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TECHNICAL REPORT

Radiopurity control in the NEXT-100 double beta decay experiment: procedures and initial measurements

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ABSTRACT: The “Neutrino Experiment with a Xenon Time-Projection Chamber” (NEXT) is intended to investigate the neutrinoless double beta decay of ^{136}Xe , which requires a severe suppression of potential backgrounds. An extensive screening and material selection process is underway for NEXT since the control of the radiopurity levels of the materials to be used in the experimental set-up is a must for rare event searches. First measurements based on Glow Discharge Mass Spectrometry and gamma-ray spectroscopy using ultra-low background germanium detectors at the Laboratorio Subterráneo de Canfranc (Spain) are described here. Activity results for natural radioactive chains and other common radionuclides are summarized, being the values obtained for some materials like copper and stainless steel very competitive. The implications of these results for the NEXT experiment are also discussed.

KEYWORDS: Radiation calculations; Time projection Chambers (TPC); Gamma detectors (scintillators, CZT, HPG, HgI etc)

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1 Introduction

The observation of neutrinoless double beta decay as a peak in the sum energy of the two emitted electrons ($Q_{\beta\beta}$), would show that neutrinos are Majorana particles and contribute to the determination of their mass hierarchy (see for instance [1]). The NEXT experiment (“Neutrino Experiment with a Xenon Time-Projection Chamber”) will search for such a decay in ^{136}Xe using a high-pressure gaseous xenon Time-Projection Chamber (TPC) with a source mass of the order of 100 kg at the Laboratorio Subterráneo de Canfranc (LSC) [2], located at the Spanish Pyrenees. The NEXT-100 detector design [3] is intended to combine, keeping the detector=source approach, the measurement of the topological signature of the event (to discriminate the signal from background) with the energy resolution optimization; this is possible thanks to the use of proportional electroluminescent (EL) amplification. As illustrated in figure 1, energy and tracking readout planes are located at opposite sides of the pressure vessel using different sensors: photomultiplier tubes (PMTs) for calorimetry (and for fixing the start of the event) and silicon multi-pixel photon counters (MPPCs) for tracking. Energy resolution below 1% FWHM at the $Q_{\beta\beta}$ energy seems reachable [3]. The expected sensitivity of NEXT is very competitive; electron neutrino effective Majorana masses below 100 meV could be explored for a total exposure of 500 kg·year [4]. The Technical Design Report of the experiment was presented in 2011, while work on prototypes is ongoing [5].

Operating in deep underground locations and using both active and passive shields is mandatory in experiments searching for rare phenomena like the nuclear double beta decay, the direct detection of dark matter and the interaction of low-energy neutrinos [6, 7]. A careful selection of radiopure materials to be used in the experimental set-up based on precise measurements of ultra-low radioactivity is also compulsory in these experiments in order to achieve, for example in NEXT, a background level of 8×10^{-4} counts $\text{keV}^{-1} \text{kg}^{-1} \text{y}^{-1}$ in the energy region of interest

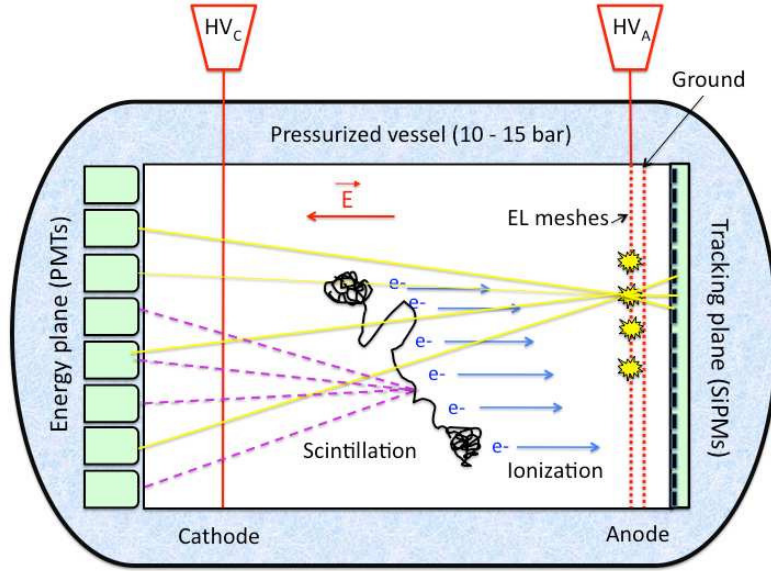


Figure 1. Concept of the NEXT experiment: light from the Xe electroluminescence (EL) generated at the anode is recorded both in the photosensor plane right behind it for tracking and in the photosensor plane behind the transparent cathode for a precise energy measurement. Primary scintillation defining the start of the event is also detected by the cathode photosensors.

(RoI) [3]. A great effort has been done by several collaborations to measure the radiopurity of many different materials (see for instance refs. [8]–[19]) and information has been compiled and made public [20, 21]. Although these data are extremely useful and can be used as a starting point in the selection of materials for any experiment, a thorough radiopurity control of the relevant components to be actually used must be always undertaken since radiopurity requirements are stringent.

The ability to discriminate signal from background is a powerful tool in NEXT. Signal events will appear uniformly distributed in the source volume of enriched xenon and will have a distinctive topology (a twisted long track, about 30 cm long at 10 bar, with blobs at both ends). Defining a fiducial volume eliminates all charged backgrounds entering the detector while confined tracks generated by neutral particles, like high-energy gammas, can be suppressed by pattern recognition. Thus, the relevance of a background source depends on its probability of generating a signal-like track in the active volume with energy around $Q_{\beta\beta}$. The neutrinoless double beta decay of ^{136}Xe would produce a peak at $Q_{\beta\beta} = 2.458 \text{ MeV}$ [22]; the main background sources at the NEXT experiment are the gamma lines at 2.615 MeV from ^{208}Tl and at 2.448 MeV from ^{214}Bi . These isotopes produced at the lower part of the natural radioactive chains of ^{232}Th and ^{238}U respectively.

- For the 2.615 MeV line, the Compton edge is well below $Q_{\beta\beta}$, but a scattered gamma can interact and produce other electron tracks close enough to the initial Compton electron, so they are reconstructed as a single object falling in the RoI. These photons can also be scattered outside the detector and then suffer photoelectric absorption inside contributing to the RoI. In addition, photoelectric electrons from the 2.615 MeV emission are produced above the RoI but can lose energy via bremsstrahlung and populate the energy window, if the emitted photons escape out of the detector.

- The gamma line at 2.448 MeV is dangerous in spite of its low intensity (1.57%) because it is very close to $Q_{\beta\beta}$.

Both ^{208}Tl and ^{214}Bi are in the progeny of Rn isotopes (^{220}Rn and ^{222}Rn respectively) which are present in air, so radon diffusion from the laboratory air as well as radon emanation from materials must be controlled. Another possible external source of background are neutrons, whether produced by natural radioactivity in the walls or shielding or as secondary products of cosmic muons; preliminary estimations seem to point that these contributions are very much below the level of concern for NEXT. Also high energy gammas can be produced in muon-induced electromagnetic cascades; although they seem to be of no importance for NEXT, they could be partially tagged by an active muon veto in the shielding.

Consequently, the activity of the lower part of ^{232}Th and ^{238}U chains (^{208}Tl and ^{214}Bi isotopes) from the components of the set-up and at the laboratory is the main concern in NEXT. According to the NEXT-100 design, the list of the main materials subject to radiopurity control and the target radiopurity for each of them were established (see table 3 in [3]). The relevant materials for the different parts of the experimental set-up to be considered include: lead and copper for shielding; stainless steel and inconel for vessel; PEEK (PolyEther Ether Ketone), polyethylene and coating materials for High Voltage (HV) and electroluminescence components; photomultipliers, windows, printed circuit boards and other electronic components (capacitors, resistors, cables...) for the energy and tracking planes. The target sensitivity was fixed following results from the literature and from the first measurements made for NEXT.

In this paper, the initial material radiopurity control performed for the NEXT experiment is presented. First, in section 2 the techniques and equipment used to carry out the measurements of activity levels are described. Then, the results obtained for all the analyzed materials are shown and discussed in section 3.

2 Techniques and equipment

The techniques employed for the radiopurity measurements for NEXT are Glow-Discharge Mass Spectrometry (GDMS) and gamma-ray spectroscopy using ultra-low background germanium detectors operated deep underground. Each one has advantages and drawbacks, which make them more or less suitable depending on the context.

2.1 GDMS

In GDMS, the sample to be analyzed forms the cathode in a plasma or discharge gas, typically argon. Argon ions are accelerated toward the sample resulting in erosion and atomization of its surface. The sputtered species are transported into the plasma where they are ionized. Ions are then extracted for mass spectrometry. This method is consequently very suitable for metals. Similarly to other techniques also based on mass spectrometry, it is fast and requires only a small sample of the material (typically, samples with a surface of $2 \times 2 \text{ cm}^2$ were prepared for NEXT measurements). However, the output given is normally only the concentration of elements while particular isotopes are not identified. Measured concentrations of U, Th and K have been converted to ^{232}Th , ^{238}U and

^{40}K activities.¹ Having no information on daughter nuclides in the chains, a possible disequilibrium cannot be detected. In any case, this technique can be very helpful to make a pre-selection of samples to be later screened with germanium detectors underground. GDMS measurements for NEXT materials were made by Shiva Technologies (Evans Analytical Group²) in France.

2.2 Germanium gamma-ray spectrometry

The most important advantage of gamma-ray spectrometry performed with germanium detectors is that, being non-destructive, the actual components to be used in experiments can be examined. Germanium detectors offer very good energy resolution and low intrinsic background. In addition, activities of isotopes dangerous for the experiment background are directly assessed. No sample pre-treatment is necessary, but massive samples and time-consuming measurements (lasting up to several weeks) are needed to quantify very low activities.

The LSC offers a Radiopurity Service³ to measure ultra-low level radioactivity using several germanium detectors operated at the Hall C for gamma-ray spectroscopy (see figure 2); it is intended to support the construction of experiments to be operated there. Since the laboratory is located at a depth of 2450 m.w.e., the cosmic muon flux is about 5 orders of magnitude lower than at sea-level surface. Radon activity in the air is between 50 and 80 Bq/m³ in the underground halls [23]. All germanium spectroscopy measurements presented here were carried out at LSC using in particular four different ~ 2.2 kg detectors from LSC and a ~ 1 kg detector from the University of Zaragoza.

- The detectors property of LSC are p-type close-end coaxial High Purity germanium detectors produced by Canberra France with 100–110% relative efficiencies⁴ and a FWHM energy resolution of ~ 2 keV at the ^{60}Co gamma line of 1332 keV. The active volume of crystals ranges from 410 to 420 cm³ and cryostats are made of ultra-low background aluminum. The data acquisition system is based on Digital Signal Processing using Canberra DSA1000 modules. Each detector has a shield consisting of 5 cm of oxygen-free copper and 20 cm of very low activity lead having < 30 mBq/kg of ^{210}Pb ; nitrogen gas is flushed inside a methacrylate box to avoid airborne radon intrusion. The detectors used for NEXT measurements are those named GeOroel, GeTobazo, GeAnayet and GeLatuca; they are in operation since 2011.
- The detector from University of Zaragoza, named Paquito, is also a p-type close-end coaxial High Purity germanium detector, but with a smaller crystal volume of 190 cm³ inside a copper cryostat. The FWHM energy resolution at 1332 keV is 2.7 keV. It is operated inside a shield made of 10 cm of archaeological lead plus 15 cm of low activity lead with nitrogen flush too. The electronic chain for data acquisition is based on standard Canberra 2020 Linear Amplifier and Canberra 8075 Analog-to-Digital-Converter modules. This detector has been used for radiopurity measurements at Canfranc for several years (more details can be found in [24]).

¹Conversion factors are: 1 ppb U = 12.4 mBq/kg ^{238}U , 1 ppb Th = 4.1 mBq/kg ^{232}Th and 1 ppm $^{\text{nat}}\text{K}$ = 31 mBq/kg ^{40}K .

²<http://www.eaglabs.com>.

³<http://www.lsc-canfranc.es/en/for-users/lsc-services/radiopurity.html>.

⁴Efficiency relative to a $3'' \times 3''$ NaI detector at 1332 keV and for a distance of 25 cm between source and detector.



Figure 2. Picture of the Hall C of the LSC where several germanium detectors inside their shieldings are operated for material radiopurity screening (courtesy of LSC). Background counting rates are reported in table 1.

Table 1. Background counting rates (expressed in counts $\text{d}^{-1} \text{kg}^{-1}$) of the germanium detectors used at LSC for the NEXT measurements. Integral rate from 100 to 2700 keV and rates at different peaks (583 keV from ^{208}Tl , 609 keV from ^{214}Bi and 1461 keV from ^{40}K) are presented. Only statistical errors are quoted.

Detector name	Mass (kg)	100–2700 keV	583 keV	609 keV	1461 keV
GeOroel	2.230	490 ± 2	0.8 ± 0.1	3.0 ± 0.2	0.41 ± 0.07
GeAnayet	2.183	714 ± 3	3.8 ± 0.2	1.7 ± 0.1	0.38 ± 0.07
GeTobazo	2.185	708 ± 3	4.0 ± 0.2	1.3 ± 0.1	0.40 ± 0.06
GeLatuca	2.187	710 ± 3	3.3 ± 0.2	5.9 ± 0.3	0.56 ± 0.08
Paquito	1	79 ± 2	0.27 ± 0.09	0.5 ± 0.1	0.25 ± 0.09

The background of each detector inside its shielding is determined by taking data with no sample for long periods of time of at least one month, due to the low counting rates. Table 1 shows the counting rates of all the detectors used in the NEXT measurements in the energy window from 100 to 2700 keV and at different peaks: 583 keV from ^{208}Tl , 609 keV from ^{214}Bi and 1461 keV from ^{40}K ; all the rates are expressed in counts per day and per kg of germanium detector and correspond to background measurements performed from May to July 2012. The complete background spectrum of GeOroel detector, registered for 38.00 days, is presented in figure 3. Lines from cosmogenic isotopes like ^{54}Mn , ^{58}Co and ^{65}Zn , having half-lives around one year, are still observable in GeOroel detector but not in the other ones.

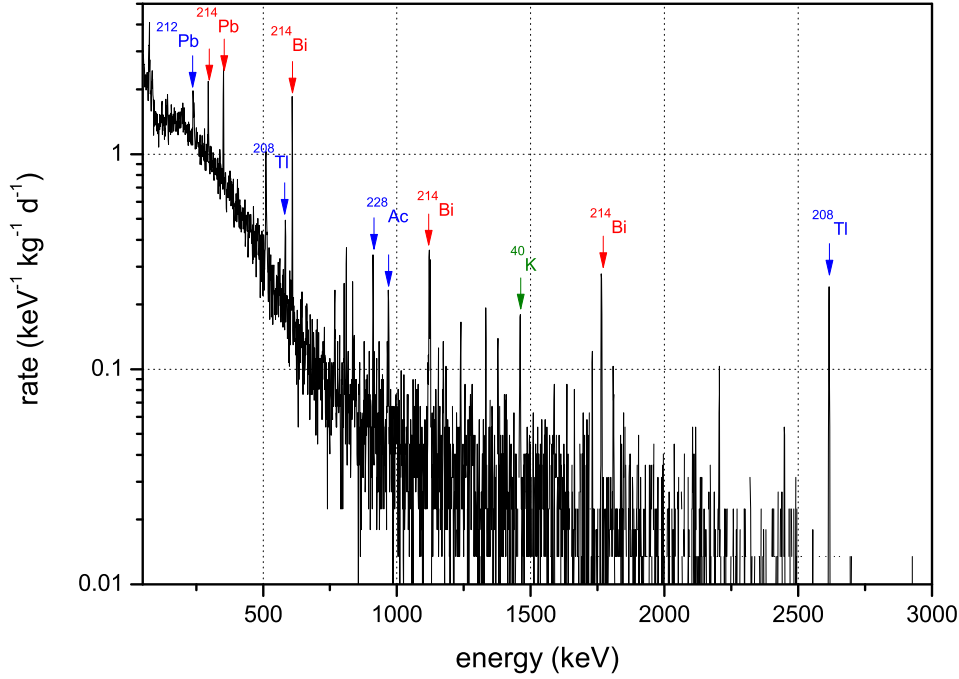


Figure 3. Energy spectrum of GeOroel detector at LSC registered in a background measurement for 38.00 days. Main gamma lines from isotopes of the ^{238}U (red) and ^{232}Th (blue) radioactive chains and from ^{40}K (green) are marked.

To derive the activity in a sample of an isotope producing a gamma emission of a certain energy, the main ingredients (together with the time of the measurement and the branching ratio of the emission) are the net signal (that is, the number of events at the gamma line due to the sample) and the full-energy peak detection efficiency at the corresponding energy.

In the context of determining ultra-low activities of a sample (at the level of mBq/kg and below), when comparing the measured energy spectrum of the sample with that of the detector background, it is not straightforward to decide if the gross signal can be considered to statistically differ from the background signal. The criteria proposed in Currie’s landmark paper [25] and more recently revised in [16, 26] have been followed here. Activities have been quantified when possible and upper limits with a 95.45% C.L. have been derived otherwise.

Concerning the estimate of the detection efficiency, Monte Carlo simulations based on the Geant4 [27] code have been performed for each sample, accounting for intrinsic efficiency, the geometric factor and self-absorption at the sample. No relevant change has been observed in the Geant4 simulation when changing version or the physical models implemented for interactions (considering the low energy extensions for electromagnetic processes based on theoretical models and on exploitation of evaluated data, G4EmLivermorePhysics class and the previous G4LowEnergy* classes [28]). Validation of the simulation has been made by comparing the efficiency curve of the detectors measured with a ^{152}Eu reference source of known activity located at 25 cm from the detector with the simulated one. Figure 4 shows the intrinsic efficiency (corrected by solid angle) obtained for GeOroel, GeTobazo, GeAnayet and GeLatuca detectors together with the correspond-

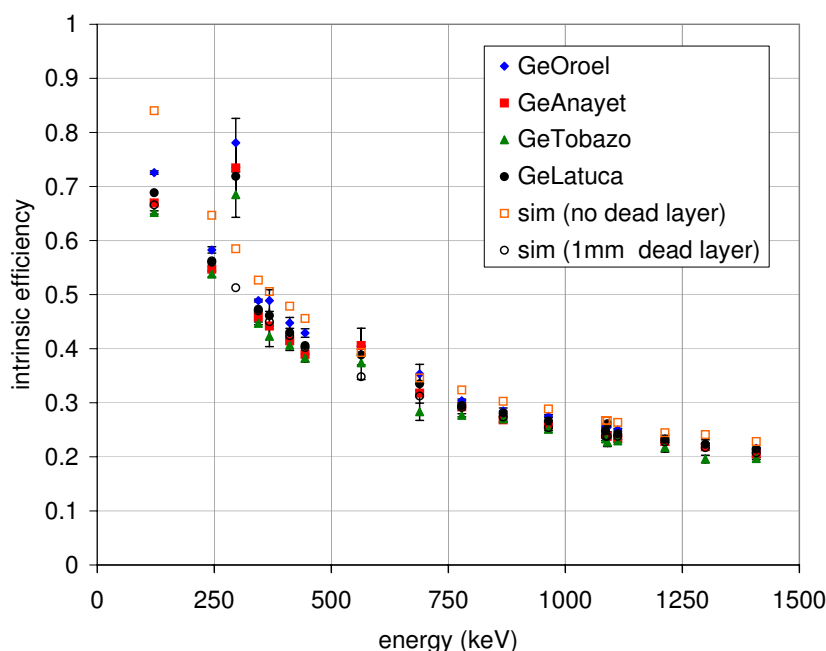


Figure 4. Intrinsic efficiency measured for different Ge detectors using a ^{152}Eu reference source and the corresponding simulation, considering a 1-mm-thick crystal dead layer or no dead layer.

ing simulations for GeTobazo/GeAnayet. The higher efficiency shown by GeOroel is due to a slightly larger volume in comparison with the other detectors. The inclusion of a 1-mm-thick dead layer in the simulation has improved the agreement with measurements, especially at low energies, reducing deviations to a level of 5%; an overall uncertainty of 10% is considered for the simulated detection efficiency of the samples and propagated to the final activity value. The larger discrepancy observed in figure 4 for 295 keV values is due to the interference in the measurements at this energy of the gamma line of ^{214}Pb , descendant of ^{222}Rn present in air (measurements with the ^{152}Eu source were carried out with the shields partially open and no background subtraction was made). The relative efficiencies derived from these measurements and also from the simulation reproduce the values specified by the manufacturer.

Activities of different sub-series in the natural chains of ^{238}U , ^{232}Th and ^{235}U as well as of common primordial, cosmogenic or anthropogenic radionuclides like ^{40}K , ^{60}Co and ^{137}Cs have been evaluated by analyzing the most intense gamma lines of different isotopes. Outgassing and chemical procedures in materials can make secular equilibrium in radioactive chains break, so information provided by germanium detectors at the different stages is very important. For ^{238}U , emissions from ^{234}Th and ^{234m}Pa are searched to quantify activity of the upper part of the chain and lines from ^{214}Pb and ^{214}Bi for the sub-chain starting with ^{226}Ra up to ^{210}Pb . For ^{232}Th chain, emissions of ^{228}Ac are analyzed for the upper part and those of ^{212}Pb , ^{212}Bi and ^{208}Tl for the lower one. Concerning ^{235}U chain, only emissions from the parent isotope are taken into account since all the other ones have very low intensities.

Table 2. Information on the germanium gamma-ray spectrometry measurements performed for NEXT at LSC: material, detector used, samples size (mass, area or number of pieces) and screening time. The corresponding row number of table 3 where the activity values obtained for each sample are reported is also quoted.

Material, Supplier	# in table 3	Detector	Sample size	Screening time (d)
Pb, Tecnibusa	5	GeAnayet	5585 g	19.44
Pb, Tecnibusa	6	GeAnayet	5585 g	35.99
Cu, Luvata	10	Paquito	681 g	39.17
Ti, SMP	11	GeOroel	121 g	38.46
Ti, SMP	12	GeTobazo	121 g	43.11
Ti, Ti Metal Supply	13	GeOroel	1804 g	47.23
304L Stainless Steel, Pfeiffer	14	Paquito	347 g	19.55
316Ti Stainless Steel, 10 mm, Nironit	15	GeTobazo	7684 g	33.00
316Ti Stainless Steel, 15 mm, Nironit	16	GeTobazo	10205 g	35.61
316Ti Stainless Steel, 50 mm, Nironit	17	GeAnayet	4816 g	34.72
Inconel 625, Mecanizados Kanter	18	GeTobazo	1004 g	27.98
Inconel 718, Mecanizados Kanter	19	GeOroel	611 g	27.93
PEEK, Sanmetal	20	Paquito	459 g	24.27
Polyethylene, IN2 Plastics	21	GeAnayet	1315 g	36.76
Semitron ES225, Quadrant EPP	22	GeOroel	1618 g	35.05
SMD resistor, Farnell	23	Paquito	50 pc	18.15
SM5D resistor, Finechem	24	Paquito	100 pc	31.45
Kapton-Cu PCB, LabCircuits	25	Paquito	260.15 cm ²	35.28
Cuflon, Polyflon	26	GeOroel	1876 g	24.29
Bonding films, Polyflon	27	GeAnayet	288 g	30.83
FFC/FCP connector, Hirose	28	Paquito	19 pc (1.23 g/pc)	6.83
P5K connector, Panasonic	29	Paquito	15 pc (0.67 g/pc)	7.58
Thermoplastic connector, Molex	30	GeLatuca	29 pc (0.53 g/pc)	17.20
Solder paste, Multicore	31	GeLatuca	457 g	44.30
Solder wire, Multicore	32	Paquito	91 g	7.74
Ta capacitor, Vishay Sprague	33	GeAnayet	277 pc (0.64 g/pc)	17.97

3 Measurements and results

Materials of very massive components and of those to be located inside the detector vessel have been selected for the first screening campaign of NEXT, trying to keep up with the schedule of construction of several subsystems like the pressure vessel, the field cage and the boards for the tracking plane. The performed measurements are described and discussed in the context of NEXT. Details concerning the detector used, the size of the sample and the time of data taking for all measurements using germanium detectors are presented in table 2. The results obtained are all independently summarized in table 3; reported errors correspond to 1σ uncertainties including both statistical and efficiency uncertainties.

Table 3. Activities measured in relevant materials for NEXT and following different techniques. GDMS results were derived from U, Th and K concentrations. Germanium gamma-ray spectrometry results reported for ^{238}U and ^{232}Th correspond to the upper part of the chains and those of ^{226}Ra and ^{228}Th give activities of the lower parts (see text).

#	Material	Supplier	Technique	Unit	^{238}U	^{226}Ra	^{232}Th	^{228}Th	^{235}U	^{40}K	^{60}Co	^{137}Cs
Shielding												
1	Pb	Cometa	GDMS	mBq/kg	0.37		0.073			<0.31		
2	Pb	Mifer	GDMS	mBq/kg	<1.2		<0.41			0.31		
3	Pb	Mifer	GDMS	mBq/kg	0.33		0.10			1.2		
4	Pb	Tecnibusa	GDMS	mBq/kg	0.73		0.14			0.91		
5	Pb	Tecnibusa	Ge	mBq/kg	<94	<2.0	<3.8	<4.4	<30	<2.8	<0.2	<0.8
6	Pb	Tecnibusa	Ge	mBq/kg	<57	<1.9	<1.7	<2.8	<22	<1.7	<0.1	<0.5
7	Cu (ETP)	Sanmetal	GDMS	mBq/kg	<0.062		<0.020					
8	Cu (C10100)	Luvata (hot rolled)	GDMS	mBq/kg	<0.012		<0.0041			0.061		
9	Cu (C10100)	Luvata (cold rolled)	GDMS	mBq/kg	<0.012		<0.0041			0.091		
10	Cu (C10100)	Luvata (hot+cold rolled)	Ge	mBq/kg		<7.4	<0.8	<4.3		<18	<0.8	<1.2
Vessel												
11	Ti	SMP	Ge	mBq/kg	<233	<5.7	<8.8	<9.5	3.4±1.0	<22	<3.3	<5.2
12	Ti	SMP	Ge	mBq/kg	<361	<6.6	<11	<10	<8.0	<15	<1.0	<1.8
13	Ti	Ti Metal Supply	Ge	mBq/kg	<14	<0.22	<0.5	3.6±0.2	0.43±0.08	<0.6	<0.07	<0.07
14	304L SS	Pfeiffer	Ge	mBq/kg		14.3±2.8	9.7±2.3	16.2±3.9	3.2±1.1	<17	11.3±2.7	<1.6
15	316Ti SS	Nironit, 10-mm-thick	Ge	mBq/kg	<21	<0.57	<0.59	<0.54	<0.74	<0.96	2.8±0.2	<0.12
16	316Ti SS	Nironit, 15-mm-thick	Ge	mBq/kg	<25	<0.46	<0.69	<0.88	<0.75	<1.0	4.4±0.3	<0.17
17	316Ti SS	Nironit, 50-mm-thick	Ge	mBq/kg	67±22	<1.7	2.1±0.4	2.0±0.7	2.4±0.6	<2.5	4.2±0.3	<0.6
18	Inconel 625	Mecanizados Kanter	Ge	mBq/kg	<120	<1.9	<3.4	<3.2	<4.6	<3.9	<0.4	<0.6
19	Inconel 718	Mecanizados Kanter	Ge	mBq/kg	309±78	<3.4	<5.1	<4.4	15.0±1.9	<13	<1.4	<1.3
HV, EL components												
20	PEEK	Sanmetal	Ge	mBq/kg		36.3±4.3	14.9±5.3	11.0±2.4	<7.8	8.3±3.0	<3.3	<2.6
21	Polyethylene	IN2 Plastics	Ge	mBq/kg	<140	<1.9	<3.8	<2.7	<1.0	<8.9	<0.5	<0.5
22	Semitron ES225	Quadrant EPP	Ge	mBq/kg	<101	<2.3	<2.0	<1.8	1.8±0.3	513±52	<0.5	<0.6
23	SMD resistor	Farnell	Ge	mBq/pc	2.3±1.0	0.16±0.03	0.30±0.06	0.30±0.05	<0.05	0.19±0.08	<0.02	<0.03
24	SM5D resistor	Finechem	Ge	mBq/pc	0.4±0.2	0.022±0.007	<0.023	<0.016	0.012±0.005	0.17±0.07	<0.005	<0.005
Energy, tracking planes												
25	Kapton-Cu PCB	LabCircuits	Ge	mBq/cm ²	<0.26	<0.014	<0.012	<0.008	<0.002	<0.040	<0.002	<0.002
26	Cuflon	Polyflon	Ge	mBq/kg	<33	<1.3	<1.1	<1.1	<0.6	4.8±1.1	<0.3	<0.3
27	Bonding films	Polyflon	Ge	mBq/kg	1140±300	487±23	79.8±6.6	66.0±4.8	60.0±5.5	832 ±87	<4.4	<3.8
28	FFC/FCP connector	Hirose	Ge	mBq/pc	<50	4.6±0.7	6.5±1.2	6.4±1.0	<0.75	3.9±1.4	<0.2	<0.5
29	P5K connector	Panasonic	Ge	mBq/pc	<42	6.0±0.9	9.5±1.7	9.4±1.4	<0.95	4.1±1.5	<0.2	<0.8
30	Thermopl. connector	Molex	Ge	mBq/pc	<7.3	1.77±0.08	3.01±0.19	2.82±0.15	<0.31	2.12±0.25	<0.022	0.27±0.03
31	Solder paste	Multicore	Ge	mBq/kg	<310	<4.9	<8.0	<6.0	<5.2	<13	<1.0	<1.6
32	Solder wire	Multicore	Ge	mBq/kg	<4900	(7.7±1.2)10 ²	<147	<14		<257	<30	<36
33	Ta capacitor	Vishay Sprague	Ge	mBq/pc	<0.8	0.043±0.003	0.034±0.004	0.032±0.003	<0.010		<0.002	<0.003

For the metal samples prepared for germanium spectrometry, a standard cleaning protocol was followed including the following steps: diamond cut, cleaning with acetone (alcohol was avoided because it may affect their mechanical properties), ultrasound bath with acid-detergent, 63% nitric acid bath, new cleaning with acetone and storing in a sealed plastic bag. Other samples were cleaned in an ultrasonic bath and with pure alcohol.

In the following, results are presented and discussed for the different screened materials grouped according to their position in the detector set-up.

3.1 Shielding

The external passive shielding for NEXT-100 will be made of lead bricks forming a 20-cm-thick lead castle and there will be also an additional 12-cm-thick inner layer of copper to attenuate the radiation originated in the vessel material [3]. Copper is expected to shield in-vessel electronics components if necessary and for the photomultipliers enclosures too.

Lead samples from different suppliers (Mifer,⁵ using two different raw materials, and Tecnibusa⁶ from Spain and COMETA⁷ from Italy) were screened by GDMS. Results are shown in rows #1–4 of table 3; it must be noted that the quantified U and Th concentrations were reported to be at the ultimate limit of detection. The results obtained for COMETA lead are in agreement with the specifications given by the company. Since a large amount of the lead bricks will be ultimately provided by Tecnibusa, two different half-brick samples ($10 \times 10 \times 5$ cm³ each) from this company were measured at LSC; results are presented in rows #5–6 of table 3. This lead has a low activity of ²¹⁰Pb at the level of some tens of Bq/kg. U and Th contamination in lead are normally not very important [6], since radioactive contaminants are effectively removed from lead together with silver [29]. For instance, for Dow Run lead produced by JL Goslar,⁸ activities of tens of μ Bq/kg were measured in [11] and even lower values have been presented by EXO [12] and GERDA [13] experiments as upper limits.

Three copper samples having different origins were also screened by GDMS. One is Electrolytic Tough Pitch (ETP) copper supplied by the Spanish company Sanmetal⁹ while the other two were made of C10100 copper from the Luvata company,¹⁰ having different production mechanism (hot versus cold rolling). The Luvata copper samples were screened together using the Paquito detector as well. All results on copper are shown in rows #7–10 of table 3. The upper bounds on activities derived from the germanium spectrometry measurement were much less stringent than those from GDMS due to its limited sensitivity; hence a new measurement, with much more mass and time and using a bigger germanium detector is foreseen. The cleanest copper we are aware of is that supplied by Norddeutsche Affinerie (Germany);¹¹ very low upper limits for its activity were set in measurements at the Gran Sasso Underground Laboratory [11] and by the EXO Collaboration [12] and even activity from the natural chains and ⁴⁰K was quantified by the XENON experiment [17], at levels of a few tens of μ Bq/kg. Although the GDMS measurement of Luvata

⁵<http://www.mifer.com>.

⁶<http://www.tecnibusa.com>.

⁷<http://www.fonderiaroma.com>.

⁸<http://www.doerun.com>, <http://www.jlgoslar.de>.

⁹<http://www.sanmetal.es>.

¹⁰<http://www.luvata.com>.

¹¹Now re-branded as Aurubis, <http://www.aurubis.com>.

copper has given information only on U and Th concentration, the upper limits derived are at the same level or even better than the results for the Norddeutsche Affinerie copper, and Luvata copper has therefore been chosen as the first option for the NEXT shield.

3.2 Vessel

The pressure vessel of NEXT-100 must be able to hold 15 bar of xenon. It consists of a cylindrical center section (barrel) with two identical torispherical heads on each end [3]. The vessel orientation is horizontal, so as to minimize the overall height. Although it will be ultimately made of stainless steel, the first considered option was titanium, so several samples of both materials have been screened. Inconel (nickel-chromium alloy) will be used to bolt the end-caps to the main body due to its excellent strength properties and therefore its radiopurity has been analyzed too.

Grade 2 titanium was initially proposed to be used for the main components of the vessel. Two samples were screened at LSC, one from a Spanish supplier, Titanio SMP¹² and the other from Titanium Metal Supply.¹³ Results are shown in rows #11–13 of table 3. The Ti SMP sample was screened using two different germanium detectors, GeOroel (row #11) and GeTobazo (row #12) as a cross-check exercise; the small differences found can be well understood taking into account the differences in the background rates of the detectors in several energy ranges. Thanks to the much larger mass available in the sample, upper limits on activities derived for the Ti Metal Supply sample are much lower than for the Ti SMP sample and it has been possible to quantify the activity of the lower part of the ²³²Th chain. Production of ⁴⁶Sc, beta emitter with $Q=2366.7$ keV and $T_{1/2}=83.8$ days, has been also observed for this sample; it must have been generated by (n,p) reactions on ⁴⁶Ti induced by fast neutrons. The LUX collaboration has carried out an exhaustive analysis of Ti samples [30], obtaining different levels of radiopurity for them; the presence of ⁴⁶Sc is usually identified.

A great deal of activity measurements for different types of stainless steel (SS) can be found in the literature (see for example refs. [10, 18, 20, 21]) showing a wide range of values. One sample of type 304L (a vacuum system piece from Pfeiffer¹⁴) was screened using the Paquito detector (see results in row #14 of table 3), obtaining the usual quite high levels of activity from natural chains. Material referenced as austenitic 1.4571 (also 316L) has been extensively studied by XENON [17] and GERDA [31] experiments, finding materials supplied by the Nironit company¹⁵ with activity levels (values or upper limits) of even tenths of mBq/kg for isotopes from the natural chains in the best cases. Three samples of 316Ti stainless steel supplied by Nironit were screened at LSC and results are presented at rows #15–17 of table 3; they have different thickness since they are intended to be used in different parts of the NEXT pressure vessel (10 mm for body, 15 mm for end-caps and 50 mm for flanges). Activities from ⁶⁰Co and ⁵⁴Mn, commonly present in steel, have been quantified for the three samples. Results for cosmogenic ⁵⁴Mn, not quoted in table 3, are 0.29 ± 0.05 , 0.55 ± 0.07 and 0.97 ± 0.14 mBq/kg for increasing thickness of sample. For the 10- and 15-mm-thick samples, upper bounds for all the other emitters investigated have been derived; for the 50-mm-thick sample, the activity of the isotopes of the ²³²Th chain has been quantified,

¹²<http://www.titaniosmp.com>.

¹³<http://www.titaniummetalsupply.com>.

¹⁴<http://www.pfeiffer-vacuum.com>.

¹⁵<http://www.nironit.de>.

pointing to secular equilibrium. It is worth noting that the sensitivity for the thickest sample was worse than for the other two, because of the lower mass available in the sample (see table 2) and the lower efficiency detection due to self-absorption. The activity values obtained for Nironit 316Ti stainless steel are of the order or below NEXT requirements; therefore, the booked batches from where the samples were taken will be used for pressure vessel construction.

Samples of inconel 718 and inconel 625 from the Spanish company Mecanizados Kanter¹⁶ were screened at LSC and results are shown in rows #18-19 of table 3. No previous results on radiopurity of this material have been found; upper limits on activities of the lower parts of the ^{238}U and ^{232}Th chains have been set at some mBq/kg for both types of inconel. Presence of ^{58}Co was identified for inconel 625.

3.3 High Voltage and electroluminescence components

The main body of the *field cage* to be placed inside the vessel will be made of high density polyethylene, with attached copper strips connected to resistors [3]; PEEK was also considered as an alternative. Wire meshes separating the different field regions of the detector, including the electroluminescence volume, will be made of stainless steel. To improve the light collection efficiency of the detector, reflector panels coated with a wavelength shifter will cover the inner part of the field cage. This *light tube* will be made of Tetratex® fixed over a 3M substrate, coated with tetraphenyl butadiene (TPB) [3]. The ArDM experiment has screened specifically polytetrafluoroethylene (PTFE) Tetratex from Donaldson Membranes¹⁷ by Inductively Coupled Plasma Mass Spectrometry (ICPMS) [32] and TPB from two different manufacturers were measured at SNOlab [18].

A sample of PEEK from Sanmetal Spanish company was screened using the Paquito detector; values obtained are shown in row #20 of table 3, pointing to a non-negligible activity. Only upper bounds on activity of PEEK were presented in [19].

Polyethylene from IN2 Plastics company¹⁸ has a very good radiopurity according to XENON results [17] and a sample of High Molecular Weight polyethylene (type PE500) was therefore chosen for screening at LSC. First results, shown in row #21 of table 3, have produced only upper limits for common radioisotopes. Semitron® ES225 plastic produced by Quadrant Engineering Plastic Products¹⁹ has been also measured and the results on its radiopurity are presented in row #22 of table 3; in this case, a quite high activity of ^{40}K has been registered.

Surface Mount Device (SMD) resistors supplied by Farnell²⁰ and by Finechem²¹ were screened at LSC (see results in rows #23–24 of table 3). Activity values obtained for Finechem resistors are much lower than for Farnell ones, and very similar to those obtained in [17] for resistors of the same type and company.

¹⁶<http://www.mecanizados-kanter.es>.

¹⁷<http://www.donaldson.com>.

¹⁸<http://www.in2plastics.com>.

¹⁹<http://www.quadrantplastics.com>.

²⁰<http://www.farnell.com>.

²¹<http://www.jfine.co.jp>.

3.4 Energy and tracking planes components

The energy measurement in NEXT is provided by the detection of the electroluminescence light by an array of photomultipliers (the so-called *energy plane*), located behind the cathode mesh. Those photomultipliers will also record the primary scintillation light that indicates the start of the event. Each photomultiplier will be sealed into individual, pressure resistant, vacuum tight copper enclosures and will be coupled to the sensitive volume through a sapphire window coated with TPB. The tracking function will be provided by a plane of MPPCs operating as sensor pixels and located behind the transparent electroluminescence gap. The MPPCs will be mounted on square boards named Dice Boards (DB); each DB will contain 8×8 sensors with a pitch of ~ 1 cm and will be coated with TPB too. The array of DBs, with ~ 7000 channels, forms the *tracking plane*. Front-end electronics boards for this tracking plane will be inside the pressure vessel, behind a copper shield. Details can be found in [3].

Although different models of photomultipliers were under consideration, Hamamatsu R11410MOD was finally chosen for the NEXT-100 detector since radiopurity specifications provided by Hamamatsu are better than for other models (~ 3.3 mBq/PMT of ^{238}U , ~ 2.3 mBq/PMT of ^{232}Th and ~ 5.7 mBq/PMT of ^{40}K). The XENON [17] and LUX [33] experiments have studied the radiopurity of this model, finding even lower activities. These levels would satisfy NEXT requirements, but the dispersion found in different measurements imposes a dedicated program to control the photomultiplier radiopurity, which will be undertaken in LSC. Sapphire windows (from Swiss Jewel Company²²) were screened by EXO [12] with activities of tenths of mBq/kg from the natural chains; a measurement of sapphire windows for NEXT is underway.

Silicon, used for the MPPCs at the tracking plane, is, as germanium, a very radiopure material with typical intrinsic activities of ^{238}U and ^{232}Th at the level of few $\mu\text{Bq/kg}$ [6]. In any case, a specific measurement of MPPCs is pending, either separately or after being mounted on the boards.

Several materials have been taken into consideration for Printed Circuit Boards (PCBs) and a large number of radiopurity measurements can be found in [21]. FR4 was disregarded for both an unacceptable high rate of outgassing and bad radiopurity (a fast measurement made with Paquito detector pointed to activities of hundreds of mBq/kg for natural chains); it seems that the glass fiber-reinforced materials at base plates of circuit boards can be a source of radioactive contamination [6]. Components made of just kapton (like cirlex) and copper offer very good radiopurity, as shown in the measurements of kapton-copper foils in [24] and in the screening of a monolayer PCB made of kapton and copper supplied by LabCircuits²³ using the Paquito detector, presented in row #25 of table 3. Also cuflon® offers low activity levels, as shown in the measurement of samples from Crane Polyflon²⁴ by GERDA [14] and at [15], using both ICPMS and Ge gamma spectroscopy. A first measurement of Polyflon cuflon made of a 3.18-mm-thick PTFE layer sandwiched by two 35- μm -thick copper sheets was made for NEXT and results are shown in row #26 of table 3. Cuflon has been chosen for the Dice Boards. Adhesive films to glue cuflon sheets are used to prepare multilayer PCBs; a sample of bonding films made of a polyolefin co-polymer and supplied also by Crane Polyflon were screened and results are presented in row #27 of table 3. Despite the high

²²<http://www.swissjewel.com>.

²³<http://www.lab-circuits.com>.

²⁴<http://www.polyflon.com>.

activities measured, the use of these films is not completely discarded since the necessary mass for each Dice Board is of only 0.59 g.

Information is also available on the radiopurity of different types of connectors [10, 18, 21]. FFC/FCP (Flexible Printed Circuit & Flexible Flat Cable) connectors were initially considered for the DBs; connectors supplied by Hirose²⁵ were selected and screened using the Paquito detector, as well as similar connectors from Panasonic²⁶ (see results in rows #28–29 of table 3). Both types of connectors show activities of at least several mBq/pc for isotopes in ²³²Th and the lower part of ²³⁸U chains and for ⁴⁰K. These values, according to [21], are of the same order than those for similar connectors supplied by Molex;²⁷ all of them are made of Liquid Crystal Polymer (LCP), thus it seems that the activity measured is related to this material. Thermoplastic connectors 503066-8011 from Molex were also screened (see results in row #30 of table 3) finding activities slightly smaller but of the same order. Since these levels are too high for NEXT sensitivity, it was concluded that this kind of connectors must be avoided at DBs and the collaboration is at the moment foreseeing a direct bonding of the FFCs/FCPs to the DBs. A sample of lead free SnAgCu solder paste supplied by Multicore (ref. 698840) has been screened and results are presented in row #31 of table 3). ^{108m}Ag, induced by neutron interactions and having a half-life of $T_{1/2} = 438$ y, has been identified in the paste, with an activity of 5.26 ± 0.40 mBq/kg, while upper limits of a few mBq/kg have been set for the common radioactive isotopes; consequently, this solder paste could be used without concern. Solder wire with similar composition from Multicore (ref. 442578) was also screened (see row #32 of table 3), finding in this case a high activity of the lower part of the ²³⁸U chain. An activity of ²¹⁰Pb of $(1.2 \pm 0.4) \times 10^3$ Bq/kg has been deduced using the bremsstrahlung emission from its daughter nuclide ²¹⁰Bi [34].

Concerning electronic components at the boards, ceramic capacitors seem to be quite radioactive [21] and have been disregarded. Tantalum capacitors (Vishay Sprague 597D²⁸) have been screened at LSC and results are presented in row #33 of table 3; activity levels are lower than for tantalum capacitors supplied by AVX,²⁹ from database in [21]. In addition to activities shown in table 3, the presence of ¹⁸²Ta (beta emitter with $Q = 1814.3$ keV and $T_{1/2} = (114.74 \pm 0.12)$ days [35], produced by neutron activation on ¹⁸¹Ta) has been identified through the observation of many of its gamma emissions.

4 Summary

A thorough radiopurity control is being performed for the construction of the NEXT-100 double beta decay experiment to be operated at LSC. Evaluation of activity levels in the most relevant materials has been carried out based on GDMS and on ultra-low background germanium gamma-ray spectrometry at LSC. Some of the results presented here represents the first measurements performed using the germanium detectors of the LSC Radiopurity Service. This kind of data have a widespread interest, being also useful for other experiments investigating rare phenomena in under-

²⁵<http://www.hirose.com>.

²⁶<http://www.panasonic-electric-works.com>.

²⁷<http://www.molex.com>.

²⁸<http://www.vishay.com>.

²⁹<http://www.avx.com>.

ground laboratories. Radiopurity information is expected to be helpful not only for the selection of materials that are radiopure enough to minimize the background level, but also for the development of the NEXT detector background model in combination with Monte Carlo simulations.

Adequate materials for the external and internal passive shields have been identified. Titanium and stainless steel samples were analyzed for the construction of the pressure vessel. The good radiopurity found for the 316Ti stainless steel supplied by the Nironit company, of the order of tenths of mBq/kg for the isotopes at the lower part of the ^{238}U and ^{232}Th chains, confirmed that this material can be used for the detector vessel in combination with an inner copper shield. The background contribution of the vessel and both internal and external shields has been studied, assuming the upper limits on the activities obtained here for Nironit 316Ti stainless steel, Luvata copper and Tecnibusa lead; the preliminary results indicate, for each material, a background level in the RoI at most of the order of 10^{-4} counts $\text{keV}^{-1} \text{ kg}^{-1} \text{ y}^{-1}$, after all cuts (fiducial and topological). Construction of shielding and vessel is thus proceeding.

Selection of in-vessel components to be used at the energy and tracking planes has been performed too, helping in the design of DB and front-end electronics boards. While resistors, capacitors and solder paste of acceptable radiopurity have been found, board-to-cable connectors containing LCP have been discarded since they have activities of few mBq per piece for isotopes of the ^{238}U and ^{232}Th chains. Cufion, being radiopure enough, has been chosen for the construction of the boards for the MPPCs.

Further radiopurity measurements using germanium detectors at LSC are scheduled before the installation and operation of the NEXT detector with the aim to control all the components that might contribute significantly to the background. Main efforts will be devoted to the careful selection of photomultipliers of the chosen model (Hamamatsu R11410MOD) and to the screening of related components (e.g., sapphire windows, bases and cans) at the energy plane and light tube materials. This task is essential to achieve the required background level in the region of interest imposed by the NEXT sensitivity goal.

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