Mass of ⁵⁷Ni[†]

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A measurement of the Q values of the ${}^{59}\text{Ni}(p, t){}^{57}\text{Ni}$ and the ${}^{58}\text{Ni}({}^{3}\text{He}, \alpha){}^{57}\text{Ni}$ reactions shows a significant discrepancy with the current tabulated mass values. We find -12738.2 ± 3.3 keV and $+8360.3 \pm 4.0$ keV, respectively, leading to a new improved mass excess for ${}^{57}\text{Ni}$ of -56078.4 ± 3.0 keV.

NUCLEAR REACTIONS ⁵⁹Ni(p,t); ⁵⁸Ni(³He, α); measured Q values, deduced mass excess for ⁵⁷Ni.

Since the 1971 Wapstra-Gove mass tables¹ were published, several reaction Q values have been found²⁻⁴ which disagree with the tables by several standard deviations. In this note, we report new results for the mass for ⁵⁷Ni found during the course of studying the ⁵⁹Ni(p, t)⁵⁷Ni and ⁵⁸Ni(³He, α)⁵⁷Ni reactions. The ⁵⁷Ni mass excess of -56104 ± 7 keV reported in the 1971 mass table is based on the end point energy for the ⁵⁷Ni-⁵⁷Co decay.⁵ A previously measured Q value of +8400 \pm 50 keV for the ⁵⁸Ni(³He, α)⁵⁷Ni reaction⁶ was not used to determine the mass excess of ⁵⁷Ni.

In the ⁵⁹Ni(p, t)⁵⁷Ni experiment, we used a 40 MeV proton beam from the Michigan State University cyclotron. The reactor produced ⁵⁹Ni target (37.4% ⁵⁸Ni, 43.0% ⁵⁹Ni, 15.2% ⁶⁰Ni, 1.0% ⁶¹Ni, 2.1% ⁶²Ni, and 1.2% ⁶⁴Ni) was a rolled foil of about 230 μ g/cm² thickness. The reaction products were detected in the focal plane of an Enge split-pole spectrograph by a position sensitive proportional counter. The resolution obtained was about 15 keV full width at half maximum (FWHM). Figure 1 shows an example of the spectra obtained. As one can see, the ground and 0.769 MeV state transitions of the ⁵⁹Ni(p, t)⁵⁷Ni reaction fall in the same region as the ground and 1.454 MeV state transitions of the ⁶⁰Ni(p, t)⁵⁸Ni reaction and the ground state transition of the ⁵⁸Ni(p,t)⁵⁶Ni reaction. These three transitions thus serve as calibration lines for the ⁵⁹Ni(p, t) Q-value determination. The spectrograph calibration method and fitting procedure is described extensively elsewhere.⁷ Since the present measurement involves only the position of the tritons from the different Ni isotopes in the target, the uncertainties attributed to the beam energy and the scattering angle and due to target thickness effects are negligible. The main sources of errors are the Q values of the calibration reactions and local nonlinearities in the positionsensitive detector. The uncertainties of both sources are estimated to be ≤ 3 keV. Using the most recent mass excess values of Jolivette *et al.*⁴ for the calibration Q values we obtain -12738.2 ± 3.3 keV for the Q value of the ⁵⁹Ni(p,t)⁵⁷Ni ground state transition. This value is 18.4 keV more negative than the -12719.8 ± 7.6 keV result calculated from the 1971 mass table.

The ⁵⁸Ni(³He, α)⁵⁷Ni experiment was performed with a 70 MeV ³He beam using a similar setup as described before. The target was prepared by vacuum evaporation of ⁵⁸Ni (isotopically enriched

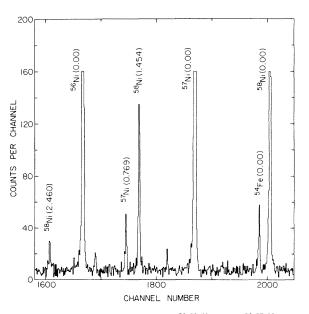


FIG. 1. Triton spectrum of the ${}^{58, 59, 60}$ Ni(p, t) ${}^{56, 57, 58}$ Ni reactions. The peaks are labeled by the final nucleus and its excitation energy.

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to 99.9%) onto a carbon backing and had a thickness of about 90 μ g/cm². In this experiment, the magnetic rigidities of the α particles from the ⁵⁸Ni(³He, α)⁵⁷Ni reaction were compared to those from the calibration reaction ${}^{52}Cr({}^{3}He, \alpha){}^{51}Cr$. The magnetic field of the spectrograph was adjusted to put the α particles from both ground state transitions on the same location of the focal plane. This method, described in detail previously,⁸ does not rely on the linearity of the detector. The dependence of the Q-value determination on the beam energy and scattering angle was minimized by choosing the calibration reaction with a very similar Q value and by measuring at very forward angles ($\theta_{1ab} \leq 10^{\circ}$). The main sources of error, here, were the uncertainties in the energy loss corrections resulting from target thickness uncertainties and the uncertainty in the Q value of the calibration reaction ${}^{52}Cr({}^{3}He, \alpha){}^{51}Cr$, again taken from Jolivette $et \ al.^4$ The Q value thus obtained for the ⁵⁸Ni(³He, α)⁵⁷Ni reaction is +8360 \pm 4.0 keV, 16 keV less positive than the published value of $+8376 \pm 8$ keV.

The results of both measurements together with

- *Research supported in part by the U.S. National Science Foundation and by the U.S. Energy Research and Development Administration under contract with Union Carbide Corp.
- ¹A. H. Wapstra and N. B. Gove, Nucl. Data <u>A9</u>, 265 (1971); N. B. Gove and A. H. Wapstra, *ibid*. <u>A11</u>, 127 (1972).
- ²J. D. Goss, C. P. Browne, and A. A. Rollefson, Phys. Rev. Lett. 30, 1255 (1971).
- ³A. Moalem and B. H. Wildenthal, Phys. Rev. C <u>8</u>, 1961 (1973).
- ⁴P. L. Jolivette, J. D. Goss, G. L. Marolt, A. A. Rollef-

TABLE I. Q-value measurements for the determination of the mass excess of 57 Ni. The mass excesses of Ref. 4 have been used throughout this work. Weighted average of the mass excess for 57 Ni: -56078.4 ± 3.0 keV.

Reaction	Q value (keV)	Mass excess of final nucleus (keV)
$^{52}Cr(^{3}He, \alpha)^{51}Cr$	$+8537.8\pm2.2$	
58 Ni(3 He, α) 57 Ni	$+8360.3\pm4.0$	-56080.4 ± 4.6
60 Ni(p,t) 58 Ni	-11904.5 ± 2.8	
${}^{58}\text{Ni}(p,t){}^{56}\text{Ni}$	-13979.0 ± 3.0	
${}^{59}{ m Ni}(p,t){}^{57}{ m Ni}$	-12738.2 ± 3.3	-56075.2 ± 3.9

the values used for calibration are collected in Table I. From the two Q values a mass excess for ⁵⁷Ni of -56078.4 ± 3.0 keV was extracted. This value is 25.6 keV more positive than the value -56104 ± 7 keV reported in the 1971 mass table. The apparent discrepancy between this 25.6 keV and the average deviation of 17.1 keV in the Q values measured is the result of the change in the measured masses of the ^{58, 59, 60}Ni isotopes since 1971.

son, and C. P. Browne, Phys. Rev. C 10, 2449 (1974).

- ⁵E. W. A. Lingeman, J. Konijn, F. Diederix, and B. J. Meijer, Nucl. Phys. <u>A100</u>, 136 (1967).
- ⁶C. M. Fou and R. W. Zurmühle, Phys. Rev. <u>140</u>, B1283 (1965).
- ⁷J. A. Nolen, Jr., G. Hamilton, E. Kashy, and I. D. Proctor, Nucl. Instrum. Methods 115, 189 (1974).
- ⁸W. Benenson, E. Kashy, I. D. Proctor, and B. M. Preedom, Phys. Lett. <u>43B</u>, 117 (1973); E. Kashy, W. Benenson, I. D. Proctor, P. Hauge, and
- G. Bertsch, Phys. Rev. C 7, 2251 (1973).