

Constraining Radon Backgrounds in LZ

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Abstract. The LZ dark matter detector, like many other rare-event searches, will suffer from backgrounds due to the radioactive decay of radon daughters. In order to achieve its science goals, the concentration of radon within the xenon should not exceed $2\mu\text{Bq/kg}$, or 20 mBq total within its 10 tonnes. The LZ collaboration is in the midst of a program to screen all significant components in contact with the xenon. The four institutions involved in this effort have begun sharing two cross-calibration sources to ensure consistent measurement results across multiple distinct devices. We present here five preliminary screening results, some mitigation strategies that will reduce the amount of radon produced by the most problematic components, and a summary of the current estimate of radon emanation throughout the detector. This best estimate totals $< 17.3\text{ mBq}$, sufficiently low to meet the detector's science goals.

INTRODUCTION

Radon and radon daughters pose significant, and often dominant, backgrounds to low-background and rare-event search experiments. These include detectors of neutrinos [1], neutrinoless double-beta decay [2], and dark matter [3, 4, 5, 6]. Within the LUX-Zeplin (LZ) detector, radon and radon daughters can diffuse into the fiducial volume within the liquid xenon. Here, radon's prompt granddaughter ^{214}Pb decays by a beta emission that may be mistaken for a nuclear recoil $\sim 0.5\%$ of the time [6]. This decay is separated in time from its parent and daughter decays with half-lives of 26.8 min and 19.9 min respectively, and it occurs without any associated gamma emission 9.2% of the time. These factors together yield a background that may be indistinguishable from a dark-matter-induced nuclear recoil, that is distributed evenly throughout the fiducial volume, and that cannot be tagged on an event-by-event basis. Events derived from radon are expected to be the single largest background in the LZ detector based on a projected activity of $1.73\mu\text{Bq/kg}$, accounting for 3.66 of the 6.29 total WIMP-like events over a 1,000 day run [6].

MITIGATION OF RADON SOURCES IN LZ

The LZ experiment will rely on two significant mitigation strategies that reduce the amount of radon produced by certain components. In addition, the low temperature of parts in contact with the liquid xenon is known to reduce the diffusion of radon through materials, which can reduce the amount of radon emanated. For sufficiently thick materials, the amount of radon diffusing out of a material is proportional to $\sqrt{D(T)\tau}$, where $\tau = 5.5$ days is the lifetime of ^{222}Rn and $D(T)$ follows the Arrhenius relation [7, 8]: $D(T) \propto \exp(-T_0/T)$ where T_0 is a temperature corresponding to the characteristic energy of diffusion for radon through a particular material. This dependence can result in significantly lower rates of radon emanation at lower temperatures in materials dominated by diffusion, like plastics (however, this has little or no effect on surface contamination including dust). For example Fluorinated Ethylene Propylene (FEP) has $T_0 \approx 6800^\circ\text{K}$ [8, 9], so the diffusion coefficient $D(T)$ at 161°K is reduced by a factor of 2.5×10^8 from room

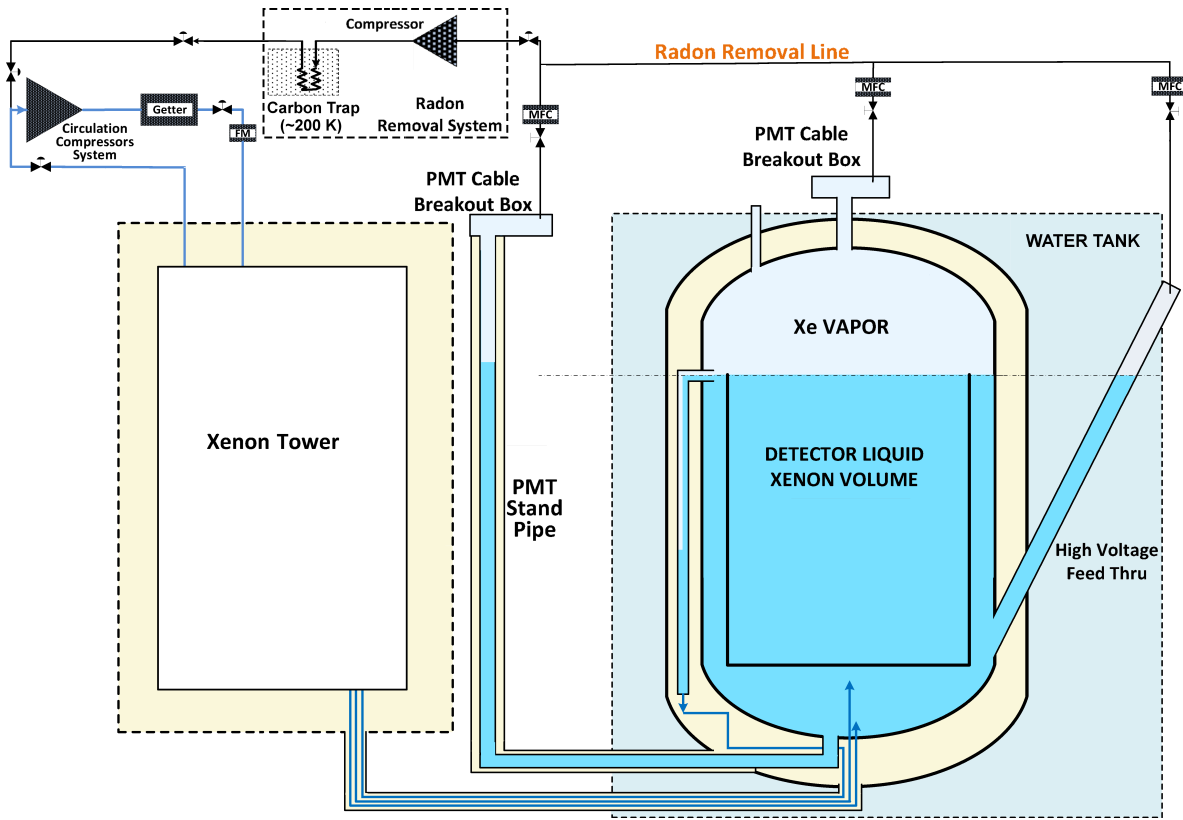


FIGURE 1. A simplified schematic of the xenon recirculation in LZ showing the primary recirculation path through the xenon tower (left). The circulation path from the top and bottom PMT breakout boxes and from the high voltage feedthrough passes through the radon removal system (top middle) before re-entering the main recirculation. Here the vacuum space is yellow, the liquid xenon is blue, the gaseous xenon is pale blue, the main liquid recirculation is dark blue lines, and the main gaseous recirculation is light blue lines. A full version can be found in [6].

temperature at 298°K, resulting in a reduction in the radon production by about four orders of magnitude. Radon emanation measurements (described below) performed at room temperature therefore significantly over-estimate the contribution of plastics that will be cold within the LZ detector. LZ will take advantage of this feature by cladding all cables in FEP, or a similar plastic, to effectively block radon produced below the cable's surface from diffusing into the xenon.

This strategy should significantly reduce the amount of radon produced by the sections of cable that are immersed in liquid xenon and held at 161°K. The ends of the cables where they meet the feedthroughs to lab space, however, are at room temperature and still produce radon at a higher rate. In order to mitigate this source of radon, LZ will employ a cooled-carbon trap, similar to [10]. While it is impractical to use such a trap to filter out the full recirculation of xenon (planned at 500 slpm), LZ is developing a device capable of removing 90% of the radon from a modest flow of only 1 slpm xenon [6]. This system will purify the xenon gas from low-flow regions of the detector before letting it re-enter the main xenon flow. As shown in Figure 1, the regions affected by this system include the gaseous spaces around the umbilical cable as well as cables from both the top and bottom PMTs, including all cable feedthroughs.

The feedthroughs carrying HV for the detector's PMTs contact the xenon in a space purified by the cooled carbon trap, so only 10% of the radon produced will affect LZ. However, the insulator on these feedthroughs is an alumina ceramic with the potential to be high in uranium, and therefore high in radon production. These feedthroughs, and potentially other components, will be encapsulated in 3 mm of MasterBond epoxy. This epoxy, demonstrated by BOREXINO to be low-radioactivity [11], will create a barrier separating the feedthrough ceramic from the xenon gas and significantly reducing its radon contribution. The effectiveness of this epoxy to prevent radon emanation is currently under measurement.

RADON EMANATION MEASUREMENTS

Four institutions within the LZ collaboration have developed systems to measure the amount of radon produced by items: University of Alabama (UA) employs a system that dissolves Rn into liquid scintillator and identifies radon by the ^{214}Bi - ^{214}Po timing coincidence; University College London (UCL) [12], University of Maryland (UMd), and South Dakota School of Mines and Technology (SDSM&T) use systems where charged radon daughters are collected on a silicon-pin diode which measures the subsequent alpha decays. These are summarized in Table 1.

With four institutions making measurements, using two different measurement techniques, ensuring consistent measurements is critical. Each system has been individually calibrated by adding known amounts of radon to a vessel via a calibrated flow-through source. In addition, LZ is engaging in a cross-calibration campaign between the various institutions in which two different samples are being measured in each emanation chamber. The first sample is a strip of rubber that produces $O(10)$ mBq (generously lent by J. Farine from work with EXO [2]), testing the systems' calibrated measurement efficiency. The second sample, a set of thoriated TIG welding rods, produces $O(1)$ mBq and tests the systems' background measurements — calibration with this sample will begin soon.

Each institution's measurements, relative to EXO's original measurements, are shown in Table 1. Most institutions agree to within uncertainties; UCL's anomalous result is under further study. The emanation and measurement process for the SDSM&T system is described briefly below.

TABLE 1. Seven vessels, at four institutions, are available to screen LZ samples for radon emanation with the throughputs shown. Most of the measurements are consistent within quoted errors. SDSM&T's 300 L and UCL's second 2.6 L vessels are still awaiting calibration.

Institution	Technology	Sample Volume	Blank Rate	Measurement (relative to EXO)	LZ Sample Throughput
UCL	Electrostatic PIN-diode	2.6 liters	0.2 mBq	1.49 ± 0.05	6 / year
		2.6 liters	0.4 mBq		
UMd	Electrostatic PIN-diode	4.7 liters	0.2 mBq	$1.13 \pm 0.06 \pm 0.18$	12 / year
SDSM&T	Electrostatic PIN-diode	13 liters	<0.3 mBq	$0.89 \pm 0.12 \pm 0.15$	18 / year
		300 liters	0.3 mBq		
UA	Liquid Scintillator Coincidence	2.6 liters	0.2 mBq	$0.83 \pm 0.15 \pm 0.08$	24 / year
		2.6 liters	0.2 mBq	$0.85 \pm 0.19 \pm 0.08$	

Sample preparation includes a hand-scrub of the entire surface with isopropyl alcohol-soaked lint-free wipes in order to remove surface dust. The sample is then sealed in a vacuum-tight emanation chamber, flushed with boil-off nitrogen, evacuated to near-vacuum, and allowed to out-gas before a measurement. This outgasing time ranges from a few days for metallic samples to weeks for porous samples — these typically absorb atmospheric radon which can subsequently diffuse out of the sample during a measurement, overwhelming the contribution intrinsic to the sample itself. Finally, the chamber is flushed with boil-off nitrogen and filled to ≈ 100 Torr. An activated charcoal trap, cooled to -77°C by a bath of isopropyl alcohol and dry ice, further reduces the radon content of the boil-off nitrogen. Use of 100 Torr rather than vacuum ensures that radon atoms produced by the nuclear recoil of a ^{226}Ra decay are captured in the gas, rather than embedding in the metallic wall of the emanation chamber. An 86.3 keV ^{222}Rn atom, produced by the decay of ^{226}Ra , has a stopping range of 5 meters at a modest vacuum of 0.1 Torr compared with 500 μm in 100 Torr nitrogen gas [13].

As shown in Figure 2 left, SDSM&T employs two electropolished stainless steel vessels for radon emanation — one 13 L, and one 300 L. Each sample is allowed to emanate in a vessel for around one week. During this time, the activity of radon within the chamber ($C(t)$) begins to approach the equilibrium activity produced by the material (C_0) as $C(t) = C_0(1 - \exp(-t/\tau))$, where $\tau = 5.5$ days is the ^{222}Rn lifetime.

After this emanation period the nitrogen gas, and the radon contained therein, is evacuated from the emanation chamber through a cold trap. During this process, the pressure inside the emanation chamber drops to ≈ 20 Torr, and is refilled back to ≈ 100 Torr, four times. This pressure cycling helps to ensure that the vast majority of radon from the chamber is captured, and that there are no pockets of gas that might remain isolated. The cold trap is a length of $\frac{3}{8}$ " electropolished stainless-steel tubing filled with brass wool; this tubing is submersed in a liquid nitrogen bath so that radon atoms adsorb to the surface of the brass wool. Since the pressure within this trap remains low throughout the process, the nitrogen carrier gas does not condense within the trap.

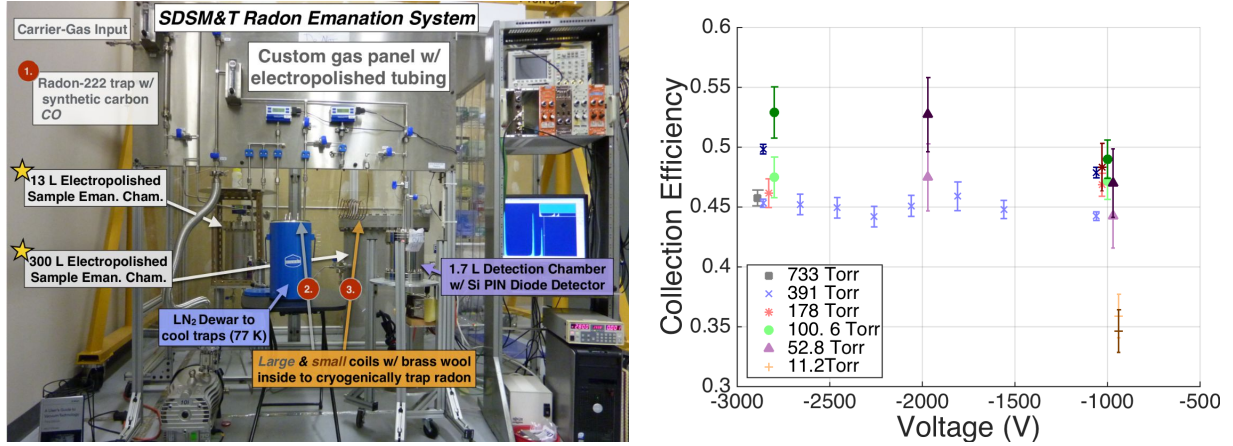


FIGURE 2. *Left:* The radon emanation measurement system at SDSM&T – components and operation described in text. *Right:* The efficiency for collecting radon daughters onto the silicon-pin diode in a nitrogen atmosphere does not vary significantly with changes in voltage or pressure. Lighter points correspond to collection efficiency for ^{218}Po , while the darker points are for ^{214}Po . SDSM&T operates at around 100 Torr and -2000 V.

Once the radon has been frozen into this first, large trap, the trap is warmed back to atmospheric pressure and a flow of fresh boil-off nitrogen gas carries the radon from this large trap into a second, smaller cold trap whose volume is small compared to that of the detection chamber. This second trap is made from $\frac{1}{8}$ " copper tubing and is also filled with brass wool and cooled with liquid nitrogen. Finally, the small copper trap is warmed to room temperature, and the 1.3 liter detection chamber is filled to 100 Torr with fresh boil-off nitrogen flowing through the trap. This process robustly and repeatably transfers at least 95% of the radon from the emanation chamber into the detection chamber.

The detection chamber is a 1.3 liter electropolished stainless-steel vessel containing a 1" silicon-pin diode held at -2000 V. Following the decay of ^{222}Rn atoms, $\approx 88\%$ of the ^{218}Po daughters are positively charged and follow the electric field lines to the diode [14]. As shown in Figure 2 right, the probability for collecting for each prompt radon daughter onto the diode is $\approx 45\%$. The subsequent alpha decays of ^{218}Po and ^{214}Po produce clear signals in the diode, which is read out by a multi-channel analyzer.

PRELIMINARY RESULTS

The radon emanation screening program for LZ is now well underway with 63 measurements completed. Table 2 lists five preliminary measurements, along with calculations of the components' total contributions within the LZ experiment with and without the mitigation strategies discussed above. The 17 km of Axon cable, used to transfer high voltage to and signals from the detector's PMTs, would produce 15.7 mBq at room temperature. This contribution alone would approach the allowable limit within the detector. Accounting for the reduction by both the low temperature of the sections within the liquid xenon and the radon removal by the cooled carbon trap results in the modest estimate of 0.8 mBq.

TABLE 2. Preliminary results from the LZ radon emanation screening program. The full contribution is the amount of radon produced by the total amount of each material in LZ, while the mitigated contribution accounts for radon reduction by the cooled carbon trap and, in the case of the Axon cables, a reduction due to temperature. Items indicated with * are expected to produce less radon by an unknown amount due to low temperature. This umbilical cable listed was rejected and will not be used.

Material	Vessel	Result	Units	Full Contribution	Mitigated Contribution
Axon Cable	SDSM&T 300 L	0.93 ± 0.27	mBq/km	15.7 ± 4.5 mBq	0.8 ± 0.23 mBq
HV Feedthrus	UA 2.6 L	0.5 ± 0.2	mBq/unit	6 ± 2.4 mBq	0.6 ± 0.24 mBq
PMT Bases	UCL 2.6 L	0.28 ± 0.17	mBq/unit	1.8 ± 1.1 mBq	1.8 ± 1.1 mBq*
PTFE	SDSM&T 300 L	< 0.015	mBq/m ²	< 1.29 mBq	< 1.29 mBq*
Umbilical Cable	SDSM&T 13 L	0.26 ± 0.06	mBq/m	2.1 ± 0.5 mBq	Rejected

TABLE 3. List of materials in contact with Xe, indicating the quantity of the material and the goal for radon emanation from the material. The estimates of radon emanation are expected usually to be conservative, as they are based on the most similar object or material for which emanation rates are available in the literature, and use only conservative models for reduction of radon emanation at LXe temperatures. Some materials are expected to emanate less radon when cold, so room-temperature emanation quantities (labeled with *) may be reduced. Expected reduction of radon by the carbon trap is included in estimates for those components affected (labeled with †). Values in **bold** are based on measurements performed by LZ.

Material	Component(s)	Quantity	Unit	Estimate (mBq)
Al ₂ O ₃ resistor	PMT Bases	9790	#	0.58*
BaTiO ₃ capacitor	PMT Bases	3010	#	0.016*
Cirlex	PMT Bases	6000	cm ²	0.37*
Titanium	Cryostat, PMT Mounts, Field Rings, Grid Supports	412,000	cm ²	0.41
PTFE	Reflectors, Field Cage	840,000	cm ²	<1.3*
PMT cabling [†]	PMT Cabling	17,000	m	0.8
PMT feedthrough [†]	PMT Feedthrough	88	#	<0.24
Steel conduit [†]	Cabling Conduit	100,000	cm ²	0.055
R11410 PMT	R11410 PMT	488	#	1.26
R8520 PMT	R8520 PMT	90	#	0.15
R8778 PMT	R8778 PMT	38	#	0.09
Polyethylene	HV Umbilical	4200	cm ²	0.10*
Tin-coated copper	HV Umbilical	11,000	cm ²	0.002
Tivar	HV Umbilical	3894	cm ²	0.004*
Acetal	HV Umbilical	195	cm ²	0.0002*
Copper	HV Umbilical	39	cm ²	0.000007
Epoxy	HV Umbilical, Feedthroughs	1000	cm ²	0.0001*
Steel	Cryostat Seals, Xe Recirculation	135,000	cm ²	0.104
Recirculation pump	Xe Recirculation	1	#	0.1
Purification getter	Xe Recirculation	2.5	kg	1.34
Transducers & Valves	Xe Recirculation	30	#	0.17
Welds	Recirculation System, Cryostat	32.3	m	0.11
Dust		1	g	10.0
Total				< 17.3

The 20 mBq requirement combined with the large total surface area exposed to xenon within LZ requires the measurements of samples with large surface area. One example is PTFE which, making up the TPC and reflector lining the inner cryostat, will have an area 84 m². The first screening sample of this material was 18 m² and fit only within the collaboration's largest emanation chamber, the 300L vessel at SDSM&T. The result of screening even this large a sample was consistent with zero, with a 90% upper limit of 0.015 mBq/m².

An additional significant contributor of radon within LZ will be dust on surfaces. With only 500 ng/cm² on xenon-wetted surfaces, the experiment will hold about 1 gram of dust. Assuming typical activity of ≈ 40 mBq/g of uranium, and a conservative estimate that 25% of the radon produced by surface dust escapes into xenon, this contaminant will contribute 10 mBq of radon to the experiment.

In order to reach its science goals, LZ must limit the activity of ²²²Rn within the 10 tonnes of xenon to 2 μ Bq/kg, or 20 mBq total. The experiment has kept an inventory of all components that will be in contact with the xenon along with an estimate of radon emanation from each [15, 16, 17, 18]. This estimation includes measurements from the screening program described above and is shown in Table 3. This radon budget helps to direct screening and mitigation efforts and informs experiment background models. The best estimates at time of writing indicate a contribution of 10 mBq from dust, and < 7.3 mBq from materials for a total of < 17.3 mBq total within the detector's xenon.

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REFERENCES

- [1] G. Bellini *et al.* (Borexino), JCAP **1308**, p. 049 (2013), arXiv:1304.7381 [physics.ins-det] .
- [2] J. B. Albert *et al.* (EXO-200 Collaboration), Phys. Rev. C **92**, p. 015503Jul (2015).
- [3] A. Bradley *et al.*, Physics Procedia **61**, 658 – 665 (2015), 13th International Conference on Topics in Astroparticle and Underground Physics, TAUP 2013.
- [4] E. Aprile *et al.* (XENON), JCAP **1604**, p. 027 (2016), arXiv:1512.07501 [physics.ins-det] .
- [5] J. Brack *et al.* (DRIFT), Journal of Instrumentation **9**, p. P07021 (2014).
- [6] B. J. Mount *et al.*, (2017), arXiv:1703.09144 [physics.ins-det] .
- [7] S. Arrhenius, *Über die Dissociationswärme und den Einfluss der Temperatur auf den Dissociationsgrad der Elektrolyte* (Wilhelm Engelmann, Leipzig, 1889), pp. 96–116.
- [8] A. Piepke and K. Pushkin, “Estimate of the EXO-200 radon production inside the TPC,” (2011), EXO Internal Memo.
- [9] S. Pauly, “Permeability and diffusion data,” Hoechst AG, Werke Kalle.
- [10] K. Abe *et al.*, Nuclear Instruments and Methods in Physics Research Section A: Accelerators, Spectrometers, Detectors and Associated Equipment **661**, 50 – 57 (2012).
- [11] C. Arpesella *et al.* (BOREXINO), Astropart. Phys. **18**, 1–25 (2002), arXiv:hep-ex/0109031 [hep-ex] .
- [12] B. Soule, “Radon emanation chamber : High sensitivity measurements for the supernemo experiment,” 08 (2013).
- [13] J. Zeigler, The stopping ranges of ions in matter, 2008.
- [14] J. Porstendörfer, Radiation Protection Dosimetry **94**, 365–373 (2001).
- [15] G. Zuzel, “Highly Sensitive Measurements of ^{222}Rn Diffusion and Emanation,” in *Topical Workshop on Low Radioactivity Techniques: LRT 2004.*, American Institute of Physics Conference Series, Vol. 785, edited by B. Cleveland, R. Ford, and M. Chen (2005), pp. 142–149.
- [16] M. Liu, H. Lee, and A. McDonald, Nuclear Instruments and Methods in Physics Research Section A: Accelerators, Spectrometers, Detectors and Associated Equipment **329**, 291 – 298 (1993).
- [17] E. Aprile *et al.* (XENON100 Collaboration), Phys. Rev. D **83**, p. 082001Apr (2011).
- [18] G. Zuzel and H. Simgen, Applied Radiation and Isotopes **67**, 889 – 893 (2009), 5th International Conference on Radionuclide Metrology - Low-Level Radioactivity Measurement Techniques ICRM-LLRMT’08.