Measurement of scintillation efficiency for nuclear recoils in liquid argon

Dan Gastler,¹ Ed Kearns,¹ Andrew Hime,² Laura C. Stonehill,² Stan Seibert,² Josh Klein,³ W. Hugh Lippincott,⁴ Daniel N. McKinsey,⁴ and James A. Nikkel⁴

¹Boston University, Boston, MA ²Los Alamos National Laboratory, Los Alamos, NM ³University of Pennsylvania, Philadelphia, PA ⁴Yale University, New Haven, CT (Dated: May 9, 2012)

The scintillation light yield of liquid argon from nuclear recoils relative to electronic recoils has been measured as a function of recoil energy from 10 keVr up to 250 keVr at zero electric field. The scintillation efficiency, defined as the ratio of the nuclear recoil scintillation response to the electronic recoil response, is $0.25 \pm 0.01 + 0.01$ (correlated) above 20 keVr.

PACS numbers:

I. INTRODUCTION

field.

A number of existing and proposed experiments use liquefied noble gases as detection media for Weakly Interacting Massive Particles (WIMPs) [1–5], a well motivated dark matter candidate [6]. Liquefied noble gases have a high scintillation yield, are relatively simple to purify of both radioactive contaminants and light absorbers, and should be easily scalable to the large masses required for very sensitive detectors. Although the best limit for the spin-independent WIMP-nucleon cross section is currently set by the germanium-based CDMS experiment [7] at $3.8 \times 10^{-44}~\rm cm^2$ for a 70-GeV WIMP mass, the XENON-10 experiment has set a comparable limit of $8.8 \times 10^{-44}~\rm cm^2$ for a 100-GeV WIMP mass [8], showing that liquefied noble gases are viable dark matter targets.

Events in a noble liquid dark-matter detector may arise from scattering off of the nucleus or atomic electrons; dark matter will only scatter off the nucleus to an appreciable extent. The ratio of the scintillation light yield for nuclear recoil events relative to electronic recoil events is defined as the scintillation efficiency or $L_{\rm eff}$.

A WIMP dark matter search requires an energy threshold on the order of tens of keV, and it is necessary to measure the scintillation efficiency down to this energy threshold so as to quantify the WIMP detection sensitivity. In order to make this measurement of $L_{\rm eff}$, a D-D neutron generator was used to produce neutrons that scattered from a liquid argon detector into an organic liquid scintillator detector used as a coincidence trigger. The organic scintillator was placed at a series of known angles, and the energies of the selected nuclear recoils in the liquid argon were kinematically determined. The scintillation efficiency was determined from the ratio of the measured electron-equivalent recoil energy at a given scattering angle to the expected nuclear recoil energy (keVr) at that angle. Details of this measurement in a 4-kg liquid argon detector are presented in this paper, along with scintillation efficiency results for nuclear recoil energies between 10 and 250 keVr at zero electric

II. REVIEW OF PHYSICAL PROCESSES AND MEASUREMENTS

Discrimination between nuclear recoil events that characterize a WIMP signal and electronic recoil events that characterize the primary backgrounds is essential in WIMP detectors, particularly for the case of liquid argon which contains the radioactive isotope ³⁹Ar. The noble liquid detectors use two methods to achieve this discrimination. Single-phase detectors use pulse-shape discrimination (PSD) based solely on scintillation light to discriminate between event types, while dual-phase detectors can collect both scintillation light and ionization, employing a combination of PSD and the relative size of the light and ionization channels to identify events. PSD is made possible because ionizing radiation in liquid noble gases results in the formation of excited diatomic molecules (excimers) that can exist in either singlet or triplet states, with very different lifetimes. In liquid argon these lifetimes are 7 ns and 1.5 μ s, respectively [9, 10] and the scintillation light is produced in the decay of these states. As different types of excitation produce different ratios of triplet to singlet molecules, the relative amplitudes of the fast and slow components can be used to determine what type of excitation occurred. The effectiveness of this PSD is directly dependent on the number of detected photoelectrons in an event, and thus the light yield for both nuclear recoils and electronic recoils sets the energy threshold for which electronic recoil backgrounds are negligible, in turn determining the ultimate sensitivity of the detector to dark-matter-induced nuclear recoils.

The excimers that provide the scintillation light are formed in two ways. An excited atom (exciton) can combine with another atom in the liquid to produce the excimer, or an ionized atom can combine with another atom to form a diatomic ion, which in turn recombines with an electron, eventually resulting in the production of the excimer. The ratio of exciton production to ion pair pro-

duction in liquid argon has been calculated to be 0.21 [11], indicating that the majority of the scintillation light in liquid argon comes from excimers formed indirectly from argon ions, rather than directly from excited argon. The average energy required to produce an electron-ion pair in liquid argon has been measured to be 23.6 ± 0.3 eV [12], and the average energy needed to produce a single photon has been calculated to be 19.5 ± 1.0 eV [13]. From this, the maximum possible scintillation yield in liquid argon is about 51 photons per keV of deposited energy, in the extreme case where the excimer formation and scintillation processes are perfectly efficient.

In actuality, the absolute light yield is reduced through a number of different mechanisms. Energy may be lost by means other than exciton and ion pair formation, the excitons may undergo non-radiative collisions, and the recombination of diatomic ions may be incomplete. The first mechanism is known to be significant for nuclear recoils, for which a significant portion of the energy is lost to atomic motion as described by Lindhard [14]. Thus, the scintillation light yield is expected to be reduced for nuclear recoil events compared to electronic recoils. Measurements of the scintillation efficiency for nuclear recoils relative to electronic recoils in liquid xenon [15–19] indicate that there is an additional reduction in the nuclear recoil scintillation yield due to collisions between free excitons that result in an ion and a ground-state atom, as described by Hitachi [20]. The rate of these biexcitonic collisions is dependent on the density of the excitations, thus the amount of quenching increases with increasing linear-energy-transfer (LET) and is most significant at larger recoil energies. This mechanism of biexcitonic quenching is expected to apply to liquid argon as well as liquid xenon, and a model for scintillation efficiency in argon, neon and xenon taking LET into account has been proposed by Mei et al. [21] A further reduction in scintillation yield can result when some fraction of the ion-electron pairs do not recombine to produce an excimer and the electrons escape instead [13].

Relative scintillation efficiencies in liquid argon have been measured for a number of different particle types. For heavy fission fragments with kinetic energy around 80 MeV, the scintillation efficiency relative to 1-MeV electrons has been measured to be 0.21 ± 0.04 [22]. For alpha particles, the scintillation efficiency has been measured to be 0.9 for 6-MeV alphas relative to 1-MeV electrons [22] and 0.4 for 5.3-MeV alphas relative to 1.2-MeV electrons [23]. The observed triplet lifetime from the 5.3-MeV alphas was 800 ns, which may indicate additional absorption due to impurities, but both values are plausible given the expectation that the scintillation efficiency for alphas should fall somewhere between that of fission fragments and unity. A previous measurement of scintillation efficiency for nuclear recoils in liquid argon by the WARP collaboration gives $0.28 \pm 10\%$, measured at 65 keV average recoil energy [3].

III. EXPERIMENTAL DETAILS

The liquid argon scintillation efficiency was measured using the MicroCLEAN detector at Yale University. The active volume is 3.14 liters of liquid argon viewed by two photomultiplier tubes (PMTs). Figure 1 shows a schematic of the central region and PMTs. The active region is defined by a teflon cylinder 200 mm in diameter and 100 mm in height, with two 3-mm-thick fused-silica windows closing the top and bottom. Two 200-mm-diameter Hamamatsu R5912-02MOD photomultiplier tubes (PMTs) are held in place by teflon rings above and below the central volume and view the active region through the windows. Because liquid argon scintillates in the vacuum ultraviolet at 128 nm [24], all inner surfaces of the teflon and fused silica are coated with a thin film of tetraphenyl butadiene (TPB) [25]. The TPB shifts the wavelength of the ultraviolet light to approximately 440 nm so that it may pass through the windows and be detected by the PMTs. Both windows are coated with 0.20 ± 0.01 mg/cm² of TPB and the teflon cylinder is coated with 0.30 ± 0.01 mg/cm² of TPB. The teflon cylinder, windows and PMTs are all immersed directly in liquid argon, contained within a 25-cm-diameter by 91-cm-tall stainless steel vessel.

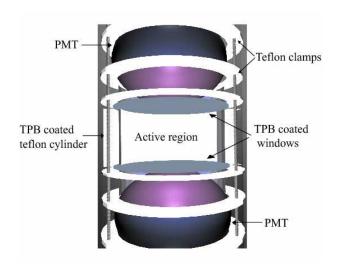


FIG. 1: Schematic representation of the scintillation volume and PMT orientation.

The stainless steel vessel is housed inside a vacuum dewar, and liquid argon is introduced though a tube on the top of the vessel. The argon is liquefied from purified gas in a copper vessel mounted to the end of a Cryomech PT805 pulse-tube refrigerator. All components that come into contact with the gas or liquid are baked to at least 60°C, and the ultra-high-purity argon gas (99.999%) is passed through a heated Omni Nupure III gas-purification getter before entering the vessel. Outgassing can cause impurities to build up in the detector, decreasing the light yield by quenching the argon excimers or absorbing the UV scintillation photons. To

avoid signal degradation, the argon is continually circulated through the getter and reliquefied at a rate greater than 2.0 slpm. No reduction in signal was observed during the run. PMT signals were ditigized using an 8-bit 500 MSPS waveform digitizer with each of the PMTs capturing both low-gain and high-gain waveforms. More details about the experimental apparatus, data acquisition, and purity measurements are available in [10].

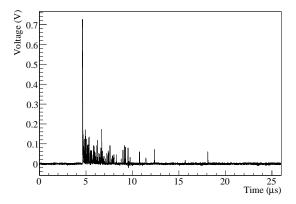


FIG. 2: Example of an electronic recoil-induced scintillation event in liquid argon.

A sample oscilloscope trace from an electronic recoil scintillation event in argon is shown in Fig. 2. A 10- μ Ci sealed ⁵⁷Co source is used for daily measurements of the scintillation light yield for electronic recoils, with a sample spectrum shown in Fig. 3 along with results from the simulation to be discussed in the next section. This source produces 122-keV, 137-keV and 14.4-keV gammarays with branching ratios of 86%, 11% and 9%, respectively. Spectra were taken for each day of data taking, with a Gaussian fit to the 122-keV peak providing a scintillation signal yield calibration for that day in units of photoelectrons per keV of energy deposited by an electronic recoil, denoted as photoelectrons/keVee. Over the course of the four-month run, the signal yield remained stable to within 5% at 4.85 ± 0.01 photoelectrons/keVee. To check the quality of the energy calibration, a 10- μ Ci ²²Na source that produces 511-keV gamma-rays is used as a second point of reference, and the 511-keV line appears at a photoelectron yield that is within 1% of the value predicted from ⁵⁷Co source calibration.

A portable Thermo Electron MP320 deuterium-deuterium (D-D) neutron generator is used as a neutron source, with an organic scintillator detector as a secondary coincidence trigger. The experimental setup can be seen schematically in Fig. 4. In the forward direction, the D-D generator produces 2.8-MeV neutrons. Some of these neutrons scatter in the liquid argon, and for a given position of the organic scintillator, only neutrons that scatter at a specific angle are selected by the coincidence trigger. By changing the angle the organic scintillator makes with the neutron generator-liquid argon detector axis, the scattering energy of the recoil nu-

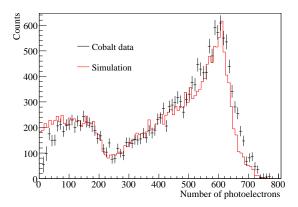


FIG. 3: (Color online) Plot of the 57 Co spectrum, along with a simulation of the expected spectrum.

cleus in the liquid argon can be varied according to the following equation:

$$E_{rec} = \frac{2m_n E_{in}}{(m_n + m_{Ar})^2} \left[m_n + m_{Ar} - m_n \cos^2(\theta) - \cos(\theta) \sqrt{m_{Ar}^2 + m_n^2 \cos^2(\theta) - m_n^2} \right],$$
(1)

where E_{in} is the incident neutron energy (2.8 MeV), A is the atomic mass number, and θ is the scattering angle of the outgoing neutron. Data were taken at 19 different angles corresponding to recoil energies between 10 keV and 250 keV. The setup also included 12 inches of poly between the neutron generator and the organic scintillator to reduce the accidental coincidence rate, although this is not shown in Figs. 4 or 5.

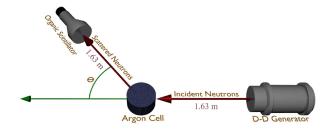


FIG. 4: (Color online) Top view of the neutron scattering setup. Shown are the neutron generator and the organic scintillator. The size of the argon cell is not representative.

IV. MONTE CARLO SIMULATIONS

In order to understand the data, we developed a Monte Carlo simulation of the argon detector, cryostat, organic scintillator and surrounding lab space. The software framework used was RAT, which combines Geant4[26], CLHEP[27], and ROOT[28] into a single simulation and

analysis package. A detailed optical model of the inner detector and PMTs is included in the Monte Carlo which allows us to estimate smearing of the detected signal. While this model gives results that are in fairly good agreement with our gamma calibrations, we add an addition smearing term for the neutron scattering analysis to take into account the lower photon yield for a given energy.

An image of the detector geometry can be seen in Fig. 5. The argon detector is in the central vertical cylinder, where the various layers of steel are set to be semitransparent so that the inner workings are visible. While there was only one organic scintillator detector at a given time in the real experiment, the simulation used many such detectors, so as to allow Monte Carlo data for half of the organic scintillator positions to be collected simultaneously and reduce computation time. In the picture, the cylinders off to the right of the argon detector represent the organic scintillator in its various positions. As the adjacent positions of the organic scintillator overlap, independent simulations were performed for each set of colored organic scintillator positions. The 12 inches of poly between the generator and the organic scintillator locations was not included in the simulation.

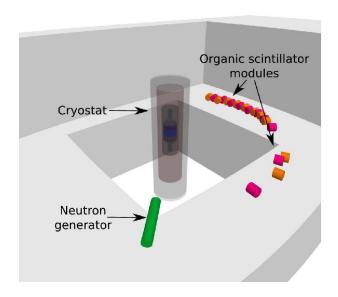


FIG. 5: (Color online) Monte Carlo detector geometry.

In addition to the neutron scattering simulation, the detector response to an external ⁵⁷Co gamma source was also modeled. The origin of the 122 keV and 137 keV gamma rays was set just outside of the outer vacuum can as in the real detector. The results of this simulation can be seen in Fig. 3 showing very good agreement with the experiment.

V. ANALYSIS

The analysis begins by combining the high-gain and low-gain waveforms for each PMT in a given event. To do this the first two microseconds of each PMT's high-gain and low-gain waveforms are separately averaged to calculate baselines. These baselines are then subtracted to give zero-offset waveforms. Each high gain waveform is then scanned to determine if the waveform digitizer is saturated. In the case of saturation, the high-gain and low-gain waveforms are aligned and the low-gain waveform's samples are inserted where the high-gain waveform was saturated.

At this point each PMT waveform is scanned to determine the trigger time of the event. The trigger time for each PMT is defined as the time where the waveform reaches 20% of its maximum height. The average of the two trigger times is taken as the start time for the event and a timing cut is applied to the two PMT waveforms to remove events where the difference in timing is greater than 20 ns. The waveforms from both PMTs are then integrated in two timing intervals, the first from 20 ns before the trigger time to 100 ns after and the second from 100 ns to 5 μ s. The region between 5 μ s and 14 μ s is scanned for single photoelectron pulses and used to determine the single photoelectron spectrum. Any region where the waveform's voltage value exceeded approximately one third of a single photoelectron's peak voltage is integrated from 10 ns preceding the crossover sample to 50 ns following it. After this procedure has been performed on every event in a run, the run's single photoelectron value is fit and used to convert the integrated waveform charges into photoelectrons.

A PMT asymmetry cut is used to remove events that are near the windows of the detector. The asymmetry is defined as the difference in the signals observed by the two PMTs divided by their sum, and events with an asymmetry of more than 60% are removed. Events with approximately 2000 photoelectrons or more can cause the saturation of one or both of the detector PMTs, and since this will cause events to have poor energy reconstruction, a cut is applied to remove events in which either PMT's output becomes greater than 2 V. This cut removed nuclear recoil events with energies above 110 keVee for runs with recoil angle below 125° degrees and nuclear recoil events with energies above 180 keVee for 125° and 142° degree runs. All cuts applied to the data up to this point are considered to be data quality cuts and were applied to both $^{57}\mathrm{Co}$ and neutron runs.

To distinguish neutron scatters from other backgrounds in the neutron data sets, two additional cuts involving the organic scintillator are applied. The first is a time-of-flight (TOF) cut which removes events in which the organic scintillator is triggered before the detector, as well as events in which the neutron arrives late due to multiple scatters. In addition, this cut also helps remove background gammas from our neutron data sets. The position and width of this cut is set by the location

of the single scattering neutrons in the Monte Carlo TOF spectrum.

The second cut associated with the organic scintillator uses the pulse shape in the organic scintillator to distinguish between neutrons and background events. We use a scatter plot of the pulse area within 100 ns of the organic scintillator trigger versus the organic scintillator waveform's maximum voltage, shown in Fig 6, to determine a quadratic curve that separates two types of events. This curve divides the scatter plot into two distinct regions: an electronic recoil band that appears for all types of runs and a nuclear recoil band that is only present in neutron runs. The separation of these regions did not appear to change with recoil energy and variation of the regions had a negligable effect on the results.

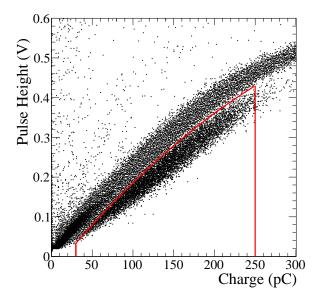


FIG. 6: (Color online) An example of the pulse shape cut applied to the organic scintillator. A scatter plot of the pulse area versus peak pulse height shows a distinct region of neutron events bounded by the red quadratic curve and lines.

We apply one final cut to remove electronic recoils from the sample, exploiting the PSD to discriminate between event types in liquid argon. Based on our previous work [10], we define a discrimination parameter, F_{prompt} , as the fraction of light arriving in a prompt time window. For comparison, the mean F_{prompt} for recoils between 5 and 32 keVee range from 0.39 to 0.28 for electronic recoils and 0.56 and 0.7 for nuclear recoils [10]. We apply a relatively loose cut, removing events with $F_{prompt} < 0.35$ from the final sample. We checked the analysis with a tighter cut of $F_{prompt} < 0.50$ and incorporated the differences in the systematic uncertainties.

To determine the scintillation efficiency of the liquid argon, the measured energy spectrum for each scattering angle is compared to the energy spectrum produced by the Monte Carlo simulation analyzed with the same asymmetry and TOF cuts. We also use the Monte Carlo simulation to calibrate the TOF cut. For each scattering angle in the Monte Carlo simulation, a TOF spectrum is produced using events that only scattered once in the detector before reaching the organic scintillator. This allows us to find the range of the TOF for single scattering neutrons for each position. After applying the same TOF and PMT asymmetry cut to the simulation as used for the data, a Monte Carlo energy spectrum is then generated for each recoil angle.

There are two convolutions applied to the Monte Carlo recoil spectra before fitting them to the data. First, to account for the variation in the single photoelectron charge, the Monte Carlo photon counts are smeared using the measured single photoelectron charge distribution from the photoelectron calibration data. Second, since the simulations were performed assuming a 100% scintillation efficiency, an additional smearing of $3.25 \times \sqrt{(1-L_{\rm eff})N_{pe}}$ is applied to the Monte Carlo to account for the difference in counting statistics between a scintillation efficiency of 100% and that obtained from the data. The proportionality constant of 3.25 empirically account for the observed broadness of the clearly resolved peaks at 191, 211 and 239 keVr scattering angles. It is well known that noble liquid detectors do not reach the ideal energy resolution predicted by Poisson photoelectron statistics, largely due to ionization-scintillation anti-correlation[1, 29].

The MINUIT fitting package is used to perform a χ^2 fit of the Monte Carlo to the data with the normalization and the scintillation efficiency as free variables. Each Monte Carlo spectrum is binned using the same binning as the corresponding recoil data and used to generate a spline for fitting. First, the entire range of the data is used in the fit. Then, a Gaussian is fit to all events in the Monte Carlo identified as singly-scattered neutrons. This fit is used to define a new fit range consisting of plus and minus three sigma around the centroid of the Monte Carlo single scattered neutron distribution. The final fits are performed over a restricted range around the single scattering Monte Carlo neutron distribution where we expect to observe our signal. This fitting procedure was checked using the Monte Carlo sample with a set $L_{\rm eff}$ value of 0.25 and was able to recover this set $L_{\rm eff}$ at each recoil angle.

The results for all organic scintillator positions are presented in Fig. 7, and the scintillation efficiency as a function of energy is shown in Fig. 8. After studying the systematic effects described in the next section, we found that the individual scintillation efficiency values were constant across the range of recoil energies studied above $20~\rm keVr$, with a mean of $0.25\pm0.01+0.01$ (correlated). However, there still existed substantial differences between the simulation and the data, giving an average $\chi^2/d.o.f.$ of 3.7 for measurments above 20 keVr. This disagreement is addressed in the next section. Below 20 keVr, our data exhibit an upturn in scintillation efficiency as the energy goes to zero, and we were unable to find an experimental cause for this upturn. It is therefore

unknown if this is a physically real effect or if we lose our ability to distinguish nuclear recoils from other backgrounds at these low energies. All observed values and uncertainties are listed in Table I.

VI. SYSTEMATIC UNCERTAINTIES

The sources of uncertainty considered are categorized as those associated with detector operation, triggering effects, Monte Carlo background normalization, TOF window and fit range effects. The uncertainties from the sources discussed in this section are all combined and the final resulting uncertainty for each scattering angle can be found in Table. I.

The first group of considered uncertainties deals with the data taking and stability of the neutron and ⁵⁷Co data runs. Since the ⁵⁷Co runs are used to calibrate the light yield of the detector, the fit error of the ⁵⁷Co peak and its stability over time directly affect the measured scintillation efficiency. These are estimated to be 2% and 1.6% respectively. There is a second uncertainty in the angle of the organic scintillator relative to the neutron generator that in turn introduces an uncertainty in the corresponding energy via Eq. 1. We have determined the uncertainty of the angular position of the organic scintillator to be 1.3° at each position.

We examined the effects of the trigger efficiency, specifically looking to address the upturn observed at low energies which could be explained by a bias introduced by the trigger level. We took data for the 22° run at three different hardware triggers and the 26° run at two different hardware triggers, and we examined the effect of hardware and software triggers on both the asymmetry cut and the final scintillation efficiency values. In all cases, the scintillation efficiency distributions did not systematically change by varying the cuts and hardware threshold. We also performed a toy Monte Carlo using the time dependence of the scintillation light [10] and the observed single photoelectron distribution to estimate possible threshold effects. This study found the effect of any threshold bias given our hardware trigger level to be less than 1%, much smaller than the other errors in the measurement. Therefore, we conclude that the triggering threshold does not explain the upturn at low energies.

A third source of uncertainty arises because the Monte Carlo simulation, as mentioned at the end of Sec. V, does not exactly reproduce the observed background shape. This can be seen by comparing the dashed and solid red lines in the large angle scattering spectra of Fig. 7. To account for this inconsistency, the data for each recoil energy were reanalyzed under the assumption that the size of the multiple scattering background in the histograms used to perform the fits varied by $\pm 50\%$ relative to that predicted by the Monte Carlo. The variations observed in this reanalysis are approximately 10% below $20~{\rm keVr}$ and 2% above $20~{\rm keVr}$ and are included in the errors listed in Table I.

To determine the uncertainty due to the TOF cuts, the TOF window was expanded separately up and down in time by 50%. This allows for recoil neutrons with smaller TOFs to be included when the window is expanded downward and larger TOF neutrons when expanded upward. The effect of this variation was mostly in the lowest three data points and allowed them to move downwards in scintillation efficiency by about 0.04.

There is an uncertainty associated with fitting the data in a limited range around the predicted single-scattered neutron peak position. To estimate this uncertainty, we expand the fit range to include ± 5 sigma around the centroid of the single-scattered neutrons, instead of the 3-sigma range used in the standard fit. The result of the wider fit range is to systematically push the determined scintillation efficiency up for energies between 20 keVr and 120 keVr. This effect appears to be caused by a disagreement in the high energy tails of the data and Monte Carlo, similar to the disagreement observed between Monte Carlo and data at lower energies for the high recoil angles. Changing the fit range adds a correlated error of +0.01 to the measured mean scintillation efficiency in Fig. 8.

Scattering angle (°)	Energy (keVr)	L_{eff}	+	-
22	11	0.41	0.10	0.08
26	15	0.35	0.07	0.06
30	19	0.28	0.05	0.04
33	22	0.26	0.03	0.03
37	28	0.23	0.03	0.03
40	33	0.25	0.02	0.02
44	39	0.26	0.02	0.03
47	45	0.26	0.02	0.02
51	52	0.27	0.02	0.02
55	59	0.24	0.02	0.02
59	67	0.23	0.02	0.02
62	72	0.25	0.02	0.02
65	79	0.24	0.02	0.02
70	91	0.23	0.02	0.02
73	98	0.24	0.02	0.02
80	114	0.28	0.03	0.04
114	191	0.29	0.02	0.02
125	211	0.26	0.02	0.02
142	239	0.25	0.02	0.01

TABLE I: Table of energies and scintillation efficiencies from Fig. 8. $L_{\rm eff}$ values for energies above 20 keVr also have an additional correlated error of +0.01. The uncertainties shown are the combined statistical and systematic uncertainties.

VII. RESULTS

The scintillation efficiency for nuclear recoils in liquid argon has been measured relative to electronic re-

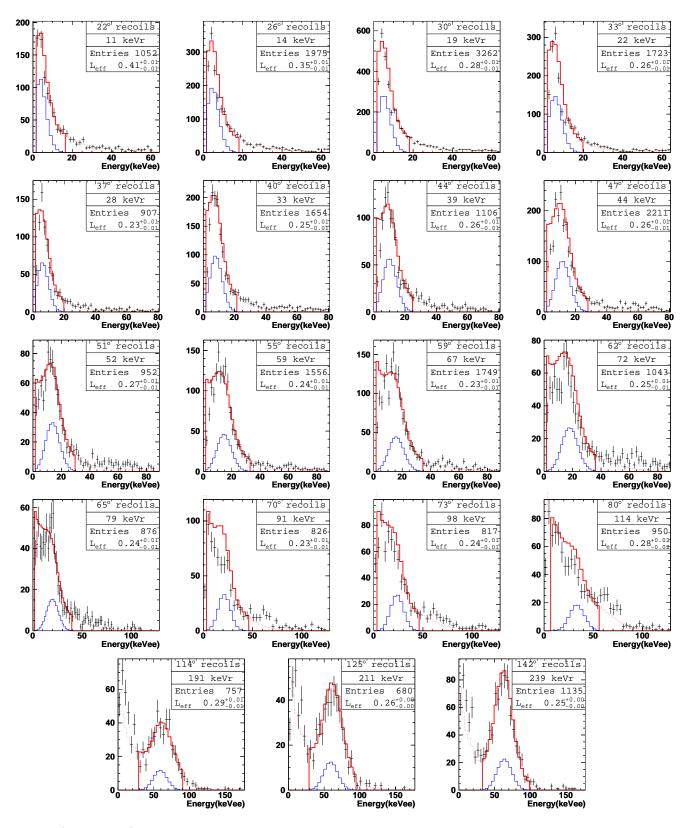


FIG. 7: (Color online) Plotted are the recoil energy spectra for the 19 organic scintillator positions used in this experiment. The data are taken with the organic scintillator located at the angle indicated in the legend of each plot with the corresponding recoil energy indicated just below. The fit value for Leff with the statistical uncertainty from the fit is also listed in the legend of each plot. In each plot, the upper (red) histogram is the output of the GEANT4 based Monte Carlo simulation of single and multiple neutron scatters in the detector. This upper histogram is fit to the data in the solid region, whereas the dotted part shows the MC simulation outside of the fit range. The lower solid histogram (blue) is the subset of the Monte Carlo events where the neutron only scatters once in the detector volume.

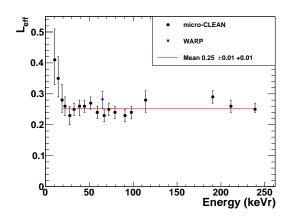


FIG. 8: (Color online) Scintillation efficiency as a function of energy from 10 to 250 keVr. The weighted mean (red line) is generated from the data above 20 keVr and puts the mean scintillation efficiency at 0.25. The value measured by WARP is 0.28 at 65 keVr [3].

coils for nuclear recoil energies from 10 keVr to 250

keVr. The scintillation efficiency values found at each of the recoil angles can be found in Table I and are plotted in Fig. 8. The ratio of the nuclear recoil scintillation response to the electronic recoil response is $0.25 \pm 0.01 + 0.01 (\text{correlated})$ for recoils above 20 keVr. An observed upturn in the scintillation efficiency below 20 keVr is currently unexplained. The scintillation efficiency for nuclear recoils should also have some dependence on any applied electric fields and this is a topic for further research.

VIII. ACKNOWLEDGEMENTS

Computer resources were supplied by Yale University Biomedical High Performance Computing Center and NIH grant: RR19895 and in part by the facilities and staff of the Yale University Faculty of Arts and Sciences High Performance Computing Center. We are also grateful for the support of the U.S. Department of Energy. This work was supported by the David and Lucille Packard Foundation.

- E. Aprile, K. L. Giboni, P. Majewski, K. Ni, M. Yamashita, R. Gaitskell, P. Sorensen, L. DeViveiros, L. Baudis, A. Bernstein, et al., New Ast. Rev. 49, 289 (2005).
- [2] D. B. Cline, Y. Seo, F. Sergiampietri, H. Wang, J. T. White, J. Gao, P. Picchi, G. Mannocchi, L. Periale, F. Pietropaolo, et al., Nucl. Phys. B (Proc. Suppl.) 124, 229 (2003).
- [3] R. Brunetti, E. Calligarich, M. Cambiaghi, F. Carbonara, A. Cocco, C. De Vecchi, R. Dolfini, A. Ereditato, G. Fiorillo, L. Grandi, et al., New Ast. Rev. 49, 265 (2005).
- [4] P. Benetti, R. Acciarri, F. Adamo, B. Baibussinov, M. Baldo-Ceolin, M. Belluco, F. Calaprice, E. Calligarich, M. Cambiaghi, F. Carbonara, et al., arXiv:astroph/0701286 (2007).
- [5] A. Rubbia, J. Phys.: Conf. Ser. 39, 129 (2006).
- [6] G. Jungman, M. Kamionkowski, and K. Griest, Phys. Rept. 267, 195 (1996).
- [7] Z. Ahmed, D. Akerib, S. Arrenberg, C. Bailey, D. Balakishiyeva, L. Baudis, D. Bauer, P. Brink, T. Bruch, R. Bunker, et al. (CDMS) (2009), 0912.3592.
- [8] J. Angle, E. Aprile, F. Arneodo, L. Baudis, A. Bernstein, A. Bolozdynya, P. Brusov, L. Coelho, C. Dahl, L. De-Viveiros, et al., arXiv:0706.0039 (2007).
- [9] A. Hitachi, T. Takahashi, N. Funayama, K. Masuda, J. Kikuchi, and T. Doke, Physical Review B 27, 5279 (1983).
- [10] W. Lippincott, D. McKinsey, J. Nikkel, K. Coakley, A. Hime, L. Stonehill, D. Gastler, and E. Kearns, Phys.Rev.C78:035801 (2008).
- [11] T. Takahashi, S. Konno, T. Hamada, M. Miyajima, S. Kubota, S. Nakamoto, A. Hitachi, E. Shibamura, and T. Doke, Phys. Rev. A 12, 1771 (1975).
- [12] M. Miyajima, T. Takahashi, S. Konno, T. Hamada,

- S. Kubota, H. Shibamura, and T. Doke, Phys. Rev. A 9, 1438 (1974).
- [13] T. Doke, A. Hitachi, J. Kikuchi, K. Masuda, H. Okada, and E. Shibamura, Jpn. J. Appl. Phys. 41, 1538 (2002).
- [14] J. Lindhard, M. Scharff, and H. Schiøtt, Mat. Fys. Medd. K. Dan. Vidensk. Selsk. 33, 1 (1963).
- [15] P. Sorensen et al., Nucl. Instrum. Meth. A601, 339 (2009), 0807.0459.
- [16] E. Aprile et al., Phys. Rev. C79, 045807 (2009), 0810.0274.
- [17] E. Aprile, K. L. Giboni, P. Majewski, K. Ni, M. Yamashita, R. Hasty, A. Manzur, and D. N. McKinsey, Phys. Rev. D 72, 72006 (2005).
- [18] V. Chepel, V. Solovov, F. Neves, A. Pereira, P. J. Mendes, C. P. Silva, A. Lindote, J. Pinto da Cunha, M. I. Lopes, and S. Kossionides, Astropart. Phys. 26, 58 (2006).
- [19] A. Manzur, A. Curioni, L. Kastens, D. McKinsey, K. Ni, and T. Wongjirad, URL http://arxiv.org/abs/0909.1063v4.
- [20] A. Hitachi, T. Doke, and A. Mozumder, Phys. Rev. B 46, 11463 (1992).
- [21] D.-M. Mei, Z. Yin, L. Stonehill, and A. Hime, Astropart. Phys. 30:12-17 (2008).
- [22] A. Hitachi, T. Takahashi, T. Hamada, E. Shibamura, N. Funayama, K. Masuda, J. Kikuchi, and T. Doke, Nucl. Instrum. Meth. 196, 97 (1982).
- [23] C. Regenfus, IDM2006 Proceedings, World Scientific 15, 32 (2007).
- [24] O. Cheshnovsky, B. Raz, and J. Jortner, J. Chem. Phys. 57, 4628 (1972).
- [25] D. N. McKinsey, C. R. Brome, J. S. Butterworth, R. Golub, K. Habicht, P. R. Huffman, S. K. Lamoreaux, C. E. H. Mattoni, and J. M. Doyle, Nucl. Instrum.

Meth. B 132, 351 (1997).

- [26] URL http://geant4.cern.ch/.
- [27] URL http://proj-clhep.web.cern.ch/.
 [28] URL http://root.cern.ch/.

[29] E. Conti, R. DeVoe, G. Gratta, T. Koffas, S. Waldman, J. Wodin, D. Akimov, G. Bower, M. Breidenbach, $R.\ Conley,\ et\ al.,\ Phys.Rev.B68:054201,2003\ (2003).$