

A Search for Short-Wavelength Neutrino Oscillation From a Nuclear Reactor

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ABSTRACT

The Precision Reactor Oscillation and SPECTrum Experiment (PROSPECT) is designed to probe short baseline oscillations of antineutrinos in search of eV-scale sterile neutrinos and precisely measure the ^{235}U reactor antineutrino spectrum from the High Flux Isotope Reactor (HFIR) at Oak Ridge National Laboratory. The PROSPECT antineutrino detector (AD) provides excellent background rejection and position resolution due to its segmented design and use of ^6Li -loaded liquid scintillator. Due to characteristics of its decay chain, ^{227}Ac was added as a calibration source that was dissolved isotropically throughout the liquid scintillator. Using the correlated production of alphas from $^{219}\text{Rn} \rightarrow ^{215}\text{Po} \rightarrow ^{211}\text{Pb}$ in the ^{227}Ac decay chain we can measure the rate of ^{227}Ac in each segment of the detector. This allows us to precisely determine the relative segment to segment volume variation to 1%. These measurements can then be applied as corrections to measurements of neutrino oscillation through the PROSPECT AD.

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These are the acknowledgements, where you express your appreciation to those who were influential and important to your success.

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CHAPTER 1

NEUTRINOS

1.1 Discovery of the Neutrino

The study of radioactive decay in the early 20th century exposed discrepancies that would lead to the postulation and eventual discovery of the neutrino. An experiment performed by Lise Meitner and Otto Hahn in 1911 offered some of the first evidence that the energy spectrum of electrons emitted by beta decay is continuous [1]. This was in stark contrast to the expected discrete spectra that had been observed in gamma and alpha emission, and suggested that three laws of conservation (energy, linear momentum, and angular momentum) were broken during beta decay. Their findings were later confirmed by experiments performed by Chadwick in 1914 [2] and Ellis and Wooster in 1927 [3].

At the time, beta decay was thought to be a two-particle decay, a process that yields a product nucleus and an electron. In 1930, Wolfgang Pauli postulated a particle he called the ‘neutron’, which would be ejected with the electron, thereby conserving energy and momentum. Describing his idea as a “desperate remedy”, this new particle would have to be neutral and non-interacting, therefore making it almost impossible to detect. In 1934, Enrico Fermi further developed the theory of beta decay, including Pauli’s particle but renaming it the neutrino, meaning “little neutral one” [4]. Due to the nature of the weakly interacting neutrino, experimental discovery would take another 20 years.

In 1956 Clyde Cowan and Fred Reines accomplished the amazing feat of discovering this elusive particle experimentally by taking advantage of the inverse beta decay (IBD) process [5]:

$$\bar{\nu} + p \rightarrow n + e^+ \quad (1.1)$$

Their idea was to place a detector near an intense source of neutrinos, fill it with an ample number of protons, and observe the resulting positrons. However, any source that generates a large enough flux of neutrinos will create large backgrounds for the experiment. As would become the challenge for every neutrino detector thereafter, Cowan and Reines had to devise a way to reduce the background such that they could obtain a measurable and believable number of neutrinos. Their original idea was to place a detector underground about 40 meters away from a fission bomb. This would create a large instantaneous flux of neutrinos providing a sufficient signal to background ratio.

After some thought, though, they realized that by detecting the neutron *and* the positron they could discriminate the IBD signal from the background with much higher success. This would allow the use of a nuclear reactor instead of a fission bomb as a neutrino source, giving them the opportunity to patiently watch for neutrinos rather than be restricted to one chance with a bomb. The final detector that would facilitate the discovery of the electron anti-neutrino, $\bar{\nu}_e$, contained 1400 liters of liquid scintillator viewed by 110 photomultiplier tubes and 200 liters of water with dissolved cadmium chloride, and was placed near the fission reactor at the Savannah River Plant in South Carolina.

The mechanisms behind the detector developed by Cowan and Reines worked as follows. Neutrinos from the reactor enter the detector and interact with protons in the water. The positron resulting from this reaction quickly collides with an electron, creating two gamma rays which Compton scatter and initiate a cascade of electrons that causes the liquid to scintillate. Simultaneously, the neutron from the initial reaction bounces around as it collides with protons until, eventually, it captures on a cadmium nucleus and releases about a 9 MeV gamma ray, also causing the liquid to scintillate. The time between the flash of light from the positron annihilation and that of the neutron capture is on the order of microseconds. It was by looking for this delayed-coincidence signature that Cowan and Reines were able to successfully detect the first neutrino, and thus pave the way for future neutrino experiments.

Only 6 years after the discovery by Cowan and Reines, Lederman, Schwartz, and Steinberger discovered the muon neutrino, ν_μ [6], but the discovery of the tau

	Fermions			Bosons	
	spin = 1/2			spin = 1	spin = 0
Generation	I	II	III	Gauge Bosons	Scalar Bosons
Quarks	u	c	t	g	H
	d	s	b	γ	
Leptons	e	μ	τ	Z	W
	ν_e	ν_μ	ν_τ	W	

Table 1.1: The Standard Model of particle physics, composed of fermions and their corresponding antiparticles, the force carries (gauge bosons), and the Higgs boson.

neutrino, ν_τ , by the DONUT (Direct Observation of the Nu Tau) experiment would not occur until 44 years later [7, 8].

1.2 Neutrinos in the Standard Model and Beyond

The Standard Model (SM) of particle physics is the result of several decades of work by many scientists. It is a field theory that describes three of the four fundamental forces (electromagnetic, strong and weak), and classifies all known elementary particles as outlined in Table 1.1.

Classified in three generations of leptons, neutrinos exist in corresponding flavors, electron neutrinos (ν_e), muon neutrinos (ν_μ), and tau neutrinos (ν_τ), to the electron (e), muon (μ), and tau (τ) leptons, respectively. For each of these flavors a corresponding antiparticle also exists, $\bar{\nu}_e, \bar{\nu}_\mu, \bar{\nu}_\tau$.

If neutrinos have exactly zero mass and travel at the speed of light, then by definition their helicity, or handedness, is a conserved property. Since experiments only measured left-handed neutrinos [9], it was assumed that all neutrinos in the SM were massless. As will be discussed in greater depth in Section 1.3, it was later experimentally discovered that neutrinos oscillate, or change flavors, indicating that at least two of the three neutrinos must have mass.

Their ability to oscillate arises from the fact that neutrinos exist as flavor eigenstates of the weak interaction, ν_α : $\alpha = e, \mu, \tau$, that are not identical to their mass states, ν_i : $i = 1, 2, 3$, which are eigenstates of the Hamilton describing the free neutrino. This means that a neutrino produced at a source has a known flavor, but its mass is described as a superposition of all possible mass eigenstates, creating three possible scenarios: (i) The mass states are the same as the flavor states, therefore,

a neutrino produced as a ν_x will remain ν_x forever. (ii) The mass states and flavor states are different, but the mass states are all the same or all zero, therefore, a neutrino produced as ν_x will still remain ν_x because the combination of mass states does not change over time. (iii) The mass states and flavor states are different and at least two of the mass states are not the same, therefore, the phase between mass states will change as the neutrino propagates through time and space causing a change of flavor state. Experimental discoveries of neutrino oscillation require that the third scenario is correct.

Given this we can define the probability that a neutrino oscillates by first describing the flavor eigenstates as:

$$|\nu_\alpha\rangle = \sum_i U_{\alpha i} |\nu_i\rangle \quad (1.2)$$

where $U_{\alpha i}$ is the unitary Pontecorvo-Maki-Nakagawa-Sakata (PMNS) mixing matrix [10]. The PMNS matrix relates flavor to mass eigenstates and can be written in factorized form as:

$$U = \begin{pmatrix} 1 & & \\ c_{23} & s_{23} & \\ -s_{23} & c_{23} & \end{pmatrix} \begin{pmatrix} c_{13} & s_{13}e^{-i\delta} \\ -s_{13}e^{i\delta} & c_{13} \end{pmatrix} \begin{pmatrix} c_{12} & s_{12} \\ -s_{12} & c_{12} \end{pmatrix} \begin{pmatrix} 1 & e^{i\alpha} \\ & e^{i\beta} \end{pmatrix} \quad (1.3)$$

where $s_{ij} = \sin \theta_{ij}$ and $c_{ij} = \cos \theta_{ij}$, $\theta_{ij} = [0, \pi/2]$, $\delta = [0, 2\pi]$ is the Dirac charge parity (CP) violation phase, and α and β are two Majorana CP violation phases.

As a neutrino, ν_α , propagates in time the mass eigenstates evolve differently (assuming that $m_1 \neq m_2 \neq m_3$), resulting in a new flavor state, ν_β . Massive neutrinos move through time and space as

$$|\nu_i(x, t)\rangle = e^{-\frac{i}{\hbar}(E_i t - \vec{p}_i \cdot \vec{x})} |\nu_i(0, 0)\rangle = e^{-i\phi} |\nu_i(0, 0)\rangle \quad (1.4)$$

The flavor state α at some point in time and space can then be defined as:

$$|\nu_\alpha(x, t)\rangle = \sum_i U_{\alpha i} |\nu_i(x, t)\rangle = \sum_i U_{\alpha i} e^{-i\phi_i} |\nu_i(0, 0)\rangle \quad (1.5)$$

Therefore, the oscillation probability that a neutrino produced as flavor ν_α will be

detected as flavor ν_β after traveling for a period of time is given by

$$\begin{aligned}
P(\nu_\alpha \rightarrow \nu_\beta) &= |\langle \nu_\alpha(0,0) | \nu_\beta(x,t) \rangle|^2 \\
&= \left| \sum_i U_{\alpha i}^* e^{-i\phi_i} U_{\beta i} \right|^2 \\
&= \sum_i \sum_k U_{\alpha i}^* U_{\beta i} U_{\alpha k} U_{\beta k}^* e^{-i(\phi_i - \phi_k)}
\end{aligned} \tag{1.6}$$

This is true for any number of neutrino generations, but for the sake of simplicity, consider the case of two neutrino oscillation. In this scenario the mixing matrix can be written as

$$U = \begin{pmatrix} U_{\alpha 1} & U_{\alpha 2} \\ U_{\beta 1} & U_{\beta 2} \end{pmatrix} = \begin{pmatrix} \cos \theta_{12} & \sin \theta_{12} \\ -\sin \theta_{12} & \cos \theta_{12} \end{pmatrix} \tag{1.7}$$

Therefore, the probability of oscillation is given by

$$\begin{aligned}
P^{2\nu}(\nu_\alpha \rightarrow \nu_\beta) &= |U_{\alpha 1}|^2 |U_{\beta 1}|^2 + |U_{\alpha 2}|^2 |U_{\beta 2}|^2 + U_{\alpha 1}^* U_{\beta 1} U_{\alpha 2} U_{\beta 2}^* (e^{i(\phi_2 - \phi_1)} + e^{-i(\phi_2 - \phi_1)}) \\
&= \sin^2 2\theta_{12} \sin^2 \left(\frac{\phi_2 - \phi_1}{2} \right)
\end{aligned} \tag{1.8}$$

Now, recall that

$$\phi_i = \frac{1}{\hbar} (E_i t - \vec{p}_i \cdot \vec{x}) \tag{1.9}$$

The mass of the neutrino is very small compared to its energy ($m_\nu \ll E_\nu$) so the momentum can be approximated as

$$p_i = \frac{1}{c} \sqrt{E_i^2 - m_i^2 c^4} = \frac{1}{c} \left(E_i - \frac{m_i^2 c^4}{2E_i} \right) \tag{1.10}$$

If it is reasonably assumed that neutrinos move at the speed of light, c , then the phase difference, $\phi_2 - \phi_1$, can be approximated as

$$\begin{aligned}
\phi_2 - \phi_1 &= \frac{1}{\hbar} \left((E_2 - E_1) \frac{L}{c} - (p_2 - p_1)L \right) \\
&= \frac{1}{\hbar} \frac{L}{c} \left(\frac{m_2^2 c^4}{2E_2} - \frac{m_1^2 c^4}{2E_1} \right) \\
&= \frac{L}{\hbar c} \frac{\Delta m_{21}^2 c^4}{2E}
\end{aligned} \tag{1.11}$$

where $t = \frac{L}{c}$, $\Delta m_{21}^2 = m_2^2 - m_1^2$ and $E_1 = E_2 = E$.

It can now be shown that the oscillation probability of a neutrino ν_α , being detected as flavor ν_β in the two neutrino mixing case is

$$P^{2\nu}(\nu_\alpha \rightarrow \nu_\beta) = \sin^2(2\theta_{12}) \sin^2\left(\frac{c^4}{4\hbar c} \frac{\Delta m_{12}^2 L}{E}\right) \quad (1.12)$$

The corresponding survival probability, the chance that a neutrino ν_α is detected as ν_α , can be described by $P^{2\nu}(\nu_\alpha \rightarrow \nu_\alpha) = 1 - P^{2\nu}(\nu_\alpha \rightarrow \nu_\beta)$.

There are several aspects of note about this probability. It can be seen that the amplitude of the oscillation probability, $\sin^2(2\theta_{12})$, depends on the mixing angle θ_{12} , while the mass splitting, Δm_{12}^2 , the energy of the neutrino, E , and the distance traveled, L , determine the frequency of oscillation. The probability is non-zero only when Δm_{12}^2 is non-zero, indicating that if an experiment observes neutrino oscillation, then at least one of the neutrinos must have mass. Finally, the dependence of oscillation on the factor $\frac{L}{E}$ allows experiments to decide the placement of neutrino detectors based on what features of neutrinos they would like to study. Theoretical models alone do not prove neutrino oscillation, however. The first experimental evidence for oscillation and neutrino mass will be described in Section 1.3.

1.3 Discovery of Neutrino Oscillation

In the late 1960's, about a decade after Cowan and Reines discovered the first neutrino, astrophysicist John Bahcall and physical chemist Raymond Davis designed an experiment to collect and count solar neutrinos, neutrinos emitted by nuclear fusion taking place in the Sun. Davis placed a 380 cubic meter tank filled with perchloroethylene (dry-cleaning fluid) 1,478 meters underground in the Homestake Gold Mine in South Dakota. Perchloroethylene was chosen because it is rich in chlorine and the tank was placed deep underground to shield the experiment from cosmic rays.

Davis was looking for the reaction



in which a neutrino would enter the tank and transform chlorine into argon. Argon is a noble gas, making it easy to chemically separate from a large amount of chlorine-rich solvent. This allowed Davis to extract and count the argon atoms, essentially counting the number of neutrinos that had been captured. The chlorine reaction has

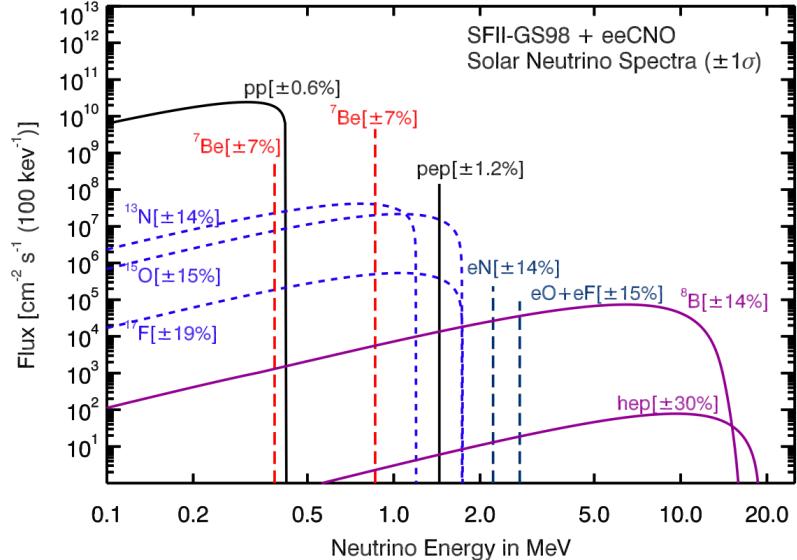


Figure 1.1: Spectrum of solar neutrino fluxes corresponding to the SFII-GS98 standard solar model. Electron capture CNO neutrinos (ecCNO) have been added in addition to standard fluxes. Electron capture fluxes are given in $\text{cm}^{-2}\text{s}^{-1}$. Figure from Ref. [11].

a threshold of 814 keV, which, according to standard solar models (see Figure 1.1), allowed the Homestake experiment to detect neutrinos created by ${}^8\text{Be}$ in the sun (and ${}^7\text{Be}$, ${}^{13}\text{N}$, and ${}^{15}\text{O}$ at lower rates).

In the end, with the Homestake experiment Davis calculated a rate of solar neutrinos that was one third of the rate predicted by calculations made by Bahcall using the Standard Model [12]. This discrepancy became known as the solar neutrino problem, and in the following years Bruno Pontecorvo wrote several theoretical papers proposing neutrino oscillation as a solution [13, 14].

Several experiments designed to measure the flux of solar neutrinos followed the Homestake Experiment including SAGE [15], GALLEX [16], and GNO [17, 18]. All three experiments built detectors based on the reaction ${}^{71}\text{Ga}(\nu, e^-){}^{71}\text{Ge}$ which has a threshold of 233 keV. This made them sensitive to neutrinos created by pp reactions in the Sun, neutrinos that the Homestake experiment was unable to measure. Even with the different energy threshold, though, all three experiments showed a deficit in neutrino flux compared to Standard Model calculations and the solar neutrino problem persisted.

Further experimental proof of neutrino oscillation came with results from the

Super-Kamiokande Experiment. Unlike previous experiments that were only sensitive to electron neutrinos, Super-K detected neutrinos through elastic scattering of electrons - a process sensitive to all neutrino flavors. Operating a high threshold of 4 MeV they observed pure ${}^8\text{Be}$ solar neutrinos, a different selection than previous solar neutrino experiments. With their large mass, good energy resolution, and ability to determine neutrino directionality, Super-K was able to confirm the solar neutrino problem effect with high statistics and place limits on the parameters of oscillation [19].

The first direct evidence for solar neutrino flavor change came from the Sudbury Neutrino Observatory (SNO) in 2001 [20, 21]. The SNO detector was an imaging Cherenkov detector using heavy water. They were able to observe neutrino flavor change through three different processes: elastic scattering of electrons, the $\nu_e - d$ charged current reaction (CC), and the $\nu_x - d$ neutral current reaction (NC). The CC reaction has a threshold of -1.4 MeV, allowing SNO to accurately measure the shape of the ${}^8\text{B}$ neutrino spectrum. The CC is also only sensitive to ν_e , whereas the NC reaction is sensitive to all three active neutrino flavors, allowing SNO to compare measured fluxes from each reaction and ultimately confirm the theory of neutrino oscillation.

Nearly a century after the initial postulation of the neutrino the scientific community has in hand experimental evidence of three neutrino flavors, mathematical models that include these neutrinos in the Standard Model of Particle Physics, and theoretical and experimental proof that neutrinos oscillate and therefore have mass. These findings, along with developments in technology and techniques, set the stage for current, and future, neutrino experiments to transform their goals from observing anomalies to making precise measurements of the physics behind the anomalies. The current best-fit values of the 3-neutrino oscillation parameters as found experimentally are shown in Table 1.2.

Parameter	Best-fit	3σ
Δm_{21}^2 [10 ⁻⁵ eV ²]	7.37	6.93 - 7.96
$\Delta m_{31(23)}^2$ [10 ⁻³ eV ²]	2.56 (2.54)	2.45 - 2.69 (2.42 - 2.66)
$\sin^2 \theta_{12}$	0.297	0.250 - 0.354
$\sin^2 \theta_{23}$	0.425 (0.589)	0.381 - 0.615 (0.384 - 0.636)
$\sin^2 \theta_{13}$	0.0215 (0.0216)	0.0190 - 0.0240 (0.0190 - 0.0242)
δ/π	1.38 (1.31)	2 σ : 1.0 - 1.9 (2 σ : 0.92 - 1.88)

Table 1.2: The current best-fit values and 3σ allowed ranges of the 3-neutrino oscillation parameters as determined experimentally [10]. The values (values in brackets) correspond to $m_1 < m_2 < m_3$ ($m_3 < m_1 < m_2$).

CHAPTER 2

REACTOR NEUTRINOS

Nuclear reactors are a pure source of electron antineutrinos, $\bar{\nu}_e$, as a result of the fission of isotopes used in the reactor fuel. The first neutrino was discovered using the nuclear reactor at the Savannah River Plant, and reactor sites continue to be popular homes for neutrino detectors. In order to perform precision reactor neutrino studies it is important to understand the reactor neutrino flux and spectrum.

2.1 Production of Reactor Neutrinos

Nuclear reactors are powered by the fission of uranium and plutonium isotopes in their cores. Specifically, in a power reactor, 99.9% of the power comes from the fission of ^{235}U , ^{239}Pu , ^{241}Pu , and ^{238}U isotopes. The reaction begins with a neutron colliding with a nucleus of one of the isotopes. This causes the nucleus to split into two fragments, usually of unequal mass, creating an unstable system. In order to reach stability neutrons have to transform into protons, a process accomplished through β decay, see Figure 2.1. Each beta decay produces an electron and corresponding electron antineutrino. In general a nuclear reactor will produce $\sim 6 \times 10^{20} \bar{\nu}_e$ per GW of thermal energy power [22].

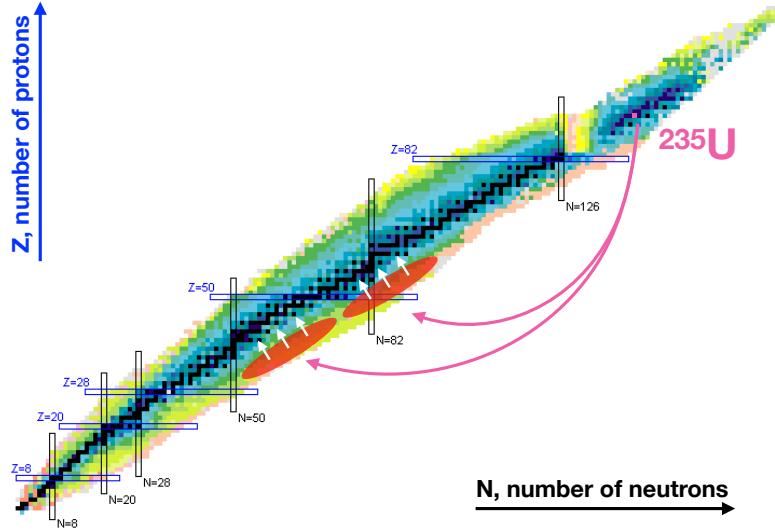


Figure 2.1: A schematic of the fission of ^{235}U [23]. After collision with a neutron ^{235}U will split into two unstable nuclei (pink arrows) which will then β decay (white arrows) until stable.

2.2 Measuring the Reactor Antineutrino Flux and Spectrum

The total $\bar{\nu}_e$ flux, $S(E_\nu)$, produced by a nuclear reactor can be expressed as the sum over the spectra of the dominant fissioning isotopes,

$$S(E_\nu) = \frac{W_{th}}{\sum_i(f_i/F)e_i} \sum_i \frac{f_i}{F} \left(\frac{dN_i}{dE_\nu} \right), \quad (2.1)$$

where f_i/F is the fission fraction for each given isotope i , W_{th} is the reactor thermal energy, e_i is the average energy released per fission by each isotope, and dN_i/dE_ν is the cumulative $\bar{\nu}_e$ spectrum of i normalized per fission.

There are two methods used to determine the $\bar{\nu}_e$ spectrum, *ab initio* summation and electron spectrum conversion. In the *ab initio* approach the spectrum is determined by summing the contributions of all β -decay branches of all fission fragments,

$$\frac{dN_i}{dE_{\bar{\nu}}} = \sum_n Y_n(Z, A, t) \sum_{n,i} b_{n,i}(E_0^i) P_{\bar{\nu}}(E_{\bar{\nu}}, E_0^i, Z), \quad (2.2)$$

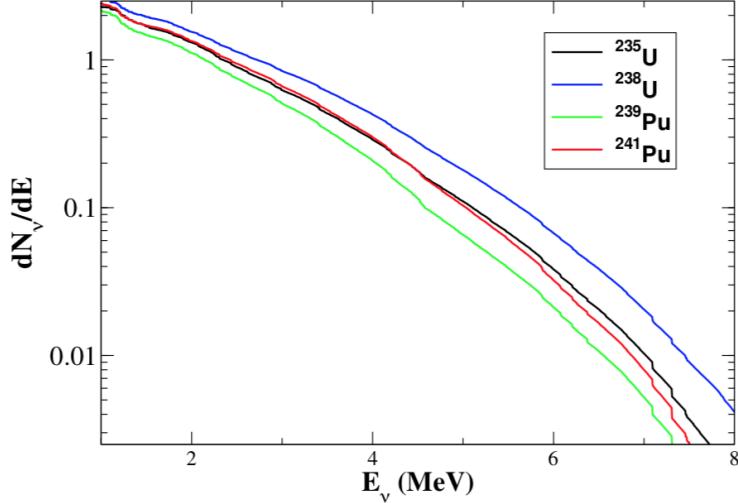


Figure 2.2: The $\bar{\nu}_e$ spectrum predicted by the summation method using the JEFF-3.1.1 database fission fragment yields and the ENDF/B-VII.1 decay library [22].

where $Y_n(Z, A, t)$ is the number of β decays of the fission fragment Z, A at time t , $b_{n,i}(E_0^i)$ are the branching ratios with endpoint energies E_0^i , and $P_{\bar{\nu}}(E_{\bar{\nu}}, E_0^i, Z)$ is the normalized $\bar{\nu}_e$ spectrum for the branch n, i . This method relies on nuclear databases, such as the Evaluated Nuclear Data File (ENDF) and Joint Evaluated Fission and Fusion (JEFF) databases, for information about the branching ratios and decay energies. The antineutrino spectrum for the four main reactor isotopes calculated using *ab initio* summation was done in Ref. [22] and the result can be seen in Figure 2.2.

Though seemingly straightforward, this approach comes with some caveats. The shear number of daughter isotopes (>1000) and individual β decay branches (>6000) make the summation non-trivial. This, along with the fact that not all branching ratios are known, and that the fission yields have been determined by several different database groups but don't always agree and have large uncertainties bring into question the validity of using only this method.

The other approach to determine the $\bar{\nu}_e$ spectrum, the conversion method, relies on converting a measured electron spectrum into an antineutrino spectrum. This involves fitting an experimentally defined total beta spectrum with individual beta spectrum according to their amplitudes, a_i ,

$$\frac{dN_i}{dE_e} = \sum_i a_i P(E, E_0^i, Z) \quad (2.3)$$

The conversion to the antineutrino spectrum is then accomplished by replacing the energy E_e in each branch by $E_0 - E_{\bar{\nu}}$, because the electron and the $\bar{\nu}_e$ share the total energy of each β -decay branch. The flux per fission is then given as the sum of $\bar{\nu}_e$ spectrum converted from each virtual β branch,

$$\frac{dN_i}{dE_{\bar{\nu}}} = \sum_i a_i P(E_0^i - E, E_0^i) \quad (2.4)$$

The electron spectra for ^{235}U , ^{239}Pu , and ^{241}Pu were measured at the Institut Laue-Langevin (ILL) reactor in Grenoble, France in the 1980s [24, 25, 26], while the spectrum of ^{238}U was more recently (2014) measured at the neutron source FRMII in Garching, Germany [27]. The ILL measurements, along with a prediction of the $^{238}\text{U} \bar{\nu}_e$ spectrum using the summation method by Vogel [28], became known as the “ILL-Vogel” flux model and was the main model used until 2011.

In 2011 Mueller *et al.* improved the prediction of the reactor antineutrino spectra by employing a method that combined information from the nuclear databases and the measured electron spectra from ILL [29]. This was followed by a further improvement by Huber who applied higher order corrections making use of the conversion method and minimizing the use of the databases as much as possible [30].

Though much work has been done to accurately model the reactor antineutrino spectra both methods are subject to uncertainties in the subdominant corrections to beta-decay. This includes radiative, weak magnetism, and finite size corrections along with uncertainties in the spectrum shape of forbidden transitions which are summarized in Ref.[22]. Besides the model uncertainties there are also experimental uncertainties that arise from knowing the thermal power of the reactor, its time-dependent fuel composition, and the fission energies of the dominant isotopes. All of these uncertainties result in a 10-20% relative uncertainty on the reactor antineutrino spectra using the *ab initio* method and $\sim 5\%$ uncertainty on the conversion approach [31].

2.3 Detection of Reactor Neutrinos

Though there are several methods that can be used to detect reactor neutrinos, including charge-current ($\bar{\nu}_e + d \rightarrow n + n + e^+$), neutral-current ($\bar{\nu}_e + d \rightarrow n + p + \bar{\nu}_e$), and antineutrino-electron elastic scattering ($\bar{\nu}_e + e^- \rightarrow \bar{\nu}_e + e^-$), the one employed by

most experiments is IBD ($\bar{\nu}_e + p \rightarrow e^+ + n$). The IBD reaction energy threshold is 1.8 MeV and the cross section is relatively high, $\sim 63 \times 10^{-44} \text{cm}^2/\text{fission}$ integrated over the entire reactor neutrino energy spectrum [31], and can be written as

$$\sigma^{(0)} \simeq 9.52 \times \left(\frac{E_e^{(0)} p_e^{(0)}}{\text{MeV}^2} \right) \times 10^{-44} \text{cm}^2 \quad (2.5)$$

where E_e and p_e are the energy and momentum of the final-state positron.

An IBD event is selected by a pair of coincident signals consisting of a positron ionization and annihilation as the prompt signal and a time delayed neutron capture on a proton or nucleus as the delay signal. The neutrino energy can be backtracked from the prompt signal as

$$E_{\bar{\nu}} = E_{prompt} + 0.78 \text{ MeV} + T_n \quad (2.6)$$

where T_n is the kinetic energy of the recoil neutron which is much smaller than the energy of the neutrino and can therefore be ignored in most cases. The IBD cross-section increases with energy, whereas the $\bar{\nu}_e$ spectrum decreases with energy creating a detected energy spectrum that peaks around 3.8 MeV and dies off after ~ 8 MeV, as seen in Figure 2.3.

In addition to great background rejection and good reconstruction of the neutrino energy, the IBD method of detecting neutrinos also allows the use of liquid scintillators and water as detection mediums.

2.4 The Reactor Antineutrino Anomaly

Several experiments have, and continue to use reactor antineutrinos as a probe of neutrino oscillation. A reactor neutrino disappearance $P(\bar{\nu}_e \rightarrow \bar{\nu}_e)$ experiment located at a distance $L \sim 1$ km can measure $\sin^2 \Delta_{31}$. At that baseline, the amplitude of the oscillation at the first maximum of $\sin^2 \Delta_{31}$ is $\sin^2 2\theta_{13}$ (recall Eq. 1.12), providing a direct measurement of θ_{13} . Three experiments were designed to make a measurement of this mixing angle, Daya Bay in China, RENO in Korea, and Double Chooz in France.

The Daya Bay Reactor Neutrino Experiment was located at the Daya Bay nuclear reactor power plant in southern China that consists of six 2.9 GW_{th} reactors. They

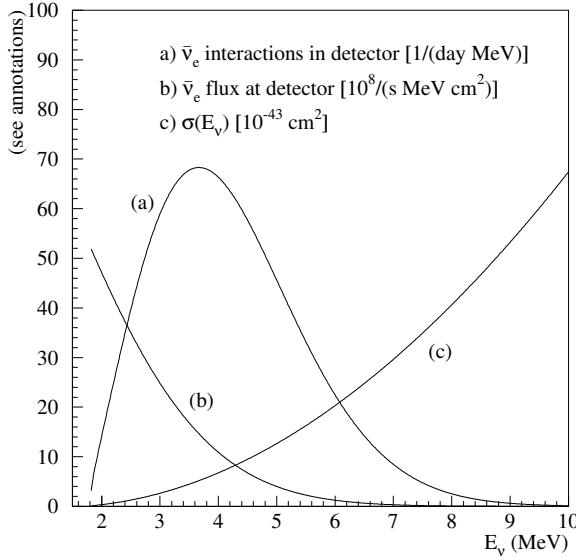


Figure 2.3: The IBD spectrum (curve (a)) measured by a 12-ton fiducial mass detector located 0.8 km from a 12-GW_{th} power reactor along with the reactor flux (curve (b)) and IBD cross section (curve (c)) as a function of energy [10].

employed two groups of near (512 m, 561 m) and one group of far (1,579 m) antineutrino detectors (AD) in order to suppress the reactor flux uncertainties [32, 33]. The IBD yield measured for each AD is shown in Figure 2.4 and it can be seen that, after correcting for small variations of fission fractions among the different sites, all rates are consistent with each other. Though results between detectors agree, the disagreement between the experimental results and most recent model calculations (Huber+Mueller) is troubling.

In order to obtain a wider picture, the Daya Bay average IBD yield at the flux-weighted baseline (573 m) of the two near detector sites was compared to measurements from nineteen other short-baseline (<1000 m) experiments as shown in Figure 2.5. The global average, including the most recent Daya Bay calculation, results in a ratio of measured to expected yield of 0.945 ± 0.007 (exp.) ± 0.023 (model) with respect to the Huber+Mueller model, a $\sim 6\%$ deficit [34]. If the model uncertainty is to be trusted this ratio suggests reactor $\bar{\nu}_e$ disappearance as close as $L < 10$ m, a phenomenon not covered in the standard 3-flavor neutrino mixing model [22]. This predicament has been labeled the “reactor antineutrino anomaly” (RAA).

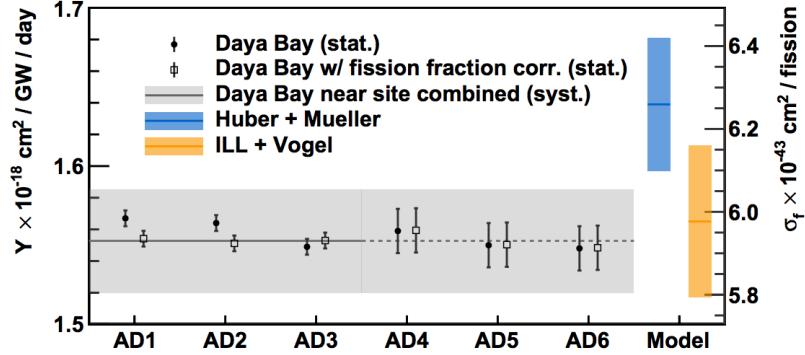


Figure 2.4: Rate of reactor antineutrino candidate events in Daya Bay’s six detectors [33]. The average of the three near detectors is shown as the gray line, extended though the far detectors as a dotted gray line. Also shown are the rates predicted using the Huber+Mueller (blue) and ILL+Vogel (orange) models.

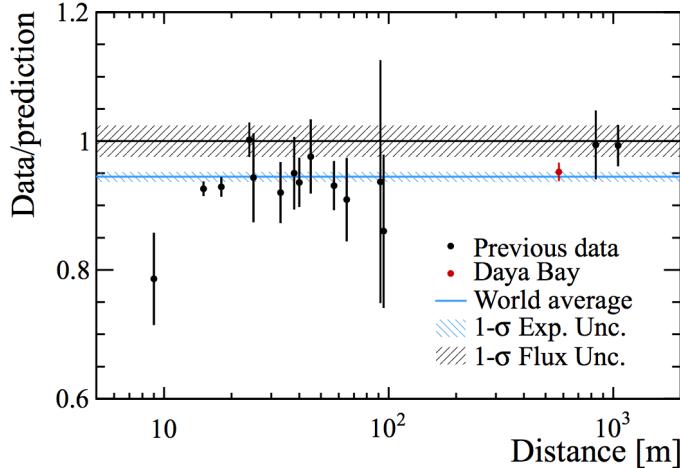


Figure 2.5: The measured reactor $\bar{\nu}_e$ rate, normalized to the Huber+Mueller model prediction, as a function of distance from the reactor [34]. The rate is corrected for 3-flavor neutrino oscillations at each baseline. The blue shaded region represents the global average and its 1σ uncertainty. The 2.7σ model uncertainty is shown as a band around unity.

One hypothesis for explaining the reactor anomaly is that reactor neutrinos are oscillating into a new type of neutrino, a sterile neutrino. A sterile neutrino is a right-handed neutrino that does not take part in weak interactions except those induced by mixing with active neutrinos [35]. Evidence for sterile neutrinos has also been observed in non-reactor neutrino experiments. Specifically, the Liquid Scintillation Neutrino Detector (LSND) measured an excess of $\bar{\nu}_e$ ($>3\sigma$) events [36] along with

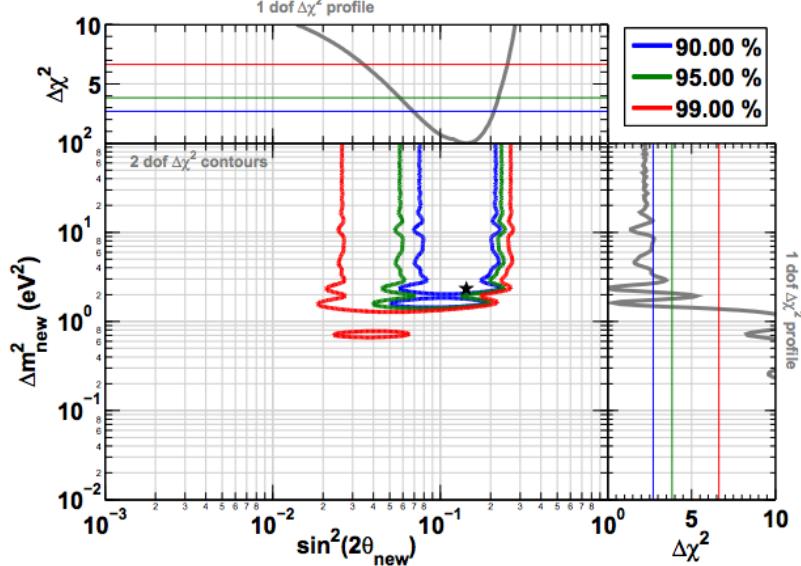


Figure 2.6: Allowed regions in the $\sin^2(2\theta_{14}) - \Delta m_{14}^2$ plane resulting from a fit of the 3+1 neutrino model to results from reactor neutrino experiments, SAGE and GALLEX, MiniBooNE, and spectrum measurements from ILL. This global fit results in the constraints $\Delta m_{14}^2 > 1.5 \text{ eV}^2$ and $\sin^2(2\theta_{14}) = 0.14 \pm 0.08$ [39].

excesses measured by the Mini Booster Neutrino Experiment (MiniBooNE) of ν_e (3.4σ) and $\bar{\nu}_e$ (2.8σ) [37]. Two solar neutrino detectors, the SovietAmerican Gallium Experiment (SAGE) and Gallium Experiment (GALLEX), have observed a deficit in electron neutrinos produced by intense artificial ^{51}Cr and ^{37}Ar radioactive sources at a significance of 3σ [38].

All of these anomalies hint at short-baseline oscillations of electron neutrinos. The simplest way to explain these discrepancies is to add onto the standard model using the 3+1 oscillation model in which there are three active neutrinos and one sterile. This would introduce three new mixing angles, θ_{14} being the one of interest in reactor neutrino experiments. A global fit of this model to neutrino data, including results from reactor experiments, SAGE and GALLEX, MiniBooNE, and spectrum measurements from ILL, result in oscillation constraints $\Delta m_{14}^2 > 1.5 \text{ eV}^2$ and $\sin^2(2\theta_{14}) = 0.14 \pm 0.08$, as shown in Figure 2.6 [39].

An alternative explanation is that the flux predictions are incorrect and have larger uncertainties than those currently applied. The idea that the calculated reactor antineutrino flux is not well understood is bolstered by results from Daya Bay [33], RENO [40], and Double Chooz [41] in which a “bump” was observed in the

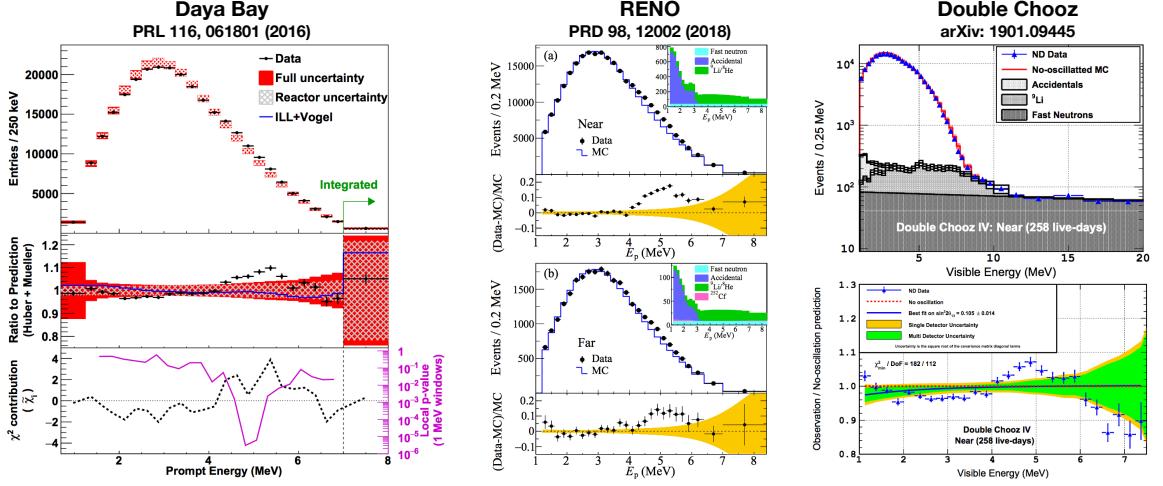


Figure 2.7: A comparison between the predicted and measured prompt energy spectra of IBD events in Daya Bay [33], RENO [40], and Double Chooz [41]. All experiments observe an excess of events above uncertainty in the model spectrum in the 4-6 MeV region.

experimentally measured antineutrino energy spectrum relative to the model spectrum, shown in Figure 2.7. Whether either or both hypothesis are true, and solve the reactor antineutrino anomaly, is a topic of great interest in the neutrino community and several experiments have been designed to address this matter.

CHAPTER 3

PROSPECT

The scientific community’s understanding of neutrinos has come a long way from Pauli’s initial proposition of its existence in 1930. Though three active neutrino flavors and their behaviors are understood and well included in the Standard Model of particle physics, recent anomalies in reactor neutrino experiment results hint at the possibility of new physics. The discovery of an eV-scale sterile neutrino would have wide ranging impacts on the field of neutrino physics and future experiments.

The Precision Reactor Oscillation and Spectrum Experiment (PROSPECT) is designed to address the reactor antineutrino anomaly by performing a reactor-model independent search for short-baseline $\bar{\nu}_e$ oscillations and making a high precise measurement of the ^{235}U $\bar{\nu}_e$ energy spectrum at a highly-enriched uranium (HEU) research reactor [42]. Located at the High Flux Isotope Reactor (HFIR) at Oak Ridge National Laboratory (ORNL) in Tennessee PROSPECT also demonstrates successful application of techniques for antineutrino detection at the surface with little overburden. PROSPECT collected data from May to December of 2018 and the first oscillation and spectrum results, with 33 and 40.3 live-days of reactor on time respectively, can be found in Ref.[43, 44].

3.1 Experimental Site

3.1.1 HFIR

HFIR is a compact research reactor that burns highly enriched uranium fuel (^{235}U), meaning that $> 99\%$ of fissions during a reactor cycle will be from ^{235}U . The HFIR

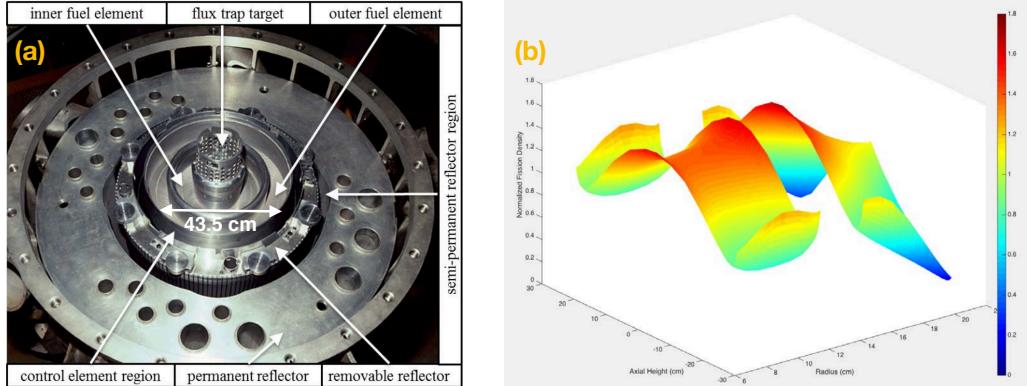


Figure 3.1: (a) The HFIR core, showing the inner and outer fuel elements and the flux trap region, as well as the control elements and Beryllium reflectors. (b) The relative fission density distribution at the start of a cycle. [45]

core consists of two concentric fuel elements with an outer diameter of 0.435 m and a height of 0.508 m, surrounded by control elements and Beryllium reflectors as shown in Figure 3.1. The reactor typically operates at 85 MW for seven 24-day cycles per year for a duty cycle of $\sim 46\%$.

3.1.2 Backgrounds at HFIR

The PROSPECT detector is located at ground level ~ 7 m from the HFIR core and separated from the reactor water pool by a 1 m thick concrete wall, show in Figure 3.2. The proximity to the reactor and lack of overburden introduces a significant level of background events from the reactor and cosmogenic sources. These events can be classified into two categories, (i) singles that are mainly due to gammas and (ii) coincident events from neutron recoil and captures. Extensive studies on the types and rate of background events at the detector site can be found in Refs.[46, 47, 48].

The largest source of gamma backgrounds was discovered to originate in the reactor pool wall, specifically an unused beam line that lies directly in front of the detector. In order to lower these backgrounds a lead shield wall (3.0 m wide, 2.1 m tall, and on average 0.10 m thick), along with shorter flanking walls on each side and a mini-wall placed at the opening of the beam line, was installed between the pool wall and the detector. Other background events were shown to come from neutron beam-lines and scattering experiments existing below the detector site, but most of these are suppressed by a concrete monolith that the detector sits on.

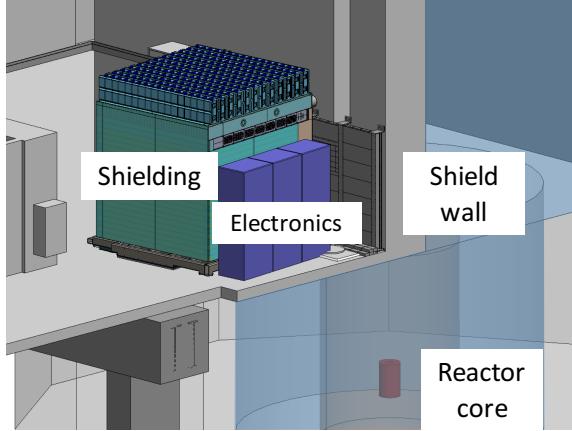


Figure 3.2: Layout of the PROSPECT experiment. The detector is installed in the HFIR Experiment Room next to the water pool and 5 m above the HFIR reactor core (red). The floor below contains multiple neutron beam-lines and scattering experiments.

The thermal neutron rate was measured to be $\sim 2/\text{cm}^2/\text{s}$ during reactor operation [46], so layers of shielding containing ^{10}B , which has a large thermal neutron cross-section and minimal gamma emission, were used in the passive shielding that surrounds the detector (see Section 3.2.2). The locally installed shield wall, along with the addition of passive shielding, resulted in a sufficient suppression of background such that a better than one-to-one signal to background ratio was achieved.

3.2 Design

The PROSPECT antineutrino detector (AD) consists of a segmented inner detector filled with ^6Li doped liquid scintillator (LiLS), contained in an acrylic and aluminum tank, and surrounded by layers of passive shielding. The active detector is made up of a 14×11 array of optically separated segments, viewed on each end by a photomultiplier tube (PMT) enclosed in an acrylic housing. The AD is placed with the segments parallel to the reactor pool wall, ~ 7 m from the reactor core, and measures ~ 3 m tall including all shielding. See Figure 3.3 for a schematic of the detector.

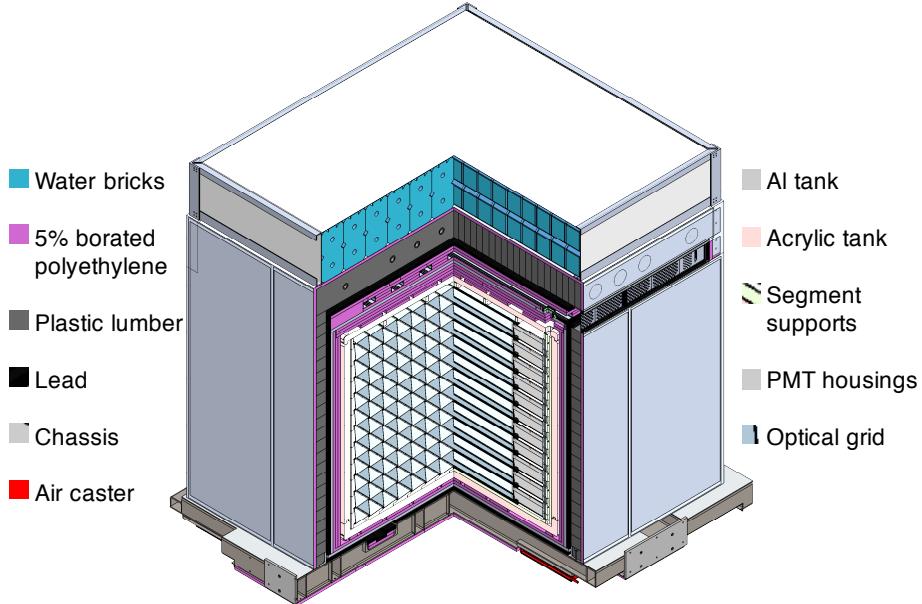


Figure 3.3: A cutaway view of the PROSPECT detector, including the inner detector, outer containment vessels, and passive shielding.

3.2.1 Active Detector

3.2.1.1 PMT Housings

A total of 308 PMTs are installed in the AD; 240 Hamamatsu R6594 SEL PMTs used in the inner segments (fiducial volume) and 68 ADIT Electron Tubes 9372KB PMTs used in the outer segments as shown in Figure 3.4. Each PMT is mounted inside a rectangular acrylic housing facing a clear 144-mm-square front window constructed from ultraviolet transmitting (UVT) acrylic, allowing them to exist inside of the LiLS. Conical reflectors were installed at the face of the housing to improve light collection efficiency in the corners. The housing is filled with optical grade mineral oil and sealed with an O-ring and a 32-mm-thick back plug. For a detailed drawing of the PMT housing module see Figure 3.5. For more information on the PMT housing design and construction see Ref.[42].

3.2.1.2 Optical Grid

The active volume of the detector, measuring $2.045\text{ m wide} \times 1.607\text{ m high} \times 1.176\text{ m long}$, is separated into 154 optically separated long segments with a $0.145\text{ m} \times 0.145\text{ m}$ square cross-sectional area. The optical grid that creates the individual segments

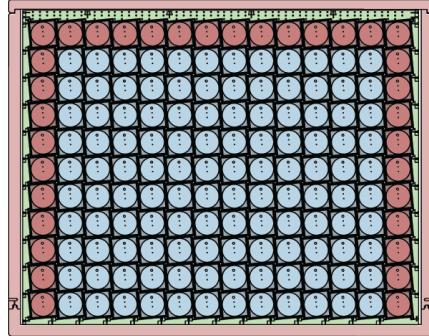


Figure 3.4: A cross-section of the inner AD showing 68 ET PMTs (red) in the outer columns and top row and 240 Hamamatsu PMTs (blue) in the remaining segments.

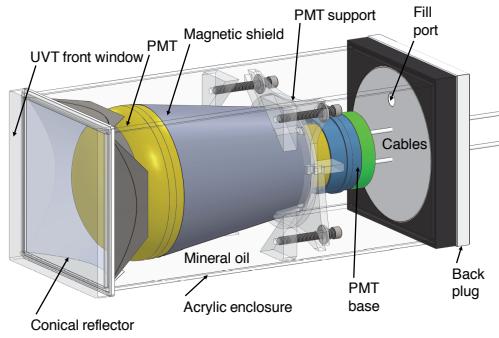


Figure 3.5: A PMT housing module.

consists of low-mass, highly specularly reflective optical separators held in position by white 3D-printed support rods. The optical separators (reflectors) are composed of a carbon fiber backbone covered on both sides with adhesive-backed 3M DF2000MA specularly reflecting film, an optically clear adhesive film, and a thin surface layer of fluorinated ethylene propylene (FEP) film. Two types of pinwheel shaped support rods were produced and strung on acrylic rods to grip the reflectors and hold them in place and separate the PMT housings from each other. The pinwheels were 3D printed using white-dyed 100-micron polylactic acid (PLA) filament and are pictured in Figure 3.6. For more details on the fabrication of the optical grid see Ref.[49].

Each segment contains a PMT housing at each end and four reflectors held in place by pinwheel rods that extend from one PMT to the other, as shown in Figure 3.7. The front windows of the PMT housings protrude ~ 1 cm into the optical grid, minimizing cross-talk between segments. Figure 3.8 shows the assembly of the top row of the detector, demonstrating the placement of the housings and optical grid.

3.2.1.3 Segment Supports

While the optical grid creates the inner volume segmentation, acrylic segment supports hold the total volume in place and determine the size of the active volume. A slab of ship-lap style acrylic underneath the bottom row of segments position them at a 5.5° tilt with a 0.146 m pitch. Horizontal planks are screwed onto the backs

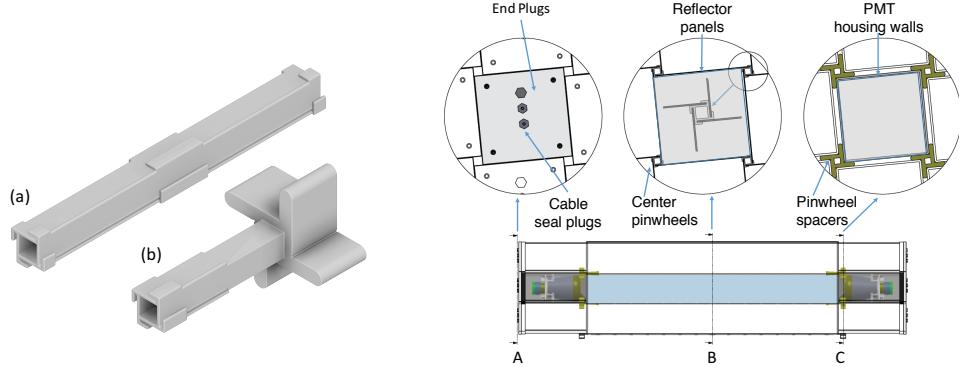


Figure 3.6: Representative pinwheel types. (a) Central pinwheel - Three tabs per side hold the optical separator in place. (b) End pinwheel - spacer arms separate the PMT housing bodies and support the pinwheel string.

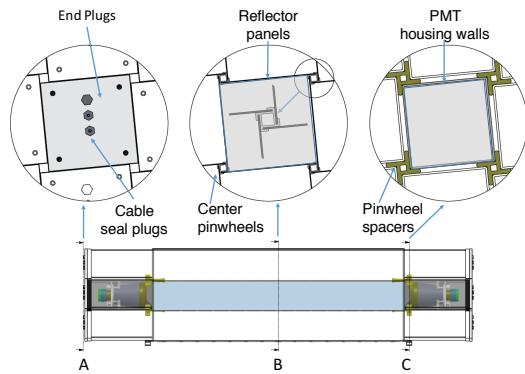


Figure 3.7: Three complete segments, including PMT housings at each end with reflectors kept in place between segments by pinwheel rods.



Figure 3.8: Assembly of the top row of the PROSPECT AD, demonstrating the placement of the PMT housings and optical grid.

of the PMT housings and attach together along the sides of the volume, while side walls constrain the outer pinwheel rows. Baffles at the top of the detector tie the four surrounding walls together and keep the top reflector layer in place. The construction of the inner detector, including all segment supports, allows the liquid scintillator to flow around all objects and therefore fill the whole space. For a photograph of the constructed inner detector see Figure 3.9.

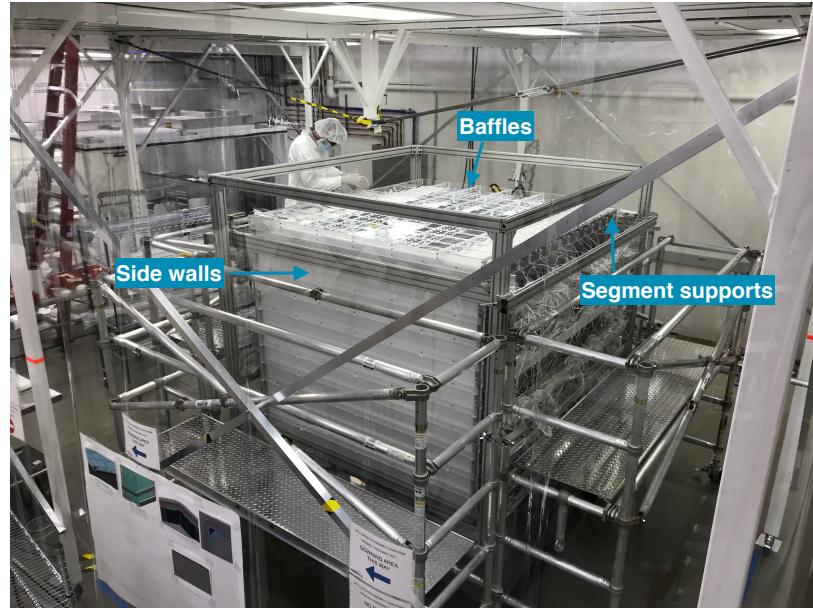


Figure 3.9: A photograph of the constructed inner detector.

3.2.1.4 Radioactive Calibration System

The radioactive calibration system is designed to measure and calibrate the energy and position response of the detector as well as to study topological effects. It does this by moving radioactive sources through a system of tubes routed throughout the active detector, the positions of which are pictured in Figure 3.10. PROSPECT deploys ^{137}Cs , ^{22}Na , ^{60}Co , ^{252}Cf , and an AmBe source, the features of which are listed in Table 3.1.

Each source is encapsulated in a small aluminum cylinder, about 12 mm long, and then connected to a timing belt as pictured in Figure 3.11. The timing belts are placed in source tubes that run along the length of the segments and are controlled

Source	Decay	γ Energy [MeV]	Purpose	Rate
^{137}Cs	β^-	0.662	segment comparison	$0.1 \mu\text{C}$
^{22}Na	β^+	$2 \times 0.511, 1.275$	positron, edge effects	$0.1 \mu\text{C}$
^{60}Co	β^-	1.173, 1.332	energy scale	$0.1 \mu\text{C}$
^{252}Cf	n (fission)	2.223 (n-H capture)	neutron response	866 n/s
AmBe	n	4.4	neutron response	70 n/s

Table 3.1: Calibration sources and their properties.

by motors mounted outside of the detector volume. For more information on the construction of the calibration system see Ref.[50].

PROSPECT also makes use of intrinsic radioactive sources for calibration and monitoring of detector characteristics. Two of these are colloquially known as “BiPo” decays, and they arise from the $^{212}\text{Bi} \rightarrow ^{212}\text{Po} + \beta \rightarrow ^{208}\text{Pb} + \alpha$ and $^{214}\text{Bi} \rightarrow ^{214}\text{Po} + \beta \rightarrow ^{210}\text{Pb} + \alpha$ decay chains which stem from naturally occurring ^{232}Th ($t_{1/2} = 14$ Gyr) and ^{238}U ($t_{1/2} = 4.5$ Gyr), respectively. Along with these, a solution of ^{227}Ac was added to the liquid scintillator to provide a source of “RnPo” decays from the chain $^{219}\text{Rn} \rightarrow ^{215}\text{Po} + \alpha \rightarrow ^{211}\text{Pb} + \alpha$. Further details on the motivation and results from adding actinium will be discussed in Chapter 5.

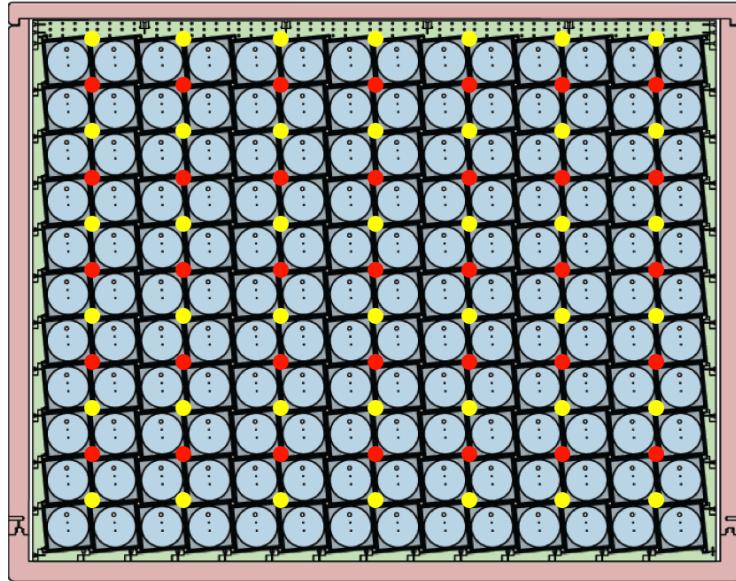


Figure 3.10: Location of the source tubes (red) routed through the active detector volume.

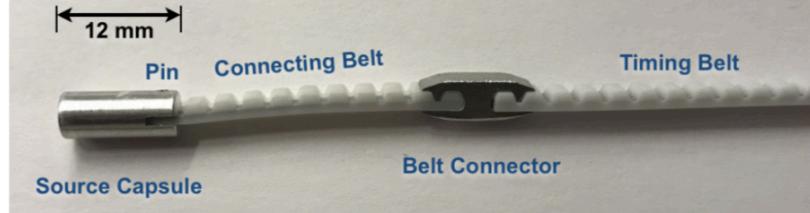


Figure 3.11: An example of a source capsule attached to the timing belt.

3.2.1.5 Liquid Scintillator

The liquid scintillator (LS) used in PROSPECT needed to accomplish two goals: (i) provide very good pulse shape discrimination (PSD) for sufficient background rejection of fast neutrons and ambient gammas and (ii) have high light yield in order to obtain good energy resolution. The LS also needed to remain stable over time and non-flammable according to facility requirements. In order to accomplish these tasks PROSPECT developed a novel lithium-doped liquid scintillator (LiLS). ${}^6\text{Li}$ was chosen as the doping agent due to it's high neutron capture cross section, which produces an α and a ${}^3\text{H}$ with about 540 keV of visible energy in the scintillator. These particles are also spatially localized, which is necessary for the compact detector size of PROSPECT.

The LiLS was created by adding a surfactant to the base LS which allowed the addition of a ${}^6\text{LiCl}$ solution to obtain a final doping of $0.082\%\pm0.001\%$ by mass. The combination of the surfactant and chloride solution forms a thermodynamically stable micro emulsion, ensuring material uniformity and allowing the addition of an actinide chloride solution (${}^{227}\text{Ac}$). A total of 5,040 liters were produced and stored in 28 separate drums. One drum was doped with a ${}^{227}\text{Ac}$ chloride solution. For more information on the fabrication of the LiLS see Ref.[51].

Prior to filling the detector, all drums of LiLS were pumped into an ISO tank storage container. Nitrogen was then bubbled through the liquid for ten days to sufficiently mix together the solution. Samples were taken from each barrel and from the mixed solution in the ISO tank with a Shimadzu UV-Vis spectrometer, the results of which can be seen in Figure 3.12. 4841 kg of LiLS was pumped into the ISO tank and a total of 4340 kg was then added to the detector. For more information on the process of filling the detector with the LiLS see Ref.[42].

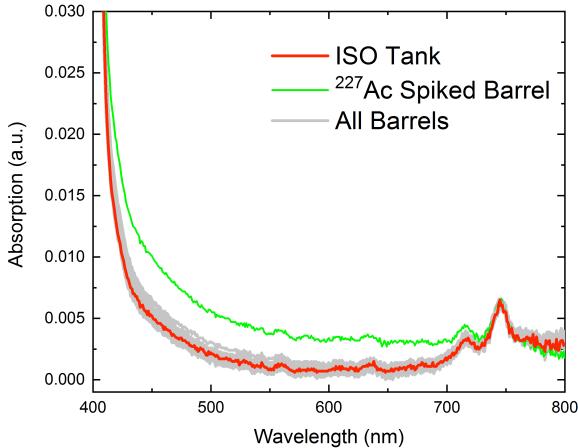


Figure 3.12: UV-Vis absorption spectra of the 28 drums if LiLS (gray+green) added to the ISO tank. The only outlier is the ^{227}Ac spiked barrel (green). The mixed sample (red) falls within the average of all individual barrels.

3.2.2 Containment Vessels and Shielding

The inner primary containment vessel is made of acrylic and holds all active detector components described in Section 3.2.1. The acrylic tank was built as three separate parts, the base, vertical walls, and lid. The inner detector was assembled on the acrylic base. After its constructions the vertical walls were lowered onto the base and secured with a Viton seal at the base.

The secondary containment vessel is made of aluminum and the top is sealed in order to control the gas environment around the detector. The space between the aluminum and acrylic tanks is filled with sheets of borated polyethylene and demineralized water for absorption of thermal neutrons.

A lead layer of $0.025\text{ m} \times 0.10\text{ m} \times 0.30\text{ m}$ interlocking brick was stacked around the perimeter of the aluminum tank. Rows of $0.10\text{ m} \times 0.10\text{ m}$ recycled polyethylene lumber were stacked on each other log cabin style and then placed around the lead layer. Polyethylene lumber was also used to create roof beams on top of the log cabin walls. The entire surface was then covered with a 0.025 m thick layer of borated polyethylene and thin aluminum sheets. To complete the passive shielding an array of water bricks were added to the top of the assembly. For a schematic of the entire construction see Figure 3.3.

3.3 Data Acquisition System

In order to perform pulse discrimination analysis of LiLS signals PROSPECT uses commercial Waveform Digitizer Modules (WFDs). A total of twenty-one CAEN V1725 WFD modules, operated in two Weiner 6023VME crates, are used to readout the 308 PMTs. Readout and control of the WFD modules is performed by two individual PCs, which are run by a control PC. A single Phillips Scientific 757D NIM Fan-In/Fan-Out module operated in a NIM bin is used for trigger signal distribution. A diagram of this system can be seen in Figure 3.13.

Acquisition of waveforms 148 samples long by all WFD channels is triggered if both PMTs in any segment exceed a signal level of 50 ADC counts above baseline (~ 100 keV) within a 64 ns coincidence window. Acquired samples from each WFD are only recorded to disk if they exceed a threshold limit of 20 ADC counts above baseline (~ 40 keV). These are labeled the segment and Zero Length Encoding (ZLE) thresholds and allow the acceptance of low energy events while maintaining a manageable data collection. For more information of the data acquisition system and data rates see Ref. [42].

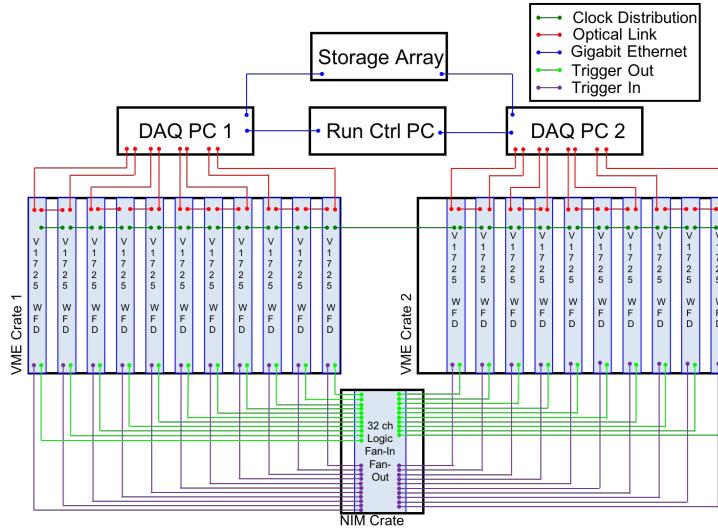


Figure 3.13: Diagram of the data acquisition system.

CHAPTER 4

PROSPECT ANALYSIS FRAMEWORK AND CALIBRATION

Events in the PROSPECT detector begin as bursts of scintillation in the liquid scintillator. In order to transform these events of light into physics data several steps have to be taken, including position reconstruction and energy calibration. This chapter will outline how these processes are performed, but before that is done a key component of the PROSPECT analysis, pulse shape discrimination, must be presented.

4.1 Pulse Shape Discrimination

Physics events in the PROSPECT detector, such as neutron captures on ${}^6\text{Li}$, produce scintillation light through ionization that is transported by way of the reflecting panels to individual PMTs. As described in Section 3.3, these signals are processed by CAEN waveform digitizers and are only accepted if they pass the segment and ZLE thresholds. Due to the nature of the liquid scintillator the shape of the digitized waveforms is defined by the ionization density of a given event. Lower ionization density events, such as electrons, have a faster scintillator decay time causing less light in the “tail” of the waveform compared to higher density events like proton recoils, as seen in Figure 4.1.

This allows the definition of a pulse shape discrimination (PSD) factor as the ratio

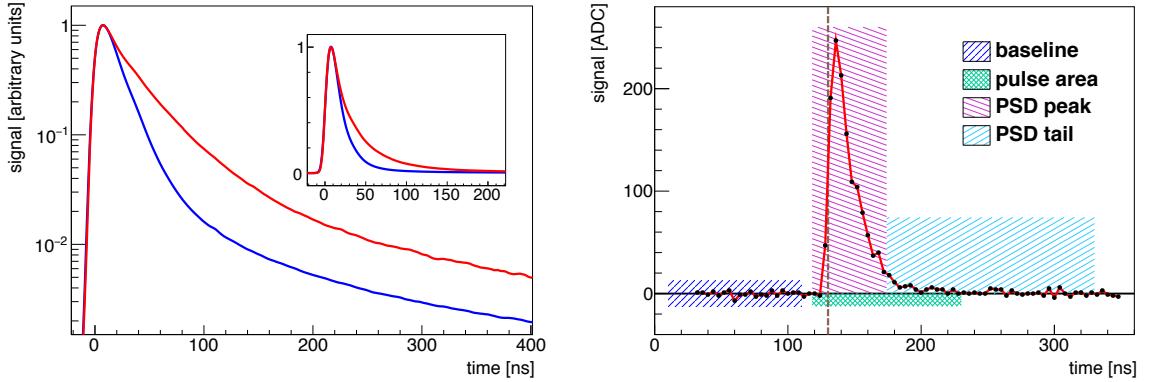


Figure 4.1: (Left) Averaged waveforms from electrons (lower, blue) and proton recoils (upper, red) [52]. The inset panel shows the same waveforms on a linear y axis. (Right) Example analysis of a typical pulse [53]. The half-height leading edge timing (dashed vertical) determines windows for baseline subtraction, pulse area, and PSD.

of the signal in the tail versus the total waveform,

$$PSD = \frac{\int_{tail:start}^{tail:end} Qdt}{\int_{-\infty}^{\infty} Qdt} \quad (4.1)$$

The tail area of a given waveform is defined as the window 44 - 100 ns after the time of the half-height leading edge. The total area is defined as the window -12 - 100 ns relative to the same leading edge time. An example of these windows on a typical pulse can be seen in Figure 4.1. Use of the PSD parameter, along with energy, provides clear separation between neutron captures on ⁶Li and other event classes such as electron recoils. An example of this for a single segment can be seen in Figure 4.2, where a pseudo-energy is calculated using the integrated pulse area of each PMT, S₀ and S₁.

4.2 Data Processing and Calibration

For simplicity consider one event contained in one segment. Both PMTs in the segment will collect photons from this event that are saved as individual waveforms. These are analyzed and information such as pulse area and height in analog-to-digital converter (ADC) units, PSD, and arrival time are saved. The next step involves combining information from both PMTs, calibrating the energy, and reconstructing the position.

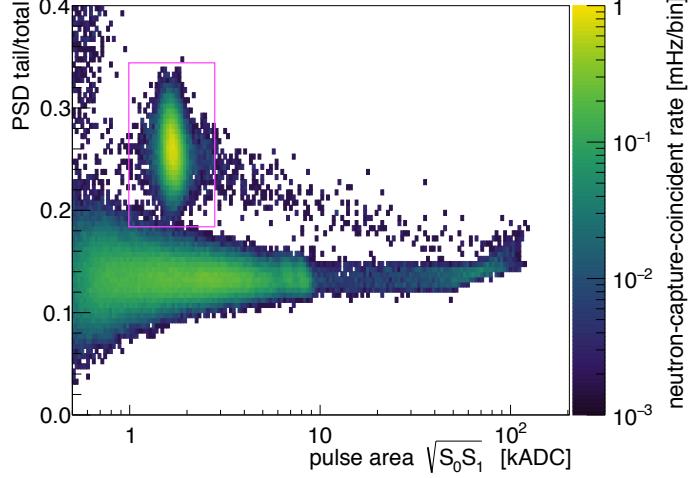


Figure 4.2: PSD vs. pseudo-energy for neutron capture coincident events in a single segment [54]. The neutron captures, outlined by the magenta rectangle, are clearly separated in PSD from electron-like events in the lower band.

4.2.1 Timing to Position Curves

Through-going muons provide a large and consistent data set which can be used to reconstruct positions based on timing. As muons travel through the pinwheel rod tabs the light transport is distorted in timing and magnitude. This creates a striping effect in the timing between the two PMTs, dt , versus the signal amplitude ($S_0 + S_1$) distribution as shown in Figure 4.3.

The striping becomes even clearer when plotting dt for events with signal amplitudes in the range $1e4 - 2e4$ ADC for a single segment as shown in Figure 4.4. For each segment this distribution is first fit with an “M”-shaped curve, $M(dt)$. Then, the residual structure is fit to a sinusoidal curve with a slowly varying phase shift term,

$$n(dt) = M(dt) \left[1 + k \cos \left(\frac{2\pi}{\delta} (a dt + b dt^3) \right) \right] \quad (4.2)$$

where $\delta = 78.5$ mm is the average spacing between pinwheel tabs, and k, a, b are fit parameters. The inner phase term provides the position calibration $z(dt) = a dt + b dt^3$, whose resulting curves are shown in Figure 4.5.

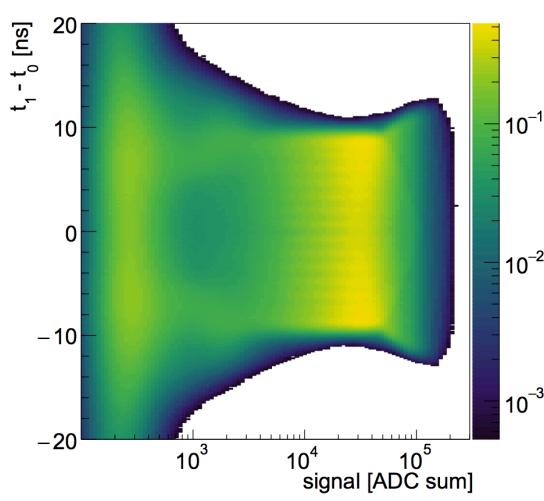


Figure 4.3: dt versus signal amplitude for through-going cosmogenic muons [55]. Striping is visible at time intervals corresponding to pinwheel placement.

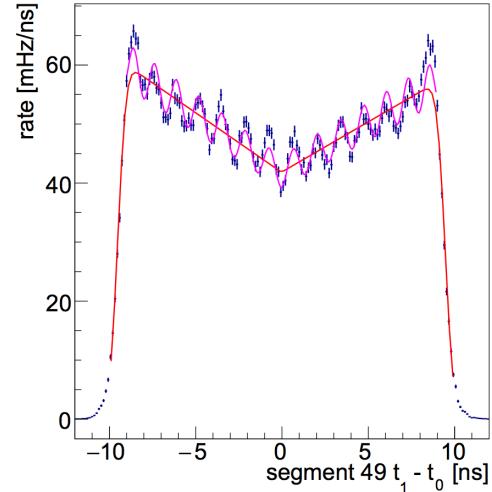


Figure 4.4: dt of through-going muons in a single segment for events with signal amplitudes in the range $1e4 - 2e4$ ADC [55]. Data (blue points) are fit with an “M”-shaped curve (red) and a sinusoidal curve (magenta).

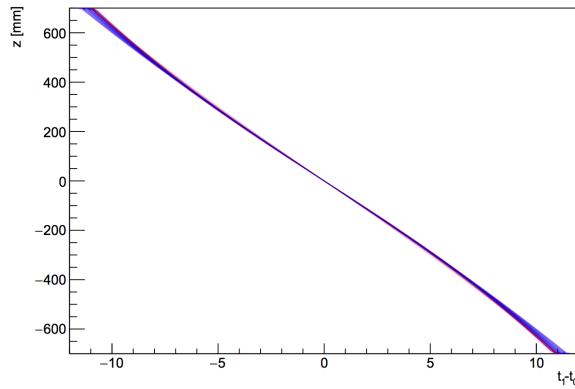


Figure 4.5: $z(dt)$ curves for all cells, extracted from fitting muon dt distributions [55].
Blue: Hamamatsu segments, red: ET segments.

4.2.2 Energy Calibration

The energy of an event is first defined as the sum of the integral of the waveforms from both segment PMTs in ADC units. The next step involves converting to energy units of MeV and correcting for longitudinal light collection variation. These procedures are performed using neutron captures on ${}^6\text{Li}$ ($n\text{Li}$) because they are well understood and always peak at a quenched energy of around 0.53 MeV electron equivalent (MeVee).

After combining the waveforms for all events, the neutron captures at the center of the segment are calibrated to 0.526 MeVee, defining the energy scale for all segments and all classes of events. Constraining the energy scale in this way creates a quadratic dependence of energy on position, as seen in the left panel of Figure 4.6. This is corrected for by calibrating a gain factor for each PMT using two metrics. The first is the neutron capture signal at cell center, $S = \sqrt{S_0 S_1}$, which is found by fitting the distribution in the left panel of Figure 4.6 with a quadratic polynomial. The second is the light ratio, $R = S_1/S_0$, at cell center which is found by fitting the distribution in the right panel of Figure 4.6 with a cubic polynomial. The gain factors for both PMTs are then defined as,

$$g_0 = \frac{S}{\sqrt{R} E_n} \quad g_1 = \frac{S \sqrt{R}}{E_n} \quad (4.3)$$

where E_n is the energy of the neutron capture peak.

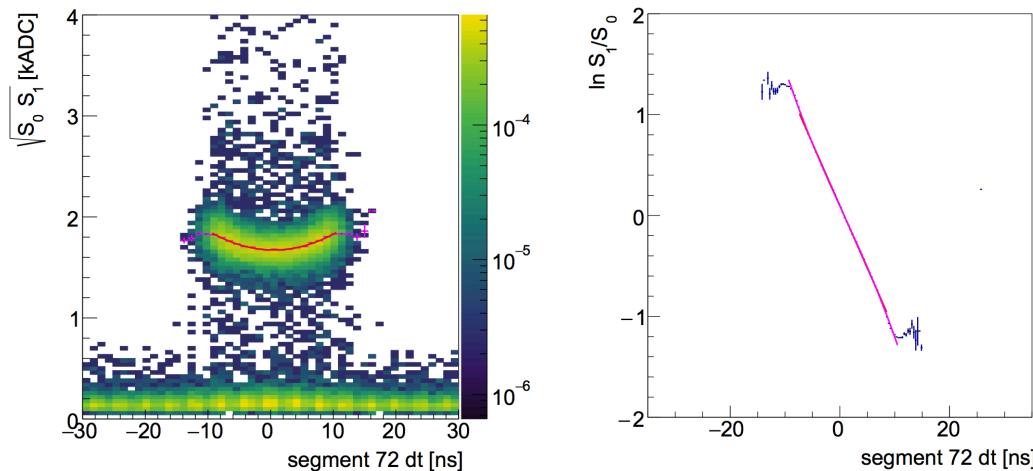


Figure 4.6: (Left) Neutron capture events in one segment versus dt , with a quadratic fit in red. (Right) The \ln signal ratio versus dt for muon events in one segment, with a cubic polynomial fit in magenta.

4.3 Monte Carlo Simulation

Due to the geometry of the PROSPECT detector and properties of the scintillator, reconstructed events are highly position and energy dependent. In order to perform an oscillation analysis it is important to understand the detector response to incoming antineutrino events. This is done by modeling the detector, including all material and scintillator properties, in a GEANT-4 based simulation package hereto referred to as PROSPECT-G4 (PG4). Monte Carlo (MC) simulations with PG4 generate a position-dependent energy response that is used to interpret real physics data. This can only be done if PG4 behaves the same way that the PROSPECT detector does. This is ensured by tuning values in PG4 to reproduce distributions measured by a variety of radioactive calibration sources and intrinsic background energy depositions.

4.3.1 Nonlinearity

The nonlinearity of the scintillator response at low energies is parameterized in PG4 using a combination of Birks' quenching model [56] and a Cherenkov radiation model. Birks' law defines the light yield per path length as a function of the energy loss per path length for a particle traveling through scintillator and at each simulation step i is applied using

$$\frac{dE_{scint}}{dx} = \frac{\frac{dE}{dx}}{1 + k_{B1} \frac{dE}{dx} + k_{B2} \left(\frac{dE}{dx} \right)^2} \quad (4.4)$$

where k_{B1} and k_{B2} are the Birks' constants which depend on the material and dE/dx is the true deposited energy.

Particles traveling faster than the speed of light in the scintillator can emit Cherenkov radiation, therefore, the number of Cherenkov photons, N , generated at each simulation step is calculated as

$$\frac{d^2N}{dxd\lambda} = \frac{2\pi\alpha z^2}{\lambda} \left(1 - \frac{1}{\beta^2 n^2(\lambda)} \right) \quad (4.5)$$

where α is the fine structure constant, z is the particle's electric charge, β is the speed of the particle, and $n(\lambda)$ is the index of refraction. The energy emitted from the Cherenkov radiation is then calculated as the summed energy of detected Cherenkov

photons,

$$E_c = k_c \sum_{\lambda} N_{\lambda} E_{\lambda} \quad (4.6)$$

where k_c is the efficiency of detecting Cherenkov light. Comparison between data and MC allows the tuning of both Birks' constants, k_{B1} and k_{B2} , and k_c .

4.3.2 Energy Response

Reconstructed energy distributions, E_{rec} , are measured using a variety of calibration sources placed at z-center positions in the detector. Similar distributions can also be created through MC simulations by setting values for k_{B1} and k_{B2} , and k_c as well as an absolute energy scale factor, A . A comparison between the data and MC allows for the tuning of these values until good agreement is shown between the two sets of distributions.

For these studies three γ -ray sources were used, ^{60}Co , ^{137}Cs , and ^{22}Na , along with γ 's resulting from neutron-Hydrogen captures using ^{252}Cf as the neutron source (γ energies listed in Table 3.1). The spectrum of ^{12}B was also measured because its β dominated energy distribution (3 MeV to 13.6 MeV) covers a similar energy range as IBD events. As well as comparing the energy spectra, ^{137}Cs and ^{22}Na are used to compare event multiplicity between MC and data. The results of these comparisons can be seen in Figure 4.7, where good agreement is seen between MC and data. The best-fit parameters determined by these studies are $(A, k_{B1}, k_{B2}, k_c) = (1.0026 \pm 0.004, 0.132 \pm 0.004 \text{ MeV/cm}, 0.023 \pm 0.004 \text{ MeV/cm}, 37 \pm 2\%)$, with a χ^2/ndf of 581.5/420.

Further agreement can be seen when comparing the peak locations for all γ -ray sources as seen in Figure 4.8. The ratio between data and MC is shown to be within $\pm 1\%$ for all sources for three different time periods.

4.3.3 Energy Resolution

A comparison of MC and data energy distributions of the γ -ray sources can also be used to define an energy resolution function. MC simulated energy distributions are smeared according to Gaussian distributions until they match data. The resolution of the smeared distributions is then compared to the true deposited energy and fit

with the function

$$\frac{\sigma_E}{E_{rec}} = \sqrt{a^2 + \frac{b^2}{E_{rec}} + \frac{c^2}{E_{rec}^2}} \quad (4.7)$$

where a , b , and c are dependent on detector geometry, photostatistics (PE/MeV), and PMT quantum efficiency respectively. The results of this can be seen in Figure 4.9, where the parameters are found to be $(a, b, c) = (1.15\% \pm 0.47\%, 4.61\% \pm 0.24\%, 0 \pm 1.3\%)$, resulting in an energy resolution at 1 MeV of $4.76\% \pm 0.2\%$.

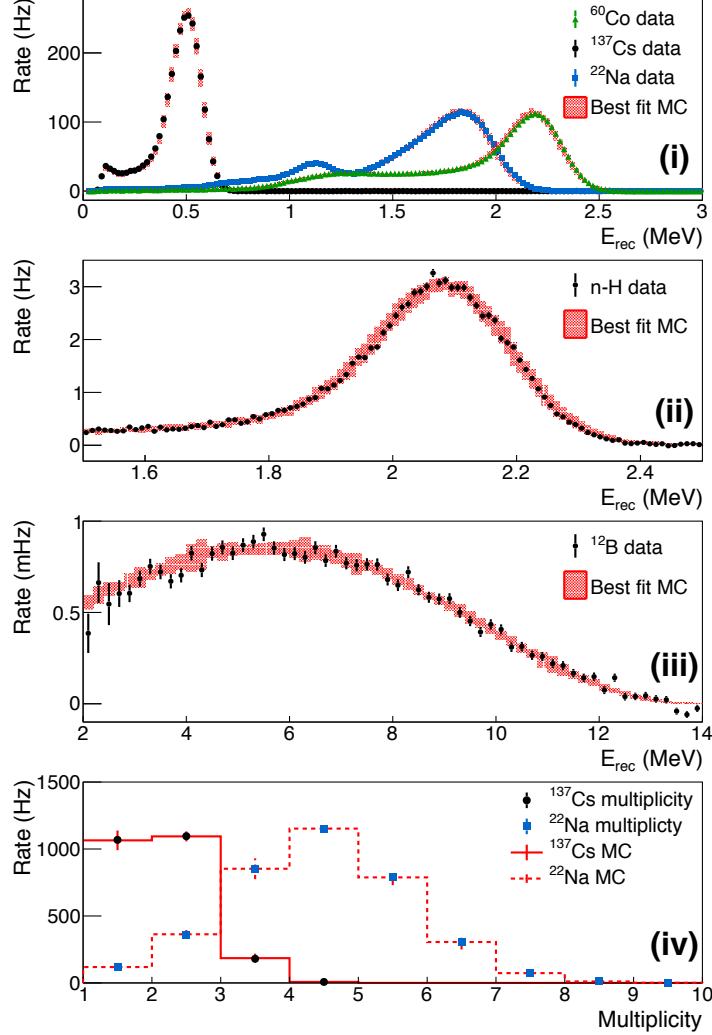


Figure 4.7: Reconstructed energy distributions for calibration data and PG4 Monte Carlo simulations [57]. (i): E_{rec} for γ -ray source deployments; (ii): E_{rec} for n-H captures from a ^{252}Cf source deployment; (iii): E_{rec} for cosmogenically produced ^{12}B ; (iv): pulse multiplicity for ^{137}Cs and ^{22}Na source deployments. Error bands indicate statistical (data) and systematic (PG4) uncertainties.

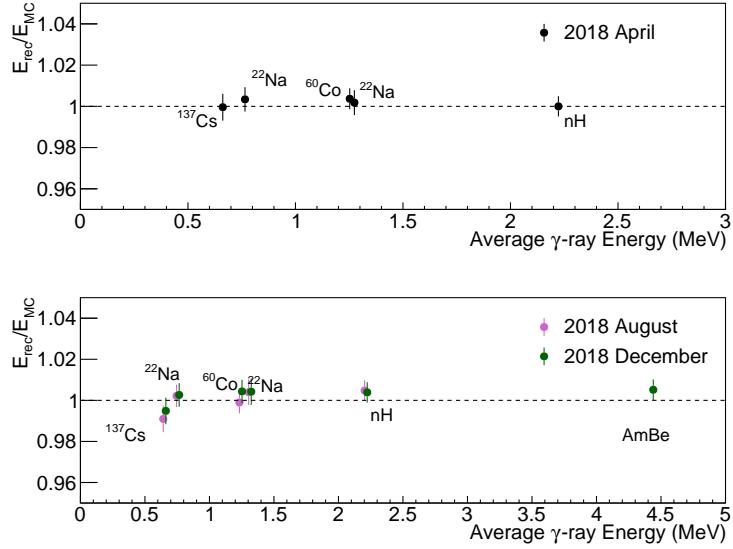


Figure 4.8: Ratio of data versus PG4 Monte Carlo simulation energy peak locations of the given γ sources, plotted versus true gamma energy for three different time periods [57]. Error bands indicate statistical and systematic uncertainties. Ratios for all datasets are within $\sim 1\%$ of unity, indicating accurate energy response modeling in PG4 for a wide energy range.

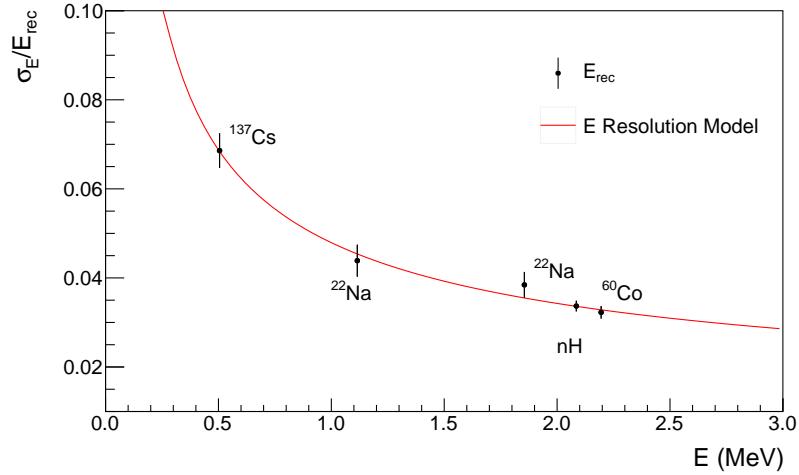


Figure 4.9: Energy resolution of PG4 Monte Carlo simulation distributions matched to data versus true gamma energy for the given calibration sources fit with the function in Eq. 4.7 [57]. Error bands indicate statistical and systematic uncertainties.

CHAPTER 5

^{227}AC AS A CALIBRATION SOURCE

5.1 Motivation

In the absence of an eV-scale sterile neutrino PROSPECT should measure IBD rates that fall like one over distance from the reactor squared. If sterile neutrino oscillation was detected, after one year PROSPECT would measure the spectrum seen in Figure 5.1, given a mass splitting of 1.78 eV^2 , . However, a situation could occur in which a sterile neutrino did not exist, or could not be measured with PROSPECT, and an oscillation is still measured. This could happen if there are segment to segment volume variations throughout the detector, mimicking an oscillation signal. Therefore, it becomes important that the product of efficiency \times volume for all segments is well known.

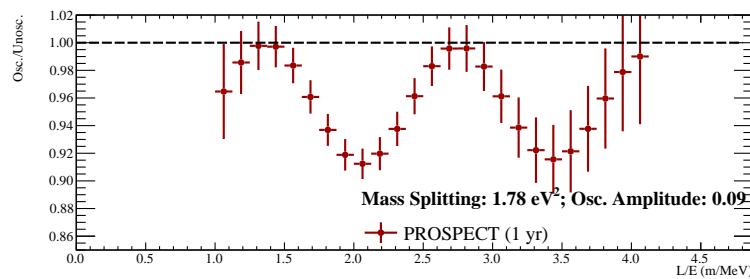


Figure 5.1: The ratio of the oscillated to un-oscillated neutrino spectrum as a function of L/E that would be observed by PROSPECT after 1 year if a sterile neutrino signal was detected [58].

This measurement can be accomplished if an event source is uniformly distributed throughout the active volume of the detector. By measuring the rate of this source in each segment the relative volumes can be compared and tracked through time. ^{227}Ac was chosen as the source and a chloride solution was prepared and dissolved in the liquid scintillator to ensure uniform distribution.

^{227}Ac was chosen for several reasons. First, because an α , α coincidence occurs in its decay chain, specifically $^{219}\text{Rn} \rightarrow ^{215}\text{Po} + \alpha \rightarrow ^{211}\text{Pb} + \alpha$, as highlighted in Figure 5.2. ^{219}Rn has a half-life of 3.96 ± 0.01 s and α -decays 100% of the time, while ^{215}Po has a half-life of 1.781 ± 0.005 ms and α -decays 99.99977(2)% of the time [59], so the α , α coincidences happen frequently. The α decay of ^{215}Po is mono-energetic at 7.39 MeV which results in a ~ 0.78 MeVee signal after quenching, well removed from nLi captures that occur around 0.5 MeVee. In addition, there are no corresponding gammas with the ^{215}Po decay, making this a very clean and well defined signal. The ^{219}Rn α decays are not so nice, as there are four dominant alpha energies and three dominant gamma energies for these decays, as listed in Table 5.1, but the use of time, energy, and PSD cuts make them easy to pair with corresponding ^{215}Po decays.

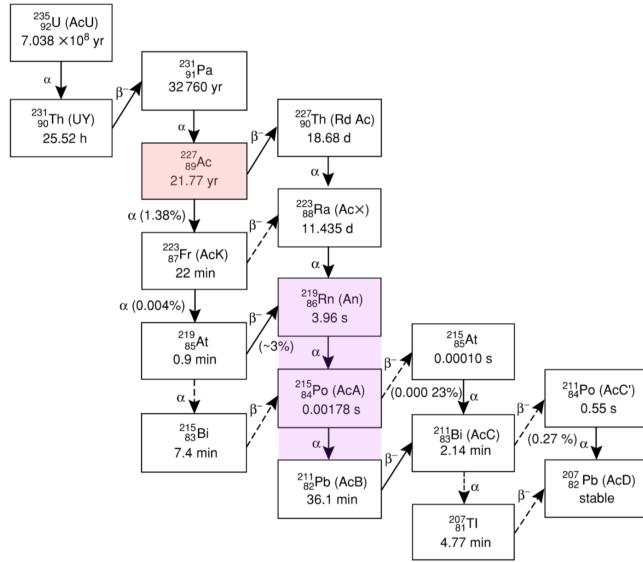


Figure 5.2: The full decay chain of ^{227}Ac (a daughter of ^{235}U), in which the α , α coincidence of interest is highlighted [60].

	E_α [keV]	I_α %	E_γ [keV]	I_γ %
^{219}Rn	6425.0(10)	7.5(6)	271.23(1)	10.8(6)
	6530(2)	0.110(10)	401.81(1)	6.6(4)
	6552.6(10)	12.9(6)	130.60(3)	0.13(9)
	6819.1(3)	79.4(10)		
^{215}Po	7386.1(8)	99.999770(20)		

Table 5.1: Energy and absolute intensity of dominant α and γ decay radiation for ^{219}Rn and ^{215}Po . Decay energies not listed here have an intensity of $<0.05\%$.

5.2 Material Compatibility

Before ^{227}Ac could be added to the PROSPECT detector, it had to be determined that ^{227}Ac and its daughters would not adsorb onto detector materials. If it was adsorbed then it would not be a uniform source in the detector, nullifying its use as a method to track relative efficiency \times volume throughout the detector. To test this six material samples were placed in vials of $^6\text{Li-LS}$ spiked with ^{227}Ac . The rate of ^{227}Ac in each sample vial and one reference vial with no material was measured and tracked over a period of 6 months. An observation of a significant decrease in rate, relative to the half-life of ^{227}Ac , would indicate that ^{227}Ac was adsorbing onto the material.

5.2.1 Material and Scintillator Preparation

The materials tested were: ultraviolet transmitting (UVT) acrylic, flourinated ethylene propylene (FEP), polylactide (PLA), polyether ether ketone (PEEK), a RG 188 cable, and viton o-rings. See Table 5.2 for a list of their uses in the detector and sample sizes. To prepare the materials they were all placed in a single beaker with ultra-pure water and cleaned ultrasonically for 30 minutes. They were then transferred to a watch glass and placed in a 50 C oven for two hours. After drying they were placed in empty 12 mL vials.

The ^{227}Ac used to spike the scintillator was obtained from Eckert and Ziegler as a solution of 3.711×10^4 Bq $\pm 1.32\%$ of ^{227}Ac in 10.22710 g of 1 M HCl, measured on September 6, 2016. 0.503 g of this solution was added to 192 g of $^6\text{Li-LS}$ on December 15, 2016. With a half-life of 21.772 ± 0.003 yrs [59], the activity of the ^{227}Ac solution before adding to the LiLS was 36788 Bq, yielding a final activity of 94.2 Bq/10 g.

This is the stock solution from which all LS was taken for the material studies and later on for spiking the detector.

Material	Detector Use	Sample Size
UVT Acrylic	Front window of PMT housing	$1.0 \times 1.15 \times 0.1 \text{ cm}^3$
FEP	Film on optical separators	$1.5 \times 1.5 \text{ cm}^2$, 3 mm thick
PLA	3D printed pinwheels	10 disks; 0.5 cm diameter, 0.1 cm thick
PEEK	Seal plugs through which the high voltage and signal cables were threaded. Screws used to bolt together segment supports. Spacers at the base of the acrylic tank.	1 Nut; ID 0.5 cm, small OD 1cm, large OD 1.1cm, thickness 0.5 cm
RG188 Cable	High voltage and signal cables	4.5" long
Viton O-ring	Seal back plugs of PMT housings and seal acrylic tank	10 O-rings; OD 6mm, ID 3mm, thickness 1.5mm

Table 5.2: Samples used to test if ^{227}Ac or its daughters would adsorb onto detector materials.

Material	Date Filled	Weight of LiLS Added (g)
Reference	12/15/2016	10.030
UVT Acrylic		9.98
FEP		9.98
PLA		9.999
PEEK	02/24/2017	9.99
RG188 Cable		9.981
Viton O-ring		10.011

Table 5.3: The weight of ^{227}Ac spiked LiLS that was added to each sample vial.

Prior to filling all sample vials the threads of each vial were wrapped with teflon tape in an effort to obtain a secure seal. The reference vial was filled on December 15, 2016 with 10.030 g of ^{227}Ac spiked LiLS from the stock solution, yielding an expected activity of 94.5 Bq. All material vials were filled on February 24, 2017 with the amount of stock solution added to each listed in Table 5.3. At the time of filling the

rate in each vial was expected to be ~ 93 Bq. For a photograph of all filled material sample vials see Figure 5.3.

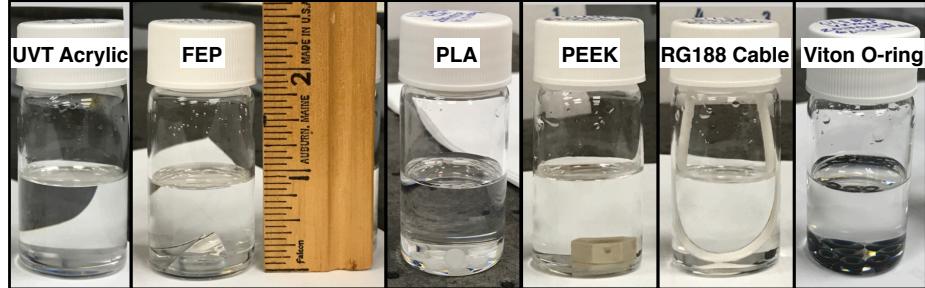


Figure 5.3: Photos of all material sample vials filled with ^{227}Ac spiked LiLS, with ruler for scale.

5.2.2 Detector

The detector consisted of a 2" photomultiplier tube coupled using optical grease to a solid cylinder of UVT acrylic painted white with an insert cut out to hold the sample vials, as shown in Figure 5.4. Placed in a dark box the PMT is cabled to a CAEN DT55xx Desktop HV Power Supply and a CAEN DT5730 8 Channel 14-bit 500 MS/s Digitizer [61]. A modified version of Wavedump 3.7.2 [62] was used to start and stop the data runs and save the waveforms of the signals.

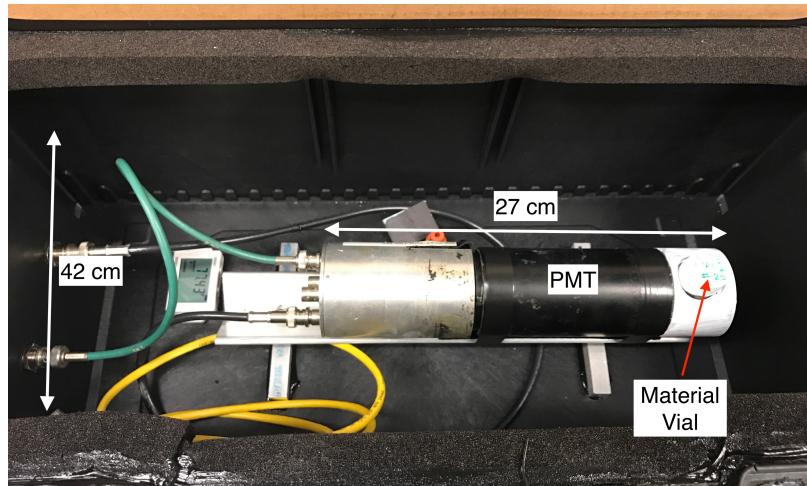


Figure 5.4: Detector used for material studies, consisting of a 2" PMT coupled to an acrylic cylinder holding the sample vials, all contained in a dark box. The PMT is cabled to a power supply and digitizer that exist outside of the box.

5.2.3 Data Analysis

The raw waveforms are analyzed to calculate the energy, PSD, and time of each signal. The energy is determined by taking an integral of the waveform in ADC units and then converted to nC by... The PSD is defined as... [DEFINE these values]

The ^{227}Ac coincident alpha events, labeled RnPo events for the remainder of this document, are found by applying a set of timing, energy, and PSD cuts along with an accidental background subtraction. Po events are found first by applying the energy and PSD cuts listed in Tabel 5.4. Rn coincidental events are found by looking in a 12.85 ms time window before a given Po event and applying the same energy and PSD cuts. This time window is 5 times the lifetime of Po, 2.57 ms, allowing the collection of all possible coincident events. Accidental events are found by looking in the same length time window, using the same energy and PSD cuts, but offset 10 Po lifetimes before a given Po event. RnPo events are then found by subtracting the accidental events from the coincident events. See Figure 5.5 for an example of typical energy and PSD distributions.

Energy	$0.01 < E < 0.055 \text{ nC}$
PSD	$0.31 < \text{PSD} < 1.0$
$\Delta t = t_{\text{delay}} - t_{\text{prompt}}$	$\Delta t < 5\tau_{\text{Po}}$

Table 5.4: Energy, PSD, and time cuts used to find RnPo events where $\tau_{\text{Po}} = 2.57$ ms. Energy and PSD cuts are applied to both prompt and delay events.

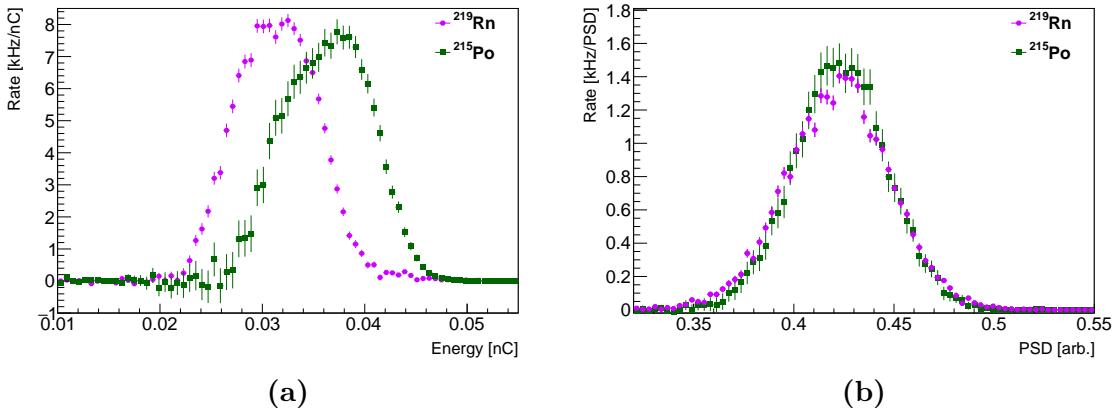


Figure 5.5: Typical energy (a) and PSD (b) distributions for RnPo events in the reference sample after accidental background subtraction.

The rate of RnPo events is measured by fitting the RnPo Δt distribution with

$$f(t) = N_0 e^{-t/\tau} \quad (5.1)$$

where N_0 and τ , the lifetime of ^{215}Po , are allowed to vary. Using the fit results, the rate is then defined as

$$R = \frac{N_0\tau}{\text{bin-width} \times \text{livetime}} \quad (5.2)$$

$$\sigma_R = R \times \sqrt{\left(\frac{\sigma_{N_0}}{N_0}\right)^2 + \left(\frac{\sigma_\tau}{\tau}\right)^2 + \frac{2\sigma_{N_0\tau}}{N_0\tau}} \quad (5.3)$$

where the livetime is measured, for each run, as the time from the beginning of the run to the last Po event. An example of a typical RnPo Δt distribution can be seen in Figure 5.6. It should be noted here that the energy and PSD cuts were made wide enough so that no efficiency correction needed to be applied.

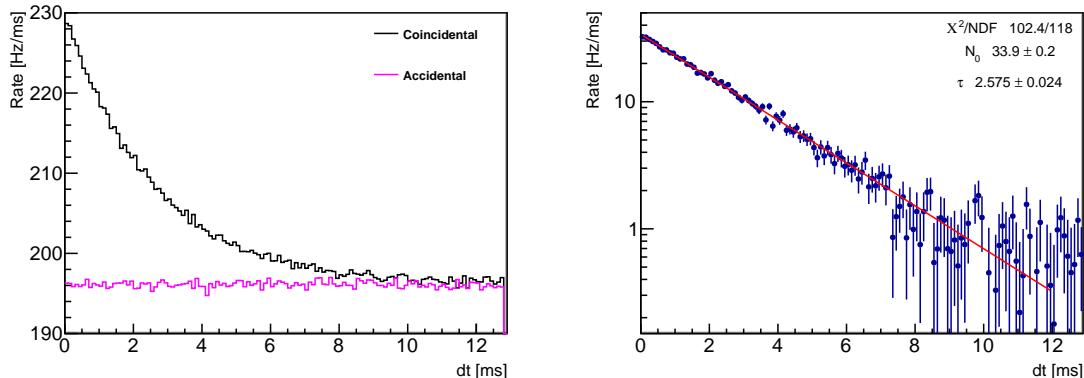


Figure 5.6: A typical example of the RnPo Δt distributions for the reference sample. Left: coincidental and accidental distributions found using the defined energy and PSD cuts. Right: the Δt distribution after subtraction of the accidental distribution, fit with Equation 5.1.

5.2.4 Results

The RnPo rate was calculated for each material sample and the reference sample over a period of about six months. These results can be seen in Figure 5.7. Though statistical errors vary from around 0.6-1%, overall rates vary at a rate of around 12%, indicating systematic variations.

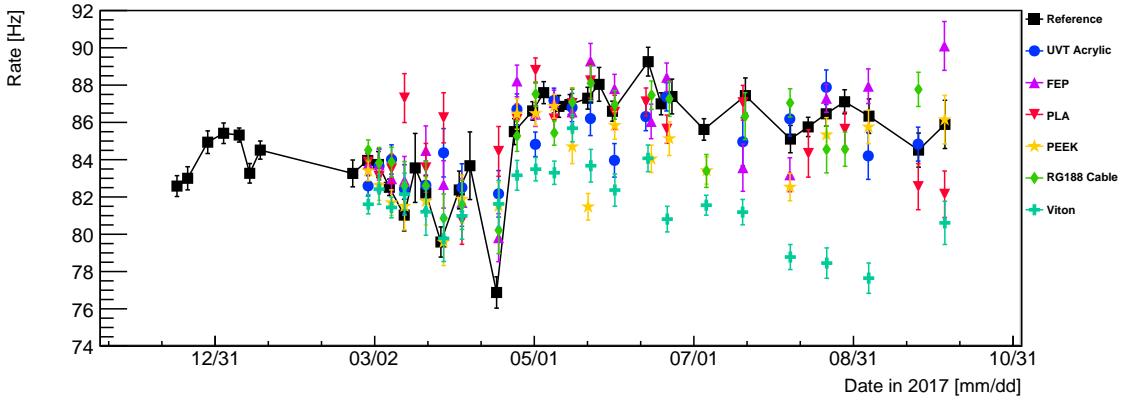


Figure 5.7: ^{227}Ac rate for each material sample. Errors are statistical.

Systematic variations can be better understood by looking at the behavior of the ^{215}Po energy distribution through time. This was done by fitting this distribution for the reference sample with a sum of two Gaussians to account for the non-Gaussian nature of the peak as demonstrated in Figure 5.8. The mean and 1σ width of each of these Gaussians versus time is show in Figures 5.9 and 5.10. It can be seen that the ^{215}Po energy mean varies about 5% and the width around 15%.

The amount of variation seen in measured rates and the ^{215}Po energy distribution indicate that the system is not repeatable and, as such, introduces significant system-

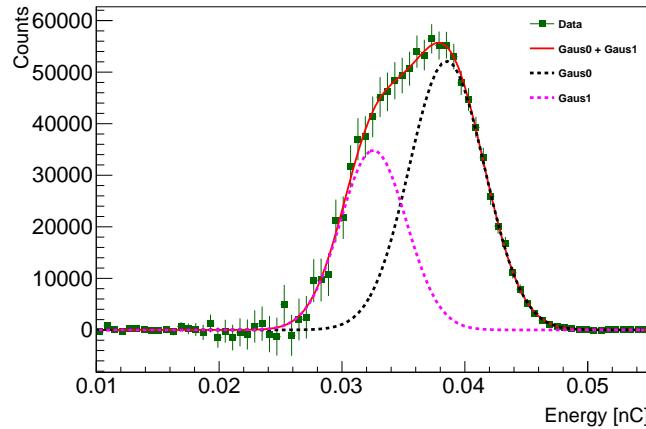


Figure 5.8: ^{215}Po energy distribution for the reference sample, fit with a sum of two Gaussians. The total fit is seen in red, while the two Gaussians are drawn as the pink and black dashed lines.

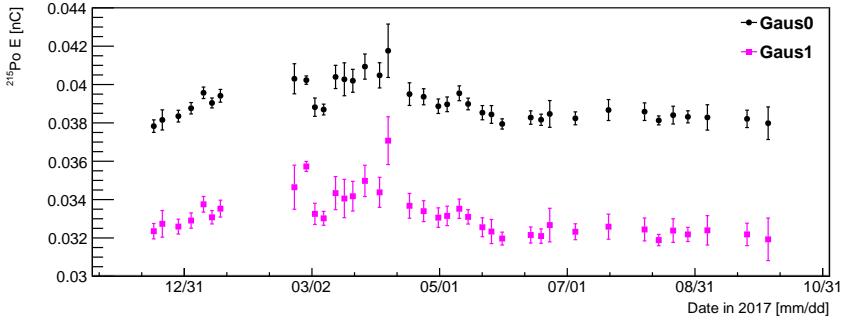


Figure 5.9: The mean of the two Gaussians fit to the ^{215}Po distribution for the reference sample.

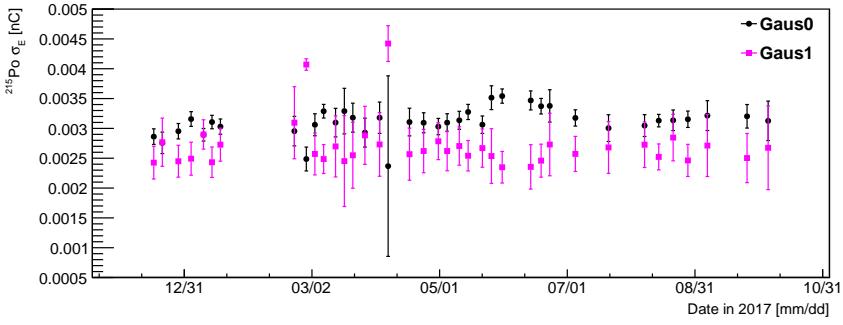


Figure 5.10: The 1σ width of the two Gaussians fit to the ^{215}Po distributions for the reference sample.

atic errors. Though the PMT and acrylic holder were glued in place, the sample vials were repeatedly removed and replaced, possibly shifting the placement of the acrylic holder and the optical grease. If the experiment were to be repeated a more robust detector system would be needed to eliminate these systematic errors.

To account for these variations all material sample rates, R_M , were compared to the reference sample rate, $R_{ref.}$. The ratio of the rates was calculated for each time bin as

$$ratio = \frac{R_M}{R_{ref.}} \quad (5.4)$$

$$\sigma_{ratio} = ratio \times \sqrt{\left(\frac{\sigma_M}{R_M}\right)^2 + \left(\frac{\sigma_{ref.}}{R_{ref.}}\right)^2} \quad (5.5)$$

and the results are shown in Figure 5.11. The ratio of rates versus time, for each material, were fit with a constant and a straight line, the results of which are tabulated

in Tables 5.5 and 5.6 respectively.

Except for the case of viton (discussed in the next section) there is no clear decrease in rate observed over the six month period for any material. Though the chi-squared results for the constant fits are not ideal, the fits to a straight line result in slopes with 50% error to greater than 100% error, indicating that a decrease in rate is not a good model. Since no obvious and significant decrease was observed it was concluded that ^{227}Ac was not adsorbing onto materials and next steps were taken to include the source in the final detector.

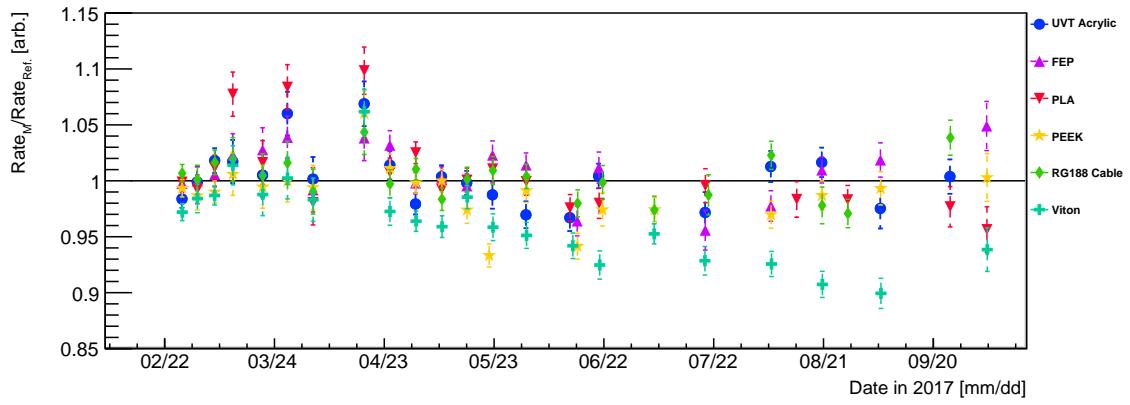


Figure 5.11: ^{227}Ac rate for each material sample, M , relative to the reference sample.

Material	Constant	χ^2/NDF
UVT Acrylic	0.997 ± 0.003	$56.5/20 = 2.82$
FEP	1.005 ± 0.003	$44.1/20 = 2.21$
PLA	1.003 ± 0.003	$80.6/20 = 4.03$
PEEK	0.985 ± 0.003	$70.9/20 = 3.55$
RG188 Cable	1.001 ± 0.003	$38.3/21 = 1.82$
Viton	0.960 ± 0.002	$136.3/21 = 6.49$

Table 5.5: The results of fitting the relative rate for each material sample with a constant.

Material	Constant	Slope [ratio/yr]	χ^2/NDF
UVT Acrylic	1.5 ± 0.8	-0.01 ± 0.02	$56.2/19 = 2.96$
FEP	1.3 ± 0.8	-0.005 ± 0.018	$44.0/19 = 2.32$
PLA	4.4 ± 0.8	-0.07 ± 0.02	$64.3/19 = 3.38$
PEEK	2.9 ± 0.8	-0.04 ± 0.02	$65.2/19 = 3.43$
RG188 Cable	2.3 ± 0.8	-0.03 ± 0.02	$35.7/20 = 1.79$
Viton	7.7 ± 0.7	-0.14 ± 0.02	$48.9/20 = 2.45$

Table 5.6: The results of fitting the relative rate for each material with a straight line.

5.2.4.1 Viton

Observation of the rate of ^{227}Ac in the viton o-ring material sample vial initially indicates a decrease in rate over time, about 10% compared to the reference vial over a six month period. Upon further inspection, though, it becomes clear that the energy spectrum, of the Rn events in particular, shift toward the system threshold as time goes on. This causes a loss of events, not due to adsorbance, but rather due to threshold effects.

Figure 5.12 shows the ^{219}Rn and ^{215}Po energy distributions in the first and last time bins for both viton and PEEK. It can be seen that at the last time bin the viton distributions sit against the threshold, compared to the PEEK distributions which approach but do not get close to the threshold. To quantify this the ^{219}Rn spectrum was fit with a sum of two Gaussians and the widths versus time are shown in Figure 5.13. It can be seen that the lower energy Gaussian becomes narrower as time goes on, indicating a loss of events due to threshold effects. Therefore, it was concluded that the decrease in ^{227}Ac rate observed in the viton o-ring sample was due to threshold effects rather than adsorption.

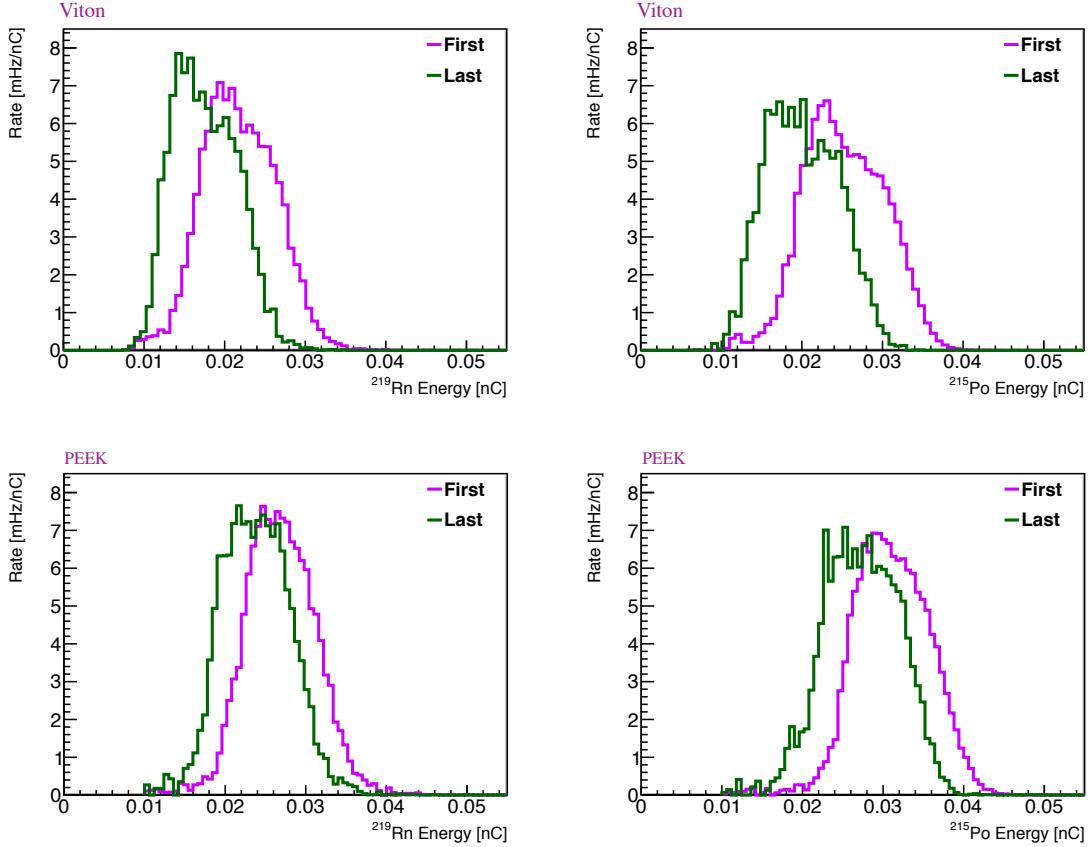


Figure 5.12: ^{219}Rn and ^{215}Po energy spectra for both the viton o-ring (top) and PEEK (bottom) material samples during the first and last time bins.

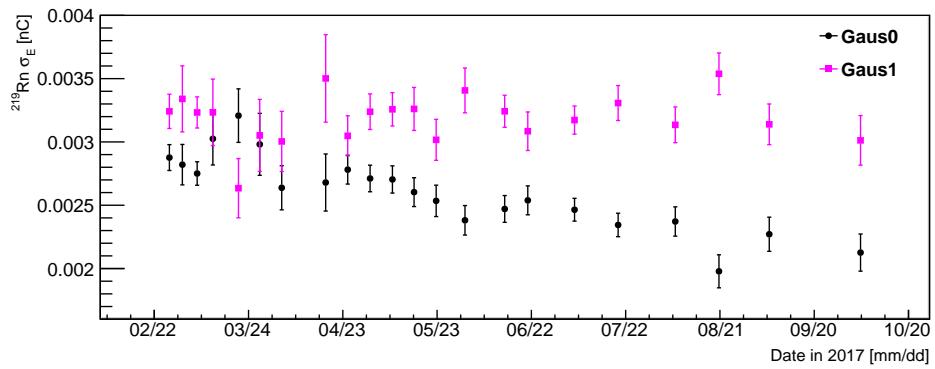


Figure 5.13: The 1σ width of two Gaussians fit to the ^{219}Rn energy spectrum versus time for the viton o-ring material sample. Black (circle): higher energy Gaussian, Magenta (square): lower energy Gaussian.

5.3 ^{227}Ac in the PROSPECT AD

Material compatibility testing determined that ^{227}Ac did not adsorb onto detector materials, confirming that it could be used as a uniformly distributed source. It will be noted here that ^{227}Ac spiked LiLS was added to a prototype detector that consisted of two stacked segments. This was done to determine that the ^{227}Ac did not degrade the performance of the liquid scintillator and that it did not introduce significant background. Initial results from the prototype concluded that it did neither and provided the last step of evidence needed before addition to the full-scale detector.

5.3.1 Spiking the LiLS

After concluding that ^{227}Ac would be added to the AD, the goal was to obtain a final activity of 0.01 Bq/segment. Assuming a total LiLS mass of 4600 kg and an active mass of 3939 kg implies a total ^{227}Ac activity of 1.8 Bq. The stock solution from which the LiLS would be spiked was the same stock that was used for the material studies, which had an activity of ~ 9.13 Bq/g on December 13, 2017, the day the spiking procedure was performed. This means that ~ 200 mg of the stock was needed to spike the LiLS.

In order to add ^{227}Ac to the total detector, a vial of spiked LiLS was added to a 55-gallon drum of LiLS prepared previously for detector filling. Before the detector was filled all drums were added to an ISO-tank and bubbled with nitrogen to ensure thorough mixing of the LiLS from all drums and the ^{227}Ac .

Spiking of the drum was done by adding the stock solution to an intermediate vial of production LiLS before spiking the vial that is added to the drum. This was done to offset the estimated 10 mg uncertainty of the balance used to weigh the vials and allows an assessment of the activity of the remaining vial. The procedure was also repeated in a second set of vials so that both vials could be measured.

The spiking procedure was performed using four vials, V0, V1, V2, and V3. The steps were:

1. Fill all four vials with production LiLS
2. Fill V1 and V2 with the ^{227}Ac spiked LiLS stock solution
3. Fill V0 with solution from V1 until desired activity is reached

4. Fill V3 with solution from V2 until desired activity is reached
5. Empty V0 into drum of production LiLS

See Figure 5.14 for a graphic of these steps along with Table 5.7 for a list of the weights of all solutions added and removed from the vials. Filling of all vials was performed using pipettes, therefore, some drops inevitably remained in the pipette at every step. Vials V1 and V2 were gently swirled after the addition of the ^{227}Ac spiked LiLS in an attempt to mix the solution.

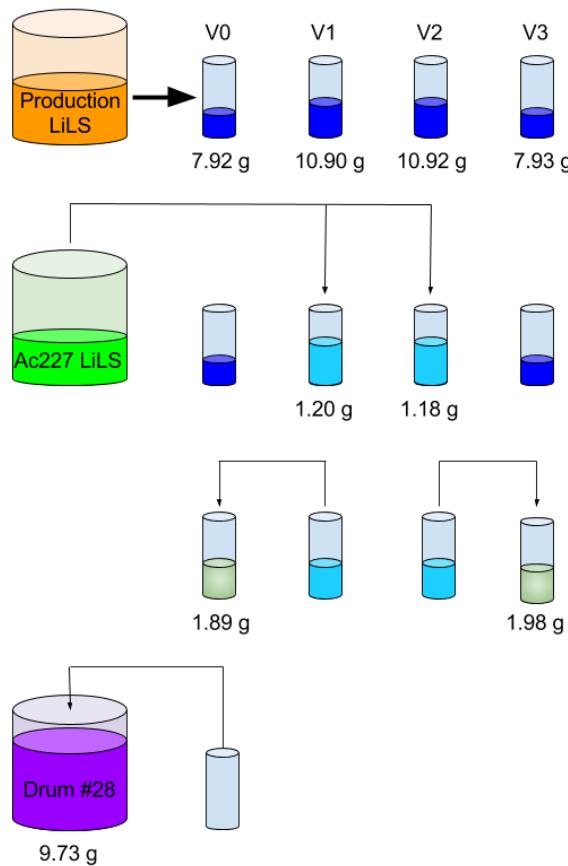


Figure 5.14: A graphic of the procedure used to spike the drum of LiLS with ^{227}Ac for filling of the AD.

The expected ^{227}Ac rate, A , in vials V0(V3) can be calculated as

$$A = C m_2 \frac{m_4}{m_1 + m_2} \quad (5.6)$$

where C is the activity of the ^{227}Ac spiked stock solution, 9.13 Bq/g, m_1 is the amount

	V0	V1	V2	V3
Production LiLS added, m_1 (g)	7.92	10.90	10.92	7.925
^{227}Ac spiked solution added, m_2 (g)		1.20	1.18	
Solution removed, m_3 (g)		2.047	2.09	
Solution added, m_4 (g)	1.885			1.98

Table 5.7: The weight of all solutions added and removed from the four vials for spiking of the LiLS with ^{227}Ac for filling of the AD.

Vial	Expected Activity (Bq)
V0	1.71
V1	9.10
V2	8.91
V3	1.76

Table 5.8: Expected ^{227}Ac activity in vials prepared for spiking the LiLS.

of production LiLS added to V1(V2), m_2 is the amount of the stock solution added to V1(V2), and m_4 is the amount of solution from V1(V2) that was added to V0(V3). The expected rate in vials V1(V2) is calculated as

$$A = C m_2 \left(1 - \frac{m_3}{m_1 + m_2} \right) \quad (5.7)$$

where m_3 is the amount of solution removed from V1(V2). The expected ^{227}Ac activity for each vial is listed in Table 5.8.

[MAKE PLOTS FOR THESE VIALS]

5.3.2 Data Set

PROSPECT began taking data in March 2018. The data set used for the ^{227}Ac analysis runs from March 5, 2018 - October 6, 2018, with a break from March 31, 2018 - April 17, 2018 when the detector was off for maintenance. The total runtime is 4048.9 hrs, with 4011.7 hrs of livetime data. 2293.7 hrs of the runtime is reactor on data, and 1755.2 hrs of reactor off.

During the data collection period used for this analysis, several PMTs exhibited abnormal behavior, including current instabilities, and are no longer in operation. Preliminary theories for the cause of this are that LiLS leaked into the PMT housings and damaged the voltage dividers, but this has yet to be confirmed. To account for

this all segments that ‘turned off’ during the data period are excluded in this analysis. The result is 90 active segments, as shown in Figure 5.15.

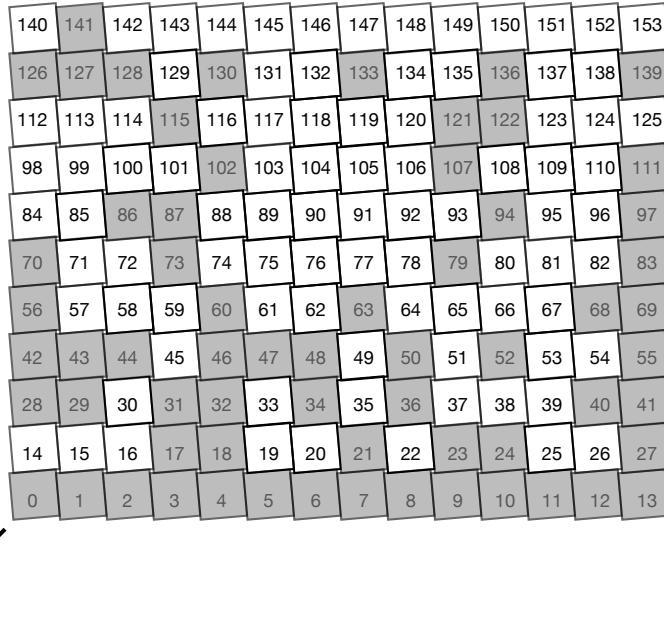


Figure 5.15: Graphic of 154 segments of the PROSPECT AD. Grayed out segments are those that ‘turned off’ during the data period and are excluded in this analysis.

5.3.3 Event Selection

RnPo events in the PROSPECT detector were found by looking at event clusters. A cluster is defined as a collection of events that occur within 20 ns of each other. Time coincident events were found by first looking for delay (Po) events. It was required that these events existed in a single multiplicity cluster and passed broad energy and PSD cuts outlined in Table 5.9.

Coincident prompt events were found by looking in a 5τ time window before the delay event, where τ is the lifetime of ^{215}Po , 2.57 ms. It was required that the highest energy event in a given cluster in that time window occurred in the same segment as the delay event and passed the energy and PSD cuts in Table 5.9. The distance between the prompt and delay event was also required to be less than 250 mm. Accidental prompt events were found by looking in a 5τ time window offset 10τ before the delay event. The same cuts applied to coincidental prompt events were applied to accidental prompt events.

Prompt Energy	$0.48 < E < 1.18 \text{ MeVee}$
Delay Energy	$0.61 < E < 1.18 \text{ MeVee}$
PSD	$0.16 < \text{PSD} < 0.36$
$\Delta z = z_{\text{delay}} - z_{\text{prompt}} $	$\Delta z < 250 \text{ mm}$
$\Delta t = t_{\text{delay}} - t_{\text{prompt}}$	$\Delta t < 5\tau \text{ ms}$

Table 5.9: Broad cuts used to find RnPo events, where τ is the lifetime of ^{215}Po , 2.57 ms.

In addition to energy, PSD, and position cuts a pileup veto was applied to all events. At the time of a trigger event all boards are signaled to begin a 592 ns acquisition window. Events arriving at the end of this window do not re-trigger the data acquisition system, thus causing truncated waveforms. In order to avoid using these truncated events, any cluster preceded by another cluster in a 800 ns window is vetoed.

The broad energy and PSD cuts listed in Table 5.9 are applied on a first pass analysis of the PROSPECT data. Changes in detector performance over time, including decreasing energy resolution and PSD, required a second pass of the data, in which the lower bounds of the energy and PSD cuts were changed to be 4σ lower than the mean of the distributions for a given time bin or segment. The PSD versus energy and ^{215}Po energy versus ^{219}Rn energy distributions, after accidental subtraction, can be seen in Figure 5.16.

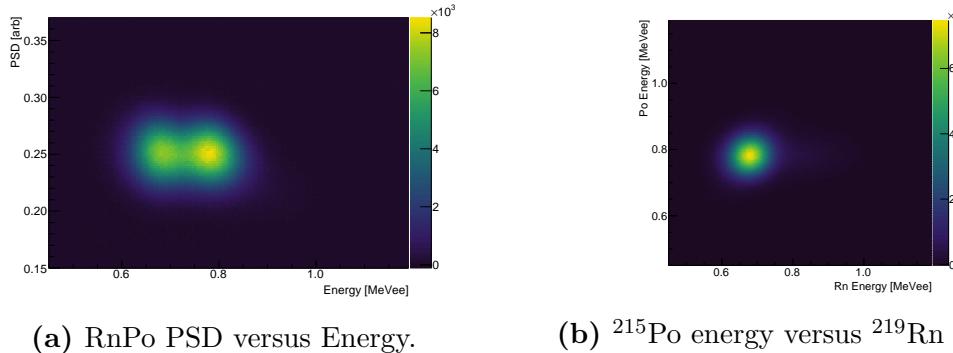


Figure 5.16: RnPo distributions for all cells integrated over all time.

5.3.4 Rate Calculation

livetime, dead time, efficiency, rate

5.3.5 Detector Stability Results

5.3.6 Volume Variation Results

CHAPTER 6

NEUTRINO OSCILLATION IN THE PROSPECT AD

CHAPTER 7

CONCLUSIONS

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APPENDIX A

SOME ADDITIONAL STUFF

Appendices go at the end.

APPENDIX B

MORE ADDITIONAL STUFF

This is another appendix.