An electron transparent proton detector for neutron decay studies

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We have developed an ultrathin (<100 nm), very strong polyimide foil which can span more than 6×6 cm² and is ideal for the fabrication of low energy proton detectors. We have produced a proton detector geometry in which protons incident on the foil with kinetic energies greater than about 25 keV produce, on average, more than ten secondary electrons in a conversion crystal evaporated on the back face of the foil. These secondary electrons can be “postaccelerated” and counted in a variety of detectors. The polyimide foils are much more durable than carbon foils previously used in similar detection geometries. LiF was chosen as the conversion crystal, which is relatively insensitive to exposure to air, improving their secondary electron yield under typical operating conditions. In addition, we describe the operation of a very simple, small scale proton accelerator and detector testing chamber capable of providing up to 10 kHz of beam with energies between 10 and 50 keV onto a biased target with a maximum ion contamination of 0.5%. © 2006 American Institute of Physics. [DOI: 10.1063/1.2186970]

I. INTRODUCTION

Measurements of the angular correlations between the decay products of free neutrons and the initial neutron spin direction provide high precision tests of the standard model without the nuclear structure corrections associated with measurements in higher mass nuclei. Recent progress in the development of cold neutron beam techniques and ultracold neutron (UCN) sources has made measurements of free neutron decay possible that are directly competitive with the neutron decay products of free neutrons and the initial neutron spin asymmetry in neutron decay because the detection of the β particle is relatively straightforward. The physics reach of free neutron decay measurements can be greatly extended by the simultaneous detection of both the decay β particle (782 keV end point energy) and the recoil proton (754 eV end point energy). Because angular correlations involving the β particle and the proton have differing sensitivities to various extensions to the electroweak standard model, one can probe physics beyond the standard model with proton detection capability. In addition to these correlations, implementing simultaneous detection provides one with a powerful tool for eliminating one of the largest systematic uncertainties in previous β asymmetry measurements: uncertainty in the absolute degree of neutron polarization. By simultaneously measuring β and proton asymmetries and taking their ratio, one determines a quantity which is independent of the neutron polarization to first order and can be used to extract the relevant physical parameters for neutron decay.

A scheme similar to the one we present here was attempted by the PERKEO II group utilizing carbon foils with MgO conversion crystals. Using this technique as a starting point, we have developed a detector technology which employs more durable and practical foils and utilizes a conversion crystal whose electron yield is not strongly degraded by exposure to air and whose performance is superior to MgO when implemented in practical experiments.

In this paper we present the construction and prototype tests of a proton conversion foil. In Sec. II, we outline the motivation for proton conversion foils as low energy proton detectors; next, some of the options we considered for foil substrates and conversion materials are presented (Secs. III and IV). In Sec. V we present measurements of the secondary yield. We then describe the proton accelerator and detector testing apparatus we constructed to evaluate our foils in Sec. VI. In Sec. VII we present calculations of the resultant proton detection efficiencies for our measured yields, and finally in Sec. VIII we present our conclusions.

II. THE PROTON DETECTION TECHNIQUE

The low energy of the recoil protons makes effective detection a technical challenge. With suitable acceleration, they can be detected with scintillators, silicon detectors, or microchannel plate detectors. Alternatively, the accelerated protons can be converted into secondary electrons.

Our approach is to permit the recoil protons from UCN...
decay to drift through a fiducial decay volume and then pass a grounded grid (see Fig. 1). The protons are then accelerated into a very thin (∼100 nm), biased foil (∼25 kV). The foil consists of a very strong substrate, coated with a high yield conversion crystal. As the protons exit the backside of the foil, they pass through a ∼10 nm thick layer of insulating crystal and liberate secondary electrons. These secondary electrons are then accelerated by the foil bias into a conventional electron detector: a gas counter, a plastic scintillator, or even a silicon detector. Electrons from neutron decay pass through the foil unhindered and can be detected by the same electron detector.

This technique allows one to use a β-decay spectrometer, such as the one already under construction by the UCNA collaboration at Los Alamos8 to measure the β-activity, such as the one already under construction by the electron detector. We found that 6F6F is a fluorinated polyimide which is synthesized through a condensation reaction between dianhydride and diamine monomers. (See Ref. 9 for an example of a synthesis process and see Ref. 10 for a summary of industry standard select properties, but note that our 6F6F probably derives its great strength from the high molecular weight synthesis process developed by one of the authors of this paper, Lozano.) 6F6F is soluble in polar organic solvents, such as tetrahydrofuran (THF) or N-methyl-2-pyrrolidone. The H2O contact angle for 6F6F foils still on a silicon substrate was 83.5±0.2°, indicating that the foils are slightly hydrophilic. These properties enable a simple and inexpensive manufacture method of very thin 6F6F foils: spin coating.

Our manufacture process is as follows: We first mixed a 1% solution, by weight, of 6F6F in high purity THF in a 20 ml borosilicate glass vial (I-CHEM C226-0020). The mixture was stirred with a teflon coated magnetic rod for 4 h and then allowed to sit over night. The mixture was then filtered twice through a 0.2 μm polytetrafluoroethylene (PTFE) membrane with a glass microfiber prefilter in a polypropylene housing (Whatman AV125EORG) and then stored in a second pre-cleaned vial.

We spun our foils onto 100 nm diameter silicon wafers. The polarity of the silicon (n or p) and crystal orientation of the surface were not critical. The surface of the wafers, as received from the manufacturer, was extremely hydrophobic. THF, a polar solvent, will not wet a hydrophobic surface, and thus, spin coating on the silicon wafers as delivered was not possible. We therefore treated the silicon surface with a mercury discharge lamp in ambient atmospheric conditions for 1 h. The UV light, in combination with the ozone generated by the UV interaction with atmospheric O2, terminated the surface with ∼10 Å of hydrophilic SiO2 and removed any residual hydrocarbon contamination.11 Before spin coating, the wafer was given a rinse with de-ionized water and blown dry with filtered research grade N2. The rinse step ensured that at least a monolayer of H2O covered the silicon surface. The 6F6F/THF solution was extracted from the storage vial into a 5 ml polyethylene/polypropylene syringe through a 0.25 mm polypropylene membrane.

TABLE I. Selected properties of prospective substrate materials taken from Refs. 29–31 and www.goodfellow.com. The minimum thickness of polyimide and 6F6F is from this research. The minimum thickness of polystyrene is from Ref. 12. Mylar® is biaxially oriented.

<table>
<thead>
<tr>
<th>Substrate</th>
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<tbody>
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<td>Cellulose nitrate</td>
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<td>6F6F</td>
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<td>320</td>
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(synthesized in one of our laboratories) offered the most promise due to its tensile strength, temperature stability, and ease of thin-film manufacture.

6F6F (CAS No. 32036-79-6 or CAS No. 29896-40-0 also known as 6FDA-4,4′-6FDA) is a fluorinated polyimide which is synthesized through a condensation reaction between dianhydride and diamine monomers. (See Ref. 9 for an example of a synthesis process and see Ref. 10 for a summary of industry standard select properties, but note that our 6F6F probably derives its great strength from the high molecular weight synthesis process developed by one of the authors of this paper, Lozano.) 6F6F is soluble in polar organic solvents, such as tetrahydrofuran (THF) or N-methyl-2-pyrrolidone. The H2O contact angle for 6F6F foils still on a silicon substrate was 83.5±0.2°, indicating that the foils are slightly hydrophilic. These properties enable a simple and inexpensive manufacture method of very thin 6F6F foils: spin coating.

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6F6F immediately parted from the wafer and floated onto the surface of the water. When sufficient polymer had floated off, we could use our other hand to take hold of the wafer under the surface of the water and continue parting the foil from the silicon. Once completely removed, the wafer was allowed to drop to the bottom of the water tank. We gently removed the 6F6F foil from the surface of the water by carefully positioning an edge of an appropriate frame next to the foil. Equally carefully, the edge of the foil was made to contact an edge of the frame, whereupon the foil would adhere to the frame. The frame was then slowly lifted from the water, perpendicular to the water surface, such that the foil spanned the opening area of the frame. The frame/foil combination was then heated at a moderate temperature (\(\sim 30 ^\circ C\)) until the water evaporated, and the foil relaxed in the frame to achieve a taut surface. The smallest foils made via this method were on circular frames with an open diameter of 0.64 cm. The largest foils were circular with a 10 cm diameter and mounted on square frames with an open side of length 5.6 cm. The strength of the resulting foils is impressive. In contrast to carbon foils, one can walk, talk, and breathe on 6F6F foils in open air without breaking them. They also routinely survived pump down in vacuum systems without requiring a lengthening of the roughing pump time.

We measured the mass thickness of the 6F6F foils by measuring the energy loss of a 45 keV proton beam through the foil. The proton accelerator is described in Sec. VI. The 6F6F foil was mounted so that it could be moved out of the beam, thus allowing the 50 mm\(^2\) silicon detector to be calibrated by the proton beam itself. Using a SRIM2000 simulation,\(^{13}\) we estimated the energy loss of a 45 keV proton beam through the 6F6F foil as a function of the foil thickness, taking into account the angular acceptance of the 50 mm\(^2\) silicon detector (\(\Delta \theta = 0.02 \text{ rad}\)). With this method, we measured the thickness of two foils to be 11.5 ± 1.5 and 8.6 ± 1.0 \(\mu g / \text{cm}^2\). The results are summarized in Table II. The two foils were manufactured using the same technique as described above, by the same experimenter, on the same day, indicating that the foil thickness is not easy to precisely reproduce.

### IV. THE CONVERSION CRYSTAL

Typical secondary electron emitters include metal oxides and alkali halides. Note that the yield from 6F6F, as that of most plastics, is relatively poor at, on average, 1 secondary per incident proton. In selecting a crystal, the overriding constraints are that the crystal must be able to be easily and cheaply deposited onto 6F6F, and must be able to survive exposure to the ambient atmosphere. Promising candidates include LiF, Al\(_2\)O\(_3\), hexatriacontane and diamond due to their high yield and surface stability. Note that although MgO has been considered in the past for similar applications, its surface is more reactive than that of NaCl to both H\(_2\)O and CO\(_2\);\(^{14}\) the secondary yield is not stable to exposure to these gases.\(^{15}\)

Al\(_2\)O\(_3\) has the distinct advantage that its surface is robust to atmospheric exposure. Alumina ceramic has an electron impact induced secondary yield between 5 and 7 at an incident energy of 1 keV. At the same energy, a sapphire crystal has a yield of 10.1.\(^{16}\) Deposition of Al\(_2\)O\(_3\) on 6F6F foils, however, may be difficult. Of the alkali halides, LiF has the most stable surface. At an incident energy of 600 eV, a vacuum cleaved crystal has a peak yield of 7.\(^{15}\) At an incident energy of 1.1 keV, a vacuum evaporated thin film has a peak yield of 8.\(^{17}\) Brief exposure of LiF to water vapor appears to reduce the electron affinity and increase the secondary yield.\(^{18}\) The principal advantage of LiF is the ease of thermal vacuum deposition. In this paper, we explore how well LiF will work as a conversion crystal. For comparison we have tested CsI, since it is a common photocathode material for x-ray and UV photon detectors.\(^{19}\)

In passing, we note two additional materials which we have identified, warranting further research due to their negative electron affinity (NEA): Hexatriacontane\(^{18}\) (HTC, \(n-C_{36}H_{74}\), CAS No. 630-06-08) and deuterium or cesium terminated diamond-like-carbon.\(^{20-22}\) NEA materials present the distinct advantage that low energy secondary electrons created in the crystal which reach the material surface are “pushed” out of the crystal instead of reflected back into the material by the more typically positive work function. In addition, these materials have long secondary electron attenuation lengths, meaning that secondaries created deep in the material have a greater chance of reaching the surface. From a practical perspective, both HTC and hydrogen terminated diamond are resilient to atmospheric exposure, and stable in UHV conditions. HTC can also be readily vacuum evaporated.

### V. MEASURED SECONDARY YIELD

We constructed proton conversion foils in the following manner. First, we manufactured 67.5 nm thick 6F6F films supported on stainless steel frames with a circular aperture of 6.3 mm. Next, we evaporated Al onto both sides of the 6F6F. The Al provides electrical conductivity to the foil, necessary for continuous proton conversion. Without breaking vacuum, we next evaporated the conversion crystal. Table III summarizes the deposition parameters. The deposition vacuum was maintained through a LN\(_2\) cold trap by an oil diffusion pump, filled with Santovac 5, backed by an oil-based rough-
ing pump. The aluminum source was a piece of 99.99% pure aluminum wire in a tantalum boat. The LiF source was a 99.9% pure powder in a tantalum boat. The CsI source was a single crystal, also in a tantalum boat. All sources were properly out-gassed before evaporation. Immediately following the evaporation, all foils had a mirrorlike quality. After 5 to 10 min of exposure to air, however, the CsI foil rapidly acquired an opaque blue appearance. After a few hours of exposure to the atmosphere, the LiF foils became crinkly. Once placed in an evacuated chamber, however, the LiF foils regained their mirrorlike appearance, indicating that either the 6F6F or the LiF layer absorbed atmospheric gases or water vapor. The CsI foil never lost the blue color, indicating a permanent change most likely due to water adsorption.

Each detector was exposed to a proton beam with a kinetic energy between 20 and 45 keV and at a rate between 100 and 1000 Hz. At each energy, the proton detectors were biased to −20 kV. The liberated secondary electrons were accelerated into a silicon detector with an efficiency of nearly 100% (see the discussion at the end of this paper). The pulse height histogram of the pulses from the silicon detector was then recorded with an Ortec PC-based multi-channel analyzer. The extraction and focus voltages of the electrode structure were adjusted to maximize the mean of the pulse height distribution (see discussion at end of this paper). Figure 2 presents representative spectra for an uncoated 6F6F foil, a CsI coated foil, and a LiF coated foil, at an incident proton energy of 45 keV. The first peak is due to electronic noise; the additional distinct peaks in the spectrum represent additional electrons liberated by the foil and detected by the silicon detector. Each secondary had an energy of 20 keV. The absolute normalization of the spectrum are scaled for clarity.

The pulse height spectrum in the silicon detector when prototype uncoated and CsI and LiF coated foils were exposed to a 45 keV proton beam. The first peak in the spectrum was due to electronic noise; the additional distinct peaks in the spectrum represent additional electrons liberated by the foil and detected by the silicon detector. Each secondary had an energy of 20 keV. The absolute normalization of the spectrum are scaled for clarity.

Second, to account for a proton field reflection, we let

\[ P_{\text{lab}}(n) = e^{-(n-\mu)^2/2\sigma^2}, \tag{2} \]

where \( n \) is the number of secondary electrons observed in the silicon detector, \( \eta \) is the probability that a 20 keV electron will backscatter (0.17), and \( P_{\text{lab}}(n) \) is the probability that \( n \) electrons will be liberated by one incident proton. We have considered two possible forms of \( P_{\text{lab}}(n) \): first, neglecting the possibility of a proton field reflection, we let

\[ P_{\text{lab}} = e^{-n(1-\eta)^N}, \tag{1} \]

The rebinned, measured electron spectra for 25 keV incident proton energy plotted with the two nonlinear fits accounting for electron backscatter alone and electron backscatter with proton reflection from the accelerating fields.

\[ P_{\text{obs}}(N) = \sum_{n=N}^{\infty} \frac{n^N}{n!(n-N)!} (1-\eta)^N \eta^{n-N} P_{\text{lab}}(n), \]
The fit parameters for the prototype LiF foil.

<table>
<thead>
<tr>
<th>Incident energy (keV)</th>
<th>Backscatter</th>
<th>Proton bounce</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>( \mu )</td>
<td>( \sigma )</td>
</tr>
<tr>
<td>25</td>
<td>10.3</td>
<td>5.6</td>
</tr>
<tr>
<td>35</td>
<td>12.2</td>
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</tr>
<tr>
<td>45</td>
<td>13.3</td>
<td>6.7</td>
</tr>
</tbody>
</table>

\[
P_{\text{lib}}(n) = (1 - b)P(n) + b \sum_{j=0}^{n} P(j)P(n-j),
\]

where \( b \) is the probability that a proton will return to an active area of the proton detector, and \( P(n) \) is the probability that a single proton traverse or impact will liberate \( n \) electrons. In this second case, \( P(n) \) is chosen to be

\[
P(n) = e^{-(n-\mu)^2/2\sigma^2}.
\]

For the two forms of \( P_{\text{lib}}(n) \), \( \mu, \sigma, \) and \( b \), are estimated on the basis of a nonlinear least-squares fit to the rebinned data. Figure 3 plots the 25 keV rebinned data and the two fits. Table IV lists the fit parameters at each proton energy. Notice that the proton field reflection fit accounts for the distribution much better than the backscatter fit alone suggesting that we saw evidence for proton field reflection. The proton return probability \( b \) is consistent with our expectations at 25 keV, but not at 35 or 45 keV. Other rare events may contribute to the probability tail, such as proton impact on the focusing electrode structure, an occurrence that would mimic a proton field reflection. Note that the difference in the estimated yield is less than 3% between the two fitting cases. Note also that to an accuracy of 5%, the average yield can be estimated on the basis of a Gaussian fit to the observed spectrum: \( \mu = \mu_{\text{obs}}/(1 - \eta) \). The width of the Gaussian distribution is much broader than expected from counting statistics alone: \( \sigma = 1.7\sqrt{\mu} \). This is probably due to energy and angle straggle of the protons through the foil.

We performed a simple aging test for the LiF prototypes in the following manner. Two prototype detectors, each coated with 50 Å of Al followed by 60 Å of LiF at a deposition rate of 0.25 Å/sec, were manufactured simultaneously. The secondary yield of one foil at 45 keV was immediately measured. After one week storage in a container filled with ambient air, the yield of the other foil at 45 keV was measured. Figure 4 plots the spectra. The average yield (corrected for electron backscatter) fell from 12.5 to 6.5. The reduced variance, \( \sigma^2/\mu \) increased from 1.7 to 2.2. Clearly, despite the stability of the LiF surface, care must be taken for long term storage of the LiF foils.

**VI. A SIMPLE PROTON ACCELERATOR**

To test the prototype conversion foils and obtain the data presented in Sec. VI, we built a simple proton accelerator (see Fig. 5) capable of providing protons at a rate of 0–10 kHz with a kinetic energy up to 50 keV and a maximum ion contamination of less than 0.5%. The accelerator consisted of the following elements in beam order: a \( \mathrm{H}_2 \) variable leak and gas handling system, a 100 W light bulb filament and associated power supply, a 60 kV electrostatic acceleration column, a dry pumping station, an electromagnet to separate the ion species, and a series of apertures to define the entrance and exit path of the beam. Note that due to the very low count rate requirements, neither steering nor focusing elements were necessary or used. Each side of the acceleration column could be biased independently of the other to \( \pm 40 \) kV. This bias scheme allowed the proton energy, determined by the potential difference across the acceleration column, to be different than the foil bias. Except for the proton source, all electronics, such as pumps, the magnet, and the electron detectors were operated at ground potential. The end of the accelerator was connected to a detector testing chamber.

Protons were liberated from \( \mathrm{H}_2 \) by the ionizing action of electrons emitted by a hot, 100 W light bulb filament suspended between two feedthroughs in the middle of a 5 cm long, 11 cm OD Al spool piece (see Fig. 6). The upstream end of this spool piece was closed off by a brass plate, opened at the edges for vacuum pump out. The downstream end was limited by an Al aperture, directly connected to the acceleration column. The exit aperture and filament were held at a fixed potential by a high voltage power supply. The cylindrical walls and brass plate were positively biased with a 300 V battery with respect to the filament and exit aperture. The bias served to extract electrons from the filament and provide sufficient energy for \( \mathrm{H}_2 \) ionization in addition to focusing the ions through the exit aperture. The filament was powered by a variac connected to a 50 kV, 250 W, isolation transformer.

Following the acceleration column, the ions passed through the exit aperture. The ions were then focused in a 60 kV cylindrical electrostatic column through the exit aperture. The ion species were separated by a magnetic field followed by a series of apertures to define the exit path of the beam. The detector was placed at the exit of the accelerator column. The beam was focused by focusing the ions through the exit aperture. The filament was powered by a variac connected to a 50 kV, 250 W, isolation transformer.

![Aging Study Spectrum](image)

**FIG. 4.** The spectra from a LiF prototype tested immediately by a 45 keV proton beam and tested after one week exposure to atmospheric conditions.

![A schematic of the proton accelerator](image)

**FIG. 5.** A schematic of the proton accelerator (secondary electron detection apparatus not shown); not to scale.
through two 3 mm diameter circular apertures punched in a 0.25 mm thick tantalum foil placed 30 cm apart at a sufficient distance from the bending magnet to ensure that the fringe field from the bending magnet was less than the Earth’s field. After the apertures, the protons passed through a 15 cm diameter, 2.5 cm high, cylindrical bending magnet chamber by making a 30° turn. The magnet chamber, biased at up to +40 kV, was insulated from the grounded magnet pole tips by a 6 mm thick sheet of ultra high molecular weight (UHMW) polyethylene. UHMW polyethylene has the highest known dielectric strength of the commercially available plastics. After the bend, the ions traveled through another two 3 mm diameter apertures located 100 cm apart.

A. Tests of the proton accelerator

At a fixed proton source pressure ($5 \times 10^{-5}$ Torr), filament power (5 W), and acceleration potential (40 kV), we recorded the count rate in a channeltron detector, positioned immediately downstream from the last aperture, as a function of the magnetic field in the bending magnet. A representative scan is plotted in Fig. 7. By also introducing He gas, we were able to positively identify four peaks due to $H_2^+; H^+_3; H_3^+$, and a peak due to the dissociation of $H_2^+$ into $H$ and $H^+$. Assuming a circular path in the bending magnet, and an equal sharing of momentum in the dissociation case, one would expect the magnetic field of the peaks $H_{dis}; H^+; H^+_2; H^+_3; He^+$ to be in the ratio $1/\sqrt{2}:1/\sqrt{2}:3/2$. We find the ratio to be $0.682:1:1.43:1.76:2.04$. The deviation from the expected ratio is less than 3%.

We realized the best performance when hydrogen gas was not leaked into the vacuum. See Fig. 8 for a scan at 25 keV. The operation proceeded as follows: first, we flooded the vacuum system with $H_2$ up to a pressure of $1 \times 10^{-3}$ Torr. We then pumped out the $H_2$ to a pressure of $1 \times 10^{-5}$ Torr before operating the accelerator. When the accelerator was used continuously, $H_2$ “treatments” were needed approximately once a day. A very conservative limit on the ion species contamination can be determined by dividing the peak height by the value of the lowest trough at fields above the $H^+$ peak. We put an upper bound on the ion contamination of 1.2% at 10 keV, 0.3% at 25 keV, and 0.1% at 40 keV.

B. The detector testing chamber

We built a detector testing chamber (see Fig. 9) which had four critical elements: an inner beam pipe to carry the beam pipe potential from outside the detector chamber...
through the inside of the chamber to the surface of the biased foil, a vertically transferrable silicon detector (25 mm$^2$ active area) to monitor the proton beam quality, a horizontal and vertical translation stage to scan the spatial response of the foils, and electrode elements to focus secondary electrons created by the conversion foil into another silicon detector (50 mm$^2$ active area) located at the end of the chamber. The 25 mm$^2$ detector was mounted on a copper cold finger connected to a LN$_2$ dewar in the vacuum system. Brass collars at the ends of the inner beam pipe minimized beam defocusing by ensuring that electric potential gradients were far from the center of the beam path. The chamber could achieve an ultimate pressure of $5 \times 10^{-7}$ Torr.

C. Secondary electron focusing elements

The energy and angle distribution of secondaries produced by proton passage through LiF or CsI is unknown. For electron$^{23}$ and ion$^{24}$ impact on Al, a cos $\theta$ weighted distribution has been observed. X-ray impact on LiF gives a peak secondary energy of $\sim 3$ eV and a maximum secondary energy of 16 eV.$^{25}$ It is expected that the energy and angle distribution of the secondaries is independent of the type or energy of the incident radiation.$^{26}$ We designed the electrode structure to optimally focus secondaries produced in a 3 mm radius circle (twice the radius of the final beam aperture) with a cos $\theta$ weighted angular distribution with energies between 0.1 and 50 eV into the 50 mm$^2$ silicon detector.

For a small diameter $\sim 20$ kV biased foil, our design consisted of an extraction plate (biased to $-19$ kV), which ensured optimal collection and prefocusing of the secondary electrons, an acceleration region, which accelerated the electrons to their full energy, and an Einzel lens, consisting of a ground cylinder, a biased cylinder ($-15$ kV), and another ground cylinder, which provided the final focus. A labeled cutaway view of the equipotential surfaces is presented in Fig. 10. The trajectory path of 0.1 and 50 eV electrons is presented in Fig. 11. A Monte Carlo simulation of the transport of $10^4$ electrons calculated a collection efficiency of 100% for both 0.1 and 50 eV electrons. For larger foils, we added a 2 mm pitch grid biased at $+200$ V with respect to the foil between the foil and the extraction plate. This grid

![Diagram of equipotential surfaces](image)

**FIG. 10.** A cutaway view of the equipotential surfaces near the focus elements as calculated by RELAX3D. Each equipotential surface represents 5% of the potential of the proton foil (typically $-20$ keV). The inner beam pipe, located immediately to the left of the proton detector, and the 200 V grid used for larger detection foils, are.

![Diagram of trajectory paths](image)

**FIG. 11.** The trajectory of 100 secondary electrons emitted from the proton foil at 0.1 and 50 eV. Note that the active area of the silicon detector was between $\pm 0.4$ cm from the beam center and along the beam axis at 28 cm. These simulations indicate the electrode structure had a collection efficiency of 100%.
There was an error in the original text. The correct text should read:

FIG. 12. A Monte Carlo energy deposition histogram in the MWPC due to secondary electrons from the proton conversion foil and due to electrons from neutron decay.

VII. SIMULATED PROTON COUNTING EFFICIENCIES

Any electron detector could be used to observe secondary electrons (SEs) from the proton conversion foil; a multiwire proportional counter (MWPC), however, offers perhaps the simplest way to distinguish $\beta$-decay electrons from secondary electrons. With a series of Monte Carlo simulations we have studied the expected response. The model MWPC consists of a 6 $\mu$m thick mylar entrance window followed by a 2.0 cm thick active region filled with 100 Torr of neopentane ($C_5H_{12}$, $\rho = 3.8 \times 10^{-4}$ g/cm$^3$) and a 6 $\mu$m thick mylar MWPC exit window. Neopentane was chosen as a prospective MWPC fill gas due to its high ion pair yield (estimated to be 1 ion pair/88 eV deposited$^{[25]}$) and low average $Z$. For analysis simplicity, we assume that the ion collection efficiency in the MWPC is 100%.

We simulated the impact of 3000 protons on a 30 kV biased conversion foil backed by the model MWPC. For each proton impact on the conversion foil, the number of liberated secondaries was sampled from a Gaussian distribution with a mean of 20 and a standard deviation of 7.6, consistent with the measured statistical distribution. Using a PENELOPE simulation,$^{[24]}$ the total energy deposition in the active region of the MWPC was modeled for this number of normal incident 30 keV electrons. For comparison, the energy deposition by electrons from $\beta$ decay in the UCNA experiment was also modeled with PENELOPE. Note that this simulation includes the effect of a magnetic field expansion region in the decay spectrometer which parallelizes the electron trajectories and reflects backscatter electrons back into the MWPC. Figure 12 plots the results of these simulations. We find that with an energy deposition threshold of 75, 100, and 125 keV, protons will be detected with an efficiency of 97%, 94%, and 90% respectively. If a 125 keV energy cut is applied to separate $\beta$-decay electrons from protons, we anticipate that only 0.02% of the $\beta$-decay electrons will be misidentified as protons in the MWPC. In the UCNA project, a plastic scintillator will be implemented behind the MWPC for $\beta$-decay detection. Thus, a proton signal in the MWPC will not be miscounted as a $\beta$-decay electron.

Lastly, the energy loss and backscatter of $\beta$-decay electrons due to passage through the conversion foil was simulated. We find that only 0.12% of the incident flux is backscattered from the conversion foil with an energy greater than 50 keV and only 0.02% is backscattered with an energy greater than 75 keV. The same simulation indicates that 99% of the transmitted $\beta$-decay electrons will lose less than 1 keV, 99.9% will lose less than 3 keV, and 99.99% will lose less than 5 keV. By comparison, 99.9% of the $\beta$-decay electrons will lose more than 1 keV on passage through the MWPC.

VIII. CONCLUSIONS

The development of electron transparent, low energy proton detectors holds the promise of greatly extending the physics reach of neutron decay experiments currently designed to detect only $\beta$ particles following neutron decay. By suitably spin coating a fluorinated polyimide (6F6F) on a silicon wafer we have been able to fabricate 600 Å thick plastic substrates which can span more than $6 \times 6$ cm$^2$. At neutron decay energies, electrons are nearly unaffected by passage through such material. Perhaps more critically, these foils are substantially stronger and more durable than carbon foils used for similar applications in the past. The addition of a thin layer of LiF enables the conversion of 25 keV recoil protons into ~10.3 secondary electrons, which when instrumented in a suitable high voltage stack, can be readily detected in a neutron decay spectrometer. Given the measured yield, we estimate a proton detection efficiency of nearly 97% when instrumented in front of a simple multiwire proportional counter. Furthermore, we have constructed a small scale proton accelerator optimized for proton detector development.

As a final note, we believe that the small scale proton accelerator we have developed for the proton detector development work may be useful to others working on related problems.

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