Direct measurement of the low-energy resonances in $^{22}Ne(\alpha, \gamma)$ ²⁶Mg reaction

Shahina,^{1,2} J. Görres,^{1,2} D. Robertson,^{1,2} M. Couder,^{1,2} O. Gomez,^{1,2} A. Gula,^{1,2} M. Hanhardt,^{3,4} T. Kadlecek,³

R. Kelmar^(D),^{1,2} P. Scholz^(D),^{1,2} A. Simon^(D),^{1,2} E. Stech,^{1,2} F. Strieder^(D),³ and M. Wiescher^(D),²

¹Department of Physics and Astronomy, University of Notre Dame, Notre Dame, Indiana 46556, USA

²The Joint Institution of Nuclear Astrophysics-Center for the Evolution of the Elements,

University of Notre Dame, Notre Dame, Indiana 46556, USA

³Department of Physics, South Dakota School of Mines and Technology, Rapid City, South Dakota 57701, USA

⁴South Dakota Science and Technology Authority, Sanford Underground Research Facility, Lead, South Dakota 57754, USA

(Received 10 May 2022; revised 8 July 2022; accepted 12 August 2022; published 22 August 2022)

The ²²Ne(α, γ) ²⁶Mg is an important reaction in stellar helium burning environments as it competes directly with one of the main neutron sources for the s-process, the ²²Ne(α, n) ²⁵Mg reaction. The reaction rate of the ²²Ne(α, γ) ²⁶Mg is dominated by the low-energy resonances at $E_{\alpha}^{lab} = 650$ and 830 keV, respectively. The $E_{\alpha}^{lab} = 830$ -keV resonance has been measured previously, but there are some uncertainties in the previous measurements. We confirmed the measurement of the $E_{\alpha}^{lab} = 830$ -keV resonance using implanted ²²Ne targets. We obtained a resonance strength of $\omega\gamma = 35 \pm 4 \mu eV$ and provide a weighted average of the present and previous measurements of $\omega\gamma = 35 \pm 2 \mu eV$ with reduced uncertainties compared to previous studies. We also attempted to measure the strength of the predicted resonance at $E_{\alpha}^{lab} = 650$ keV directly for the first time and found an upper limit of $\omega\gamma < 0.15 \mu eV$ for the strength of this resonance. In addition, we also studied the $E_{\mu}^{lab} = 851$ -keV resonance in ²²Ne(p, γ) ²³Na and obtained a resonance strength of $\omega\gamma = 9.2 \pm 0.7$ eV with significantly lower uncertainties compared to previous measurements.

DOI: 10.1103/PhysRevC.106.025805

I. INTRODUCTION

The slow neutron capture process (s-process) is responsible for the creation of half the elements beyond iron. It takes place when the timescale between successive neutron captures is much larger than their lifetime of β decay [1]. Hence, less neutron-rich nuclei are produced along the line of β stability. The s-process can be categorized into two types: the weak s-process responsible for the creation of elements over the range 60 < A < 90, and the main s-process which creates nuclei with A > 90 [2]. The temperature range of interest is 0.2 to 0.6 GK for both scenarios. Both processes require a sufficiently large neutron flux. However, the source of these neutrons is still under debate. In asymptotic giant branch (AGB) stars, the main s-process nucleosynthesis occurs during the late stages of stellar evolution when the star has a degenerate C-O core, a thin radiative He intershell, and an expanded convective envelope. During the AGB phase, the star experiences a series of flashes called thermal pulses which are triggered by the sudden activation of the 3α process at the base of the He intershell under highly degenerate gas conditions [2]. In this intershell environment, two important reactions are activated, ${}^{13}C(\alpha, n) {}^{16}O$ prior to the thermal pulse and ²²Ne(α , n)²⁵Mg during the thermal pulse, which act as the main sources of free neutrons for the s-process. Both 22 Ne(α , n) 25 Mg and 13 C(α , n) 16 O provide the neutron flux in low-mass AGB stars, ${}^{13}C(\alpha, n) {}^{16}O$ being the dominant contributor. The ¹³C buildup occurs in a thin zone (the ¹³C pocket) via the capture of mixed-in hydrogen on the abundant ${}^{12}C$ via ${}^{12}C(p, \gamma) {}^{13}N(\beta^+\nu) {}^{13}C$. ${}^{13}C(\alpha, n) {}^{16}O$ burns radiatively for an extended time, releasing a high neutron flux when the temperature exceeds 0.8×10^8 K [2]. The second dominant neutron source, ${}^{22}Ne(\alpha, n) {}^{25}Mg$, has a negative Q value of -478 keV and thus requires a sufficiently high temperature. Such conditions are reached only during the advanced thermal pulses in low-mass AGB stars and the core helium burning phase of massive stars.

 22 Ne(α , n) 25 Mg also acts as the main neutron source for the weak s-process in core helium burning of massive stars [3]. The 22 Ne is produced from the large amount of 14 N left at the end of the CNO cycle in the Helium burning core via the following reaction chain:

¹⁴N(
$$\alpha, \gamma$$
)¹⁸F(β^+, ν)¹⁸O(α, γ)²²Ne. (1)

However, because of its negative Q value the temperature during core helium burning is not sufficient to trigger the ignition of ${}^{22}\text{Ne}(\alpha, n) {}^{25}\text{Mg}$ at once. Only towards the end of the helium burning phase with the stellar core gradually contracting, high temperature and density conditions sufficient for the above reaction are reached. Hence, the ${}^{22}\text{Ne}(\alpha, n) {}^{25}\text{Mg}$ reaction is operative only during the peak of the helium flash and at the end of core helium burning.

At lower temperature the neutron-producing role of ${}^{22}\text{Ne}(\alpha, n){}^{25}\text{Mg}$ is complicated by the competing ${}^{22}\text{Ne}(\alpha, \gamma){}^{26}\text{Mg}$ reaction. The ${}^{22}\text{Ne}(\alpha, \gamma){}^{26}\text{Mg}$ reaction has a positive Q value = 10614.74 ± 0.03 keV and, therefore, starts operating at relatively low temperature before ²²Ne(α , n) ²⁵Mg can kick in. As a consequence, it is important to investigate the reaction rate of ²²Ne(α , γ) ²⁶Mg in order to constrain the neutron production for the weak s-process.

At stellar helium burning temperature, low-energy resonances at $E_{\alpha}^{\text{lab}} = 650$ and 830 keV lie within the Gamow window and dominate the $^{22}\text{Ne}(\alpha, \gamma)^{26}\text{Mg}$ reaction rate. $^{22}\text{Ne}(\alpha, \gamma)^{26}\text{Mg}$ proceeds through the resonant capture reaction mechanism, thus populating natural parity states in the compound nucleus ^{26}Mg [4]. Above T = 0.3 GK, the rate is dominated by the $E_{\alpha}^{\text{lab}} = 830$ -keV resonance which has been observed in both (α, n) and (α, γ) direct measurement experiments.

The ²²Ne(α , γ) ²⁶Mg reaction was first measured directly by Wolke *et al.* [5] for $E_{\alpha}^{\text{lab}} = 0.71$ to 2.25 MeV. They found 15 new resonances in the energy range covered, with the lowest energy resonance at 830 keV. They concluded that the reaction rate is strongly impacted by the 830-keV resonance, whose resonance strength was measured to be $\omega \gamma =$ $36 \pm 4 \mu eV$. For this measurement, they used a differentially pumped gas target system with neon gas enriched to 99% in ²²Ne. The γ -ray transitions were detected with Ge(Li) detectors. A recent direct measurement of this resonance was also performed by Hunt et al. [6] who obtained its resonance strength to be $\omega \gamma = 46 \pm 12 \ \mu \text{eV}$. Unlike Wolke *et al.* [5], they used implanted ²²Ne targets to determine the resonance strength from the thick target yield curve. They detected the entire cascade of γ rays using a $\gamma\gamma$ -coincidence spectrometer, consisting of a high-purity germanium (HPGe) detector surrounded by a NaI(Tl) annulus.

Complementary to the direct measurements, Talwar et al. [4] performed indirect measurements using (⁶Li, d) α -transfer and (α, α') inelastic scattering techniques to populate states that are most likely to appear as resonances in the ²²Ne(α, γ)²⁶Mg reaction. They observed a strong transition in both (⁶Li, d) and (α, α') measurements at $E_x = 11.167$ MeV, which corresponds to a resonance in the ²²Ne + α system at $E_{\alpha}^{\text{lab}} = 650$ keV. They assigned a spin-parity of $J^{\pi} = (1^{-}, 2^{+})$ to this state and obtained a spectroscopic factor of $S_{\alpha} = 0.36$ (corresponding to $\Gamma_{\alpha} =$ 0.18 μ eV). Based on this, they suggested a resonance strength of $\omega \gamma = 0.54 \pm 0.07 \ \mu \text{eV}$. They concluded that this state completely dominates the ${}^{22}Ne(\alpha, \gamma){}^{26}Mg$ reaction rate between $T \simeq 0.2$ GK and $T \simeq 0.4$ GK and would also potentially dominate the 22 Ne(α , n) 25 Mg reaction rate below 0.2 GK. Therefore, it is essential to measure this low-energy resonance in ${}^{22}Ne(\alpha, \gamma) {}^{26}Mg$ in order to accurately predict the total neutron flux for the weak s-process.

In this work, we present the direct measurements of these two low-energy resonances at $E_{\alpha}^{lab} = 650$ and 830 keV that play an important role in the reaction rate of $^{22}Ne(\alpha, \gamma)$ ^{26}Mg . Several resonances are expected below the 650-keV resonance [4]. However, the penetrability rapidly drops with beam energy and the corresponding α widths are at least 2 or 3 orders of magnitude lower. Hence, the resulting resonance strength would be well below the sensitivity of this experiment. For this reason, we concentrated on the search for the 650-keV resonance, which would have the largest impact on



FIG. 1. Absolute γ -ray detection efficiency as a function of energy for the Ge detector used for the determination of target stoichiometry.

the neutron production for the weak s-process if the predicted strength is confirmed [4].

II. EXPERIMENTAL SETUP

A. Target preparation and characterization

The ²²Ne targets were made and tested at the University of Notre Dame using the 5U St. Ana Pelletron accelerator. Ne gas enriched in ²²Ne to 99% was used to produce a ²²Ne beam with an energy of 200 keV, which was implanted onto thick Ta backings, $3.8 \text{ cm} \times 3.8 \text{ cm}$ in size. Tantalum was used as a backing, because it has a high concentration of ²²Ne at implantation saturation [7] and beam-induced reactions with the backing are minimized. The ²²Ne beam shape was defined by slits in front of the target, and the beam was wobbled in both x and y directions to have uniform implantation of 22 Ne onto Ta. To measure the saturation curve several ²⁰Ne targets with increasing implantation dosages of 30, 60, 90, 120, 150, 180, 225, 270, and 300 mC were produced. The saturation was achieved near 300 mC, and hence all the implanted ²²Ne targets used during the experiment had a 300 mC dosage over a target area of 3 cm^2 .

The implantation profile of the ²²Ne targets was investigated using the 1278-keV resonance in the 22 Ne (p, γ) 23 Na reaction [8]. The efficiency of the emitted γ rays was measured by using a p-type coaxial germanium detector operating at +4000 V. The detector was placed at an angle of 55° with respect to the beam direction to minimize angular distribution effects. The distance of the detector from the target was 20 cm to reduce the summing effects. Moreover, a 1-mm-thick lead sheet was attached to the front face of the detector to reduce the γ -ray flux from inelastic scattering from the Ta backing. The absolute full-energy peak efficiency was measured using calibrated radioactive sources like ⁵⁶Co and ⁶⁰Co. The sources were placed exactly at the target position, 20 cm from the detector to have the same geometry. The efficiency curve was extended to 10.76 MeV using the well-known resonance at 992 keV in the ${}^{27}\text{Al}(p,\gamma){}^{28}\text{Si}$ reaction [9]. The absolute efficiency curve for the Ge detector is shown in Fig. 1. It was obtained by normalizing the efficiency at higher energies, to the efficiency of the 1779-keV γ -ray in the



FIG. 2. Yield curve for the $E_P^{\text{lab}} = 1278$ keV resonance in the ${}^{22}\text{Ne}(p, \gamma){}^{23}\text{Na}$ reaction.

²⁷Al (p, γ) ²⁸Si resonance, taking into account the well-known branching ratios and angular distributions [9]. The relative efficiency was converted into absolute efficiency using the yield value of $(1.08 \pm 0.06) \times 10^{-9} \gamma$ rays (1779 keV) per incident proton [9], which corresponds to a resonance strength of 1.93 ± 0.13 eV [10].

The stoichiometry of the ²²Ne targets was determined using the well-known resonance at 1278 keV in ${}^{22}Ne(p, \gamma){}^{23}Na$ [8] using the thick-target yield formula [11],

$$\omega\gamma = \frac{2Y\epsilon_{\rm eff}}{\lambda^2\eta},\tag{2}$$

where *Y* is the yield, η is the efficiency, λ is the deBroglie wavelength of the incident particle, and ϵ_{eff} is the effective stopping power of protons given by

$$\epsilon_{\rm eff} = \frac{M_{\rm Ne}}{M_P + M_{\rm Ne}} \Big[\epsilon_{\rm Ne} + \Big(\frac{N_{\rm Ta}}{N_{\rm Ne}}\Big) \epsilon_{\rm Ta} \Big], \tag{3}$$

where $N_{\text{Ta}}/N_{\text{Ne}}$ is the initial target stoichiometry, M_P and M_{Ne} are the masses of the projectile and target nuclei (in atomic mass units), while ϵ_{Ne} and ϵ_{Ta} denote the stopping power of the protons in Ne and Ta, respectively, which were obtained using SRIM [12]. The quoted uncertainties of these values for the relevant proton energy are 1.6% and 3% for Ne and Ta, respectively.

The yield curve of the 1278-keV resonance was measured with protons from the St. Ana accelerator utilizing the same target station that was used for the Ne implantation. The yield curve is shown in Fig. 2, which is obtained after taking the average of the absolute yield calculated using the following γ -ray transitions: 2540, 2980, 3914, 6102, 7036, and 9576 keV. The yield from the individual γ rays was corrected with the branching ratio obtained from Ref. [13] and the angular distribution obtained from Ref. [14]. The yield curve in Fig. 2 depicts the distribution of the implanted ²²Ne in the Ta backing. In order to obtain the stoichiometry of the implanted target, Eq. (2) was first used to calculate the effective stopping power of the incident protons, using the resonance strength of $\omega \gamma = 10.8 \pm 0.7$ eV from Ref. [8]. The effective stopping power thus obtained was then substituted in Eq. (3) to compute the stoichiometry of the target at maximum yield, which turned out to be $N_{\text{Ta}}/N_{\text{Ne}} = 2.7 \pm 0.3$. The total number *n* of Ne nuclei implanted onto the Ta backing was thereafter obtained using the following formula [11],

$$n = \frac{2A_Y}{\lambda^2 \omega \gamma},\tag{4}$$

where A_Y is the area under the ²²Ne(p, γ)²³Na resonance yield curve. A_Y was obtained by numerical integration of the excitation curve shown in Fig. 2. The total number of Ne nuclei implanted, n, was found to be $(6.21 \pm 0.37) \times 10^{17}$ atoms/cm². The error of the integration and the statistical errors are small compared to the uncertainty of the resonance strength. The value obtained for the total number of implanted Ne nuclei is consistent with the implantation dose within uncertainty and indicates an implantation efficiency of nearly 100%.

B. CASPAR accelerator

The measurement of the low-energy resonances in 22 Ne(α, γ) 26 Mg was performed at CASPAR (Compact Accelerator System for Performing Astrophysical Research), located on the 4850-ft level of the Sanford Underground Research Facility in Lead, South Dakota, at the former site of the Homestake Gold Mine [15,16]. The rock above acts as an effective shield from the cosmic-ray-induced background, thus helping to measure the weak resonances that have a low signal-to-background ratio. The 1-MV JN positive ion accelerator at CASPAR was used to deliver protons and α beams onto the target with intensity ranging from 10 to 100 μ A. The energy calibration of the beam was performed by measuring the front edges of the resonances in ${}^{18}O(\alpha, \gamma)^{22}Ne$ at 750 and 767 keV [17] and scanning the well-known resonances in ${}^{27}\text{Al}(p, \gamma){}^{28}\text{Si}$ [18]. The beam energy uncertainty amounted to ± 2 keV. The beam was sent through an analyzing magnet that bent it by 25° before transporting it to the experimental end station. The beam was defocused and delivered to the target. The size of the beam spot was nearly 1 cm in diameter. The target was mounted inside the High EffiCiency TOtal absorption spectrometeR (HECTOR) [19] at the end of the beamline, as shown in Fig. 3. The setup is identical to the setup used for the measurement of ${}^{27}\text{Al}(p, \gamma){}^{28}\text{Si}$ resonances [20] and the details of the setup are shown in Fig. 2 of Ref. [20]. The electrically isolated target chamber acted as a Faraday cup, which was used for the integration of the beam current. To limit the buildup of carbon on the target, a copper tube cooled to liquid nitrogen temperature was placed inside the beamline, extending from the cold trap up to the target surface. The cold trap was electrically isolated and biased to -300 V in order to suppress the secondary electrons. The target was also water cooled in order to reduce the damage caused by heating due to high beam intensities.

C. *y*-summing detector: HECTOR

HECTOR consists of an array of 16 NaI(Tl) crystals with two photomultiplier tubes attached to each segment. Each



FIG. 3. Experimental setup showing HECTOR placed at the end of the beamline at the CASPAR underground laboratory. The target was placed at the center of the detector cube.

NaI(Tl) crystal is housed in an aluminum casing. The crystals are arranged in a frame to form a cube $16 \times 16 \times 16$ in. in dimension. The center of the detector array consists of a 60-mm borehole that allows the target to be placed at the center giving a 4π total coverage for the detection of γ rays. The advantage of using HECTOR for the ²²Ne(α, γ)²⁶Mg measurement is its high detection efficiency. The signal from each photomultiplier tube is recorded individually and then gain-matched and summed off-line to improve the detector resolution. A digital data acquisition system is used to readout the data from the 16 NaI(Tl) detectors. A detailed description of the HECTOR and its data acquisition can be found in Ref. [19]. In this work, known level schemes were used to determine the summing efficiency of HECTOR for the measured resonances. This approach was introduced and tested for the ${}^{27}Al(p, \gamma){}^{28}Si$ resonances in the recent work of Ref. [20] and was applied in the analysis of the ¹⁸O(α , γ)²²Ne reaction [21].

III. EXPERIMENTAL PROCEDURE AND ANALYSIS

The low-energy resonances at $E_{\alpha}^{\text{lab}} = 650$ and 830 keV in the ²²Ne(α, γ)²⁶Mg reaction were measured using the implanted ²²Ne targets (as discussed in Sec. II A) at CAS-PAR [15]. The stability of the implanted ²²Ne targets was monitored throughout the course of the experiment using the well-known resonances at $E_p^{\text{lab}} = 479$ keV and 851 keV in the ²²Ne(p, γ)²³Na reaction [8]; the 1278-keV resonance of this reaction is not accessible with the CASPAR accelerator. Figure 4 shows the yield curve for the 479-keV resonance of ²²Ne(p, γ)²³Na for one of the ²²Ne targets. The yield



FIG. 4. Yield curve for the $E_p^{\rm lab} = 479$ -keV resonance in the $^{22}\text{Ne}(p, \gamma)^{23}\text{Na}$ reaction. The yield curve is obtained from the sum peak in HECTOR [19]. The solid-circle (blue) and open-circle (orange) data points show the difference in yield at the beginning and at end of the experiment. The statistical error in the data is $\simeq 1\%$, which is why the error bars are not visible.

is obtained using the sum peak in HECTOR [19] at $E_{\Sigma} = E_{\rm CM} + Q = 9252.27$ keV (where $E_{\rm CM}$ is the center-of-mass energy of the projectile target system and Q is the reaction Q value). The solid circle (blue) and open circle (orange) data points show the difference in the yield at the beginning and the end of the experiment, after accumulating a charge of nearly 4 C. The targets were fairly stable and the yield decreased only by 5% at the end of the experiment, as shown in Fig. 4. No heat damage or blistering was caused by the high intensity of the beam current. As a consequence, we changed each target after it has been exposed to an accumulated charge of around 4 C.

To optimize the experimental yield of the narrow resonance at 830 keV in 22 Ne(α, γ) 26 Mg, it has to be measured at the maximum of the ²²Ne distribution. For this purpose, the energy loss of protons from the resonance energy of 479 keV till the maximum of the ²²Ne distribution (as shown in Fig. 4) in 22 Ne (p, γ) 23 Na was converted to the energy loss of α particles in ²²Ne(α, γ)²⁶Mg using the relative stopping powers from SRIM [12]. This implies that the maximum of the ²²Ne distribution for the α -particle beam would correspond to an α energy of 910 keV for the $E_{\alpha}^{\text{lab}} = 830$ keV resonance. The ²²Ne concentration is constant to 10% within ± 30 keV of this energy. To confirm the position of the top of the thick-target yield curve/plateau, a scan of the $E_{\alpha}^{\text{lab}} = 830$ -keV resonance was performed using large energy steps. Thus data were acquired at the top of the plateau, i.e., at $E_{\alpha}^{\text{lab}} = 910$ keV, with a total accumulated charge of Q = 2.94 C on the ²²Ne target. The beam-induced background was measured using an implanted ²⁰Ne target at the same α -particle energy. The sum peak spectrum for the 830-keV resonance is shown in Fig. 5(a). The solid (blue) histogram shows the on-resonance spectrum taken with a ²²Ne target, where the sum peak is located at an energy of $E_{\Sigma} = 11.3$ MeV. The dashed (orange) histogram shows the



FIG. 5. (a) Sum spectra for the $E_{\alpha}^{\text{lab}} = 830\text{-keV}$ resonance. The solid (blue) histogram shows the sum spectrum from the $^{22}\text{Ne}(\alpha, \gamma)^{26}\text{Mg}$ reaction and the sum peak is located at $E_{\Sigma} = E_{\text{CM}} + Q = 11.3$ MeV. The dashed (orange) histogram shows the beam-induced background using the ^{20}Ne target. The dash-dotted (brown) histogram shows the simulated spectrum obtained from GEANT4 [22]. The thick solid (green) line shows the Gaussian plus a linear function fit to the sum peak. (b) The dashed (orange) histogram is the sum of beam-induced background and simulation. It shows excellent agreement with the experimental spectrum shown in solid (blue).

background spectrum obtained using a ²⁰Ne target. The sum peak from the background ²⁰Ne(α, γ)²⁴Mg reaction is located at $E_{\Sigma} = 10.02$ MeV, which is well separated from the sum peak of ²²Ne(α, γ)²⁶Mg.

Computation of the strength of the 830-keV resonance requires the experimental yield (discussed above) as well as the efficiency of the sum peak, as evident from Eq. (2). The efficiency of the 830-keV resonance sum peak was obtained using GEANT4 [22] simulations. The level scheme and the branching ratio for the reaction were taken from the previous measurements performed in Refs. [5,6]. The entire γ cascade was simulated and the resultant simulated spectrum is shown in Fig. 5(a), as the dash-dotted (brown) histogram. Adding the beam-induced background to the simulation shows excellent agreement with the ²²Ne(α , γ)²⁶Mg experimental spectrum as shown in Fig. 5(b).

The sum peak was fitted with a Gaussian plus a linear function, shown as a thick solid (green) line in Fig. 5, and was integrated within the $E_{\Sigma} - 3\sigma$ to $E_{\Sigma} + 3\sigma$ range to get the total number of events under the peak. The simulated sum peak was fitted using the same procedure. The efficiency is given by the ratio of the number of counts in the simulated sum peak to that of the total simulated cascade events.

The strength of the 830-keV resonance in 22 Ne(α , γ) 26 Mg was determined relative to the well-known 479-keV resonance



FIG. 6. Sum spectra for the $E_{\alpha}^{\text{lab}} = 650$ -keV resonance. The solid (blue) histogram depicts the sum spectrum from the ²²Ne(α, γ) ²⁶Mg reaction, whereas the dashed (orange) histogram shows the beam-induced background using the ²⁰Ne target. The two solid vertical lines show the region of integration for the sum peak. The background (orange) histogram was normalized to the same accumulated charge.

in ${}^{22}\text{Ne}(p, \gamma){}^{23}\text{Na}$ [8] using the expression for the maximum yield of an infinitely thick target [11],

$$\frac{\omega\gamma_1}{\omega\gamma_2} = \left(\frac{\epsilon_{r,1}}{\epsilon_{r,2}}\right) \left(\frac{\lambda_{r,2}^2}{\lambda_{r,1}^2}\right) \left(\frac{Y_1}{Y_2}\right) \left(\frac{\eta_2}{\eta_1}\right),\tag{5}$$

where the subscripts 1 and 2 correspond to the resonance of interest and the standard resonance. In this approach, only relative values of the stopping powers and efficiencies are needed. The efficiency of the 479-keV resonance in 22 Ne (p, γ) 23 Na (η_2) was obtained in the same way as the 830-keV resonance, using the level scheme and branching ratio from Ref. [23]. The error in the resonance strength $\omega \gamma_1$ depends on the accuracy of $\omega \gamma_2$, and the uncertainties of the ratios of stopping powers, efficiencies, and yields. Using this relative approach, we obtained $\omega \gamma = 35 \pm 4 \ \mu eV$ for this resonance. The uncertainty in the resonance strength is dominated by the uncertainty of $\omega \gamma_2$ ($E_P^{\text{lab}} = 479$ -keV resonance), which is 6.4%, and the relative stopping power of α 's and protons in Ta. The uncertainty in the stopping power for α 's in Ta is 6% and for protons in Ta the uncertainty is 3.4%. The statistical uncertainty associated with the sum peak integral is 3.5% for the 830-keV resonance and 1.4% for the 479-keV resonance. The error of the relative efficiency is 5%, mainly resulting from the uncertainty in the branching ratio of the 830-keV resonance in 22 Ne(α , γ) 26 Mg.

The other low-energy resonance in ²²Ne(α, γ)²⁶Mg at $E_{\alpha}^{\text{lab}} = 650 \text{ keV}$ was measured in the same way as the $E_{\alpha}^{\text{lab}} = 830 \text{ keV}$ resonance. Data were collected at $E_{\alpha}^{\text{lab}} = 711 \text{ keV}$, with a total accumulated charge of Q = 22.9 C on top of the resonance. To determine the beam-induced background, data were taken with a ²⁰Ne target at the same α -particle energy, with a total accumulated charge of Q = 6.52 C. The sum spectrum for the $E_{\alpha}^{\text{lab}} = 650 \text{ keV}$ resonance is shown in Fig. 6. The solid (blue) histogram shows the on resonance spectrum

taken with a ²²Ne target, where the sum peak is located at an energy of $E_{\Sigma} = 11.16$ MeV. The dashed (orange) histogram shows the background spectrum that was taken with the ²⁰Ne target. The width of the region of integration was taken to be the same as that of the 830-keV resonance sum peak, shifting only by the center-of-mass energy of the incident α -particle beam. The two vertical lines show the region of integration for the 650-keV-resonance sum peak. The background on the low-energy side of the peak is arising mainly from neutrons, generated by (α, n) reactions induced by α particle decay from the uranium and thorium present in the surrounding rock. The other possible source of background of high-energy γ rays is from the ¹¹B(α , γ) ¹⁵N reaction [24] which has a Q value of $Q = 10\,991.18$ keV. ¹¹B is usually present in the Ta backing as a contaminant. A narrow resonance in the ${}^{11}B(\alpha, \gamma){}^{15}N$ reaction [24] is known at an energy of 606 keV, which corresponds to an excitation energy of 11.44 MeV.

The resonance strength of the $E_{\alpha}^{\text{lab}} = 650\text{-keV}$ resonance in $^{22}\text{Ne}(\alpha, \gamma)^{26}\text{Mg}$ was determined relative to the $E_{\alpha}^{\text{lab}} =$ 830-keV resonance using Eq. (5). The efficiency of the sum peak for the $E_{\alpha}^{\text{lab}} = 650\text{-keV}$ resonance was first calculated taking the same decay scheme as the $E_{\alpha}^{\text{lab}} = 830\text{-keV}$ resonance, because the decay scheme is unknown for this resonance. Using this relative approach, we obtained an upper limit of $\omega\gamma < 0.11 \ \mu\text{eV}$ for the $E_{\alpha}^{\text{lab}} = 650\text{-keV}$ resonance. However, to calculate the influence of the choice of the involved γ cascades on the upper limit, we simulated various cascades with different γ -ray multiplicities and γ energies. These resulted in an upper limit of 0.15 μeV independent of the actual branching ratio of this resonance.

During the course of the experiment, we repeatedly measured the $E_P^{\text{lab}} = 851$ -keV resonance in ${}^{22}\text{Ne}(p, \gamma) {}^{23}\text{Na}$. This resonance was used to monitor the target stability during the experiment, primarily because it is one of the strongest low-energy resonances in ${}^{22}Ne(p, \gamma){}^{23}Na$ [25]. However, the strength of the resonance is poorly known ($\pm 36\%$). For this reason, we determined its resonance strength relative to the known $E_P^{\text{lab}} = 479$ -keV resonance ($\omega \gamma = 0.594 \pm$ 0.038 eV) [8] using Eq. (5). The level scheme and branching ratio to calculate the efficiency of the 851-keV-resonance sum peak were taken from Ref. [13]. We obtained a resonance strength of $\omega \gamma = 9.2 \pm 0.7$ eV for this resonance. This value is in agreement with the previously reported value of $\omega \gamma =$ 7.0 ± 2.5 eV [25] but with significantly lower uncertainty. With this improved accuracy this strong low-energy resonance can now be used to trace Ne implanted in various materials with high sensitivity without the need for proton energies past 1 MeV.

IV. DISCUSSION

The present resonance strength for the 830-keV resonance is compared with the results of previous measurements in Table I. Wolke *et al.* [5] used one passively shielded Ge detector, whereas Jaeger [26] used one passively and actively shielded Ge detector. Both of these measurements used a windowless gas cell target with enriched ²²Ne gas. Hunt *et al.* [6] also used one passively and actively shielded detector and the

TABLE I. Comparison of the previous literature resonance strength values with the present work for the $E_{\alpha}^{\text{lab}} = 830\text{-keV}$ resonance.

Work	ωγ (μeV)
Wolke <i>et al.</i> [5]	36 ± 4
Jaeger (Thesis) [26]	33 ± 4
Hunt <i>et al</i> . [6]	46 ± 12
This work	35 ± 4
Weighted average ^a	35 ± 2

^aCommon uncertainties of the individual values are negligible (for details see text).

target was produced by implantation of ²²Ne onto titanium. In contrast to the previous measurements, our present experiment utilized a γ -summing detector that was located on the 4850-ft level of the Sanford Underground Research Facility in Lead, South Dakota [15], which reduced cosmic-ray-induced background significantly. Here, the targets consisted of ²²Ne implanted onto Ta sheets. A common uncertainty of these experiments is the α stopping power obtained from SRIM [12]. The uncertainty in the α stopping power for pure ²²Ne gas is 2%, which is small compared to the total error of 10% of the two gas target experiments [5,26]. Titanium backings were used by Hunt et al. [6] that accounted for an uncertainty of 5% for the α stopping power, which is small compared to the total error of 26%. The present experiment uses Ta backings that have an uncertainty of 5% for the α stopping power. Our present measurement is relative to the ${}^{22}Ne(p, \gamma){}^{23}Na$ 479-keV resonance and we never make use of the absolute value of the stopping power. Only the ratio of α to proton stopping power is required. For these reasons common uncertainties of the individual experimental values are negligible. This leads to a weighted average of the previous and present measurements of $\omega \gamma = 35 \pm 2 \ \mu eV$.

While we confirm the previous studies of the resonance strength for the 830-keV resonance in ${}^{22}\text{Ne}(\alpha, \gamma){}^{26}\text{Mg}$, the strength of the corresponding resonance in the ${}^{22}\text{Ne}(\alpha, n){}^{25}\text{Mg}$ still carries large uncertainties. Previous direct studies of this resonance in ${}^{22}\text{Ne}(\alpha, n){}^{25}\text{Mg}$ [26–28] suggests values for its resonance strength which differ substantially (up to a factor of 4) amongst each other. These measurements also differ from the prediction of the coincidence studies of the decay of this level using the ${}^{22}\text{Ne}({}^{6}\text{Li}, d\gamma)$ and ${}^{22}\text{Ne}({}^{6}\text{Li}, dn)$ reactions [29]. Unlike the direct measurements the coincidence studies suggest a comparable strength between the (α, n) and (α, γ) decay modes. Further studies of the (α, n) reactions are needed to constrain the resonance strengths.

For the lowest-energy resonance at $E_{\alpha}^{\text{lab}} = 650$ keV, the observed yield only provides an upper limit for the resonance strength ($\omega\gamma < 0.15 \ \mu\text{eV}$). This limit is a factor of 4 lower than suggested by Ref. [4] on the basis of a $J^{\pi} = 1^{-}$ assignment obtained from the angular distribution analysis of the $^{22}\text{Ne}(^{6}\text{Li}, d)$ ^{26}Mg and $^{26}\text{Mg}(\alpha, \alpha')$ ^{26}Mg studies. Low-energy studies of the $^{22}\text{Ne}(^{6}\text{Li}, d)$ ^{26}Mg and $^{22}\text{Ne}(^{7}\text{Li}, t)$ ^{26}Mg reactions [30] on the other hand observed the $E_x = 11.3$ -MeV ($E_{\alpha}^{\text{lab}} = 830$ -keV) state but did not

populate the state at $E_x = 11.17$ MeV ($E_{\alpha}^{\text{lab}} = 650$ keV). The measurement performed by Talwar *et al.* [4] was at about twice the center-of-mass energy of Ota et al. [29], whereas Jayatissa et al. [30] did the measurement at sub-Coulomb energies. For this reason it could be possible that Talwar et al. [4] observed a high-spin state while this is unlikely for the low-energy experiments performed by Ota et al. [29] and Jayatissa et al. [30]. This might suggest that the state at $E_x = 11.17$ MeV $(E_{\alpha}^{\text{lab}} = 650 \text{ keV})$ is a high-spin state, e.g., $J^{\pi} = 4^+$, which cannot be excluded from angular distributions [4]. The single-particle width $(\Gamma_{\rm sp} = 5 \times 10^{-10} \text{ eV})$ is 3 orders of magnitude lower than that of a $J^{\pi} = 1^{-}$ state ($\Gamma_{\rm sp} = 5 \times 10^{-7} \text{ eV}$), predicted in Ref. [4]. Several states have been observed in this excitation range of ²⁶Mg, which cannot be resolved in the transfer and scattering studies discussed here, so it cannot be excluded that the α elastic scattering populates a different state than the α -transfer reactions. The α -capture data discussed here indicate that the state at $E_x = 11.17$ MeV is a high-spin state and for this reason should have a resonance strength distinctively smaller than that predicted in Ref. [4] based on a 1⁻ assignment.

If the resonance at $E_{\alpha}^{\text{lab}} = 650 \text{ keV}$ would indeed correspond to a high-spin state, 4⁺, then its contribution to the $^{22}\text{Ne}(\alpha, \gamma)^{26}\text{Mg}$ reaction rate would be reduced by 2 orders of magnitude because of the reduction of the penetrability. In this case its contribution to the total (α, γ) reaction rate would be less significant and would only play an important role in the temperature range around 0.2 GK (see, e.g., Fig. 8 of Ref. [4]). At higher temperatures the reaction rate would be mostly dominated by the resonance at $E_{\alpha}^{\text{lab}} = 830 \text{ keV}$ and towards lower temperatures by resonances corresponding to neutron bound states in ^{26}Mg . As a consequence, the depletion of ^{22}Ne during the lower-temperature burning phases between helium flashes in the AGB phase and the cooler phases of core helium burning would be reduced. This in turn would translate into an enhanced production of s-process elements that would be increased compared to the network calculation of Talwar *et al.* [4] (see, e.g., Figs. 14 and 17 of Ref. [4]).

V. SUMMARY

In summary, we measured the two low-energy resonances in ²²Ne(α, γ)²⁶Mg, which dominate the reaction rate at stellar helium burning temperatures around 0.3 GK. The measurement of the $E_{\alpha}^{\rm lab} = 830$ -keV resonance confirms the previous results with considerably smaller uncertainties. Based on our present measurement we provide a new weighted average value of the resonance strength, which is $35 \pm 2 \mu eV$. We performed the first direct measurement of the lowest-energy resonance at $E_{\alpha}^{\text{lab}} = 650$ keV. We provided an upper limit for the resonance strength which is considerably lower than suggested by Ref. [4]. For future measurements, either an extended gas target could be used in an underground environment, which would improve the yield substantially compared to using the implanted ²²Ne target or an experiment could be performed using inverse kinematics and a recoil separator. However, considering the discussion of the properties of this resonance, a direct measurement might not be able to observe this resonance if it is indeed a high-spin state. In addition, we also measured the $E_p^{\text{lab}} = 851$ -keV resonance in 22 Ne (p, γ) 23 Na and obtained a resonance strength of $\omega \gamma =$ 9.2 ± 0.7 eV, with significantly lower uncertainties compared to previous measurements.

ACKNOWLEDGMENTS

The authors thank the staff of the Sanford Underground Research Facility (SURF) for their hospitality and technical support given during the course of the experiment. This research was funded by the National Science Foundation (NSF) through Grants No. PHY-2011890 (University of Notre Dame, Nuclear Science Laboratory), No. PHY-1430152 (the Joint Institute for Nuclear Astrophysics, Center for the Evolution of the Elements), and No. PHY-1913746 (The Compact Accelerator System for Performing Astrophysical Research at the Sanford Underground Research Facility); the Sanford Underground Research Facility (SURF, Grant No. DE-SC0020216); and NSF Grant No. PHY-1614442.

- C. E. Rolfs, W. S. Rodney, and D. D. Clayton, Am. J. Phys. 57, 188 (1989).
- [2] S. Bisterzo, R. Gallino, F. Kppeler, M. Wiescher, G. Imbriani, O. Straniero, S. Cristallo, J. Grres, and R. J. deBoer, Mon. Not. R. Astron. Soc. 449, 506 (2015).
- [3] F. Kaeppeler, M. Wiescher, U. Giesen, J. Görres, I. Baraffe, M. El Eid, C. M. Raiteri, M. Busso, R. Gallino, M. Limongi, and A. Chieffi, Astrophys. J. 437, 396 (1994).
- [4] R. Talwar, T. Adachi, G. P. A. Berg, L. Bin, S. Bisterzo, M. Couder, R. J. deBoer, X. Fang, H. Fujita, Y. Fujita, J. Görres, K. Hatanaka, T. Itoh, T. Kadoya, A. Long, K. Miki, D. Patel, M. Pignatari, Y. Shimbara, A. Tamii *et al.*, Phys. Rev. C **93**, 055803 (2016).
- [5] K. Wolke, H. W. Becker, C. Rolfs, U. Schröder, H. P. Trautvetter, V. Harms, K. L. Kratz, J. W. Hammer, M. Wiescher, and A. Wöhr, Z. Phys. A 334, 491 (1989).

- [6] S. Hunt, C. Iliadis, A. Champagne, L. Downen, and A. Cooper, Phys. Rev. C 99, 045804 (2019).
- [7] E. Selin, S. Arnell, and O. Almn, Nucl. Instrum. Methods 56, 218 (1967).
- [8] R. Depalo, F. Cavanna, F. Ferraro, A. Slemer, T. Al-Abdullah, S. Akhmadaliev, M. Anders, D. Bemmerer, Z. Elekes, G. Mattei, S. Reinicke, K. Schmidt, C. Scian, and L. Wagner, Phys. Rev. C 92, 045807 (2015).
- [9] A. Antilla, J. Keinonen, M. Hautala, and I. Forsblom, Nucl. Instrum. Methods 147, 501 (1977).
- [10] B. Paine and D. Sargood, Nucl. Phys. A 331, 389 (1979).
- [11] C. Iliadis, *Nuclear Physics of Stars* (Wiley-VCH Verlag, Weinheim, Germany, 2007).
- [12] J. F. Ziegler, M. Ziegler, and J. Biersack, Nucl. Instrum. Methods Phys. Res., Sect. B 268, 1818 (2010).
- [13] E. Bakkum and C. Van Der Leun, Nucl. Phys. A 500, 1 (1989).

- [14] M. Viitasalo, M. Piiparinen, and A. Anttila, Z. Phys. A 250, 387 (1972).
- [15] D. Robertson, M. Couder, U. Greife, F. Strieder, and M. Wiescher, Eur. Phys. J. Web Conf. 109, 09002 (2016).
- [16] Compact accelerator system for performing astrophysical research, https://caspar.nd.edu/, accessed 2022-04-01.
- [17] R. B. Vogelaar, T. R. Wang, S. E. Kellogg, and R. W. Kavanagh, Phys. Rev. C 42, 753 (1990).
- [18] P. M. Endt and C. van dèr Leun, Nucl. Phys. A **214**, 1 (1973).
- [19] C. S. Reingold, O. Olivas-Gomez, A. Simon, J. Arroyo, M. Chamberlain, J. Wurzer, A. Spyrou, F. Naqvi, A. C. Dombos, A. Palmisano, T. Anderson, A. M. Clark, B. Frentz, M. R. Hall, S. L. Henderson, S. Moylan, D. Robertson, M. Skulski, E. Stech, S. Y. Strauss *et al.*, Eur. Phys. J. A 55, 77 (2019).
- [20] O. Olivas-Gomez, A. Simon, D. Robertson, A. C. Dombos, F. Strieder, T. Kadlecek, M. Hanhardt, R. Kelmar, M. Couder, J. Grres, E. Stech, and M. Wiescher, Eur. Phys. J. A 58, 57 (2022).
- [21] A. C. Dombos, D. Robertson, A. Simon, T. Kadlecek, M. Hanhardt, J. Görres, M. Couder, R. Kelmar, O. Olivas-Gomez, E. Stech, F. Strieder, and M. Wiescher, Phys. Rev. Lett. 128, 162701 (2022).
- [22] S. Agostinelli, J. Allison, K. Amako, J. Apostolakis, H. Araujo, P. Arce, M. Asai, D. Axen, S. Banerjee, G. Barrand, F. Behner, L. Bellagamba, J. Boudreau, L. Broglia, A. Brunengo, H. Burkhardt, S. Chauvie, J. Chuma, R. Chytracek, G.

Cooperman *et al.*, Nucl. Instrum. Methods Phys. Res., Sect. A **506**, 250 (2003).

- [23] R. Longland, C. Iliadis, J. M. Cesaratto, A. E. Champagne, S. Daigle, J. R. Newton, and R. Fitzgerald, Phys. Rev. C 81, 055804 (2010).
- [24] T. R. Wang, R. B. Vogelaar, and R. W. Kavanagh, Phys. Rev. C 43, 883 (1991).
- [25] P. M. Endt, Nucl. Phys. A 521, 1 (1990).
- [26] M. Jaeger, Thesis, University of Stuttgart, 2001, doi: 10.18419/opus-4673.
- [27] V. Harms, K.-L. Kratz, and M. Wiescher, Phys. Rev. C 43, 2849 (1991).
- [28] U. Giesen, C. P. Browne, J. Grres, S. Graff, C. Iliadis, H. P. Trautvetter, M. Wiescher, W. Harms, K. L. Kratz, B. Pfeiffer, R. E. Azuma, M. Buckby, and J. D. King, Nucl. Phys. A 561, 95 (1993).
- [29] S. Ota, G. Christian, G. Lotay, W. N. Catford, E. A. Bennett, S. Dede, D. T. Doherty, S. Hallam, J. Hooker, C. Hunt, H. Jayatissa, A. Matta, M. Moukaddam, G. V. Rogachev, A. Saastamoinen, J. A. Tostevin, S. Upadhyayula, and R. Wilkinson, Phys. Lett. B 802, 135256 (2020).
- [30] H. Jayatissa, G. Rogachev, V. Goldberg, E. Koshchiy, G. Christian, J. Hooker, S. Ota, B. Roeder, A. Saastamoinen, O. Trippella, S. Upadhyayula, and E. Uberseder, Phys. Lett. B 802, 135267 (2020).