Nuclear emulsion with molybdenum filling for 2β -decay observation



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Application of the nuclear emulsion as a detector surrounding the 2β -decay target

□ J.H.Fremlin et al. Pros.Phys.Soc. V.65, p.911 (1952) For 16 isotopes the lower limits were measured relative to (0v) – mode ~10¹⁵ – 10¹⁷ years.

□ A.S.Barabash et al. Preprint ITEP 104 – 88 (1988) The limit for $2\beta(2\nu)$ from ${}^{96}Zr > 0.7x10^{17}$ years was established.

Proposed method

 \checkmark

- In the stage of production the nuclear emulsion is filled with fine powder of the required isotope. Thus, the nuclear emulsion is simultaneously target and detector
- The main advantage of this approach to the 2β-decay study is the visualization of events and the possibility of all the decay characteristics measurement: the total energy, the single electron energies
 - and their angle of divergence.

Testing of proposed method

R&D with use of molybdenum fine powder of industrial production were held.

There were made 10 plates (9x12x0.0075) cm³, Vem =8.1cm³ with 1.43 grams Mo (4.6% of the dry emulsion weight).

Testing of proposed method

ESTABLISHED

Molybdenum does not "spoil" of the nuclear emulsion properties and the powder does not interfere with scanning and measurements on microscope



Fig.1. Emulsion image with different size of Mo grains

In the emulsion polymerization process, the partial deposition of large Mo grains take place in the field of gravity



Testing of proposed method

- It is necessary to eliminate Mo grains larger than 6-8 microns and periodically turn over the emulsion layers while drying.
- **II**. Visual evaluation shows that that the emulsion charge with molybdenum powder can be increased in 1.5-2 times.

Estimation for the experiment with 1 kg of Molybdenum

- For 1 gram of Mo it should be taken 5.6 cm³ of dry emulsion;
- > For 1 kg. \rightarrow 5.6 liters (21.3 kg. of dry emulsion).
- > For 1 liter of dry emulsion \rightarrow 115 kg. of gel.
- > For test with 1 kg of $^{100}Mo \rightarrow 65.5$ kg of gel.
- > 860 emulsion layers (9x12x0.06) cm³ with filling increased by factor of 1.5-2 give
 570 – 430 emulsion layers (10 -12 emulsion chambers).
- Measurements on three units with scanning speed 1 layer per day will take for about a year.

Background estimation

- With zero background and 70% detection efficiency for $0\nu\beta\beta^{100}$ Mo during one year one can get result ~1.5x10²⁴ years. The first stage of the test: ~ 100 gr. ¹⁰⁰Mo for background estimation and measurement of ~10³ 2v2\beta-decays.
- Nuclear emulsion does not have temporal resolution. Consecutive, not simultaneous escape of 2 electrons from grain will simulate 2β decay of ¹⁰⁰Mo.

Background estimation

Bad purification of ¹⁰⁰Mo and two decays of ⁴⁰K in gelatin near the grain ($T_{1/2} = 1.28 \times 10^9$ years, $E_{\beta} = 1.312$ MeV)

⁹⁰Sr decay near the grain ⁹⁰Sr₃₈ \rightarrow ⁹⁰Y₃₉ +e⁻+v_e (T_{1/2} =28.8 years, E_e = 0.549 MeV) ⁹⁰Y₃₉ \rightarrow ⁹⁰Zr₄₀ +e⁻+v_e (E_e = 2.28 MeV)

Total energy 2.829 MeV.



Background estimation



Fig.4. Image of real emulsion with Mo-conglomerates and simulation of flight two electrons with different energy value

There were considered positron-nuclear interactions in which produced relativistic particles are escaping from the interaction point with various angles. The pairs of particles with a divergence angle $\boldsymbol{\varphi}$ were selected and precision of their trajectory convergence to the interaction point were determined (schematically looks as intersecting straight lines). As the intersection point "**d**" the minimum distance between them is taken.

Position of "d" within the angle φ does not exceed the limits of conglomerate (grain of Mo) and 80% of "d" is concentrated in the area of ~1micron. <d>=(0.60±0.03) micron is about the emulsion grain size, and no correlations between d and φ are found.

Fig.4. Distribution of value d, minimal distance between two tracks, generated the particles from point of collision positron and emulsion nuclei



Fig.5. Distribution of value d, minimal distance between two ¹⁸⁰ tracks, with regard to angle φ_{160} between two tracks



WE assume that grain size is <R_k>=3 micron and its "danger area" is about ~d (0.6 micron).

In this case, the exposure with 1 kg of $^{100}\text{Mo}(5.6 \text{ liters of emulsion})$ the number of background decays will be suppressed by a factor of $\sim 1.5 \times 10^{-2}$

Clean up potassium up to $\sim 10^{-8}$ g/g gives the number of decays 40 K in a "dangerous zone" $\sim 0.7 \times 10^{-5}$ decay/year*grain and two electron observing $\sim 5 \times 10^{-11}$. This probability should be reduced by the accuracy of determining the escape of two electrons from one point.

Thus, due to 40 K we have $\beta\beta$ -decay: ~ 5 events a year in exposure of 1 kg 100 Mo and less than one event in the energy range of (3±0.3) MeV.

In the strontium disintegration both electrons are emitted from a single point and can be perceived as $\beta\beta$ -decay particles.

When the strontium activity is ~1 mBq/kg and the amount of emulsion is 21.3 kg (1 kg of 100 Mo), about 1 event a year can be detected in the "danger area" with the energy of electrons (~0.5 and ~2.3 MeV),

> 2.8 MeV (with energy resolution 10%).

This background can be reduced due to the "symmetrical" $\beta\beta$ -decay.

Background from natural radioactive elements: Thorium, Uranium, Radium and Actinium series

In a typical, not extreme, case **1 cm³** of emulsion contains about 20 decays, forming **3-5 ray stars of** α-particles.

The decay chain always begin with a successive emission of α -particles, and only at the end of the chain the elements emitting β and gamma particles arise.

Path of α -particles in emulsion is **10 – 50 micron** (E_{α} = 3 ÷ 9 MeV). The α -stars are detected in the emulsion with about ~100% efficiency.

"Double events": grain of 100 Mo and α -star allow to exclude the background of electrons from natural radioactive elements.

Emulsion "star" from consecutive α-decay of Thorium radioactive elements



Nuclear Track Emulsion Workshop Romania 2013

SUMMARY

Result

10 – 12 emulsion chambers, filled with 1 kg 100 Mo, are processed on three scanning microscopes for 1 year. The expected result of the decay period measurement is ~1.5x10²⁴ years.

Background (ββ)

Background from ⁴⁰K is ~5events/year and <1 event in the region (3±0.3) MeV.

Background from ⁹⁰Sr is about ~1 event/year with the electron energy >2.8 MeV. Background from natural radioactivity is almost completely excluded.

Accuracy of energy measurement of charged particles by their range in nuclear emulsion. The energy of muons from $\pi \rightarrow \mu \nu$ decay is monochromatic one. According to our measurements $\mathbf{F} = (4, 12 \pm 0, 1)$ MeV

 $E_{\mu} = (4.12 \pm 0.1) \text{ MeV}$

Thank vou for attention



Background slides

Next generation experiments

 Main goal: Reaching sensitivity
 ~0.01 - 0.1 eV
 Strategy: investigation more than one isotopes > 2-3; use different strategy

Experiments are going to be realized in the nearest ~ 3÷10 years

CUORE (¹³⁰Te)
 GERDA (⁷⁶Ge)
 MAJORANA (⁷⁶Ge)
 EXO (¹³⁶Xe)
 SuperNEMO (⁸²Se)
 KamLAND (¹³⁶Xe)
 SNO+ (¹⁵⁰Nd)



GERDA - Majorana





- · 'Bare' enrGe array in liquid argon
- Shield: high-purity liquid Argon / H₂O
- Phase I (~ 2011): ~ 18 kg (HdM/IGEX diodes)
- Phase II (~ 2013): add ~ 20 kg new detectors Total ~ 40 kg



- Modules of enrGe housed in high-purity electroformed copper cryostat
- Shield: electroformed copper / lead
- Initial phase: R&D prototype module (~ 2013) Total ~ 40 kg

Joint Cooperative Agreement:

- Open exchange of knowledge & technologies (e.g. MaGe, R&D)
- Intention to merge for 1 ton exp. Select best techniques developed and tested in GERDA and Majorana

Starts with 1000 kg ~ 2016÷2017

Tracking

- concept: scale Gotthard experiment adding Ba tagging to suppress background (¹³⁶Xe→¹³⁶Ba⁺⁺ +2e)
- single Ba⁺ detected by optical spectroscopy
- two options with 63% enriched Xe
 - High pressure Xe TPC
 - LXe TPC + scintillation calorimetry + tracking
- expected bkg only by ββ-2ν
 ▶ energy resolution σ_E = 2%

$\begin{array}{c} \mathbf{P}_{1/2} \\ \mathbf{493} \text{ nm} \\ \mathbf{493} \text{ nm} \\ \mathbf{10} \\ \mathbf{$

Present R&D

EXO

- Ba⁺ spectroscopy in HP Xe / Ba⁺ extr.
- energy resolution in LXe (ion.+scint.)
- Prototype scale:
- 200 kg enriched L¹³⁶X e without taggin
- all EXO functionality except Baid
- operate in WIPP for ~two years
 Protorype goals:
- Test all technical aspects of EXO (except Baid)
- Measure 2v mode
- Set decent limit for 0v mode (probe Heidelberg- Moscow)

Full scale appariment at WIPP or SNOLAB
■10 t (for LXe \Rightarrow 3 m ³)
$b = 4 \times 10^{-3} \text{ c/keV/ton/y}$
T ₁₁₂ > 1.3×10 ²⁸ y in 5 years
<m_> < 0.013 ÷ 0.037 eV</m_>

LXe TPC



A module			20 modules		
Submodule Solore and Laibenster		ever and	OODdaN		
Tracker In (essembled, "0.5m between source and calerimeter)		4			
Tracker	Demonstrator module	20 Modules		2012	
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source : ⁸² Se Drift chambers for tracking	Demonstrator module 7 kg 2 000	20 Modules 140 kg 40 000	Demonstrator SuperNEMO	~2013 ~2015	
source : ⁸² Se Drift chambers for tracking Electron calorimeter	Demonstrator module 7 kg 2 000 500	20 Modules 140 kg 40 000 10 000	Demonstrator SuperNEMO	~2013 ~2015	
Tracker In (essembled, "0.5m between source and calerimeter) Source : ⁸² Se Drift chambers for tracking Electron calorimeter γ veto (up and down)	Demonstrator module 7 kg 2 000 500 100	20 Modules 140 kg 40 000 10 000 2 000	Demonstrator SuperNEMO	~2013 ~2015	
Tacker (essembled, "0.5m between source and colorimeter) Source : ⁸² Se Drift chambers for tracking Electron calorimeter γ veto (up and down) T _{1/2} sensitivity	Demonstrator 7 kg 2 000 500 100 6.6 10 ²⁴ y (No background)	20 Modules 140 kg 40 000 10 000 2 000 1. 10 ²⁶ y	Demonstrator SuperNEMO	~2013 ~2015	





scintillator filling