Characterization of the CATRiNA neutron detector system

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Abstract

The Compound Array for Transfer Reactions in Nuclear Astrophysics (CATRiNA) is an array of 16 liquid deuterated-benzene (C₆D₆) neutron detectors. The fast-response and pulse-shape-discrimination (PSD) capabilities of liquid scintillators allow for the extraction of neutron energies via time-of-flight and for the identification and separation of neutron/gamma interactions. In addition to these properties, deuterated liquid scintillators have a structured pulse-height spectrum with promising capabilities for neutron spectroscopy. In this paper, we report on the development and characterization of the CATRiNA neutron detector system at Florida State University (FSU). Monte-Carlo simulations and measurements with neutron and γ -ray sources, as well as mono-energetic neutrons (E_n = 2.3 - 7.3 MeV) from in-beam measurements using the ⁷Li(p,n) reaction, were performed to characterize the detectors. CATRiNA is envisioned to be used to measure several nuclear reactions like (³He,n), (d,n), (p,n), relevant for nuclear structure and nuclear astrophysics studies.

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1 1. Introduction

Neutron detection systems are becoming increasingly necessary in nuclear structure and nu-3 clear astrophysics research. Radioactive beam 4 facilities are probing exotic neutron-rich and 5 neutron-deficient nuclei, where the detection of 6 neutrons in reactions using radioactive beams pro-7 vides additional information about the properties 8 of such exotic nuclei. Ideally, neutron detection 9

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systems are expected to perform neutron spectroscopy and use neutrons to 'tag' other reaction by-products. Efforts in the field have led to the development of large sets of neutron detector systems, for example Refs. [1–3]. Organic scintillators with hydrogen-based compounds (e.g. Anthracene, Stilbene, Benzene) are traditionally the standard for neutron detection due to the high cross section between neutron-proton (n-p) elastic scattering. In particular, liquid scintillators like the NE213 (C₈H₁₀) [4], are useful due to their fast re-

sponse time (about a few ns), capability to distin- 51 21 guish neutron/gamma (n/γ) interactions within the 52 22 scintillating material based on the signal's pulse 53 23 shape, and for extraction of the neutron energy us-24 ing the Time-of-Flight (ToF) technique. 25

Deuterated-benzene (C_6D_6) scintillators have re-26 cently generated great interest in basic nuclear 27 physics research and related applications [1, 5-28 7]. Deuterated-benzene scintillators have equiva-29 lent response time and pulse-shape discrimination 30 (PSD) capabilities to the standard hydrogen-based 31 scintillators (like the NE213), but with a distinc-32 tive peak observed in the light-output spectrum ob-33 tained by integrating over the scintillator's light-34 vield signal when responding to incident mono-35 energetic-neutrons. This behavior is due to the 36 asymmetry in the cross section of n-d elastic scat-37 tering, peaking at backward angles with a maxi-38 mum energy transfer of 8/9 incident neutron en-39 ergy (E_n) [5, 6]. This peak is not distinguishable 40 in hydrogen-based detectors due to the isotropic 41 cross section for n-p elastic scattering [5, 6]. The 42 response of the deuterated scintillators to the light-43 output spectrum opens the possibility to identify 44 mono-energetic neutron groups without relying ex-45 clusively on ToF information. 46

Deuterated scintillator arrays are being used, 47 for example, at Canada's particle accelerator cen-48 tre (TRIUMF) and University of Michigan (UM). 49 TRIUMF is using the Deuterated Scintillator Ar-50

ray for Neutron Tagging (DESCANT), which consists of 70 deuterated benzene liquid scintillators [1]. DESCANT has been designed to couple with the γ -ray spectrometers TIGRESS and GRIFFIN to enable studies of fusion-evaporation using neutronrich nuclei by neutron tagging. Additionally, DES-CANT will be used to study neutron-emission probabilities of short-lived states above the neutron threshold, which are important for nuclear astrophysical studies. At UM, the University of Michigan Deuterated Scintillator Array (UM-DSA), an array of several deuterated-benzene detectors has been developed [7]. It has been shown that the UM-DSA can be used for neutron spectroscopic studies by only relying on the back-scatter peak information on the light-output spectra to extract neutron energies.

In this paper we describe the development and characterization of the Compound Array for Transfer Reactions in Nuclear Astrophysics (CATRiNA) at Florida State University. CATRiNA is a neutron detector array composed of 16 liquid scintillator detectors based on deuterated-benzene as scintillating material, which has been designed to study nuclear reactions relevant for nuclear structure and nuclear astrophysics studies.

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Figure 1: Schematics of the EJ-315 CATRiNA detectors [8]. The C_6D_6 liquid material is contained in a 4" by 2" Al cylinder.

77 2. CATRINA

78 2.1. Detectors and Design

The CATRINA detectors are of the type EJ-315 109 79 manufactured by Eljen Technology [8]. Each EJ- 110 80 315 detector consists of C₆D₆ scintillating mate- 111 81 rial, which is contained in a 4" diameter \times 2" ₁₁₂ 82 deep cylinder embedded in an aluminum capsule 113 83 coated with TiO₂ to reduce scintillating light loss 114 84 and a Pyrex@glass window on the optical end. A 115 85 nitrogen gas bubble occupies 3% of the volume 116 86 within the the aluminum capsule along with the liq- 117 87 uid scintillating material to reduce light quenching. 118 88 An acrylic light guide couples the glass window 119 89 to a ET Enterprise 9821B Photo-Multiplier Tube 120 90 (PMT) [9]. The PMT has a spectral range of 285- 121 91 630 nm and the EJ-315 detector has a maximum 122 92 emission of 450 nm. A schematic of the detectors 123 93 is shown in Fig. 1. 94 124

A 16 channel WIENER MPOD Minicrate High-¹²⁵
Voltage (HV) power supply [10] provides about ¹²⁶
-1700V to each PMT and can be controlled re-¹²⁷

motely. This allows for fast access to monitor and adjust the HV of the detectors during experiments.The anode output signal from the PMT is sent to a data acquisition system for data analysis.

2.2. Data Acquisition System (DAQ)

For the present characterization, CATRiNA used an analog data acquisition system where signals from the anode of each detector's PMT are sent to a 16 channel Mesytec MCFD module [11]. The MCFD module gives an amplitude and a fast logic signal for each detector. The amplitude signals are delayed by 10.6 m ribbon cable. Both amplitude and logic signals are then split and connected to the two 16-channel banks of a Mesytec MQDC module [12]. The MQDC module allows to specify charge integration time in each bank for PSD, which range from 24 to 219 ns. A logic OR of the fast logic signals is used as a master-trigger and it is sent to a VME controller. The individual fast logic signals are further delayed by about 100 ns and can be used as individual inputs to a CAEN V775 TDC [13] module. To allow coincidence measurements of neutrons, charge particles and/or gammarays, this delayed signal can also be used to couple CATRiNA to other detector systems since scintillator detectors are much faster than conventional silicon and germanium detectors. An independent timing-reference signal is derived from the accelerator radio-frequency (RF) signal, which has a 165 ns period and is used as a common stop for the

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Figure 2: Layout of the data acquisition system used with CATRiNA. All 16 detector's anode signals are sent to the MCFD module. The MCFD splits the input signal and sends as an output the timing and amplitude component to the MQDC module for independent pulse integration. Timing signals from the MCFD are also sent to a CAEN TDC module. The MCFD's OR signal is fed to a LOGIC unit in order to 141 generate a master trigger for the VME controller.

TDC module. The schematics of the electronics 128 is shown in Fig. 2. 129

3. Detector Characterization 130

3.1. γ -ray source measurements 131

The light response of the detectors was cali- 149 132 brated using two gamma-ray sources, ⁶⁰Co and ¹⁵⁰ 133 ¹³⁷Cs. Due to a low photo-peak cross section of liq- 151 134 uid scintillators, maximum energy transfer is dom- 152 135 inated by the Compton effect [4]. The gamma- 153 136 rays from the radioactive sources will Compton 154 137 scatter with the electrons of the scintillating ma- 155 138 terial in the detector, producing a Compton edge 156 139 when maximum gamma energy is transferred to 157 140



Figure 3: Experimental gamma-ray spectra measured with 137Cs and 60Co sources used for calibration of the CATRINA detector's light-output.

the electrons. A standard calibration unit, "keVee" (keV-electron-equivalent), has been adopted. Here, 1,000 keVee equals the total light-output produced by a 1,000 keV recoiling-electron [14]. A spectrum of the Compton edges measured in the detectors with ⁶⁰Co and ¹³⁷Cs gamma-ray sources is shown Fig. 3.

The location of the Compton edge is broadened due to the detector's size and resolution. Placement of the Compton edge was done by simulating the response of the detector with and without detector resolution [15]. For this, a GEANT4 simulation was developed to understand the light-output of the CATRiNA detectors Simulations using the G4OpticalPhysics, [16]. G4EmStandardPhysics, HadronElasticPhysicsHP, and G4DecayPhysics package consisted of a 2"

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Figure 4: MC simulated spectra for 60 Co and 137 Cs gamma-ray sources on C₆D₆ material using GEANT4. These simulations do not include detector resolution.

long by 4" diameter cylinder of C_6D_6 material, $_{174}$ 158 embedded in a 1.5 mm thick aluminum outer cas-159 ing. Simulated scintillation photons from ⁶⁰Co and 175 160 ¹³⁷Cs gamma-ray sources were compared to the ex- 176 161 perimental measurements. The simulations of the 177 162 gamma-ray source spectra without detector resolu- 178 163 tion are shown in Fig 4. A simulation of a ¹³⁷Cs ₁₇₉ 164 spectrum is shown in Fig. 5 with and without de- 180 165 tector resolution. Following the procedure of Ref. 181 166 [15], the location of the Compton edge was placed 182 167 at the point at which both spectra intersect at the 183 168 upper edge of the peak (indicated by the black ar- 184 169 row in Fig. 5), usually reported as a percentage of 185 170 the maximum height of the spectrum with resolu- 186 171 tion. In this case, the Compton edge was placed at 187 172 80% maximum height. 188 173



Figure 5: MC simulated spectrum of ¹³⁷Cs using GEANT4 with and without detector resolution. The intersection at the upper edge of the two peaks (black arrow) is chosen as the location for the Compton edge, which is about 80% maximum height.

3.2. ²⁵²Cf source measurements

A pulse-shape discrimination analysis using the charge integration method [17] was performed to separate neutron from gamma-ray interactions in the detectors. A spectrum of the scintillator's light-output can be obtained by integrating over the pulse signals from the PMT's anode. The rise and decay times of the light-output signal emitted from the PMT's anode depends entirely on the scintillating particle type [4]. The decay time of the pulse produced by a gamma interaction is faster in comparison to that of a neutron pulse signal. This is shown in Fig. 6 where different timing gates used for the analysis are also shown. The gates labelled Short, Long and Tail correspond to the integrated



Figure 6: Example of pulse signals emitted by the PMT which are produced by neutron and gamma interactions. The different gate integration times used to discriminate n/γ interactions ²¹⁷ are also shown. ²¹⁸

area (V·ns) and are used to perform PSD in or- 220 189 der to distinguish neutron and gamma interactions. 190 The integration times for the Short and Long gates ²²¹ 191 are programmed into the two banks of the MQDC 192 module. The Short gate is chosen to integrate the 193 223 rise of the pulse, while the Long gate integrates the 194 224 whole pulse. The Tail integration gate is obtained 195 225 by subtracting the Short gate from the Long gate. 196 226 To find the most favorable integration times for 227 197 the MQDC module, a ²⁵²Cf neutron source was 198 used. By varying the integration times, we found 229 199 that the optimal values were 32 ns and 187 ns for 200 the Short and Long gate, respectively. The PSD 230 201 plots shown in the present study were generated 231 202 by taking the ratio of the gates (Short/Long) and 232 203 plotting it versus the Long integration value of the 233 204

pulse. A typical example of a PSD obtained with 205 the 252 Cf source can be seen in Fig. 7(a) where 206 neutron and gamma-ray interactions are separated 207 in two well defined groups. It is important to note 208 that the integration times strongly depend on the 209 integration window. For example, if the window 210 opens before the pulse rise, additional time must 21 be given to the Short and Long gate to compensate 212 for this delay. 213

To quantify the performance of the detectors when separating neutron and gamma interactions, a Figure-of-Merit (FOM) is used. The FOM for each detector was obtained from the 1 dimensional projection of the Short/Long parameter at a given value for the Long parameter, as shown in Fig. 7(b). The FOM is defined as [18, 19]:

$$FOM = \frac{\Delta_C}{\Delta_n + \Delta_\gamma} \tag{1}$$

where Δ_C is the separation between the centroids of the neutron and gamma peaks and Δ_n and Δ_γ are the full-width at half-maximum (FWHM) of the peaks, respectively. A FOM greater than 1 is indicative of a good separation. For the test with a ²⁵²Cf source, a FOM of 1.3 for an energy threshold of 250 keVee was achieved for all 16 CATRiNA detectors.

3.3. Measurement of the $^{7}Li(p,n)^{7}Be$ reaction

Further study of CATRiNA's performance was done by measuring the 7 Li(p,n) 7 Be reaction at the John D. Fox accelerator laboratory at FSU.

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Figure 7: (a) Typical PSD plot obtained with a 252 Cf source. 252 Here, the ratio of the gates (Short/Long) is plotted against the 253 Long gate spectrum and is used for n/ γ identification. The 254 neutrons appear to the left due to their longer decay time, which corresponds to a larger value for the Long integration. (b) Extraction of FOM. Figure-of-Merit are extracted by projecting the PSD plot onto the x-axis with a given threshold and 257 calculating the width and separation of the n/ γ groups accord- 258 ing to Eq. 1. For the data shown, a 250 keVee threshold was used and a FOM of 1.3 was extracted, indicating a good n/ γ separation.

Experimental data on the excitation function of 234 7 Li(p,n)⁷Be reaction and the relative cross section 235 of the neutrons from the ground and first excited 236 states of ⁷Be as a function of proton energy is available in the literature [20] and was used for comparison with the data measured with CATRiNA allowing for the extraction of an experimental efficiency curve of the CATRiNA detectors. In the present experiment, a proton beam from the FN Tandem accelerator was tuned at different energies from 4 to 9 MeV, in 1 MeV increments. The proton beam was pulsed to short bunches of around ~ 2 ns duration with 82.5 ns in between pulses. The proton beam was used to bombard a ~ 200 μ g/cm² thick ⁷Li target, which was prepared by evaporating pure ⁷Li onto a polyvinyl formal resin (formvar) backing at 10-20 Å/s and transported under vacuum to minimize oxidation. Contributions from formvar backing can be neglected since it evaporates once the 252 beam strikes the target. Neutrons from the ground state and first excited state in ⁷Be populated via the (p,n) reaction were measured in the detectors. The ground state Q-value for the reaction is Q = -1.64MeV. The first excited state is located at $E_{ex1} = 429$ keV above the ground state. Contributions of the neutrons from the second and higher excited states in ⁷Be are negligible in the measurements due to the high excitation energy of the second excited 26 state in ⁷Be ($E_{ex2} = 4570 \text{ keV}$). 262

²⁶³ In the present configuration, the detectors are





Figure 9: Experimental setup for the measurement of the 7 Li(p,n)⁷Be at the John D. Fox Accelerator Laboratory at FSU.

Figure 8: Schematic of the CATRiNA array. The 16 detectors are placed inside a plastic case that can slide and lock to the frame.

285 placed in two layers, in an array of aluminum ma-264 286 terial as shown in Fig. 8. Each layer can hold eight 265 detectors with 5° spacing in between. Individual 266 detectors are enclosed in a plastic case and strapped²⁸⁸ 267 on to the array where they can easily slide and lock ²⁸⁹ 268 between 1 - 2 m from the reaction target to optimize²⁹⁰ 269 ToF information. CATRiNA has an angular cover-²⁹¹ 270 age of 40° which allows to obtain angular distribu-²⁹² 27 tions in a wide range. At the 1 m position, the solid ²⁹³ 272 angle coverage of each detector is about 7.8 msr.²⁹⁴ 273 For the present experiment, CATRiNA was located ²⁹⁵ 274 in target room 1 at the John D. Fox Accelerator 296 275 Laboratory [21] where the array was suspended by ²⁹⁷ 276 a crane next to the target chamber. The beam-stop 277 was extended from the reaction chamber to mini-278 mize neutron background. A silicon detector was 299 279 also placed in the reaction chamber and was used 300 280 as a monitor to normalize the beam current using 301 281 elastic scattering. The silicon detector was added 302 282

to the electronics as discussed in Sec. 2.2 via an additional CAEN ADC module. To calculate the dead-time, the before- and after-processing signals were sent to a CAEN scaler module. The experimental setup is shown in Fig. 9.

Neutron energies were extracted using the ToF information from the TDC modules by obtaining the time difference (Δ ToF) of the prompt-gamma and neutron peaks coming from the target. To achieve this, the accelerators RF signal was used as a 'stop' signal for the TDC, with the 'start' signal being events registered in the CATRiNA detectors. Energy of the neutrons was then calculated from the target to detector distance and the measured ToF.

4. Results and Discussion

A typical PSD spectrum obtained during the ⁷Li(p,n)⁷Be experiment is shown in Fig. 10 where a clear separation between neutrons and gamma rays is visible. The integration parameters used



Figure 10: Typical PSD Spectrum of one of the CAT-RiNA detectors obtained during the reaction measurement of 7 Li(p,n) 7 Be for proton energy of 7 MeV. Groups of n/ γ events are clearly separated for further analysis.

were optimized during the measurements with neu-303 tron and gamma sources described in Sec. 3.2. 304 From the PSD spectra at each proton energy, the 305 FOM for each detector was extracted as described 306 in Sec. 3.2 for cut-off thresholds of 180 and 250 307 keVee. The FOM for each neutron energy group of 308 one of the detectors used in the experiment (detec-309 tor at 10°) is shown in Fig. 11, where they average 310 1.3 for all the energies covered which indicates a 311 good n/γ separation. 312

For each detector, the ToF information was gated with the neutron and gamma gates from the PSD spectrum. This allowed to extract the energy of the neutrons via the time difference between the prompt gamma-ray from the interaction of the beam with the target and neutron signals. A typical ToF spectrum for one of the detectors at each



Figure 11: 1-Dimensional plots for different neutron energies of a detector placed at 10° obtained by projecting PSD plots to the x-axis with a 250 keVee threshold. The FWHM and centroid for both neutron peak (left) and gamma peak (right) are calculated by fitting a Gaussian curve. All detectors showed similar behavior.

energy used in the present experiment is shown in Fig. 12.

The neutron ToF spectra are obtained by apply-322 ing a cut to the neutron group in the PSD plot. The 323 prompt gamma signal is superimposed as reference 324 for ToF calculations. For the first couple of pro-325 ton energies ($E_p = 4$ and 5 MeV), neutrons from 326 the ground and first excited state in ⁷Be are well 327 resolved and the individual cross sections were ex-328 tracted. For higher proton energies ($E_p = 6, 7, 8$ 329 and 9 MeV), the cross sections for neutrons from 330 the ground and first excited state in ⁷Be could not 331 be separated and are extracted together. Back-332 ground neutrons appeared mainly at higher ener-333 gies coming from the Faraday cup were also ob-334 served in the spectra. 335

One of the most important aspects of the char-336 acterization of CATRiNA was to explore the pulse-337 height spectra as a function of neutron energy. Af-338 ter the n/γ separation is performed in the PSD spec-339 tra, the events in the neutron gate are projected on 340 to the Long integration axis (y-axis) along with a 341 ToF condition, which eliminates most gamma in-342 teractions as well as scattered low-energy neutron 343 events. The 1-dimensional projections of the neu-344 tron events in one of the detectors for different neu-345 tron energies are shown in Fig. 13. It can be seen 346 that there is a distinctive peak that shifts to the 347 right as neutron energy increases. A Gaussian fit to 348 the peaks show that neutron groups with 500 keV 349



Figure 12: Time-of-Flight spectra of neutrons from the ⁷Li(p,n)⁷Be reaction for a CATRiNA detector placed 10° from the beam-axis at forward angles. Neutrons from reactions which populated the ground state of ⁷Be are marked with a black line and the energies are labelled. For reactions with ground state neutrons of $E_n = 4.3-7.3$ ($E_p = 6-9$ MeV), neutrons from the first excited state cannot be resolved. Additionally, neutrons from the target chamber's Faraday cup are seen at higher energies ($E_n = 6.3-7.3$ MeV). Superimposed is the prompt-gamma signal in red, which was used as reference for ToF calculations.



Figure 13: Light-output response of the CATRINA detectors to quasi-mono-energetic neutrons from the ⁷Li(p,n) reaction. A distinctive shoulder which shifts to the right as neutron energy increases is observed. For each peak, the energy of the $_{363}$ neutron groups is highlighted.

difference can be well separated. A light-output curve as a function of the energy deposited (8/9 E_n) for the CATRiNA detectors was extracted and is shown in Fig. 14. The light-output curve is fitted with a standard exponential form given by [22]:

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$$L = aE_{dep} - b(1 - e^{-cE_{dep}}),$$
 (2) 371

where L is the light-output, E_{dep} is the energy de- ³⁷³ posited and a, b and c are adjustable parameters. The parameters used in the fit were a = 0.8666, b = 11.2839 and c = 0.0609 and covers all the data ³⁷⁵ points measured.

The pulse-height resolution of the CATRiNA de- 377
 tectors to neutron events was determined as follows 378



Figure 14: Measured light-output curve for the CATRiNA detectors fitted with a standard exponential form [22].

according to Ref. [15],

$$R = \frac{1.5(L_{1/2} - L_{max})}{L_{1/2}}.$$
(3)

Here, $L_{1/2}$ is the location along the edge of the peak where the value is 50% of the maximum and L_{max} is the location of the maximum. The measured resolution for the neutron peaks observed in Fig. 13 as a function of the pulse-height is shown in Fig. 15. The resolution function is expected to follow a $1/\sqrt{pulseheight}$ trend, as reported by other studies [22, 23], fitting the following functional form to the data:

$$R = \sqrt{\alpha^2 + \frac{\beta^2}{L} + \frac{\gamma^2}{L^2}},\tag{4}$$

where L is the light-output, α , β and γ are adjustable parameters that describe the transmission of the scintillation light through the detector as well as the electronics noise. In this work, $\alpha = 0.1311$, β

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Figure 15: Measured energy resolution of the CATRiNA detectors to neutron events. The resolution is fitted with a functional form given by eq. 4 and shows a comparable trend to works done with similar detector types and sizes [22, 23].

 $_{379} = 0.1181$ and $\gamma = 0.0128$ are the fitted parameters. $_{397}$ The results showed a similar behavior to other ex- $_{398}$ periments performed with EJ-315 detectors of sizes $_{399}$ $_{382}$ 7.36x7.36 cm² [23] and 7.62x5.08 cm² [22]. The 400 differences found in resolution are attributed to dif- 401 ferences in detector size. 402

A second MC simulation was performed us- 403 385 ing the neutron-transport code "Monte-Carlo N- 404 386 Particle Transport Code (MCNP6)" [24]. MCNP6 405 387 was used to calculate the efficiency of the detectors 406 388 and compare to those obtained experimentally. A 407 389 point source of mono-energetic neutrons was sim- 408 390 ulated and the threshold was set using a cut-off on 409 39 the recoil-energy of the deuteron. A conversion be- 410 392 tween the energy deposited by the neutrons to the 411 393 deuteron and light-output (in MeVee) is shown in 412 394



Figure 16: MCNP6 simulated efficiency curve of monoenergetic neutrons at different keVee thresholds.

Fig. 14. To obtain the energy deposit information in MCNP6, the PTRAC class was used which provides energy information of every neutron after each collision in the scintillator material. The efficiency of a neutron scintillator detector is threshold dependent, therefore different values were used to understand its behavior. Simulated efficiencies for different thresholds are shown in Fig. 16.

An experimentally measured efficiency curve was extracted by comparing the calculated cross section from the present experiment to data in the literature [20]. Two different thresholds, 180 keVee and 250 keVee, were set on the neutrons in the PSD spectrum. The experimental efficiency curves at 180 and 250 keVee threshold are shown in Fig. 17 and are compared to the MC simulated efficiency curves, finding them in overall good agreement within error bars. The energy spread is less

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Figure 17: Efficiency curves simulated with MCNP6 for one of the CATRiNA detectors at 180 keVee and 250 keVee threshold plotted along with the extracted experimental efficiencies. ⁴³⁹

than 8% for the energy range covered, which was
calculated by considering the timing resolution of
the bunched beam, the thickness of the detectors as
well as the uncertainty in the position of the array.

417 5. Summary and Outlook

We have developed and characterized the CAT- 447 418 RiNA neutron detector array at FSU. CATRINA 448 419 consist of 16 liquid scintillator detectors based on 449 420 EJ-315 deuterated benzene (C_6D_6) as scintillating 450 421 material. Measurements with gamma and neutron 451 422 sources were performed to optimize the parame- 452 423 ters of the MCFD and MQDC modules needed 453 424 for PSD analysis based on the charge-integration 454 425 method. The optimized integration times for PSD 455 426 were found to be 32 ns (Short gate) and 187 ns 456 427 (Long gate). FOM greater than 1 were consis- 457 428 tently obtained for a 250 KeVee threshold with a 458 429 ²⁵²Cf source and 180 KeVee threshold during in- 459 430

beam experiment for all detectors. Two Monte-431 Carlo simulations were performed to simulate the 432 response of the CATRINA detectors and an effi-433 ciency curve as a function of neutron energy. A 434 measurement of the The ${}^{7}Li(p,n){}^{7}Be$ reaction was 435 carried out to study the performance of the detec-436 tors to quasi-mono-energetic neutrons and to vali-437 date the MC simulations. 438

Pulse-height analysis showed distinct neutron energy groups for different neutron energies. The analysis and experimental results showed that different neutron energies can be identified without fully relying in ToF if neutron energy differences are larger than ~ 500 keV for the energy ranged covered. These results opens great possibilities in neutron spectroscopy studies where multiple correlations in PSD, ToF and pulse-height analysis could be combined to optimize and enhance the use of neutron detectors for nuclear reactions relevant in nuclear structure and nuclear astrophysics studies such as (³He,n), (d,n), (p,n). Further developments include the digitization of the data acquisition system and the exploration of different techniques for unfolding the pulse-height signals [7, 22] to improve the resolution of the detectors. The use of CATRINA in coincidence with other detector systems at FSU, like the Split-Pole Spectrograph as well as with arrays of charge-particles and gamma-rays detectors is also being explored.

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- 471 [1] P. E. Garrett. Descant the deuterated scintilla- 509
 472 tor array for neutron tagging. *Hyperfine Interactions*, 510
 473 225(1):137–141, Jan 2014. 511
- [2] W.A. Peters, S. Ilyushkin, M. Madurga, C. Matei, 512 474 S.V. Paulauskas, R.K. Grzywacz, D.W. Bardayan, C.R. 513 475 Brune, J. Allen, J.M. Allen, Z. Bergstrom, J. Blackmon, 514 476 477 N.T. Brewer, J.A. Cizewski, P. Copp, M.E. Howard, 515 R. Ikeyama, R.L. Kozub, B. Manning, T.N. Massey, 516 478 M. Matos, E. Merino, P.D. O'Malley, F. Raiola, C.S. 517 479 Reingold, F. Sarazin, I. Spassova, S. Taylor, and D. Wal- 518 480 ter. Performance of the versatile array of neutron de- 519 481 tectors at low energy (vandle). Nuclear Instruments 520 482 and Methods in Physics Research Section A: Acceler- 521 483 ators, Spectrometers, Detectors and Associated Equip- 522 484 ment, 836:122 - 133, 2016. 485 523
- 486 [3] L.T. Baby, S Kuvin, Ingo Wiedenhoever, M Anasta- 524
 487 siou, David Caussyn, K Colbert, N Quails, and D Gay. 525
 488 Resoneut: A detector system for spectroscopy with (d,n) 526
- 489 reactions in inverse kinematics. *Nuclear Instruments* 527
- 490 and Methods in Physics Research Section A: Acceler- 528
- 491ators, Spectrometers, Detectors and Associated Equip-529492ment, 877, 09 2017.530
- 493 [4] F.D. Brooks. Development of organic scintillators. Nu- 531
- 494 *clear Instruments and Methods*, 162(1):477 505, 1979. 532

- [5] V. Bildstein, P.E. Garrett, J. Wong, D. Bandyopadhyay, J. Bangay, L. Bianco, B. Hadinia, K.G. Leach, C. Sumithrarachchi, S.F. Ashley, B.P. Crider, M.T. McEllistrem, E.E. Peters, F.M. Prados-Estévez, S.W. Yates, and J.R. Vanhoy. Comparison of deuterated and normal liquid scintillators for fast-neutron detection. *Nuclear Instruments and Methods in Physics Research Section A: Accelerators, Spectrometers, Detectors and Associated Equipment*, 729:188 – 197, 2013.
- [6] Chris C. Lawrence, Andreas Enqvist, Marek Flaska, Sara A. Pozzi, and F. D. Becchetti. Comparison of spectrum-unfolding performance of (ej315) and (ej309) liquid scintillators on measured 252cf pulse-height spectra. Nuclear Instruments and Methods in Physics Research, Section A: Accelerators, Spectrometers, Detectors and Associated Equipment, 729:924–929, 10 2013.
- [7] M. Febbraro, F. D. Becchetti, R. O. Torres-Isea, J. Riggins, C. C. Lawrence, J. J. Kolata, and A. M. Howard. (d, n) proton-transfer reactions on ⁹Be,¹¹ B,¹³ C,^{14,15} N, and ¹⁹F and spectroscopic factors at $E_d = 16$ MeV. *Phys. Rev. C*, 96:024613, Aug 2017.
- [8] Eljen Technology. Deuterated ej315. https://eljentechnology.com/products/ liquid-scintillators/ej-315, Accessed: 2018-04-15.
- [9] ET Enterprises Ltd. Photomultipliers. http: //www.et-enterprises.com/photomultipliers, Accessed: 2018-04-15.
- [10] W-IE-NE-R. Mpod lv/hv. http://www.wiener-d. com/sc/power-supplies/mpod--lvhv/, Accessed: 2018-04-15.
- [11] Mesytec. Mcfd-16. https://www.mesytec.com/ products/nuclear-physics/MCFD-16.html, Accessed: 2018-04-15.
- [12] Mesytec. Mqdc-32. https://www.mesytec.com/ products/nuclear-physics/MQDC-32.html, Accessed: 2018-04-15.
- [13] CAEN. V775. https://www.caen.it/products/

495

| 533 | | v775/, Accessed: 2018-04-15. | 571 |
|-----|------|--|-----|
| 534 | [14] | Mark A. Norsworthy, Marc L. Ruch, Michael C. Hamel, | 572 |
| 535 | | Shaun D. Clarke, Paul A. Hausladen, and Sara A. Pozzi. | 573 |
| 536 | | Light output response of ej-309 liquid organic scintil- | 574 |
| 537 | | lator to 2.86-3.95 mev carbon recoil ions due to neu- | 575 |
| 538 | | tron elastic and inelastic scatter. Nuclear Instruments | 576 |
| 539 | | and Methods in Physics Research Section A: Acceler- | |
| 540 | | ators, Spectrometers, Detectors and Associated Equip- | |
| 541 | | ment, 884:82 - 91, 2018. | |
| 542 | [15] | G. Dietze and H. Klein. Gamma-calibration of ne 213 | |
| 543 | | scintillation counters. Nuclear Instruments and Methods | |
| 544 | | in Physics Research, 193(3):549 – 556, 1982. | |
| 545 | [16] | S. Agostinelli et al. Geant4-a simulation toolkit. Nu- | |
| 546 | | clear Instruments and Methods, 506:250 - 303, 2003. | |
| 547 | [17] | F.D. Brooks. A scintillation counter with neutron and | |
| 548 | | gamma-ray discriminators. Nuclear Instruments and | |
| 549 | | Methods, 4(3):151 – 163, 1959. | |
| 550 | [18] | R.A. Winyard, J.E. Lutkin, and G.W. McBeth. Pulse | |
| 551 | | shape discrimination in inorganic and organic scintilla- | |
| 552 | | tors. i. Nuclear Instruments and Methods, 95(1):141 - | |
| 553 | | 153, 1971. | |
| 554 | [19] | R.A. Winyard and G.W. McBeth. Pulse shape discrimi- | |
| 555 | | nation in inorganic and organic scintillators, ii. Nuclear | |
| 556 | | Instruments and Methods, 98(3):525 – 533, 1972. | |
| 557 | [20] | R. R. Borchers and C. H. Poppe. Neutrons from proton | |
| 558 | | bombardment of lithium. Phys. Rev., 129:2679-2683, | |
| 559 | | Mar 1963. | |
| 560 | [21] | Florida State University. John d. fox accelerator facil- | |
| 561 | | <pre>ity. http://fsunuc.physics.fsu.edu/, Accessed:</pre> | |
| 562 | | 2018-04-15. | |
| 563 | [22] | Chris C. Lawrence, Andreas Enqvist, Marek Flaska, | |
| 564 | | Sara A. Pozzi, A.M. Howard, J.J. Kolata, and F.D. Bec- | |
| 565 | | chetti. Response characterization for an ej315 deuterated | |
| 566 | | organic-liquid scintillation detector for neutron spec- | |
| 567 | | troscopy. Nuclear Instruments and Methods in Physics | |
| 568 | | Research Section A: Accelerators, Spectrometers, De- | |
| 569 | | tectors and Associated Equipment, 727:21 – 28, 2013. | |
| 570 | [23] | Haitang Wang, Donald Carter, Thomas N. Massey, and | |
| | | | 15 |
| | | | |

Andreas Enqvist. Neutron light output function and resolution investigation of the deuterated organic liquid scintillator ej-315. Radiation Measurements, 89:99 -106, 2016.

[24] T. Goorley et al. Initial mcnp6 release overview. Nuclear Technology, 180:298 - 315, 2012.