

Bolometric Search for Neutrinoless Double Beta Decay

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SMU

OUTLINE

- * Neutrino physics & DBD0v
- * DBD0v sensitivity
- * DBD0v bolometric detectors
 - * CUORE & CUORICINO
 - * Background studies
- * Radon-induced surface contaminations
- * Effects on large mass experiments & solutions
- * Conclusions

Neutrino physics

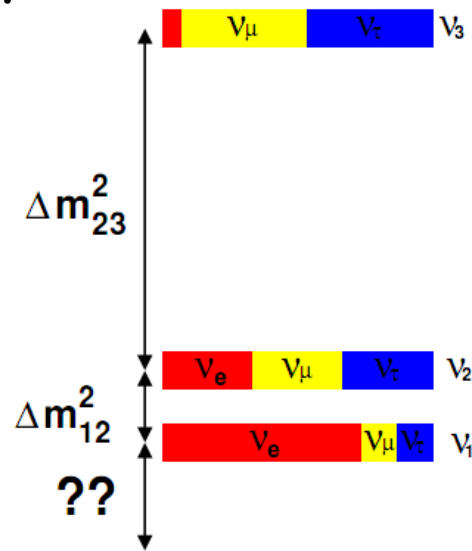
known

- Neutrinos oscillate
=> Non-zero mass
- Some oscillation parameters are known

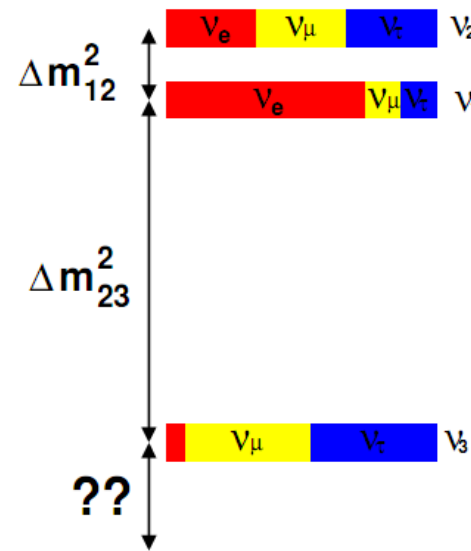
unknown

- Mass hierarchy
- Absolute mass scale
- Dirac or Majorana particle

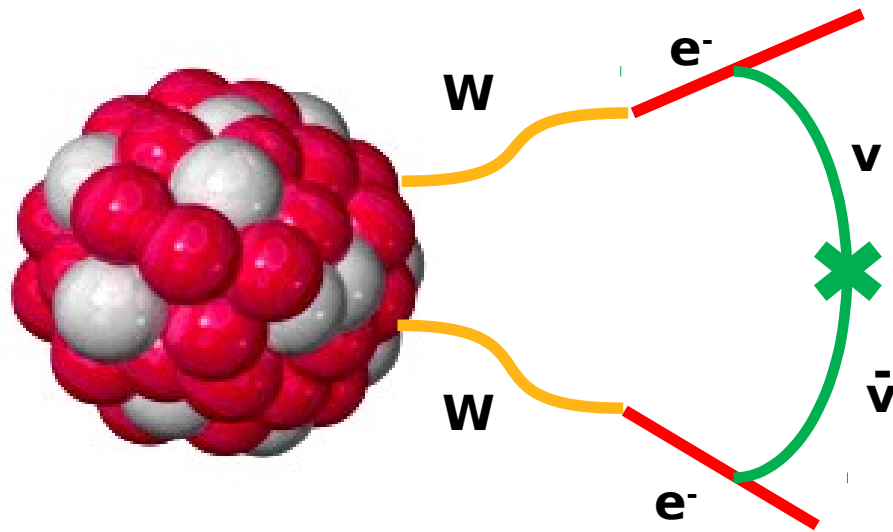
Normal hierarchy



Inverted hierarchy



DBDO ν



- 2nd order nuclear weak decay
- not allowed in the SM
- $t_{1/2}$ expected $> 10^{25}$ y

If observed:

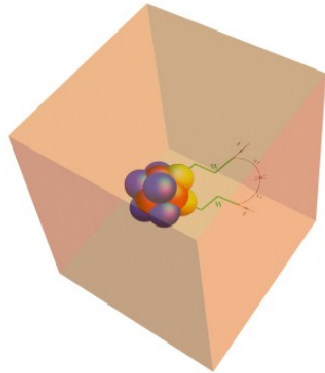
- the neutrino is a Majorana particle: $\nu_e \equiv \bar{\nu}_e$

- $\Delta L = 2$, lepton number violation

- ν mass measurement: $(T_{1/2}^{0\nu})^{-1} = G^{0\nu}(Q, Z) |M^{0\nu}|^2 \frac{|\langle m_{\beta\beta} \rangle|^2}{m_e^2}$

Design of a DBDOv experiment

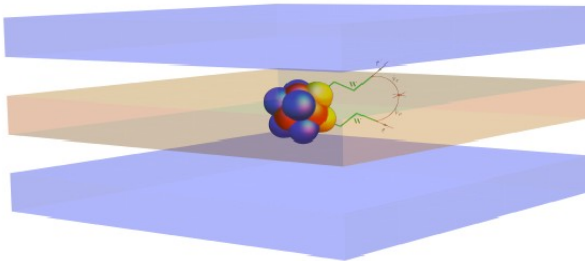
Source = Detector



Calorimetry

- * High efficiency
- * Good energy resolution
- * Large mass source

Source $\neq$ Detector

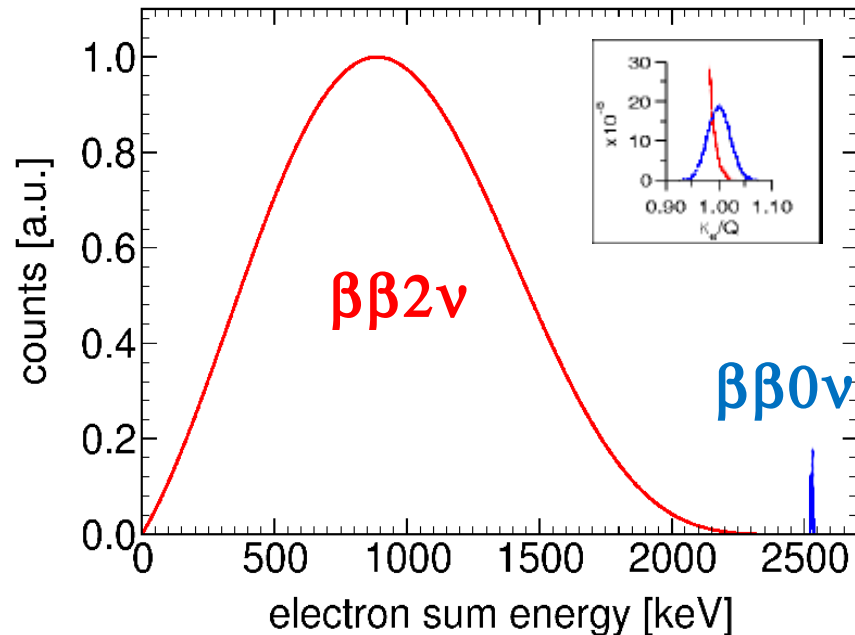


Scintillation

Tracking

- * Large mass source
- * Particle identification

What are we looking for ??



Monochromatic signal
@ Q-value ($\sim$ MeV range)

$$S_{0\nu} \propto \varepsilon \frac{i.a.}{A} \sqrt{\frac{M \cdot T}{\Delta E \cdot b}}$$

$b \neq 0$

ε : efficiency

$i.a.$: isotopic abundance

A : atomic mass number

M : source mass

T : live time

ΔE : FWHM in the ROI

b : bkg in the ROI

...

DBDO ν Sensitivity

$$S_{0\nu} \propto \varepsilon \frac{i.a.}{A} \sqrt{\frac{M \cdot T}{\Delta E \cdot b}}$$

ε : efficiency $\Rightarrow \sim 100\%$ for bolometers

i.a.: isotopic abundance $\Rightarrow$ Natural
Enrichment $\Rightarrow$ \$\$\$\$
radiopurity

M: detector mass [kg] $\Rightarrow$ Next generation ~ 1 ton

T: time [y] $\Rightarrow \sim 5$ y

ΔE : energy resolution [keV] $\Rightarrow \sim 5$ keV for bolometers

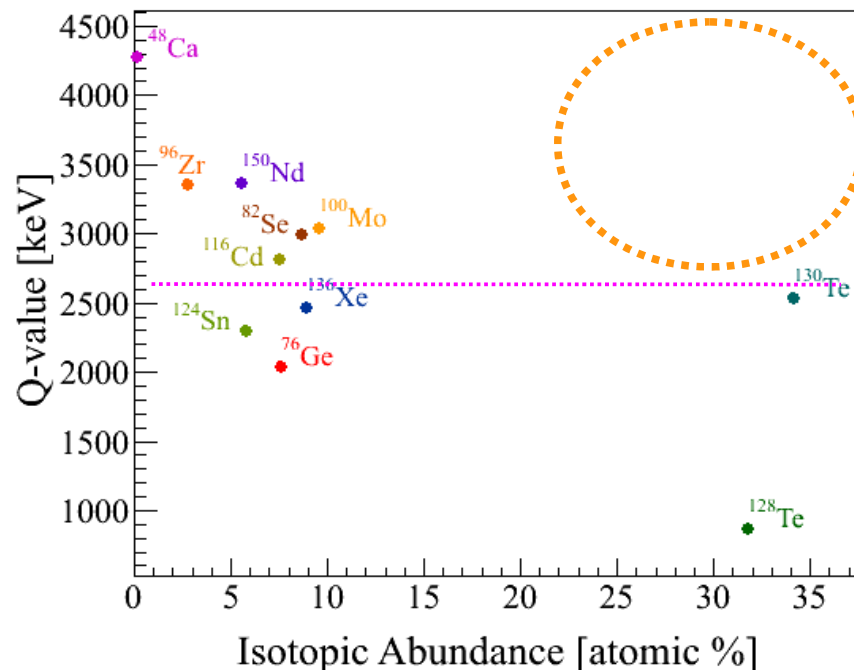
b: background [c/keV/kg/y] $\Rightarrow \sim 0.01$ c/keV/kg/y CHALLENGE

Isotope choice

The calorimetric technique allows a wide selection of possible DBDOv candidates:

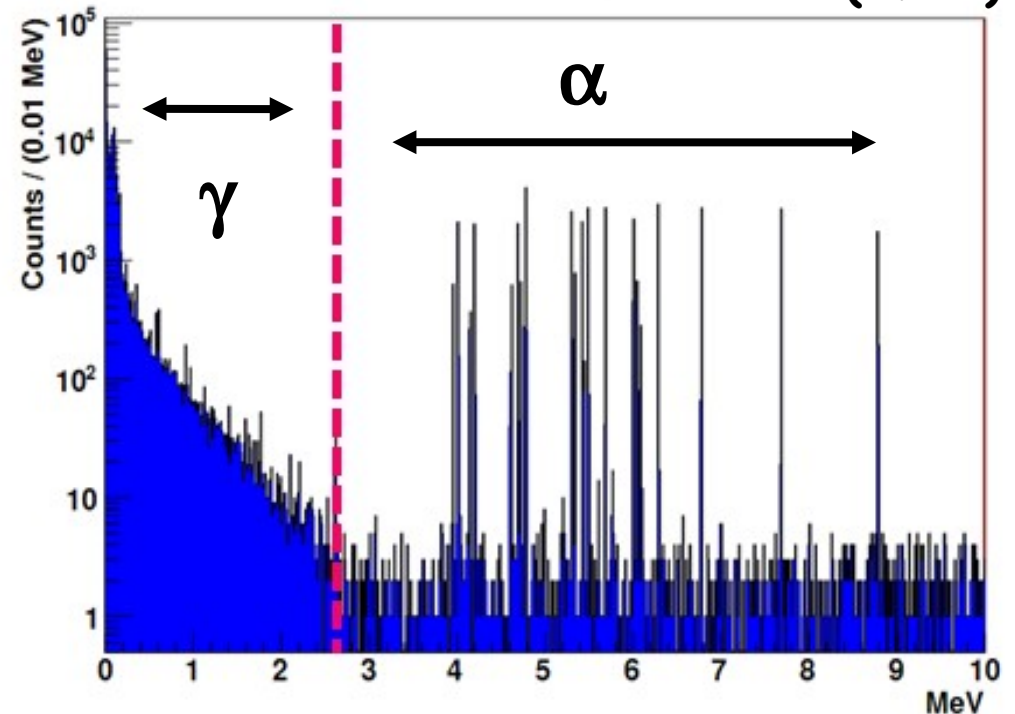
No golden isotope: High Q-value & High i.a.

$$S_{0\nu} \propto \varepsilon \frac{i.a.}{A} \sqrt{\frac{M \cdot T}{\Delta E \cdot b}}$$



²⁰⁸Tl highest natural
gamma line @ 2.6 MeV

Spectrum of a germanium bolometer calibrated
with natural radioactive sources (U/Th)



Bolometer background sources

Due to the limited detector energy resolution and the fact that they are sensitive to all types of radiation, there are various background sources for DBDOv experiment:

- Neutrons** =>
 - neutron activation: (n,g) reactions
 - appropriate shielding for n moderation and absorption
- Muons** =>
 - energy deposit in the ROI
 - installation under a mountain or in a mine / coincidence
- Gammas** =>
 - natural radioactivity / multiCompton events
- Betas** =>
 - appropriate isotope choice / material selection
- Alphas** =>
 - degraded alphas emitted by surfaces
 - surface cleaning / particle discrimination

CUORE & Cuoricino

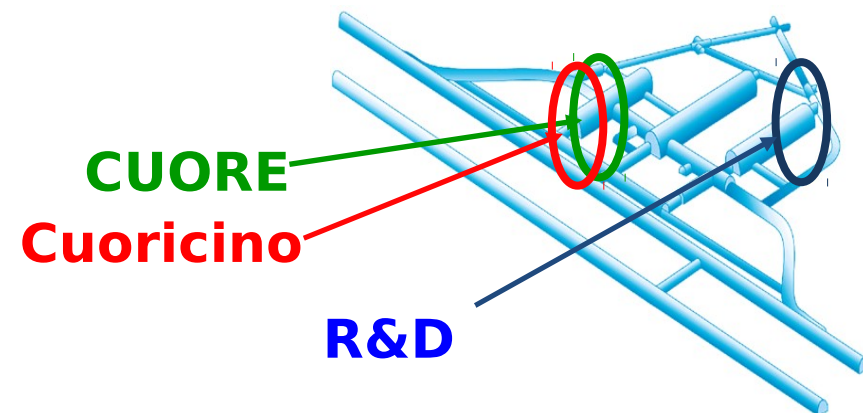
CUORE & Cuoricino use the bolometric technique to search for DBD0v in ^{130}Te

CUORE will be able to span low neutrino mass region (tens of meV)

Cuoricino was a small CUORE prototype, about 5 years of data taking starting in 2003

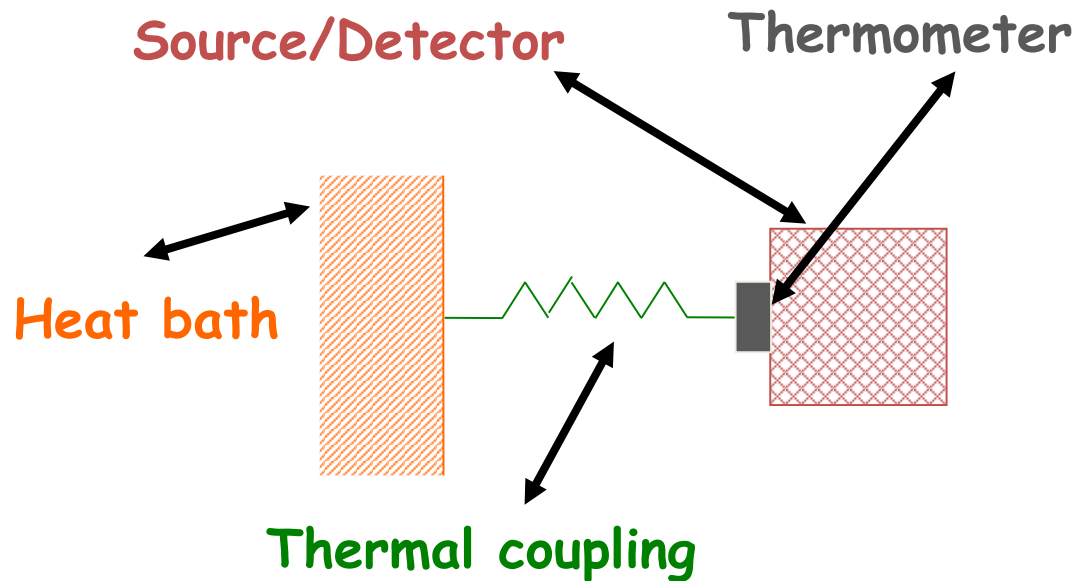
Experiments located underground in the Laboratori Nazionali del Gran Sasso @ 3400 m.w.e.

- Muon flux: $(3.2 \pm 0.2) 10^{-8} \mu/\text{s}/\text{cm}^2$
- Neutron flux: $10^{-6} - 10^{-7} \text{ n}/\text{s}/\text{cm}^2$
- Rn level: 50-100 Bq/m³



Bolometric Technique

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



e.g. TeO_2

$$\begin{aligned} M &= 750 \text{ g} \\ C &\cong 2 \text{ nJ/K} \\ &\text{@10mK} \\ \Rightarrow &1 \text{ MeV} / 100 \text{ } \mu\text{K} \end{aligned}$$

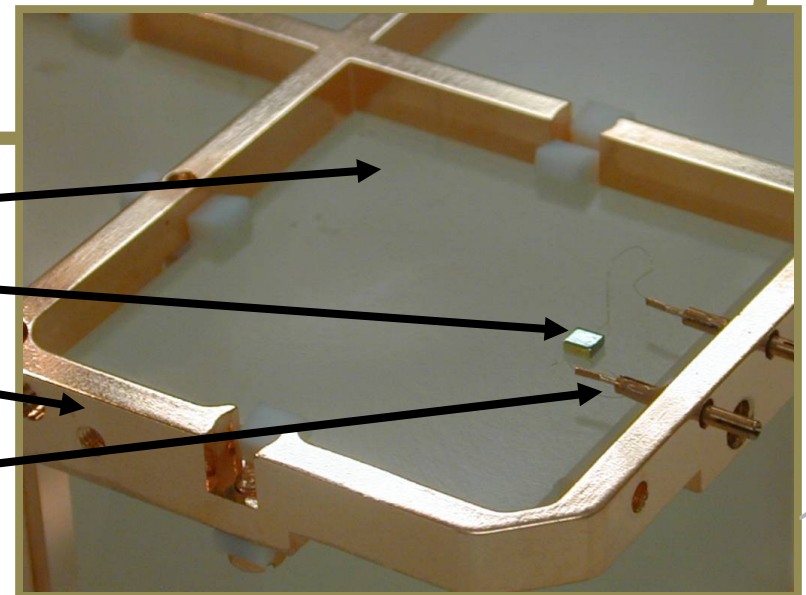
$$\begin{aligned} R &\cong 100 \text{ M}\Omega \\ dR/dT &\cong 0.1 \text{ M}\Omega/\mu\text{K} \\ \Rightarrow &1 \text{ mV} \end{aligned}$$

TeO_2 -> Source/Detector

NTD-Ge -> Thermometer

Copper -> Heat bath

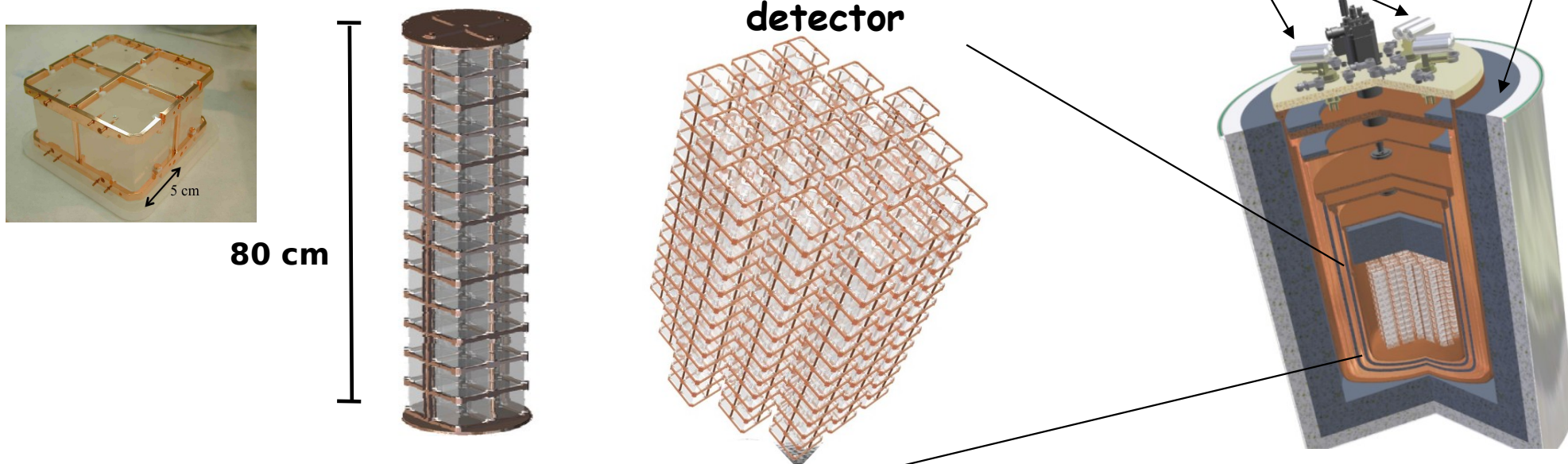
NTD-Ge gold wires -> Thermal coupling



The CUORE experiment

Cryogenic **U**nderground **O**bservatory for **R**are **E**vents

Array of 988 detectors. 19 CUORICINO-like towers $M = 0.741$ ton of TeO_2 (~ 200 kg ^{130}Te) to measure DBD0v of ^{130}Te with bolometric detector.



Sensitivity: $1.6 \cdot 10^{26}$ y [T: 5y; ΔE : 5 keV; b: 0.01 c/keV/kg/y]

$$m_{\beta\beta} = 35-82 \text{ meV}$$

NME from F.Simkovic et al. Phys.Rev. C77 - J.Suhonen et al. Int.Jou.Mod.Phys. E17 -
J.Menendez et al. Nucl. Phys.A818 - J.Barea et al. Phys. Rev. C79

Cuoricino

Cuoricino results :

- no evidence of DBD0v
- Statistics : $\sim 19.75 \text{ kg} \cdot \text{y}$ of ^{130}Te
- Bkg @ ROI: $0.18 \pm 0.02 \text{ c/keV/kg/y}$
- *FWHM @ 2615 keV (^{208}Tl) $\rightarrow 6.3 \text{ keV}$

Shieldings

Internal:

$\rightarrow$ 1cm low activity Pb
($A < 4 \text{ mBq/Kg}$ in ^{210}Pb)

External:

$\rightarrow$ 20cm Pb
 $\rightarrow$ 20cm Borated Polyethylene
 $\rightarrow$ Anti-Rn box: Nitrogen overpressure

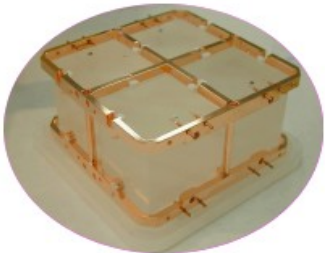
A tower of 62 TeO_2 crystals

11 floors made of 4 crystals

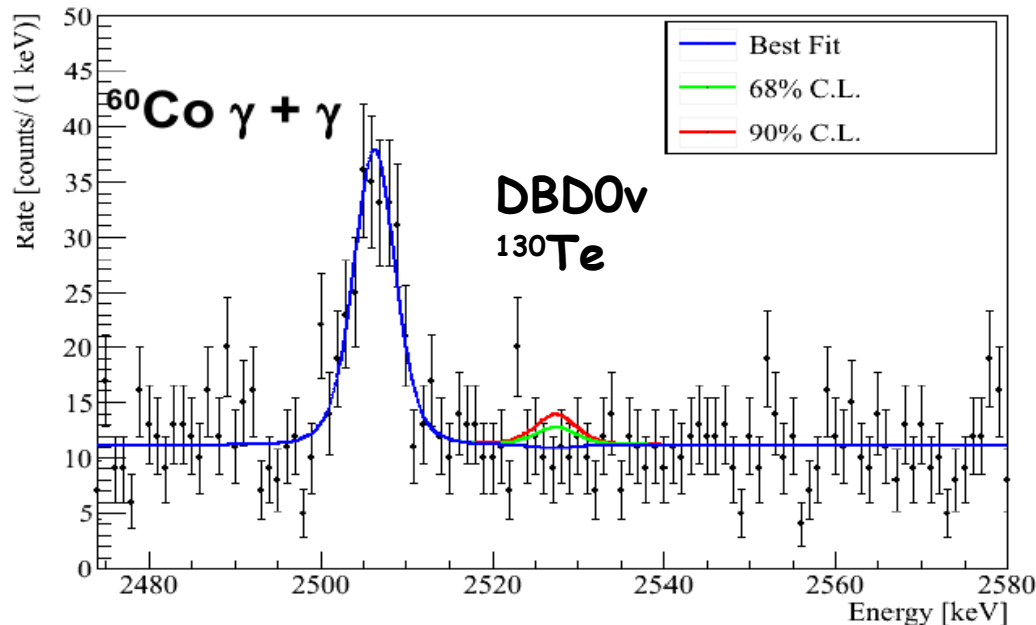
- not enriched
- Mass: 790g
- Dimensions: $5 \times 5 \times 5 \text{ cm}^3$

2 floors made of 9 crystals:

- Mass: 330g
- Dimensions: $3 \times 3 \times 6 \text{ cm}^3$
- 2 enriched in ^{128}Te (82%)
- 2 enriched in ^{130}Te (75%)



Total mass: 40.7 Kg (11.3 Kg in ^{130}Te)



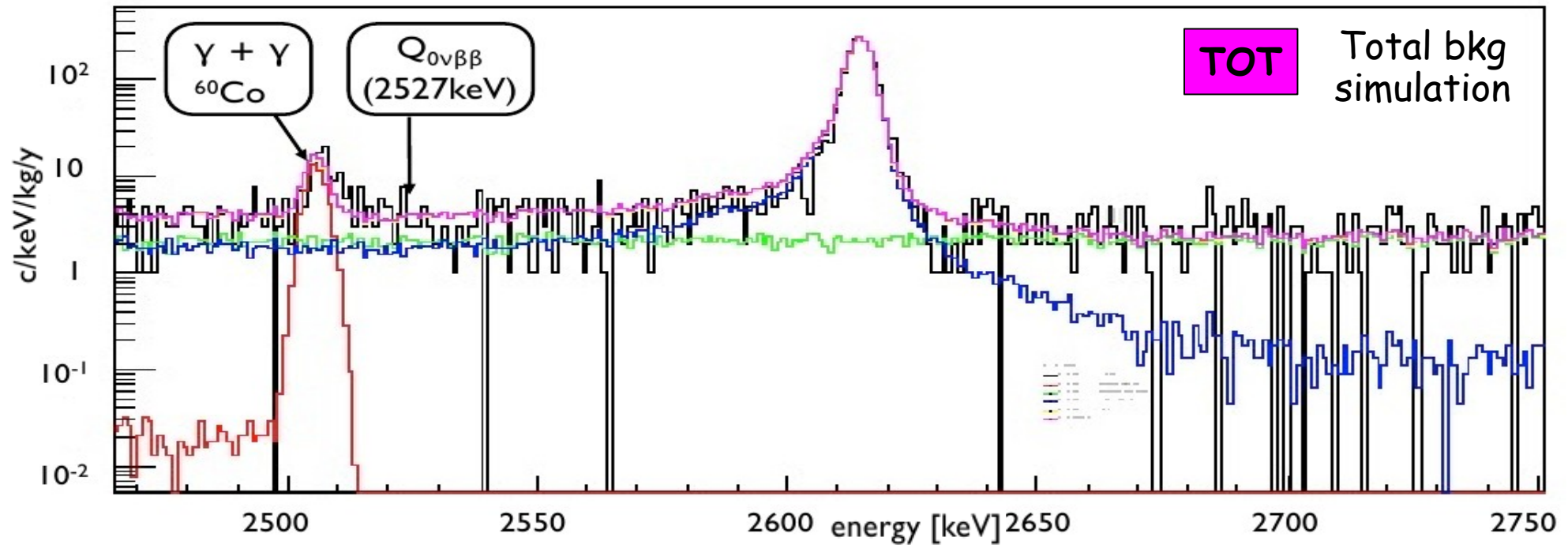
$$T_{1/2}^{0\nu} > 2.8 \cdot 10^{24} \text{ y @ 90\% C.L.}$$

Astropart. Phys. 34 (2011) 822–831 E. Andreotti et al. (LP)

$$m_{\beta\beta} \leq 0.3 - 0.7 \text{ eV}$$

- 1 Šimkovic et al.,
PRC 77 (2008) 045503
- 2 Civitarese et al.,
JoP:Conference series
173 (2009) 012012
- 3 Menéndez et al.,
NPA 818 (2009) 139
- 4 Barea and Iachello,
PRC 79 (2009) 044301

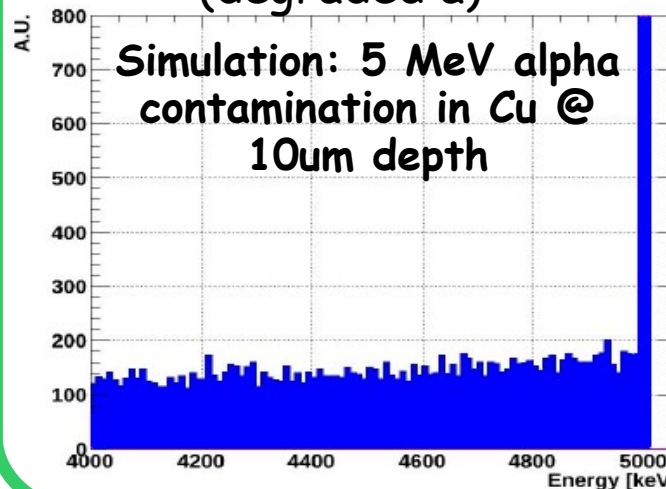
Cuoricino



35% ^{232}Th in
cryostat (g/b)

^{208}Tl multiCompton
+ Thorium betas

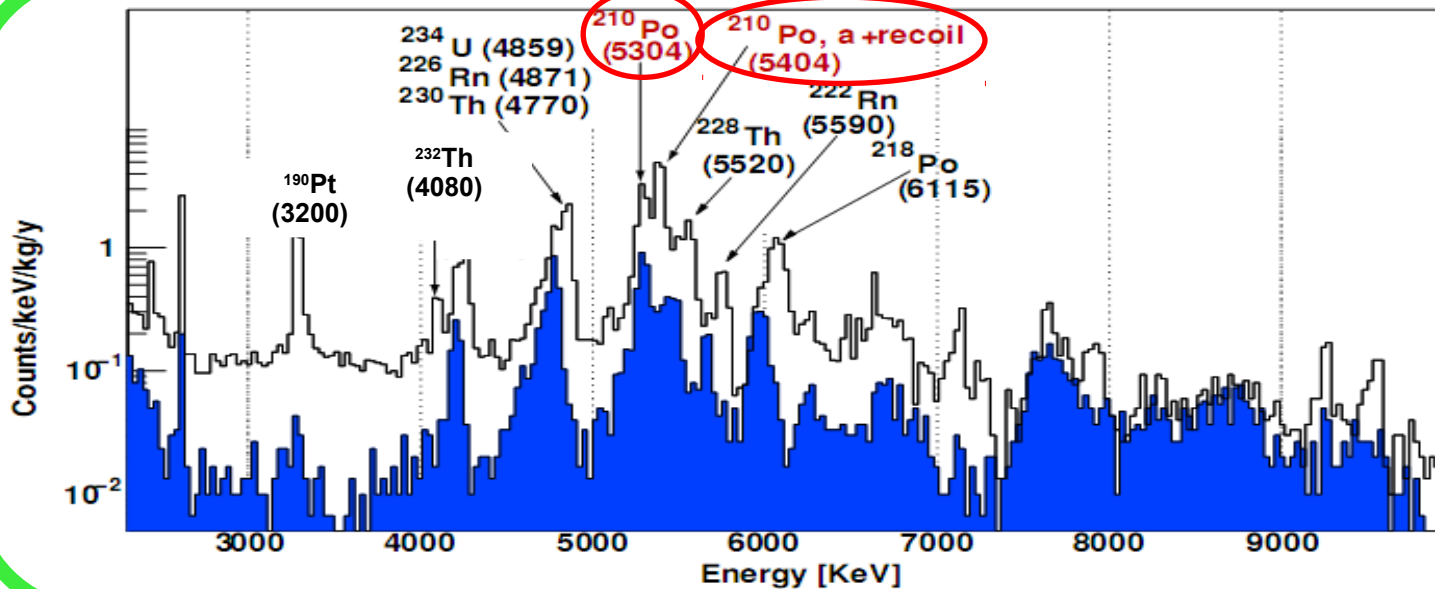
65% $^{232}\text{Th} + ^{238}\text{U}$ in Copper
and TeO_2 surfaces
(degraded a)



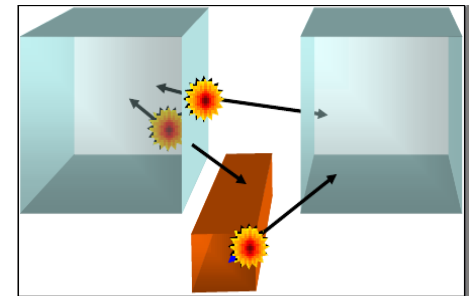
0% ^{60}Co in Copper

@ 2505 keV
Cosmogenic
activation

Alpha contaminations in Cuoricino



- Multiplicity 1 events
- Multiplicity 2 events



Various and delocalized alpha contaminations

- Production { • ^{190}Pt
- Cleaning { • ^{238}U and daughters
- Recontaminations { • ^{232}Th and daughters

- Bulk of crystals } 
- Copper surfaces } 
- Crystal surfaces }

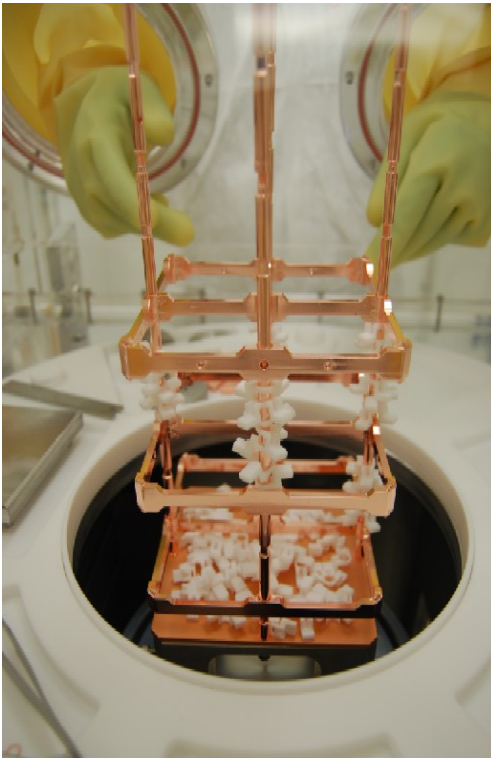
^{210}Po

Surface contamination studies

Copper cleaning R&D

TTT

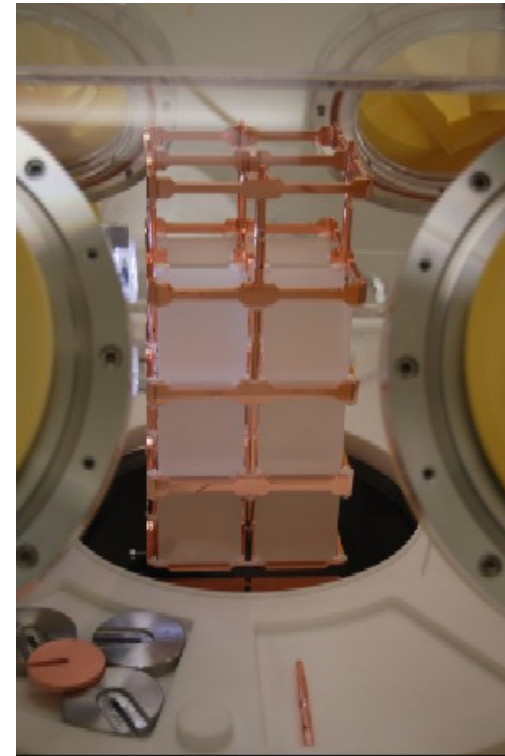
Specific measurements were performed in order to identify the best copper surface treatment for reducing the background in the ROI



TeO₂ surface radiopurity studies

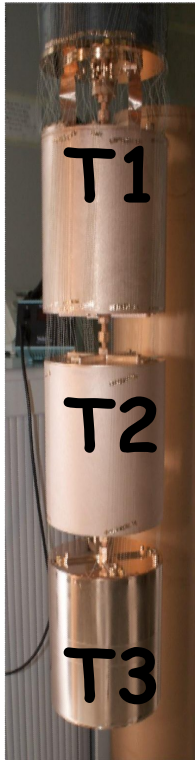
CCVR

Systematic control of bulk and surface contaminations of TeO₂ crystals



TTT detector

TTT (Three Tower Test): large mass detector for testing Cu surface contaminations with 3 different cleaning procedures inside Cuoricino cryostat (same bkg and operational conditions and reprocessed Cuoricino crystals)



- Test the final design of the CUORE single module (reduction of inert material facing the crystals);
- Test the best Cu "cleaning recepies";
- Bkg study and abatement;

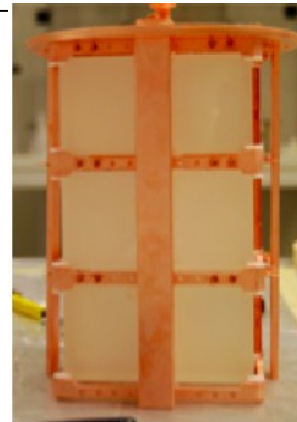
T1
Polyethylene
Cleaning:

- * Soap
 - * Passivation
 - * 7 layers (70um)
- Polyethelene complete coverage



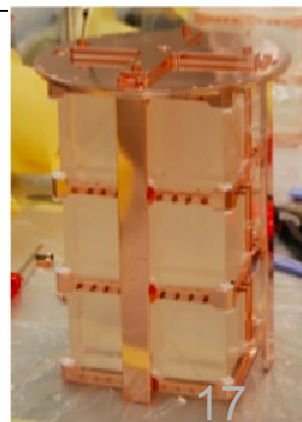
T2
Chemical New
Cleaning:

- * Soap
- * Acid electro erosion
- * Etching
- * Passivation



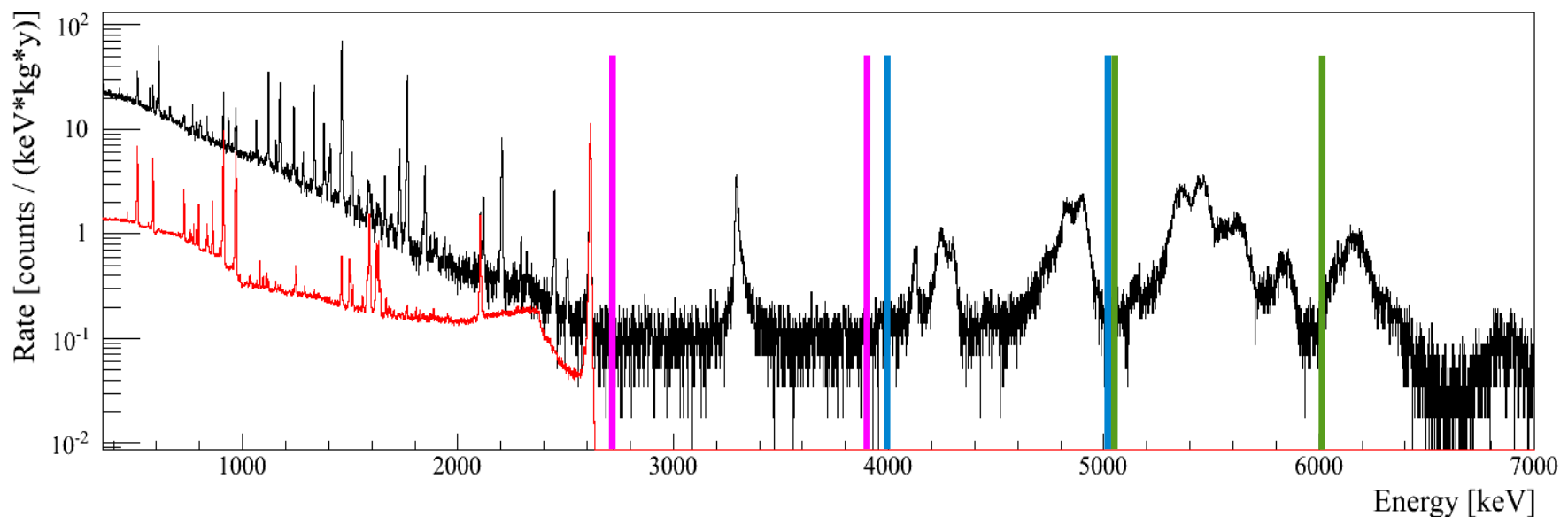
T3
Plasma
Cleaning:

- * Tumbling
- * Chemical etching
- * Electrochemical etching
- * Plasma



TTT detector performances and results

Tower	Live Time [day*crystal]	FWHM [keV] @ 2615 keV	* <u>[2.7-3.9] MeV</u> : in between g and a region. Suitable region for degraded a studies
T1	942.25	4.6	* <u>[4-5] MeV</u> : main a peaks of $^{232}\text{Th}+^{238}\text{U}$
T2	610.32	4.1	* <u>[5-6] MeV</u> : ^{210}Po (^{210}Pb) contaminations
T3	601.73	4.7	



TTT results

Tower	continuum 2.7-3.9 MeV	U/Th 4-5 MeV	Po/Pb 5-6 MeV
T1	0.068 ± 0.006	0.27 ± 0.01	1.25 ± 0.04
T2	0.120 ± 0.012	0.36 ± 0.03	1.82 ± 0.11
T3	0.072 ± 0.008	0.25 ± 0.02	1.80 ± 0.09
Cuoricino	0.116 ± 0.002	0.576 ± 0.003	0.962 ± 0.004

Rates
[c/keV/kg/y] for
the TTT and
Qino, no
coincidence cuts
are applied
on TTT

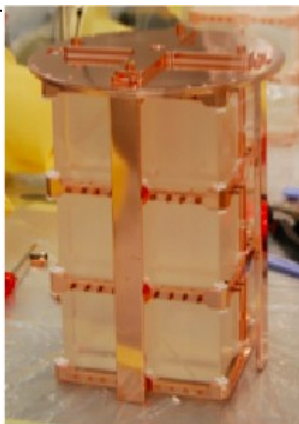
T1
Polyethylene
Cleaning:

- * Soap
 - * Passivation
 - * 7 layers (70um)
- Polyethelene complete coverage



T3
Plasma
Cleaning:

- * Tumbling
- * Chemical etching
- * Electrochemical etching
- * Plasma

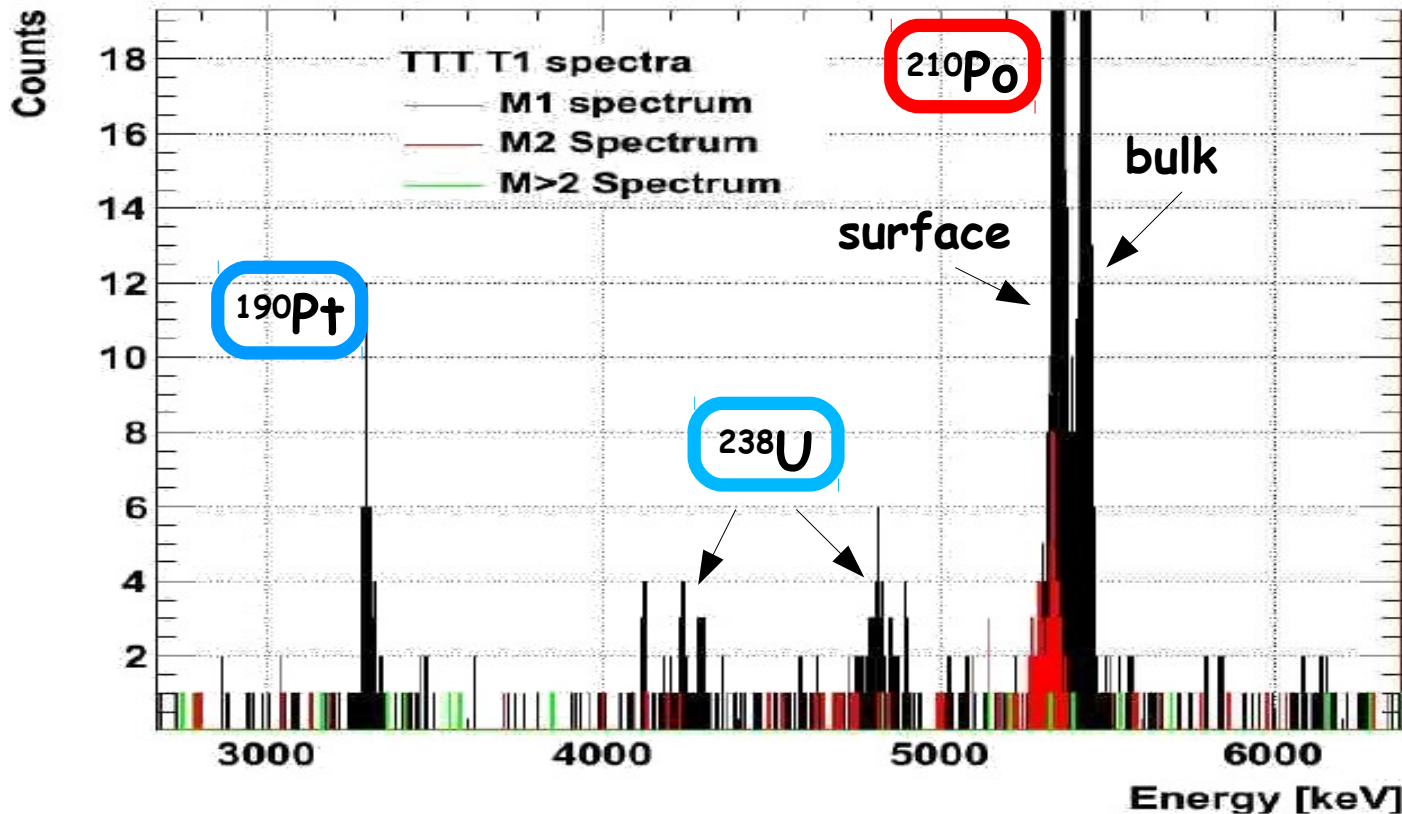


* T1 and T3 are compatible and show better bkg's compared to Cuoricino;

* Uranium and Thorium surface/bulk contaminations are reduced by a factor of ~2;

* No strong control of Polonium (Lead) re-contaminations

TTT-T1 results



* M1 shows ^{210}Po bulk crystal contaminations and surface contaminations (?crystals?, ?Copper/Polyethelene?)

* M2 tells us that crystals can explain a part of the overall ^{210}Po contamination. By exclusion Copper/Polyethelene will explain the rest

^{210}Po

CCVR: CUORE Crystals Validation Runs

The CCVRs are dedicated cryogenic set-ups planned to test the various batches of ready-to-use produced crystals.

CUORE crystal production started
in 2008 in Jaiding (CHINA):

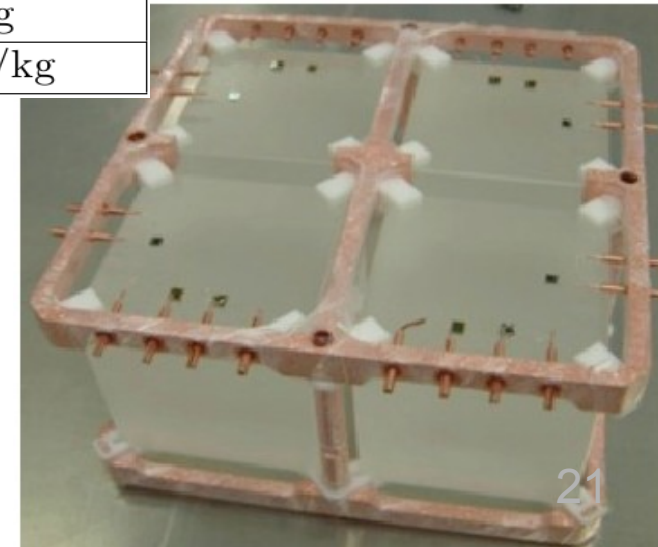
- * ~30 crystals/month
- * 700 crystals already delivered

- * Bolometer performances: essentially energy resolution
- * Bulk a contaminations
- * Surface contaminations

Isotope	Allowed Contamination
^{238}U	$< 3 \cdot 10^{-13} \text{ g/g}$
^{232}Th	$< 3 \cdot 10^{-13} \text{ g/g}$
^{210}Pb	$< 1 \cdot 10^{-5} \text{ Bq/kg}$
^{210}Po	$< 0.1 \text{ Bq/kg}$
^{40}K	$< 1 \cdot 10^{-3} \text{ Bq/kg}$

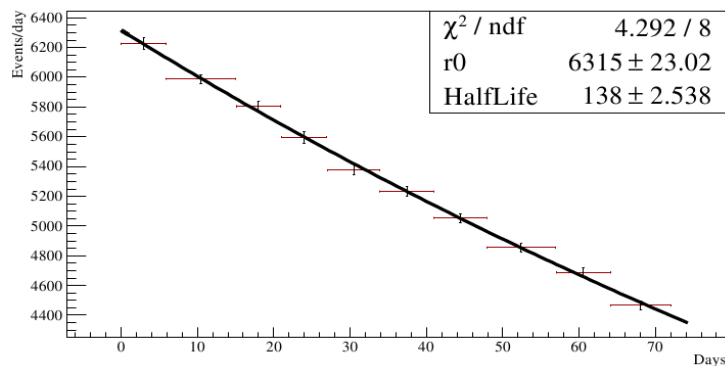
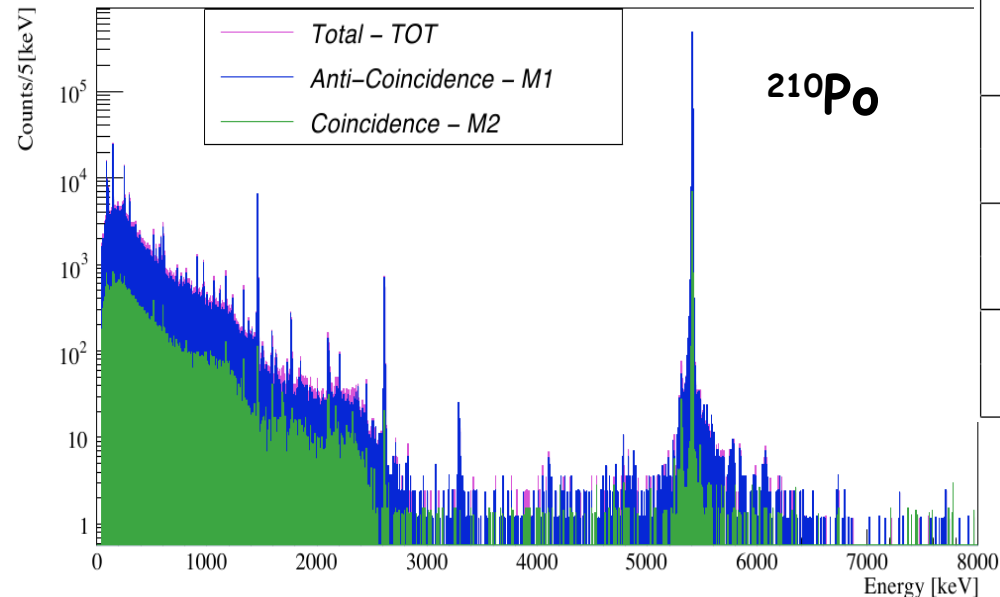
Up to now 7 CCVR tests were performed.

Detector	Livetime [d]	Crystal tested
CCVR1	59.9	7, 11, 39, 41
CCVR2	19.4	7, 11, 76, 97
CCVR3	43.05	180, 190, 229, 236
CCVR4	25.8	313, 340, 354, 380
CCVR5	30.3	416, 421, 436, 455



CCVR results

Total Live Time: 713 days
Total Mass: 15 kg of TeO₂



		Continuum (2700, 3900) [keV]	U/Th (4000, 5000) [keV]	²¹⁰ Po (5000, 6000) [keV]
CCVR	M1	0.09±0.02	0.13±0.01	-
	M2	0.015±0.007	0.014±0.003	-
TTT	M1	0.052 ±0.008	0.28±0.02	1.30±0.07
	M2	0.009±0.003	0.0018±0.005	0.09±0.01
Cuoricino	M1	0.104 ±0.002	0.522±0.003	0.846±0.004
	M2	0.009± 0.001	0.084±0.001	0.173±0.002

*** Continuum higher in CCVR-M1 because of ²¹⁰Po tail (from MC simulation is 20%)**

*** CCVR has lower bulk and surface U/Th contaminations compared to Qino**

*** High ²¹⁰Po contamination (vanishing). Po and Te are chemically affined;**

CCVR final results

Total CCVR bulk contaminations

Chain	Nuclide	Upper limit [Bq/kg]	Upper limit [g/g]
^{238}U	^{238}U	2.5E-07	2.0E-14
	^{234}U	4.7E-07	3.6E-14
	^{230}Th	5.7E-07	4.4E-14
	^{226}Ra	6.7E-07	5.3E-14
	^{218}Po	1.6E-07	1.3E-14
^{232}Th	^{232}Th	1.3E-07	3.1E-14
	^{212}Bi	8.4E-07	2.1E-13

Total CCVR surface contaminations

Depth	Nuclide	Upper limit 90% C.L. [Bq/cm ²]
1 μm	^{238}U	3.7E-09
	^{226}Ra	8.9E-09
	^{232}Th	1.9E-09

Total ^{210}Pb contamination:

Bulk 3.3E-6 Bq/kg
Surf @ 1 μm 9.2E-9 Bq/cm²

(Qino reprocessed) TTT-T1 Surf @
1 μm 6.1E-8 Bq/cm²

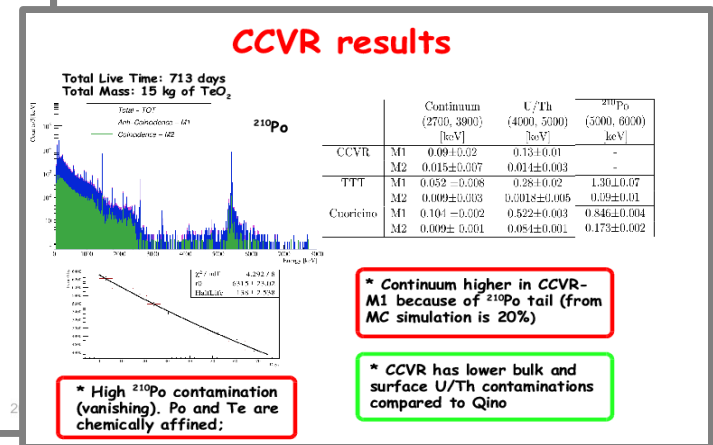
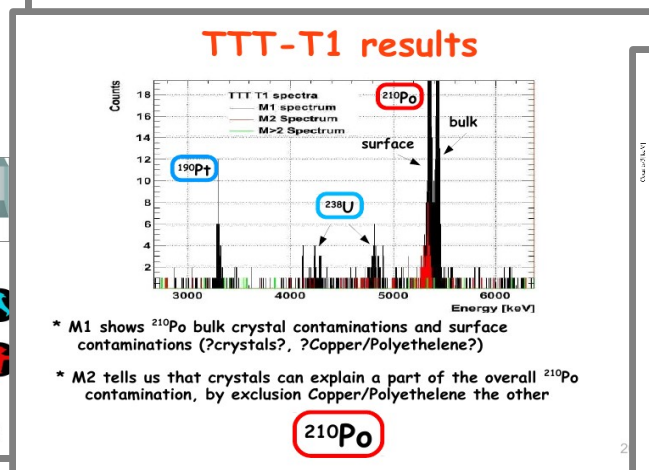
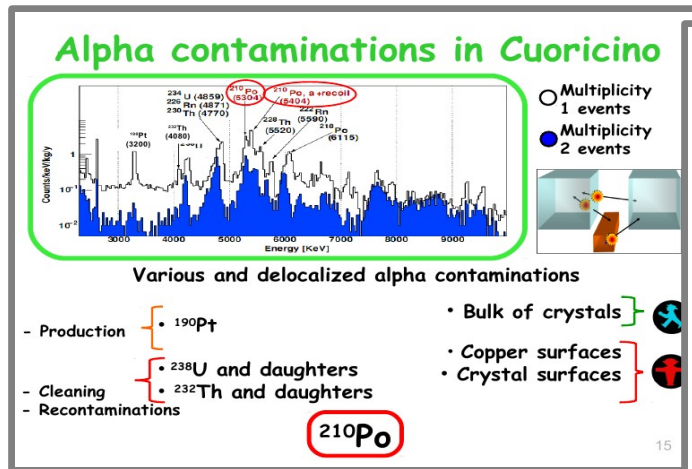
Qino: $\langle \text{FWHM} \rangle_{2615 \text{ keV}} = 6.3 \pm 2.5 \text{ keV}$

CCVR: $\langle \text{FWHM} \rangle_{2615 \text{ keV}} = 4.6 \pm 1.9 \text{ keV}$

CCVR: $\langle \text{FWHM} \rangle_{5.4 \text{ MeV}} = 5.6 \pm 2.1 \text{ keV}$

Total contribution to CUORE in the
ROI: $5.5 \cdot 10^{-3} \text{ c/keV/kg/y}$

Qino + TTT + CCVR results



* Surface contaminations are a limitation to the sensitivity of low background experiments.

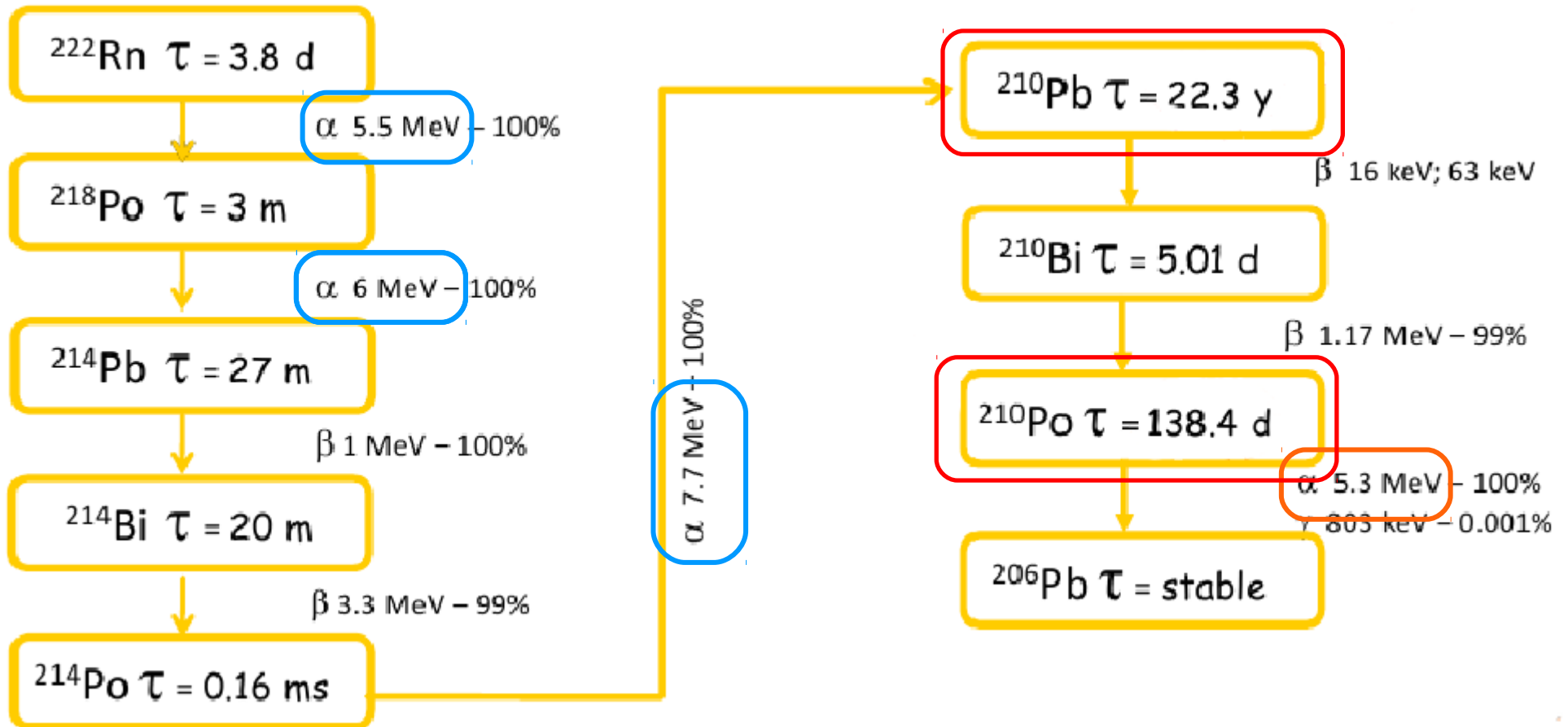
* The main source of re-contamination for Copper (TTT) and TeO_2 crystals (CCVR) seems to be induced by ^{210}Po (^{210}Pb daughter).

^{210}Pb and ^{210}Po are produced by ^{222}Rn decay

Is ^{222}Rn a direct background source for DBD0v bolometric experiments ?

Can a detector exposed to ^{222}Rn be contaminated ?

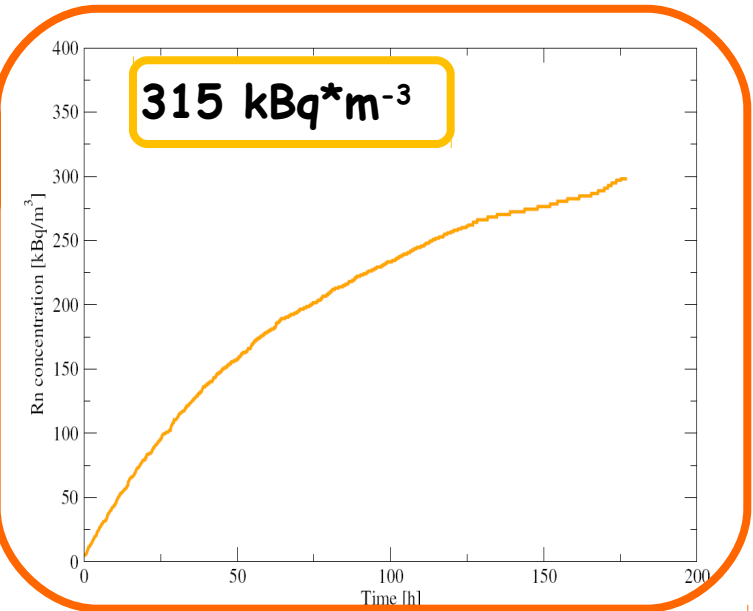
Radon decay chain



Rn-induced contaminations (1)



Rn concentration in the radon-box

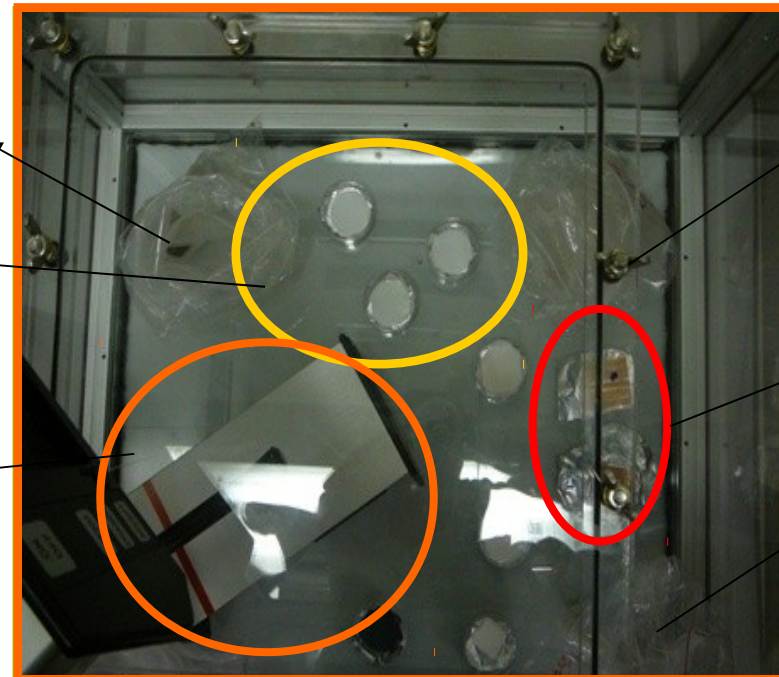


Sample	Material	Exposure [d]
Cu_OFHC	Cu	56
Rame_BaseMen	Cu	63
TeO2_1Mis	TeO ₂	49
TeO2_2Mis	TeO ₂	73
Cu_OFHC_brv	Cu	16
TeO2_brv	TeO ₂	14

²³⁸U source

TeO₂ slabs

Rn detector



²³⁸U source

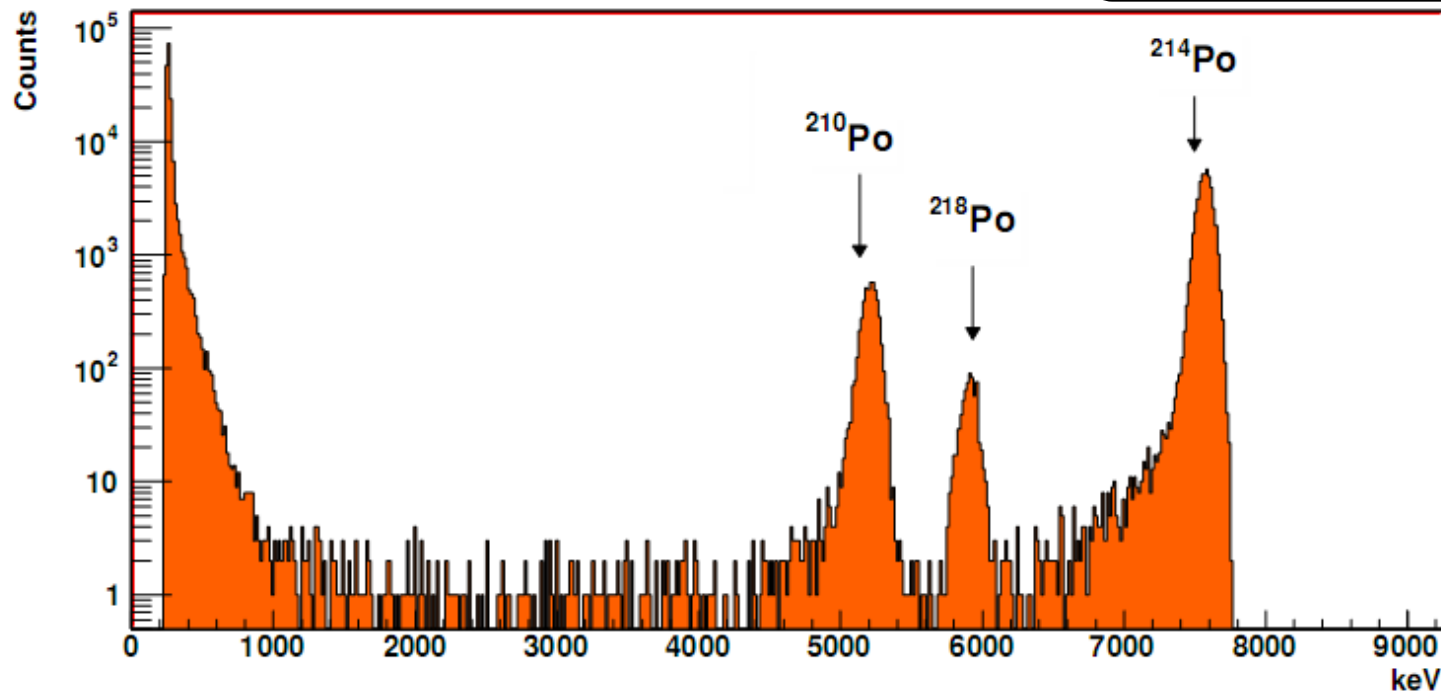
Cu slabs

²³⁸U source

Rn-induced contaminations (2)

Which contaminants?

Copper sample



Detector:

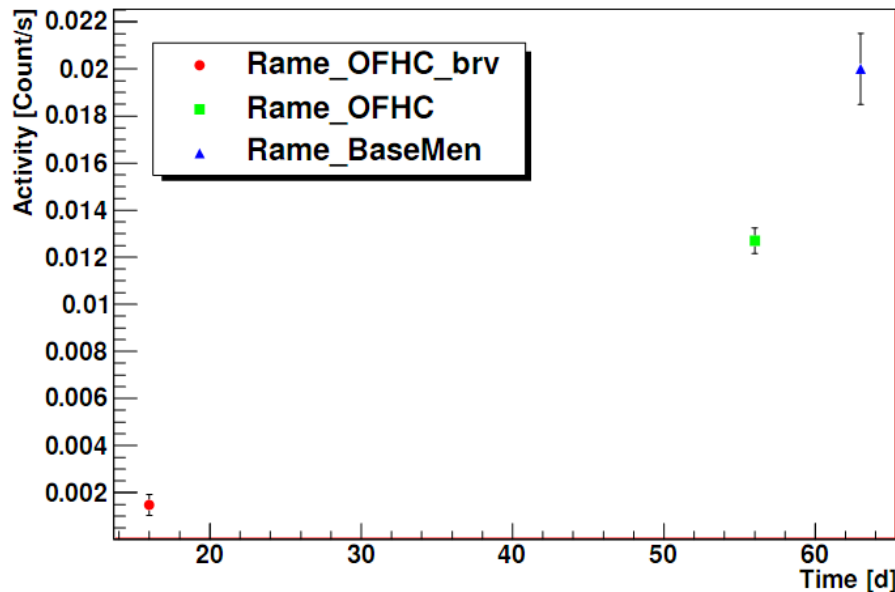
SBD of 1200 mm²
in vacuum (7 μm Hg)

- Detector exposure to Rn directly induces Polonium contaminations
- Radon does not diffuse inside the sample

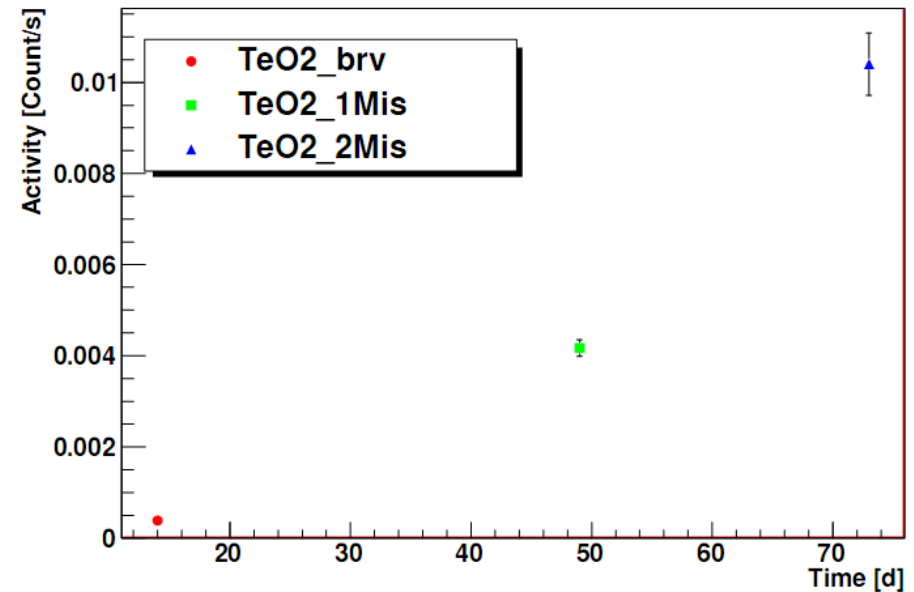
Analysis of contaminants

^{210}Po activity evaluated under the peak

Copper samples activity after ^{222}Rn exposure



TeO_2 samples activity after ^{222}Rn exposure



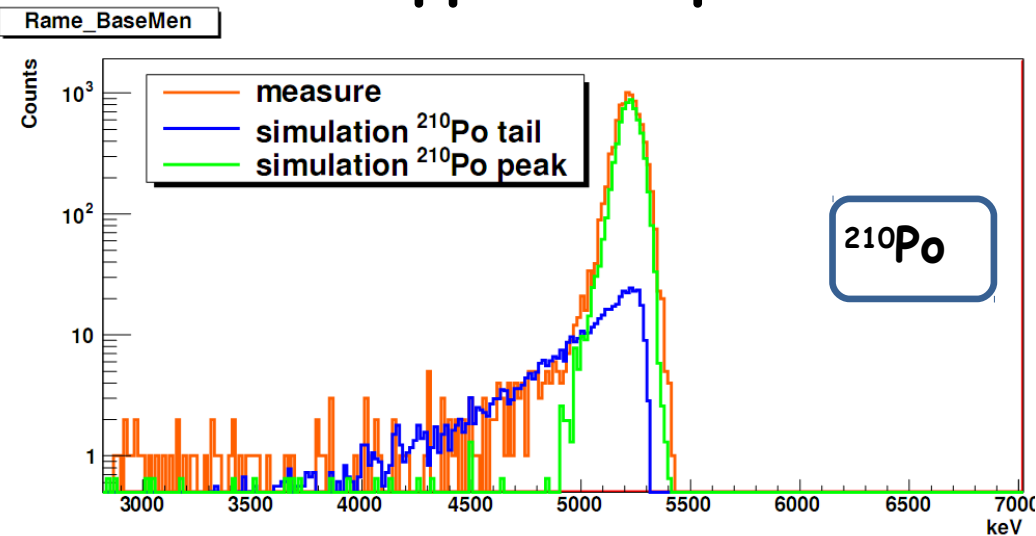
Eur. Phys. J. C (2011) 71:1805 M. Clemenza et al. (LP)

^{210}Po contaminations are strictly related to the time exposure of the samples (Cu & TeO_2) to Radon.

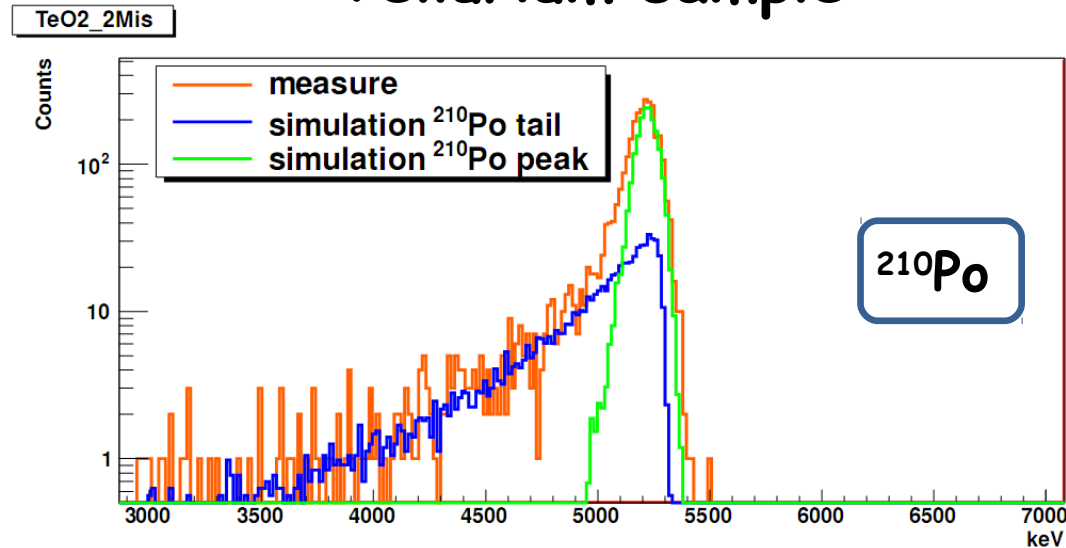
N.B.: ^{210}Po ($T_{1/2}=138$ d) is not in equilibrium with ^{210}Pb ($T_{1/2}=22.4$ y)

^{210}Po diffusion

Copper sample



Tellurium sample



$$Y = A \cdot \exp(x \cdot \Lambda)$$

Sample	Material	Exposure time	Diffusion depth
Rame_BaseMen	Copper	63 days	430 ± 20 nm
Rame_OFHC	Copper	56 days	410 ± 20 nm
Rame_OFHC_brv	Copper	16 days	***
TeO2_2Mis	Tellurium Oxide	73 days	940 ± 20 nm
TeO2_1Mis	Tellurium Oxide	49 days	500 ± 20 nm
TeO2_brv	Tellurium Oxide	14 days	***

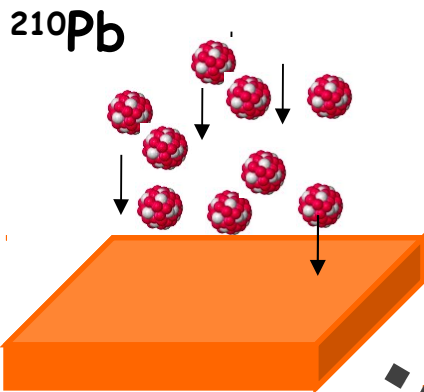
^{214}Po and ^{210}Po diffusion in TeO_2 and in Cu

Who produces ^{210}Po ?

$$A = \frac{N_1}{\tau_1} = \frac{N_2}{\tau_2} = \frac{N_3}{\tau_3} = \dots$$

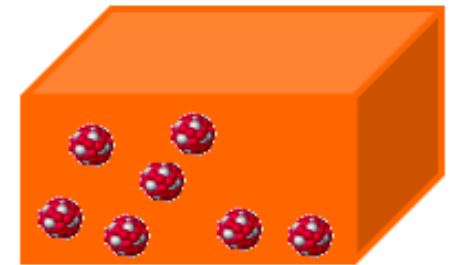
An element concentration in a closed system is proportional to its life-time

Prompt Pb production mechanism



Delayed Pb production mechanism

^{218}Po & ^{214}Po



^{210}Pb is the element most concentrated in the Rn-box

^{210}Po $\tau = 138.4$ d

Detectable during the prompt ^{210}Po decays

Detectable during the delayed ^{210}Po decays

^{210}Pb contaminations

^{210}Pb evaluation starting from ^{210}Po contamination
“prompt” ($t \sim \text{h}$) and “delayed” ($t \sim 300 \text{ d}$)

$$A_{Po} = A_{Pb}^0 \frac{\lambda_{Pb}}{\lambda_{Po} - \lambda_{Pb}} (e^{-\lambda_{Pb}t} - e^{-\lambda_{Po}t})$$

$$\frac{\text{Delayed } A_{Pb}^0}{\text{Prompt } A_{Pb}^0} \sim 6$$

~ 85% of ^{210}Pb contamination is generated by delayed decays of Rn fast daughters (^{218}Po , ^{214}Po).

=> Informations need for the design of a clean room
It is important to reduce Rn concentration more than the Pb one. 31

Sticking Factor

The Rn sticking factor is evaluated on ^{210}Pb (starting from the ^{210}Po contaminations):

$$\Sigma = \frac{A_{210\text{Pb}}^0 \cdot \tau_{210\text{Pb}}}{\Gamma \cdot S \cdot t_{\text{exp}}}$$

Γ : Rn nuclei flux

S : sample surface

t_{exp} : time exposure to Rn

Sample	Material	Exposure time	Σ_{Rn}
Rame_BaseMen	Copper	63 days	$8.39 \cdot 10^{-9} \pm 4.53 \cdot 10^{-10}$
Rame_OFHC	Copper	56 days	$3.15 \cdot 10^{-9} \pm 3.15 \cdot 10^{-11}$
Rame_OFHC_brv	Copper	16 days	$2.32 \cdot 10^{-9} \pm 1.25 \cdot 10^{-10}$
TeO2_2Mis	Tellurium Oxide	73 days	$1.69 \cdot 10^{-9} \pm 5.36 \cdot 10^{-11}$
TeO2_1Mis	Tellurium Oxide	49 days	$1.41 \cdot 10^{-9} \pm 9.46 \cdot 10^{-11}$
TeO2_brv	Tellurium Oxide	14 days	—

Effects on CUORE

Effects on CUORE sensitivity due to an exposure of Cu to Radon

Cu exposed for 1 year @ 100 mBq/m³

Σ for Cu = 1.85×10^{-4} @ 315 kBq/m³

Hypothesis : Linearity @ 100 mBq/m³



$$A_{po \text{ Total}} = 1.1 \times 10^{-7} \text{ Bq/cm}^2$$



Monte Carlo simulation

$$B \leq 3.4 \times 10^{-3} \text{ counts/keV/kg/y}$$



@ 10^{-3} Counts/keV/kg/y

Is there a way out ?

Polonium (Pb) is one of the source of surface contaminations, but does not explain the background in the ROI.

Even if we completely understand the mechanisms inducing surface contaminations (degraded alpha), is it really possible to prevent re-contaminations of radiopure materials?

Will we ever have a 0 alpha background?

SOLUTION

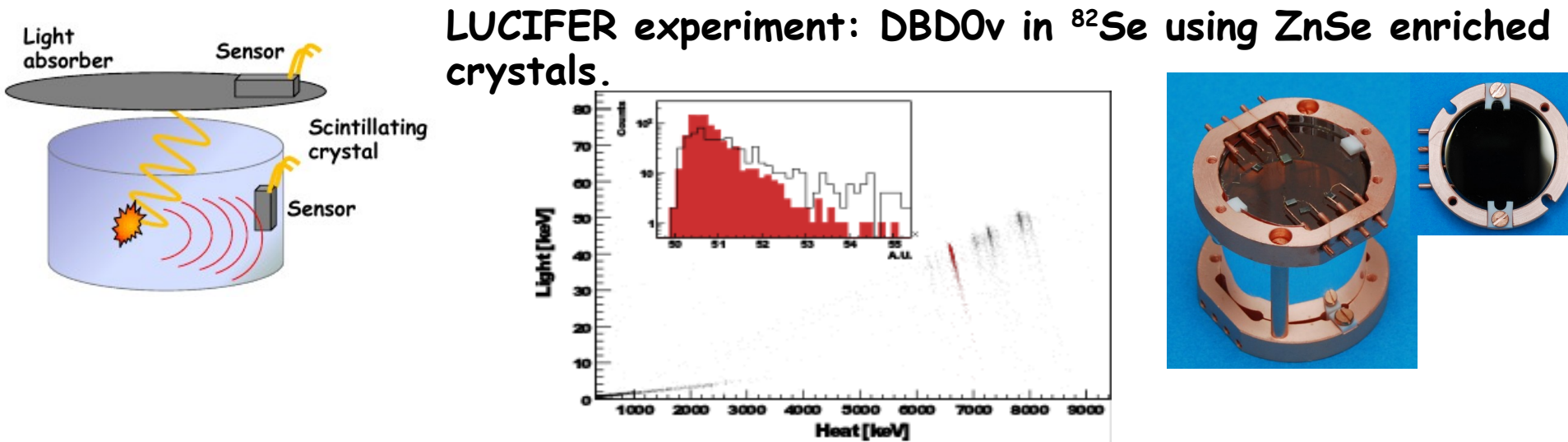
α / β discrimination

Reduce the alpha background in the DBD0v
ROI to a 0-level

α / β discrimination

There are different proposed solutions:

* Scintillating bolometers: combination of heat and light for particle interaction discrimination:



* Cherenkov light emission: [arXiv:1106.6286]: Discrimination of alpha and beta/gamma interactions in a TeO_2 bolometer.

* Pulse shape analysis: identification of α and β interaction in the detector through pulse shape analysis of signals.

ZnMoO₄ bolometers allow α / β discrimination just on the heat signal

[arXiv:1011.5415] Gironi et al.

Conclusions

- * Polonium-210 (Lead-210) are the most intense source of surface contaminations for DBD0v bolometric experiments;
- * Radon-induced contaminations are one mechanism of surface contaminations, but they do not explain the whole problem;
- * ^{222}Rn It is real daughters, but they do not explain the whole problem;
- * Particle discrimination seems to be a solution to the problem, but it still needs to be proved the scale to large mass detector;

The end

