



LAWRENCE
LIVERMORE
NATIONAL
LABORATORY

A New Measurement of the 60 keV Emission from Am-241 using Metallic Magnetic Calorimeters

G. B. Kim, S. T. P. Boyd, R. H. Cantor, L. A.
Bernstein, S. Friedrich

September 3, 2019

18th International Workshop on Low Temperature Detectors
(LTD-18)
Milan, Italy
July 22, 2019 through July 26, 2019

Disclaimer

This document was prepared as an account of work sponsored by an agency of the United States government. Neither the United States government nor Lawrence Livermore National Security, LLC, nor any of their employees makes any warranty, expressed or implied, or assumes any legal liability or responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by trade name, trademark, manufacturer, or otherwise does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States government or Lawrence Livermore National Security, LLC. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States government or Lawrence Livermore National Security, LLC, and shall not be used for advertising or product endorsement purposes.

A New Measurement of the 60 keV Emission from Am-241 using Metallic Magnetic Calorimeters

G.B. Kim, S.T.P. Boyd, R.H. Cantor, L.A. Bernstein,
S. Friedrich

the date of receipt and acceptance should be inserted later

Abstract We report a new measurement of the 60 keV transition from ^{241}Am . It uses a metallic magnetic calorimeter (MMC) gamma-ray detector calibrated in the region around 60 keV by four high accuracy X-rays and gamma-rays from the decay of ^{169}Yb . We determine an energy of $59,539.3 \pm 0.3$ (stat) ± 0.3 (syst) eV, which is 1.6 ± 0.4 eV lower than the current literature value of $59,540.9 \pm 0.1$ eV. We discuss the sources of this uncertainty and approaches to address them.

Keywords Am-241, gamma-ray spectroscopy, nuclear data, metallic magnetic calorimeter, microcalorimeter

1 Introduction

The 60 keV transition in the decay of ^{241}Am produces one of the most widely used calibration lines for low-energy gamma-ray detectors. ^{241}Am is long-lived and widely available, the transition has a high branching ration of 35.9 %, and the gamma energy has an uncertainty of only ± 0.1 eV, significantly smaller than that of most other isotopes [1, 2]. The current literature value of 59,540.9(1) eV has been determined from its difference to the ^{161}Tb gamma-ray at 48,915.62(14) eV [2, 3], which in turn is based on a measurement with a bent-crystal spectrometer [4]. The ^{241}Am - ^{161}Tb spectra were measured repeatedly with one Si(Li) and two high-purity Ge detectors, and the top portion of the peaks were fit to a Gaussian function over a channel range of little more than one FWHM. The differences between the ^{241}Am and the ^{161}Tb centroids varied by 1 eV for the three detectors, and their weighted average had an uncertainty of 0.1 eV. Note that the only statistical uncertainties were included in [3]. Systematic errors, e.g. due to the low-energy tail of the detector response, the shape of the

G.B. Kim, S. Friedrich
Lawrence Livermore National Laboratory, Livermore, CA 94550, USA
S.T.P. Boyd
University of New Mexico, Albuquerque, NM 87131, USA
R.H. Cantor
Star Cryoelectronics, Santa Fe, NM 87508, USA
L.A. Bernstein
Lawrence Berkeley National Laboratory, Berkeley, CA 94720, USA
E-mail: kim90@llnl.gov

Compton background, the choice of the fit range or the non-linearity of the MCA, have not been taken into account. While this is usually justified, the importance of ^{241}Am to calibrate low-energy gamma spectra make it desirable to measure its decay radiation independently.

Metallic magnetic calorimeter (MMC) gamma-ray detectors provide an order of magnitude higher energy resolution than semiconducting gamma detectors and can therefore increase the accuracy of centroid measurements accordingly. They also have a predictable response function that is mostly linear with energy with only a small and reproducible second-order correction [5, 6]. This makes them well-suited to re-measure the energy of the 60 keV transition in the decay of ^{241}Am accurately. Among the possible calibration sources, ^{169}Yb is the best choice because its absolute gamma-energies have been measured with an accuracy of ≤ 0.1 eV with a double-flat Si crystal spectrometer whose lattice spacing has been referenced to the Cs reference scale for frequencies [2, 7]. The K-shell X-rays of the ^{169}Yb daughter ^{169}Tm have been measured with similar accuracy [8, 9]. Since ^{169}Yb is not commercially available, we have produced it in a (d,2n) reaction by irradiation of a monoisotopic ^{169}Tm target with 15 MeV deuterons. This paper discusses our initial measurements of the 60 keV decay in ^{241}Am with an MMC gamma detector calibrated by ^{169}Yb .

2 Experiment

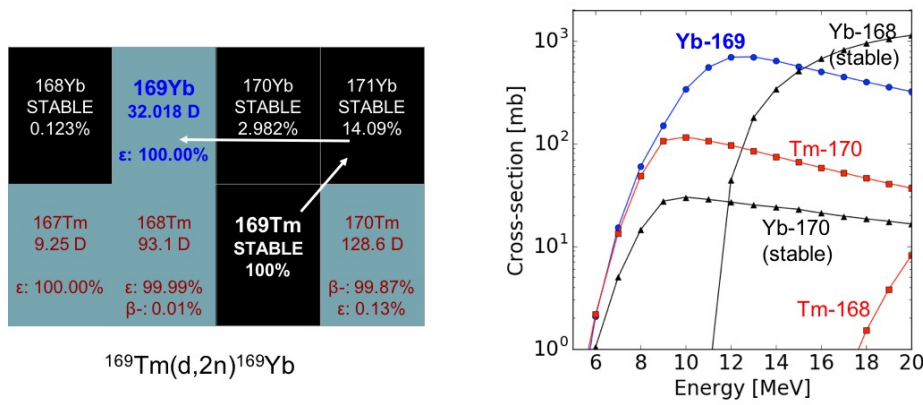


Fig. 1 (Left) Table of nuclides around ^{169}Yb [10]. The arrows indicate the way ^{169}Yb is produced in the (d,2n) nuclear reaction on ^{169}Tm . (Right) Calculated cross-sections for a ^{169}Tm target as a function of deuteron energy [11]. ^{168}Tm is a radioactive isotope whose production should be minimized. (Color figure online)

2.1 Yb-169 Calibration Source

Among all isotopes used for gamma detector calibration, the gamma-rays of ^{169}Yb are known with the highest accuracy in the energy range below 300 keV [2, 7]. Specifically, the decay of ^{169}Yb produces a gamma-ray at 63,120.44(3) eV with a branching ratio of 43.62 % that is just above the energy to the ^{241}Am emission of interest. In addition, the ^{169}Yb decay generates strong Tm K X-rays whose energies are just below the ^{241}Am line

and have been measured with similarly high accuracy [8, 9]. ^{169}Yb is therefore the ideal isotope to calibrate an MMC detector for an accurate measurement of the ^{241}Am gamma-ray at 60 keV. Since ^{169}Yb is not commercially available, we made a 0.25 diameter target out of a commercial 100 μm thick thulium foil, which consists to 100 % of the isotope ^{169}Tm , and irradiated it at the 88" Cyclotron at Lawrence Berkeley National Laboratory with a 15 MeV deuteron beam. Simulations show that 15 MeV deuterons lose on average of 2.7 MeV in Tm over 100 μm . The beam energy was then chosen to produce ^{169}Yb with a cross section of ~ 500 mbarn throughout the foil in the reaction $^{169}\text{Tm}(d,2n)^{169}\text{Yb}$ while minimizing the amount of ^{168}Tm , whose decay is accompanied by several strong gamma-rays. In addition, 15 MeV deuterons produce the stable isotopes ^{168}Yb and ^{170}Yb and the radioactive ^{170}Tm , whose decay produces an additional albeit weak calibration line at 84.25474(8) keV. For a beam current of 750 nA, roughly a third of which hit the Tm target, we initially produced ~ 1.8 μCi of ^{169}Yb during one hour of irradiation.

2.2 Metallic Magnetic Calorimeters

The experiment used an MMC gamma detector array designed at the University of New Mexico (UNM) and fabricated at STAR Cryoelectronics [12, 13]. The MMCs consist of 30 μm thick Au absorbers with an area of $(500 \mu\text{m})^2$ that are supported on paramagnetic Ag:Er sensors by eight Au posts. They are operated in a cryogen-free dilution refrigerator with a base temperature < 7 mK. Gamma signals are amplified with a two-stage SQUID from STAR Cryoelectronics, with the first-stage SQUID being integrated on the same chip as the MMC. The array has 14 MMC pixels and has an energy resolution of as high as 38 eV FWHM at 60 keV [12]. The MMC response is very consistent with that of MMC gamma detectors developed at Heidelberg University and read out with a two-stage SQUID preamplifier by Magnicon [14]. This gives us some confidence that deviations from literature values are not due to peculiarities in the MMC response. For this experiment, we did not chemically separate the ^{169}Yb from the Tm target, but attached the irradiated target to a 1/16" Cu foil to reduce low energy Tm L X-rays and mounted it in front of the MMC gamma detector at a distance of 5 mm. An external ^{241}Am source is periodically added outside the cryostat for repeated measurements with and without ^{241}Am source to measure the background and check consistency.

2.3 Analysis Procedure

We capture the full gamma-induced waveforms with a four-channel 14-bit GaGe digitizer and write them to disc for subsequent off-line analysis. Only two pixels of the MMC array are selected for this measurement to avoid degrading the energy resolution due to Joule heating of the MMC by power dissipation of the SQUIDs, which are located on the same substrate as the MMC [15]. We have found that a trapezoidal filter produces spectra with comparable energy resolution as an optimal filter that is typically used to process microcalorimeter signals [5] and prefer it because it is faster than the optimal filter and does not require template signals or noise spectra. Filtering parameters are set to a peaking time of 1 ms and a gap time of 10 μs . We correct for a drift of signal amplitudes due to slow temperature fluctuations by averaging 50 amplitudes of the 177.21307 keV line of ^{169}Yb and correcting all amplitudes by this scale factor.

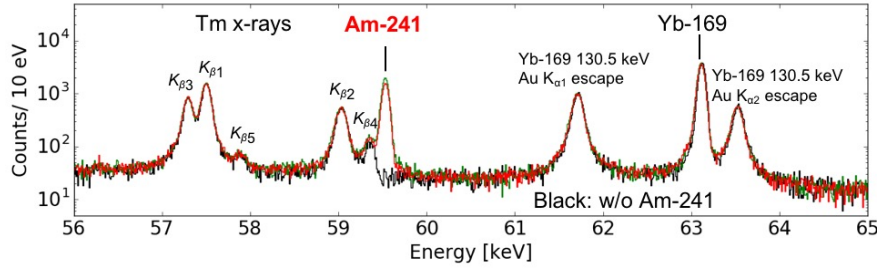


Fig. 2 Calibrated spectra from two MMC pixels. The measurement is repeated with (red, green) and without (black) the ^{241}Am source. (Color figure online)

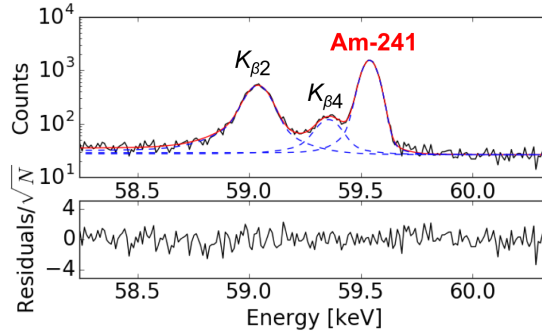


Fig. 3 Spectral fit to three peaks in the 60 keV region. Fits to individual peaks (dashed, blue) and the sum (solid, red) are shown. (Color figure online)

3 Results

The combined gamma-ray spectra of the ^{169}Yb and ^{241}Am sources in the region around 60 keV are shown in Fig. 2. Despite the high energy resolution of 80 eV in this run, there is some line overlap of the ^{241}Am line of interest with the Th $K_{\beta 4}$ X-ray at 59.3554(3) keV. It is this line overlap made us take the spectra both with and without the ^{241}Am source so that we could accurately determine the spectral background without the ^{241}Am line. Since the MMC response function is constant, we can determine the line shape from a strong gamma-ray and apply it to other lines in the spectrum. We chose to extract the line shape from the isolated ^{169}Yb gamma-ray at 177.2130 keV and fit it to a Gaussian function with small tails due to pile-up as before [2, 7]. The small step due to small-angle scattering in the Cu filter is fit by a complementary error function centered at the peak position. We then keep the ratio between the Gaussian and the tail constant and only vary the centroid position, the peak width and the step height to fit the other gamma-rays in the spectrum. For X-rays, we convolve the MMC response function with the natural linewidth (Voigt function) before the fit.

Fig. 3 shows the fits in the 60 keV region that includes the ^{241}Am gamma-ray of interest and the two Tm $K_{\beta 2}$ and $K_{\beta 4}$ X-rays. They agree with the measured spectra within the statistical accuracy of the measurement, although the energy of the Tm $K_{\beta 4}$ X-ray has to be shifted by 152(3) eV from the literature value of 59.2038(10) keV [9]. This suggests that

there is no other significant Tm X-ray peak in the 60 keV region except the interfering Tm $K_{\beta 4}$ line.

The spectra are calibrated using only those X-rays and gamma-rays whose literature values are known with very high accuracy. This includes the Tm $K_{\beta 1}$ X-ray at 57.50876(15) keV [9], the ^{169}Yb gamma-ray at 63.12044(3) keV [16], and the two Au K X-ray escape lines from the ^{169}Yb gamma-ray at 130.52293(4) keV [16]. For a Au $K_{\alpha 1}$ energy of 68.80450(18) keV and a Au $K_{\alpha 2}$ energy of 66.99073(22) keV [9], these lines are seen at energies of 61.71843(18) and 63.5322(22) keV, respectively, with uncertainties and linewidths set by the Au K X-rays. Calibration uncertainties are obtained as before [5, 6] by successively varying the centroid energies by their statistical and literature uncertainties and calculating different calibration curves for each set of calibration points. The calibration uncertainty as a function of energy is then given by the standard deviation of these calibration curves (Fig 4, shaded area).

Fig. 4 shows the residuals of the energy calibration, i.e., the difference between the measured average energies from the two MMC pixels and their literature values. While the values of the calibration points are consistent with the literature values within the uncertainty of the measurement, the energy of the ^{241}Am gamma-ray is slightly lower. It is measured as 59.5393(4) keV, which is 1.6 eV lower than the current literature value of 59.5409(1) keV. This deviation is in the direction of an earlier evaluated value of 59.537(1) keV for the ^{241}Am emission [3]. Although the accuracy of the measurement is currently still limited by an error of 0.4 eV, the deviation is 4 σ , which suggests that the current literature value may be slightly off.

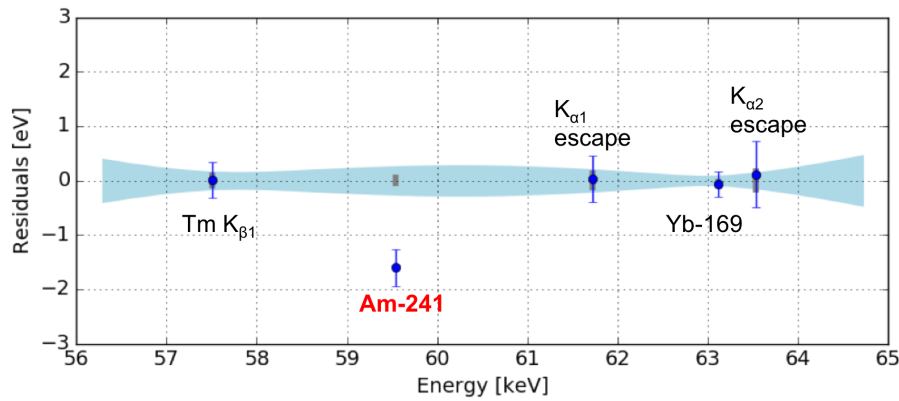


Fig. 4 Residuals of the energy calibration with statistical errors (*blue bars*), literature errors (*gray bars*) and total calibration errors (*shaded area*). While the residuals of the four calibration lines are consistent with zero, i.e. the measured energies are consistent with the literature values, the energy of the ^{241}Am emission is 1.6(4) eV smaller than the literature value of 59.5409(1) keV. (Color figure online)

4 Discussion

MMC gamma-ray detectors are well-suited for accurate measurements of nuclear data due to their high energy resolution, their good linearity and their reproducible response function. We have combined MMCs with the most accurate low-energy gamma calibration source

available, ^{169}Yb , to re-measure the energy of the important calibration line from the decay of ^{241}Am . We find an energy of 59.5393(4) keV that is 1.6(4) eV lower than the current literature value of 59.5409(1) keV. At this point, the 0.4 eV uncertainty of our measurement is still too high to firmly ascertain a discrepancy, and the experiment should therefore be repeated with more detectors and better statistics. In addition, there are systematic uncertainties of 0.3 eV in our measurement that have not been considered in the earlier characterization of the ^{241}Am decay. These include uncertainties due to detector drift, the choice of the fit function and range as well as binning effects. Finally, the precision of cryogenic detectors is reaching a point where the non-linearity of ADC can make a non-negligible contribution to the observed nonlinearity. We are currently investigating ADC non-linearities carefully [17], and may have to introduce an ADC correction to account for this effect. Once we understand all systematic errors quantitatively, it would ultimately be desirable to repeat this measurement at several institutions to assess if the systematic errors are confirmed by independent data sets. We are currently setting up a collaboration for that purpose.

Acknowledgements This work was funded by DOE NA-22 under grant LL16-MagMicro-PD2La. It was performed under the auspices of the U.S. DOE by LLNL under Contract DE-AC52-07NA27344

References

1. M. Basunia, Nuclear Data Sheets **107**(8), 2323 (2006). DOI <https://doi.org/10.1016/j.nds.2006.07.001>
2. R. Helmer, C. van der Leun, Nucl. Instr. and Meth. A **450**(1), 35 (2000). DOI [https://doi.org/10.1016/S0168-9002\(00\)00252-7](https://doi.org/10.1016/S0168-9002(00)00252-7)
3. R. Helmer, Nucl. Instr. and Meth. A **330**(3), 434 (1993). DOI [https://doi.org/10.1016/0168-9002\(93\)90572-Y](https://doi.org/10.1016/0168-9002(93)90572-Y)
4. B. Jeckelmann, et al., Nucl. Instr. and Meth. A **241**(1), 191 (1985). DOI [https://doi.org/10.1016/0168-9002\(85\)90532-7](https://doi.org/10.1016/0168-9002(85)90532-7)
5. C.R. Bates, et al., Appl. Phys. Lett. **109**(2), 023513 (2016). DOI 10.1063/1.4958699
6. G.B. Kim, et al., J. Low Temp. Phys. **193**(5), 1236 (2018). DOI 10.1007/s10909-018-1978-0
7. E. Kessler, et al., Nucl. Instr. and Meth. **160**(3), 435 (1979). DOI [https://doi.org/10.1016/0029-554X\(79\)90197-6](https://doi.org/10.1016/0029-554X(79)90197-6)
8. G. Borchert, Zeitschrift Naturforschung Teil A **31**, 102 (1976)
9. R.D. Deslattes, et al., Rev. Mod. Phys. **75**, 35 (2003). DOI 10.1103/RevModPhys.75.35
10. National nuclear data center. URL <https://www.nndc.bnl.gov/>
11. Tendl nuclear data library (2017). URL https://tendl.web.psi.ch/tendl_2017/tendl2017.html
12. S.T.P. Boyd, R. Hummatov, et al., J. Low Temp. Phys. **193**(3), 435 (2018). DOI 10.1007/s10909-018-2017-x
13. R. Hummatov, J.A. Hall, et al., J. Low Temp. Phys. **193**(5), 752 (2018). DOI 10.1007/s10909-018-1946-8
14. G.B. Kim, et al., J. Radioanal. Nucl. Chem. **318**(1), 803 (2018). DOI 10.1007/s10967-018-6182-9
15. S.T.P. Boyd, et al., This Special Issue (2019)
16. C.M. Baglin, Nuclear Data Sheets **109**(9), 2033 (2008). DOI <https://doi.org/10.1016/j.nds.2008.08.001>
17. S. Friedrich, et al., This Special Issue (2019)