

${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ cross section at low energies

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The flux of ${}^7\text{Be}$ and ${}^8\text{B}$ neutrinos from the Sun and the production of ${}^7\text{Li}$ via primordial nucleosynthesis depend on the rate of the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ reaction. In an extension of a previous study showing cross section data at 127–167 keV center-of-mass energy, the present work reports on a measurement of the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ cross section at 106 keV performed at Italy's Gran Sasso underground laboratory by the activation method. This energy is closer to the solar Gamow energy than ever reached before. The result is $\sigma = 0.567 \pm 0.029_{\text{stat}} \pm 0.016_{\text{sys}}$ nb. The data are compared with previous activation studies at high energy, and a recommended $S(0)$ value for all ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ activation studies, including the present work, is given.

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I. INTRODUCTION

The ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ and ${}^3\text{He}({}^3\text{He}, 2p){}^4\text{He}$ reactions compete in the proton-proton (p - p) chain of solar hydrogen burning. The ratio of their rates at the temperature of the solar center determines how much the ${}^7\text{Be}$ and ${}^8\text{B}$ branches of the p - p chain contribute to solar hydrogen burning. The ${}^3\text{He}({}^3\text{He}, 2p){}^4\text{He}$ cross section being comparatively well known [1], the predicted flux of solar neutrinos from ${}^7\text{Be}$ and ${}^8\text{B}$ decay [2] depends on the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ cross section: The 9% uncertainty in its extrapolation to the solar Gamow energy (23 keV) obtained in a global analysis [3] contributes 8% [4] to the uncertainty in the predicted fluxes for solar ${}^7\text{Be}$ and ${}^8\text{B}$ neutrinos, in both cases the major nuclear contribution to the total uncertainty. The flux of solar ${}^8\text{B}$ neutrinos has been measured in the SNO and SuperKamiokande neutrino detectors [5,6], with a total uncertainty as low as 3.5% [6]. The solar ${}^7\text{Be}$ neutrino flux is planned to be measured in the Borexino and KamLAND neutrino detectors.

The production of ${}^7\text{Li}$ in big-bang nucleosynthesis (BBN) is also highly sensitive to the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ cross section in the energy range $E \approx 160$ – 380 keV [7]. A recent compilation for the purpose of BBN adopts 8% uncertainty [8] for the cross section. Based on the baryon to photon ratio from observed anisotropies in the cosmic microwave background [9], nucleosynthesis network calculations predict primordial ${}^7\text{Li}$ abundances [10] that are significantly higher than observations of old stars [11,12]. Either a completely new interpretation of

the stellar abundance data (e.g., Ref. [13]) or a dramatically lower ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ cross section at relevant energies may explain this discrepancy.

Since the cross section of the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ reaction is of the order of attobarn at $E = 23$ keV, the cross-section data from experiments carried out at higher energies are parametrized by the astrophysical S factor $S(E)$ defined as

$$S(E) = \sigma(E)E \exp[2\pi\eta(E)],$$

where $2\pi\eta(E) = 164.12E^{-0.5}$ is the Sommerfeld parameter [14], and E the center-of-mass energy in keV. The S factor is then used to extrapolate the data to the low energies of astrophysical interest, and often its extrapolation to zero energy, $S(0)$, is quoted.

The ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ reaction has a Q value of 1.586 MeV [15], and at low energy it proceeds via radiative capture into the ground state and the first excited state of ${}^7\text{Be}$ (Fig. 1). The final ${}^7\text{Be}$ nucleus decays with a half-life of 53.22 ± 0.06 days to ${}^7\text{Li}$, emitting a 478-keV γ ray in $10.44 \pm 0.04\%$ of the cases [16]. The cross section can be measured by detecting either the induced ${}^7\text{Be}$ activity (activation method) or the prompt γ rays from the reaction (prompt- γ method).¹

¹An experiment based on a third method to measure the cross section, namely the detection of ${}^7\text{Be}$ nuclei in a recoil mass separator, is in progress at the ERNA facility [17].

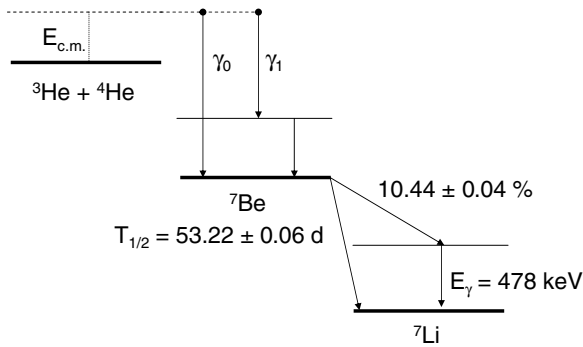


FIG. 1. Level diagram of the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ reaction ($Q = 1.586$ MeV) and the decay of ${}^7\text{Be}$.

Previous studies of the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ reaction [18–21] that used the activation technique cover the energy range $E = 420$ – 2000 keV and are briefly recalled here.

Osborne *et al.* [18] have measured the cross section by the activation technique at two energies, $E = 945$ and 1250 keV. A ${}^3\text{He}$ gas cell closed by a Ni + Cu window was bombarded by an α beam, and the activity of ${}^7\text{Be}$ implanted into a Ta catcher foil was measured with a Ge(Li) detector.

A similar experimental technique (gas cell with Ni window, Au catcher foil, and Ge(Li) detector) has been used by Robertson *et al.* [19]. The beam intensity was measured by current integration as well as by Rutherford backscattering (RBS) from a gold foil. An attempt was made to study the loss of ${}^7\text{Be}$ from the catcher, giving a 20% upper limit. The cross section was determined at $E = 987$ keV, with both direct (${}^3\text{He}$ gas cell and α beam) and inverse (${}^4\text{He}$ gas cell and ${}^3\text{He}$ beam) kinematics yielding consistent results.

Volk *et al.* [20] measured the energy-integrated cross section using a 0.8-bar ${}^3\text{He}$ gas cell in which the α beam stopped. The created ${}^7\text{Be}$ was collected onto an Al foil. The energy dependence of the cross section from a previous prompt- γ study was adopted to derive an $S(0)$ value.

Recently, Nara Singh *et al.* [21] carried out a precise activation experiment at $E = 420$ – 950 keV. A ${}^3\text{He}$ gas cell closed with a Ni window was bombarded with an α beam. The beam intensity was measured by both current integration and RBS. The produced ${}^7\text{Be}$ was collected on a Cu catcher and the activity was measured by a HPGe detector.

Cross-section measurements by the prompt γ -ray method [18,22–27] cover the energy range $E = 107$ – 2500 keV, although with limited precision at low energies. A global analysis of all available experimental data [3] indicates that S factor data obtained with the activation method are systematically 13% higher than the prompt- γ results.

Theoretical calculations reproduce the global shape of the S factor curve rather well (e.g., Refs. [28–30]). However, the slope of this curve has been questioned [29] for $E \leq 300$ keV, where there are no high-precision data.

The aim of the present activation study is to provide high-precision data at energies that are low enough to effectively constrain the extrapolation to solar energies and high enough to be relevant for big-bang nucleosynthesis. To study the solar interior [4,31,32], to investigate the low-energy slope

of the S factor curve [29], and to sharpen big-bang ${}^7\text{Li}$ abundance predictions [7,33], such precision ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ measurements have been recommended. In the present work, a new experimental cross-section number is reported at $E = 106$ keV, lower than ever before reached by direct experiment. In addition, cross-section data at $E = 127$ – 169 keV that have been published previously in abbreviated form [34] are presented with full detail here. A new $S(0)$ for the activation method based on all available experimental data is recommended.

II. EXPERIMENT

A. The accelerator

The Laboratory for Underground Nuclear Astrophysics (LUNA) [35] in Italy’s Gran Sasso underground laboratory (LNGS) has been designed for measuring small nuclear cross sections for astrophysical purposes. Its low laboratory background [36] has made it possible to study several reactions of astrophysical relevance [1,37–40].

The irradiations for the present study have been carried out at the 400-kV LUNA2 accelerator [41] at energies $E_\alpha = 250$, 300, 350, and 400 keV, with a typical current of $200 \mu\text{A}$ ${}^4\text{He}^+$. The beam energy is obtained with an uncertainty as low as 300 eV from a precision resistor chain calibrated through radiative-capture reactions, and it exhibits an energy spread of less than 100 eV [41].

The beam intensity is measured by using a beam calorimeter with constant temperature gradient similar to the one described previously [42], and a precision of 1.5% is obtained from the difference between the calorimeter power values with and without incident ion beam, by taking into account the calculated energy loss in the target gas [43]. The calorimeter has been calibrated at various beam energy and intensity values using the evacuated gas target chamber as a Faraday cup, with a proper secondary electron suppression voltage applied.

B. The gas target setup

The ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ reaction takes place in a differentially pumped windowless gas target (Fig. 2) filled with enriched ${}^3\text{He}$ gas (with an isotopic purity $>99.95\%$, a pressure of 0.7 mbar, and a corresponding target thickness of 8–10 keV). The exhaust from the first pumping stage (2050 m^3/h Roots pump) and the second pumping stage (three 1000 1/s turbomolecular pumps) is compressed by a 500 m^3/h Roots blower and an oil-free forepump, cleaned in a getter-based gas purifier and recirculated into the target. After passing the three pumping stages [the closest of which to the target being shown in Fig. 2(a)] and a connection pipe (b), the ion beam from the accelerator enters the target chamber (d) through an aperture of 7 mm diameter (c) and is finally stopped on a disk (h) of 70 mm diameter that serves as the primary catcher for the produced ${}^7\text{Be}$ and as the hot side of the beam calorimeter just described.

The pressure in the gas target chamber has been monitored continuously during the irradiations at two positions with

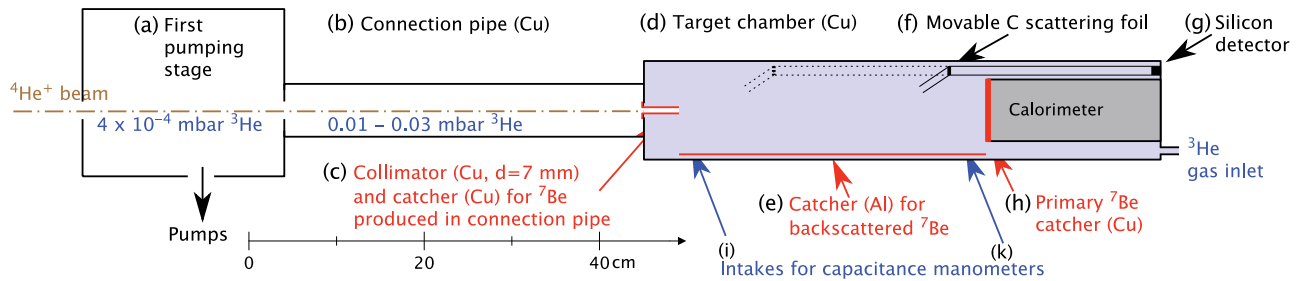


FIG. 2. (Color online) Schematic view of the target chamber used for the irradiations. See text for details.

capacitance manometers [Figs. 2(i)–2(k)]. The pressure and temperature profiles (Fig. 3) have been measured without the ion beam in a chamber of the same dimensions as the actual gas target chamber but with several ports along the beam path for pressure and temperature sensors. The pressure has been found to be equal to better than 0.25% at the different positions. The temperature profile has been observed to vary monotonically between the water-cooled collimator (15°C) and the hot side of the calorimeter (67°C). Linear interpolations have been used to calculate pressure and temperature between the measured positions. To reflect the uncertainty from the linear interpolation, a relative uncertainty of 13% has been assigned to the part of the target thickness contained in the 7 mm collimator (which comprises 5% of the total target thickness and where the pressure drop is significant), resulting in 0.7% uncertainty for the total target thickness. Combining this uncertainty with the 0.25% manometer precision and with the 0.3% uncertainty from the temperature measurement gives a precision of 0.8% for the target thickness without ion beam.

The thinning of the target gas through the beam heating effect [44] and the fraction of gases other than ^3He have been measured to obtain the effective target thickness. For this purpose, a 100- μm -thick silicon detector [Fig. 2(g)] detects projectiles that have been elastically scattered first in the target gas and subsequently in a movable 15 $\mu\text{g}/\text{cm}^2$ carbon foil

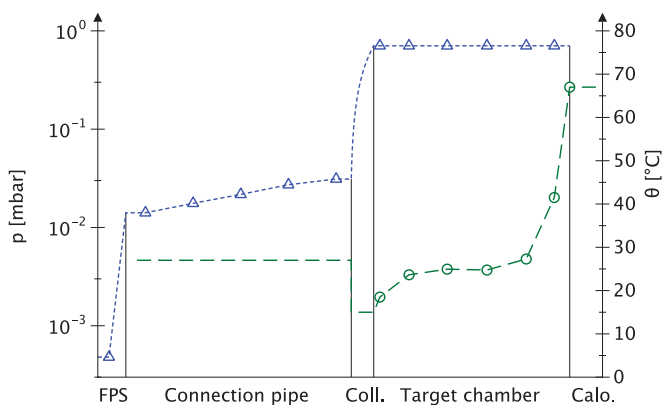


FIG. 3. (Color online) Measured pressure (p , blue triangles) and temperature (θ , green circles) profile inside the target chamber and adjacent regions: first pumping stage (FPS), connection pipe, collimator (Coll.), target chamber, and calorimeter (Calo.). The dashed lines indicate the interpolated profile adopted where there are no data.

(f). The beam heating effect has thus been investigated at several positions along the chamber in a wide beam energy and intensity range, and the average corrections shown in Table III were found. The amount of contaminant gases (mainly nitrogen) is monitored with the silicon detector during the irradiations (Fig. 4), kept below $1.0 \pm 0.1\%$ and corrected for in the analysis. Further details of the elastic scattering measurements are described elsewhere [45,46].

C. Sample irradiation

The catchers are irradiated with charges of 60–200 C, accumulating ^7Be activities of 0.03–0.6 Bq. Table I shows details of the irradiations. Calculations for the straggling of the ^4He beam and of the produced ^7Be nuclei in the ^3He gas and for the emission cone of ^7Be (opening angle 1.8° – 2.1°) have been carried out and show that 99.8% of the ^7Be produced inside the target chamber, including the 7 mm collimator, reaches the primary catcher.

D. Offline ^7Be counting

After the irradiation, the catcher is dismantled and counted subsequently with two 120% relative efficiency HPGe detectors called LNGS1 (Fig. 5) and LNGS2 (Fig. 6), both properly shielded with copper and lead, in the LNGS

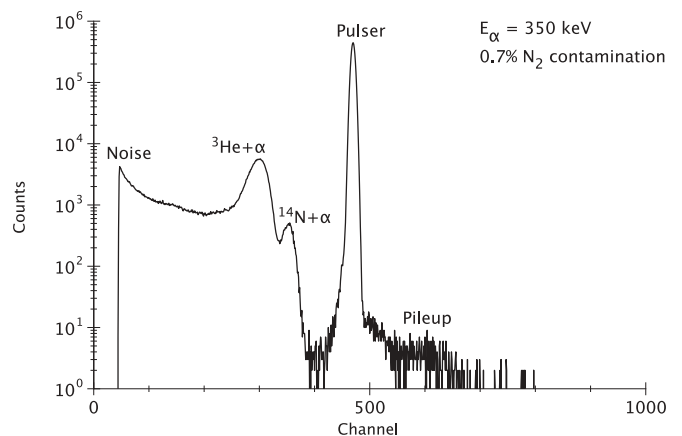


FIG. 4. Elastic scattering spectrum taken with the silicon detector at $E_\alpha = 350$ keV, showing a contamination of 0.7% N_2 in the ^3He gas.

TABLE I. Details of the irradiations. In all cases the target pressure was 0.7 mbar.

Sample	E_α (keV)	Target gas	Irradiation (days)	Charge (coulombs)	Average current (μ A)
D	249.8	^3He	6.5	83	149
B	298.8	^3He	10.5	215	237
A	348.4	^3He	9.5	203	248
F	398.2	^3He	2.9	63	250
E	400.2	^4He	6.5	104	187

underground counting facility [47]. Detector LNGS1 is additionally equipped with an antiradon box, and its laboratory background is two orders of magnitude lower than with equivalent shielding overground [47].

The samples have been counted in close geometry (e.g., in the case of detector LNGS1 the distance between sample and detector endcap was 5 mm). To obtain a precise efficiency calibration at this close distance, three homogeneous ^7Be sources of 200–800 Bq activity and 8 mm active diameter were prepared with the $^7\text{Li}(p, n)^7\text{Be}$ reaction. Thin layers of LiF ($\approx 20 \mu\text{g}/\text{cm}^2$) evaporated onto Ta backings and protected by evaporated gold layers ($\approx 5 \mu\text{g}/\text{cm}^2$) have been irradiated by 2.5-MeV protons from the ATOMKI Van de Graaff accelerator. The absolute activity of the calibration sources was subsequently determined in far geometry with two HPGe detectors at ATOMKI and with one HPGe detector, called LNGS3, at LNGS. The absolute efficiency of each of these three detectors has been determined by using a set of commercial γ -ray calibration sources. The three source kits used for calibrating the detectors were mutually independent. All three measurements gave consistent results, and the activities of the ^7Be calibration sources have been determined with a final uncertainty of 1.8%.

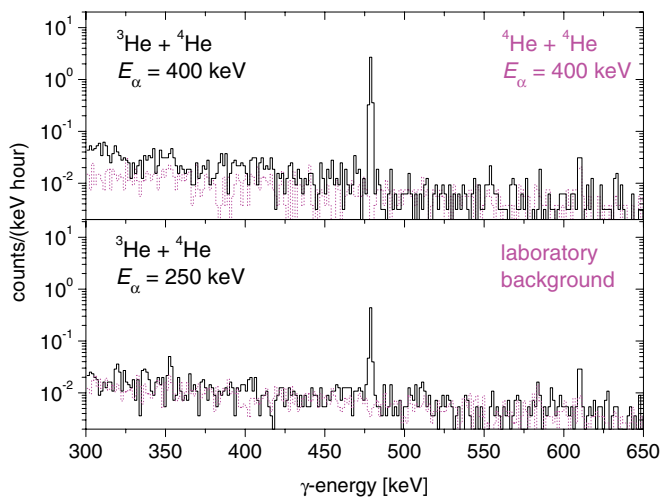


FIG. 5. (Color online) Offline γ -counting spectra from detector LNGS1. Solid black line: ^3He gas bombarded at $E_\alpha = 400$ keV (top panel, sample F) and 250 keV (bottom panel, sample D), respectively. Dotted red line, top panel: ^4He gas bombarded at $E_\alpha = 400$ keV (sample E). Dotted red line, bottom panel: laboratory background.

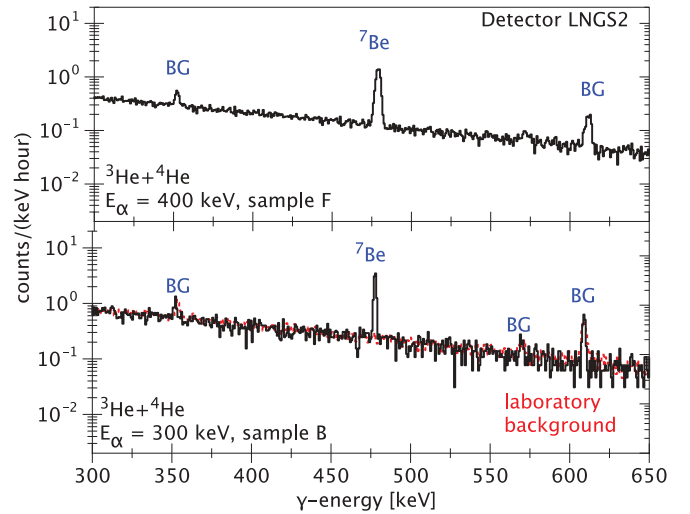


FIG. 6. (Color online) Offline γ -counting spectra from detector LNGS2. Solid black line: ^3He gas bombarded at $E_\alpha = 400$ keV (sample F) and 300 keV (sample B), respectively. Dotted red line, bottom panel: laboratory background.

The three ^7Be calibration sources were then used to calibrate detectors LNGS1 and LNGS2 at close geometry. Owing to the relatively low activities of the calibration sources, random coincidence summing effect and dead-time correction were negligible. In the case of detector LNGS2, which has a horizontal geometry, calibration sources and samples were placed horizontally in front of the detector, and not vertically on top of it as in the case of detector LNGS1. The impact of statistical submillimeter variations in the distance between source/sample and detector endcap resulting from the horizontal geometry of detector LNGS2 has been evaluated by moving the calibrated sources. An additional uncertainty of 1.2% resulting from this effect is included in the final statistical uncertainties given for detector LNGS2.

The ^7Be distribution in the catchers has been calculated from the ^7Be emission angle and straggling, and GEANT4 [48] simulations give $0.8 \pm 0.4\%$ to $1.5 \pm 0.4\%$ correction for the γ -ray efficiency because of the tail of the distribution at high radii.

Without aiming for high precision, we have determined the half-life of ^7Be in the Cu host material from the decay curve measured on the present, weak activated samples (Fig. 7). The weighted average of all measured samples gives a half-life of 52.2 ± 1.5 days, compatible with 53.22 ± 0.06 days from a recent compilation [16]. The value from Ref. [16] has been used for the data analysis.

The activity values referring to the end of the irradiations measured with the two HPGe detectors are in all cases in good agreement (Table II). For each sample, the weighted average of the activity values has been used for the data analysis.

E. Parasitic ^7Be production in the primary catcher

Oxygen-free high-conductivity (OFHC) copper has been studied as a possible material for the primary catcher. This material has good heat conductivity (required for the beam

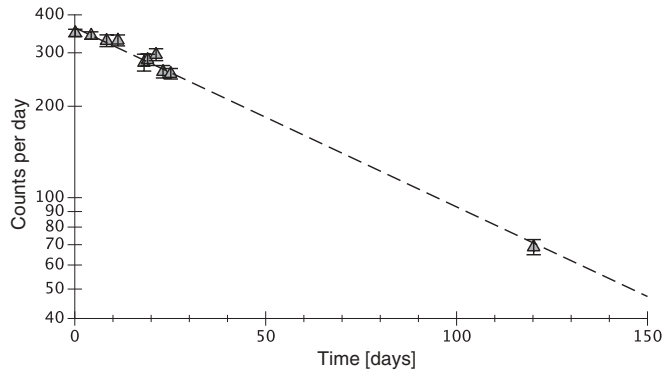


FIG. 7. Counting rate of sample A on detector LNGS2 as a function of time. The dashed line is an exponential fit to the data.

calorimeter described here), sustains high α doses without blistering, and has a relatively low charge number to limit backscattering of ${}^7\text{Be}$ nuclei. Since possible ${}^6\text{Li}$ or ${}^{10}\text{B}$ impurities in the catcher material can give rise to unwanted ${}^7\text{Be}$ production through the ${}^6\text{Li}(d, n){}^7\text{Be}$ and ${}^{10}\text{B}(p, \alpha){}^7\text{Be}$ reactions induced by traces of ${}^2\text{DH}_2^+$ or even ${}^2\text{D}_2^+$ in the ${}^4\text{He}^+$ beam, this material was studied in detail prior to adopting it for the main experiment.

Samples from the material to be used for the catcher were irradiated with 700-keV protons and deuterons at the ATOMKI Van de Graaff accelerator. After the irradiations, the γ activity of the samples has been observed, showing no γ peak from ${}^7\text{Be}$ decay. Based on the known cross section of the ${}^6\text{Li}(d, n){}^7\text{Be}$ [49–51] and ${}^{10}\text{B}(p, \alpha){}^7\text{Be}$ [52,53] reactions, an upper limit of 3 ppm has been determined for the concentration of both ${}^6\text{Li}$ and ${}^{10}\text{B}$. By varying the settings of the LUNA2 analyzing magnet and taking the isotopic abundance of deuterium into account, an order-of-magnitude upper limit of $10^{-7} d/\alpha$ has been obtained. By combining this fraction with the aforementioned upper limits for ${}^6\text{Li}$ and ${}^{10}\text{B}$ contaminations and the ${}^6\text{Li}(d, n){}^7\text{Be}$ and ${}^{10}\text{B}(p, \alpha){}^7\text{Be}$ cross sections at lower energy, the induced ${}^7\text{Be}$ activity from parasitic reactions is shown to be six orders of magnitude less than the activity from the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ reaction expected by using the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ cross section from Ref. [28].

The only excess activity detected on the samples irradiated at ATOMKI was ${}^{24}\text{Na}$ produced by the ${}^{23}\text{Na}(d, p){}^{24}\text{Na}$

TABLE II. Counting times τ and activities measured for the different catchers with the two HPGe detectors. Uncertainties are purely statistical; in the case of detector LNGS2 it also includes the 1.2% repositioning uncertainty discussed in the text.

Sample	LNGS1		LNGS2		Adopted activity (mBq)
	τ (days)	Activity (mBq)	τ (days)	Activity (mBq)	
D	16	25.3 ± 1.3	—	—	25.3 ± 1.3
B	12	208 ± 6	21	203 ± 6	205 ± 4
A	6	472 ± 14	22	495 ± 11	486 ± 9
F	10	319 ± 11	11	310 ± 8	313 ± 6
E	16	$<0.21 (2\sigma)$	—	—	$<0.21 (2\sigma)$

reaction. However, the half-life of ${}^{24}\text{Na}$ (15 h) is short compared with that of ${}^7\text{Be}$, so in the main experiment the offline γ -counting spectra taken immediately after the irradiation was concluded were compared with spectra taken several days later, when any possible ${}^{24}\text{Na}$ traces have decayed out. Thus, Compton background from the 2.754-MeV γ ray following the decay of ${}^{24}\text{Na}$ has been ruled out as a significant contributor of background in the offline γ counting.

Based on these considerations, OFHC copper was finally selected as material for the primary catcher. To rule out not only ${}^6\text{Li}(d, n){}^7\text{Be}$, ${}^{10}\text{B}(p, \alpha){}^7\text{Be}$, and ${}^{23}\text{Na}(d, p){}^{24}\text{Na}$ but any possible source of parasitic ${}^7\text{Be}$, during the main experiment for one catcher the enriched ${}^3\text{He}$ target gas was replaced with 0.7-mbar ${}^4\text{He}$. This catcher was then bombarded at the highest available energy of $E_\alpha = 400$ keV. Despite the high applied dose of 104 C, in 16 days counting time no ${}^7\text{Be}$ has been detected (Fig. 5, top panel, and Table II), establishing a 2σ upper limit of 0.1% for parasitic ${}^7\text{Be}$.

F. ${}^7\text{Be}$ losses

${}^7\text{Be}$ losses by backscattering from the primary catcher and by incomplete collection were studied experimentally at $E_\alpha = 400$ keV and with Monte Carlo simulations at 250, 300, 350, and 400 keV. For the backscattering study, parts of the inner surface of the chamber were covered by aluminum foil functioning as secondary catcher [Fig. 2(e)], and the foil sample was subsequently counted on detector LNGS1 (Fig. 8, upper panel). It was found that $1.3 \pm 0.5\%$ of the created ${}^7\text{Be}$ is lost from backscattering, consistent with the 1.5% value obtained in a GEANT4 [48] simulation using a SRIM-like multiple scattering process [54]. At lower energies, the simulation result of up to 2.9% was used as backscattering correction (Table III, column 7), with an adopted uncertainty of 0.5%. Incomplete ${}^7\text{Be}$ collection occurs since 3.5% of the

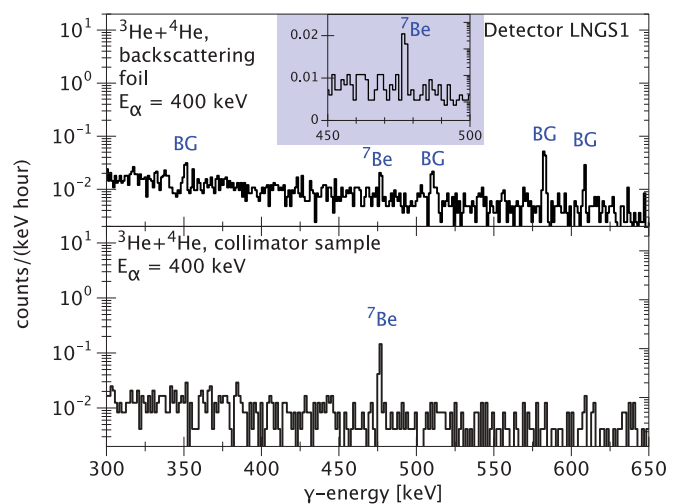


FIG. 8. (Color online) Offline γ -counting spectra from detector LNGS1. Collimator and backscattering foil counted to study ${}^7\text{Be}$ losses. The peaks marked with BG in the upper panel are background lines from impurities in the Al foil. The inset in the upper panel has a linear ordinate.

TABLE III. Experimental (exp) and calculated (calc) corrections for irradiation and collection for each run. Effective energy and cross section are given in the final two columns. Only the statistical uncertainty is given. See Table V for the adopted systematic uncertainty.

Run	E_α (keV)	Target gas	Corrections				E^{eff} (keV)	$\sigma(E^{\text{eff}})$ (10^{-9} b)
			Beam heating	Contaminant gases	Backscattering	Collection losses		
D	249.8	^3He	2.9%	0.5% _{exp}	2.9% _{calc}	2.4% _{calc}	105.6	0.567 ± 0.029
B	298.8	^3He	4.9%	0.3% _{exp}	2.2% _{calc}	2.3% _{calc}	126.5	1.87 ± 0.04
A	348.4	^3He	5.4%	0.3% _{exp}	1.8% _{calc}	2.2% _{calc}	147.7	4.61 ± 0.07
F	398.2	^3He	5.7%	1.0% _{calc}	1.3% _{exp}	2.6% _{exp}	168.9	9.35 ± 0.19

total ^3He target thickness is in the connecting pipe, and a part of the ^7Be created there does not reach the primary catcher but is instead implanted into the 7-mm collimator [Fig. 2(c)]. At $E_\alpha = 400$ keV, a modified collimator functioning as secondary catcher was used and counted on detector LNGS1 (Fig. 8, lower panel). A $2.6 \pm 0.4\%$ effect was observed, consistent with a simulation ($2.1 \pm 0.4\%$). For $E_\alpha = 250\text{--}350$ keV, incomplete ^7Be collection was corrected for based on the simulation (with up to a 2.4% correction and an adopted uncertainty of 0.5%).

Sputtering losses of ^7Be by the ^4He beam were simulated [43], showing that for the present beam energies sputtering is 10^4 times less likely than transporting the ^7Be even deeper into the catcher, so it has been neglected.

All Monte Carlo calculations mentioned in Secs. II D–II F have been carried on until a statistical uncertainty of 0.2% or better was reached; this value is negligible compared to the systematic uncertainties discussed in the appropriate section.

III. RESULTS

The effective center-of mass-energy E^{eff} has been calculated by assuming a constant S factor over the target length [14]. The uncertainties of 0.3 keV in E_α [41] and of 4.4% in the energy loss [43] lead to 0.16-keV uncertainty in E^{eff} and thus contribute 0.5% (at $E^{\text{eff}} = 169$ keV) to 1.1% (at $E^{\text{eff}} = 106$ keV) to the S factor uncertainty.

The effective energy and cross-section results for each sample are shown in the last two columns of Table III. The systematic uncertainties are summarized in Table IV, giving a total value of 3%. For the present low energies an electron screening enhancement factor f [55] of up to 1.016 has been calculated in the adiabatic limit, but this has not been corrected for (Table V).

The present data (Table V and Fig. 9) touch the energy range relevant to big-bang ^7Li production. Their uncertainty of at most 6% (systematic and statistical combined in quadrature)

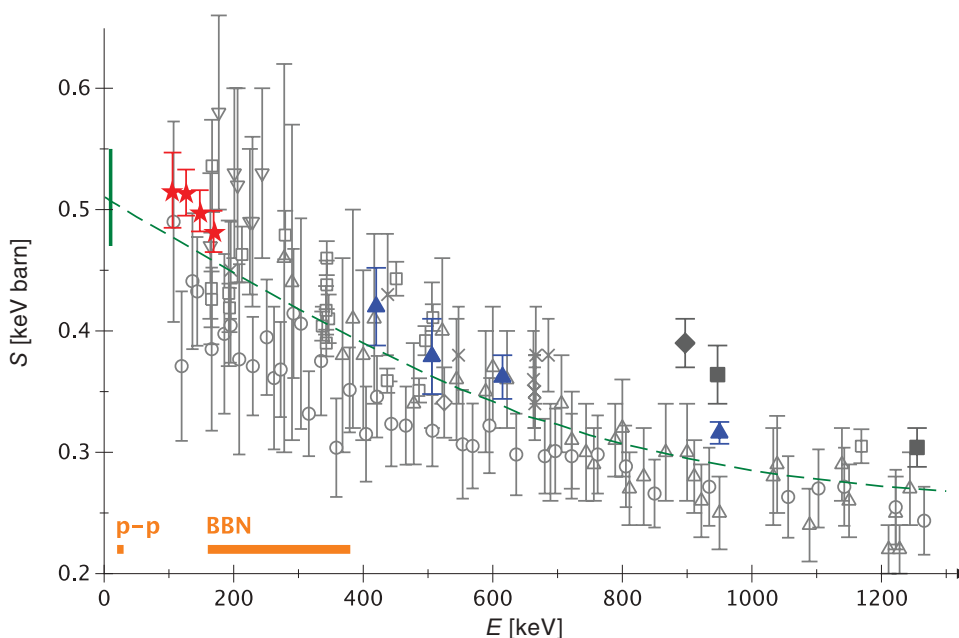


FIG. 9. (Color online) Astrophysical S factor for $^3\text{He}(\alpha, \gamma)^7\text{Be}$. Activation data: filled squares [18], filled diamonds [19], filled triangles [21], and stars (present work). Prompt- γ data: triangles [23], inverted triangles [24], circles [25] (renormalized by a factor 1.4 [27]), squares [18], diamonds [26], and crosses [27]. Dashed line: previously adopted R -matrix fit [8]. Horizontal bars: energies relevant for solar p - p hydrogen burning and for BBN.

TABLE IV. Systematic uncertainties in the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ astrophysical S factor. The uncertainty resulting from the effective energy affects only the S factor result, not the cross section.

Source	Uncertainty
${}^7\text{Be}$ counting efficiency	1.8%
Beam intensity	1.5%
Beam heating effect	1.3%
Effective energy	0.5–1.1%
Target pressure and temperature without beam	0.8%
Incomplete ${}^7\text{Be}$ collection	0.5%
${}^7\text{Be}$ backscattering	0.5%
${}^7\text{Be}$ distribution in catcher	0.4%
478-keV γ -ray branching [16]	0.4%
${}^7\text{Be}$ half-life [16]	0.1%
N_2 contamination in target gas	0.1%
Parasitic ${}^7\text{Be}$ production	0.1%
Total:	3.0–3.1%

is comparable to or lower than previous activation studies at high energy and lower than prompt- γ studies at comparable energy.

TABLE V. Cross-section and S factor results, relative uncertainties, and electron screening [55] enhancement factors f .

E^{eff} (keV)	$\sigma(E^{\text{eff}})$ (10^{-9} b)	$S(E^{\text{eff}})$ (keV b)	$\Delta S/S$		f
			stat.	syst.	
105.6	0.567	0.516	5.2%	3.1%	1.016
126.5	1.87 ^a	0.514	2.0%	3.0%	1.012
147.7	4.61 ^a	0.499	1.7%	3.0% ^b	1.009
168.9	9.35 ^a	0.482	2.0%	3.0% ^b	1.008

^aCross section previously published in abbreviated form in Ref. [34].

^bSystematic uncertainty 0.1% higher than the one given in Ref. [34]. The conclusions of Ref. [34] are unaffected.

TABLE VI. Extrapolated S factor $S(0)$ from activation studies of ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$.

	Ref.	$S(0)$ (keV b)
Osborne <i>et al.</i>	[18]	0.535 ± 0.040
Robertson <i>et al.</i>	[19]	0.63 ± 0.04
Volk <i>et al.</i>	[20]	0.56 ± 0.03
Nara Singh <i>et al.</i>	[21]	0.53 ± 0.02
present work		0.547 ± 0.017
Weighted average, all activation studies		0.553 ± 0.012
Weighted average, all prompt- γ studies	[3]	0.507 ± 0.016

IV. CONCLUSION

To obtain a recommended $S(0)$ value for the activation method, following Ref. [3] it is instructive to list the extrapolated $S(0)$ values for the different activation studies together with their quoted uncertainty (Table VI). For the present data, by adopting the curve shape from Ref. [8] an extrapolated $S(0) = 0.547 \pm 0.017$ keV b is obtained. The weighted average of all activation studies, including the present work, is found to be 0.553 ± 0.012 keV b, significantly higher than the weighted average of all prompt- γ studies, 0.507 ± 0.016 keV b [3].

With the addition of the new data, the systematic difference in normalization between prompt- γ and activation studies of ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ is now smaller than in Ref. [3]. However, it is still significant and much larger than the uncertainty required to match, for example, the 3.5% precision of the solar ${}^8\text{B}$ neutrino data [6].

In conclusion, prompt- γ experiments with precision comparable to the present activation data are called for to verify the normalization of the prompt- γ data.

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