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Charge-state distribution of Li ions from the β decay of laser-trapped ⁶He atoms

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The accurate determination of atomic final states following nuclear β decay plays an important role in several experiments. In particular, the charge state distributions of ions following nuclear β decay are important for determinations of the $\beta - \nu$ angular correlation with improved precision. Beyond the hydrogenic cases, the decay of neutral ⁶He presents the simplest case. Our measurement aims at providing benchmarks to test theoretical calculations. The kinematics of Li^{*n*+} ions produced following the β decay of ⁶He within an electric field were measured using ⁶He atoms in the metastable (1s2s, ³S₁) and in the (1s2p, ³P₂) states confined by a magneto-optical trap. The electron shake-off probabilities were deduced including their dependence on ion energy. We find significant discrepancies on the fractions of Li ions in the different charge states with respect to a recent calculation.

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I. INTRODUCTION

Atomic and molecular degrees of freedom can play an important role in precision nuclear beta-decay experiments. In nuclear beta decays, two energetic leptons (an electron and an anti-neutrino) are emitted, while the daughter nucleus recoils. The nucleus is usually in an atom or molecule, and the sudden change of its charge and its recoiling motion may cause electron excitations, shake-offs and molecular excitations. The final state of the recoil ion affects the shape of the β -energy spectrum. For example, in measurements of the β -energy spectrum near the end-point from molecular tritium, the final electronic state distribution [1-4] can affect the determination of the mass of anti-neutrinos. The helicity properties of the weak interaction imply correlations between the momenta of the outgoing particles [5]. Thus, precise measurements of the $\beta - \nu$ angular correlation can be used to search for new interactions [6–9]. In such experiments the $\beta - \nu$ angular correlation coefficient is deduced from the kinematics of the recoil ion, which can depend on the molecular binding[7, 8, 10].

The beta decay of ⁶He presents a good opportunity to determine the $\beta - \nu$ correlation. Because of the large endpoint and the relatively light mass of the nucleus, the $\beta - \nu$ correlation has a significant effect on the kinematics

of the charged particles from the decay. A measurement performed in 1963 [11] was one of several landmark experiments that determined the V - A nature of the weak interaction: the charged weak currents are of vector and axial vector type. Fundamental measurements of this kind have renewed interest in the context of searching for hints of new physics as deviations from the expectations based on the Standard Model [12]. An ongoing experiment is aiming at a measurement of the $\beta - \nu$ angular correlation in ⁶He decay with improved precision [13, 14]. In this experiment, the momentum of the recoil ion emitted from a cold and dilute cloud of laser-cooled 6 He atoms confined in a magneto-optical trap (MOT) [14] is determined through a full kinematics reconstruction in a strong electric field. Due to the sudden change in nuclear charge, the electrons do not always find the corresponding orbits in the Li atom and can be shaken off. Thus, the ⁶Li ion can have electric charges between +1 and +3. The fraction in a given charge state not only depends on the overlap of the initial electronic wave function and the final continuum states, but also on the ion energy, so a proper extraction of the $\beta - \nu$ correlation coefficient requires understanding the shake-off effect.

Quantitative comparisons of charge distributions have been presented for two heavier systems, ³⁵Ar [15] and ²¹Na [16]. The calculations in ²¹Na did not take into account a potentially important cancellation factor in the recoil-energy-dependent fraction that we describe below. The comparisons in ³⁵Ar showed agreement at the 1% level. As we indicate below the ⁶He system invites for higher precision comparisons, because the electronic wave functions for helium can be calculated with high accuracy. The case of ⁶He also presents a nice benchmark to test aspects of the calculations that are relevant for other

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problems, like the role of electron-electron interactions, the use of the sudden approximation, and methods for calculating charge distributions after the shake-off process. Several calculations have been performed for ⁶He [17–19]. A confirmation of the calculated fraction of Li^{3+} was recently performed with the hydrogen-like system of ⁶He ions [20]. Here, we address the case of the ⁶He neutral atom, which presents additional ingredients associated with the two electrons.

TABLE I: Comparison of calculated versus measured 6 Li ion charge fractions (in %) for 6 He decays from the ^{1}S atomic ground state.

Ion	Theory ^a [19]	Previous
		Experiment [21]
Li^+	88.99(2)	89.6(2)
Li^{2+}	9.7(1)	10.4(2)
Li^{3+}	1.2(1)	0.042(7)

^a In Ref. [19] the energy-dependent shake-off probabilities are modeled as $P = A + B \times E_{ion}$ and the values of parameters Aand B are presented for all charge states. We calculated the average ion energy $\langle E_{ion} \rangle$, and used it to calculate the theoretical energy-integrated charge state fraction. When calculating $\langle E_{ion} \rangle$, the $\beta - \nu$ correlation coefficient is assumed to be -1/3, and no β -energy threshold or directional restrictions on the emitted leptons are applied. In this case, $\langle E_{ion} \rangle = 0.723$ keV.

The charge distribution of Li ions from the decay of 6 He in its electronic ground state was measured by Carlson et al. [21]. Table I shows a comparison to the most recent calculation of Schulhoff and Drake [19]. As can be observed, there are significant discrepancies for Li-ion fractions in the different charge states. The main aim of the calculations in Ref. [19] was to study the dependence of charge distribution on the Li-ion energy, but a clear prediction is also given for the overall fractions in different charge states.

We report here the first measurement of the electron emission probabilities following the β decay of ⁶He atoms confined via a magneto-optical trap working between the metastable (1s2s, ³S₁) and the (1s2p, ³P₂) states. We present data with both the trapping lasers off (pure ³S₁) and on (approximate 50%/50% mixture of ³S₁ and ³P₂) and compare with the calculations of Schulhoff and Drake [19] assuming decay from the ³S₁ state.

II. EXPERIMENTAL METHOD

The ⁶He atoms were produced via the ⁷Li(d, t)⁶He reaction. Up to 2×10^{10} ⁶He atoms per second were produced by bombarding a lithium target with an 18 MeV deuteron beam, delivered by the tandem Van de Graaff accelerator at the University of Washington [22].

The ⁶He atoms were pumped into a RF-discharge tube where a fraction ($\sim 10^{-5}$) of the ⁶He atoms were brought to the ${}^{3}S_{1}$ metastable atomic state. The forward-going metastable atoms were then transversely-cooled, slowed by a Zeeman slower, and trapped in a magneto-optical trap, all based on resonant excitation of the ${}^{3}S_{1}$ to ${}^{3}P_{2}$ electronic transition via 1083 nm laser light. Due to the small efficiency for pumping ⁶He atoms to the metastable state there was a considerable amount of ground-state ⁶He in the chamber hosting the MOT. To reduce events from non-trapped ⁶He atoms, the atoms in the MOT were periodically pushed by a laser beam into a measurement chamber through a 5 mm-diameter 30-mm-long aperture tube for differential pumping and recaptured by a second MOT. This ⁶He trap contained over 2100 atoms on average.

During the measurement, the trapping lasers of this MOT were alternatively switched on and off with a 1 : 1 duty cycle and a period of 100 μ s. At the beginning of each off-cycle of 50 μ s, the ~50% fraction of the atoms previously excited to the ${}^{3}P_{2}$ level quickly decay back to the ${}^{3}S_{1}$ level within the 100 ns lifetime of the excited state. Time correlation of the decay events with the switching cycle thus allowed us to isolate decays from ⁶He purely in the ${}^{3}S_{1}$ state (laser off) from decays with an ~50% admixture of the ${}^{3}P_{2}$ excited state (laser on). Meanwhile, the fast switching provided sufficient confinement of the atom cloud.

We studied the charge-state distributions of the recoil ⁶Li ions by analyzing their time-of-flight (TOF) and energy spectra, which required detecting the β particle and the recoil ion in coincidence. The configuration of the detection chamber is shown in Fig. 1.

A β -telescope, consisting of a multi-wire proportional chamber (MWPC) and a scintillation detector, is placed above the trap. The scintillation detector measures the energy of the β particle (E_{β}). The MWPC detects the entrance position of the particles and strongly suppresses γ -ray backgrounds triggering the scintillator when applying an appropriate coincidence gate between the two β detectors. The MWPC runs with 1 atm Ar – CO₂ (9:1 by volume) gas and is separated from the MOT vacuum by a 127 μ m thick 3.81-cm diameter beryllium window.

A Micro Channel Plate (MCP) detector [23] is placed below the trap for detecting recoil ions and determining their hit positions with a resolution of 190 μ m (FWHM). Electrodes are installed in-between the β -telescope and the MCP detector to create an electric field of $E \approx$ 1.3 kV/cm and accelerate the recoil ions emitted from the trap towards the MCP detector. This enhances the ion-collection solid angle so that ~ 85% of the ⁶Li⁺ and 100% of the ⁶Li²⁺ and ⁶Li³⁺ ions are collected within the 75 mm diameter active area of the MCP. It also allows for the ions to have enough energy to trigger the MCP detector with a maximal detection efficiency (~50%), independent of their initial charge state and recoil energy.

The TOF measurement of the recoil ion is started by a scintillator signal and stopped by a MCP signal. The res-

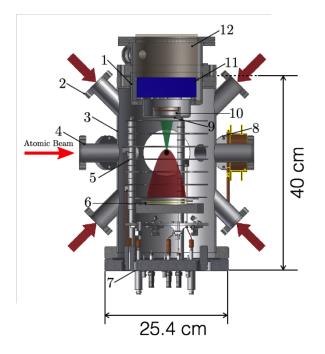


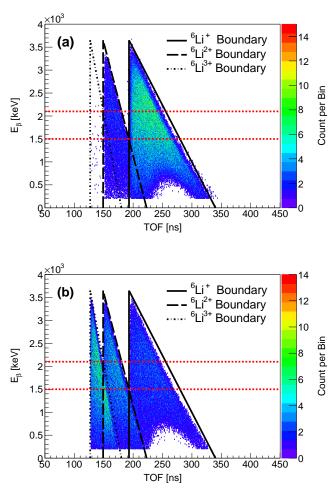
FIG. 1: Cross-section view of the detector system mounted on the measurement chamber. 1) re-entrant β -telescope housing, 2) trapping laser ports, 3) main chamber, 4) ⁶He transfer port, 5) electrode assembly, 6) micro-channel plate (MCP) recoil-ion detector, 7) 10 inch custom feedthrough flange for HV and MCP connections, 8) trap monitoring ports, 9) 127 μ m Be foil, 10) multi-wire proportional chamber (MWPC), 11) plastic scintillator, 12) lightguide to photo-multiplier tube.

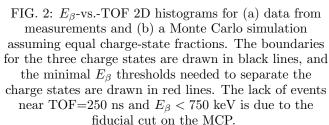
olution of the TOF measurement is 820 ps (FWHM). ⁶Li ions in different charge states have different accelerations in the applied electric field, and are thus partially separated in TOF as shown in Fig. 2. The overlaps between charge states can be avoided by applying a high enough threshold to the β -energy as indicated by the horizontal dashed lines.

To fully reconstruct the initial energy of the recoil ion, the MOT position must also be determined. We ionize the trapped ⁶He atoms periodically (at ~20 Hz) using a 2-ns pulsed beam of ultraviolet (337 nm) nitrogen laser that has sufficient photon energy to photo-ionize the ³P₂ state but not the ³S₁ state. The vertical coordinate (along the electric field direction) of the MOT is determined through the TOF of the photo-ions with respect to the laser pulse.

A unique feature of the trapping of metastable He atoms is that it allows for monitoring the horizontal shape of the MOT via "Penning-ions". The latter are generated by collisions between neutral atoms in the non-perfect vacuum and metastable ⁶He atoms. The uncertainty of the start position of each ⁶Li recoil ion is limited in this data set by the size of the MOT which is 1.4 mm (FWHM). With the precisely measured MOT position,

electric field strength, recoil-ion TOF and hit positions, the initial momenta of recoil ions are determined. The detector responses are calibrated daily, and the electric field and MOT position are monitored while the data are taken. The details of the construction, calibration and performance of the detector system are described in Ref. [24].





Decays from non-trapped ⁶He atoms in the decay chamber and scattering of β particles prior to their detection generate an undesired background. To suppress this background, we reconstructed the momentum of the emitted anti-neutrino using the measured β -momentum and ion-momentum, and evaluated the *Q*-value (the total kinetic energy released from the decay), which should be 3.5 MeV. Decays outside the MOT yield incorrect Q-values because the reconstruction assumes the events originate at the MOT. Our Q-value cut (Q-cut) accepted events with $3.072 \leq Q \leq 3.858$ MeV, so that ~ 90% of the events from non-trapped decays were eliminated while the total data loss for events from trapped decays was less than 0.1% [24]. The width of this cut was limited by the energy resolution in our scintillator. In order to determine the contribution of the remaining events from non-trapped decays, we introduced large amounts of ground-state ⁶He atoms via a bypass pipe.

Fig. 3 shows a comparison of the TOF spectra. In Fig. 3a the data from the non-trapped ⁶He is normalized and overlaid with the data taken with the trap on. The shortest TOF for a ⁶Li ion emitted from the MOT is $T_{LE3} = 127$ ns which is the TOF leading edge for ⁶Li³⁺. Events with TOF < T_{LE3} must be from non-trapped ⁶He decays occurring closer to the MCP. Therefore, the normalization was chosen to match the TOF spectra without the *Q*-cut in the TOF region from 10 ns to 110 ns. The same normalization factor was applied to the TOF and ion-energy spectra after the *Q*-cut (shown in Fig. 3b) and then these spectra were subtracted from those from the corresponding data run to remove the remaining events from untrapped ⁶He atoms.

III. DATA ANALYSIS AND RESULTS

As apparent from the data shown in Fig. 2a and Fig. 3b, we clearly observe some shake-off fraction yielding ${}^{6}\text{Li}^{2+}$ ions, however, we do not observe ${}^{6}\text{Li}^{3+}$ ions above background and can therefore only set an upper limit. In the following, we will treat these two cases separately.

We first address the total fraction of ${}^{6}\text{Li}^{3+}$ ions (P_3) , regardless of their energies. P_3 is the ratio $N_3/N_{\rm all}$, where N_3 is the number of observed ⁶Li³⁺ ions, and N_{all} is the number of observed ⁶Li ions in all charge states. It is important to choose an appropriate E_{β} threshold to ensure a correct determination of $N_{\rm all}$. As shown in Fig. 2, some of the ⁶Li⁺ events with small E_{β} are lost because they have too high transverse momentum to hit the active area of the MCP. Therefore, the E_{β} threshold should be set high enough so that ⁶Li ions in all charge states above the E_{β} threshold fall in the active area of the MCP. We chose 1 MeV as the E_{β} threshold, and under this condition the event loss due to the finite size of the MCP fiducial area is then less than 0.1% according to the Monte Carlo simulation. However, an E_{β} threshold at 1 MeV is not high enough to separate ⁶Li³⁺ from ⁶Li²⁺ ions. Therefore, to obtain the correct N_3 , we determined the number of events below the leading edge $(T_{LE2} =$ 149 ns) of ${}^{6}\text{Li}^{2+}$ TOF spectrum, and then corrected the ${}^{6}\text{Li}^{3+}$ counts to include the counts overlapping with ${}^{6}\text{Li}^{2+}$ events using a Monte Carlo simulation. Given the E_{β} threshold at 1 MeV, the percentage of ${}^{6}\text{Li}^{3+}$ ions that are below T_{LE2} over all ⁶Li³⁺ ions is 85.5%. (Alternatively, it is also possible to choose an E_β threshold at 2.1 MeV

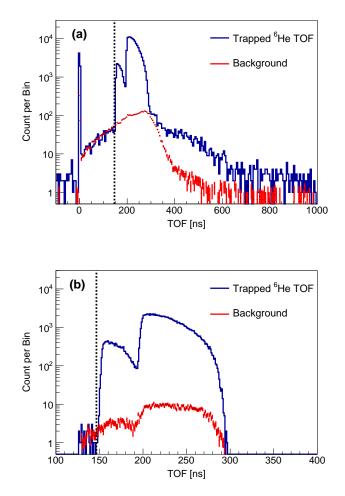


FIG. 3: TOF spectra for laser-off data (a) without and (b) with the Q-cut. The normalized spectra from non-trapped atoms (red curves) for each case are overlaid. E_{β} threshold was set to 1 MeV when generating these plots. The TOF region between 10 and 100 ns on (a) was used to normalize the curves. The sharp peak at time ≈ 0 in spectra (a) corresponds to electrons which backscattered hitting both the MCP and β detectors. The TOF leading edge for ⁶Li²⁺ is shown in black dashed lines in (a) and (b).

so that the ${}^{6}\text{Li}^{3+}$ ions are completely separated from the ${}^{6}\text{Li}^{2+}$ ions, but such a high threshold rules out $\sim 75\%$ of the data, yielding poor statistics so it was not adopted.)

The TOF spectrum with the Q-cut for the laser-off data is plotted in Fig. 3b. The event counts in the ${}^{6}\text{Li}^{3+}$ TOF region (TOF $< T_{LE2}$) are dominated by the background generated by the non-trapped ⁶He decays. A summary of results is shown in Table. II. The measurements are consistent with $P_3 = 0$ within 1 standard deviations, and the 90% confidence levels are calculated only in the physical region where the count of events are greater than 0.

Next, we studied the fraction of ${}^{6}\text{Li}^{2+}$ ions and its

TABLE II: Measured ${}^{6}\text{Li}^{3+}$ ion charge fractions for ${}^{6}\text{He}$ decays from atomic excited states, under the conditions

that $E_{\beta} > 1$ MeV. The negative P_3 values originate from the background subtraction. The 90% confidence levels are calculated only in the physical region where $P_3 > 0$.

Laser	$P_3 \times 10^5$	$\Delta P_3 \times 10^5$	Upper limit $\times 10^5$
status			90% C.L.
On	-1.2	6.5	10
Off	-0.6	6.6	11

dependence on the recoil-ion energy. In this study, the initial energy of the recoil ions (E_{Ion}) are reconstructed for each event, so it is necessary to determine the charge state of each ion with no ambiguity. Therefore, the minimal threshold on E_{β} , 1.5 MeV, was applied so that ${}^{6}\text{Li}^{2+}$ ions are completely separated from ${}^{6}\text{Li}^{+}$ in TOF as shown by the lower of the two horizontal red dashed lines in Fig. 2. As discussed above, the upper limit for the probability of having a ${}^{6}\text{Li}^{3+}$ ion is at the 10^{-4} level, approximately three orders of magnitude lower than the probability of having a ${}^{6}\text{Li}^{2+}$ ion. Therefore, we neglected ${}^{6}\text{Li}^{3+}$ ions in this study assuming that all events below the leading edge ($T_{LE1} = 193$ ns) of ${}^{6}\text{Li}^{+}$ TOF spectrum correspond to ${}^{6}\text{Li}^{2+}$ ions. The E_{Ion} spectra for ${}^{6}\text{Li}^{+}$ and ${}^{6}\text{Li}^{2+}$ with their corresponding normalized backgrounds are plotted in Fig. 4.

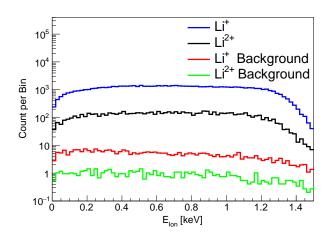


FIG. 4: ⁶Li recoil ion initial energy distributions for Li⁺ and Li²⁺. Background spectra are also plotted.

Theoretically, the probability distribution functions (PDFs) of E_{Ion} in the two charge states are

$$PDF_i(E_{Ion}) = \Phi(E_{Ion}) \times P_i(E_{Ion}) \times \eta_i.$$
(1)

where i = 1, 2 indicates the charge state, and P_i is the probability of having an ion in such charge state. $\Phi(E_{Ion})$ represents the part of the probability distribution function that is independent of charge state and depends only 5

on the β -decay dynamics, *i.e.* the $\beta - \nu$ correlation coefficient $a_{\beta\nu}$. The detection efficiency, η_i , depends on the detection geometry, detector response functions and event reconstruction parameters and conditions. The expected energy dependence for $P_i(E_{Ion})$ is [19]:

$$P_i = A_i + B_i E_{Ion}, \qquad (2)$$

where A_i and B_i are parameters. Taking the ratio of PDF₂ and PDF₁, one gets

$$R(E_{Ion}) = \frac{(A_2 + B_2 E_{Ion}) \times \eta_2}{(A_1 + B_1 E_{Ion}) \times \eta_1}.$$
 (3)

The $\Phi(E_{Ion})$ functions in the numerator and the denominator cancel each other and $R(E_{Ion})$ no longer depends on $a_{\beta\nu}$ explicitly. With the $E_{\beta} = 1500$ keV threshold there is no event loss due to MCP fiducial area for both charge states q = 1 and q = 2. In principle, the ion detection efficiency η_i depends on the charge state be-cause ${}^{6}\text{Li}^{+}$ and ${}^{6}\text{Li}^{2+}$ have very different final energies $(\approx 13 \text{ keV versus} \approx 26 \text{ keV})$, and their position distributions on the MCP are also different. However, the difference in gain of the MCP for these two charge states is only 2.5%. Due to a very low MCP charge threshold, it results in less than 2×10^{-5} efficiency difference. Therefore, we made the assumption that $\eta_2 = \eta_1$, so they cancel each other in Eq. (3). (The systematic uncertainty generated by this approximation was studied using Monte Carlo simulations including the measured 0.82% (RMS) variation of the MCP efficiency over the surface of the MCP.) Because ⁶Li³⁺ ions are neglected, the sum of $P_1(E_{Ion})$ and $P_2(E_{Ion})$ is 1, and thus Eq. (3) becomes

$$R(E_{Ion}) = \frac{A_2 + B_2 E_{Ion}}{1 - A_2 - B_2 E_{Ion}}.$$
 (4)

We take the ratio of the measured E_{Ion} spectra (with background subtraction applied) for the charge states q = 2 and q = 1, and fit the ratio histogram to Eq. (4), as shown in Fig. 5. The fit region is chosen to be from 0.1 keV to 1.1 keV, in order to make the systematic uncertainties introduced by the assumptions made above negligible compared to the statistical uncertainty. A_2 and B_2 are fit parameters, and the fit results are listed in Table III. We also obtain the fractions of ${}^{6}\text{Li}^{+}$ and ${}^{6}\text{Li}^{2+}$ regardless of their energies by counting the total number of events above and below T_{LE1} . The results of the fractions for all three charge states of ⁶Li ions are summarized and compared to theoretical calculations [19] in Table IV. All the fractions determined in this experiment are obtained with an E_{β} threshold. Applying an E_{β} threshold, which is inevitable in this experiment, changes the energy spectrum of the recoil ions, and thus affects the fractions of ⁶Li ions in different charge states due to their dependences on the ion energies. Therefore, in Table IV the theoretical fractions are calculated based on the results of Ref. [19] and applied with the same E_{β} threshold as in our experiment.

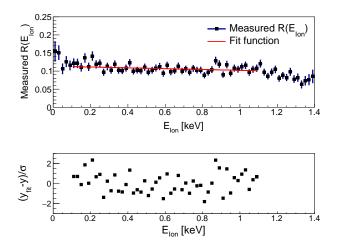


FIG. 5: Ratio between the E_{Ion} spectra for the charge state 2 and 1, fit to Eq. (4) in the range $0.1 \text{ keV} < E_{Ion} < 1.1 \text{ keV}$ for the laser-off data. Normalized residuals inside the fit region are plotted in the lower panel.

TABLE III: Fit results for A_2 and B_2 in Eq. (4).

Laser	$A_2 \times 100$	$\frac{B_2 \times 10^4}{(\text{keV}^{-1})}$	$\chi^2/{ m dof}$
On	10.1 ± 0.3	-36 ± 42	45/48
Off	10.2 ± 0.3	-94 ± 42	49/48

TABLE IV: Comparison of calculated versus measured 6 Li ion charge fractions (in %) for 6 He decays from $^{3}S_{1}$ atomic metastable state.

Ion	Theory ^a [19]	This work
Li^+	88.63(2)	90.5(1)
Li^{2+}	9.5(1)	9.5(1)
Li ³⁺	1.9(1)	≤ 0.01

^a When calculating the theoretical charge fractions for ⁶Li⁺ and ⁶Li²⁺, the averaged ion energy is determined using a 1.5 MeV β -energy threshold as used in the experiment. In this case, $\langle E_{ion} \rangle = 0.74$ keV. For ⁶Li³⁺, β -energy threshold is 1 MeV, and $\langle E_{ion} \rangle = 0.74$ keV as well.

In order to understand the systematic shifts of A_2 and B_2 caused by approximations like assuming η_1 and η_2 to be identical, we ran Monte Carlo simulations with the values of A_2 and B_2 from Ref. [19]. In the simulations, we modeled the experimental parameters, such as the detector geometry and response functions, and the efficiencies and electric field, as close as possible to the experimental setup. The simulated data were processed in the same way as that used for the experimental data. There is no significant deviation of the extracted B_2 value from its input value, while the extracted A_2 deviates from its input by $-7(4) \times 10^{-4}$. Implementation of the 0.82% spatial

variation of the MCP efficiency does not result in significant deviations of A_2 and B_2 . We also studied how the fit values of A_2 and B_2 change with respect to parameters used in the ion-energy reconstruction, and the systematic uncertainties associated with these parameters. The corresponding non-negligible systematic uncertainties of A_2 and B_2 are listed in Table V, and in total are smaller than the statistical uncertainties listed in Table III. Note that these systematic uncertainties are all related to the ion-energy calculation, so they do not affect the determination of the energy-integrated charge-state fraction listed in Table IV.

TABLE V: Systematic shifts and uncertainties of A_2 and B_2 .

	Shift $A_2 \times 100$	$\Delta A_2 \times 100$	$\Delta B_2 \times 10^4$
			(keV^{-1})
Approximations ^a	-0.07	0.04	<6
Vertical MOT Pos.		0.066	6.1
Electric field		0.018	3.1
TOF Origin		0.055	4.6
Total	-0.07	0.1	10

^a Uniform electric field, uniform efficiency of the MCP over its surface, point-like MOT.

IV. DISCUSSION

A comparison of our measurements to calculations for the decay of ⁶He from its atomic metastable state is shown in Table IV. The ⁶Li³⁺ and ⁶Li⁺ fractions measured in this experiment have small (~ 2%) but significant (~ 19 σ for ⁶Li³⁺ and ⁶Li⁺) discrepancies with the theoretical calculations of Ref. [19]. The calculation for the ⁶He decays from the atomic ground state similarly over-predicts the ⁶Li³⁺ fraction measured by Carlson *et al.* [21] as shown in Table I. It is possible that there is a missing consideration in the calculation that systematically leads to a higher ⁶Li³⁺ fraction for both initial atomic states. In Ref. [19] the charge state of the ⁶Li ion is determined solely by the final energy of the two orbital electrons, $E_{tot} = E_1 + E_2$, with respect to the ionization energies for 1 and 2 electrons, E_{ion-1} and E_{ion-2} :

if
$$E_{\text{tot}} \leq E_{\text{ion}-1} \rightarrow {}^{6}\text{Li}^{+}$$

if $E_{\text{ion}-1} \leq E_{\text{tot}} \leq E_{\text{ion}-2} \rightarrow {}^{6}\text{Li}^{2+}$
if $E_{\text{tot}} > E_{\text{ion}-2} \rightarrow {}^{6}\text{Li}^{3+}$.

In considering possible sources for the discrepancy we note, for example, that the condition $E_{\text{tot}} > E_{\text{ion}-2}$ could be met without double ionization if one of the electrons takes away a significant fraction of the energy as kinetic energy, leaving the other electron bound. This could lead to the systematic overestimation of the ⁶Li³⁺ probability as observed.

In contrast to the integrated charge-state fractions, the ion-energy dependencies of the ⁶Li ion charge-state fractions (the B parameters) are of concern for the determinations of the $\beta - \nu$ correlation coefficient. The experiment of Ref. [21] was performed with the same method and apparatus used in Ref. [11] to determine the $\beta - \nu$ correlation coefficient. As shown in Table VI, there is a significant discrepancy between the B parameters measured by Carlson *et al.* and the ones calculated in Ref. [19]. While we find a plausible explanation for some of the differences between theory and experiment for the overall Li-ion fractions, as stated above, it is more difficult to understand how the B factors could be a factor of ~ 7 smaller in the calculations. The B values measured by Carlson *et al.* are close to a naive prediction ignoring a cancellation that takes place between nS and nP final state configurations shown in detail in Ref. [19]. Johnson et al. were aware of the cancellation and the fact that their measured B factors disagreed with the more accurate calculation [11], but at the time they reported concerns about the difficulty of including a large enough set of states in their calculation. There is an implicit suggestion that the discrepancy should not be taken seriously because the calculation was incomplete. No such concerns exist for the recent calculation of Schulhoff and Drake [19], so we conclude that this suggests either an unaccounted-for experimental issue or a failure of the framework for the calculation. Changing the B parameters from their measured value to zero affects the determination of $a_{\beta\nu}$ only by 0.6% which is smaller than the 1% uncertainty claimed by Ref. [11]. In the present context of trying to achieve more precise determinations, the issue is more important.

TABLE VI: Comparison of calculations [19] to previous measurements [21] of charge-distribution probabilities from the electronic ground state of ⁶He.

	Theory [19]		Experiment [21]	
Ion	$A\times 100$	$B \times 10^4$	$A\times 100$	$B\times 10^4$
		(keV^{-1})		(keV^{-1})
Li^+	89.03(2)	-6.17(2)	89.9(2)	-45(7)
Li^{2+}	9.7(1)	+5.8(1)	10.1(2)	+42(7)
Li^{3+}	1.2(1)	+0.34(14)	0.018(15)	+0.33(13)

Our results for the A and B parameters for the ⁶He decays from the atomic metastable state and the corresponding theoretical calculations are shown in Table VII. Unfortunately, our results don't have the statistical power to claim a precision test of the calculations, in particular for the B parameter. We also calculated the parameters A_2 and B_2 for decays from initial atomic state ³P₂, based on the A_2 and B_2 values in Table III for the laser-on case (mixture of 50% ³S₁ and 50% ³P₂) and laser-off case (pure ³S₁). The results together with the final results for the ³S₁ initial states are summarized in Table VIII. The results for these two initial states are

not significantly different from each other.

TABLE VII: Same as Table VI for the decay of ⁶He from its metastable state from this work.

	Theory [19]		This work ^a	
Ion	$A\times 100$	$B \times 10^4$	$A\times 100$	$B\times 10^4$
		(keV^{-1})		(keV^{-1})
Li^+	88.711(3)	-11.06(0)	89.9(3)(1)	94(42)(10)
Li^{2+}	9.42(7)	+10.39(7)	10.1(3)(1)	-94(42)(10)
Li^{3+}	1.86(7)	+0.74(148)	-	-

^a Systematic shifts for A_2 is included. The number in the first parenthesis is the statistical uncertainty, and the number in the second parenthesis is the systematic uncertainty.

TABLE VIII: Results for parameters A_2 and B_2 , for 3S_1 and 3P_2 initial states.

Initial State	$A_2 \times 100$	$B_2 \times 10^4$
		(keV^{-1})
${}^{3}S_{1}$	10.1(3)(1)	-94(42)(10)
${}^{3}P_{2}$	10.0(7)(1)	22(94)(10)

^a The number in the first parenthesis is the statistical uncertainty, and the number in the second parenthesis is the systematic uncertainty.

Additionally, in measurements of the $\beta - \nu$ correlation. where the TOF spectrum is fitted to templates generated by Monte Carlo simulations and the charge state groups are not completely separated in TOF, both the A and B parameters can affect the fit result. We have used a Monte Carlo simulations to study how the uncertainties of A_2 and B_2 translate into the uncertainty of the $\beta - \nu$ correlation coefficient. Based on the experimental uncertainties listed in Table VII, A_2 results in a 0.3% relative uncertainty of $a_{\beta\nu}$, and B_2 results in a 0.6% relative uncertainty. Therefore, the results from this paper are sufficient for an experiment aiming at determining $a_{\beta\nu}$ to 1%. Because the charge-state analysis and the $a_{\beta\nu}$ analysis can use the same data set, the uncertainties of A_2 and B_2 will be improved as more data are taken to achieve better than 1% uncertainties on $a_{\beta\nu}$.

V. CONCLUSIONS

We have measured the ⁶Li ion charge-state fractions for ⁶He decays from atomic metastable state ³S₁. The overall fractions for ⁶Li⁺ and ⁶Li³⁺ (Table IV) show small but significant disagreement with the recent theoretical calculation of Ref. [19]. We discuss a plausible explanation.

We also point out that there is no satisfactory explanation for a large discrepancy between the same calculation and the results of Carlson et al.[21] for the Li-ion energy dependence of the fractions from decays of the electronic ground state, suggesting either an unaccounted-for experimental issue or a failure of the framework for the calculation.

The A and B parameters in the ion-energy dependent charge-state fraction expression (Eq. (3)) were also determined (Table. VIII), and the precision is sufficient for a determination of $a_{\beta\nu}$ with 1% relative precision. The precisions of the A and B parameters can be improved as more data for the $\beta - \nu$ correlation measurement are taken.

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