New direct study of the ${}^{19}\text{Ne}(p,\gamma){}^{20}\text{Na}$ reaction cross section

M. Couder, C. Angulo, E. Casarejos, P. Demaret, P. Leleux, and F. Vanderbist

Institut de Physique Nucléaire et Centre de Recherches du Cyclotron, Université catholique de Louvain, Louvain-la-Neuve, Belgium (Received 19 November 2003; published 23 February 2004)

The ¹⁹Ne $(p, \gamma)^{20}$ Na reaction is part of the chain of reactions linking the hot-CNO cycles and the rp-process in hydrodynamic hydrogen and helium burning. The resonance strength of the 448 keV level has been measured in the recoil separator ARES. An upper limit of 15 meV (90% C.L.) has been obtained. This result slightly restricts a prior determination using different methods.

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In hydrodynamic H- or He-burning occurring in novae and X-ray bursts, hot-CNO cycles can be escaped to the rp-process through several chains of reactions starting from ${}^{18}\text{F}.{}^{18}\text{F}(p,\alpha){}^{15}\text{O}(\alpha,\gamma){}^{19}\text{Ne}(p,\gamma){}^{20}\text{Na}$ or ${}^{18}\text{F}(p,\gamma){}^{19}\text{Ne}(p,\gamma){}^{20}\text{Na}$ or ${}^{18}\text{F}(p,\gamma){}^{19}\text{Ne}(p,\gamma){}^{20}\text{Na}$ or ${}^{18}\text{F}(p,\gamma){}^{20}\text{Na}$ or ${}^{18}\text{F}(p,\gamma){}^{20}\text{Na}$ for ${}^{18}\text{F}(p,\gamma){}^{20}\text{Na}$ are action is of great interest. In the temperature domain of relevance in the above environments, the reaction rate is dominated by a resonant level at 448 keV above threshold, at an excitation energy of 2.643 MeV in ${}^{20}\text{Na}$.

This level was produced previously in nuclear reactions induced by stable beams, like ${}^{20}Ne({}^{3}He,t){}^{20}Na$ [3–6] or 20 Ne $(p,n){}^{20}$ Na [7]. Angular distributions of the tritons suggested a $J^{\pi}=1^+$ assignment, while the (p,n) reaction data was consistent with a 3⁺ assignment. This level was not observed in the ²⁰Mg delayed β -decay [8–10]; as Gamow-Teller β -decay from a 0⁺ state strongly favors 1⁺ final states, a 1⁺ assignment would require the 2.643 MeV level to be an intruder state. Resonance strengths of 6 meV [4] and 7 meV [5] were calculated. In a microscopic three-cluster model calculation [11], a $J^{\pi}=1^{-}$ was attributed to this level, and a resonance strength of 33 meV was calculated. Finally, Brown et al. [12] suggested the 2.643 MeV level to be the analog of the 2.966 MeV 3⁺ state in ²⁰F; from the shellmodel-calculated lifetime (τ) of 3.5 fs for the mirror level, they obtained a resonance strength of 80 meV for the 2.643 MeV state. Adopting the most recent experimental result, i.e., $\tau \leq 12$ fs [13], a lower limit of 16 meV was deduced for the resonance strength [14].

Several investigations of the ¹⁹Ne $(p, \gamma)^{20}$ Na reaction were performed in Louvain-la-Neuve in the last decade. An upper limit of 21 meV (90% C.L.) was set on the resonance strength of the 2.643 MeV level, when combining the data of two experiments detecting α decays, following β^+ feeding of high levels in ²⁰Ne [15]: the detectors used were doublesided silicon strip detectors, and solid-state nuclear track detectors, respectively. Later on, ¹⁹Ne beams of higher energies induced the (p, γ) reaction to higher levels above threshold; positrons were detected in a stack of scintillators at the end of a magnetic spectrometer, and resonance strengths were also deduced [16]. All results were summarized and correctly renormalized, and astrophysical consequences were drawn, in a subsequent publication [17].

The above measurements had two drawbacks, i.e., a low detection efficiency, of the order of 1.5%, and the presence

of a background from the ¹⁹Ne $(d, n)^{20}$ Na competing reaction, induced by the beams on the deuterium present in the CH₂ target. The latter imposed an additional measurement on a CD₂ target to be made after each (p, γ) measurement.

ARES, the Astrophysics REcoil Separator, was built to measure radiative capture reactions of astrophysical interest, in inverse kinematics. ARES is coupled to CYCLONE44 [18], a cyclotron which is used as a postaccelerator to bring radioactive ions to the required energy. The aim in ARES is to separate product ions from beam ions and to detect the former. This separation proceeds in three steps (Fig. 1): (i) the most abundant charge state of the product ions (and of the beam ions at the same time) is selected in a dipole magnet; (ii) a Wien filter or velocity filter transmits the product ions and deflects the beam ions; (iii) a $\Delta E - E$ identification is performed in a ΔE gas ionization chamber and a E silicon detector of the PIPS type. Extensive tests had been realized for the ${}^{19}F(p, \gamma){}^{20}Ne$ "mirror" reaction [19]: a resonant state at 635 keV center of mass (c.m.) energy, of 6.2 keV total width and of 1.6 eV resonance strength, had been selected. The main results of these tests are the following: a transmission of 11.5% was measured for the ²⁰Ne⁷⁺ product ions (the most abundant charge state); the charge distribution of ¹⁹F and of ²⁰Ne ions behind the CH₂ target was measured and was found in good agreement with the Shima et al. [20] calculation; the measured energy distribution of the product ions was fairly well reproduced by a GEANT simulation containing specific energy losses of ¹⁹F and ²⁰Ne ions in CH₂ (the target material) and Isobutane (the gas of the ΔE counter), which had been previously measured.

In summary a measurement of the ¹⁹Ne(p, γ)²⁰Na reaction in ARES should benefit from two improvements as compared with previous methods: a slightly larger transmission efficiency (~4%, from the product of the percentage of the ²⁰Na⁷⁺ beyond the target and the transmission of ²⁰Na⁷⁺ in ARES) and a suppression of the background from the (d,n) reaction. Simulations of the ¹⁹Ne(d,n) reaction in ARES showed indeed a null transmission of ²⁰Na⁷⁺ ions from this reaction.

In the ¹⁹Ne(p, γ)²⁰Na reaction measurement, a 9.8 MeV ¹⁹Ne³⁺ radioactive beam accelerated by CYCLONE44, was focused on a 80 μ g/cm² CH₂ target. Two PIPS monitors located at forward angle (±15°) recorded charged particles scattered by or recoiling from the target. Their time-of-flight with respect to the cyclotron RF and their energy were dis-



FIG. 1. Layout view of ARES coupled to CY-CLONE44.

played in a two-dimensional spectrum permitting particle identification [Fig. 2(a)]. The recoil proton spectra are of particular interest [Fig. 2(b)]. As the resonance width is smaller than the lower limit (~ 0.5 keV, see Ref. [21]) able to distort in a significant way the classical Coulomb pattern, proton spectra were fitted with a Rutherford cross section. The fit provided with the product of the Hydrogen content in the target and the integrated beam intensity. Working with a low intensity radioactive beam, no hydrogen loss is expected and a 2-to-1 stoecheometry was assumed throughout the experiment (different targets were used), leading to an average ¹⁹Ne³⁺ beam on target equal to $(1.08\pm0.11)10^8$ pps. Another specific part of the proton spectra, i.e., the upper edge, allowed to deduce the incident beam energy on target (let us recall that high-energy protons are produced by the beam in the front face of the target). Regarding the radioactive beam itself, two other characteristics were measured as follows.

(i) *The beam purity*. It is well known that radioactive beams produced by the ISOL-method can be contaminated by a stable isobar. The ¹⁹Ne beam with a reduced intensity was transported through ARES and identified in the ΔE -*E* telescope. The contamination in ¹⁹F was at most 0.7%.

(ii) *The beam energy distribution on target*. This information is an important input to the simulation code, which will eventually furnish the transmission efficiency of the ²⁰Na product ions. A full width at half maximum (FWHM) of the ¹⁹Ne beam equal to 260 keV was measured in a PIPS detector at the target place. This number is much larger than the FWHM of a stable ¹⁹F beam at the same energy (170 keV), indicating that the tuning of CYCLONE44 to maximize both the ¹⁹Ne-¹⁹F separation and the ¹⁹Ne acceleration efficiency leads to a degradation of the beam quality.

The beam energy had been adjusted so that 20 Na ions from the resonant state were produced at about 60 μ g/cm² from the front face of the target; the thickness that they had to cross before leaving the target was sufficient to reach the equilibrium of the charge states.

In order to transmit properly ²⁰Na ions through ARES, their energy behind the target has to be known. Discrepancies among stopping power tables lead us to measure the stopping power for ²⁰Ne and ²³Na in CH₂, and also in isobutane, as we did before for ¹⁹F and ²⁰Ne in the test reaction.

Coming now to the ¹⁹Ne(p, γ) measurement, ARES was

tuned as follows: the ¹⁹Ne⁷⁺ beam was transported through ARES, all the elements being optimized successively. Typical transmission of 60% was obtained. Then, knowing the energy of ²⁰Na ions behind the target, the dipole field and the Wien filter magnetic field were changed accordingly to transport ²⁰Na ions to the ΔE -E detector. About 500 ¹⁹Ne⁷⁺ were transmitted per second as well, the rejection factor of the ¹⁹Ne beam being typically equal to 5×10^{-6} . This is roughly a factor of 10 worse than the one obtained in the ¹⁹F(p, γ)²⁰Ne test reaction. The lower quality of the radioactive beam is probably the reason for that.

The counting rate of the ΔE -E coincidence was limited to a few hundred Hz, in order to limit pile-up of ¹⁹Ne events extending into the region of the ΔE vs $E + \Delta E$ plane where ²⁰Na events were expected. The ΔE gas counter was operated at a pressure of 4 mb isobutane. Figure 3 presents a two-dimensional spectrum of the ΔE vs $E + \Delta E$ events. The dark area is the region of the 19Ne ions. A solid line surrounds the region of the ²⁰Na ions, which was determined from a simulation of ²⁰Na events tracked through ARES from the target to the ΔE -E detector. The measured stopping power of ²³Na ions in CH₂ and in isobutane was part of the simulation. Two regions of equal surface were selected right and left to the region of interest (ROI). Pile-up events, occurring in these regions at a level of 10^{-7} of the leaky beam events, were summed and the mean of the counts in both regions was subtracted from the counts in the ROI.

The remaining counts were corrected for the efficiency of the setup, which is the product of two factors, the amount of $^{20}Na^{7+}$ ions after the target, and the transmission of $^{20}Na^{7+}$ in ARES. The first factor (0.37) was obtained from Shima's tables [20], which had been checked for ^{19}F and ^{20}Ne ions, and had been found in agreement with our measurements, at least for the most abundant state, i.e., 7^+ [19]. The second factor was deduced from a simulation of $^{20}Na^{7+}$ ions tracked from the target to the ΔE -E telescope. The global efficiency was 2.7%. It is justified to suppose that the observed yield is from the 2.643 level in view of the very small contribution expected from direct capture [16]. The resonance strength $\omega\gamma$ is then given by

$$\omega \gamma = \frac{Y}{I} \frac{2}{\lambda^2} \epsilon_{lab} \frac{m_t}{m_t + m_p},$$



FIG. 2. (a) Two-dimensional spectrum, time-of-flight vs energy, for particles recorded in a silicon detector at +15°, resulting from interactions of ¹⁹Ne beam ions and a CH₂ target. Regions corresponding to scattered ¹⁹Ne and recoil ¹²C are clearly visible in the high-energy region. The proton region is encircled. A parasitic ¹⁹Ne beam of low energy and low intensity ($\leq 1\%$ of the main beam) accelerated by the cyclotron appears to the right of the proton region. The low-energy pulses showing no correlation with the cyclotron RF are positron events. (b) The projection of the proton region to the energy axis (solid histogram) is shown. The dotted histogram is a fit.

where *Y* is the number of ²⁰Na events corrected for the efficiency, *I* is the integrated ¹⁹Ne beam intensity, λ is the c.m. wave length (in cm²), ϵ_{lab} is the stopping power of ¹⁹Ne ions in CH₂ (in 10⁻¹⁵ eV cm²/at), $m_t(m_p)$ is the mass of the target (of the projectile), in a.m.u.

The final result is $\omega \gamma = -1.0 \pm 9.6$ meV. To the 9.6 meV statistical error a systematic uncertainty should be added. The latter comprises the uncertainty to the integrated beam and to the transmission of ARES: the first one was estimated above, while the second one was obtained by changing in the simulation code the slits positions by 0.5 mm (dipole exit and Wien filter exit) and taking extreme values of the efficiency. A systematic uncertainty of 0.6 meV was deduced.



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FIG. 3. Two-dimensional spectrum, ΔE vs $E+\Delta E$, in the end detectors of ARES for the ¹⁹Ne $(p, \gamma)^{20}$ Na reaction. The count scale is logarithmic in intensity. The dark area consists in ¹⁹Ne leaky beam events. The leaky ¹⁹F contaminant beam events are visible just below the ¹⁹Ne region. The framed area limited by solid lines is the expected location for ²⁰Na events from the 2.643 MeV level. Adjacent regions limited by dashed lines are used to estimate the background. See text for more details. This figure is a typical run. During the experiment, changes of target, of ARES settings and of ΔE gas pressure resulted in different locations for ²⁰Na events.

Both types of uncertainties were combined in quadrature, the positive part of the Gaussian distribution of the result was renormalized to 1, and an upper limit of 15.2 meV (90% C.L.) was obtained. This value reduces slightly the upper limit from our first experiment [17], it is in agreement with the first prediction of 6 meV or 7 meV for a 1⁺ state [4,5], and does not contradict (at a 2σ level) the most recent calculated lower limit of 16 meV for a 3⁺ state [14].

In summary, a new measurement of the resonance strength of the 2.643 MeV level was performed with a recoil mass separator. This measurement restricts a previous one obtained with quite different methods. Significant improvements of the ARES transmission and efficiency are required to yield more than an upper limit to the 2.643 MeV level. Some of them are presently examined.

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