Search for resonances in the 22 Na $(p, \gamma)^{23}$ Mg reaction

J. Görres, M. Wiescher, and S. Graff Department of Physics, University of Notre Dame, Notre Dame, Indiana 46556

R. B. Vogelaar, B. W. Filippone, C. A. Barnes, S. E. Kellogg, and T. R. Wang W. K. Kellogg Radiation Laboratory, California Institute of Technology, Pasadena, California 91125

B. A. Brown

Department of Physics and Astronomy, Michigan State University, East Lansing, Michigan 48824 (Received 1 September 1988)

The reaction 22 Na $(p,\gamma)^{23}$ Mg has been investigated in the energy range $E_p = 0.4 - 1.27$ MeV using a radioactive 22 Na target. Upper limits for the γ yield have been determined for transitions to states in 23 Mg up to $E_x = 4.4$ MeV. The upper limits for the expected resonances in this energy range are more than an order of magnitude weaker than previously predicted. These upper limits are compared with shell model calculations.

I. INTRODUCTION

The understanding of the rp-process nucleosynthesis¹ in the Ne-Na-Mg region is of particular interest for the interpretation of ²²Ne anomalies in meteoritic inclusions² as the decay product of 22 Na ($t_{1/2} = 2.6$ yr). Several reaction sequences, depending on the temperature and density conditions, may lead to the formation of ²²Na. Most favorable is the Ne-Na cycle³

20
Ne $(p, \gamma)^{21}$ Na $(\beta^+ \nu)^{21}$ Ne $(p, \gamma)^{22}$ Na,

PHYSICAL REVIEW C

where ²²Na is formed by well-understood proton capture where ^{-1}Na is formed by well-understood proton capture reactions. 4,5 The abundance of ^{22}Na , however, also depends critically on the rates of possible depletion reactions, the most important being $^{22}Na(p,\gamma)^{23}Mg$. Another mechanism for producing ^{22}Na , more likely to occur in high-temperature and high-density conditions, 1,6 is

$$^{20}{\rm Ne}(p,\gamma)^{21}{\rm Na}(p,\gamma)^{22}{\rm Mg}(\beta^+\nu)^{22}{\rm Na}$$
 .

In this case the production rate also depends on the rate of the competing proton capture rate on 22 Mg. While the reaction rate for 22 Mg $(p, \gamma)^{23}$ Al has recently⁷

been deduced from a measurement of its resonance energy coupled to a shell-model calculation of its resonance strength, only theoretical estimates for the 22 Na $(p, \gamma)^{23}$ Mg reaction rate exist. Little was known about the structure of the compound nucleus ²³Mg above the proton threshold at Q = 7.58 MeV when the first estimate was published. Following this, the $^{25}Mg(p,t)^{23}Mg$ (Ref. 9) and 24 Mg(3 He, α) 23 Mg (Ref. 10) reactions were used to identify proton unbound levels in ²³Mg (Fig. 1). However, these experiments did provide neither unique spin and parity assignments for the levels nor information about the partial widths necessary to calculate the resonance strengths for the 22 Na (p, γ) reaction. Therefore the direct measurement is highly desirable.

Because of the relatively long half-life of ²²Na $(t_{1/2}=2.6 \text{ yr})$, the use of a radioactive target is more favorable than the use of a radioactive beam. 11 The radioactive ²²Na target, however, produces an extremely large flux of 511 and 1275 keV γ rays. To circumvent this problem, Rolfs and Kavanagh¹² have developed a new type of detector, which is not sensitive to γ radiation below $E_{\gamma} = 2.2$ MeV. A disadvantage of this detector is the lack of energy resolution, which requires ultraclean ²²Na targets. We therefore chose a more traditional experimental approach for the direct measurement of the 2 Na $(p,\gamma)^{23}$ Mg reaction above a proton energy of $E_p = 0.400 \text{ MeV}.$

II. EXPERIMENTAL EQUIPMENT AND SETUP

Proton beams of $10-30 \mu A$ were provided by the 3 MV Tandem Pelletron at the Kellogg Radiation Laboratory over the energy range of 0.400-1.275 MeV. The proton beam passed through a Ta collimator and was directed onto the target. The targets were mounted normal to the beam and directly watercooled. A liquid-nitrogen-cooled Cu tube was extended to within 10 cm of the target to reduce carbon deposition. During the experiment a slowly varying (0.1 Hz) ramping voltage (0 to +8 kV) was applied to the electrically insulated target chamber. This allowed us to measure the excitation function over an energy range of 8 keV per beam-energy step. The accumulated charge was measured continuously as a function of the ramping voltage and could be corrected for leakage current. This leakage current, due to the water cooling, was $\leq 1 \,\mu A$.

The production and properties of the ²²Na target have been described in detail elsewhere, ¹³ and will be discussed here only briefly. The ²²Na targets were prepared from an aqueous ²²NaCl solution (Amersham Corporation) with a specific activity of 1.3 mCi/ml. Three to five drops of the solution were applied to the backing and slowly dried under a heat lamp. This procedure was repeated until the desired activity was reached. The backing consisted of 0.5 mm Ta with a 0.05 mm deep reces-

8

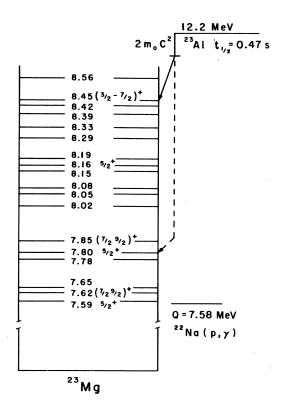


FIG. 1. Level structure of ²³Mg above the proton threshold (Refs. 9, 10, and 14).

sion (diameter 6 mm) to physically constrain the activity. The Ta previously was coated with a 50 μ g/cm² Ni layer to minimize the diffusion of the Na into the backing (see Ref. 13). The total activity of the target was limited by the counting rate in the γ detector to 60 μ Ci. This activity, distributed over an area of 0.28 cm², corresponds to a ²²Na areal density of 8.9×10^{14} atoms/cm² (see Table I and Ref. 13).

Independent tests indicated that the loss of the target material due to proton bombardment could be kept to $\leq 10\%$ over a period of 1 week by defocusing the beam of 20 μ A to produce a uniform beam spot (4–5 mm diameter). Activity scans of the targets before and after runs show no significant change in the ²²Na distribution (see Ref. 13).

Contaminants in the 22 NaCl solution were measured using known, strong (p, γ) resonances and are listed in Table I; for details see Ref. 13. The large amount of Si in the solution arises from radiation damage to the glass

TABLE I. Measured composition of the ²²Na target (Ref. 13).

Element	n (atoms/cm ²)
²² Na	$(8.9\pm0.4)\times10^{14}$
²³ Na	$(1.1\pm0.2)\times10^{15}$
Cl	$(1.2\pm0.6)\times10^{15}$
Mg	$(3.2\pm0.8)\times10^{15}$
Si	$(1.1\pm0.1)\times10^{17}$
Z > 18	$(\simeq 10^{17})$

container in which the aqueous solution was contained. Heavier elements, $Z \ge 18$, could not be measured directly with this method, because there are no strong resonances in the energy region investigated. Ca, Cu, and Fe were indicated by the supplier, but they produce no significant beam-induced γ background and contribute only to the total target thickness.

Because 22 Na and 23 Na have the same chemical properties, the 23 Na contaminant was used to measure the depth profile and the total thickness of the 22 Na target using the narrow ($\Gamma < 0.07 \text{ keV}$) $E_R = 0.677 \text{ MeV}$ resonance in 23 Na(p, γ) 24 Mg. 14 Prior tests showed that the amount of 23 Na on the backing itself is negligible. 13 The yield curve over the resonance is shown in Fig. 2. The observed resonance energy of 0.677 MeV agrees well with the known value, proving that the Na is on the surface of the target, as expected from previous tests. 13 The observed target thickness is \simeq 3 keV at $E_p = 0.677$ MeV with 1 keV of the thickness arising from the energy loss of the protons in the Si, and the remaining target thickness attributed to the heavier elements mentioned above. This measurement was repeated several times during the course of the experiment to monitor the target stability.

The γ radiation was observed with a 35% Ge detector at 0° with respect to the beam. The high γ -ray flux from the decay of ²²Na made it necessary to shield the detector from the target in order to keep the counting rate below 25 kHz (\simeq 40% computer dead time). To achieve this with the highest possible detection efficiency for the expected high-energy γ rays ($E_{\gamma} \simeq 8$ MeV) several tests were made. Mounting an Al target in the same geometry, the γ -ray yield of the known 0.992 MeV resonance in ²⁷Al(p,γ)²⁸Si (Ref. 14) for the transition $R \to 1.78$ MeV was measured relative to the yield of the $E_{\gamma} = 1.78$ MeV transition from the decay of the first excited state of ²⁸Si to its ground state. The optimal configuration is one which preferentially absorbs the low-energy γ rays while maintaining high efficiency for the high-energy γ rays. These measurements were made for several combinations of Pb thickness and target-

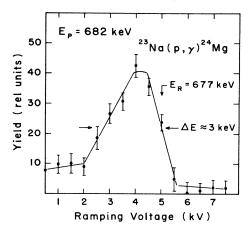


FIG. 2. Yield curve of the 23 Na $(p,\gamma)^{24}$ Mg resonance at $E_R = 677$ keV; for details see text.

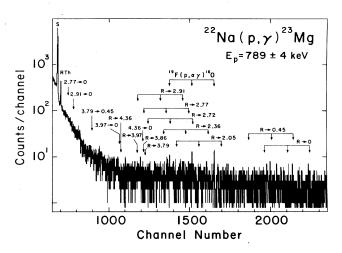


FIG. 3. Typical γ spectrum obtained at $E_p = 789 \pm 4$ keV. Indicated are the positions of analyzed γ transitions expected in ²³Mg; summing peaks are identified by the letter S.

detector distance. The best results were obtained with 22 mm of Pb between the target and the detector and a total detector distance of 35 mm.

A commercially available amplifier (ORTEC 572) was used to reject pileup events. The measured energy resolution of the detector at high rates ($\simeq 25~\mathrm{kHz}$) was 3.5 keV at $E_\gamma = 1.33~\mathrm{MeV}$. To determine the γ efficiency, the $^{22}\mathrm{Na}$ target was placed close to an Al target (100 $\mu\mathrm{g/cm^2}$) mounted in the target chamber to reproduce the high counting rate. The thick target γ -ray yield of the strong $^{27}\mathrm{Al}(p,\gamma)^{28}\mathrm{Si}$ resonance at 0.992 MeV was measured and from the known resonance strength 15 and branching ratios of the γ -ray decay, 16 the absolute detector efficiency was determined for γ energies between 2.4 and 11 MeV. At lower energies no reaction lines could be observed due to the high $^{22}\mathrm{Na}$ activity.

The γ -ray pulse height was stored along with the ramping voltage in a 4096 \times 16 data array for off-line

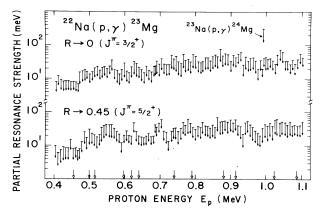


FIG. 4. Excitation function of the transition to the ground state (top) and the first excited state (bottom) of ²³Mg. Only every third point is plotted in the smooth regions of the excitation function. The open arrows at the bottom of the graph indicate the position of expected resonances.

analysis. The dead time was determined by standard pulser techniques.

III. DATA ANALYSIS AND RESULTS

A comparison with the γ branching ratios of states in the mirror nucleus $^{23}{\rm Na}$, in the same excitation energy range, suggests a decay mainly to the ground and first excited ($E_x=0.45~{\rm MeV}$) states of $^{23}{\rm Mg}$. The $E_\gamma=0.45~{\rm MeV}$ transition cannot be observed due to the intense γ background from the decay of $^{22}{\rm Na}$. Thus the two primary transitions were monitored in an on-line analysis during the course of the experiment. In a subsequent off-line analysis all possible transitions above the pileup peak of the 1.275 MeV γ ray were analyzed. A typical γ spectrum is shown in Fig. 3.

For the analysis of the data the 4096×16 data arrays were broken up into three arrays and the events were

TABLE II. Experimental upper limits of the partial resonance strengths (in meV) for the indicated transitions at the energies where states have been observed (Refs. 9 and 10).

E_p (lab)	$R \rightarrow E_x \text{ (MeV)}$									
(MeV)	0.0	0.45	2.05	2.36	2.72	2.77	3.79	3.86	3.97	4.36
0.456	9	9	11	13	19	17	11	9	12	8
0.499	14	25	35	32	28	22	19	16	15	21
0.515	23	21	17	19	23	22	11	14	13	13
0.597	30	20	78	25	25	23	14	14	19	15
0.619	17	21	650 ^a	50	30	43	21	18	17	32
0.640	18	18	53	35	35	42	14	18	13	14
0.740	30	44	46	42	48	55	50	50	48	29
0.791	39	44	23	32	30	30	30	30	29	26
0.853	55	35	56	39	59	59	36	79	34	43
0.881	72	68	330 ^a	220 ^a	300^{a}	117	100	168	150	100
0.915	56	43	145	430 ^a	100	100	62	60	64	65
1.024	27	70	60	60	100	63	57	48	50	45
1.087	40	39	150	99	280a	105	130	100	190	96
1.234	44	43	109	300 ^a	220 ^a	130	130	114	155	97
1.271	41	28	110	400 ^a	400 ^a	89	100	260 ^a	256a	110

^aOverlap with background line.

TABLE III. Observed and predicted strengths for resonances in 22 Na(p, γ) 23 Mg; for details see text.

E_x^a (MeV)	$J^{\pi\mathrm{b}}$	E_p (lab) (MeV)	Resonance st Expt.	trength (eV) Ref. 8°
8.014	$\frac{3}{2}$	0.456	$\leq 1.8 \times 10^{-2}$	2.4×10^{-1}
8.055	$\frac{\frac{5}{5}}{\frac{5}{2}}$ +	0.499	$\leq 3.9 \times 10^{-2}$	3.8×10^{-1}
8.071	5 +	0.515	\leq 4.4 \times 10 ⁻²	1.5
8.149	$\frac{3}{2}$ +	0.597	\leq 5.0 \times 10 ⁻²	4.3×10^{-1}
8.170	$\frac{5}{2}$ +	0.619	$\leq 3.8 \times 10^{-2}$	1.5
8.190	$\frac{7}{2}$	0.640	$\leq 3.6 \times 10^{-2}$	4.6×10^{-1}
8.285	-	0.740	\leq 7.4 \times 10 ⁻²	
8.334		0.791	$\leq 8.3 \times 10^{-2}$	
8.393		0.853	$\leq 9.0 \times 10^{-2}$	
8.420		0.881	$\leq 1.4 \times 10^{-1}$	
8.453		0.915	$\leq 9.9 \times 10^{-2}$	
8.557		1.024	$\leq 9.7 \times 10^{-2}$	
8.617		1.087	$\leq 7.9 \times 10^{-2}$	
8.758		1.234	$\leq 8.7 \times 10^{-2}$	
8.793		1.271	$\leq 6.9 \times 10^{-2}$	

^aReference 10 for $E_x \le 8.2$ MeV; Ref. 9 for $E_x \ge 8.2$ MeV.

projected on to the E_{γ} axis. The resulting three spectra represent the γ spectra for three consecutive ramping ranges of $\frac{1}{3}$ of the total ramping energy of 8 keV. The energies of possible primary transitions were calculated using the median energy for each energy step and a Q value of 7.578 MeV. The Doppler shift and γ recoil were included in these calculations. Regions of ± 2 channels ($\simeq \pm 7$ keV) around the expected positions were analyzed. This width includes the errors in the Q value (1.9 keV), energy calibration (2 keV), and incident proton energy (1 keV), which result in a total uncertainty of $\simeq 1$ channel. The width of high-energy γ -ray lines is estimated to 3 channels from the observed width of strong γ lines of $E_{\gamma} \simeq 8$ MeV, which were observed in the reactions 23 Na $(p,\gamma)^{24}$ Mg and 30 Si $(p,\gamma)^{31}$ P.

Since at the excitation energies of 23 Mg under investigation here, $E_x = 7.97 - 8.63$ MeV, no particle channels except protons are open, it can be assumed that the width Γ of possible resonances in 22 Na $(p, \gamma)^{23}$ Mg is smaller than the observed target thickness of ≈ 3 keV. This assumption is supported by the width of the corresponding states

in the analog nucleus 23 Na. 14 The γ yields can then be converted into partial resonance strengths $\omega \gamma_i$ using the well-known relation for the thick target yield of a narrow resonance: 19

$$\omega \gamma_i = \frac{2\epsilon}{\lambda^2} \frac{m_t}{m_p + m_t} \frac{I_{\gamma}}{(1 - \mathrm{DT}) \eta_{\gamma} N_p} \ ,$$

with ϵ the stopping power, λ the center-of-mass proton wave length, $m_t(m_p)$ the mass of the target (projectile), I_γ the intensity of the γ line, DT the dead time of the computer ($\simeq 40\%$), η_γ the γ efficiency and N_p the total number of incident particles. The stopping power ϵ was calculated from the ratio of the target thickness ΔE_t and the number of target atoms per cm². The target thickness as a function of the proton energy was determined from the excitation functions of the strong $^{29,30}\text{Si}(p,\gamma)^{30,31}\text{P}$ resonances at $E_R=0.416, 0.620$, and $0.983 \text{ MeV},^{14}$ and of the $^{23}\text{Na}(p,\gamma)^{24}\text{Mg}$ resonance at $E_R=0.677 \text{ MeV}$ (Sec. II). The results have not been corrected for possible angular distribution effects. Systematics of angular distributions of reactions with similar Q values 14 suggest a maximum correction of 50%.

No resonance was observed in any of the analyzed transitions for proton energies of $E_p = 0.400 - 1.100$ MeV. In addition, no resonances were observed for the states at $E_x = 8.758$ and 8.793 MeV. ¹⁰ The upper limits of the resonance strength for the ground-state transition and the transition to the first excited state are shown in Fig. 4. The resulting upper limits (1 standard deviation) for the partial resonance strengths for the transitions to the ground and first ten excited states in ²³Mg are listed in Table II, for proton energies that correspond to known states in ²³Mg. ^{9,10}

IV. DISCUSSION

To assign upper limits for the total resonance strengths of the expected resonances, assumptions about the dominant γ decay of these states have to be made. As already pointed out by Wiescher and Langanke, a comparison with the mirror nucleus ²³Na suggests γ decay mainly to the ground and first excited state. Adopting this assumption yields the upper limits for the total resonance strengths listed in Table III. For comparison, the estimates by Wiescher and Langanke are also given. Their values are corrected for slight changes in the excitation energy to by scaling the proton widths with the relative change of the Coulomb penetrabilities. The resulting

TABLE IV. Comparison of shell-model calculations and experiment (Ref. 14) for the transition to the ground state and to the first excited state in the analog nucleus ²³Na.

E_{i}		$\Gamma_{_{\mathcal{V}}}$ (eV) (calc.)	Γ_{ν} (eV) (expt.; Ref. 14)		
(MeV)	J^{π}	g.s.	0.44	g.s.	0.44	
2.08	$\frac{7}{2}$ +	2.7×10^{-3}	1.5×10^{-2}	$(1.5\pm0.2)\times10^{-3}$	$(1.5\pm0.2)\times10^{-2}$	
2.39	$\frac{1}{2}$ +	4.2×10^{-4}	2.1×10^{-4}	$(5.3\pm1.0)\times10^{-4}$	$(2.9\pm0.5)\times10^{-4}$	
2.98	$\frac{3}{2}$ +	6.5×10^{-2}	4.0×10^{-2}	$(7.6\pm1.5)\times10^{-2}$	$(5.5\pm1.1)\times10^{-2}$	
3.91	5 +	2.6×10^{-2}	1.1×10^{-3}	$(4.9\pm0.9)\times10^{-2}$	$(4.8\pm1.1)\times10^{-3}$	

^bTentative spin assignment adopted from Ref. 8.

^cCorrected for new values of E_x (Ref. 9).

TABLE V. Shell-model predictions of the γ transition width $(2J+1)\Gamma_{\gamma}$ for the transition to the ground state and to the first excited state in ²³Mg in the excitation range of $E_x = 8-9$ MeV.

	$(2J+1)\Gamma_{\gamma}$ (eV)			
J^{π}	$E_i \rightarrow g.s.$	$E_i \rightarrow 0.45$		
$\frac{1}{2}$ +	0.7 -0.8	$3 \times 10^{-4} - 7 \times 10^{-3}$		
$\frac{3}{2}$ +	0.4 -0.8	$5 \times 10^{-2} - 2$		
$\frac{3}{2}$ + $\frac{5}{2}$ + $\frac{7}{2}$ +	2 -4	$2 \times 10^{-3} - 4$		
$\frac{7}{2}$ +	0.06-0.3	$2 \times 10^{-1} - 4$		
$\frac{9}{2}$ +		$1 \times 10^{-5} - 3 \times 10^{-1}$		

upper limits (Table III) are in all cases but one $(E_R = 0.881 \text{ MeV})$ below 100 meV and consequently at least a factor of 10 smaller than the estimates⁸ in the overlapping energy range. This result is somewhat surprising since many $(p, \dot{\gamma})$ reactions with similar Q values in this mass region exhibit at least one resonance^{14,17} with a strength of 0.5–1.0 eV in the same proton energy range as covered in this experiment, e.g., $^{22}\text{Ne}(p, \gamma)^{23}\text{Na}, ^{23}\text{Na}(p, \gamma)^{24}\text{Mg}$. One exception²⁰ is the reaction $^{26}\text{Al}(p, \gamma)^{27}\text{Si}$, where the resonance strengths for all observed resonances are smaller than 100 meV.

At proton energies above 0.4 MeV, the proton width is, in general, larger than the corresponding γ width. Therefore the present results indicate much smaller γ widths than previously assumed. Since no experimental information exists, Wiescher and Langanke assumed γ widths deduced from the average of the observed transition-strength distributions $(E1=3\times10^{-3})$ W.u., M1=0.3 W.u., E2=3 W.u.). However, these distributions cover a range of 5 orders of magnitude and it is therefore entirely possible that the choice of the averages of these distributions could overestimate the actual γ widths by as much as 2 orders of magnitude. Furthermore, most of the known values for Γ_{γ} are for transitions between states below an excitation energy of 4 MeV. At these excitation energies the nuclear states have a

predominantly single-particle structure, while at the higher excitation energies of interest here, $E_x = 8-9$ MeV, the states are more likely of 2p-2h nature, indicating smaller γ transition strengths.

To obtain a more realistic estimate for the γ widths, shell-model calculations for an A = 23 nucleus were performed. Using the code OXBASH (Ref. 22) and Wildenthal's universal sd interaction²³ for the mass range A = 17-39, the energies of the positive-parity states $(J^{\pi} \le \frac{9}{2}^+)$ were calculated up to an excitation energy of 12 MeV. Subsequently, the reduced γ transition probabilities for M1 and E2 transitions for all of these states to the ground and the first excited state were calculated. For the analog nucleus ²³Na, the shell-model results are in reasonably good agreement with experimental values (Table IV), which were derived for states with known half-lifes and γ branching ratios. ¹⁴ To compare the theoretical values with the present experimental upper limits, $(2J+1)\Gamma_{\gamma}$ was calculated for the transitions to the ground and first excited states. The shell-model results range from 0.06 eV to 4 eV for the ground-state transition and from 1×10^{-5} eV to 4 eV for the transition to the first excited state (Table V). The experimental upper limits for the corresponding transitions are $\leq 0.13 - \leq 1.0$ eV and $\leq 0.13 - \leq 1.0$ eV, respectively. Both experimental and theoretical values are, on the average, an order of magnitude lower than previously predicted.

To obtain reliable experimental stellar reaction rates further experiments with an improved sensitivity are necessary. This can be achieved either by the use of improved targets, e.g., isotope-separated, implanted targets, or by the use of the recently developed heavy water (D_2O) detector.¹²

ACKNOWLEDGMENTS

The authors wish to thank R. W. Kavanagh and R. E. Azuma for many useful discussions and B. Leist for help during the course of the experiment. This work was supported by National Science Foundation Grants PHY84-21302, PHY85-05682, and PHY87-14432.

¹R. K. Wallace and S. E. Woosley, Astrophys. J. Suppl. **45**, 389 (1981)

²P. Eberhardt, M. H. A. Jungck, I. O. Meier, and F. Niederer, Astrophys. J. Lett. 234, L169 (1979).

³J. B. Marion and W. A. Fowler, Astrophys. J. **125**, 221 (1957).

⁴C. Rolfs, W. S. Rodney, M. A. Shapiro, and H. Winkler, Nucl. Phys. **A241**, 460 (1970).

⁵J. Görres, H. W. Becker, L. Buchmann, C. Rolfs, P. Schmalbrock, H. P. Trautvetter, A. Vlieks, J. W. Hammer, and T. R. Donoghue, Nucl. Phys. A408, 372 (1983).

⁶M. Wiescher, J. Görres, F. K. Thielemann, and H. Ritter, Astron. Astrophys. 160, 56 (1986).

⁷M. Wiescher, J. Görres, B. Sherrill, M. Mohar, J. S. Winfield, and B. A. Brown, Nucl. Phys. A484, 90 (1988).

⁸M. Wiescher and K. H. Langanke, Z. Phys. A **325**, 309 (1986).

⁹H. Nann, A. Saha, and B. H. Wildenthal, Phys. Rev. C 23, 606 (1981).

¹⁰C. P. Browne, M. Wiescher, A. A. Rollefson, R. W. Tarara, P. Schmalbrock, V. Wijekumar, T. R. Donoghue, and H. J. Hausman, University of Notre Dame, Biennial Report, 1987, p. 158; P. Schmalbrock, T. R. Donoghue, H. J. Hausman, M. Wiescher, V. Wijekumar, C. P. Browne, and A. A. Rollefson, in Capture Gamma-Ray Spectroscopy and Related Topics (Holiday Inn-World's Fair, Knoxville, Tennessee), Proceedings of the Fifth International Symposium on Capture Gamma-Ray Spectroscopy and Related Topics, AIP Conf. Proc. No. 125, edited by S. Raman (AIP, New York, 1984), p. 785.

¹¹C. Rolfs, H. P. Trautvetter, and W. S. Rodney, Rep. Prog. Phys. **501**, 3 (1987).

¹²C. Rolfs and R. W. Kavanagh, Nucl. Instrum. Methods A244, 507 (1986).

¹³M. Wiescher, J. Görres, K. L. Kratz, B. Leist, K. H. Chang, B. W. Filippone, L. W. Mitchell, M. J. Savage, and R. B.

- Vogelaar, Nucl. Instrum. Methods A267, 242 (1988).
- ¹⁴P. M. Endt and C. van der Leun, Nucl. Phys. **A310**, 1 (1978).
- ¹⁵P. B. Lyons, J. W. Toevs, and D. S. Sargood, Nucl. Phys. A130, 1 (1969).
- ¹⁶A. Antilla, J. Keinonen, M. Hautala, and I. Forsblom, Nucl. Instrum. Methods 147, 501 (1977).
- ¹⁷M. A. Meyer and J. J. Smit, Nucl. Phys. **A205**, 177 (1973).
- ¹⁸A. H. Wapstra and G. Audi, Nucl. Phys. **A432**, 55 (1985).
- ¹⁹C. Rolfs and A. E. Litherland, in *Nuclear Spectroscopy and Reactions*, edited by J. Cerny (Academic, New York, 1974), p.
- 143.
- ²⁰L. Buchmann, M. Hilgemeier, A. Krauss, A. Redder, C. Rolfs, H. P. Trautvetter, and T. R. Donoghue, Nucl. Phys. A415, 93 (1984).
- ²¹P. M. Endt, At. Data Nucl. Data Tables **23**, 3 (1979).
- ²²B. A. Brown, A. Etchegoyen, and W. D. M. Rae, Michigan State University Cyclotron Laboratory Report 524, 1985.
- ²³B. H. Wildenthal, in *Progress in Particle and Nuclear Physics*, edited by D. H. Wilkinson (Pergamon, Oxford, 1984), Vol. 11, p. 5.