

Measurement of Cross Sections for the $^{63}\text{Cu}(\alpha, \gamma)^{67}\text{Ga}$ Reaction from 5.9-8.7 MeV

M. Shamsuzzoha Basunia¹, Eric B. Norman^{1*}, Howard A. Shugart²,
Alan R. Smith¹, Michelle J. Dolinski², Brian J. Quiter³

¹ Nuclear Science Division, Lawrence Berkeley National Laboratory, Berkeley, California 94720

² Department of Physics, UC Berkeley, Berkeley, California 94720

³ Department of Nuclear Engineering, UC Berkeley, Berkeley, California 94720

Abstract. We have measured cross sections for the $^{63}\text{Cu}(\alpha, \gamma)^{67}\text{Ga}$ reaction in the 5.9-8.7 MeV energy range using an activation technique. Natural Cu foils were bombarded with alpha beams from the 88" Cyclotron at Lawrence Berkeley National Laboratory (LBNL). Activated foils were counted using gamma spectrometry system at LBNL's Low Background Facility. The $^{63}\text{Cu}(\alpha, \gamma)^{67}\text{Ga}$ cross-sections were determined and compared with the latest NON-SMOKER theoretical values. Experimental cross sections were found to be in agreement with theoretical values.

INTRODUCTION

Cross-section measurements for charged-particle capture reactions on nuclei heavier than iron are important for nucleosynthesis studies [1] and for testing statistical model predictions. The inner zones of supernovae, where temperatures exceed 10^9 K are places where proton and alpha particle reactions on medium to heavy nuclei may be important in determining the mix of elements and isotopes that emerge from such stellar explosions. Within the last few years, some proton capture cross sections in the $A=90$ -100 mass region [2-4] and alpha capture on ^{144}Sm , ^{70}Ge , and ^{96}Ru isotopes [5-7] have been reported. Experimental alpha capture cross sections on ^{96}Ru and ^{144}Sm were reported to be about 2.5 and 5-7 times lower, respectively, than the latest theoretical values. An earlier measurement of alpha capture on ^{40}Ca also found cross sections to be about 3-5 times lower than the theoretical predictions [8]. However, the experimental S-factor values for the $^{70}\text{Ge}(\alpha, \gamma)^{74}\text{Se}$ reaction were in agreement with statistical model calculations [6]. It is important to investigate alpha capture cross sections for different mass regions to test the theoretical models. Rapp *et al.* [7] indicated a possible deficiency in the theoretical treatment of the alpha channels for the mass region around 100 and suggested additional alpha induced cross section

measurements over a wider mass range for understanding and improving the situation.

Here we report the measured cross sections for the $^{63}\text{Cu}(\alpha, \gamma)^{67}\text{Ga}$ reaction in the 5.9-8.7 MeV energy range using an activation technique. Experimental procedure and comparison of the measured data with the latest theoretical values are presented and discussed.

EXPERIMENTAL PROCEDURE

Natural Cu foil of thickness ~ 1 mg/cm² used in this experiment were purchased from ACF-Metals, Tucson, Arizona. The foils were floated on water from glass slides and mounted on circular aluminum holders. Three stacks of targets each having four ^{nat}Cu and one ^{nat}Ti foil of thickness 2.7 mg/cm² were prepared. The target stacks were mounted on a thick water-cooled copper block that also served as a beam stop. Two stacks were irradiated, each for an hour, with alpha beams of energies 8.8 MeV and 7.9 MeV from the 88" Cyclotron at LBNL. The beam current was 1 μA . The third stack was irradiated for 6 hours with 7.0 MeV beam energy and 0.1 μA current. The uncertainty in the beam energy was about 1%. The incident alpha beam energy on the successive foils

* Present address: Lawrence Livermore National Laboratory

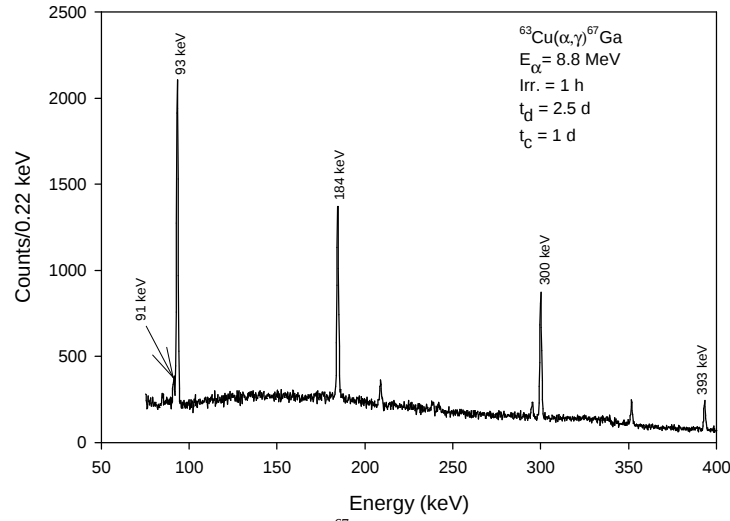


FIGURE 1. A partial HPGe γ -ray spectrum of ^{67}Ga characteristic γ lines (t_d and t_c = decay and counting time).

was calculated based on the energy loss through Cu foils using dE/dx values estimated using the TRIM (the Transport of Ions in Matter) code [9]. On average, the loss per Cu foil was about 300 keV. The beam current was integrated using a Brookhaven Instruments Corporation Integrator.

The titanium foil, at the end of each stack, was used for beam current calibration using the $^{48}\text{Ti}(\alpha,n)^{51}\text{Cr}$ reaction and for catching the recoil ^{67}Ga radioisotopes from the preceding copper foil to estimate the recoiled fraction.

Following each irradiation, the copper targets were counted immediately using an HPGe detector to measure the ^{68}Ga activity, produced through the $^{65}\text{Cu}(\alpha,n)^{68}\text{Ga}$ reaction. All the copper foils were later recounted for longer periods of time to measure the ^{67}Ga activity using another HPGe detector, 80% relative efficiency, at LBNL's Low Background Facility (LBF). The energy resolution of the HPGe detector was 1.9 keV at $E_\gamma = 1332.5$ keV. Gamma ray energy spectra were accumulated in 16384 channels using an ORTEC PC-based acquisition system. A partial HPGe γ -ray spectrum collected at the LBF is shown in Figure 1 for the characteristic γ -energies of ^{67}Ga . The ^{67}Ga radioactivity in samples bombarded with the two highest beam energies was sufficiently high to count at 25 cm and 15 cm away from the detector end cap. However, for the lowest beam energy, samples were counted at the end cap surface of the HPGe detector. Efficiency calibration of the HPGe detectors was done using calibrated point sources of ^{22}Na , ^{54}Mn , ^{57}Co , ^{60}Co , ^{109}Cd , ^{133}Ba , and ^{137}Cs purchased from Isotope Products Laboratories.

The efficiency curve for the surface counting position was generated from the peak efficiency data at 25 cm using count ratios of single gamma sources at surface position and 25 cm [10]. Single gamma lines 88.0 keV, 320.1 keV, 661.4 keV, and 834.8 keV from ^{109}Cd , ^{51}Cr , ^{137}Cs , and ^{54}Mn , respectively, were used. The ^{51}Cr source was available from the current experiment.

All gamma spectra were analyzed using ORTEC Gamma Vision software. The 91 keV and 93 keV gamma lines of ^{67}Ga were slightly overlapped in the tail. The combined area of these two peaks was used together to determine the $^{63}\text{Cu}(\alpha,\gamma)^{67}\text{Ga}$ cross section. The cross sections were deduced from the well known activation equation:

$$A_o = nsf(1 - e^{-\lambda t}) \quad (1)$$

Where, A_o = ^{67}Ga activity at the end of irradiation (disintegration/sec), n = number of ^{63}Cu nuclei ($\#/\text{cm}^2$), σ = cross section (cm^2), ϕ = number of incident α particles ($\#/\text{sec}$), and $(1 - e^{-\lambda t})$ = growth factor for a decay constant λ and irradiation time t .

The activity, A_o , at the end of irradiation was deduced from the measurement using the following equation:

$$A_o = IN_o = IC / \{I_g e^{-(t_{cs}-t_{ie})} - e^{-\lambda(t_{ce}-t_{ie})}\} \quad (2)$$

Where, N_o = number of ^{67}Ga nuclei at the end of irradiation, t_{cs} , t_{ce} , t_{ie} = counting start, counting end, and irradiation end times, respectively, C = net area

TABLE 1. Nuclear data of the product radioisotopes used in this experiment [11]

Nuclear reaction	Half life	E _g (keV) (I _g %)	
		uncertainty for the least significant digit(s)	
⁶³ Cu(α,γ) ⁶⁷ Ga	3.26 d	91.3(3.16 9), 93.3(39.2 10), 184.6(21.2 3), 300.2 (16.8 22), 393.5(4.68 6)	
⁶⁵ Cu(α,n) ⁶⁸ Ga	67.63 min	1077.4 (3.0 3)	
⁴⁸ Ti((α,n) ⁵¹ Cr	27.7 d	320.1 (9.92 5)	

under the peak for a counting duration ($t_{cs} - t_{ce}$), I_g = gamma ray intensity, and e = detector peak efficiency.

Cross sections of the ⁶³Cu(α,γ)⁶⁷Ga reaction were deduced using all ⁶⁷Ga γ-rays and found to be statistically consistent to each other. Nuclear data for the product nuclei used in this experiment is presented in Table 1. Cross sections for the ⁶³Cu(α,γ)⁶⁷Ga reaction reported in this paper are deduced using the 184 keV γ-ray. In all gamma spectra, this peak had smooth tailing on both sides with statistically reasonable peak area. Absolute γ-ray intensities of ⁶⁷Ga are deduced in this work, considering the recent ⁶⁷Ga decay data [12] and using relative γ-ray intensities from Ref. [13]. We used 184 keV γ-ray intensity of $20.7 \pm 0.1\%$, about 2 percent lower than the value in Ref. [11]. There was an overlapping bombarding energy for the last foil of the 1st stack and the 1st foil of the 2nd stack. The agreement between these two cross sections for the common energy was excellent. This served as a cross check for the two different sets of irradiation for the ⁶³Cu(α,γ)⁶⁷Ga reaction measurement.

Titanium foils were counted after about 7 days at the LBF for the ⁵¹Cr and recoiled ⁶⁷Ga activities using the HPGe spectrometry system. This length of decay period allowed the 91.3 keV and 93.3 keV ⁶⁷Ga peaks to appear in the spectra. Recoiled ⁶⁷Ga activity was determined using equation (1) and (2) and was found to be about 10%-14% in this experiment. Assuming a uniform ⁶⁷Ga recoil out of the successive foils in the stack, a correction of 12% was made for the first Cu foil ⁶⁷Ga activity in each stack.

Measured cross sections for the ⁴⁸Ti(α,n)⁵¹Cr reaction were compared with the published experimental data [14] for beam current calibration. The comparison provided very reliable current integration of the Brookhaven Instruments Corporation Integrator for the 8.8 MeV and 7.9 MeV beams. For the 7.0 MeV beam, the comparison was incomplete using the ⁴⁸Ti(α,n)⁵¹Cr reaction, because in this case published cross sections were partially available for the interacting alpha energy range through the titanium foil. However, employing other cross checks, such as simultaneous ⁶⁵Cu(α,n)⁶⁸Ga cross sections measurement and comparison with known experimental

results, we are confident on the current integrator reading for the 7.0 MeV beam.

Considering all uncertainties of detector efficiency calibration, target foil thickness, beam current, counting statistics, decay data, and recoil fraction, we report 15% uncertainties for the measured cross sections.

RESULTS AND DISCUSSION

Measured cross sections for the ⁶³Cu(α,γ)⁶⁷Ga reaction are presented in Table 2. In Figure 2, measured values are presented along with the latest theoretical values of the NON-SMOKER statistical model [15]. Theoretical data points were obtained using the finite range droplet model (FRDM) masses from the Uniform Resource Locator (URL) [15]. These data points were not available in regular intervals in the studied energy range. However, from Figure 2, it can be seen that the agreement between the experimental and theoretical data are reasonably good for the ⁶³Cu(α,γ)⁶⁷Ga reaction cross sections.

TABLE 2. Measured cross sections for the ⁶³Cu(α,γ)⁶⁷Ga reaction

Energy (Lab) (MeV)	Cross section (mb)
8.65 ± 0.09	1.08 ± 0.16
8.37 ± 0.08	1.04 ± 0.16
8.08 ± 0.08	0.93 ± 0.14
7.80 ± 0.08	0.69 ± 0.10
7.54 ± 0.08	0.41 ± 0.06
7.24 ± 0.07	0.26 ± 0.04
6.99 ± 0.07	0.18 ± 0.03
6.88 ± 0.07	0.13 ± 0.02
6.56 ± 0.07	0.07 ± 0.01
6.22 ± 0.06	0.026 ± 0.004
5.88 ± 0.06	0.012 ± 0.002

The comparison of measured ⁶⁵Cu(α,n)⁶⁸Ga cross-sections in this work with those of Stelson *et al.* [16] were found to be excellent. This agreement provides an indication of the experimental integrity for the reported ⁶³Cu(α,γ)⁶⁷Ga cross-section measurement.

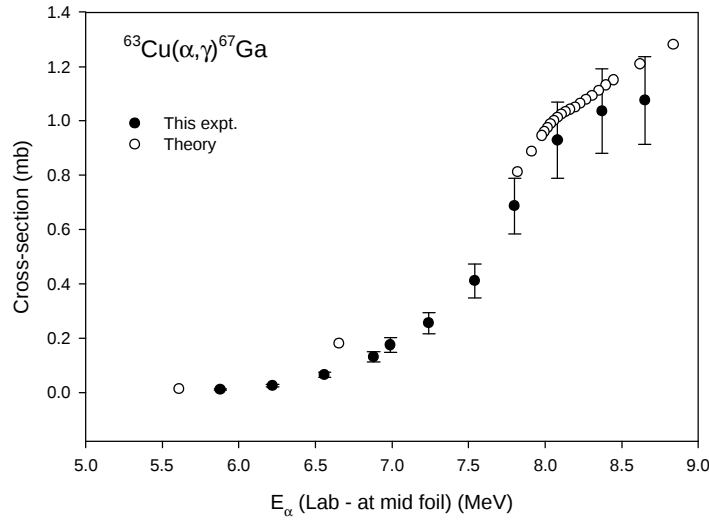


FIGURE 2. Experimental and theoretical cross sections for the $^{63}\text{Cu}(\alpha, \gamma)^{67}\text{Ga}$ reaction.

Based on the present results and those from studies of the $^{70}\text{Ge}(\alpha, \gamma)^{74}\text{Se}$ reaction [6], it appears that the theoretical calculations of (α, γ) cross sections in the mass region of $A=60-70$ are in reasonably good agreement with the experimental data.

ACKNOWLEDGMENTS

We wish to acknowledge our gratitude to W. S. Tiffany and G. A. Seaman for technical assistance during experimental setup, C. R. Siero and R. K. Thatcher for the 88" cyclotron operation. This work is sponsored by Office of Science, U.S. Department of Energy, under contract DE-AC03-76SF00098.

REFERENCES

1. Hoffman, R. D., Woosley, S. E., and Qian, Y.-Z., *Astrophys. Journal* **482**, 951-962 (1997).
2. Sauter, T., and Käppeler, F., *Phys. Rev. C* **55**, 3127-3138 (1997).
3. Bork, J., Schatz, H., Käppeler, F., and Rauscher, T., *Phys. Rev. C* **58**, 524-535 (1998).
4. Chloupek, F. R., Murphy, A. St. J., Boyd, R. N., Cole, A. L., Gorres, J., Guray, R. T., Raimann, G., Zach, J. J., Rauscher, T., Schwarzenberg, J.V., Tischhauser, P., and Wiescher, M. C., *Nucl. Phys A* **652**, 391-405 (1999).
5. Somorjai, E., Fülöp, Zs., Kiss, Á. Z., Rolfs, C., Trautvetter, H-P., Greife, U., Junker, M., Arnould, M., Rayet, M., and Oberhummer, H., *Nucl. Phys. A* **621**, 293c-296c (1997).

6. Fülöp, Zs., Kiss, Á. Z., Somorjai, E., Rolfs, C. E., Trautvetter, H-P, Rauscher, T., and Oberhummer, H., *Z. Phys. A* **355**, 203-207 (1996).
7. Rapp, W., Heil, M., Hentschel, D., Käppeler, F., and Reifarth, R., *Phys. Rev. C* **66**, 015803-11, (2002).
8. Cooperman, E. L., Shapiro, and M.H., Winkler, H., *Nucl. Phys. A* **284**, 163-176 (1977).
9. Ziegler, J. F., and Biersack, J. P., *The Stopping and Range of Ions in Solids*, New York, Pergamon Press, 1985. URL: <http://www.srim.org/>.
10. Gmuca, Š, and Ribanský, I., *Nucl. Inst. and Meth.* **202**, 435-437 (1982).
11. Firestone, R. B., Shirley, V. S., *Table of Isotopes*, 8th Edition, New York, John Wiley & Sons, Inc., 1996.
12. Simpson, B. R. S., and Ntsoane, T. P., *App. Rad. Iso.* **52**, 551-556 (2000).
13. Meyer, R. A., Prindle, A. L., Myers, W. A., Hopke, P. K., Dieterly, D., and Koops, J. E., *Phys. Rev. C*, **17**, 1822-1831 (1978).
14. Morton, A. J., Tims, S. G., Scott, A. F., Hansper, V. Y., Tingwell, C. I. W., and Sargood, D. G., *Nucl. Phys. A* **537**, 167-182 (1992).
15. Rauscher, T., and Thielemann, F.-K., *Atomic Data Nuclear Data Tables* **79**, 47-64 (2001). URL: <http://quasar.physik.unibas.ch/~tommy/nosmo.html>.
16. Stelson, P. H. and McGowan, F. K., *Phys. Rev.* **133**, B911-B919 (1964).