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**Practice for use of a radiochromic film  
dosimetry system**

Pratique de l'utilisation d'un système dosimétrique à film  
radiochromique

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## Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75% of the member bodies casting a vote.

ASTM International is one of the world's largest voluntary standards development organizations with global participation from affected stakeholders. ASTM technical committees follow rigorous due process balloting procedures.

A pilot project between ISO and ASTM International has been formed to develop and maintain a group of ISO/ASTM radiation processing dosimetry standards. Under this pilot project, ASTM Subcommittee E10.01, Dosimetry for Radiation Processing, is responsible for the development and maintenance of these dosimetry standards with unrestricted participation and input from appropriate ISO member bodies.

Attention is drawn to the possibility that some of the elements of this International Standard may be the subject of patent rights. Neither ISO nor ASTM International shall be held responsible for identifying any or all such patent rights.

International Standard ISO/ASTM 51275 was developed by ASTM Committee E10, Nuclear Technology and Applications, through Subcommittee E10.01, and by Technical Committee ISO/TC 85, Nuclear Energy.



## Standard Practice for Use of a Radiochromic Film Dosimetry System<sup>1</sup>

This standard is issued under the fixed designation ISO/ASTM 51275; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision.

### 1. Scope

1.1 This practice covers the handling, testing, and procedure for using a radiochromic film dosimetry system to measure absorbed dose in materials irradiated by photons or electrons in terms of absorbed dose in water.

1.2 This practice applies to radiochromic film dosimeters that can be used within part or all of the specified ranges as follows:

1.2.1 The absorbed dose range is 1 Gy to 100 kGy.

1.2.2 The absorbed dose rate is  $1 \times 10^{-2}$  to  $1 \times 10^{13}$  Gy/s (1-4).<sup>2</sup>

1.2.3 The radiation energy range for both photons and electrons is 0.1 to 50 MeV.

1.2.4 The irradiation temperature range is  $-78$  to  $+60^\circ\text{C}$ .

1.3 This practice applies to radiochromic films of various formats, including small pieces used to measure a single dose value, strips used for one-dimensional dose-mapping, and sheets used for two-dimensional dose-mapping. Three-dimensional dose-mapping may be achieved by proper placement of any of these formats.

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:

E 170 Terminology Relating to Radiation Measurements and Dosimetry<sup>3</sup>

E 178 Practice for Dealing with Outlying Observations<sup>4</sup>

E 275 Practice for Describing and Measuring Performance of Ultraviolet, Visible, and Near Infrared Spectrophotometers<sup>5</sup>

E 456 Terminology Relating to Quality and Statistics<sup>4</sup>

E 668 Practice for Application of Thermoluminescence-

Dosimetry (TLD) Systems for Determining Absorbed Dose in Radiation-Hardness Testing of Electronic Devices<sup>3</sup>

E 925 Practice for the Periodic Calibration of Narrow Band-Pass Spectrophotometers<sup>5</sup>

E 958 Practice for Measuring Practical Spectral Bandwidth of Ultraviolet-Visible Spectrophotometers<sup>5</sup>

E 1026 Practice for Using the Fricke Reference Standard Dosimetry System<sup>3</sup>

#### 2.2 ISO/ASTM Standards:

51204 Practice for Dosimetry in Gamma Irradiation Facilities for Food Processing<sup>3</sup>

51205 Practice for Use of a Ceric-Cerous Sulfate Dosimetry System<sup>3</sup>

51261 Guide for Selection and Calibration of Dosimetry Systems for Radiation Processing<sup>3</sup>

51707 Guide for Estimating Uncertainties in Dosimetry for Radiation Processing<sup>3</sup>

2.3 *International Commission on Radiation Units and Measurements (ICRU) Reports:*<sup>6</sup>

ICRU Report 14 Radiation Dosimetry: X-Rays and Gamma Rays with Maximum Photon Energies Between 0.6 and 50 MeV

ICRU Report 17 Radiation Dosimetry: X-Rays Generated at Potentials of 5 to 150 kV

ICRU Report 34 The Dosimetry of Pulsed Radiation

ICRU Report 35 Radiation Dosimetry: Electron Beams with Energies Between 1 and 50 MeV

ICRU Report 60 Radiation Quantities and Units

### 3. Terminology

#### 3.1 Definitions:

3.1.1 *analysis wavelength*—wavelength used in a spectrophotometric instrument for the measurement of optical absorbance.

3.1.2 *calibration curve*—graphical representation of the dosimetry system's response function.

3.1.3 *dosimeter batch*—quantity of dosimeters made from a specific mass of material with uniform composition, fabricated in a single production run under controlled, consistent conditions and having a unique identification code.

3.1.4 *dosimetry system*—a system used for determining absorbed dose, consisting of dosimeters, measurement instruments and their associated reference standards, and procedures for the system's use.

3.1.5 *measurement quality assurance plan*—a documented program for the measurement process that ensures on a

<sup>1</sup> This guide is under the jurisdiction of ASTM Committee E10 on Nuclear Technology and Applications and is the direct responsibility of Subcommittee E10.01 on Dosimetry for Radiation Processing, and is also under the jurisdiction of ISO/TC 85/WG 3.

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<sup>2</sup> The boldface number in parentheses refer to the bibliography at the end of this practice.

<sup>3</sup> *Annual Book of ASTM Standards*, Vol 12.02.

<sup>4</sup> *Annual Book of ASTM Standards*, Vol 14.02.

<sup>5</sup> *Annual Book of ASTM Standards*, Vol 03.06.

<sup>6</sup> Available from the International Commission on Radiation Units and Measurements, 7910 Woodmont Ave., Suite 800, Bethesda, MD 20814, USA.



continuing basis that the overall uncertainty meets the requirements of the specific application. This plan requires traceability to, and consistency with, nationally or internationally recognized standards.

3.1.6 *net absorbance*,  $\Delta A$ —change in measured optical absorbance at a selected wavelength determined as the absolute difference between the pre-irradiation absorbance,  $A_0$ , and the post-irradiation absorbance,  $A$ , as follows:

$$\Delta A = |A - A_0|. \quad (1)$$

3.1.6.1 *Discussion*—In practice, an average pre-irradiation absorbance,  $\bar{A}_0$ , may be used to determine net absorbance.

3.1.7 *radiochromic film-dosimeter*—specially prepared film containing ingredients that undergo change in optical absorbance under ionizing radiation. This change in optical absorbance can be related to absorbed dose in water.

3.1.8 *response function*—mathematical representation of the relationship between dosimeter response and absorbed dose for a given dosimetry system.

3.1.9 *specific net absorbance* ( $\Delta k$ )—Net absorbance,  $\Delta A$ , at a selected wavelength divided by the optical pathlength,  $d$ , through the dosimeter material as follows:

$$\Delta k = \Delta A/d. \quad (2)$$

3.2 Definitions of other terms used in this standard that pertain to radiation measurement and dosimetry may be found in ASTM Terminology Standard E 170. Definitions in ASTM E 170 are compatible with ICRU 60; that document, therefore, may be used as an alternative reference.

#### 4. Significance and Use

4.1 The radiochromic film dosimetry system provides a means of determining absorbed dose in materials. Under the influence of ionizing radiation, chemical reactions take place in the radiochromic film creating or enhancing, or both, optical absorption bands. Absorbance is determined within these radiation-induced absorption bands using a spectrophotometer or photometer (See 5.1.2).

4.2 In the application of a specific dosimetry system, absorbed dose is determined by use of a calibration curve traceable to national standards.

4.3 The absorbed dose determined is usually specified in water. Absorbed dose in other materials may be determined by applying the conversion factors discussed in ISO/ASTM Guide 51261.

NOTE 1—For comprehensive discussion of various dosimetry methods applicable to the radiation types and energies discussed in this practice, see ICRU Reports 14, 17, 34, and 35.

4.4 Radiochromic film dosimetry systems are commonly applied in the industrial radiation processing of a variety of products, for example, sterilization of medical devices and processing of foods (11, 13).

#### 5. Apparatus

5.1 *Components of the Dosimetry System*—The following shall be used to determine absorbed dose with radiochromic film dosimetry systems:

5.1.1 *Radiochromic Film Dosimeters*:

5.1.2 *Spectrophotometer or Photometer*, having documentation covering analysis wavelength range, accuracy of wavelength selection, absorbance determination, spectral bandwidth, and stray light rejection. Examples of appropriate wavelengths for analysis for specific dosimetry systems are provided by the manufacturer and in Refs. (3-14, 19).

5.1.3 *Holder*, to position the dosimeter reproducibly in and perpendicular to the measuring light beam.

5.1.4 *Calibrated Thickness Gage*, with a precision of  $\pm 2\%$  of the film thickness (at a 95 % confidence level), if the film's thickness is to be measured.

NOTE 2—Documentation provided by the manufacturer of the radiochromic film dosimeter with regard to the film thickness and its variability may be substituted for direct measurement of thickness by the user. This information should be verified by the user by analyzing a representative sample of films.

NOTE 3—Some radiochromic film dosimeters contain a substrate which is not radiochromic. With such dosimeters the thickness is not measured.

5.1.5 Packaging materials for radiochromic films, where applicable.

#### 6. Performance Check of Instrumentation

6.1 The performance of the photometer or spectrophotometer shall be checked as specified in Section 7.4, and documented.

6.1.1 When using a spectrophotometer, check and document the accuracy of the wavelength scale and absorbance scale at or near the analysis wavelength(s) at intervals not to exceed one month during periods of use and as specified by the end-user's internal procedures.

6.1.2 When using a photometer, check and document the accuracy of the absorbance scale at intervals not to exceed one month during periods of use and as specified by the end-user's internal procedures.

6.1.3 Compare the information obtained in 6.1.1 or 6.1.2 with the original instrument specifications to ensure adequate performance.

6.2 Check the thickness gage prior to first use and periodically thereafter to assure reproducibility and lack of zero drift. Check and document the calibration of the gage at intervals not to exceed six months. Use gage blocks, traceable to national standards for this purpose.

#### 7. Calibration of the Dosimetry System

7.1 Prior to use, the dosimetry system shall be calibrated in accordance with the user's documented procedure that specifies details of the calibration process and quality assurance requirements. This calibration procedure shall be repeated at regular intervals to ensure that the accuracy of the absorbed dose measurement is maintained within required limits. Detailed calibration procedures are provided in ISO/ASTM Guide 51261.

7.2 *Calibration of Dosimeters*—Irradiation is a critical component of the calibration of the dosimetry system. Calibration irradiations may be performed in several ways, including irradiating the dosimeters using:

7.2.1 a calibration facility that provides an absorbed dose or an absorbed-dose rate having measurement traceability to



nationally or internationally recognized standards, or

7.2.2 an in-house calibration facility that provides an absorbed dose or an absorbed-dose rate having measurement traceability to nationally or internationally recognized standards, or

7.2.3 a production or research irradiation facility together with reference or transfer-standard dosimeters that have measurement traceability to nationally or internationally recognized standards.

7.3 When the radiochromic film dosimeter is used as a transfer-standard dosimeter, the calibration irradiation may be performed only as stated in 7.2.1, and not as stated in 7.2.2 or 7.2.3.

7.4 *Instrument Calibration*—Calibrations of the individual instruments used in the analysis of the dosimeters shall be verified at periodic intervals. These calibrations shall be traceable to nationally or internationally recognized standards where available. For example, if an optical absorbance-measuring instrument such as a spectrophotometer or densitometer is used, then appropriate standards shall be used to verify the accuracy of the optical absorbance at a specified wavelength(s). (See ASTM Practices E 275, E 925, and E 958.)

## 8. Procedure

### 8.1 *Examination and Storage Procedure:*

8.1.1 Ultraviolet radiation may cause the film to change color. Perform tests to assure that the handling/reading environment does not cause measurable color development. If needed, place ultraviolet filters over fluorescent lights or windows to reduce color development (see also 9.3.1).

8.1.2 Handle dosimeters according to manufacturer's instructions. Avoid handling dosimeter surfaces with bare fingers.

8.1.3 Visually inspect the films for imperfections. Gently dust the dosimeters if necessary. Discard any dosimeters that show imperfections such as scratches that could give rise to erroneous readings.

8.1.4 Identify the dosimeters.

8.1.5 Store the dosimeters according to the manufacturer's written recommendations.

### 8.2 *Irradiation Procedure:*

8.2.1 Determine the pre-irradiation absorbance,  $A_0$ , for each dosimeter film at the selected analysis wavelength. This may be done for each dosimeter or by use of an average  $\bar{A}_0$ , determined by reading several dosimeters and documenting the variation.

NOTE 4—If the orientation of the dosimeter has an effect on the absorbance reading, keep track of the dosimeter orientation (for example, by marking a corner or edge of the dosimeter).

8.2.2 If necessary, package the dosimeters to provide controlled environmental conditions during irradiation.

8.2.3 Mark the packaged dosimeters appropriately for identification.

8.2.4 Irradiate the dosimeters.

NOTE 5—The dosimeters may be irradiated in the product undergoing processing or in a medium of similar composition, or water, of appropriate dimensions so as to approximate electron equilibrium conditions. Such

equilibrium conditions may not exist within dosimeters placed throughout the product under actual processing conditions. This particularly is the case near interfaces of different materials. Irradiation under nonequilibrium conditions, such as on the surface of a product package, is often used to monitor the absorbed dose delivered to the product and may be related to absorbed dose within the product by correction factors under certain conditions. For a detailed discussion of this subject, see ISO/ASTM Guide 51261.

### 8.3 *Analysis Procedure:*

8.3.1 Avoid any exposure to ultraviolet radiation that may induce coloration of the dosimeter film (see 8.1.1).

8.3.2 Determine the post-irradiation absorbance,  $A$ , at the selected analysis wavelength(s).

8.3.3 If appropriate for the dosimeter being used, measure the thickness of each dosimeter film, or use the average thickness and document the variation.

NOTE 6—Certain films may be too thin to allow the accurate determination of thickness using conventional gage technology. In such cases, statistical methods may be employed in order to provide the uncertainty values discussed in Section 12.

8.3.4 Calculate the absorbed dose based on the pre- and post-irradiation absorbances, the thickness, if appropriate, and the calibration curve or response function.

NOTE 7—Examples of radiochromic film dosimeter applications that employ reflectance techniques or alternative methods for determining net and specific net absorbance appear in the literature. The user may choose to refer to these techniques (15, 16).

## 9. Characterization of Each Batch of Dosimeters

### 9.1 *Reproducibility of Specific Net Absorbance:*

9.1.1 For each batch of dosimeters, the reproducibility of specific net absorbance should be obtained by analyzing the data from the sets of dosimeters irradiated during the calibration at each dose value.

9.1.2 Use the sample standard deviation ( $S_{n-1}$ ) determined during calibration to calculate coefficients of variation ( $CV$ ) for each absorbed dose value as follows:

$$CV = \frac{S_{n-1}}{\Delta \bar{k}} \times 100 (\%) \quad (3)$$

9.1.3 Document these coefficients of variation and note any that are unusually large.

NOTE 8—In general, if the coefficients of variation values are greater than 2 %, then a redetermination of the data should be considered, or the batch of dosimeters should be rejected.

### 9.2 *Post-Irradiation Characterization:*

9.2.1 Some types of dosimeters may fade or may continue color development after irradiation. This effect may depend on post-irradiation storage conditions such as temperature, humidity, or atmosphere. In order to determine if this is significant in a given application, measure the absorbance at the selected wavelength(s) over the period of anticipated analysis and over the range of expected storage conditions (7, 8, 14, 15, 20).

9.2.2 If absorbances measured in 9.2.1 are found to vary significantly with post-irradiation storage time, then apply correction factors for such time-dependent variations, taking into account the calibration curve for that batch of dosimeters,



in order to minimize dosimetric errors during routine application.

9.2.3 For a given set of irradiation conditions, this procedure needs to be performed only once for a given batch of dosimeters.

### 9.3 Other Factors:

9.3.1 The effects of temperature, humidity, absorbed dose rate, incident energy spectrum, electron equilibrium and background ultraviolet radiation shall be taken into account. See Refs 2-6 and 11-15 for examples of the types and magnitudes of the effects. Appropriate written information regarding the magnitude and effect(s) upon the measurement made by the dosimetry system may be obtained from the scientific literature, dosimeter manufacturer, distributor, irradiation facility operator, or a qualified testing organization.

## 10. Application of Dosimetry System

10.1 The number of dosimeters required for the measurement of absorbed dose on or within a material is determined by the precision of the dosimetry system and the required precision associated with the application. Appendix X3 of Practice E 668 describes a statistical method for determining this number.

10.2 Follow the procedures in accordance with 8.2 and 8.3.

10.3 Determine the absorbed dose from the mean specific net absorbance values and the system calibration curve that results from following the procedures in accordance with Section 7.

10.4 Record the calculated absorbed dose and all other relevant data as outlined in Section 11.

## 11. Minimum Documentation Requirements

11.1 Record the dosimeter manufacturer, type, and batch number (code).

11.2 Record or reference the date of calibration, calibration source, and associated instruments used.

11.3 Record or reference the irradiation environmental conditions for the dosimeters, including temperature, pressure (if other than atmospheric), relative humidity, and composition of the surrounding atmosphere (if other than air).

11.4 Record the date of irradiation and the dates on which the non-irradiated and irradiated dosimeters are analyzed.

11.5 Record the absorbance and thickness data, if appropriate, and the resulting absorbed dose values.

11.6 Record or reference the uncertainty in the value of the absorbed dose.

11.7 Record or reference the measurement quality assurance plan used for the dosimetry system application.

## 12. Measurement Uncertainty

12.1 To be meaningful, a measurement of absorbed dose shall be accompanied by an estimate of uncertainty. Components of uncertainty shall be identified as belonging to one of two groups:

- (A) those which are evaluated by statistical methods, or
- (B) those which are evaluated by other means.

Additional information is given in ISO/ASTM Guide 51707 and references 17 and 18, where these components are referred to as Type A and Type B, respectively. In reporting uncertainty, other classifications such as precision and bias may be useful.

12.2 If this practice is followed, the estimate of the combined uncertainty of an absorbed dose determined by this dosimetry system should be within about  $\pm 6\%$  at a 95% confidence level (corresponding to a coverage factor of 2).

NOTE 9—The identification of Type A and Type B uncertainties is based on the methodology adopted in 1993 by the International Organization for Standardization (ISO) for estimating uncertainty. This is different from the way uncertainty has been traditionally expressed in terms of “precision” and “bias,” where precision is a measure of the extent to which replicate measurements made under specified conditions are in agreement, and bias is a systematic error (see ASTM Terminologies E 170 and E 456).

The purpose of using the method of expressing uncertainties as Type A and Type B recommended in the ISO Guide to the Expression of Uncertainty in Measurement (18) is to promote an understanding of how uncertainty statements are arrived at and to provide a basis for the international comparison of measurement results.

NOTE 10—ISO/ASTM Guide 51707 defines possible sources of error in dosimetry performed in radiation processing facilities and offers procedures for estimating the resulting magnitude of the uncertainties in the measurement results. Basic concepts of measurement, estimate of the measured value of a quantity, “true” value, error and uncertainty are defined and discussed. Components of uncertainty are discussed and methods are given for evaluating and estimating their values. Their contributions to the standard uncertainty in the reported values of absorbed dose are considered and methods are given for calculating the combined standard uncertainty and an estimate of overall (expanded) uncertainty.

## 13. Keywords

13.1 absorbed dose; dose measurement; dosimeter; dosimetry; electron beam; gamma radiation; ionizing radiation; quality control; radiation processing; radiochromic dosimetry; radiochromic film; ICS 17.240



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