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Scintillating bolometers based on ZnMoO₄ and Zn¹⁰⁰MoO₄ crystals to search for $0\nu2\beta$ decay of ¹⁰⁰Mo (LUMINEU project): first tests at the Modane Underground Laboratory

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Abstract

The technology of scintillating bolometers based on zinc molybdate (ZnMoO₄) crystals is under development within the LUMINEU project to search for $0\nu2\beta$ decay of 100 Mo with the goal to set the basis for large scale experiments capable to explore the inverted hierarchy region of the neutrino mass pattern. Advanced ZnMoO₄ crystal scintillators with mass of ~ 0.3 kg were developed and Zn¹⁰⁰MoO₄ crystal from enriched 100 Mo was produced for the first time by using the low-thermal-gradient Czochralski technique. One ZnMoO₄ scintillator and two samples (59 g and 63 g) cut from the enriched boule were tested aboveground at milli-Kelvin temperature as scintillating bolometers showing a high detection performance. The first results of the low background measurements with three ZnMoO₄ and two enriched detectors installed in the EDELWEISS set-up at the Modane Underground Laboratory (France) are presented.

Keywords: Double beta decay, Scintillating bolometer, ZnMoO₄ crystal scintillator, Low counting experiment

1. Introduction

Scintillating bolometers — cryogenic detectors with a heat-light double read-out — can play a crucial role in next-generation experiments to study neutrino properties and weak interaction via investigating neutrinoless double beta $(0v2\beta)$ decay, as discussed in Refs. [1, 2]. This technique is extensively developing now within the LUCIFER [3, 4], the AMORE [5, 6], and the LUMINEU [7] $0v2\beta$ projects. This paper describes the recent achievements in the framework of the LUMINEU programme (Luminescent Underground Molybdenum Investigation for NEUtrino mass and nature).

LUMINEU is devoted to the development of a technology based on zinc molybdate (ZnMoO₄) scintillating bolometer as a basis for the realization of a high-sensitivity $0\nu2\beta$ experiment. The good prospects of this material for the bolometric technique are clearly shown

in recent investigations [1, 8–13]. An important point in the realization of LUMINEU is concerned with the technology of growing high-quality radiopure large mass (0.3-0.5 kg) ZnMoO₄ single crystals with the aim to produce scintillators enriched in 100 Mo (Zn 100 MoO₄). Here we report a significant progress in the development of ZnMoO₄ crystal scintillators using deeply purified compounds (containing molybdenum with natural isotopic composition and enriched in 100 Mo). We also present results of both aboveground and underground low temperature tests of new scintillating bolometers based on natural ZnMoO₄ and enriched Zn 100 MoO₄ crystal scintillators in light of their possible application to next-generation $0\nu2\beta$ decay experiments.

2. Development of zinc molybdate based scintillating bolometers

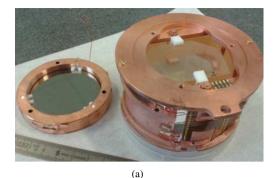
A precursor of the LUMINEU programme, a slightly yellow colored 313 g ZnMoO₄ sample with irreg-

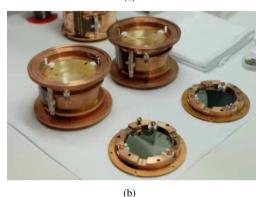
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ular shape, was produced from the first large volume ZnMoO₄ crystal boule grown by the low-thermal-gradient Czochralski (LTG Cz) technique [14, 15] in the Nikolaev Institute of Inorganic Chemistry (NIIC, Novosibirsk, Russia). The second sample (with mass 329 g) produced from this boule was tested as a scintillating bolometer at the Gran Sasso National Laboratories (LNGS, Assergi, Italy) [11].

Advanced ZnMoO₄ crystal boules with mass of ~ 1 kg have been produced recently at the NIIC by using the LTG Cz growth technique and molybdenum purified by sublimation in vacuum and double recrystallization from aqueous solutions [13]. The crystals were recrystallized to improve quality of the material, and two colorless ZnMoO₄ cylindrical samples (with size ⊘50×40 mm and mass 336 and 334 g) were produced from them. Moreover, a zinc molybdate crystal boule (with mass 171 g) enriched in 100 Mo to 99.5% was developed for the first time at the NIIC [16, 17], and two scintillation elements (with mass 59 and 63 g) were cut from the boule. The enriched molybdenum was purified by sublimation and recrystallization from aqueous solutions. It is worth noting the high yield of the Zn¹⁰⁰MoO₄ crystal boule from the initial charge (84%) and low level of total irrecoverable losses of enriched material (4%) achieved in the frame of this R&D [16]. Some coloration of the crystal (in contradiction with the practically colorless samples produced from natural molybdenum) can be explained by remaining traces of iron in the enriched molybdenum and by crystallization procedure performed only one time [16].

In order to construct scintillating bolometers, all the above described samples were held inside Copper holders by using PTFE clamps. Both Zn¹⁰⁰MoO₄ crystals were mounted in one Copper holder. The crystal scintillators were surrounded by a reflector foil (3M VM2000/2002) to improve light collection. Thin ultrapure Ge wafers (⊘50×0.25 mm) were used for detecting scintillation light. The 313 g crystal was viewed by two light detectors fixed on the opposite sides. The ZnMoO₄ / Zn¹⁰⁰MoO₄ crystals and the Ge photodetectors were instrumented with Neutron Transmutation Doped (NTD) Ge thermistors used as temperature sensors. All the crystals were also assembled with an individual heating element based on a heavily-doped silicon meander. Such devices provide a stable resistance value and are used to inject periodically a certain amount of thermal energy with the aim to control and stabilize the thermal bolometric response. All the detector modules are shown in Fig. 1.





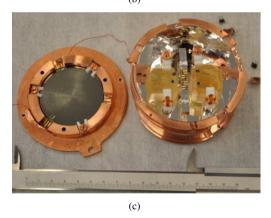


Figure 1: Photographs of scintillating bolometers based on ultra-pure Ge photodetectors and ZnMoO₄ precursor with mass of 313 g (a), advanced quality (see text) ZnMoO₄ crystals with masses of 336 and 334 g (b), and enriched Zn¹⁰⁰MoO₄ crystals with masses of 59 and 63 g (c).

3. Aboveground low temperature tests

The 313 g ZnMoO₄ precursor and both Zn¹⁰⁰MoO₄ crystals were tested in aboveground cryogenic facilities of the Centre de Sciences Nucléaires et de Sciences de la Matière (CSNSM, Orsay, France) with "wet" and "dry" ³He/⁴He dilution refrigerators, respectively.

Both cryostats are surrounded by passive shield made of low activity lead to minimize signals pile-up caused

by environmental gamma rays due to a slow time response of the bolometers (hundreds millisecond). The stream data were recorded by a 16 bit ADC with a sampling frequency of 30 kHz and 10 kHz for natural and enriched detectors, respectively. The ZnMoO₄ precursor was operated at 17 mK during the measurements (over 38 h), while the Zn¹⁰⁰MoO₄ array was tested at 13.7 mK (18 h), 15 mK (5 h), and 19 mK (24 h) base temperatures. Both detectors were irradiated by gamma quanta from a weak ²³²Th source, while the photodetectors were calibrated with the help of ⁵⁵Fe sources fixed close to the Ge slabs.

The data treatment (here and below) was performed by using the optimum filtering [18]. The spectrometric performances of the precursor-based bolometer were deteriorated by the pile-ups effect due to considerably high counting rate ≈ 2.5 Hz (e.g. see in Table 1 the energy resolution of the 2615 keV γ peak). In spite of this, the test shows normal operability of the detector and allows us to estimate the scintillation light yield for the registered $\gamma(\beta)$ events and muons, as well as the possibility of particle discrimination between $\gamma(\beta)$ and α events due to the quenching of scintillation for α particles. All these data are reported in Table 1.

Both enriched crystals demonstrate similar performance at all the temperatures [16]. The 2-dimensional histogram obtained from the heat-light double read-out of the 59 g Zn¹⁰⁰MoO₄ bolometer at 13.7 mK is shown in Fig. 2 (a). The light and the heat signals detected simultaneously allow to get a clear discrimination between α and $\gamma(\beta)$ particles. The absence in Fig. 2 (a) of peculiarities related with the detection of α events (except a small structure possibly caused by ²¹⁰Po, as often occurs in scintillators) indicates on encouraging radiopurity of the tested enriched crystals. Good spectrometric properties of the enriched detectors, even at aboveground conditions, are well visible from Fig. 2 (b), while some further information about their performances is presented in Table 1.

4. Underground cryogenic measurements

The 313 g detector was moved deep underground (\approx 4800 m w.e.) to the Modane Underground Laboratory (Laboratoire Souterrain de Modane, LSM, France) and tested during the EDELWEISS-III commissioning runs. The ZnMoO₄ bolometer together with fifteen ultra-pure Ge detectors (0.8 kg each) fully covered with interleaved electrodes (FID) were installed inside the 3 He/ 4 He inverted dilution refrigerator with a large experimental volume (50 l) [19]. The EDELWEISS setup, located inside a clean room (ISO Class 4) and sup-

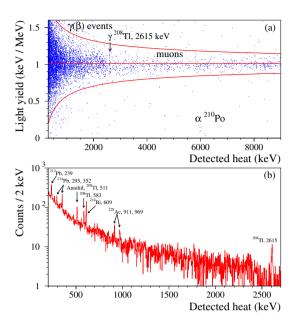


Figure 2: (a) The scatter plot of the light-to-heat signal amplitude ratio as a function of the heat signal amplitude accumulated at 13.7 mK in aboveground test with the 59 g Zn 100 MoO $_4$ scintillating bolometer during 18 h of calibration measurements with the 232 Th source. The visible band is related to $\gamma(\beta)$ events (below 2.6 MeV) and cosmic muons. Three sigma intervals of the light yield for the $\gamma(\beta)$ band are shown by solid red curves together with the median value. (b) The energy spectrum built from the data presented in the upper plot. The peaks observed in the energy spectrum belong to the 232 Th source and environmental gamma's (daughters of 226 Ra). The energy of marked peaks are given in keV.

plied by deradonized ($\approx 30 \text{ mBq/m}^3$) air flow, is surrounded by a massive shield made of low background lead (20 cm thick) and polyethylene (50 cm). The setup is surrounded by a 5 cm thick plastic scintillator muon veto (95% coverage), and equipped by neutron and radon counters.

The triggered signals were recorded by a 14 bit ADC in 2 s window with 2 kHz sampling rate (the half of the window contains the baseline data). The base temperature was stabilized around 19 mK. One light detector was very sensitive to microphonic noise and could not be used for measurements. The energy scale of the ZnMoO₄ detector has been measured in calibration runs with $^{133}{\rm Ba}$ and $^{232}{\rm Th}~\gamma$ sources, performed over 546 h and 70 h, respectively. The background data were accumulated over 305 h.

The powerful discrimination capability achieved with the 313 g ZnMoO₄ scintillating bolometer is well illustrated in Fig. 3 (a), which shows a full separation of $\gamma(\beta)$ -induced events from populations of α particles caused by trace impurity by radionuclides from U/Th chains (mainly, ²¹⁰Po, see below). The energy spectrum

Table 1: List of achieved performances with ZnMoO₄ and Zn¹⁰⁰MoO₄ detectors tested in aboveground and underground measurements. We report the energy resolution for the heat channels (FWHM — Full Width at the Half of Maximum) estimated as filtered baseline and measured for γ quanta and α particles of internal ²¹⁰Po. We report also the light yield for $\gamma(\beta)$ events (LY $_{\gamma(\beta)}$) and quenching factor for α particles (QF $_{\alpha}$).

Detector		FWHM (keV)					$LY_{\gamma(\beta)}$	QF_{α}
Crystal	Mass (g)	Baseline	¹³³ Ba	²¹⁴ Bi	²⁰⁸ Tl	²¹⁰ Po	(keV/MeV)	
			356 keV	609 keV	2615 keV	5407 keV		
ZnMoO ₄	313	1.4(1)	6.4(1)	6(1)*	24(2)* / 9(2)	19(1)	0.77(11)*	0.15(2)* / 0.14(1)
	336	1.5(2)	6(1)	_	_	29(4)	_	_
	334	1.06(3)	3.8(4)	-	_	15(1)	_	0.19(2)
Zn ¹⁰⁰ MoO ₄	59	1.4(1)*	_	5.0(5)*	11(3)*	_	1.01(11)*	≈ 0.15*
	63	1.8(1)*	_	10(1)*	_	_	0.93(11)*	≈ 0.15*

^{* -} results based on the aboveground measurements

accumulated with the ²³²Th gamma source (see Fig. 3 (b)) demonstrates high spectrometric properties of the detector. An overview of the detector's performances during underground measurements is given in Table 1.

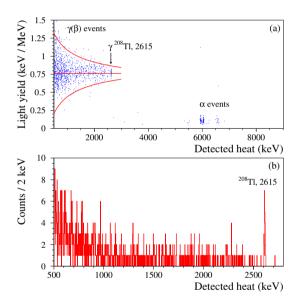


Figure 3: (a) Plot reporting the light-to-heat signal amplitude ratio as a function of the heat signal amplitude for the 313 g ZnMoO₄ detector installed in the EDELWEISS set-up. The detector was cooled down to 19 mK and irradiated over 51 h by γ quanta from the 232 Th source. Two visible bands correspond to $\gamma(\beta)$ events and α particles. The positions of the α events are shifted from the nominal values due to thermal energy overestimation for α particles in case of using calibration data for γ 's. Three sigma intervals of the light yield for the $\gamma(\beta)$ band and its median value are drawn. (b) The energy spectrum of the 232 Th source measured by the 313 g ZnMoO₄ scintillating bolometer during 51 h of underground cryogenic run.

After completing the EDELWEISS-III commissioning runs, other two ZnMoO₄-based scintillating bolometers ($\oslash 50 \times 40$ mm) and the Zn¹⁰⁰MoO₄ array together with 36 FID Ge detectors were assembled. The EDELWEISS set-up was also upgraded: a) a polyethylene shield at the 1 K plate was added; b) new ultra ra-

diopure NOSV Copper [20] screens were installed; c) all detectors were provided with individual low background Copper-Kapton cables. In addition, a pulser system to assist to the calibration of the thermal response of the $ZnMoO_4$ / $Zn^{100}MoO_4$ detectors will be implemented soon.

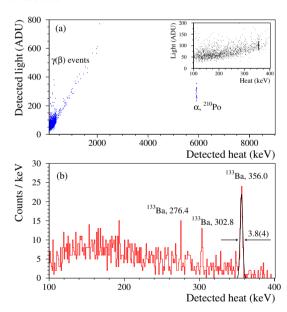


Figure 4: (a) Scatter plot of the light versus the heat signals measured by the 334 g ZnMoO₄ scintillating bolometer in a 15 h calibration run with the 133 Ba gamma source in the EDELWEISS set-up. A cluster of events located far from the $\gamma(\beta)$ population corresponds to α particles of 210 Po. The data for the light channel are presented in ADU (Analogue-to-Digital Unit). (Insert) Part of the scatter plot corresponding to the energy range of the used source. (b) The energy spectrum of the 133 Ba source measured over 15 h by the 334 g ZnMoO₄ scintillating bolometer.

After the upgrade of the set-up the data are recorded by a 16 bit ADC with 1 kHz sampling rate (the length of pulse profile is 2 s with the half of the window for the baseline data). The working temperature is stabilized at 18 mK. The energy scale of the detectors was measured with the ¹³³Ba gamma source (the measurements with the ²³²Th source are foreseen).

The set-up is still under optimization, especially as far as the control of the vibration-induced noise is concerned. Therefore, we discuss here, as an illustrative example, only the results achieved with the 334 g natural ZnMoO₄ scintillating bolometer. This detector exhibits full $\alpha/\gamma(\beta)$ separation, as shown in Fig. 4 (a), as well as excellent spectrometric properties, as demonstrated in Fig. 4 (b). Other relevant information about performances of $\oslash 50 \times 40$ mm ZnMoO₄ detectors are reported in Table 1.

5. Radiopurity of ZnMoO₄ and Zn¹⁰⁰MoO₄ crystals

The radiopurity level of the ZnMoO₄ crystals was estimated by analysis of the α events selected from the underground runs, while the data of the aboveground measurements were used in case of the Zn¹⁰⁰MoO₄ samples. The position of the 5.4 MeV α peak of the internal ²¹⁰Po, clearly visible in the data for the natural crystals, was used to stabilize the thermal response of the detectors. For instance, the spectra of the α events registered by the detectors based on 313 g (a) and 334 g (b) ZnMoO₄ crystals over 851 h and 527 h, respectively, are shown in Fig. 5.

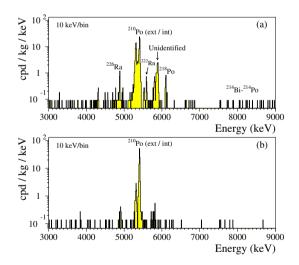


Figure 5: The α spectra collected in the low background measurements in the EDELWEISS set-up with the ZnMoO₄ scintillating bolometers based on the 313 g precursor (a) and the 334 g advanced sample (b) operated over 851 h and 527 h, respectively. The origin of the α events providing the highest rate are indicated.

The crystals are slightly polluted by 210 Po detected through 5.4 MeV α peak confirming a broken equilibrium in the radioactive chain. 226 Ra (and its daughters 222 Rn, 218 Po, and 214 Bi- 214 Po events), and 228 Th (with

daughter 224 Ra¹) were detected in the 313 g crystal, while the ZnMoO₄ scintillators produced by recrystallization have shown a much better level of radiopurity, particularly in 226 Ra. It is also evident a higher surface contamination by 210 Po of the 313 g crystal or/and of the bolometer components close to it (a peak at 5.3 MeV corresponds to E_{α} of 210 Po). In addition, excess counts around 5.8 MeV also indicate a possible surface contamination but its origin has not been identified.

The activity of internal 210 Po was derived from the fit of the 5.4 MeV peak, while 3σ intervals (according to the energy resolution of the internal 210 Po — see Table 1) centered at the Q_{α} value were used for the calculation of the area of the peaks of other radionuclides from U/Th chains. The background contribution was evaluated in two energy regions (3.3–4 and 4.35–4.7 MeV) with a flat α continuum in which no peaks are expected. The number of counts excluded with 90% C.L. were calculated by using the Feldman-Cousins procedure [21].

Table 2: Radioactive contamination of the ZnMoO₄ and Zn¹⁰⁰MoO₄ crystals tested as scintillating bolometers in aboveground and underground conditions. The mass of the crystals and the total time of the accumulated data are also presented. The results for the large mass ZnMoO₄ crystal which was operated as the scintillating bolometer at the LNGS (Italy) [11] are given for comparison. The uncertainties are given with 68% C.L., while all the limits are at 90% C.L.

Nuclide	Activity (mBq/kg)									
	Zn ¹⁰⁰ MoO ₄		ZnMoO ₄							
	59 g	63 g	336 g	334 g	313 g	329 g [11]				
	42 h	42 h	291 h	527 h	851 h	524 h				
²²⁸ Th	≤ 0.25	≤ 0.21	≤ 0.024	≤ 0.007	0.010(3)	≤ 0.006				
²³⁸ U	≤ 0.26	≤ 0.21	≤ 0.008	≤ 0.002	≤ 0.008	≤ 0.006				
²²⁶ Ra	≤ 0.26	≤ 0.31	≤ 0.021	≤ 0.009	0.26(5)	0.27(6)				
²¹⁰ Po	0.9(3)	1.1(3)	0.94(5)	1.02(7)	0.62(3)	0.70(3)				

Data (or limits) on radioactive contamination of the $ZnMoO_4$ and $Zn^{100}MoO_4$ scintillators are summarized in Table 2, where the results for another $ZnMoO_4$ sample, produced from the same boule as the 313 g crystal was, are presented for comparison. As it is seen from Table 2, the improved purification and crystallization procedure adopted for the LUMINEU crystals of 334 and 336 g has lead to a significant reduction of the internal contamination, especially for ^{226}Ra which is not detectable now while it was clearly present in both precursor crystals (313 and 329 g). In particular, the radiopurity levels ($\leq 0.01 \text{ mBq/kg}$) achieved for ^{228}Th and

¹Taking into account a short half-life of ²¹⁶Po (≈145 ms), which is comparable with the time response of the 313 g detector (hundreds ms), subsequent α decays of ²²⁰Rn-²¹⁶Po give pile-ups and therefore were discarded from the data by the pulse-shape analysis.

 226 Ra are fully compatible with next-generation $0v2\beta$ experiments capable to explore the inverted hierarchy region of the neutrino mass pattern [1, 2].

6. Conclusions

A significant progress is achieved in development of ZnMoO₄ crystal scintillators for the LUMINEU project. Large volume crystal boules (~ 1 kg each) were grown by the low-thermal-gradient Czochralski technique from deeply purified molybdenum. A Zn¹⁰⁰MoO₄ crystal boule with a mass of 0.17 kg was produced from enriched 100 Mo (to 99.5%) for the first time. Three natural (~ 0.3 kg) and two enriched (~ 0.06 kg) scintillation elements were produced for low temperature studies. Production of large volume Zn¹⁰⁰MoO₄ crystal scintillators from enriched 100 Mo is in progress.

The cryogenic scintillating bolometric tests of the natural and enriched crystals showed a high performance of the detectors. The deep purification of molybdenum and recrystallization significantly improve the radioactive contamination of ZnMoO₄ crystals by 228 Th and 226 Ra to the level of \leq 0.01 mBq/kg requested by the LUMINEU project.

The results of this study clarify the excellent prospects of $ZnMoO_4$ scintillating bolometers for the next generation $0\nu2\beta$ experiments aiming to approach the inverted hierarchy region of the neutrino mass pattern.

7. Acknowledgments

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