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^{222}Rn emanation measurements for the XENON1T experiment

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Abstract The selection of low-radioactive construction materials is of utmost importance for the success of low-energy rare event search experiments. Besides radioactive contaminants in the bulk, the emanation of radioactive radon atoms from material surfaces attains increasing relevance in the effort to further reduce the background of such experiments. In this work, we present the ^{222}Rn emanation measurements performed for the XENON1T dark matter experiment. Together with the bulk impurity screening campaign, the results enabled us to select the radio-purest construction materials, targeting a ^{222}Rn activity concentration of $10\ \mu\text{Bq/kg}$ in 3.2 t of xenon. The knowledge of the distribution of the ^{222}Rn sources allowed us to selectively eliminate problematic components in the course of the experiment. The predictions from the emanation measurements were compared to data of the ^{222}Rn activity concentration in XENON1T. The final ^{222}Rn activity concentration of $(4.5 \pm 0.1)\ \mu\text{Bq/kg}$ in the target of XENON1T is the lowest ever achieved in a xenon dark matter experiment.

1 Introduction

Many cosmological observations suggest that a large fraction of the total matter density of the Universe is made up of an unknown form of dark matter [1]. However, despite a large experimental effort, dark matter has not yet been discovered. XENON1T [2] was the largest and most sensitive so far in the series of XENON direct dark matter search experiments [3, 4]. Its successor XENONnT started data-taking in 2020. The primary aim of these experiments is the detection of weakly interacting massive particles (WIMPs), a promising dark matter candidate [5]. As in other astroparticle physics experiments [6–8] liquid xenon is used as an efficient target for particle detection. The XENON detectors are dual-phase time projection chambers (TPCs) with a gaseous layer of xenon on top of the target. Particles interacting with xenon nuclei or the atomic electrons create scintillation light and

ionization electrons. The light is detected by two arrays of UV-sensitive photomultiplier tubes (PMTs) on top and bottom of the detector. The ionization electrons are drifted upwards to the liquid-gas interface, where they are extracted and create a second, delayed scintillation light signal. Both signals are used to gain information about the location and energy of the interaction. They are also used to reject background either by the event multiplicity (multi-scatter versus single-scatter events) or by the type of interaction (electronic recoil versus nuclear recoil events).

XENON1T operated for two years, starting from December 2016. Similarly to other astroparticle physics experiment looking for rare events, it required an extremely low background level. Throughout the different generations of the XENON experiments, external background sources have been suppressed, e.g. by an improved external shield and xenon self-shielding and by the mitigation of radioactivity from materials. Their level was marginal in XENON1T and intrinsic background sources became dominant. Among them, the radioactive isotope ^{222}Rn induced the leading background component [9]. Its long-lived mother nucleus ^{226}Ra ($T_{1/2} = 1600$ years) is part of the primordial ^{238}U decay chain and thus present in most materials. Once ^{226}Ra decays, the created noble gas isotope ^{222}Rn may emanate from inner surfaces into the xenon volume.

As ^{222}Rn has a relatively long half-life ($T_{1/2} = 3.8$ days), it can reach the active dark matter target, where the β -decays of its daughter isotope ^{214}Pb can mimic signal events. To achieve the scientific goal of XENON1T, a ^{222}Rn activity concentration of $10\ \mu\text{Bq/kg}$ was required [9]. Other radon isotopes may also lead to background events. However, their contribution was strongly suppressed due to their small abundance in the detector and much shorter half-lives, that did not allow for their dispersion within the target volume. XENON1T has performed a comprehensive bulk impurity screening campaign to select radio-pure materials using high purity germanium (HPGe) spectroscopy and Inductively coupled plasma mass spectrometry (ICP-MS) [10]. However, the measured ^{226}Ra bulk activity can in general not be used to predict how much ^{222}Rn emanates from the material, because surface impurities may become dominant. This made dedicated ^{222}Rn emanation measurements necessary, which are described in this article.

There are two ways for a radon atom to leave the material in which it is produced: by recoil or by diffusion [11]. In the first case, the decay occurs directly on or below the material's surface. The recoil energy, which the radon nucleus receives during the α -decay of its mother radium nuclide (85 keV [12] in the case of ^{222}Rn), is sufficient to eject it from the material. In the second case, the radon atom diffuses inside the bulk

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of the material. If it reaches a boundary surface before its decay, it will be emanated. Data on radon diffusion in metals are rare, but its diffusion coefficient is even smaller than the minuscule one of xenon in metals [13]. Thus, it is reasonable to assume that radon diffusion plays a significant role only in soft or porous materials. As a consequence, a radon emanation measurement of metals is mostly a test of surface impurities.

The article is structured as follows. Section 2 discusses the employed ^{222}Rn assay techniques. In Sect. 3, we present the screening results of investigated materials, most of which were eventually used for the construction of XENONIT. Section 4 describes the overall ^{222}Rn emanation measurements of the assembled XENONIT detector and gives a summary on the inferred ^{222}Rn budget, as well as a comparison to XENONIT data. We also describe radon reduction methods that were applied during detector operation. We close with a summary and outlook in Sect. 5.

2 The ^{222}Rn assay techniques

For XENONIT we used two methods to study ^{222}Rn emanation. The most sensitive method applied ultra-low background miniaturized proportional counters [14], developed for the GALLEX solar neutrino experiment [15]. These devices are made of high purity synthetic quartz with an iron cathode and a thin tungsten-wire (13 μm diameter) anode. The active volume of the counters is around 1 cm^3 and the counting gas consists of 90 % argon and 10 % methane to which the radon to be measured is added. ^{222}Rn atoms decay by α -disintegration, followed by two further α -decays from ^{218}Po and ^{214}Po until the long-lived ^{210}Pb breaks the secular equilibrium. Cosmic muons as well as environmental β - and γ -radiation cannot deposit energies above 50 keV in the miniaturized counters. In contrast, α -decays exhibit a larger energy deposition, allowing for their clear identification. The detection efficiency for the three α -decays is not equal, because the gaseous ^{222}Rn is distributed in the entire counter volume, while polonium ions deposit on surfaces. On average $(49.3 \pm 2.0)\%$ of all α -decays are detected, yielding an expectation value of (1.48 ± 0.06) counts per ^{222}Rn decay. The background count rate above 50 keV scatters around one count per day for the different proportional counters. Thus, a minimum detectable activity of $\sim 20\ \mu\text{Bq}$ can be achieved.

Prior to a measurement, the emanated ^{222}Rn atoms had to be collected from the samples, concentrated and mixed with the counting gas. For this purpose, the samples were placed in gas-tight emanation vessels made of glass or stainless steel. We ensured that the pieces of a sample were not stacked in order to let radon escape freely from all surfaces. Ambient air was removed by pumping or flushing with radon-free carrier gases (in most cases helium). Then, the vessel was filled with

the carrier gas and sealed and the emanated ^{222}Rn accumulated until the ^{222}Rn activity reached a sizable fraction of its equilibrium value. After typically one week, the mixture of carrier gas and ^{222}Rn atoms was pumped or flushed through a gas purifier to remove gaseous impurities. The ^{222}Rn atoms were trapped in an activated carbon column at liquid nitrogen temperature afterwards.

Larger samples, such as subsystems of the entire XENONIT experiment, could not be placed in emanation vessels. In these cases, the carrier gas was usually filled directly into the gas-tight system. Such samples took the role of both, the emanation vessel and the investigated ^{222}Rn source, whereas the rest of the procedure remained unchanged. Sometimes only a fraction of the filled carrier gas could be extracted due to limited pumping power. In such cases, the quoted activities were corrected for this reduced extraction efficiency, assuming that the radon was homogeneously distributed in the gas.

The concentrated ^{222}Rn sample was further processed in a sample purification system. The same system was also used to fill the proportional counters [16]. It featured several cold traps and a non-evaporable hot getter pump to remove unwanted trace-impurities that could impair the counter performance. In the final step, the gas sample was mixed with the counting gas and pushed into the counter by means a mercury column.

In some cases, measurements had to be done on samples that were previously exposed to xenon. The subsequent xenon out-gassing inhibited the use of miniaturized proportional counters due to their small volume and the difficulty to separate radon from xenon. In such cases we used electrostatic radon monitors with a significantly larger active volume [17, 18]. The positively charged ^{222}Rn daughters were collected on a silicon PIN diode biased with a negative high voltage with respect to the vessel's walls. All subsequent α -decays were recorded by the diode and the signal from ^{214}Po was evaluated as it has the highest detection efficiency (approximately 30 %). We used two different monitors with a vessel size of 1 liter and 4 liters, respectively. The background due to self-emanation of the monitor was negligible compared to the signal in all measurements. Even though the sensitivity of the radon monitors was about four times worse compared to the measurements with proportional counters, it was sufficient for our applications in XENONIT.

All results in this article are given with a combined uncertainty σ including statistical and systematical errors. If the result is compatible with zero within $1.645\ \sigma$, a 90% C.L. upper limit is given instead. Whenever a sample was additionally screened by γ -ray spectroscopy, we quote the result obtained in [10] and refer to the identifier used in that work as Radioassay-ID (RID).

Table 1 ^{222}Rn emanation measurements of grade-1 titanium samples. The normalization to surface area and, where relevant, to weld seam length is also given. For comparison the ^{226}Ra bulk activity taken from [10] is quoted and referred to as RID (Radioassay-ID) defined there

ID	Sample	Supplier	Description	Treatment	^{222}Rn emanation rate	^{226}Ra activity [10]
#1	Ti grade-1	Supra Alloys	14 plates 20.3 cm \times 20.3 cm \times 0.25 cm Total: 1.2 m ² /6.9 kg	Untreated	(120 \pm 30) μBq (101 \pm 28) $\mu\text{Bq}/\text{m}^2$	RID #40 < 0.6 mBq/kg
#1a			Same as sample #1	Etched for 2 h in 1.8% HNO_3	(160 \pm 50) μBq (140 \pm 40) $\mu\text{Bq}/\text{m}^2$	
#2	Ti grade-1	Nironit	13 plates 20 cm \times 20 cm \times 0.2 cm Total: 1.1 m ² /4.6 kg 17.4 m of TIG-weld seam	Untreated	(540 \pm 60) μBq (510 \pm 5) $\mu\text{Bq}/\text{m}^2$ (31 \pm 3) $\mu\text{Bq}/\text{m}$	RID #39 < 1.3 mBq/kg
#2a			7 plates of sample #2 Total: 0.6 m ² /2.4 kg 9.4 m of TIG-weld seam	Etched for 2 h in 0.6% HF/7.6% HNO_3 ("pickling")	(73 \pm 28) μBq (130 \pm 50) $\mu\text{Bq}/\text{m}^2$ (7.8 \pm 2.9) $\mu\text{Bq}/\text{m}$	
#3	Ti grade-1	Nironit	13 plates 20 cm \times 20 cm \times 0.2 cm Total: 1.1 m ² /4.6 kg	Untreated	(2980 \pm 200) μBq (2810 \pm 190) $\mu\text{Bq}/\text{m}^2$	RID #38 (1.1 \pm 0.4) mBq/kg
#3a			6 plates of sample #3 Total: 0.5 m ² /2.1 kg	Electropolished 30 μm surface removed	< 39 μBq < 80 $\mu\text{Bq}/\text{m}^2$	

3 Material screening

This section presents the results of samples which were measured during the preparation and construction phase of XENON1T. We also list the supplier of the samples as it was not always possible to identify the manufacturer of the raw material.

3.1 Metal samples

3.1.1 Titanium

In an early phase of the project a cryostat made of grade-1 titanium was considered for XENON1T and the ^{222}Rn emanation rate of several titanium samples was measured. The results are given in Table 1.

Sample #1 was from Supra Alloys and samples #2 and #3 were from Nironit. There were 17.4 m of TIG-welded¹ seam on the plates of sample #2 to test the influence of welding on the ^{222}Rn emanation rate. We performed surface cleaning tests for all samples. Sample #1a was treated for 2 h in a 1.8% nitric acid (HNO_3) solution. Titanium itself is not soluble in nitric acid, but the acid may remove trace impurities, in particular ^{226}Ra , from the surface. In contrast, sample #2a was pickled for 2 h in a 0.6% hydrogen fluoride (HF) and 7.6% nitric acid solution. This procedure dissolved titanium and consequently removed the upper few micrometers

of the sample. In case of sample #3a, 30 μm of the surface were removed by electro-polishing. All results are given as both, an absolute ^{222}Rn emanation rate and a rate normalized to the surface of the sample. For samples #2 and #2a we also normalize the result to the length of the weld seam. Note, however, that we could not distinguish whether ^{222}Rn came from the weld seam or from the surface. Thus, both normalizations cannot be true simultaneously and must be considered as upper limits.

Before any purification, the ^{222}Rn emanation rates of the grade-1 titanium samples exhibited large variations, which did not show up in the ^{226}Ra activity obtained from γ -ray spectroscopy (last column in Table 1), hinting at a surface contamination. The nitric acid treatment did not show any improvement (#1a). Pickling, in contrast, reduced the ^{222}Rn emanation rate. Under the assumption that the contamination was equally distributed among the 13 plates, we obtained a factor 4.0 ± 1.6 reduction by comparing the result of sample #2 to sample #2a. The most interesting sample was #3 which showed a particularly high ^{222}Rn emanation rate. It disappeared completely after electro-polishing (#3a). Evidently, a major part of the total ^{226}Ra activity of the sample was located on the surface. Thus, the true bulk activity must have been lower than reported in [10] (RID #38), since γ -ray spectroscopy cannot resolve the spatial distribution of the radio-impurities in a sample and usually assumes that all activity is in the bulk. Sample #3 and sample #3a nicely illustrate the complementarity of ^{222}Rn assay technique and γ -ray spectroscopy, and the importance to apply both methods.

¹ TIG: tungsten inert gas.

The results of the titanium ^{222}Rn screening campaign suggested that electro-polishing can suppress the ^{222}Rn emanation rate of titanium to a negligible level. However, our titanium samples showed a too high uranium bulk contamination. Therefore, the XENON1T cryostat was made of stainless steel [2].

3.1.2 Stainless steel

Stainless steel was mainly used for vessels and pipes in the XENON1T inner detector system. Parts of the TPC were also made of stainless steel, however, their surface area were small and the contribution to ^{222}Rn emanation was thus expected to be minor. Table 2 summarizes the results of all investigated stainless steel samples. Samples #4 to #9 were screened to investigate the contribution from stainless steel welds. Samples #10 and #11 were bellows which could be a potential ^{222}Rn source in stainless steel pipes. Samples #12 to #14 were related to heat exchangers and sample #15 was the packing material of the xenon distillation column [19]. As in the case of titanium, we could not distinguish whether the measured ^{222}Rn emanation rate of TIG-welded samples originated from the surface or from the weld. Thus, in Table 2 we give both normalizations in addition to the absolute ^{222}Rn emanation rate.

Sample #4 and sample #5 had a similar ^{222}Rn emanation rate, although the weld seam length of sample #4 was significantly longer. This pointed to a subdominant contribution of the welds. That result was further supported by a test of sample #6. It had no welds at all, but showed a similar ^{222}Rn emanation rate as the welded sample #5 (normalized to its surface). Subsequent electro-polishing of sample #5 reduced the measured ^{222}Rn emanation rate by a factor 3.4 ± 1.2 (#5a).

We further investigated stainless steel samples TIG-welded by Lamm's Machine Inc., the company that built the cryogenic system for XENON1T. We did not test any unwelded samples, but three welded samples with different surface treatments (#7 – #9). The results normalized to the surface area were in agreement with samples #5 and #6 from ALCA Technology S.r.l. Note that an upper limit was found for the untreated sample (#7). Hence, a further ^{222}Rn reduction by cleaning attempts was not measurable within our sensitivity. Again it was confirmed that stainless steel TIG-welds do not represent a notable additional source of ^{222}Rn , which is in tension with findings from other experiments [14,20].

The measurements of the bellows were motivated by the relatively high ^{222}Rn emanation rate of the cryogenic pipe (#52 in Table 8). This was a six-fold stainless steel pipe-in-pipe system that connected the XENON1T cryostat to the cryogenic system (see Sect. 4.1 and [2]). It contained about 10 m of stainless steel bellows in the pipes to compensate for thermal shrinkage. We tested two spare bellows from

the same supplier with 35 mm and 100 mm inner diameter, respectively (#10 and #11). Although we found a small positive signal for sample #10, the result indicated that the bellows did not constitute the main ^{222}Rn source of the cryogenic pipe.

We also tested two heat exchangers, which were used to evaporate and re-condense xenon in the purification loop. They were made of stainless steel plates brazed with a copper alloy. We measured the larger one (#12) prior to any purification and found a ^{222}Rn emanation rate of (510 ± 50) mBq. Subsequently, we cleaned it by exposing all internal surfaces to a 1.8 % nitric acid solution for about 12 h. The same treatment was done for a smaller heat exchanger of the same type, which, however, was not measured before. After the treatment both heat exchangers were measured together (#13) and yielded a result of (134 ± 24) μBq . Clearly, the treatment was effective despite the rather weak concentration of nitric acid. The heat exchangers were combined with a high purity electrical heater (#14). Its ^{222}Rn emanation rate was found to be (70 ± 27) μBq .

The last sample (#15) in Table 2 was from the XENON1T xenon distillation column for krypton removal [19]. The column was filled with structured stainless steel packings to increase the contact surface between the gaseous and liquid phase of xenon. Since the same column was used for radon removal tests [21] (see Sect. 4.3), we were interested in its ^{222}Rn emanation rate. Indeed, all 55 packings together only emanated (48 ± 20) μBq , which was an excellent result for the rather large sample surface (5.2 m^2).

3.2 Gas purification system

3.2.1 Gas purifiers

In order to maintain its ultra-high chemical purity, the xenon in XENON1T was continuously cleaned by SAES gas purifiers. Two of them were used in parallel to provide the required purification efficiency. Each purifier contained a porous, highly chemically-active zirconium-alloy in two cartridges. The larger was operated at 400°C , while the smaller one was kept at room temperature and acted as a dedicated hydrogen removal unit. Altogether, ~ 4 kg of active material was used in each purifier.

We measured the two gas purifiers of XENON1T (#16 and #17) and two smaller models (#18 and #19 with about eight times smaller active mass) which used the same alloy, but which had no hydrogen removal unit. The results are presented in Table 3. We measured the ^{222}Rn emanation rate in two different thermal conditions; at room temperature (cold state) and at operating temperature (hot state). While only the hot state was relevant for the experiment, the measured rate of the cold state could have given insight into the ^{222}Rn emanation process relevant for these porous materials. An enhanced

Table 2 ^{222}Rn emanation results of various stainless steel (SS) samples. Where relevant, the normalization to surface area and to weld seam length is also given

ID	Sample	Supplier	Description	Treatment	^{222}Rn emanation rate
#4	Plates with welds (304 or 316L)	CRIOTEC Impianti S.p.A.	5 plates w. 8.25 m TIG-weld seam 17.9 cm \times 17.6 cm \times 1 cm Total: 0.33 m ² /12.9 kg	Untreated	(190 \pm 30) μBq (560 \pm 100) $\mu\text{Bq}/\text{m}^2$ (23 \pm 4) $\mu\text{Bq}/\text{m}$
#5	Tubes with welds (316L)	ALCA Technology S.r.l.	3 tubes w. 4.8 m TIG-weld seam Outer diameter: 10.18 cm Thickness of wall: 0.34 cm Length: 30 cm Total: 0.56 m ² /7.2 kg	Untreated	(214 \pm 26) μBq (380 \pm 50) $\mu\text{Bq}/\text{m}^2$ (45 \pm 5) $\mu\text{Bq}/\text{m}$
#5a	Tubes with welds (316L)	ALCA Technology S.r.l.	Same as sample #5	Electro-polished	(47 \pm 19) μBq (80 \pm 30) $\mu\text{Bq}/\text{m}^2$ (10 \pm 4) $\mu\text{Bq}/\text{m}$
#6	Tubes (316L)	ALCA Technology S.r.l.	3 tubes Outer diameter: 10.18 cm Thickness of wall: 0.34 cm Length: 30 cm Total: 0.56 m ² /7.2 kg	Untreated	(160 \pm 50) μBq (290 \pm 90) $\mu\text{Bq}/\text{m}^2$
#7	Tube with welds (304L)	Lamm's machine Inc.	Tube with 3.6 m TIG-weld seam Outer diameter: 10.16 cm Thickness of wall: 0.2 cm Length: 34 cm Total: 0.19 m ² /1.5 kg	Untreated	$\leq 38 \mu\text{Bq}$ $\leq 200 \mu\text{Bq}/\text{m}^2$ $\leq 11 \mu\text{Bq}/\text{m}$
#8	Tube with welds (304L)	Lamm's machine Inc.	Tube with 3.6 m TIG-weld seam Outer diameter: 10.16 cm Thickness of wall: 0.2 cm Length: 34 cm Total: 0.19 m ² /1.5 kg	Electro-polished	(52 \pm 23) μBq (270 \pm 120) $\mu\text{Bq}/\text{m}^2$ (14 \pm 6) $\mu\text{Bq}/\text{m}$
#9	Tube with welds (304L)	Lamm's machine Inc.	Tube with 3.6 m TIG-weld seam Outer diameter: 10.16 cm Thickness of wall: 0.2 cm Length: 34 cm Total: 0.19 m ² /1.5 kg	Etched and electro-polished	(57 \pm 17) μBq (300 \pm 90) $\mu\text{Bq}/\text{m}^2$ (16 \pm 4) $\mu\text{Bq}/\text{m}$
#10	Stainless steel bellow	Streas S.r.l.	Bellow with one CF40 flange 1 m long, inner diameter 35 mm	Untreated	(130 \pm 40) μBq
#11	Stainless steel bellow	Streas S.r.l.	Bellow with one CF100 flange 0.4 m long, inner diameter 100 mm	Electro-polished	$\leq 160 \mu\text{Bq}$
#12	Large heat exchanger	GEA Group	Type FG5X12-60: 60 SS plates (338 mm \times 130 mm) brazed with copper alloy	Untreated	(510 \pm 50) μBq
#13	Small heat exchanger combined with large heat exchanger	GEA Group	Type FG3X8-20: 20 SS plates (226 mm \times 86 mm) brazed with copper alloy plus type FG5X12-60 (see above)	Etched for \sim 12h with 1.8% HNO_3	(134 \pm 24) μBq
#14	High-purity electrical heater with large heat transfer surface	D.A.T.E. (Developpement et Applications des Techniques de L'Energie)	Custom-designed heater complying with very high purity standards	Etched for \sim 15min with 2% HNO_3	(70 \pm 27) μBq
#15	Stainless steel packing material	Sulzer Ltd.	55 structured packings, type EX 0.095 m ² /piece, total surface: 5.2 m ²	Untreated	(48 \pm 20) μBq (9 \pm 4) $\mu\text{Bq}/\text{m}^2$

^{222}Rn release rate at elevated temperature would have given evidence for diffusion-driven emanation. The two gas purifiers used in XENON1T differed significantly in their hot state emanation rate (#16 emanated almost five times more

^{222}Rn than #17). The two smaller gas purifiers showed a significantly lower ^{222}Rn emanation rate. For sample #16, the cold state emanation was reduced by a factor 1.91 ± 0.27 in comparison with its hot state. In contrast, for sample #18

Table 3 ^{222}Rn emanation rates of four noble gas purifiers from the company SAES. PS4-MT50-R535 is identical to PS4-MT50-R2, but received a new commercial label

ID	Model	Mass of active material	Cold state activity [mBq]	Hot state activity [mBq]	Used in XENON1T
#16	MonoTorr PS4-MT50-R2	~ 4 kg	0.61 ± 0.04	1.17 ± 0.15	Yes
#17	MonoTorr PS4-MT50-R535	~ 4 kg	–	0.24 ± 0.03	Yes
#18	MonoTorr PS4-MT3-R2	~ 0.5 kg	0.09 ± 0.03	0.09 ± 0.03	No
#19	MonoTorr PS3-MT3-R2	~ 0.5 kg	–	≤ 0.015	No

Table 4 ^{222}Rn emanation measurements of xenon recirculation pumps

ID	Sample	Description	^{222}Rn emanation rate [mBq]	Used in XENON1T	Comment
#20	QDrive pump 2S132CX	Serial no. C204	5.2 ± 0.2	No	
#20a	QDrive pump 2S132CX	Serial no. C204 after repair	2.5 ± 0.1	Yes	
#21	QDrive pump 2S132CX	Serial no. C205	3.5 ± 0.2	No	
#22	QDrive pump 2S132CX	Serial no. C206	4.5 ± 0.2	Yes	
#23	Magnetically coupled piston pump [24]		0.29 ± 0.09	Yes	Replacing all QDrive pumps

no difference between the hot and cold state was observed within the measurement uncertainty.

The ^{222}Rn emanation rate of the gas purifiers is not fully understood, but the large difference for identical models suggested that it depends on the purity of the raw materials. That opens up the possibility for further ^{222}Rn reduction by material screening.

3.2.2 Recirculation pumps

XENON1T used customized QDrive piston pumps from Chart Industries for xenon gas recirculation [22]. Three pumps (#20, #21 and #22) were measured for their ^{222}Rn emanation rate. The results are summarized in Table 4. After a mechanical failure of QDrive pump C204, it was sent back to the manufacturer for repair. Afterwards, its ^{222}Rn emanation rate was lower by more than a factor two (#20a), probably due to the replacement of a dirty polyester resin (see text below and Table 6).

Since the contribution of the recirculation pumps presented a major fraction of XENON1T's total ^{222}Rn budget (see Sect. 4.2), we performed further investigations to understand the origin of the observed ^{222}Rn emanation rate. For this purpose, we screened most of the individual components of a yet unassembled QDrive pump (see Table 5). We found that the stators of the pump's electrical motor (#24) were the dominant ^{222}Rn sources, followed by the pistons (#25) and the magnet cores (#26).

To study the origin of the identified ^{222}Rn sources, we further investigated their constituent parts (see Table 6). First, we noticed that the bare magnets did not emanate a lot of ^{222}Rn

(#30). The ^{222}Rn emanation rate rather originated from the epoxy coating as can be seen from the comparison to sample #31. Note, however, that the dirty coating of sample #31 was only used for prototype pumps. In the pumps of XENON1T it was already replaced by an alternative much cleaner coating (#32), which we identified by our screening effort. The stator was a ring-shaped structure of silicon steel with four electromagnets, each formed by a copper coil which was held in place by polyester resin. The wire used for the copper coil showed no measurable ^{222}Rn emanation rate (#33), whereas a significant emanation rate was found for the silicon steel (#34) and the polyester resin (#35). The latter one was responsible for a large fraction of the stator's ^{222}Rn emanation rate. Later, we identified a cleaner alternative (#36). The new resin was applied in the repaired C204 pump. It may explain the observed ^{222}Rn reduction (#20 and #20a), if one considers that the surface to volume ratio of the resin inside the pump was larger than for the aliquots of sample #35 and #36. Moreover, the used amount of resin in a pump was hard to quantify and fluctuated among different pumps.

Motivated by the relatively large ^{222}Rn emanation rate of the QDrive pumps, we followed an approach of EXO-200 [23] to build a cleaner magnetically coupled piston pump. The new pump was developed within the XENON collaboration together with groups from the nEXO collaboration [24]. Its ^{222}Rn emanation rate was found to be (0.29 ± 0.09) mBq (#23 in Table 4), an order of magnitude lower than the results obtained for the QDrive pumps. The new pump was successfully installed for the last data taking phase of XENON1T and its impact on the ^{222}Rn budget is discussed in Sect. 4.3.

Table 5 ^{222}Rn emanation measurements of all parts of a QDrive recirculation pump prior to assembly

ID	Sample	Description	^{222}Rn emanation rate
#24	2 stators	Silicon steel frame, Cu wire winding, polyester impregnate	(2.99 ± 0.15) mBq
#25	2 pistons	Made of brass	(0.68 ± 0.06) mBq
#26	2 magnet cores	Each equipped with 8 magnets	(0.28 ± 0.11) mBq
#27	4 flexure assemblies	Silicon steel, stainless steel	(0.132 ± 0.021) mBq
#28	Polyester lacing	To fix Cu wire winding	< 0.053 mBq
#29	Remaining small items	3 other types of epoxy, screws, nuts, spacers	< 0.020 mBq

Table 6 Results of ^{222}Rn emanation measurement to identify and replace the dominant ^{222}Rn sources in the QDrive pump

#30	Bare magnets	6 pieces: 1.55 cm \times 1.88 cm \times 1.88 cm each Total surface: 112 cm ²	< 0.021 mBq < 1.9 mBq/m ²
#31	Magnets with epoxy coating	8 pieces: 1.55 cm \times 1.88 cm \times 1.88 cm each Total surface: 150 cm ²	(0.37 ± 0.04) mBq (24.8 ± 2.9) mBq/m ²
#32	Alternative epoxy coating for magnets (from Magnet Coating Engineering)	Cured on Cu substrate Surface: 618 cm ²	(0.093 ± 0.017) mBq (1.51 ± 0.28) mBq/m ²
#33	1.1 mm copper wire for magnet winding	1.68 kg	< 0.097 mBq
#34	Wingard & Co. silicon steel	6 plates (9.3 cm \times 8.9 cm \times 0.05 cm) Fulfills ASTM A677 standard Total: 167 cm ² /29.4 g	(0.240 ± 0.040) mBq (14.1 ± 2.2) mBq/m ²
#35	Dolphon CC-1105 HTC	Solventless polyester resin 396 cm ² /95.4 g	(0.70 ± 0.05) mBq (17.7 ± 1.2) mBq/m ²
#36	Elantas GRC 59-25 (alternative to Dolphon CC-1105 HTC)	Low viscosity hermetic varnish 398 cm ² /51.9 g	(0.027 ± 0.012) mBq (0.76 ± 0.30) mBq/m ²

3.3 Other samples

This section presents ^{222}Rn emanation measurements of TPC components and other samples measured for the XENON1T experiment. The complexity of the XENON1T TPC made it impossible to screen every component. Therefore, we focused on samples which either cover a large surface area inside the TPC or are known to be potential ^{222}Rn sources. We investigated the light sensors (#37 and #38), their cables and connectors (#39 to #43), the potting material for the cable feedthroughs (#44), the PTFE reflectors of the TPC (#45), copper of the field shaping rings (#46) as well as a ^{220}Rn calibration source (#47). A detailed description of their usage in XENON1T can be found in [2]. The results are summarized in Table 7.

XENON1T used Hamamatsu R11410-21 photomultiplier tubes (PMTs) [25] as light sensors (#37). Their ^{222}Rn emanation rate was measured in a helium-free environment and we used neon as carrier gas. Helium would have penetrated into the PMTs, creating an unacceptably high rate of afterpulses [26]. We also measured the ^{222}Rn emanation rate of the PMT

high voltage divider circuits (base) used to read out the signal and to supply the high voltage (#38). They consisted of a printed circuit board made of Cirlex, and several resistors and capacitors soldered onto it. It should be noted that after the measurement a different type of resistors was selected for the boards used in XENON1T. It proved to have a comparable ^{226}Ra activity, but no dedicated ^{222}Rn emanation test was performed. With the results given in Table 7 and taking into account that XENON1T used 248 PMTs, we estimated a total contribution of (1.08 ± 0.26) mBq from the PMTs and their bases. Note that the ^{226}Ra activity in the PMTs (RID #69 in [10]) was much higher, which means that less than a percent was emanated.

We also investigated the cables, which were used inside the detector. We measured the Kapton-insulated high voltage cable (#39) and three types of coaxial cable, one with Kapton insulation (#40) and two with PTFE insulation (#41 and #42). The ^{222}Rn emanation rate for all cables was below the detection limit. The ^{226}Ra activity of the cables, [10] normalized to their mass was at least ten times larger than its ^{222}Rn emanation rates for each cable sample. This indicated that the

Table 7 ^{222}Rn emanation measurements of various other samples. For comparison the ^{226}Ra bulk activity taken from [10] is quoted and referred to as RID (Radioassay-ID) defined there

ID	Sample	Supplier	Description	^{222}Rn emanation rate	^{226}Ra activity [10]
#37	R11410 PMTs	Hamamatsu	29 pieces, mixture of low performance PMTs and mechanical samples	$(58 \pm 28) \mu\text{Bq}$ $(2.0 \pm 1.0) \mu\text{Bq/PMT}$	RID #69 $(600 \pm 100) \mu\text{Bq/PMT}$
#38	PMT bases	Fralock	55 pieces, made of Cirlex with soldered resistors and capacitors	$(129 \pm 25) \mu\text{Bq}$ $(2.4 \pm 0.5) \mu\text{Bq/piece}$	RID #94 $(15 \pm 2) \mu\text{Bq/piece}$
#39	Kapton single wire cable	Accu-Glass	30 AWG solid core wire 100 m/0.076 kg for high voltage supply (used in XENON1T)	$\leq 35 \mu\text{Bq}$ $\leq 0.35 \mu\text{Bq/m}$ $\leq 460 \mu\text{Bq/kg}$	RID #57 $(4000 \pm 1000) \mu\text{Bq/kg}$
#40	Kapton coaxial cable	Accu-Glass	30 AWG, 50 Ω cable 100 m/0.55 kg for signal readout (not used in XENON1T)	$\leq 25 \mu\text{Bq}$ $\leq 0.25 \mu\text{Bq/m}$ $\leq 45 \mu\text{Bq/kg}$	
#41	PTFE coaxial cable	Huber & Suhner	RG196 78 m/0.594 kg for signal readout (not used in XENON1T)	$\leq 44 \mu\text{Bq}$ $\leq 0.56 \mu\text{Bq/m}$ $\leq 74 \mu\text{Bq/kg}$	RID #59 $(1000 \pm 300) \mu\text{Bq/kg}$
#42	PTFE coaxial cable	koax24	RG196 182 m/1.59 kg for signal readout (used in XENON1T)	$\leq 58 \mu\text{Bq}$ $\leq 0.32 \mu\text{Bq/m}$ $\leq 36 \mu\text{Bq/kg}$	RID #55 and #56 $(400 \pm 200) \mu\text{Bq/kg}$
#43	D-sub type contact pins	Accu-Glass	1200 male and female pieces made of Cu/Be and Cu/bronze	$\leq 47 \mu\text{Bq}$ $\leq 0.039 \mu\text{Bq/piece}$	
#44	Epoxy for potting	Reliable Hermetic Seals	3 discs 9.5 mm thick, 95 mm diameter 506 cm ² /191 cm ³ /0.47 kg	$\leq 51 \mu\text{Bq}$ $\leq 1000 \mu\text{Bq/m}^2$	
#45	PTFE panels	Amsler & Frey	67 pieces, total: 4.06 m ² /31.9 kg Length: 19 cm–24.5 cm Width: 13 cm–19 cm Thickness: 0.5 cm–1.6 cm	$(97 \pm 21) \mu\text{Bq}$ $(24 \pm 5) \mu\text{Bq/m}^2$ $(3.0 \pm 0.7) \mu\text{Bq/kg}$	RID #50 $< 120 \mu\text{Bq/kg}$
#46	Copper rods	Luvata	150 pieces, each 15 cm long and 2 cm diameter 1.5 m ² /7069 cm ³ /56.7 kg	$\leq 25 \mu\text{Bq}$ $\leq 17 \mu\text{Bq/m}^2$ $\leq 0.44 \mu\text{Bq/kg}$	
#47	^{220}Rn calibration source [27] ^a	PTB	Electro-deposited ^{228}Th on stainless steel disc	$\leq 46 \mu\text{Bq}$	

^aThe limit presented here differs slightly from the result published in [27] ($\leq 55 \mu\text{Bq}$) due to a data re-evaluation

^{222}Rn sources were located in the inner part of the cables. The impact of the cables on the ^{222}Rn budget is discussed in more detail in Sect. 4.1. Different high voltage cable sections were connected at the top of the TPC and just in front of the vacuum feedthroughs with D-sub type pins which showed no measurable ^{222}Rn emanation rate (#43). The cables were fed to the air-side via potted feedthroughs. The epoxy used for this potting (#44) was measured. For about half a kilogram of material, we found an upper limit of $\leq 51 \mu\text{Bq}$.

The reflective walls of the XENON1T TPC were formed by diamond-shaved PTFE panels. We measured (non diamond-shaved) leftover pieces from the fabrication of these

panels, corresponding to a surface area of $\geq 50\%$ of the PTFE surface in XENON1T. The results (see #45) indicated that PTFE was a sub-dominant ^{222}Rn source inside the TPC.

We also measured copper rods from the same batch as the TPC field-shaping rings (#46), for which we found no detectable signal. Finally, we measured the ^{222}Rn release rate of a ^{220}Rn source that was used as a calibration source for XENON1T (#48) [27]. If the source had also released some ^{222}Rn , it would have required a long waiting time after each usage until the ^{222}Rn had decayed. Our measurement showed that its ^{222}Rn emanation rate was below the detection limit and negligible with respect to the other ^{222}Rn sources.

4 XENON1T results

4.1 Measurements of subsystems

In the following, we briefly describe the XENON1T inner detector systems and report on the ^{222}Rn emanation results of either individual subsystems or of combined measurements among them. More details on the measurements themselves can be found in [18, 28]. We only assayed those subsystems that were continuously in contact with xenon during the data acquisition periods and, therefore, contributed to the ^{222}Rn budget. A schematic view of the subsystems most relevant for ^{222}Rn emanation is shown in Fig. 1.

The TPC was hosted in a double-walled, vacuum-insulated stainless steel cryostat, which was closed at the top by a dome. The dome was in turn connected to the cryogenic pipe, which also contained the cables and guided them to the electric feedthrough. Evaporated xenon which reached the cryogenic system got liquefied and was returned back to the cryostat. In a second loop, the xenon gas passed through the purification system that contained the recirculation pumps and the gas purifiers. Note that a major part of the XENON1T infrastructure will be re-used for its upgrade XENONnT, so the obtained results will be relevant for the future as well [29].

The results of the measurements are listed in Table 8. After electro-polishing its inner surface, the cryostat (#48) was measured at the fabrication site (ALCA Technology S.r.l.). From the result of sample #5a (see Table 2) we predicted a ^{222}Rn emanation rate of $(80 \pm 30) \mu\text{Bq}/\text{m}^2$ for electro-polished stainless steel under the assumption of no contribution from the weld seam. Thus, we expected $(610 \pm 230) \mu\text{Bq}$ for the 7.6 m^2 surface of the cryostat. This was about one third of what we have measured (#48). The discrepancy may come from the fact that a large vessel cannot be cleaned as easily as small-size samples.

The cable pipe contained not only the cables for all 248 XENON1T channels, but also about 200 extra channels foreseen for the upgrade to XENONnT. Altogether, there were 4.1 km of PTFE insulated coaxial PMT signal cables and 4.5 km of Kapton insulated high voltage cables in the detector. From the results of sample #39 and sample #42 presented in Sect. 3.3, we derived an upper limit of $\leq 2.9 \text{ mBq}$ for the cables alone. For the entire cable pipe we measured a positive number of $(2.7 \pm 0.2) \text{ mBq}$ (#49), which is compatible with that limit. Similar contributions to the overall ^{222}Rn budget were found to originate from the cryogenic system (#50) and from the cable feedthroughs (#51), respectively.

The cryogenic pipe (#52) consisted of a 250 mm diameter pipe enclosed in a 400 mm vacuum jacket. The 250 mm pipe itself contained the cable pipe (100 mm diameter) and five thinner pipes. The following components contributed to the ^{222}Rn emanation signal of sample #52: the inner surface of

the 250 mm pipe, the outer surface of the cable pipe, as well as the inner and outer surfaces of the included thin pipes. This means the cables and the inner surface of the cable pipe were not included in this measurement. All stainless-steel pipes were electro-polished, except for the bellows. The cryogenic pipe could not be measured separately. Instead, its ^{222}Rn emanation rate was obtained by measuring it simultaneously with the inner vessel of the cryostat. By subtracting the known result of the latter one (#48), we inferred a ^{222}Rn emanation rate of $(9.4 \pm 1.0) \text{ mBq}$ for it.

Finally, we were interested in the ^{222}Rn emanation rate of the TPC. From now on we will use the term ‘inner detector volume’ for all subsystems illustrated in Fig. 1 except the purification system. The TPC contribution could be obtained by subtracting the signal of the inner detector volume after and before TPC installation. The latter one was found to be $(18.4 \pm 1.0) \text{ mBq}$ (see Table 8) by summing up the contributions from sample #48 to #52 and adding the heat exchangers (#13) and the heater (#14). Due to the size and complexity of the inner detector volume with the TPC, only a fraction of the carrier gas could be extracted and the final result was obtained by up-scaling as described in Sect. 2. Up-scaling is only appropriate in case of a homogeneous ^{222}Rn distribution in the carrier gas. Thus, we thoroughly mixed the sample gas immediately before the extraction by adding ^{222}Rn -free carrier gas. Moreover, we extracted from various ports to ensure that locations with possibly different ^{222}Rn activity concentrations were averaged out. More details on the procedure can be found in [18]. From the obtained result of $(19.3 \pm 2.1) \text{ mBq}$ we calculated the activity of the TPC alone (#54) which turned out to be compatible with zero. The result could be converted to an upper limit of 4.4 mBq at 90% C.L. This was in agreement with the known ^{222}Rn sources of the TPC as quoted in Sect. 3.3.

4.2 Overall ^{222}Rn budget in XENON1T

The measurements of the subsystems allowed us to calculate the expected ^{222}Rn budget for XENON1T. Apart from the inner detector volume (#53), the xenon was always in contact with the xenon recirculation pumps and the gas purifiers in the purification system. The measurements of their ^{222}Rn emanation rates were presented in sect. 3.2. The two gas purifiers together emanated $(1.41 \pm 0.15) \text{ mBq}$ (see Table 3). During the main dark matter search phases, the so-called science run 0 (SR0 [30]) and science run 1 (SR1 [31]), three QDrive pumps were used: C204 (after repair, #20a), C206 (#22) and C207. Their ^{222}Rn emanation rates can be taken from Table 4 except for pump C207, which was not measured. Its signal could be estimated by taking the average of the highest (#20) and the lowest (#21) emanation rate measured for the QDrive pumps with an enlarged systematic error to cover both results within the uncertainty range. With that

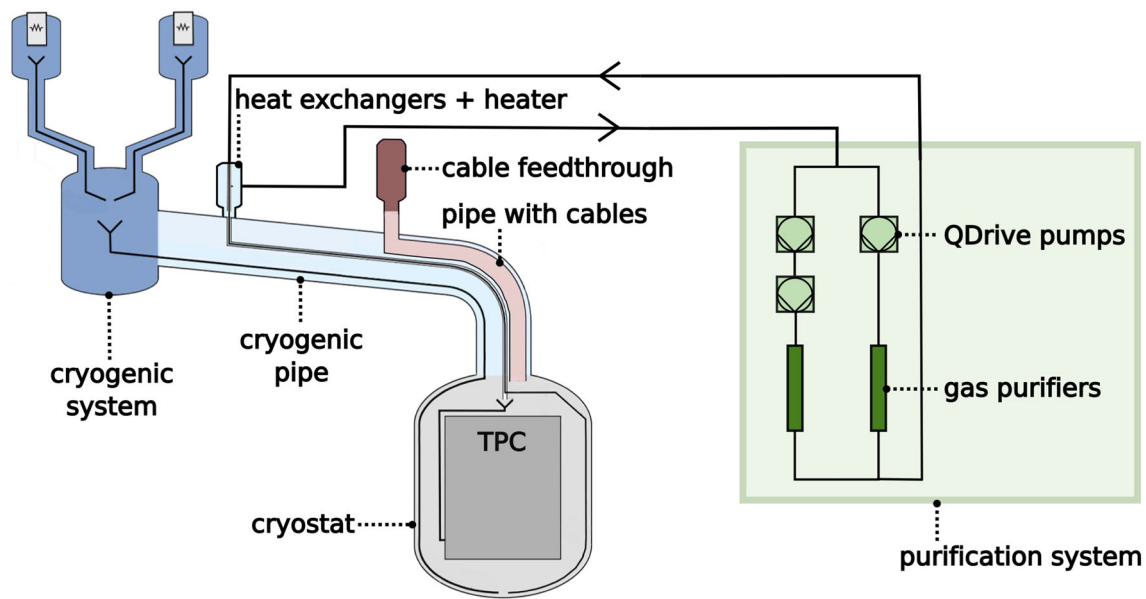


Fig. 1 Schematic setup of XENON1T (not to scale). The colors indicate different sub-components and are also used in Fig. 2, which shows their individual contributions to the overall ^{222}Rn rate

Table 8 Results from measurements of several subsystems of the XENON1T setup. The cryogenic pipe and the TPC were not measured directly. Their ^{222}Rn emanation rates were inferred indirectly by subtracting the results from two measurements, respectively

ID	Component	Activity [mBq]	Comment
#48	Cryostat (inner vessel)	1.8 ± 0.3	
#49	Pipe with cables	2.7 ± 0.2	
#50	Cryogenic system	2.4 ± 0.3	
#51	Cable feedthroughs	1.9 ± 0.2	
#52	Cryogenic pipe (without cable pipe and cables)	9.4 ± 1.0	Indirect measurement
#13 and #14	heat exchangers and heater	0.20 ± 0.04	From Table 2
	Inner detector volume without TPC (sum of all items above)	18.4 ± 1.0	
#53	Inner detector volume with TPC	19.3 ± 2.1	
#54	TPC	≤ 4.4	Indirect measurement: difference between Inner detector volume with and without TPC

assumption, we obtained (4.4 ± 0.9) mBq for pump C207. Thus, the estimated ^{222}Rn emanation rate of all three pumps together was (11.3 ± 0.9) mBq.

Summing up all components, we obtained a total ^{222}Rn budget for XENON1T of (32.0 ± 2.3) mBq. Under the assumption of a homogeneous radon distribution, we expected a ^{222}Rn activity concentration of (10.0 ± 0.7) $\mu\text{Bq/kg}$ with a total xenon mass of 3.2 t in XENON1T meeting the design goal. The pie chart in Fig. 2 presents the relative contribution of all components. The dominant elements were the QDrive pumps and the cryogenic pipe which together accounted for about two-thirds of the entire ^{222}Rn budget.

α -decays of ^{222}Rn (5.5 MeV) and ^{218}Po (6.0 MeV) in the TPC could be selected by their reconstructed energy [32–

34]. The $\sim 1\%$ energy resolution of the detector at these high energies was sufficient to allow for a clear separation of the two α -peaks. Other background sources in that energy range were subdominant and could be ignored. Therefore, the α -analysis represented a reliable way to measure the ^{222}Rn and ^{218}Po activity concentrations in the detector. In Fig. 3, we show the α -rate evolution during the science runs. An average activity concentrations of (13.3 ± 0.5) $\mu\text{Bq/kg}$ and (12.7 ± 0.5) $\mu\text{Bq/kg}$ for ^{222}Rn and ^{218}Po , respectively, was found for SR0 and SR1, excluding the distillation period for radon removal discussed in Sect. 4.3. ^{218}Po is often positively charged after its creation [35]. Thus, ion drift as well as convective motion may transport ^{218}Po out of the analysis volume and cause its deposition on TPC surfaces. This may

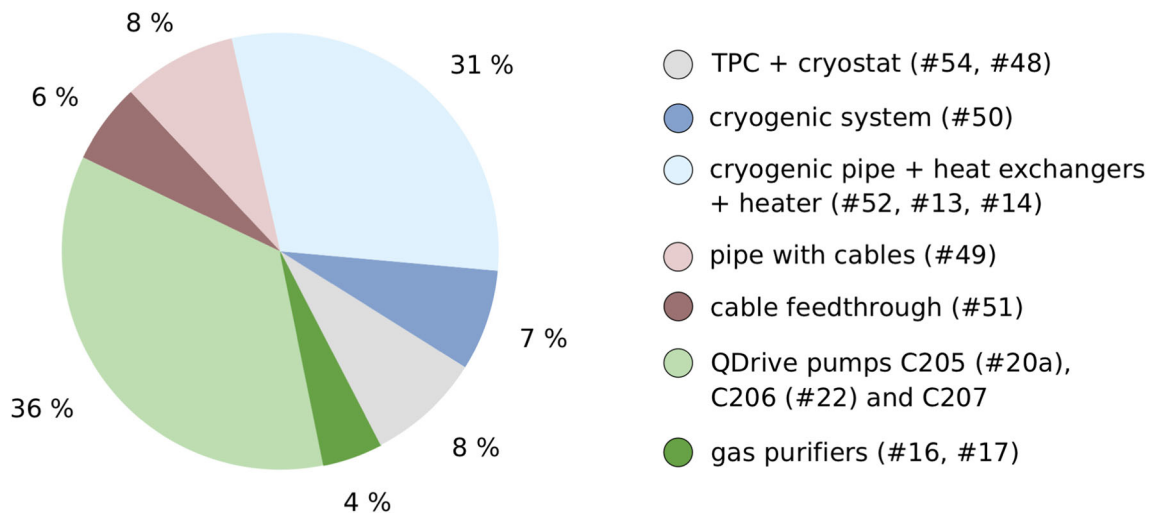


Fig. 2 The different sub-system contributions to the overall ^{222}Rn emanation rate in XENON1T. The colors correspond to those used in Fig. 1. The numbers in the brackets refer to the item numbers. QDrive pump C207 was not measured. Its ^{222}Rn emanation rate was estimated (see text)

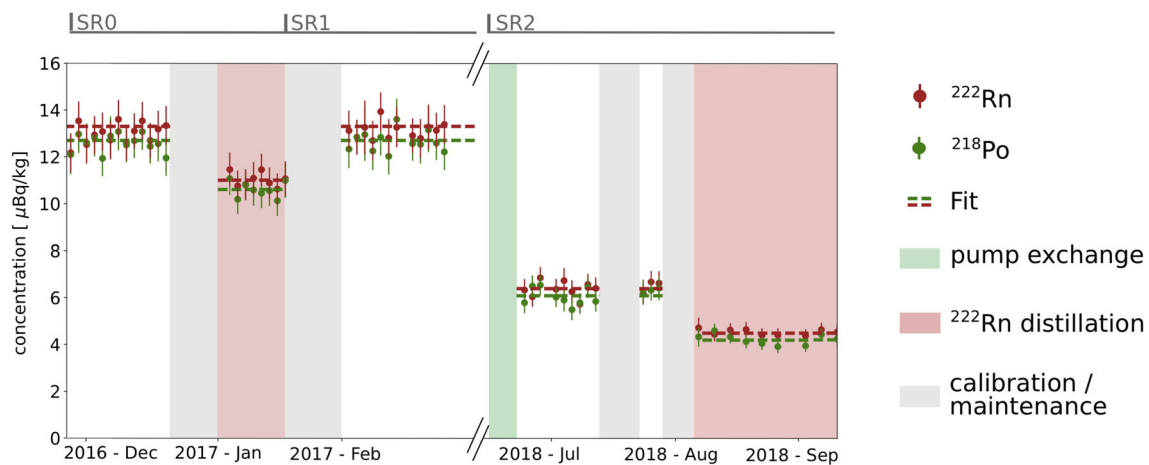


Fig. 3 The activity concentration evolution of ^{222}Rn and ^{218}Po during XENON1T data-taking. In science run 0 and 1 (SR0 and SR1) the activity concentrations were stable over the entire time and we only show the initial period here. Xenon distillation campaigns to remove ^{222}Rn

as well as the exchange of the recirculation pumps lead to a reduction of the ^{222}Rn and ^{218}Po activity concentration. They gray regions indicate periods of detector calibration or maintenance

explain the slightly lower observed ^{218}Po activity concentration with respect to ^{222}Rn .

The discrepancy between the $(13.3 \pm 0.5) \mu\text{Bq/kg}$ deduced from α -analysis and the expectation of $(10.0 \pm 0.7) \mu\text{Bq/kg}$ from emanation measurements corresponds to $(10.6 \pm 2.8) \text{mBq}$ inside 3.2 t of liquid xenon. A possible explanation for the discrepancy was the ^{222}Rn release from the QDrive recirculation pumps. As most samples in this work, they were measured at room temperature. However, during operation, they heated up and the diffusion-driven ^{222}Rn emanation could have been enhanced at elevated temperature. In addition, it could have been that the unmeasured QDrive pump C207 emanated more than expected.

4.3 Reduction of ^{222}Rn

There are several possibilities to further reduce the ^{222}Rn budget. The best option is to remove ^{226}Ra , the mother nucleus of ^{222}Rn , from the experiment. This was achieved by replacing the QDrive pumps in science run 2 (SR2). They were exchanged for the magnetically coupled piston pump described in Sect. 3.2.2 [24]. The decrease of the ^{222}Rn activity concentration in liquid xenon before and after the pump exchange is shown in Fig. 3. It corresponded to an absolute reduction of $(19.2 \pm 1.0) \text{mBq}$ in 3.2 t of xenon. From the room temperature emanation measurements, presented in Table 4, one would have expected a reduction

of only (11.0 ± 0.9) mBq.² The observed difference of (8.2 ± 1.3) mBq agreed within the uncertainty with the difference between ²²²Rn emanation measurements and α -analysis (see Sect. 4.2). This supported the hypothesis that the ²²²Rn emanation rate of the QDrive pumps was larger than estimated. By exchanging the recirculation pumps, the largest ²²²Rn source was successfully removed from the experiment yielding a reduced background level [36]. This is very promising for XENONnT, where magnetically coupled piston pumps will be employed.

Another possibility for ²²²Rn reduction is an online radon removal system in the xenon purification loop, which separates xenon from radon and retains the latter until its disintegration. Such a radon removal system based on cryogenic distillation was pioneered by the XENON collaboration [21, 37]. In XENON1T, we realized radon removal by employing the cryogenic distillation system built for krypton removal from xenon and operating it in reverse direction [38]. As shown in Fig. 3, the distillation led to a ²²²Rn reduction of $\sim 20\%$ during SR0, although only a small fraction of the recirculation flow was distilled. A compatible absolute ²²²Rn reduction was achieved in a second xenon distillation run performed in SR2, after the pump exchange. The finally accomplished ²²²Rn activity concentration in XENON1T was (4.5 ± 0.1) μ Bq/kg.

As a consequence of the promising results, the XENON collaboration realized two new purification systems for XENONnT which complement the existing ones and significantly improve the xenon purity. The first one is a novel liquid purification system which is able to produce and maintain ultra-pure liquid xenon at a very fast flow rate. The second one is a dedicated distillation column which was designed [38] and built for radon removal and which takes advantage of the large flow rate enabled by the liquid purification system.

The impact of the new radon removal system on the experiment's background can be further maximized if ²²²Rn is flushed out of the detector before it enters the TPC. A detailed mapping of the ²²²Rn sources in XENON1T was obtained by the ²²²Rn emanation measurements of the various sub-systems, presented in this work. Thus, a targeted xenon flow pattern optimization could be studied. Such a flow pattern optimization with respect to the radon removal system will be applied in the XENONnT experiment [29].

5 Summary and outlook

The background rate of current xenon dark matter detectors is dominated by ²²²Rn-induced events and it is expected that

² The sum of the signals of the three replaced QDrive pumps minus the signal of the newly mounted magnetically coupled piston pump.

²²²Rn daughters will remain an essential background component in future experiments. Therefore, ²²²Rn emanation measurements become increasingly important and provide complementary information to the bulk radioactivity screening efforts. In this article, we presented the results of a comprehensive material screening campaign for ²²²Rn emanation carried out for the XENON1T experiment, using state-of-the-art counting techniques. We selected construction materials with the lowest possible ²²²Rn emanation rate and we were able to identify and locate the remaining ²²²Rn sources in the experiment.

The predicted activity concentration from these measurements was in agreement with the target ²²²Rn activity concentration of 10 μ Bq/kg in 3.2 t of xenon [9]. Results from an α -analysis of the XENON1T data were about 30% higher than the prediction. The discrepancy could be understood by an underestimation of the recirculation pump's ²²²Rn emanation rate. With the exact knowledge of the distribution of ²²²Rn sources in XENON1T, it became possible to selectively eliminate problematic items. The ultimately measured ²²²Rn activity concentration of (4.5 ± 0.1) μ Bq/kg is the lowest ever achieved in a xenon dark matter experiment. Significant improvements are possible in XENONnT and further projects, for instance, by continuous xenon distillation.

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