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AN INVESTIGATION OF THE ASTROPHYSICALLY IMPORTANT $^{14}\text{N}(p, \gamma)^{15}\text{O}$ REACTION

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AN INVESTIGATION OF THE ASTROPHYSICALLY IMPORTANT $^{14}\text{N}(p, \gamma)^{15}\text{O}$ REACTION

A Dissertation

Submitted to the Graduate School
of the University of Notre Dame
in Partial Fulfillment of the Requirements
for the Degree of

Doctor of Philosophy

by
Bryce Alan Frentz

Ani Aprahamian, Director

Graduate Program in Physics
Notre Dame, Indiana
February 2020

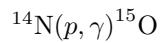
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Abstract

by

Bryce Alan Frentz



What do you even put in an abstract for a thesis? Seems a little ridiculous... But I suppose that this is where I'll write a page or so about the final results and try to summarize everything I've done in the last six years. And now I think this is long enough to show that it is double spaced for a formatting check.

Bananas is fruits. Tacos is sandwiches.

NEW DEDICATION NAME

To Laura, without whose support I probably couldn't have accomplished anything.

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PREFACE

When nothing is done, nothing is left undone. - Lao Tzu

ACKNOWLEDGMENTS

I would like to acknowledge ... Thanks everyone!

CHAPTER 1

INTRODUCTION

1.1 Overview

1.1.1 General overview

Nuclear physics data, in both the forms of theoretical predictions and experimental results, is an ever richer trove of information useful in solving many astrophysical problems. The partnership between the fields of nuclear physics and astrophysics was actually born in the early days of nuclear physics when Sir Arthur Eddington postulated that recent laboratory discoveries could explain the energy generation of the sun and stars [34], and the relationship between the two has only deepened in the years since. Today, the field is known as nuclear astrophysics and is understood to be the interdisciplinary branch of physics concerned with the nucleosynthetic processes occurring in stars and their detection on Earth. As many astrophysical phenomena are governed by underlying nuclear physics, developments in one area naturally link to increased knowledge in the other - a scientific point and counterpoint. So, as a nuclear physicist, it is through the study of the microscopic aspects of the universe that we also elucidate its behavior in the grandest scales imaginable.

The general questions at heart of nuclear astrophysics, as defined by the community in a recent White Paper [51], mirror a subset of the fundamental questions from the 2012 Decadal Survey of Nuclear Physics [49] as follows:

- How did visible matter come into being and how does it evolve?
- How does subatomic matter organize itself and what phenomena emerge?
- How can the knowledge and technological progress provided by nuclear science best be used to benefit society?

Re-formed through the lens of nuclear astrophysics, these can be summarized into an overarching goal: to understand and observe both the processes by which the chemical elements are formed (in either quiescent or explosive scenarios) and also those mechanism(s) responsible for their relative abundance distribution observed today. Crucially, stellar evolution is cyclic: stars are born, evolve, and ultimately die, seeding the interstellar medium with the collective material they've created

over their life in order to form the building blocks for the next stellar generation. As each of the different stellar phases are guided in part by the underlying nuclear physics, investigations into these questions provide a rich ink between humans and our place in the cosmos.

The synthesis of the chemical elements we see began with the universe as the Big Bang created hydrogen, helium, and trace amounts of lithium [5]. Shortly after the Big Bang, the temperature and density of the Universe dropped too low for any significant nuclear fusion to occur and overcome the mass 5 and mass 8 barriers. This was the end of Big Bang Nucleosynthesis (BBN), as after this point, the elemental abundances were nearly stable, with the only changes coming from the radioactive decay of ${}^3\text{He} \rightarrow t$ and ${}^7\text{Be} \rightarrow {}^7\text{Li}$ [27].

From this point, generations of stellar evolution created nearly all other elements up to iron, with those beyond iron being formed in exotic phenomena, leading to the currently observed elemental abundance distribution, shown in Figure 1.1. The synthesis of elements heavier than BBN products, commonly referred to as “metals” by astronomers, began with the formation of the first stars. Cold gas clouds, composed primarily the BBN products of hydrogen and helium (and metals in subsequent stellar generations), coalesce under their mutual gravitational attraction, converting their gravitational potential energy to heat – increasing their pressure, density, and temperature. Upon passing a critical threshold for these physical quantities, typically temperatures >1 MK, thermonuclear fusion reactions begin in the core [54]. This transition marks the “birth” of a star, having reached a point where nuclear reactions provide a sufficient heat source to balance energy losses from radiation.

1.1.2 Stellar evolution

Every star is unique and their intrinsic quantities when forming, such as mass or initial abundance distribution, prescribe the nuclear reactions that can occur throughout their lifetimes. Despite their differences, correlations emerge in studies of their observational properties , like luminosity, mass distribution, temperature, color, etc. The most evident patterns appeared when classifying stars by luminosity (or magnitude) vs temperature (or color), named the *Hertzsprung-Russel* (HR) diagram [23, 34, 54], distilling the most important relationship between stellar properties. An example HR diagram for stars in the solar neighborhood is displayed in Figure 1.2. Since HR diagrams represent a snapshot of the evolutionary track of a set of stars, the most densely populated areas of such plots are those in which individual stars necessarily spend most of their lives. Conversely, as there is a lower probability of observing a star in a short-lived phase, sparsely populated areas of HR diagrams

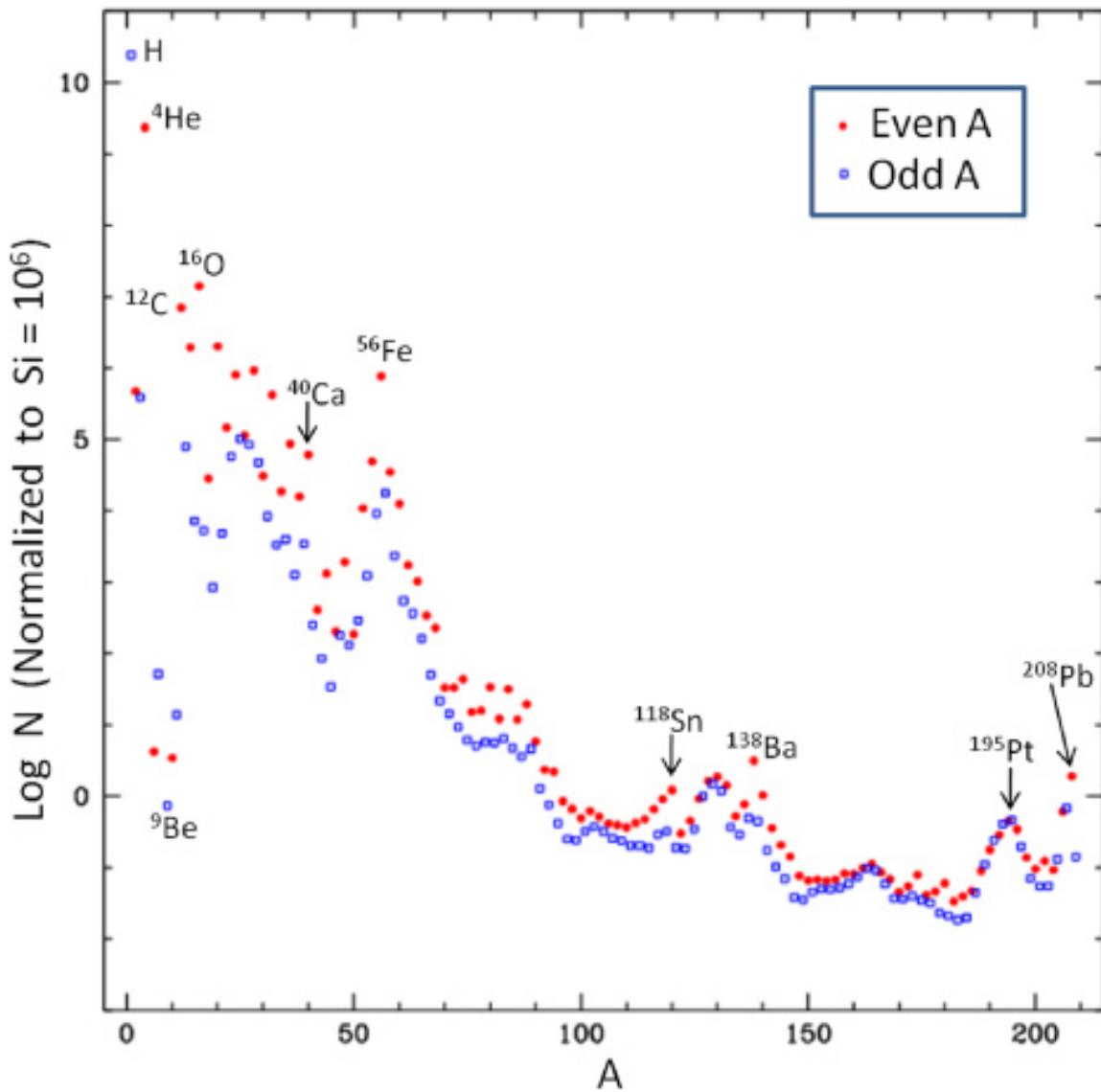


Figure 1.1. The current solar isotopic abundance pattern vs. mass number for even and odd nuclei [37]. Abundances are based on both meteoritic samples and observations of the solar photosphere.

detail such phases of stellar burning.

The vast majority of stars lie on the diagonal band stretching from the upper left (hot, bright stars) to the lower right (cool, faint stars) of the diagram. This feature is called the main sequence. These stars support themselves through core hydrogen burning – implying that stars spend most of their lives in this long period of quiescent hydrogen burning in the *pp chains* and the CNO cycles, detailed in subsequent sections. Other prominent features of typical HR diagrams are the clusters of stars in the upper quadrants (very bright stars), commonly referred to as the giant branch, the lower left quadrant (hot, faint stars), where the white dwarf stars reside, and the lower right quadrant beyond the main sequence, occupied by brown dwarfs. Each classification corresponds to different nuclear reactions, with stars evolving off the main sequence being characterized by phases of burning of heavier elements, like helium or carbon.

Any single star's evolutionary track will follow different paths depending on its initial mass [34, 54]. For cosmic objects of mass $\lesssim 0.08 M_{\odot}$, where M_{\odot} is the mass of the sun, core temperatures never rise high enough to sustain hydrogen fusion, and thus they evolve directly into brown dwarf stars. In the range of mass $0.08 M_{\odot} \lesssim M \lesssim 0.4 M_{\odot}$, stars undergo core hydrogen fusion through the *pp chains*, but do not reach temperatures sufficient to initiate helium burning after exhausting hydrogen in the core. Ultimately, stars in this classification become white dwarfs. The next mass grouping consists of Sun-like stars, with masses $0.4 M_{\odot} \lesssim M \lesssim 8.0 M_{\odot}$, which evolve off the main sequence and continue growing and burning (both in the stellar core and shells in their stellar envelope). Such stars achieve temperatures high enough to create final abundance distribution composed of primarily carbon and oxygen in their core. After losing their stellar envelopes, these stars are carbon-oxygen white dwarfs. However, if such a star has a companion of sufficient mass, it will eventually explode in a type Ia supernova. Lastly, are stars of mass $8.0 M_{\odot} \lesssim M$, which go through sequences of carbon, neon, oxygen, and silicon burning to produce a dominantly iron-nickel core. At this point, as the stars are more massive than the Chandrasekhar limit, the gravitational attraction will exceed the support of electron degeneracy pressure causing a core collapse (type II) supernova. Depending on the star's mass at this stage, it will become either a neutron star or black hole. While not the topic of this thesis, such explosive scenarios are currently an area of intense research as they are the location of many unresolved issues of nuclear astrophysics, including what the National Research Council identified in 2003 as one of the eleven greatest unanswered questions of the century, namely the origin of the elements heavier than iron [62].

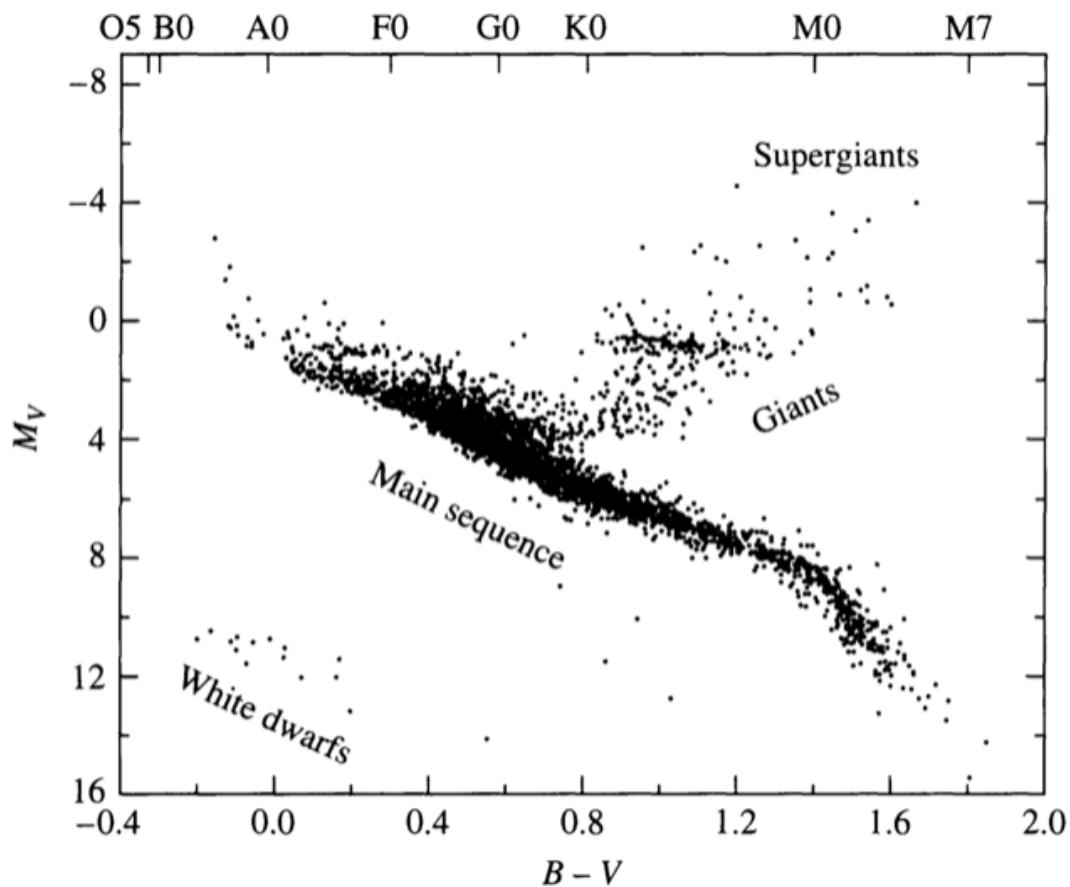


Figure 1.2. An example of a Herzprung-Russell diagram. The horizontal axis gives the temperature of the star while the vertical shows the stellar luminosities [23].

1.1.3 Hydrogen Burning

The lifetimes of stars along the main sequence vary drastically, from millions of years to trillions of years, depending on the star's properties when forming. Likewise, the dominant type of hydrogen burning that occurs during the main sequence phase is also highly dependent on stellar characteristics like mass and initial composition. Broadly, there are two sets of nuclear processes that comprise the hydrogen burning phases in stellar cores, the *pp chains* and the CNO cycles, with each ultimately functions in a similar way, to convert four hydrogen nuclei to one helium nucleus, releasing ~ 26 MeV of energy [52].

The *pp chains* are the main energy generator in smaller stars with core temperatures below 20 MK [54], like the sun. There are three variations of the *pp chains* (denoted ppI, ppII, and ppIII), shown below.

<u>ppI</u>	<u>ppII</u>	<u>ppIII</u>
$p(p, e^+ \nu) d$	$p(p, e^+ \nu) d$	$p(p, e^+ \nu) d$
$d(p, \gamma)^3\text{He}$	$d(p, \gamma)^3\text{He}$	$d(p, \gamma)^3\text{He}$
$^3\text{He} (^3\text{He}, 2p) ^4\text{He}$	$^3\text{He} (^4\text{He}, \gamma) ^7\text{Be}$	$^3\text{He} (^4\text{He}, \gamma) ^7\text{Be}$
	$^7\text{Be} (e^- \nu) ^7\text{Li}$	$^7\text{Be} (p, \gamma) ^8\text{B}$
	$^7\text{Li} (p, \alpha) ^4\text{He}$	$^8\text{B} (e^+ \nu) ^8\text{Be}$
		$^8\text{Be} (\alpha) ^4\text{He}$

Between the three branches there are clear similarities and differences. The energy released between each of ppI, ppII, and ppIII are 26.20 MeV, 25.66 MeV, and 19.75 MeV, respectively, with the difference coming from the amount of energy carried away by the electrons, positrons, and neutrinos. Of these chains, ppI is the most prominent, occurring $\sim 85\%$ of the time, where the remainder branches off as a ^3He nucleus will fuse with a ^4He nucleus, resulting in the ppII or ppIII chain, occurring $\sim 14.99\%$ and $\sim 0.1\%$ of the time, respectively [54]. Regardless of the branching for the *pp chains*, each starts with the $p(p, e^+ \nu) d$ reaction. This reaction is governed purely by the weak interaction, and thus is about 20 orders of magnitude slower than those proceeding via the strong interaction [52], like the reactions in the chains that follow. Therefore, energy production, nucleosynthesis, and stellar evolution timescales for stars dominated by the *pp chains* are restricted

by the $p(p, e^+ \nu) d$ reaction.

However, for most stars the CNO cycles are the dominant energy source. Stellar energy generation depends sensitively on the temperature in the star's core, which is closely tied to its mass. For stars of mass $M \gtrsim M_\odot$, where cores contain carbon and oxygen seed nuclei and temperatures exceed 20 MK, the CNO cycles' energy production overtakes that of the *pp chains*. Thus, the knowledge of the overall rate of this cycle is important for the study of their evolution.

The CNO cycles are the collection of four similar energy producing cycles broadly characterized by their synthesis of helium from hydrogen with a carbon catalyst, seeded by earlier generations of stars. There are four CNO cycles, listed below with their relationships shown in Fig. 1.5.

<u>CNO1</u>	<u>CNO2</u>	<u>CNO3</u>	<u>CNO4</u>
$^{12}\text{C}(p, \gamma)^{13}\text{N}$	$^{14}\text{N}(p, \gamma)^{15}\text{O}$	$^{15}\text{N}(p, \gamma)^{16}\text{O}$	$^{16}\text{O}(p, \gamma)^{17}\text{F}$
$^{13}\text{N}(\beta^+\nu)^{13}\text{C}$	$^{15}\text{O}(\beta^+\nu)^{15}\text{N}$	$^{16}\text{O}(p, \gamma)^{17}\text{F}$	$^{17}\text{F}(\beta^+\nu)^{17}\text{O}$
$^{13}\text{C}(p, \gamma)^{14}\text{N}$	$^{15}\text{N}(p, \gamma)^{16}\text{O}$	$^{17}\text{F}(\beta^+\nu)^{17}\text{O}$	$^{17}\text{O}(p, \gamma)^{18}\text{F}$
$^{14}\text{N}(p, \gamma)^{15}\text{O}$	$^{16}\text{O}(p, \gamma)^{17}\text{F}$	$^{17}\text{O}(p, \gamma)^{18}\text{F}$	$^{18}\text{F}(\beta^+\nu)^{18}\text{O}$
$^{15}\text{O}(\beta^+\nu)^{15}\text{N}$	$^{17}\text{F}(\beta^+\nu)^{17}\text{O}$	$^{18}\text{F}(\beta^+\nu)^{18}\text{O}$	$^{18}\text{O}(p, \gamma)^{19}\text{F}$
$^{15}\text{N}(p, \alpha)^{12}\text{C}$	$^{17}\text{O}(p, \alpha)^{14}\text{N}$	$^{18}\text{O}(p, \alpha)^{15}\text{N}$	$^{19}\text{F}(p, \alpha)^{16}\text{O}$

As with the *pp chains*, the net result of each of the CNO cycles is the conversion of four protons into helium, releasing ~ 26 MeV of energy in the process [54]. Additionally, all of the heavier elements (carbon, nitrogen, oxygen, and fluorine) are only catalysts, with their overall abundance unchanged through the cycles while only the hydrogen is consumed. Therefore, this implies that the CNO cycles can occur even if the amount of heavy elements is relatively small, in the case of only being seeded by a single prior generation of stars [34]. Of the cycles, however, the main CNO1 cycle contributes $\sim 99\%$ of the CNO energy production [2], so hereafter CNO1 will be referred to as simply the CNO cycle, while the others will be designated as needed.

Within the CNO cycle, the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction is the slowest and thus the rate-limiting step of the whole process, ruling the energy production and the overall time spent in this burning phase, evidenced recently by the adjustment of the estimation of globular cluster ages [35].

The relative energy production between the *pp chains* and CNO cycles are shown in Fig. 1.4. It can be seen that the CNO cycle accounts for $< 1\%$ of solar energy production [1, 2]. However, it

produces 1.6 % of solar neutrinos through the β -decay of ^{13}N and ^{15}O [14], which can be used to test assumptions of the Standard Solar Model (SSM), as these rates are one of the largest sources of uncertainty in model predictions [58]. Efforts to measure the neutrino fluxes from the β -decays of ^{13}N and ^{15}O are either ongoing or planned at the Borexino, Super-Kamiokande, and SNO+ laboratories [37].

These laboratories are well known for their contributions in solving the *solar neutrino problem* [37] (and references therein). However, after this solar problem was solved, a new problem arose, named the solar abundance problem [1, 59]. The problem is as follows, solar models describe the structure, composition, and evolution of a $1M_{\odot}$ star from ignition through the sun's current age and are constrained by the plethora of helioseismology data from the Sun. Recent analyses of the Sun's photosphere lead to a reduced metallicity estimate, destroying the agreement between helioseismology and solar models [10]. In discussing potential solutions to this problem, it was proposed to use the neutrino fluxes from the β -decay of CNO isotopes ^{13}N and ^{15}N to address this problem, as these quantities depend linearly on their abundance within the solar core [14, 15]. Therefore, with more precise measurements of these neutrino fluxes (like the example given in Fig. 1.3) imminent, it is ever more important to have precise knowledge of the relevant nuclear cross-sections, as the fluxes from CNO neutrinos are one of the largest uncertainties present.

Due to the importance of the $^{14}\text{N}(p,\gamma)^{15}\text{O}$ reaction, it is the focus of this work. Presented first, in Sec. 1.2 and Sec. 1.3 are an overview of nuclear reactions and reaction theory relevant for nuclear astrophysics. An extension of these theories using the R -matrix formalism will be presented in Sec. 1.4. The current state of knowledge surrounding the reaction, its uncertainties, and previous measurements are discussed in Sec. 1.5. Finally, an outline of the remainder of this work will be presented in Sec. 1.6.

1.2 Thermonuclear reaction rates

Stars are fueled by the energy released during nuclear reactions. Recall that a nuclear reaction can be written as

$$a + A \rightarrow b + B + Q \quad \text{or} \quad A(a,b)B^* \quad (1.1)$$

where a denotes the projectile, A the target nucleus, b the ejectile, B the reaction product, and Q the energy released (or absorbed) during the reaction, typically present as kinetic energy distributed among the reaction products. For this reaction to occur, the initial nuclei must have enough energy

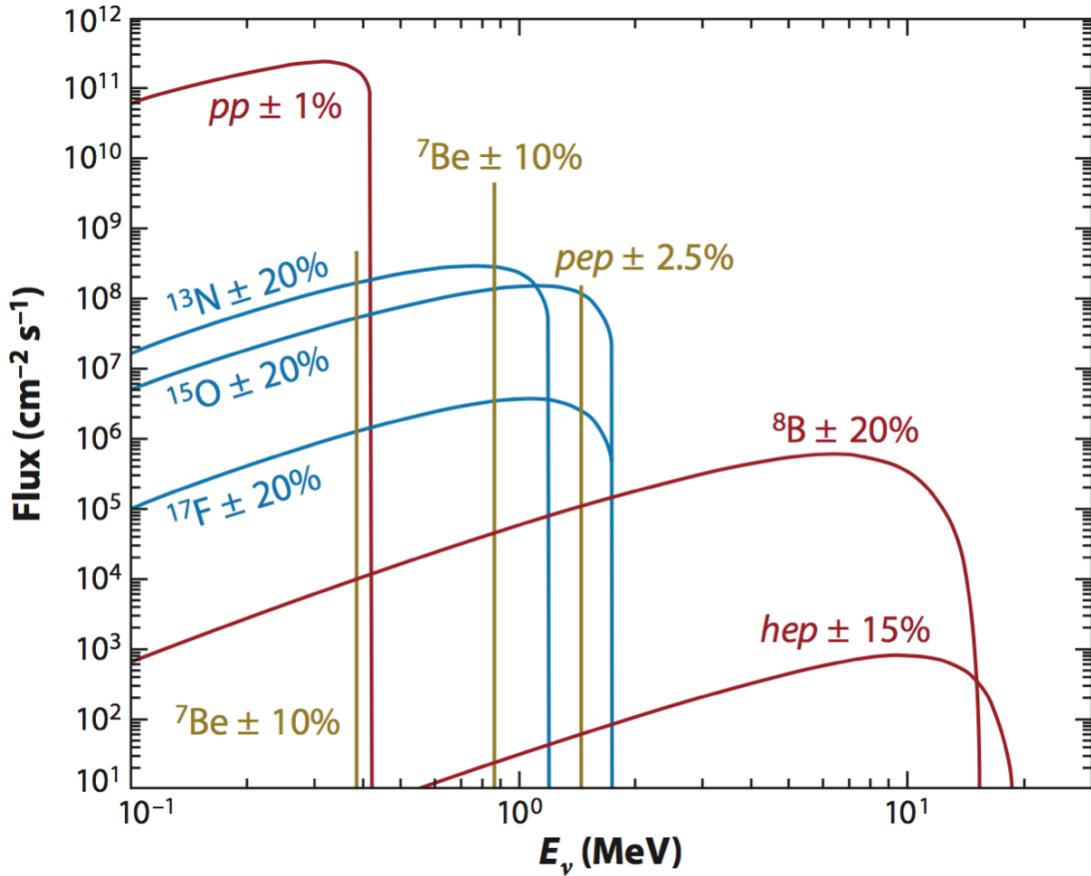


Figure 1.3. Predicted solar neutrino flux spectrum, taken from [64] and produced from data in [13]. As measurements of this spectrum are on the way, it is critical to understand the nuclear physics inputs, particularly those coming from the β -decay of ^{15}O , in order to resolve discrepancies between solar models and helioseismology.

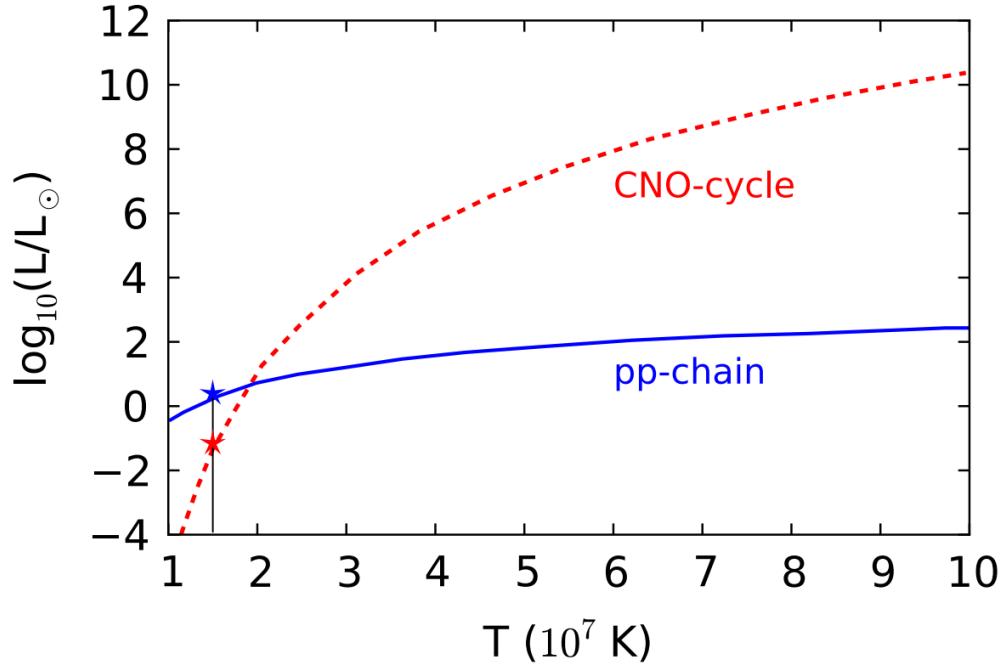


Figure 1.4. Energy dependence of the *pp chains* and the CNO cycles as a function of stellar temperature assuming a solar abundance distribution of CNO elements [20]. The sun’s temperature is marked with the star in the figure. This shows that for lower mass stars like the sun the *pp chains* are the dominant mode of energy production in main sequence burning. However, for hotter, more massive stars, the CNO cycle quickly dominates.

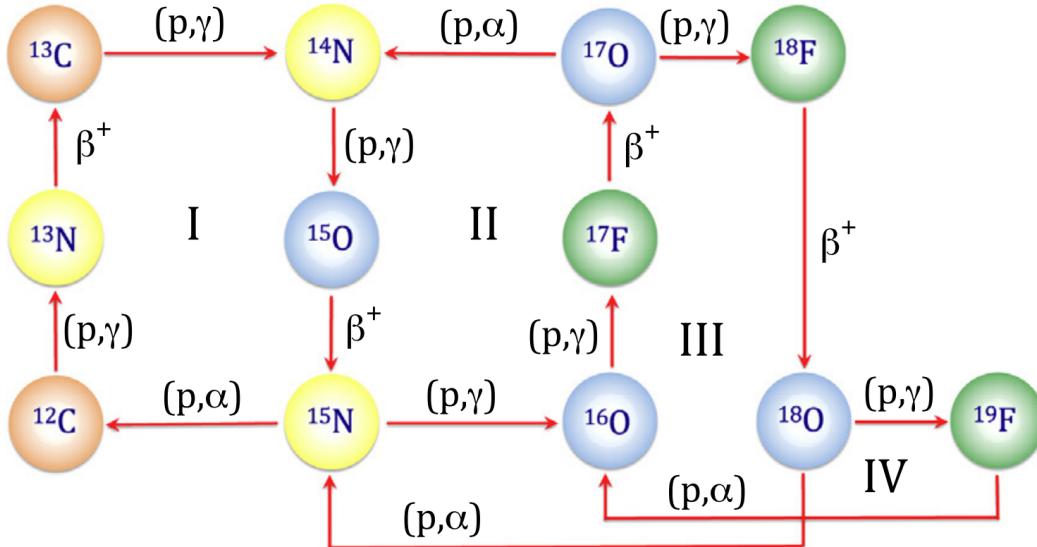


Figure 1.5. A depiction of the CNO cycles for hydrogen burning in stars [20]. Cycles II, III, and IV only contribute meaningfully at very high temperatures or explosive burning scenarios.

to overcome the Coulomb repulsion created by their constituent protons. The probability of their interaction, called their cross section (σ), is therefore energy dependent. As stars are powered by thousands of such reactions at a variety of energies, the understanding of nuclear cross-sections are crucial components to the subsequent understanding of stellar evolution and nucleosynthesis.

Consider a stellar environment filled with two types of nuclei with number densities (nuclei / cm³) N_1 and N_2 , respectively, and relative velocity v . The rate of reaction for these two species per unit volume would then be proportional to the probability that a given pair of particles would react multiplied by the number of pairs that interact per unit time. Mathematically, the reaction rate R_{12} would therefore be

$$R_{12} = \frac{N_1 N_2}{1 + \delta_{12}} v \sigma(v) \quad (1.2)$$

with δ_{12} to prevent overcounting if the particles are identical. In stellar environments, the relative velocity between the two particles can take a range of values related to the temperature, T . The probability distribution of the potential relative velocities, $P(v)$, is described by the Maxwell-

Boltzmann distribution [54]:

$$P(v) = 4\pi v^2 \left(\frac{\mu}{2\pi kT} \right)^{3/2} \exp \left(-\frac{\mu v^2}{2kT} \right) \quad (1.3)$$

where k is the Boltzmann constant, relating the kinetic energy of a particle to the temperature of its environment, and $\mu = m_1 m_2 / (m_1 + m_2)$ is the reduced mass of the two nuclei. Explicitly, the probability that a given pair of nuclei having a relative velocity between v and $v + dv$ is $P(v)dv$ and satisfies the unity expression

$$\int_0^\infty P(v)dv = 1. \quad (1.4)$$

Therefore, the reaction rate per particle pair $\langle \sigma v \rangle$ with relative velocity v is the average value of v multiplied with $\sigma(v)$ for a given temperature,

$$\langle \sigma v \rangle = \int_0^\infty v P(v) \sigma(v) dv. \quad (1.5)$$

Using a few useful substitutions, this formulation allows us to rewrite Equation 1.2, the general reaction rate, as a function of energy instead of velocity. To do this, recall that the kinetic energy is $E = \frac{1}{2}\mu v^2$ and its derivative is $dE/dv = \mu v$. By explicitly writing the Maxwell-Boltzmann distribution into and replacing the appropriate terms in Equation 1.5 leads to

$$\langle \sigma v \rangle = \left(\frac{8}{\pi \mu} \right)^{1/2} \left(\frac{1}{kT} \right)^{3/2} \int_0^\infty \sigma(E) E \exp \left(-\frac{E}{kT} \right) dE. \quad (1.6)$$

It is important to note that the only unknown in the entirety of the reaction rate is the expression of the nuclear cross section as a function of energy, $\sigma(E)$. Therefore, to understand the reaction rates for stellar environments, $\sigma(E)$ needs to be determined.

Recall that earlier, the cross section, $\sigma(E)$, is the probability of two nuclei reacting if brought together at a specific energy, E . While this quantity is an expression of the probability of reaction, it is analogous to the classical, cross-sectional area of the nucleus, so it is expressed in units of area (specifically barns, where $1 \text{ b} = 10^{-24} \text{ cm}^2$). An insightful description is to think of the cross section coming from throwing two balls at each other with random perturbations off of a common axis, classically. If either ball's size changes (cross-sectional area), the chances of hitting will also change in a manner that is directly proportional to the alteration of the respective sizes of the balls since the total cross-sectional area is $\sigma = \pi(R_1 + R_2)^2$. For our quantum mechanical nuclei, the nuclear

size needs to be replaced with the deBroglie wavelength, resulting in $\sigma = \pi\lambda$. Translating this into another formulation that is more familiar, the cross section is the number of reactions occurring divided by the number of potential reactions, given as

$$\sigma = \frac{N_R}{(N_i N_t)/A}, \quad (1.7)$$

where N_R is the number of reactions that take place per unit time, N_i is the number of incident nuclei per unit time, N_t is the number of target nuclei within the area of incidence, A .

In astrophysical environments, nuclei typically do not have enough energy to overcome the Coulomb repulsion provided by the protons within the nucleus and combine with the attractive nuclear strong force. A diagram of the combined nuclear and Coulomb potentials is given in Fig. 1.6. The repulsive Coulomb interaction causes the cross section to drop quickly at low center-of-mass energies [34]. Thus, for such reactions to occur, nuclei must actually tunnel through the Coulomb barrier.

The Coulomb penetrability, P , can be approximated as the leading term of the s -wave ($\ell = 0$) barrier transmission coefficient:

$$P = \exp(-2\pi\eta), \quad (1.8)$$

where η is the Sommerfeld parameter,

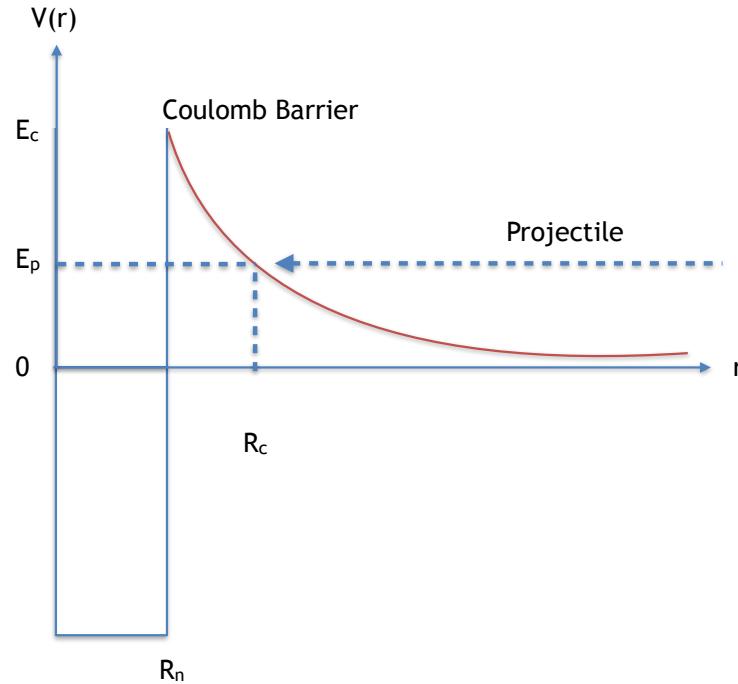
$$\eta = \frac{Z_1 Z_2 e^2}{4\pi\epsilon_0 \hbar v} = \alpha Z_1 Z_2 \sqrt{\frac{\mu c^2}{2E_{cm}}}, \quad (1.9)$$

where Z_i is the respective proton number of each nucleus, v is the incident center-of-mass velocity, $\alpha \approx 1/137$ is the fine-structure constant, μ is the reduced mass of the system, and E_{cm} is the center of mass energy of the interacting nuclei. Numerically, it is convenient to evaluate the quantity $2\pi\eta$, known as the Gamow factor, as

$$2\pi\eta = 0.989534 Z_1 Z_2 \left(\frac{\mu}{E_{cm}} \right)^{1/2}, \quad (1.10)$$

where E_{cm} is expressed in units of MeV and μ in atomic mass units (amu or simply, u).

By factoring the Coulomb penetrability out of the cross section, we can separate the contributions



1

Figure 1.6. Schematic plot of the combined attractive nuclear and repulsive Coulomb potentials where R_n is the nuclear radius, R_c is the classical turning point due to Coulomb repulsion, E_c is the height of the Coulomb barrier, and E_p is the projectile's energy. Because this is a quantum system, the projectile has a chance to tunnel through the Coulomb barrier and make it into the attractive nuclear potential when it has energy $E_p < E_c$.

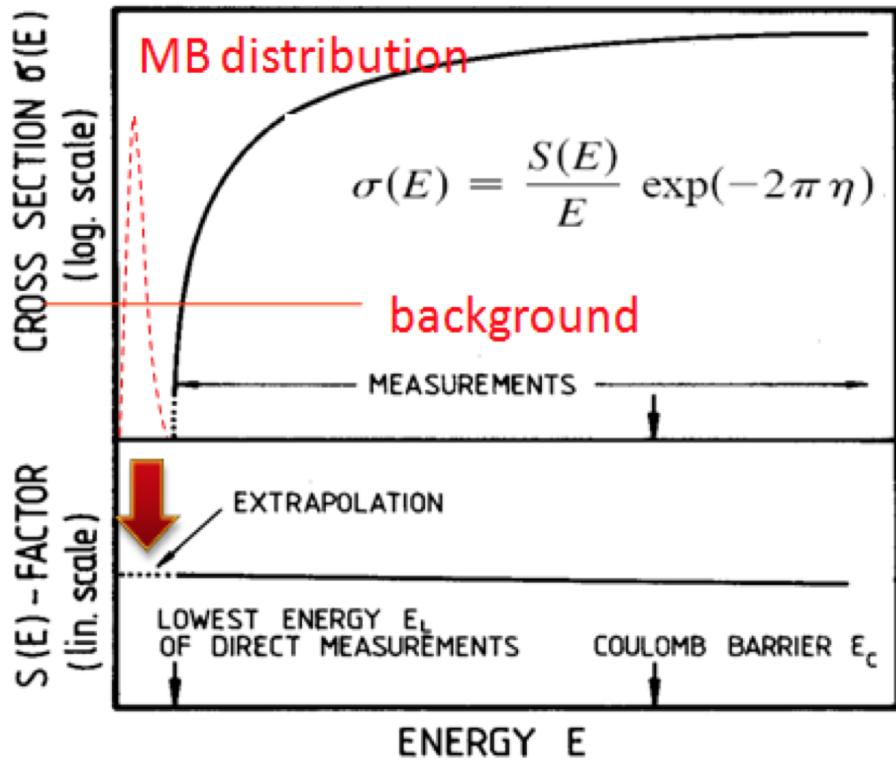


Figure 1.7. The energy dependence of the nuclear cross section and astrophysical S-factor [52]. The cross section drops dramatically due to Coulomb repulsion as the incident energy drops, however the S -factor, by removing the Coulomb penetrability varies slowly towards the Gamow window (barring the absence of nuclear resonances).

from the Coulomb barrier to isolate the nuclear physics in the astrophysical S -factor, $S(E)$,

$$\sigma(E) = \frac{1}{E} S(E) e^{-2\pi\eta}. \quad (1.11)$$

The astrophysical S -factor is slowly varying with energy, unlike the nuclear cross section. Fig. 1.7 provides a comparison of the energy dependence of both the cross section and S -factor, where you can see that the cross section drops by orders of magnitude in an energy regime where the S -factor is relatively constant. This implies that the Coulomb effects can obscure important nuclear physics effects. Additionally, as the S -factor is slowly varying, extrapolations of its behavior towards zero energy from higher energy data are significantly more reliable than those made from cross section data.

Now, after defining the S -factor, the reaction rate can be written finally as

$$\langle \sigma v \rangle = \left(\frac{8}{\pi \mu} \right)^{1/2} \left(\frac{1}{kT} \right)^{3/2} \int_0^\infty S(E) \exp \left(-\frac{E}{kT} - \frac{b}{E^{1/2}} \right) dE. \quad (1.12)$$

where b has units of $(\text{MeV})^{1/2}$ and is given as

$$b = 0.989534 Z_1 Z_2 \mu^{1/2}. \quad (1.13)$$

Recall that one of the most useful features of $S(E)$ is that it is typically smoothly varying with respect to energy, meaning that the reaction rate in total is dependent primarily on the exponential term inside the integrand of Eq. 1.12. The two components of this term arise from the Coulomb penetrability and the Maxwell-Boltzmann velocity distribution described in Eq. 1.3. The convolution of these two quantities can be used to estimate the energy range where the reaction rate should be maximized for given stellar temperatures [34].

The Coulomb penetrability and Maxwell-Boltzmann velocity distribution have opposite behaviors in energy regimes with the penetrability dropping to zero at low energies while the probability distribution for the velocities is maximized at low energies and vice-versa at high energies. The product of the terms is therefore negligible for nearly all energies. However, for a small set of energies this product, and therefore the reaction rate, maximizes. This behavior, as well as their product, commonly called the Gamow Peak, is presented in Fig. 1.8.

The Gamow peak's maximum can be obtained with basic calculus on the exponential term in Eq. 1.12 by setting the derivative with respect to energy equal to zero and solving for the energy value. This results in the maximum energy of the reaction rate being

$$E_0 = \left(\frac{bkT}{2} \right)^{2/3}. \quad (1.14)$$

For a given stellar temperature, T , the average effective energy at which the reaction occurs is E_0 . Additionally, this also allows for an approximation of the range of energies over which the reaction primarily occurs, provided that the S -factor is smoothly varying (which is not true if resonances are present, which will be discussed in the next section). This range of energies is known as the Gamow Window, Δ :

$$\Delta = \frac{4}{\sqrt{3}} \sqrt{E_0 k T}. \quad (1.15)$$

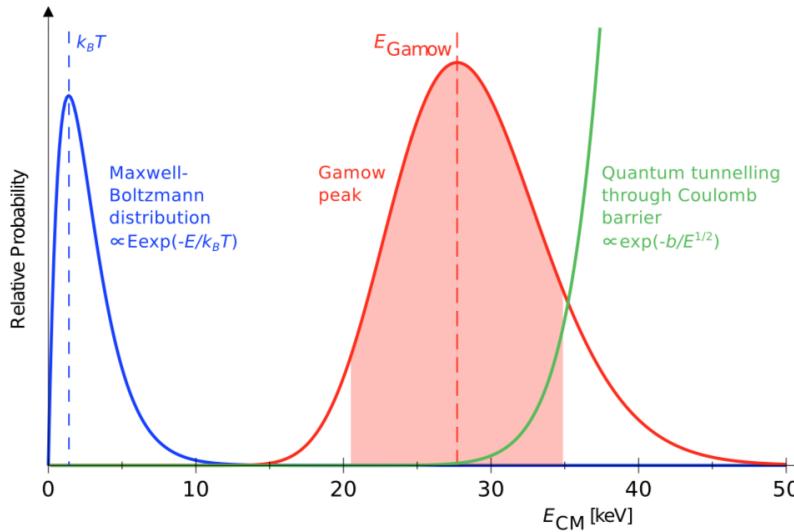


Figure 1.8. The two dominant terms of the reaction rate calculation and their convolution (Gamow peak) for the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction at $T = 0.016$ GK [16]. The y-axis is given in arbitrary units.

The Gamow window identifies the range of energies over which experimental data should be taken to provide a complete understanding of stellar processes and the interplay of the Maxwell-Boltzmann velocity distribution with the Coulomb penetrability to create the Gamow window is presented in Fig. 1.8. As an example, the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction in the sun has a Gamow peak energy at $E_0 = 26.6$ keV and $\Delta = 13.6$ keV, meaning that measuring the cross section in the energy range of 19.8 - 33.4 keV is crucial in the understanding of this reaction.

Unfortunately, though, measuring at energies this low is not feasible, experimentally. This is because at these energies the signal-to-noise ratio of observables drops dramatically and the signals of interest are obscured within background noise. To provide accurate measurements of such situations, experiments must necessarily then occur in the timescale of years to measure a single datum. This is simply unfeasible, practically, and our understanding must rely on extrapolations from higher-energy data. As the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction is not alone in having a Gamow window at such low energies, extrapolation techniques must be employed to describe the cross section and S-factor for nearly all astrophysically important reactions at the Gamow window. This, then, in a nutshell highlights the primary goal of experimental nuclear astrophysics: to a) directly measure relevant cross-sections down to astrophysical energies when possible, or b) measure as low in energy as possible to provide the best constraint for theoretical extrapolations to Gamow energies.

1.3 Nuclear reactions

For the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction, both resonant and direct capture contributions are important. Resonant reactions are those in which an incoming particle combines with a target to form an excited state in a compound nucleus and then subsequently decay as part of a two-step process. Alternatively, direct capture reactions are those in which the particle pair proceeds directly to a final, bound nuclear state in one step. An energy level diagram of the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction is shown in Fig. 1.9.

1.3.1 Direct capture reactions

The other side of the proverbial coin from resonant reactions is direct capture, or nonresonant reactions. Such cases are defined as those in which the cross section and S -factor vary smoothly with energy, being the result of a one-step, electromagnetic process where the capture proceeds from the initial state of two incident particles to the final, bound state of the system. This process is also highlighted as a part of Fig. 1.9.

With nonresonant reactions, the S -factor varies smoothly with energy. This means that the earlier characterization of the S -factor, reaction rate, and Gamow energy/window parameters in Equations 1.6 - 1.15 are applicable for the case of direct reactions. In extrapolating experimental data to stellar energy ranges, it is often useful to express an effective S -factor, S_{eff} from the Taylor expansion of the component functions [34],

$$S_{\text{eff}} = S(0) \left(1 + \frac{5kT}{36E_0} \right) + S'(0)E_0 \left(1 + \frac{35kT}{36E_0} \right) + \frac{1}{2}S''(0)E_0^2 \left(1 + \frac{89kT}{36E_0} \right), \quad (1.16)$$

where E_0 is the Gamow peak energy from Equation 1.14. Armed with this expansion, the S -factor can be successfully extrapolated to low, stellar energies in cases where the S -factor varies slowly with energy. This means that for nonresonant reactions, $S(E)$ can be approximated by its zero-energy value, $S(0)$, and only small corrections given by its first and second derivatives, $S'(0)$ and $S''(0)$, respectively.

1.3.2 Resonant reactions

The presence of a resonance enhances the cross section at its specific center-of-mass energy and means that the S -factor no longer varies smoothly with respect to energy. The specific increase is highly dependent on the situation since, as a two-step process, it depends on the probability of both

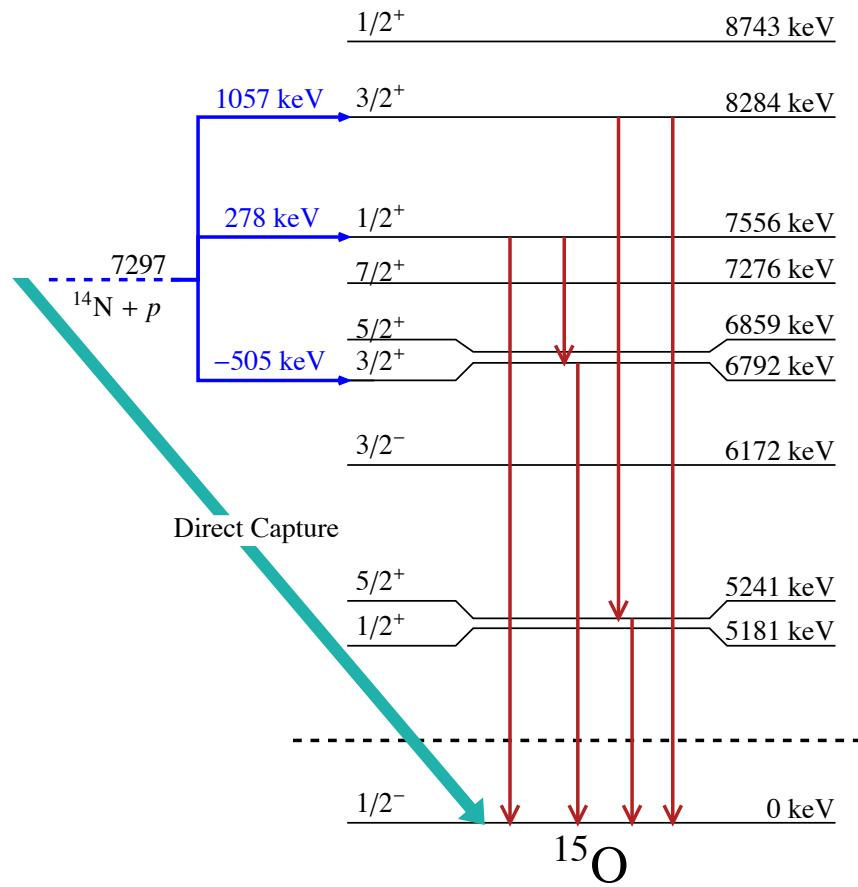


Figure 1.9. An energy level diagram of the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction. This shows the pathways of decay to the ground state of the ^{15}O nucleus. The paths shown in red indicate a resonance reaction, with the formation of the compound nucleus in an excited state, while the path in green shows a direct capture process, proceeding immediately to the final state (ground in this example) from any energy above the proton threshold.

forming the compound nucleus and decaying via a specific pathway, also shown in Fig. 1.9. To understand this mathematically, consider an isolated, narrow resonance at energy E_R ; the reaction cross section under the influence of such a resonance is described by the Breit-Wigner formula:

$$\sigma_{BW}(E) = \pi\lambda^2 \frac{2J_R + 1}{(2J_1 + 1)(2J_2 + 1)} \frac{\Gamma_{in}\Gamma_{out}}{(E - E_R)^2 + \Gamma_T^2/4} \quad (1.17)$$

where λ is the deBroglie wavelength corresponding to the incident momentum, $p = \hbar/\lambda$, J_1 and J_2 are the spins of the target and projectile, J_R is the spin of the resonance in the compound nucleus, Γ_{in} and Γ_{out} are the entrance and exit reaction channel widths (essentially the probability for a particular part of the process to occur), and Γ_T is the total width ($\Gamma_t = \Gamma_1 + \Gamma_2 + \dots + \Gamma_n$). The total width is related to the mean lifetime, τ , of the excited state by

$$\tau\Gamma_T = \hbar. \quad (1.18)$$

For most nuclear states, the widths can be determined experimentally, allowing for the understanding of the total width, Γ_T . However, due to many of the earlier described constraints, this is not always easy in practice. Additionally, the fact that different resonances have different strengths and can potentially dominate the contributions of others adds another complication. On the other hand, this implies that the total width will be dominated by that specific decay's width. If the dominant decay is one of the primary entrance or exit channels astrophysically, this will have a proportionally large effect on the overall cross section and reaction rate. In the case of the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction, there are two states near the proton threshold, one above and one below in energy, as shown in Fig. 1.9, and these two provide the largest contributions to the overall astrophysical reaction rate.

To characterize this, we replace the Breit-Wigner formula from Equation 1.17 into the reaction rate equation 1.6 [34]. This leads to

$$\langle\sigma v\rangle_{BW} = \left(\frac{8}{\pi\mu}\right)^{1/2} \left(\frac{1}{kT}\right)^{3/2} \int_0^\infty \sigma_{BW}(E) E \exp\left(-\frac{E}{kT}\right) dE. \quad (1.19)$$

Because we are considering a narrow energy range for this resonance, both the energy, E , and the exponential term provided by the Maxwell-Boltzmann velocity distribution, $\exp(-E/kT)$, are constant over the integration. This allows us to rewrite the integral as

$$\langle\sigma v\rangle_{BW} = \left(\frac{8}{\pi\mu}\right)^{1/2} \left(\frac{1}{kT}\right)^{3/2} E_R \exp\left(-\frac{E_R}{kT}\right) \int_0^\infty \sigma_{BW}(E) dE. \quad (1.20)$$

The previous integral is proportional to the area under the resonance cross section and is known as the resonance strength. By substituting the Breit-Wigner formula, the integral becomes

$$\int_0^\infty \sigma_{\text{BW}}(E) dE = \pi \lambda_R^2 \left(\frac{2J_R + 1}{(2J_1 + 1)(2J_2 + 1)} \right) (1 + \delta_{12}) \Gamma_{\text{in}} \Gamma_{\text{out}} \int_0^\infty \frac{1}{(E - E_R)^2 + (\Gamma_T/2)^2} dE, \quad (1.21)$$

which is solved analytically as

$$\int_0^\infty \sigma_{\text{BW}}(E) dE = \pi \lambda_R^2 \left(\frac{2J_R + 1}{(2J_1 + 1)(2J_2 + 1)} \right) (1 + \delta_{12}) \frac{\Gamma_{\text{in}} \Gamma_{\text{out}}}{\Gamma_T}. \quad (1.22)$$

For simplicity, the statistical factor, ω , is now defined as

$$\omega = \frac{2J_R + 1}{(2J_1 + 1)(2J_2 + 1)} (1 + \delta_{12}) \quad (1.23)$$

and the width ratio, γ , is

$$\gamma = \frac{\Gamma_{\text{in}} \Gamma_{\text{out}}}{\Gamma_T}, \quad (1.24)$$

leading to

$$\int_0^\infty \sigma_{\text{BW}}(E) dE = 2\pi^2 \lambda_R^2 \omega \gamma. \quad (1.25)$$

The product $\omega\gamma$ is called the resonance strength and is very important for reaction rate calculation, since the rate is dependent primarily on it and the resonance energy, E_R . This can be seen by implementing this integral calculation with the earlier derivation in Equation 1.19, resulting in

$$\langle \sigma v \rangle = \left(\frac{2\pi}{\mu kT} \right)^{3/2} (\omega\gamma) \hbar^2 \exp \left(-\frac{E_R}{kT} \right). \quad (1.26)$$

1.4 R-matrix theory

In many cases, the nuclear cross-section landscape is quite complicated, with the combined effects of multiple resonances and direct capture contributions. With all of this overlap, the Breit-Wigner formulation starting in Equation 1.17 is no longer valid. Therefore, a reliable theory accounting for all contributions and interferences is needed to describe relevant information and perform meaningful extrapolations. For these purposes, *R*-matrix theory was developed and is now commonly used in

nuclear astrophysics calculations; details of its formal derivation can be found in References [11, 39].

The core concept of *R*-matrix theory is to describe the wave function of the system by matching individual components of the wavefunction in both the region of the compound nucleus where the nuclear potential is present and also to an exterior region in which only the Coulomb potential contributes (and nucleons are treated as individual particles). At the boundary of these two regions, known as the channel surface and occurring at the channel radius, a_c , the wave functions and their derivatives must match. The *R*-matrix relates these wave functions for each different reaction channel and is defined as

$$\mathbf{R}_{c'c} = \sum_{\lambda} \frac{\gamma_{\lambda c'} \gamma_{\lambda c}}{E_{\lambda} - E}, \quad (1.27)$$

where c and c' are the respective entrance and exit channels of the given reaction and the γ_{λ} is the reduced width, given as

$$\gamma_{\lambda}^2 = \hbar^2 / (2ma_c) u_{\lambda}^2(a_c), \quad (1.28)$$

with the u_{λ} forming a complete set of basis states that satisfy the Schrödinger equation. The reaction channels are defined as

$$c = \alpha s \ell, \quad (1.29)$$

where the α 's identify the interacting particle pair, the ℓ 's are the respective orbital angular momenta of the particles, the s 's are their respective spins.

While the theory is named for the *R*-matrix, the cross section is actually calculated through other intermediate quantities, namely the collision matrix, \mathbf{U} , and the transition matrix, \mathbf{T} . The collision matrix relates specifically the incoming channels, y_c , to those of the outgoing particles, x_c , defined as

$$x_c = - \sum_{c'} \mathbf{U}_{c'c} y_{c'}. \quad (1.30)$$

Ultimately, \mathbf{U} is proportional to the *R*-matrix as

$$\mathbf{U} \propto |\mathbf{R}|^2. \quad (1.31)$$

The transition matrix, on the other hand, is defined through the collision matrix as

$$\mathbf{T}_{c'c} = e^{2i\omega_c} \delta_{c'c} - \mathbf{U}_{c'c}, \quad (1.32)$$

where ω_c represents the Coulomb phase shift and $\delta_{c'c}$ is the Dirac delta function relating channels c' and c . The angle integrated cross section is then

$$\sigma_{\alpha'\alpha} = \frac{\pi}{k_\alpha^2} \sum_{J\ell'\ell s's} \omega |\mathbf{T}_{c'c}^J|^2 \quad (1.33)$$

where J is the total angular momentum of the system and ω is the statistical spin factor as defined by Equation 1.23.

In this work, the multichannel, multilevel R -matrix code AZURE2 [11] was used to perform the calculations. A user inputs the spin-parity of the nuclear states, initial energies, partial widths of the excited states, the asymptotic normalization coefficient's (ANC's) of bound states (which describe the tail of the interaction below the particle threshold), and any external cross section or S -factor data into AZURE2. The computation of S -factor and cross section are done via a simple GUI interface. The results are truncated to finite eigenstates behind the scenes to allow the code to operate quickly. As such, a background pole is typically included at high-energy to account for contributions from states not included directly in the calculations. The theory is also referred to as phenomenological R -matrix because each of the γ_λ 's and E_λ 's in Equation 1.27 are free fit parameters.

1.5 The $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction

The $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction has been investigated many times in recent years by a wave of campaigns spurred by new facilities, equipment, and analyses [17–19, 24, 29, 30, 35, 36, 40, 42–45, 47, 53, 56, 57, 61, 63, 65]. Collectively, these experiments represent an effort to understand the astrophysical S -factor at stellar energies through both direct and indirect methods. The indirect approaches have two faces: measuring the width of the subthreshold state at excitation energy $E_x = 6792$ keV ($E_R = -505$ keV) and the Asymptotic Normalization Coefficient (ANC) of the ground state, while the direct approach is simply that - measuring the cross section to low energies and extrapolating from that data to astrophysical energies, usually with the help of theories like R -matrix. The level scheme of the reaction product of ^{15}O is shown in Fig. 1.9.

1.5.1 Reaction cross section

Cross section measurements of the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction have been made and the contributions of the different γ -ray transitions have been disentangled to characterize the behavior of the S -factor and provide more reliable low-energy extrapolations. Measurements have ranged as low in energy as 70 keV (center-of-mass), still far above the solar Gamow window at 27 keV. Five different capture transitions contribute to the cross section at low energies: those proceeding through the DC/R \rightarrow gs (Direct capture or resonant capture to the ground state), the DC/R \rightarrow 6.79 MeV primary transition, the DC/R \rightarrow 6.17 MeV primary transition, the DC/R \rightarrow 5.24 MeV primary transition, and the DC/R \rightarrow 5.18 MeV primary transition. In evaluating the total S -factor, each of these transitions contributes and must therefore be known in turn.

The first comprehensive measurement of the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction was performed by Schröder *et al.* in 1987 [56]. They studied the reaction with protons ranging in energy from 0.2 - 3.6 MeV and found that the DC/R \rightarrow gs transition accounted for near 50 % of the total S -factor when extrapolated to zero energy, which itself was determined to be $S(0) = 3.2 \pm 0.54$ keV b. The dominant uncertainty the authors found was caused by the interference of the subthreshold state at 6.79 MeV (505 keV below the proton threshold) with the ground state capture, assigning the state a width of $\Gamma^{6.79} = 6.3$ eV. These results were then used in further reviews [2] and the NACRE tabulations of reaction rate data [7].

In 2001, however, Angulo *et al.* revisited the data from Schröder *et al.* [56] using the R -matrix formalism and recommended an $S(0)$ value lower by nearly a factor of two [6]. This owed primarily to two discoveries in the analysis: a new value for the contribution from the ground state capture, revising it to $S_{gs}(0) = 0.08$ keV b, and the dominance that the width of the 6.79 MeV level shows on the behavior of $S(0)$. Consequently, they recommended new values for the total S -factor of $S(0) = 1.77 \pm 0.20$ keV b and a gamma width of $\Gamma^{6.79}=1.75$ eV, nearly a factor of four change. The authors also identified the contribution of the DC/R \rightarrow 6.17 MeV primary transition to be the other important transition, with all others accounting for < 5% of the total astrophysical S -factor. This study, similar to its subject reaction, set off a chain of new measurements and re-analyses of data with R -matrix techniques. The vast majority of these studies were done by the groups at TUNL and LUNA, with the results of all studies tabulated in Table 1.1.

The most recent measurements were published by Li *et al.* in 2016 [42] and Wagner *et al.* in 2018 [63]. These experiments differed in their aim. The Li experiment focused on higher-energy contributions and determined radiative capture data for capture to the 6.79 MeV level in the 1.5 - 3.4

Year	Reference	Astrophysical $S(0)$ (keV b)			
		R/DC \rightarrow 0.00	R/DC \rightarrow 6.79	R/DC \rightarrow 6.17	Others ^d
1987	Schröder <i>et al.</i> [56]	1.55 \pm 0.34	1.41 \pm 0.02	0.14 \pm 0.05	0.1
2001	Angulo <i>et al.</i> ^a [6]	0.08 ^{+0.13} 0.08 ^{-0.06}	1.63 \pm 0.17	0.06 ^{+0.01} 0.06 ^{-0.02}	—
2003	Mukhamedzhanov <i>et al.</i> [48]	0.15 \pm 0.07	1.40 \pm 0.20	0.133 \pm 0.02	0.02
2004	Formicola <i>et al.</i> [29]	0.25 \pm 0.06	1.35 \pm 0.05 (stat) \pm 0.08 (sys)	0.06 ^{+0.01b} 0.06 ^{-0.02}	0.04
2005	Imbriani <i>et al.</i> [36]	0.25 \pm 0.06	1.21 \pm 0.05	0.08 \pm 0.03	0.07
2005	Runkle <i>et al.</i> [53]	0.49 \pm 0.08	1.15 \pm 0.05	0.04 \pm 0.01	—
2005	Angulo <i>et al.</i> [8]	0.25 \pm 0.08	1.35 \pm 0.04	0.06 \pm 0.02	0.04
2006	Bemmerer <i>et al.</i> [17]	—	—	—	—
2008	Marta <i>et al.</i> [43]	0.20 \pm 0.05	—	0.09 \pm 0.07	—
2010	Azuma <i>et al.</i> [11]	0.28	1.3	0.12	0.11
2011	Adelberger <i>et al.</i> [1]	0.27 \pm 0.05	1.18 \pm 0.05	0.13 \pm 0.06	0.08
2016	Li <i>et al.</i> [42]	0.42 \pm 0.04 (stat) $^{+0.09}_{-0.19}$ (sys)	1.29 \pm 0.06 (stat) \pm 0.06 (sys)	—	—
2018	Wagner <i>et al.</i> [63]	0.19 \pm 0.01 (stat) \pm 0.05 (sys)	1.24 \pm 0.02 (stat) \pm 0.11 (sys)	—	—

Table 1.1: A summary of previous S -factor determinations. a) R -matrix analysis on available data, not a measurement. b) Adopted from Angulo *et al.* [6]. c) Measured S -factor at 70 keV. d) Calculated difference of the total $S(0)$ from the other transitions.

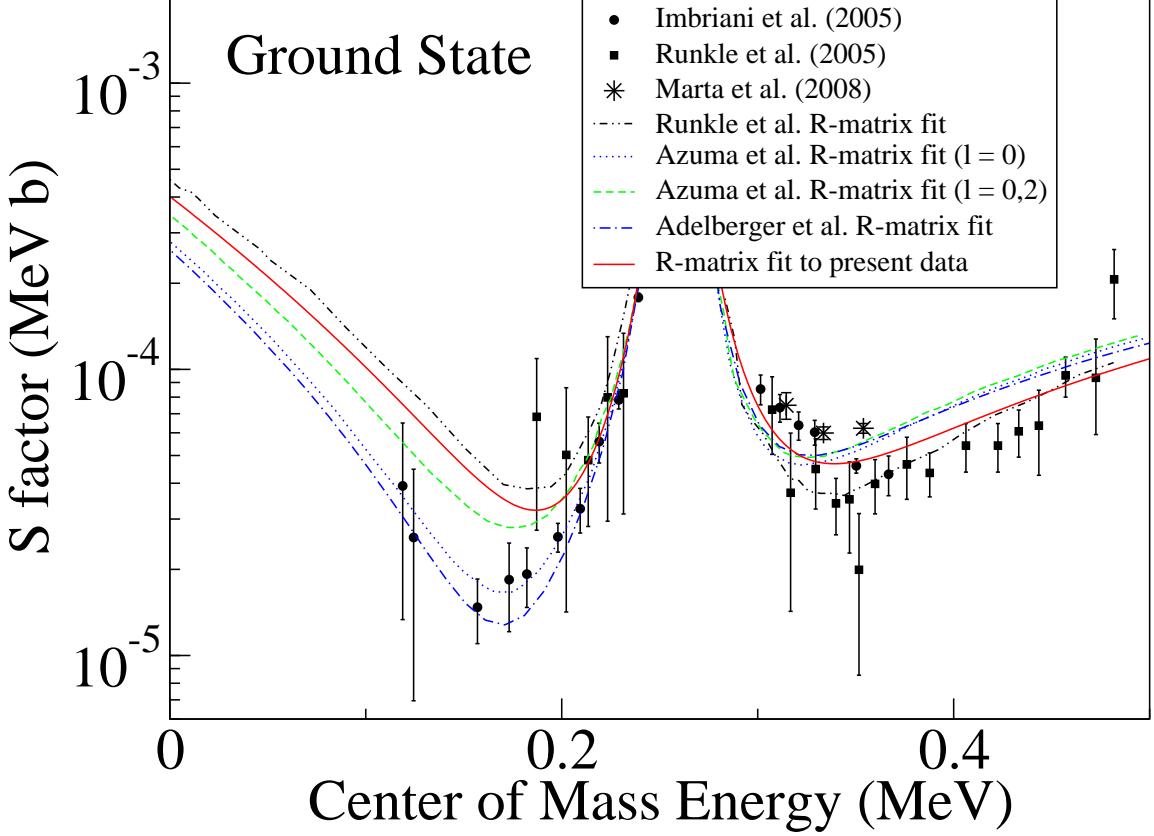


Figure 1.10. *R*-matrix extrapolation of the *S*-factor given in [42]. The authors identified the major sources of uncertainty as the uncertainty as the DC/R \rightarrow gs primary transition, the DC/R \rightarrow 6.17 MeV level primary transition, and the Γ width of the 6.79 MeV state.

Of these, the latter is the dominant term. This figure highlights the effects that these uncertainties have on low-energy extrapolations, varying by nearly a factor of two.

MeV proton energy range and the proton energy range of 0.6 - 3.4 MeV for the DC/R \rightarrow gs transition from 0.6 to 3.4 MeV. The authors also reported angular distributions for their measurements. The authors identified the primary, remaining sources of uncertainty as the DC/R \rightarrow gs primary transition, the DC/R \rightarrow 6.17 MeV level primary transition, and the Γ width of the 6.79 MeV state. Fig. 1.10 shows the effects of these uncertainties in the authors' *R*-matrix calculations and how it differs from other calculations. Due to the astrophysical importance of this reaction detailed in Section 1.1, an accurate determination of this extrapolated *S*-factor is of the utmost importance. Critically, the data reported do not overlap with the low energy measurements of the LUNA and TUNL groups and their absolute scale does not align, also leading to further inconsistent extrapolation and significant error in the deduced $S(0)$ value.

The Wagner *et al.* study attempted to remedy this, providing an independent cross check of

the ground state and 6.79 MeV capture transitions in the proton energy range of $E_p = 366\text{--}1289$ keV, overlapping with the low-energy data sets. In the study, the authors found S -factors elevated significantly above those of any previous measurement for both of the transitions measured. Due to this, their results for the extrapolated $S(0)$ and contributions from the DC/R \rightarrow 6.79 MeV transition agree with those from Li *et al.*, but they are unable to draw conclusions about the ground state transition or any of the other, weaker transitions. Ultimately, this means that their results were unable to address any of the major uncertainties left plaguing the situation.

1.5.2 Lifetime

The Γ width of the 6.79 MeV subthreshold state is the dominant uncertainty on the total S -factor at zero energy, as illustrated by Fig. 1.10. The literature to date on the lifetimes/widths of the important levels in ^{15}O are summarized in Table 1.2. Some of the values present in the table are those extracted as a fit parameter from R -matrix analyses, instead of experimental measurements [1, 6, 11, 43, 53, 56?]. By having a dominant term of the calculation be a floating fit parameter, accuracy and reliability in the extrapolations are lost. Therefore, determination of this quantity experimentally is critical.

There are five measurements of the lifetime or width of the 6.79 MeV state "directly" [18, 30, 47, 57, 65], highlighted in red in Table 1.2, and they are unfortunately disparate in their extracted values and have relatively large errors.

The first measurement was performed in 2001 by Bertone *et al.*, using the Doppler Shift Attenuation Method (DSAM) to determine the lifetime of the state. This technique will be discussed in further detail later. The authors populated the state through resonant capture of protons in the $E_p = 278$ keV state and observed the decaying γ -rays at three different angles. The measured lifetime for the 6.79 MeV state was determined to be $\tau = 1.60_{-0.72}^{+0.75}$ fs, corresponding to a width of $\Gamma = 0.41_{-0.13}^{+0.34}$ eV (determined by Equation 1.18).

At RIKEN, in 2004, Yamada *et al.* performed a measurement of the radiative width via Coulomb Excitation experiment [65] as an independent verification of the Bertone result. For this, ^{15}O nuclei were produced by fragmentation of an ^{16}O beam incident on a ^9Be target and then inelastically scattered off of a thick lead target. The de-excitation γ -rays were detected with an array of NaI (Tl doped) scintillation detectors. In order to disentangle the 6.79 MeV peak of interest from the 6.86 Mev peak in their spectrum, the authors compared the experimentally obtained spectrum from with the results of a Monte Carlo simulation. Due to the poor energy resolution of the detector system,

	$\tau_{6.79}$ fs	$\tau_{6.17}$ fs	$\tau_{5.18}$ fs	$\Gamma^{6.79}$ (eV)
Gill <i>et al.</i> [31]	< 28	< 2.5	8.2 ± 1.0	--
Schröder <i>et al.</i> [56]	--	--	--	6.3
Bertone <i>et al.</i> ^a [18]	$1.60^{+0.75}_{-0.72}$	$2.10^{+1.33}_{-1.32}$	$9.67^{+1.34}_{1.24}$	$0.41^{+0.34b}_{-0.13}$
Angulo <i>et al.</i> [6]	--	--	--	1.75 ± 0.60^c
Yamada <i>et al.</i> [65]	$0.69^{+1.10}_{-0.69}$	--	--	$0.95^{+0.60}_{-0.95}$
Formicola <i>et al.</i> [29]	--	--	--	0.8 ± 0.4^c
Schürmann <i>et al.</i> [57]	< 0.77	< 0.77	8.4 ± 1.0	> 0.85
Azuma <i>et al.</i> [11]	--	--	--	2.38^c
Adelberger <i>et al.</i> [1]	--	--	--	3.25^c
Michelagnoli ^{a,d} [47]	< 1.0	--	--	> 0.66
Galinski <i>et al.</i> ^b [30]	< 1.8	< 2.5	--	> 0.44

TABLE 1.2

A SUMMARY OF PREVIOUS LIFETIME/WIDTH MEASUREMENTS FOR
 IMPORTANT STATES IN ^{15}O . A) MEASUREMENTS REPORTED WITH 90%
 CONFIDENCE LIMIT. B) MEASUREMENTS REPORTED WITH 68.3%
 CONFIDENCE LIMIT. C) REPORTED VALUE CALCULATED FROM R -MATRIX
 ANALYSIS. D) REPORTED IN PH.D. THESIS, NOT PEER REVIEWED.

the authors ultimately had difficulty separating the peaks and only reported an upper bound on the width, $\Gamma = 0.95^{+0.60}_{-0.95}$ eV, and a corresponding lifetime of $\tau = 0.69^{+1.10}_{-0.69}$ fs.

Four years later, Schürmann *et al.* published a new measurement of the lifetime using the DSAM technique and an improved experimental setup [57]. This measurement used a single high-purity germanium (HPGe) detector on a rotating track and measuring at many more angles ranging from $40 - 116^{\text{deg}}$, allowing for a check of asymmetries around the 90^{deg} point and reducing systematic uncertainties from the use of different detectors. Similar to the Bertone measurement [18], the authors populated the state of interest via the resonance at $E_p = 278$ keV. In contrast to the Bertone result and similar to the Yamada work, the authors in this case could not recommend a finite value for the measured lifetime, obtaining a lifetime upper limit of $\tau < 0.77$ fs.

In an attempt to provide another experimental technique for determining the lifetime, Galinski *et al.* performed another DSAM measurement at the TRIUMF facility [30]. The major difference this time, however, was that the reaction occurred in inverse kinematics (i.e. one where the heavier reaction input is accelerated towards the stationary, lighter one), emulating a result from over 50 years ago [31]. The ${}^{15}\text{O}$ in this experiment were produced via the ${}^3\text{He}({}^{16}\text{O}, \alpha){}^{15}\text{O}$ reaction with a ${}^3\text{He}$ -implanted gold-foil and the de-excitation γ -rays were detected with a clover HPGe detector (a single detector containing four separate Ge crystals) and filtered in coincidence with the reaction product α 's. The authors then determined the lifetime with a Bayesian line-shape analysis of the spectra and found a most probable lifetime of $\tau = 0$ fs and an upper limit given by experimental uncertainties of $\tau < 1.8$ fs with a 68.3% confidence limit. The authors, however, acknowledge that their result was statistics limited.

The final measurement of this lifetime comes from an unpublished doctoral dissertation by Michelagnoli in 2013 [47]. Similar to Galinski *et al.*, this measurement was done in inverse kinematics, with the $d({}^{14}\text{N}, {}^{15}\text{O})n$ reaction. The γ rays from this measurement were detected with the state-of-the-art AGATA Demonstrator array using γ -ray tracking for precision angular data and a lineshape analysis of the data was performed by comparing with the results of a Monte Carlo simulation in the Geant4 software. This work also obtained only an upper-bound on the lifetime, $\tau < 1.0$ fs with a 68.3% confidence limit. While this result is consistent with other literature values, it should nonetheless be taken with a grain of salt as it has not been peer-reviewed.

In aggregate, these results are unsatisfactory, with an error that is still uncomfortably large. Particularly, the DSAM results (with the exception of Galinski [30]) neglected the systematic uncertainties from the stopping power of the targets in their ultimate lifetime calculation. The forward

kinematics reactions [18, 57] both used nitrogen-implanted tantalum-foils for the experiment (saturated to a composition of Ta_2N_3), but calculated the lifetimes assuming target densities and stopping powers equal to those of pure tantalum. However, in this lifetime range, the deceleration of the ^{15}O nuclei occurs in the region where the nitrogen is present in the lattice. Specifically, this means that the presence of the nitrogen has a significant effect on the actual stopping power. According to the authors of the respective works, by using accurate densities and stopping powers the inferred lifetimes obtained in their works would have changed by 100% [18] and 2% [57], respectively. In both, the authors also contend that their calculations are appropriate because they used an identical method to calculate the lifetime of the 5.18 MeV state, agreeing with literature values.

1.6 Objective

The results of a series of experiments and subsequent analyses performed to better understand the low energy behavior of the $^{14}N(p, \gamma)^{15}O$ will be presented herein. These include the measurement of the $^{14}N(p, \gamma)^{15}O$ reaction at the University of Notre Dame Nuclear Science Laboratory (NSL) and the Compact Accelerator System for Performing Astrophysical Research (CASPAR), located in the Sanford Underground Research Facility in the renovated Homestake Gold Mine in South Dakota, as well as a measurement of the lifetime of the 6.79 MeV level in ^{15}O also at the NSL. These data will then be used in an *R* Matrix calculation to provide a new determination of the $^{14}N(p, \gamma)^{15}O$ reaction rate.

An outline of the experimental techniques used in these measurements will be presented in Chapter 2, including specific details about each phase of the measurement. There will be a brief overview of the standard equipment common to each of the measurements. Following this, the specific features of the different types of measurements and the facilities in which they were made will be presented in further detail.

Chapters 3 - 4 will present details on the different facets of the data reduction and analysis. The first of these, Chapter 3, shows the data for $^{14}N(p, \gamma)^{15}O$ taken at the NSL and CASPAR while detailing the process by which the cross-section and astrophysical *S*-factor were extracted. Shifting focus, Chapter 4 will focus on the nuclear lifetimes, describing the codes that were developed and used to analyze the Doppler Shift in the γ -ray data to determine the lifetime of the states in ^{15}O . The validity of the method will be checked against other known levels populated in the reaction, like the states at 5.18 MeV and 6.17 MeV, respectively.

The cross-section information and newly measured lifetime will be utilized in a new *R*-matrix

analysis of the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction, presented in Chapter 5. The new constraints these data provide on the low-energy behavior of the extrapolated *S*-factor will be provided, alongside a description of their impact to relevant astrophysical quantities.

Finally, Chapter 6 will present an overview of the respective impacts of the measurements paired with future outlook.

CHAPTER 2

EXPERIMENTAL SETUP AND PROCEDURES

2.1 Introduction

In this chapter, the details of the different experimental conditions for each phase of the dissertation are presented. Following that, sections are also included to highlight some techniques and equipment commonly used in low-energy nuclear physics, with an emphasis on those used in this work. The discussion is presented from the lens of the general to the specific, introducing general concepts for beam production and transport, acceleration systems, and radiation detection. Following the discussion of the scientific principles, even more detail about the facilities and equipment is given, highlighting differences between the facilities and techniques employed when compared to common equipment for nuclear physics experiments.

2.2 Experimental Overview

2.2.1 Setup: cross section

To measure the cross section of the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction, a campaign of experiments was performed at the University of Notre Dame Nuclear Science Laboratory (NSL) and at the CASPAR facility, part of the Sanford Underground Research Facility in Western South Dakota. More data about these facilities can be found in Sections 2.4.1 and 2.4.2, respectively.

First, measurements were performed at the NSL in January 2018 to provide an above ground data set with which to compare to the underground measurements made at CASPAR throughout 2018. A schematic diagram of the setup used in these measurements is given in Fig. 2.6 while pictures of the setup are shown as part of Fig. 2.11 and Fig. 3.3. The schematic diagram, however, purposefully lacks the lead castle placed around the detector to shield from radiogenic background as it created a more confusing image. The effects of this lead shielding can be clearly seen in Fig. 2.13. Additionally, these pictures are of the setup at the CASPAR facility, specifically. The equipment used and its placement was entirely the same for the data taken at either the NSL or CASPAR, less

a few lead bricks. Therefore, these pictures are sufficient to present the setup for all phases of the experiment.

At the NSL, the 5MV 5U single-ended Van de Graaf electrostatic accelerator was used to produce beams of ^1H ions, created with the electron cyclotron resonance ion source, at energies ranging between 800 keV and 1200 keV. This energy range was chosen as it provides an overlap to the CASPAR facilities capabilities and lower energies were prevented due to stability issues in the accelerator at the time. At the CASPAR facility, on the other hand, the ^1H ions were created with a radiofrequency ion source and accelerated with a 1MV JN single-ended Van de Graaf electrostatic accelerator at energies ranging from 270 keV up to 1060 keV. This energy range is the entirety of the JN accelerator's operating range. While it is technically possible to push the accelerator down to voltages below 200 kV (and corresponding proton beam energies of <200 keV), doing so required a complete reconfiguration of the corona stack inside the accelerator and was deemed too insignificant a gain in measured data for the time that such a change necessitated. More detail on the operating principles of these accelerators and ion sources are given in Sections 2.3.1 and 2.3.2, respectively.

In both facets of the experiment, a magnetic dipole was used to select the desired beam after production. A combination of electrostatic and magnetic steering and focusing elements guided the beam to the target station through the beamline under vacuum. The principles upon which these systems operate are given in Section 2.3.3.

The nitrogen targets for these experiments were both evaporated TiN and sputtered ZrN targets, all produced in Bochum, Germany. The TiN targets were backed with Ta while the ZrN targets were backed with W. All backings were machined to 2 inch squares before the addition of the active target material, after which point the corners were cut off so that the target would fit inside the holder. The targets were produced to nominal thicknesses of 50 nm (ZrN) and 100 nm (both ZrN and TiN). Targets were held in place during the experiment by securing them inside a target holder that was attached to the end of the beam line. To protect the targets from damage, the holder was connected to a water chiller which passed water through the back of the holder, cooling the targets constantly throughout the experiment. The use of rubber O-rings and hoses made the target holder electrically isolated from the rest of the beam line, allowing it to be used as a measure of the charge delivered directly, without the use of external Faraday cups. Additionally, to reduce carbon buildup on the target through the course of the runs, a cold trap was employed in front of the target for the measurements. This can be seen in Fig. 2.6 and is detailed in Section 2.3.3. The liquid nitrogen reservoir was filled every two hours, keeping the copper finger sufficiently cold throughout

the duration of the measurements.

In tuning, the beam spot on target was viewed with a quartz viewer or a MgO scintillation target, ensuring good quality of the delivered beam. In keeping the beam focus and centered through the runs, some of which lasted hours, pairs of upstream slits were utilized at both laboratories. These slits were moved to the edge of the beam at the end of the tuning process. When used together, a pair of slits defines a central line through which the beam passes to the center of the target. If the beam drifts or loses focus one (or more) of the slits will record a current. Thus, these enable a consistent delivery of beam to target throughout running. The locations of the slits, tuning elements, and the target stations at the NSL are presented in Figs. 2.9, 2.5, and 2.10 while that information for the CASPAR facility is depicted in Fig. 2.12. Additionally, the beam delivered to the target at the NSL was in the range of 5 - 10 μA whereas at CASPAR it was on the order of 50 - 120 μA .

In this measurement, the γ -rays were observed with a coaxial p-type HPGe detector (manufactured by Canberra) rated with 130% relative efficiency (relative to the efficiency of a 3" \times 3" NaI detector with 1332 keV γ -rays). The HPGe detector was biased with +4000 V from an Ortec 660 Power Supply module. 1.5 mm of lead shielding was placed in front of the crystal's face to reduce the number of low-energy x-rays entering the detector (< 100 keV photon energy) because they increase the dead time in the electronics decreasing processing time for signals of interest. A lead castle was built around the detector at each location to reduce radiogenic background from the target rooms. The arrangement of the lead around the detector was not the same between the two locations due to availability restrictions, but they provided similar background reduction at both locations. The detector was placed at an angle of 55° in order to minimize any angular distribution effects on the cross section, as that is the minimum of the second order Legendre polynomial which describes the angular behavior of the cross section. Additionally, the detector was placed on a sliding, isolating platform, enabling its distance to the target to be easily changed. Measurements were primarily taken with the detector in close geometry, with only 2 mm between the target and the center of the lead shielding on the face of the detector (closer would have resulted in them touching on the corner because of their different orientation) but some were measured in far geometry, up to 25.4 cm between the target and lead shielding. This was done to provide a reference for the effects of summing on our detector response at high energies, which will be discussed further in Section 3.4.

The electronics necessary for these measurements were minimal, since the HPGe detector has a pre-amplifier for signal processing attached to the charge recorder. Therefore, it was connected to an Ortec 672 Spectroscopic Amplifier with coarse gain set to 5, fine gain set to 10, Gaussian

shaping, automatic pole zero adjustment, and a shaping time of $6\ \mu\text{s}$. Following this, the signals were sent to an Ortec 927 ASPEC Multichannel Analyzer (dual input model) which digitized the signals and was used in concert with the MAESTRO analysis software for recording the spectra.

2.2.2 Setup: lifetimes

The goal of this measurement was to determine the lifetime of the 6.79 MeV state with a variety of target materials and nitrogen contents in order to identify any systematic effects which were unidentified in previous measurements (see Section 1.5.2). In August 2019, this measurement was undertaken at the NSL using the Doppler Shift Attenuation Method (DSAM) in forward kinematics with the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction. Details of the DSAM can be found in Section 2.5.1. A schematic diagram of the setup used in these measurements, which was the same as used during the cross section experiments, is shown in Fig. 2.6. At the NSL, the 5MV 5U single-ended Van de Graaf electrostatic accelerator was used to produce beams of ^1H ions, created with the electron cyclotron resonance ion source, at energies $E_p = 1020\ \text{keV}$ and $E_p = 1570\ \text{keV}$, which provided good signal-to-background discrimination for the states of interest. At $E_p = 1570\ \text{keV}$, the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction also produced ^{15}O nuclei in the 5.24 MeV and 6.18 MeV excited states, which have known lifetimes of $8.2 \pm 1.0\ \text{fs}$ and $< 2.5\ \text{fs}$, respectively [4]. Both of these states are also within the range of DSAM to measure, allowing additional verification of results for the measurement of the 6.79 MeV state in ^{15}O .

After creation and acceleration of the ^1H ion beam, a magnetic dipole selected the desired beam product and energy. A combination of electrostatic and magnetic steering and focusing elements guided the beam to the target station through the beamline under vacuum. The principles upon which these systems operate are given in Section 2.3.3.

In tuning, the beam spot on target was viewed with a quartz viewer ensuring good quality of the delivered beam. In keeping the beam focus and centered through the runs, some of which lasted hours, pairs of upstream slits were utilized at both laboratories. These slits were moved to the edge of the beam at the end of the tuning process. When used together, a pair of slits defines a central line through which the beam passes to the center of the target. If the beam drifts or loses focus one (or more) of the slits will record a current. Thus, these enable a consistent delivery of beam to target throughout running. The locations of the slits, tuning elements, and the target stations at the NSL are presented in Figs. 2.9, 2.5, and 2.10. Additionally, the beam delivered to the target fell in the range of 20 - 50 μA .

For this experiment, implanted nitrogen targets were produced in July 2019 at the NSL and utilized for the measurement. Section 2.5.2 details the creation process and properties of the various fabricated targets. From these, the targets ultimately selected for use in the experiment were the high-dose Ta, W, and Mo implanted targets to highlight differences in backing materials while the low-dose Ta target was measured at the end of the experiment to provide a comparison on the nitrogen content in the targets. Specifically, these were selected to highlight the desired effects and be completed within the experimental time constraint. If more time had been available, all of the generated targets would have been studied for a complete mapping of the parameter space. The results of this undertaking as well as the data reduction and analysis necessary to arrive at those conclusions will be detailed in Chapter 4.

In this measurement, the γ -rays were observed with one of the GEORGINA detectors, a coaxial n-type HPGe detector rated with 100% relative efficiency (relative to the efficiency of a 3" \times 3" NaI detector with 1332 keV γ -rays). 1.5 mm of lead shielding was placed in front of the crystal's face to reduce the number of low-energy x-rays entering the detector (<100 keV photon energy) because they increase the dead time in the electronics. The detector was placed a rotating table with angular precision such that the detector's center could be placed within $\pm 0.5^\circ$ of the intended angle. Angles used in the course of the experiment were 0° , 45° , 60° , 75° , 90° , 111° , 135° , and -90° . Measurements were taken with the detector in far geometry, with 25.4 cm between the target and lead shielding at the crystal's face. At this distance, the entire crystal face spanned an angular range of $\pm 9^\circ$. This measurement was also performed on the solid target beam line within the Sta. Ana target room, marked as the GEORGINA position in Fig. 2.10. Unlike the cross-section measurement, however, the yield produced in the reaction is unimportant, so there was no need to account for efficiency or summing in this case. Additionally, to prevent any artificial peak shifting arising from gain shifts in the electronics to enter into the final product, the data were saved and the data acquisition restarted hourly at each angle. This allowed us to monitor for gain shifting throughout the experiment prevent it from obfuscating results. Ultimately, no evidence of gain shifting was found throughout the experiment. A CeBr detector was fixed at a backward angle of -135° close geometry in order to provide a consistency check throughout the experiment.

The HPGe detector used for this measurement was biased to -4500 V with an Ortec 660 power supply module. It, like the detector used in the cross section measurements, comes with a pre-amplifier for signal processing. For this measurement, though, the Mesytec MDPP-16 module proved invaluable. It acted as the signal amplifier, analyzer, and digitizer. Through it, the settings

applied were shaping time = 480, signal rise time = 40, pole-zero adjustment = 5000, gain = 1.24, and threshold = 320. This module also comes with its own, proprietary analysis software for recording and saving the data files, which we also employed.

2.3 Experimental Equipment

2.3.1 Accelerators

First and foremost, low-energy nuclear physics (generally understood to be the regime in which incident kinetic energies for reactions are below 1 GeV, often even as low as sub-MeV) requires particle acceleration systems in order to bring beams and targets together. The most common techniques for accelerating ions are 1) to use strong electrostatic fields for a straight-line acceleration, such as in the Van de Graaff accelerator, or 2) by using a combination of electric and magnetic fields to accelerate the particles in cyclotron motion, aptly named a cyclotron. Van de Graaff accelerators are the most prominent of all accelerators, though, and are the type used in this work.

The basic idea of a Van de Graaff accelerator is shown in Fig. 2.1. The governing principle is that a beam of ions produced in an ion source (Section 2.3.2) are manipulated by additional electromagnetic elements (Section 2.3.3) into an area of high electric potential. This high voltage is produced and maintained by continuously transporting positive charge from ground to a Faraday cage with a field free interior (referred to as the terminal, with voltage V_T). The positively charged ions, being in close proximity to the positively charged terminal feel a repulsive force, accelerating the ions out of the terminal and through the machine. Upon exiting the accelerator, the ions have energy

$$E_{ion} = qV_T = \frac{1}{2}m_{ion}v_{ion}^2 \quad (2.1)$$

where V_T again is the terminal voltage and q is the charge state of the ion going through the accelerator, m_{ion} is the mass of the ion, and v_{ion} is the ion's velocity. When the terminal voltage is in MV, the beam energy is then provided in MeV. Specifically for the 5U accelerator, though, Equation 2.1 must be modified with an additional term. Inside the ion source, there is an electrostatic extractor, which transports the beam of ions out of the terminal and into the acceleration tube. With this correction, the beam energy for the 5U is given as

$$E_{ion} = q(V_{extractor} + V_T). \quad (2.2)$$

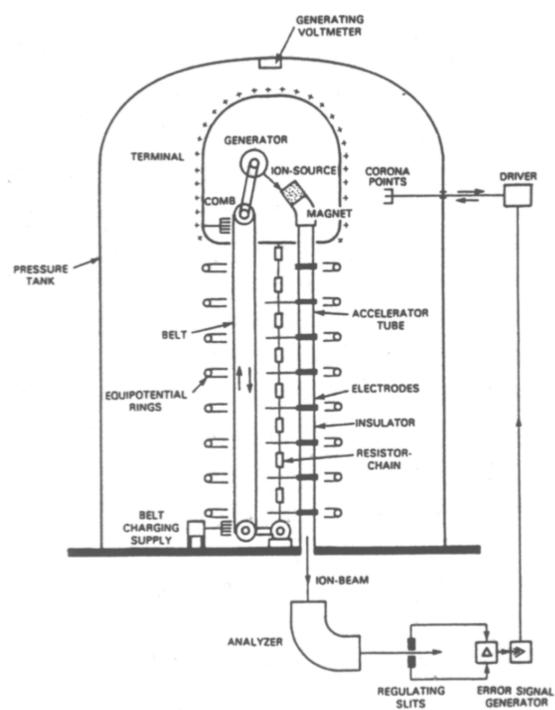


Figure 2.1. A diagram of a Van de Graaff accelerator [52]. The charge transportation in this figure is shown with a belt, like the JN accelerator model at CASPAR, while the 5U accelerator at the NSL uses a Pelletron chain.

The charge is delivered to the terminal via a mechanical delivery system. The 5U Sta. Ana accelerator at the NSL uses four Pelletron chains [33], each with its own power supply, while the JN Van de Graaff accelerator at CASPAR utilizes a rotating belt. In either case, the entire charging system is contained inside of a pressurized tank of insulating gas, SF₆ for the 5U and CO₂ for the JN, reducing the chance of a terminal discharge to ground from dielectric breakdown. This is not the only element to ensure the operational stability of the machine. The charging systems supply charge continuously to the terminal. However, charge is always being removed from the terminal via three primary pathways: 1) the beam of ions passing through the accelerator, 2) the resistive column, and 3) the corona system. In order to maintain stability, the currents that pass through these systems must always be in balance according to Kirchoff's Rule,

$$I_{chains} = I_{beam} + I_{column} + I_{corona}. \quad (2.3)$$

The column is actually a series of equipotential hoops connected by equal resistors. Each conducting hoop provides a small step down in voltage from the terminal, ensuring that the beam of ions feel a smooth potential gradient throughout their acceleration. These can be seen in Fig. 2.2, showing an interior view of Sta. Ana accelerator tank.

The corona system is the real workhorse maintaining the stability of the accelerator's charging systems. As small irregularities can be present between different links in the Pelletron chains or the belt itself, the corona system is responsible for precise second-by-second compensation. The mechanical part of the system consists of an arm with sharp metal points on the end which can be moved towards and away from the terminal to draw away current. On the back end, however, it also is controlled by a voltage feedback system to adjust the current draw, keeping the terminal at a constant voltage. The feedback for the system is provided either by an external reference through a generating voltmeter or from the beam itself as it passes through an analyzing dipole magnet at the exit of the accelerator. As the beam passes through the magnet it will pass through a set of slits. If V_T changes slightly, the energy of the beam will subsequently change, impacting the balance of the beam on the exit slits. This change in current will be fed back to the corona system, which in turn will compensate by changing the charge delivered to the terminal in order to maintain the slit balance. With this, V_T can be kept incredibly steady, within roughly 1 kV for every MV on the terminal. This is the biggest advantage of a Van de Graaff accelerator over a cyclotron. Intrinsically, this acceleration system makes nuclear experiments possible through monoenergetic beams.



Figure 2.2. View inside the Sta. Ana accelerator tank, showing the equipotential hoops as part of the column.

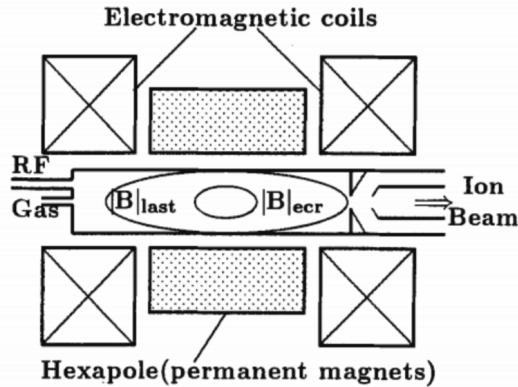


Figure 2.3. Depiction of the interior of an ECR ion source with magnetic components labeled [46]. This example shows a source operating from left to right, with the gas and RF generator inputs depicted as well as the exit for the ions. As discussed in the text, the combination of magnets confines the plasma in the center as it is pushed from left to right and out of the source.

2.3.2 Ion sources

A key item for the successful operation of any accelerator is the ion source, which produces the beams of charged particles used in experiments. Generating such beams is not a trivial task and, as such, many methods have been developed for achieving this. The two types relevant to this work, however, are the Electron-Cyclotron Resonance (ECR) ion source, used in the Sta. Ana accelerator at the NSL, and the radio frequency (RF) ion source, utilized in the JN accelerator at CASPAR. Both systems are housed within their respective accelerator terminals, which acts as a Faraday cage and shields the internal components from external fields. Additionally, both systems operate on the same basic principle, namely that a given gas is ionized through the use of electron collisions in the source, but the mode of electron excitation is different.

For the specifics of the ECR system, the 5U employs a Pantechnik Nanogen 14.5 GHz ECR source. This utilizes the cyclotron resonance of electrons (as the name obviously implies) to produce and maintain a plasma.

During operation, neutral gas (which includes the element to be accelerated as a component) is injected into a cavity surrounded by magnets. These magnets provide a superposition of axial and multipole fields in order to achieve a minimum at the center of the cavity increasing radially outwards with the goal of confining and stabilizing the generated ions. A cartoon depiction of such a system is presented in Fig. 2.3. In order to generate the ions from neutral gas, a current is

supplied to the chamber by a negative bias probe providing a constant supply of charge. From this, then, electrons are stripped and continually accelerated in cyclotron motion following the familiar cyclotron frequency relation:

$$\omega_{ecr} = \frac{e|B|_{ECR}}{m_e} = \omega_{rf} \quad (2.4)$$

where ω_{ecr} is the frequency for the cyclotron resonance of electrons in a given magnetic field B_{ECR} , e and m_e are the charge and mass of the electron, respectively, and ω_{rf} is the frequency supplied by a radio radio-frequency (RF) power supply. This last term is included to highlight that during operation this is supplied externally and controlled in the use of the source in order to produce the plasma. After ionization, the magnetic configuration confines the plasma radially, while the axial magnetic field (provided by the extractor) pushes it through the cavity towards the exit of the source. Upon exiting the chamber and the Faraday cage of the terminal, the positive ions are next to a the positively charged terminal at high voltage, thus accelerating the ions.

An ECR ion source has many advantages of over other common ion source designs. The primary advantage lies in its ability to deliver high-intensity beams (as much as hundreds of μA). For use in nuclear astrophysics experiments, such as the ones performed in this work, low cross sections are a problem to overcome, which is mitigated through this higher production. Additionally, due to the relatively simple restrictions on the inputs, these sources can provide a wide array of species, as they can produce beams of nearly anything that can be made into a gas, and exhibit long term stability, reliability, and longevity [46]. On the other hand, however, ECR ion sources also have some disadvantages compared to other common sources. One main drawback is that the ions are produced through the production of a plasma, meaning they are not directly controlled by any operational parameter [46]. This leads to complicated tuning and sensitive operation, experimentally. Additionally, such sources are only capable of generating positive beams and can therefore only be utilized in single stage accelerators. Finally, due to the size of these ion sources and their necessary components, they are often impractical. In order to employ one in the Sta. Ana accelerator, the accelerator design had to be vertical. If the accelerator had been horizontal (like the other accelerators at the NSL) the torque generated by the ECR ion source would have shattered the acceleration column. Vertically, however, the source is stable inside the accelerator. Unfortunately, the choice to build and use the accelerator vertically is not one that every lab can make.

The radio frequency (RF) ion source utilized at CASPAR operates on a very similar principle to the ECR ion source inside the Sta. Ana accelerator. A depiction of the RF ion source's components

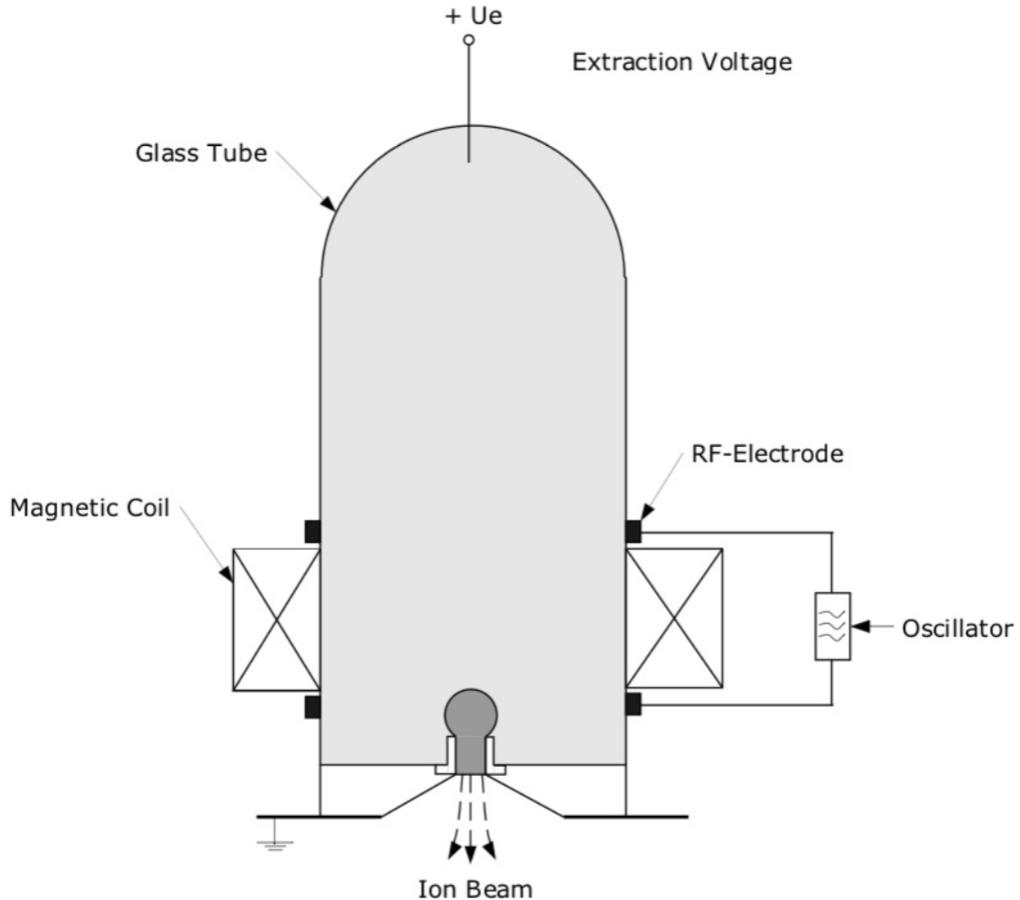


Figure 2.4. Depiction of the interior of a RF ion source with magnetic components labeled [41]. The coils on either side provide the field to ionize the gas, whereas the probe on top provides the field necessary to extract the ions from the chamber.

is given in Fig. 2.4. In this case, the source chamber is a glass tube filled with a monoatomic gas, at CASPAR the only options are ^1H and ^4He . Electromagnetic coils are placed on both sides of this tube and generate RF waves in the tube, between the rings. In response to these waves, the electrons oscillate through and collide with the source gas, ionizing it. At the rear of the glass tube, an electrode supplies a constant, positive voltage (with respect to the terminal), forcing any ionized atoms towards the exit of the source. Upon leaving the source, as with the ECR ion source, the ions are accelerated because of their repulsion to the positively charged terminal.

The RF ion source at CASPAR was also recently refurbished before installation, leaving it capable of beam production at even higher intensities than the ECR ion source inside the Sta.

Ana accelerator. It can produce beams up to $150 \mu A$, enabling similar high-intensity experiments. Another similarity to the ECR ion source is that it provides excellent operational stability and reliability. It does, however have significant drawbacks as well. The first and foremost of these is the degradation of the ion source canal. Operating the source causes a buildup on the exit canal of the source tube (higher intensity operation accelerates this process), necessitating significant maintenance. Additionally, the glass tube housing the plasma is quite sensitive. If power is supplied to the source when gas is not present, significant damage to the tube can occur. This is only mitigated with vigilant operation. The other major drawback is that the source is only capable of providing beams of two species (typically hydrogen and helium). In order to change the delivered species, a full tank opening and source alignment is required.

Altogether, though, these sources are ideal for this work. They provide the necessary ion beams at high intensity, enabling the experiments.

2.3.3 Beam transport

Upon exiting the accelerator, the beam envelope contains a distribution of different masses, charges, and energies because the ion sources are indiscriminate in their production methods. For example, an ion source might produce some ionized diatomic gas, instead of monoatomic, like producing ${}^1H + {}^1H$ due to insufficient breakup of the source gas. However, all particles created in the same ionization state will have the same energy, dictated by Equation 2.2. In order to filter such contaminants out of the beam, analyzing magnets are employed. These are typically 90° dipole magnets, also utilizing the cyclotron motion of ions in a magnetic field in order to select a given type of particle at a specific energy. These magnets are typically located at the exit of accelerators, and, due to their fixed size, have a fixed radius, R_{am} , through which a beam can pass. Therefore, for a given particle of mass m , energy E , and charge state q , the magnetic field, B_{am} , inside the analyzing magnet can be tuned to allow through only a specific particle/energy combination by

$$B_{am} = \frac{\sqrt{2mE}}{qR_{am}} = \frac{\sqrt{2m}}{qR_{am}}\sqrt{E} = k\sqrt{E} \quad (2.5)$$

, where k is a constant for a unique combination of particle and charge state. This separates the in-beam contaminants from the ions of experimental interest.

In order to deliver the beam of ions to the target chamber and experimental hall, a series of steering and focusing elements are placed along the beam pipes. As the beam is self-repulsive due to it all being the same charge, these are necessary to contain the beam and keep its intensity at

acceptable levels for experimental purposes. They also allow adjustment of the beam's position in order to deliver it to a specific experimental setup and allowing for simple transitions between different target locations. A schematic of the Sta. Ana and its beam transportation equipment is provided in Fig. 2.5.

Vacuum systems are important components of the beam production and experimental operation. All experimental components, like the ion source chamber, acceleration tubes, target chamber, and connecting beam pipes need to be free of residual gas. If not, the beam quality would suffer dramatically, collisions of the beam with the remaining atmosphere would cause significant intensity and energy losses, preventing transport and, ultimately, the measurement. In order to maintain an experimental vacuum level (typically $\sim 10^{-7}$ Torr), a series of metal pipes run from the ion source to the target chamber. These are connected by air-tight gaskets and have isolating valves employed with high-vacuum pumps (cryogenic or turbomolecular) along the line. These enable easier maintenance or changing of experimental conditions without destroying the vacuum created through the entire beam line. Without high-vacuum, there would be additional adverse effects on the target quality. Contaminants, typically hydrocarbons, present in residual gas can accumulate on the surface of the target due to the beam heating the surface. This is unwanted, as it not only provides an additional layer of energy loss for the beam before interacting with the target but also a significant contaminant in experimental results due to carbon's high cross-section.

Besides maintaining clean, high-vacuum near a target, another common method for reducing the carbon build-up on experimental surfaces is to utilize a so called cold trap in front of a target. A cold trap is a liquid nitrogen cooled copper pipe designed to be nearly the same size and shape as the beam pipe to prevent it from interfering with an experiment. An example of a cold trap like the one used in this work is shown in Fig. 2.6. The cold trap works much like a cryogenic pump. By cooling the copper pipe with liquid nitrogen, the residual hydrocarbons in the beam pipe and target chamber will condense on the surface of the pipe, trapping them before most can reach the target and cause carbon buildup. A common practice, which we also employed in this work, is to apply a negative voltage to the cold trap, forcing electrons scattered by the collision of the beam with the target back to the end of the chamber and providing an accurate reading of the amount of beam delivered to the target.

5U Beamlime Schematic

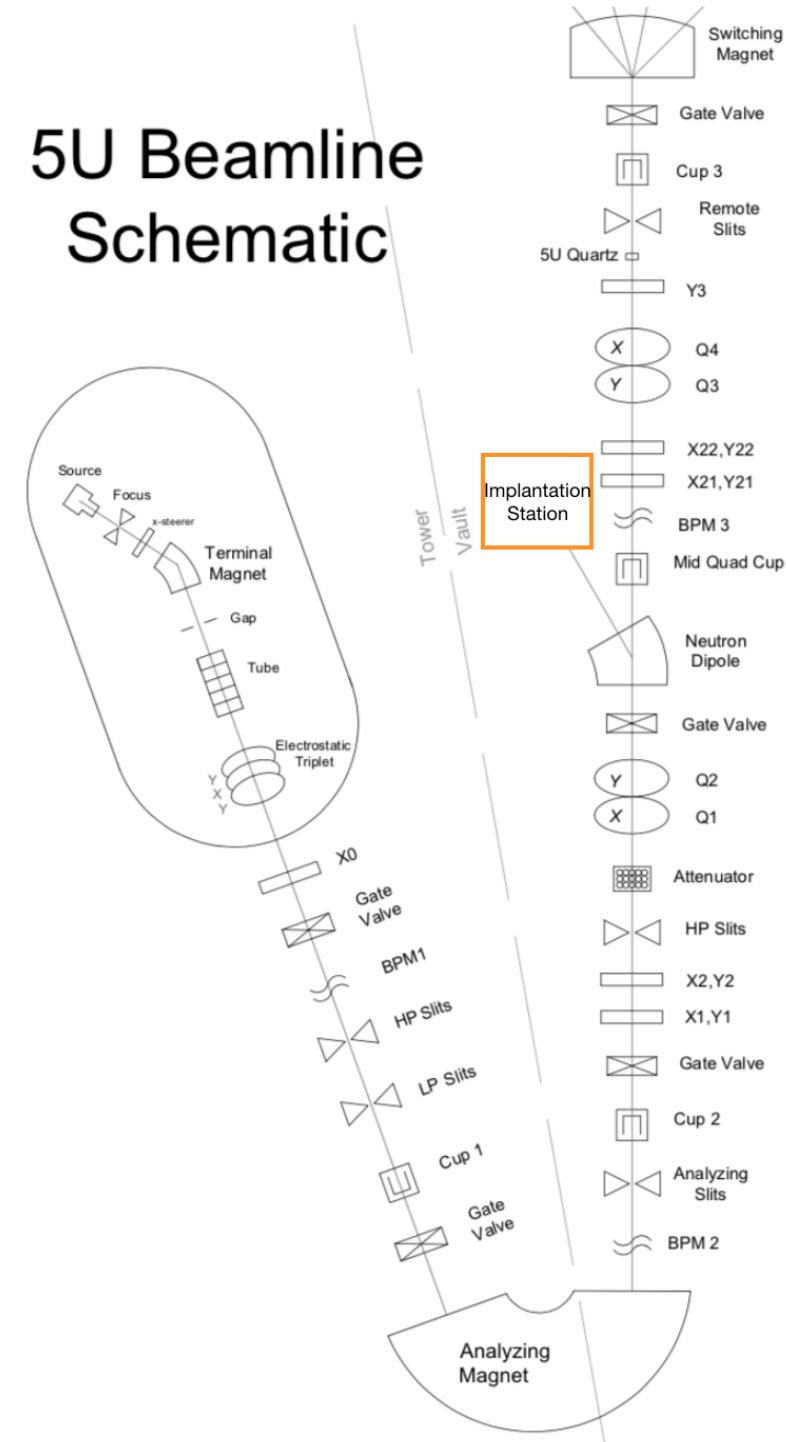


Figure 2.5. Schematic of the Sta. Ana accelerator at the NSL and its associated beam line. The target room containing experimental equipment is located after the switching magnet and is highlighted later (see Section 2.4.1). For the implanted target production (see Section 2.5.2) occurred at the implantation station off the neutron dipole, shown in this figure.

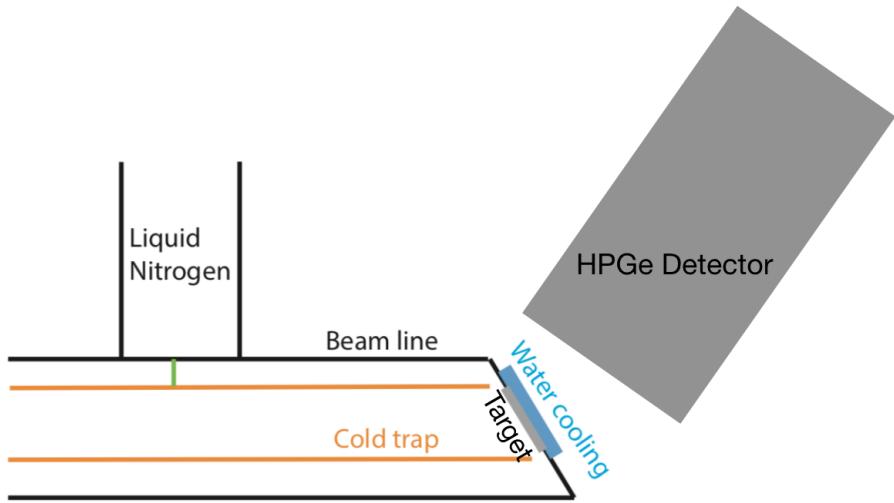


Figure 2.6. Depiction of the experimental setup used in this work, including the angled target holder, HPGe detector, and cold trap.

2.3.4 Radiation detection

Generally speaking, the first step in the design of an experiment is understanding the necessary detection system to match the reaction. This is because, like a child trying to reconstruct a broken Lego set, measurements are done by measuring the products of a reaction and trying to work backwards to understand its form and function. In nuclear physics experiments, there are three primary classes of radiation to consider as the reaction products: 1) energetic charged particles, 2) neutrons, and 3) electromagnetic radiation (photons). Of these, the first class can be further subdivided into heavy ions, like the proton, α particle, or other nuclei, versus light particles, namely electrons. For all classifications of particles, they are detected primarily through energy deposition into another medium, with the specific interaction dependent on the type of particle. These methods will be introduced below, but the reader is referred to [38] for a complete discussion of radiation interaction with matter and subsequent detection techniques.

For example, for charged particles in class 1, the dominant mechanism for energy deposition is through collisions with atomic electrons mediated by the electromagnetic force. As the particles pass through a medium, it generates an electromagnetic field in the surrounding space, which induces excitation in atomic electrons and even ionization if the energy is large enough. Each such interaction results in energy transfer from the incident particle to its neighborhood, slowing it to an eventual stop (usually). In many cases, the particle will stop inside the material, meaning that all

of the energy was deposited. On the other hand, it is possible for the particle to pass outside of the medium due to its trajectory, meaning that only a fraction of the total energy from the particle will be deposited in the material. The path a particle traces through a medium is dependent primarily on its own properties, like mass, as heavier particles are more resistant to changes in momentum from interactions with atomic electrons, while an electron's path in a medium is typically chaotic, with significant scattering from interactions with other electrons. The detection medium thus plays little part in the detection of charged radiation.

Neutrons, by contrast, interact significantly differently with matter because of their electrically neutral state. Because they cannot react via the electromagnetic interaction, they can only exchange energy with their surrounding through the strong nuclear interaction. This implies that they will only have meaningful interactions with atomic nuclei inside of the detection material. This presents a significant challenge, however, as atomic nuclei occupy only a minuscule volume of space. This makes the medium in which the neutrons are detected much more important. In order to increase the probability of detection, the medium would ideally have a higher density, increasing the number of potential interaction sites for the neutron. Unfortunately, this can introduce another problem, as higher density materials are often those with higher mass nuclei as constituents. While such material will have a greater chance to interact with the incident neutrons, these heavy nuclei will not acquire significant energy in collisions due to conservation of energy and momentum, similar to how the momentum of heavy charged particles is largely unaffected by interactions with atomic electrons. As such, a neutron traveling in a high-density material with heavy nuclei may bounce chaotically many times before exiting the medium without depositing enough energy to produce a strong enough signal to be detected by the system's electronics. Therefore, in the detection of neutrons, the medium is almost always chosen to be composed of light elements, which are more favorable for the detection of neutrons as each collision will produce a more significant exchange of energy with the medium. Materials with a high neutron cross section, like boron on ${}^3\text{He}$, are typically chosen as detection media. Due to the challenges of detecting high-energy (several MeV or greater) neutrons, a common technique for experiments involving neutrons is time-of-flight. In these cases, it is extremely unlikely to capture all of the neutron energy in a detection medium, so by measuring the time between neutron emission and interaction in a detector at a known distance the velocity, and kinetic energy, of the neutron is determined. This, however, necessitates a detailed understanding of the timing characteristics of a detector medium.

Finally, for quanta of electromagnetic radiation, or photons the interaction proceeds through

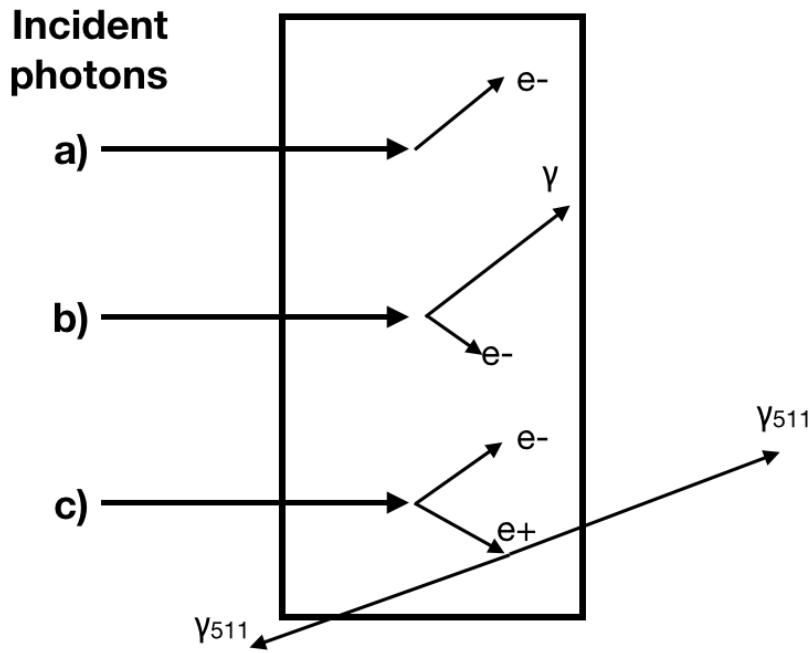


Figure 2.7. Depiction of the modes for photons to interact with matter, showing a) the photoelectric effect, b) Compton scattering, and c) electron-positron pair production.

yet different means. As an electrically neutral particle, photons do not interact with the medium in general and only at discrete points through the atomic electrons. As such, the interactions are much more probable than those of the neutron while being less than those of charged particles. Similar to charged particles, though, it is highly likely that photon interactions with matter result in a much higher chance to deposit all of the photon's energy in the medium. Clearly, by increasing the elemental number of the detection material's constituents, the probability of interaction with photons increases by increasing the number of possible interaction sites. As such, higher Z materials are preferred for the detection of photons.

On balance, there are three primary modes for photons to interact with matter: 1) the photoelectric effect, 2) Compton scattering, and 3) electron-positron pair production [38]. Fig. 2.7 illustrates these possible interactions and Fig. 2.8 depicts the ways in which these effects manifest in measured spectra.

In photoelectric absorption, the photon transfers its energy to an atomic electron, leaving it in an excited state or ionizing it completely, where the electron deposits its energy like any other

charged particle. If ejected, the electron will have energy E_e :

$$E_e = E_\gamma - E_b \quad (2.6)$$

where E_γ is the incident photon energy and E_b is the binding energy of the electron. In these cases, the electron is most likely ejected from the innermost atomic shell. For low-energy photons (below ~ 1 MeV), this is the dominant interaction process [38].

Compton scattering is a similar process in that it also ends up producing an energetic electron inside the detection medium. In this type of interaction, a photon scatters inelastically from an atomic electron, imparting some of its energy to the electron and changing the momentum of both. Unlike the photoelectric effect, not all of the photon's energy is absorbed by the electron. As such, any of the interactions can occur again in the medium, like subsequent scatterings, full photoelectric absorption, or pair production, or the scattered photon could leave the material entirely. For the former conditions, the full energy of the photon will still (likely) be absorbed in the material. However, in the event that a subsequent scattered photon leaves a detection medium, this will manifest in the total energy measured in this event being lower than the true incident photon energy, artificially reducing the true signal and increasing the background. This is known as the Compton edge, which is shown in Fig 2.8. This effect is also the dominant effect in cases where the photon has a middle range of energies (between ~ 1 - 6 MeV) [38].

The final mechanism by which photons interact with matter is through electron-positron pair production, often shortened to just pair production. This occurs when a photon traveling spontaneously transforms into a matter-antimatter pair, the electron and positron, respectively. This effect is induced by an interaction with a virtual photon from the electromagnetic field in the space the real photon is traveling. For this to occur, the photon's energy must be greater than the rest mass-energy of the pair, namely 1.022 MeV for electron-positron. After creation, the pair will share the total original energy (less their rest mass-energy) and momentum of the photon in their subsequent kinetic energies and momenta. From this point, the electron will continue to scatter and deposit energy in the normal manner for charged particles, while the positron will annihilate with one of the atomic electrons in the surrounding material, creating two diametrically opposed gamma-rays at 511 keV. With these, there are three potential outcomes: 1) the two new photons will interact with the surrounding medium through the processes already detailed, depositing their energy, and resulting in the total deposition of the initial photon's energy in the detection medium; 2) one of the new 511 keV photons will leave the detector without interacting and result in a deposition of the total

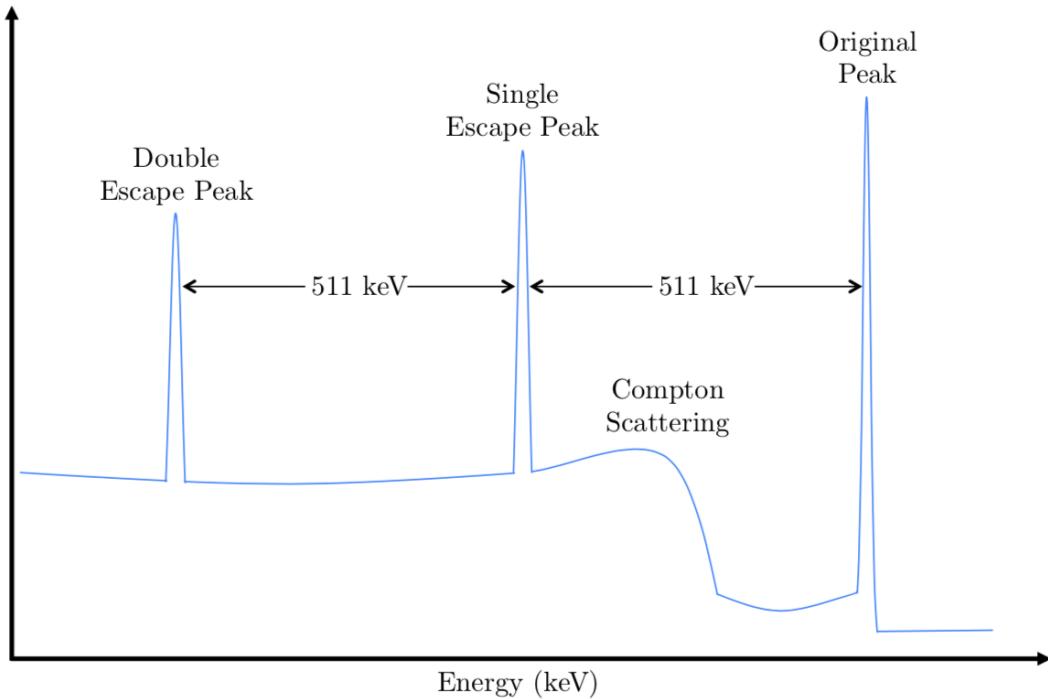


Figure 2.8. Cartoon depiction of an example experimental spectrum for photon interactions [32]. Artifacts from the ways in which photons interact with matter are highlighted in the spectrum and are common in experimental gamma spectra.

initial energy less 511 keV; 3) both of the subsequent 511 keV photons leave the detector, resulting in an energy deposition of the total initial photon's energy minus 1.022 MeV. The two situations where a subsequent photon leaves the material are more common with higher initial photon energies, manifesting in their own peaks in a measured spectrum. These are called escape peaks, shown in Fig. 2.8, with the single escape peak being when one photon leaves the medium and the double arising when both 511 keV photons escape the system. Whereas the full deposition of the photons energy is also shown in that spectrum and known as the full-energy peak.

This work is concerned ultimately with measuring photons as reaction products and employed high-purity germanium (HPGe) semiconducting detectors for that purpose. These are characterized by having a relatively small band gap between the electrons within the valence and conducting bands. When photons enter the Ge crystal, they excite electrons across the band gap from the valence to the conducting band, leaving behind vacancies, known as holes, in the valence band. These excited electron-hole pairs feel an electric field supplied by the detector's bias voltage, causing them to

drift to the charge collection electrodes at the end of the crystal, resulting in an electric current pulse. Since HPGe has a small band gap, around 0.7 eV (depending on impurities) [38], it requires constant cooling of the medium by liquid nitrogen in order to reduce the number of electron-hole pairs created by thermal excitations in the valence band. However, by cooling these detectors, HPGe becomes extremely useful because of the large number of charge carries produced by the small band gap in an energy deposition event, giving superior energy resolution. It is for this reason that HPGe detectors are ubiquitous in nuclear physics laboratories around the world - even those shown in movies. Comparatively though, HPGe detectors are more expensive and (typically) less efficient than NaI detectors, which are the other primary type of photon detector, so they are not without downsides to use as well.

Thus, in every experiment it is important to anticipate reaction outcomes when planning what type of radiation detection system to employ. As stated, this work was concerned with the measurement of high-energy photons from a nuclear reaction, so we chose to use HPGe detectors for their superior resolution.

2.4 Cross-section measurements

This set of data was taken over the course of five separate experiments. The first occurred at the University of Notre Dame's Nuclear Science Laboratory (NSL) in January of 2018 and covered the proton energy range of $E_p = 800 - 1200$ keV. The experiment was then continued at the Compact Accelerator System for Performing Nuclear Astrophysics (CASPAR) facility at the Sanford Underground Research Facility located in Lead, South Dakota in three increments: February 2018, May 2018, and August / September 2018. These measurements covered the energy range from $E_p = 270 - 1100$ keV, in order to measure the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction cross-section to compare the performance of the CASPAR facility to an above-ground laboratory.

2.4.1 Experiment at Notre Dame

The goal of this experiment was to measure the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction cross-section in a low energy range of the Sta. Ana accelerator, proton energy range of $E_p = 800 - 1200$ keV, to provide a self consistent data set with which to compare the results of a measurement at CASPAR. Owing to the difficult experimental environment at CASPAR, this also allowed a dry-run of the procedures and an opportunity to work out kinks with the equipment before moving underground. The Notre Dame Nuclear Science Laboratory's (NSL) layout is shown in Fig. 2.9, with this phase of the

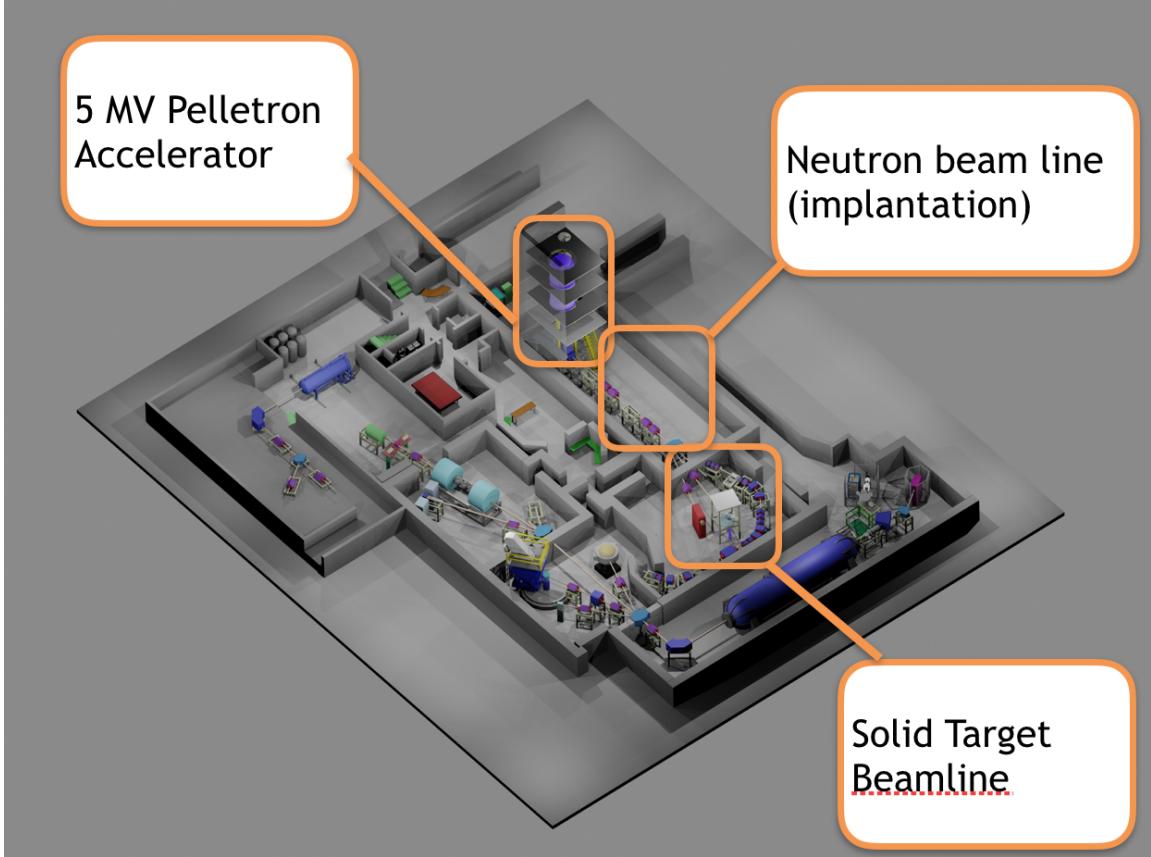


Figure 2.9. The Notre Dame Nuclear Science Laboratory, with experimental areas important for this work identified.

measurement occurring on the Solid Target beam line, a schematic of which is shown in Fig. 2.10. A diagram of the specific setup used in this experimental phase is depicted in Fig. 2.6. In order to reduce confounding effects, it was also employed identically at CASPAR.

2.4.2 The CASPAR facility

In an attempt to extend the data range down to the lowest possible energies and provide a proof-of-principle for the facility, a second phase of the experiment was performed at the CASPAR facility. CASPAR is the Compact Accelerator System for Performing Astrophysical Research, located on the 4850 ft level of the Sanford Underground Research Facility, and is the first underground accelerator facility in the United States. This facility was originally the Homestake Gold Mine in Western South Dakota, just outside (and underneath) the town of Lead. It is also the location of the famous Ray Davis and John Bahcall underground neutrino experiment in which the solar neutrino problem

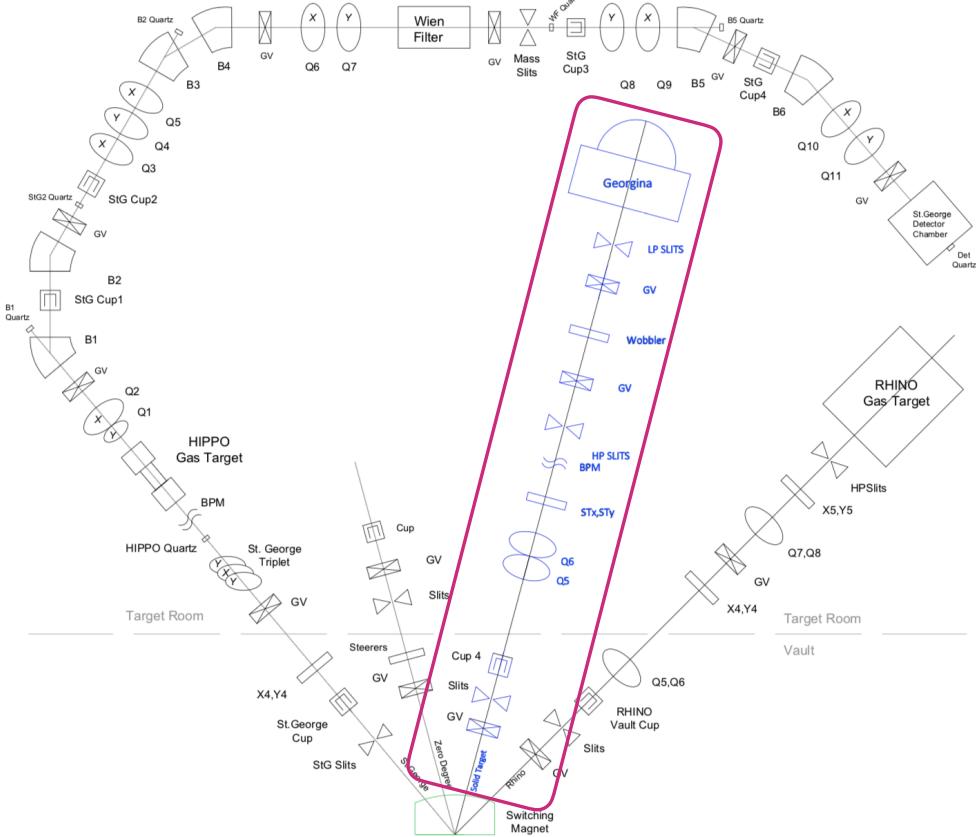


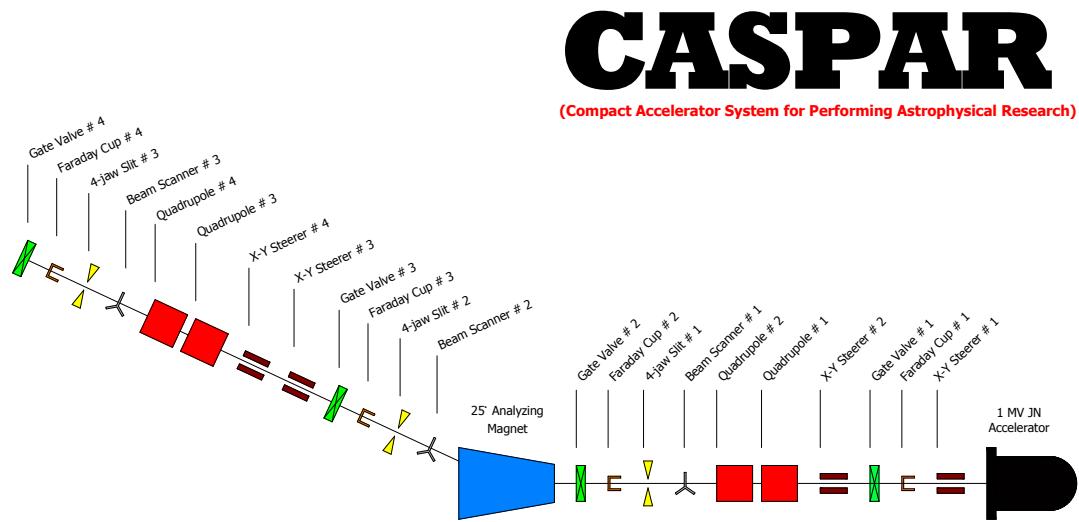
Figure 2.10. Schematic of the Sta. Ana accelerator's target room, showing all experimental stations. The solid target beam line used in this experiment is emphasized for clarity. For these experiments, the target chamber and detectors were placed at the location named *Georgina*.



Figure 2.11. Pictures of the CASPAR facility.

was identified [12]. The accelerator located at the facility is the recently refurbished JN Van de Graaff, which had been employed at the NSL since its arrival in the early 2000's. Fig. 2.11 shows some pictures of the facility while Fig. 2.12 provides a detailed schematic of the equipment available at the CASPAR facility.

Some of the problems with performing low-energy nuclear astrophysics measurements were detailed in Section 1.2. The most significant of these is the diminishing cross section at astrophysically relevant energies due to Coulomb repulsion. There are a number of techniques for mitigating this problem: 1) extrapolating from higher energy with techniques like *R*-matrix (see Section 1.4), 2) increasing the reaction yield with high-intensity accelerators and targets, 3) performing long term measurements to enable measurements of ridiculously low cross section reactions, and 4) drastically reduce cosmic background to increase the signal to background ratio and prevent these events from hiding a signal of interest. Of these, CASPAR is equipped to employ the solutions of 2, 3, and 4. In Section 2.3.2, the RF ion source utilized inside the JN accelerator was described, but the useful



Updated on: 11-July-2018

Figure 2.12. Schematic of the equipment employed at the CASPAR facility. In this case, the figure is read right-to-left, with beam generated in the JN accelerator on the right of the figure.

feature is that it can deliver proton and α beams up to $150 \mu\text{A}$, increasing the experimental yield dramatically. Additionally, without the constraints and competition of multiple running groups, which is the reality at larger labs like the NSL or the National Superconducting Cyclotron Laboratory at Michigan State University, the facility is able to measure a reaction as long as necessary (five weeks in this case). Finally, with nearly a mile of rock cover above the facility provided by the mountains, the facility has excellent passive shielding. The background reduction compared to a surface laboratory like the NSL is shown in Fig. 2.13. Strictly speaking, due to the high content of the uranium and thorium decay chains in the surrounding rock, the radiogenic background (that at energies below ~ 2.6 MeV) is actually higher at the CASPAR facility than the surface. This, however, can be (and is) addressed by adding additional lead shielding around the detector crystal. Taken together, these features advance CASPAR beyond the capabilities of other laboratories performing nuclear astrophysics experiments and make it particularly useful. It's for these reasons that it was chosen to perform the low-energy phase of this work.

Additionally, the JN accelerator at CASPAR expands on the potential measurement range compared to the LUNA II facility, the other preeminent underground nuclear astrophysics laboratory in the world [28]. The two have similar background reduction due to their environments and equal capacity to perform extensive measurements. However, until the LUNA-MV facility is completed [22, 26], CASPAR's expanded energy regime makes it the only facility with the capability to overlap with those measurements performed at surface laboratories, making it the ideal facility for low-energy nuclear astrophysics.

The measurements at CASPAR were undertaken over the course of five weeks in February 2018, May 2018, and August / September 2018. These covered the proton energy range from $E_p = 270 - 1060$ keV, overlapping with the measurements at the NSL. As stated in Section 2.4.1, the equipment like detector, electronics, and overall setup used in this campaign was the same as what was used at the NSL. The only significant different was the different targets used in this experiment. These were sputtered ZrN targets of varying thickness (50 - 100 nm sputtered layer thickness, 11 - 22 keV thick for protons at 278 keV) produced specifically for this experiment in Bochum, Germany. The targets were produced by sputtering a layer of Zr onto a W backing in a nitrogen atmosphere. Given the high intensity of the beam delivered at CASPAR, these were constantly monitored for degradation and carbon buildup to reduce target effects on the measurement.

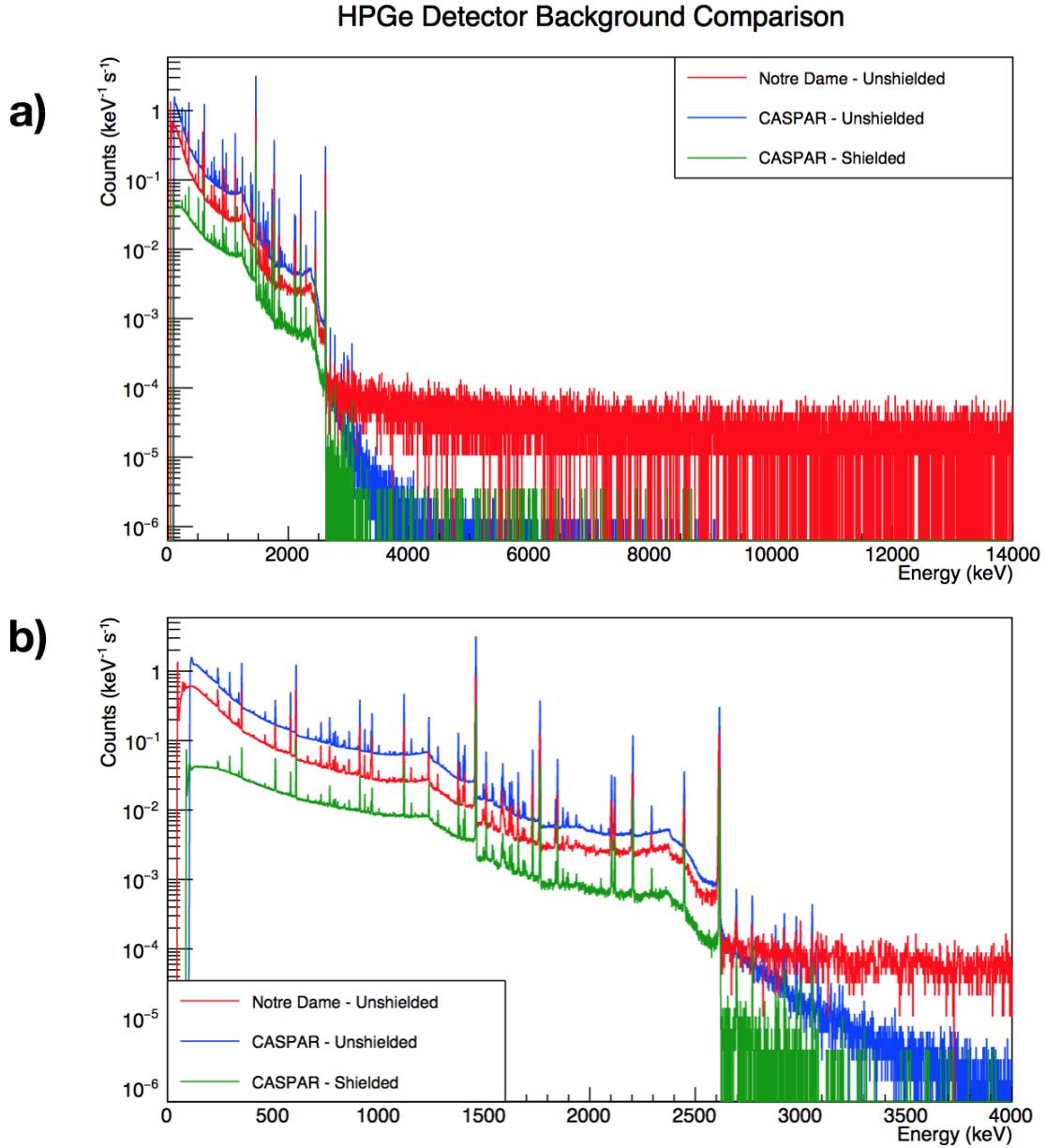


Figure 2.13. Background γ spectrum taken at the NSL, CASPAR, and CASPAR utilizing passive lead shielding, with a) showing the full spectrum and b) showing a subset of the spectrum at low energies to highlight the radiogenic background, which is higher at CASPAR without lead shielding. Clearly, the facility provides a prodigious reduction of cosmogenic background, making the facility advantageous for low-energy nuclear astrophysics experiments.

2.5 Lifetime measurement

The goal of this measurement was to determine the lifetime of the 6.79 MeV state in ^{15}O with a variety of target materials and nitrogen contents in order to identify any systematic effects which were unidentified in previous measurements. Targets for this measurement were produced at the NSL in the summer of 2019. The process of fabricating these targets is presented in Section 2.5.2. When measuring the lifetime, the ^{15}O nuclei were generated using the $^{14}\text{N}(p,\gamma)^{15}\text{O}$ reaction at energies ranging from 1020 keV to 1540 keV. At these energies, the excited states in ^{15}O at 5.24 MeV and 6.18 MeV, with known lifetime, are also populated. The details of how these measurements were made are given in Section 2.5.3 By measuring all of these, these additional states can be used as a check to ensure the validity of the measurement technique used herein: the Doppler Shift Attenuation Method.

2.5.1 The Doppler-Shift Attenuation Method for lifetime measurements

Nuclear lifetimes are important observables to measure because they provide a link to understanding the strong nuclear force. They are necessary to determine the reduced transition probabilities which are one of the most important probes of nuclear structure and the forces governing the behavior of nuclei. The Doppler Shift Attenuation Method (DSAM) is a well characterized method for determining the lifetimes of excited nuclear states that decay via gamma emission in the range of $10^{-11} - 10^{-15}$ s [21]. The ranges where DSAM is applicable are compared with other common methods for determining nuclear lifetimes in Fig. 2.14.

The basic idea of DSAM is to produce an excited nucleus inside of a dense target in which the reaction product will decay. The de-excitation of the nucleus then takes place either while the nucleus is slowing down or after it stops, depending on the lifetime of the specific state and the stopping within the target. Then, as is true with the classical Doppler shift, the energy of the emitted γ -rays are shifted depending on their emission angle relative to the motion of their source, the decaying nucleus. Thus, γ 's emitted at different velocities during the slow-down and detected at given angles will have a spread of energies. Specifically, for a given nucleus decaying at a time t with a speed of $v(t)$, the Doppler shifted energy of the gamma ray, E_γ , is

$$E_\gamma = E_0 \left(1 + \frac{v(t)}{c} \cos(\theta) \right), \quad (2.7)$$

where E_0 is the energy of the decay for a nucleus at rest, c is the speed of light, and θ is the lab angle

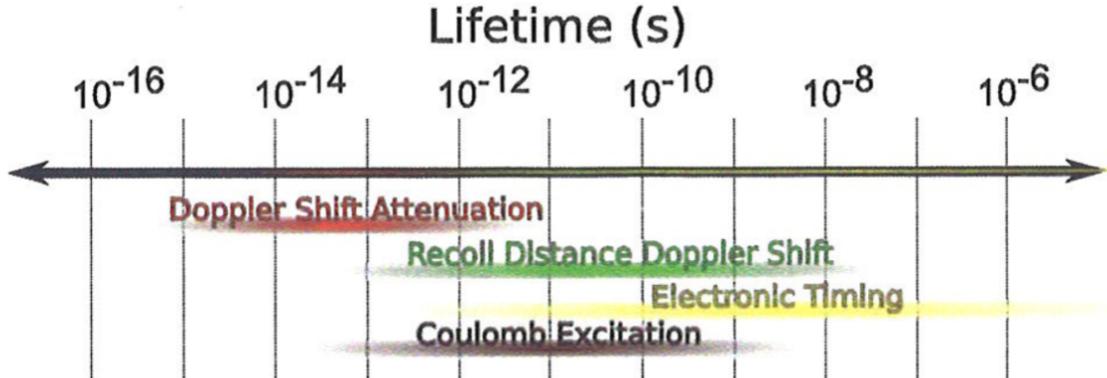


Figure 2.14. Ranges over which the different lifetime measuring techniques are valid (Obtained via personal communication with M.K. Smith in 2017).

between the momentum vectors of the decaying nucleus and gamma ray, respectively. A depiction of this scenario is given in Fig. 2.15.

As the nuclear decay process is statistical in nature, the measured spectrum of gamma energies for a particular transition is a continuous distribution of energies from the unshifted, rest energy peak to the maximally Doppler shifted peak. As Equation 2.7 was specific to a single transition, it is necessary to consider the entire sample of decays measured. By defining the average velocity of all nuclei at the time of decay, \bar{v} , the initial velocity of all recoiling nuclei, v_0 , and their ratio, $F(\tau)$, called the attenuation factor, it is possible to relate the behavior of the entire distribution to the lifetime of the state. This is

$$E_\gamma = E_0 (1 + F(\tau) \beta_0 \cos(\theta)), \quad (2.8)$$

where $\beta_0 = v_0/c$ is the relativistic velocity factor. $F(\tau)$ therefore provides an analytical relation between the nuclear lifetime and the measured Doppler shift, given by Blaugrund in 1966 [21] as

$$F(\tau) \beta_0 = \frac{1}{\tau} \int_0^\infty \beta(t) \exp\left(\frac{t}{\tau}\right) dt, \quad (2.9)$$

where $\beta(t) = v(t)/c$ is the time dependent relativistic velocity factor. The devil, however, is in the details of calculating $\beta(t)$, since it is reliant on stopping powers of the nuclei in the target material, implying an accurate knowledge of the target composition and population pattern of the nucleus. In some cases the feeding is not well known and for nearly all cases, there is an estimated uncertainty

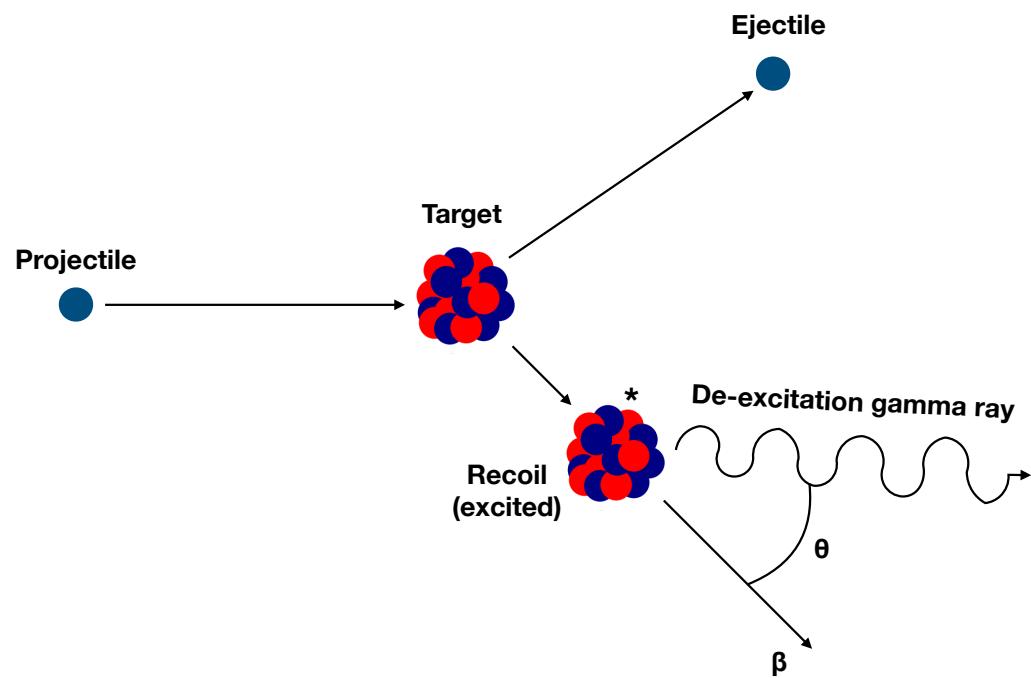


Figure 2.15. A generic nuclear reaction showing the creation of an excited nucleus and subsequent decay, causing a Doppler shift in the measured energy of the γ ray.

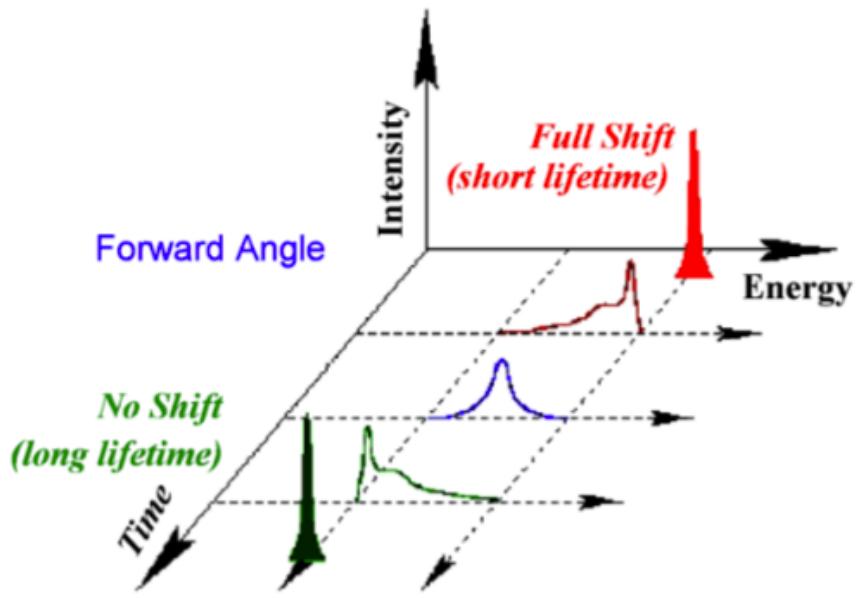


Figure 2.16. The effect of the nuclear lifetime on the measured gamma spectrum for a given experimental scenario [55]. In forward angles, the γ 's energy is shifted to higher values, while the measured energy of the γ is lower at backward angles, with the maximum shift in either direction coming when the γ is emitted in a direction parallel to the motion of the decaying nucleus. A γ emitted at 90° is unshifted and unbroadened.

in the assessment of stopping powers of materials to be at least 10%.

It is important to note that the Doppler shift can either increase or decrease the energy of the measured γ ray, depending on whether the emitted gamma ray is at forward or backward angles relative to the nucleus' motion. Fig. 2.16 shows the Doppler effect on a measured γ spectrum, shifting the energy according to the measured angle. It is evident that the lifetime of a particular excited state has a dramatic effect on the location and shape of this distribution, for the longer a specific lifetime is, the broader its measured distribution will be (and vice versa) while the shift in the centroid of the peak will be lower than that of short lifetimes where the nucleus has a relatively higher velocity upon decay.

Therefore, by performing a lineshape analysis on the distribution of the measured γ 's and a mapping of the distribution's shift with angle one can determine the nuclear lifetime. The procedure for the Doppler shift determination is particularly straightforward. By plotting the resulting centroids measured for the gamma distribution vs the $\cos(\theta)$, one should obtain a straight line, the slope of which is the product $E_0\beta_0F(\tau)$. So, for a given experimental approach with the same reaction and

targets, this slope and therefore attenuation factor should match. This was not the case for the Bertone *et al.* and Schürmann *et al.* works [18, 57], where the authors used the same reaction on ostensibly the same targets and reached incongruous $F(\tau)$ values.

2.5.2 Target production

To take advantage of the Doppler Shift Attenuation Method, therefore, the active layer of the target must be dense to slow down the recoiling nucleus in an appreciable time and must have a lot of target nuclei since the 6.79 MeV state in ^{15}O is populated weakly. As such, implanted nitrogen targets were utilized in this work. As the previous measurements utilized enough nitrogen to saturate tantalum to a composition of Ta_2N_3 [18, 57], it was concluded that Ta backings should be utilized in this work too, albeit with a lower nitrogen content. Additionally, to examine any systematic effects that the backing material could have on the ultimate extraction of a lifetime, backings of tungsten and molybdenum would also be implanted with nitrogen for comparison. With each backing material, targets were produced with three different levels of nitrogen to isolate the effect of nitrogen saturation on the observed Doppler Shift.

The properties of implanted Ta-N targets were studied in Seithe et al. [60]. In that work, various nuclides, including ^{14}N , were implanted into Ta and Au with energy up to 200 keV and subsequently studied with (p, γ) reactions. In this work, target saturation occurred with nitrogen doses creating the Ta_2N_3 compound. The authors identified this to be a reliable method for producing isotopically enriched targets capable of withstanding high intensity beam loads. Therefore, the procedure for fabricating implanted nitrogen targets used herein was based off their work.

Before the implantation, the target backings were cut and cleaned, both chemically and with oxygen plasma, according to standard procedures at the NSL. The targets were then produced in July 2019 with the Sta. Ana accelerator at the NSL by accelerating a ^{14}N beam into the various backings at the implantation station we created off of the neutron dipole, shown in Fig. 2.5. For operational stability, the ^{14}N was implanted at 350 keV, in contrast to the 200 keV by [60]. As such, the targets produced in this work had a comparatively broader distribution of nitrogen, creating a larger active volume. Nominally, for the different levels of nitrogen saturation, the aim was to implant doses equivalent to an atomic percentage of 10%, 20%, and 30% nitrogen in the compound. These goals were chosen as they provided a significantly different nitrogen content from the targets used in Refs. [18, 57] but were expected to also highlight any systematic differences in nitrogen content. Going to higher dose implantations, like the 60% necessary for saturation, was deemed a

diminishing return compared to the longer time for production.

The actual amount of nitrogen implanted in the targets was assessed by scanning the 1057 keV resonance in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction. This was chosen as it gives off three strong γ -rays in its decay: 3043, 5241, and 8284 keV, respectively. The area under the resonance scan is directly related to the number of active atoms in the target:

$$A_\gamma = n \times \frac{\lambda_r^2}{2} \times \omega\gamma \quad (2.10)$$

where A_γ is the area under the yield curve, n is the number of target atoms, λ_r is the De Broglie wavelength of the resonance, and $\omega\gamma$ is the resonance strength. The process is made simpler if a target with known characteristics is available for comparison. When relating the areas of the yield curves for the two targets, the resonance features cancel, leaving:

$$\frac{A_\gamma^1}{A_\gamma^2} = \frac{n_1}{n_2}. \quad (2.11)$$

Thus, by comparing the area of the yield curves with a known target, the unknown number of target atoms is easily determined. In this work, the reference target was a sputtered TiN target used by Li *et al.* [42] and produced in Bochum, Germany, with $3.34 \times 10^{17} \text{ }^{14}\text{N}$ atoms/cm².

A typical spectrum from the implanted targets produced at the NSL is shown in Fig. 2.17, specifically from the low-dose Ta target. The yield curves for all the implanted targets produced in this work are presented in Fig. 2.18, while a comparison of the high-dose yield curves with that of the reference target is provided in Fig. 2.19. As the reference target's nitrogen layer is on the target's surface, the resonance occurs at the nominal proton energy (1057 keV) while the resonance is seen at higher proton energies for the implanted targets due to the energy loss in the material before reaching the layer containing the high concentrations of nitrogen. From these, it is evident that the implanted targets produced in this work have significantly higher concentrations of nitrogen than the sputtered TiN target used for reference. This is ideal for the lifetime measurements due to the weak population of the 6.79 MeV state of interest. A complete summary of the implanted targets fabricated in this way and their actual nitrogen contents are shown in Table 2.1.

2.5.3 Measurement at Notre Dame

In August 2019, the Doppler Shift Attenuation Method was used to measure the lifetime of the 6.79 MeV state in ^{15}O with some of the implanted targets described in the previous section. The

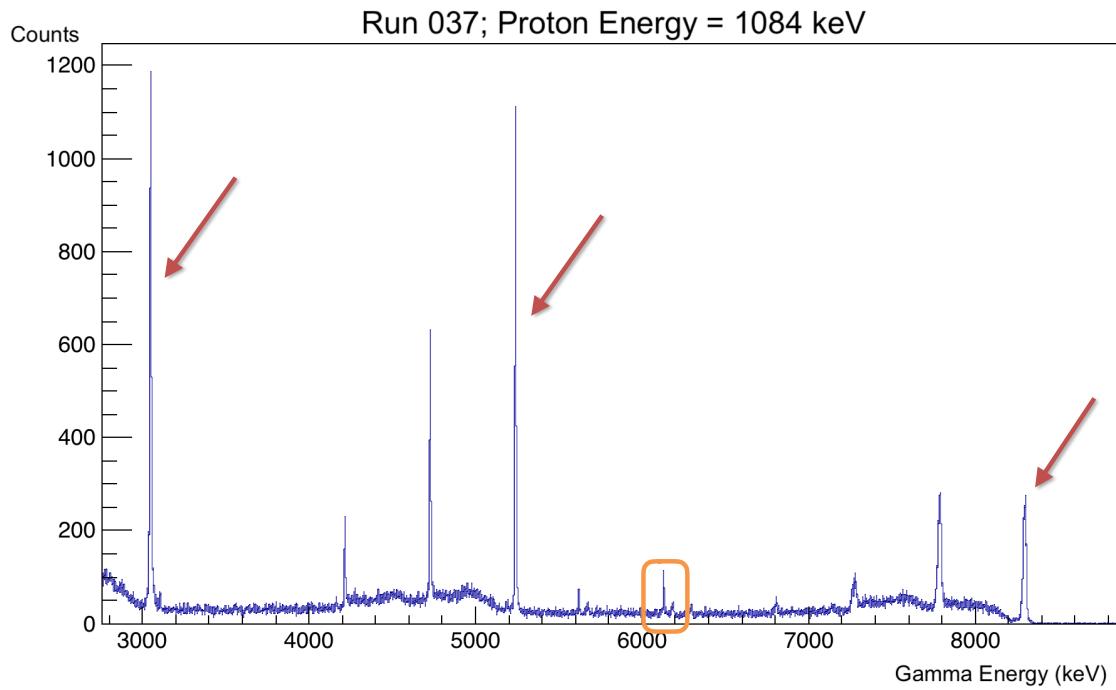


Figure 2.17. A spectrum from the low-dose tantalum backed target at an on-resonance proton energy with features typical of the implanted targets produced at the NSL. The three prominent decay peaks at 3043, 5241, and 8284 keV, respectively, are shown with the arrows while the contaminant peak from $^{19}\text{F}(p, \alpha\gamma)^{16}\text{O}$ at 6190 keV is in the orange box.

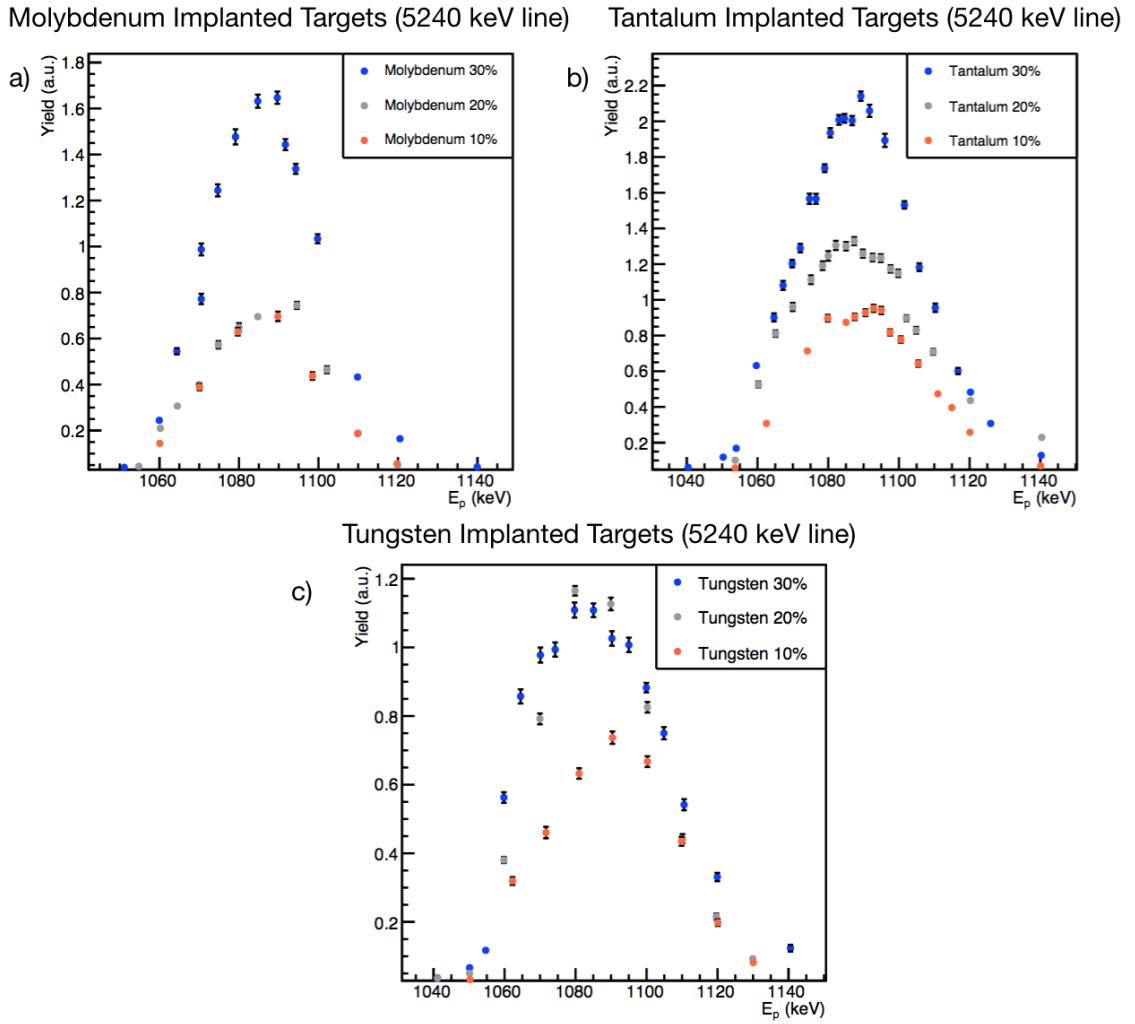


Figure 2.18. Yield curves (showing the 5240 keV line) for the different implanted targets produced as a part of this work with each backing and nitrogen dose: a) Mo targets, b) Ta targets, and c) W targets. The legend of each only gives the nominal goal percentage of nitrogen in the targets, not the actual implanted does (see text for details).

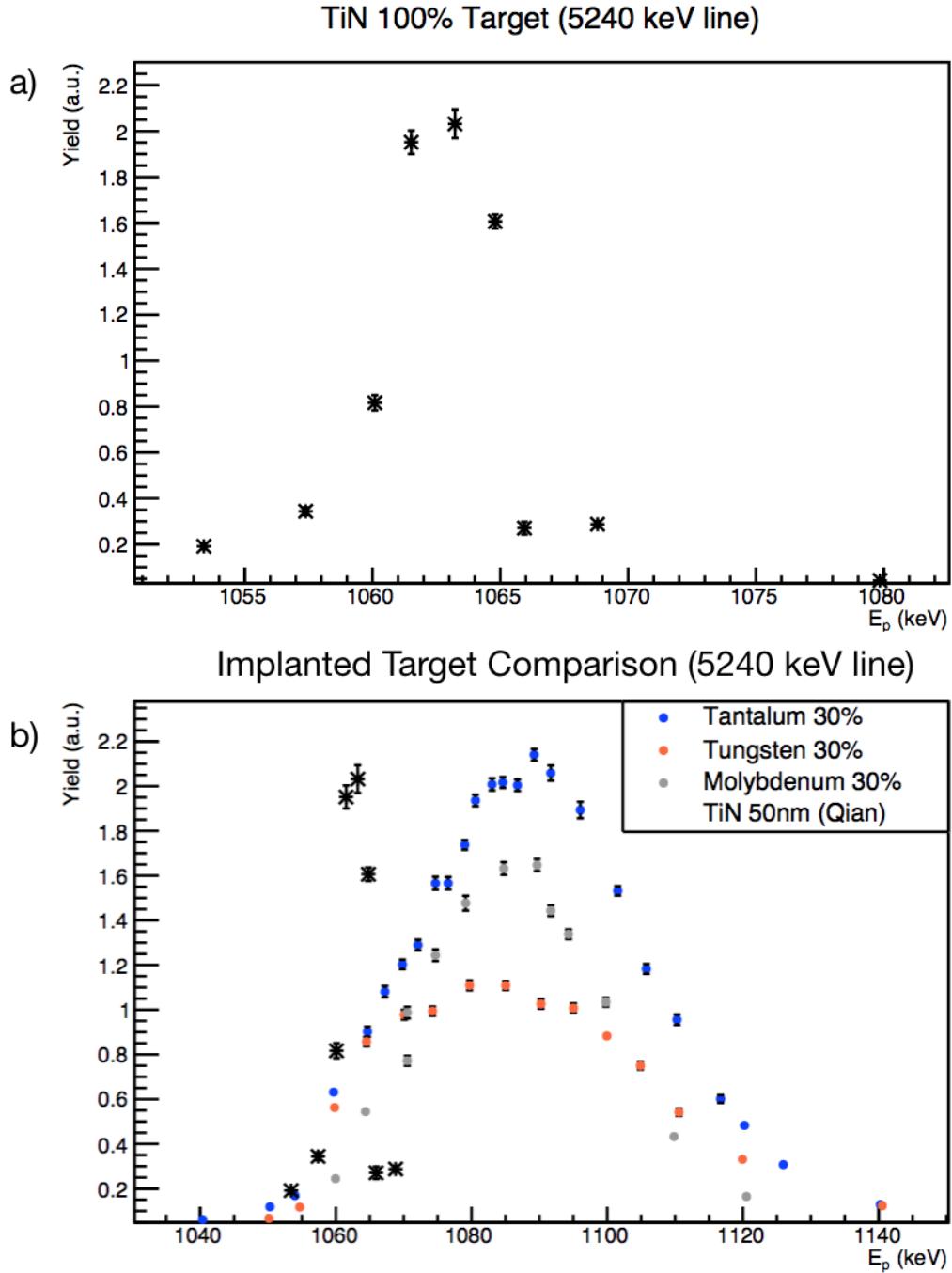


Figure 2.19. Yield curves of the 5140 keV line for the target comparisons: a) yield curve for the TiN target alone and b) yield curve for the high dose implanted targets plotted with that of the TiN target. Percentages quoted in the legend are for identification and differentiation only (see text for details).

Target	Nitrogen atoms ($10^{17}/\text{cm}^2$)	Atomic percentage
Mo (low)	5.46 ± 0.11	11 ± 2
Mo (mid)	6.08 ± 0.12	14 ± 2
Mo (high)	13.10 ± 0.66	26 ± 5
Ta (low)	9.63 ± 0.29	17 ± 3
Ta (mid)	14.37 ± 0.57	26 ± 4
Ta (high)	21.29 ± 1.28	36 ± 6
W (low)	7.33 ± 0.15	13 ± 2
W (mid)	11.62 ± 0.35	19 ± 3
W (high)	13.38 ± 0.54	22 ± 4

TABLE 2.1

ALL TARGETS PRODUCED VIA THE IMPLANTATION METHOD FOR THIS WORK AND THEIR RESPECTIVE NITROGEN CONTENT.

^{15}O nuclei were created with the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction. The astrophysical S factor for the R/DC $\rightarrow 6.79$ MeV transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction (as measured by Schröder *et al.* [56]) is shown in Fig. 2.20. The only resonance which populates the state of interest is that at $E_{c.m.} = 259$ keV, which is unfortunately below the minimum voltage threshold for the Sta. Ana accelerator. This indicates that, for the purposes of this experiment, the 6.79 MeV state must be populated through direct capture which is a significantly weaker process.

As such, multiple energies were chosen at which to measure the reaction, $E_p = 1020$ keV and $E_p = 1570$ keV. At $E_p = 1570$ keV, there is a minimum in the background from the strong 6.12 MeV line coming from the $^{19}\text{F}(p, \alpha\gamma)^{16}\text{O}$ reaction. The reaction at 1570 keV also produced ^{15}O nuclei in the 5.24 MeV and 6.18 MeV excited states, which have known lifetimes of 8.2 ± 1.0 fs and < 2.5 fs, respectively [4]. Both of these states are also within the range of DSAM to measure, allowing cross-validation of results for the measurement of the 6.79 MeV state. An example spectrum from the measurement showing the presence of all described states is given in Fig. 2.21.

As the energy at which the peak is observed for each detector angle is of critical importance for determining the overall Doppler shift (and therefore measured lifetime), the relative position of the peaks within each spectrum is also highly salient. To prevent any artificial peak shifting arising

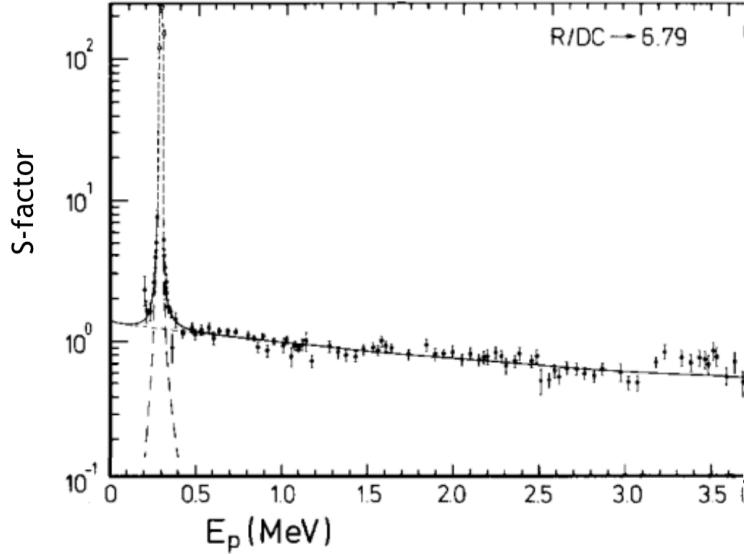


Figure 2.20. Astrophysical S factor for the $R/DC \rightarrow 6.79$ MeV transition from [56]. With the exception of the resonance at $E_{c.m.} = 259$ keV, the S factor is remarkably flat, even decreasing at high energies, indicating that the state is weakly populated in all energy ranges accessible with the Sta. Ana accelerator used herein.

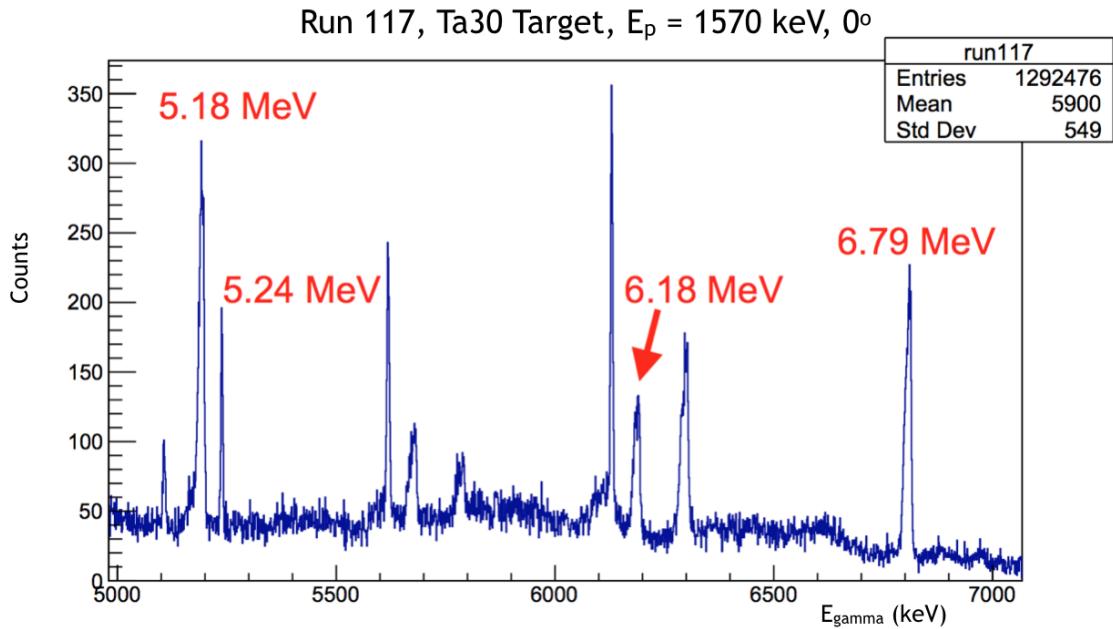


Figure 2.21. Example spectrum from the measurement of the 6.79 MeV state lifetime. This was produced with the high-dose Ta implanted target, proton energy $E_p = 1570$ keV, and the detector at 0° . The excited states of ^{15}O are indicated, where the 5.24, 6.18, and 6.79 MeV states are all available to measurement with DSAM.

from gain shifts in the electronics to enter into the final product, the data were saved and the data acquisition restarted hourly at each angle. This allowed us to monitor for gain shifting throughout the experiment prevent it from obfuscating results. Ultimately, no evidence of gain shifting was found throughout the experiment.

CHAPTER 3

CROSS SECTION DATA REDUCTION AND ANALYSIS

3.1 Introduction

In aggregate, the data taken at both the CASPAR and NSL experiments consists of γ -ray energy data from the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction, observed with a single, 130% HPGe detector placed at 55° relative to the beam direction. These data were collected for reactions over the combined proton energy range of 270 - 1200 keV. The primary interest of these experiments was monitoring the R/DC \rightarrow GS transition and the R/DC \rightarrow 6.79 MeV + 6.79 MeV \rightarrow GS transition sequence. As such, the energies of the concerned photons ranged in energy from \sim 600 keV up to \sim 8500 keV. This chapter details the processes by which this data is gathered and turned into an experimental cross section.

3.2 Angular corrections

The angular distribution of a cross section, W , can be described by

$$W_{\text{exp}} = a_0 \left(1 + \sum_{i=1}^n a_i Q_i P_i(\cos(\theta)) \right) \quad (3.1)$$

where a_i are the angular distribution coefficients, Q_i are correction factors due to the finite size of a given detector, and the $P_i(\cos(\theta))$ are the Legendre polynomials of order i . For the conditions of this work, odd numbered terms as well as those of order higher than 2 give negligible contribution to the angular distribution. Therefore, the resulting angular distribution of this reaction is of the form

$$W_{\text{exp}} = a_0 + a_2 Q_2 P_2(\cos(\theta)). \quad (3.2)$$

Experimentally, to address any effects arising from an angular distribution of this form, the detector was placed at 55° . This is the zero of the 2nd order Legendre polynomial, thereby minimizing

any effects on the cross section arising from the detector's angle. Simultaneously, this means that no correction of the data is necessary.

3.3 Energy calibration

It is important to establish the true relationship between γ -rays of different energies incident in a detector and the signal they produce during data acquisition. This connection is determined by calibrating the system with γ -rays of well-known energy from room background, given radioactive sources, like ^{137}Cs or ^{60}Co , and well-studied nuclear reactions, like $^{27}\text{Al}(p, \gamma)^{28}\text{Si}$. The reactions are also used in calibration because no natural sources of radioactivity provide γ 's with energy higher than 3.6 MeV, whereas the $^{27}\text{Al}(p, \gamma)^{28}\text{Si}$ reaction provides γ ray energies up to 10.7 MeV, ensuring that the detector is well calibrated over the entire energy range for γ 's that will be seen in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction. The exact relationship between the channel number in the acquisition system analog-to-digital converter (ADC) and the incident photon energy, E_γ , is characterized by the standard linear relationship

$$E_\gamma = m \times \text{Channel} + b \quad (3.3)$$

where m is simply the slope and b the offset of the fit. For a HPGe detector, a linear relationship is sufficient and appropriate to describe the ADC response. However, this process was redone for every phase of the experiment because slightly different gains applied to the ADC's and signal amplifiers provide a different relationship in the electronics. Therefore, despite using the same detector, each phase of the experiment required its own energy calibration, an example of which is shown in Fig. 3.1 for the CASPAR setup.

3.4 Efficiency

Efficiency is a term that can, in different contexts, have many different meanings. However, in experimental physics it is known to be the ratio for the response of an instrument to the actual physical quantity for which it is employed to measure. In these cases, it is the ratio between the photons recorded and those emitted in a given event. For this experiment, both the total efficiency, η^{tot} , and the full-energy peak (FEP) efficiency, η^{FEP} , were required. The total efficiency is the probability that the γ ray enters and deposits any amount of energy within the detector, while, on the other hand, the FEP efficiency is the probability that the full energy of an emitted γ ray will

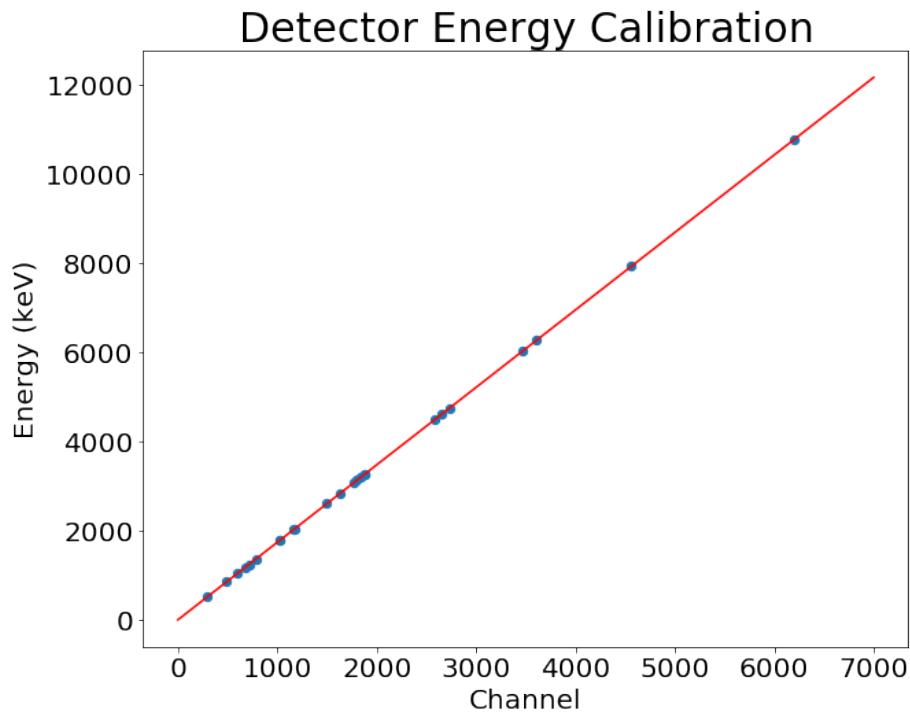


Figure 3.1. Energy calibration curve for the HPGe detector taken at CASPAR. The calibration incorporated natural background, radioactive sources, and the products from the $^{27}\text{Al}(p, \gamma)^{28}\text{Si}$ reaction.

be deposited within the detector. As both efficiency types are dependent on the physical geometry of the system, they are determined for each experimental setup in turn.

3.4.1 Total efficiency

The total efficiency of a detector / source geometry is the probability that a given photon from the source will enter and deposit any amount of energy within the detector. For extremely simple geometries, the total efficiency can be calculated following the approach laid out in Debertin and Helmer [25],

$$\eta^{tot} = \frac{1}{4\pi} \int (1 - e^{-\mu x}) d\Omega \quad (3.4)$$

where μ is the energy and absorber dependent attenuation coefficient, x is the position from the detector face, and the integration takes place over the solid angle of the detector from the front to the back. This formulation makes it abundantly clear that η^{tot} is so highly dependent on the geometry of the setup.

However, in practice, there is no analytic formulation of the total efficiency because one needs to account for any potential scattering of γ -rays off of any surrounding material, like the shielding or target chamber, for example. Additionally, when measuring the total efficiency for a given setup, the presence of multiple decays from physical sources provides additional complications. Therefore, most commonly the total efficiency is determined using single-line radioactive sources, like ^{137}Cs . In this case, when accounting for the ever-present background, the total efficiency is the total number of counts in a spectrum divided by the total number of decays occurring during the measurement time, which can be easily calculated with the radioactive decay law

$$N(t) = N_0 e^{-\lambda t} A(t) = A_0 e^{-\lambda t}, \quad (3.5)$$

where $N(t)$ is the number of nuclei remaining after time t , N_0 is the initial number of radioactive nuclei, λ is the radioactive decay constant for a given nucleus, $A(t)$ is the activity of the source at time t compared to the activity at time $t = 0$ of A_0 . The decay constant, λ , can be calculated from either the nuclear lifetime, τ , or the decay half-life, $t_{1/2}$, as

$$\lambda = \frac{1}{\tau} \lambda = \frac{\ln(2)}{t_{1/2}}. \quad (3.6)$$

This is how the total efficiency for the detector systems was determined at 662 keV, the γ energy

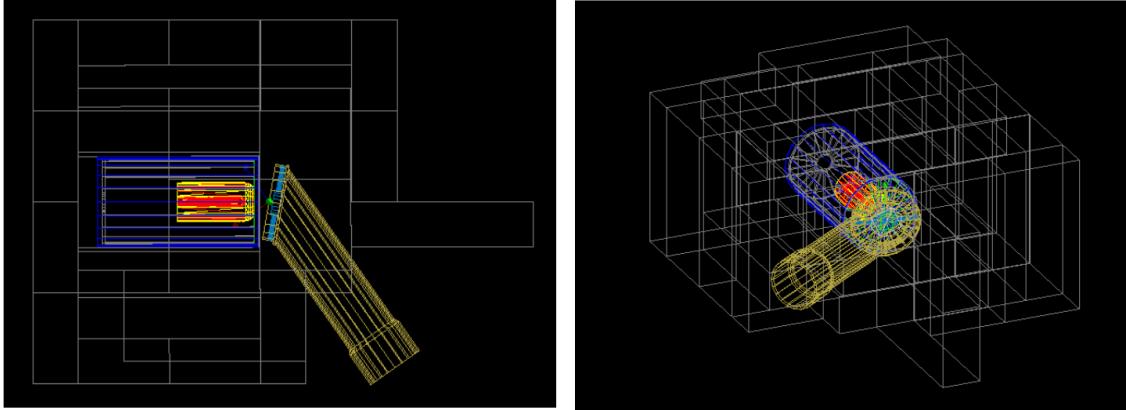


Figure 3.2. Simulation of the experimental setup at CASPAR, including the target chamber (in gold), water cooling (in light blue), HPGe detector (in dark blue), and lead bricks for shielding (in gray). Both pictures are of the same experimental setup from different angles to convey the layout. Simulating this setup is necessary to determine the total efficiency of the experimental setup. In this determination, a single-line γ source is placed at the target location and its decays simulated for a variety of energies, allowing a determination of the total efficiency at each.

from the decay of ^{137}Cs .

To extend η^{tot} to the whole range of energies relevant to the experiments, the experimental setup (including the target chamber, water cooling, and shielding) was built and simulated in Geant4 [3], an example of the simulation for the CASPAR setup is shown in Fig. 3.2 where the actual setup is given in Fig. 3.3 for comparison. Then, a single-emission source was placed at the target location and simulated 10,000,000 decays, recording the deposited energy inside the detector. The energy of the source was changed and simulated at energies of 662, 1300, 2000, 3000, 4000, 6000, 7000, 8000, and 10000 keV to cover the entire range of γ ray energies present in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction. For each of the different decay energies, the spectrum of recorded events in the detector was analyzed in the exact same way as the actual ^{137}Cs data to produce the total efficiency at each energy. This simulated data was used in conjunction with the measured ^{137}Cs total efficiency data point to determine an accurate curve of the total efficiency across all experimentally relevant energies, shown in Fig. 3.4.



Figure 3.3. A picture of the experimental setup at CASPAR, showing the lead shielding around the detector and target chamber. This figure is provided for comparison to the simulated setup to prove its accuracy.

3.4.2 Full energy peak efficiency

The full energy peak (FEP) efficiency, on the other hand, is the probability that the full energy of a given photon will be recorded within the detector. Mathematically, this is

$$\eta^{FEP} = \frac{n(E)}{R(E)} \quad (3.7)$$

where $n(E)$ is the count rate in the measured peak at an energy, E , where $R(E)$ is the rate at which photons of energy E are emitted from the source [25]. From this point, if not specified, efficiency as a term is used to denote the FEP efficiency whereas the total efficiency will always be named explicitly.

Determining the efficiency for a given setup is necessary for each geometry since, like the total efficiency, it is related to the specific source-detector geometry; it is not an intrinsic property of a given detector. In this campaign of experiments, the efficiency was determined each time with a combination of radioactive sources, like ^{137}Cs , ^{60}Co , and ^{152}Eu , and well-known nuclear resonances, like $^{14}\text{N}(p, \gamma)^{15}\text{O}$ at $E_p = 278$ keV and the $^{27}\text{Al}(p, \gamma)^{28}\text{Si}$ resonance at $E_p = 992$ keV. These reactions were chosen because, much like the energy calibration, these extend the detector's measured efficiency to a much higher energy range (up to 10.7 MeV) compared to radioactive sources.

From radioactive sources, the detector's efficiency at a given γ ray energy, $\eta(E_\gamma)$, can be deter-

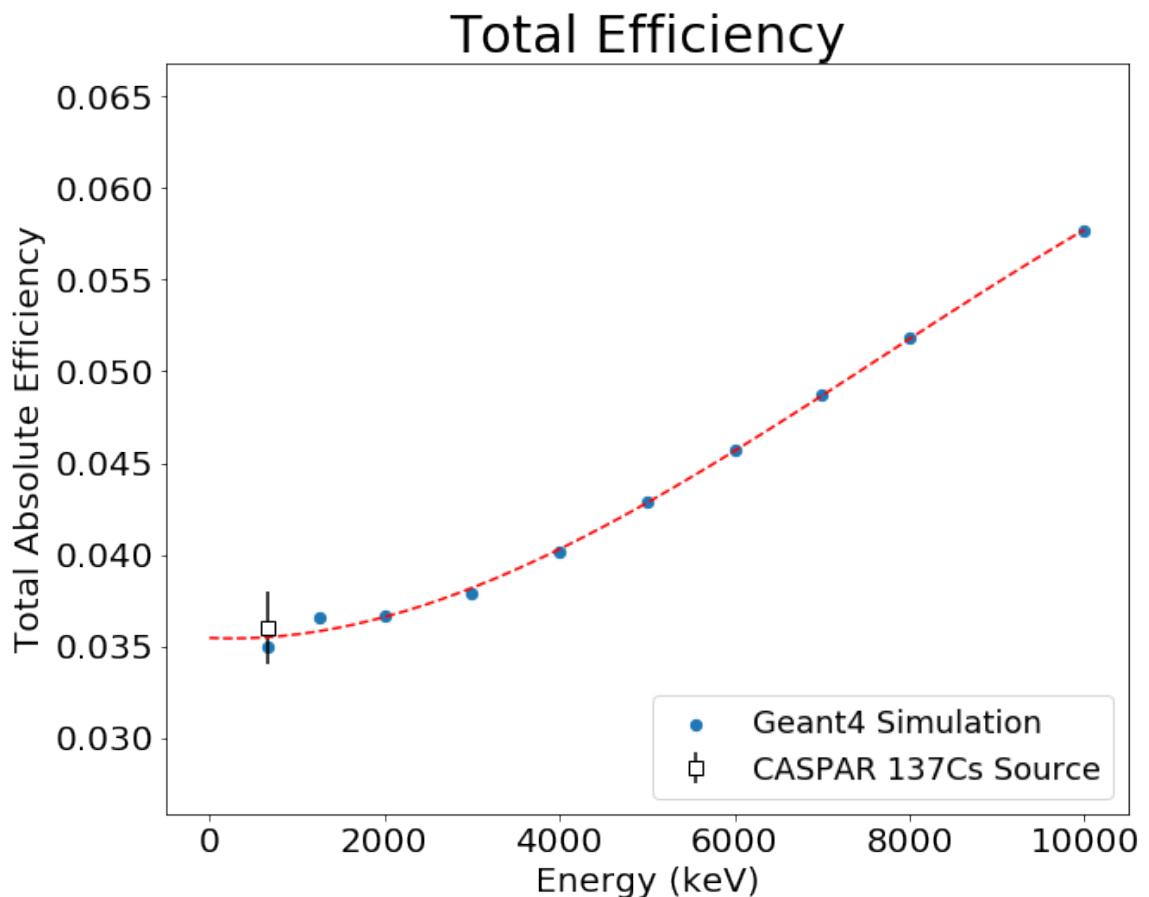


Figure 3.4. The total efficiency of the HPGe in the CASPAR setup across all relevant energies. The points from the Geant4 simulation are taken raw without scaling, showing the efficacy of this technique and accuracy of the simulation.

mined with

$$\eta(E_\gamma) = \frac{N_\gamma}{A(t)B_\gamma t_{live}C_i}, \quad (3.8)$$

where N_γ is the measured number of counts in the full energy peak, $A(t)$ is the source's activity at the time of the measurement, B_γ is the branching ratio for the specific decay, t_{live} is the live time of the measurement, and C_i are the correction factors, if necessary. For nuclear resonances, on the other hand, $\eta(E_{gamma})$ is

$$\eta(E_\gamma) = \frac{N_\gamma}{qB_\gamma RC_i}, \quad (3.9)$$

where q is the collected charge and

$$R = \frac{\lambda_r^2}{\pi} \frac{\omega\gamma}{\epsilon_{sp}} \frac{M+m}{M} \tan^{-1}\left(\frac{\Delta E}{\Gamma}\right) \quad (3.10)$$

with λ_r is the proton's de Broglie wavelength at the resonance energy, $\omega\gamma$ is the resonance strength (as defined in Chapter 1), ϵ_{sp} is the stopping power of the target at the resonance energy, M is the target atomic mass, m is the projectile atomic mass, ΔE is the thickness of the target, and Γ is the resonance width.

For this determination, the branching ratios for the 278 keV resonance in $^{14}\text{N}(p, \gamma)^{15}\text{O}$ were taken from Daigle *et al.* [24] where for the 992 keV resonance in $^{27}\text{Al}(p, \gamma)^{28}\text{Si}$ they were obtained from Antilla *et al.* [9] and those from the 406 keV resonance in $^{27}\text{Al}(p, \gamma)^{28}\text{Si}$ are taken from Powell *et al.* [50]. The efficiencies were then fit with

$$\eta = \exp\left(\sum_{i=0}^3 a_i (\ln(E_\gamma))^i\right), \quad (3.11)$$

where the a_i are fit coefficients and E_γ is the γ -ray energy and the independent variable in the fit. An example of this determination for the CASPAR setup in both the close and far geometries can be seen in Fig. 3.5. In both calculations, only statistical errors were included in the fit and the highest energy point was excluded due to summing effects, which will be discussed in more detail in the next section. This can be clearly seen as the highest energy point at close distance (1.1 cm from target) is off by an inordinately large factor. For a reference, measurements were also taken at 25.24 cm away from the target chamber, making the summing effects negligible. In this case, the summing effects provide very little impact (if any) on the highest energy point, which was still

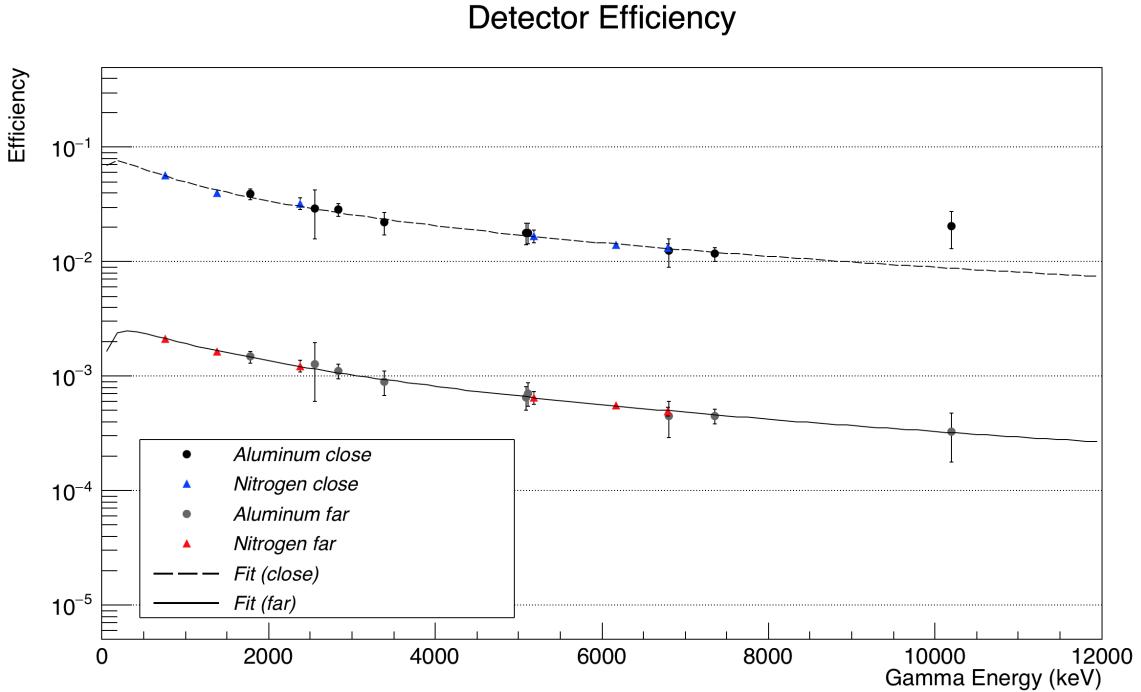


Figure 3.5. Detector efficiency as a function of γ -ray energy for the measurement at CASPAR in both close and far geometries. The highest energy point in the close geometry was excluded from the fit because it suffers from summing-in effects artificially inflating the count rate of this measured peak (see text for details). At the far distance between the detector and the target, summing effects are negligible. This can be seen as the highest-energy data point now lies perfectly with the fit from the other data.

excluded from this fit. However, for the far distance it follows the trend of the other data nearly perfectly.

3.5 Summing corrections

Summing corrections need to be applied to data due to data-recording phenomena manifesting in measured spectra. There are two different ways in which the recorded data can be affected, the so-called summing-in and summing-out effects, and these are more pronounced when measurements happen with the detector and source in close proximity or with simple reaction product decay structures, as in the case in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction.

For a simple nuclear decay, like that in Fig. 3.6, summing manifests in a measured spectrum in two different ways. If any portion of the energy from γ_1 is deposited in the detector simultaneous with γ_2 , the the resultant recorded event will have an energy higher than either individual γ -ray

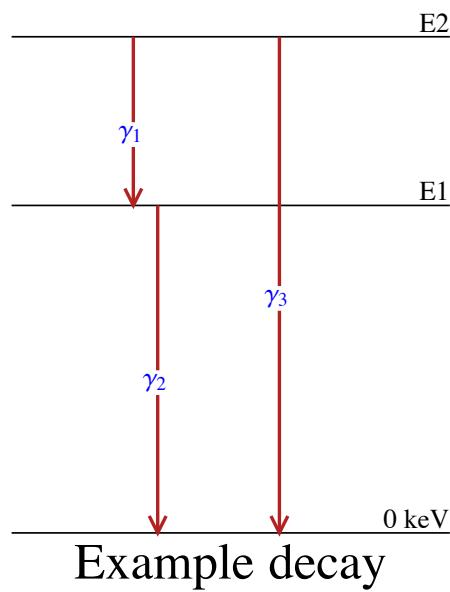


Figure 3.6. Example cascade nuclear decay for demonstrating summing effects (see text for details).

and, therefore, will appear at a different energy in the spectrum. Consequently, γ_2 's count rate will be artificially lowered. By the same logic, the count rate for γ_1 will also be artificially dropped. This loss of counts in the peaks recorded at energies E_1 and E_2 is known as the summing-out effect. As an extension of this thought experiment, if the full energy of both γ_1 and γ_2 is deposited in the detector at the same time the resulting measured energy will match E_3 , artificially inflating the count rate of that peak. This increase is called the summing-in effect.

To correct for these effects, the probability of measuring a different energy than expected must be taken into account. For a situation where there is no summing, the rate of γ rays is simply the yield

$$Y_i = RB_i\eta_i, \quad (3.12)$$

where R is the normalization from Equation 3.10, B is the branching ratio, and η is the FEP efficiency. However, when summing-out occurs, the yield is artificially dropped by the factor $RB\eta_1\eta_2^{tot}$, which is the probability of counting the full energy of γ_1 along with any part of γ_2 . Mathematically, this observed yield for both γ_1 and γ_2 is

$$Y_1 = RB_1\eta_1 - RB_1\eta_1\eta_2^{tot} \quad (3.13)$$

$$= RB_1\eta_1 (1 - \eta_2^{tot}) \quad (3.14)$$

$$Y_2 = RB_2\eta_2 (1 - \eta_1^{tot}). \quad (3.15)$$

Then, in order to correct for this, the ratio of the true emitted γ 's to the observed rate for γ_1 is

$$\frac{Y_1}{Y_1^{obs}} = \frac{1}{1 - \eta_2^{tot}} = C_1, \quad (3.16)$$

where the C_i is the correction factor. The same relation holds for γ_2 . This makes clear why the total efficiency is so important: without it, summing-out is not an effect which can be corrected. For γ_3 , however, the situation requires the total energy deposition of γ_1 and γ_2 at the same time, so correcting for summing-in requires the FEP efficiency. This is

$$Y_3 = RB_3\eta_3 + RB_1\eta_1\eta_2 \quad (3.17)$$

where the second term includes B_1 because the decay chain has to start with γ_1 . By the same token,

the correction factor for summing-in is

$$C_3 = \frac{Y_3}{Y_3^{obs}} = \frac{1}{1 + \left(\frac{B_1\eta_1\eta_2}{B_3\eta_3} \right)}. \quad (3.18)$$

With a more complicated nuclear level scheme, the summing corrections become more difficult to apply accurately. A more detailed treatment of the necessary methods can be found in [25]. This is primarily due to more complicated feeding and the presence of multiple decays from a given level. In general, the correction is for the i th level is

$$C_i = 1 + C_i^{in} - C_i^{out} \quad (3.19)$$

$$C_i^{in} = \sum_{m,n} \frac{B_m p_n \eta_m \eta_n}{B_i \eta_i} \quad (3.20)$$

$$C_i^{out} = \sum_{j,k} p_k \eta_k^{tot} + \frac{B_j p_i \eta_j^{tot}}{B_i} \quad (3.21)$$

where the p_i are the probability of feeding. Fortunately, for the ^{15}O nucleus, all intermediate decays have a branching ratio of 100%, as can be seen in Fig. 3.7. The summing out corrections in each case can then be

$$C_i^{out} = 1 - \eta_j^{tot}, \quad (3.22)$$

where the i and j denote the γ rays feeding into and decaying from the state, respectively (like γ_1 and γ_6 for example). For this measurement, the summing-out effect for every transition is relatively small, falling in the range of 3.6 - 5.2 % for all of the relevant γ 's. Fig. 3.8 shows this, depicting the summing out correction across the range of relevant gamma energies in this experiment.

Conversely, only the R/DC→GS transition needs to be corrected for summing-in effects as the total energy of the other decay branches matches the energy of γ_7 , and, as such, should be a larger effect since the true branch to the ground state is $\sim 4\%$. The observed yield of γ_7 in this case will be

$$Y_7 = RB_7\eta_7 + RB_1\eta_1\eta_6 + RB_2\eta_2\eta_5 + RB_3\eta_3\eta_4. \quad (3.23)$$

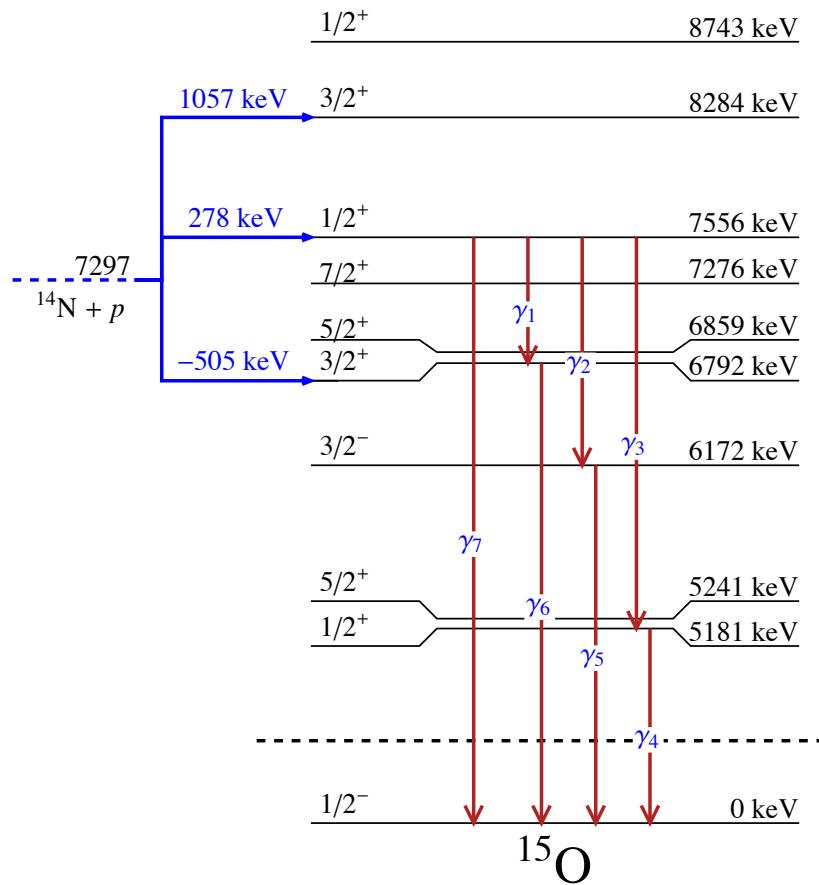


Figure 3.7. All decays from the 278 keV proton resonance in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction. Other than the resonance level, all decays are sequential, with the levels having only one decay directly to ground state. Of note, this then means that the total energy for each decay sequence matches the direct transition, γ_7 , so it must be corrected for summing-in effects from all the other branches.

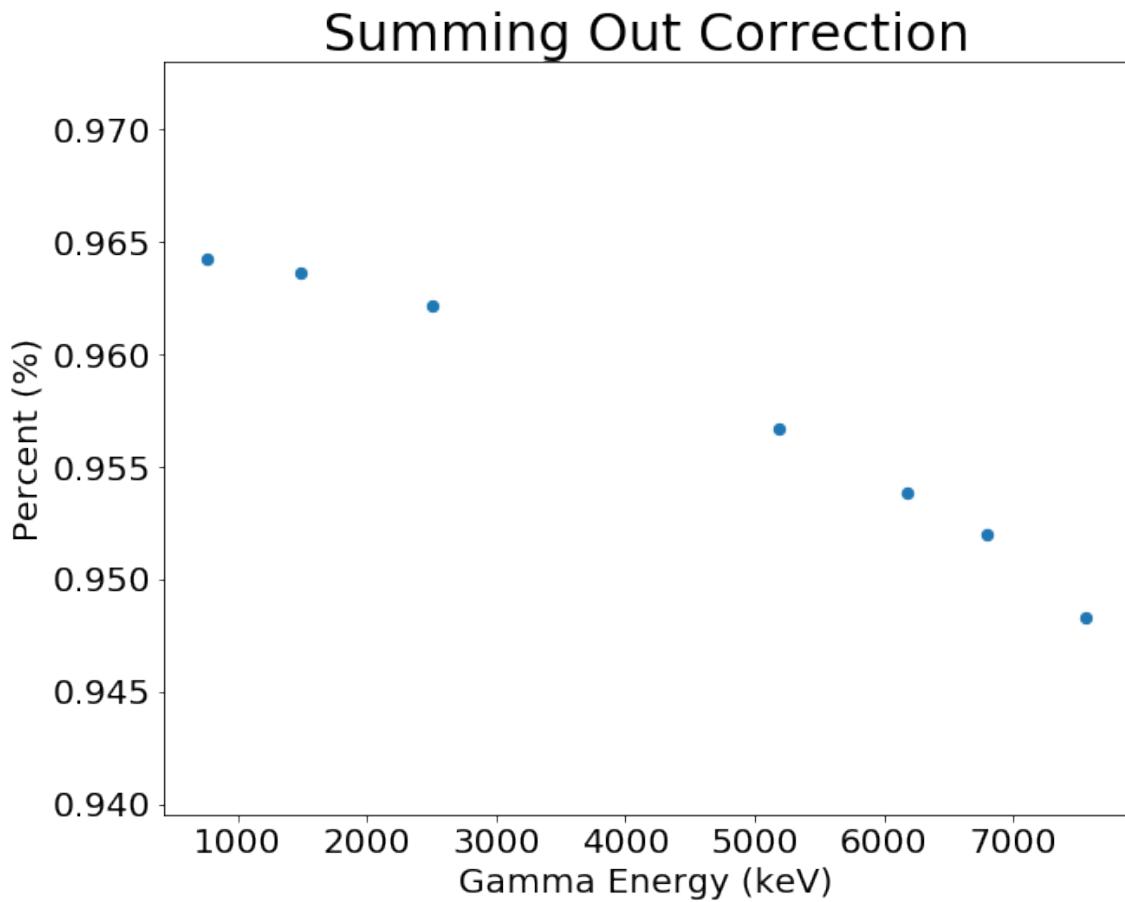


Figure 3.8. The summing out correction as a function of incident γ -ray energy. This figure shows the percentage of gammas recorded to the true incident number.

From this, then, the correction factor for the ground state transition will necessarily be

$$C_7^{in} = 1 + \frac{B_1\eta_1\eta_6 + B_2\eta_2\eta_5 + B_3\eta_3\eta_4}{B_7\eta_7}. \quad (3.24)$$

In this measurement, this correction is quite large. It was determined to be a 61% correction to the ground state transition using this method.

There is a second method for calculating the summing-in correction which was also employed. This relies on the measured efficiency being taken at different distances to the source of the γ rays. In close geometry, the solid angle of the detector is much larger when viewed from the source and summing is present (which can be seen from the highest energy data point in the detector efficiency plot, Fig. 3.5). However, at a larger distance, in this case 25.4 cm, the solid angle of the detector is significantly smaller but summing-in effects are not present in the detector. Therefore, the ratio of efficiencies between the close and far distance measurements when accounting for the difference in detector solid angle is the necessary summing-in correction. For this method, the summing-in correction was found to be 57% for the ground state transition.

The fact that these two methods give nearly the same value for the summing-in effect is encouraging to the efficacy of these results. When incorporating this result into the final cross section determination, the mean of these two correction values was taken, at 59%.

3.6 Target characterization

In order to ultimately determine the nuclear cross section and the astrophysical S -factor for a given reaction, the number of active atoms in the target must be known. As a reminder, the nuclear cross is a probability of interaction. The number of possible reaction sites must be known. The targets used in these measurements were sputtered TiN and ZrN targets, backed by Ta and W, respectively. By performing an energy scan over a known, thin resonance with the targets it is possible to determine the number of active atoms in the target.

Fig. 3.9 presents a typical example of a resonance scan, which is just the measured excitation function over a resonance. This one was taken at CASPAR with the 278 keV resonance in $^{14}\text{N}(p, \gamma)^{15}\text{O}$ with ZrN target (# 5) by monitoring the $R \rightarrow 6.79$ MeV transition. Even for a monoenergetic beam, there is still a small gaussian distribution of energies encompassed in the beam (see Chapter 2 for more details). When the high energy tail of this distribution reaches the low energy side of the resonance, the measured yield from that resonance will appear. Further increasing the beam energy implies that more of the energy distribution of the beam reaches the resonance

energy and the yield continues to increase. This low-energy side of the scan is referred to as the front edge and contains the intrinsic information about the resonance as well as the beam distribution. The true resonance energy, E_r is found halfway up the front edge, 50% of the maximum yield, because at this point a symmetric beam distribution would be half above and half below the resonance in energy. Similarly, the slope of the front edge details the energy spread in the beam, with a broader distribution corresponding to a lower slope and a tighter distribution providing a sharp front edge.

For the incident beam particles that are above the resonance energy, they will lose energy traveling through the target and drop to energies within the resonance width and proceed through resonant capture. The maximum yield for such a scan will occur when the entire beam distribution is at an energy such that either a maximum of the ions are within the resonance width or the energy loss in the target brings a maximum number of the ions into that same range. This maximum is known as the plateau and is the flat portion at the top of the curve in Fig. 3.9. As a target's thickness increases the thicker this plateau becomes, since the ions will have a greater energy loss within the active target material, allowing them to decrease to resonance energy from higher and higher starting energies. For an infinitely thick target, the resonance plateau would never decrease again on the high side, because the beam would always lose enough energy to fall back into the resonance while still in the active target material.

When targets are not infinitely thick, however, the beam energy can be increased to a point where the energy loss of the ions will eventually be such that the energy of the ions is still above the resonance width when leaving the active target material, meaning the reaction no longer proceeds through the resonant capture mechanism. When this happens, the yield starts to decrease from the level of the plateau. By a similar argument to the beam distribution leading to increased yield on the front edge, the yield on the back edge (where the plateau decreases again) drops because of the same distribution effects. The slope of this decrease is affected by the target material (leading to different energy losses) and the homogeneity (or lack thereof) of the active material. The full-width half-maximum (FWHM) of this excitation curve represents the target thickness in terms of energy loss, ΔE .

For such a thin resonance occurring at E_r , the stopping power of the target, $\epsilon(E)$, the de Broglie wavelength, λ , and the partial widths Γ_i do not change appreciably over the resonance energy. This allows the maximum yield, at E_0 , to be calculated using [34]

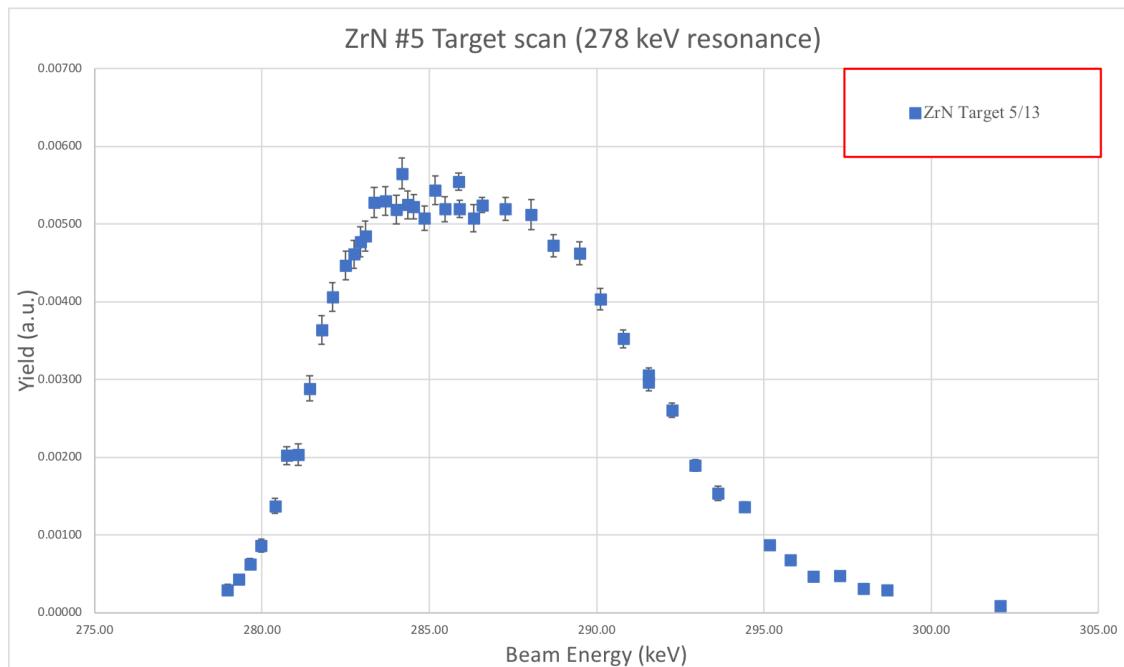


Figure 3.9. Resonance scan (measured Excitation curve over a resonance) taken at CASPAR with the 278 keV resonance in $^{14}\text{N}(p, \gamma)^{15}\text{O}$ with ZrN target (# 5). This shows typical behavior of an energy scan over a thin resonance and from a measurement like this, the number of nitrogen atoms in the target can be determined (see text for details).

$$Y(E_0) = \int_{E_0 - \Delta E}^{E_0} \frac{1}{\epsilon(E)} \frac{\lambda^2}{4\pi} \omega \frac{\Gamma_a \Gamma_b}{(E_0 - E)^2} dE = \frac{\Gamma^2 / 4}{\Gamma^2 / 4} dE \quad (3.25)$$

$$= \frac{\lambda_r^2 \omega \gamma}{2\pi \epsilon_r} \left[\tan^{-1} \left(\frac{E_0 - E_r}{\Gamma/2} \right) - \tan^{-1} \left(\frac{E_0 - E_r - \Delta E}{\Gamma/2} \right) \right]. \quad (3.26)$$

Then, with some further algebraic reduction [34], one can show

$$E_0 = E_r + \frac{\Delta E}{2} \quad (3.27)$$

$$Y_{max} = \frac{\lambda_r^2 \omega \gamma}{\pi \epsilon_r} \arctan \left(\frac{\Delta E}{\Gamma} \right) \quad (3.28)$$

$$FWHM = \sqrt{\Gamma^2 + \Delta E^2} \quad (3.29)$$

$$n = \frac{\Delta E}{\epsilon_r}, \quad (3.30)$$

where n is the number density of target atoms, atoms per unit area. When the target thickness, ΔE is much bigger than the resonance width, Γ , the FWHM of the resonance scan is simply the target thickness. Thus, by selecting thin resonances and determining the energy loss from their scans, like that in Fig. 3.9, the number density of active nuclei in the target can be determined with Equation 3.30. This method is useful because it does not rely on any external efficiency calibrations or corrections (since such a correction would apply to the entire curve equally, not changing the FWHM and, therefore, the thickness). Thus, this method is effective and appropriate for a uniform target material with known stoichiometry.

The stopping powers used for these determinations were calculated using the James F. Ziegler's industry standard SRIM program, [66]. The output from this program provides the discrete values for the stopping power of a user specified target material over a range of user specified energies. Using the stopping powers at the resonance energy in units of (eV / (atoms/cm²)) and the overall uncertainty in the calculations of 4.6% [66] the number of target atoms can be determined for each resonance scan. Table 3.1 summarizes the results for this calculation and gives the initial number density of nitrogen atoms for the targets used throughout this experiment.

However, by delivering more beam to the targets, more of the active material is removed and the targets degrade. In order to account for this effect, multiple resonance scans were taken using the different targets at different points in the experiments in order to determine the number of active target atoms throughout the measurement. By taking the initial scans as baseline values and

Target	$n \left(10^{17} \text{ atoms / cm}^2\right)$
TiN	7.070 ± 0.566
ZrN #5	5.623 ± 0.450
ZrN #1	5.826 ± 0.466
ZrN #12	10.480 ± 0.834

TABLE 3.1

ACTIVE TARGET ATOMS FOR EACH OF THE TARGETS USED AS A PART OF
THIS MEASUREMENT.

comparing the target thickness as a function of charge delivered a linear relationship can be deduced. Fig. 3.10 shows the deterioration of the targets as a function of the delivered charge throughout the experiment and the linear fits to these data. These relationships can then be used to independently correct the individual measured runs for the accumulated charge on the target up to that particular point.

3.7 Cross-section determination

In the following analysis, the differential cross section determination for each of the measured transitions follows the same procedure with the different individual correction factors for each of the transitions. Therefore, without loss of generality, they all occur according to the following procedure.

The yield of each transition, Y_i , is the number of reactions per incident projectile (protons, in this case). It was calculated by

$$Y_i = \frac{N_\gamma C_{in}}{\eta N_p C_{out}}, \quad (3.31)$$

where N_γ is the measured number of γ -rays at the energy corresponding to the transition of interest, C_{in} and C_{out} are the summing in and out corrections, respectively, N_p is the number of protons measured on target throughout the run, and η is the detector efficiency at the measured γ ray energy, E_γ . The uncertainty in this yield arises from the uncertainties in the counts of the γ -ray peak and the efficiency. In the case of the off-resonance region of the cross section, the uncertainty

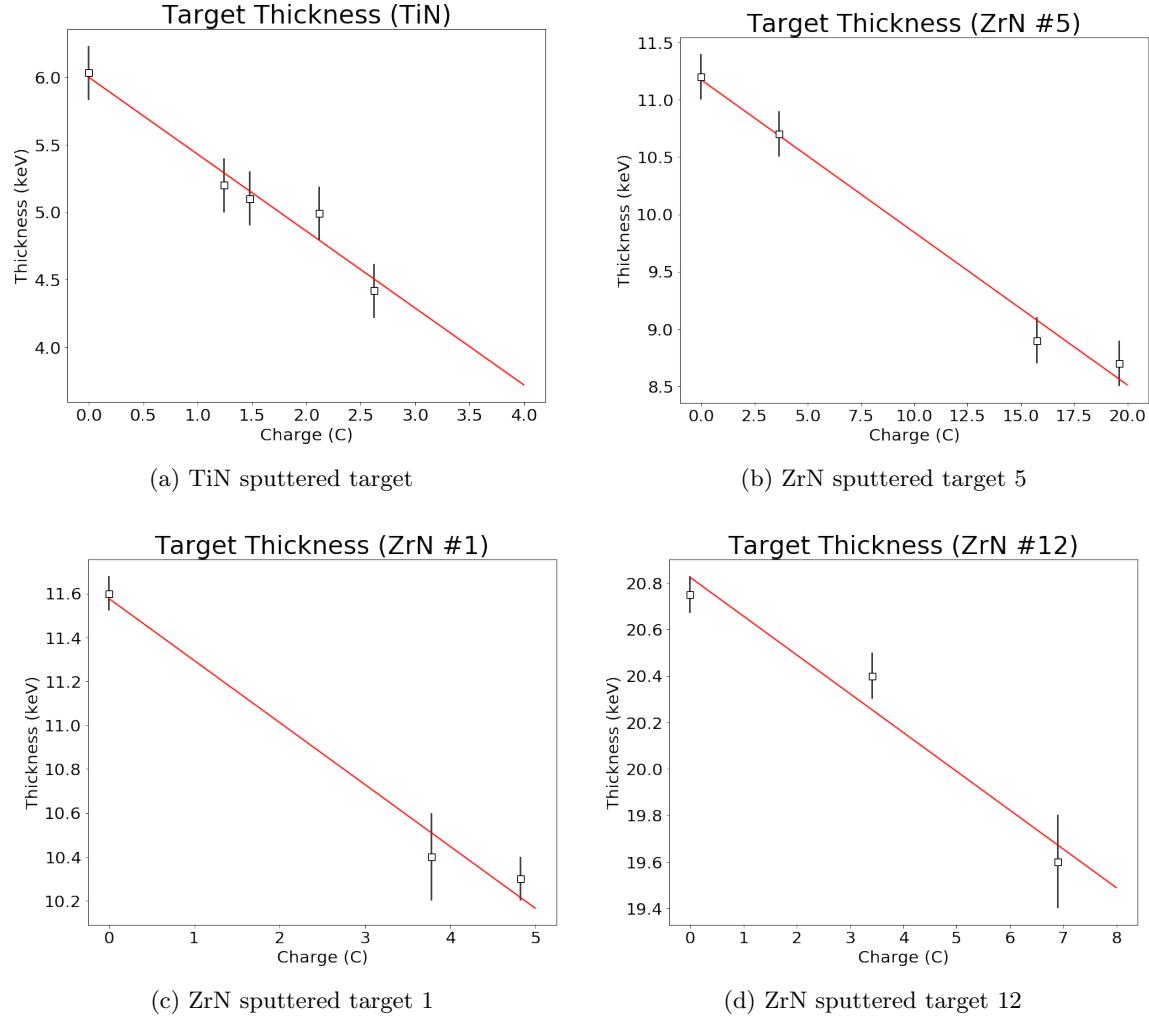


Figure 3.10. The thicknesses of the various targets used in this experiment as a function of the charge delivered. This shows the deterioration of the targets through the course of the experiment where the thickness was monitored by scanning known resonances.

in the number of counts is the dominant term. These uncertainties are propagated through the yield calculation for each individual data point.

The experimental yield at a given incident proton energy, $Y(E_p)$, corresponds to the differential cross section, $\frac{d\sigma}{d\Omega}$, integrated over the target thickness, ΔE :

$$Y(E_p) = \int_{E_p - \Delta E}^{E_p} \frac{d\sigma(E)}{d\Omega} \frac{1}{\epsilon(E_p)} dE, \quad (3.32)$$

where, again, ϵ is the target's stopping power at the incident proton energy.

For a thin target, where the cross section is constant over the target thickness, this reduces to the familiar

$$\frac{d\sigma}{d\Omega} = \frac{Y}{N_T} = \frac{N_\gamma C_{in}}{\eta N_p C_{out} N_T}, \quad (3.33)$$

where N_T is the number of active target nuclei (in this case, nitrogen). This is the method by which the differential cross section of the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction was determined for the different transitions. The uncertainty on this differential cross section was calculated for each individual point by propagating the uncertainty in the number of nitrogen atoms with the uncertainty in the calculated yield. This also took into account the loss of nitrogen within the targets over the course of the experiment (see Sec. 3.6 for details). The resulting differential cross sections for the 6.79 MeV and ground state transitions are given in Figs. 3.11 and 3.12, respectively. From these calculations, the astrophysical S -factor was subsequently determined. These are shown, alongside literature data from [36, 42, 53, 56], in the series of Figs. 3.13 - 3.19. Since the literature data is given as total S -factor, their data was scaled down by a factor of 4π to give a differential S -factor with which to compare our measured data.

For the 6.79 MeV transition, the points with large errors (specifically at 900 keV and 1000 keV) are energies at which the 6.79 MeV peak of interest interfered with a background peak (from Ti). The large error point at 1060 is only in the Notre Dame data and comes from not running long enough to see appreciable counts in this transition. The resonance at this energy populates the ground state strongly and that was the focus of the measurement at those energies. This same lack of statistics is why the data points at 800 keV also have comparatively larger uncertainties. These features can be seen highlighted in Fig. 3.15.

For the ground state transition, the trend in our data indicates that our measurement agrees well with the Schröder data [56] and Li data [42] coming in from higher energies, with our measured

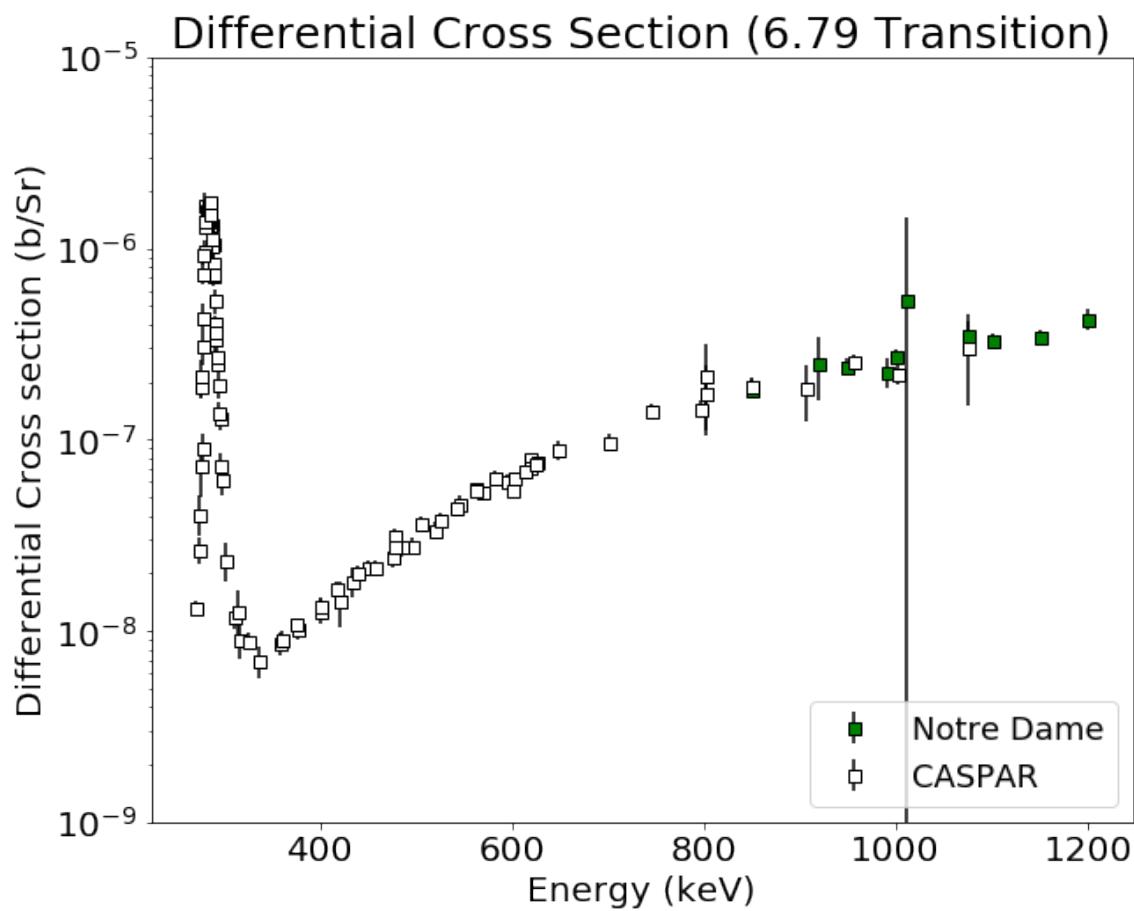


Figure 3.11. Differential cross section for the 6.79 MeV transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction.

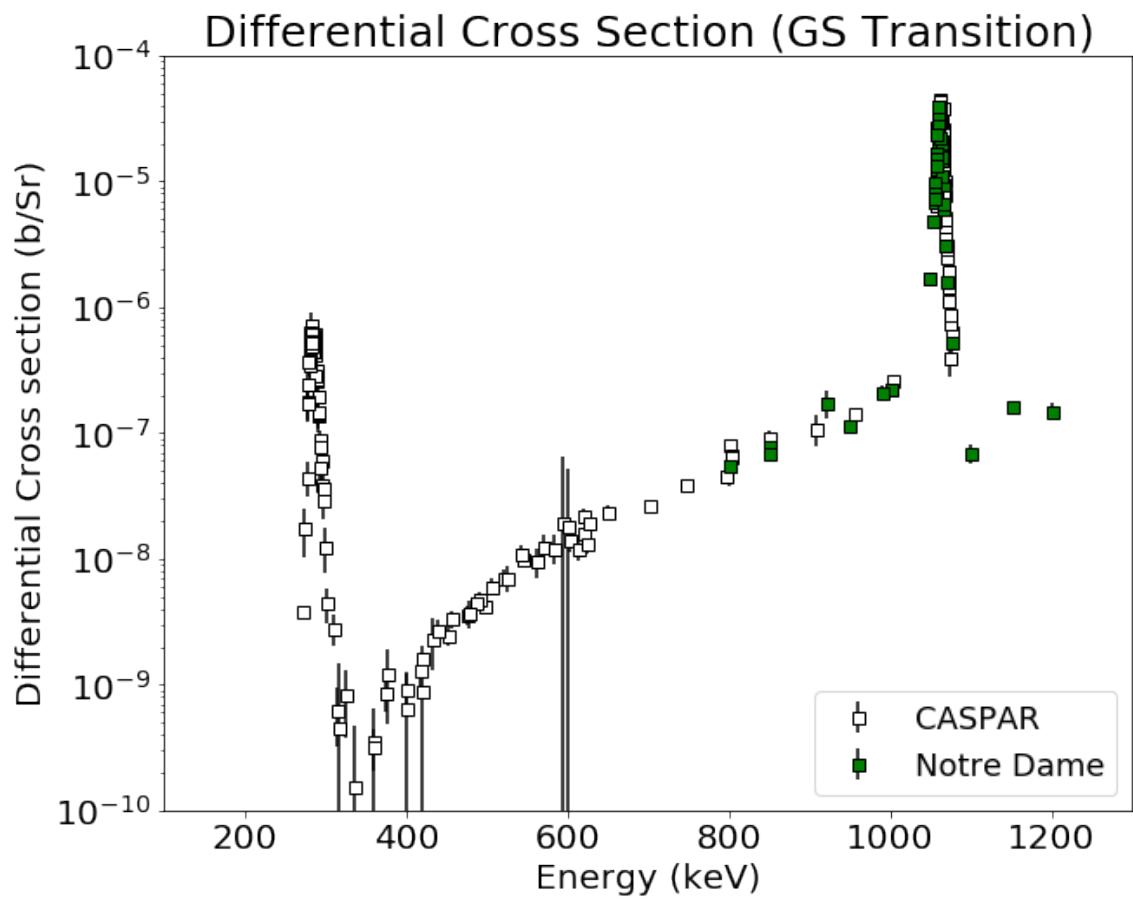


Figure 3.12. Differential cross section for the ground state transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction.

S -factor dropping in the interference region between 300-400 keV to agree with the LUNA [36] and Runkle [53] data there. Our overall scale seems to be higher in the off resonance region, however, and only agrees with the highest energy data from the TUNL experiment [53]. Overall, this data shows significantly higher uncertainties at lower energies than the 6.79 MeV transition data. This is because this is a much more weakly populated transition and it was not feasible to run long enough to achieve appreciable statistics in the γ -ray data at these energies. Therefore, they have uncertainties dominated by the rate of measured γ 's in the experiment.

In order to move from differential measurements to a total cross section or total S -factor, as well as extrapolate to the Gamow window, an R -matrix calculation was performed combining the results from this measurement with the results from the lifetime measurement, described later in Chapter 4. The results from this R -matrix calculation, extrapolation, and implications from this measurement are presented in Chapter 5.

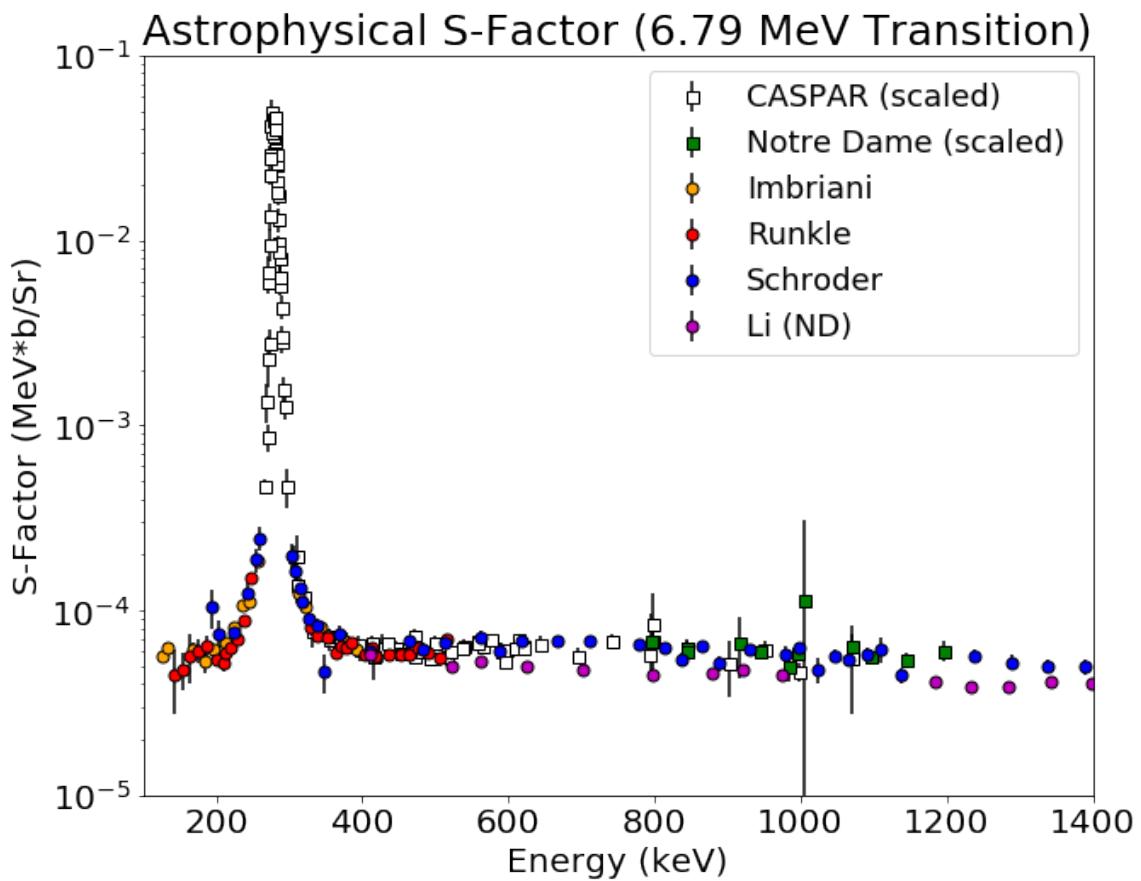


Figure 3.13. Differential S -factor for the 6.79 MeV transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction for the entire energy range measured in this experiment. Also plotted are data from [36, 42, 53, 56] for comparison.

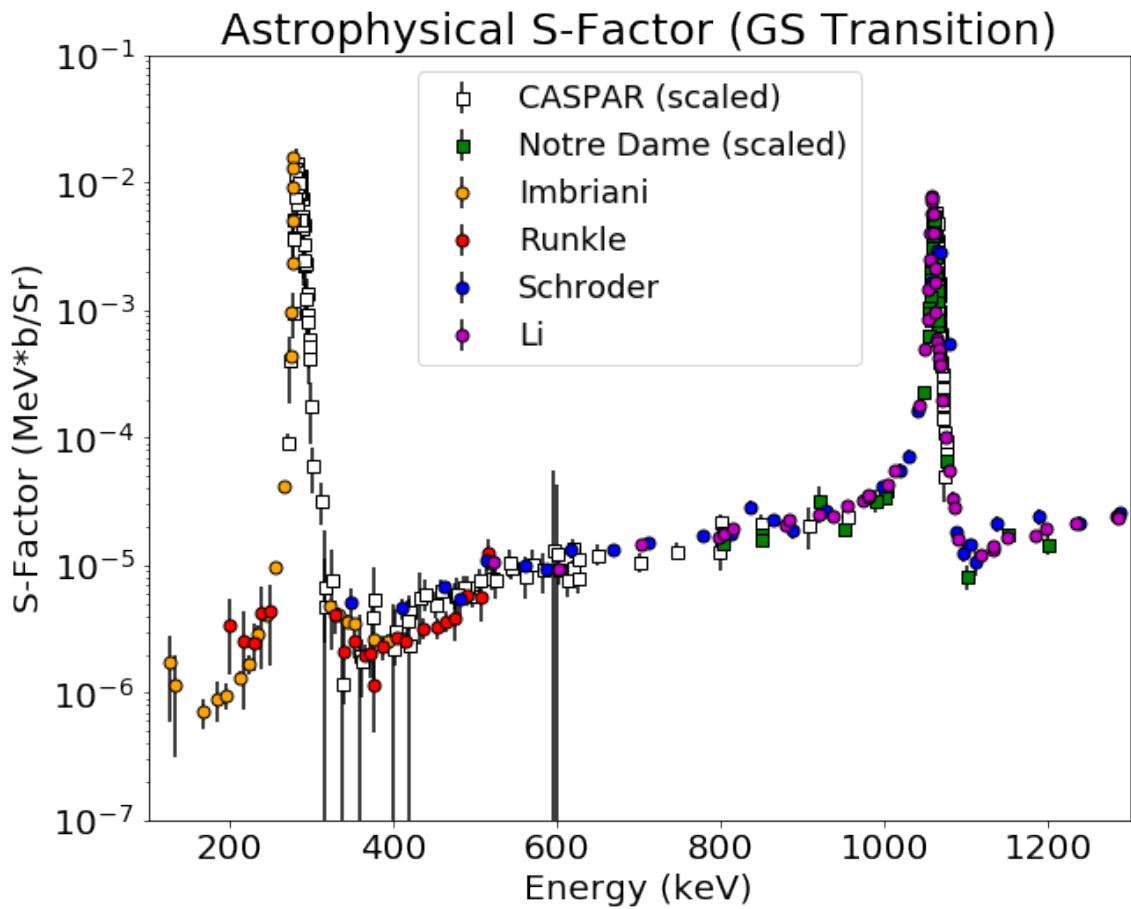


Figure 3.14. Differential S -factor for the ground state transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction for the entire energy range measured in this experiment. Also plotted are data from [36, 42, 53, 56] for comparison.

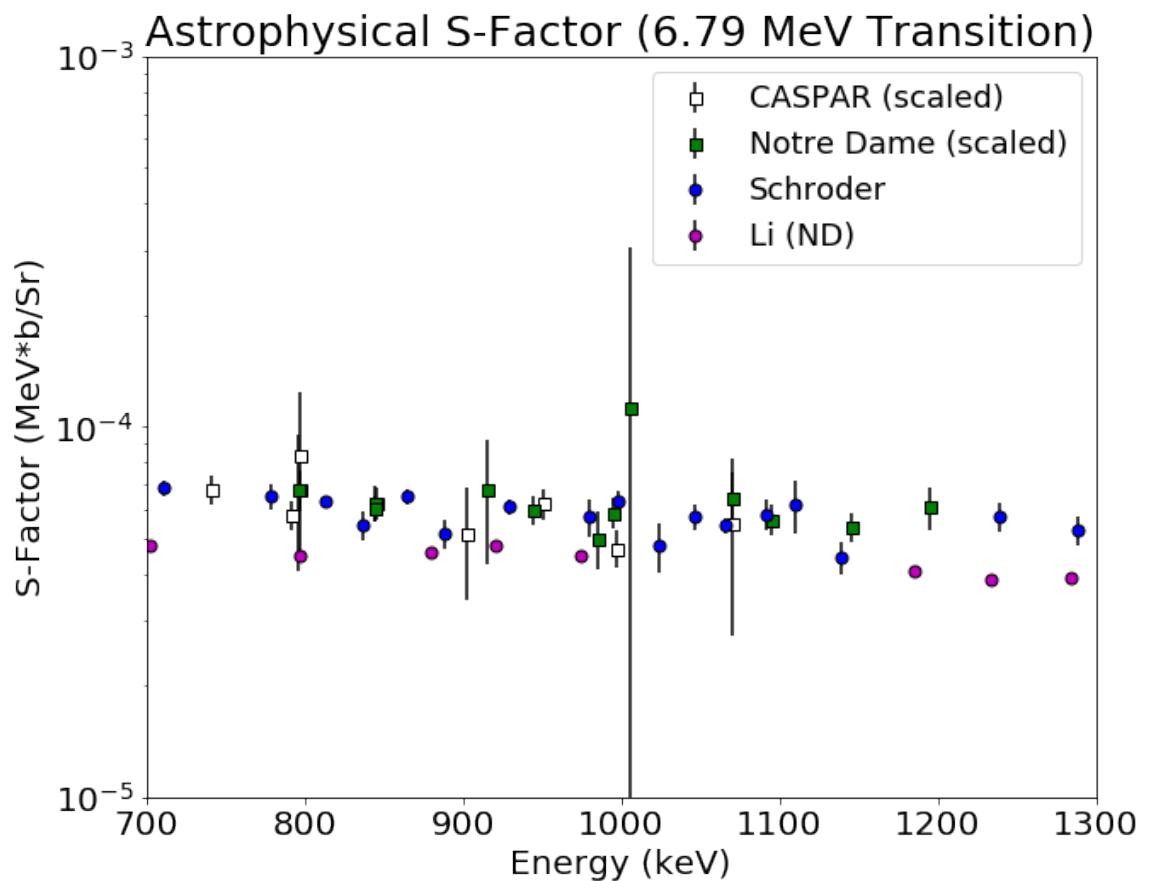


Figure 3.15. Differential cross section for the 6.79 MeV transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction, highlighting the high energy region to compare the data taken at the NSL to CASPAR, showing the efficacy of the facility. Also plotted are data from [42, 56] for comparison.

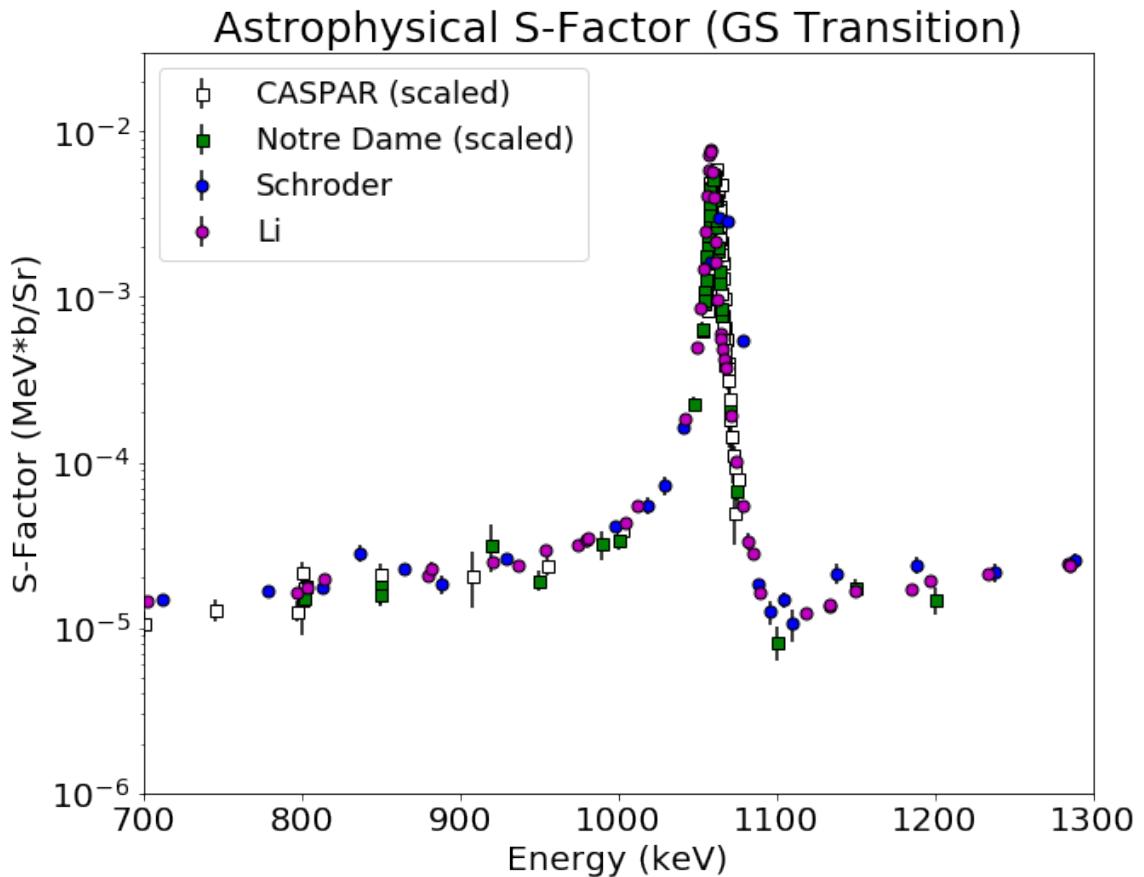


Figure 3.16. Differential cross section for the 6.79 MeV transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction highlighting the high energy region to compare the data taken at the NSL to CASPAR, showing the efficacy of the facility. Also plotted are data from [42, 56] for comparison.

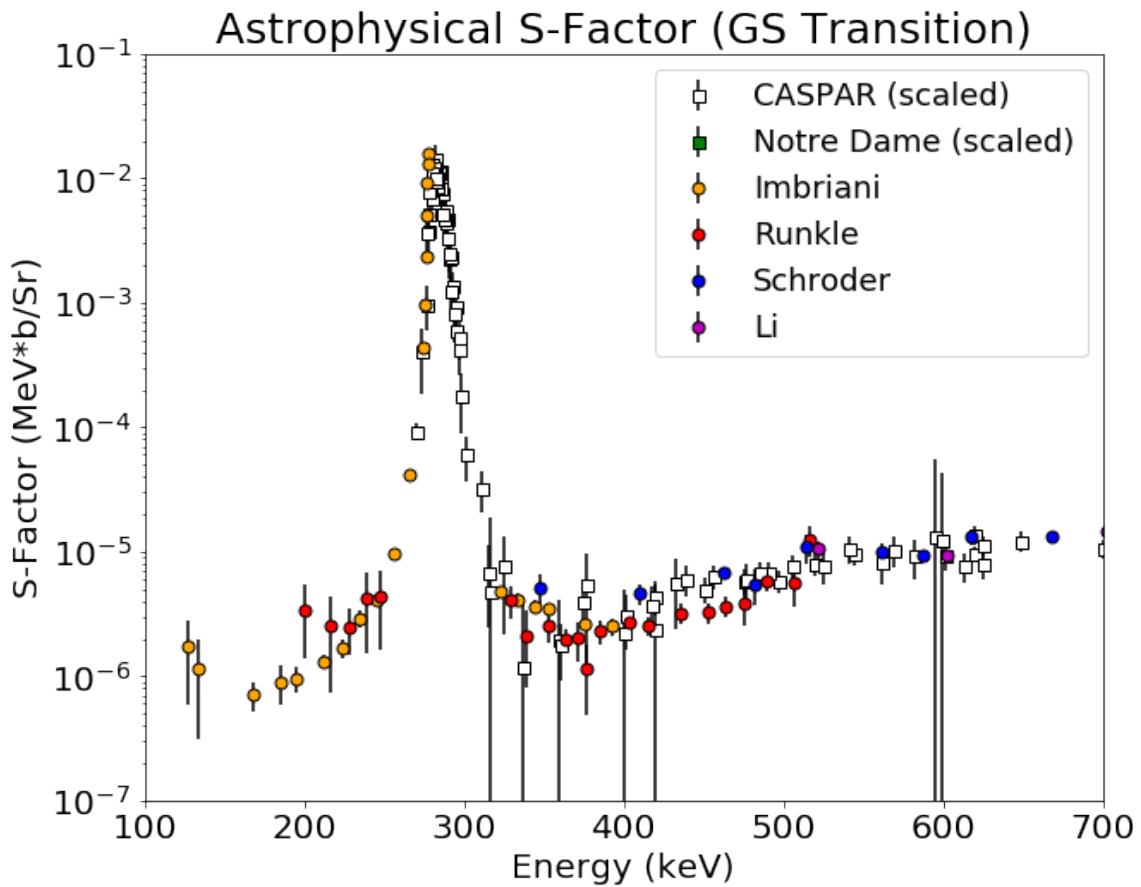


Figure 3.17. Differential cross section for the ground state transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction, primarily detailing the off resonance region measured at CASPAR in this experiment. Also plotted are data from [36, 42, 53, 56] for comparison.

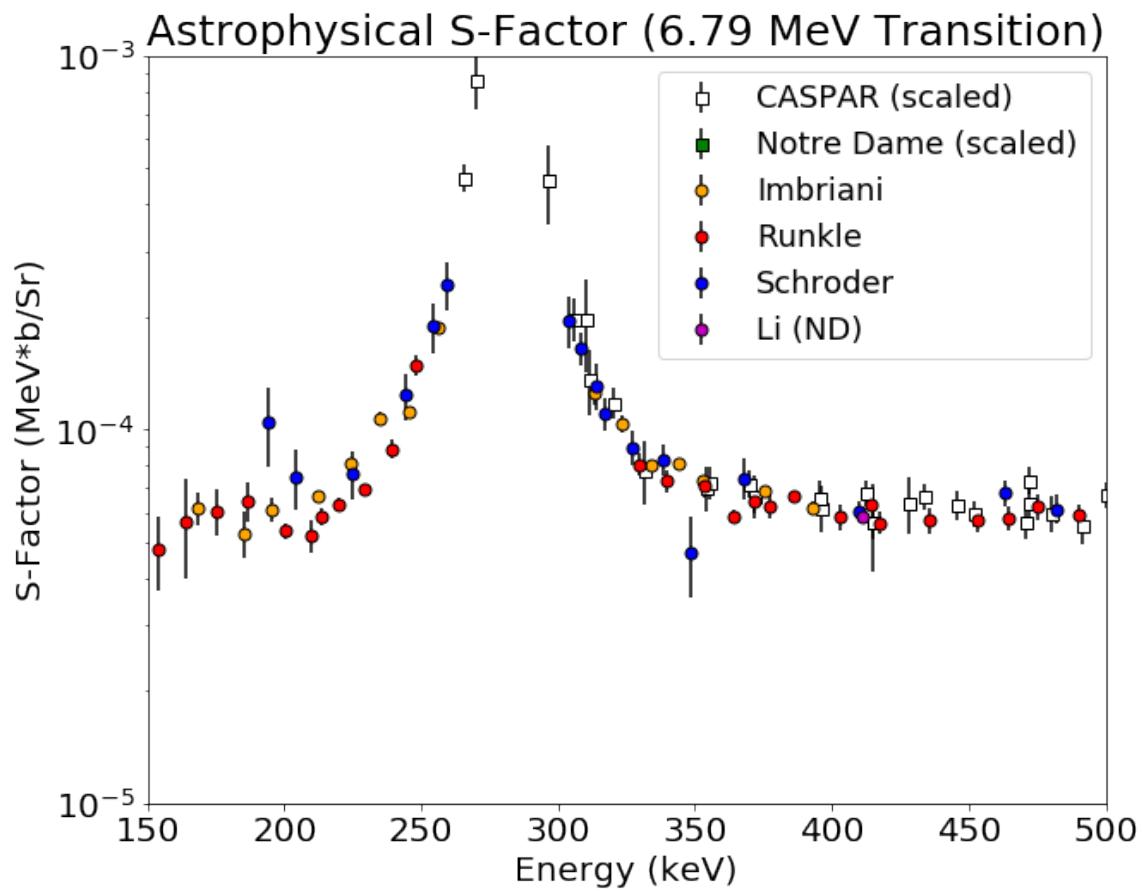


Figure 3.18. Differential cross section for the 6.79 MeV transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction, showing the behavior at low energy. Also plotted are data from [36, 42, 53, 56] for comparison.

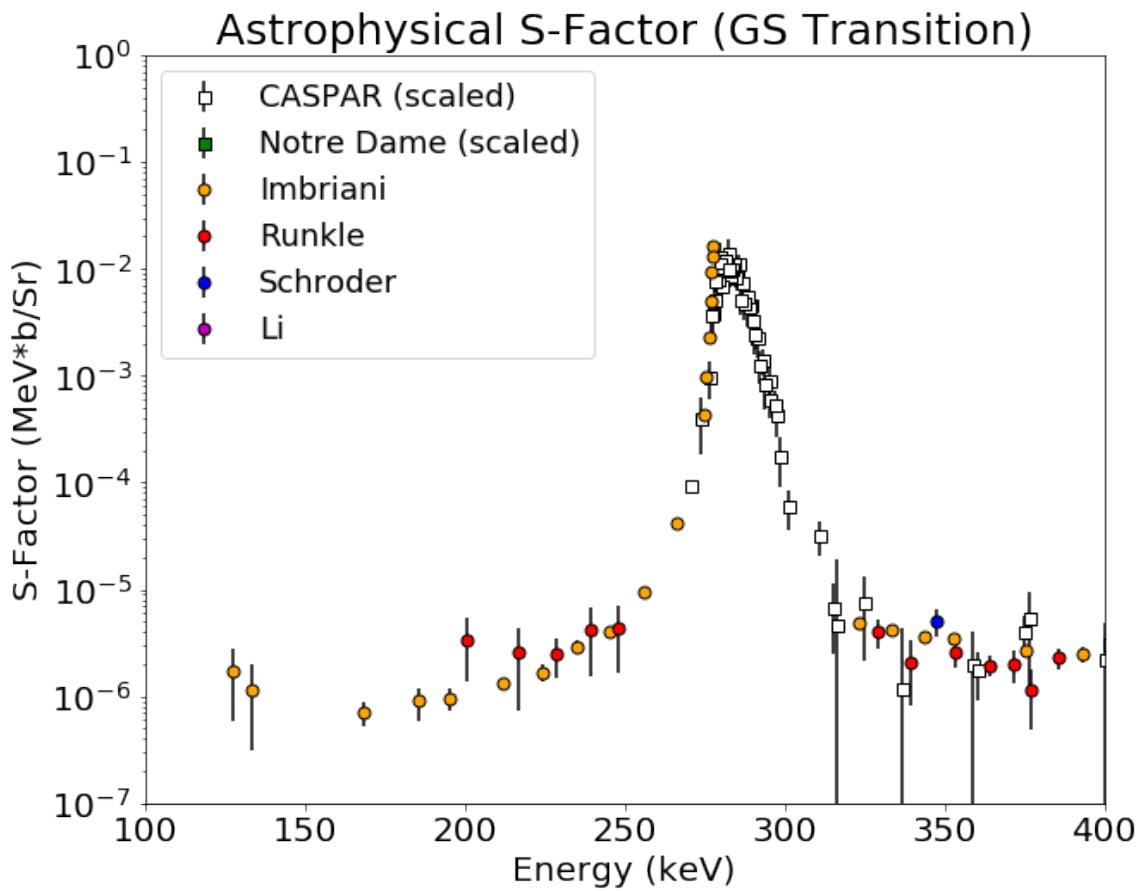


Figure 3.19. Differential cross section for the ground state transition in the $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction, showing the behavior of the S -factor at low energy. Also plotted are data from [36, 42, 53, 56] for comparison.

CHAPTER 4

MEASURED LIFETIMES

- 4.1 Determining a nuclear lifetime
- 4.2 Lifetime of the 5.18 MeV state in ^{15}O
- 4.3 Lifetime of the 6.17 MeV state in ^{15}O
- 4.4 Lifetime of the 6.79 MeV state in ^{15}O

CHAPTER 5

R-MATRIX ANALYSIS

5.1 Fits to the capture data

5.2 Inclusion of the new lifetime

CHAPTER 6

RESULTS AND CONCLUSIONS

6.1 Reaction rates

6.2 $^{14}\text{N}(p, \gamma)^{15}\text{O}$ reaction summary and outlook

APPENDIX A

ANALYSIS CODES

A.1 Definitions

Several codes used within this analysis are reproduced herein for posterity and ease of access.

```

1  ****
2  * The calibration program from the
3  * 14N(p,g)150 6.79 MeV state lifetime
4  * measurement (data replay and calibration)
5  * Usage with root:
6  *      .L calibrate.cpp
7  *      Run(runNumber) - for individual run
8  *      addData() - for all runs (need to change loop)
9  *
10 * The way I wrote this was to first "RECREATE" the calibratedSpectra.root file
11 * so I interactively do Run(1), then I change this file so that the run() function
12 * now updates the TFile you create. After that point, simply load into root and use
13 * the runAll() function, after changing your run number bounds, to create and add
14 * the
15 * remaining histograms to the new storage file.
16 *
17 *
18 *
19 * Bryce Frentz (bfrentz@nd.edu)
20 ****
21
22 #include <cstring>
23 #include <iostream>
24 #include <fstream>
25 #include <cstdio>
26 #include "TCanvas.h"
27 #include "TRandom2.h"
28 #include "TChain.h"
```

```

29 #include "TH1F.h"
30 #include "TH2F.h"
31 #include "TFile.h"
32 #include "TCut.h"
33 #include "TTreeReader.h"
34 #include "TTreeReaderValue.h"
35 #include "TTreeReaderArray.h"
36
37 using namespace std;
38
39
40 // Run function
41 // Inputs: integer runNum
42 // Outputs:
43 // This method takes a run number and takes the data from the given run, calibrates
44 // the spectra,
45 // and outputs the resulting histogram to a new file (calibratedSpectra.root).
46 void Run(int runNum){
47
48     // Create a random # generator in range [0,1]
49     TRandom2 myRandom(0);
50
51     //Optimize cache and disk reading because this isn't default
52     //in this version of ROOT.
53     TChain chain("evtTree");
54     chain.SetCacheSize(1E8);
55     //chain.AddBranchToCache("*");
56
57     // For adding files in chain
58     ifstream fRuns;
59     char dataf[255];
60
61
62     // Get data
63     sprintf(dataf, "./lifetime_%03d.root", runNum);
64     TFile *f = new TFile(dataf);
65     cout << "Data File: " << dataf << endl;
66     chain.Add(dataf);
67
68

```

```

69 // Create an output file for the histograms
70 TFile file("./calibratedSpectra.root","UPDATE");
71
72
73 // Create histogram
74 char title[255];
75 sprintf(title, "Georgina Spectrum run%03d", runNum);
76 TH1D *h0 = new TH1D("h0", title, 8192, -0.5, 9999.5);
77
78 char hname[7];
79 sprintf(hname, "run%03d", runNum);
80
81 // Centroid of 1460.83: 8718.83
82 // Centroid of 6129.59: 36609.6
83 // Calibration slope and offset:
84 Double_t a = 0.167393;
85 Double_t b = 1.381436;
86 // Event energy
87 Double_t ecal;
88
89 // Get event tree
90 TTreeReader reader(&chain);
91 TTreeReaderArray<UShort_t> event_e(reader,"event.e");
92
93 ULong64_t nEvent = chain.GetEntries();
94 cout << nEvent << " events to be analyzed." << endl;
95
96
97 // Start the event loop
98 while(reader.Next()) {
99     if(event_e[0] > 0) {
100         // Calibrate from channel to energy and fill output run histogram
101         ecal = a * (event_e[0] + myRandom.Rndm() -0.5) + b;
102         h0->Fill(ecal);
103     }
104     else ecal = 0;
105 }
106
107 h0->SetName(hname);
108 h0->Write(hname, TObject::kWriteDelete);
109

```

```
110 //file.Write();
111 file.Close();
112
113 cout << hname << " has been added." << endl;
114 cout << endl;
115
116 f->Close();
117
118 }
119
120
121 // Main method
122 // Inputs:
123 // Outputs:
124 // This method loops over all files, invoking the Run method in order to add all the
125 // data to one file
126 void addData(){
127
128     for(int i = 192; i < 241; i++){
129         Run(i);
130     }
131
132     cout << endl;
133     cout << "Finished adding all runs." << endl;
134     cout << "Have a nice day." << endl;
135     cout << endl;
136 }
```

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