Identification of a new low-lying state in the proton drip line nucleus 19Na

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The second excited state ($J^{\pi} = 1/2^+$, $\ell = 0$) of the proton-rich nucleus ¹⁹Na has been observed for the first time using the elastic scattering reaction ${}^{1}H({}^{18}Ne,{}^{1}H){}^{18}Ne$. An intense (4×10⁶ pps) ¹⁸Ne radioactive beam was produced at the CRC-RIB facility at Louvain-la-Neuve to bombard a polyethylene target. The recoil protons were detected at 20 different angles in the LEDA segmented silicon detector. The resulting elastic scattering cross sections have been analyzed using a global *R*-matrix fit. We find that the second excited state in ¹⁹Na lies at the center-of-mass energy $E_{cm} = 1.066 \pm 0.003$ MeV with respect to the ¹⁸Ne+*p* threshold, and has a proton width $\Gamma_p = 101 \pm 3$ keV. The Coulomb shift between the $1/2^+$ mirror levels in ¹⁹O and ¹⁹Na is 0.73 MeV, among the largest values currently observed in light exotic nuclei.

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I. INTRODUCTION

The structure of nuclei near the drip lines is one of the major current interests in nuclear physics. Proton-rich light nuclei are a remarkable case since the level scheme is not known for many species. For example, almost no spectroscopic information is available on 19 Na, which is proton unbound by 321 ± 13 keV [1]. Even though it is not far from stability, very little is known experimentally about its structure and level scheme: there are no spin assignments, and only the location of the ground state and of the first excited state have been determined by nuclear reactions involving stable beams [2,3]. The 19 Na nucleus has also some interest for nuclear astrophysics. The ¹⁸Ne(2p, γ)²⁰Mg reaction may play a role on long-lived waiting point isotopes near the proton drip lines for the *rp*-process reaction flow, although very high densities would be required $[4]$. However, only the *Q* value of the ¹⁸Ne(p , p)¹⁸Ne reaction is of relevance as the width of its resonances does not enter the reaction rate because production and decay of 19 Na are in an equilibrium $|5|$.

The intense low-energy 18 Ne radioactive beam at the Louvain-la-Neuve CRC-RIB facility has made possible the study of the low-lying states of 19 Na. The aim of the present experiment was to measure the location and the width of the second excited state of 19 Na, which according to the mirror nucleus ¹⁹O ($E_x = 1.4717 \pm 0.0004$ MeV) should have a spin

 $J^{\pi}=1/2^+$ [1]. The parameters of the ground and the first excited states, whose predicted widths are of the order of eV $[1]$, could not be obtained by the present technique and their measurement was therefore beyond the scope of this work.

The experimental method is described in Sec. II. The data analysis is presented in Sec. III, followed by the conclusions in Sec. IV.

II. EXPERIMENTAL METHOD

We have used the inverse elastic scattering technique by bombarding a proton-rich target with a 18 Ne beam at different laboratory energies. The elastic scattering technique has been widely applied in the past for nuclear spectroscopy studies $[6-9]$. It makes use of the sensitivity of the protons to the presence of a resonant state in the compound nucleus. When the resonant state is scanned with the appropriate ion beam and target, the recoil proton spectra show spectacular changes when detected at forward laboratory angles. These spectra contain precise information on the resonance energy, angular momentum, and width provided all experimental effects, mainly the energy resolution and angular resolution of the detectors, are properly taken into account $[10]$.

Post-accelerated radioactive beams of $^{18}Ne^{2+}$ at E_{lab} = 21 and 23.5 MeV, and ¹⁸Ne³⁺ at E_{lab} = 28 MeV, respectively, were provided by the CYCLONE110 cyclotron. The ¹⁸Ne atoms were produced through the ¹⁹F($p, 2n$)¹⁸Ne reaction by bombarding a LiF target with an intense 30-MeV proton beam produced by the CYCLONE30 cyclotron and were ionized to the 2^+ or to the 3^+ state in a ECR source, before being post-accelerated. A detailed description of the production of the 18 Ne radioactive beam can be found in

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Ref. $[11]$. The average intensity of the ¹⁸Ne beams on target was of the order of 4×10^6 pps, which was maintained for periods of several days, allowing us to obtain excellent statistics for each of the three beam energies (the statistical error of the proton events per energy bin of 25 keV at each angle ranges between 1% and 8% for all energies). The target consisted of a 0.5-mg/cm² polyethylene $(CH_2)_n$ foil with a very thin Au coating evaporated on the upstream face of the target. We obtained a target thickness of $(520 \pm 10) \mu g/cm^2$ by measuring the energy loss of the 28-MeV ¹⁸Ne beam [12]. The target thickness and the three different beam energies have been chosen such that the energy range of interest could be scanned and the expected interference pattern could clearly show up. A thicker $(CH_2)_n$ target could be used to scan the energy range of interest using a single beam energy. However, the energy straggling of the beam and of the recoil protons would have been larger, introducing an additional uncertainty in the data. The thickness of the Au layer, used for normalization purposes (see below), was obtained by bombarding the targets with a 5.6-MeV 7 Li beam. The scattered ⁷Li on Au and the recoil protons from the ${}^{1}H(^{7}Li,{}^{1}H)$ reaction were detected simultaneously in a PIPS detector located at 60° with respect to the beam axis. The number of Au atoms in the target N_{Au} can be calculated from the relation

$$
N_{\rm Au} = N_{\rm p} \frac{C_{\rm p}}{C_{\rm Li}} \frac{(d\sigma/d\Omega)_{\rm Li+p}}{(d\sigma/d\Omega)_{\rm Li+Au}},\tag{1}
$$

where N_p is the number of H atoms in the target, C_p and C_{Li} are the number of detected recoil protons and scattered Li, respectively, and $(d\sigma/d\Omega)_{\text{Li+p}}$, $(d\sigma/d\Omega)_{\text{Li+Au}}$ are the differential cross sections for ${}^{1}H({}^{7}Li,{}^{1}H){}^{7}Li$ and 197 Au(7 Li, 7 Li)¹⁹⁷Au, respectively. The $1H(TLi,1H)$ ⁷Li cross section value was obtained from literature $[13]$. We have assumed a pure Rutherford cross section for $197Au(^{7}Li,^{7}Li)$ ¹⁹⁷Au. We derived a Au layer thickness of $8.0 \pm 0.5 \ \mu g/cm^2$.

The recoil protons were detected in two LEDA segmented silicon detector arrays $[14]$ situated at about 62 and 12 cm from the target. Due to the kinematical variation of the recoil proton energy with laboratory angle, only the 20 most forward angles were used at $\theta_{\text{lab}}=4.9^{\circ}-11.7^{\circ}$ ($\theta_{\text{c.m}}$) $= 170.2^{\circ} - 156.6^{\circ}$ and $\theta_{\text{lab}} = 22.6^{\circ} - 29.9^{\circ}$ ($\theta_{\text{c.m}}$ $= 131.0^{\circ} - 120.2^{\circ}$, with angular resolution $\Delta \theta_{\rm lab} \approx 0.2^{\circ}$ and 0.9°, respectively. An 18- μ m-thick Mylar foil was situated in front of all LEDA sectors, except one, to stop the high energy scattered 18 Ne and most of the recoiled 12 C particles, and therefore to reduce the data acquisition dead time. The ''uncovered'' LEDA sector was used for monitoring the number of incoming beam particles by detecting the 18 Ne ions scattered on the thin Au layer, assuming a purely Rutherford elastic cross section for $^{197}Au(^{18}Ne,^{18}Ne)^{197}Au$ at the measured energies. The advantage of this method is that one obtains an intrinsic normalization simultaneously to the recoil proton measurement.

For each LEDA strip, not only the energy but also the time-of-flight (TOF) of the particle with respect to the cyclotron rf were recorded and displayed in a two-dimensional spectrum. Figure $1(a)$ shows a typical two dimensional spec-

FIG. 1. (a) Summed TOF spectra at $\theta_{lab} = 4.9^{\circ}$ obtained with a 23.5-MeV 18 Ne beam on a "nominal" 0.5-mg/cm² polyethylene target. The recoil protons are marked by a selection on the figure. (b) The proton signal is shown as a function of the proton laboratory energy.

trum obtained with a 23.5-MeV 18 Ne beam on a 0.5-mg/cm² polyethylene target during 2 days of running time. The spectrum was obtained by adding the calibrated singles spectra from several LEDA strips at the smallest measured laboratory angle $\theta_{lab} = 4.9^\circ$. This spectrum corresponds also to the angle with lowest accumulated statistics (smallest solid angle). The recoil protons, marked by a selection on the figure, are well separated from the recoiled ^{12}C , not completely stopped in the mylar foil, and the uncorrelated β background. In Fig. $1(b)$, only the proton signal is shown as a function of the proton laboratory energy. The interference pattern between a *s*-wave resonance and the Rutherford contribution is clearly present. Similar spectra were obtained at all measured angles and at beam energies of 21.0, 23.5, and 28.0 MeV, covering laboratory proton energies from 2.1 to 5.5 MeV. Data obtained at overlapping energy regions are in excellent agreement.

III. DATA ANALYSIS AND DISCUSSION

In reverse kinematics, the c.m. energy $E_{\text{c.m.}}$ is related to the laboratory energy of the protons E_p , recoiling at the laboratory angle θ_{lab} by

$$
E_{\rm c.m.} = E_{\rm p} \times \frac{M_{\rm b} + 1}{4M_{\rm b}\cos^2\theta_{\rm lab}},
$$
 (2)

where M_b is the mass number of the beam. The tranformation of the scattering angle and the differential cross section to the c.m. frame is given by

and

$$
\theta_{\rm c.m.} = \pi - 2 \,\theta_{\rm lab} \tag{3}
$$

$$
\frac{d\sigma}{d\Omega_{\rm c.m.}}(E_{\rm c.m.}, \theta_{\rm c.m.}) = \frac{d\sigma}{d\Omega_{\rm lab}}(E_{\rm lab}, \theta_{\rm lab})/4 \cos \theta_{\rm lab}. \tag{4}
$$

The proton spectra are slightly degraded by two experimental effects: (i) the opening angle of the detectors ($\Delta \theta_{lab}$), introducing an uncertainty in the proton energy ΔE_{θ} , given by $\lfloor 7 \rfloor$

$$
\Delta E_{\theta} = 2E_{\text{p}} \tan \theta_{\text{lab}} \Delta \theta_{\text{lab}}, \qquad (5)
$$

and (ii) the energy straggling of the protons ΔE_{stras} in the target and in the mylar foil used to stop the heavier ions. A good estimate of ΔE_{strag} is obtained by using the Bohr formula [15]. The total energy broadening (ΔE) is obtained by adding quadratically ΔE_{θ} and ΔE_{strag} . Typical values of ΔE are 12 keV for a laboratory angle of $\theta_{\rm lab} = 4.9^{\circ}$, 17 keV for $\theta_{\rm lab} = 11.7$ ° ($\Delta E_{\rm drag}$ is the most important contribution in both cases), and 40 keV for $\theta_{\rm lab} = 29.9^{\circ}$ (ΔE_{θ} is the most important contribution).

From the recoil proton spectra we have obtained differential cross sections, for each of the 20 laboratory angles and for c.m. energies in the range $E_{\text{c.m.}}=0.7-1.5$ MeV, by correcting the number of counts for the solid angle of the detectors, the number of protons in the target, and the total number of incident beam particles (obtained by the normalization on Au, as explained above). Corrections for energy loss in the target and in the mylar foil have also been applied $[12]$. Figure 2 shows the differential cross section versus c.m. energies for three typical c.m. angles (a) 170.2° , (b) 156.6° , and (c) 120.2°. The total error includes the statistical error and the uncertainties due to the normalization and the energy calibration. The dotted curve is the Rutherford cross section.

We have used the *formalism* $[16]$ *to fit the dif*ferential cross sections in the $120.2^{\circ} - 170.2^{\circ}$ angular range and the 0.7–1.5-MeV c.m. energy range. In the *R*-matrix theory, the nuclear phase shift is defined by

$$
\delta^{\ell} = \delta_{\rm HS}^{\ell} + \delta_{\rm R}^{\ell},\tag{6}
$$

where $\delta_{\text{HS}}^{\ell}$ and δ_{R}^{ℓ} are the hard-sphere and the *R*-matrix phase shifts, respectively. Here, we have used hard-sphere phase shifts for all partial waves (ℓ values up to $\ell_{\text{max}}=2$ have been taken into account) except for $\ell=0$, where we have applied the *R*-matrix phase shift given by

$$
\delta_{\mathbf{R}}^0 = \arctan \frac{P_0 R^0}{1 - S_0 R^0},\tag{7}
$$

where P_0 and S_0 are the $\ell=0$ penetration and shift factors, respectively, and R^0 is the *R*-matrix defined with a single pole. The fitted parameters are the pole parameters converted to the resonance energy E_R and the proton width Γ_p of the $1/2^+$ state in ¹⁹Na (in *R*-matrix notation, E_R and Γ_p are "observed" parameters). In order to account for the experimental effects described above, we have convoluted the calculations

FIG. 2. Elastic cross section versus c.m. energies for the system ¹⁸Ne + p for c.m. angles (a) 170.2° , (b) 156.6° , and (c) 120.2° . The thicker solid curves are *R*-matrix fits ($a=5$ fm) with parameters of Table I, that includes an energy correction for opening angle and beam straggling. The thinner solid curve are the fits obtained without ΔE folding. The dotted curves are the Rutherford cross sections.

with a Gaussian distribution, with a full width at half maximum ΔE , which was set equal to the total energy broadening as calculated above.

We have tested the sensitivity of the fit to the *R*-matrix channel radius *a* by performing global fits for three different values $a=4,5$, and 6 fm. The results of these fits are presented in Table I. The quoted errors include an uncertainty of 3 keV due to the data bining. We have adopted the values for $a=5$ fm, which provides the lowest χ^2/N value, χ^2/N = 0.44 for E_R = 1.066 ± 0.003 MeV and Γ_p = 101 ± 3 keV. The small errors are due to the strong sensitivity of χ^2 with respect to E_R and Γ_p . The thicker solid curve in Fig. 2 is the *R*-matrix fit ($a=5$ fm) for the adopted values (Table I) and includes the correction for opening angle and beam strag-

TABLE I. Results of the *R*-matrix fits $(N=367)$.

	$a=4$ fm	$a=5$ fm	$a=6$ fm
E_R (MeV)	1.067 ± 0.003	1.066 ± 0.003	1.064 ± 0.003
	104 ± 3	101 ± 3	95 ± 3
Γ_p (keV) χ^2/N	0.53	0.44	0.49
$\theta_{\rm p}^2$ (%)	29.8	22.9	21.6

gling discussed above. The thinner solid curve [unresolved] for (a)] corresponds to $\Delta E = 0$. *R*-matrix fits using other spin assignments strongly disagree with the experimental data. The dimensionless reduced width θ_p^2 represents more than 20% of the Wigner limit. This value is similar to or even larger than the reduced width of $\ell=0$ states in other nuclei such as ¹³N ($E_x = 2.37$ MeV, $\theta_p^2 = 21.8\%$) or ¹⁴O (E_x $= 5.17$ MeV, $\theta_p^2 = 21.4\%$).

The isobar diagram for the mirror nuclei 19 O and 19 Na is shown in Fig. 3, including the new $1/2^+$ state in ¹⁹Na. The

FIG. 3. Isobar diagram for 19 O and 19 Na.

large θ_p^2 value is also consistent with the important Coulomb shift (shown by an arrow in the figure). In ^{19}O , the excitation energy of the mirror state is $E_x = 1.472$ MeV [1] whereas it is $E_r = 0.745 \pm 0.013$ MeV for ¹⁹Na. This Coulomb shift (-0.73 MeV) is among the largest values observed in light nuclei. It is typical of deformed states, found in nuclei near the drip lines.

IV. CONCLUSIONS

Using a low-energy ¹⁸Ne radioactive beam and a thin polyethylene target we have measured the elastic scattering reaction 1 H(18 Ne, 1 H) 18 Ne. From a global *R*-matrix fit for c.m. energies $E_{\text{c.m.}}$ = 0.7–1.5 MeV and c.m. angles $\theta_{\text{c.m.}}$ $=120.2^{\circ} - 170.2^{\circ}$, we have determined the energy and the width of the $1/2^+$ state in ¹⁹Na, $E_R = 1.066 \pm 0.003$ MeV and Γ_p =101±3 keV. This state corresponds to the mirror of the $E_x = 1.472$ MeV state in ¹⁹O. We find a very large reduced width, typical of deformed states. Consequently the Coulomb shift of this state with respect to the mirror one in ^{19}O is among the largest values observed in light nuclei.

According to the mirror nucleus 19 O, several states should be present in the c.m. energy range 2–3 MeV. In order to obtain a global picture of the low-energy states of ¹⁹Na, further investigations are planned in the near future.

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