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Renner, J.; Gehman, VM.; Goldschmidt, A.; Matis, HS.; Miller, T.; Nakajima, Y.; Nygren, D.... (2015). Ionization and scintillation of nuclear recoils in gaseous xenon. Nuclear Instruments and Methods in Physics Research Section A: Accelerators, Spectrometers, Detectors and Associated Equipment. 793:62-74. doi:10.1016/j.nima.2015.04.057.



The final publication is available at http://dx.doi.org/10.1016/j.nima.2015.04.057

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Additional Information

Ionization and scintillation of nuclear recoils in gaseous xenon

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Ionization and scintillation produced by nuclear recoils in gaseous xenon at approximately 14 bar have been simultaneously observed in an electroluminescent time projection chamber. Neutrons from radioisotope α -Be neutron sources were used to induce xenon nuclear recoils, and the observed recoil spectra were compared to a detailed Monte Carlo employing estimated ionization and scintillation yields for nuclear recoils. The ability to discriminate between electronic and nuclear recoils using the ratio of ionization to primary scintillation is demonstrated. These results encourage further investigation on the use of xenon in the gas phase as a detector medium in dark matter direct detection experiments.

I. INTRODUCTION

Xenon has been the detection medium of choice in multiple experiments searching for rare physics events due to its favorable properties as a detection medium [1, 2] including the availability of two channels of energy measurement, scintillation and ionization, that can be accessed simultaneously in a single detector. In par-

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ticular, recent experiments have employed liquid xenon in searching for interactions of WIMP (Weakly Interacting Massive Particle) dark matter [3–5], and neutrinoless double-beta $(0\nu\beta\beta)$ decay [6]. Both of these processes have strong implications in fundamental physics. WIMPs are strong candidates to be a possible constituent of cold dark matter (see for example [7]), thought to make up the majority of matter in the universe. The observation of $0\nu\beta\beta$ decay (see for example [8]) would establish the Majorana nature of the neutrino and provide information on the absolute value of the neutrino masses and the neutrino mass hierarchy.

WIMPs interact via the electroweak force, allowing them to scatter off nuclei, and so the signature of a WIMP in a pure xenon detector would be the recoil of a xenon nucleus, in which the energetic nucleus excites and ionizes xenon atoms to produce primary scintillation photons and electron-ion pairs. Nuclear recoils have been observed and well-characterized in liquid xenon. In particular, it is known that the scintillation and ionization yields of nuclear recoils are lower, or quenched, relative to those of energetic electrons (electronic recoils) of the same kinetic energy. A model that predicts these yields based on the existing measurements has been constructed in [9]. Liquid xenon has also clearly shown that the amount of quenching in both scintillation and ionization is not the same, enabling one to discriminate between electronic recoils and the nuclear recoil signals of interest to dark matter detection by using the ratio of ionization to primary scintillation (see for example [10– [12]).

The use of xenon in the gas phase may provide several advantages that would imply greater sensitivity in searches for dark matter and $0\nu\beta\beta$ decay. In particular, the gas phase offers improved energy resolution [13] over the liquid phase, largely believed to be due to the observed significant fluctuation in energy deposition between the ionization and scintillation channels [14] in liquid Xe. Though this can be corrected by combining both channels to recover some of the lost energy resolution in the liquid phase, as is done in [6], the inability to achieve light collection efficiencies beyond $\sim 20\%$ (the LUX experiment [3] achieved an average detection efficiency of 14% for primary scintillation) limits overall resolution via Poisson fluctuations in the detection of a relatively small signal. Better energy resolution could lead to improved electron/nuclear recoil discrimination. Under the right conditions and possibly with the addition of a molecular additive to the pure xenon gas, the amount of electronion recombination in nuclear recoil tracks may depend on the orientation of the drift electric field, thereby providing information about the direction of a WIMP scatter [15, 16].

Here we report data on the ionization and scintillation of nuclear recoils in gaseous xenon. Further details of this study can be found in [17] and [18]. In addition, scintillation and ionization of nuclear recoils were previously presented in [19]. The experiment was performed with a high pressure xenon gas time projection chamber (TPC) constructed as a prototype for the NEXT (Neutrino Experiment with a Xenon TPC) experiment, the NEXT prototype for research and development towards detection of neutrinoless Double Beta and Dark Matter (NEXT-DBDM). NEXT intends to search for $0\nu\beta\beta$ decay with an electroluminescent TPC containing 100 kg of enriched (91% $^{136}\mathrm{Xe}$) xenon. Should potential advantages be found in using gaseous xenon to search for dark matter, one could comtemplate a simultaneous $0\nu\beta\beta$ and dark matter search with a ton-scale gaseous xenon detector. A clear understanding of nuclear recoils in gaseous xenon is a critical first step in this direction.

II. EXPERIMENTAL SETUP AND CALIBRATION

A. Detector Hardware and Operation

The NEXT-DBDM detector is described in detail in [20]. Here we summarize this description and describe the modifications made for this study. Figure 1 gives an overview of the experimental setup and source locations.

The main hardware of the TPC consists of a stainless steel cylindrical pressure vessel (20 cm diameter, 33.5 cm length) with one end closed in an ellipsoidal shape and the other sealed via a ConFlat flange to a stainless steel lid to which the internal components forming the TPC are attached. The internal hardware consists of a hexagonal field cage separated into a drift (active) region of length 8 cm and an amplification region of length 5 mm by grids of wire mesh stretched tightly across metal The active region is enclosed by teflon panels with copper strips attached to their outer surfaces which are connected via resistors to grade the drift field. The panels are supported by thin plastic frames, and the teflon surfaces facing the active region were coated with tetraphenyl butadiene (TPB) by dissolving the TPB in toluene and spraying it directly onto the surface using an airbrush. An array of 19, 1-in. diameter Hamamatsu photomultiplier tubes (PMTs) arranged in a hexagonal pattern is located at the end opposite to the amplification region. High voltages for the wire meshes are passed into the pressure vessel through the lid via commercial feedthroughs rated to 20 kV at 17 bar and connected via teflon-coated wire to the mesh frames. The lid is connected by a long tube to a stainless steel octagon with 8 ConFlat ports, several of which are occupied by multi-pin feedthroughs through which PMT high voltages are input to the interior of the detector and through which the PMT signals are output. An opening of diameter 1.7 cm extends through the center of the octagon and down the tube to a 2 mm source entrance window to the interior of the pressure vessel. An external sodium iodide (NaI) scintillator coupled to a PMT was often used to detect gamma rays emitted in coincidence with the neutrons or gamma rays of interest.

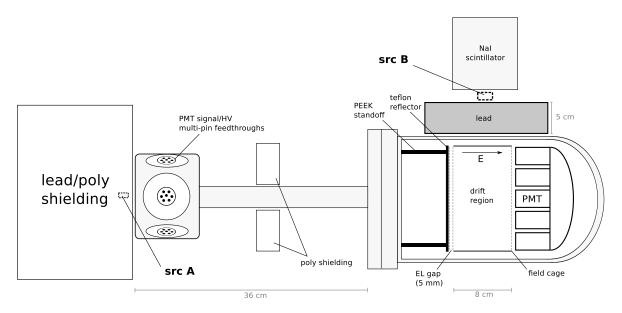


FIG. 1. Schematic of experimental configuration (not drawn to scale) for datasets used in this study. When the source was at position A, both the NaI scintillator and the source were enclosed in the lead/poly shielding.

The pressure vessel is connected to a gas system allowing for full system pump-down to pressures on the order of 5×10^{-5} Torr. The gas system also permitted reclamation/reintroduction of the xenon gas used in operation to/from a steel cylinder and constant recirculation of the xenon gas during operation through a heated zirconium-based getter, to remove impurities such as O_2 , H_2O , and N_2 . Typically after servicing the TPC, the entire system is pumped to $\sim 5 \times 10^{-5}$ torr, flushed with Ar gas, pumped a second time, and filled with xenon gas. The gas must be purified for several days before optimal electron lifetimes (of order 10s of milliseconds) are realized.

A typical event in the detector occurs when an energetic particle interacts in the xenon medium, producing primary scintillation in the form of $\lambda \sim 170$ nm photons, and ionization [1]. The primary scintillation is detected immediately by the PMTs and constitutes the signal denoted as S1. The ionization is drifted via an electric field to the amplification region, a narrow region of high field in which the electrons are accelerated to energies sufficient to repeatedly excite but not ionize the xenon atoms in the medium. Each excitation results in the emission of a xenon UV photon so that each individual electron traversing the high field region produces a number of photons g equal to [21, 22]

$$g = 140(E/p - 0.83)p\Delta x, (1)$$

where Δx is the thickness in cm of the region of high field E, given in kV/cm, and p is the gas pressure in bar. This process, called electroluminescence, provides a means of amplification with relatively lower fluctuations than amplification based on electron avalanches and results in a light signal denoted as S2 that is proportional to the number of ionization electrons produced in the event.

The TPB on the walls surrounding the active region shifts a significant fraction of the xenon scintillation light to the visible regime ($\lambda \sim 430$ nm) at which the quantum efficiency of the PMTs is higher. Relative to results obtained before the TPB was applied to the walls of the field cage, the TPB provided about a factor of 2 increase in light collection efficiency and was essential to observing the low S1 signals produced by nuclear recoils discussed in Sec. III.

All PMT signals are sampled continuously using a Struck SIS3302 digitizer and buffered in waveforms of 16384 samples that are stored in digitizer memory when a trigger constructed from a network of NIM modules is activated. The events are read out to a desktop computer in blocks of 512 and processed using an automated data management and analysis system based on ROOT [23] and FMWK [24]. In the analysis, the signals from the 19 PMTs are baseline subtracted and summed, and peaks corresponding to PMT pulses are located and integrated to give a number of detected photons (see Fig. 4 for an example waveform). Each peak is considered as a candidate S1 or S2 pulse based on its width and, when the location of S1 in the waveform was fixed due to tagging with an external scintillator, its location. Once a single S1 pulse and one or more S2 pulses have been identified, the drift time of the ionization produced in the event can be determined as the difference in time between the beginning of the S1 pulse and the centroid in time of the S2 pulse. One or more S2 peaks are possible in a single event if the ionization track has multiple components that arrive at the amplification region at distinct times. This could occur, for example, if a xenon fluorescent x-ray were produced and traveled several cm before making a distinct ionization track of its own. Key quantities such as the integrated numbers of photons in the S1 and S2 signals,

the drift time, and pulse integration and timing information from the external NaI scintillator are all stored in a ROOT tree recorded on an event-by-event basis at the end of the analysis to facilitate access.

B. Sources and Configuration

The data discussed in this study were acquired using radiation emitted from four different radioactive sources - two producing only gamma radiation and two radioisotope neutron sources producing energetic ($\sim 1\text{-}10~\text{MeV}$) neutrons. The neutron-producing $^9\text{Be}(\alpha, n)^{12}\text{C}$ reaction that occurs within the sources often leads to the emission of a high-energy gamma ray in coincidence with the emitted neutron. This gamma ray has an energy of either 4.4 MeV or 7.7 MeV. To more efficiently isolate neutron-induced events in this study, the detection of a 4.4 MeV gamma ray using a NaI scintillator was included as one of the trigger conditions when data was taken with a neutron source. Further details on the neutron production mechanisms and spectra of the radioisotope neutron sources are discussed in detail in the Appendix.

The configurations of the sources used in this study are itemized in the following subsections. Of the 19 PMTs in the energy plane, 18 were operational during the acquisition of all datasets, and all datasets were taken with a drift field of $370~\rm V/cm$ at a gas pressure of approximately 14 bar.

1. 241 Am/Be Neutron Source

An americium-beryllium neutron source containing a mass of ²⁴¹Am with approximately 56 mCi of activity was positioned in source location A as shown in Fig. 1, just in front of the long tube connecting to the pressure vessel lid. The source was surrounded by a layer of lead about 2 in. thick and a layer of polyethylene also about 2 in. thick, such that it was only exposed in the direction along the tube. Additional polyethylene shielding was placed along the tube to collimate the incident neutron flux to the cross-sectional area of the tube. The NaI scintillator was placed within the shielding nearby the source to tag 4.4 MeV gamma rays emitted from it. In this configuration, neutrons were emitted far enough from the xenon volume that their interactions could be distinguished from those induced by gamma rays. This was done by using the time-of-flight measured as the time difference in the arrival of the S1 produced in the TPC and the gamma ray tagged in the NaI scintillator.

2. ²³⁸Pu/Be Neutron Source

A plutonium-beryllium neutron source containing a mass of 238 Pu with approximately 10 mCi of activity was positioned at the side of the pressure vessel between the

NaI scintillator and a lead brick at source location B of Fig. 1. The presence of the lead greatly reduced the background due to gamma rays emitted as products of the neutron-generating (α, n) reaction.

3. ²²Na Gamma Source

A sodium source containing a mass of 22 Na with approximately 10 μ Ci of activity was placed at the side of the pressure vessel, similar to the configuration shown for source B in Fig. 1, though no lead block was present, the source was positioned about 16 cm from the side of the pressure vessel, and the NaI scintillator was positioned several centimeters away from the source to avoid pileup. Two collinear 511 keV gamma rays are emitted from the source. One gamma ray was tagged with the NaI scintillator while the other was incident on the xenon volume.

4. 137 Cs Gamma Source

A cesium source containing a mass of ¹³⁷Cs with approximately 1 mCi of activity was contained in a lead enclosure to which a cylindrical lead collimator with an opening of diameter 3.5 mm was fitted at one end. The resulting collimated beam of 662 keV gamma rays was placed at source location A as shown in Fig. 1 and pointed down the tube connecting the octagon and pressure vessel through the 2 mm thick source entrance window.

C. Detector Calibration with a ¹³⁷Cs Source

Before discussing the characterization of nuclear recoils, we establish a gamma-based calibration of the detector using a $^{137}\mathrm{Cs}$ source (configuration 4 of Sec. II B). Gamma rays of energy 662 keV were directed axially through the center of the TPC lid, and the S1 and S2 signals were examined to determine key xenon properties such as the amount of energy required to produce a primary scintillation photon W_{sc} , the relation used to correct for the z-dependence (lower detection probability of photons produced farther from the PMT array) of S1, and the light collection efficiency ϵ at the EL plane. We first assume a value for the amount of energy required to produce an ionization electron $W_{\mathrm{i}}=24.7\pm1.1~\mathrm{eV}$ (see Sec. 3.2 of [21]).

Events corresponding to full-energy depositions of 662 keV gamma rays were isolated as a peak in the S2 distribution produced by the source. A central fiducial cut was made according to the weighted average (x, y) location of the event determined using the distribution of electroluminescent light produced on the PMT plane, and the events were corrected for electron attachment by multiplying by a z-dependent exponential factor corresponding

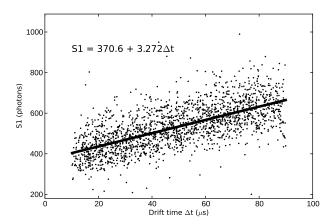


FIG. 2. S1 vs. drift time (effectively the z-coordinate) for events determined to fall in the 662 keV peak in a 137 Cs calibration run. The linear fit was used to determine a correction factor to compute the S1 value for each event as if it occurred at the EL plane (z = 0).

to an electron lifetime of $\tau_e \approx 8.3$ ms. The ¹³⁷Cs S2 photopeak was found to lie at $S_2 = 561617 \pm 92$ photons. Operating at EL gain $g = 734 \pm 89$ photons/e⁻, we calculate for the light collection efficiency at the EL plane,

$$\epsilon = \frac{S_2 \cdot W_i}{g \cdot E_{\gamma}} = 0.0285 \pm 0.0037,$$
 (2)

where $E_{\gamma} = 661657 \pm 3 \text{ eV}$ [25] is the energy of the gamma ray. The geometrical z-dependence of S1 detection efficiency was determined by plotting the integrated S1 values of the events in the ¹³⁷Cs S2 photopeak vs. event drift time (see Fig. 2). A linear dependence is observed,

$$S_1 = S_{1,0} + k\Delta t,\tag{3}$$

where Δt is the drift time and $S_{1,0}$ is a constant corresponding to the number of S1 photons detected at the EL plane for the 662 keV energy deposition. It was determined that $S_{1,0}=370.6\pm4.0$ photons and $k=3.272\pm0.076$ photons/ μ s, so that the S1 signal detected for an event with drift time Δt , denoted as $S_1(\Delta t)$, could be corrected for its z-dependence as

$$S_1' \equiv S_{1,0} \cdot \frac{S_1(\Delta t)}{S_{1,0} + k\Delta t} = \frac{S_1(\Delta t)}{1 + (k/S_{1,0})\Delta t}.$$
 (4)

Knowing how to correct the S1 signal to its value at the EL plane, and knowing the light collection efficiency at the EL plane, we can compute the energy required to produce an S1 photon $W_{\rm sc} = (E_{\gamma}/S_1)\epsilon'$, where $\epsilon' = \epsilon/\Omega$ and Ω is a factor accounting for the optical effect of the EL wire mesh grids. This correction is necessary because ϵ is calculated using S2 light produced in the EL gap between two grids while the S1 light is produced

in the active region. The factor Ω was determined by a Monte Carlo simulation in which photons were generated within the EL gap and just in front of the EL gap, and the resulting efficiencies in the two cases were compared. The grids in the simulation were given the nominal transparency of the physical mesh grids, equal to 88% at 0° incidence angle, and from the Monte Carlo, the relative light collection efficiency was determined to be described by $\Omega=0.83\pm0.08$, assuming 10% errors. Using Eq. 2, we find

$$W_{\rm sc} = \frac{S_2 \cdot W_{\rm i}}{S_1 \cdot g \cdot \Omega} = 61.4 \pm 18.0 \text{ eV},$$
 (5)

with an applied drift field of 370 V/cm, and note we have assumed an additional systematic error of 15 eV obtained from Monte Carlo studies (see Sec. III D). Because the Monte Carlo study attempted to construct a consistent picture from electronic recoil yields, EL gain, and optical/geometrical effects in the TPC, this additional error could originate from discrepancies in any of these areas. The obtained $W_{\rm sc}$ is lower than those obtained in other references including $W_{\rm sc}=76\pm12~{\rm eV}$ [26] and $W_{\rm sc}=111\pm16~{\rm eV}$ [27], though these experiments were carried out under significantly different operating conditions and in one case [26] in a mixture of 90% Xe / 10% He gas.

We also note that both S1 and S2 pulses appeared to possess long tails, a property which was not present during previous operation before TPB was placed on the walls of the field cage. TPB itself is not expected to produce such an effect, and the present effect may be due to an agent present in the toluene-based solution employed in the coating process. While the S2 light is produced over a timespan of several microseconds to tens of microseconds, the S1 light arrives over a short timescale, allowing for the characterization of the tail accompanying a single, fast pulse of light. The S1 pulses were found to be well described by a two component exponential, one with a short decay time constant of $\tau_s \approx 100$ ns, and one with a longer decay constant of $\tau_l \approx 1.4\,\mu s$.

III. NUCLEAR RECOILS IN HIGH PRESSURE XENON GAS

A. Spectrum of Emitted Neutrons

In this section, we present data taken with the radioisotope neutron sources, and we first show the expected spectrum of emitted neutrons from such sources. The neutron spectrum from a ²⁴¹Am/Be source is shown in Fig. 3 and was calculated assuming the source consisted of a volume filled with beryllium throughout which the isotope ²⁴¹Am was uniformly distributed. Further details and a description of the relevant calculations can be found in the Appendix.

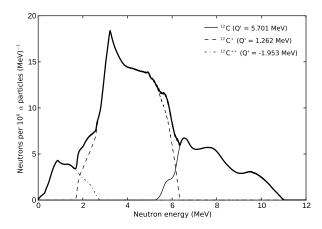


FIG. 3. Calculated neutron emission spectrum for a 241 Am/Be neutron source considering only the neutrons produced in the 9 Be(α , n) 12 C reaction. The spectrum is divided into three components for which the resulting carbon nucleus is found in the ground (solid line), first excited (dashed line), or second excited (dot-dashed line) state. The bold solid line shows the sum of all three components. The neutrons of greatest interest to this study are those for which the carbon nucleus is left in the first excited state and decays emitting a 4.4 MeV gamma ray.

B. Analysis of Experimental Data

The neutrons emitted from the radioisotope sources used may be accompanied by gamma rays, and often the neutrons themselves scatter inelastically off nuclei in the environment (in the xenon or surrounding detector hardware), resulting in the emission of various gamma rays in the de-excitation of the nuclei. Therefore, careful analysis is necessary to isolate the nuclear recoil events, and to do so, we make a sequence of analysis cuts on the data to remove gamma-induced events and finally identify a band of events at low values of S1 and S2 that consists primarily of nuclear recoils.

The data presented here, consisting of 3682304 events in total, was acquired over about 577 hours, during which time the acquisition system was actively recording events at a rate of about 3 Hz for about 346 hours. The rate of acquisition was stable throughout the majority of the run, and 6 blocks of 512 events were removed due to anomalous rates. Most of the downtime was due to the readout and storage of events from the digitizer. Events occurring within 5 minutes of occasional high voltage breakdown within the detector were discarded. This eliminated about 3.8% of the events, thus slightly lowering the effective live-time.

The analysis cuts were performed in a sequence consisting of: tagging and single-pulse event identification, time-of-flight, diffusion, and radial cuts.

A typical tagged neutron candidate event is shown in Fig. 4. The S1 signal and NaI pulses were fit to a function of the form

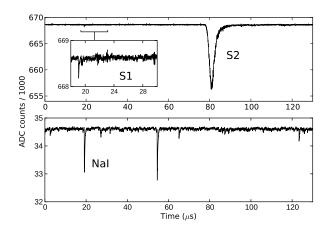


FIG. 4. A typical candidate neutron event. Note the S1 in coincidence with a pulse in the PMT coupled to the NaI scintillator, followed by the single, Gaussian-like S2 pulse indicative of a pointlike energy distribution in the TPC. The slight bulge in the right tail of the S2 pulse is due to extended light emission from the presence of the TPB coating on the teflon walls of the field cage.

$$f(x) = p_0 + \frac{p_1 e^{-(x-p_2)/p_4}}{1 + e^{-(x-p_2)/p_3}},$$
 (6)

where x is the sample value and the p_i are fit parameters. The value of p_2 for each type of pulse was taken to be its initial arrival time. An event was required to contain at least one pulse in the NaI scintillator and an S1 pulse, both with arrival times in a sample region near the expected S1 arrival time defined by the trigger. The integrated area of an NaI pulse was required to lie within a broad region chosen to correspond to the spectrum of a 4.4 MeV gamma deposition. If multiple NaI pulses met the required criteria on arrival time and charge, the NaI pulse closest in start time to that of the S1 signal was selected. The event was also required to contain a single S2 pulse, that is, one integrated pulse as defined by the pulse-finding and integration algorithm with charge q_2 nearly equal to the total integrated S2 charge Q_2 , more specifically $0.95q_2 < Q_2 < 1.05q_2$. An event for which all of these conditions are met is considered to have passed the tagging and single-pulse identification cuts.

The time-of-flight τ for a given event is determined for properly tagged events as the difference between the arrival times of the S1 and NaI pulses and is given by

$$\tau = \left[\frac{1 + E_0/K}{\sqrt{1 + 2E_0/K}}\right] (d/c) + \Delta t_{\text{off}} \tag{7}$$

where E_0 and K are the rest and kinetic energies of the particle, $d \approx 50$ cm is the distance traveled by the particle from the source to the point of detection, c is the speed of light, and $\Delta t_{\rm off}$ is an offset time due to the

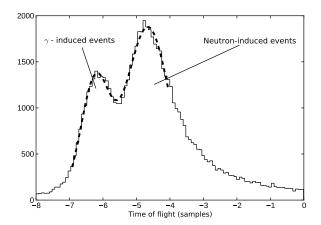


FIG. 5. Time-of-flight distribution (1 sample = 10 ns) for properly tagged events with S2 > 175000 photons and 30 photons < S1 < 500 photons (corrected for z-dependence). The left peak in the distribution is due to gammas originating from the source, and the right peak is due to events induced by neutrons from the source. The fit to a sum of two Gaussians is shown for which $\mu_1 = -6.24$, $\sigma_1 = 0.429$ (left) and $\mu_2 = -4.66$, $\sigma_2 = 0.647$ (right).

trigger and electronics. The tagging procedure serves to eliminate a significant amount of gamma background, and a clear peak in time-of-flight due to events induced by gammas from the source is not evident in the overall time-of-flight distribution. However, by examining only higher energy depositions consisting of a relatively higher fraction of gamma-induced events, two peaks are apparent in the time-of-flight distribution. Figure 5 shows the relevant regime of the time-of-flight distribution for high-S2 events, in which the leftmost peak corresponds to events produced by gammas emitted by the source. The fit shown to a sum of two Gaussians gives for the component corresponding to the left peak, a mean of $\mu_{\gamma} \equiv \mu_{1} = -6.24$ and sigma $\sigma_{\gamma} \equiv \sigma_{1} = 0.429$. We can then solve using Eq. 7 with $E_0 = 0$, $c \approx 30$ cm/ns and $\tau = \mu_{\gamma}$ for $\Delta t_{\rm off} = \mu_{\gamma} - d/c \approx -64$ ns. Since for an active region of length 8 cm all gammas emitted from the source should arrive within a time interval of 8 cm / c = 0.27 ns, which is significantly less than $\sigma_{\gamma} = 4.29$ ns, we can use σ_{γ} as a measure of the time-of-flight resolution, noting that for nuclear recoil events we expect the resolution to be poorer due to their lower S1 signals. These S1 signals may consist of only several photons arriving within a time interval relatively large compared to the duration of a single photoelectron signal in a PMT, thereby producing inconsistencies in the fit to Eq. 6.

Figure 6 shows the time-of-flight distribution for lowenergy S2 events. Here the time-of-flight has been calculated in nanoseconds and the offset $\Delta t_{\rm off}$ removed. Cuts were made according to the maximum and minimum times-of-flight generated in a Monte Carlo using the calculated neutron source spectrum described in Sec. III A, allowing for an additional 8 ns (approximately $2\sigma_{\gamma}$) on both ends of the cut range. Any event with time-of-flight

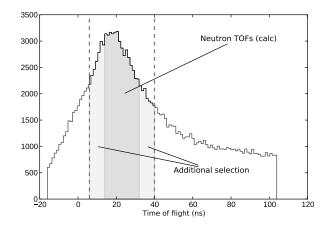


FIG. 6. Time-of-flight distribution for properly tagged events with S2 < 40000 photons and S1 < 50 photons (corrected for z-dependence). The shaded region shows the time-of-flight cuts applied to select 6 ns < τ < 40 ns.

within the selected region shown in Fig. 6 was considered to have passed the time-of-flight cuts.

Electron diffusion provides an additional means of confirming the validity of an event with a single S2 pulse. If the selected S1 truly corresponds to the primary scintillation produced by the ionization collected as S2, the width of the S2 pulse will increase with the drift time of the ionization approximately as [28]

$$\sigma^2 = \sigma_0^2 + 10D_L^2 \bar{t} / v_d, \tag{8}$$

where σ_0^2 is a constant determined by the drift time of the electron across the electroluminescent (EL) gap, D_L is the longitudinal diffusion constant in mm/ $\sqrt{\text{cm}}$, v_d is the electron drift velocity in mm/ μ s, and \bar{t} is the drift time in μ s. Note that this relation is an approximation as it assumes a Gaussian S2 pulse, which may not be strictly the case for events originating near the EL gap. Events for which S1 was properly selected will fall in a band described approximately by Eq. 8, shown in Fig. 7, in which the S2 pulse width was determined by a Gaussian fit. The centroid and $\pm 2\sigma$ lines of the band were determined using a procedure similar to the one applied in [29]. Using a maximum drift time of about 95 μ s and a drift length of 8 cm, we have $v_d = 0.84 \text{ mm}/\mu\text{s}$, and using the constant and linear terms of the centroid fit line, we find $\sigma_0 = 0.89~\mu \text{s}$ and $D_L = 0.37~\text{mm}/\sqrt{\text{cm}}$. All events within the $\pm 2\sigma$ lines are considered to have passed the diffusion cuts.

Using the pattern of light cast upon the PMT plane during S2 production, an average (x,y) location for each event can be constructed. Though this method does not provide the precision of a finer-granularity tracking plane, it allows for some fiducialization and, therefore, elimination of events that originate from the walls of the TPC. The average x and y coordinates are calculated by weighting the position of each PMT by the amount of S2 signal

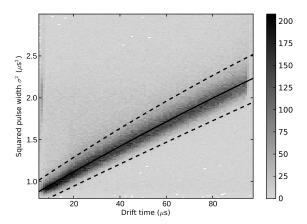


FIG. 7. Pulse width σ^2 from a Gaussian fit to the S2 pulse in each event plotted against drift time for events near and within the diffusion band. The centroid of the band is marked with a solid line, and dashed lines define the region of ± 2 standard deviations from the centroid. The equation of the centroid fit shown is $\sigma^2 = 0.801 + 0.0161\bar{t} - 0.0000143\bar{t}^2$.

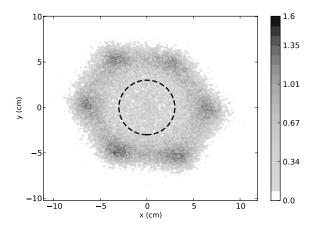


FIG. 8. The distribution of average reconstructed (x,y) locations for events with S2 < 40000 photons. The shading is done on a \log_{10} scale. The dashed circle defines a fiducial cut of r < 3 cm.

observed by that PMT. The resulting pattern is shown in Fig. 8 and has been scaled and shifted so that it is centered upon (0,0) and its dimensions match those of the physical dimensions of the PMT plane. This required a shift of all reconstructed points by (0.21, -0.20), scaling in x by a factor of 8.71, and scaling in y by a factor of 12.6. The scaling and shift procedure was necessary due to the uniformity of the light pattern cast on the PMTs and uncertainties in the individual PMT single-photon responses. A fiducial cut of r < 3 cm is superimposed, and events lying inside the selected region pass the cut. The tight fiducial cut is made here to show more clearly where the nuclear recoils lie in S1-S2 space, but this cut is varied in the forthcoming discussions to obtain increased statistics at the expense of more background events.

From the known characteristics of nuclear recoil signals

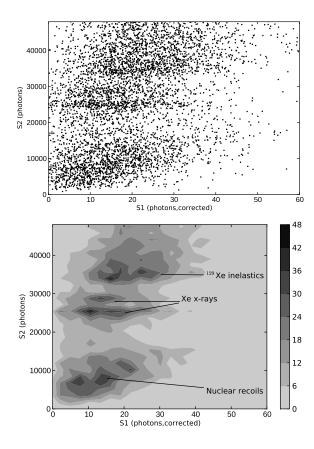


FIG. 9. S1 (corrected for z-dependence) vs. S2 signals for events passing tagging, single-pulse, time-of-flight, diffusion, and radial cuts, shown in scatter (above) and contour (below) formats. Events produced by neutron inelastic scattering on ¹²⁹Xe (approx. 40 keV), by xenon fluorescent x-rays (approx. 29 keV and 34 keV), and by nuclear recoils lie in distinct bands on the plot.

in liquid xenon, one suspects a class of events with low S1 and S2 and a different S2/S1 slope than the electronic recoil events. Figure 9 shows the relevant region of (S1, S2) space, including the low-energy nuclear recoil band, for all tagged events passing the single-pulse, time-of-flight, diffusion, and radial cuts (r < 3 cm) described above.

C. Electronic and Nuclear Recoil Discrimination

For the purpose of dark matter searches, one is especially interested in the ability to discriminate between nuclear recoils (the potential signals) and electronic recoils (background). In liquid xenon, this can be done by examining the ratio of the S2/S1 signals produced in an event, as this ratio is significantly lower on average for nuclear recoils. Because the light collection efficiency in the experiments producing the present data is low (of order 3-5%) and, therefore, the observed nuclear recoil events occupy a region of low S1 and S2 in which large fluctuations exist and the detection efficiency \neq 1, these

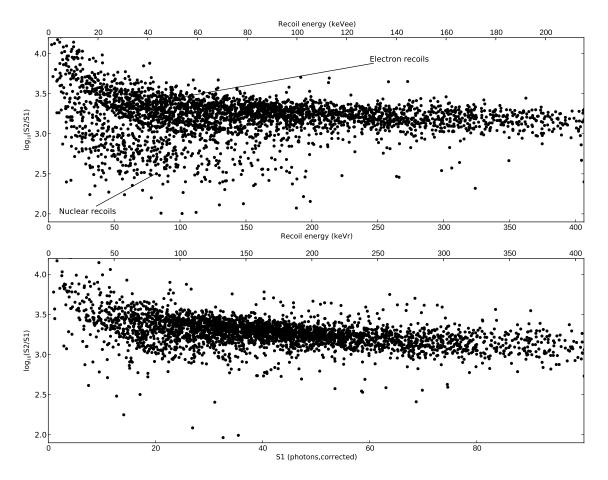


FIG. 10. The logarithm of the ratio of S2/S1 signals plotted (with equivalent scales) against S1 corrected for z-dependence for data taken with the 241 Am/Be source (above) and a 22 Na source (below). The S1 values are calibrated to their corresponding electron recoil energies (keV) and nuclear recoil energies (keVr) using the W-values W_i and W_{sc} , EL gain g, and light collection efficiency ϵ from section II C and the estimated nuclear recoil quenching factors α_{S1} and α_{S2} from section III D.

data are not ideal for characterizing the full potential of the discrimination power of xenon gas. However, they can be used to demonstrate that S2/S1 discrimination is possible and has potential in the gas phase.

Figure 10 shows the logarithm of the ratio S2/S1 plotted against S1 for data taken with the ²⁴¹Am/Be neutron source and with a ²²Na source (configuration 3 of Sec. IIB). The events shown passed the single-pulse, time-offlight, diffusion, and fiducial cuts discussed in Sec. III B. The number of events considered from the $^{241}\mathrm{Am/Be}$ dataset was reduced so that the two datasets could be compared with similar statistics. S1-S2 selection cuts were not applied so that events from both electronic and nuclear recoils could be shown. The nuclear recoil events form a band clearly distinguishable from the electronic recoil events, though some background events lie in the nuclear recoil band in the ²²Na data. This is due to several reasons, one being that at low S1, the small number of photons detected are subject to more significant Poisson fluctuations, resulting in poorer resolution and greater likelihood of yielding an abnormally large S1 for a given S2. Furthermore, it is possible to produce S1 in a

region of the TPC from which S2 cannot be collected, for example in the small gap between the PMT plane and the wire mesh that defines the beginning of the drift region. Often a low-energy event is accompanied by additional gamma rays that escape the active region, in which case an event with a single-pulse S2 could consist of additional gamma energy deposited but only seen in S1, yielding a range of possible S1 signals for a given S2.

D. Monte Carlo and Estimated Ionization and Scintillation Yields

The present results do not include information on the absolute energy of each nuclear recoil on an event-by-event basis, and so the ionization and scintillation yields for nuclear recoils can only be determined in principle by using the measured recoil spectrum as a whole and comparing it to expectations based on calculation and Monte Carlo simulation. For example, the peaks at larger angles in the neutron elastic scattering cross section (see Appendix) should lead to the presence of an identifi-

able feature in the recoil energy spectrum near 80 keVr. The presence of this feature is not statistically significant enough in the present data to make a strong definite claim, though its possible presence is investigated in [17] along with fits of experimental data to Monte Carlo spectra, and the results are used to obtain estimated nuclear recoil yields for S1 and S2. However, many uncertainties were present, including inaccuracies in the modeling of detector threshold effects and the energy dependence of the S1 and S2 yields, known to be non-trivial in liquid xenon [9] and for which no previously published data in gaseous xenon is known to the authors.

In this study rather than detailing a particular method of extracting information on the nuclear recoil yields, we choose values for the nuclear recoil yields (informed by the results obtained in [17]), assume a constant energy dependence, and then show that these assumptions are reasonable by comparison of experimental and Monte Carlo spectra. For this comparison we use data acquired with a 238 Pu/Be neutron source (positioned as in experimental setup 2 described in Sec. IIB) and a corresponding full Geant4 [30] Monte Carlo simulation. The electronic recoil yields were chosen by demanding consistency with the results obtained in Sec. II. $W_i = 24.7 \text{ eV}$ was assumed along with EL gain q = 734, and the PMT quantum efficiency Q was adjusted so that the 662 keV S2 peak in simulation was consistent with that found in experimental data, yielding $Q \approx 17\%$. $W_{\rm sc}$ was adjusted until the intercept $S_{1,0}$ matched that of Fig. 2, yielding $W_{\rm sc} = 45.69 \text{ eV}$. The difference between this value of $W_{\rm sc}$ and that calculated in Sec. II was used to assign an additional systematic uncertainty of 15 eV (see Eq. 5). For all recoil energies, the nuclear recoil yields are set to be equal to the electronic recoil yields multiplied by a quenching factor α , where we have chosen for S1 $\alpha_{S1} = 0.53$ and for S2 $\alpha_{S2} = 0.17$, corresponding to nuclear recoil yields of $Y_1 = \alpha_{\rm S1}/W_{\rm sc} = 11.6~{\rm ph/keV}$ for primary scintillation and $Y_2 = \alpha_{\rm S2}/W_{\rm i} = 6.9 \; {\rm e}^{-1}/{\rm keV}$ for ionization.

The Monte Carlo included the pressure vessel, teflon walls of the field cage and teflon reflector, PMT array, mesh grids, and lead block. A detailed model for the wavelength shifting introduced by the TPB was not implemented, however the teflon reflectivities on the walls and the teflon reflector were selected such that the zdependence of S1 matched that of Fig. 2 in a simulation using axially-incident 662 keV gamma rays, similar to the experimental setup using the ¹³⁷Cs source described in Sec. II. The walls were taken to be 100% reflective and the back reflector to be 79% reflective. Though these values are not consistent with the expected teflon reflectivity in xenon gas (from [31] expected to be 50-60%), they properly reproduced the linear geometric dependence of S1. The source was modeled by emitting neutrons from a single point behind the lead block in a random direction, accompanied by a 4.4 MeV gamma ray also emitted in a random direction. One neutron and one gamma ray were emitted per event, and the spectrum of emitted neutron energies was taken to be that calculated for neutrons produced with a carbon nucleus in the first excited state ¹²C* (similar to that shown in Fig. 3). The neutron interactions were modeled using the Geant4 high-precision (HP) neutron models, which use neutron cross section data from evaluated nuclear data libraries. The consistency of the Monte Carlo and calculated spectra of nuclear recoils can be seen by examining Fig. 14.

In the simulation, each photon produced via primary scintillation was tracked throughout its entire trajectory, and ionization electrons were produced in groups called ionization clusters. The final location of each ionization cluster after drift was calculated, taking into account diffusion, and the electroluminescent process was modeled by producing a number of photons equal to the EL gain g. Each EL photon was not tracked individually but its detection probability was looked up in a table indexed by its (x,y) production location in the 2D EL plane. This table was produced in an independent Monte Carlo run in which 10^6 photons were generated per point on a grid dividing the EL plane and the detection probability for each PMT recorded based on the number of photons collected out of the 10^6 generated. From the record of photons detected at each time in the event, realistic waveforms were constructed for each PMT matching the noise characteristics of the experimental waveforms and adding exponential pulses for each photoelectron detected. The resulting waveforms were passed through a nearly identical analysis to that of the experimental data to give results that could be compared directly to experiment. To simulate the experimental gamma-ray tagging procedure, only neutron-induced events, those in which a neutron scattered inelastically on any material in the simulation or elastically on xenon, were considered in the analysis. To roughly match the detector threshold effects observed at low S1 values in experiment, a peak-finding threshold was chosen appropriately in the Monte Carlo analysis.

Figure 11 shows the S2 signals plotted against S1 corrected for z-dependence in both data and Monte Carlo. Smearing was applied to the S1 and S2 values determined in Monte Carlo to account for geometric dependencies in the detection that degraded the experimental energy resolution, which appeared to influence more heavily S1. The analysis included tagging, single-pulse, diffusion, and radial cuts. Similar cuts were made in the corresponding Monte Carlo run, and the good qualitative agreement, particularly in the location of the nuclear recoil band, shows that the nuclear recoil quenching factors employed in the simulation are reasonable. One significant discrepancy in the Monte Carlo results is the relatively fewer number of events in the band lying at approximately $35000~\mathrm{S2}$ photons which is due to $40~\mathrm{keV}$ gamma rays produced by neutron inelastic scattering on ¹²⁹Xe. This is likely due to the fact that only neutrons accompanied by a 4.4 MeV gamma (energies 2-6 MeV) are considered in the Monte Carlo, while in the experimental data some fraction of the neutrons accompanied by the 7.7 MeV gamma (energies < 3 MeV) are also included. This is

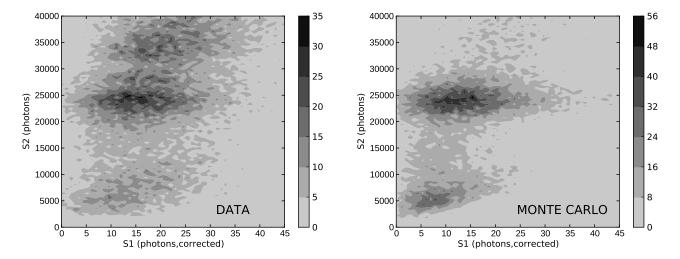


FIG. 11. S2 vs. S1 for low-energy events in experiment and Monte Carlo simulations generated based on the selected constant nuclear recoil yields.

because the 7.7 MeV gamma may not fully deposit its energy in the NaI scintillator and therefore may give an energy in the window of allowed NaI energies (2.4-5.0 MeV for the experimental run described in this section, and similar for that described in Sec. III B - III D). The inelastic n + ¹²⁹Xe scattering cross section is significantly higher [32] for these lower energy neutrons, so one should expect relatively more 40 keV gamma rays in the experimental data. However, the nuclear recoils produced by these lower energy neutrons will be lower in energy and therefore should not contribute significantly to the nuclear recoil spectrum at the energies observable in the data.

IV. CONCLUSIONS

We have simultaneously observed ionization and scintillation produced by nuclear recoils in gaseous xenon. It was confirmed that xenon in the gas phase, as in the liquid phase, is capable of distinguishing nuclear and electronic recoils based on the ratio of observed scintillation to ionization. It appears that relative to the corresponding yields for electronic recoils, the primary scintillation yield for nuclear recoils is quenched by a factor of roughly 2, and the ionization yield by a factor of roughly 5. Further investigation is required to determine precise nuclear recoil yields and fully investigate the potential advantages offered by gas phase operation. The greatest impact of the present results is the ability to inform future measurements of nuclear recoils in gaseous xenon.

The presence of gamma depositions in regions from which S2 cannot be collected provides an evident background and is known to also be of concern in the liquid phase. This is likely to be of even greater concern in the gas phase, as gamma rays can travel farther, and is evident in the inability to eliminate events consisting only of a single fluorescent x-ray. In such events, x-rays produced outside of the active region were able to travel into the active region and interact there. Such events can in principle be eliminated with a fiducial cut, but this would require a larger detector than the one used in the present experiment.

Though only estimates of the nuclear recoil yields were given, they can be used to predict the necessary light collection efficiency required to observe recoils of a given energy of interest. With 3% light collection efficiency at the EL plane, we were not able to see many recoils with energy less than approximately 30 keV. Therefore a $\sim 10\%$ light collection efficiency at least would be necessary to perform a stronger measurement of the nuclear recoil yields down to near 10 keV. The higher photon statistics will also help in characterizing the electronic/nuclear recoil discrimination power based on S2/S1 that is possible in the gas phase. The TPB was found to be a necessity for achieving enough light collection efficiency to identify the nuclear recoils, and in the future more sophisticated ideas such as the use of light guides coupled to PMTs [15] may be necessary to achieve the light collection regime of interest.

Appendix: Radioisotope Neutron Sources

1. Neutron Production Mechanism

The neutron sources used in this study all consist of an α -emitting radioactive isotope mixed with beryllium (${}^{9}\text{Be}$) and generate neutrons based on the (α, n) reaction [33]

$$\alpha + {}^9 \text{ Be} \rightarrow {}^{12} \text{ C} + \text{n.}$$
 (A.1)

The Q-value of the reaction is Q = 5.701 MeV, and this

TABLE I. Selected radioactive decay products of ²³⁸Pu and ²⁴¹Am. All data shown is from [25]. Both isotopes emit alpha particles with an average energy of approximately 5.5 MeV. Gamma ray emission from ²³⁸Pu is negligible, while some lowenergy gamma rays are emitted by ²⁴¹Am. In particular, the 60 keV gamma ray is likely to produce significant background without sufficient shielding of the source.

Isotope, $\tau_{1/2}$, Q	Product	Energy (keV)	% BR
²³⁸ Pu	α	5357.7	0.105
$\tau_{1/2} = 87.7 \pm 0.3 \text{ yr}$	α	5456.3	28.98
$Q = 5593.20 \pm 0.19$	α	5499.03	70.91
$^{241}\mathrm{Am}$	γ	13.946	9.6
$\tau_{1/2} = 432.2 \pm 0.7 \text{ yr}$	γ	59.5412	35.9
$Q = 5631.81 \pm 0.12$	α	5388.23	1.6
	α	5442.80	13.0
	α	5485.56	84.5

energy is released in the form of neutron kinetic energy, neglecting any carbon recoil kinetic energy, unless the carbon nucleus is left in an excited state, in which case some of the energy is emitted in the form of a gamma ray in coincidence with the neutron [34]. If left in the first excited state ¹²C*, a gamma ray of energy 4.439 MeV is emitted, and if left in the second excited state ¹²C**, a gamma ray of energy 7.654 MeV is emitted [35]. In this study we detect the coincident gamma and include it in the acquisition trigger to tag neutron-emitting decays and thereby significantly reduce the number of background events acquired.

We describe here how to calculate the neutron spectra of radioisotope sources under the assumption that their active regions consist of a uniform volume of ⁹Be throughout which the α -emitting isotope is uniformly distributed. We also assume that the total number of α -emitting isotope atoms present is much less than the total number of beryllium atoms in the mixture, so that each emitted α can be considered to interact with only atoms of beryllium. In addition, we do not consider neutrons produced due to the break-up reaction $\alpha + {}^{9}\text{Be}$ $\rightarrow \alpha' + {}^{9}\text{Be}^{*}, {}^{9}\text{Be}^{*} \rightarrow {}^{8}\text{Be} + \text{n [36]}.$ These neutrons lie at lower energies $\lesssim 3$ MeV, and they will not be observed in the adopted trigger scheme as no coincident gamma ray is emitted. Our calculations follow those of [36], [37], and [38], and make use of (α, n) cross sections from the Japanese Evaluated Nuclear Data Library (JENDL) [39] and neutron-xenon scattering cross sections from the Evaluated Nuclear Data File (ENDF) [32]. The cross sections were processed using the tools developed in [40]. Table I gives information on the decays of isotopes ²³⁸Pu and $^{241}\mathrm{Am}$ used as α -emitters in the sources used in this study.

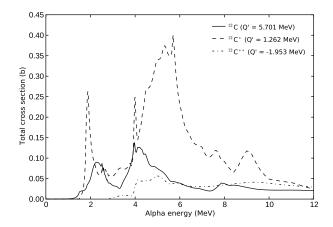


FIG. 12. Total cross section for the reaction ${}^{9}\mathrm{Be}(\alpha,\mathrm{n}){}^{12}\mathrm{C}$. All data shown in this figure are from JENDL [39].

2. Spectrum of Emitted Neutrons

The neutron production process proceeds as follows. The α -emitting isotope decays, yielding an α particle of mass m_{α} that travels through the surrounding medium of beryllium atoms of mass m_b , losing energy and often stopping completely without undergoing the reaction shown in Eq. A.1. However, some alpha particles (of order 1 in 10^4) will undergo the neutron-producing reaction of interest at an energy of E_{α} and generate a neutron of mass m_n at some angle θ in the center of mass frame of the interaction that can be directly related to the emitted neutron energy E_n via [37]

$$\cos \theta = E_n (m_n + m_c)^2 - E_{\alpha} (m_b m_c + m_n m_{\alpha}) - Q' m_c (m_n + m_c) \frac{-Q' m_c (m_n + m_c)}{2[E_{\alpha} m_{\alpha} m_n (E_{\alpha} m_b m_c + Q' m_c (m_n + m_c))]^{1/2}},$$
(A.2)

where Q' in this case is the energy released in the reaction in the form of neutron kinetic energy and may be equal to the full Q-value of 5.701 MeV or the full Q-value minus the energy of excitation left with the resulting carbon nucleus that is emitted in the form of a gamma ray. The angular distribution of neutrons for a given alpha energy E_{α} can thus be expressed in terms of the neutron energy E_n , and the number of neutrons emitted with energy in an interval $(E_n, E_n + dE_n)$ can be written as

$$G(E_n; E_{\alpha})dE_n = \frac{1}{\sigma_T(E_{\alpha})} \frac{d\sigma(E_n; E_{\alpha})}{d\Omega} \cdot \frac{4\pi}{E_n(\theta = 0) - E_n(\theta = \pi)} dE_n,$$
(A.3)

where $\sigma_T(E_\alpha)$ is the total cross section of the (α, \mathbf{n}) reaction for alpha energy E_α , shown in Fig. 12, and $d\sigma/d\Omega$ is the differential cross section. The number of neutrons

with energy E_n produced for emitted alpha i of energy $E_{\alpha,i}$ can then be determined by multiplying $G(E_n; E_{\alpha})$ by the probability of interaction at alpha energy E_{α} along the track, that is, $\rho \sigma_T(E_{\alpha}) dx = \rho \sigma_T(E_{\alpha}) (dx/dE_{\alpha}) dE_{\alpha}$, and integrating over all energies from 0 to $E_{\alpha,i}$,

$$N_{i}(E_{n}) = \int_{0}^{E_{\alpha,i}} \frac{4\pi [d\sigma(E_{n}; E_{\alpha})/d\Omega]}{dE_{\alpha}/(\rho dx)[E_{n}(\theta=0) - E_{n}(\theta=\pi)]} dE_{\alpha}.$$
(A.4)

The final spectrum will be a sum of such integrals over the emitted α energies of the source weighted by their branching ratios x_i ,

$$N(E_n) = \sum_{i} x_i N_i(E_n). \tag{A.5}$$

Figure 3 of Sec. III A shows the spectrum for a 241 Am/Be source calculated assuming the alpha particle energies and branching ratios from Table I. Note that this calculation is only valid for $dx \ll \lambda_{\alpha,n}$, where $\lambda_{\alpha,n}$ is the mean interaction length for the (α,n) . However, the average length of the track produced by a 5.5 MeV alpha particle in beryllium (density $\rho=1.23\times10^{23}$ cm⁻³ [41]) can be calculated using the alpha stopping power from [42] as $\int (dx/dE)dE \approx 28\mu\text{m}$. Using total cross section $\sigma_T < 0.4$ barns (see Fig. 12), $\lambda_{\alpha,n} = (\rho\sigma_T)^{-1} > 20$ cm, and therefore this condition holds.

3. Resulting Spectrum of Nuclear Recoils

From a spectrum of emitted neutrons, one can construct a spectrum of nuclear recoils produced when those neutrons are incident on a volume of xenon atoms by knowing the cross section for neutron elastic scattering (differential with respect to solid angle). Similar to Eq. A.2, we can express the energy of a nuclear recoil in terms of the cosine of the scattering angle in the center-of-mass frame of the neutron-nucleus collision as

$$E_{x'} = \frac{2E_n m_n m_{x,a}}{(m_n + m_{x,a})^2} (1 - \cos \theta),$$
 (A.6)

where $m_{x,a}$ is the mass of the target xenon nucleus and a is an index corresponding to the isotope of the nucleus.

Similar to Eq. A.3, we can use this relation to write the angular distribution of scattered neutrons in terms of the recoil energy, and determine the number of neutron interactions per unit recoil energy $N_{r,a}(E_{x'}; E_n)$ yielding a nuclear recoil with energy in the interval $(E_{x'}, E_{x'} + dE_{x'})$ for a neutron of energy E_n incident on a xenon nucleus of isotope a,

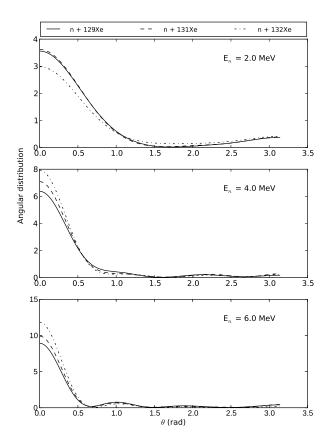


FIG. 13. Angular distributions (data from [32]) for elastic neutron scattering, $n + Xe \rightarrow n' + Xe'$, for three different neutron energies incident on three different xenon isotopes. Note the strong peak in the forward direction, followed by one or several peaks at higher angles which are responsible for the structure in the nuclear recoil spectrum at higher energies.

$$N_{r,a}(E_{x'}; E_n) = \frac{(m_n + m_{x,a})^2}{m_n m_{x,a}} \cdot \frac{\pi}{E_n} \left[\frac{d\sigma(E_{x'}; E_n)}{d\Omega} \right] \rho \Delta x, \tag{A.7}$$

where ρ is the density of the xenon volume and Δx is the thickness of xenon traversed by the neutron. Figure 13 shows the angular distribution for neutrons scattered off of several different xenon isotopes. Note that because we will be interested in the normalized spectrum, and $\rho\Delta x$ is a constant independent of the isotope, it can be absorbed into an overall normalization factor in the final spectrum, which is a weighted superposition of individual nuclear recoil spectra summed over xenon isotopes and integrated over neutron energies,

$$N(E_{x'}) = \sum_{a} f_a \int_0^\infty N(E_n) N_{r,a}(E_{x'}; E_n) dE_n. \quad (A.8)$$

where the f_a are the fractional compositions of natural xenon for each isotope and $N(E_n)$ is the spectrum of emitted neutrons from Eq. A.5.

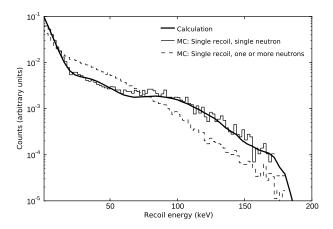


FIG. 14. Spectrum of nuclear recoils assuming neutrons emitted from a $^{238}{\rm Pu/Be}$ source in which the carbon nucleus resulting from the $^9{\rm Be}(\alpha,{\rm n})^{12}{\rm C}$ reaction was left in the first excited state. The spectrum was obtained from a calculation using Eq. A.7 and Eq. A.8 (thick solid line) and from a Geant4 Monte Carlo. Two curves are shown from the same Monte Carlo run. The solid line is the spectrum considering only events containing a single xenon nuclear recoil and a single neutron. The dashed line is the spectrum of events containing a single xenon nuclear recoil, but in which the event may have contained multiple neutrons produced by interaction of the incident neutron with the lead. The presence of these secondary neutrons significantly distorts the spectrum of xenon nuclear recoils.

Figure 14 shows the calculated spectrum of nuclear recoils assuming the neutron spectrum is that emitted by the source with a coincident 4.4 MeV gamma ray, corresponding to the dashed curve in Fig. 3. A target of natural xenon is assumed with fractional isotopic composition taken from the "representative" values reported in [43].

Isotopes ¹²⁴Xe and ¹²⁶Xe are excluded from the calculation due to lack of cross section data but have negligible natural abundances. The spectrum is compared with results from a Geant4 Monte Carlo of the experimental setup in which a source emitting the same neutron spectrum input to the calculation was positioned behind a 2-in. thick lead block. The calculated and Monte Carlo recoil spectra agree well when considering only events in which a single xenon recoil occurred and only one neutron was present throughout the entire event. Including also events in which the emitted neutron interacted in the lead to produce additional neutrons gives a significantly altered spectrum.

ACKNOWLEDGMENTS

This work was supported by the following agencies and institutions: the Director, Office of Science, Office of Basic Energy Sciences, of the U.S. Department of Energy, and the National Energy Research Scientific Computing Center (NERSC), supported by the Office of Science of the U.S. Department of Energy, both under Contract No. DE-AC02-05CH11231; the European Research Council under the Advanced Grant 339787-NEXT; the Ministerio de Economía y Competitividad of Spain under Grants CONSOLIDER-Ingenio 2010 CSD2008-0037 (CUP), FPA2009-13697-C04-04, FPA2009-13697-C04-01, FIS2012-37947-C04-01, FIS2012-37947-C04-02, FIS2012-37947-C04-03, and FIS2012-37947-C04-04; and the Portuguese FCT and FEDER through the program COMPETE, Projects PTDC/FIS/103860/2008 and PTDC/FIS/112272/2009. J. Renner acknowledges the support of a Department of Energy National Nuclear Security Administration Stewardship Science Graduate Fellowship, grant number DE-FC52-08NA28752.

E. Aprile, A. E. Bolotnikov, A. I. Bolozdynya, and T. Doke, *Noble Gas Detectors* (Wiley VCH, Verlag GmbH & Co. KGaA, Weinheim, 2006).

^[2] E. Aprile and T. Doke, Rev. Mod. Phys. 82, 2053 (2010).

^[3] D. S. Akerib *et al.* (LUX Collaboration), Phys. Rev. Lett. 112, 091303 (2014).

^[4] E. Aprile (XENON100 Collaboration), Phys. Rev. Lett. 109, 181301 (2012).

^[5] D.Yu. Akimov et al. (ZEPLINIII Collaboration), Phys. Lett. B 709, 14 (2012).

^[6] M. Auger et al. (EXO Collaboration), Phys. Rev. Lett. 109, 032505 (2012).

^[7] G. Bertone, D. Hooper, and J. Silk, Physics Reports 405, 279 (2005).

^[8] J. J. Gómez-Cadenas, J. Martín-Albo, M. Mezzetto, F. Monrabal, and M. Sorel, Riv. Nuovo Cim. 35, 29 (2012).

^[9] M. Szydagis, N. Barry, K. Kazkaz, J. Mock, D. Stolp, M. Sweany, M. Tripathi, S. Uvarov, N. Walsh, and M. Woods, JINST 6, P10002 (2011),

http://nest.physics.ucdavis.edu/site.

^[10] E. Aprile, K. L. Giboni, P. Majewski, K. Ni, and M. Yamashita, Phys. Rev. B 76, 014115 (2007).

^[11] P. Sorensen et al., Nucl. Instrum. Methods A 601, 339 (2009).

^[12] M. Horn et al. (ZEPLINIII Collaboration), Phys. Lett. B 705, 471 (2011).

^[13] A. Bolotnikov and B. Ramsey, Nucl. Instrum. Methods A 396, 360 (1997).

^[14] E. Conti et al., Phys. Rev. B 68, 054201 (2003).

^[15] D. R. Nygren, J. Phys. Conf. Ser. 460, 012006 (2013).

^[16] D. C. Herrera et al., International Conference on Technology and Instrumentation in Particle Physics (TIPP), (2014), (Proceedings to be published).

^[17] J. Renner, Ph.D. thesis, University of California, Berkeley (2014).

^[18] J. Renner, V. Gehman, A. Goldschmidt, C. Oliveira, and D. Nygren, in 13th International Conference on Topics in Astroparticle and Underground Physics (TAUP) (2013) (Proceedings to be published).

- [19] J. White, Presentation at PPC 2008: 2nd International Workshop on Interconnection Between Particle Physics and Cosmology.
- [20] V. Álvarez et al. (NEXT Collaboration), Nucl. Instrum. Methods A 708, 101 (2013).
- [21] D. Nygren, Nucl. Instrum. Methods A 603, 337 (2009).
- [22] C. M. B. Monteiro et al., JINST 2, P05001 (2007).
- [23] ROOT, http://root.cern.ch/drupal.
- [24] FMWK, https://cdcvs.fnal.gov/redmine/projects/nusoft/wiki/FMWK.
- [25] L. P. Ekström and R. B. Firestone, WWW Table of Radioactive Isotopes, database version 2/28/99, http://ie.lbl.gov/toi.
- [26] A. Parsons, T. Edberg, B. Sadoulet, S. Weiss, J. Wilkerson, K. Hurley, R. Lin, and G. Smith, IEEE Trans. Nucl. Sci. 37, 541 (1990).
- [27] S. J. C. do Carmo et al., JINST 3, P07004.
- [28] V. Álvarez *et al.* (NEXT Collaboration), JINST 8, P05025 (2013).
- [29] C. E. Dahl, Ph.D. thesis, Princeton University (2009).
- [30] Geant4 Collaboration, Nucl. Instrum. Methods A 506, 250 (2003).
- [31] C. Silva et al., J. Appl. Phys. 107, 064902 (2010).
- [32] "Evaluated Nuclear Data File (ENDF)," http://www.nndc.bnl.gov/exfor/endf00.jsp.
- [33] G. F. Knoll, Radiation Detection and Measurement, 3rd ed. (Wiley, Hoboken, N.J., 2000).

- [34] The mean lifetimes of the excited carbon nuclear states of interest here are so short (< 0.1 ps) compared to the measurable time scale that the gamma can be considered to be emitted simultaneously with the neutron for practical purposes.
- [35] "Evaluated Nuclear Structure Data File (ENSDF)," http://www.nndc.bnl.gov/ensdf.
- [36] K. Geiger and L. V. D. Zwan, Nucl. Instrum. Methods 131, 315 (1975).
- [37] A. Vijaya and A. Kumar, Nucl. Instrum. Methods 111, 435 (1973).
- [38] A. Kumar and P. Nagarajan, Nucl. Instrum. Methods 140, 175 (1977).
- [39] T. Murata et al., "JENDL (α ,n) Reaction Data File 2005," http://wwwndc.jaea.go.jp/ftpnd/jendl/jendl-an-2005.html.
- [40] C. M. Mattoon et al., Nuclear Data Sheets 113, 3145 (2012), https://ndclx4.bnl.gov/gf/project/gnd/.
- [41] J. Beringer et al. (Particle Data Group), Phys. Rev. D 86, 010001 (2012).
- [42] NIST, "ASTAR: stopping power and range tables for helium ions," http://physics.nist.gov/PhysRefData/Star/Text/ASTAR.html. Accessed July 19, 2013.
- [43] P. D. Biévre and P. Taylor, Int. J. Mass Spec. Ion Processes 123, 149 (1993).