6$	$\sigma_{n,\alpha} < 170$	$\sigma_{n,\alpha} < 2E-5$	$\sigma_{n,\alpha} 1610 + 1040$	β^- $\beta^- 0.9; 1.3 ...$ $\gamma 515 ... (362 ...)$ $\sigma_{n,\alpha} 3500$

- ^{162}Er irradiation at **ILL nuclear reactor** (Grenoble, France)
 - ▶ thermal neutron flux at **ILL**: $1.3 \times 10^{15} \text{ n/cm}^2/\text{s}$
 - ▶ **burn up** $^{163}\text{Ho}(\text{n},\gamma)^{164}\text{Ho}$: $\sigma_{\text{burn-up}} \approx 200\text{b}$ (preliminary result from **PSI** analysis)
 - ▶ $^{165}\text{Ho}(\text{n},\gamma)$ (mostly from $^{164}\text{Er}(\text{n},\gamma)$) $\rightarrow ^{166\text{m}}\text{Ho}$ (β , $\tau_{1/2} = 1200\text{y}$) $\rightarrow A(^{163}\text{Ho})/A(^{166\text{m}}\text{Ho}) = 100 \sim 1000$
- chemical pre-purification and post-separation at **PSI** (Villigen, CH)
- **HOLMES needs $\approx 200 \text{ MBq}$ of ^{163}Ho**
 - with reasonable assumptions on the (unknown) global embedding process efficiency...

HOLMES source production



- **enriched Er_2O_3** samples* irradiated at **ILL** and pre-/post-processed at **PSI**
 - ▶ 25 mg irradiated for 55 days (2014) → $A(\text{Ho}^{163}) \approx 5 \text{ MBq}$ ($A(\text{Ho}^{166m}) \approx 10 \text{ kBq}$)
 - ▶ 150 mg irradiated for 50 days (2015) → $A(\text{Ho}^{163}) \approx 38 \text{ MBq}$ ($A(\text{Ho}^{166m}) \approx 37 \text{ kBq}$)
- **Ho chemical separation** with ion-exchange resins in hot-cell at **PSI**
 - ▶ efficiency $\geq 79\%$ (preliminary)
- **540 mg of 25% enriched Er_2O_3** irradiated 50 days at **ILL** early in 2017
 - ▶ $A(\text{Ho}^{163})_{\text{theo}} \approx 130 \text{ MBq}$ (enough for R&D and 500 pixels) ($A(\text{Ho}^{166m}) \approx 180 \text{ kBq}$)



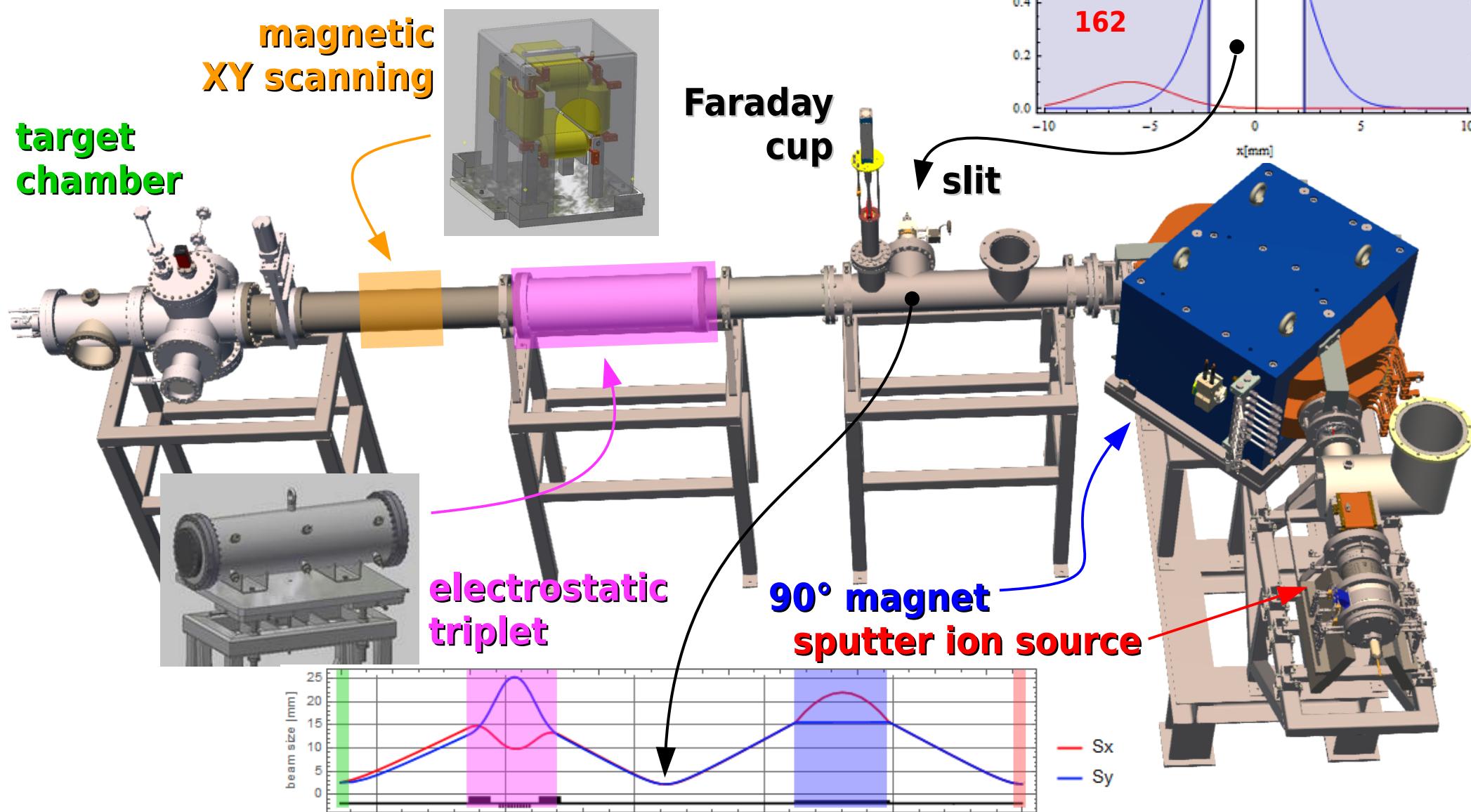
* from INFN and CENTRA (Lisbon)

A. Nucciotti, PSI, Villigen (CH), May 23rd, 2017

HOLMES mass separation and ion implantation



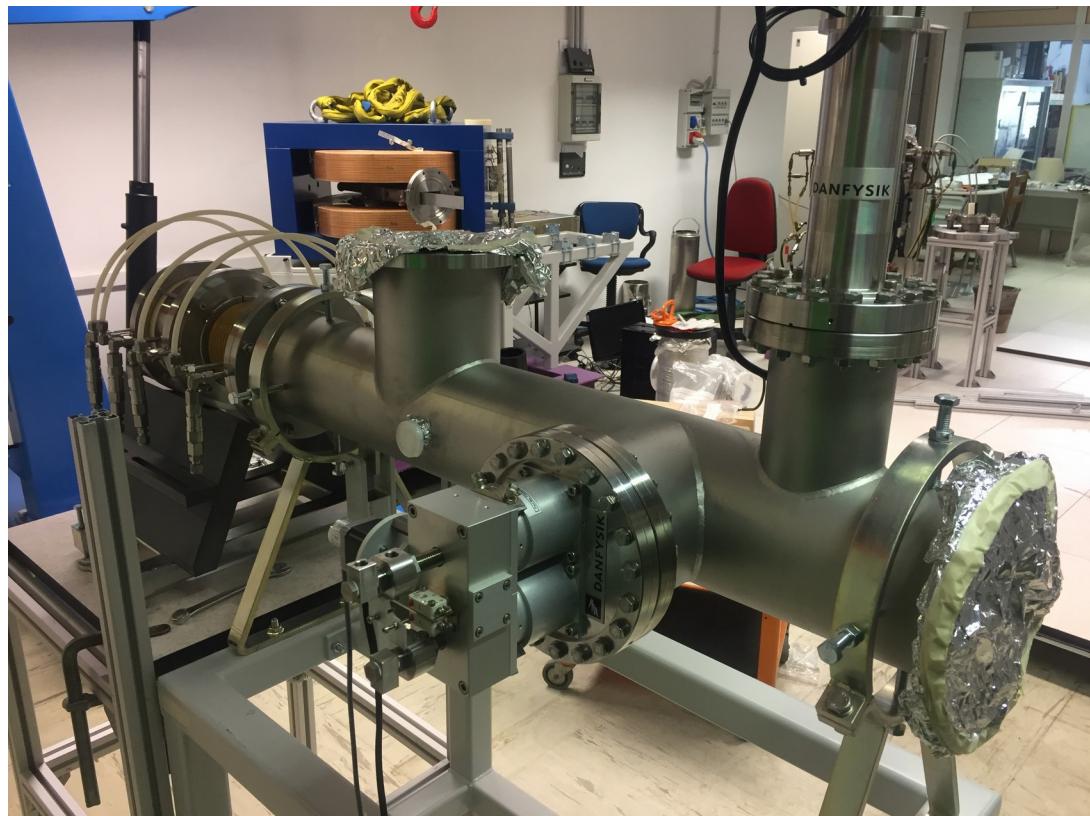
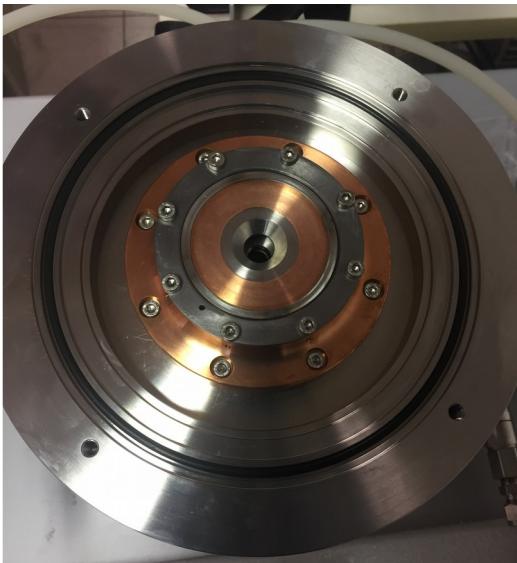
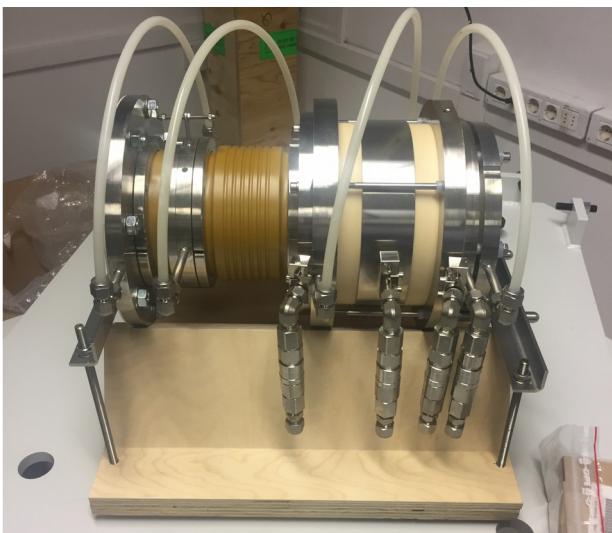
- extraction voltage 30-50 kV → 10-100 nm implant depth
- ^{163}Ho / $^{166\text{m}}\text{Ho}$ separation better than 10^5
- **ion source, magnet** and **slit** delivered end **2016**



HOLMES ion implantation system / 2



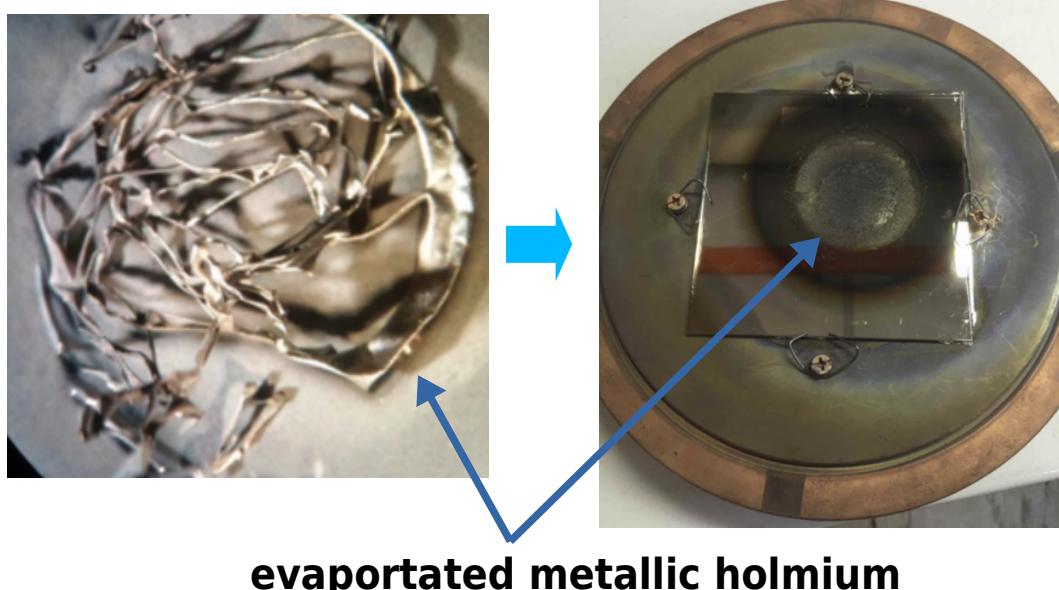
testing the ion source



Ion source sputter target production / 1



- Metallic holmium sputter target for implanter ion source
- 30% enriched $\text{Er}_2\text{O}_3 \rightarrow \text{Ho}_2\text{O}_3$
- thermoreduction/distillation in furnace
 - ▶ $\text{Ho}_2\text{O}_3 + 2\text{Y(met)} \rightarrow 2\text{Ho(met)} + \text{Y}_2\text{O}_3$ at $T > 1600^\circ\text{C}$
- new furnace set-up in 2016
- work in progress to
 - ▶ optimize the process
 - ▶ measure efficiency ($\approx 70\%$, preliminary)



evaporated metallic holmium

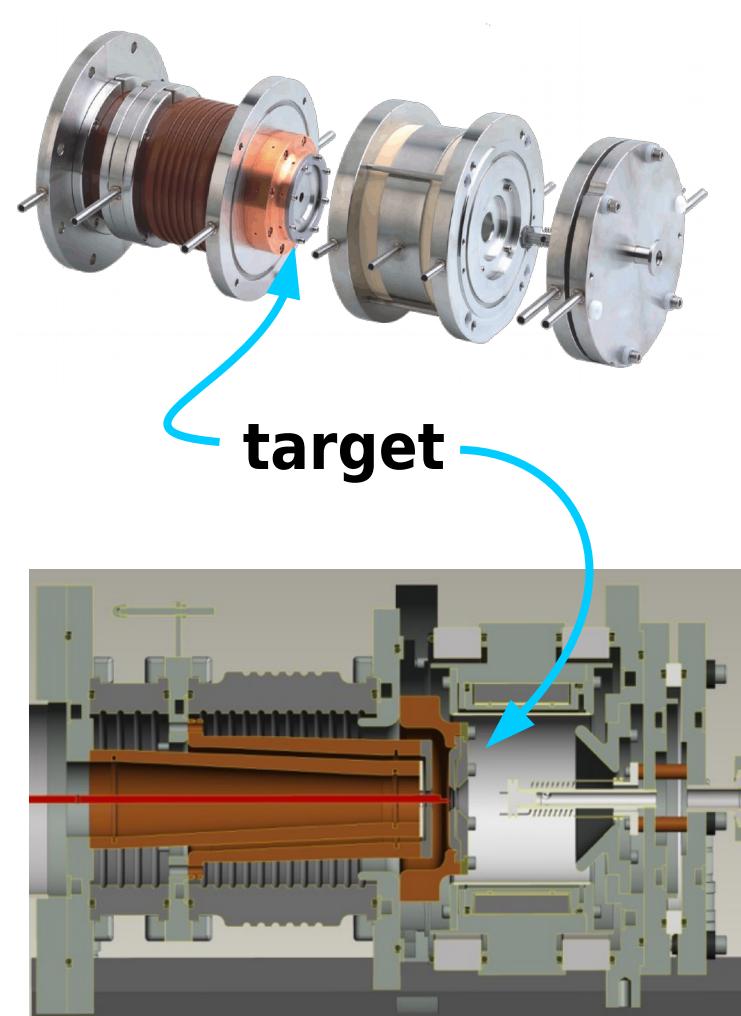


Ion source sputter target production / 2



■ Metallic holmium sputter target for implanter ion source

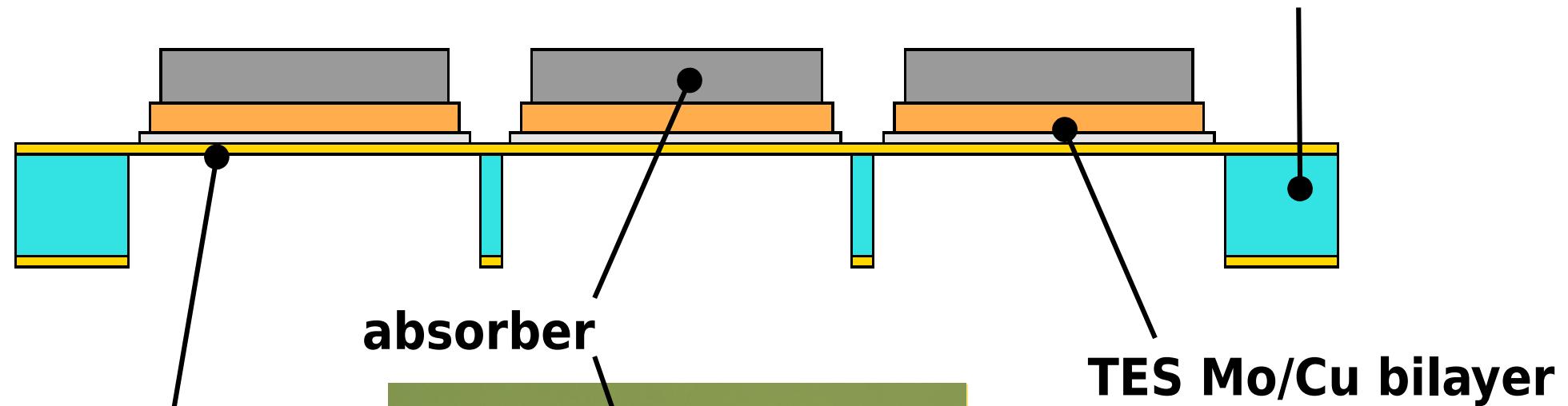
- ▶ work is in progress to produce the sputter target
- ▶ sintering Ho with other metals



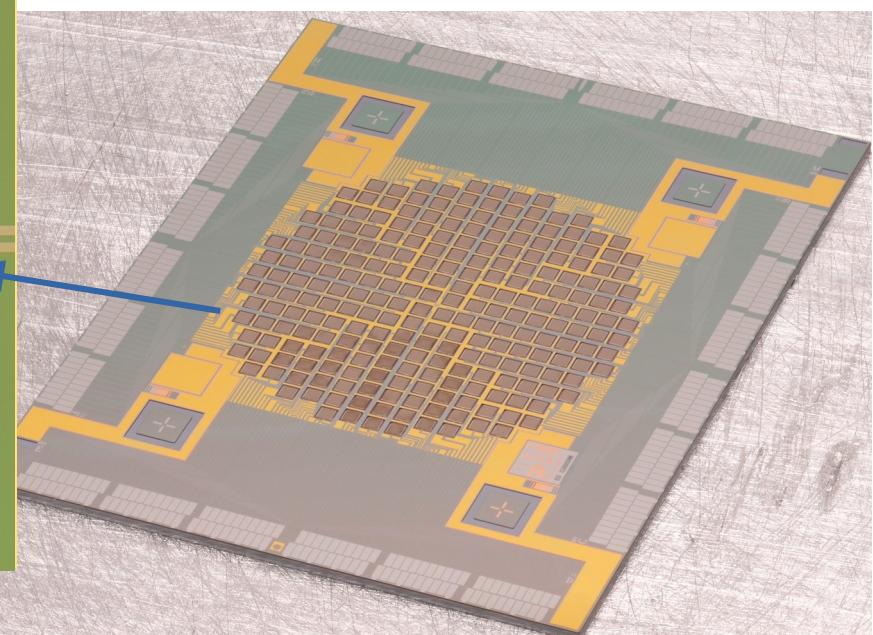
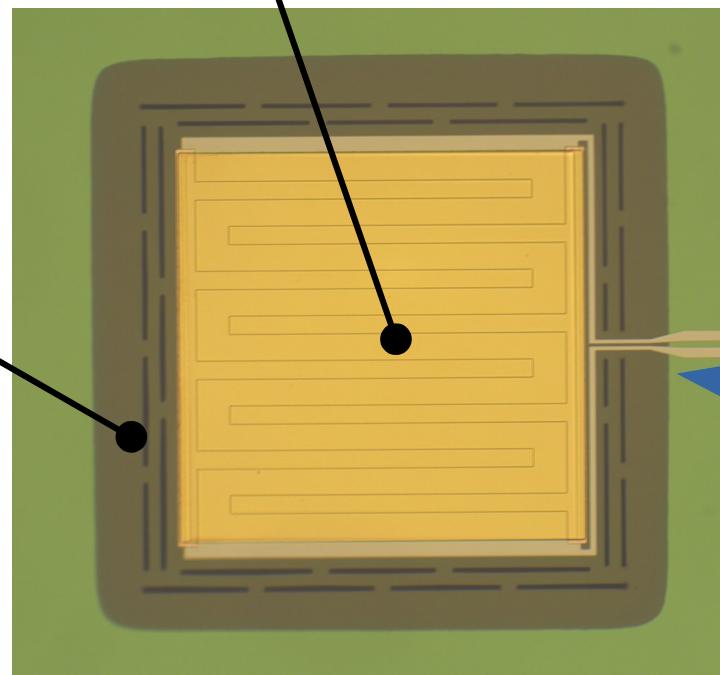
TES microcalorimeters



**micromachined
silicon substrate**



SiN_x membrane

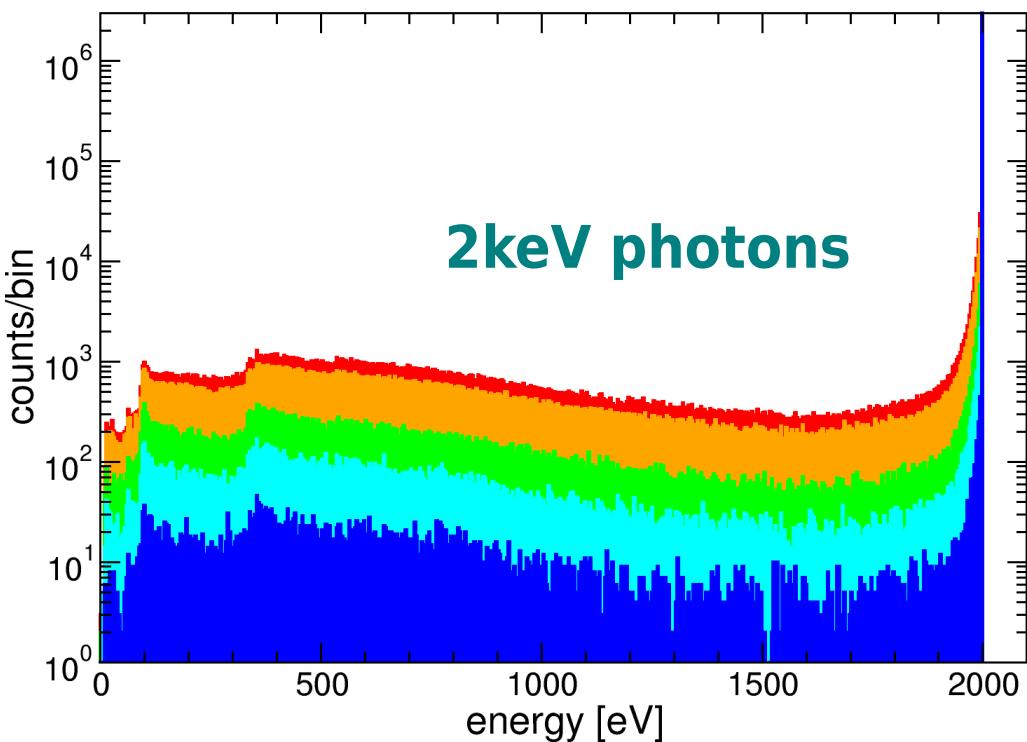
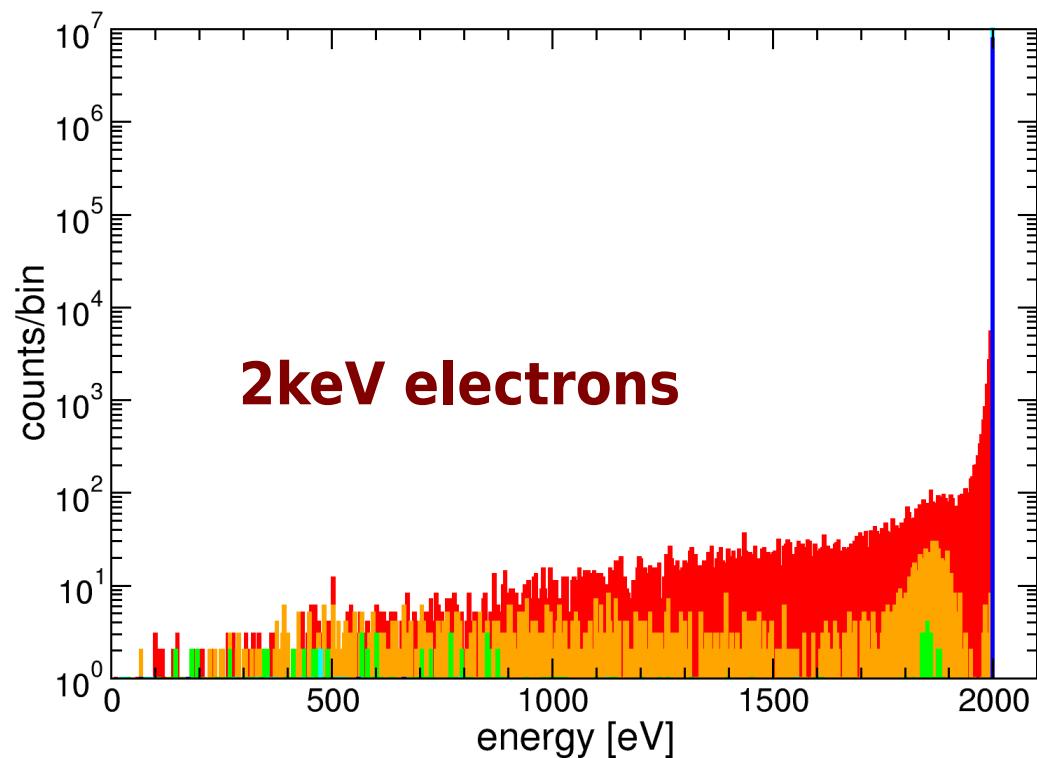
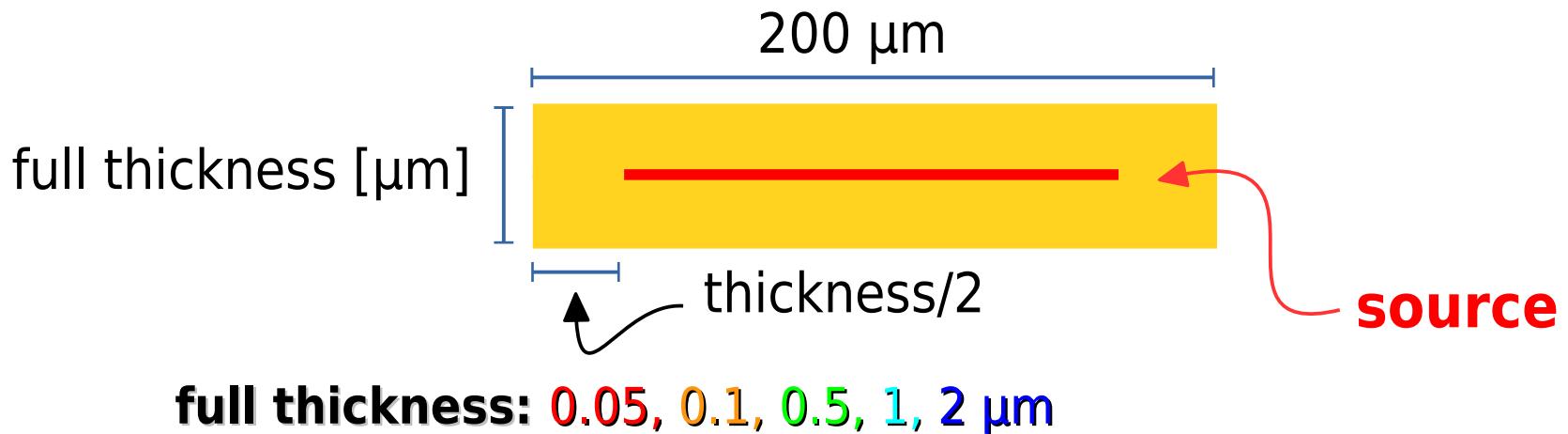


NIST TES array for X-ray spectroscopy

TES absorber design: stopping EC radiation / 1



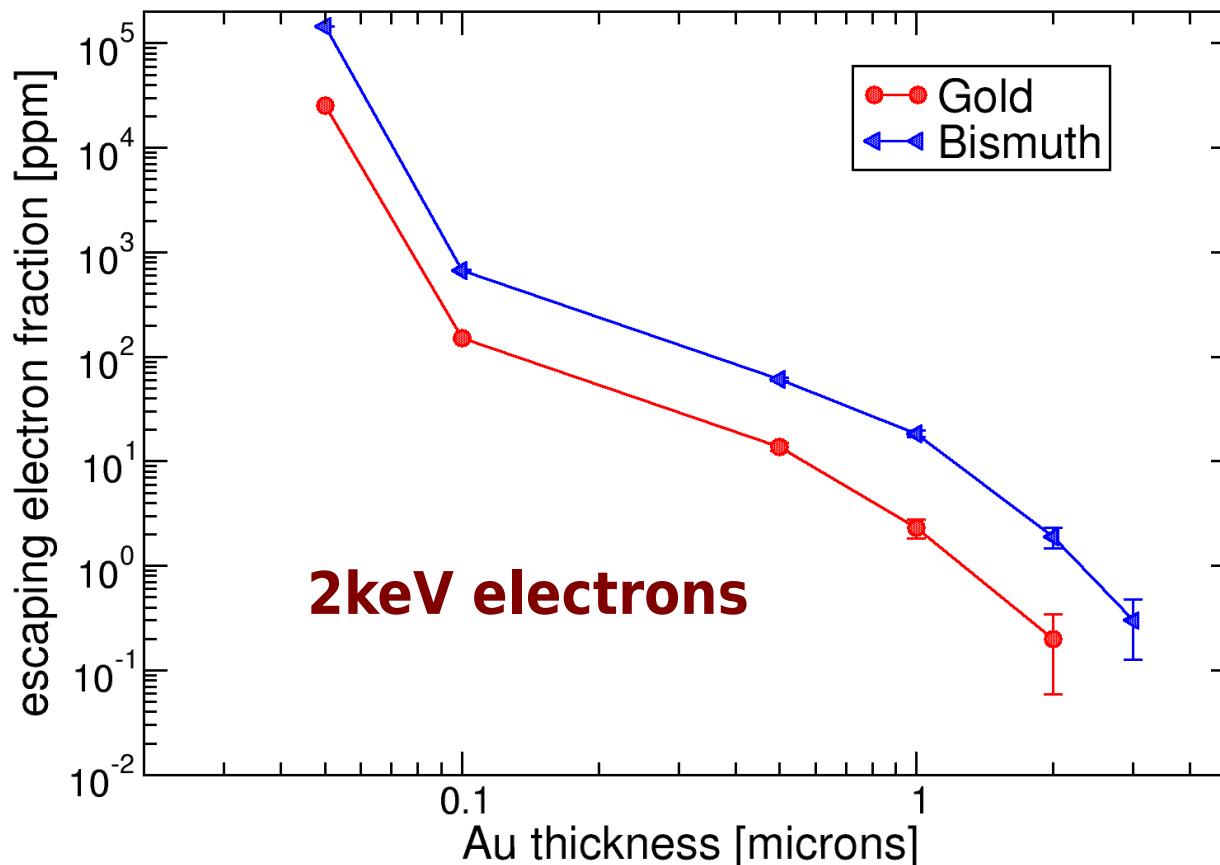
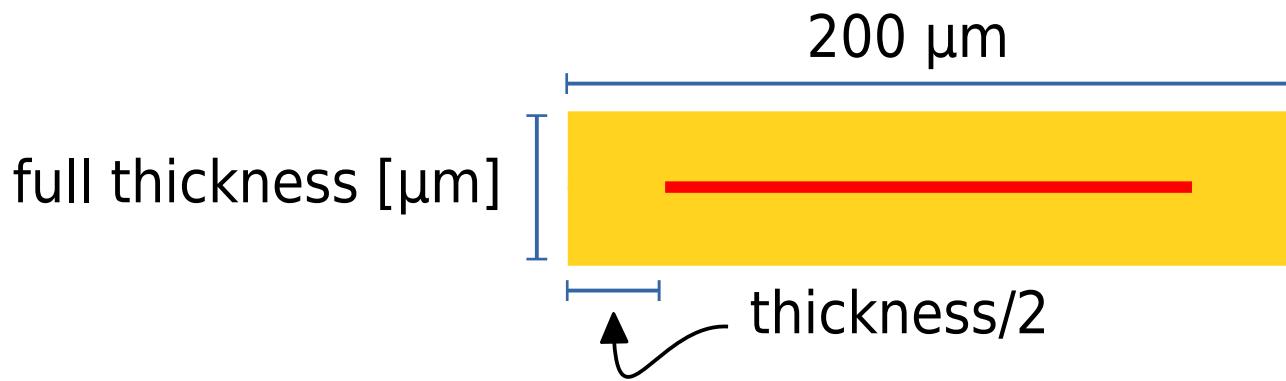
Geant4 + LowEnergyEM MC simulation



TES absorber design: stopping EC radiation / 2



Geant4 + LowEnergyEM MC simulation



Multiplexed TES array read-out

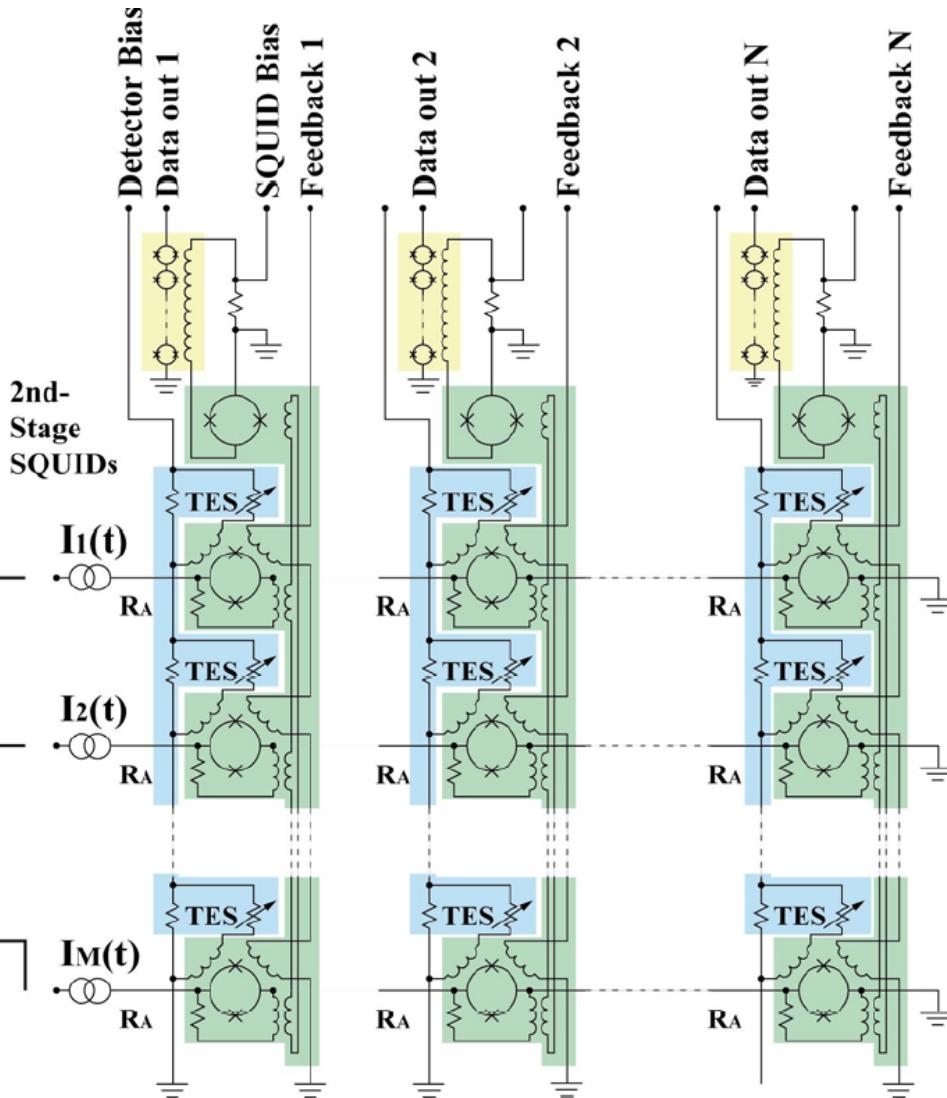
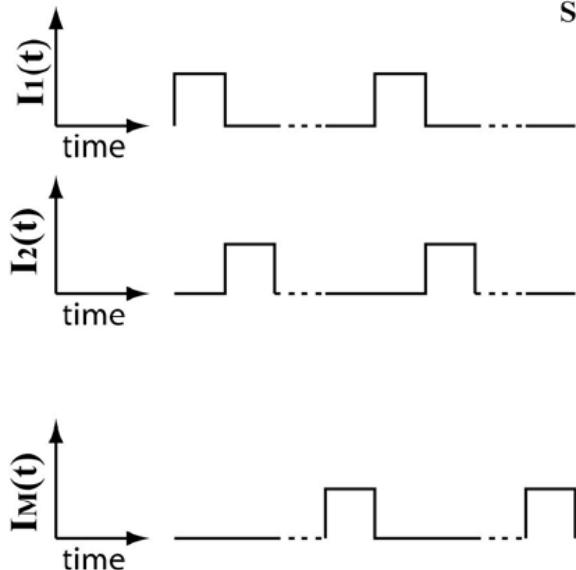


TDM Time Division Multiplexing

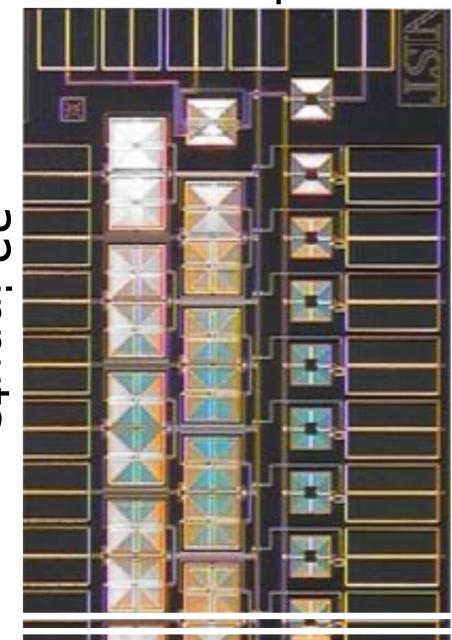
$N \times N$ TES array

without *mux* $\rightarrow N \times N$ readout channels
with *mux* $\rightarrow N$ addresses + N outputs

Boxcar Modulation Functions



column output + ...

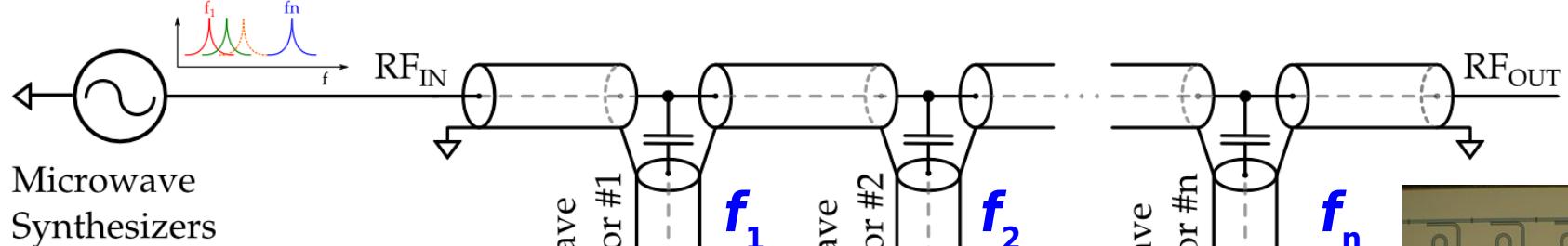


Series Array SQUID (4K)

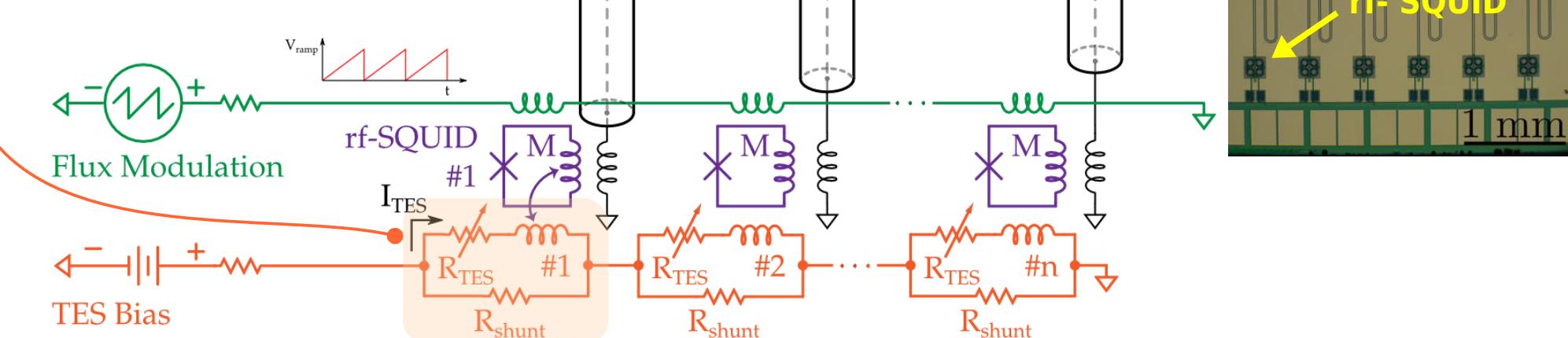
Multiplexer chip(s)

Detector chip(s)

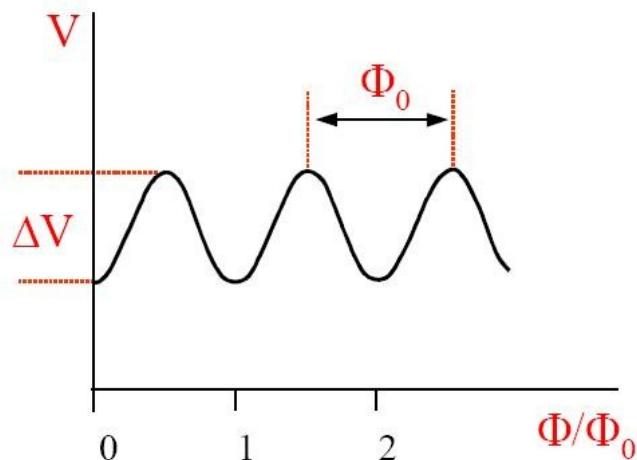
HOLMES array read-out: rf-SQUID μwave mux



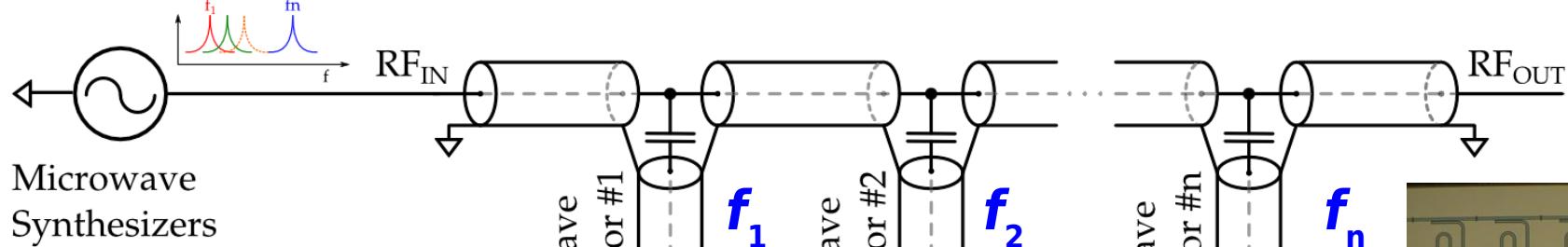
dc biased TES sensor



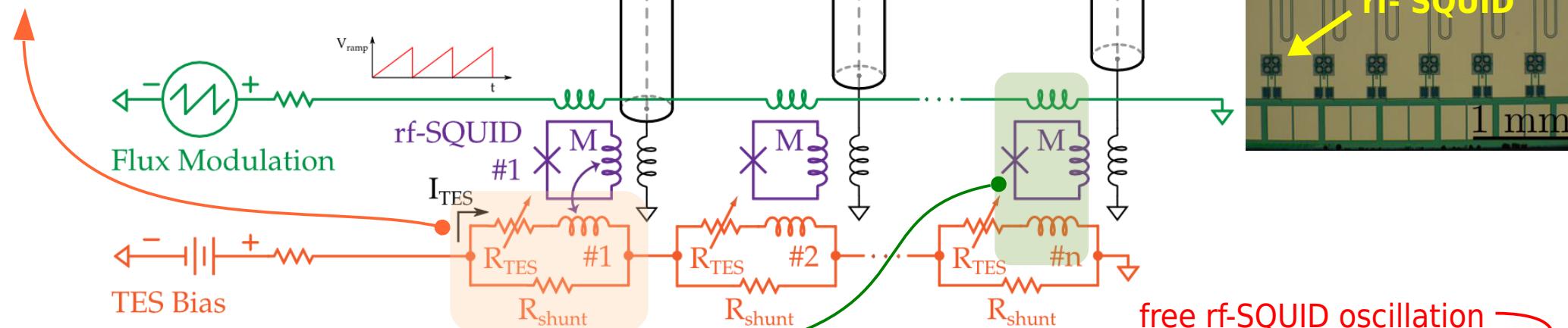
periodic
SQUID
response



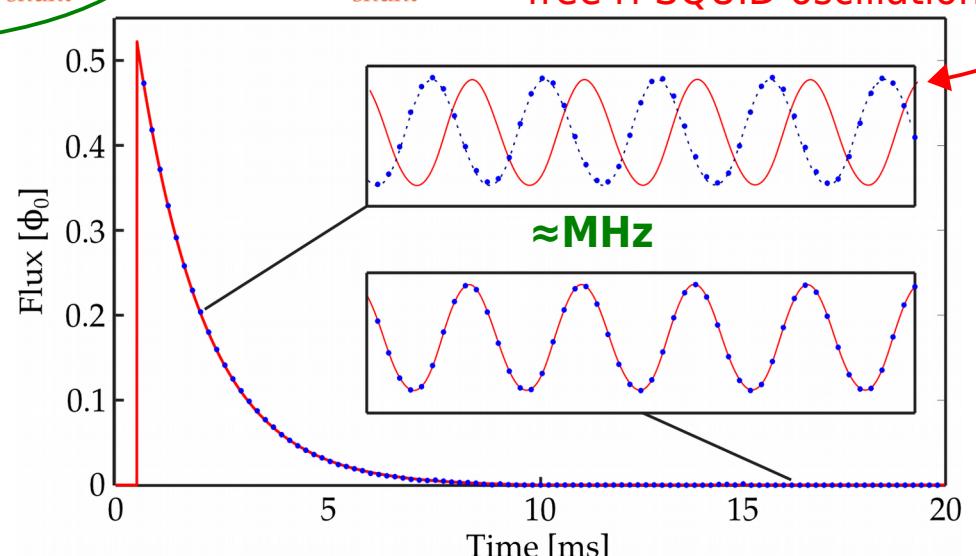
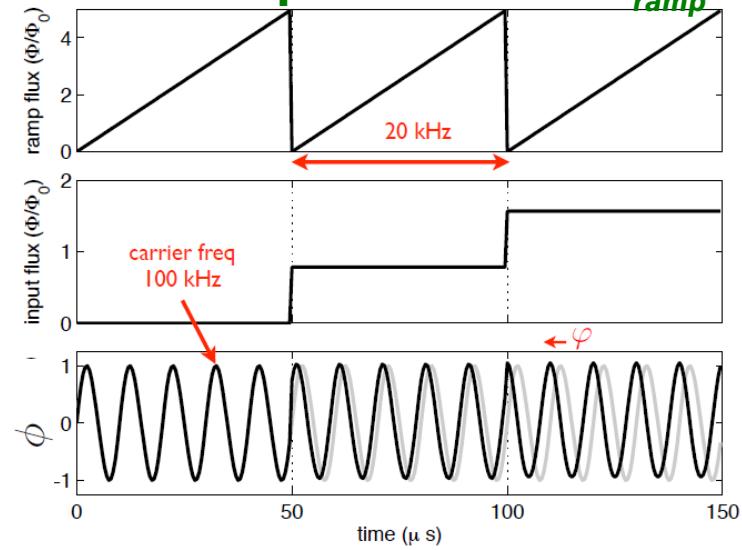
HOLMES array read-out: rf-SQUID μwave mux



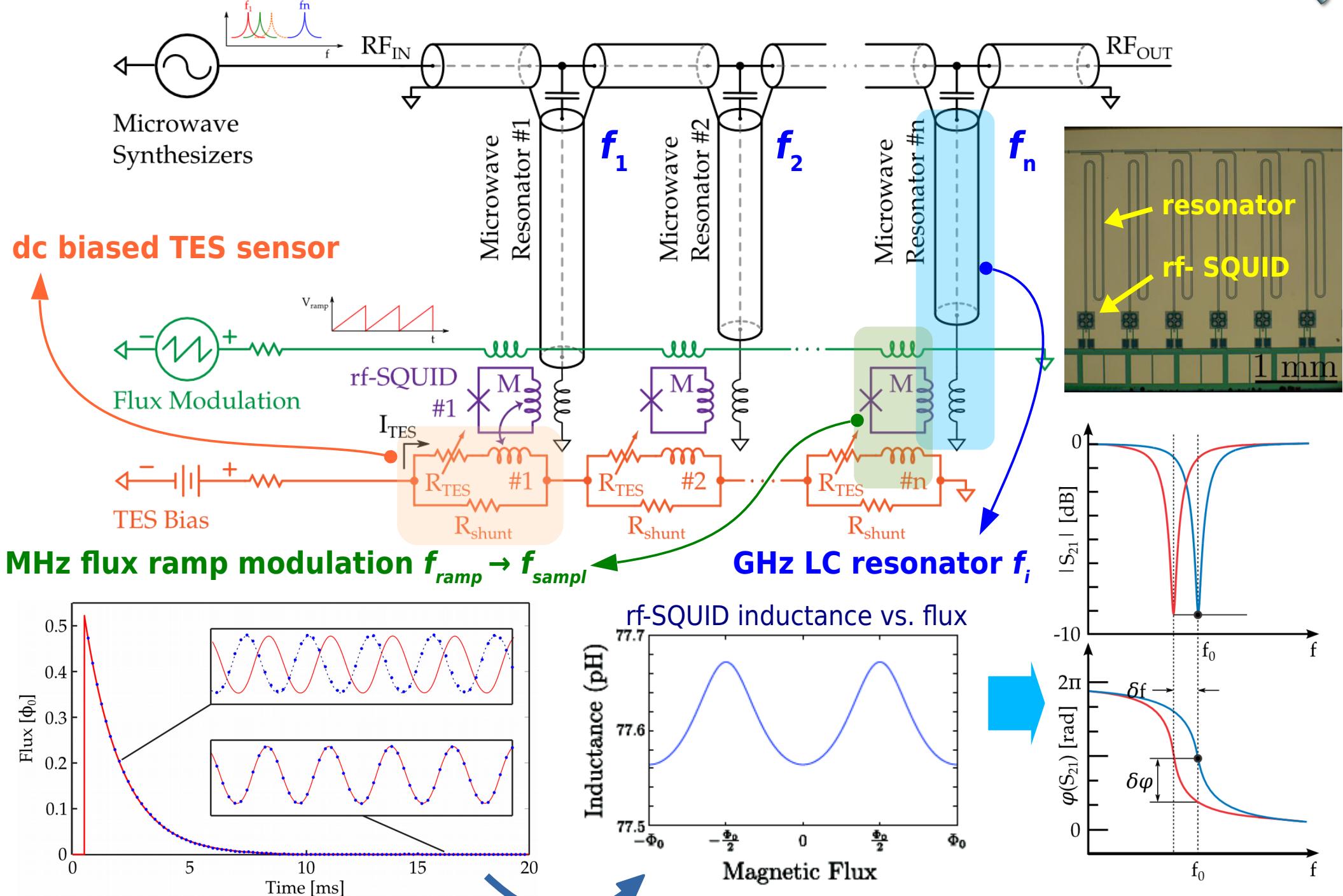
dc biased TES sensor



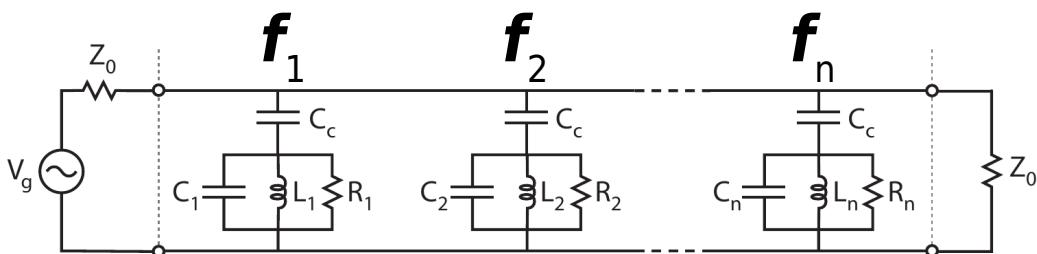
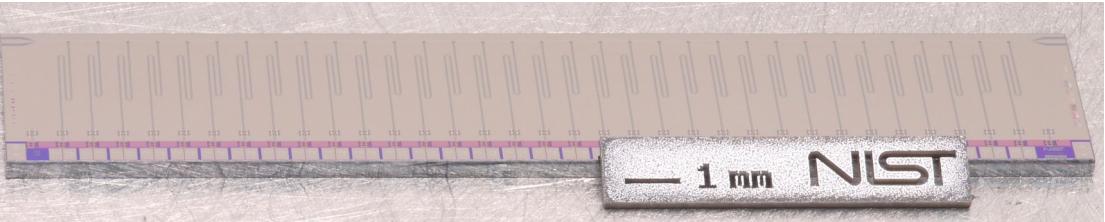
MHz flux ramp modulation $f_{ramp} \rightarrow f_{sample}$



HOLMES array read-out: rf-SQUID μwave mux

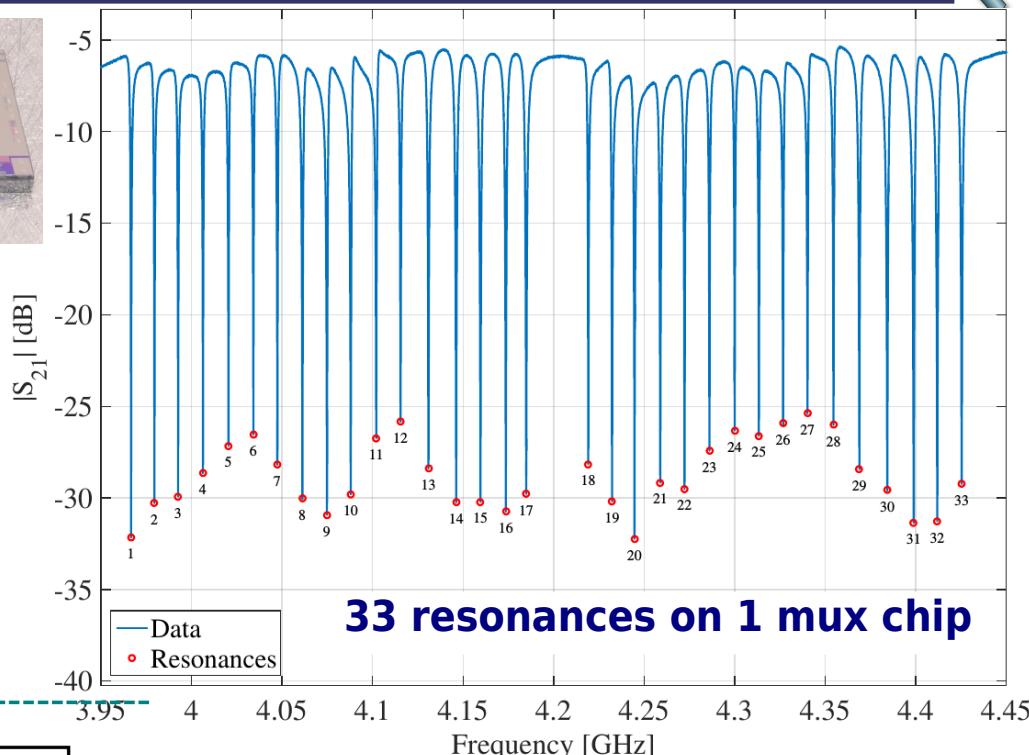
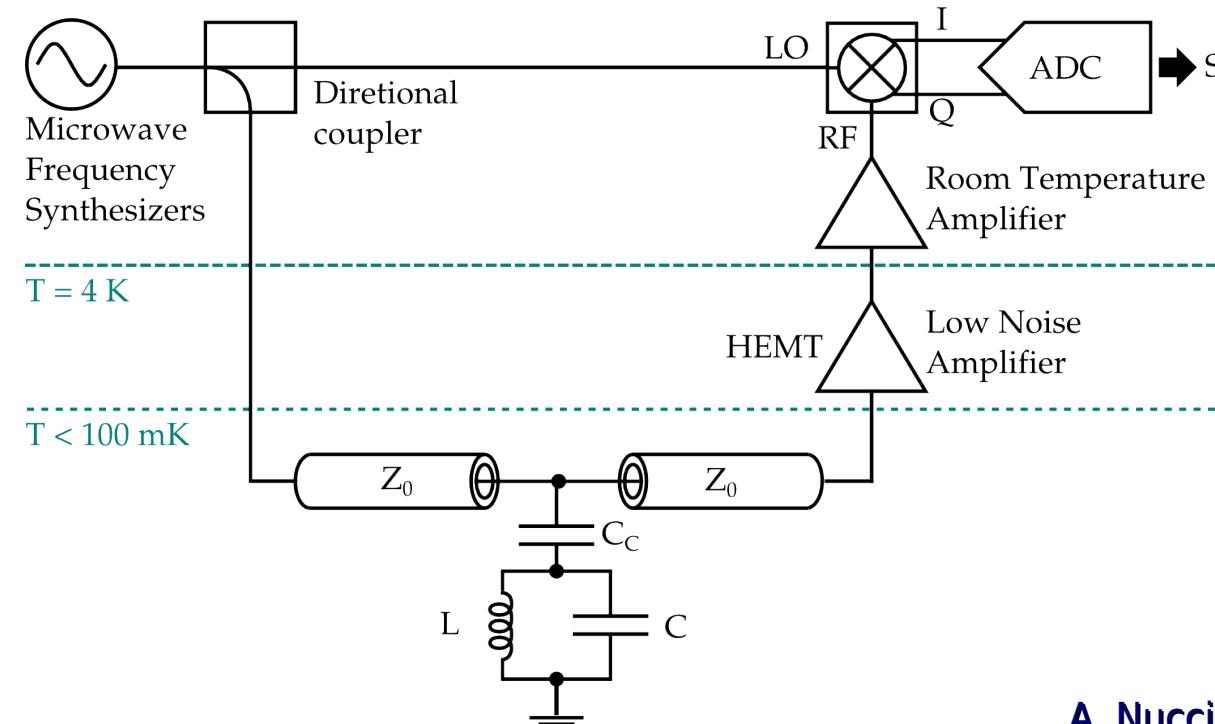


μ wave with RF carrier homodyne read-out

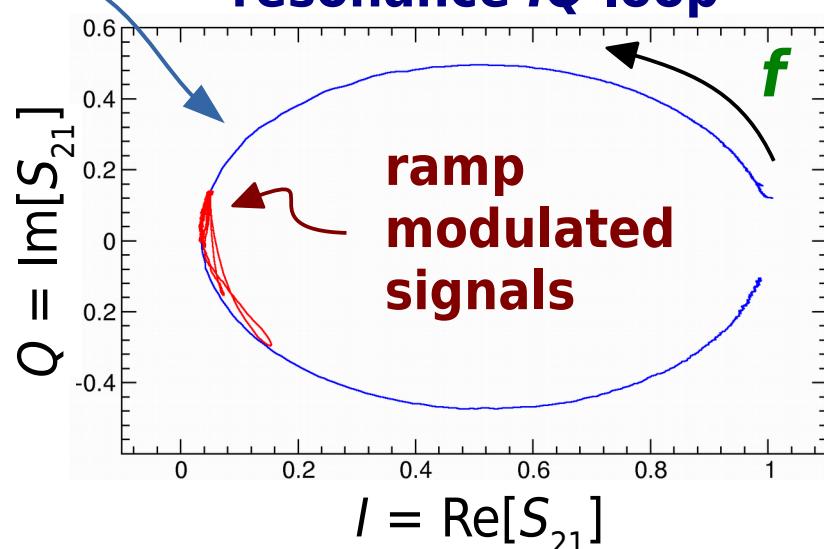


device i is read out tuning the RF carrier to f_i

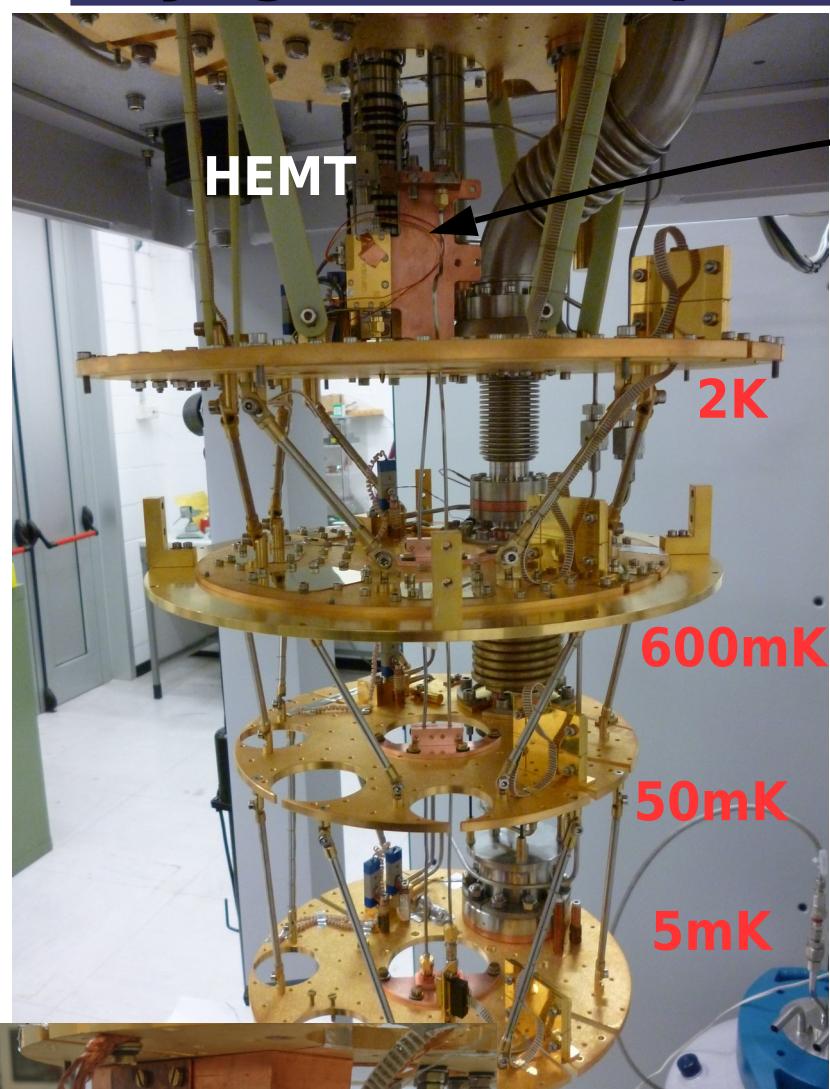
$T = 300 \text{ K}$



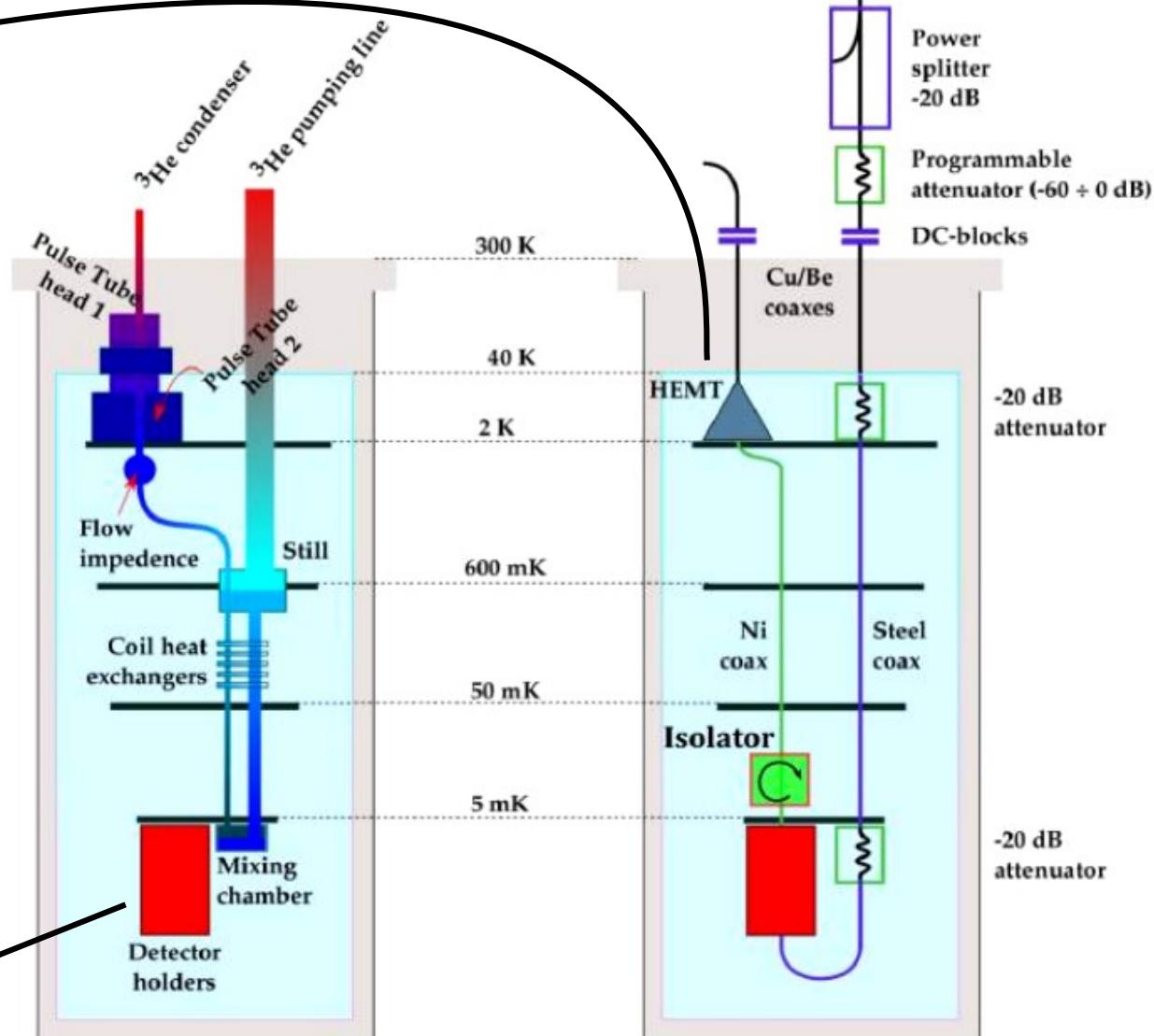
resonance IQ loop



Cryogenic set-up

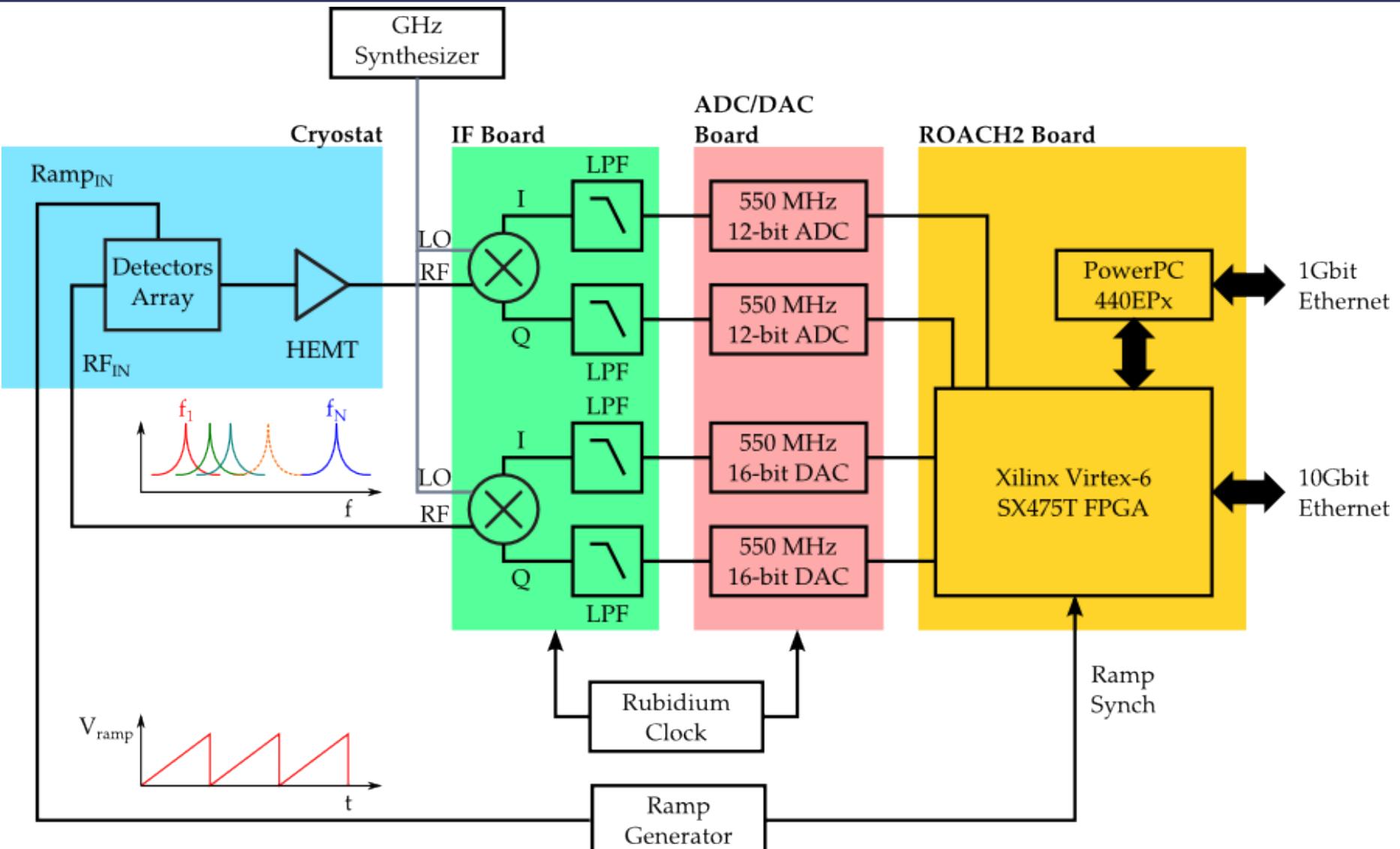


LHe-free dilution refrigerator



detector holder

HOLMES DAQ: Software Defined Radio



multiplexing factor n_{TES}

f_{BW} required bandwidth per channel $\approx 1/\tau_{rise} \rightarrow$

$$n_{TES} \approx \frac{f_{ADC}}{10 f_{BW}}$$

Detector time resolution

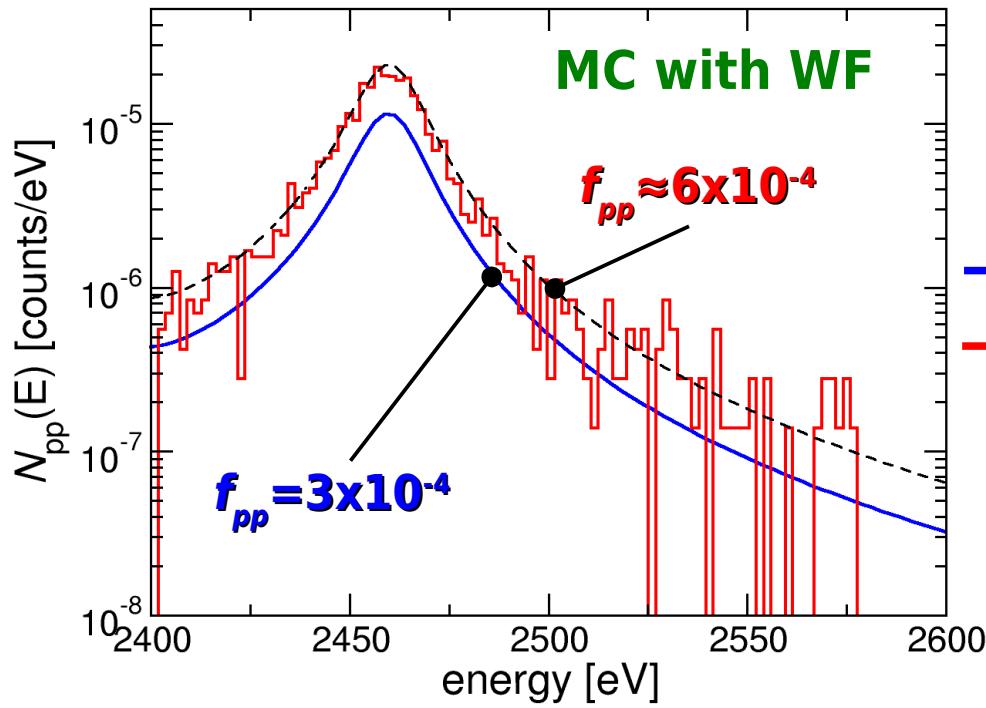


- for subsequent (Δt) events with energy E_1 and E_2 : time resolution $\tau_R = \tau_R(E_1, E_2)$

$$N_{pp}(E) = A_{EC} \int_0^{\infty} \tau_R(E, \epsilon) N_{EC}(\epsilon) N_{EC}(E - \epsilon) d\epsilon$$

- Montecarlo pile-up spectrum simulations

- event pairs with $E_1 + E_2 \in [2.4 \text{ keV}, 2.6 \text{ keV}]$ (drawn from ^{163}Ho spectrum), $\Delta t \in [0, 16\mu\text{s}]$
- pulse shape and noise from NIST TES model, sampled with f_{samp} , record length, and n bit
- process with pile-up detection algorithms:
 - Wiener Filter WF or Single Value Decomposition SVD**
- evaluate **effective time resolution τ_{eff}** from **pile-up detection efficiency $\eta(\Delta t)$**



$$f_{pp} = A_{EC} \Delta t_{max} \left[1 - \int_0^{\Delta t_{max}} \frac{\eta(x)}{\Delta t_{max}} dx \right] = A_{EC} \tau_{\text{eff}}$$

— $f_{pp} N_{EC}(E) \otimes N_{EC}(E)$ with $A = 300 \text{ Bq}$ and $\tau_R = 1 \mu\text{s}$
— WF simulation with $f_{\text{samp}} = 1 \text{ MHz}$, $\tau_{\text{rise}} \approx 10 \mu\text{s}$, and $A_{EC} = 300 \text{ Bq}$

best time resolution

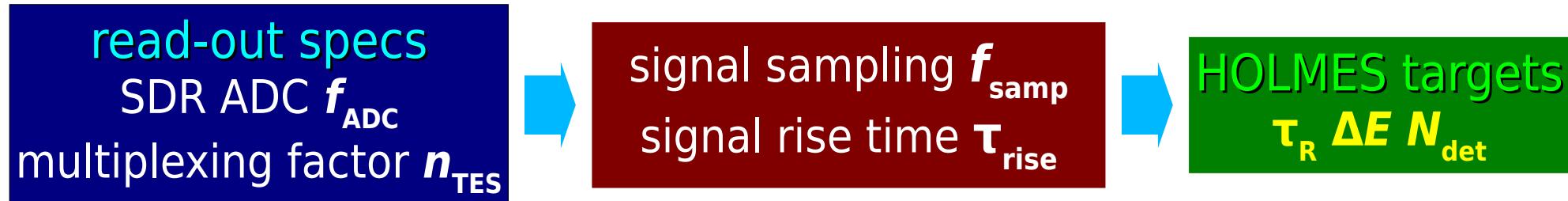
SVD ($f_{\text{samp}} = 0.5 \text{ MHz}$, $\tau_{\text{rise}} \approx 10 \mu\text{s}$) $\rightarrow \tau_{\text{eff}} \approx 1 \mu\text{s}$

HOLMES detector design



design mostly driven by **read-out bandwidth** requirements

- TES microwave multiplexing with rf-SQUID ramp modulation + Software Defined Radio (SDR)



$$f_{\text{samp}} \geq \frac{R_d}{\tau_{\text{rise}}} \approx \frac{5}{\tau_{\text{rise}}} \quad \text{detector signal sampling (signal BW)}$$

$$f_{\text{res}} \geq 2n_{\Phi_0}f_{\text{samp}} \quad \text{flux ramp modulated signal BW (resonator BW)}$$

$$f_n \geq g_f f_{\text{res}} = \frac{2R_d g_f n_{\Phi_0}}{\tau_{\text{rise}}} \quad \text{microwave tones separation } (g_f \gtrsim 10)$$

multiplexing factor

$$n_{\text{TES}} = \frac{f_{\text{ADC}}}{f_n} \leq \frac{f_{\text{ADC}} \tau_{\text{rise}}}{2 R_d g_f n_{\Phi_0}} \approx \frac{f_{\text{ADC}} \tau_{\text{rise}}}{200}$$

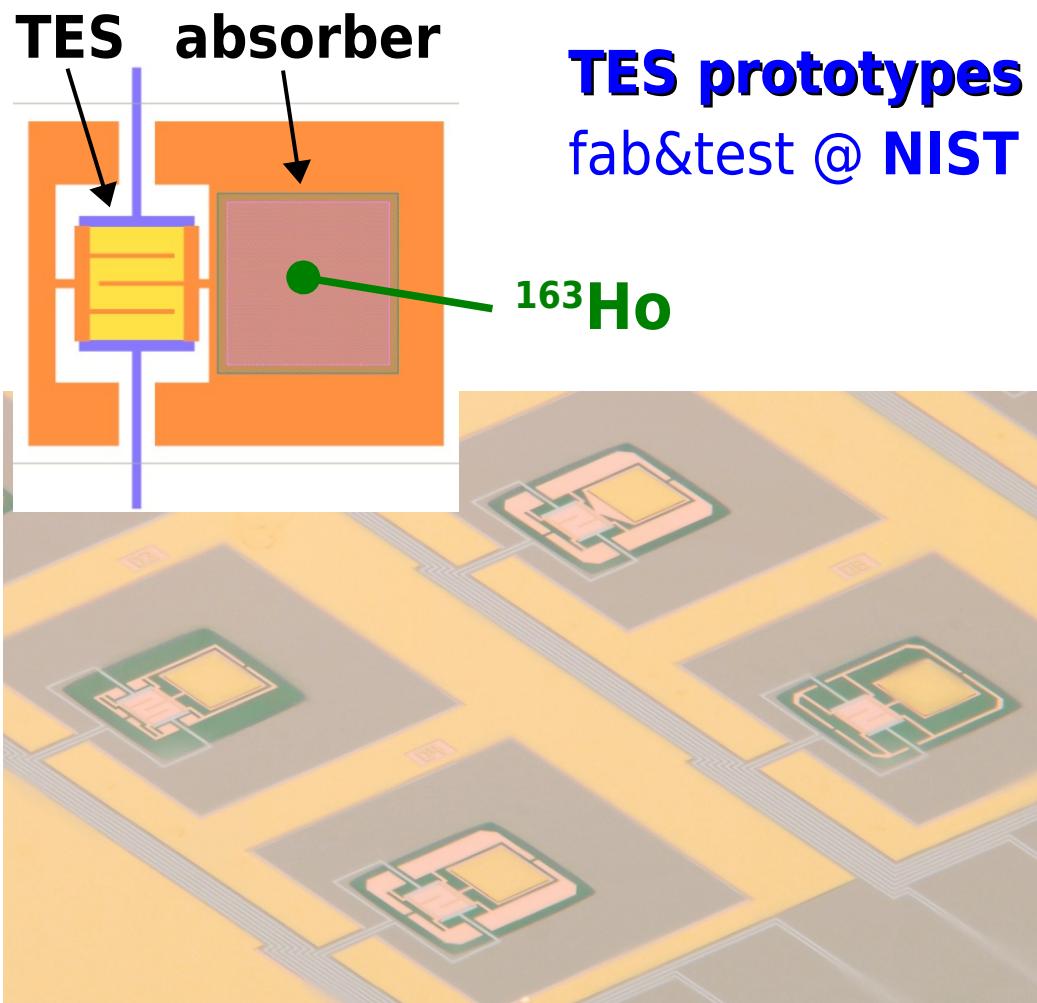
for fixed $f_{\text{ADC}} = 550 \text{MHz}$ and $n_{\text{TES}} \approx 30 \leftrightarrow \tau_{\text{rise}} \approx 10 \mu\text{s}$ with $f_{\text{samp}} = 0.5 \text{MHz}$

→ check for τ_R and $\Delta E...$

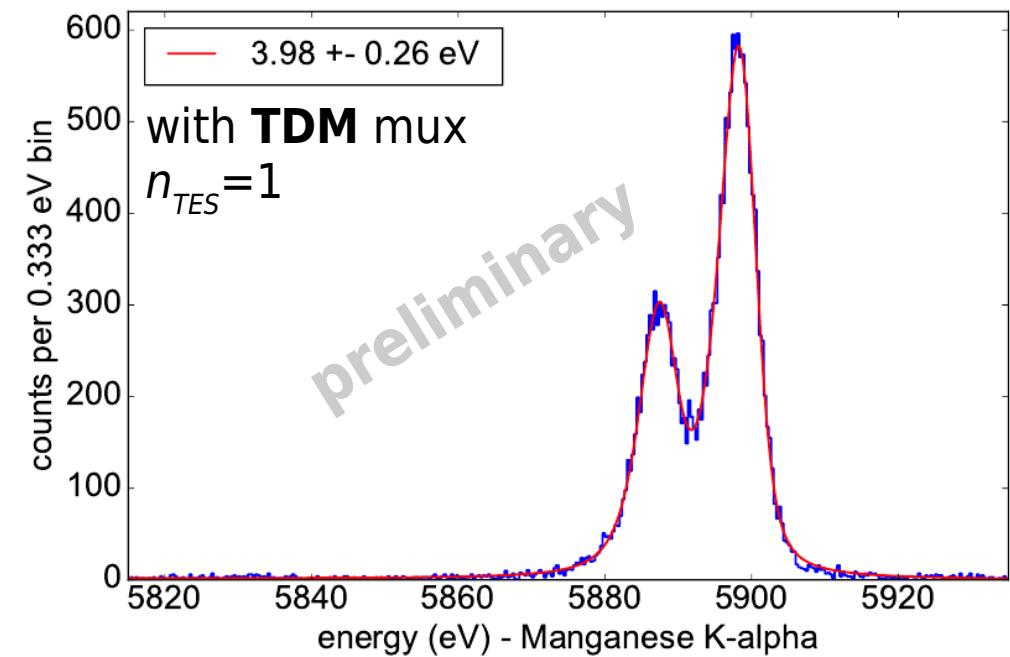
HOLMES pixel design and test



- optimize design for speed and resolution
 - ▷ specs @3keV : $\Delta E_{FWHM} \approx 1\text{eV}$, $\tau_{rise} \approx 10\mu\text{s}$, $\tau_{decay} \approx 100\mu\text{s}$
- **2 μm Au** thickness for *full* electron and photon absorption
 - ▷ GEANT4 simulation: 99.99998% / 99.927% full stopping for 2 keV electrons / photons
- **side-car** design to avoid TES proximitation and G engineering for τ_{decay} control

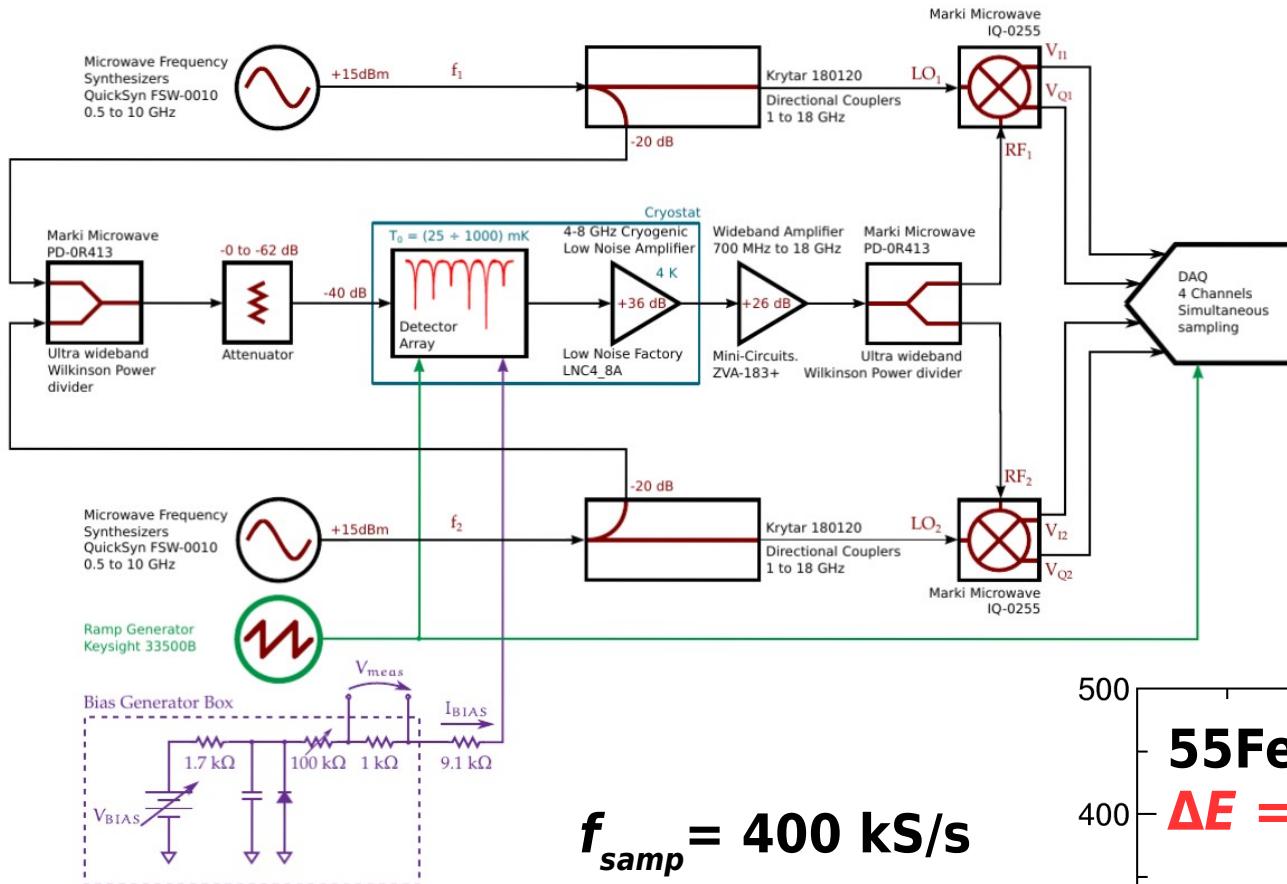


TES prototypes
fab&test @ **NIST**



- ▷ $\Delta E_{FWHM} \lesssim 4 \text{ eV}$ @ 6 keV ($\rightarrow \approx 3 \text{ eV}$ @ Q_{EC})
- ▷ $\tau_{rise} \approx 6 \mu\text{s}$ (with $L=38\text{nH}$ \rightarrow to be slowed)
- ▷ $\tau_{decay} \approx 130 \mu\text{s}$ (still tunable)

Detector testing with homodyne read-out

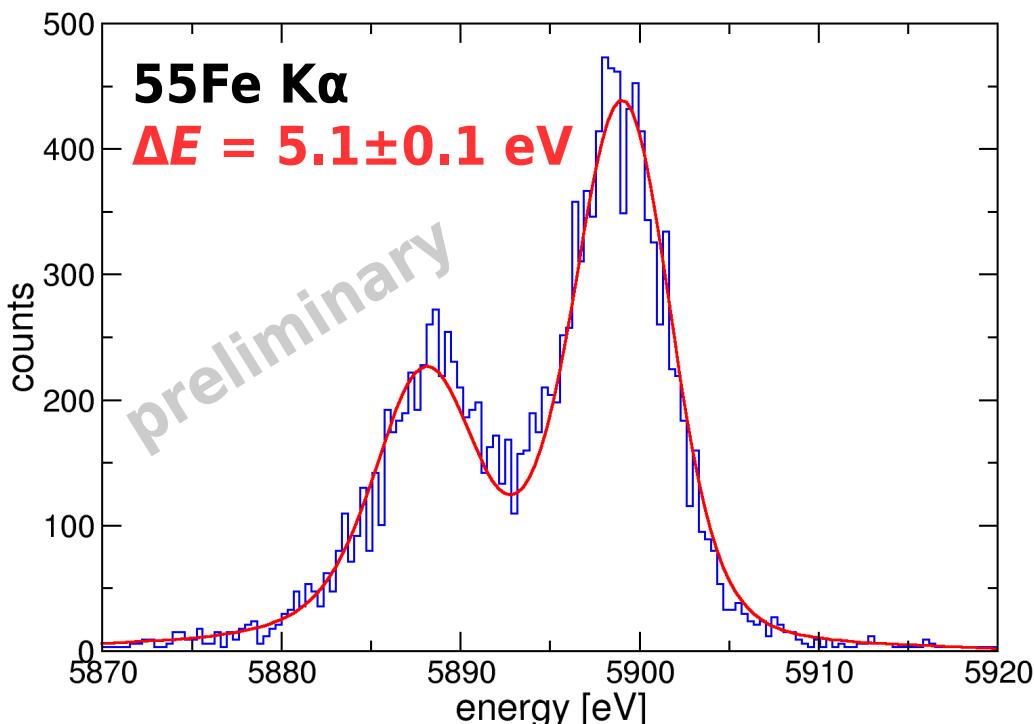
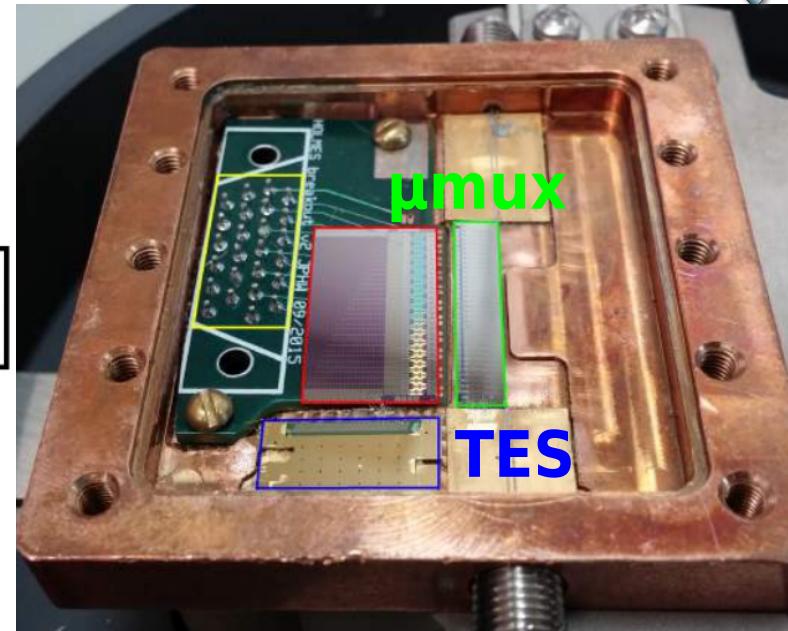


$$f_{\text{samp}} = 400 \text{ kS/s}$$

$$\Delta E_0 = 4.0 \text{ eV}$$

$$\tau_{\text{rise}} = 35 \mu\text{s}$$

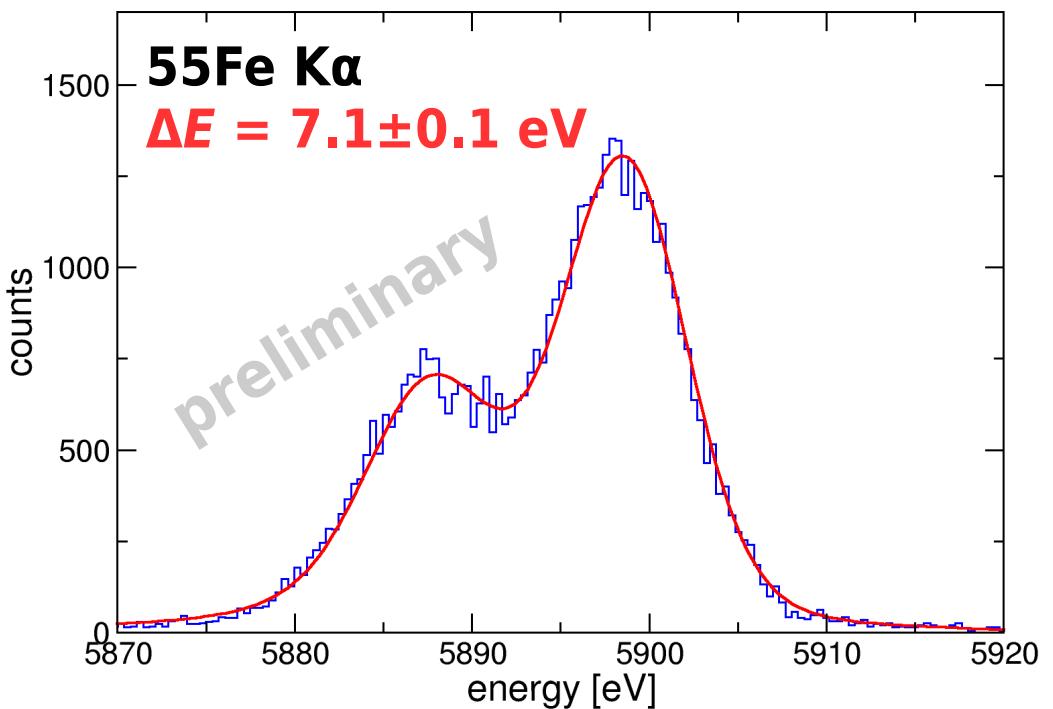
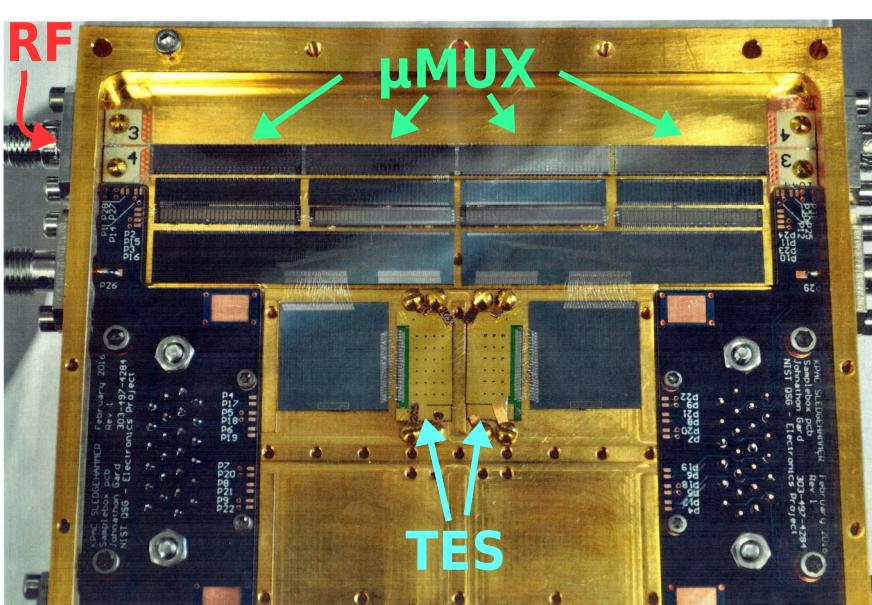
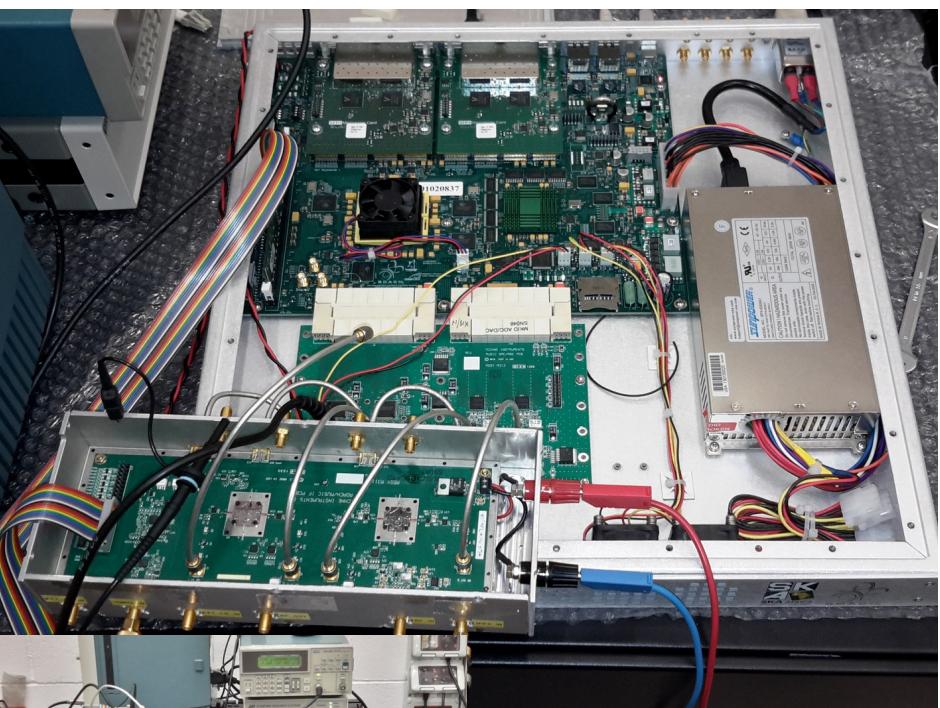
$$\tau_{\text{decay}} = 141 \mu\text{s}$$



Detector testing with HOLMES DAQ



ROACH-2 based Software Defined Radio



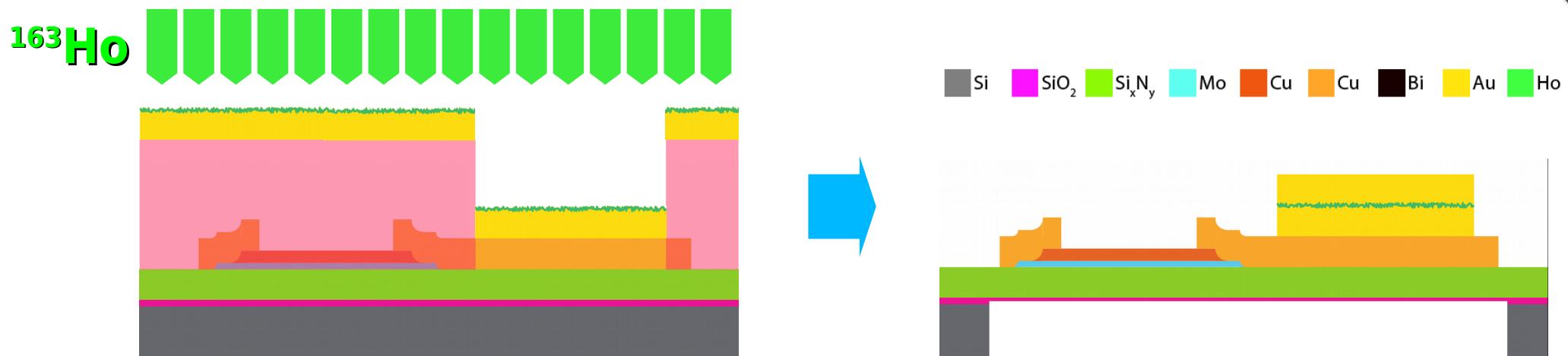
$$f_{\text{samp}} = 500 \text{ kS/s}$$

$$\Delta E_0 = 5.6 \text{ eV}$$

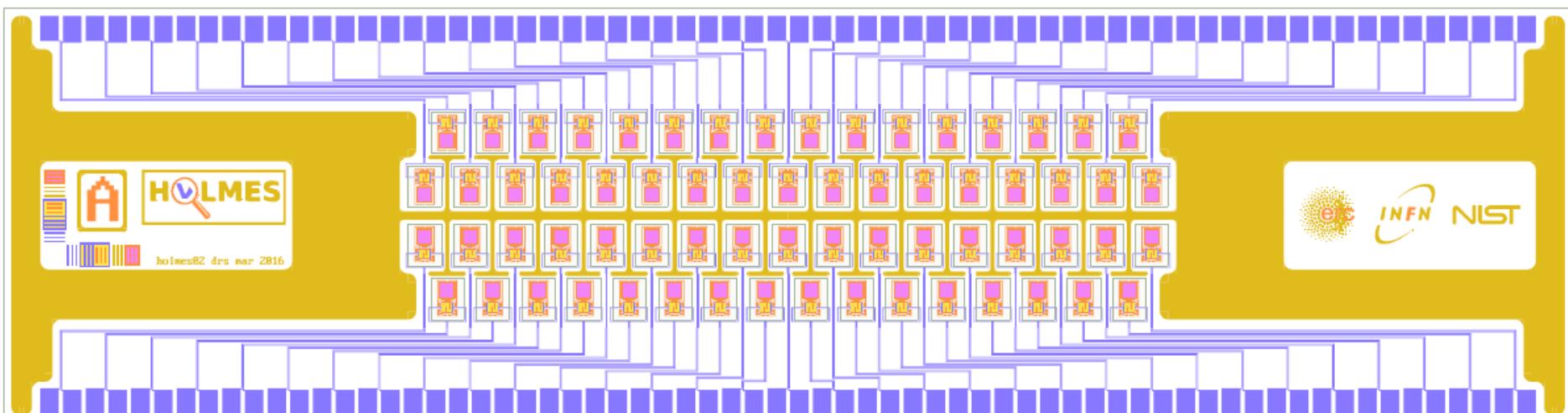
$$\tau_{\text{rise}} = 6.5 \mu\text{s}$$

$$\tau_{\text{decay}} = 67 \mu\text{s}$$

HOLMES detector design and fabrication



- TES array fabricated at **NIST**, Boulder, CO, USA
- ^{163}Ho implantation at **INFN**, Genova, Italy
- 1 μm **Au** final layer deposited at INFN Genova
- fabrication process definition in progress
- **HOLMES 4×16 linear sub-array** for low parasitic L and high implant efficiency

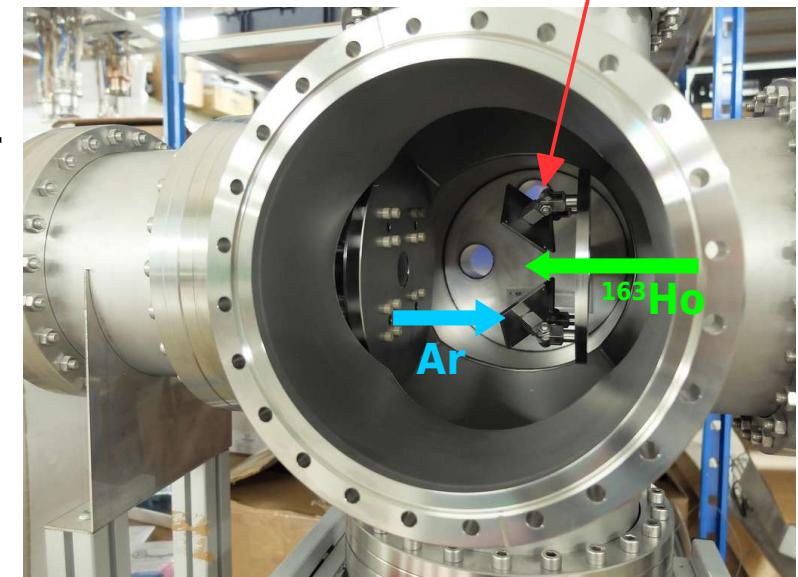
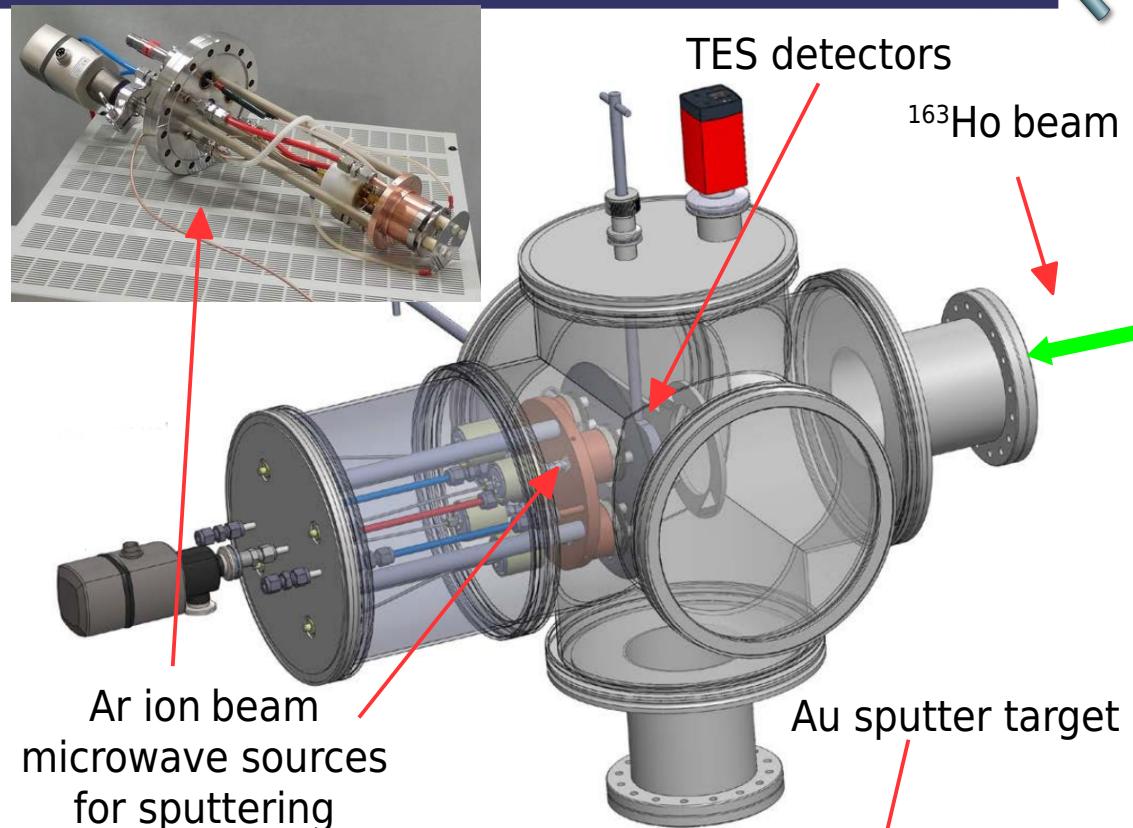
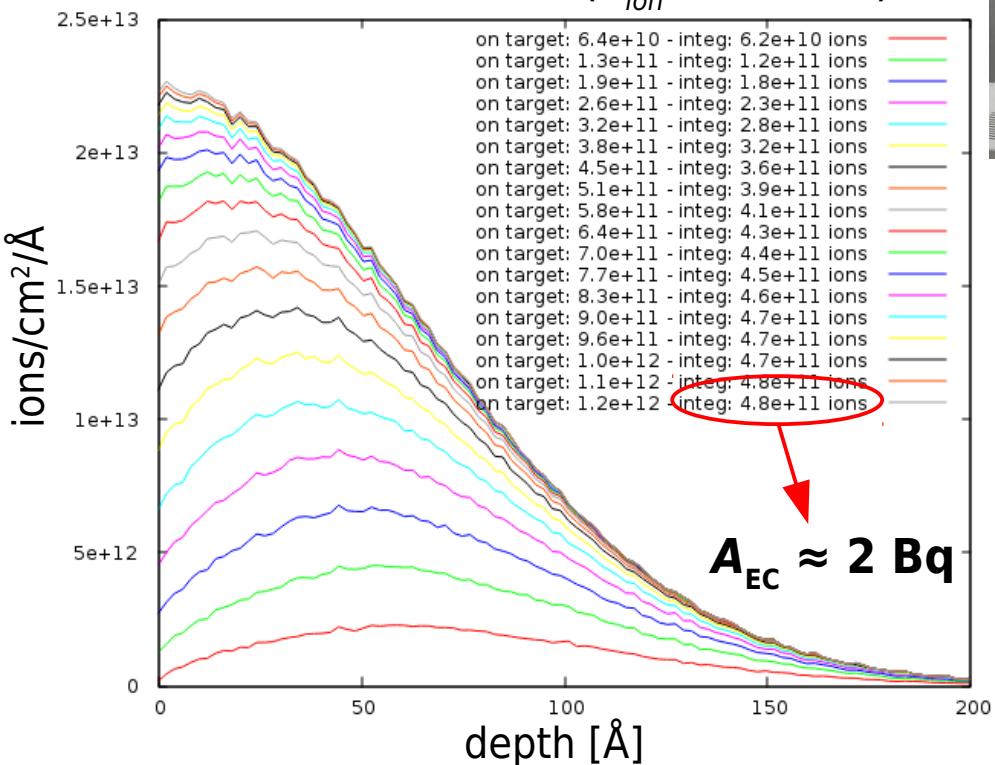


Target chamber for absorber fabrication



ion implant simulation with SRIM2013

^{163}Ho ions on Au ($E_{ion} = 50 \text{ keV}$)



- ^{163}Ho ion beam sputters Au from absorber
 - ▶ ^{163}Ho concentration in absorber saturates
 - ▶ compensate by Au co-evaporation
- final 1 μm Au layer in situ deposition
- tests are in progress

HOLMES schedule and conclusions



Project Year	2015	2016		2017		2018	
Task	S2	S1	S2	S1	S2	S1	S2
Isotope production							
TES pixel design and optimization							
Ion implanter set-up and optimization							
Full implanted TES pixel fabrication							
ROACH2 DAQ (HW, FW, SW)							
32 pix array 6mo measurement							
Full TES array fabrication							
HOLMES measurement							

▪ **HOLMES project status**

- many technical challenges are being addressed in parallel
- design phase is complete
- ion implanter setting up is in progress
- first ^{163}Ho implantation coming shortly
- spectrum measurements will begin late in 2017

Backup slides

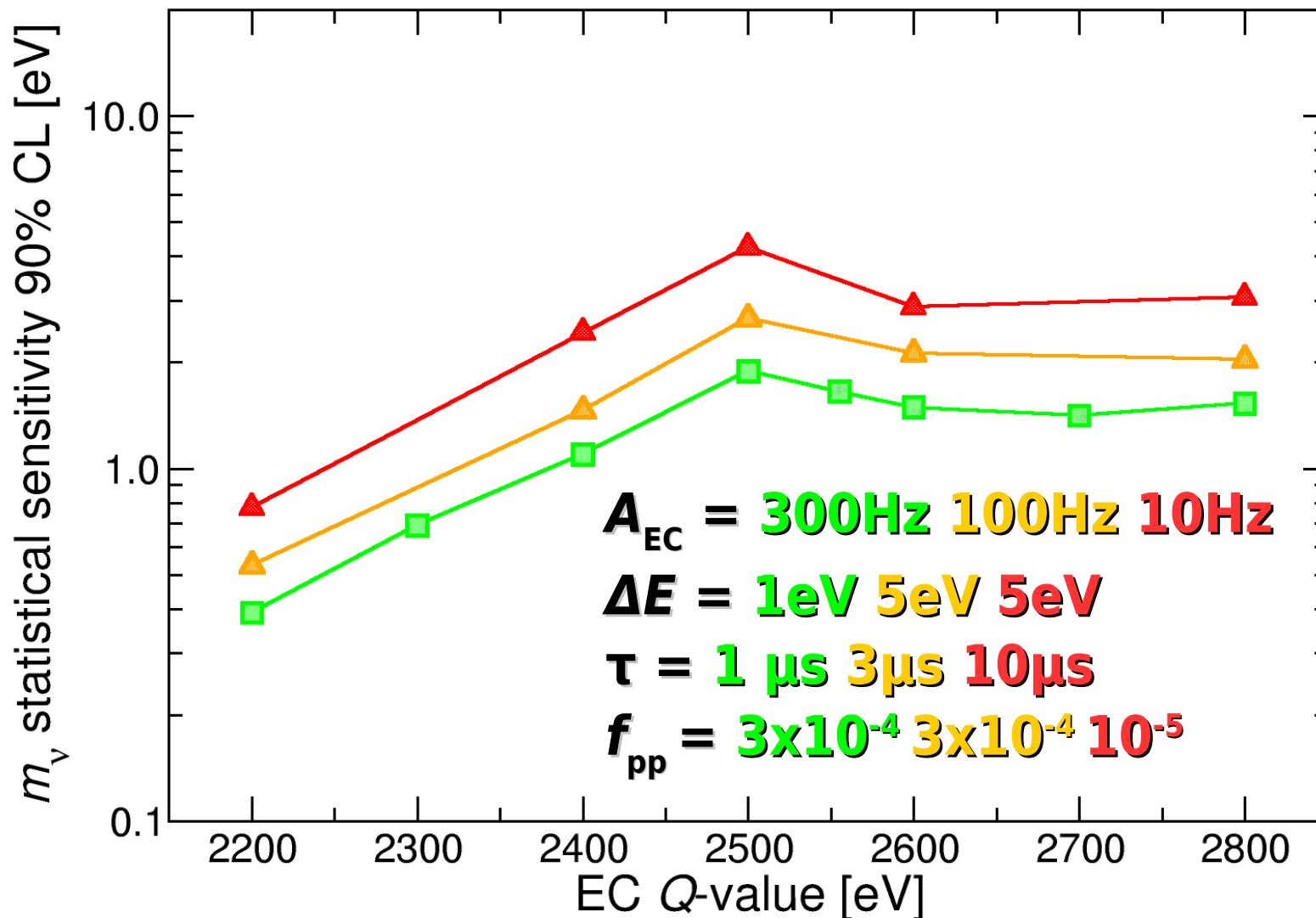


...

Worst case scenarios...



1000 channels
 $t_M = 3$ years



^{163}Ho production and embedding



■ ^{163}Ho production by nuclear reaction

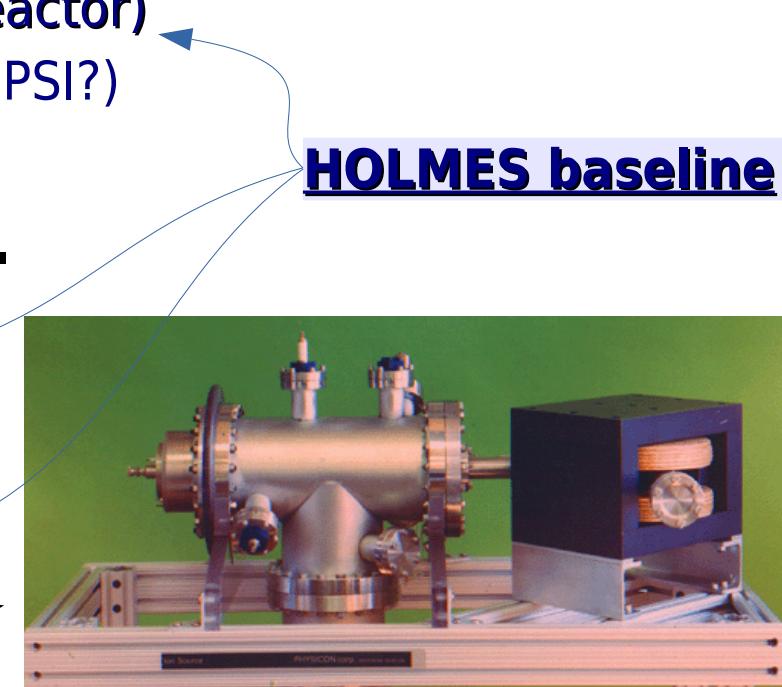
- ▶ high yield
- ▶ low by-products contaminations (in particular $^{166\text{m}}\text{Ho}$, $\beta \tau_{1/2} = 1200\text{y}$)
- ▶ not all cross sections are well known
 - neutron activation of enriched ^{162}Er (nuclear reactor)
 - $^{163}\text{Dy}(p,n)^{163}\text{Ho}$ $E_p > 10 \text{ MeV}$ (direct, low yield → PSI?)
 - $^{\text{nat}}\text{Dy}(\alpha, xn)^{163}\text{Er}$ and $^{159}\text{Tb}(^7\text{Li}, 3n)^{163}\text{Er}$

■ ^{163}Ho Separation from Dy, Er and more ...

- ▶ radiochemistry (before and/or after irradiation)
- ▶ magnetic mass separation
- ▶ resonance ionization laser ion source (RILIS)?

■ ^{163}Ho embedding in detector absorber

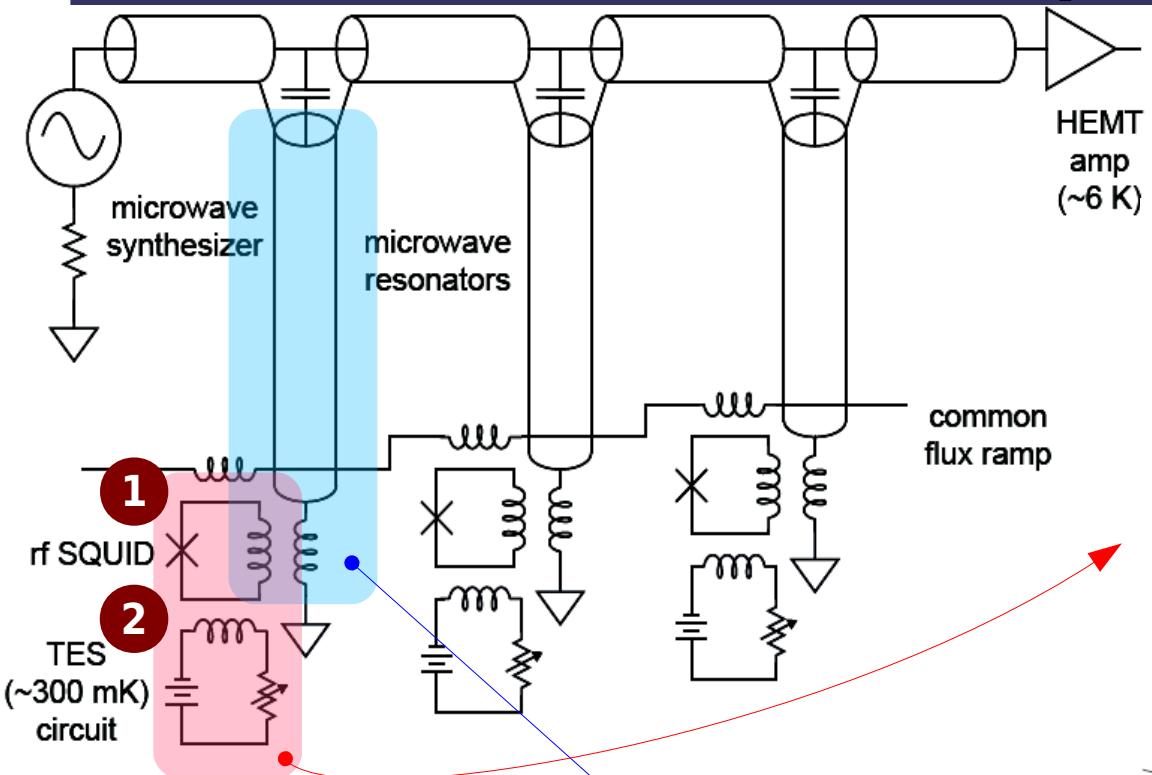
- ▶ implantation (+magnetic separation)
- ▶ Au film deposition for full containment



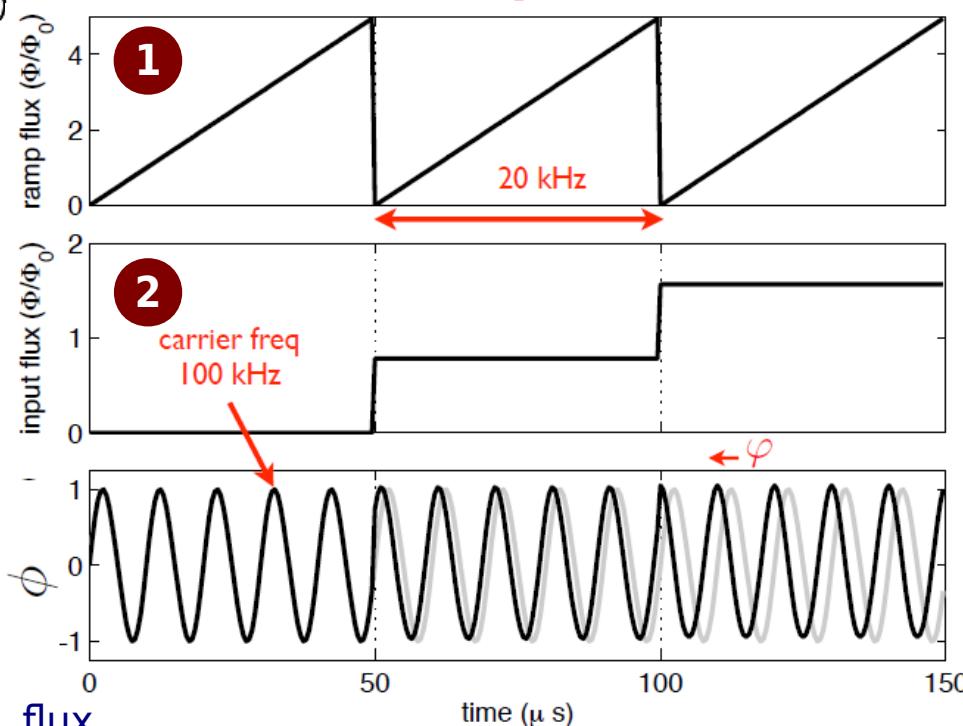
J.W. Engle et al., NIM B 311 (2013) 131-138

particle	p	$n 10^{14} \text{ n/cm}^2/\text{s}$	p 16 MeV 80 μA	p 24 MeV 240 μA	α 40 MeV 30 μA
target	W/Ta	^{162}Er (40%)	$^{\text{nat}}\text{Dy}$ 200mg/cm ²	$^{\text{nat}}\text{Dy}$ 20g	$^{\text{nat}}\text{Dy}$ “thick”
^{163}Ho prod rate [nuclei/h]	10^{14}	$10^{13-15}/\text{mg } ^{162}\text{Er}$	10^{14}	10^{15}	10^{13}

rf-SQUID microwave multiplexing

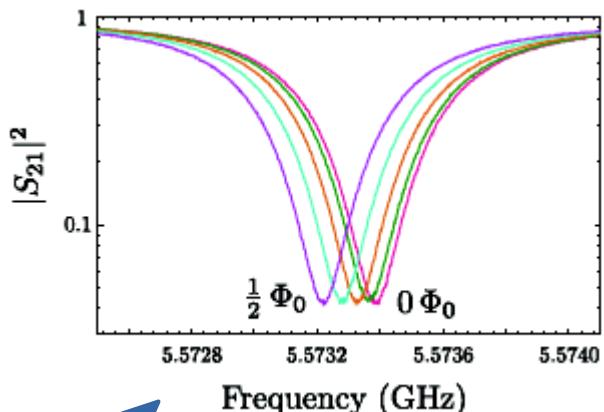
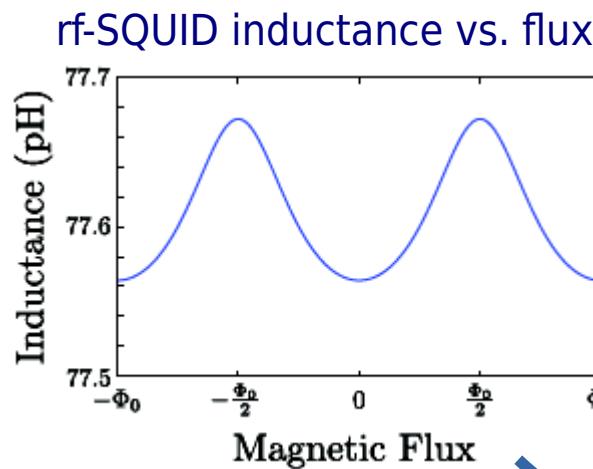


MHz flux ramp modulation



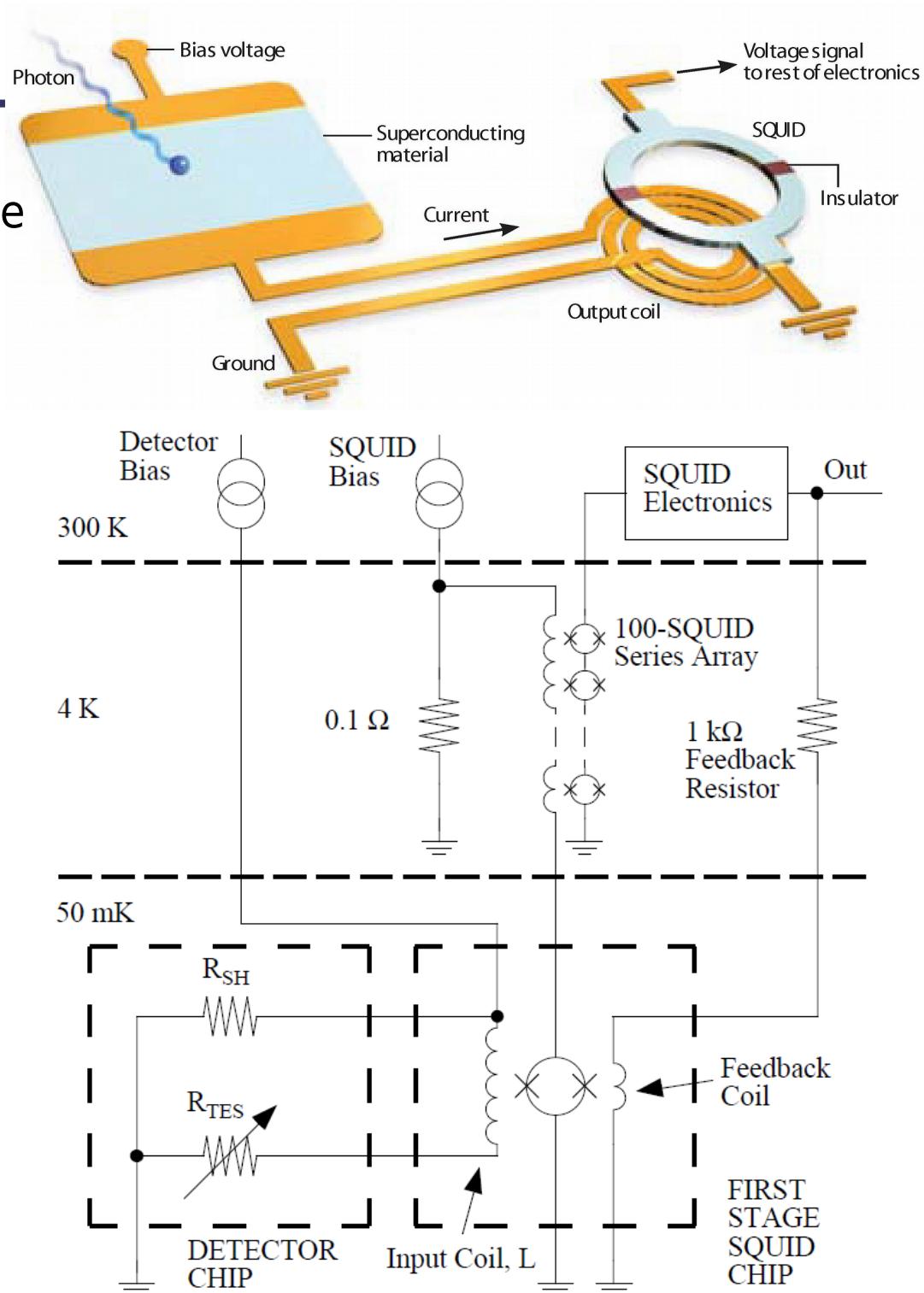
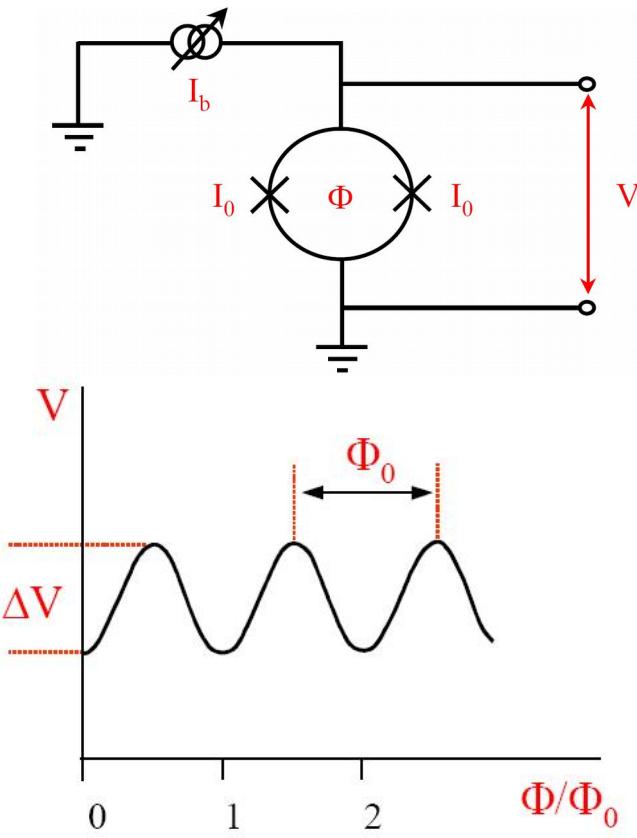
GHz LC resonator

resonance frequency vs. flux



TES read-out: SQUID

- low impedance suitable for multiplexable dc-SQUID magnetometers
- current amplifier configuration
▷ $\Delta I \rightarrow \Delta\Phi \rightarrow \Delta V$
- feedback linearized response



The holmium experiments



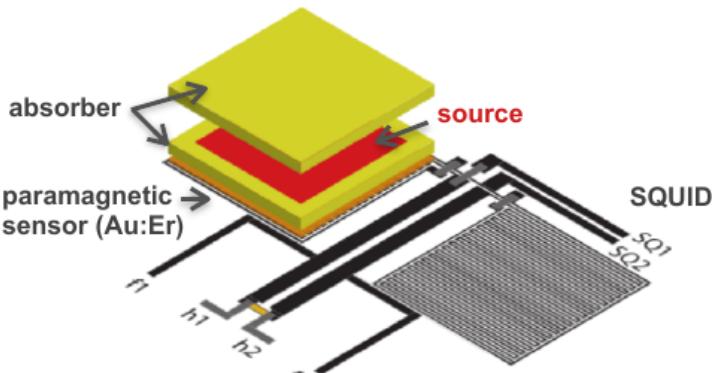
	ECHo	HOLMES	NUMECS
^{163}Ho production	$^{162}\text{Er} (n, \gamma)$	$^{162}\text{Er} (n, \gamma)$	$\text{Dy} (p, nx)$
Absorber	Gold	Gold	Nanoporous gold
Sensor	Au:Er magnetic	TES Mo/Cu	TES Mo/Cu
Present status			
ΔE at M1 peak [eV]	12	—	43 (incl. Γ_{M1})
τ_{rise} [μs]	0.13	—	—
A_{EC} [Bq]	0.2	—	0.1
Projected ($E_0 = 2800$ eV)			
N_{det}	100	1000	4096
ΔE [eV]	<5	1	—
τ_{rise} [μs]	<1	1	—
A_{EC} [Bq/detector]	10	300	100
f_{pp}	10^{-6}	3×10^{-4}	—
t_M [y]	1	3	1
$\Sigma_{90}(m_{\nu_e})$ [eV]	10	1.5	1



The ECHo Experiment

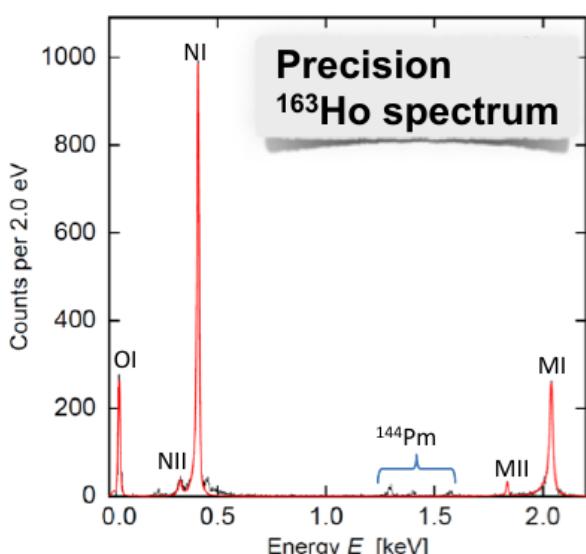


Heidelberg (Univ., MPIK), U Mainz,
U Tübingen, TU Dresden,
U Frankfurt, HU Berlin, ILL Grenoble,
PNPI St Petersburg, U Bratislava,
IIT Roorkee, Saha Inst. Kolkata



Technology

- magnetic micro-calorimeter (MMC) arrays with microwave squid multiplexing readout
- fast rise time (~ 130 ns) and excellent linearity & resolution ($\Delta E \sim 5$ eV)
- isotope production: $^{162}\text{Er}(n,\gamma)^{163}\text{Ho}$ offline mass separation



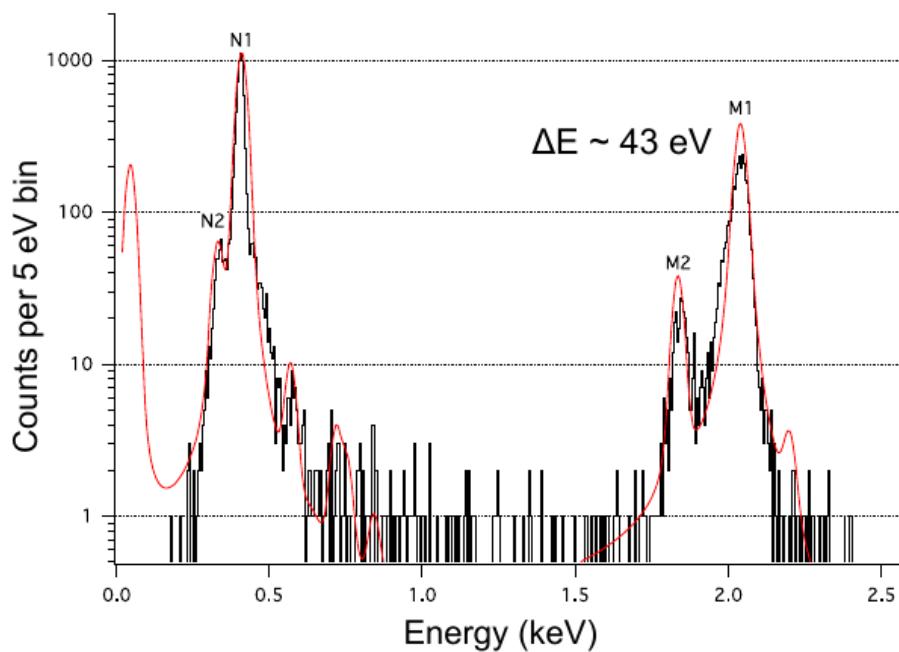
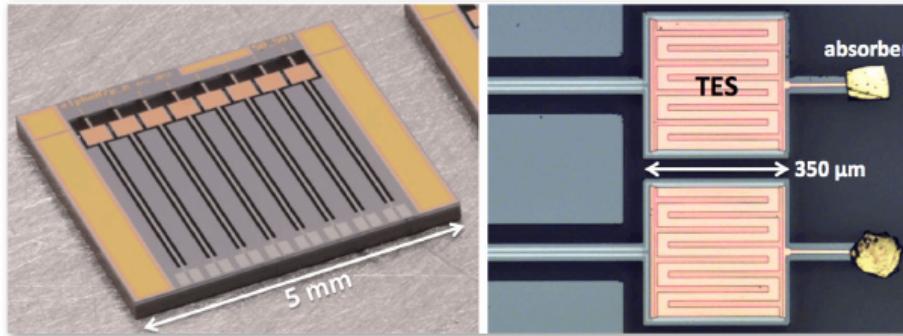
- new improved detectors
 improved implantation with purified source
 proof of multiplexed readout
 first underground meas.

Timeline

- **Phase I: ECHo-1k**
to be completed by 2018
2 x 50 pixel x 10 Bq,
4 months: $m(v_e) < 10$ eV
- **Phase II: ECHo-1M**
array of 10^5 detectors
50 x 2000 pixel x 10 Bq,
2 years: sub-eV sensitivity



The NuMECS Experiment



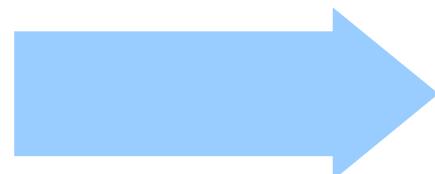
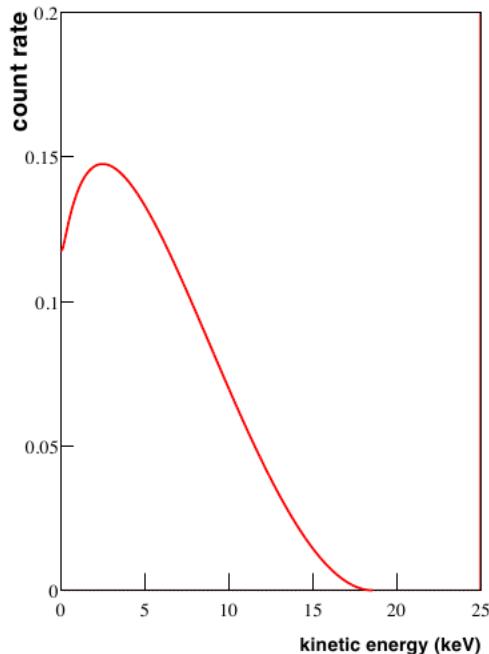
Technology

- transition-edge sensors (TES, Mo-Cu bilayer)
- gold foil absorbers with dried solution containing ^{163}Ho
- ^{163}Ho production by proton irradiation of natural dysprosium

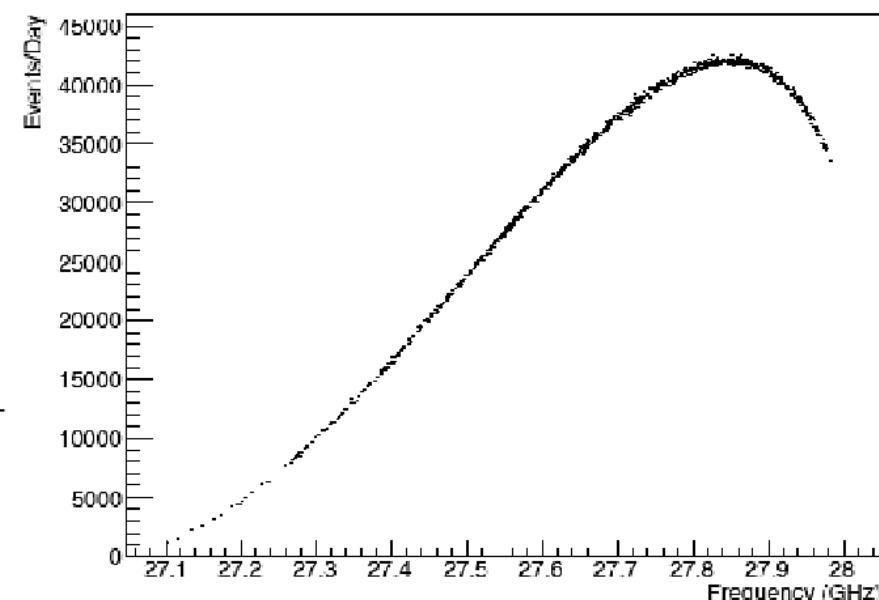
^{163}Ho spectrum

- obtained after systematic improvement of absorber production
- still limited statistics (40 hrs, 0.1 Bq) and resolution

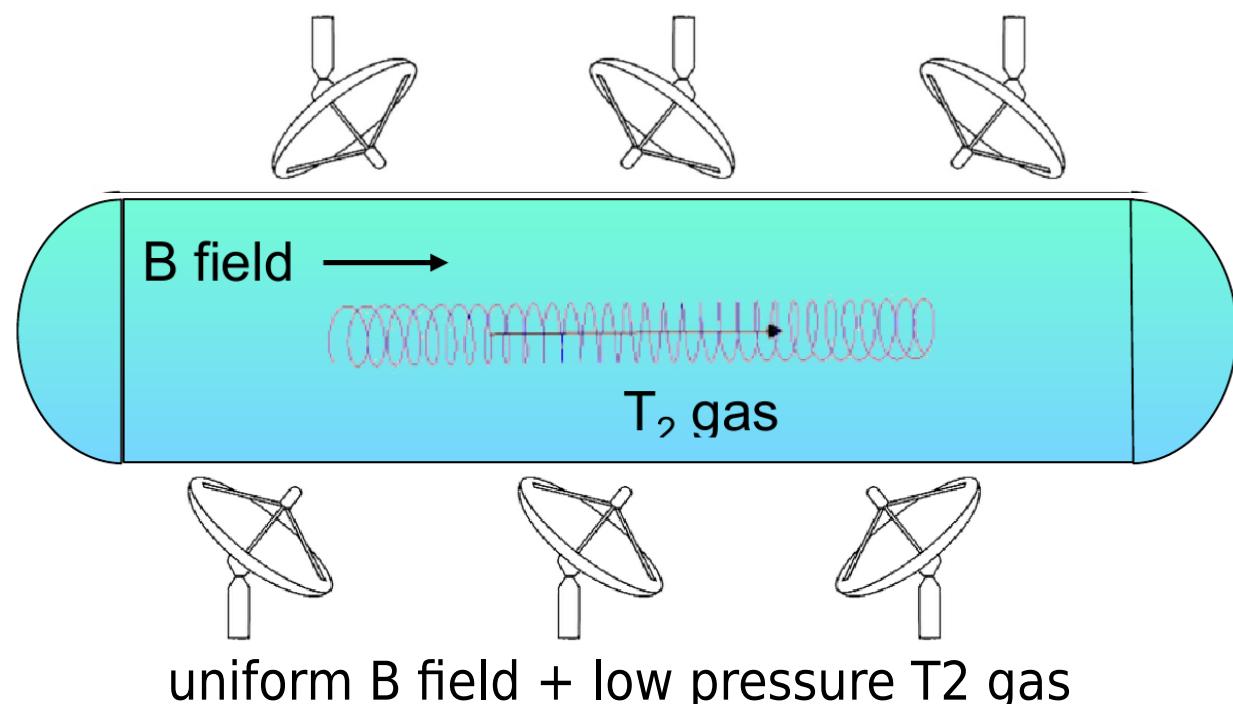
Beyond spectrometers: Project8 / 1



$$\omega(\gamma) = \frac{\omega_0}{\gamma} = \frac{eB}{K + m_e}$$



GHz cyclotron radiation detection to measure K

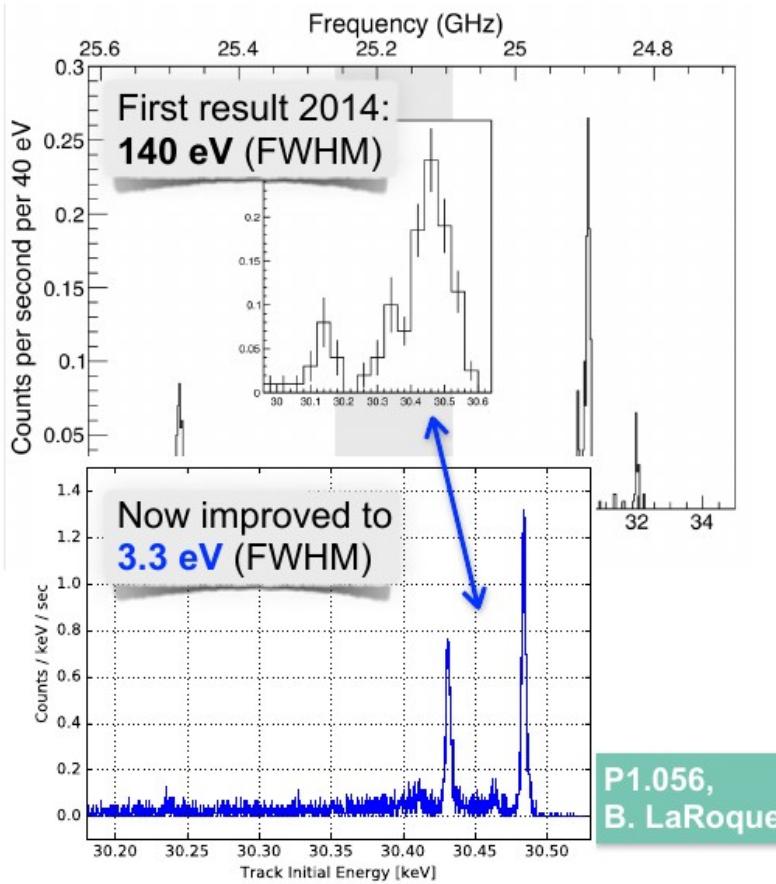




Project 8 – next goals

- **Phase I (2010-2016)**

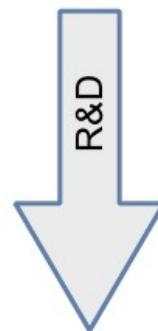
- Demonstration of CRES method
- Conversion electron lines from ^{83m}Kr



- **Phase II (2015-2017)**

- Spectroscopy of continuous T_2 spectrum
- Systematics, energy resolution

P1.057,
N. Oblath



- **Phase III (2016-2020)**

- 10-20 cm³ eff. source volume (1 yr)
- phased-array antenna
- sensitivity goal: **2 eV** (90% CL)

P4.047,
B. VanDevender

- **Phase IV**

- Large-scale exp., with atomic tritium source,
- for sub-eV sensitivity (hierarchy scale)

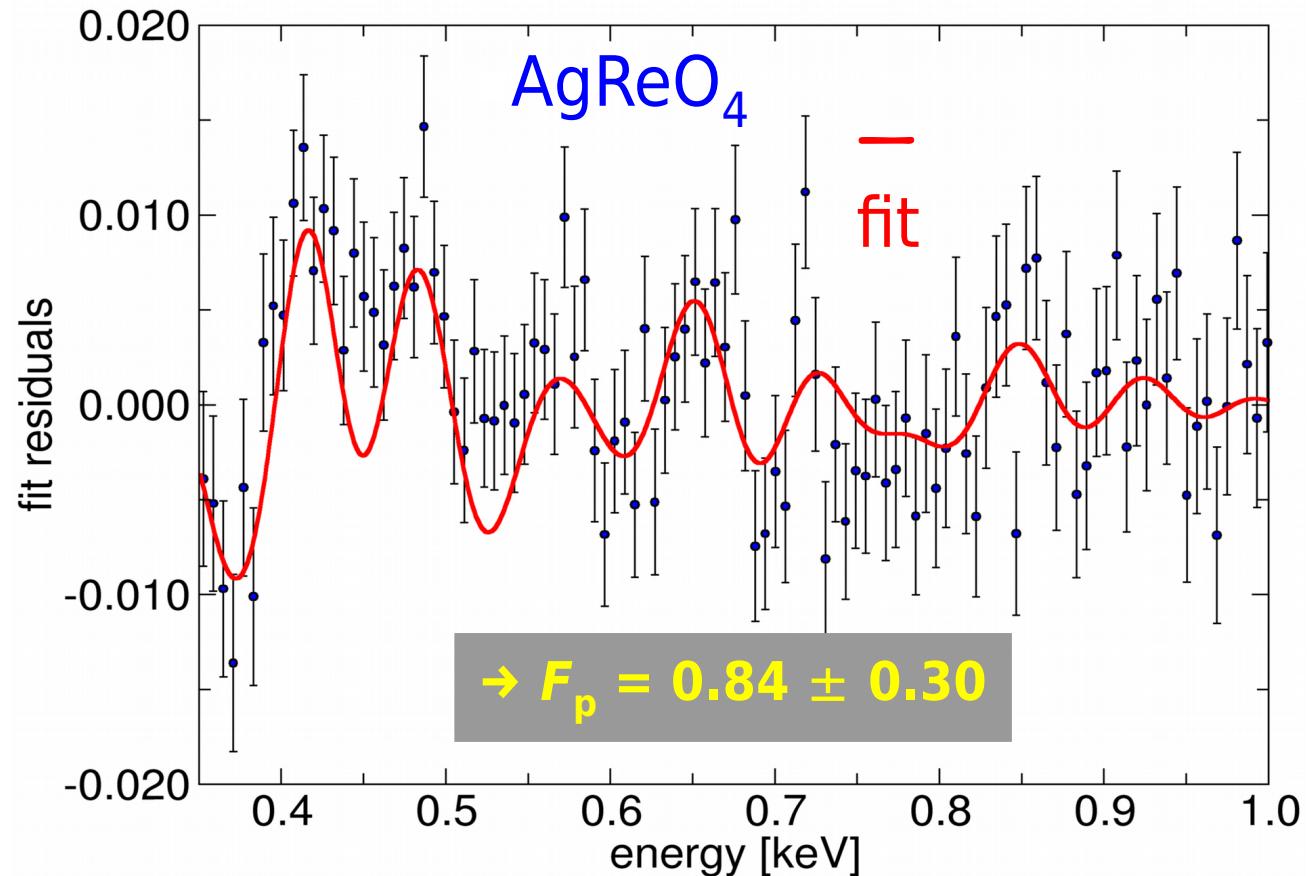
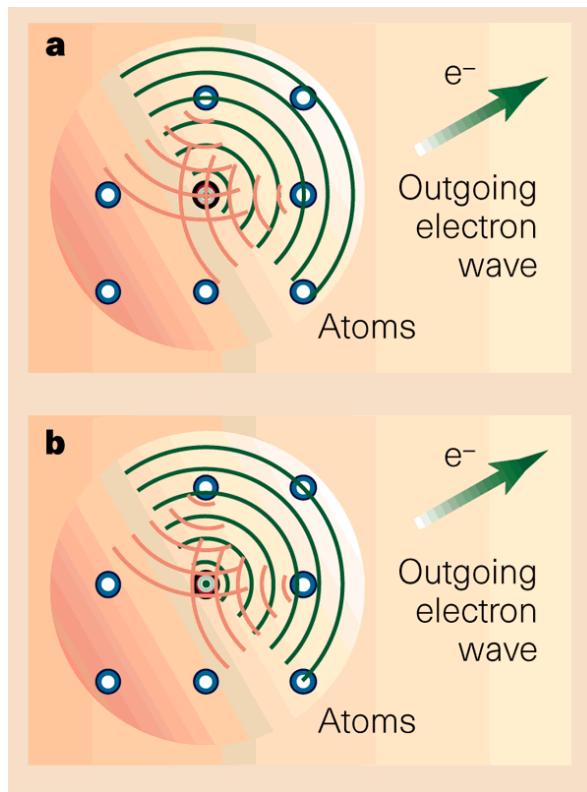
BEFS: Beta Environmental Fine Structure



Modulation of the electron emission probability due to the atomic and molecular surrounding of the decaying nucleus: it is explained by the wave structure of the electron (analogous of EXAFS)

$$\chi_{BEFS}(k_e) = F_s \chi_{EXAFS}^{l=0} + F_p \chi_{EXAFS}^{l=1}$$

$$\chi_{EXAFS}^l(k_e) = (-1)^l \sum_{n=1}^N B_{nl}(k_e, R_n) e^{-2k_e^2 \sigma_n^2} \sin(2k_e R_n + \delta_{0l} + \delta_{nl})$$

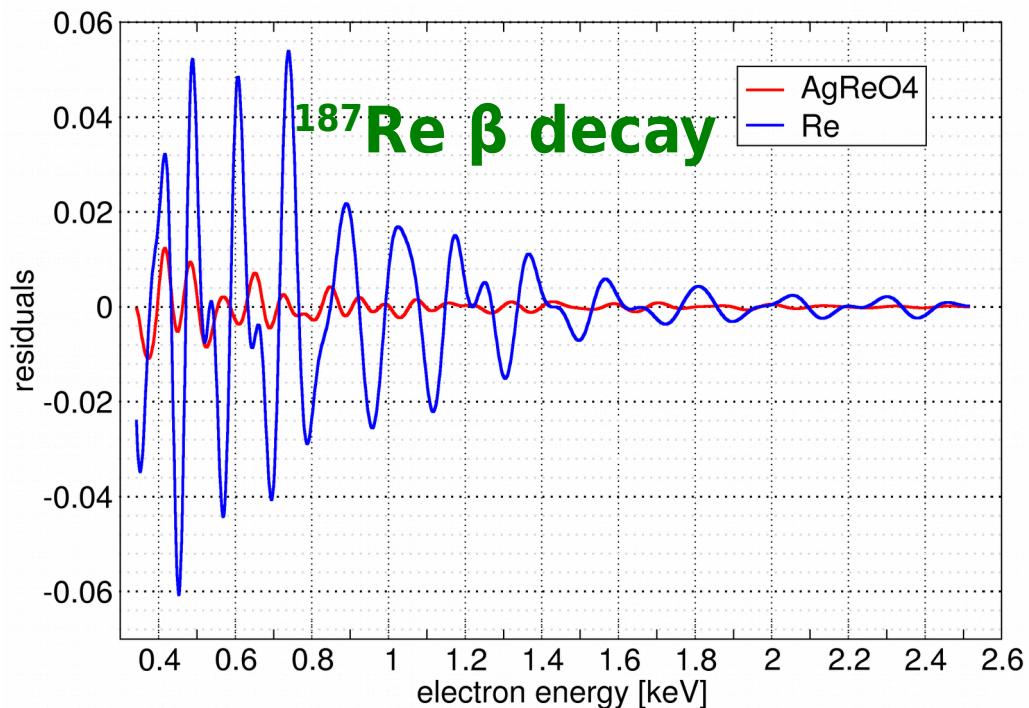
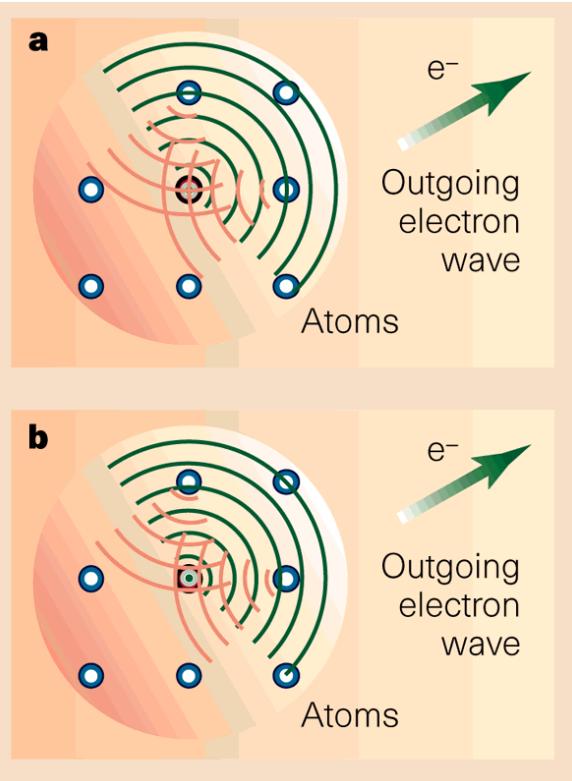


Beta Environmental Fine Structure in ^{163}Ho ?



$$\chi_{BEFS}(k_e) = F_s \chi_{EXAFS}^{l=0} + F_p \chi_{EXAFS}^{l=1}$$

$$\chi_{EXAFS}^l(k_e) = (-1)^l \sum_{n=1}^N B_{nl}(k_e, R_n) e^{-2k_e^2 \sigma_n^2} \sin(2k_e R_n + \delta_{0l} + \delta_{nl})$$

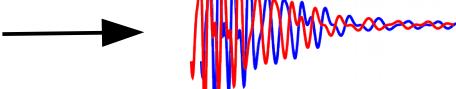


- what about BEFS in ^{163}Ho spectra?

▷ E_c deposited by cascade processes → convolution?



▷ different transition sequences → cancellation?



▷ smeared position of ^{163}Ho in host lattice → attenuation?





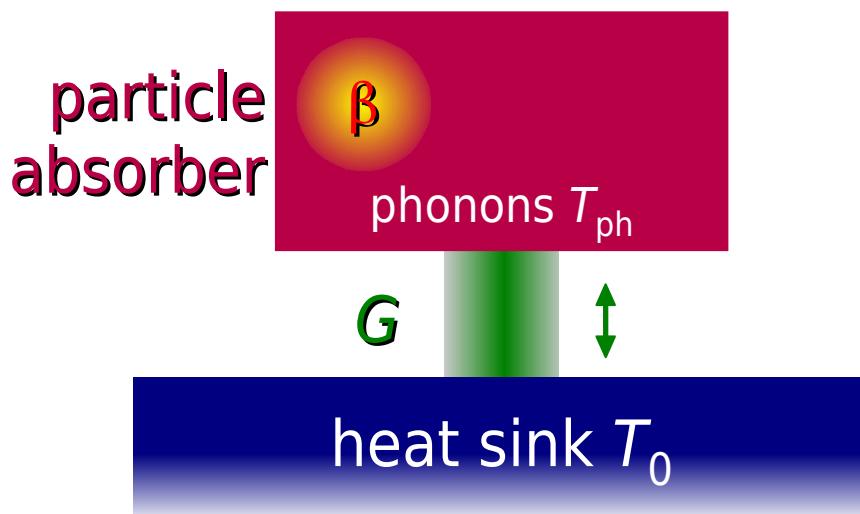
Resolution limit: LTD vs. ionization detectors

Ionization detectors

- measure only the energy that goes into ionization ($\sim 1/3$)
 - in semiconductors: energy to create an e-h pair $W_0 \approx 3 \text{ eV} \Rightarrow N_{eh} = E/W_0$
 - statistical fluctuations on N_{eh} limit the energy resolution $\sigma_E = \sqrt{F N_{eh}} W_0 = \sqrt{F E W_0}$
 - in practice: $\Delta E_{FWHM} \approx 115 \text{ eV}$ at 6 keV for silicon
- other limitations from electron transport properties (material restriction, purity...)

Cryogenic detectors

- measure the energy that goes into heat (100%)
 - no branching \Rightarrow no statistical fluctuations
 - resolution limit: random energy flow through G
 - statistical fluctuations of internal energy $U = \langle U \rangle \pm \Delta U_{rms}$



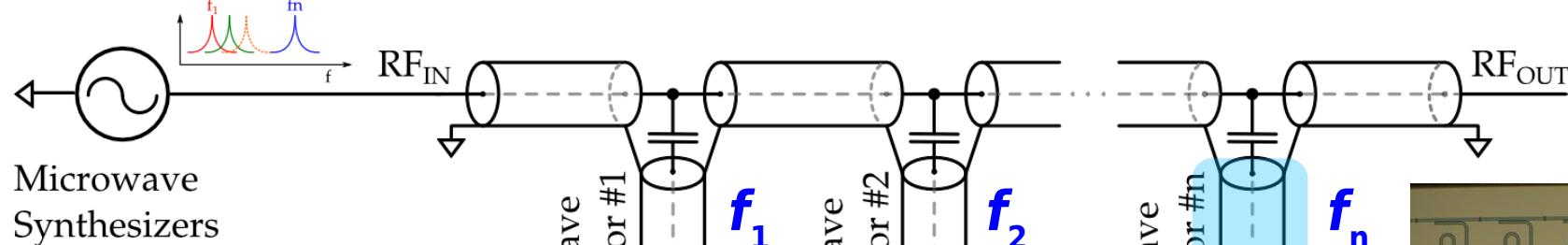
$$N_{ph} = \frac{\langle U \rangle}{\langle E_{ph} \rangle} = \frac{CT}{k_B T}$$

$$\Delta U_{rms} = \sqrt{N_{ph}}(k_B T) = \sqrt{k_B T^2 C}$$

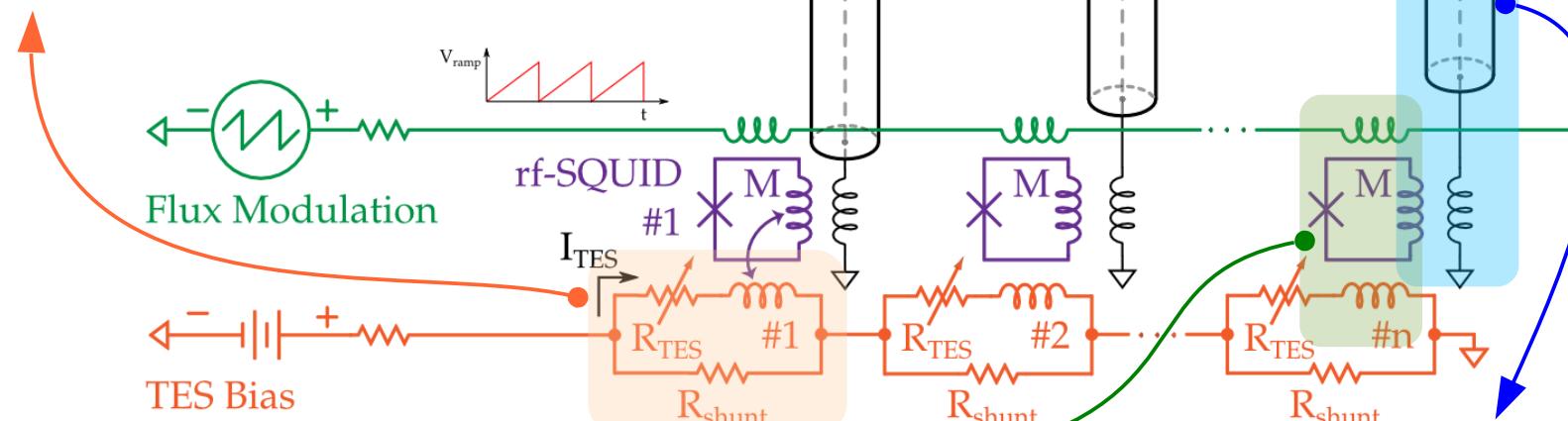
1 mg of Re @ 100 mK

$$C \sim 10^{-13} \text{ J/K} \Rightarrow \Delta U_{rms} \sim 1 \text{ eV}$$

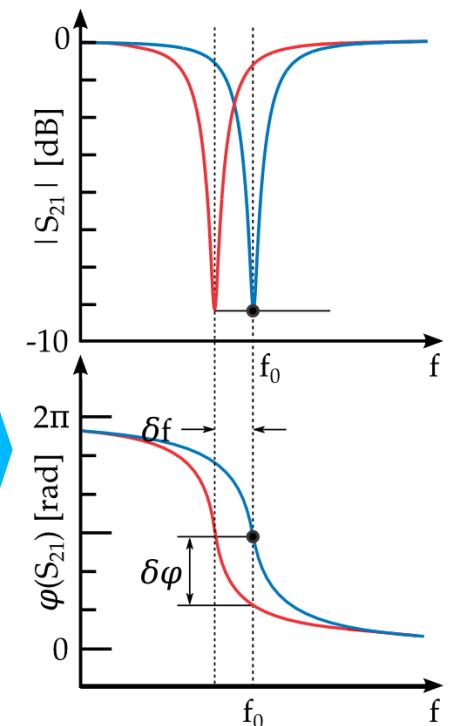
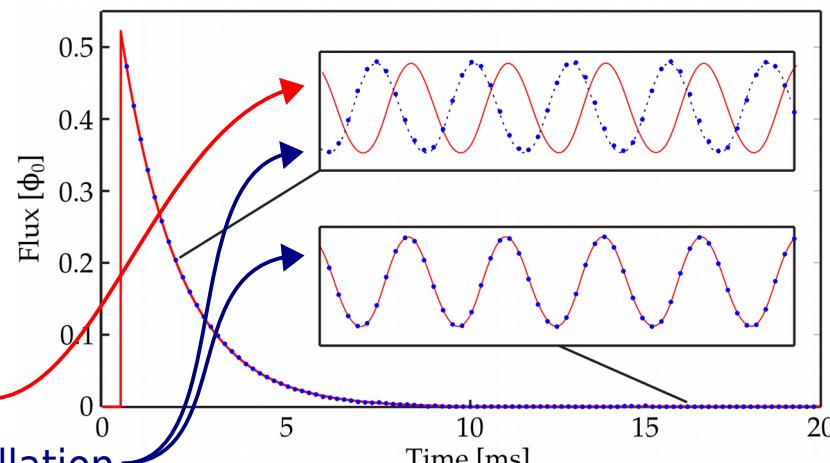
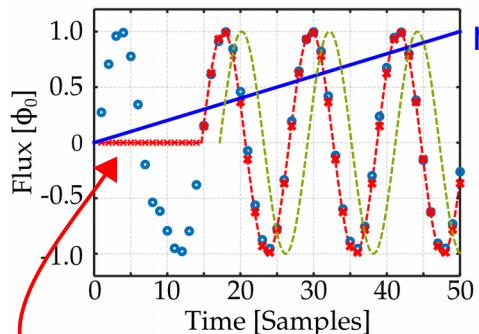
HOLMES array read-out: rf-SQUID μwave mux



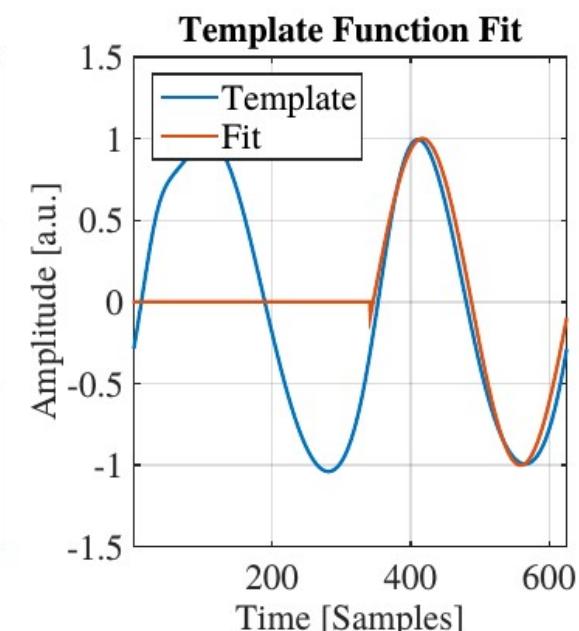
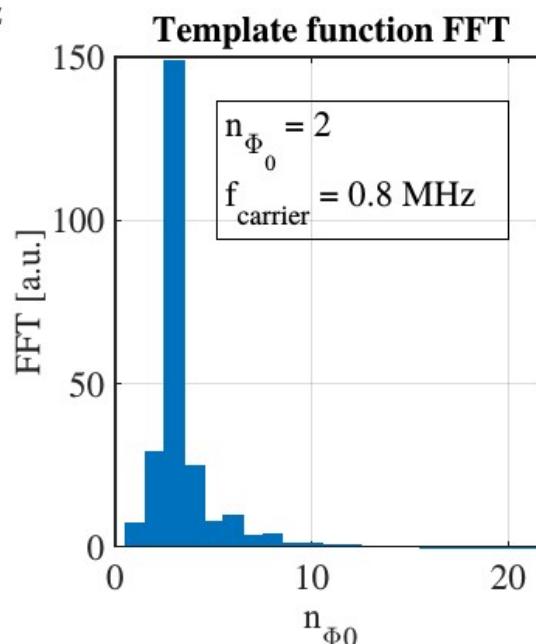
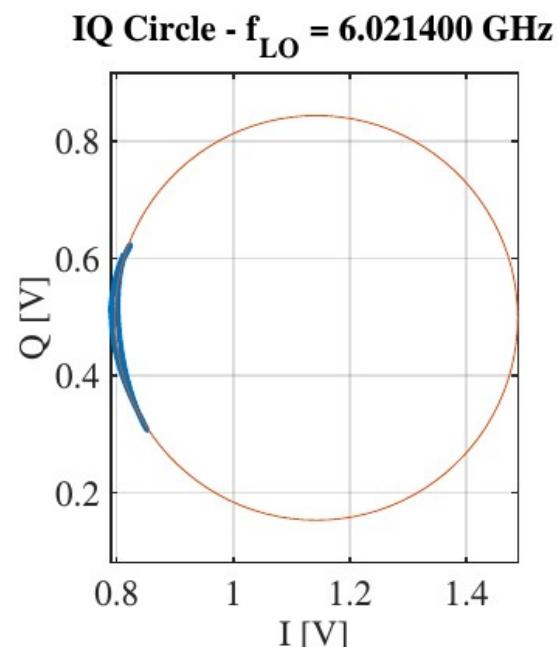
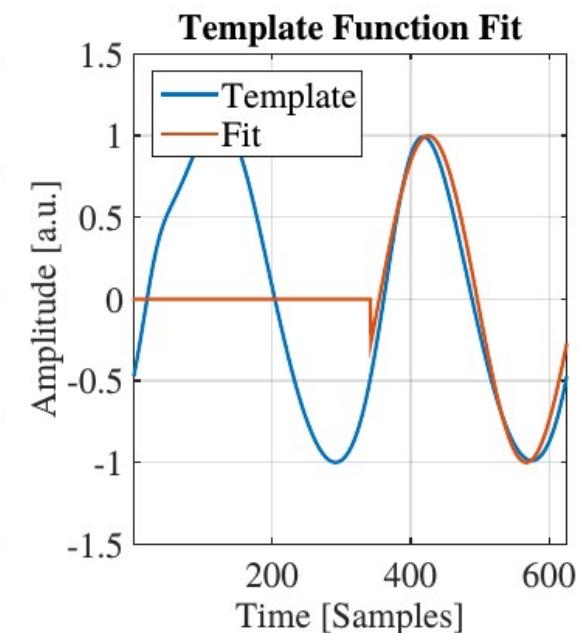
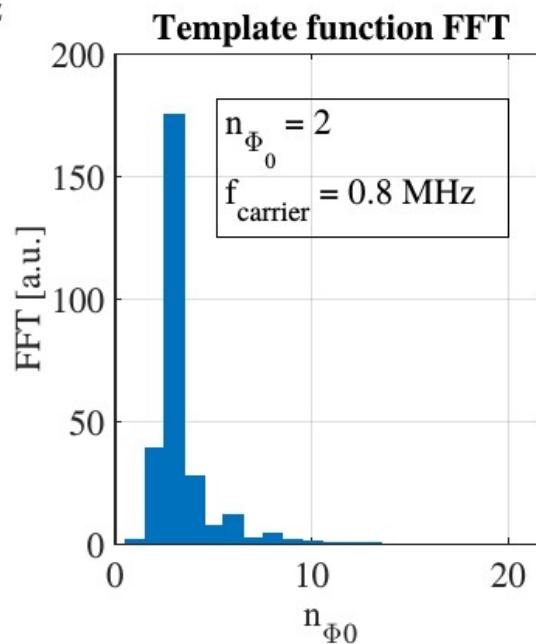
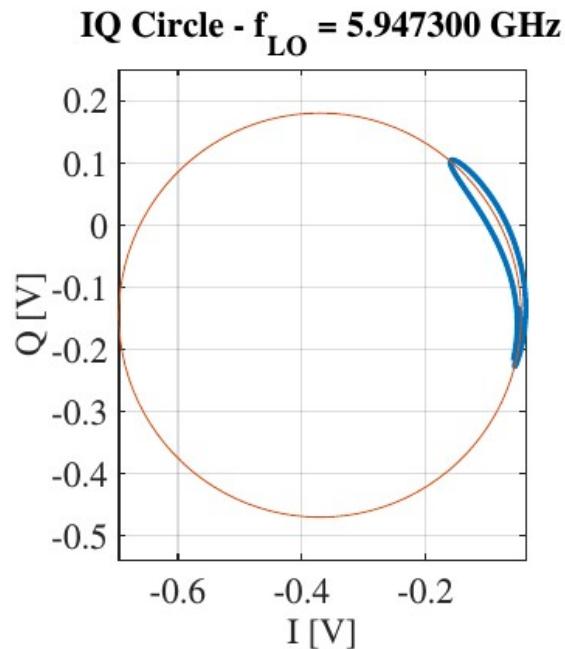
dc biased TES sensor



MHz flux ramp modulation $f_{ramp} \rightarrow f_{sample}$



Ramp demodulation



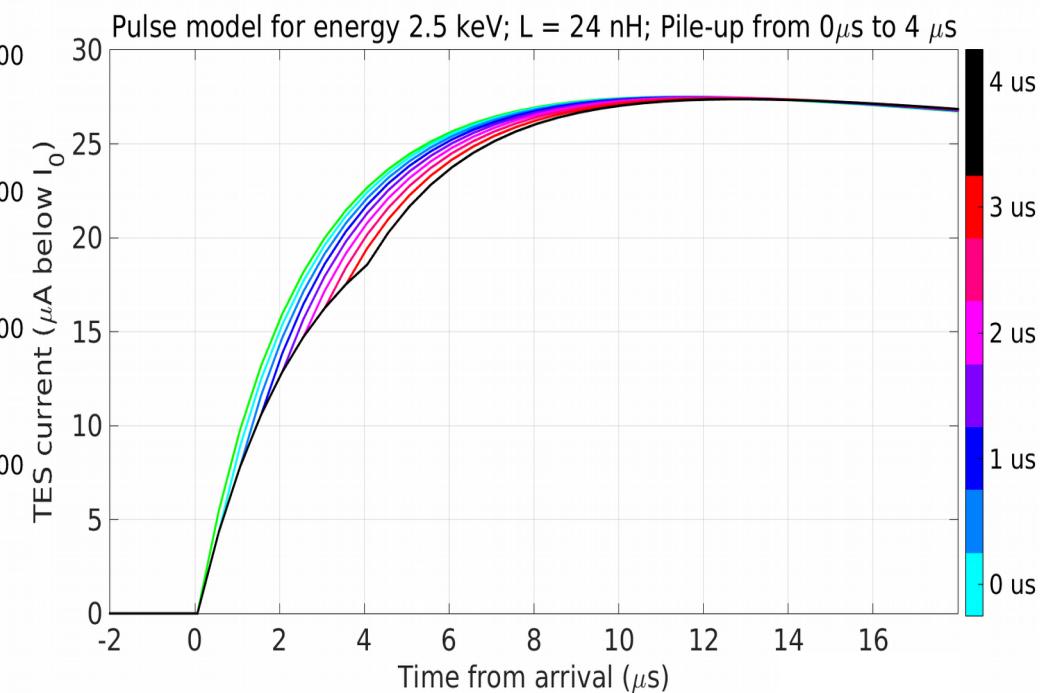
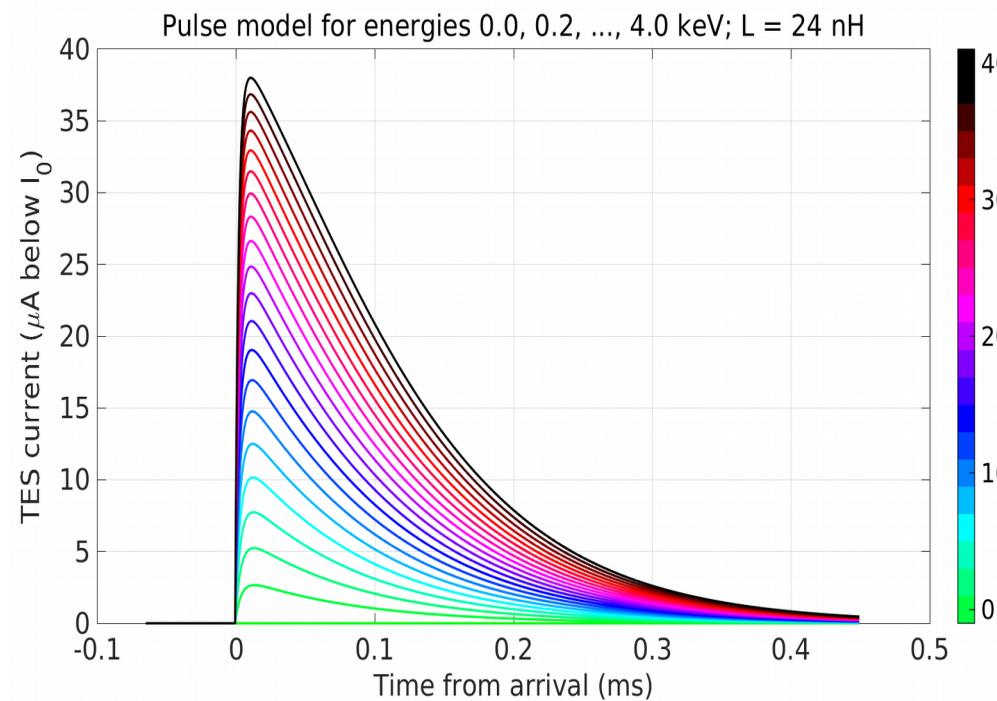
Rise time pile-up



2 pulses with:

- $\tau_{\text{rise}} = 1.5 \text{ ms}$
- $\tau_{\text{decay}} = 10 \text{ ms}$
- $A_2/A_1 = 0.5$

$$A(t) = A \left(e^{-t/\tau_{\text{decay}}} - e^{-t/\tau_{\text{rise}}} \right)$$



HOLMES signal processing and in-line analysis



- normal data taking (permanent RAID storage)
 - ▶ save only n -tuples (6×4 byte words) *
 - ▶ high threshold ($E_{th} \approx 2.022\text{keV}$, $E_{M1} = 2.041\text{keV}$, $Q_{EC} = 2.8\text{keV}$, 21% of spectrum) *
 - ▶ about 150TB in 3 years (un-compressed)
- periodic minimum bias samples (temporary storage)
 - ▶ tune parameters for real time pulse processing
 - ▶ full waveform (512 samples at 12 bit) for immediate off-line analysis *
 - ▶ full spectrum → 20TB/day
 - ▶ combined with high threshold data
- lower threshold is possible with compression

ROACH2 FW real-time

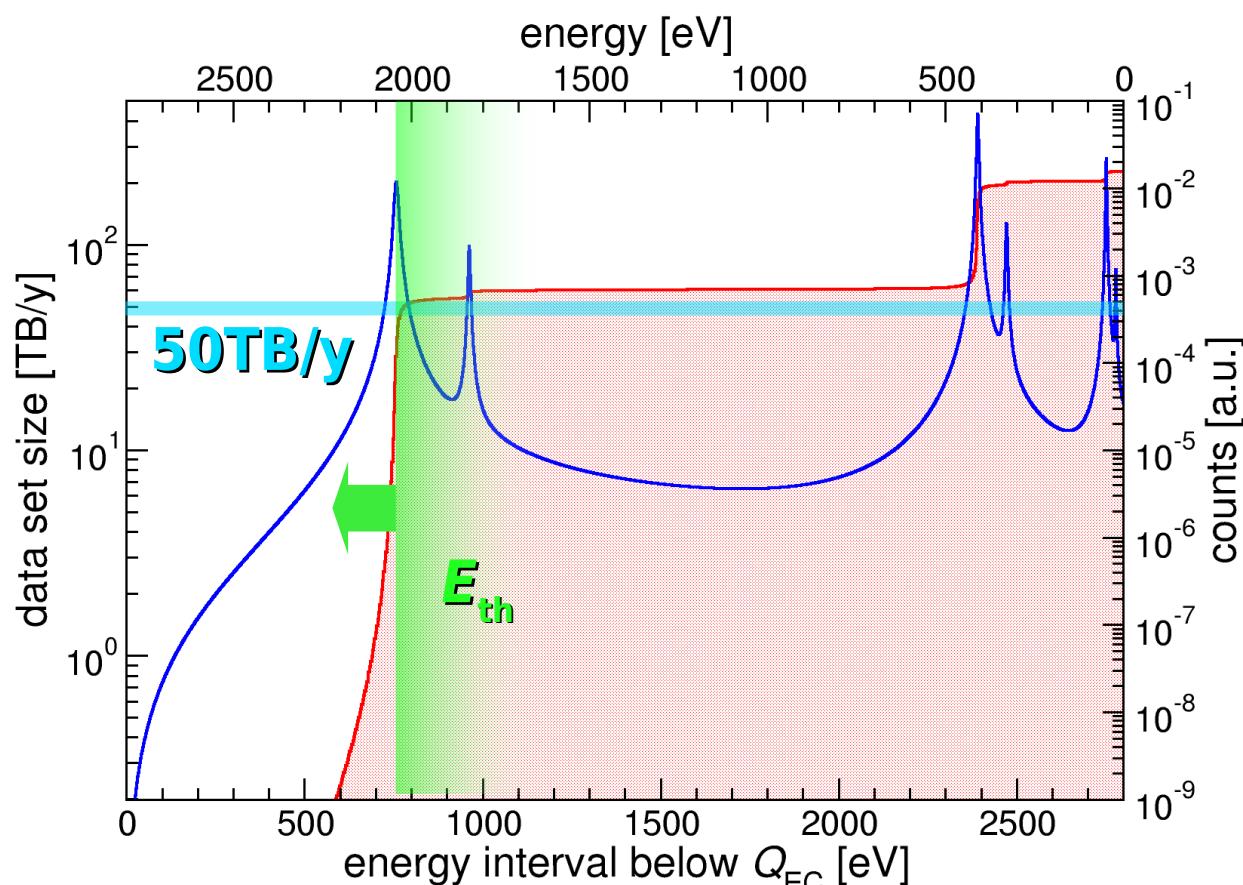
pulse processing:

- threshold cut
- ...

SERVER quasi real-time

pulse processing:

- OF analysis → n -tuples
- pile-up detection
- ...

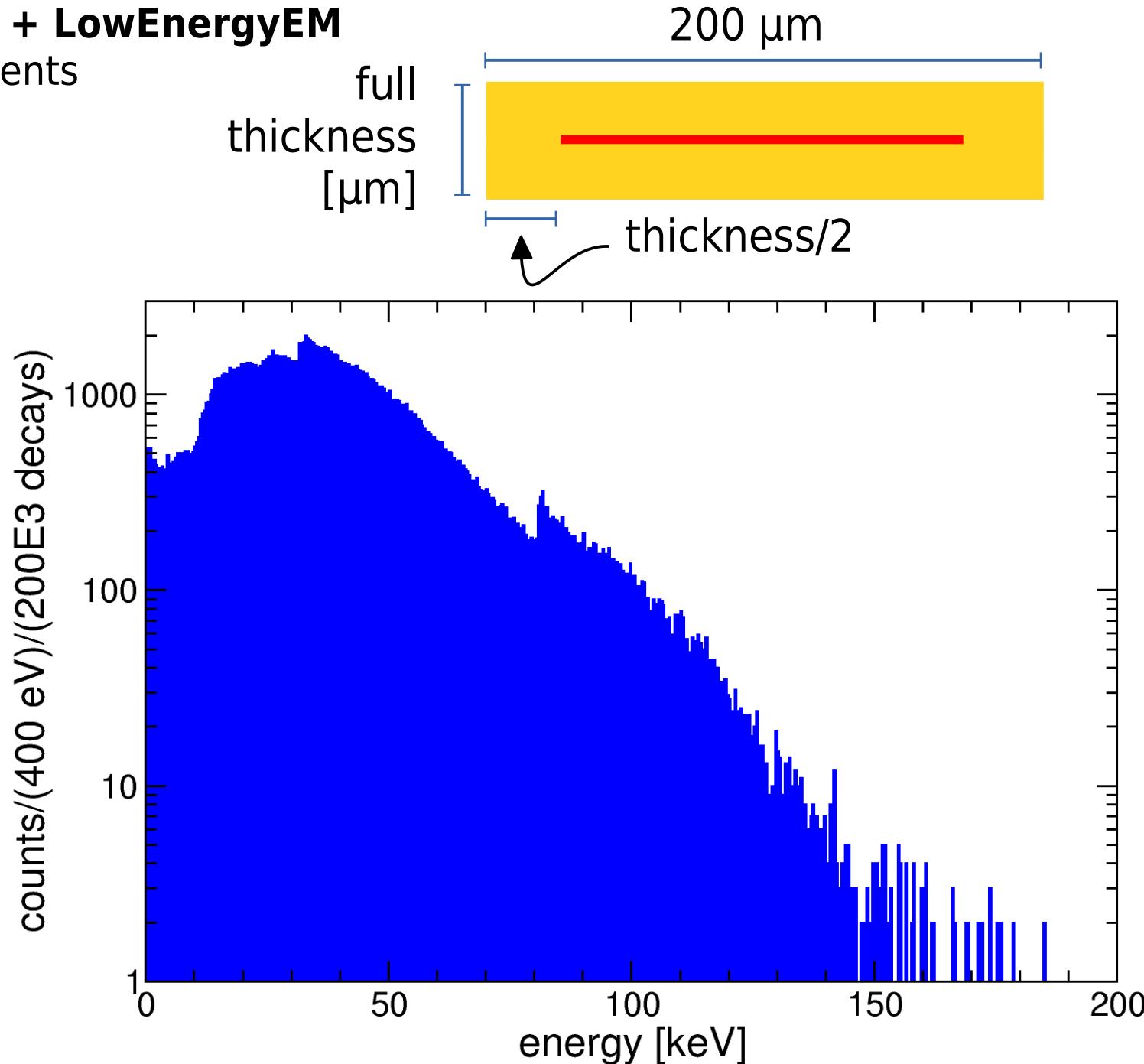


^{166m}Ho background



Geant4 + LowEnergyEM

2×10^5 events



Low energy background sources / 2

