

BRIEF REPORTS

Brief Reports are short papers which report on completed research or are addenda to papers previously published in the Physical Review. A Brief Report may be no longer than four printed pages and must be accompanied by an abstract.

Thick-target yields of iodine isotopes from proton interactions in Te, and the double- β decays of $^{128,130}\text{Te}$

M. T. F. da Cruz,^{1,*} D. W. Bardayan,² Y. Chan,¹ A. García,¹ M. M. Hindi,² R.-M. Larimer,¹ K. T. Lesko,¹ E. B. Norman,¹ Della F. Rossi,¹ R. G. Stokstad,¹ F. E. Wietfeldt,^{1,3} and I. Žlimen^{1,†}

²Physics Department, Tennessee Technological University, Cookeville, Tennessee 38505

³Physics Department, University of California, Berkeley, California 94720

(Received 26 May 1993)

We measured thick-target yields of $^{126,128,130}\text{I}$ from bombardments of natural Te targets with 15-, 30-, 45-, and 50-MeV protons, together with iodine production cross sections for 1.85- and 5.0-GeV protons. Using these data, we have estimated the relative cosmic-ray induced production of ^{126}Xe , ^{128}Xe , and ^{130}Xe in Te ores. These quantities are significantly different from those used previously in a determination of the ratio of the double- β decay half-lives of ^{130}Te and ^{128}Te , and as a result the cosmic-ray correction is smaller than previously assumed. A revised correction of cosmic-ray produced xenon can change the half-life ratio by about 6%. This quantity is of importance because it can be used to set a limit on the $0-\nu$ double- β decay mode.

PACS number(s): 25.40.-h, 23.40.-s, 27.60.+j

The first evidence of ^{130}Te double- β decay was obtained more than 40 years ago by Inghram and Reynolds [1], who performed a geochemical experiment, showing that ^{130}Te decays to ^{130}Xe . The geochemical method is based on the measurement of the number of double- β decay daughter atoms that are trapped in the parent mineral. The accuracy of this technique depends on several factors. First, large differences in the chemical properties of the parent and daughter species facilitate the separation and counting of the relatively small number of daughter atoms. Second, contaminations of the parent mineral with the daughter species at the time of formation of the mineral can limit the technique's accuracy. Third, the mobility of the daughter species in the mineral and the diffusion times will affect the measurement. Besides simple diffusion of the atoms, seismic and thermal effects must be considered. Fourth, the ore must be precisely dated and be of sufficient age that an adequate concentration of the daughter species has accumulated to permit accurate measurement. Finally, the measured quantity of double- β decay candidate nuclei must be corrected for other processes that can also generate the daughter species. Processes such as fission, charged-particle capture, and cosmic-ray spallation reactions must be considered. So far, the geochemical method has been successfully applied only to those double- β decay candidates

which lead to noble-gas daughter species: $^{82}\text{Se} \rightarrow ^{82}\text{Kr}$ and $^{128,130}\text{Te} \rightarrow ^{128,130}\text{Xe}$. These results have been recently reviewed by Manuel [2]. The ^{82}Se case is of particular interest because double- β decay has been measured for this system both by the geochemical method and by direct counting. The agreement of the half-lives determined by these two very different techniques provides an excellent consistency check of both methods.

The double- β decay half-lives of $^{128,130}\text{Te}$ have been recently remeasured by Bernatowicz *et al.* [3], using the geochemical method. A recent review by Moe [4] points out a possible discrepancy between the absolute half-life values obtained by Bernatowicz *et al.* and those from previous results [2], where the half-life of ^{130}Te decay was measured relative to that of ^{82}Se , from ores containing minerals of both elements. However, the half-life ratio $t_{1/2}(^{130}\text{Te})/t_{1/2}(^{128}\text{Te})$ is independent of the age of the minerals and of the retention of the daughter atoms. Bernatowicz *et al.* measured the number of atoms, and isotopic abundances of Xe present in samples collected from several tellurium ores. Deviations from the isotopic abundances of atmospheric Xe, in the form of excesses of some of the isotopes, were found. As can be seen from Fig. 1, several processes besides double- β decay can produce Xe isotopic excesses: (i) fission, (ii) neutron capture on Te, (iii) $^{127}\text{I}(n, \gamma)$, (iv) α -particle-, (v) neutrino-, (vi) cosmic-ray muon- and secondary-proton reactions. After correcting the measured excesses for the mechanisms (i)-(v), there still remained a small excess of ^{126}Xe , which is not a double- β decay product, that Bernatowicz *et al.* attributed to muon- and proton-induced reactions.

Cosmic rays observed at sea level consist mostly (\approx

*On leave from Instituto de Física, Universidade de São Paulo, Caixa Postal 20516, 01498 São Paulo, SP, Brasil.

†On leave from Rugjer Bošković Institute, Zagreb, Croatia.

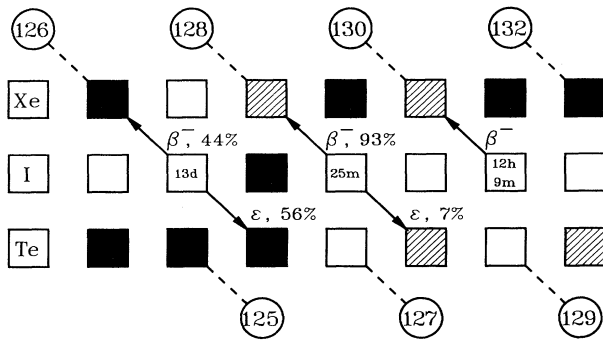


FIG. 1. Region of the chart of the nuclides around Te-I-Xe. The circles contain the mass number of the isotopes in the corresponding diagonal, the filled squares represent stable isotopes, and the dashed ones are the double- β decay parent ($^{128,130}\text{Te}$) and daughter ($^{128,130}\text{Xe}$) nuclei. The radioactive iodine isotopes relevant for the problem have their half-lives and decay modes indicated.

75%) of muons, with a vertical flux of about $1.3 \times 10^2 \text{ m}^{-2} \text{ s}^{-1}$ [5]. The muon energy spectrum falls steeply with energy, $\propto E^{-2}$, and faster above a few TeV. Muons are highly penetrating particles; they are observed in several underground experiments, with a total flux decreasing approximately exponentially as a function of the depth [6,7].

Bernatowicz *et al.* considered muon-induced reactions, and also those caused by secondary protons (i.e., protons produced in the muon-induced nuclear spallation in the rock), showing that their contribution does not affect the value inferred for the ^{130}Te double- β decay half-life, because they account for only 10^{-5} of the ^{130}Xe measured excesses. On the other hand, the same calculations show an effect as large as 20% for the ^{128}Te half-life, as observed in one of Bernatowicz *et al.*'s samples. However, the correction to the ^{128}Xe excess for this mechanism depends critically on how one scales the ^{128}Xe excess from that of ^{126}Xe , the "cosmic-ray detector." Because these corrections are model dependent, we decided to address them by measuring the yields of iodine from proton reactions on tellurium targets, to shed light on the contribution of the secondary protons underground, and estimate that of direct muon reactions.

The Te targets were made of 99.99% purity metallic tellurium, purchased from Johnson Matthey. Tellurium pieces were crushed and pressed at 120°C for 4 min, to form the targets. For the low-energy activations (≤ 50 MeV) the beam was stopped in the targets, which were disks, 1.9 cm in diameter and 1.0 cm thick, with densities of about 6.01 g/cm^3 (96–97% of the crystalline tellurium density). The targets for the high-energy experiments were (i) a disk, 3.0 cm^3 in diameter and 0.7 cm thick for the 1.85-GeV activation, with a density of 6.04 g/cm^3 , and (ii) a square, $5.1 \times 5.1 \times 1.0 \text{ cm}^3$ for the 5.0-GeV activation, with a density of 5.66 g/cm^3 .

The low-energy proton activations were performed at the 88-Inch Cyclotron at Lawrence Berkeley Laboratory; those at high-energy were performed at the LBL's Bevatron accelerator. At the Bevatron, the targets were held in air, with the Te slabs assembled in a stack, to-

gether with Polycast acrylic plastic sheets (polymethyl methacrylate, $[\text{C}_5\text{O}_2\text{H}_8]_n$). These plastic sheets served to monitor the integrated beam exposure, through the production of ^{11}C , from the C and O contents of the plastic [8,9]. The 15-, 30-, 45-, and 50-MeV proton activations were done with bombardment times of 10 min for the first two, and 1 h for the last one, each having an integrated current of $10 \mu\text{C}$. The 1.85- and 5.0-GeV proton experiments were done with bombardment times of approximately 1 h, with integrated currents of 60 and 5 nC, respectively. To check for iodine losses during the activations, due to the heating of the target, one of the activated targets was heated to $\approx 100^\circ\text{C}$ on a hot plate, for a period corresponding to the bombardment time. The radioactivity lost was less than 1%.

Following the irradiations, the targets were γ counted with a 100-cm^3 coaxial HPGe detector inside a 5-cm-thick lead shielding. Due to the widely different half-lives of the iodine isotopes under study: ^{126}I (13 days), ^{128}I (25 min), ^{130m}I (9 min), and ^{130g}I (12 h), three different sizes of time bins were used for counting: 5-min bins during the first hour, 1-h bins during the next 24 h, and then several 6-h bins. Some of the targets were measured 2 weeks later, to confirm the amount of radioactivity attributed to the 13-day component. All photopeaks in the spectra could be attributed to reactions on Te.

The data analysis was carried out by fitting the photopeaks of characteristic γ rays of each isotope, with emission probabilities greater than 1% for all spectra, using the program GELFIT [10]. The time-dependent yields of each γ -ray line were fit to determine initial activities. The decay of the 9-min isomer of ^{130}I has no pure, intense transitions (i.e., with an emission probability $>1\%$) that are not produced in the decay of ^{130g}I . We then performed a two-component fit of the 536-keV photopeak time-dependent yields to deduce the separate contributions of ^{130m}I and ^{130g}I . All half-lives measured were in agreement with the values found in the literature [11], indicating no loss of iodine from the targets during the measurements.

The γ -ray detection efficiency curves for the thick-target yields were determined as follows. We measured the detection efficiency for standard γ -ray calibration sources placed in front and behind our targets for several locations on the target surface. These scans of the target surface were then averaged for the front and back measurements separately, to approximate the extended-source geometry. We then corrected the efficiency data for coincident γ -ray summing effects using the program KORSUM [12], and fit the front and back efficiencies as a function of the photon energy. Because the high-energy measurements were done with target thicknesses much smaller than the corresponding proton range, we used the geometric average of the previous two efficiency curves as our effective efficiency. This corresponds to a first-order correction of the self-attenuation and solid-angle difference between the front and back efficiencies. For proton energies of 50 MeV and below, the targets completely stopped the beam, so we had to do a different correction for the self-attenuation. For this, we fit the total photon-absorption cross section for tellurium in the en-

ergy range of interest, with data taken from the work of Hubbell [13]. Then we took the efficiency curve determined for the front of the target and folded it with an energy-dependent term of the type $\exp[-\mu(E) \times \frac{R}{2}]$, where $\mu(E)$ is the energy-dependent photon-absorption coefficient, and R is the range of the protons. We also performed a solid-angle correction to obtain the effective detection efficiency. We verified that these effective efficiencies reproduced the correct relative γ -ray intensities for all decays studied. Figure 2 shows a spectrum obtained from the 50-MeV activation, with the principal γ rays identified.

To make a statement about the proton-induced xenon excesses, Bernatowicz *et al.* used a simple model. They assumed that only (p, n) reactions take place, and that they all have the same cross section per nucleon. Then, the relative yields of the resulting Xe isotopes would depend only on the Te isotopic abundances in the ores and on the β -decay branching ratios of the produced iodine to xenon. While for sufficiently low-energy protons it is true that only (p, n) reactions are possible, it can be seen from Table I that the Q values for the $^{128,130}\text{Te}(p, 3n)^{126,128}\text{I}$ and $^{130}\text{Te}(p, 5n)^{126}\text{I}$ reactions are not terribly high. The energy spectrum of protons underground is not well characterized. Thus it is reasonable to expect that these additional reactions could also contribute to the observed $^{126,128}\text{Xe}$ excesses. One can generalize the model of Bernatowicz *et al.* to include these reactions by assuming that, where they are energetically allowed, they have the same cross sections as those of the (p, n) reactions. The relative yields predicted by this model are shown in Table II.

The low-energy bombardments we did provide information on the separate contributions of the (p, n) , $(p, 3n)$, and $(p, 5n)$ reactions. Table III shows our results for the iodine thick-target yields, and the production cross sections for the high-energy experiments. The integrated beam current of the 50-MeV activation suffered a systematic error. Thus, for this activation we only quote relative yields. Table IV contains the relative Xe yields inferred from the data in Table III. We also present data

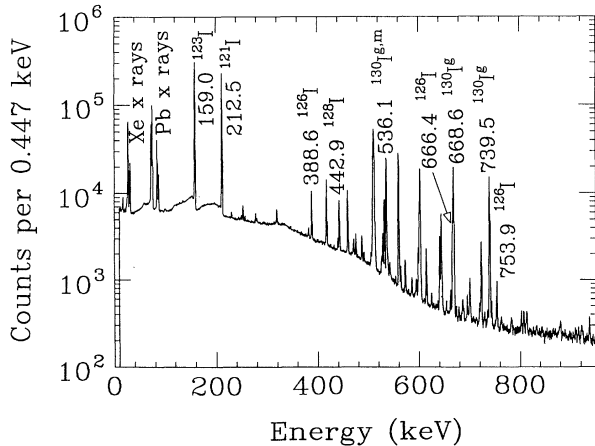


FIG. 2. Partial γ -ray spectrum observed from the 50-MeV proton activation.

TABLE I. Q values, in MeV, for the (p, xn) reactions of interest.

Target	(p, n)	$(p, 3n)$	$(p, 5n)$
^{126}Te	-2.9		
^{128}Te	-2.0	-18.0	
^{130}Te	-1.2	-16.5	-32.5

calculated from thick-target iodine yields of Roughton *et al.* [14].

Based on their model, Bernatowicz *et al.* inferred $N(^{128}\text{Xe})/N(^{126}\text{Xe})=3.47$ (this becomes 3.56 if we assume equal cross sections instead of equal cross sections per nucleon, and take the Te isotopic abundances and I β^- -decay branches from Ref. [11]). If we assume that (p, xn) reactions are the sole source of iodine and xenon in the ore, we would expect the relative Xe yields to be roughly given by the experimental ratios in Table IV. However, the appropriate value to use is obviously dependent on the proton energy spectrum. Table IV shows that even for energies between 3 and 15 MeV, the 3.56 ratio is only achieved in a very narrow energy range, a more appropriate value lying between ≈ 1 and ≈ 3 .

The muon reaction $\mu^\pm + ^A\text{Te} \rightarrow \mu^\pm + \pi^\mp + ^A\text{I}$ is the equivalent of a (p, n) reaction. Bernatowicz *et al.* estimated that proton- and muon-induced reactions contribute roughly equal amounts of Xe. We can take into account these direct muon reactions keeping the ratios of ^{128}Xe and ^{130}Xe with respect to ^{126}Xe as 3.56 and 4.08, respectively (see Table II), for their contribution. For the corresponding ratios of xenon isotopes produced from secondary proton reactions, we take 1.5 and 0.4, which are values that lie within the range of our experimental results. This yields a total ratio of 2.0, assuming that muon- and proton-induced reactions produce the same amount of xenon. In this case, the half-life ratio becomes $(3.74 \pm 0.10) \times 10^{-4}$, which is consistent with the result obtained by Bernatowicz *et al.* for the deeply buried Kalgoolie krennerite sample.

In conclusion, we find that the ratio of cosmic-ray produced $^{128}\text{Xe}/^{126}\text{Xe}$ is energy dependent. The simple model developed by Bernatowicz *et al.*, and generalized by us, fails to reproduce the measured ratios at any energy. Combining these results with the fact that the energy spectrum of underground protons is not well known, leads to a reduction in the cosmic-ray correction to the ratio of double- β decay half-lives of $^{128,130}\text{Te}$. Our experimental results and our generalization of Bernatowicz *et al.*'s model suggest that this correction could drop from the 12% value used by Bernatowicz *et al.* to 6%. With our half-life ratio of 3.74×10^{-4} and the data from Bernatowicz *et al.* for ^{130}Te half-life, we calculate a half-

TABLE II. Relative production of Xe in a simple model in which it is assumed that, where they are energetically allowed, all (p, xn) reactions have the same cross section.

	$x = 1$	$x = 1, 3$	$x = 1, 3, 5$
$N(^{128}\text{Xe})/N(^{126}\text{Xe})$	3.56	2.76	1.65
$N(^{130}\text{Xe})/N(^{126}\text{Xe})$	4.08	1.53	0.92

TABLE III. Our results for the $^{126,128,130}\text{I}$ thick-target yields, together with the (p, xn) cross sections of the high-energy activations.

E_p (MeV)	^{126}I	^{128}I	^{130}I
	(10 ⁻⁴ nuclei/proton or mb) ^a		
15	1.31(2)	1.08(1)	0.42(1) <i>g</i> ^b
			0.70(1) <i>m</i>
30	13.5(1)	11.0(1)	1.04(3) <i>g</i>
			0.98(2) <i>m</i>
45	23.6(5)	13.8(1)	1.1(1) <i>g</i>
			0.90(6) <i>m</i>
1850	6.9(23)	5.7(4)	3.0(1) ^c
5000	15(6)	10(2)	4.6(9) ^c

^aThe values quoted for the 1850- and 5000-MeV activations are cross sections, in mb, for the (p, xn) reactions on natural Te.

^b“*g*” and “*m*” identify the separate yields for the ground and metastable states of ^{130}I .

^cThe decay of the 9-min isomer was not observed in the high-energy experiments. These cross sections were calculated from the observed activities of the ground state only, being a combination of the value for the ground state, plus 83% of that for the isomer, corresponding to the branching ratio for the isomeric transition.

life of $(7.2 \pm 0.3) \times 10^{24}$ yr for ^{128}Te . The values for the upper limits of the nonstandard parameters are no more than 5% different from those obtained by Bernatowicz *et al.*, i.e., 2.4 eV for the effective neutrino mass, $\langle m_\nu \rangle$, 6.0×10^{-8} for the mixing of left- and right-handed weak currents, $\langle \eta \rangle$, and 4.9×10^{-6} for the pure right-handed weak-current term $\langle \lambda \rangle$, as calculated in the manner of Suhonen *et al.* [15]. If we calculate the neutrino mass limit by taking both, $\langle \eta \rangle$ and $\langle \lambda \rangle$ as zero, we then get

TABLE IV. Relative Xe production from the thick-target I yields of Ref. [14] (upper part) together with our data (lower part), $^{126}\text{Xe} \equiv 1$. The numbers in parentheses are the uncertainties in our measurements.

E_p (MeV)	^{128}Xe	^{130}Xe
3.029	0.75	2.15
3.346	3.70	4.43
3.934	2.34	3.61
4.292	2.93	4.07
4.645	2.00	2.92
5.049	1.75	2.80
5.687	2.42	3.86
6.082	2.81	3.17
6.360	1.87	3.51
15	1.76(9)	1.95(10)
30	1.74(8)	0.34(2)
45	1.25(6)	0.19(2)
50	0.94(22)	0.17(5)
1850	1.4(5)	0.39(14)
5000	1.1(5)	0.28(13)

$$\langle m_\nu \rangle \leq 1.5 \text{ eV.}$$

We wish to thank T. Bernatowicz for useful discussions and for providing the results of his measurements prior to publication. We also wish to thank the Di-Lepton Spectrometer Group for their assistance in the high-energy proton activations. This work was supported by the U.S. Department of Energy, Nuclear Physics Division, via Grant Nos. DE-AC03-76SF00098 and DE-FG05-87ER40314. M. T. F. da Cruz was supported by Fundação de Amparo à Pesquisa do Estado de São Paulo, FAPESP, São Paulo, Brasil. D. F. Rossi was supported under the DOE-TRAC Fellowship Program from Allen D. Nease High School, St. Augustine, Florida.

- [1] M.G. Inghram and J.H. Reynolds, Phys. Rev. **76**, 1265 (1949); **78**, 822 (1950).
- [2] O.K. Manuel, J. Phys. G **17**, S221 (1991), and references therein.
- [3] T. Bernatowicz, J. Brannon, R. Brazzle, R. Cowsik, C. Hohenberg, and F. Podosek, Phys. Rev. Lett. **69**, 2341 (1992); Phys. Rev. C **47**, 806 (1993).
- [4] M.K. Moe, University of California at Irvine Report UCI-NEUTRINO 93-1, 1993.
- [5] M. Aguilar-Benitez *et al.*, The Particle Data Group, Phys. Rev. D **45**, S1 III.23 (1992).
- [6] H.V. Klapdor-Kleingrothaus, Max-Planck Institut für Kernphysik Report MPI-H-V 34-92, 1992.
- [7] H. Bilokon, A. Castellina, B. D’Ettore Piazzoli, G. Mannocchi, P. Picchi, and S. Vernetto, Nucl. Instrum. Methods Phys. Res. A **303**, 381 (1991).
- [8] A.R. Smith, J.B. McCaslin, J.V. Geaga, J.C. Hill, and J.P. Vary, Phys. Rev. C **28**, 1614 (1983).
- [9] D.L. Olson, B.L. Berman, D.E. Greiner, H.H. Heckman, P.J. Lindstrom, and H.J. Crawford, Phys. Rev. C **28**, 1602 (1983).
- [10] The fitting program GELIFIT obtained from Argonne National Laboratory, and D.C. Radford, private communication.
- [11] E. Browne and R.B. Firestone, *Table of Radioactive Isotopes* (Wiley, New York, 1986).
- [12] K. Debertin and R.G. Helmer, *Gamma- and X-Ray Spectrometry with Semiconductor Detectors* (North-Holland, Amsterdam, 1988).
- [13] J.H. Hubbell, At. Data **3**, 241 (1971). From this work we took the total photon-absorption cross sections for Sn and Xe, and used their geometric average as the Te cross section.
- [14] N.A. Roughton, M.R. Fritts, R.J. Peterson, C.S. Zaidins, and C.J. Hansen, At. Data Nucl. Data Tables **23**, 177 (1979).
- [15] J. Suhonen, S.B. Khadkikar, and A. Faessler, Nucl. Phys. **A535**, 509 (1991).