



LIBRARY Michigan State University

This is to certify that the

dissertation entitled

Decay Studies of the Neutron-Rich Nuclides 19 20 N and N

presented by

Don William Anthony

has been accepted towards fulfillment of the requirements for

Ph.D. degree in <u>Chemical Ph</u>ysics

Date July 18, 2000

MSU is an Affirmative Action/Equal Opportunity Institution

0-12771

PLACE IN RETURN BOX to remove this checkout from your record.

TO AVOID FINES return on or before date due.

MAY BE RECALLED with earlier due date if requested.

DATE DUE	DATE DUE	DATE DUE

11/00 c:/CIRC/DateDue.p65-p.14

Decay Studies of the Neutron-Rich Nuclides ¹⁹N and ²⁰N

By

Don William Anthony

A DISSERTATION

Submitted to
Michigan State University
in partial fulfillment of the requirements
for the degree of

DOCTOR OF PHILOSOPHY

Department of Chemistry

2000

ABSTRACT

DECAY STUDIES OF THE NEUTRON-RICH NUCLIDES 19N AND 20N

By

Don William Anthony

Secondary, radioactive ion beams of ¹⁹N and ²⁰N were produced at the National Superconducting Cyclotron Laboratory at Michigan State University by fragmenting a primary beam of ²²Ne (80 MeV/A) in a beryllium target. The isotopes of interest were implanted within a plastic scintillator and their decay was monitored. Beta events were recorded by the implantation scintillator, delayed-neutron emission was observed by the neutron bar array as well as three ⁶Li inorganic scintillators, and gamma-ray events were recorded by two germanium detectors. Quantitative measurements of the branching ratios for delayed-neutron and gamma ray emission were made for both ¹⁹N and ²⁰N, and the half-lives of the species were deduced.

The ¹⁹N data suggests that there are twelve delayed-neutrons emitted during the decay comprising 50.9(62)% of the total decay strength. The gamma-ray spectroscopy yields several branches to bound, excited levels of ¹⁹O contributing 39.4(38)% of the total decay strength. There is also evidence of significant delayed-neutron emission to bound excited states of ¹⁸O. The half-life is determined to be 299(19) milliseconds and the ground state spin and parity is restricted to (1/2- or 3/2-). From the delayed-neutron data, the resonant capture strength for the ¹⁸O(n, γ)¹⁹O reaction has been estimated and found not to compete well with the direct capture component except at the highest temperatures possible during the Inhomogeneous Big Bang scenario. The measured

ranching ratios and inferred properties of the populated intermediate states do not compare well with shell model predictions.

The ²⁰N data suggests eight delayed-neutron branches with energies in excess of 1.1 MeV. Above this threshold, delayed neutron emission accounts for 34.0(40)% of the total decay strength. The gamma-ray analysis leads to 27.0(46)% of the decay proceeding through bound excited states of the ²⁰O daughter. There is also evidence that delayed-neutron emission to ¹⁸O populates excited bound states rather than the ground state. The half-life has been re-measured to be 121(15) milliseconds and the ground state spin and parity of ²⁰N is restricted to be low-spin, most likely (1-).

Dedicated to the memory of my grandparents

Bill and Mary Anthony

George and Edna Friess

ACKNOWLEDGEMENTS

I would first like to thank my supervisor and friend, Dave Morrissey, for his guidance and support through this process. Without his good counsel and eleventh-hour heroics, this experiment could not have been successful. Second, I want to thank Paul Mantica for permitting me to take part in his experimental program, and for his continual willingness to help me learn to become an experimentalist.

I also wanted to express my gratitude to the graduate students, past and present, at the NSCL. I think that I learned more with and from them than any other source. I look forward to continuing the lifelong friendships that have been forged here over the last four years.

To my parents, Dave and Marcelle Anthony and my brother and sister, David and Dawna, you all must know that your support and love have made it possible for me to meet with success in my life. I owe you all more than I could express with mere words.

To my wife, Paola, thank you for everything. There is absolutely no doubt that these achievements are as much yours as they are mine. I can't imagine life without you and I love you more than anything in the world. You have been the most blessed gift in my life and everything I have done, or will do, is simply because of you.

And finally, Abigail, you have changed my life. Now I have a sense of scale, of what is important, and of what "unconditional love" means. I am forever yours.

TABLE OF CONTENTS

LIST OF FIGURES	ix
LIST OF TABLES.	xiiv
CHAPTER 1 INTR	ODUCTION1
1.1 General	Introduction1
1.1.1	Structure
1.1.2	Nucleosynthesis
1.2 Literatur	re Search – ¹⁹ N, ²⁰ N
1.2.1	Prior Measurements of ¹⁹ N Decay Properties6
1.2.2	Prior Measurements of ²⁰ N Decay Properties9
1.2.3	Literature Search Summary10
1.3 Shell Me	odel Predictions for the Decay of ¹⁹ N12
1.4 Experim	nental Goals13
CHAPTER 2 EXPE	ERIMENTAL SETUP14
2.1 Producti	on of Secondary Radioactive Beams14
2.2 Experim	nental End-Station15
2.2.1	Implantation System16
2.2.2	Neutron Bar Array17
2.2.3	⁶ Li Glass Scintillators19
2.2.4	High Purity Germanium Detectors19
2.3 Electron	ics and Data Acquisition20
2.4 Seconda	ry Beam Purity21

CHAPTER 3 EXPE	RIMENTAL RESULTS	27
3.1 Overview	v of Analysis Procedure	27
3.2 Half-Life	Measurement of ¹⁹ N	28
3.3 Half-Life	Measurement of ²⁰ N	29
3.4 Total Nu	mber of Events	30
3.4.1	Dead Time Considerations	31
3.4.2	Summary	35
3.5 Neutron	Bar Array	36
3.5.1	Time Calibration of the Neutron Bar Array	37
3.5.2	Neutron Bar Array Efficiency Calibration	45
3.5.3	Neutron Bar Results – ¹⁹ N and ²⁰ N	49
3.5.4	¹⁹ N Delayed Neutron Spectrum	49
3.5.5	²⁰ N Delayed Neutron Spectrum	52
3.6 ⁶ Li Inorga	anic Scintillators	54
3.6.1	Efficiency Calibration	57
3.6.2	¹⁹ N Results – ⁶ Li Detectors	60
3.7 Gamma-l	Ray Data	63
3.7.1	Gamma-ray Spectra of the Isotopes of Interest	66
3.7.2	Background Gamma Rays	71
3.7.3	Ratio Of Efficiencies	71
3.7.4	Gamma Ray Segregation	72
3.7.5	Gamma-Ray Half-Life Measurements	73
376	Gamma Ray Assignments ¹⁹ N. ²⁰ N Secondary Beams	75

3.7.7 Efficiency Calibration – Standard Sources	77
3.7.8 Efficiency Calibration – On-Line Sources	80
3.7.9 Efficiency Calibration – Germanium Coincidences	82
3.7.10 Efficiency Calibration – Summary	83
3.7.11 Gamma Ray Intensities – ¹⁹ N	87
3.7.12 Intensities of Higher Energy ¹⁹ N Gamma Rays	88
3.7.13 Gamma Ray Intensities – ²⁰ N	90
CHAPTER 4 INTERPRETATION	93
4.1 Interpretation of ¹⁹ N Gamma Rays	93
4.2 Interpretation of ¹⁹ N Delayed Neutrons	96
4.3 Delayed Neutron Assignments ¹⁹ N	99
4.4 Interpretation of ²⁰ N Gamma Rays	101
4.5 Interpretation of ²⁰ N Neutron Spectrum	104
4.6 ¹⁹ N Shell Model Considerations	106
4.7 Beta Decay of ²⁰ N	108
4.8 Astrophysics Results ¹⁹ N	110
4.8.1 Direct Capture Part of the $^{18}O(n,\gamma)^{19}O$ Reaction Rate	111
4.8.2 Resonant Capture Part of the ¹⁸ O(n,γ) ¹⁹ O Reaction Rate	114
4.8.3 Comparison of Direct and Resonant Capture Strengths	116
CHAPTER 5 SUMMARY AND CONCLUSIONS	118
5.1 Summary and Conclusions – ¹⁹ N	118
5.2 Summary and Conclusions – ²⁰ N	121
I IST OF REFERENCES	122

LIST OF FIGURES

FIGURES IN THIS DISSERTATION ARE PRESENTED IN COLOR

Figure	1 Energy level diagram of the mass 19 system pertinent to the decay of ¹⁹ N	7
Figure	2 Energy level diagram of the mass 20 system pertinent to the decay of ²⁰ N1	. 1
Figure	3 Schematic diagram of the implantation detection system	17
Figure	4 Schematic diagram of the Neutron Bar Array	8
Figure	5 Schematic diagram of the entire detector end-station	20
Figure	6 Electronics diagrams of the PPAC, Real-Time clock and RF pulsing control	
	elements of the end-station	22
Figure	7 Electronics diagrams for the neutron bars, ⁶ Li detectors and germanium detectors	23
Figure	8 Electronics diagrams for the components of the Implantation system2	<u>?</u> 4
Figure	9 Fit to the half-life spectrum of ¹⁹ N	29
Figure	10 Fit to the half-life spectrum of ²⁰ N	0
Figure	11 Delayed neutron spectrum of ¹⁶ C showing the gates used to reflect back onto the half-life spectrum for the dead-time analysis	33
Figure	12 Best fit curves to the neutron-gated half-life spectra of ¹⁶ C	14
Figure	13 Fit to the full half-life spectrum recorded by the implantation scintillator during the decay of ¹⁶ C	35
Figure	14 Spectrum collected by one of the array bars from the precision time calibrator.	38
Figure	15 Delayed-neutron spectrum collected by one of the array bars during the decay of ¹⁶ C	4 0
Figure	16 Plot of inverse velocity versus channel centroid for the calibration of the neutron bar array	44

Figure	Aggregate TOF delayed-neutron spectrum recorded by the neutron bar array uring the decay of ¹⁶ C47
Figure	Aggregate TOF delayed-neutron spectrum recorded by the neutron bar array uring the decay of ¹⁷ N
Figure	Calibration curve for the total efficiency of the neutron bar array49
Figure	Delayed-neutron TOF spectrum collected by the neutron bar array uring the decay of ¹⁹ N
Figure	Delayed-neutron TOF spectrum collected by the neutron bar array luring the decay of ²⁰ N
Figure	2 Delayed-neutron TOF spectrum collected by ⁶ Li detector #3 during ne decay of ¹⁶ C
Figure	B Delayed-neutron TOF spectrum collected by ⁶ Li detector #3 during ne decay of ¹⁷ N
Figure	Time calibration curve for ⁶ Li detector #3 from the ¹⁶ C and ¹⁷ N decays56
Figure	5 Calibration curve for the total efficiency of ⁶ Li detector #3
Figure	Parameters necessary to determine the geometric efficiency of an perature
Figure	7 Energy spectrum of delayed neutrons recorded by ⁶ Li detector #1 uring the decay of ¹⁹ N
Figure	3 Energy spectrum of delayed neutrons recorded by ⁶ Li detector #2 uring the decay of ¹⁹ N
Figure	Energy spectrum of delayed neutrons recorded by ⁶ Li detector #3 uring the decay of ¹⁹ N
Figure	Gamma-ray spectrum collected in Ge#2 from the ⁶⁰ Co calibration sourc65
Figure	Energy calibration curves for Ge#1 and Ge#265
Figure	2 Gamma-ray spectrum collected by Ge#1 during the decay of ¹⁶ C67
Figure	Gamma-ray spectrum collected by Ge#2 during the decay of ¹⁶ C67
Figure	Gamma-ray spectrum collected by Ge#1 during the decay of ¹⁷ N68

Figure	35 Gamma-ray spectrum collected by Ge#2 during the decay of ¹⁷ N68
Figure	36 Gamma-ray spectrum collected by Ge#1 during the decay of ¹⁹ N69
Figure	37 Gamma-ray spectrum collected by Ge#2 during the decay of ¹⁹ N69
Figure	38 Gamma-ray spectrum collected by Ge#1 during the decay of ²⁰ N70
Figure	39 Gamma-ray spectrum collected by Ge#2 during the decay of ²⁰ N70
Figure	40 Ratio of counts detected in Ge#1 to Ge#2 during the beam-off period of the ¹⁹ N experimental runs
Figure	41 Ratio of counts detected by Ge#2 over all times to beam-off during the decay of ¹⁹ N
Figure	42 Gamma-ray gated half-life spectra for two transitions observed in Ge#175
Figure	43 Gamma-ray spectrum of ⁶⁰ Co collected by the NaI start detector79
Figure	44 Gamma-ray spectrum collected in Ge#2 gated on the ⁶⁰ Co 1332 keV line in the NaI start detector
Figure	45 Data points taken for the efficiency determination of Ge#284
Figure	46 Best fit to the efficiency calibration data for Ge#285
Figure	47 Interpolation and extrapolation error bars for the efficiency calibration of Ge#2
Figure	48 Partial energy level diagram of ¹⁹ O showing the bound state transitions observed in this experiment during the decay of ¹⁹ N94
Figure	49 The beta decay of ¹⁹ N showing the transitions that can be ascribed to ¹⁹ N on the basis of the experimental data101
Figure	50 Partial energy level diagram of ²⁰ O showing the bound state transitions observed in this experiment during the decay of ²⁰ N
Figure	51 The beta decay of ²⁰ N showing the transitions that can be ascribed to ²⁰ N on the basis of the experimental data108
Figure	52 Comparison of the direct capture and resonant capture cross sections

Figure 53 The beta decay of ¹⁹ N showing the transitions that may be conclusively identified from the experimental data	.122
Figure 54 The beta decay of ²⁰ N showing the transitions that may be conclusively identified from the experimental data	.125

LIST OF TABLES

Table 1 Literature search for experimental measurement of ¹⁹ N and ²⁰ N decays10
Table 2 Beam-on and Beam-off pulsing times for each nuclide investigated15
Table 3 Beam purity at the A1200 focal plane and the experimental end-station for those secondary nuclides isolated in this experiment
Table 4 Total number of decays recorded by the implantation detector for each of the isotopes of interest. Also given is the calculated number of decays from the implantation rate and scaler measured dead-times
Table 5 PMT slopes for the neutron bar array as deduced from the precision time calibrator data
Table 6 Calibration delayed neutrons of ¹⁶ C and ¹⁷ N given by energy and branching ratio
Table 7 Centroid channels of the neutron peaks utilized in the calibration of the neutron bar array
Table 8 Centroid channels of the prompt beta peaks utilized in the calibration of the neutron bar array
Table 9 Parameters derived from the delayed-neutron energies and their centroid channels necessary to calibrate the neutron bars
Table 10 Summary of the calibration fits for each of the array bars. The parameters of and offset are generated from the M and B results
Table 11 Data points used to generate an efficiency calibration for the neutron bar array. The Monte-Carlo points are from reference [93Ha]48
Table 12 Delayed-neutron lines observed during the decay of ¹⁹ N. The TOF is given along with the neutron energies in both the lab and Center of Mass frames51
Table 13 Delayed-neutron branching ratios for those neutron lines observed in the neutron bar array during the decay of ¹⁹ N
Table 14 Summary of the delayed neutrons observed during the decay of ²⁰ N. Listed are the centroid channels and the corresponding neutron energy in both the lab and Center of Mass frames
Table 15 Calculated branching ratios for the delayed-neutron lines observed by the neutron bar array during the decay of ²⁰ N

Table 16 Calibration Parameters for the lithium detectors in the seven bit time domain. The inverse velocity of a neutron in (ns/m) is equal to the TDC channel times the slope plus the offset
Table 17 Calibration points utilized to determine the total efficiency of each lithium detector. The source neutrons are from the decay of ¹⁶ C and ¹⁷ N
Table 18 Calculated value of the geometric efficiency and best-fit value of the scaling parameter for each of the lithium detectors
Table 19 Branching ratio determination for delayed neutrons observed by the lithium detectors during the decay of ¹⁹ N. Also tabulated are the parameters necessary for the branching ratio calculation along with the associated errors
Table 20 Gamma rays attributed to background71
Table 21 Gamma ray assignments for transitions observed during the beam-off period of the ¹⁹ N production runs
Table 22 Gamma ray assignments for transitions observed during the beam-off period of the ²⁰ N production runs
Table 23 Cross table for ⁶⁰ Co. For N gamma rays emitted at "column" energy there are N times "cell value" gamma rays emitted at "row" energy in coincidence78
Table 24 Cross table for ²⁰⁷ Bi. For N gamma rays emitted at "column" energy there are N times "cell value" gamma rays emitted at "row" energy in coincidence
Table 25 Cross table for ¹³⁴ Cs. For N gamma rays emitted at "column" energy there are N times "cell value" gamma rays emitted at "row" energy in coincidence78
Table 26 Off-line source calibration data for determining the efficiencies of the Ge detectors. The emboldened columns are the data necessary to generate a calibration curve
Table 27 On-line data pertinent to the efficiency calculation of the Ge detectors82
Table 28 Ge detector efficiencies determined using the coincidence technique83
Table 29 Gamma-ray intensities for transitions assigned to the beta decay of ¹⁹ N87
Table 30 Literature values for the de-excitation branching of the 3945 keV excited state of ¹⁹ O
Table 31 Calculated intensities of the de-excitation gamma rays from the 3945 keV excited state of ¹⁹ O

Table 32 Final gamma-ray intensities for those gamma rays attributed to the decay of 19N
Table 33 Calculated relative intensities of gamma rays emitted during the decay of ²⁰ N. The intensities are relative to the 1674 keV line that is set to 10091
Table 34 Absolute intensity of gamma rays ascribed to the decay of ²⁰ N92
Table 35 Intensity of the gamma-ray lines attributed to the decay of ²⁰ N92
Table 36 Gamma-ray transitions identified as being a result of ¹⁹ N beta decay to bound excited states of ¹⁹ O
Table 37 Summary of beta decay strength to individual bound levels in ¹⁹ O from the decay of ¹⁹ N
Table 38 Summary of observed delayed-neutron transitions from ¹⁹ N by energy96
Table 39 Gamma rays characteristic of de-excitation in ¹⁸ O. The half-lives of the gamma rays are listed along with the intensity of the lines assuming the ¹⁹ N parent97
Table 40 Values for the (β+n) feeding strength to each of the pertinent ¹⁸ O excited states98
Table 41 Neutrons in coincidence with selected ¹⁸ O gamma rays. Listed are those neutrons that may be in coincidence on the bases of the gated spectrum99
Table 42 Cross table for the assignment of intermediate states of ¹⁹ O populated during the decay of ¹⁹ N
Table 43 Gamma-ray transitions identified as being a result of ²⁰ N beta decay feeding bound excited states of ²⁰ O
Table 44 Bound excited levels of ²⁰ O populated during the decay of ²⁰ N. Listed are the branching ratios for the beta transitions as determined from the measured gamma-ray data
Table 45 Energy and branching ratio data for delayed neutrons observed during the decay of ²⁰ N. Also given is the P _n value for delayed neutrons emitted above 1.1 MeV in energy
Table 46 Calculated log(ft) and B(GT) values for the possible beta transitions in the decay of ¹⁹ N. The transition probability is the branching ratio divided by 100109

Table 47 Comparison of the ¹⁹ O energy levels from this experiment with made using the shell model. The subscripts on the energies indicate the cho ¹⁹ O level due to the uncertainty in the neutron decay	ices for the
Table 48 Calculated log(ft) and B(GT) values for specific decays assigned to of ²⁰ N. The transition identifier segregates beta decay to bound states from neutron emitting level	he delayed
Table 49 Summary of transitions observed during the decay of ¹⁹ N that definitively assigned to intermediate levels in ¹⁹ O	
Table 50 Summary of transitions observed during the decay of ²⁰ N that definitively assigned to intermediate levels in ²⁰ O	cannot be

CHAPTER 1

INTRODUCTION

1.1 GENERAL INTRODUCTION

The advent of radioactive ion beam technologies has opened numerous areas of inquiry for practitioners of nuclear science. Until recently, investigation of the atomic nucleus and its properties had been restricted to the 300 or so nuclides in the valley of β-stability as opposed to the some 7000 nuclides that are predicted to be particle bound. The new experimental tools that permit the production and isolation of radioactive species have empowered theorists and experimentalists to engage in studies that would not be possible in their absence. Among the many fields that have been broached in the near past are studies of nuclear structure, nuclear reactions, the nature of astrophysical cycles, and the cosmological processes postulated to have initiated the universe.

The degree to which we understand the nucleus is often measured by the ability of theoretical models to predict fundamental properties such as isotopic masses, half-lives, β -decay systematics, delayed particle emission, and reaction strengths. Structure models, such as the shell model, are continually tested and refined by experimental data that restrict or fix the input parameters; and models of stellar and cosmological nucleosynthesis are similarly affected by both decay and reaction measurements. What follows is a brief summary of the importance of studying the decay of ¹⁹N and ²⁰N by experiment for their ability to illuminate specific aspects of nucleosynthesis and their

relevance with regard to the predictions and refinement of the structure models in this mass region.

1.1.1 STRUCTURE

Measuring the beta decay properties of a nuclide provides information about the quantum ground state of that isotope and states in the daughter that are populated during the decay. Though the energy spectrum of the emitted beta particles is continuous, observation of delayed particles (gamma rays, protons, neutrons, alphas) whose spectra are discreet typically permits quantitative determination of the daughter level fed by the beta decay.

If the relative energy of the parent and daughter ground states are known then the energy of the populated level can be deduced from the secondary particle. This then allows for the Q-value of the beta decay to be calculated by difference. Further, if the strength of the specific transition, also known as the branching ratio, can be measured quantitatively then the energy of the transition coupled with the strength of the transition can be utilized to calculate the log(ft) value which is the logarithm of the partial half-life times a spectral correction factor and proportional to the nuclear matrix element. Knowing the log(ft) value facilitates the assignment, or at least a restriction, on the spin and parity of the intermediate level in the daughter that was fed by the beta transition.

The most salient features predicted by models of nuclear structure, particularly the shell model, are the energy and quantum properties of excited states in nuclei. Although there are other properties, the spins, parities and energies are the most easily tested by experiment and, for the most part, are quantities upon which other inferred

properties are based. The utility of the shell model is that the model can be applied to systems that may not be accessible to experimental inquiry. In order to inspire confidence in such extrapolation, it is necessary for the model to be proven valid in systems that are testable by experiment.

For the specific cases of studying ¹⁹N and ²⁰N, the decay measurements will provide insight into the structure of ¹⁹O and ²⁰O respectively. By restricting the spin and parity of the excited levels populated during the decay, the predictions of the shell model for these nuclides will be tested. Since these isotopes are relatively light the shell model should accurately describe the properties in this region prior to extrapolation to more exotic species.

1.1.2 NUCLEOSYNTHESIS

The generation of elements from the basic nuclear constituents is believed to have occurred during the early stages of the so-called big bang and within the interior of stellar objects — which is an ongoing process. The process is driven by the fusion of seed nuclei with the simplest objects; neutrons, protons and in some cases, alphas. Many of these reactions and their cross sections are known for the fusion of stable isotopes with the lighter particles, however, most of the reaction cross sections for capture by unstable species have not been measured.

To make the network calculations for the production of elements, and hence the isotopic abundances, it is necessary to know or estimate the reaction cross sections for many radioactive species with the simple species. This is true of both the primordial and stellar nucleosynthesis networks. Although it would be desirable to measure the reaction cross sections for all nuclei with the low mass subset, such voluminous,

comprehensive assessments are not currently required. Typically, specific reactions are targeted as being particularly important to measure experimentally while other, less perturbing reaction rates may be estimated systematically. One reaction cross section that has been propounded as important for both inhomogeneous models of the big bang (IMB) and stellar nucleosynthesis is the neutron capture on ¹⁸O producing ¹⁹O and gamma rays.

The standard model of the big bang predicts that elements beyond lithium will not be produced with any appreciable strength. The IMB model, where the number of neutrons is larger than protons, predicts that elements up to and beyond mass (A=20) may be generated [94Ra]. The production of elements beyond neon in this model proceeds through the generation of 18 O via the 14 C(α , γ) and 17 O(n, γ) reactions. Here mass may either be recycled through the 18 O(p, α) 15 N reaction or mass may be built up through the 18 O(n, γ) 19 O reaction. For material to be processed out of the CNO range into the mass (A=20) region, the strength of the 18 O(n, γ) reaction must exceed that of the 18 O(n, α) reaction [96Me]. Thus, 18 O represents a potential end-point for the accumulation of mass in the IMB scenario; rendering the 18 O(n, γ) 19 O a quantity of primary importance to measure.

With regard to network calculations in astrophysics: the ¹⁸O seed nuclide is utilized in the mass build-up occurring in the helium core burning stage of asymptotic giant branch stars. In these objects, ¹⁸O is manufactured from alpha capture on ¹⁴N and processed through to ²²Ne by the ¹⁸O(α , γ) reaction. These stars are propounded to be the main locales for the slow neutron capture process (s-process); where the source of the neutrons is the ²²Ne(α ,n) reaction. If the ¹⁸O(n, γ) reaction strength is relatively

high, the neutron production (and thus s-process rates) will be dramatically altered. The reason being two-fold: first, if the $^{18}O(n,\gamma)$ reaction competes well with the $^{18}O(\alpha,\gamma)$ then neutron production will be quenched by removing the ^{22}Ne seed from the $^{22}Ne(\alpha,n)$ reaction, and second, a strong $^{18}O(n,\gamma)$ will remove neutron material from the system that would otherwise initiate the s-process on different seed nuclei. Consequently, an experimental insight into the $^{18}O(n,\gamma)$ reaction is of critical importance as an input parameter for the network calculations of this stellar environment.

Information may be gained about the $^{18}O(n,\gamma)^{19}O$ reaction by measuring the decay properties of ^{19}N . By measuring the beta branching strength of ^{19}N to levels in ^{19}O and calculating the log(ft) values, the spins and parities of the populated levels can be restricted. From these level properties, at least those levels above the neutron emission threshold, the resonant neutron capture strength of the $^{18}O(n,\gamma)$ reaction can be calculated.

1.2 LITERATURE SEARCH - ¹⁹N, ²⁰N

Numerous publications appear in the literature reporting experimental studies of the decay modes of ¹⁹N and ²⁰N. In fact, there have been half-life studies, measurements of the total neutron emission probability, and an experiment designed to record the γ-ray spectra following the decay of ¹⁹N. However, rather than clarifying the decay properties of these nuclides, the published quantities are disparate - rendering the inferred decay modes somewhat uncertain. What follows is an overview of the published experimental studies relevant to the decay properties of these neutron rich nuclides.

1.2.1 PRIOR MEASUREMENTS OF ¹⁹N DECAY PROPERTIES

The energy level scheme of the mass A=19 system pertinent to the decay of 19 N is shown in figure 1 [95Ti] with the level energies given in MeV relative to the ground state of 19 O. The scheme indicates that the β -decay Q-value is about 12.5 MeV (see also [93Au]), while the maximum energy available for β -delayed neutron emission is about 8.5 MeV. In addition, β -delayed alpha particle emission is energetically possible by about 3.5 MeV and, although this particular channel has not been observed experimentally, similar decay channels have been observed during the decay of other nitrogen isotopes [93Bu, 94Do].

There are eight known levels in the ^{19}O daughter above the ground state that are bound with respect to neutron emission. Therefore, any beta decay feeding to these levels will result in the emission of one or more de-excitation γ -rays. What is more, there are some forty known levels in ^{19}O above the neutron emission threshold through which β -delayed neutron emission is, at least, energetically possible. Given that there is so much energy available for β -delayed neutron emission and that there are so many levels through which such a transition may occur, it should not be surprising that β -delayed neutron emission comprises a significant fraction of the decay strength of this nuclide.

Experimental study of the decay of ¹⁹N began at GANIL in 1986 with an experiment by Dufour *et al.* where β -delayed γ -ray spectroscopy was performed [86Du]. The results of the study found the half-life of ¹⁹N to be 0.32(10) seconds and they were also able to identify three γ -ray transitions which they ascribed to levels in ¹⁹O. The characteristic energies of the three gamma lines were found to be 96.0, 709.2,

12.54					
¹⁹ N			11.58 ()		
IV	11.25 ()				
			10.66 7/2-		
	10.21 7/2-				
	9.93 ()		9.98 () 9.9 7/2-		
	9.6 7/2- 9.43 ()		9.56 ()		
	9.253 ()		9.324 () 9.064 ()		8.996
	9.022 () 8.916 ()		8.923 ()		45
	8.561 ()		8.591 () 8.45 ()		$^{15}\text{C} + \alpha$
	8.247 () 8.048 ()		8.132 ()		
	0.0 10 ()				
			7.508 ()		
	7.242 () 6.98 8 ()		7.118 ()		
			6.903 ()		
	6.583 () 6.4058 ()		6.4662 (7/2)		
	6.1916 ()		6.2693 7/2- 6.1169 3/2+		
	5.7046 7/2+				
	5.5035 ()		5.54 3/2+ 5.384 (9/2)		
	5.1484 (5/2+) 5.007 (3/2)		5.082 1/2- 4.9683 (5/2)		
	4.7026 5/2+		4.582 3/2-		
	4.4025 (3/2)		4.3281 (3/2)		
3.957	4.1093 3/2+		3.9449 3/2-		
¹⁸ ∩ ⊥ n					
™O + n	3.2316 (1/2-)		7.1575 5 /0 .		
	3.0674 (3/2+)		3.1535 5/2+		
			2.779 7/2+		
	2.3715 9/2+				
			1.4717 1/2+		
			7 - .		
	0.096 3/2+		0 5/2+	←	19
			0 3/27	•	

Figure 1 Energy level diagram of the mass 19 system pertinent to the decay of ¹⁹N.

and 3137.8 keV with relative intensities of 100(10), 63(21), and 76(21), respectively. The energies imply a cascade from the 3.9449 MeV level through the 3.2316 and 0.0960 MeV levels, however, the errors on the intensities confound any serious attempt to determine specific branching strengths to individual levels in the ¹⁹O daughter.

Indeed, the Dufour result is in itself an enigma because other studies [70Fi,71Hi] where the 3.9449 MeV level has been populated (through transfer reactions) suggest that this state relaxes to the ground state, 0.0960 and 1.4717 MeV excited states with approximately the same probability [95Ti]. The text of Dufour *et al.* indicates that their experiment was sensitive to transitions in this energy range - thus the absence of these lines in their report is puzzling.

There have been two experiments conducted in the past to measure the total neutron emission probability (P_n) of ¹⁹N. The first was performed at GANIL by Mueller *et al.* [88Mu], and the second by Reeder and collaborators at LAMPF [91Re]. The GANIL study re-measured the half-life to be 0.210(+200 –100) seconds and found P_n to be 0.33 (+34 -11), while the Reeder group determined the half-life to be 0.329(19) seconds and the total neutron emission probability to be 0.624(26) of the decay strength. Though the two measurements of the P_n value agree within error, the Mueller result has a large uncertainty and, what is more, the Reeder value has since been retracted by private communication [97Gi] and in conference proceedings [94Re]. The latter report gives an updated analysis and reports a half-life of 0.255(10) seconds, and the revised P_n as 0.487(21) of the total decay strength. Consequently, the only remaining refereed value in the literature for the P_n strength of ¹⁹N is that reported by Mueller.

To date there has been no experiment to make a spectroscopic measurement of delayed neutrons emitted by ^{19}N . The only spectroscopic evidence appears in a publication by Ozawa and collaborators in 1995 [95Oz] where ^{19}N appears as a contaminant in their experiment designed to measure the delayed neutron spectrum of ^{19}C . They were able to attribute one peak in the neutron TOF spectrum as being due to the decay of ^{19}N . This neutron appeared at an energy of 0.59 MeV in the lab and was attributed to the ^{19}N β -decay populating the 4.582 MeV state in ^{19}O . Their interpretation was based on the change in intensity of this peak as a function of time relative to the other observed neutrons.

1.2.2 PRIOR MEASUREMENTS OF ²⁰N DECAY PROPERTIES

The data in the literature pertinent to the decay of 20 N is less by comparison to 19 N. The portion of the mass (A=20) system for the decay of 20 N is shown in figure 2 with the level energies given in MeV relative to the ground state of 20 O. There have been two measurements of the half-life and two measurements of the P_n value of its decay. Once again, the work was done by the Mueller and the Reeder collaborations [88Mu, 91Re]. The Mueller result for the half-life is 0.100(+30-20) seconds and the P_n is 0.53(+11-7), while the Reeder publication of 1991 lists the half-life as 0.142(19) seconds and the total neutron emission probability as 0.661(50) as a fraction of the total decay strength. Again, the Reeder values were updated in 1995 [94Re] and the new values are given as 0.129(8) for the half-life and 0.520(33) for the P_n value. There is no data available with respect to γ -ray emission following the β -decay, and no publication involving a spectroscopic measurement of delayed neutrons emitted by 20 N.

1.2.3 LITERATURE SEARCH SUMMARY

In total, there have been six half-life measurements of ^{19}N . However those with the better statistical accuracy do not agree to within two standard deviations. The value of the P_n strength that remains in the literature is largely uncertain, and the delayed γ -ray data do not agree with other, related measurements on the excited states of ^{19}O . The ^{20}N experiments have better agreement in half-life and P_n value but independent confirmation of the Mueller result [88Mu] is still desirable. There is no spectroscopic or branching strength data available for ^{20}N regarding beta, delayed neutron or γ -ray emission. A summary of the experimental data currently in the literature, regarding the decay of ^{19}N and ^{20}N , is given in table 1.

Table 1 Literature search for experimental measurement of the ¹⁹N and ²⁰N decay.

QUANTITY	¹⁹ N	²⁰ N	REF.
Q _β (MeV)	12.54	17.97	93Au
Half-Life (ms)	329 (19)	142 (19)	91Re
	210 (+200)(-100)	100 (+30)(-20)	88Mu
	235 (32)	-	88Sa
	300 (80)	-	88Du
	320 (100)	-	86Du
	255 (10)	129 (8)	94Re
Neutron Emission	62.4 (26)	66.1 (50)	91 R e
Probability (%)	33 (+34)(-11)	53 (+11)(-7)	88Mu
	48.7 (21)	52.0 (33)	94Re
Decay γ-Rays (keV)	96, 709, 3138	N/A	86Du

Figure 2 Energy level diagram of the mass 20 system relevant to the decay of ²⁰N.

1.3 SHELL MODEL PREDICTIONS FOR THE DECAY OF 19N

In 1988, following the appearance in the literature of the Dufour experimental results [88Du], a shell model study of the 19 N decay was published by Warburton [88Wa]. Essentially, the paper included calculations of the B(GT) strength for each of the allowed and first forbidden transitions to levels in 19 O. In this model the ground-state spin and parity of 19 N was assumed to be (1/2-), which is consistent with the known spin and parity of the other odd-mass nitrogen isotopes, and the program OXBASH was used to determine the spin, parity and energy of the 19 O levels. The salient features of the shell model results are as follows. First, only about 7% [6%] of the beta strength populates γ -emitting allowed [first forbidden] final states in 19 O. Second, that the fraction of the decay leading to neutron emission should be about 0.87 (or 87%) of the total decay strength. Third, that the half-life of 19 N should be about 0.54 seconds. And fourth, that less than 5% of the allowed strength populates levels in 19 O above the (15 C + α) threshold.

Taking the theoretical and experimental results together, it may be concluded that the calculated half-life over-predicts the observed half-life by about a factor of two, and that the total β -delayed neutron emission probability, P_n , is predicted to be significantly higher than that which has been measured. Also, the model predicts significant β -strength to bound levels that would yield (either directly or through cascade) the 1376 keV γ -ray characteristic of the transition between the 1472 and 96 keV levels in ^{19}O – a transition that was not observed in the Dufour experiment [88Du].

Warburton comments that, if the calculated B(GT) values to the allowed states under-predict the actual values by and order of magnitude, the missing transition

strength would by reduced to a degree where the experimental study [88Du] may not have been sensitive to it. Increasing the B(GT) values in this way would also bring the model half-life in line with experiment, however, the P_n value under those conditions would be reduced to about 0.3 or 30% of the total branching strength. Indeed, Warburton summarizes that the predicted P_n value is very large given the energy range and number of bound states available in ^{19}O , and suggests that experimental measurement of the total neutron and γ -ray emission probabilities may be the only way to measure the absolute β -strengths to low lying levels in ^{19}O .

1.4 EXPERIMENTAL GOALS

In light of the shortcomings and potential errors in the measured values of the $^{19.20}$ N decay modes and the general disagreement between experiment and theory in the case of 19 N, an experiment to re-measure the decay properties of 19 N and 20 N was proposed at the National Superconducting Cyclotron Laboratory. The goals of the experiment were to thoroughly establish the decay properties of these nuclides by making spectroscopic measurements of the delayed neutrons and γ -rays emitted during their decay. From these measurements, the predictions of the shell model may be compared and contrasted with experiment, and the importance of the 18 O(n, γ) reaction in both the IMB and stellar nucleosynthesis networks can be estimated.

CHAPTER 2

EXPERIMENTAL SETUP

2.1 PRODUCTION OF SECONDARY RADIOACTIVE ION BEAMS

The only naturally occurring sources of the isotopes of interest are the hot, dense environments associated with stellar objects. When this is taken together with the fact that the half-lives of ¹⁹N and ²⁰N are reportedly less than one second [95Ti], it should not be surprising that there are no terrestrial samples of these nuclides upon which experiments may be performed. As a result, a facility capable of artificially generating these isotopes was required.

One facility capable of producing ¹⁹N and ²⁰N in yields commensurate with the experimental goals is the National Superconducting Cyclotron Laboratory at Michigan State University. In this study, the necessary radioactive ion beams (RIBs) were generated by fragmenting an intense primary beam of ²²Ne in a beryllium target. The primary beam of ²²Ne was accelerated in the K1200 Cyclotron and extracted at an energy of 80 MeV/A. The ²²Ne beam then struck a thin beryllium foil (564mg/cm²) resulting in the production of a wide variety of secondary fragments near or below mass (A=22).

The nuclides of interest were selected, sequentially, using the A1200 fragment mass analyzer [91Sh]. The fragmentation products were separated on the basis of their mass to charge ratio (A/Z) by the first set of A1200 dipoles. The specific nuclides were

further separated from other reaction products by passing them through an achromatic wedge (241 mg/cm² Al equivalent) and the second pair of A1200 dipoles. The wedge induced momentum shifts, based on the atomic charge of the recoil products, allowing unambiguous separation among nuclides at the final focus of the A1200.

Once purified, the RIBs were transported, using various electromagnetic devices, to the low background experimental end-station located in the N3 vault at the NSCL complex. Generally speaking, the transmission of the secondary beam from the A1200 focus to the experimental end-station was about 80 percent over the course of the study. For this experiment, the primary beam was pulsed to allow for efficient and conclusive data recording. The beam-on and beam-off intervals were selected on the basis of the half-life of the nuclide being transported to the experimental end-station. The beam pulsing sequences for each nuclide studied are given in table 2.

Table 2 Beam-On and Beam-Off pulsing times for each nuclide investigated.

Nuclide	Experimental Runs	Beam-On Time	Beam-Off Time
17 _N	17 - 23	8 s	12 s
¹⁶ C	46 - 50	1.5 s	3 s
¹⁹ N	59 - 90	1 s	3 s
²⁰ N	93 - 105	0.5 s	0.8 s
²⁰ N	106 - 116	0.375 s	0.375 s

2.2 EXPERIMENTAL END STATION

The radioactive beams of ¹⁶C, ¹⁷N, ¹⁹N, and ²⁰N initially passed through a parallel plate avalanche counter (PPAC) located at the inlet of the 92" chamber and immediately upstream of the main detector array. The PPAC was most useful during the preliminary beam tuning as it provided position information on the incoming beam.

Downstream of the PPAC, the RIBs passed through a thin Zirconium window separating the beam-line vacuum (10⁻⁶ torr) from the end station that was at atmospheric pressure. The detection system consisted of four dependent sub-systems that will be referred to as the implantation system, neutron bar array, ⁶Li array, and Germanium pair.

2.2.1 IMPLANTATION SYSTEM

After transmission through the PPAC, the ion beam entered the first detection sub-system called the implantation system. The implantation system consisted of two transmission mounted, silicon surface barrier detectors, a series of aluminum degrader foils, and a thin plastic scintillator. The silicon detectors were 300 µm thick and had surface areas of 300mm². The scintillator was made of BC412 plastic and had the dimensions of 2.5cm x 2.5cm x 0.3cm deep. The two silicon detectors were placed one on the upstream side, and one on the downstream side of the plastic scintillator. The upstream silicon was referred to as the dE detector, the downstream silicon as the VETO detector, and the plastic scintillator was referred to as the START detector. The components of the implantation system and the detector designations are shown in Figure 3.

During the beam-on intervals, the beam passed first through the dE detector, was degraded in the aluminum foils, and then implanted into the START scintillator. The dE detector provided a useful real-time monitor of the purity of the secondary beam, while the dE detector in concert with the START detector and the VETO detector provided data on the fraction and purity of the secondary beam actually implanted within the scintillator.

During the beam-off intervals, the scintillator generated a signal caused by transmitted beta particles from the decay of the implanted species. The coincident signals from the photomultiplier tubes (PMT's) were used for half-life measurements, the determination of the total number of decay events, and also provided a start signal for the delayed neutron time-of-flight spectroscopy.

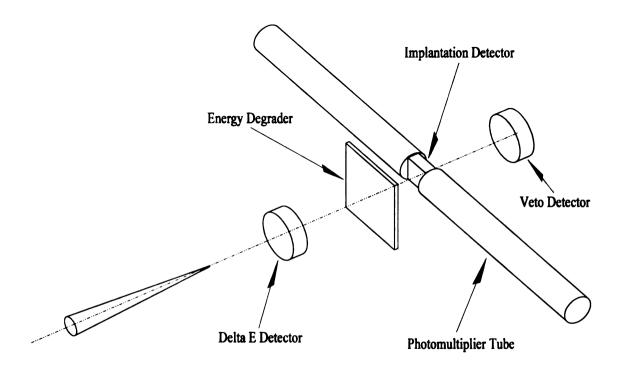


Figure 3 Schematic diagram of the implantation detection system.

2.2.2 NEUTRON BAR ARRAY

The second detector system utilized in this study was the so-called Neutron Bar Array. The neutron bar array was originally constructed by Dr. Richard Harkewicz during the period of his graduate study at Michigan State University [92Ha]. This array consists of sixteen plastic scintillator bars, composed of BC412 plastic, with the approximate dimensions 157cm x 7.6cm x 2.54cm thick. The scintillator bars were

shaped such that the radius of curvature of every bar is equal to one meter. The sixteen neutron bars are housed in an aluminum frame with the bars arranged such that they are concentric with a single point in space. At this point, the START detector is situated to enable a time-of-flight technique for the determination of delayed neutron energies.

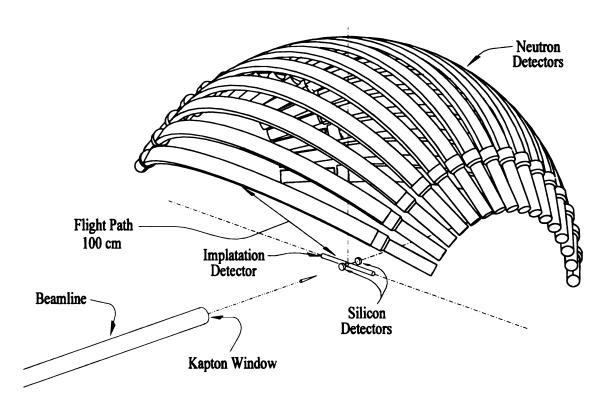


Figure 4 Schematic diagram of the Neutron Bar Array.

The beta decay in the implantation scintillator provides a common start while a neutron event detected within an array bar provides the individual stop. The time difference between the events yields the time-of-flight and, knowing the flight path, the neutron energy. The neutron bar array was selected for its ability to efficiently detect delayed neutrons in the energy range between one MeV and ten MeV. A schematic diagram of the neutron bar array is shown in Figure 4.

2.2.3 ⁶Li GLASS SCINTILLATORS

To detect beta-delayed neutrons emitted with energies lower than the threshold of the neutron bar array, the detector end-station contained three inorganic scintillators containing enriched quantities of ⁶Li. The detectors were right-cylinders with radii equal to 2.54 cm. Two of the three were 2.54 cm deep while the depth of the third crystal was equal to 1 cm. The three detectors were placed about the START scintillator without shadowing the bars of the array, and with as much separation as possible to minimize scattering among the devices. The two thicker ⁶Li glass scintillators were placed about 25cm from the source activity while the thinnest detector was situated about 10cm from the source activity.

The ⁶Li glass scintillators also provided spectroscopic information on delayed neutron emission using a TOF technique. Again, the implantation scintillator provided a start signal from the beta decay and the detection of a neutron by a ⁶Li glass scintillator gave an individual stop signal.

2.2.4 HIGH PURITY GERMANIUM DETECTORS

To detect gamma rays emitted during the decay of the isotopes of interest, the detector end-station included two, high purity germanium crystals. The detectors were 120% and 80% efficient relative to a standard sodium iodide detector and were placed as close as possible to the source activity given the physical constraints of the neutron bar array and its supporting apparatus. The detectors were situated upstream of the source activity to minimize the effect of the opening angle with respect to the beampipe. The angle between the detectors and the implantation position was about eighty degrees. The more efficient (Ge#1) and the less efficient (Ge#2) were placed at about

17cm and 22cm away from the source activity, respectively. Figure 5 is a schematic representation of the end station showing the implantation system, the neutron bar array, ⁶Li detectors and germanium detectors.

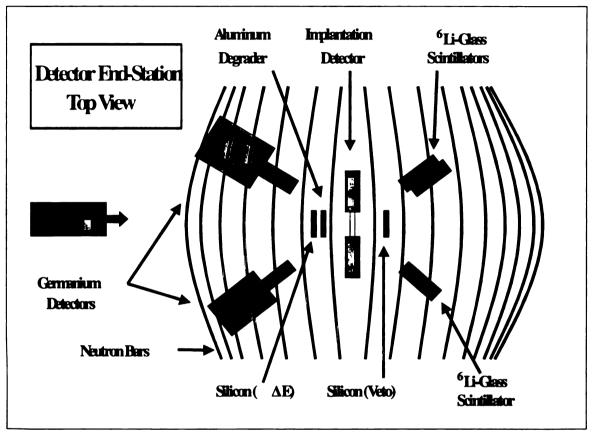


Figure 5 Schematic diagram of the entire detector end-station (top view).

2.3 ELECTRONICS AND DATA ACQUISTION

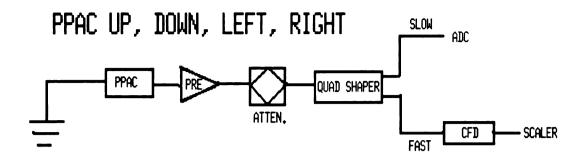
Data taking for the experiment employed standard electronics modules available within the NSCL electronics pool for all systems including the CAMAC portion. The beam pulsing signals were supplied by a programmable dual gate generator where a TTL signal was input to the cyclotron RF control. The phase of one of the RF electrodes was shifted by this signal and the beam was cut-off in the accelerator.

For both the beam-on and beam-off intervals, a master gate was caused by a coincident event in the two PMTs attached to the implantation scintillator. This event in conjunction with the beam-status bit (on or off) triggered read instructions for the electronics for the detectors of interest. The neutron detectors were disabled during the beam-on period, as was the real-time clock, however, the germanium detectors were enabled during both the beam-on and beam-off intervals. Figures 6-8 show the electronics set-up utilized in this experiment.

The data were recorded on an event-by-event basis and written to 8mm tape for subsequent offline analysis. The in-house SARA analysis code was utilized for both online and offline purposes along with the associated XAMINE display package. The data taking was divided into runs of approximately two hours duration to ensure manageability of the files during analysis.

2.4 SECONDARY BEAM PURITY

In this experiment, four radioactive ion beams were generated by the fragmentation of a ²²Ne beam in a beryllium foil. Two of the beams were produced for calibration purposes (¹⁶C and ¹⁷N) and two of the beams were the isotopes of interest for the experiment (¹⁹N and ²⁰N). The purity of the secondary beams was established at two points. The first was during beam tuning through the A1200 fragment mass analyzer (which is independent of the user defined set-up in the experimental vaults) and the second was by analysis of the data recorded downstream by the implantation system at the end-station.



BEAM PULSING AND REAL-TIME CLOCK

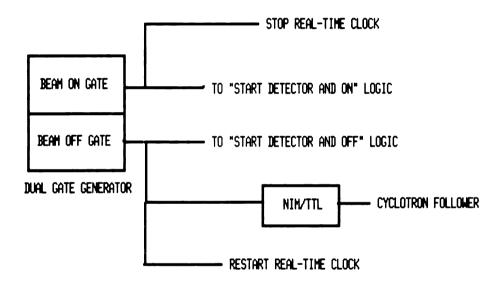
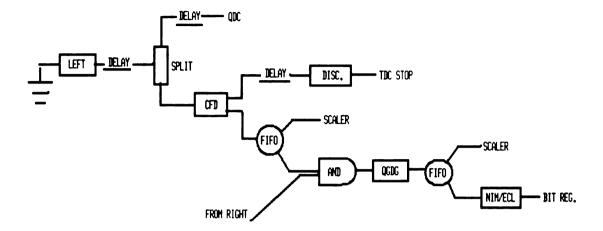
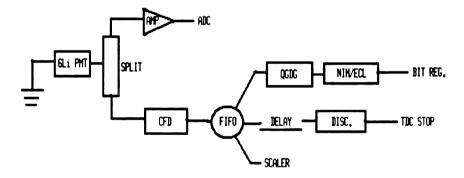


Figure 6 Electronics diagrams of the PPAC, Real-Time clock and RF pulsing control elements of the end-station.

Neutron Bars



6Li Scintillators



GERMANIUM DETECTORS

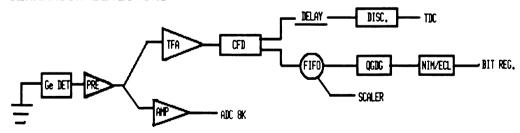


Figure 7 Electronics diagrams for the neutron bars, ⁶Li detectors and germanium detectors.

START DETECTOR

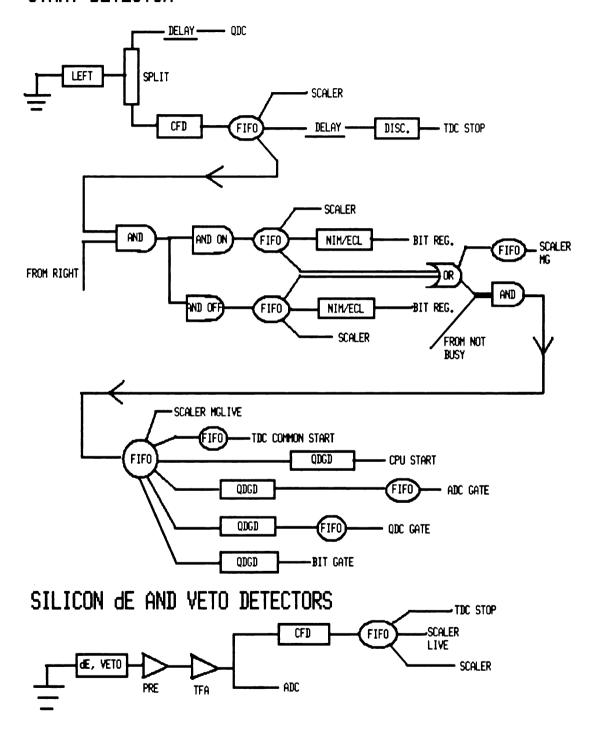


Figure 8 Electronics diagrams for the components of the Implantation system.

During beam tuning, the purity of the secondary product of interest is determined by monitoring both the fragment TOF through the separator, and the energy loss of the fragments in two silicon PIN detectors (typically 300 microns thick). The optimum settings are selected for variables such as the momentum acceptance of the spectrometer, the width and breadth of the momentum slits, the thickness of the achromatic wedge, the thickness of the production target and, occasionally, the composition of the target itself. Prior to transmitting the secondary beam to the desired experimental vault, a purity run is performed and the data stored for later reference.

For this experiment, the secondary beams were also monitored for purity at the end-station using the implantation system. The yield and composition of the secondary beam entering the vault is determined by the dE detector and the RF time-of-flight from the cyclotron while the yield and composition of the beam actually implanted into the start detector is determined from the dE vs. RF time-of-flight vetoed by the second silicon detector located downstream of the implantation scintillator. The results of the purity analysis for both the spectrometer analysis and end-station response is shown in table 3 for the four secondary beams of 16 C, 17 N, 19 N, and 20 N.

Table 3 Beam purity at the A1200 focal plane and the experimental end-station for those secondary nuclides isolated in this experiment.

SECONDARY	A1200	IMPLANTED	CONTAMINANT	FRACTION
BEAM	% PURE	% PURE		%
¹⁶ C	99.3 (9)	99.70(6)	¹⁹ N, ¹⁸ N	0.7
¹⁷ N	>99.9	99.75(4)	N/A	N/A
¹⁹ N	99.5(10)	99.5(1)	¹⁷ C	0.5
20 N	98.8(17)	94.4(21)	¹⁸ C	1.1

From the data collected it is clear that the calibration beams of ¹⁶C and ¹⁷N, as well as the ¹⁹N nuclide of interest, are better than 99% pure, while the secondary ²⁰N beam is about 95% pure. The lower purity of the ²⁰N beam is not unexpected due to the nature of the process through which ²⁰N is produced from a primary beam of ²²Ne. The reaction required a nucleon transfer or a charge exchange with the target material and hence the momentum distribution of this particular isotope covers a larger range. As a consequence, there is a greater probability of momentum overlap with other secondary products and thus the purity is reduced relative to the fragmentation products.

The implantation rate taken together with the scaler values for the dead-time during the beam-on and beam-off intervals permits a determination of the total number of decays for each of the isotopes of interest that would have been observable during the beam-off period. The results of these calculations are given later in table 4 in comparison to the more rigorous approach described in section 3.4.2.

CHAPTER 3

EXPERIMENTAL RESULTS

3.1 OVERVIEW OF ANALYSIS PROCEDURE

The analysis of the experimental data can be grouped into three parts. The first is the analysis of the beta particle emission leading to evaluations of the half-lives and to the total number of decays of the isotopes of interest (N_{β}) . The second is with respect to delayed neutron emission by analyzing the data recorded in the neutron bar array and the ⁶Li inorganic scintillators to determine the delayed-neutron energies and intensities. The third is an analysis of the gamma-rays emitted during the decays as recorded by the germanium detectors.

In this chapter, the information that can be gleaned strictly from the experimentally recorded data is presented. What follows is a detailed analysis of the methods by which the efficiencies of the detection systems were established, the calculations for the intensities of the delayed-neutron and gamma-ray lines, the determination of the decay half-lives of ¹⁹N and ²⁰N, and the calculations for the total number of decays observed for each of the isotopes. The order of presentation is first, the beta decay, second the delayed-neutron emission and third, the gamma-ray analysis.

In Chapter 4, the experimental results are interpreted to yield the decay schemes of ¹⁹N and ²⁰N. To do so, the gamma-ray data and delayed neutron results are viewed in concert with the idea of accounting for gamma-ray cascades and to consider the possibility of delayed neutron and gamma coincidences.

3.2 HALF-LIFE MEASUREMENT OF 19N

The first priority in the experimental study of ¹⁹N was to determine the half-life of the nuclide. Due to the high secondary yield of ¹⁹N the data acquisition experienced a high degree of dead time. To minimize the contribution of this effect to the half-life determination, the primary ²²Ne beam was attenuated by a factor of about 30 during the run in which the half-life was measured.

The beta decay of ¹⁹N populates states either in ¹⁹O or ¹⁸O depending on whether the exit channel is pure beta or delayed neutron emission. As such, the half-life fit to the recorded data must include a term for the decay of ¹⁹O. Fortunately, the half-life of ¹⁹O (26.9 seconds) is well established [95Ti] and is also roughly two orders of magnitude longer than the ¹⁹N parent. This large difference, taken together with the fact that the total time of the half-life run was much longer than either half-life (~2 hours), permits a simplifying assumption where the two decays can be treated as independent quantities and the feeding term characteristic of secular decay modes may be eliminated.

The time sequence for the pulsing of the primary beam during the half-life run was set such that the beam-on period (implantation) was one second and the beam-off period (data acquisition) was three seconds. The half-life determination was made on the basis of the spectrum collected as a function of the real-time clock (Ortec model RC014) during the beam-off periods. The full-scale time of the clock was set to sixteen seconds.

The delayed neutron daughter, ¹⁸O is stable and therefore a secular component for this decay is not necessary. Figure 9 shows the half-life fit to the data collected in

the implantation scintillator for the decay of ^{19}N . Shown are the raw data (black), the ^{19}N component (blue), the background due to ^{19}O (black), and the aggregate fit to the spectrum (red). The result of the fit indicates that the half-life of ^{19}N is $299 \pm 3 \pm 16$ milliseconds. This result is in close agreement with references [88Du, 91Re].

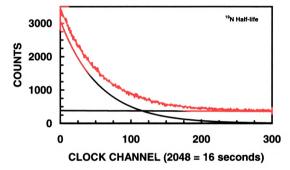


Figure 9 Fit to the half-life spectrum of 19N.

3.3 HALF-LIFE MEASUREMENT OF ²⁰N

The production rate of ²⁰N from the primary ²²Ne beam was orders of magnitude lower than that realized in the ¹⁹N experiment. As a result, there was little or no dead time associated with monitoring the decay of ²⁰N. The half-life fit included a term for the decay of both ²⁰O and ²⁰F, the daughter and granddaughter, but once again these half-lives are orders of magnitude longer than that of ²⁰N – permitting the same simplifying assumption as with the ¹⁹N case.

During the ²⁰N runs, the beam-off time (where the real-time clock is active) was changed from 0.8 seconds to 0.375 seconds to improve the efficiency for recording ²⁰N events (see table 2). Both series, long and short respectively, permit half-life determinations for ²⁰N. Figure 10 shows the fit to the long beam-off data with the ²⁰N component (blue), the sum of the daughter contributions (black) and the aggregate fit to the data (red).

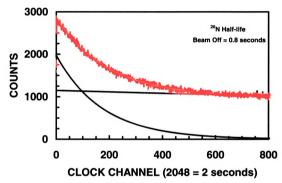


Figure 10 Fit to the half-life spectrum of ²⁰N.

The results of the long and short beam-off time data for the half-life of 20 N are $118 \pm 6 \pm 10$ and $124 \pm 7 \pm 10$ milliseconds, respectively. These results are in reasonably good agreement with the previously reported measurements in the literature [94Re].

3.4 TOTAL NUMBER OF EVENTS

The total number of decay events for each nuclide was determined by the response of the implantation scintillator to beta particles. The total number of events is a necessary quantity to calculate the efficiency of the neutron detectors and to determine the absolute branching ratios for the ¹⁹N and ²⁰N decay channels. The specific procedure used to solve for the total number of decays was to integrate the half-life spectrum collected for each nuclide.

3.4.1 DEAD TIME CONSIDERATIONS

Due to the high yield of secondary fragments, the data acquisition system experienced considerable dead-time. To account for the effect of dead-time analytically, the procedure outlined by Knoll [89Kn] and Leo [94Le] for non-paralyzable dead-time was employed. In this technique, the relationship between the actual and observed activity can be described by the equation:

$$m = n/(1+n\tau)$$
 [1]

Where m is the observed activity, n is the actual activity and τ is a dead-time related constant. It is noteworthy that the parameter, τ , will depend on the properties of the particular nuclide under scrutiny (half-life, implantation rate, daughter properties) thus τ is not general and must be solved for each secondary product of interest.

To elucidate an accurate value for the parameter, τ , subset half-life spectra were generated by applying tight gates around the peaks of the well-separated neutron transitions recorded by the neutron bar array, and reflecting back onto the half-life spectrum. This protocol maximized the signal/background ratio and also ensured that the resultant half-life spectrum was as pure as possible – containing only one radioactive component. The caveat to this technique is that the dead-time formula is no longer strictly applicable. This is due to the fact that τ (though it is a constant over all channels) is dependent on the total instantaneous activity and not on any subset.

However, the fraction of counts missed due to the computer being busy will be a constant at each channel for all subsets of the total half-life spectrum. Mathematically, the value of:

$$1/(1+n\tau)$$
 [2]

will be a different constant at each channel for all gated and ungated half-life spectra. Clearly, n, the actual number of counts at any channel, varies among the gated subsets, thus the solution for τ will vary based on the inverse of n. However, since at channel zero (or any specific channel for that matter) the fraction of counts missed by the computer is the same for all subsets of the total half-life spectrum, the *product* of the value of τ with the *initial* activity will be the same for all subsets. For our purposes, it is more convenient to solve for the activity-times- τ constant (A τ con) at channel zero, and for the initial gated activity for each subset of the total half-life histogram and compare across subsets for reliability.

For the calibration beams of ¹⁶C and ¹⁷N, the protocol was as follows: first, the neutron-gated half-life spectra were fit with MINUIT and MINOS where the decay constant was fixed to the known half-lives and Atcon and the initial activities were allowed to vary freely – as was the background. Second, the total half-life spectrum was fit in MINUIT/MINOS in a similar way except that the value of Atcon was restricted to be within the range defined by the solution of the fits to the gated spectra. Third, from the minimized fit, the total number of decays for each isotope was extracted. For the ¹⁹N and ²⁰N nuclides, this same procedure was followed except that, in addition to fixing the half-lives to the literature or independently determined values, the procedure was repeated in its entirety with the half-lives permitted to vary freely in

the minimization. As an example, figure 11 shows the aggregate delayed neutron spectrum of ¹⁶C along with the neutron gates used to reflect back onto the half-life spectrum.

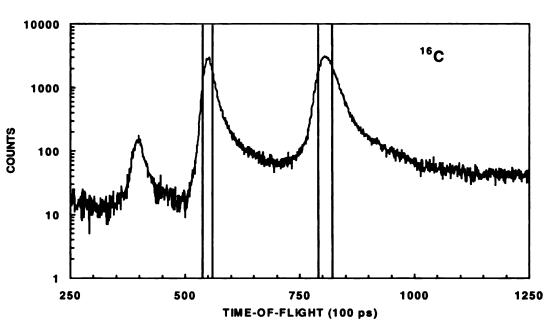


Figure 11 Delayed-neutron spectrum of ¹⁶C showing the gates used to reflect back onto the half-life spectrum for the dead-time analysis.

The gated half-life spectra were fit using the equation given by Knoll [89Kn] and Leo [94Le] where the quantity n was of the form:

$$n = A_1 \operatorname{Exp}[-\lambda_1 \operatorname{CH}] + A_2 \operatorname{Exp}[-\lambda_2 \operatorname{CH}] + \operatorname{constant}$$
 [3]

where λ_1 was fixed to the known decay constant of 16 C in inverse channels, A_1 is the initial activity of 16 C in counts/channel, A_2 and λ_2 were parameters included to allow for the possibility of a second radioactive component, and CH is the channel of the half-life spectrum. In addition to these parameters, the value of the total initial activity times the dead-time parameter τ (A τ con) was allowed to vary freely for each gated spectrum.

Figure 12 shows the neutron-gated half-life spectra and the best-fit curves to those data as determined in MINUIT/MINOS.

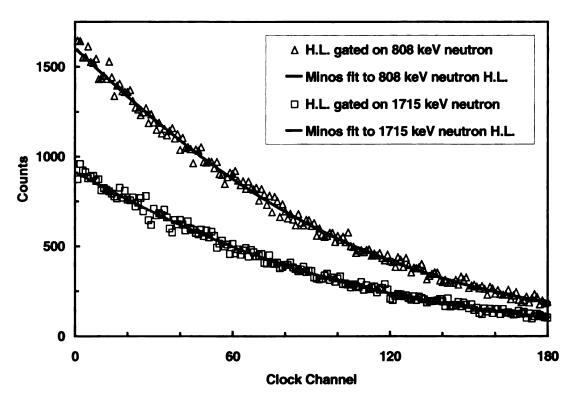


Figure 12 Best fit curves to the neutron-gated half-life spectra of ¹⁶C.

From these fits, the value of Atcon is deduced to be 0.64(5) for the ¹⁶C data, which is the weighted mean of the best-fit results for this parameter in MINUIT/MINOS. This value of Atcon was input into the parameter list for the fit to the total half-life spectrum and permitted to vary between the bounds defined by the one-sigma range. Again, the half-lives of ¹⁶C and the beta daughter ¹⁶N were fixed to their literature values, while the initial activities and a constant background were permitted to vary freely. Figure 13 shows the total half-life spectrum collected for the ¹⁶C calibration runs and the best fit to those data using the procedure outlined above.

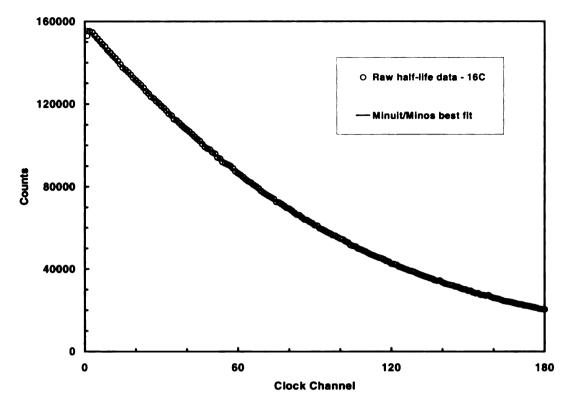


Figure 13 Fit to the full half-life spectrum recorded by the implantation scintillator during the decay of ¹⁶C.

The best-fit value of Atcon was 0.642(5), which is within the range defined by the gated data. Integrating the ¹⁶C component with the dead time included leads to a total number of ¹⁶C decays recorded in the experiment as 1.25(5) x 10⁷ events.

3.4.2 SUMMARY

A similar procedure was followed for all isotopes of interest. In all cases, the reduced chi-squared of the fits was better than 1.15 and the value for Atcon derived from the gated spectra reflected the value obtained for the total half-life data. Table 4 lists the final values for the total number of decays of each of the observed isotopes in this experiment. Also given in table 4 are the results for a determination of the total number of decays of the primary isotopes as calculated from the implantation rates and the scalar dead-times.

Table 4 Total number of decays recorded by the implantation detector for each of the isotopes of interest. Also given is the calculated number of decays from the implantation rate and scalar-measured dead times.

Secondary Beam	Isotope	Total # of Decays	Total # of Decays
		From Analysis	from Implantation
16C	¹⁶ C	1.25(5) x 10 ⁷	1.3(3) x 10 ⁷
¹⁶ C	¹⁶ N	$3.54(10) \times 10^5$	N/A
¹⁷ N	¹⁷ N	$2.3(1) \times 10^7$	$2.4(5) \times 10^7$
¹⁹ N	¹⁹ N	$4.64(10) \times 10^7$	$5.3(12) \times 10^7$
¹⁹ N	¹⁹ O	$4.4(2) \times 10^7$	N/A
²⁰ N	20 N	$7.60(10) \times 10^5$	$7.4(21) \times 10^5$

3.5 NEUTRON BAR ARRAY

The sixteen detectors of the neutron bar array were utilized to determine the energy of delayed neutrons emitted during the decay of the nuclides of interest and also the intensity or strength of the observed branches. Therefore, the individual detectors required calibration for both energy and detection efficiency. Since neutron energies were determined using a time-of-flight technique, the energy calibration is effectively a time calibration for the neutron bars. To calibrate the array in time, two techniques were combined: first, the slope of the responses from the TDC modules were calibrated using a precision time calibrator and second, the known delayed neutron lines of ¹⁶C and ¹⁷N were monitored to deduce the TDC offset. For the efficiency calibration, the decay of ¹⁶C and ¹⁷N provided transitions with known intensities over a range of neutron energies while Monte-Carlo simulations allow for the determination of the

array's efficiency for higher energy neutrons. Each of these techniques is discussed in the following sections.

3.5.1 TIME CALIBRATION OF NEUTRON BAR ARRAY

The energy of neutrons emitted following the beta decay of ¹⁶C, ¹⁷N, ¹⁹N, and ²⁰N were determined using a time-of-flight technique. As a consequence, it was desirable to find the time response of the time-to-digital-converters (TDCs) accurately and independent of the detectors' intrinsic responses. This was accomplished using a precision time calibrator (Ortec model 462) where the calibrator start was split and sent to the implantation detector CFDs, while the calibrator stop was input, sequentially, to the CFD channel of each photomultiplier tube (PMT) in the experimental set-up. The period of the time calibrator was set to ten nanoseconds and the range was set to 320 nanoseconds. These were chosen knowing that the TDC modules' full scales were set by hand to be approximately 200 nanoseconds. Due to the fact that each bar of the neutron array contained two PMTs, the time response of both TDC channels was determined independently. A representative spectrum obtained using the time calibrator is shown in figure 14.

This technique was used to determine the slope of the time response of each photomultiplier tube utilized for time-of-flight measurements. In total there were 32 such PMTs – two for each of the sixteen neutron bars. Table 5 gives the result of the TAC calibrator measurement for the slope of the PMTs.

The slopes determined by this method were incorporated into pseudo parameters within the SARA analysis code. The event time for a detected neutron in one of the

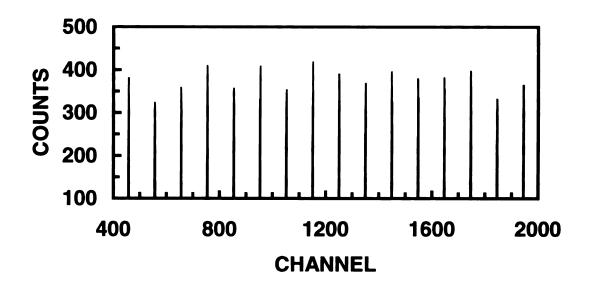


Figure 14 Spectrum collected by one of the neutron array bars from the precision time calibrator.

Table 5 PMT slopes for the neutron bar array as deduced from the precision time calibrator data.

PMT Designation	SLOPE Channels/ns	PMT Designation	SLOPE Channels/ns
BAR 01 R	9.960(4)	BAR 01 L	9.891(2)
BAR 02 R	9.899(5)	BAR 02 L	9.910(3)
BAR 03 R	10.002(1)	BAR 03 L	10.275(2)
BAR 04 R	10.278(3)	BAR 04 L	10.317(2)
BAR 05 R	10.506(2)	BAR 05 L	10.541(3)
BAR 06 R	10.354(3)	BAR 06 L	10.376(2)
BAR 07 R	10.357(4)	BAR 07 L	10.511(5)
BAR 08 R	10.300(4)	BAR 08 L	10.262(4)
BAR 09 R	10.313(2)	BAR 09 L	10.133(2)
BAR 10 R	10.317(3)	BAR 10 L	10.054(4)
BAR 11 R	10.253(3)	BAR 11 L	10.230(2)
BAR 12 R	10.603(5)	BAR 12 L	10.601(3)
BAR 13 R	10.609(4)	BAR 13 L	10.611(5)
BAR 14 R	10.571(3)	BAR 14 L	10.553(4)
BAR 15 R	10.578(4)	BAR 15 L	10.611(3)
BAR 16 R	10.336(4)	BAR 16 L	10.766(4)

array bars was defined as the average TDC channel recorded by the two phototubes of each array bar relative to the start signal of the implantation detector. The incorporation of the slopes into the pseudo code results in a histogram for each bar where the channels reflect a time duration of 100 picoseconds.

From these histograms having slopes of 100ps per channel, the energy/time response of the array bars was determined, on-line, by observing the delayed neutron emission from known sources. The decay of ¹⁶C and ¹⁷N are known to exhibit beta delayed neutron emission. The energies of the delayed neutrons emitted from these sources serve as useful on-line calibration beams for the neutron bar array. Table 6 gives the delayed neutron lines from these sources and the reference from which they are drawn.

Table 6 Calibration delayed neutrons of ¹⁶C and ¹⁷N given by energy and branching ratio.

SOURCE	ENERGY CM keV	ENERGY LAB keV	INTENSITY Percent	REFERENCE
¹⁷ N	407.6	384	38.0 (13)	76Al
¹⁷ N	1245.4	1172	50.1 (13)	76Al
¹⁷ N	1808.2	1702	6.9 (5)	76Al
¹⁶ C	859	808	84.4 (17)	93Ti
¹⁶ C	1829	1715	15.6 (17)	93Ti

Secondary beams of ¹⁶C and ¹⁷N were generated in a manner completely analogous with that described in section 2.1 and monitored under experimental conditions identical to those of the ¹⁹N and ²⁰N runs. The exception being that the beam pulsing sequence was adjusted according to the half-lives of the respective nuclides.

The spectra collected for ¹⁶C and ¹⁷N delayed neutrons were analyzed on a bar-by-bar basis to determine the peaks' centroids as a function of neutron energy.

To determine the centroid of the calibration peaks, the slope-adjusted spectra were fit to functional forms containing peak and background components. The peak shape was approximated by a combination of gaussian and Breit-Wigner functions. Below the centroid the shape of the peak is gaussian while at channels above the centroid the shape is Breit-Wigner. This functional form proved to be the most reliable estimation of the response function of the array detectors. The background term was simply a broad asymmetric gaussian plus a constant to account for random events and neutron back scattering from the end-station environment.

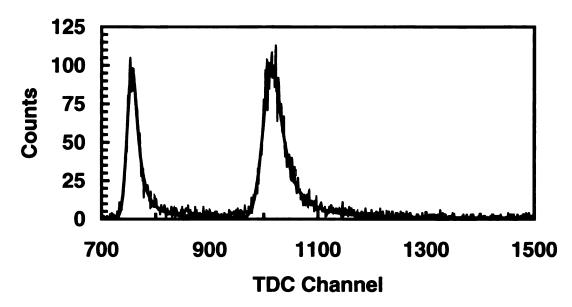


Figure 15 Delayed-neutron spectrum collected by one of the array bars during the decay of ¹⁶C.

As an example, figure 15 shows the delayed neutron spectrum collected in one of the array bars during the decay of ¹⁶C. The fit to the data is indicated in red where the best fit was determined by minimizing the total Chi-squared using the program

MINUIT from the Cern library. Typically, the reduced Chi-squared was between 0.9 and 1.2 for the fits to the individual bars for both the ¹⁶C and ¹⁷N data.

Table 7 gives the results of the fits to all of the calibration neutron peaks.

Tabulated are the centroids of the calibration lines (in TDC channels) as a function of delayed-neutron energy.

Table 7 Centroid channels of the neutron peaks utilized in the calibration of the neutron bar array.

Array Bar	1715 keV peak Centroid	808 keV peak Centroid	1702 keV peak Centroid	1172 keV peak Centroid
1	754.9	1009.8	759.1	874.7
2	753.7	1011.7	757.5	874.6
3	776.8	1034.3	781.6	898.5
4	745.9	1005.6	749.0	864.5
5	814.4	1081.2	819.7	937.9
6	796.6	1060.3	803.0	919.9
7	810.6	1069.0	816.6	931.8
8	756.4	1013.5	760.4	877.5
9	737.1	997.3	740.4	858.5
10	752.1	1009.8	757.5	873.1
11	718.7	975.3	725.5	838.3
12	776.7	1036.2	782.0	897.5
13	748.1	1004.4	751.7	868.0
14	732.5	989.1	735.4	853.4
15	729.1	981.9	732.7	851.4
16	773.3	1033.4	778.1	895.1

Unfortunately the individual array bars were not efficient enough at low energy to detect the 384 keV neutron from the decay of ¹⁷N. However, in addition to the known neutron transitions, the plastic bars were able to record events due to the beta particles.

These peaks provide additional calibration points in the time spectrum of each array detector. Table 8 gives the MINUIT result for the centroid of the prompt beta transition on a bar-by-bar basis.

Table 8 Centroid channels of the prompt beta peaks utilized in the calibration of the neutron bar array.

Array Bar	β-Prompt ¹⁶ C Centroid	β-Prompt ¹⁷ N Centroid
1	234.0	234.6
2	232.6	233.4
3	256.2	257.9
4	226.3	226.6
5	294.0	295.2
6	276.2	277.3
7	292.4	293.0
8	238.7	239.4
9	221.4	220.8
10	233.9	234.5
11	200.2	200.6
12	258.3	258.5
13	229.4	229.7
14	210.2	210.8
15	201.8	202.3
16	247.7	248.7

To calibrate the individual detectors, the neutron and beta energies were converted into velocities with the dimension of metres/(100 picoseconds). The unusual unit was chosen to take advantage of the known 100 picosecond wide bins of the PMT slope corrected histograms. For the neutrons, the velocity was determined from the general equation:

$$E = \frac{1}{2} m\beta^2 c^2$$
 [4]

where E is the laboratory energy listed in table 6. For the neutrons, no relativistic correction was included as the energies were low enough to treat them classically. For the β -particles, the energy used to calculate the velocity was for the most probable energy in the distribution. This was taken to be 0.33 of the β -decay energy - yielding most probable β energies of 1.265 and 1.522 MeV for ¹⁷N and ¹⁶C, respectively. These were deduced by weighting the known decay branches by their normalized probability and summing over the possible transitions. The calculated velocity of the representative β particles, unlike the neutron case, included a relativistic correction.

The inverse of the calculated particle velocity was fit to the observed histogram channel using the general form:

$$1/v = 1/d \cdot (TDCChannel - Offset)$$
 [5]

where v is the velocity (m/100ps), d is the distance between the position of the source activity and the array bar (metres), and the offset is measured in channels or 100ps. Expanding the equation leads to a simple linear relationship:

$$1/v = 1/d \cdot TDCChannel - 1/d \cdot Offset$$
 [6]

or equivalently:

$$y = M \cdot x + B \tag{7}$$

By way of example, table 9 lists the particle energies, their calculated velocities, and respective peak centroids for the determination of the parameters M and B (leading to d and Offset) for one of the sixteen array bars.

Table 9 Parameters derived from the delayed neutron energies and their centroid channels necessary to calibrate the neutron bars.

Particle Type	Energy MeV	Velocity m/100ps	1/v 100ps/m	Peak Centroid Channel
Neutron	1.702	0.00180	554.2	759.1
Neutron	1.172	0.00149	667.8	874.7
Neutron	1.715	0.00181	552.1	754.9
Neutron	0.808	0.00124	804.3	1009.8
¹⁶ C β	1.522	0.02823	35.4	234.0
¹⁷ N β	1.265	0.02742	36.5	234.6

These data, specifically columns four and five of table (above) were fit to a line yielding best fit values for M and B along with their associated interpolation errors. Figure 16 shows the data and the fit for neutron bar number one where the centroid is plotted on the X-axis and the inverse velocity is plotted on the Y-axis.

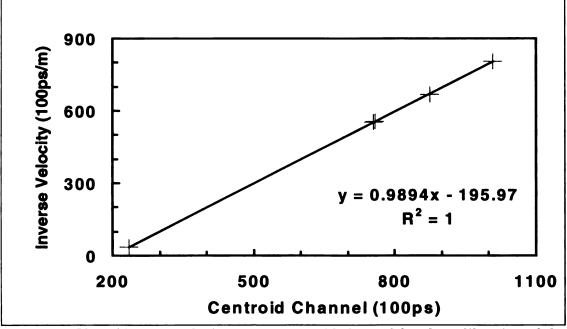


Figure 16 Plot of inverse velocity versus centroid channel for the calibration of the neutron bar array.

From the best-fit values of M and B determined on a bar-by-bar basis, values for the parameters d and Offset can be calculated. When known, the constants d and Offset can be incorporated into new SARA pseudo parameters that will result in histograms that have normalized the time-of-flight to a flight path of one metre. The channel number will be a measure of the inverse velocity in units of (100ps/metre) or, equivalently, time-of-flight in hundreds of picoseconds. What is perhaps most important is that, following this transformation, the time-of-flight spectra of the individual array bars are gain matched, if you will, which allows for a cumulative neutron TOF spectrum to be recorded. This is advantageous generally, but in particular it ensures that the experiment as a whole is as sensitive to lower-energy delayed neutrons as possible. Table 10 lists the values of the best-fit parameters and the calculated values of d and Offset for each of the sixteen array bars.

3.5.2 NEUTRON BAR ARRAY – EFFICIENCY CALIBRATION

The time calibrated spectra of the individual bars were summed to generate the final delayed neutron spectra with the caveat that bar #15 was excluded due to a failure of one of the phototubes during the experiment. The summed spectra collected for each of the calibration isotopes are shown in figures 17 and 18. Included in the figures are the raw data, the individual peak contributions (blue), the two-component background (black), and the aggregate fit to the data (red). The best fit is the minimum in the total χ^2 which was determined using the minimization package MINUIT. The error in the fit for each parameter was calculated within the MINUIT routine using the associated MINOS package that removes the parameter correlation and hence delivers a more true evaluation of the individual parameter errors.

Table 10 Summary of the calibration fits for each of the array bars. The parameters d and Offset are generated from the M and B results.

Array Bar	M Value	B value	Distance d	Offset
	(from fit)	(from fit)	metres	Channels
1	0.989(2)	-196(1)	1.011(2)	198(1)
2	0.987(2)	-193(1)	1.013(2)	196(1)
3	0.988(2)	-217(2)	1.012(2)	220(2)
4	0.989(2)	-187(1)	1.011(2)	189(2)
5	0.981(4)	-252(3)	1.019(4)	257(4)
6	0.983(3)	-235(3)	1.018(4)	239(3)
7	0.990(2)	-254(2)	1.010(2)	256(2)
8	0.992(3)	-200(2)	1.008(3)	202(2)
9	0.992(5)	-181(3)	1.008(5)	183(3)
10	0.991(2)	-196(1)	1.009(2)	197(2)
11	0.991(2)	-163(1)	1.009(2)	164(1)
12	0.989(2)	-219(2)	1.011(2)	221(2)
13	0.992(1)	-192(1)	1.008(1)	193(1)
14	0.986(2)	-172(1)	1.014(2)	174(1)
15	0.980(4)	-162(3)	1.020(4)	165(3)
16	0.978(2)	-206(1)	1.022(2)	211(1)

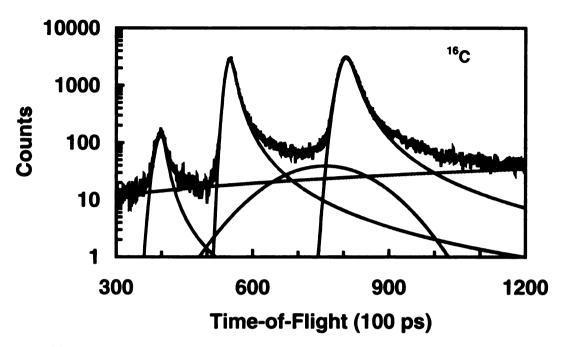


Figure 17 Aggregate TOF delayed-neutron spectrum recorded by the neutron bar array during the decay of 16 C.

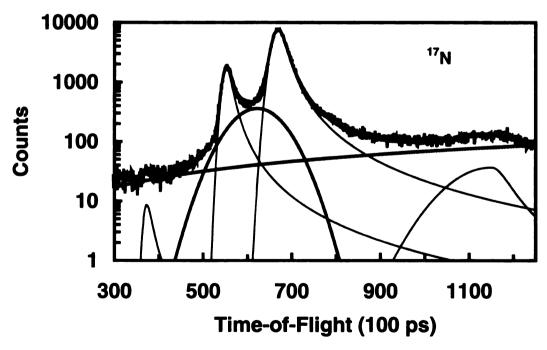


Figure 18 Aggregate TOF delayed-neutron spectrum recorded by the neutron bar array during the decay of ^{17}N .

To generate an efficiency calibration curve, the known transitions of the calibration isotopes (¹⁶C, ¹⁷N) were integrated. Knowing the individual branching ratios for the delayed neutron transitions [93Ti, 76Al] and the total number of parent decays, several calibration points for the total efficiency were deduced. For higher energy neutrons, the efficiency of the array has been approximated using a Monte-Carlo method [93Ha]. Table 11 gives the calculated efficiency points from the on-line data and figure 19 shows the final efficiency calibration curve including the Monte-Carlo data and the spline fit. The error in the efficiency determination has been shown to be about ten percent of the extracted value [93Ha].

Table 11 Data points used to generate and efficiency calibration for the neutron bar array. The monte-carlo points are from reference [93Ha].

Source	E _n (keV)	Branch (%)	Parent	Peak Area	Efficiency
			Decays		Total
¹⁷ N	384	38.0 (13)	2.3(1)E7	6.1(5)E3	6.98(55)E-4
¹⁷ N	1172	50.1 (13)	2.3(1)E7	3.33(4)E5	2.89(10)E-2
¹⁷ N	1702	6.9 (5)	2.3(1)E7	5.07(32)E4	3.20(31)E-2
¹⁶ C	808	84.4 (17)	1.25(5)E7	1.50(3)E5	1.42(5)E-2
¹⁶ C	1715	15.6 (17)	1.25(5)E7	8.51(16)E4	4.37(49)E-2
Monte Carlo	1800	N/A	N/A	N/A	3.35(34)E-2
Monte Carlo	3000	N/A	N/A	N/A	3.16(32)E-2
Monte Carlo	4500	N/A	N/A	N/A	2.86(29)E-2

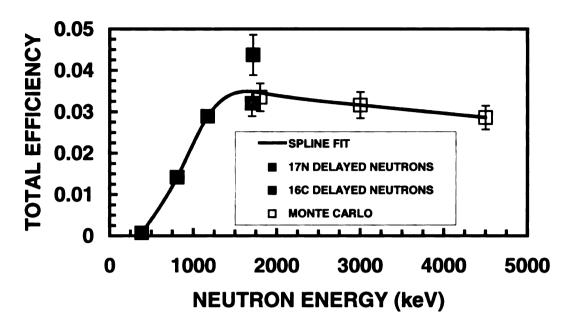


Figure 19 Calibration curve for the total efficiency of the neutron bar array.

3.5.3 NEUTRON BAR RESULTS - 19N AND 20N

The delayed neutron spectra of the isotopes of interest (¹⁹N, ²⁰N) were recorded under experimental conditions identical with those of the calibration beams of ¹⁶C and ¹⁷N. The spectra collected by the individual bars were summed to generate cumulative delayed neutron spectra suitable for integration. The raw data were fit to peaks having the same functional form as with the calibration spectra and were minimized using the MINUIT/MINOS package.

3.5.4 19N DELAYED NEUTRON SPECTRUM

Figure 20 shows the cumulative neutron TOF spectrum collected by the neutron bar array during the decay of ¹⁹N. The background components of the fit are shown in black, the individual peaks are shown in blue, and the sum of the components is shown in red. The TOF scale is given in bins having 100 ps width.

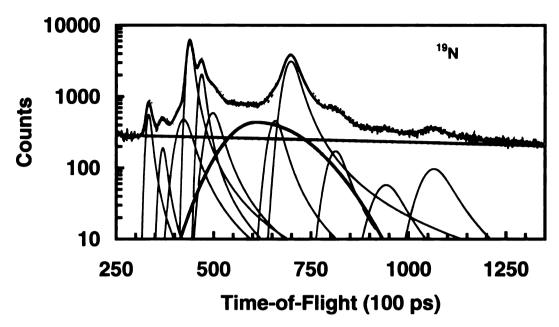


Figure 20 Delayed-neutron time-of-flight spectrum collected by the neutron bar array during the decay of ¹⁹N.

As is evidenced by the figure, the delayed neutron spectrum of ¹⁹N is a rich one. The best-fit to the data indicates that there are eleven delayed-neutron lines with eight of these being reasonably well-separated, and three appearing as shoulders on larger peaks. From the best-fit centroids and knowing the time of flight scale, the energy of the neutron lines may be calculated. Table 12 lists the peak centroids, and calculated delayed neutron energy in both the laboratory and center-of-mass reference frames.

From the integrated areas of the eleven observed neutron peaks, the total efficiency calibration, and the sum of the parent beta decays, the branching ratios of the transitions can be calculated. Table 13 lists the ¹⁹N observed delayed neutrons by energy, the values of the parameters necessary for the branching ratio calculation, and the corresponding branching ratios.

Table 12 Delayed neutron lines observed during the decay of ¹⁹N. The TOF is given along with the neutron energies in both the lab and center of mass frames.

Peak Designation	Time of Flight	Neutron Energy	Neutron Energy
	(100 ps)	Lab. (MeV)	C.o.M. (MeV)
1	335.24	4.65(8)	4.91(8)
2	371.55	3.79(6)	4.00(6)
3	424.68	2.90(5)	3.06(5)
4	441.01	2.69(4)	2.84(4)
5	471.07	2.36(4)	2.49(4)
6	499.75	2.09(4)	2.21(4)
7	659.86	1.20(3)	1.27(3)
8	699.93	1.07(3)	1.13(3)
9	814.76	0.79(3)	0.83(3)
10	944.24	0.59(3)	0.62(3)
11	1066.70	0.46(3)	0.49(3)

Table 13 Delayed neutron branching ratios for those neutron lines observed in the neutron bar array during the decay of ¹⁹N.

Neutron	$E_n(MeV)$	Peak	Parent	Total	Branching
Peak	СоМ	Area	Decays	Efficiency	Ratio (%)
1	4.91(8)	1.19(3)E4	4.64(10)E7	2.85(29)E-2	0.90(10)
2	4.00(6)	5.13(32)E3	4.64(10)E7	3.00(30)E-2	0.37(4)
3	3.06(5)	2.91(36)E4	4.64(10)E7	3.18(32)E-2	1.97(32)
4	2.84(4)	1.44(3)E5	4.64(10)E7	3.22(33)E-2	9.6(11)
5	2.49(4)	5.34(12)E4	4.64(10)E7	3.29(33)E-2	3.5(4)
6	2.21(4)	3.50(78)E4	4.64(10)E7	3.36(34)E-2	2.2(4)
7	1.27(3)	2.60(62)E4	4.64(10)E7	2.96(30)E-2	1.9(2)
8	1.13(3)	1.88(2)E5	4.64(10)E7	2.50(25)E-2	16.2(17)
9	0.83(3)	1.25(7)E4	4.64(10)E7	1.33(14)E-2	2.0(3)
10	0.62(3)	6.48(29)E3	4.64(10)E7	6.51(66)E-3	2.2(3)
11	0.49(3)	1.14(4)E4	4.64(10)E7	2.84(29)E-3	8.7(10)

3.5.5 ²⁰N DELAYED NEUTRON SPECTRUM

The data collected by the neutron bar array during the decay of ²⁰N was summed in a manner consistent with the previously detailed analyses. Unfortunately, the production of ²⁰N from a primary beam of ²²Ne requires a particle transfer or charge-exchange mechanism rather than the more probable nucleon removal. As a result, the intensity of the ²⁰N secondary beam was dramatically reduced relative to the other secondary isotopes studied. Because of this reduction in intensity, the data collected by the neutron bar array was compressed by a factor of two to better examine the features of the plot and to improve the fits to the spectrum.

Once again, the individual bar data were summed to produce and aggregate delayed-neutron time-of-flight spectrum for the decay of ²⁰N. The time bins, in this case, had widths of 200 ps. Figure 21 is the aggregate neutron bar spectrum and shows the raw data, background terms (black), individual peaks (blue) and the cumulative fit to the spectrum (red). It is noted that this spectrum has the y-axis on a linear scale rather than on a logarithmic scale as for the previous spectra.

The cumulative fit from MINUIT supports a delayed neutron spectrum consisting of eight transitions. Six of these are well separated in energy while the other two appear as shoulders of larger peaks. From the best-fit centroids, the energy of the delayed neutrons can be calculated knowing that each channel represents 200 ps in time. Table 14 lists the ²⁰N peak centroids, and the corresponding energies in both the laboratory and center of mass reference frames.

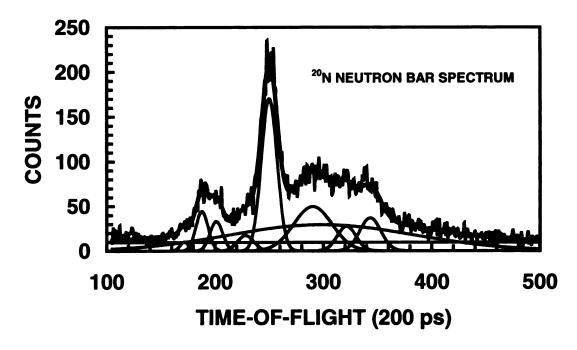


Figure 21 Delayed neutron spectrum recorded by the neutron array during the decay of $^{20}\rm{N}$.

Table 14 Summary of delayed neutrons observed during the decay of ²⁰N. Listed are the centroid channels and the corresponding neutron energy in both the lab and center of mass frames.

Peak Designation	Centroid 1ch = 200 ps	Neutron Energy Laboratory (MeV)	Neutron Energy C.o.M. (MeV)
1	171.9	4.41(7)	4.64(7)
2	188.1	3.68(6)	3.88(6)
3	201.4	3.21(5)	3.38(5)
4	228.4	2.50(4)	2.63(4)
5	250.3	2.08(4)	2.19(4)
6	290.6	1.55(3)	1.63(3)
7	321.7	1.26(3)	1.33(3)
8	343.6	1.11(3)	1.16(3)

From the integrated peak areas, the measured number of parent ²⁰N decays, and the efficiency calibration, the branching ratio for each transition was calculated. Table

15 is a summary of the branching ratio calculation for ²⁰N delayed neutrons that includes the values of the parameters necessary to elucidate the decay strength.

Table 15 Calculated branching ratios for the delayed neutron lines observed by the neutron bar array during the decay of ²⁰N.

Neutron Peak	E _n (MeV) CoM	Peak Area	Parent Decays	Total Efficiency	Branching Ratio (%)
1	1.16(3)	900(30)	7.6(1)E5	2.65E-02	4.5(5)
2	1.33(3)	600(38)	7.6(1)E5	3.12E-02	2.5(3)
3	1.63(3)	2200(57)	7.6(1)E5	3.47E-02	8.3(9)
4	2.19(4)	3087(56)	7.6(1)E5	3.37E-02	12.1(13)
5	2.63(4)	329(33)	7.6(1)E5	3.26E-02	1.3(2)
6	3.38(5)	486(25)	7.6(1)E5	3.12E-02	2.1(3)
7	3.88(6)	624(26)	7.6(1)E5	3.03E-02	2.7(3)
8	4.64(7)	102(19)	7.6(1)E5	2.88E-02	0.5(1)

3.6 ⁶Li INORGANIC SCINTILLATORS

In addition to the neutron bar array, the experimental set-up included three ⁶Li glass scintillators to detect delayed neutron events. The TDC modules for these devices were calibrated in a manner completely analogous with the procedure outlined for the bars of the array. The slopes of the TDC channels were determined by the use of a precision time calibrator set to 10ns intervals and a 320ns range.

To complete the calibration, the decay of the isotopes ¹⁶C and ¹⁷N were monitored by the glass scintillators. Figures 22 and 23 show the spectra collected by one of the three ⁶Li detectors (detector #3) in the time domain for each of the ¹⁶C and ¹⁷N calibration beams. Included in the plot is the best fit to the data assuming a gaussian peak shape below the peak centroid and a Briet-Wigner shape above the centroid. The error bars are the statistical errors of the points only. It is noted that, due

to the low total efficiency of the ⁶Li devices, the spectra are compressed to seven bits to better illuminate the salient features of the presented time-of-flight plots.

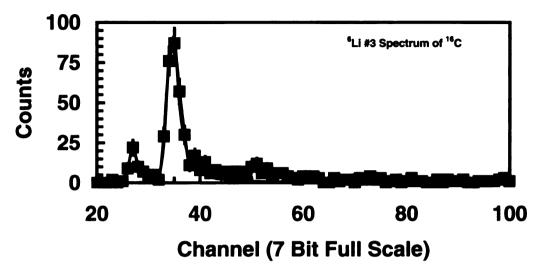


Figure 22 Delayed neutron spectrum collected in ⁶Li detector #3 during the decay of ¹⁶C.

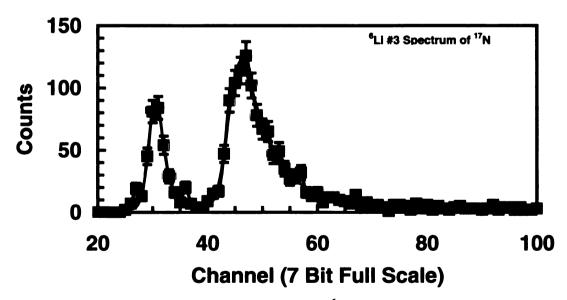


Figure 23 Delayed neutron spectrum collected in ⁶Li detector #3 during the decay of ¹⁷N.

As with the neutron bars, a calibration curve was constructed on the basis of the observed centroid channel of the delayed neutrons and their known energies. The parameters of the fit were the same as outlined in section 3.5.1 and will not be repeated here. Figure 24 shows the calibration curve for one of the ⁶Li detectors (#3) generated using the compressed data. The y-axis value is inverse velocity in (ns/m) and the x-axis is the TDC channel of the neutron peak centroid.

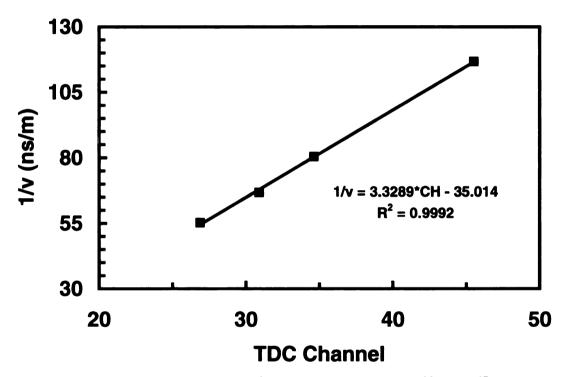


Figure 24 Time calibration curve for ⁶Li detector #3 from the ¹⁶C and ¹⁷N decays.

The procedure was repeated for each of the three ⁶Li detectors in the seven-bit time domain. The final values of the slope and offset parameters relating the TDC channel to the inverse velocity are given in table 16. The inverse velocity in (ns/m) will be equal to the channel multiplied by the slope parameter plus the offset parameter. The energy of the neutron in MeV will be 5227.083 times the squared velocity in ns/m.

Table 16 Calibration parameters for the lithium detectors in the seven bit time domain. The inverse velocity of a neutron in (ns/m) is equal to the TDC channel times the slope plus the offset.

Detector	Slope	Offset
⁶ Li #1	3.631	-36.406
⁶ Li #2	7.651	-114.15
⁶ Li #3	3.329	-35.014

3.6.1 EFFICIENCY CALIBRATION

To determine the efficiency of the ⁶Li glass detectors as a function of energy, the four delayed neutrons emitted by the calibration isotopes, ¹⁷N and ¹⁶C were integrated in each of the spectra. From these data, the known branching ratios of the neutron lines, and the total number of decays extracted from the half-life spectra, a measure of the ⁶Li glass detectors' efficiencies at the corresponding energies was calculated. Table 17 gives the measured total efficiency as a function of neutron energy for the three ⁶Li detectors.

Table 17 Calibration points utilized to determine the total efficiency of each lithium detector. The source neutrons are from the decay of ¹⁶C and ¹⁷N.

E _n keV	⁶ Li #1 ε–total	⁶ Li #2 ε–total	⁶ Li #3 ε–total
384	1.68(15) E-4	2.62(16) E-4	1.35(9) E-4
808	4.06(39) E-5	6.65(61) E-5	3.13(31) E-5
1172	3.81(35) E-5	7.96(49) E-5	3.01(22) E-5
1715	3.39(62) E-5	5.70(94) E-5	3.08(58) E-5

The neutron capture cross section of ⁶Li is well known over a large range of incident neutron energies and they have been tabulated in the common nuclear

databases [00NN]. The molar composition of the inorganic material was supplied by the manufacturer [97Bi] and the isotopic fraction of ⁶Li was also made available. From the known neutron-capture cross section and the physical characteristics of the scintillators, the intrinsic efficiency of the three ⁶Li detectors was calculated following the procedure outlined in [81Fr]. The center of mass energies were converted to lab energy for the cross-sectional data and the intrinsic efficiency curve was scaled to fit the measured total efficiencies supplied by the calibration beams of ¹⁶C and ¹⁷N. Figure 25 shows the scaled intrinsic efficiency for ⁶Li detector #3 and the data to which the cross-sectional curve was scaled.

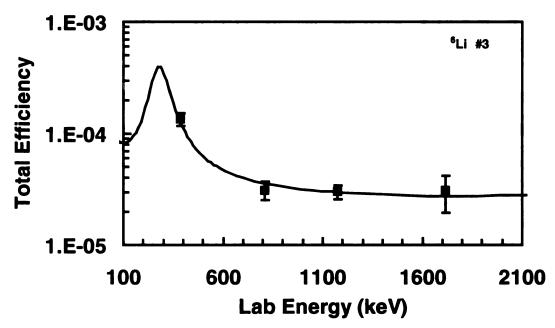


Figure 25 Calibration curve for the total efficiency of ⁶Li detector #3.

The scaling factor derived for the three ⁶Li glass scintillators should be representative of the detectors' geometric efficiencies. The solid angle coverage of the

detectors was calculated from the measured physical geometry and crystal dimensions.

The necessary parameters to calculate the geometric solid angle are shown in figure 26.

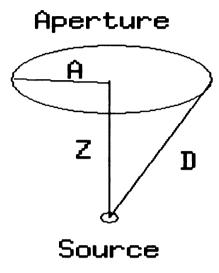


Figure 26 Parameters necessary to determine the geometric efficiency of an aperture.

Here, A is the detector radius (2.54cm), Z is the linear source-to-aperture distance, and D is the distance between the source activity and the edge of the aperture. From these values, and assuming that the source/detector separation is much larger than the source spread, the geometric solid angle (G_p) may be calculated from the expression:

$$G_p = 1/2(1 - Z/D)$$
 [8]

Table 18 lists the geometric solid angle for each of the ⁶Li detectors calculated in this manner along with the scaling parameter for the efficiency fits based on the reported cross-sectional data. The error in the scaling parameter reflects the change in its value necessary to produce a $\Delta \chi^2$ of one unit in the fit.

Table 18 Calculated value of the geometric efficiency and best fit value of the scaling parameter for each of the lithium detectors.

Detector	Geometric Efficiency	Scaling Factor
⁶ Li #1	4.11(12) x 10 ⁻³	3.57(37) x 10 ⁻³
⁶ Li #2	1.46(8) x 10 ⁻²	1.57(13) x 10 ⁻²
⁶ Li #3	2.64(7) x 10 ⁻³	2.89(25) x 10 ⁻³

Of note here is the good agreement between the solid angle estimates from the physical geometry and the best-fit value for the scaling parameter, particularly for detectors two and three. The agreement provides a greater measure of confidence in the total efficiency derived for the ⁶Li glass scintillators and thus for the branching ratio determinations for the ¹⁹N isotope of interest.

3.6.2 ¹⁹N RESULTS – ⁶Li DETECTORS

Unfortunately, the intensity of the ²⁰N secondary beam taken together with the total efficiency of the ⁶Li detectors resulted in too few events recorded by the devices to be of use. As a consequence, the results presented here are restricted to those for ¹⁹N.

The goal behind including the ⁶Li detectors was to enable the observation of low energy delayed neutrons from the decay of ¹⁹N. To accentuate the low energy peaks in the ⁶Li spectra, the time calibration was utilized to re-bin the data into the energy domain. The following three figures 27-29 show the spectra collected by each of the ⁶Li detectors during the decay of ¹⁹N where the data have been converted into an energy scale and redistributed into bins of 40 keV width. Also shown are the fits to the peaks where the peak shapes used were simple gaussians and the background is of the form 1/x².

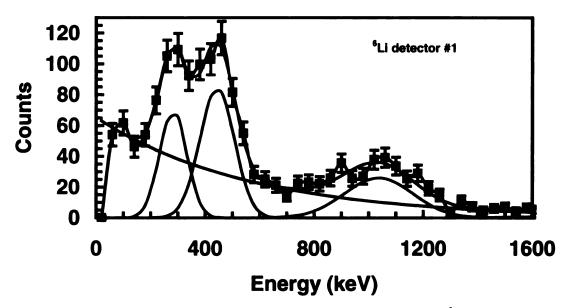


Figure 27 Energy spectrum of delayed neutrons recorded by ⁶Li detector #1 during the decay of ¹⁹N.

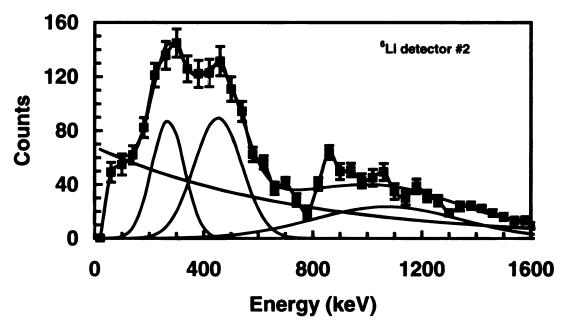


Figure 28 Energy spectrum of delayed neutrons recorded by ⁶Li detector #2 during the decay of ¹⁹N.

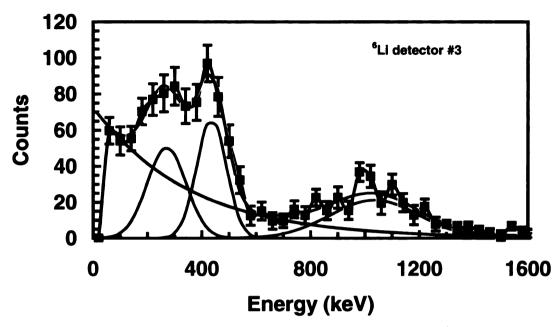


Figure 29 Energy spectrum of delayed neutrons recorded by ⁶Li detector #3 during the decay of ¹⁹N.

From the integrated area of the peaks, the detection efficiency calibration, and the total number of ¹⁹N decay events, the branching ratio can be deduced for the transitions observed in each of the three devices. The dominant sources of error in the branching ratio determination come from the integrated area of the peaks and the total efficiency. Due to the uncertainty in the shape of the ⁶Li energy spectrum background, the error was estimated from a series of fits to the data with different background shapes including exponentials, lines, and functions of the form 1/x². All fits were done in MINUIT/MINOS and the large uncertainties in the peak areas reflect differences in the best-fit values of the peak parameters over the various functional forms of the background.

The error in the efficiency comes from two sources; first, the error in the scaling parameter imparts an error of nearly ten percent in all three cases and, second, the error

in the centroid energy from the fits drives an additional uncertainty in the efficiency, particularly for low energy neutrons where the slope of the efficiency curve is extremely steep. Consequently, taking the energy error into account yields large fractional errors for the ~280 and ~460 keV transitions. Table 19 gives the necessary parameters and the calculated branching ratios for the delayed neutrons by detector.

Table 19 Branching ratio determination for delayed neutrons observed by the lithium detectors during the decay of ¹⁹N. Also tabulated are the parameters necessary for the branching ratio calculation along with the associated errors.

Detector	Neutron Energy (keV)	Peak Area	¹⁹ N Total Decays	Efficiency (Total)	Branching Ratio (%)
⁶ Li #1	280(80)	331(115)	4.64(10)E7	4.9(13)E-4	1.5(7)
⁶ Li #1	450(40)	449(115)	4.64(10)E7	1.01(16)E-4	9.6(29)
⁶ Li #1	1040(40)	305(100)	4.64(10)E7	3.86(41)E-5	17.0(59)
⁶ Li #2	270(80)	483(155)	4.64(10)E7	8.6(14)E-4	1.2(4)
⁶ Li #2	460(40)	649(155)	4.64(10)E7	1.63(24)E-4	8.6(20)
6Li #2	1070(40)	590(195)	4.64(10)E7	6.61(69)E-5	19.2(67)
⁶ Li #3	280(80)	306(80)	4.64(10)E7	3.93(65)E-4	1.7(6)
⁶ Li #3	440(40)	306(80)	4.64(10)E7	8.5(19)E-5	7.8(27)
⁶ Li #3	1040(40)	267(50)	4.64(10)E7	3.13(34)E-5	18.4(40)

The branching ratio results for the corresponding transitions among the three ⁶Li detectors are in excellent agreement. The weighted means of the measurement yield branching ratios of 1.4(22) %, 8.6(10) % and 18.2(23) % for the delayed neutrons having laboratory energies of about 280 keV, 460 keV and 1050 keV respectively.

3.7 GAMMA RAY DATA

The experimental end-station included two HPGe detectors to monitor gammarays emitted during the decay of the isotopes of interest. The detectors were energy calibrated with in-house sources of ¹³⁴Cs, ²⁰⁷Bi, and ⁶⁰Co - all of which have known decay schemes and well-established gamma cascades. The detectors were energy calibrated over the full 8192 channel spectra.

The in-house sources were monitored by affixing the source's solid support backing to the implantation scintillator. However, unlike the radioactive beam runs, the implantation scintillator was removed from the trigger and replaced by a NaI detector "start" signal. This substitution for the implantation scintillator "start" permitted conclusive identification of the gamma ray causing the "start" signal in the electronics which later facilitated a determination of the HpGe detectors' efficiency at various known energies. The NaI detector was placed directly beneath the implantation scintillator to provide the maximum possible solid angle coverage which averaged out angular correlations. As an example of the spectra collected in the HPGe detectors using the calibration sources, Figure 30 shows the response of one of the gamma-ray detectors to the ⁶⁰Co activity.

The peaks observed in the Germanium detectors were fit to symmetric gaussian shapes plus linear backgrounds. Using these data, a calibration line for each of the two germanium detectors was determined by plotting the centroid channel versus the known transition energy. The data points were fit to a line using EXCEL. The raw data and the results of the straight line fits are shown in figure 31. The slope and offset for the energy calibration of Ge1 are 0.4919(2) keV/CH and 11.51(31) keV, respectively, while the slope and offset for the full-scale energy calibration of Ge2 are 0.7309(3) keV/CH and 9.02(29) keV.

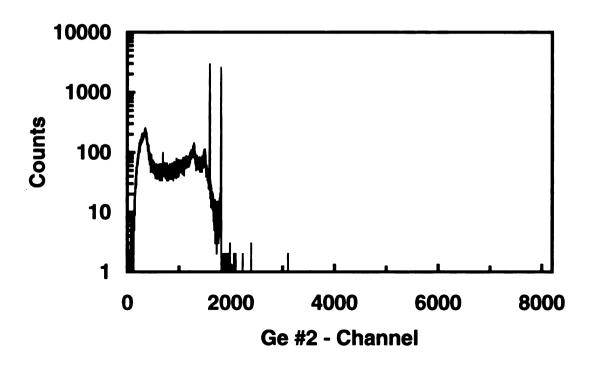


Figure 30 Gamma-ray spectrum collected in Ge #2 from the ⁶⁰Co calibration source.

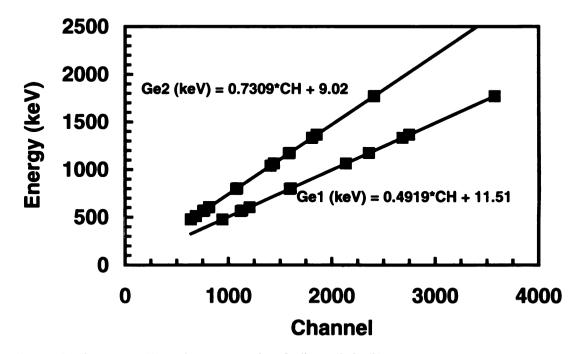


Figure 31 Energy calibration curves for Ge#1 and Ge#2.

3.7.1 GAMMA-RAY SPECTRA OF THE ISOTOPES OF INTEREST

The experiment proceeded by implanting secondary beam activity within the start detector for a set time followed by a beam-off interval where the real-time clock was in play. The germanium detectors, however, were supported by an electronics algorithm that permitted data to be collected during both the implantation period and the beam-off period. This was done to allow for the possibility of observing gamma rays of lower intensity. For the branching ratio determinations outlined in section 4.1, the included data are restricted to the beam-off interval as this is the portion of the sequence from which the beta activity may be conclusively extracted.

Figures 32 through 39 show the spectra collected in the germanium detectors during the beam-off sequences of the data runs for the four secondary beams (16 C, 17 N, 19 N, 20 N). The spectra are plotted on a logarithmic scale of counts versus energy in keV as calculated from the calibration formulae indicated in figure 31.

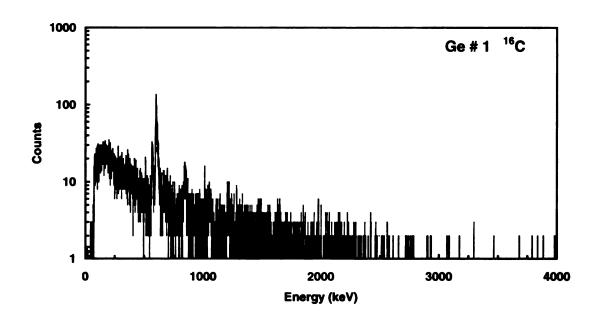


Figure 32 Gamma-ray spectrum collected by Ge#1 during the decay of ¹⁶C.

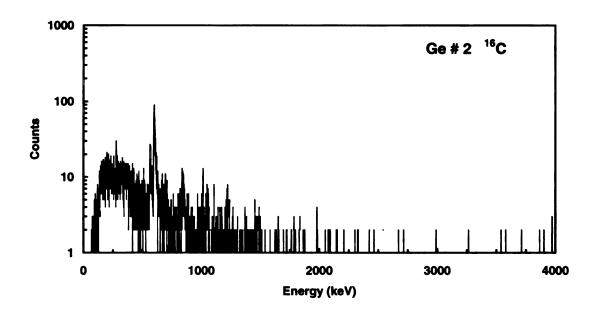


Figure 33 Gamma-ray spectrum collected by Ge#2 during the decay of ¹⁶C.

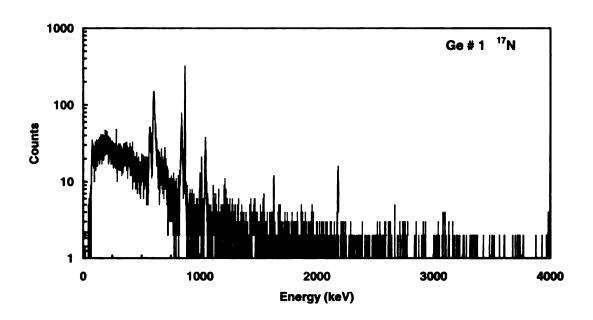


Figure 34 Gamma-ray spectrum collected by Ge#1 during the decay of ¹⁷N.

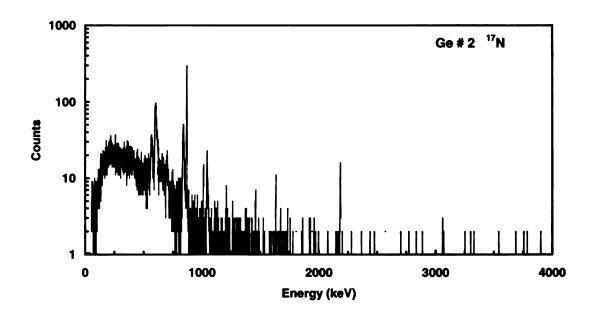


Figure 35 Gamma-ray spectrum collected by Ge#2 during the decay of ¹⁷N.

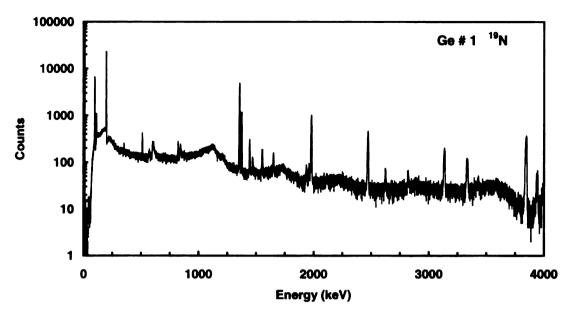


Figure 36 Gamma-ray spectrum collected by Ge#1 during the decay of ¹⁹N.

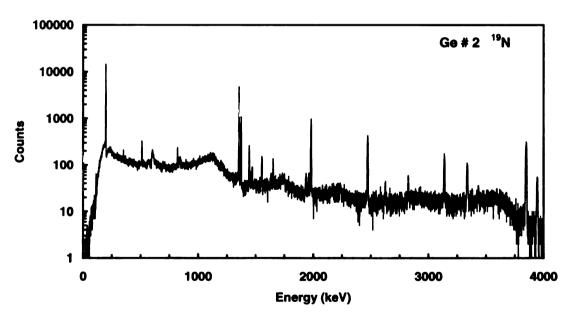


Figure 37 Gamma-ray spectrum collected by Ge#2 during the decay of ¹⁹N.

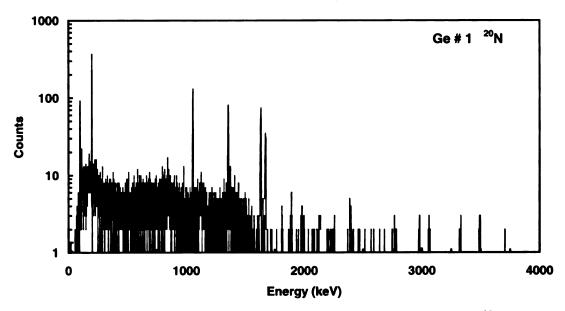


Figure 38 Gamma-ray spectrum collected by Ge#1 during the decay of 20 N.

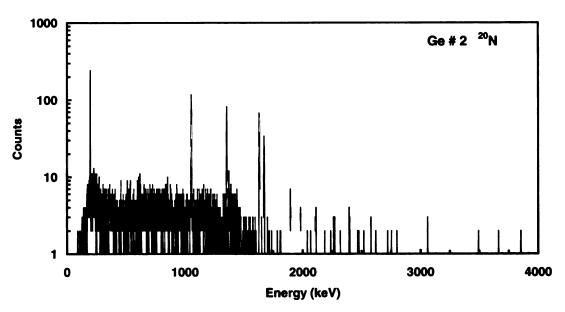


Figure 39 Gamma-ray spectrum collected by Ge#2 during the decay of ²⁰N.

3.7.2 BACKGROUND GAMMA RAYS

Gamma-ray lines that appeared in the spectra of any three of the four isotopes of interest were assigned as being due to background. Table 20 lists, by detector, the transitions satisfying this criterion. As expected, there is a direct correspondence between the two detectors for the gamma rays identified in this way. The exception to the background assignment, of course, is the 511 keV transition which is not necessarily due to background exclusively.

Table 20 Gamma rays attributed to background.

Ge # 1	Ge # 2
Energy (keV)	Energy (keV)
352	351
420	419
442	441
511	511
569	571
605	606
844	840
1014	1014
1273	1273
1367	1368
1808	1809

3.7.3 RATIO OF EFFICIENCIES

To determine the detection efficiency ratio of Ge1 to Ge2, and to detect systematic problems that may have developed over time in the germanium detectors, the ratio of counts in Ge1 to Ge2 for non-background transitions was calculated. The ratio of Ge1 to Ge2 intensities should be a constant for all gamma transitions originating at the implantation position provided that both germanium detectors were stable over the course of the experiment. Furthermore, if constant, the ratio of the counts is equal to

the ratio of the total detection efficiencies. Figure 40 shows a plot of the ratio of counts in Ge1 to counts in Ge2 (during the beam-off period) versus the energy of the gammaray line for those transitions observed during the ¹⁹N production runs.

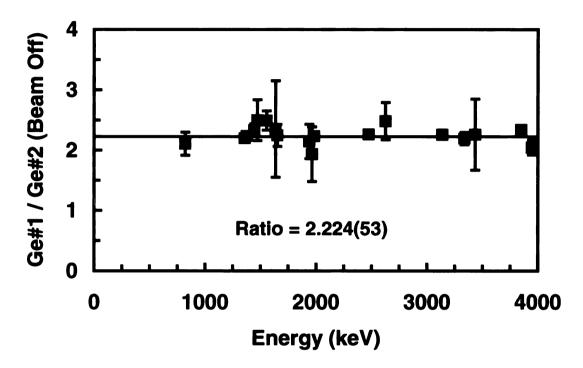


Figure 40 Ratio of counts detected in Ge#1 to counts in Ge#2 during the beam-off period of the ¹⁹N experimental runs.

The figure suggests that the ratio of counts is indeed a constant. The value of the constant is 2.225(53) determined from a weighted linear fit to the data yielding a best-fit slope consistent with zero. Thus, the total detection efficiency of Ge1 is equal to this constant multiplied by the total detection efficiency of Ge2 over the energy range indicated.

3.7.4 GAMMA RAY SEGREGATION

As noted earlier, the germanium detectors were active during both the implantation and beam-off intervals. By taking the ratio of counts detected at each

gamma line over all times to the counts detected during the beam-off period, it was possible to segregate the observed gamma rays into groups. This was useful for assigning gamma rays as being due to the decay of the parent or the decay of the daughter.

The implantation and beam-off periods were set to maximize the counts detected from the isotope of interest. However, there may be significant total activity due to the beta decay of the daughter. The intensity of counts detected over all times to beam-off will not be the same for the daughter(s) as it will be for the isotope of interest – due to the difference between the half-lives. Thus, if the statistics are good enough, a plot of the counts over all times to beam-off counts should filter gammas due to the daughter's decay from those due to the implanted species. Figure 41 shows an example of this for ¹⁹N where the ratio of counts in Ge2 for all times to beam-off is plotted against transition energy.

From figure 41 it appears that four of the observed gamma rays (blue) are not from the same source as the others (red). This is not unexpected as, in addition to gamma rays from the decay of ¹⁹N, there should also be gamma rays present following the beta decay of the ¹⁹N beta daughter, ¹⁹O. This result was useful for the assignment of previously unobserved gamma rays.

3.7.5 GAMMA RAY HALF-LIFE MEASUREMENTS

In addition to the detection ratios, the gamma rays were assigned on the basis of their half-life. In most cases, the intensity of the transitions was sufficiently strong to permit an analysis by this technique, however, it is noted that for the weakest transitions this method was not conclusive due to statistics and to the unfortunate fact that the general background, whether due to Compton edges or betas or..., tended to have a half-life consistent with the dominant source implanted.

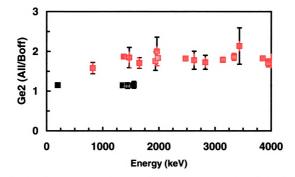


Figure 41 Ratio of counts detected in Ge #2 over all times to beam-off period during the decay of ¹⁹N.

Unlike the determination of the beta half-lives, the approach taken here was not rigorous. To circumnavigate the dead-time, the half-life spectra were fit over the final 80% of the clock channels rather than utilizing the more complicated dead-time/t formalism outlined in section 3.4.1. Fortunately, the high purity of the secondary beams and the dramatic difference between the half-lives of the parent and daughter isotopes permitted a conclusive separation of gamma rays even using this less rigorous approach. It is necessary to note that the best-fit value of the half-lives from the gamma gating were systematically longer than those from the beta decay analysis – not

surprisingly. As an example, figure 42 shows two representative half-life spectra from gamma rays observed during the implantation of a ¹⁹N secondary beam.

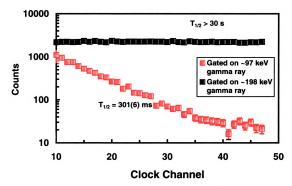


Figure 42 Gamma-ray gated half-life spectra for two transitions observed in Ge #1.

3.7.6 GAMMA RAY ASSIGNMENTS 19N, 20N SECONDARY BEAMS

The gamma gated half-life spectra and the ratio of counts analysis facilitate the assignment of gamma rays to specific sources and as originating from specific levels. From the half-life data, the identity of the beta-decaying isotope may be extracted and from the energy of the gamma ray, the specific transition can be determined by comparison with daughter levels and in some cases, known transition energies given in the literature. Table 21 give the assignment of the observed, non-background gamma rays for the ¹⁹N secondary beam runs and table 22 for the ²⁰N secondary beam runs.

Table 21 Gamma ray assignments for transitions observed during the beam-off period of the $^{19}{\rm N}$ production runs.

R.I.B.	γ-Ray	β-Decay	γ-Ray	E-initial	E-final
	keV	Source	Source	level	level
¹⁹ N	96	¹⁹ N	¹⁹ O	96	0
¹⁹ N	110	¹⁹ O	¹⁹ F	110	0
¹⁹ N	197	¹⁹ O	¹⁹ F	197	0
¹⁹ N	511	-	-	-	-
¹⁹ N	821	¹⁹ N	¹⁸ O	4456	3634
¹⁹ N	1357	¹⁹ O	¹⁹ F	1554	197
¹⁹ N	1374	¹⁹ N	¹⁹ O	1472	96
¹⁹ N	1444	¹⁹ O	¹⁹ F	1554	110
¹⁹ N	1472	¹⁹ N	¹⁹ O	1472	0
¹⁹ N	1554	¹⁹ O	¹⁹ F	1554	0
¹⁹ N	1633				
¹⁹ N	1652	¹⁹ N	¹⁹ O	3634	1982
¹⁹ N	1938	¹⁹ N	¹⁸ O	3920	1982
¹⁹ N	1960	¹⁹ N	¹⁹ O	-	-
¹⁹ N	1980	¹⁹ N	¹⁸ O	1982	0
¹⁹ N	2473	¹⁹ N	¹⁹ O	3945	1472
¹⁹ N	2623	¹⁹ N	¹⁹ O	-	-
¹⁹ N	2822	¹⁹ N	¹⁹ O	-	-
¹⁹ N	3136	¹⁹ N	¹⁹ O	3232	96
¹⁹ N	3338	¹⁹ N	¹⁹ O	-	-
¹⁹ N	3434	¹⁹ N	¹⁹ O	-	-
¹⁹ N	3849	¹⁹ N	¹⁹ O	3945	96
¹⁹ N	3945	¹⁹ N	¹⁹ O	3945	0

Table 22 Gamma ray assignments for transitions observed during the beam off period of the $^{20}{\rm N}$ production runs.

R.I.B.	γ-Ray	β-Decay	γ-Ray	E-initial	E-final
	keV	Source	Source	level	level
²⁰ N	96	²⁰ N	¹⁹ O	96	0
²⁰ N	110	¹⁹ O	¹⁹ F	110	0
²⁰ N	197	¹⁹ O	¹⁹ F	197	0
^{20}N	1057	20 O	$^{20}\mathbf{F}$	1057	0
²⁰ N	1127				
^{20}N	1357	¹⁹ O	¹⁹ F	1554	197
^{20}N	1374	^{20}N	¹⁹ O	1472	96
^{20}N	1444	¹⁹ O	¹⁹ F	1554	110
^{20}N	1634	²⁰ F	²⁰ Ne	1634	0
20 N	1674	^{20}N	20 O	1674	0
²⁰ N	1896	20 N	20 O	3570	1674

3.7.7 EFFICIENCY CALIBRATION – STANDARD SOURCES

The efficiency of detectors Ge1 and Ge2 at specific gamma-ray energies was established utilizing both on-line and off-line data. The bulk of the efficiency determination arises from standard sources that were placed at the implantation position and monitored by the germanium detectors with a sodium iodide serving as the trigger for the electronics. This is the off-line portion.

The technique involved selecting one gamma ray in the NaI detector and calculating the number of gamma rays at a different energy that would be emitted in coincidence with the first one. The sources used were ¹³⁴Cs, ²⁰⁴Bi, and ⁶⁰Co which all have gamma cascades as a part of their decay scheme [96Fi]. The following cross tables, for each of the isotopes (tables 23-25), detail the coincidence strength for gamma ray pairs in a cascade. The column headers indicate the energy of the gamma ray detected by the NaI trigger. The row headers indicate the energy of a second gamma ray that is in coincidence with the trigger – available to be recorded by one of the germanium devices. The table value is the fraction of the column-energy counts that will be emitted at the row energy. The efficiency of each germanium detector at a particular energy was determined by gating on one of the gamma rays detected in the NaI crystal. The trigger peak was integrated to determine the total number of start signals generated by the trigger gamma ray. As an example, figure 43 shows the relevant portion of the NaI spectrum collected for the ⁶⁰Co calibration source and the fits to the two peaks.

Table 23 Cross-table for ⁶⁰Co. For N gamma rays at "column" energy there are N times "cell value" gamma rays emitted at "row" energy in coincidence.

⁶⁰ Co	1173	1332
1173	0	0.99945
1332	1	0

Table 24 Cross-table for ²⁰⁷Bi. For N gamma rays at "column" energy there are N times "cell value" gamma rays emitted at "row" energy in coincidence.

²⁰⁷ Bi	1770	1442	1063	898	328	570
1770	0	0	0	0	0	6.91e-2
1442	0	0	0	1	1	7.10e-6
1063	0	0	0	0	0	8.42e-1
898	0	9.94e-1	0	0	0	0
328	0	5.50e-3	0	0	0	7.1e-6
570	1	5.50e-3	1	0	1	0
j						

Table 25 Cross-table for ¹³⁴Cs. For N gamma rays at "column" energy there are N times "cell value" gamma rays emitted at "row" energy in coincidence.

¹³⁴ Cs	1365	802	569	1039	475	796	1168	563	605
1365	0	0	0	0	0	0	0	0	3.10e-2
802	0	0	0	0	0	0	8.57e-1	8.57e-1	7.30e-2
569	0	0	0	0	0	1.81e-1	0	0	1.58e-2
1039	0	0	0	0	0	0	0	0	1.00e-2
475	0	0	0	0	0	0	1.43e-1	1.43e-1	1.20e-2
796	0	0	0	0	0	0	0	0	8.69e-1
1168	0	1.77e-1	1	0	1.77e-1	0	0	0	0
563	0	8.23e-1	0	0	8.23e-1	0	0	0	9.00e-2
605	1	8.23e-1	1	1	8.23e-1	1	0	1	0

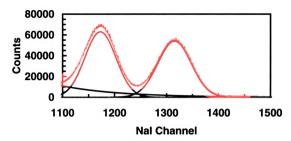


Figure 43 Gamma-ray spectrum of 60 Co collected by the NaI start detector.

The energy of the trigger gamma ray, in this case 1332 keV, was designated the column of the appropriate cross-table. The energy of a gamma ray detectable by either germanium in coincidence with the trigger was found in the cross-table (by row). Multiplying the integrated number of events in the trigger peak by the value in the cell defined by the row and column energies determined the total number of transitions at the row energy that could possibly have been detected by the germanium detectors.

Setting a gate around the NaI peak and reflecting back onto the individual germanium spectra allowed for a measurement of the coincidence spectrum. The peak in the germanium spectrum corresponding to the row energy was then integrated to determine the intensity at that energy. As an example, figure 44 shows the spectrum accumulated by Ge2 while gated on the 1332 keV peak in the NaI detector.

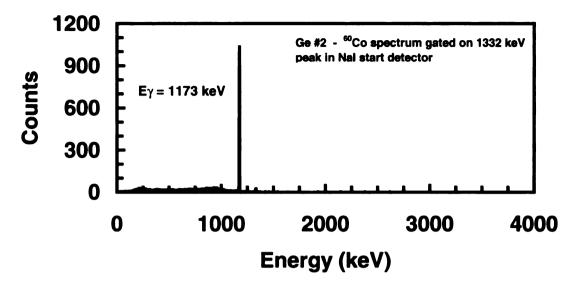


Figure 44 Gamma-ray spectrum collected in Ge # 2 gated on the ⁶⁰Co 1332 keV line in the NaI start detector.

The efficiency of the specific germanium detector at the row energy is then equal to the integrated area of that peak in the germanium detector divided by the number that could possibly have been detected under these conditions. Table 26 lists the results of the efficiency determinations for Ge1 and Ge2 using this technique. Tabulated are the calibration source, the NaI peak energy and integration, the Ge1 (Ge2) peak energy and integration, fraction as taken from the cross-tables, and calculated Ge1 (Ge2) efficiency.

3.7.8 EFFICIENCY CALIBRATION – ON-LINE SOURCES

In addition to the off-line source measurements, there are data available within the experimental runs that provide a measure of the germanium detector efficiencies at various energies. The decay of ¹⁷N provides two known gamma-ray lines for which the branching is known [93Ti] and the decay of ¹⁹O provides five gamma-ray lines with known branching ratios [95Ti]. From the integrated half-life spectra, the total number

of decays due to each isotope may be determined. Taking the individual branching ratios and the total number of decays in concert yields the number of gamma rays emitted at the specific energies. The efficiency of the germanium detector at this energy is then simply the observed intensity of the line divided by the total number emitted. Table 27 lists the quantities necessary to deduce the germanium detector efficiencies at the tabulated energy.

Table 26 Off-line source calibration data for determining the efficiencies of the Ge detectors. The emboldened columns are the data necessary to generate a calibration curve.

Source	Nal gate	Counts	Gel Peak	Counts	Fraction	Efficiency
	(keV)		(keV)		(x-table)	Ge # 1
60Co	1173	4.74E+06	1332	11294	1	2.38E-03
60Co	1332	4.29E+06	1173	10795	0.99945	2.52E-03
207Bi	570	1.20E+07	1063	22077	8.42E-01	2.19E-03
207Bi	570	1.20E+07	1770	1663	6.91E-02	2.01E-03
207Bi	1063	5.28E+06	570	17629	1	3.34E-03
207Bi	1770	3.41E+05	570	1044	1	3.06E-03
134Cs	795	1.25E+06	604	4131	1	3.31E-03
134Cs	795	1.25E+06	569	722	1.81E-01	3.19E-03
134Cs	605	2.05E+06	563	704	9.00E-02	3.81E-03
134Cs	605	2.05E+06	796	5374	8.69E-01	3.01E-03
Source	NaI gate	Counts	Ge2 Peak	Counts	Fraction	Efficiency
	(keV)		(keV)		(x-table)	Ge # 2
⁶⁰ Co	1173	4.74E+06	1332	5222	1	1.10E-03
⁶⁰ Co	1332	4.29E+06	1173	5055	0.99945	1.18E-03
²⁰⁷ Bi	570	1.20E+07	1063	11177	8.42E-01	1.11E-03
²⁰⁷ Bi	570	1.20E+07	1770	801	6.91E-02	9.70E-04
207 Bi	1063	5.28E+06	570	9238	1	1.75E-03
²⁰⁷ Bi	1770	3.41E+05	570	581	1	1.70E-03
¹³⁴ Cs	795	1.25E+06	604	2076	1	1.66E-03
¹³⁴ Cs	795	1.25E+06	569	364	1.81E-01	1.61E-03
¹³⁴ Cs	605	2.05E+06	563	318	9.00E-02	1.72E-03
¹³⁴ Cs	605	2.05E+06	796	2506	8.69E-01	1.40E-03

Table 27 On-Line data pertinent to the efficiency calculation of the germanium detectors.

Εγ	Branch	# of βs	# of γs	Gel	Ge2	εGe1	εGe2
keV	%	(/ le7)		Counts	Counts	x1000	x1000
19	$O \rightarrow {}^{19}F^*$	+ β-					
110	2.54(10)	4.4(2)	1.12(6)E6	3.1(1)E3	N/A	2.77(15)	N/A
197	95.9(21)	4.4(2)	4.2(3)E7	9.96(4)E4	3.64(2)E4	2.36(12)	0.86(5)
1357	50.4(11)	4.4(2)	2.2(2)E7	5.57(3)E4	2.54(2)E4	2.51(13)	1.14(6)
1444	2.64(6)	4.4(2)	1.16(6)E6	2.84(8)E3	1.22(5)E3	2.44(14)	1.05(7)
1554	1.39(3)	4.4(2)	6.1(4)E5	1.77(7)E3	7.1(4)E2	2.89(18)	1.16(9)
17	$N \rightarrow {}^{17}O*$	+ β-					
871	3.34(50)	2.3(1)	7.7(12)E5	2.58(6)E3	1.15(4)E3	3.36(53)	1.50(24)
2184	0.34(6)	2.3(1)	7.8(15)E4	150(14)	7.2(10)E1	1.92(44)	0.92(21)

3.7.9 EFFICIENCY CALIBRATION – GERMANIUM COINCIDENCES

In addition to the on-line calibrations described in the previous section, it was possible to generate efficiency calibration points for Ge#1 at low energy using a coincidence technique. The energy levels of the ¹⁹O and ¹⁹F daughters are well known and some of the properties of the decay of each level are also known or may be inferred. For example, the observation of a 1376 keV line during the decay of ¹⁹N indicates the transition from the 1472 keV excited state in ¹⁹O to the 96 keV excited state. It must be true that for every 1376 keV gamma ray there is a corresponding emission of a 96 keV

gamma ray. Therefore, the efficiency of Ge#1 at 96 keV can be determined by gating on the 1376 keV line in Ge#2 and reflecting back onto the Ge#1 energy spectrum. The efficiency of Ge#1 at 96 keV will be equal to the number of observed 96 keV counts in the gated spectrum divided by the total number of 1376 keV counts in Ge#2. The statistics of such an analysis are necessarily poor due to the multiplicative aspect of the germanium efficiencies. However, in the absence of another method to measure the efficiency of at least one of the germanium detectors at 96 keV, the poor statistics will have to be absorbed. Table 28 gives the results of the germanium coincidence analysis and the calculated efficiency at the points taken.

Table 28 Germanium detector efficiencies determined using the coincidence technique.

Ge#2 Peak	Ge#1 Peak	Ge#1 Peak	Ge#2 Peak	Efficiency at
Gating (keV)	Energy (keV)	Area	Area	Ge#1 Energy
3849	96	14(3.7)	6604(84)	2.12(57)E-3
1376	96	18(4.3)	10412(109)	1.73(41)E-3
1357	197	82(9.1)	29080(175)	2.82(32)E-3
1444	110	2(1.4)	1384(72)	1.45(102)E-3
2473	96	11(3.3)	6401(85)	1.72(52)E-3
2473	1376	16(4)	6401(85)	2.50(63)E-3

3.7.10 EFFICIENCY CALIBRATION – SUMMARY

Taking the data acquired from the on-line and off-line sources in concert, efficiency curves for detectors Ge1 and Ge2 were constructed. Figure 45 shows the comprehensive raw efficiency data for detector Ge2.

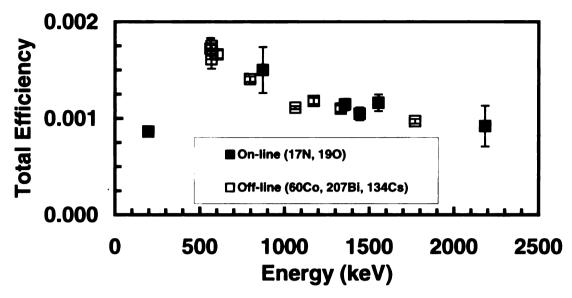


Figure 45 Data points taken for the efficiency determination of Ge#2.

The data above 500 keV were fit to a functional form given in [88De]. The equation of the fit is given as:

$$\varepsilon = c \cdot (E/Eo)^{-\delta}$$
 [9]

where ε is the efficiency, c is a dimensionless variable parameter, E is the gamma-ray energy in keV, Eo is a constant set equal to 1 keV to eliminate the energy dimension, and δ is a constant based on the volume of the germanium crystal that is given by:

$$\delta = 2.14 - 0.629 \log V \tag{10}$$

Where V is the volume of the germanium detector in cm³. In this experiment, the volume of detector Ge1 was 542 cm³ and the volume of Ge2 was 411 cm³ where these quantities were deduced from the specification sheets provided by the manufacterer [00Ro]. Figure 46 shows the same calibration data for Ge2 and the best fit to the efficiency data using the above formulae.

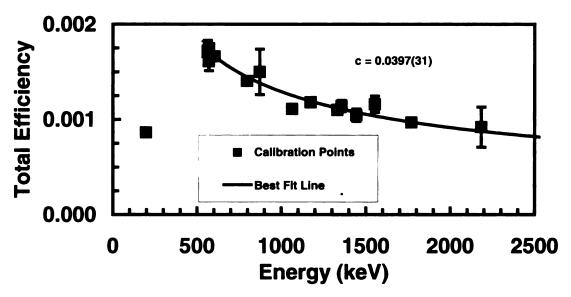


Figure 46 Best fit to the efficiency calibration data for Ge#2.

Since there is only one variable in the fit and therefore no correlation problem, the error on parameter c was deduced by the change in its value necessary to produce an increase in the total chi-squared of one unit. This error is valid over the interpolation region ranging from about 560 keV to 2180 keV in energy. To estimate the error associated with extrapolating the fit to higher energies, the high efficiency at 560 keV (c + σ) and the low efficiency at 2180 keV (c - σ) were fit to the same functional form. This was repeated for the low efficiency at 560 keV (c - σ) and the high efficiency at 2180 keV (c + σ). The two curves diverge above the interpolation range and hence give significantly higher error on the efficiency in the high-energy extrapolation region. Figure 47 shows the efficiency data for Ge2, the best-fit line to the data, the error range in the interpolation region, and the error range in the extrapolation region. It is noted that the percent error in the interpolation region is roughly 8% of the efficiency whereas at 3200 keV, the percent error has increased to approximately 14% of the efficiency.

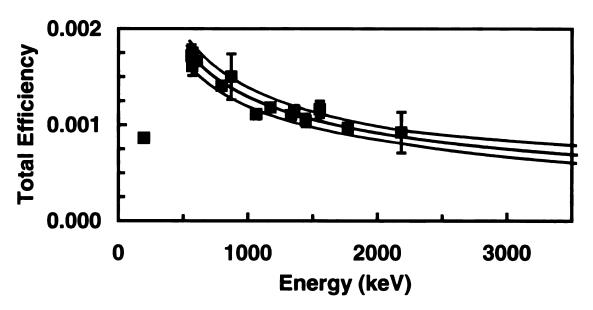


Figure 47 Interpolation and extrapolation error bars for the efficiency determination of Ge#2.

The calibration data points cover a range having an upper limit of 2200 keV or thereabouts. As a consequence, the efficiency calibration for the germanium detectors is suspect for the higher energy gamma rays (>3.0 MeV) that were observed from the decay of ¹⁹N (see figures 36, 37). The efficiencies at these gamma ray energies would be extrapolations over an extremely large span. However, the reliability of the extrapolation was checked at a somewhat higher energy by comparing the extrapolation against data recorded by the same germanium detectors during a different NSCL experiment [00Pr] having a calibration point at 2615 keV. The shapes of the calibration curves above 500 keV and including the 2615 keV point were found to be in excellent agreement. This agreement between the experiments improved the confidence for efficiency extrapolation up to at least 2600 keV.

3.7.11 GAMMA RAY INTENSITIES – ¹⁹N

The branching ratios for gamma ray emission as a function of energy can be calculated for those transitions that were assigned as following the beta decay of ¹⁹N. The integrated area of the gamma lines divided by the detector efficiency leads to the total number of gamma rays emitted over the course of the data runs. This value divided by the total number of decays (from the integrated half-life spectra) of the respective isotopes gives the intensity of the lines per decay of the parent isotope. Table 29 gives the energy of the gamma line, number of decays observed in each of Ge1 and Ge2, the efficiency of each detector at that energy, the total number of source decays, and the calculated intensity of each gamma line.

Table 29 Gamma-ray intensities for transitions assigned to the beta decay of ¹⁹N.

Energy	Gel	Ge2	β-Source	εGel	εGe2	Iγ Ge l	Iγ Ge2
(keV)	Counts	Counts	(x 1e-7)	(x 1000)	(x 1000)	(x 1000)	(x 1000)
96	25155	N/A	4.64(10)	1.86(51)	N/A	291(80)	N/A
821	1220	579	4.64(10)	2.75(22)	1.42(11)	10(1)	9(1)
1374	12455	5571	4.64(10)	2.22(20)	1.10(9)	121(13)	109(11)
1472	824	330	4.64(10)	2.15(18)	1.06(9)	8(1)	7(1)
1652	1426	635	4.64(10)	2.05(15)	1.01(7)	15(2)	14(2)
1938	845	394	4.64(10)	1.91(14)	0.93(7)	10(2)	9(2)
1980	15043	6735	4.64(10)	1.89(14)	0.92(7)	172(15)	158(14)
2473	7946	3513	4.64(10)	1.73(15)	0.82(8)	99(10)	92(10)
3136	3617	1602	4.64(10)	1.56(19)	0.72(9)	50(7)	48(7)
3849	8454	3614	4.64(10)	N/A	N/A	N/A	N/A
3945	1249	611	4.64(10)	N/A	N/A	N/A	N/A

3.7.12 INTENSITIES OF HIGHER ENERGY ¹⁹N GAMMA RAYS

For the case of the 3232 keV gamma line, there is no alternative to the efficiency determination other than the extrapolation. Therefore, the entry in table 29 for the 3232 keV line is based on that portion of the analysis even though it is significantly higher in energy than the more reliable interpolation range. However, the 3945 keV level in ¹⁹O has been populated in the past and the subsequent de-excitation of this level has been monitored [70Fi, 71Hi]. From the independence hypothesis, the de-excitation of a level should not be a function of the manner by which it is populated – provided there is no polarization component. Therefore, since the de-excitation gamma rays and their intensities have been recorded in the past, this provides an opportunity to determine the intensities by other means. Table 30 lists the data from the literature relevant to the relaxation of the 3945 keV level.

Table 30 Literature values for the de-excitation branching of the 3945 keV excited state of ¹⁹O.

Initial Level	Final Level	Gamma Energy	Branch (%)	Branch (%)
(keV)	(keV)	(keV)	[70Fi]	[71Hi]
3945	0	3945	24(8)	33(8)
3945	96	3849	48(8)	39(8)
3945	1472	2473	28(4)	28(4)

The detection efficiency for the 2473 keV line is considered reliable due to the fact that this energy is below 2615 keV which was the highest energy point taken as a calibration line for the germanium detectors. Accepting the efficiency at 2473 keV as

valid (and thus the calculated intensity), and utilizing the relative de-excitation strength of the 3945 keV level from the literature, extrapolating the efficiency of the germanium detectors for the 3945 and 3849 keV levels is obviated. Table 31 gives the calculated intensities for the 3945 and 3849 keV levels on the basis of the literature data and the subsequent beta decay branching ratio to the 3945 keV level.

Table 31 Calculated intensities of the de-excitation gamma rays from the 3945 keV excited state of ¹⁹O.

Ge #1	Ιγ 2473 keV	Ιγ 3945	Ιγ 3849
	(fixed)	Calculated	Calculated
Ref. [70Fi]	99(10)	85(32)	170(41)
Ref. [71Hi]	99(10)	117(35)	138(37)
Ge #2			
Ref. [70Fi]	92(10)	79(30)	158(39)
Ref. [71Hi]	92(10)	108(33)	128(35)

The compilation by Tilley *et al.* of the previously published experimental results, states that the de-excitation values given in [71Hi] are considered to be more reliable that that of reference [70Fi]. Due to the lack of confidence in the efficiency extrapolation, the value quoted for the intensity of the de-excitation gamma rays at 3945 and 3849 keV from the bound excited state at 3945 keV will be that derived on the basis of the experimental results in [71Hi]. Table 32 lists the final values for the absolute intensity of the gamma rays ascribed as being due to the decay of ¹⁹N.

Table 32 Final gamma-ray intensities for those gamma rays attributed to the decay of ¹⁹N.

Energy	Iγ Ge l	Iγ Ge2
(keV)	(x 1000)	(x 1000)
96	291(80)	N/A
821	10(1)	9(1)
1374	121(13)	109(11)
1472	8(1)	7(1)
1652	15(2)	14(2)
1938	10(2)	9(2)
1980	172(15)	158(14)
2473	99(10)	92(10)
3136	50(7)	48(7)
3849	138(37)	128(35)
3945	117(35)	108(33)

3.7.13 GAMMA RAY INTENSITIES - 20N

Due to the low production rate of ²⁰N relative to the other RIBs, the potential for missing gamma rays during the beam off period was substantial. To limit this effect, the relative intensities of the gamma rays ascribed to the decay of ²⁰N were determined using the data collected in the germanium detectors over all times. The inherent assumption in this portion of the analysis was that the relative efficiency of the germanium detectors as a function of energy is not affected by monitoring over all times.

Table 33 lists the peak areas, efficiencies, and calculated relative intensities for gamma rays observed in each of the germanium detectors over all times and assigned to the decay of ²⁰N. The intensities are relative to the 1674 keV line which is set to one hundred.

Table 33 Calculated relative intensities of gamma rays emitted during the decay of ²⁰N. The intensities are relative to the 1674 keV line that is set to 100.

Εγ	Ge #1	Ge #2	ε-Ge #1	ε-Ge #2	Iγ Ge #1	Iγ Ge #2
keV	Counts	Counts	(x 1000)	(x 1000)	Relative	Relative
96	968(32)	N/A	1.86(51)	N/A	88(24)	N/A
1374	234(39)	93(32)	2.22(20)	1.10(9)	18(3)	15(5)
1674	1206(45)	565(26)	2.05(15)	1.00(7)	100(8)	100(8)
1814	74(13)	48(15)	1.97(20)	0.96(8)	6(1)	9(3)
1895	174(37)	74(12)	1.93(18)	0.94(8)	15(4)	14(3)
2394	155(26)	59(10)	1.76(16)	0.84(9)	15(3)	12(3)
2782	47(19)	12(5)	1.65(15)	0.78(9)	5(2)	4(2)

The results for the relative intensities of the lines are in close agreement between the two germanium detectors. The gamma lines at 96 and 1374 keV are characteristic of transitions in ¹⁹O. These gamma rays indicate beta delayed neutron emission from ²⁰N leading to excited levels of ¹⁹O rather than to the ground state.

To determine the absolute intensities, the stronger peaks that were integrable in the beam-off spectra collected by the germanium detectors were analysed. This permits the calculation of the absolute intensity at these energies due to the fact that it is during this time period when the beta decay of ²⁰N can be monitored quantitatively. Also, the half-life gated on the 96 and 1374 keV lines as they appear in the beam-off spectra confirm them as being due to the decay of ²⁰N. Table 34 gives the absolute intensities for some of the ²⁰N gamma ray lines using this method.

Table 34 Absolute intensity of gamma rays ascribed to the decay of ²⁰N.

Εγ	Ge #1	Ge #2	20 N β -	ε-Ge #1	ε-Ge #2	Iγ Ge #1	Iγ Ge #2
keV	Counts	Counts	/10 ⁵	(x 1000)	(x 1000)	(x 1000)	(x 1000)
96	347(21)	N/A	7.6(1)	1.86(51)	N/A	246(69)	N/A
1374	74(16)	40(9)	7.6(1)	2.22(20)	1.10(9)	44(11)	48(12)
1674	421(23)	205(16)	7.6(1)	2.05(15)	1.00(7)	270(25)	270(29)
1895	32(8)	N/A	7.6(1)	1.93(18)	0.94(8)	22(6)	N/A
2394	57(15)	N/A	7.6(1)	1.76(16)	0.84(9)	43(12)	N/A

Taking the relative gamma intensities in concert with the smaller sub-set of measured absolute intensities, the absolute intensities for all of the lines for both detectors can be calculated. Since the 1674 keV line has the lowest error, this line's intensity was utilized to calculate the absolute intensity of the others. Table 35 gives the final evaluated intensities of the gamma lines due to 20 N.

Table 35 Intensity of the gamma-ray lines attributed to the decay of ²⁰N.

Eγ keV	Iγ Ge #1 Measured	Iγ Ge #1 Calculated	Iγ Ge #2 Measured	Iγ Ge #2 Calculated	Ιγ Final
	(x 1000)	(x 1000)	(x 1000)	(x 1000)	(x 1000)
96	246(69)	-	-	N/A	246(69)
1374	44(11)	-	48(12)	-	46(9)
1674	270(25)	-	270(29)	-	270(19)
1814	-	16(4)	-	24(9)	18(4)
1895	22(6)	-	-	38(10)	26(6)
2394	43(12)	-	-	32(10)	37(8)
2782	-	13(6)	-	11(5)	12(4)

CHAPTER 4

INTERPRETATION

4.1 INTERPRETATION OF ¹⁹N GAMMA RAYS

To determine the branching ratios for the beta decay of ¹⁹N to bound excited states of ¹⁹O, the transitions listed in table 32 of section 3.7.12 were assigned as either being due to transitions between states in ¹⁹O or between states in ¹⁸O. Table 36 lists those transitions assigned to ¹⁹O along with their intensities and the characteristic levels of the transition. The intensities are given per 1000 parent decays.

Table 36 Gamma-ray transitions identified as being a result of ¹⁹N beta decay to bound excited states of ¹⁹O.

Energy	¹⁹ O Initial	¹⁹ O Final	Iγ Ge l	Iγ Ge2
(keV)	Level (keV)	Level (keV)	(x 1000)	(x 1000)
96	96	0	291(80)	N/A
1374	1472	96	121(13)	109(11)
1472	1472	0	8(1)	7(1)
2473	3945	1472	99(10)	92(10)
3136	3232	96	50(7)	48(7)
3849	3945	96	138(37)	128(35)
3945	3945	0	117(35)	108(33)

From these data, it is possible to construct an energy level diagram illustrating the observed transitions. This will show which gamma rays must be in coincidence with each other and permit a determination of which levels are fed directly from the

beta decay and which gamma rays are consequences of cascades. Figure 48 shows the known energy level scheme for the ground state and bound excited states of ¹⁹O. The levels are given in keV relative to the ground state at zero and are adapted from reference [95Ti].

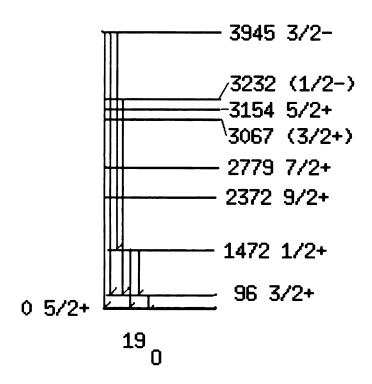


Figure 48 Partial energy level diagram of ¹⁹O showing the bound-state transitions observed in this experiment during the decay of ¹⁹N.

To deduce the branching ratios for the beta decay of ¹⁹N to bound states of ¹⁹O, the intensities of those transitions that are not fed by a cascade remain unchanged. Those transitions that may be due to both beta feeding and as a result of one or more gamma-ray cascades require that their intensities be adjusted for the cascade feeding. In this case, the only transitions that may be due to both beta feeding and cascade feeding are the 1472-0, 1472-96, and 96-gs transitions. Unfortunately, the experimental

conditions prevented a determination of the germanium detector efficiency near 96 keV for detector Ge#2. Consequently, there can be no redundancy for the calculation of the beta feeding to the 96 keV level during the decay of ¹⁹N.

From the observed intensities and the derived decay scheme, the branching ratios for pure beta feeding to bound excited states of ¹⁹O can be calculated. The sum of the three transitions proceeding from the 3945 keV level is a measure of the beta strength to this level. The intensity of the 3232-1472 transition is a measure of the beta decay branching ratio to the 3232 keV level, and the sum of the 1472-gs and 1472-96 transitions, less the feeding through cascades, is a measure of the beta decay to the 1472 keV excited state. Similarly, the intensity of the 96 keV gamma ray less the cascade feeding from multiple sources will be a measure of the beta branch to the 96 keV level. Table 37 gives the calculated direct beta-decay branching ratios to bound excited states of ¹⁹O where soluble. The last row of the table gives the total branching ratio to bound excited states of ¹⁹O above and not including the 96 keV level.

Table 37 Summary of beta decay strength to individual bound levels in ¹⁹O from the decay of ¹⁹N.

Beta Decay Level	Branching Ratio (%)	Branching Ratio (%)	B.R. (%)
Populated Directly	Germanium #1	Germanium #2	Mean Value
3945 3/2-	35.4(52)	32.9(51)	34.2(37)
3232 (1/2-)	5.0(10)	4.8(7)	4.9(6)
1472 1/2+	0.30(18)	0.24(16)	0.27(12)
96 3/2+	< 4.3	N/A	<4.3
Ground State 5/2+	N/A	N/A	N/A
TOTAL B.R.	40.7(54)	38.0(53)	39.4(38)

The branching ratio determination for direct beta feeding to the ¹⁹O first excited state at 96 keV is given as an upper limit. The branching ratio calculated from the 96 keV line's intensity less the feeding is consistent with zero. The upper limit represents subtracting the minimum possible cascade contribution (mean less one sigma) from the maximum possible intensity (mean plus one sigma) of the 96 keV line. This experiment was not sensitive to the beta decay of ¹⁹N feeding the ground state of ¹⁹O directly.

4.2 INTERPRETATION OF ¹⁹N DELAYED NEUTRONS

Between the neutron bars and the ⁶Li detectors, the experiment was able to record twelve beta-delayed neutrons emitted during the decay of ¹⁹N. Table 38 summarizes the observed delayed neutrons and their calculated branching ratios.

Table 38 Summary of observed delayed-neutron transitions from ¹⁹N by energy.

Neutron Energy (MeV)	B.R. (%) from Array	B.R. (%) from ⁶ Li glass
4.91(8)	0.90(10)	-
4.00(6)	0.37(4)	-
3.06(5)	1.97(32)	-
2.84(4)	9.6(11)	-
2.49(4)	3.5(4)	-
2.21(4)	2.2(4)	-
1.27(3)	1.9(2)	-
1.13(3)	16.2(17)	18.2(23)
0.83(3)	2.0(3)	-
0.62(3)	2.2(3)	-
0.49(3)	8.7(10)	8.6(10)
0.30(8)	-	1.4(3)

To determine the beta branching to specific excited levels of ¹⁹O that subsequently emit a neutron, it is necessary to determine if the neutron emission populates the ¹⁸O ground state or one of the excited states. There is about 8.5 MeV available for delayed neutron emission and over twenty excited states in ¹⁸O that could possibly be populated following the emission of a neutron from an excited level of ¹⁹O.

The analysis of the gamma-ray data collected by the germanium detectors showed that there were four gamma rays characteristic of transitions in ¹⁸O. These gamma rays could be indicative of beta-delayed neutron emission populating excited states of ¹⁸O or they could simply be due to the beta decay of an ¹⁸N contaminant though there is no evidence of this (see table 3). If the ¹⁸O gamma rays are due to ¹⁹N delayed neutron emission then the corresponding half-lives of the gamma rays should be equal to the half-life of ¹⁹N (~300ms) and if the ¹⁸O gamma rays are due to the beta decay of ¹⁸N then the gated half-lives will reflect the half-life of ¹⁸N (~620ms). Table 39 gives the measured half-lives of the four ¹⁸O gamma rays and the intensity of the lines for the measurements made by each of the germanium detectors. The intensities of the gamma rays are given per 1000 ¹⁹N decays.

Table 39 Gamma rays characteristic of de-excitation in ¹⁸O. The half-lives of the gamma rays are listed along with the intensity of the lines assuming the ¹⁹N parent.

Eγ keV	E _i level	E _f level	Half-Life (ms)	Iγ Ge #1	Iγ Ge #2	Ιγ Mean
1982	1982	0	300(10)	172(15)	158(14)	166(10)
1938	3920	1982	273(17)	10(2)	9(2)	10(2)
1652	3634	1982	305(14)	15(2)	14(2)	15(2)
821	4456	3634	364(45)	10(1)	9(1)	10(2)

On the basis of the measured gamma-ray half-lives, it may be concluded that the observed 18 O gamma rays are indeed due to the decay of 19 N and thus reflect delayed neutron emission to excited states of 18 O. Accounting for the cascade feeding in the same manner as outlined in section 4.1, and incorporating level schemes from [95Ti], the branching ratios for delayed neutron emission populating the specific excited states of 18 O may be calculated. Table 40 gives the total strength of the (β +n) feeding of the 18 O excited levels.

Table 40 Values for the $(\beta + n)$ feeding strength to each of the pertinent ^{18}O excited states.

¹⁸ O Excited State Populated	Beta + Neutron Feeding Strength
(keV)	(% of ¹⁹ N decays)
1982	13.7(11)
3634	0.1(3)
3920	1.2(3)
4456	1.5(4)

To determine the beta feeding to specific levels in ¹⁹O prior to the neutron or gamma ray emission, it is desirable to find which delayed neutrons are in coincidence with the ¹⁸O gamma rays that were observed. An effort was made to measure, directly, which neutrons and gammas correspond to the same transition by gating on the gamma rays of table 40 and reflecting back onto the (summed) delayed neutron spectrum recorded by the neutron bar array. Table 41 gives the results of the coincidence measurement for each of the gamma rays gated.

Table 41 Neutrons in coincidence with selected ¹⁸O gamma rays. Listed are those neutrons in coincidence and those neutrons that may be in coincidence on the basis of the gated spectrum.

Gamma-Ray Gate (keV)	Neutron in Coincidence (keV) Definite	Neutron in Coincidence (keV) Possible
1982	2840	490
	830	2210
		1270
1653	1270	
1938	830	
821	None	None

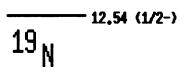
On the basis of these results, it can be concluded that the 830 keV neutron is emitted from a level in ¹⁹O that populates the 3920 keV excited state of ¹⁸O due to the fact that this neutron appears to be in coincidence with both the 1938 and 1982 keV gamma lines. Also, the 1270 keV neutron appears to be in coincidence with the 1653 keV gamma ray – suggesting that the beta delayed neutron populates the 3634 keV excited state or perhaps the 4456 keV excited state. It is not possible to distinguish between the two given the statistics of the (821 keV gamma ray gated) neutron spectrum. Also, the strong 2840 keV neutron is in coincidence with the 1982 keV gamma ray suggesting that this neutron is emitted by a level in ¹⁹O that populates the first excited state of ¹⁸O. The 2210 and 490 keV neutrons may or may not be in coincidence with the 1982 keV gamma line. The low statistics of the coincidence spectra prevent conclusive assignment of these neutrons to the gamma ray.

4.3 DELAYED NEUTRON ASSIGNMENTS 19N

From the delayed neutron branching ratios, observed ¹⁸O gamma rays and the measured neutron + gamma ray coincidences, a cross table for the possible delayed

neutron transitions may be constructed (Table 42). The cross table cells represent the energy of the intermediate ¹⁹O level populated by the beta decay and emitting the neutron. The columns define the level of ¹⁸O populated following the emission of the neutron. The energies are given in keV for all entries and the cell values are relative to the ground state of ¹⁹O. Those cells that are blank have been eliminated as possible transitions either on the basis of the total energy (greater that that available for delayed neutron emission) or on the basis of the neutron branching ratio relative to the ¹⁸O gamma-ray intensities. Since there is no evidence of any ¹⁸O levels being populated other than those listed in table 41, the cross table (42) will be limited to those excited states plus the ¹⁸O ground state.

Of the twelve delayed neutron transitions, only five can be conclusively assigned to specific levels in ¹⁹O. These delayed neutron energies and corresponding ¹⁹O level are underlined in the cross table. The remaining seven cannot be fixed as they may proceed through different intermediate states on the basis of the measured data. From the delayed neutron branching ratios and the definitive assignments it can be concluded that the small possible branch for delayed neutrons populating the 3634 keV excited level in ¹⁸O is either in fact zero (with which the measurement is consistent) or is a result of a delayed neutron that was not measured in our experiment. Figure 49 shows the transitions due to the beta decay of ¹⁹N that can be conclusively identified on the basis of the data recorded in this experiment. The energies are given in keV and are relative to the ground state of ¹⁹O. The excited levels in ¹⁸O are relative to the ¹⁸O ground state.



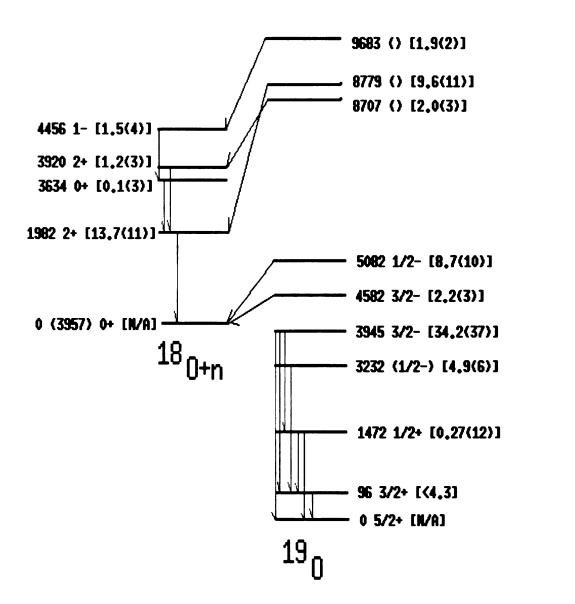


Figure 49 The beta decay of ¹⁹N showing the transitions that can be ascribed to ¹⁹N on the basis of the experimental data.

Table 42 Cross table for the assignment of intermediate states of $^{19}{\rm O}$ populated during the decay of $^{19}{\rm N}$.

E-Neutron	E- ¹⁹ O level				
C.o.M.	(¹⁸ O gs)	(¹⁸ O 1982)	(¹⁸ O 3634)	(¹⁸ O 3920)	(¹⁸ O 4456)
4910	8867	10849			
4000	7957	9939			
3060	7017	8999			
<u>2840</u>		<u>8779</u>			
2490	6447	8429			
2210	6167	8149			
<u>1270</u>					<u>9683</u>
<u>1130</u>	<u>5087</u>				
<u>830</u>				<u>8707</u>	
<u>620</u>	<u>4577</u>				
490	4447	6429			
300	4257	6239			

4.4 INTERPRETATION OF ²⁰N GAMMA RAYS

To determine the branching ratios for the beta decay of ²⁰N to bound excited states of ²⁰O, the transitions labeled in table 32 of section 3.7.12 were assigned as either

being due to transitions between states in ²⁰O or between states in ¹⁹O. Table 43 lists those transitions indicative of beta decay to bound states of ²⁰O.

Table 43 Gamma-ray transitions identified as being a result of ²⁰N beta decay feeding bound excited states of ²⁰O.

Gamma Energy	²⁰ O Initial	²⁰ O Final	Intensity
keV	Level	Level	(x 1000)
1674	1674	0	270(19)
1814	5387	3570	18(4)
1895	3570	1674	26(6)
2394	4072	1674	37(8)
2782	4456	1674	12(4)

From these data it is possible to construct and energy level diagram illustrating the observed transitions among the known ²⁰O levels. As with the ¹⁹N data, this will make it possible to determine the cascade feeding and thus lead to an assessment of the branching ratios for the feeding of specific levels in ²⁰O during the beta decay. Figure 50 shows the gamma transitions in ²⁰O observed in this experiment.

From the observed intensities and the derived decay scheme, the branching ratios for pure beta feeding to specific bound levels in ²⁰O can be calculated. Table 44 gives the result of the branching ratio determination for the decay of ²⁰N to bound excited levels in ²⁰O.

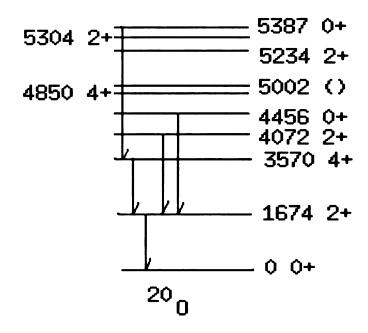


Figure 50 Partial energy level diagram of 20 O showing the bound-state transitions observed in this experiment during the decay of 20 N.

Table 44 Bound excited levels of ²⁰O populated during the decay of ²⁰N. Listed are the branching ratios for the beta transitions as determined from the measured gamma-ray data.

²⁰ O Excited Level Populated Directly	Branching Ratio of Direct β-Feeding (Percent)	
Energy of Level (keV)		
5387 0+	1.8(4)	
4456 0+	1.2(4)	
4072 2+	3.7(8)	
3570 4+	0.8(8)	
1674 2+	19.5(22)	
G.S. 0+	N/A	
Total Branching Ratio	27.0(46)	

4.5 INTERPRETATION OF ²⁰N NEUTRON SPECTRUM

Analysis of the delayed neutron spectrum collected by the neutron bar array suggests eight beta delayed neutrons emitted by the ²⁰N isotope. Table 45 lists the energy of those delayed neutrons and their corresponding branching ratios. Taking the data together, the delayed neutron emission probability from ²⁰N is 34.0(40) percent of the decay strength for delayed neutrons above 1.1 MeV in energy. The low rate and decreasing detection efficiency of the array below this cut off prevent measurement of delayed neutrons at lower energy.

Table 45 Energy and branching ratio for delayed neutrons observed during the decay of 20 N. Also given is the P_n value for delayed neutrons emitted above 1.1 MeV in energy.

Energy of ²⁰ N Delayed Neutron	Branching Ratio
(Center-of-Mass MeV)	(Percent)
1.16(3)	4.5(5)
1.33(3)	2.5(3)
1.63(3)	8.3(9)
2.19(4)	12.1(13)
2.63(4)	1.3(2)
3.38(5)	2.1(3)
3.88(6)	2.7(3)
4.64(7)	0.5(1)
TOTAL	34.0(40)

The gamma-ray spectra collected during the decay of ²⁰N showed two gamma rays at 96 and 1374 keV - which are characteristic of transitions in ¹⁹O. The first a

transition from the 96 keV first excited state to the ground state and the second a transition from the 1472 keV second excited state to the 96 keV level. These gamma rays were shown to have half-lives consistent with that of ²⁰N indicating that there is beta delayed neutron emission from ²⁰N populating excited levels in ¹⁹O rather than the ground state. The branching ratios for these transitions, with cascade feeding taken into account, are 20.0(70) and 4.6(9) percent for the 96 and 1374 keV transitions, respectively.

With regard to delayed neutron emission from ²⁰N, it is not possible to assign beta decay branching ratios to specific excited levels of ²⁰O above the neutron emission threshold. This is due to the following considerations: first, several possibilities exist for the final ¹⁹O level populated rendering the evaluation of the level energy inconclusive, second, the implantation rate of ²⁰N coupled with the detection efficiencies prevent neutron-gamma coincidence measurements that could restrict the ²⁰O level energy, third, only seven ²⁰O excited levels above the neutron emission threshold are known with but four of these being above the lowest experimentally observed neutron energy, and fourth, the extent of the above-threshold levels is below 10.5 MeV relative to the ground state of ²⁰O while the Q-value is about 18 MeV.

The one exception to this is the 2.19 MeV delayed neutron peak. The intensity of this transition is strong enough that the coincidence between it and the 96 keV should have been observed. The spectrum, however, does not indicate a coincidence between the two - suggesting that the delayed neutron emission populates the ground state of ¹⁹O rather than the 96 keV excited state. Furthermore, one of the known ²⁰O levels is at 9.77 MeV (~2.17 MeV above the neutron emission threshold) which is consistent in

energy with the 2.19 MeV neutron observed. Thus, this transition is tentatively assigned as proceeding through the 9.77 MeV level in ²⁰O with a branching ratio of 12.1(13) percent. Figure 51 shows a portion of the mass 20 level scheme where the transitions resulting from the decay of ²⁰N that can be definitively assigned are presented. The energies of the levels in ²⁰O are given in keV relative to the ²⁰O ground state and the excited levels of ¹⁹O are given in keV relative to the ¹⁹O ground state.

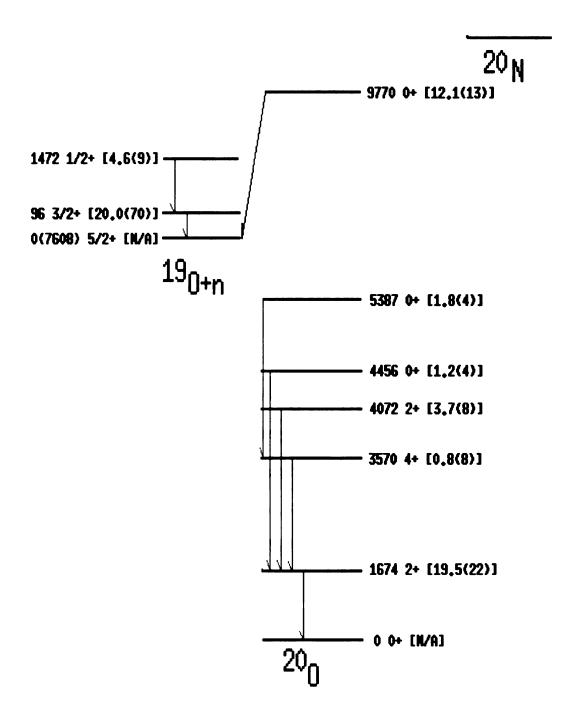


Figure 51 The beta decay of ²⁰N showing the transitions that can be conclusively identified on the basis of the experimental data.

4.6 19N SHELL MODEL CONSIDERATIONS

From the energies of the ¹⁹O intermediate levels that may be populated directly in the ¹⁹N beta decay and the measured branching ratios of the transitions, the log(ft) values of the specific transitions can be calculated. The log (ft) values provide insight into the nature of the decay (allowed etc...) and also lead to a determination of the B(GT) transition strength. The log (f) values are taken from the literature [71Go] while the t value is simply equal to the half-life of ¹⁹N in seconds divided by the measured branching ratio of the specific channel expressed as a transition probability (BR/100). The B(GT) strength is equal to (6177/ft).

Table 46 lists all transitions observed in this experiment (delayed neutron as well as beta decay to bound states of 19 O), the experimentally derived transition strength, the log (ft) values and the calculated B(GT) strength. The column labeled "transition identifier" gives the energies of the delayed neutrons for the (β +n) transitions while "to bound" indicates a transition to the bound 19 O level listed in the second column. For those cases where the delayed-neutron emitting state is not conclusive, all possibilities are shown based on the delayed neutron energy.

Shell model calculations to predict the Gamow-Teller beta decay strengths of ¹⁹N to levels in ¹⁹O were calculated by Warburton in reference [88Wa]. In the model, the even parity states of ¹⁹O were calculated within the (0s)⁴(0p)¹²(2s,1d)³ model space, the ¹⁹N states were assumed to arise from the (0s)⁴(0p)¹¹(2s,1d)⁴ model space, and the odd parity ¹⁹O levels required consideration of both the (0s)⁴(0p)¹¹(2s,1d)⁴ component and the (0s)⁴,(0p)¹²(2s,1d)²(0f,1p)¹ component to account for all possible 1hω excitations. The model uses a modified Millener-Kurath interaction to calculate the

odd-parity levels in ¹⁹O and the Chung-Wildenthal or Universal (2s1d) interaction for the low-lying even-parity levels.

Table 46 Calculated log(ft) and B(GT) values for the possible beta transitions in the decay of ¹⁹N. The transition probability is the branching ratio divided by 100.

Transition	¹⁹ O level	Transition	Log (ft)	B(GT)
Identifier	Populated	Probability		
(β+n) 4910	8867	0.009	4.65	1.37E-01
(β+n) 4910	10849	0.009	3.20	3.87
$(\beta + n) 4000$	7957	0.0037	5.49	2.00E-2
$(\beta + n) 4000$	9939	0.0037	4.39	2.52E-1
$(\beta + n) 3060$	7017	0.0197	5.13	4.55E-2
$(\beta + n) 3060$	8999	0.0197	4.25	3.45E-1
$(\beta + n) 2840$	8779	0.0964	3.68	1.28
$(\beta+n) 2490$	6447	0.0349	5.08	5.09E-2
$(\beta + n) 2490$	8429	0.0349	4.30	3.07E-1
$(\beta+n)$ 2210	6167	0.0224	5.37	2.65E-2
$(\beta + n) 2210$	8249	0.0224	4.57	1.67E-1
$(\beta+n)$ 1270	9683	0.0189	3.87	8.32E-1
(β+n) 1130	5087	0.1618	4.83	9.18E-2
(β+n) 830	8707	0.0202	4.40	2.45E-1
(β+n) 620	4577	0.0215	5.83	9.04E-3
(β+n) 490	4447	0.0869	5.27	3.33E-2
$(\beta + n) 490$	6429	0.0869	4.70	1.24E-1
$(\beta + n) 300$	4257	0.014	6.11	4.78E-3
$(\beta + n) 300$	6239	0.014	5.55	1.74E-2
to bound	3945	0.342	4.79	9.95E-2
to bound	3232	0.049	5.81	9.64E-3
to bound	1472	0.0027	7.39	2.54E-4
to bound	96	0.043	6.42	2.33E-3
to bound	0	n/a	n/a	n/a

Table 47 compares the transitions observed in this experiment with the predictions of the shell model calculations of Warburton. The table, where necessary, identifies transitions by subscript that cannot be conclusively ascribed to the specific level in ¹⁹O. Tabulated are the branching ratio, log(ft), B(GT), and level spins. The B(GT) values have been multiplied by a factor of 1000.

Table 47 Comparison of the ¹⁹O energy levels from this experiment with predictions made using the shell model [88Wa]. The subscripts on the energies indicate the choices for the ¹⁹O level due to the uncertainty in the neutron decay.

	This	s Experir	nent	<u> </u>		Shell M	1odel Pre	dictions	
Ex	Jπ	BR%	Logft	B(GT)	Ex	Jπ	BR%	Logft	B(GT)
96	3/2+	<4.3	6.42	2.3	96	3/2+	0.49	7.30	
1472	1/2+	0.27	7.39	0.25	1472	1/2+	4.88	6.06	
3232	1/2,3/2	4.9	5.81	9.6	3232	1/2-	1.1	6.77	1.0
3945	1/2,3/2	34.2	4.79	99.5	3945	3/2-	6.2	5.86	8.5
4257 ₁	1/2,3/2	1.4	6.11	4.8					
44472	1/2,3/2	8.7	5.27	33.3					
4577	1/2,3/2	2.2	5.83	9.0	4582	3/2-	3.4	5.97	6.6
5087	1/2,3/2	16.2	4.83	245.0	5082	1/2-	39.2	4.77	103.6
61673	1/2,3/2	2.2	5.37	26.5					
6239_{1}	1/2,3/2	1.4	5.55	17.4					
6429_2	1/2,3/2	8.7	4.7	124.0					
64474	1/2,3/2	3.49	5.08	50.9	-				
					6755	3/2-	12.7	4.75	109.3
7017 ₅	1/2,3/2	2.0	5.13	45.5					
					7119	3/2-	4.3	5.08	50.6
					7509	3/2-	12.2	4.49	200.1
					7622	1/2-	1.3	5.42	23.3
					7843	3/2-	3.7	4.87	82.7
7957 ₆	1/2,3/2	0.37	5.49	20.0					
					8196	3/2-	5.4	4.55	172.6
82493	1/2,3/2	2.24	4.57	167.0					
84294	1/2,3/2	3.49	4.30	307.0					
					8505	1/2-	3.6	4.57	166.9
					8506	3/2-	2.5	4.74	111.2
8707	1/2,3/2	2.02	4.40	245.0					
8779	1/2,3/2	9.64	3.68	1280					
8867 ₇	1/2,3/2	0.9	4.65	137.0					
8999 ₅	1/2,3/2	1.97	4.25	345.0					
9683	1/2,3/2	1.89	3.87	832.0					
9939 ₆	1/2,3/2	0.37	4.39	252.0					
108497	1/2,3/2	0.9	3.20	3870					

4.7 BETA DECAY OF ²⁰N

For the beta decay of 20 N feeding bound excited levels of 20 O and the one delayed neutron transition assignment, it is possible to determine both the log(ft) and B(GT) values for the specific transitions. The Q value of the 20 N ground state to 20 O

ground state transition is known to be 17.97 MeV. From this quantity, the branching ratios and the half-life measurement, the log(ft) and B(GT) values were calculated. Table 48 gives the results of these calculations.

Table 48 Calculated log(ft) and B(GT) values for specific decays assigned to the decay of log(ft). The transition identifier segregates beta decay to bound states from the delayed neutron emitting level.

Transition	²⁰ O level	Transition	Log(ft)	B(GT)
Identifier	keV	Probability		(x 1000)
$(\beta + n) 2190$	9770	0.121(13)	4.7	123
To bound	1674	0.195(22)	5.9	8
To bound	3570	0.008(8)	7.1	0.5
To bound	4072	0.037(8)	6.3	3
To bound	4456	0.012(4)	6.8	1
To bound	5387	0.018(4)	6.4	2.5

The log(ft) values are consistent with first forbidden beta transitions. This leads to the conclusion, since 0+ and 2+ levels are populated, that the spin and parity of the ²⁰N ground state is low (1- or 2-) and most likely equal to (1-).

4.8 ASTROPHYSICS RESULTS 19N

To deduce the probability and relative strengths of astrophysical reactions, and thus the rates of isotopic production, the most important quantity to determine, on a reaction-by-reaction basis, is the reaction rate per particle pair, $\langle \sigma v \rangle$. This quantity incorporates, into the reaction rate, changes in the cross section as a function of energy where the energy, in itself a temperature dependent distribution, is based on the particular astrophysical environment. Mathematically, $\langle \sigma v \rangle$ is given by the expression:

$$\langle \sigma v \rangle = (8/\pi\mu)(kT)^{-3/2} \int_0^\infty \sigma(E)E \exp(-E/kT)dE$$
 [11]

where σ is the cross section, v is the velocity, μ is the reduced mass, k is the Boltzmann constant, and T is the temperature of the system [88Ro, 95Ra, 67Fo].

The above expression is completely general though, typically, reaction cross sections are separated into direct capture and resonant capture components. The separation is permissible due to the fact that all of the non-trivial differences between the direct and resonant components resides in the $\sigma(E)$ term of the integral. Thus, rewriting the total cross section $\sigma(E)$ as a sum of the $\sigma(E)$ components affects the equation only by expanding the integral into a sum of component integrals. The total cross section $\sigma(E)$, then, may be expressed as [95Oz, 67Fo]:

$$\langle \sigma v \rangle = \langle \sigma v \rangle_{dc} + \langle \sigma v \rangle_{res}$$
 [12]

where the subscripts dc and res refer to direct capture and resonant capture, respectively.

4.8.1 DIRECT CAPTURE PART OF THE $^{18}O(n,\gamma)^{19}O$ REACTION RATE

For neutron induced astrophysical reactions, such as the (n,γ) reaction that is of interest here, determining the direct capture component to the overall rate is somewhat simpler than for rates involving charged particles. The reason this is so is due to the fact that penetrability considerations, generally associated with coulomb interactions or tunneling phenomena, are either non-existent or greatly reduced where neutrons are involved. For example, the absorbtion of $(\ell=0)$ or s-wave neutrons by a target nucleus is not hindered by a barrier to its penetration (penetrability = 1). Of course, even higher order neutron partial waves $(\ell>0)$ do not experience a Coulomb barrier, however, they

are weakly hindered by an angular momentum barrier which is a function of the system's reduced mass and the interaction energy. Even so, the angular momentum barrier is effectively much lower than the barrier to charged particles and furthermore, can be approximated by a simple mass and energy relation. For p-wave (ℓ =1) neutrons, the penetrability is approximated by:

$$P_1 = \mu^{1/2} E^{3/2}$$
 [13]

The direct capture cross section $\langle \sigma v \rangle_{dc}$ can be solved as a Maclaurin series expansion in $E^{1/2}$ about zero energy. The result in terms of $\langle \sigma v \rangle_{dc}$ takes on the form [67Fo, 88Ro]:

$$\langle \sigma v \rangle_{dc} = S(0) \left[1 + 0.3312 \frac{S'(0)}{S(0)} T_9^{1/2} + 0.06463 \frac{S''(0)}{S(0)} T_9 \right]$$
 [14]

where the constant S and its derivatives are parameters of the expansion and can be determined experimentally [67Fo], the derivatives of S are with respect to $E^{1/2}$, and T_9 is the temperature in billions of degrees Kelvin.

For our particular reaction of interest, the $^{18}O(n,\gamma)^{19}O$ reaction, the values of the constant S and its derivatives are tabulated in [94Ra]. Using these values, the direct capture cross section $\langle \sigma v \rangle_{dc}$ can be calculated for the $^{18}O(n,\gamma)^{19}O$ reaction. The result yields:

$$\langle \sigma v \rangle_{dc} \simeq 4.60 \cdot 10^{-21} T_9 \ cm^3 / sec$$
 [15]

for temperatures of $(T_9 > 0.01)$. However, it is noted that the values of the constants given in [67Fo] are the arbitrary estimates made by the author. These estimates were made prior to the measurement of the thermal neutron absorption cross section of 18 O and are reported with an uncertainty of a factor of two.

In addition to the method of calculating the direct capture cross section as above, the direct capture cross section $\langle \sigma v \rangle_{dc}$ may be approximated by the sum of the s-wave $(\ell=0)$ and p-wave $(\ell=1)$ components such that [94Ra]:

$$\langle \sigma v \rangle_{dc} = \langle \sigma v \rangle_{dc,s} + \langle \sigma v \rangle_{dc,p}$$
 [16]

Determining the s-wave part of the total cross section $\langle \sigma v \rangle_{dc,s}$ takes advantage of the so-called 1/v law where it holds that the product $\sigma_n v_n$ is a constant over all energies [88Ro]. Therefore, knowing the cross section at *any* energy provides knowledge of the cross section at *all* energies for the s-wave part of the interaction. Since the thermal neutron absorption cross section is known for ¹⁸O [94Ra], the s-wave component $\langle \sigma v \rangle_{dc,s}$ can be found from the expression [67Fo]:

$$\langle \sigma v \rangle_{dc.s} = 2.20 \cdot 10^{-19} \sigma(thermal) cm^3/sec$$
 [17]

taking the experimental value of σ (thermal) to be 1.60 x 10⁻⁴ barns, the calculated s-wave portion of the direct capture cross section becomes:

$$\langle \sigma v \rangle_{dc,s} = 3.52 \cdot 10^{-23} \ cm^3 / sec$$
 [18]

To determine the p-wave component of the direct capture cross section, $\langle \sigma v \rangle_{dc,p}$, the expression:

$$\langle \sigma v \rangle_{dc,p} = 1.787 \cdot 10^{-16} S_n T_9 cm^3 / \text{sec}$$
 [19]

from Rauscher *et al.* [94Ra] was used. In this expression, the S_n term is the S-factor for p-wave neutrons given in units of (barn MeV^{-1/2}). This term incorporates all of the strictly nuclear properties of the cross section into one expression where the cross section and the S-factor are related by:

$$\sigma(E) = S_n(\mu E)^{1/2}$$
 [20]

The approximation for the penetrability of p-wave neutrons (equation 13) has been folded into the expression. The value of S_n for the $^{18}O(n,\gamma)^{19}O$ reaction has been measured experimentally and is listed in table 7 of reference [94Ra]. Given that:

$$S_n = 52.4 \ \mu barns / MeV^{1/2}$$
 [21]

the p-wave component of the direct capture cross section can be calculated as:

$$\langle \sigma v \rangle_{dc,p} = 9.36 \cdot 10^{-21} T_9 cm^3 / sec$$
 [22]

Thus, the total direct capture cross section $\langle \sigma v \rangle_{dc}$ for the ¹⁸O(n, γ)¹⁹O reaction as calculated from the sum of the s-wave and p-wave components is then:

$$\langle \sigma v \rangle_{dc} = \langle \sigma v \rangle_{dc,s} + \langle \sigma v \rangle_{dc,p} = 3.52 \cdot 10^{-23} + 9.36 \cdot 10^{-21} T_9 cm^3 / sec$$
 [23]

The large p-wave cross section relative to the s-wave is most probably due to direct p-wave capture into the first excited, 96 keV level (3/2⁺) of ¹⁹O.

Because the parameters given in [67Fo] are arbitrarily assigned and because the parameters listed in [94Ra] are determined through experiment, the value of $\langle \sigma v \rangle_{dc}$ calculated in the above equation is considered more reliable than that of equation 15. Thus, the result generated by equation 23 will be the value incorporated into the expressions for the total reaction cross section from here forward.

4.8.2 RESONANT CAPTURE PART OF THE ¹⁸O(n, y) ¹⁹O REACTION RATE

In addition to the direct capture of neutrons by a nuclide, there is the possibility of neutron absorption through resonances in the reaction product. The strength of resonant capture depends on the relative energy of the level with respect to the unbound system, the temperature or energy distribution of the incident neutrons, the resonant widths of the entrance and exit channels and, the spin/parity of the resonance and target

nuclide. The functional form of the equation used to determine the resonant part of the cross section, $\langle \sigma v \rangle_{res}$, is given as [95Oz, 88Ro, 94Ra, 67Fo]:

$$\langle \sigma v \rangle_{res} = 2.557 \cdot 10^{-13} (\omega \gamma)_r A^{-3/2} T_9^{-3/2} Exp(-11.605 E_r / T_9) cm^3 / sec$$
 [24]

where A is the reduced mass in atomic mass units, E_r is the center of mass energy of the resonance in MeV, and $(\omega \gamma)_r$ is the resonance strength in MeV.

The resonance strength $(\omega \gamma)_r$ is the product of a statistical factor and the gamma width of the resonance (Γ_{γ}) . Mathematically, the value is given by:

$$(\omega \gamma)_r = (2J_r + 1)/[(2J_T + 1)(2J_P + 1)] \bullet \Gamma_r$$
 [25]

Where the J values are for the resonance, target nuclide ground state, and projectile, and the gamma width (Γ_{γ}) is in units of eV. For the case of the $^{18}O(n,\gamma)^{19}O$ reaction rate, the target spin is 0 (^{18}O ground state spin), the neutron projectile spin is ½ (incident neutron), and the gamma width is expected (Γ_{γ}) to be less than 1 eV from the systematics [93En, 95Oz]. Consequently, the value for $(\omega \gamma)_r$ can be calculated to be:

$$(\omega \gamma)_r \le (2J_r + 1)/2 \bullet 10^{-6} MeV$$
 [26]

with J_r being the spin of the resonance in ¹⁹O absorbing the incident neutron.

To assess the upper limit on the resonance contribution to the $^{18}O(n,\gamma)^{19}O$ reaction proceeding through the states observed in this experiment, all observed neutrons were assumed to populate the ground state in ^{18}O . This minimizes the resonance energy and thus maximizes the potential contribution to the reaction. Also, the beta transitions were assumed to be of the allowed type (which is consistent with the log(ft) values) and the ground state spin of ^{19}N was assumed to be $\frac{1}{2}$ -, which is consistent with the other known odd-mass nitrogen isotopes. Assuming the allowed

decay and the spin of the ¹⁹N ground state, the spin and parity of the intermediate levels in ¹⁹O are restricted to being either ½- or 3/2-, with 3/2- being the choice that will maximize the resonant contribution (see equation 26).

On the basis of these assumptions, the measured energies of the delayed neutrons, and with reference to equation 24, the strength of the $^{18}O(n,\gamma)^{19}O$ reaction rate through each excited level of ^{19}O can be reduced to a functional form having but one parameter – the temperature. Equation 27 gives the functional form of the resonant contribution for the delayed neutron emitting states observed in this experiment.

$$<\sigma v>_{res} = 5.55 \bullet 10^{-19} T_9^{-3/2} \times$$

$$[Exp(-3.48/T_9) + Exp(-5.69/T_9) + Exp(-7.20/T_9) + Exp(-9.63/T_9) + Exp(-13.11/T_9) + Exp(-14.74/T_9) + Exp(-25.65/T_9) + Exp(-28.90/T_9) + Exp(-32.96/T_9) + Exp(-35.51/T_9) + Exp(-46.42/T_9) + Exp(-56.98/T_9)] [27]$$

Clearly, the last several terms will not contribute much to the total resonant capture strength due to the large negative exponent. This is a consequence of the high relative energy of the state with respect to the $(^{18}O + n)$ energy.

4.9.3 COMPARISON OF DIRECT AND RESONANT CAPTURE STRENGTHS

The direct capture strength $\langle \sigma v \rangle_{dc}$ is given as a function of temperature in equation 23 while the resonant capture strength is given as a function of temperature in equation 27. A plot of the two components of the $^{18}O(n,\gamma)^{19}O$ reaction over all temperatures will illuminate the relative probability for the two branches. Figure 52 shows such a plot where the direct capture component is shown in red and the resonant capture component is shown in blue. Also shown is the temperature range normally associated with the inhomogeneous big bang model [94Ra].

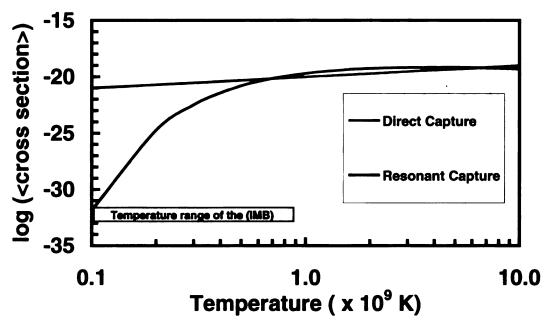


Figure 52 Comparison of the direct capture and resonant capture cross sections for the $^{18}O(n,\gamma)^{19}O$ reaction as derived from this work.

The important feature of the plot is that the direct capture cross section dominates the resonant capture cross section over most of the temperature range of interest for the inhomogeneous big bang model. In fact, the direct capture component is many orders of magnitude stronger up to a temperature of about 0.4 T₉. Above a temperature of 0.8 T₉ however, the resonant capture is comparable to the direct capture and thus could be the dominant component of the reaction rate at the upper edge of the postulated temperature range of the IMB model.

CHAPTER 5

SUMMARY AND CONCLUSIONS

5.1 SUMMARY AND CONCLUSIONS - 19N

In this experiment, quantitative measurement of the ^{19}N decay was made including a determination of the half-life, beta decay to bound excited states of ^{19}O and delayed neutron emission to levels in ^{18}O . The half-life of ^{19}N was determined to be 299 ($\pm 3 \pm 16$) milliseconds; which is in agreement with reference [91Re].

Seven gamma rays characteristic of ¹⁹O were observed accounting for 39.4(38)% of the total decay strength. The gamma rays suggest direct beta feeding to three, possibly four, bound excited levels of ¹⁹O. The levels populated are the 3945 (3/2-), 3232 (1/2-), 1472 (1/2+) and possibly the 96 (3/2+) level, with branching ratios of 34.2(37), 4.9(6), 0.27(12) and 4.3(UL), respectively. The results of the gamma ray spectra are not consistent with those published in [88Duf] as the 709 keV gamma ray reported there was not observed in this experiment.

Twelve delayed neutrons were observed by the neutron bar array and the ⁶Li glass detectors. The neutrons ranged in energy from 300 keV up to nearly five MeV and in branching ratio from 0.37(4)% up to 16.2(17)% of the total decay strength. In addition, gamma rays from ¹⁸O were determined to be from the ¹⁹N source suggesting delayed neutron emission to bound excited states of ¹⁸O rather than direct population of the ground state. The total delayed neutron emission probability for ¹⁹N was found to be 50.9(62)% of the decay strength; which is in agreement with reference [94Re]. The

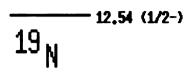
branching ratio for delayed neutron emission to bound excited states of ¹⁸O was found to be 16.5(21)% of the total. Figure 53 shows the beta feeding to specific levels in ¹⁹O, where possible, and table 49 shows the transitions that cannot be assigned on the basis of these experimental data.

Table 49 Summary of transitions observed during the decay of ¹⁹N that cannot be definitively assigned to intermediate levels in ¹⁹O.

Transition Type	Neutron Energy (CoM)	Branching Ratio (%)
$(\beta + n)$	300 keV	1.4(3)
$(\beta + n)$	490 keV	8.7(10)
$(\beta + n)$	2210 keV	2.2(4)
$(\beta + n)$	2490 keV	3.5(4)
$(\beta + n)$	3060 keV	1.97(32)
$(\beta + n)$	4000 keV	0.37(4)
$(\beta + n)$	4910 keV	0.90(10)

A comparison of the shell model predictions with the data recorded in this experiment is not particularly favorable. The model derived half-life is a factor of two larger than experiment and, with the exception of the lower energy levels being comparable in energy, the delayed neutron emitting excited states do not appear to track between experiment and theory even when the transition uncertainty is included in the mix. It is hoped that consideration of these new data will facilitate improvements in the model parameters to better approximate the reality of this nuclear system.

From the observed neutron energies and their log(ft) values, a functional upper limit on the resonant neutron capture cross section of ¹⁸O was derived. The resonant



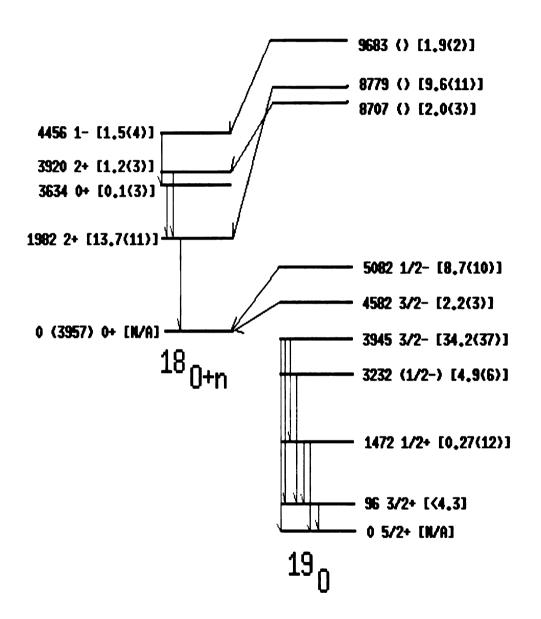


Figure 53 The beta decay of ¹⁹N showing the transitions that may be conclusively identified from the experimental data.

cross section was compared with the direct capture cross section and found to be orders of magnitude lower, over most of the temperature range of interest, for inhomogeneous big bang models of nucleosynthesis. It is noted, however, that at about $(T_9 = 0.8)$, the upper limit of the resonant part is comparable to the direct capture cross section.

From the beta decay log(ft) values and the known spin and parity of the ¹⁹O levels populated, the ground state spin and parity of ¹⁹N is restricted to be either (1/2-) or (3/2-). This result is consistent with the common-sense assignment of (1/2-) that is characteristic of all other odd-mass nitrogen isotopes where the ground state properties are known. It is noted that this experiment was not sensitive to the ¹⁹N ground-state to ground-state decay, or to delayed alpha emission which is energetically possible.

5.2 SUMMARY AND CONCLUSIONS – ²⁰N

As with 19 N, the 20 N data permitted re-evaluation of the beta decay half-life, measurement of the delayed neutron spectrum, and observation of beta feeding to bound excited states of the daughter 20 O. The half-life of 20 N was found to be 121 (\pm 5 \pm 10) milliseconds; which is in agreement with the results published in [94Re, 88Mu].

Analysis of the gamma ray data illuminated five transitions characteristic of ²⁰O indicating direct beta feeding to bound levels of the daughter. From the gamma ray energies and the inferred cascades the beta decay can be demonstrated to populate four, possibly five, bound excited levels of ²⁰O. The populated levels are the 5387 (0+), 4456 (0+), 4072 (2+), 1674 (2+) with branching ratios of 1.8(4), 1.2(4), 3.7(8) and 19.5(22) percent, respectively. The fifth possible level is at 3570 (4+) with a branching ratio of 0.8(8) percent (which is also consistent with zero percent). From these spin/parity

values and the calculated log(ft) of the beta decay, the ground state of ²⁰N is most consistent with a (1-) spin and parity assignment.

The neutron bar array permitted identification of eight delayed neutrons ranging in energy from 1.16 to 4.46 MeV and ranging in branching ratio from 0.5(1) to 12.1(13) percent of the total decay strength. The probability for neutron emission is calculated to be 34.0(40)% of the total decay strength for neutrons above 1.1 MeV in energy. From the gamma ray data, two transitions characteristic of ¹⁹O are observed and assigned as following directly from the decay of ²⁰N. These are the 96 keV and 1374 keV lines and their observation suggests delayed neutron emission from ²⁰N populating excited levels of ¹⁹O rather than the ground state.

Due to the lack of information regarding higher energy levels in ²⁰O, and the uncertainty in the decay characteristics, it is only possible to assign one beta transition to a specific neutron-emitting level of ²⁰O. Figure 54 shows the beta feeding to specific levels in ²⁰O, where possible to assign, and table 50 shows the transitions that cannot be assigned on the basis of these experimental data.

Table 50 Summary of transitions observed during the decay of ²⁰N that cannot be assigned to specific intermediate levels of ²⁰O.

Transition Type	Neutron Energy (CoM)	Branching Ratio (%)
$(\beta + n)$	1160 keV	4.5(5)
$(\beta + n)$	1330 keV	2.5(3)
$(\beta + n)$	1630 keV	8.3(9)
$(\beta + n)$	2630 keV	1.3(2)
$(\beta + n)$	3380 keV	2.1(3)
$(\beta + n)$	3880 keV	2.7(3)
$(\beta + n)$	4640 keV	0.5(1)

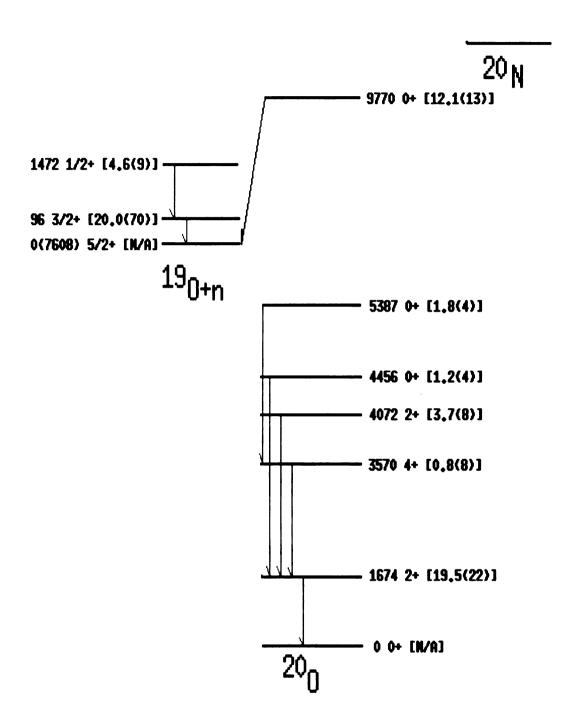


Figure 54 The beta decay of ²⁰N showing the transitions that may be conclusively identified from the experimental data.

REFERENCES

- [67Fo] W.A. Fowler et al., Ann. Rev. Astr. AP 5, 525 (1967)
- [70Fi] Fintz et al., Nucl. Phys. A150, 49 (1970)
- [71Hi] Hibou et al., Nucl. Phys. A171, 603 (1971)
- [71Go] N.B. Gove and M.J. Martin, Nucl. Data Tables 10, 205 (1971)
- [76Al] D.E. Alburger and D.H. Wilkinson, Phys. Rev. C13, 835 (1976)
- [81Fr] Nuclear and Radiochemistry; G. Friedlander et al., John Wiley and Sons, New York (1981)
- [8201] J.W. Olness et al., Nucl. Phys. A378, 539 (1982)
- [83Ga] C.A. Gagliardi et al., Phys. Rev. C27, 1353 (1983)
- [86Du] J.P. Dufour et al., Z. Phys. A324, 487 (1986)
- [87Al] D.E. Alburger et al., Phys. Rev. C35, 1479 (1987)
- [88De] Gamma- and X-Ray Spectrometry with Semiconductor Detectors; K. Debertin and R.G. Helmer, Elsevier Science Publishers, Amsterdam, (1988)
- [88Du] J.P. Dufour et al., Phys. Lett. **B206**, 195 (1988)
- [88Kr] <u>Introductory Nuclear Physics</u>; K.S. Krane, John Wiley and Sons, New York (1988)
- [88Mu] A.C. Mueller et al., Z. Phys. A330, 63 (1988)
- [88Ro] <u>Cauldrons in the Cosmos</u>; C.E. Rolfs and W.S. Rodney, University of Chicago Press, Chicago (1988)
- [88Sa] M. Samuel et al., Phys. Rev. C37, 1314 (1988)
- [88Wa] E.K. Warburton, Phys. Rev. C38, 935 (1988)

[89Kn] Radiation Detection and Measurement; G.F. Knoll, John Wiley and Sons, New York (1989)

[91Sh] B.M. Sherrill et al., Nucl. Inst. Meth. **B56**, 1106 (1991)

[91Re] P.L. Reeder et al., Phys. Rev. C44, 1435 (1991)

[92Ha] The Beta Decay Branching Ratios of the Neutron-Rich Nucleus ¹⁵B; R.

Harkewicz, Ph.D. Thesis, Michigan State University (1992)

[93Au] Audi and Wapstra, Nucl. Phys. A565, 1 (1993)

[93Bu] L. Buchmann et al., Phys. Rev. Lett. 70, 726 (1993)

[93En] P.M. Endt, Atomic Data and Nucl. Data Tables 55, 171 (1993)

[93Sc] The Beta Delayed Neutron Decay of the Exotic Nuclei ¹⁸N, ¹⁷C, and ¹⁸C; K.

Scheller, Ph.D. Thesis, University of Notre Dame, (1993)

[93Ti] D.R. Tilley et al., Nucl. Phys. **A565**, 1 (1993)

[94Do] M. Dombsky et al., Phys. Rev. C49, 1867 (1994)

[94Le] Techniques for Nuclear and Particle Physics Experiments; W.R. Leo, Springer-

Verlag, New York (1994)

[94Ra] T. Rauscher et al., APJ. 429, 499 (1994)

[94Re] P.L. Reeder et al., Int. Conf. Nucl. Data for Science and Tech. – Gatlinburg (1994)

[95Oz] A. Ozawa et al., Nucl. Phys. A592, 244 (1995)

[95Sc] K. Scheller et al., Nucl. Phys. A582, 109 (1995)

[95Ti] D.R. Tilley et al., Nucl. Phys. **A595**, 1 (1995)

[96Fi] Table of Isotopes; R.B. Firestone, John Wiley and Sons, New York, (1996)

[96Me] J. Meissner et al., Phys Rev C53, 459 (1996)

[97Bi] K. Blackmon, Bicron Technical Data Sheet, Private Communication (1997)

[97Gi] U. Giesen to D. Anthony, Private Communication (1997)

[98Po] Study of the d(⁷Be, ⁸B)n Reaction; C.F. Powell, Ph.D. Thesis, Michigan State University (1998)

[00Nn] National Nuclear Data Center; www.nndc.bnl.gov, (2000)

[00Pr] J.I. Prisciandaro to D. Anthony, Private Communication (2000)

[00Ro] R. Ronningen to D. Anthony, Private Communication (2000)

