

Designation: E523 - 16

Standard Test Method for Measuring Fast-Neutron Reaction Rates by Radioactivation of Copper¹

This standard is issued under the fixed designation E523; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon (ε) indicates an editorial change since the last revision or reapproval.

1. Scope

- 1.1 This test method covers procedures for measuring reaction rates by the activation reaction $^{63}\text{Cu}(n,\alpha)^{60}\text{Co}$. The cross section for ^{60}Co produced in this reaction increases rapidly with neutrons having energies greater than about 4.5 MeV. ^{60}Co decays with a half-life of 1925.27 days (±0.29 days)(1)² and emits two gamma rays having energies of 1.1732278 and 1.332492 MeV(1). The isotopic content of natural copper is 69.17 % ^{63}Cu and 30.83 % $^{65}\text{Cu}(2)$. The neutron reaction, $^{63}\text{Cu}(n,\gamma)^{64}\text{Cu}$, produces a radioactive product that emits gamma rays (1.34577 MeV (E1005)) which might interfere with the counting of the ^{60}Co gamma rays.
- 1.2 With suitable techniques, fission-neutron fluence rates above $10^9~{\rm cm^{-2}\cdot s^{-1}}$ can be determined. The $^{63}Cu(n,\alpha)^{60}Co$ reaction can be used to determine fast-neutron fluences for irradiation times up to about 15 years, provided that the analysis methods described in Practice E261 are followed. If dosimeters are analyzed after irradiation periods longer than 15 years, the information inferred about the fluence during irradiation periods more than 15 years before the end of the irradiation should not be relied upon without supporting data from dosimeters withdrawn earlier.
- 1.3 Detailed procedures for other fast-neutron detectors are referenced in Practice E261.
- 1.4 This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.

2. Referenced Documents

- 2.1 ASTM Standards:³
- E170 Terminology Relating to Radiation Measurements and Dosimetry
- E181 Test Methods for Detector Calibration and Analysis of Radionuclides
- E261 Practice for Determining Neutron Fluence, Fluence Rate, and Spectra by Radioactivation Techniques
- E844 Guide for Sensor Set Design and Irradiation for Reactor Surveillance, E 706 (IIC)
- E944 Guide for Application of Neutron Spectrum Adjustment Methods in Reactor Surveillance, E 706 (IIA)
- E1005 Test Method for Application and Analysis of Radiometric Monitors for Reactor Vessel Surveillance
- E1018 Guide for Application of ASTM Evaluated Cross Section Data File, Matrix E706 (IIB)

3. Terminology

- 3.1 Definitions:
- 3.1.1 Refer to Terminology E170.

4. Summary of Test Method

- 4.1 High-purity copper (<1 ppm cobalt) is irradiated in a neutron field, thereby producing radioactive 60 Co from the 63 Cu(n, α) 60 Co reaction.
- 4.2 The gamma rays emitted by the radioactive decay of ⁶⁰Co are counted in accordance with Test Methods E181 and the reaction rate, as defined by Practice E261, is calculated from the decay rate and irradiation conditions.
- 4.3 The neutron fluence rate above about 4.5 MeV can then be calculated from the spectral-weighted neutron activation cross section as defined by Practice E261.

¹ This test method is under the jurisdiction of ASTM Committee E10 on Nuclear Technology and Applications and is the direct responsibility of Subcommittee E10.05 on Nuclear Radiation Metrology.

Current edition approved Dec. 1, 2016. Published January 2017. Originally approved in 1976. Last previous edition approved in 2011 as E523 – 11. DOI: 10.1520/E0523-16.

² The boldface numbers in parentheses refer to a list of references at the end of this standard.

³ For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

5. Significance and Use

- 5.1 Refer to Guide E844 for the selection, irradiation, and quality control of neutron dosimeters.
- 5.2 Refer to Practice E261 for a general discussion of the measurement of fast neutron fluence rate with threshold detectors. The general shape of the 63 Cu(n, α) 60 Co cross section is also shown in Fig. 1 (3, 4) along with a comparison to the current experimental database (5). This figure is for illustrative purposes only to indicate the range of the response of the 63 Cu(n, α) 60 Co reaction. Refer to Guide E1018 for descriptions of recommended tabulated dosimetry cross sections.

Note 1—The cross section appropriate for use under this standard is ENDF/B-VI release 8 library since it contains a covariance matrix. The ENDF/B-VII library has been released, but it does not contain a covariance matrix for this reaction. For dosimetry applications, an uncertainty metric expressed as a covariance matrix is required. See Guide E1018.

- 5.3 The major advantages of copper for measuring fast-neutron fluence rate are that it has good strength, is easily fabricated, has excellent corrosion resistance, has a melting temperature of 1083°C, and can be obtained in high purity. The half-life of ⁶⁰ Co is long and its decay scheme is simple and well known.
- 5.4 The disadvantages of copper for measuring fast neutron fluence rate are the high reaction apparent threshold of 5 MeV, the possible interference from cobalt impurity (>1 $\mu g/g$), the reported possible thermal component of the (n, α) reaction, and the possibly significant cross sections for thermal neutrons for

⁶³Cu and ⁶⁰Co (that is 4.50 and 2.0 barns, respectively),(6) which will require burnout corrections at high fluences.

6. Apparatus

- 6.1 NaI(Tl) or High Resolution Gamma-Ray Spectrometer—Because of its high resolution, the germanium detector is useful when contaminant activities are present or when it is necessary to analyze before the 12.701-h half-life ⁶⁴Cu has decayed away.
- 6.2 Precision Balance, able to achieve the required accuracy.

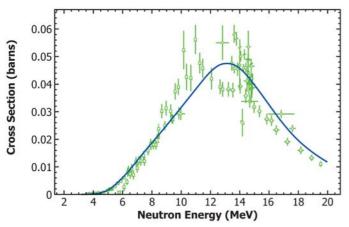


FIG. 1 63 Cu(n, α) 60 Co Cross Section with EXFOR Experimental Data

7. Materials

- 7.1 *Copper Metal*—Pure copper metal in the form of wire or foil is available.
- 7.1.1 The metal should be tested for impurities by a neutron activation technique. If the measurement is to be made in a thermal-neutron environment, there must be no cobalt impurity (<1 μ g/g) because the reaction 59 Co(n, γ) 60 Co produces the same product as produced in the subject reaction. To reduce this interference, the use of a thermal-neutron shield during irradiation would be advisable if cobalt impurity is suspected.
- 7.2 Encapsulating Materials—Brass, stainless steel, copper, aluminum, quartz, or vanadium have been used as primary encapsulating materials. The container should be constructed in such a manner that it will not create significant flux perturbation and that it may be opened easily, especially if the capsule is to be opened remotely (see Guide E844).

8. Procedure

- 8.1 Decide on the size and shape of the copper sample to be irradiated, taking into consideration the size and shape of the irradiation space. The mass and exposure time are parameters that can be varied to obtain a desired disintegration rate for a given neutron fluence rate level (see Guide E844).
 - 8.2 Weigh the sample.
- 8.3 Irradiate the sample for the predetermined time period. Record the power level and any changes in power during the irradiation, the time at the beginning and end of the irradiation, and the relative position of the monitors in the irradiation facility.
- 8.4 A waiting period of about 6 days is recommended between termination of the exposure and analyzing the sample for ⁶⁰Co content. This allows the 12.701-h ⁶⁴Cu(1) to decay so that there is no interference from the gamma rays emitted by ⁶⁴Cu, that is, the 0.511 and 1.34577 MeV gamma rays(1). However, analysis may be performed sooner if a suitable gamma-ray or peak analysis technique is used.
- 8.5 Check the sample for activity from cross-contamination by other irradiated materials. Clean, if necessary and reweigh.
- 8.6 Analyze the sample for ⁶⁰Co content in disintegrations per second using the gamma-ray spectrometer (see Test Methods E181 and E1005).
- 8.7 Disintegration of ⁶⁰Co nuclei produces 1.173228 MeV and 1.332492 MeV gamma rays with probabilities per decay of 0.9985 and 0.999826 respectively.(1) When analyzing either peak in the gamma-ray spectrum, a correction for coincidence summing may be required if the sample is placed close to the detector (10 cm or less) (see Test Methods E181).

9. Calculations

9.1 Calculate the saturation activity A_s , as follows:

$$A_{s} = \frac{A}{(1 - e^{-\lambda t_{i}}) \cdot e^{-\lambda t_{w}}} \tag{1}$$

where:

 $A = {}^{60}$ Co disintegrations per second measured by counting,



 λ = decay constant for ${}^{60}\text{Co} = 4.16697 \times 10^{-9} \text{ s}^{-1}$,

 t_i = irradiation duration s,

t_w = elapsed time between the end of irradiation and counting, s.

Note 2—The equation for $A_{\rm s}$ is valid if the reactor operated at essentially constant power and if corrections for other reactions (for example, impurities, burnout, etc.) are negligible. Refer to Practice E261 for more generalized treatments.

9.2 Calculate the reaction rate, R_s , as follows:

$$R_{s} = A_{s}/N_{o} \tag{2}$$

where:

 $A_{\rm s}$ = saturation activity, and

 N_0 = number of ⁶³Cu atoms.

9.3 Refer to Practice E261 and Guide E944 for a discussion of fast-neutron fluence rate and fluence.

10. Report

10.1 Practice E261 describes how data should be reported.

11. Precision and Bias

Note 3—Measurement uncertainty is described by a precision and bias statement in this standard. Another acceptable approach is to use Type A and B uncertainty components(7,8). This Type A/B uncertainty specification is now used in International Organization for Standardization (ISO) standards and this approach can be expected to play a more prominent role in future uncertainty analyses.

- 11.1 General practice indicates that disintegration rates can be determined with a bias of \pm 3 % (1 σ %) and with a precision of \pm 1 % (1 σ %).
- 11.2 When this measured activity is used in conjunction with a neutron spectrum to determine a neutron fluence, the energy-dependent uncertainty of the spectrum and the cross section are important considerations. The energy-dependent uncertainty, expressed as a percentage, for the 63 Cu(n, α) 60 Co cross section is shown in Fig. 2(3)
- 11.3 Test results have been reported in well characterized neutron benchmark fields.

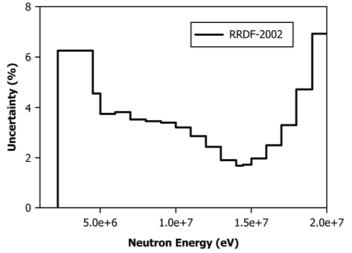


FIG. 2 Energy-dependent Uncertainty (%) for the 63 Cu(n,a) 60 Co Cross Section

11.3.1 In the 252 Cf spontaneous fission reference neutron field, the measured cross section is 0.6887 b \pm 1.9 % (9) and the calculated cross section using the RRDF-2002 cross section is 0.69248 b with a spectrum integrated cross section uncertainty of 1.399 % (3) and a spectrum characterization uncertainty of 1.37 %. This results in a calculated-to-experimental (C/E) ratio of 1.0055 \pm 2.73 %.

11.3.2 In the 235 U thermal neutron field, the measured cross section is 0.4935 b \pm 4.9 % (10) and the calculated cross section using the RRDF-2002 cross section is 0.53294 b with a spectrum integrated cross section uncertainty of 1.461 % (3) and a spectrum characterization uncertainty of 6.042 %. This results in a calculated-to-experimental (C/E) ratio of 1.08 \pm 7.92 %.

12. Keywords

12.1 activation; activation reaction; copper; cross section; dosimetry; fast-neutron monitor; neutron metrology; pressure vessel surveillance; reaction rate; 63 Cu(n, α) 60 Co

REFERENCES

- (1) Update of X-ray and Gamma Ray Decay Data Standards for Detector Calibration and Other Applications: Vol 1: Recommended Decay Data, High Energy Gamma Ray Standards and Angular Correlation Coefficients, International Atomic Energy Agency, Vienna, report STI/PUB/1287, 2007.
- (2) Nuclear Wallet Cards, compiled by J. K. Tuli, National Nuclear Data Center, April 2005.
- (3) Zolotarev, K. I., Ignatyuk, A. V., Mahokhin, V. N., Pashchenco, A. B., "RRDF-98 Russian Reactor Dosimetry File, report IAEA-NDS-193, March 1999. The last full release was in 1998. Updated versions corresponding to the RRDF-2002 library have been incorporated into the IRDF-2002 dosimetry library.
- (4) International Reactor Dosimetry File 2002 (IRDF-2002, IAEA Technical Report Series No. 452, International Atomic Energy Agency, Vienna, Austria, 2006. Available at: http://www-nds.iaea.org/irdf2002/index.html.
- (5) "EXFOR Formats Description for Users (EXFOR Basics)," report

- IAEA-NDS-206, International Atomic Energy Agency, Vienna, Austria, June 2008. On-line database available at URL: http://www-nds.iaea.org/indg-nexp.html. Data here as present on January 2011.
- (6) Mughabghab, S. F., Atlas of Neutron Resonances: Resonance Parameters and Thermal Cross Sections Z=1-100, Elsevier, Amsterdam, 2006.
- (7) Guide to the Expression of Uncertainty in Measurement, International Organization for Standardization, 1995, ISBN 92-67-10188-9.
- (8) Taylor, B. N. and Kuyatt, C. E., Guidelines for Evaluating and Expressing the Uncertainty of NIST Measurement Results, NIST Technical Note 1297, National Institute of Standards and Technology, Gaithersburg, MD, 1994.
- (9) Mannhart, W., Validation of Differential Cross Sections with Integral Data, Report INDC(NDS)-435, pp. 59-64, IAEA, Vienna, September 2002.
- (10) Mannhart, W., Progress Report INDC(Ger)-045, pp. 40-43, 1999.



ASTM International takes no position respecting the validity of any patent rights asserted in connection with any item mentioned in this standard. Users of this standard are expressly advised that determination of the validity of any such patent rights, and the risk of infringement of such rights, are entirely their own responsibility.

This standard is subject to revision at any time by the responsible technical committee and must be reviewed every five years and if not revised, either reapproved or withdrawn. Your comments are invited either for revision of this standard or for additional standards and should be addressed to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee, which you may attend. If you feel that your comments have not received a fair hearing you should make your views known to the ASTM Committee on Standards, at the address shown below.

This standard is copyrighted by ASTM International, 100 Barr Harbor Drive, PO Box C700, West Conshohocken, PA 19428-2959, United States. Individual reprints (single or multiple copies) of this standard may be obtained by contacting ASTM at the above address or at 610-832-9585 (phone), 610-832-9555 (fax), or service@astm.org (e-mail); or through the ASTM website (www.astm.org). Permission rights to photocopy the standard may also be secured from the Copyright Clearance Center, 222 Rosewood Drive, Danvers, MA 01923, Tel: (978) 646-2600; http://www.copyright.com/