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1 **Resonant α scattering of ${}^6\text{He}$: limits of clustering in ${}^{10}\text{Be}$**

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Abstract

20

21 The structure of ^{10}Be was studied via resonant α -particle scattering of a neutron-rich ^6He beam.
22 A time projection chamber, PAT-TPC, was operated in an active-target mode to provide a gaseous
23 ^4He target and trace the beam and reaction products traversing its active tracking volume. This
24 significantly lowered the detection threshold of reaction products at low energies. Elastic scattering,
25 inelastic scattering to the ^6He 2^+ state, and the $^6\text{He}(\alpha, 2n)^8\text{Be}$ reaction were measured below an
26 energy of 6 MeV in the center-of-mass frame. Continuous spectra of excitation functions and
27 angular distributions were obtained from unambiguously-identified recoiling α particles for the
28 elastic and inelastic channels. While a resonance of the 4^+ state at 10.15 MeV in ^{10}Be previously
29 reported was confirmed, no other resonances were identified in the elastic channel over the measured
30 energy region. The results are in line with antisymmetric molecular dynamics calculations that
31 predict the limits of α clustering in high-spin states due to a spin-orbit force.

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33 I. INTRODUCTION

34 Clustering of α particles is a unique aspect of nuclear correlations. It is known that α
35 clustering often occurs in light nuclei along the $N = Z$ line as already suggested since 1930s
36 to explain their level schemes [1–3]. Archetypal examples are the ground state of ${}^8\text{Be}$ and
37 the second 0^+ state of ${}^{12}\text{C}$ (referred to as “Hoyle” state) [4, 5], manifesting well-developed 2α
38 and 3α clusters, respectively, and likewise further heavier systems such as 4α states in ${}^{16}\text{O}$ [6]
39 or 6α states in ${}^{24}\text{Mg}$ [7]. These cluster states, which exclusively consist of α particles, have
40 been providing a unique playground to discuss bosonic condensations [6, 8, 9] or geometries
41 of multiple quantum objects [3, 7] in femtometer-scale systems.

42 The structure of the nucleus, a quantum many-body system, can drastically change with
43 addition or removal of nucleons. How, if at all, do these nucleons affect α clustering in nuclei?
44 The structure of ${}^{11}\text{B}$, the nucleus with one proton removed from ${}^{12}\text{C}$, was recently studied
45 via the (d, d') reaction [10]. The large monopole strength extracted for the $3/2^-$ state at
46 8.56 MeV indicates a well-developed $2\alpha + t$ cluster structure, suggesting that the clustering
47 nature of the Hoyle state in ${}^{12}\text{C}$ persists in the presence of a proton hole. Theoretical studies
48 on neutron-rich beryllium isotopes that trace the cluster evolution away from ${}^8\text{Be}$ also predict
49 the persistence of 2α clusters in the neutron-excess systems ${}^{10}\text{Be}$ [11–24] and ${}^{12}\text{Be}$ [25, 26].
50 It is further suggested that the unique correlation of neutrons and 2α clusters diversifies the
51 evolution of structure. The formation of predicted structures, such as neutron-molecular
52 orbitals [15, 16, 25, 26] or a di-neutron pair around 2α cores [24], plays an important role
53 to stabilize 2α cores [16, 25, 26], break the axial symmetry [24], or possibly quench the shell
54 gap at the magic number 8 [25–28]. However, there are limited experimental data to support
55 such interplay between α clusters and valence neutrons.

56 In the present paper, we report on the excitation properties of ${}^{10}\text{Be}$ via resonant α
57 scattering of ${}^6\text{He}$. The neutron-rich ${}^{10}\text{Be}$ nucleus is a simple system consisting of ${}^8\text{Be}$ plus
58 $2n$. Nevertheless, ${}^{10}\text{Be}$ involves a complex level scheme that features three rotational bands
59 built on the 0^+ ground state, the 1^- state at 5.96 MeV, and the second 0^+ state at 6.18 MeV.
60 Theoretically, the origin of these bands is explained in terms of molecular orbitals of valence
61 neutrons [15, 16]. In this picture, these neutrons are delocalized over the 2α cores and
62 occupy different orbital levels, thus different intrinsic states. There has been a number of
63 studies conducted in search of experimental signatures of the predicted intrinsic structures.

64 Particularly high-spin members of the $0_{\text{g.s.}}^+$ and 0_2^+ bands have been searched using different
 65 reaction probes. The 4^+ member of the 0_2^+ band was first speculated for the state at 10.2 MeV
 66 found in a study of the ${}^7\text{Li} + {}^7\text{Li}$ reaction [29]. While this state was later confirmed at
 67 10.15(2) MeV using the ${}^7\text{Li} + {}^{6,7}\text{Li}$ reactions, a spin-parity of 3^- was assigned from the
 68 angular correlation of α and ${}^6\text{He}$ particles following the α decay [30]. Another angular
 69 correlation study using the ${}^6\text{He} + {}^{6,7}\text{Li}$ reactions, however, gave a conflicting assignment
 70 of 4^+ , indicating that the method depends on the model assumed in reaction analyzes [31].
 71 A spin-parity of 4^+ was assigned without such assumptions in a recent measurement of α
 72 scattering from ${}^6\text{He}$, in which the 10.15-MeV state in ${}^{10}\text{Be}$ was resonantly populated and
 73 the angular distribution was characteristic of the angular momentum 4 [32]. The associated
 74 large width for α emission was interpreted as a strong indication of the predicted molecular
 75 structure. The 4^+ member of the $0_{\text{g.s.}}^+$ band is considered the 4^+ state at 11.76 MeV [33]. The
 76 α clustering of the ground state band is naïvely assumed from its level scheme being nearly
 77 equal to ${}^8\text{Be}$, having the 2^+ state at 3.37 MeV (${}^8\text{Be}$ 2_1^+ 3.03 MeV) and the 4^+ state at 11.76
 78 MeV (${}^8\text{Be}$ 4_1^+ 11.4 MeV) [33]. This assumption has been supported by theoretical studies
 79 that describe well the level scheme within the molecular orbital picture [15, 16, 18, 19, 21,
 80 22, 24]. The possibility of a shell-model-like structure was, however, recently discussed in a
 81 study of ${}^{10}\text{Be}$ via inelastic scattering with ${}^{12}\text{C}$ [34], where the excitation to the 4^+ member
 82 of the $0_{\text{g.s.}}^+$ band was not observed. This suggests there is a delicate balance between the
 83 persistence and the dissociation of α clusters in ${}^{10}\text{Be}$, and merits further elucidation.

84 In this study, we measured scattering of ${}^6\text{He}$ on α particles at low energies to resonantly
 85 populate states in ${}^{10}\text{Be}$. The resonance strength, which is related to the decay width for
 86 α emission, is an important indicator of the degree of α clusterization [32, 35]. Elastic α
 87 scattering of ${}^6\text{He}$ was measured in a few previous studies [32, 36–38]. The measurement by
 88 Ter-Akorpian *et al.* was performed at a center-of-mass energy ($E_{\text{c.m.}}$) of 60.3 MeV [36], while
 89 those of Raabe *et al.* were at 11.6 and 15.9 MeV [37, 38]. At these higher energies, the
 90 data are well reproduced by direct reaction analyses based on the coupled-discretized con-
 91 tinuum channels formalism [39] and the coupled-reaction channel formalism [40]. Freer *et al.*
 92 recently measured scattering at low energies below 5 MeV, and identified a resonance orig-
 93 inating from the 4^+ state at 10.15 MeV [32]. Their measurement was, however, performed
 94 at only three center-of-mass energies, namely $E_{\text{c.m.}} = 2.44, 3.00$ and 4.44 MeV. Therefore,
 95 there was no continuous excitation spectrum, which hampers the ability to thoroughly scan

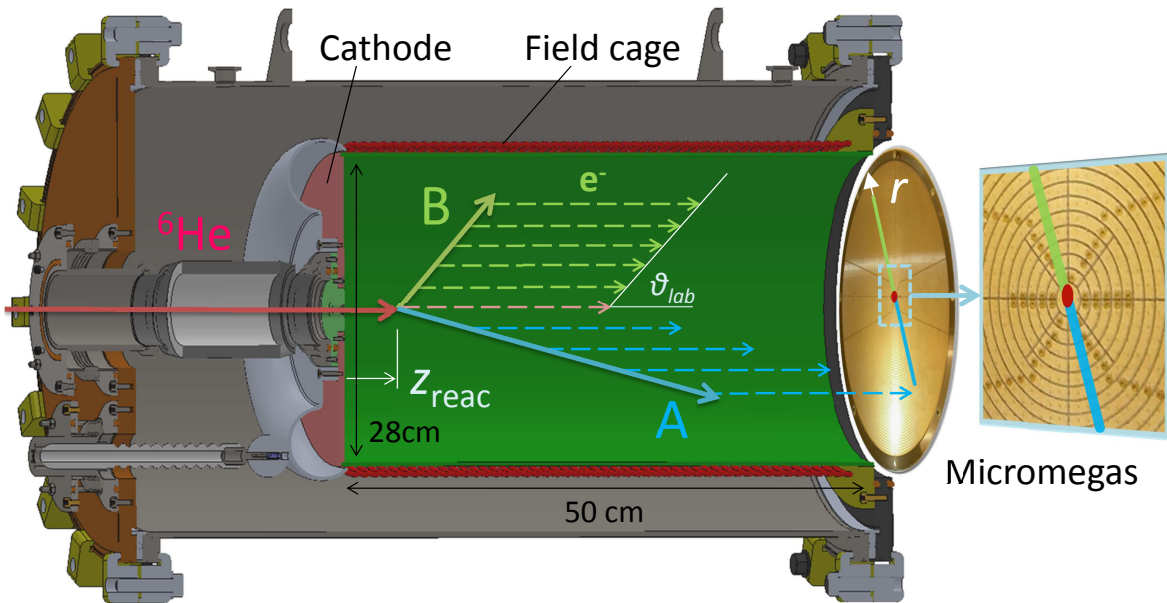
96 resonances over a given excitation energy region. In addition, the reconstruction of the reac-
97 tion kinematics had an ambiguity because recoiling α particles and scattered ${}^6\text{He}$ particles
98 were not differentiated, while the obtained angular distribution was correctly analyzed using
99 simulations that took this ambiguity into account. In the present study, we measured contin-
100 uous excitation functions over a wide energy range of $E_{\text{c.m.}} = 2\text{--}6$ MeV and reconstructed
101 angular distributions from unambiguously-identified recoiling α particles. This allowed us to
102 survey resonances over the energy domain where the 4^+ members of the ground-state band
103 and the second 0^+ band are located, as well as the region where some other resonances have
104 been predicted [14, 23].

105 An experimental challenge is the detection of low-energy helium particles from scattering,
106 which quickly lose their energy in the target medium. We used a time projection chamber
107 (TPC) in the “active-target” mode to address this issue, where the gas of the TPC serves
108 simultaneously as the α -particle target as well as the tracking medium of reaction products.
109 This allows measurements of the energy and scattering angle of particles which stop inside
110 the target. This setup is capable of measuring not only elastic scattering, but also inelastic
111 scattering and neutron-emission channels such as ${}^6\text{He}(\alpha, 2n){}^8\text{Be}$, in which negative reaction
112 Q values further lower the energy of the reaction products. This new method also enables
113 the determination of reaction energies from the direct measurement of reaction positions. In
114 the thick target method [41], the established and widely-used method to measure continuous
115 excitation functions in inverse kinematics, the reaction energy is indirectly obtained from
116 the energy of recoiling particles assuming a given kinematical scenario. This usually requires
117 several different settings for beam energies to avoid mixing elastic and inelastic scattering. In
118 our method, the unambiguous identification of the reaction energy allows us to differentiate
119 reaction channels, thus enabling us to cover a wide energy range in a single measurement.

120 II. EXPERIMENT

121 The experiment was performed at the *TwinSol* radioactive nuclear beam facility [42]
122 at the University of Notre Dame. Scattering of ${}^6\text{He}$ on α particles was measured using
123 the Prototype Active-Target Time-Projection Chamber (PAT-TPC) [43]. A secondary ${}^6\text{He}$
124 beam with an energy of 15 MeV was produced using the ${}^7\text{Li}(d, {}^3\text{He}){}^6\text{He}$ reaction. A stable
125 ${}^7\text{Li}$ beam was accelerated to 29.2 MeV by the FN Tandem accelerator and impinged on a

126 gaseous deuterium target. The pressure of the deuterium gas was 1200 mm-Hg on average.
 127 The gas cell had windows consisting of two 4.2-mg/cm² Havar foils 2.5 cm apart along
 128 the beam axis. The effective path length in the cell was longer by a few millimeters due to
 129 bowing of the foils under pressure. Outgoing reaction products were collected and focused by
 130 a pair of superconducting solenoidal magnets of the *TwinSol* device [42]. A 6-mg/cm² CH₂
 131 foil was placed at the cross-over point between the two magnets, which significantly slowed
 132 down higher Z contaminants, namely Li, from the primary beam, and greatly improved the
 133 purity of the ⁶He beam. During the beam tuning, a silicon E - ΔE telescope was used in
 134 front of the PAT-TPC as a beam monitor. The beam was predominantly composed of ⁶He
 135 and the main contaminant was ⁴He with an energy of about 22 MeV. The telescope was
 136 then removed from the beam line once the beam tuning was completed.



137

138 FIG. 1. (Color online) Schematic drawing of the experimental setup of the PAT-TPC. The inset
 139 is a magnified view of the segmented anode pad plane of the Micromegas detector near the beam
 140 axis.

141 The PAT-TPC was installed 1.5 m downstream of the end of *TwinSol*. A schematic
142 drawing of the setup is shown in Fig. 1. The detector encompasses a cylindrical field cage of
143 50 cm in length and 28 cm in diameter, which is centered on the beam axis. The cage was
144 filled with a He:CO₂ 90:10 mixture gas at atmospheric pressure, which serves simultaneously
145 as the tracking medium as well as the reaction target containing ⁴He. To limit impurities,
146 the gas was kept continuously flowing at a rate of 4.5 cm³/s, which replaces the whole volume
147 of gas every 2 hours. The ⁶He particles entering through an entrance window travel down
148 along the symmetry axis of the field cage and induce nuclear reactions with the gas nuclei.
149 Both the beam particles and outgoing charged particles from the reactions ionize gas atoms
150 while traveling across the gas volume. Their paths are reconstructed from the ionization
151 electrons, which are transported downstream by an electric field parallel to the beam axis,
152 and multiplied by the gas electron amplifier, Micromegas [44]. The latter was fabricated by
153 the SEDI/IRFU, CEA-Saclay. An electric field of 0.8 kV/cm was generated by a negative
154 potential of -40 kVDC applied to a cathode plate at the upstream end of the cage. The
155 initial potential was stepped down by a series of equipotential rings toward the Micromegas.
156 The resulting electron drift velocity was measured to be about 2.4 cm/ μ s [43]. The field
157 cage is gas tight and is surrounded by nitrogen gas at atmospheric pressure, which provides
158 a high dielectric strength to mitigate the risk of discharge to the chamber walls at ground
159 potential.

160 The secondary beam was focused on the entrance window of the PAT-TPC. The spot
161 size of the ⁶He beam at the window was estimated to be 25 mm in diameter from solenoid
162 optics calculations, while the entrance aperture is circular with a diameter of about 8 mm.
163 The beam was thus collimated by the window, which changed the intensity and purity of
164 ⁶He in the TPC. These quantities were estimated using signals from the micromesh of the
165 Micromegas, of which a detailed description is given later. The ⁶He particles are decelerated
166 in the gas and stop at the end of the field cage about 50 cm downstream of the beam
167 window, fully depositing 15 MeV of energy. In contrast, ⁴He has higher energy and punches
168 through the TPC while losing only 6 MeV in the detector volume. This results in a pulse-
169 height difference in the micromesh signals, thus allowing identification and counting of the
170 respective nuclei. The average intensity of the collimated ⁶He beam was 2×10^3 counts per
171 second with a purity of 90% at a primary ⁷Li(3⁺) beam intensity of 0.5 electric μ A.

172 The Micromegas consists of an anode readout plane and a micromesh stretched over it.

173 Electron avalanches occur in the narrow amplification gap of $128\ \mu\text{m}$ defined by the anode
174 and the micromesh. The micromesh was biased at $-320\ \text{V}$ to create a high field gradient
175 over the gap, while the anode plane was grounded. The gas gain was measured to be about
176 75 as also reported elsewhere [43]. The anode plane of the Micromegas has a circular-shaped
177 active area of $250\ \text{mm}$ in diameter. It is segmented into multiple pads to read out and locate
178 the avalanche electrons. Specifically, the anode pads consist of a 5-mm -diameter central
179 pad and 2-mm -pitch coaxial strips, each spaced by a gap of $0.25\ \text{mm}$ (inset of Fig. 1). The
180 coaxial strips are divided into quadrants and the five innermost strips into octants. The
181 beam particles travel approximately perpendicular to the anode plane. The central pad and
182 its neighboring coaxial strips record the energy deposit profile as a function of time, and
183 the position along the beam axis (z) is deduced from the drift time of ionization electrons.
184 The particles emitted from a reaction travel away from the center to the perimeter. Each
185 particle traverses a series of coaxial strips in a given quadrant. This set of strips thus gives
186 the energy deposit profile as a function of the radius r of the strip as well as z , providing
187 the polar angle and the range of the reaction products. Examples of energy deposit profiles
188 are found in our previous report [43].

189 A total of 253 signals from the anode plane were read out by a combined preamplifier/shaper/wave-
190 digitizer system developed for the neutrino-flux monitor of the T2K experiment [45, 46],
191 which is referred to as T2K electronics hereafter. The T2K electronics were set to record
192 the waveform of input signals at $12.5\ \text{MHz}$ over the full time range of $40\ \mu\text{s}$. A time bin
193 width thus corresponds to $80\ \text{ns}$, or a spatial size of $2\ \text{mm}$ along the beam axis given the
194 electron drift velocity of $2.4\ \text{cm}/\mu\text{s}$. To generate external triggers for the T2K electronics,
195 signals from the micromesh as well as from some coaxial anode strips were routed to an
196 auxiliary circuit consisting of NIM-standard modules. Two sets of data were taken with
197 different triggers. The first trigger was optimized for the elastic and inelastic scattering
198 of ${}^6\text{He}$ on α particles. The primary signature of these reactions is the observation of two
199 helium isotopes, ${}^4\text{He}$ and/or ${}^6\text{He}$, involved in the final state. Because of the low atomic
200 number ($Z = 2$), both particles have a long range and traverse many strips away from the
201 center. The trigger was thus designed to record events which fire specific off-center strips in
202 different quadrants. The four coaxial strips, each belonging to different quadrants, located
203 at a radius of $18\ \text{mm}$ were used as trigger sources. The data acquisition was triggered on
204 tracks that extend $18\ \text{mm}$ or more in radius in two or more quadrants. For monitoring

205 purposes, the trigger from one quadrant, or multiplicity equal to one, was also added after
 206 downscaling by a factor of 32. The second trigger was used to record ${}^6\text{He} + \alpha$ reactions
 207 involving beryllium in the final state. These reactions produce a large energy deposit in a
 208 short distance due to the high Z of Be and its low recoil velocity. The time structure of
 209 the micromesh signal was used to generate a trigger when the signal had (a) a higher Bragg
 210 peak and (b), later times with respect to the Bragg peak of the fully stopped ${}^6\text{He}$ beam.
 211 The late times imply that the observed Bragg peak is located closer to the beam entrance,
 212 making it a good indicator of the occurrence of a reaction during the deceleration of the
 213 beam. The average rate of Trigger 1 was 15 Hz, while that of Trigger 2 was 50 Hz. The
 214 live-time ratio of the data acquisition was 70% and 40%, respectively, for these two triggers.

215 III. ANALYSIS

216 A. Elastic and inelastic scattering

217 In the following section, we describe the analysis procedure of the elastic and inelastic
 218 scattering data taken with Trigger 1 (optimized for these processes). Part of the analysis
 219 procedure, particularly the tracking of the ${}^6\text{He}$ beam and reaction products (${}^6\text{He}$ and/or
 220 ${}^4\text{He}$), is the same as reported in Ref. [43]. Energy deposition was calculated using the SRIM
 221 code [47].

222 The total kinetic energy (TKE) and the emission angle (θ_{lab}) of reaction products were
 223 obtained by analyzing the set of information on charge Q_i , radial position r_i and signal
 224 timing t_i from the coaxial strips, denoted by i , which the particle tracks traveled over. First,
 225 the angle θ_{lab} , which is defined with respect to the beam axis, or the axis of time projection
 226 (Fig. 1), was determined from the slope of the t vs. r plot. A linear fitting function was
 227 adopted and defined as:

$$t = \frac{r}{v_{\text{drift}} \tan \theta_{\text{lab}}} + t_1 \quad (1)$$

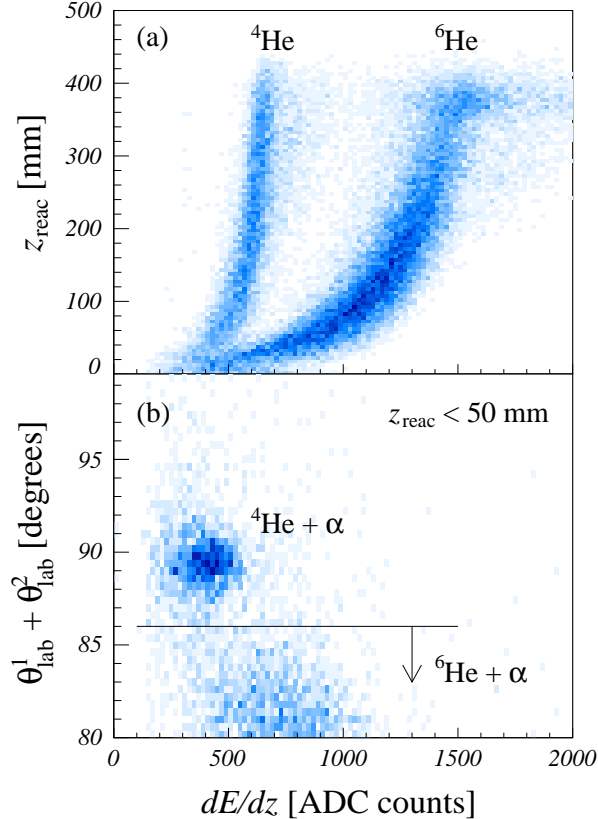
228 with v_{drift} being the electron drift velocity and t_1 the time at $r = 0$. Once θ_{lab} was determined
 229 from the fit, the Q vs. r plot, which represents the energy deposition profile in the radial
 230 direction, was compared to calculated curves at a given TKE using the θ_{lab} previously
 231 obtained. The optimal TKE was determined to minimize χ^2 with respect to the experimental
 232 data. Some of the particles escaped from the active volume of the PAT-TPC, particularly

233 the ones with large or small scattering angles where the energies are higher. To obtain
 234 the acceptance over a wide angular range, we also analyzed the events involving particle
 235 escapes. While the range cannot be determined for these particles due to the missing position
 236 information of the Bragg peak, the amplitude and slope of the energy deposit profile towards
 237 the Bragg peak are still sensitive to the TKE, which allows TKE determination by the same
 238 χ^2 minimization procedure. The quality of the TKE determination of escaping particles
 239 will be discussed later when the reconstruction of the excitation energy spectrum of ${}^6\text{He}$
 240 is presented. When a given particle stops inside the active volume, ${}^6\text{He}$ and α particles
 241 can be differentiated from the amplitude of the tail of Bragg peak as demonstrated in our
 242 previous report [43]. In the present analysis, however, we differentiated ${}^6\text{He}$ and ${}^4\text{He}$ from
 243 the reaction kinematics as detailed later.

244 Reconstruction of reaction kinematics by the missing mass method requires knowing
 245 the energy of the beam particles. In the present measurement, the energy continuously
 246 decreases as the beam particle travels along the beam axis. To determine the reaction
 247 energy, the energy deposition before reaching the reaction position (z_{reac}) needs to be taken
 248 into account. In this analysis, z_{reac} was determined from the recorded drift times of ionization
 249 electrons released at the positions of the reaction vertex and the beam entrance. The drift
 250 time corresponding to the reaction position was given by that of the reaction vertex t_1
 251 previously deduced in the analysis of the trajectory of the reaction products. The drift time
 252 of electrons released at the entrance (t_0) was obtained by analyzing the waveform of anode
 253 signals near the central region, which represents the energy deposit profile of beam particles
 254 along the beam axis. The anode signals of the central pad and its neighboring coaxial strips
 255 were summed when the beam charge spreads over multiple pads. The difference between
 256 the two times $t_0 - t_1$ was then translated into z_{reac} using the electron drift velocity. The
 257 corresponding energy loss was calculated and subtracted from the initial beam energy to
 258 define the energy at the moment of the reaction (E_{reac}).

259 The ${}^6\text{He}$ ions and the contaminant ${}^4\text{He}$ ions in the secondary beam were differentiated
 260 from the energy deposit per unit length (dE/dz) averaged over the track from t_0 to t_1 . The
 261 value of dE/dz provides a good measure of isotope separation as the ${}^6\text{He}$ particles have
 262 lower energies (3.8 MeV/ u or less) compared to the ${}^4\text{He}$ contaminants (4.4–5.6 MeV/ u).
 263 Figure 2(a) shows the scatter plot between dE/dz and z_{reac} , where two loci corresponding
 264 to ${}^6\text{He}$ and ${}^4\text{He}$ are clearly separated. The separation of the loci, which becomes smaller at

265 shorter travel lengths, is uncertain below $z_{\text{reac}} = 50$ mm, where the finite value of the shaping
 266 time ($0.5 \mu\text{s}$) smears the difference in dE/dz . In this region, the properties of the reaction
 267 kinematics were further used to eliminate elastic scattering events of ${}^4\text{He}$ on α particles. A
 268 gate was set on the sum of the angles $\theta_{\text{lab}}^{(i)}$ of the two reaction products labeled $i = 1$ and 2.
 269 Figure 2(b) shows the $\theta_{\text{lab}}^{(1)} + \theta_{\text{lab}}^{(2)}$ vs. dE/dz plot for the region of $z_{\text{reac}} < 50$ mm. A cluster
 270 is seen at 90 degrees. This corresponds to ${}^4\text{He}$ beam scattered by the ${}^4\text{He}$ in the gas. The
 271 opening angle of two identical particles from elastic scattering always equals 90 degrees in
 272 the laboratory frame regardless of the center-of-mass scattering angle. The gate displayed
 273 in the figure was adopted to exclude these ${}^4\text{He}$ -beam events.



274

275 FIG. 2. (Color online) Identification of the secondary beam particles. (a) Scatter plot between the
 276 energy loss per unit length of beam path dE/dz and the reaction position z_{reac} . (b) Scatter plot
 277 between dE/dz and the opening angle of reaction products $\theta_{\text{lab}}^{(1)} + \theta_{\text{lab}}^{(2)}$ for the events at $z_{\text{reac}} < 50$ mm.
 278 The gate to select ${}^6\text{He}$ scattering is indicated.

279 The excitation energy of ${}^6\text{He}$ (E_x) and the center-of-mass scattering angle ($\theta_{\text{c.m.}}$) were

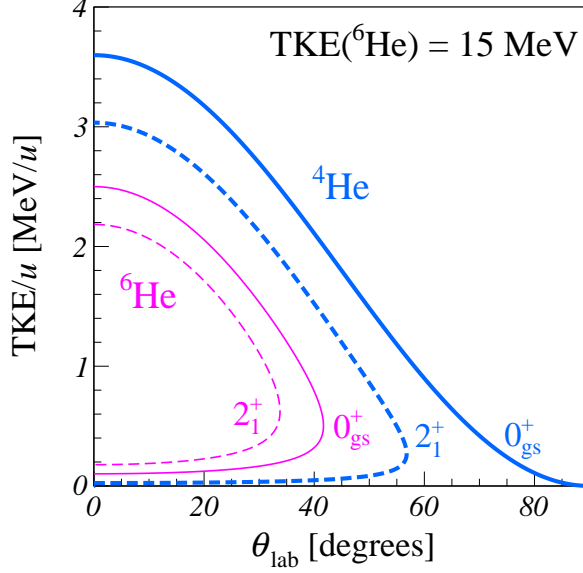
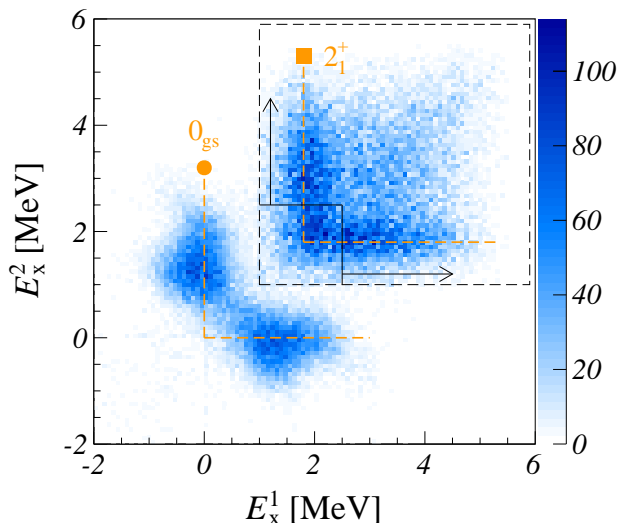


FIG. 3. (Color online) TKE/ u vs. θ_{lab} plot for ${}^6\text{He} + \alpha$ scattering at 15 MeV. Elastic scattering and inelastic scattering to the first 2^+ state of ${}^6\text{He}$ are denoted by the solid and dashed lines, respectively. Calculated curves are shown both for the recoiling α (thick blue) and scattered ${}^6\text{He}$ (thin magenta) particles.

280 obtained by the missing-mass method using E_{reac} as well as TKE and θ_{lab} of the recoiling
 281 α particles. An inherent issue in deducing excitation-energy spectra from ${}^6\text{He} + \alpha$ -particle
 282 scattering is that another α particle is produced in the final state when inelastically scattered
 283 ${}^6\text{He}$ decays via $2n$ emission. All excited states of ${}^6\text{He}$ are unbound above the $2n$ separation
 284 energy at 0.972(1) MeV [48]. It is therefore necessary to correctly select the α particle
 285 recoiling from the target. The information on kinematical properties of the reaction was
 286 used to eliminate the ambiguity in identifying the recoiling particle. Figure 3 shows the
 287 calculated plots of TKE per nucleon (TKE/ u) vs. θ_{lab} for elastic and inelastic scattering
 288 to the 2^+ state at a beam energy of 15 MeV. It is found that TKE/ u , or the velocity of
 289 ${}^6\text{He}$, is always smaller than that of the recoiling α particle at a given laboratory angle
 290 except for the very forward region below 10 degrees in the center-of-mass frame, which is
 291 outside the present detector acceptance. Therefore, the range of the α particle following
 292 $2n$ emission decay tends to be shorter than that of the recoiling α particle. According to
 293 these characteristics, we adopted the following procedure. First, two excitation energies
 294 ($E_x^{(i)}$) were obtained individually from two reaction products by assuming the particle i as

295 the recoiling α particle. The set of $E_x^{(i)}$ were then compared. If the ${}^6\text{He}$ particle, or the α
 296 particle after $2n$ emission decay, is identified as the recoiling α particle, its shorter range
 297 results in an underestimate of TKE and thus gives higher excitation energies than the true
 298 value. Accordingly, we adopted the particle with a smaller E_x as the recoiling α particle.



299

300 FIG. 4. (Color online) Scatter plot between $E_x^{(1)}$ and $E_x^{(2)}$. The events to the left of the solid lines
 301 were excluded from the analysis. The contents of the region inside the dashed lines are weighted
 302 by a factor of 3 for presentation purposes.

303 Figure 4 shows the scatter plot between $E_x^{(1)}$ and $E_x^{(2)}$, where the indices 1 and 2 were
 304 randomly assigned. Loci corresponding to the ground state and the first 2^+ state at 1.79 MeV
 305 are visible. It is evident in the locus of elastic scattering that the smaller $E_x^{(i)}$ value of a
 306 given pair is mostly found near 0 MeV and well separated from the larger $E_x^{(i)}$ which is
 307 wrongly estimated from the scattered ${}^6\text{He}$. The result is similar for the 2^+ state, of which
 308 the locus appears above 1.9 MeV only. However, the separation in the region close to the
 309 $E_x^{(1)} = E_x^{(2)}$ line is not as clear as that of the ground state because of the velocity shift due to
 310 particle emission and the contribution of breakup events. Therefore, above $E_x = 1$ MeV, the
 311 events with $E_x^{(1)}$ and $E_x^{(2)} < 2.5$ MeV were eliminated from the analysis to ensure a sufficient
 312 difference in energy for properly selecting α particles.

313 The resulting excitation-energy spectrum of ${}^6\text{He}$ is shown in Fig. 5(a). Two peaks
 314 visible near 0 and 2 MeV correspond to the 0^+ ground state and the first 2^+ state at

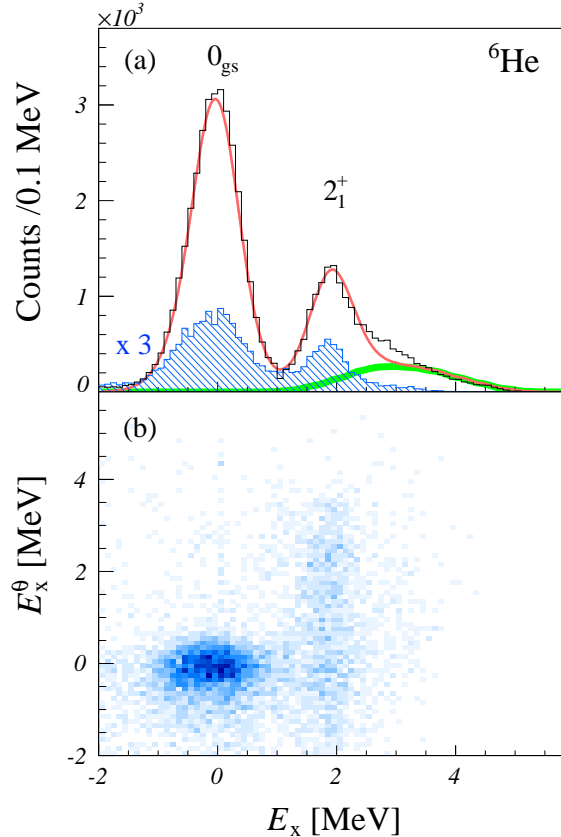


FIG. 5. (Color online) (a) Excitation-energy spectrum of ${}^6\text{He}$ (unshaded histogram). The best-fit result (thin red line) is shown together with the contribution of the background from the ${}^6\text{He}$ breakup reaction (thick green line). The spectrum in the shaded histogram was made by selecting backward scattering ($\theta_{\text{c.m.}} > 90^\circ$) with particle escape from the TPC. (b) Scatter plot of E_x and E_x^θ . E_x^θ is the excitation energy of ${}^6\text{He}$ obtained from the angles of the two reaction products.

315 1.797(25) MeV [49]. A fit to the spectrum gives excitation energies of $-0.01(1)$ and
 316 1.90(1) MeV, respectively, where the quoted errors are statistical. The FWHM resolu-
 317 tion is about 0.85 MeV. The fitting function consisted of a Gaussian function for the ground
 318 state, a Voigt function [50] for the unbound 2^+ state, and background from the breakup
 319 reaction of ${}^6\text{He}$ involving the final state of an α particle and 2 neutrons. The Voigt function
 320 is a convolution of a Breit-Wigner function and a Gaussian, which allows the natural width
 321 of an unbound state and the resolution of the detection system to be taken into account.
 322 The width of the 2^+ state was set to the adopted value, 0.113(20) MeV [49]. The background
 323 shape was simulated by the Monte-Carlo method, where the α particle and neutrons in the

324 final state were generated in a uniform phase space. The procedure of identification based on
 325 the comparison between a pair of $E_x^{(i)}$ was taken into account. The best-fit curve shown in
 326 the figure well reproduces the experimental data. The background of the breakup reaction
 327 denoted by the thick line accounts for the tail towards higher energies.

328 In the present analysis, TKE for an input to the missing-mass method was deduced
 329 from the slope of the Bragg curve when recoiling α particles escape from the active region
 330 and hence the Bragg peak is unavailable. Without the precise position information of the
 331 Bragg peak, the analysis might result in inaccurate excitation energies. The gated spectrum
 332 for one-particle escape events is shown in the shaded spectrum of Fig. 5(a). The spectrum
 333 clearly shows a two peak structure, confirming good reconstruction of the reaction kinematics.
 334 To further confirm the results, the adopted excitation energies are compared to excitation
 335 energies (E_x^θ) that were obtained from the correlation of laboratory angles, $\theta_{\text{lab}}^{(1)}$ and $\theta_{\text{lab}}^{(2)}$,
 336 between the two reaction products. This method does not require the knowledge of the
 337 TKE of reaction products. Thus we can correctly extract excitation energies from the
 338 escape events as long as the scattered ${}^6\text{He}$ particle is stable against particle emission and
 339 keeps its initial angle unchanged after scattering. The scatter plot of E_x^θ against E_x is shown
 340 in Fig. 5(b). A gate was set to select backward scattering angles of $\theta_{\text{c.m.}} > 90$ degrees, where
 341 most of the escaping particles are recoiling α nuclei. The presence of a cluster centered at
 342 $E_x^\theta = E_x = 0$ MeV indicates that most of the events involving the ground state have E_x
 343 well correlated with E_x^θ . The ratio of the number of events in the cluster is nearly 90% with
 344 respect to that of the peak in the shaded spectrum in Fig. 5(a), limiting the systematic
 345 uncertainty in yields to 10%. The locus for the 2^+ state widely spreads along the vertical
 346 axis of E_x^θ because the angles of scattered particles shift after the $2n$ emission.

347 B. Two-neutron emission channel

348 To investigate branching to the two-neutron emission ${}^6\text{He}(\alpha, 2n){}^8\text{Be}$ channel, the trigger
 349 was set for reactions having a peak in the energy loss profile which was taller than the
 350 Bragg peak of the ${}^6\text{He}$ beam particles (Trigger 2). In the data analysis a peak was observed
 351 for Bragg amplitudes that corresponded closely to $4.5 \text{ MeV mg}^{-1} \text{ cm}^2$, the value expected
 352 for two highly-correlated α particles, such as originating from the decay of ${}^8\text{Be}$. Since this
 353 amplitude is lower than $5.9 \text{ MeV mg}^{-1} \text{ cm}^2$ for ${}^9\text{Be}$ ions with a higher Z recoiling from

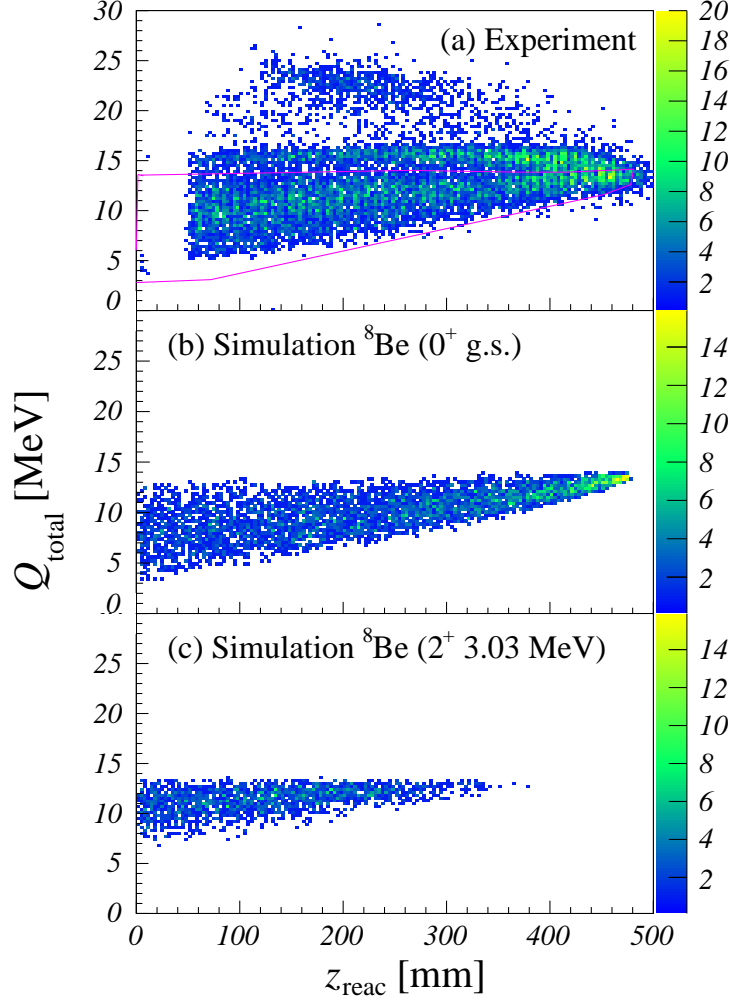


FIG. 6. (Color online) Total charge collected Q_{total} as a function of the location of the reaction vertex z_{reac} . (a) Experimental data. The elastic events are clearly distinguishable near $Q_{\text{total}} = 15$ MeV. Events above the locus of elastic scattering indicate pile up of beam particles. Only events inside the contour line were used in calculating the excitation function. (b) Simulated results for the decay into the 0^+ ground state and (c) the 2^+ state at 3.03 MeV of ^8Be .

354 the $^6\text{He}(\alpha, n)^9\text{Be}$ reaction, we concluded the main decay channel that contributed was
 355 $^{10}\text{Be} \rightarrow 2n + ^8\text{Be}$, with ^8Be unbound to α decay by 92 keV. Beam contaminants were
 356 eliminated offline by comparing the particle's energy loss profile near the beginning of its
 357 track to the experimentally-measured energy-loss profile for ^6He . The location of the reaction
 358 vertex z_{reac} was determined from the time where the enhancement of the energy deposits
 359 starts due to the creation of reaction products. Since the beam particles and the charged

360 reaction products were stopped in the detector, we could use the total charge collected in the
 361 detector (Q_{total}) as the energy sum signal, which was obtained by summing all anode signals.
 362 Q_{total} is plotted as a function of the location of the reaction vertex z_{react} in Fig. 6(a). The total
 363 charge is converted to energy by normalizing the experimental data to the allowed energies
 364 from a kinematical simulation of the sequential decay of $^{10}\text{Be} \rightarrow 2n + ^8\text{Be} \rightarrow 2n + 2\alpha$
 365 described later. The events shown are selected by the Bragg amplitude for two- α correlated
 366 events. The events in the $z_{\text{react}} < 50$ mm region were rejected because it was difficult to
 367 distinguish ^8Be events from beam contaminants, particularly ^7Li , when the incident energy
 368 loss curve was relatively short. As can be seen in Fig. 6(a), there is an accumulation of
 369 events near 15 MeV for the charge collected for full beam energy deposition corresponding
 370 to elastic scattering. Below this line is a broad range of energies deposited in the detector.
 371 These events are attributed to the $^6\text{He}(\alpha, 2n)^8\text{Be}$ reaction since the neutrons were not
 372 detected in our experiment and the energy taken away by them will be missing in the total
 373 energy signal. We simulated events corresponding to the decay of ^{10}Be into ^8Be with the
 374 unobserved two neutrons, for which we assumed isotropic emission in the center-of-mass
 375 system and phase-space sharing of the available energy between the two neutrons. It is also
 376 assumed that the decay of $^{10}\text{Be} \rightarrow 2n + ^8\text{Be} \rightarrow 2n + 2\alpha$ is sequential. The decay into the
 377 2^+ state at 3.03 MeV of ^8Be , which is energetically allowed at higher energies, is simulated
 378 as well. The results for the ground state and the 2^+ state are shown in Fig. 6(b) and (c),
 379 respectively. The kinematical simulation for the ground state is seen to agree well with the
 380 data, confirming the interpretation of the events. The simulation also served as a guide
 381 for defining the contour line seen in Fig 6(a). Only the events inside the contour line were
 382 subsequently used to calculate the excitation function.

383 IV. RESULTS

384 A. Elastic and inelastic scattering

385 To obtain excitation functions and angular distributions, cross sections ($d\sigma/d\Omega$) were
 386 deduced for a given bin of the reaction position z_{react}^i and the scattering angle $\theta_{\text{c.m.}}^j$ specified
 387 by the indices (i, j), respectively. Yields ($Y^{(i, j)}$) of the ground state and the 2^+ state were
 388 obtained by fitting excitation energy spectra using the same function as adopted for the fit

389 to the angle-integrated spectrum of Fig. 5(a). $Y^{(i, j)}$ was then translated into $d\sigma/d\Omega^{(i, j)}$
 390 following the expression:

$$d\sigma/d\Omega^{(i, j)} = Y^{(i, j)} / \left(\epsilon_{eff}^{(i, j)} \cdot \epsilon_{DAQ} \cdot N_{6\text{He}} \cdot N_{\text{He}}^{(i)} \cdot \Delta\Omega^{(j)} \right), \quad (2)$$

391 where the indices i and j denote that the associated parameter depends on z_{reac} and $\theta_{\text{c.m.}}$,
 392 respectively. The detection efficiency $\epsilon_{eff}^{(i, j)}$ was estimated by simulations, where the geome-
 393 tries of the detector, the condition of the trigger, and the energy losses of beam and reaction
 394 products were taken into account. The live-time ratio of the data acquisition ϵ_{DAQ} was 70%
 395 on average. The integrated count of ${}^6\text{He}$ beam particles $N_{6\text{He}}$ was estimated using the counts
 396 of micromesh signals measured by a scaler. This was about 2×10^8 after taking into account
 397 the beam purity of 90%. The number of He atoms in the gaseous target $N_{\text{He}}^{(i)}$ was calculated
 398 from the density of the He:CO₂ 90:10 gas at atmospheric pressure (0.33 mg/cm³) and the
 399 bin size of z_{reac} . The solid angle $\Delta\Omega^{(j)}$ was calculated from the bin size of $\theta_{\text{c.m.}}$.

400 The resulting excitation functions for elastic scattering are shown in Fig. 7. The beam
 401 energy in the laboratory frame, E_{reac} , was converted into the center-of-mass energy by
 402 $E_{\text{c.m.}} = 0.4 \times E_{\text{reac}}$. Given the separation threshold of an α particle at 7.42 MeV [48],
 403 $E_{\text{c.m.}}$ is related to the excitation energy in ${}^{10}\text{Be}$ by $E_x = E_{\text{c.m.}} + 7.42$ MeV. The systematic
 404 error of $E_{\text{c.m.}}$ is estimated to be ± 0.1 MeV, which arises from the deviation of E_{reac} with
 405 respect to the energy sum of the ${}^6\text{He}$ and α particles from elastic scattering. A strong en-
 406 hancement of cross sections is observed around 2.7 MeV in the angle-integrated spectrum of
 407 $\theta_{\text{c.m.}} = 65^\circ - 135^\circ$ in Fig. 7(a). A peak consistently exists around this energy in the different
 408 angular regions shown in Figs. 7(b)–(h), indicating the presence of a resonance. A fit with a
 409 Voigt function [50] and a linear background gives a resonance energy of $E_{\text{c.m.}} = 2.57(15)$ MeV,
 410 where the quoted uncertainty is the sum of the statistical and systematic errors. This is
 411 consistent with the resonance of the state at $E_x = 10.15(2)$ MeV, or $E_{\text{c.m.}} = 2.73(2)$ MeV,
 412 observed in a previous measurement of elastic scattering [32]. No peak is visible in the
 413 excitation functions except for this strong resonance. The absence of sizable resonances
 414 indicates that no other states have a large decay width for α emission within the energy
 415 window of the present study ($E_{\text{c.m.}} = 2 - 6$ MeV).

416 The excitation functions for inelastic scattering are shown in Fig. 7. A sizable decay
 417 width to the 2^+ state of ${}^6\text{He}$ was inferred for the 2.73-MeV resonance in the analysis of
 418 the previous study [32]. However, this could not be measured in the present study due to

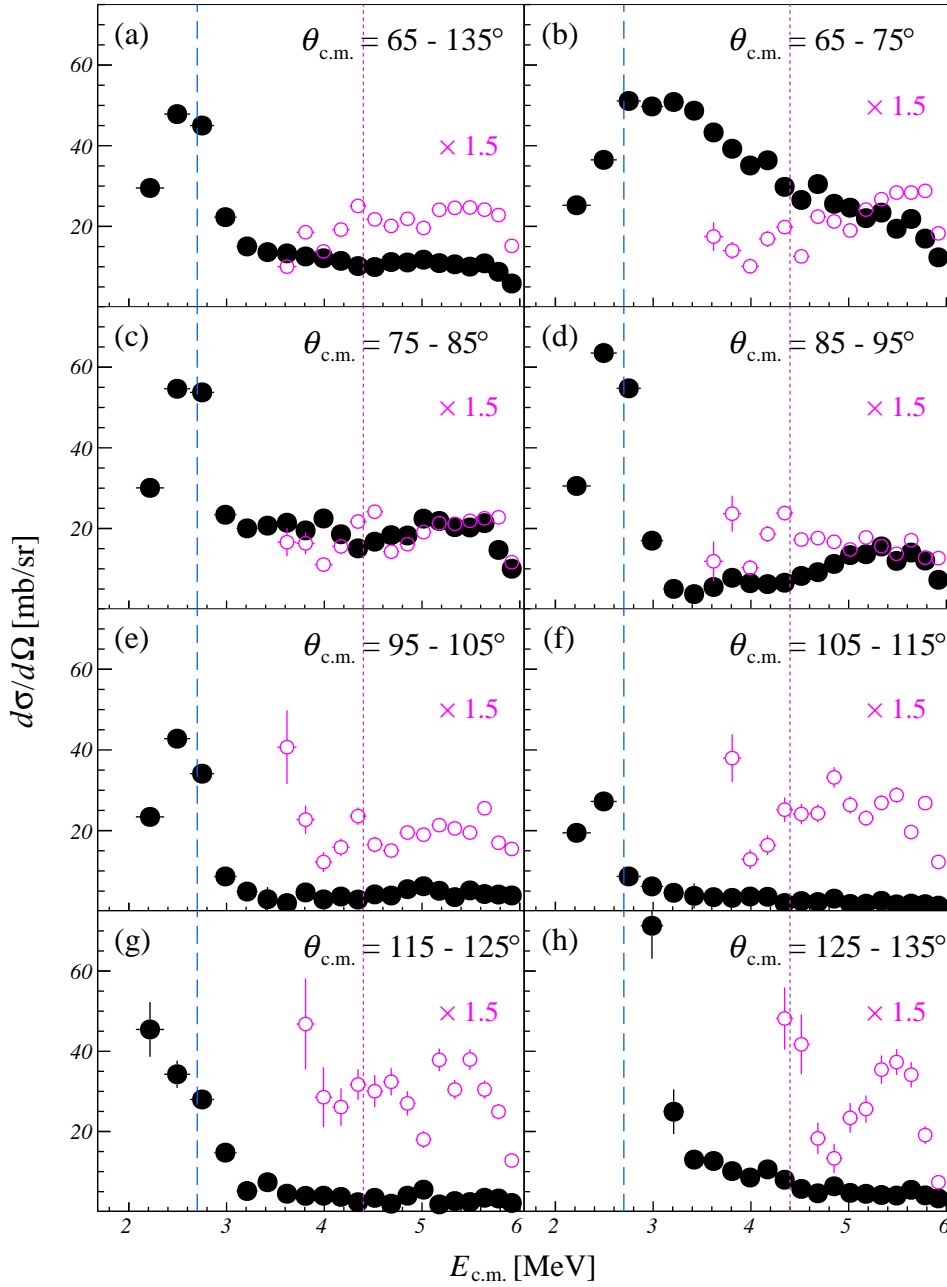


FIG. 7. (Color online) Excitation functions of ${}^6\text{He} + \alpha$ scattering: (a) angle-integrated spectrum over $\theta_{\text{c.m.}} = 65^\circ - 135^\circ$, and (b–h) spectra for 10° angular bins. The data for elastic scattering are shown by the filled circles, while those for inelastic scattering to the 2^+ state are shown by the open circles. The inelastic scattering data are scaled by a factor of 1.5. The dashed and dotted lines denote $E_{\text{c.m.}} = 2.7$ and 4.4 MeV, respectively.

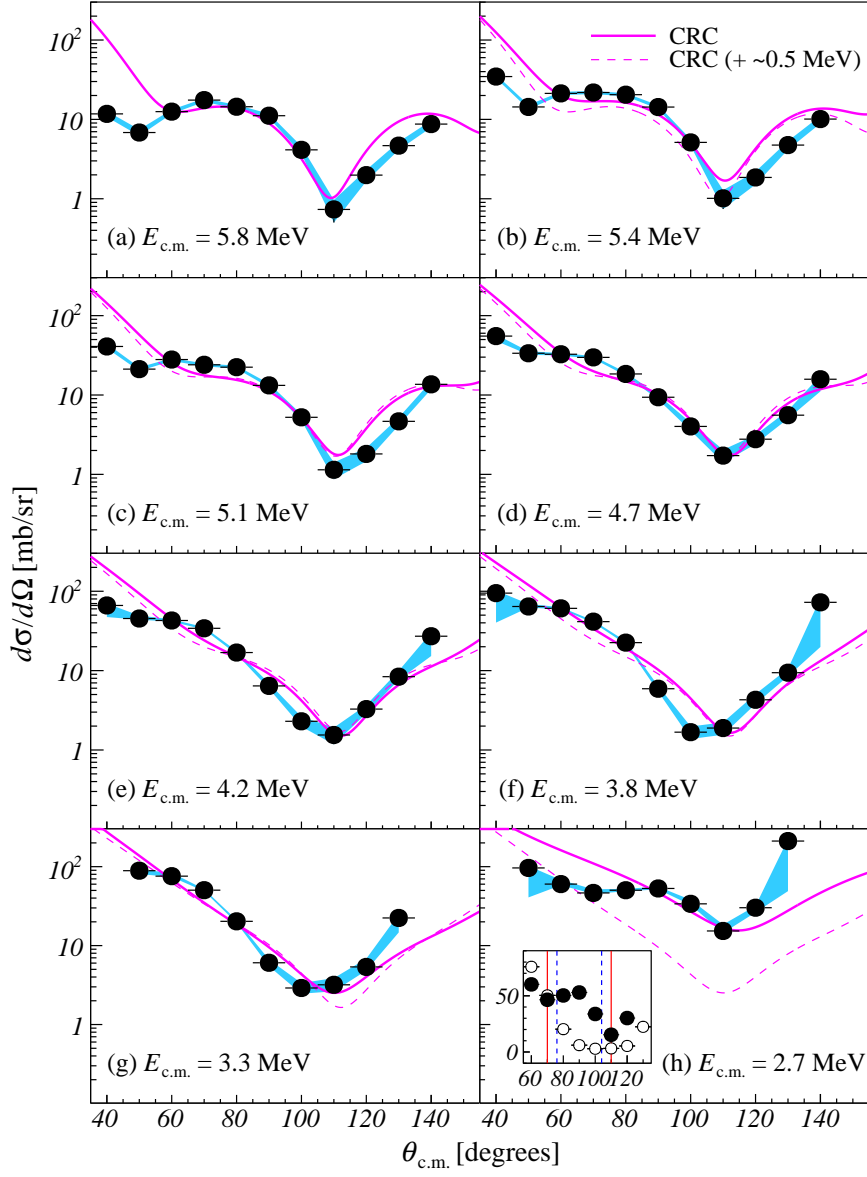


FIG. 8. (Color online) Angular distributions of elastic ${}^6\text{He} + \alpha$ scattering. The results of CRC calculations (solid lines) are compared to the data. The CRC results at the higher energy bins are also shown for reference with the dashed lines. The blue shaded areas denote the systematic errors from the ambiguities in the beam angle. The inset (h) shows the data at $E_{\text{c.m.}} = 2.7$ (full circles) and 3.3 MeV (open circles) on a linear scale. The angles where the Legendre polynomials $P_L(\cos \theta_{\text{c.m.}})$ for $L = 4$ and 6 become zero are denoted by the solid and dashed lines, respectively.

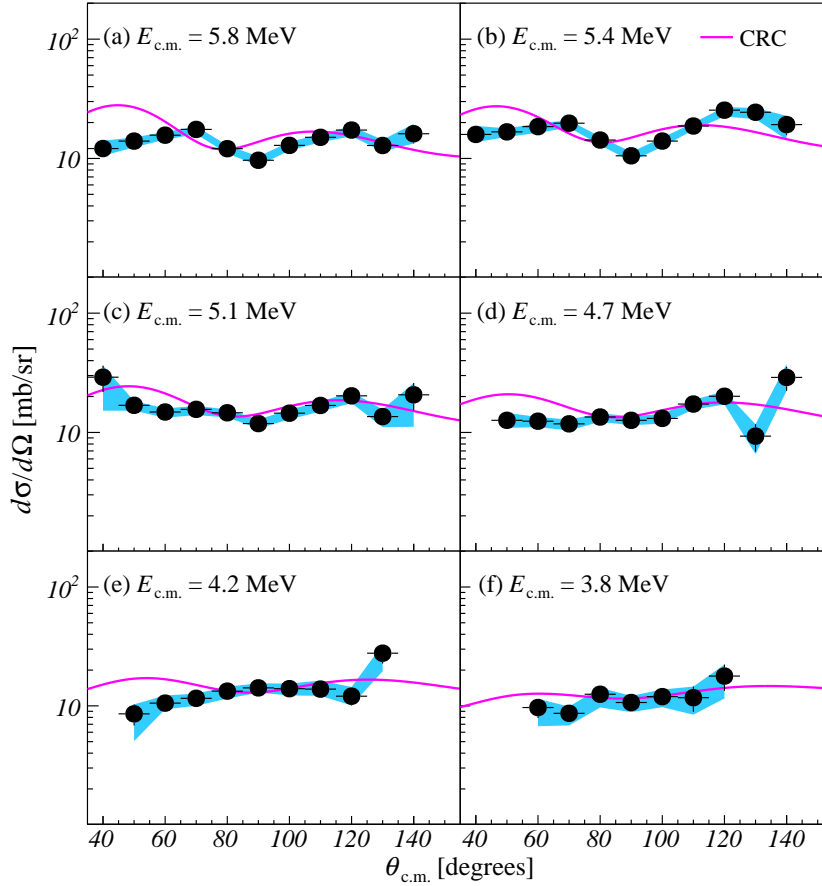


FIG. 9. (Color online) Angular distributions of inelastic ${}^6\text{He} + \alpha$ scattering to the ${}^6\text{He} 2^+$ state. The results of CRC calculations (solid lines) are compared with the data. The blue shaded areas denote the systematic errors from the ambiguities of the beam angle.

419 the lack of acceptance at this energy. While there are no resonances as strong as the one
 420 observed in the elastic channel, a small peak is visible at 4.4 MeV in the inelastic channel
 421 consistently over the angular region of $\theta_{\text{c.m.}} = 65^\circ - 105^\circ$, which might be due to a resonance.

422 The differential cross sections were deduced for eight different incident energies to study
 423 angular distributions. The resulting data for elastic scattering are displayed in Fig. 8. The
 424 vertical error bars are only statistical, and the horizontal bars denote the size of the angular
 425 bins. The leading contributions of the systematic error are the uncertainty of yields for
 426 the escape events (10%) and the detection efficiency. The latter mostly comes from the
 427 uncertainties of the beam angle, which could be inclined by about 1 degree with respect to
 428 the central axis. The blue shaded areas in Figs. 8 denote the variation of cross sections when

429 the incident beam angle varies by 1 degree. As seen in the figures, a series of differential
 430 cross sections for elastic scattering show a gradual and continuous change in their angular
 431 distributions as a function of $E_{\text{c.m.}}$ until the cross sections are enhanced at $E_{\text{c.m.}} = 2.7$ MeV.
 432 In the inset of Fig. 8(h), the on-resonance data at $E_{\text{c.m.}} = 2.7$ MeV (filled circles) are
 433 compared to the off-resonance data at $E_{\text{c.m.}} = 3.3$ MeV (open circles) shown on a linear scale.
 434 At $E_{\text{c.m.}} = 2.7$ MeV, the cross section clearly peaks at 90° and dips at 60° and 110° . In
 435 contrast, the data at $E_{\text{c.m.}} = 3.3$ MeV lack such a steep rise and drop, indicating that the peak
 436 and dip structures seen in the 2.7-MeV data are due to the resonance. Angular distributions
 437 of resonance scattering primarily follow the square of the Legendre polynomial $[P_L(\cos \theta_{\text{c.m.}})]^2$
 438 corresponding to the angular momentum (L) involved. The diffractive pattern observed is
 439 therefore a useful measure to identify L . The presence of a peak at 90° rules out the odd
 440 angular momenta, of which the corresponding Legendre polynomials always become zero at
 441 this angle. The dips at 70° and 110° more agree with the polynomial of $L = 4$ than that of
 442 $L = 2$ having zeros at 55° and 125° , or $L = 6$ at 76° and 104° . The dip angles of $L = 4, 6$
 443 are shown for reference by the solid and dashed lines, respectively, in the inset of Fig. 8(h).
 444 The diffractive pattern of the experimental data is seen closer to that of $L = 4$ and clearly
 445 deviated from that of $L = 6$ despite the larger angular bins and the lower statistics compared
 446 to a previous measurement [32]. We therefore assign $L = 4$ for the resonance observed. Since
 447 the initial and final states involve spinless particles only, the spin-parity of the corresponding
 448 resonant state in ^{10}Be is assigned as 4^+ . The present results thus confirm the resonance of
 449 the 4^+ state at $E_x = 10.15$ MeV in ^{10}Be observed in a previous measurement of elastic
 450 $^6\text{He} + \alpha$ scattering [32], and rule out the spin-parity of 3^- claimed in an angular correlation
 451 study using the $^7\text{Li} + ^6,7\text{Li}$ reactions [30]. The angle-integrated cross section ($\sigma_{4\pi}$) can be
 452 translated into the partial α decay width (Γ_α) via the relation [52]:

$$\sigma_{4\pi} = (2L + 1) \frac{\pi}{k^2} \frac{\Gamma_\alpha^2}{(E_{\text{c.m.}}^0 - E_{\text{c.m.}})^2 + (\Gamma/2)^2}, \quad (3)$$

453 with k being the wavenumber, $E_{\text{c.m.}}^0$ the resonance energy, and Γ the total decay width. The
 454 total width of 296(15) keV measured in a previous study [30] was adopted for Γ . $\sigma_{4\pi}$ was
 455 estimated from the cross section of the resonance in the angular range $\theta_{\text{c.m.}} = 65^\circ - 135^\circ$ in
 456 Fig. 7(a) given the $P_4(\cos \theta_{\text{c.m.}})^2$ dependence of the angular distribution. The resulting ratio
 457 of Γ_α/Γ is 0.49(5), which corresponds to $\Gamma_\alpha = 145(15)$ keV. This Γ_α/Γ ratio agrees with
 458 0.45(3) from the previous study [32].

TABLE I. Best-fit values of the optical model potential parameters and the $\langle {}^6\text{He}|{}^4\text{He} + 2n\rangle$ overlap spectroscopic factor. See text for details.

$E_{c.m.}$ (MeV)	N_R	$W(r)$ (MeV)	r_W (fm)	a_W (fm)	$SA_{g.s.}$
5.82	1.0	60.0	1.26	0.27	0.80
5.45	1.0	60.0	1.27	0.25	0.80
5.06	1.0	60.0	1.27	0.25	0.80
4.64	1.0	60.0	1.27	0.25	0.80
4.21	1.0	60.0	1.27	0.25	0.80
3.76	1.0	60.0	1.28	0.25	0.80
3.26	1.0	65.0	1.28	0.32	0.80
2.69	1.0	65.0	1.38	0.20	0.80

459 To assess the contribution of direct reactions, coupled reaction channels (CRC) calcula-
460 tions were performed using the code FRESKO [53]. These calculations are similar to those of
461 Khoa and von Oertzen [40], although some simplifications were made in light of their results.
462 We adopted the same basic model as Ref. [40] for the optical potentials, i.e. a double-folded
463 real potential and a Woods-Saxon imaginary potential, giving a potential of the form:

$$U(r) = N_R V_{DF}(r) + iW(r) + V_C(r), \quad (4)$$

464 where $V_C(r)$ is the usual Coulomb potential with radius $R_C = 1.25(A_1^{1/3} + A_2^{1/3})$ in fm,
465 $V_{DF}(r)$ is the double-folded real potential with normalization parameter N_R and $W(r)$ is the
466 imaginary potential of depth W in MeV, radius $R_W = r_W(A_1^{1/3} + A_2^{1/3})$ in fm and diffuseness
467 a_W in fm. The parameters N_R , $W(r)$, r_W and a_W were adjusted to obtain the best agreement
468 with the data at each energy. The double-folding potentials were calculated with the code
469 DFPOT [54] using the M3Y effective interaction [55]. The ${}^4\text{He}$ nuclear-matter density was
470 derived from the three-parameter Fermi distribution charge density of Ref. [56] by unfolding
471 the proton charge distribution as described in Ref. [57], assuming that $\rho_n = (N/Z)\rho_p$, and
472 the ${}^6\text{He}$ nuclear-matter density was the FC6 model density of Ref. [58]. Inelastic excitation
473 of the 1.8-MeV 2^+ state of ${}^6\text{He}$ was included in a similar way as detailed in Ref. [40]. The
474 real part of the inelastic scattering form factor was calculated using the double-folding
475 model, the proton and neutron parts of the transition density being calculated using the

476 Tassie Model [59], and the proton and neutron deformation lengths taken from Ref. [40].
 477 The imaginary part of the inelastic form factor was obtained by deforming the Woods-
 478 Saxon imaginary potential using the isoscalar deformation length of Ref. [40]. Following the
 479 conclusions of Ref. [40], we adopted the simplifying assumption of a pure $(1p_{3/2})^2$, S -wave
 480 ($J = L = 0$) configuration for the two neutrons in the ${}^6\text{He}$ ground state, where J is the
 481 internal angular momentum of the $2n$ cluster and L its orbital angular momentum with
 482 respect to the ${}^4\text{He}$ core. The $\langle {}^6\text{He} | {}^4\text{He} + 2n \rangle$ overlaps were calculated using the Bayman-
 483 Kallio method [60] and the spectroscopic amplitude was adjusted to give the best description
 484 of the data. Also following the conclusions of Ref. [40], the two-step sequential (${}^6\text{He}$, ${}^5\text{He}$; ${}^5\text{He}$,
 485 ${}^4\text{He}$) and indirect (${}^6\text{He}$, ${}^6\text{He}_{2+}^*$; ${}^6\text{He}_{2+}^*$, ${}^4\text{He}$) transfer paths were omitted as having negligible
 486 influence at the incident energies concerned. Test calculations for the $E_{\text{c.m.}} = 5.82$ MeV data
 487 supported this conclusion. Values of N_{R} , $W(r)$, r_W , a_W and the spectroscopic amplitude for
 488 the $\langle {}^6\text{He} | {}^4\text{He} + 2n \rangle$ overlap $\text{SA}_{\text{g.s.}}$ are given in Table I. The best-fit value of $\text{SA}_{\text{g.s.}}$ is somewhat
 489 smaller than the expected value of about 1.25 quoted in Khoa and von Oertzen [40], but is
 490 close to 0.85, the smallest empirical value they obtained.

491 The calculated differential cross sections of elastic scattering are compared to the exper-
 492 imental data in Fig. 8. To display the variation of the angular distribution as a function of
 493 energy, the CRC calculations for the previous energy bins are shown together by the dashed
 494 lines. Except for the last energy bin at 2.7 MeV, the calculated cross sections reproduce the
 495 data. The calculated cross sections vary smoothly as a function of $E_{\text{c.m.}}$, while the reaction
 496 parameters are nearly constant. At 2.7 MeV, the cross sections are suddenly enhanced and
 497 the radius and diffuseness parameters of the imaginary potential drastically change. This
 498 result indicates that elastic scattering predominantly proceeds via direct reactions outside
 499 of the region near $E_{\text{c.m.}} = 2.7$ MeV. It supports the conjecture made from the excitation
 500 functions that there is no sizable resonance in the region from 9.4 to 13.4 MeV in E_{x} of ${}^{10}\text{Be}$
 501 except for the 4^+ state at 10.15 MeV.

502 The resonance of another 4^+ state at 11.76 MeV, therefore, was not identified at the
 503 corresponding energy of $E_{\text{c.m.}} = 4.4$ MeV. The upper limit of the partial decay width was
 504 estimated by searching the limit where the resonance can be identified as a peak over the
 505 contribution of non-resonant scattering in the spectrum. The resulting value is $\Gamma_{\alpha}/\Gamma < 0.15$,
 506 or $\Gamma_{\alpha} < 20$ keV for the known Γ value of 121(10) keV [51].

507 The differential cross sections of inelastic ${}^6\text{He} + \alpha$ scattering to the ${}^6\text{He}$ 2^+ state are

508 shown in Fig. 9 together with the CRC calculations denoted by the solid lines. The CRC
 509 calculations for the magnitude of the cross sections are in good agreement with the experi-
 510 mental data, suggesting the validity of the framework of reaction analysis and the adopted
 511 optical-model potential parameters. The diffractive pattern is slightly out of phase with
 512 respect to the CRC predictions in the energy bin of $E_{\text{c.m.}} = 4.2$ MeV, where a resonance is
 513 inferred from the excitation function spectra. The experimental angular distribution peaks
 514 at 90° and symmetrically decreases towards 60° and 120° . This is characteristic of an angular
 515 momentum of $L = 2$.

516 B. Two-neutron emission channel

517 The angle-integrated excitation function was deduced for the ${}^6\text{He}(\alpha, 2n){}^8\text{Be}$ channel.
 518 The reaction vertex was converted to reaction energy E_{reac} after correcting for the energy
 519 loss using a program based on the SRIM code [47]. The cross section of the reaction as
 520 a function of center-of-mass energy $E_{\text{c.m.}}$ is shown in Fig. 10. Note that above 3 MeV,
 521 the spectrum includes also the contribution from the ${}^8\text{Be}$ 2^+ state. The contribution of
 522 the ground state, however, more likely dominates as the $z_{\text{reac}}-Q_{\text{total}}$ data (Fig. 6(a)) agree
 523 well with the simulation for the ground state (Fig. 6(b)). These two components would be
 524 differentiated by analyzing the correlations between two α particles that were not measured
 525 in this study. The vertical error bars are statistical only, while the horizontal error bars
 526 denote the sum of statistical and systematic errors. The latter comes from the ambiguity in
 527 locating the reaction vertex which is estimated to be ± 10 mm. The cross section increases
 528 from the reaction threshold of the ${}^6\text{He}(\alpha, 2n){}^8\text{Be}$ reaction at 1.06 MeV toward higher energies.
 529 It is clear that no significant resonance strength exists in the energy region covered in
 530 the present study. There is a minor peak structure at 2.5 MeV. It is difficult to judge
 531 whether this is due to a resonance or to a statistical fluctuation, particularly without the
 532 corresponding information from an angular distribution. In the former case, it would be
 533 possible that it originates from the 10.15-MeV 4^+ state with nearly the same resonance
 534 energy. A fit with a Voigt function [50] was made to estimate the possible partial width
 535 Γ_{sBe} . Three different backgrounds, namely linear, quadratic, and exponential functions, were
 536 tested. The resonance energy was set to the result from the elastic channel (2.56 MeV) and
 537 varied within the error (0.15 MeV), while the experimental resolution was fixed to 0.25 MeV

538 rms, which arises from the uncertainty in reaction energy (0.1 MeV) and that in vertex
 539 determination (0.2 MeV). The resulting $\Gamma_{s_{\text{Be}}}/\Gamma$ value is 0.09(5) and this gives an upper
 540 limit of $\Gamma_{s_{\text{Be}}}/\Gamma \sim 0.15$ for this possible decay branch.

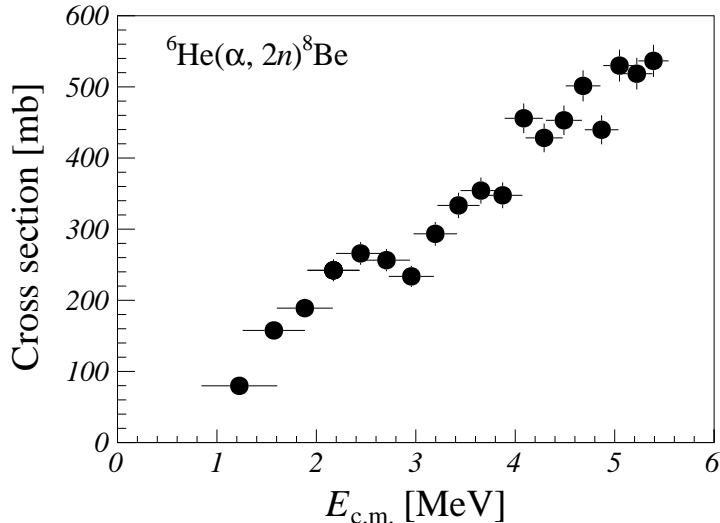


FIG. 10. Angle-integrated excitation function for the ${}^6\text{He}(\alpha, 2n){}^8\text{Be}$ channel.

542 V. DISCUSSION

543 The present study identified a 4^+ state with a large α decay width $\Gamma_{\alpha}/\Gamma = 0.49(5)$ at
 544 9.98(15) MeV in ${}^{10}\text{Be}$. The observed state most likely corresponds to the known 4^+ level
 545 at 10.15(2) MeV [31, 32] given the observed excitation energy and spin-parity. In previous
 546 studies [24, 32, 33], this state is considered the 4^+ member of a rotational band built on the
 547 second 0^+ state at 6.1793(7) MeV [51]. The excitation energies of ${}^{10}\text{Be}$ states are plotted
 548 against $J(J+1)$ in Fig. 11. The linear extrapolation from the 0_2^+ state and the 2^+ state
 549 at 9.560(20) MeV [51] indeed nicely agrees with the 10.15-MeV state in energy. The large
 550 moment of inertia from the narrow level spacing of the band members is well explained by
 551 the σ -type molecular orbital structure from both cluster-model calculations [21, 22, 25] and
 552 microscopic calculations based on the antisymmetric molecular dynamics (AMD) method
 553 [15, 24, 26]. In this picture, the valence neutrons are delocalized over the two clusterized α
 554 cores and the extension along the α cores' axis gives strong deformative characteristics to
 555 ${}^{10}\text{Be}$. The large decay width for α emission indicates a high degree of clusterization in this

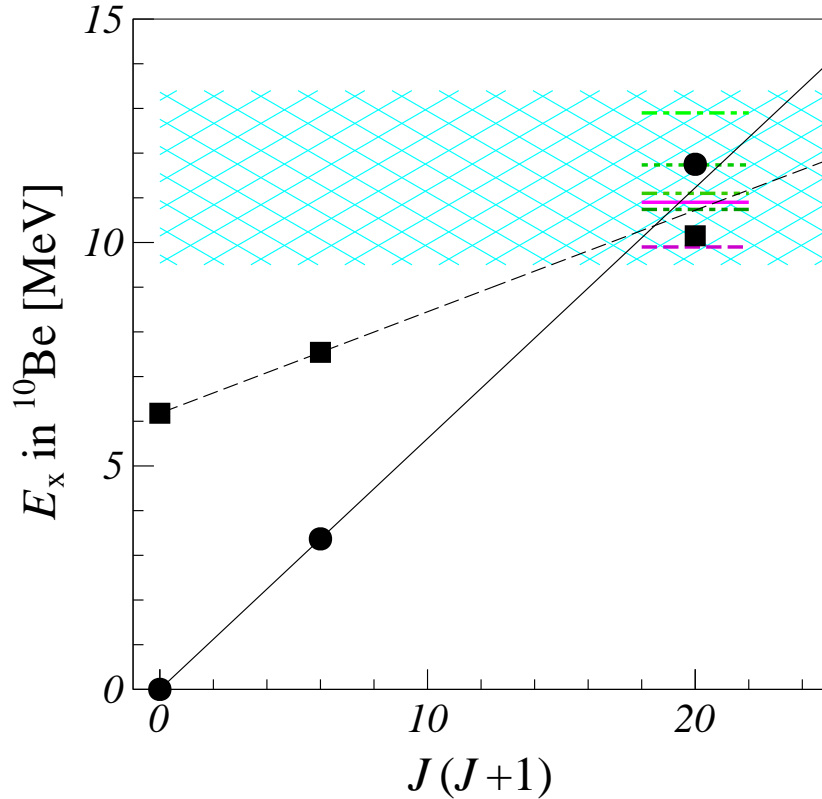


FIG. 11. (Color online) Plot of E_x vs. $J(J+1)$ for ^{10}Be . The band members of the ground and the second 0^+ states are shown by the circles and squares, respectively. The linear extrapolation using the 0^+ and 2^+ states is shown for each band. The horizontal lines at $J = 4$ denote predicted level energies of the 4^+ member of the ground state band from the β - γ constraint AMD method [24] (solid line), the variational AMD method [15] (dashed line), the four-body cluster model [21] (dotted line), the molecular orbital model [16] (dot-dashed line), the semi-microscopic algebraic cluster model [18], (double-dot-dashed line), and the multi-cluster generator coordinate method [19], (triple-dot-dashed line). The data of Refs.[16, 21] were obtained from the calculated values with respect to the threshold energy of $2\alpha + 2n$ at 8.386 MeV. The shaded area denotes the energy domain covered by the present study.

556 4^+ state and supports this type of cluster structure. An α spectroscopic factor of 3.1(2) is
557 estimated in a recent analysis of the measured partial width [61]. This value is as large
558 as the spectroscopic factors of about 1.5 for the ground-state band members of ^8Be with
559 well-developed two α clusters [61, 62].

560 In addition to the 0_2^+ state, theoretical studies [15, 16, 22, 24] predict a π -type cluster
 561 structure for the 0^+ ground state, in which valence neutrons are extending perpendicular
 562 to the axis of the two α cores. Given the 2^+ state at 3.37 MeV, the 4^+ state of the $0_{\text{g.s.}}^+$
 563 band is anticipated at around 11 MeV as seen in the linear extrapolation shown in Fig. 11.
 564 In previous studies [24, 33], the 4^+ state at 11.76(2) MeV is considered the most likely
 565 candidate for the 4^+ member of the $0_{\text{g.s.}}^+$ band because of its excitation energy and spin-parity.
 566 In the present study, however, there was no resonance observed around $E_x = 11.8$ MeV
 567 ($E_{\text{c.m.}} = 4.4$ MeV). This is in stark contrast with the significant resonance strength of the
 568 4^+ state of the 0_2^+ band at 10.2 MeV. The α decay width of the 11.8-MeV state is estimated
 569 less than 20 keV and is much smaller than $\Gamma_\alpha = 145(15)$ keV deduced for the 10.2-MeV
 570 state. Such a difference is unexpected as both 4^+ states belong to the rotational bands of
 571 the clusterized 0^+ states. Nearly the same spectroscopic amplitudes of ${}^6\text{He} + \alpha$ are predicted
 572 for these 4^+ states in the microscopic $2\alpha + 2n$ four-cluster-model [21]. The present result
 573 does not agree with this prediction. The small spectroscopic amplitude of the 4^+ member is
 574 also unlike the ground state 0^+ band of ${}^8\text{Be}$, despite what appears to be a similar moment
 575 of inertia. The α spectroscopic factors are predicted to be equally large in all 0^+ , 2^+ , and
 576 4^+ states in ${}^8\text{Be}$ [62], which is supported by the folding potential model that well describes
 577 the level energies and widths of these states [63].

578 There are two possible scenarios to account for the hindered strength of the 4^+ member
 579 of the $0_{\text{g.s.}}^+$ band. First is the possibility that the 4^+ state at 11.8 MeV does not belong to the
 580 $0_{\text{g.s.}}^+$ band, and the real band member exists outside the energy window of the present study
 581 ($E_{\text{c.m.}} = 2\text{--}6$ MeV or $E_x = 9.4\text{--}13.4$ MeV). This scenario implies an unusual level spacing
 582 for the ground state band. On the contrary, regardless of the framework, most theoretical
 583 studies [15, 16, 18, 19, 21, 24] predict the 4^+ state of the $0_{\text{g.s.}}^+$ band in the energy range
 584 $E_x = 10\text{--}13$ MeV (Fig. 11), the region anticipated from the proportionality to $J(J + 1)$.
 585 The second scenario is that the 4^+ state does belong to the $0_{\text{g.s.}}^+$ band, but with a reduced
 586 degree of clusterization that hinders the decay branch for α emission. The weakening of
 587 clustering in the 4^+ state is pointed out by an early AMD study of ${}^{10}\text{Be}$ [15]. In this
 588 prediction, the 0^+ ground state of ${}^{10}\text{Be}$ is highly clusterized, as confirmed in another later
 589 AMD study [64]. However, the component of two- α cores in the rotational band members
 590 is gradually reduced as the total spin increases. The origin of the dissociation of α clusters
 591 is attributed to the nuclear spin-orbit force. A stronger spin-orbit force yields a smaller

592 amplitude of two α cores in the 2^+ and 4^+ states of the $0_{\text{g.s.}}^+$ band. This scenario naturally
 593 explains why the four cluster model of $2\alpha + 2n$ overpredicts the spectroscopic amplitude of
 594 ${}^6\text{He} + \alpha$. In the molecular model, α clusters are assumed as the basis of wave functions,
 595 and thus the model does not incorporate the effects of their dissociation. On the other hand,
 596 the AMD method adopts Slater determinants where the spatial part of each single-particle
 597 wave function is a Gaussian function, and does not require the assumption of inert cluster
 598 cores [65]. This allows one to study the formation and dissociation of α clusters within one
 599 framework. The absence of resonance strength of the 4^+ state at 11.8 MeV agrees more with
 600 the interpretation of AMD calculations, which predict the limits of α clustering in higher
 601 spin members of the $0_{\text{g.s.}}^+$ band due to the spin-orbit force.

602 It is known that the spin-orbit force dissociates α clusters in high-spin states of ${}^{20}\text{Ne}$ [66–
 603 68] or ${}^{44}\text{Ti}$ [67]. Such an effect is considered to be hindered in ${}^8\text{Be}$ as it is a lighter nucleus
 604 with a weaker spin-orbit force [67]. On the other hand, recent AMD studies on ${}^{12}\text{C}$ [64, 69],
 605 an isotone of ${}^{10}\text{Be}$, point out that the effect of the spin-orbit force is pronounced by the
 606 sub-shell closure at $N = 6$ and this explains the small degree of clusterization for its ground
 607 state. In ${}^{10}\text{Be}$, this effect might be more important for the $0_{\text{g.s.}}^+$ band, which primarily
 608 consists of p -shell configurations [15], than for the intruder 0_2^+ band. The present result for
 609 ${}^{10}\text{Be}$ implies a delicate competition between the persistence of the clusters in ${}^8\text{Be}$ and their
 610 dissociation by the spin-orbit force as the number of neutrons increases toward the neutron
 611 drip line.

612 Finally, we discuss the present results in comparison with theoretical calculations of res-
 613 onant α scattering of ${}^6\text{He}$ in the framework of the four-body cluster model [14, 23]. The
 614 study using the resonating-group method predicts drastic changes in the scattering phase
 615 shift of $L = 3$ at $E_{\text{c.m.}} = 3.7$ MeV and $L = 0$ around 4.5 MeV for elastic scattering [14]. The
 616 angular distribution at 3.7 MeV is predicted to show enhanced cross sections with an oscil-
 617 lation characteristic of $L = 3$. However, in the present measurement, sizable resonances are
 618 clearly absent in the corresponding energy region in the elastic channel. Inelastic scattering
 619 to the ${}^6\text{He}$ 2^+ state was studied in a more recent work based on the generalized two-center
 620 cluster approach, where the reaction of ${}^6\text{He} + \alpha$ and the structure of ${}^{10}\text{Be}$ are described in a
 621 unified manner [23]. Excitation functions for $L = 1, 2$ were calculated and a few resonances
 622 are proposed in the energy region of $E_{\text{c.m.}} = 2\text{--}7$ MeV. The possible resonance of $L = 2$
 623 inferred at 4.4 MeV in the present study is in line with the $L = 2$ resonance predicted at

624 4 MeV. It originates from a 0^+ state in ^{10}Be with a large amplitude of the $\alpha + {}^6\text{He}(2^+)$ com-
 625 ponent, thus having a sizable strength in the inelastic channel. Further theoretical studies,
 626 particularly on excitation functions for the elastic channel that are not presented in Ref. [23],
 627 might be useful to interpret the inferred resonance.

628 VI. CONCLUSION

629 Scattering of neutron-rich ${}^6\text{He}$ nuclei on α particles was studied at the *TwinSol* facil-
 630 ity [42] at the University of Notre Dame. A time projection chamber, PAT-TPC [43], using
 631 He:CO₂ 90:10 gas at atmospheric pressure was operated in the active-target mode to pro-
 632 vide a gaseous ${}^4\text{He}$ target and track the beam and reaction products traversing the target.
 633 This unique capability significantly lowered the threshold for the energy of helium nuclei,
 634 allowing a continuous measurement of the excitation functions over a wide energy range
 635 with the unambiguous reconstruction of angular distributions, which are difficult to obtain
 636 in conventional studies using non-active targets.

637 Excitation functions and angular distributions were thus measured for elastic scattering
 638 and inelastic scattering to the ${}^6\text{He } 2^+$ state below a center-of-mass energy of 6 MeV. The
 639 present system also allowed us to measure the excitation function of the ${}^6\text{He}(\alpha, 2n){}^8\text{Be}$
 640 channel, which requires the detection of low-energy α particles following the decay of un-
 641 bound ${}^8\text{Be}$. The resulting excitation function of elastic scattering shows a resonance at
 642 $E_{\text{c.m.}} = 2.56(15)$ MeV, or an excitation energy $E_x = 9.98(15)$ MeV in ^{10}Be . A spin-parity
 643 of 4^+ was assigned from the angular distribution. The partial α decay width was estimated
 644 to be $\Gamma_\alpha/\Gamma = 0.49(5)$. These results obtained from the unambiguously-identified recoiling
 645 α particles confirm the previous results obtained for the state at 10.15(2) MeV without
 646 differentiating the ${}^6\text{He}$ and α particles, but with considerably better statistics [32]. The
 647 assignment of 4^+ supports the claim of Ref. [32] that the 10.15-MeV state corresponds to a
 648 member of the 0_2^+ band.

649 The large partial width for α decay indicates a highly-developed α cluster structure in this
 650 4^+ state. It is consistent with most theoretical predictions proposing the σ -type molecular
 651 orbital structure for the 0_2^+ band. On the contrary, no resonance strength for another 4^+
 652 state was observed in this energy region, where the 4^+ member of the ground state 0^+ band
 653 is expected. Except for the region near $E_{\text{c.m.}} = 2.6$ MeV, the angular distributions vary

654 smoothly as a function of energy, which is well reproduced by coupled-reaction channels
655 calculations. The hindered resonance strength is incompatible with theoretical calculations
656 done in the framework of the four-body cluster model, which predict a sizable spectroscopic
657 amplitude of ${}^6\text{He} + \alpha$ due to a π -type molecular orbital structure [21]. However, it is in
658 line with the results from an AMD study where the α clusters of the 0^+ ground state are
659 gradually dissociated by the nuclear spin-orbit force as the spin increases to its 2^+ and 4^+
660 rotational band members [15]. The present results support the limits of clustering in ${}^{10}\text{Be}$
661 due to the spin degree-of-freedom, and calls for more detailed spectroscopy of individual
662 cluster states in ${}^{10}\text{Be}$ and related microscopic theoretical studies.

663 The excitation function of the inelastic channel shows a slight enhancement at $E_{\text{c.m.}} = 4.4$ MeV
664 that may be due to a resonance. An angular momentum of $L = 2$ is suggested from the
665 angular distribution. This resonance may be related to a 0^+ state with a large ${}^6\text{He}(2^+) + \alpha$
666 component, which is predicted around 4 MeV [23].

667 The angle-integrated excitation function was obtained for the ${}^6\text{He}(\alpha, 2n){}^8\text{Be}$ channel,
668 with no strong resonance observed in the region of $E_{\text{c.m.}} = 1\text{--}5$ MeV. While the cross sec-
669 tion is slightly enhanced around $E_{\text{c.m.}} = 2.5$ MeV, it is uncertain whether this is a statistical
670 fluctuation or a possible resonance originating from the 4^+ state at 10.15 MeV. As mea-
671 sured in this study, this 4^+ state has a large partial decay width of $\Gamma_\alpha/\Gamma = 0.49(5)$ to the
672 ${}^6\text{He}(0^+) + \alpha$ final state, which is considered to be an indication of α clustering. The result of
673 the ${}^6\text{He}(\alpha, 2n){}^8\text{Be}$ channel, in contrast, gives an upper limit of $\Gamma_{s_{\text{Be}}}/\Gamma \sim 0.15$ for the decay
674 to ${}^8\text{Be}$, a nucleus with a pronounced 2α structure. It will be interesting to see if the partial
675 width to such a three body system (${}^8\text{Be} + 2n$) is sensitive to various α -cluster structures
676 predicted for ${}^{10}\text{Be}$.

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- 686 [1] J.A. Wheeler, Phys. Rev. **52**, 1083 (1937).
- 687 [2] D.M. Dennison, Phys. Rev. **96**, 378 (1954).
- 688 [3] H. Morinaga, Phys. Rev. **101**, 254 (1956).
- 689 [4] F. Hoyle, Astrophys. J. Suppl. Ser. **1**, 121 (1954).
- 690 [5] C.W. Cook, W.A. Fowler, C.C. Lauritsen, and T. Lauritsen, Phys. Rev. **107**, 508 (1957).
- 691 [6] T. Wakasa *et al.*, Phys. Lett. B **653**, 173 (2007).
- 692 [7] A.H. Wuosmaa, R.R. Betts, B.B. Back, M. Freer, B.G. Glagola, Th. Happ, D.J. Henderson, P.
693 Wilt, and I.G. Bearden, Phys. Rev. Lett. **68**, 1295 (1992).
- 694 [8] A. Tohsaki, H. Horiuchi, P. Schuck, and G. Röpke, Phys. Rev. Lett. **87**, 192501 (2001).
- 695 [9] Y. Funaki, T. Yamada, H. Horiuchi, G. Röpke, P. Schuck, and A. Tohsaki, Phys. Rev. Lett.
696 **101**, 082502 (2008).
- 697 [10] T. Kawabata *et al.*, Phys. Lett. B **646**, 6 (2007).
- 698 [11] M. Seya, M. Kohno, and S. Nagata, Prog. Thor. Phys. **65**, 204 (1981).
- 699 [12] Y. Kanada-En'yo, H. Horiuchi, and A. Ono, Phys. Rev. C **52**, 628 (1995).
- 700 [13] W. von Oertzen, Z. Phys. A **354**, 37 (1996).
- 701 [14] K. Fujimura, D. Baye, P. Descouvemont, Y. Suzuki, and K. Varga, Phys. Rev. C **59**, 817
702 (1999).
- 703 [15] Y. Kanada-En'yo, H. Horiuchi, and A. Dote, Phys. Rev. C **60**, 064304 (1999).
- 704 [16] N. Itagaki and S. Okabe, Phys. Rev. C **61**, 044306 (2000).
- 705 [17] Y. Ogawa, K. Arai, Y. Suzuki, and K. Varga, Nucl. Phys. A **673**, 122 (2000).
- 706 [18] L. Hernández de la Peña, P.O. Hess, G. Lévai, and A. Algora, J. Phys. G **27**, 2019 (2001).
- 707 [19] P. Descouvemont, Nucl. Phys. A **699**, 463 (2002).
- 708 [20] N. Itagaki, S. Hirose, T. Otsuka, S. Okabe, and K. Ikeda, Phys. Rev. C **65**, 044302 (2002).
- 709 [21] K. Arai, Phys. Rev. C **69**, 014309 (2004).
- 710 [22] M. Ito, K. Kato, and K. Ikeda, Phys. Lett. B **588**, 43 (2004).
- 711 [23] M. Ito, Phys. Lett. B **636**, 293 (2006).
- 712 [24] T. Suhara and Y. Kanada-En'yo, Prog. Theo. Phys. **123**, 303 (2010).

- 713 [25] N. Itagaki, S. Okabe, and K. Ikeda, Phys. Rev. C **62**, 034301 (2000).
- 714 [26] Y. Kanada-En'yo and H. Horiuchi, Phys. Rev. C **68**, 014319 (2003).
- 715 [27] D. Suzuki *et al.*, Phys. Rev. Lett. **103**, 152503 (2009).
- 716 [28] D. Suzuki, Eur. Phys. J A **48**, 130 (2012).
- 717 [29] N. Soić *et al.* Europhys. Lett. **34**, 7 (1996).
- 718 [30] N. Curtis, D.D. Caussyn, N.R. Fletcher, F. Maréchal, N. Fay, and D. Robson, Phys. Rev. C
719 **64**, 044604 (2001).
- 720 [31] M. Milin *et al.*, Nucl. Phys. A **753**, 263 (2005).
- 721 [32] M. Freer *et al.*, Phys. Rev. Lett. **96**, 042501 (2006).
- 722 [33] H.G. Bohlen, T. Dorsch, Tz. Kokalova, W. von Oertzen, Ch. Schulz, and C. Wheldon, Phys.
723 Rev. C **75**, 054604 (2007).
- 724 [34] S. Ahmed *et al.*, Phys. Rev. C **69**, 024303 (2004).
- 725 [35] H. Yamaguchi *et al.*, Phys. Rev. C **83**, 034306 (2011).
- 726 [36] G.M. Ter-Akopian *et al.*, Phys. Lett. B **426**, 251 (1998).
- 727 [37] R. Raabe *et al.*, Phys. Lett. B **458**, 1 (1999).
- 728 [38] R. Raabe *et al.*, Phys. Rev. C **67**, 044602 (2003).
- 729 [39] K. Rusek and K.W. Kemper, Phys. Rev. C **61**, 034608 (2000).
- 730 [40] D.T. Khoa and W. von Oertzen, Phys. Lett. B **595**, 193 (2004).
- 731 [41] K.P. Artemov, M.S. Golovkov, V.Z. Goldberg, V.V. Pankratov, A.E. Pakhomov, I.N. Serikov,
732 and V.A. Timofeev, Sov. J. Nucl. Phys. **55**, 1460 (1992).
- 733 [42] F. Becchetti, M.Y. Lee, T.W. O'Donnell, D.A. Roberts, J.J. Kolata, L.O. Lamm, G. Rogachev,
734 V. Guimarães, P.A. DeYoung, and S. Vincent, Nucl. Instr. and Meth. Phys. Res., Sect. A **505**,
735 377 (2003).
- 736 [43] D. Suzuki *et al.*, Nucl. Instr. and Meth. Phys. Res., Sect. A **691**, 39 (2012).
- 737 [44] Y. Giomataris, Ph. Rebourgeard, J.P. Robert, and G. Charpak, Nucl. Instr. and Meth. Phys.
738 Res., Sect. A **376**, 29 (1996).
- 739 [45] P. Baron, D. Calvet, E. Delagnes, X. de la Broise, A. Delbart, F. Druillolle, E. Mazzucato,
740 E. Monmarthe, F. Pierre, and M. Zito, IEEE Transactions on Nuclear Science NS-55, 1744
741 (2008).
- 742 [46] Y. Kudenko, Representing the T2K Collaboration, Nucl. Instr. and Meth. Phys. Res., Sect. A
743 **598**, 289 (2009).

- 744 [47] J.F. Ziegler, M.D. Ziegler, and J.P. Biersack., Nucl. Instr. and Meth. Phys. Res., Sect. B **268**,
745 1818 (2010).
- 746 [48] G. Audi, A.H. Wapstra, C. Thibault, Nucl. Phys. A **729**, 337 (2003).
- 747 [49] D.R. Tilley, C.M. Cheves, J.L. Godwin, G.M. Hale, H.M. Hofmann, J.H. Kelley, C.G. Sheu,
748 and H.R. Weller, Nucl., Phys. A **708**, 3 (2002).
- 749 [50] W. Voigt, Münch. Ber., 603 (1912).
- 750 [51] D.R. Tilley, J.H. Kelley, J.L. Godwin, D.J. Milener, J.E. Purcell, C.G. Sheu, and H.R. Weller,
751 Nucl., Phys. A **745**, 155 (2004).
- 752 [52] A.M. Lane and R.G. Thomas, Phys. Mod. Phys. **30**, 257 (1958).
- 753 [53] I.J Thompson, Comput. Phys. Rep. **7**, 167 (1988).
- 754 [54] J. Cook, Comput. Phys. Commun. **25**, 125 (1982).
- 755 [55] G. Bertsch, J. Borysowicz, H. McManus, and W.G. Love, Nucl. Phys. A **284**, 399 (1977).
- 756 [56] J.S. McCarthy, I. Sick, and R.R. Whitney, Phys. Rev. C **15**, 1396 (1977).
- 757 [57] G.R. Satchler and W.G. Love, Phys. Rep. **55**, 183 (1979).
- 758 [58] J.S. Al-Khalili, J.A. Tostevin, and I.J. Thompson, Phys. Rev. C **54**, 1843 (1996).
- 759 [59] L.J. Tassie, Australian J. Phys. **9**, 407 (1956).
- 760 [60] B.F. Bayman and A. Kallio, Phys. Rev. **156**, 1121 (1967).
- 761 [61] H.T. Fortune and R. Sherr, Phys. Rev. C **84**, 024304 (2011).
- 762 [62] D. Kurath, Phys. Rev. C **7**, 1390 (1973).
- 763 [63] P. Mohr, H. Abele, V. Kölle, G. Staudt, H. Oberhummer, and H. Krauss, Z. Phys. A **349**,
764 339 (1994).
- 765 [64] N. Itagaki, S. Aoyama, S. Okabe, and K. Ikeda, Phys. Rev. C **70**, 054307 (2004).
- 766 [65] Y. Kanada-En'yo, Phys. Rev. Lett. **81**, 5291 (1998).
- 767 [66] T. Tomoda and A. Arima, Nucl. Phys. A **303**, 217 (1978).
- 768 [67] T. Yamada, Phys. Rev. C **42**, 1432 (1990).
- 769 [68] M. Kimura, Phys. Rev. C **69**, 044319 (2004).
- 770 [69] Y. Kanada-En'yo, Prog. Theo. Phys. **117**, 655 (2007).