
**Practice for use of calorimetric
dosimetry systems for dose
measurements and dosimetry system
calibration in electron beams**

*Pratique de l'utilisation des systèmes dosimétriques calorimétriques
pour pour des mesures de dose délivrée par un faisceau d'électrons et
pour l'étalonnage de dosimètres*

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Contents

	Page
1 Scope	1
2 Referenced Documents	1
3 Terminology	2
4 Significance and use	2
5 Interferences	3
6 Apparatus	3
7 Calibration procedures	4
8 Dose measurement procedures	5
9 Calibration of other dosimetry systems	6
10 Documentation	7
11 Measurement uncertainty	7
12 Keywords	7
Annexes	7
Figure 1 Example of a polystyrene calorimeter used for routine measurements at a 10-MeV industrial electron accelerator	3
Figure 2 Absorber (phantom) for irradiation at 10 MeV electron irradiation facility of routine and transfer-standard dosimeters (10). Material: Polystyrene	4
Figure 3 Example of measurements of temperature of a graphite calorimeter before and after irradiation (7)	6
Figure 4 Example of on-line measurements of a graphite calorimeter (5)	6
Table 1 Measurement uncertainties of routine polystyrene calorimetric dosimetry systems from Risø high dose reference laboratory (in percent, at $k = 2$) (9)	7
Table A1.1 Results for alanine and calorimeter dose measurements	8
Table A2.1 Thickness and size of several graphite calorimetric bodies designed at NIST for use at specific electron energies	8

Foreword

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Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at www.iso.org/members.html.

ISO/ASTM 51631:2020(E)



Standard Practice for Use of Calorimetric Dosimetry Systems for Dose Measurements and Routine Dosimetry System Calibration in Electron Beams¹

This standard is issued under the fixed designation ISO/ASTM 51631; 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 preparation and use of semi-adiabatic calorimetric dosimetry systems for measurement of absorbed dose and for calibration of routine dosimetry systems when irradiated with electrons for radiation processing applications. The calorimeters are either transported by a conveyor past a scanned electron beam or are stationary in a broadened beam.

1.2 This document is one of a set of standards that provides recommendations for properly implementing dosimetry in radiation processing, and describes a means of achieving compliance with the requirements of ISO/ASTM Practice 52628 for a calorimetric dosimetry system. It is intended to be read in conjunction with ISO/ASTM Practice 52628.

1.3 The calorimeters described in this practice are classified as Type II dosimeters on the basis of the complex effect of influence quantities. See ISO/ASTM Practice 52628.

1.4 This practice applies to electron beams in the energy range from 1.5 to 12 MeV.

1.5 The absorbed dose range depends on the calorimetric absorbing material and the irradiation and measurement conditions. Minimum dose is approximately 100 Gy and maximum dose is approximately 50 kGy.

1.6 The average absorbed-dose rate range shall generally be greater than 10 Gy·s⁻¹.

1.7 The temperature range for use of these calorimetric dosimetry systems depends on the thermal resistance of the calorimetric materials, on the calibration range of the temperature sensor, and on the sensitivity of the measurement device.

1.8 *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, health, and environmental practices and determine the applicability of regulatory limitations prior to use.*

¹ This practice is under the jurisdiction of ASTM Committee E61 on Radiation Processing and is the direct responsibility of Subcommittee E61.02 on Dosimetry Systems, and is also under the jurisdiction of ISO/TC 85/WG 3.

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appropriate safety, health, and environmental practices and determine the applicability of regulatory limitations prior to use.

1.9 *This international standard was developed in accordance with internationally recognized principles on standardization established in the Decision on Principles for the Development of International Standards, Guides and Recommendations issued by the World Trade Organization Technical Barriers to Trade (TBT) Committee.*

2. Referenced Documents

2.1 ASTM Standards:²

E666 Practice for Calculating Absorbed Dose From Gamma or X Radiation

E668 Practice for Application of Thermoluminescence-Dosimetry (TLD) Systems for Determining Absorbed Dose in Radiation-Hardness Testing of Electronic Devices

E3083 Terminology Relating to Radiation Processing: Dosimetry and Applications

2.2 ISO/ASTM Standards:²

51261 Practice for Calibration of Routine Dosimetry Systems for Radiation Processing

51649 Practice for Dosimetry in an Electron Beam Facility for Radiation Processing at Energies Between 300 keV and 25 MeV

51707 Guide for Estimating Uncertainties in Dosimetry for Radiation Processing

52628 Practice for Dosimetry in Radiation Processing

2.3 International Commission on Radiation Units and Measurements (ICRU) Reports:³

ICRU Report 34 The Dosimetry of Pulsed Radiation

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

ICRU Report 80 Dosimetry Systems for use in Radiation Processing

ICRU Report 85a Fundamental Quantities and Units for Ionizing Radiation

² For referenced ASTM and ISO/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.

³ Available from the Commission on Radiation Units and Measurements, 7910 Woodmont Ave., Suite 800, Bethesda, MD 20814, U.S.A.



2.4 *Joint Committee for Guides in Metrology (JCGM) Reports:*

JCGM 100:2008, GUM 1995, with minor corrections, Evaluation of measurement data – Guide to the Expression of Uncertainty in Measurement⁴

JCGM 200:2012, VIM International vocabulary of metrology – Basic general concepts and general terms⁵

3. Terminology

3.1 Definitions:

3.1.1 *primary-standard dosimetry system*—dosimetry system that is designated or widely acknowledged as having the highest metrological qualities and whose value is accepted without reference to other standards of the same quantity.

3.1.2 *reference standard dosimetry system*—dosimetry system, generally having the highest metrological quality available at a given location or in a given organization, from which measurements made there are derived.

3.1.3 *transfer standard dosimetry system*—dosimetry system used as an intermediary to calibrate other dosimetry systems.

3.1.4 *type II dosimeter*—dosimeter, the response of which is affected by influence quantities in a complex way that cannot practically be expressed in terms of independent correction factors.

3.2 Definitions of Terms Specific to This Standard:

3.2.1 *adiabatic*—no heat exchange with the surroundings.

3.2.2 *calorimeter*—assembly consisting of calorimetric body (absorber), thermal insulation, and temperature sensor with wiring that, when irradiated, exhibits increase in the absorber temperature that can be related to absorbed dose. This language parallels that of dosimeter.

3.2.3 *calorimetric body*—mass of material absorbing radiation energy and whose temperature is measured.

3.2.4 *calorimetric dosimetry system*—dosimetry system consisting of calorimeter, measurement instruments and their associated reference standards, and procedures for the system's use.

3.2.5 *endothermic reaction*—chemical reaction that consumes energy.

3.2.6 *exothermic reaction*—chemical reaction that releases energy.

3.2.7 *heat defect (thermal defect)*—amount of energy released or consumed by chemical reactions caused by the absorption of radiation energy.

3.2.8 *specific heat capacity*—amount of energy required to raise 1 kg of material by the temperature of 1 K.

⁴ Document produced by Working Group 1 of the Joint Committee for Guides in Metrology (JCGM/WG 1). Available free of charge at the BIPM website (<http://www.bipm.org>).

⁵ Document produced by Working Group 2 of the Joint Committee for Guides in Metrology (JCGM/WG 2). Available free of charge at the BIPM website (<http://www.bipm.org>).

3.2.9 *thermistor*—electrical resistor with a well-defined relationship between resistance and temperature of the thermistor.

3.2.10 *thermocouple*—junction of two metals producing an electrical voltage with a well-defined relationship to junction temperature.

3.3 Definitions of other terms used in this standard that pertain to radiation measurement and dosimetry may be found in ASTM Terminology E3083. Definitions in E3083 are compatible with ICRU Report 85a; that document, therefore, may be used as an alternative reference.

4. Significance and use

4.1 This practice is applicable to the use of calorimetric dosimetry systems for the measurement of absorbed dose in electron beams, the qualification of electron irradiation facilities, periodic checks of operating parameters of electron irradiation facilities, and calibration of other dosimetry systems in electron beams. Calorimetric dosimetry systems are most suitable for dose measurement at electron irradiation facilities utilizing conveyor systems for transport of product during irradiation.

NOTE 1—For additional information on calorimetric dosimetry system operation and use, see ICRU Report 80. For additional information on the use of dosimetry in electron accelerator facilities, see ISO/ASTM 51649, and ICRU Reports 34 and 35, and Refs (1-3).⁶

4.2 The calorimetric dosimetry systems described in this practice are not primary standard dosimetry systems. The calorimeters are classified as Type II dosimeters (ISO/ASTM 52628). They might be used as internal standards at an electron beam irradiation facility, including being used as transfer standard dosimetry systems for calibration of other dosimetry systems, or they might be used as routine dosimeters. The calorimetric dosimetry systems are calibrated by comparison with transfer standard dosimeters.

4.3 The dose measurement is based on the measurement of the temperature rise (dosimeter response) in an absorber (calorimetric body) irradiated by an electron beam. Different absorbing materials are used, but the response is usually defined in terms of dose to water.

NOTE 2—The calorimetric bodies of the calorimeters described in this practice are made from low atomic number materials. The electron fluences within these calorimetric bodies are almost independent of energy when irradiated with electron beams of 1.5 MeV or higher, and the mass collision stopping powers are approximately the same for these materials.

4.4 The absorbed dose in other materials irradiated under equivalent conditions can be calculated. Procedures for making such calculations are given in ASTM Practices E666 and E668, and Ref (1).

4.4.1 Calorimeters for use at industrial electron accelerators have been constructed using graphite, polystyrene or a Petri dish filled with water as the calorimetric body (4-10). The thickness of the calorimetric body should be less than the range of the incident electrons.

4.4.2 Polymeric materials other than polystyrene might also be used for calorimetric measurements. Polystyrene is used

⁶ The boldface numbers in parentheses refer to the bibliography at the end of this practice.

because it is known to be resistant to radiation (11) and because almost no exo- or endothermic reactions take place (12).

5. Interferences

5.1 *Extrapolation*—The calorimetric dosimetry systems described in this practice are not adiabatic, because of the exchange of heat with the surroundings or within the calorimeter assembly. The maximum temperature reached by the calorimetric body is different from the temperature that would have been reached in the absence of that heat exchange. The temperature drifts before and after irradiation should be extrapolated to the midpoint of the irradiation period in order to determine the true temperature increase due to the absorbed dose.

5.2 *Heat Defect*—Chemical reactions in irradiated material (resulting in what is called the heat defect or thermal defect) might be endo- or exothermic and might lead to measurable temperature changes (3).

5.3 *Specific Heat Capacity*—The specific heat capacity of some materials used as a calorimetric body might change with accumulated absorbed dose, thereby affecting the response of the calorimeters. This is notably the case for polymers, such as polystyrene, and it will therefore be necessary to recalibrate calorimetric dosimetry systems at intervals that will depend on the total accumulated dose.

NOTE 3—For calorimeters using polystyrene as material for the calorimetric body, the change in specific heat capacity might be in the order of 1 % per accumulated dose of 1 MGy. It can therefore be useful to track accumulated dose for polystyrene calorimeters.

5.4 *Influence Quantities*—The response of the calorimetric dosimetry systems to absorbed dose does not depend on ambient relative humidity and temperature.

5.5 *Temperature Effects from Accelerator Structure*—The calorimeters are often irradiated on a conveyor used for passing products and samples through the irradiation zone. Recognizing that the thermal insulation around the calorimetric body is not perfect, there is possibility that, for example, radiated heat from the mechanical structures of the irradiation facility and from the conveyor might contribute to the measured temperature increase in the calorimeters.

5.6 *Thermal Equilibrium*—The most reproducible results are obtained when the calorimeters are in thermal equilibrium with their surroundings before irradiation.

5.7 *Foreign Materials*—The temperature sensors, wires, etc. of the calorimeter represent foreign materials, which might influence the temperature rise of the calorimetric body. These components should be as small as possible.

5.8 *Dose Gradients*—Dose gradients will exist within the calorimetric body when it is irradiated with electrons. These gradients must be taken into account, for example, when other dosimeters are calibrated by comparison with calorimetric dosimetry systems.

6. Apparatus

6.1 *A Typical Graphite Calorimeter* is a disc of graphite placed in a thermally insulating material such as foamed plastic (4-6). A calibrated thermistor or thermocouple is embedded

inside the disc. Some typical examples of graphite disc thicknesses and masses are listed in Annex A2 (2).

6.2 *A Typical Water Calorimeter* is a sealed polystyrene Petri dish filled with water and placed in thermally insulating foamed plastic (4). A calibrated temperature sensor (thermistor) is placed through the side of the dish into the water. The shape and size of the water calorimeter can be similar to the shape and size of the polystyrene calorimeter (see 6.3).

6.3 *A Typical Polystyrene Calorimeter* is a polystyrene disc placed in thermally insulating foamed plastic. A calibrated thermistor or thermocouple is imbedded inside the disc. The dimension of the polystyrene disc might be similar to that of the graphite and water calorimeters (9). See Fig. 1 as an example of a polystyrene calorimeter designed for use at 10 MeV electron irradiation (13).

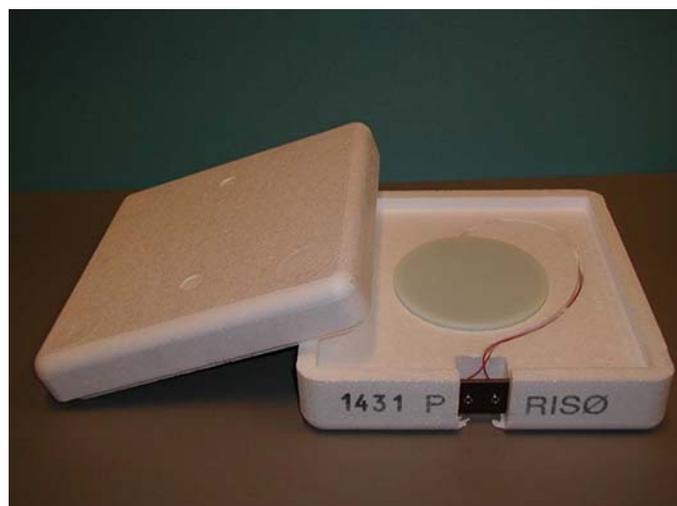
6.4 The thickness of the calorimetric body should be less than the range of the irradiating electrons, typically not exceeding $\frac{1}{3}$ of the range of the incident electrons. This will limit the effects of variation of the dose gradients within the calorimetric body.

6.5 Radiation-resistant components should be used for the parts of the calorimeter that are exposed to the electron beam. This also applies to insulation of electrical wires.

6.6 Good thermal contact must exist between the temperature sensor and the calorimetric body. For graphite and polystyrene calorimeters, this can be assured by adding a small amount of heat-conducting compound when mounting the temperature sensor.

6.7 *Measurement*—The response of the calorimeters is the temperature rise of the calorimetric body. This temperature rise is usually registered by thermistors or thermocouples.

6.7.1 *Thermistor*—A high-precision ohm-meter can be used for measurement of thermistor resistance. The meter should have a reproducibility of better than $\pm 0.1\%$ ($k=1$) and a combined uncertainty of better than $\pm 0.2\%$ ($k=1$). It should preferably be equipped for four-wire type resistance



Courtesy of Risø High Dose Reference Laboratory.

FIG. 1 Example of a polystyrene calorimeter used for routine measurements at a 10-MeV industrial electron accelerator

measurements, especially if the thermistor resistance is less than 10 kΩ. With the four-wire measurement technique, the effects of resistance in the measurement wires and electrical contacts are minimized.

6.7.2 Other appropriate instrumentation may be used for the thermistor resistance measurement, for example, a resistance bridge or commercially calibrated thermistor readers (5). It is important for both ohm-meters and resistance bridge measurements to minimize the dissipated power in the thermistor, preferably below 0.1 mW, in order to avoid self-heating of the thermistor during measurement.

6.7.3 *Thermocouple*—A high-precision digital voltmeter, or other dedicated instrument (2), can be used for the measurement. The reproducibility of the voltmeter should be better than 0.1 μV ($k=1$), and a combined uncertainty of better than ±0.2 % ($k=1$).

6.7.4 *Suppliers*—Some commercial suppliers of calorimetric dosimetry systems are listed in Annex A3.

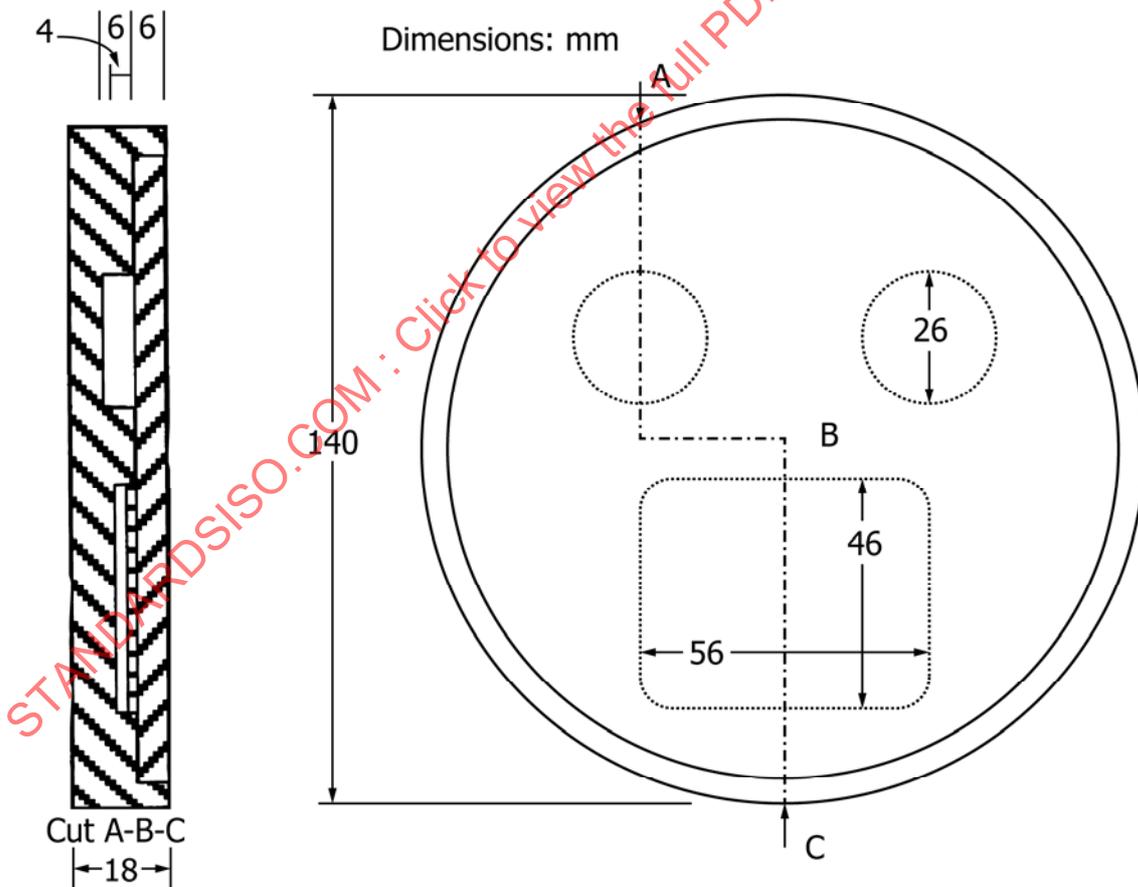
7. Calibration procedures

7.1 Prior to use, the calorimetric dosimetry system (consisting of calorimeter and measurement instruments) shall be calibrated in accordance with the user’s documented procedure

that specifies details of the calibration process and quality assurance requirements. This calibration process shall be repeated at regular intervals to ensure that the accuracy of the absorbed dose measurement is maintained within required limits. Calibration methods are described in ISO/ASTM Guide 51261.

7.2 Graphite, water or polystyrene calorimetric dosimetry systems should be calibrated by comparison with transfer standard dosimetry systems from an accredited calibration laboratory by irradiating the calorimeter(s) and transfer-standard dosimeters sequentially (or simultaneously) at an electron irradiation facility. The radiation field over the cross-sectional area of the calorimetric body shall be uniform over the time required to irradiate the calorimeters and the transfer-standard dosimeters. Any non-uniformity should be taken into account when evaluating and comparing dose to calorimeter and dose to transfer-standard dosimeter.

7.3 It must be assured that the transfer-standard dosimeters and the calorimeters are irradiated to the same dose. Specially designed absorbers (phantoms) are needed for irradiation of the transfer-standard dosimeters, see for example Fig. 2.



NOTE 1—All dimensions are in mm. Alanine transfer standard dosimeters in cylindrical flat holders (diameter 25 mm, thickness 6 mm) to be placed in the round cut-outs. Routine dosimeters (thin film dosimeters) to be placed in rectangular cut-outs. The centres of both dosimeters are placed in the same depth in the absorber.

FIG. 2 Absorber (phantom) for irradiation at 10 MeV electron irradiation facility of routine and transfer-standard dosimeters (10). Material: Polystyrene



7.4 The specific heat capacities of polystyrene and of graphite are functions of temperature, while the specific heat capacity of water is almost constant within the temperature range normally employed in electron beam calorimetry. The calibration curves of the calorimetric dosimetry systems are therefore expected to be functions of the average temperature of the calorimetric body (see Note 4).

7.4.1 For graphite calorimetric dosimetry systems, the calibration curve might take the following form:

$$\text{Dose} = (T_2 - T_1 - T_a) \cdot c_G \cdot (S_{e/\rho})_w / (S_{e/\rho})_G \cdot k$$

where:

- T_1 = temperature before irradiation,
 T_2 = temperature after irradiation,
 T_a = temperature rise from irradiation facility components,
 c_G = specific heat capacity of graphite,
 $(S_{e/\rho})_w$ = electronic mass stopping power of water,
 $(S_{e/\rho})_G$ = electronic mass stopping power of graphite, and
 k = calibration constant to be determined during calibration verification.

NOTE 4—Repeated measurements of specific heat of various types of graphite have been carried out over the range of 0 to 50°C, indicating a value for the specific heat capacity of graphite c_G ($J \cdot \text{kg}^{-1} \cdot ^\circ\text{C}^{-1}$) = $644.2 + 2.86 T$, where T is the mean temperature ($^\circ\text{C}$) of the graphite. This value must, however, not be considered a universal value (6).

7.4.2 For polystyrene calorimetric dosimetry systems, the calibration curve might take the following form:

$$\text{Dose} = (T_2 - T_1 - T_a) \cdot F(T) \cdot k$$

where:

- T_1 = temperature before irradiation,
 T_2 = temperature after irradiation,
 T_a = temperature rise from irradiation facility components,
 $F(T)$ = function representing specific heat capacity of polystyrene, and
 k = calibration constant to be determined during calibration verification.

NOTE 5—The function $F(T)$ takes the form $F(T) = C1 + C2 \cdot T$, where $C1$ and $C2$ are constants and T is the mean temperature ($^\circ\text{C}$) of the calorimetric body following irradiation. The values of $C1$ and $C2$ depend on the type of polystyrene used for making the calorimetric absorber.

NOTE 6—The value of T_a depends on the facility where the calorimeter is used. T_a can be determined by passing a calorimeter through the irradiation zone shortly after the electron beam has been switched off, and measuring the temperature increase of the calorimetric absorber.

NOTE 7—The sensitivity of water calorimetric dosimetry systems is approximately $3.4 \text{ kGy} \cdot ^\circ\text{C}^{-1}$ and for polystyrene calorimetric dosimetry systems it is approximately $1.4 \text{ kGy} \cdot ^\circ\text{C}^{-1}$. For graphite calorimetric dosimetry systems, the sensitivity is approximately $0.75 \text{ kGy} \cdot ^\circ\text{C}^{-1}$.

7.5 Calibration of all types of calorimetric dosimetry systems used as routine dosimetry systems should be verified by the user by comparison with reference standard or transfer standard dosimetry systems at a frequency determined by the user.

7.6 It is recommended that calibration of the calorimetric dosimetry system be carried out by irradiation at the user's facility in order for the effect of influence quantities to be minimized.

7.7 Calorimetric dosimetry systems can be calibrated by irradiation at a calibration laboratory. The calibration obtained in this way must be verified by irradiation of the calorimeters and transfer-standard dosimeters together at the user's facility.

NOTE 8—Calibration curves provided by manufacturers of calorimeters are typically not obtained by irradiation at the user's facility. Such calibration curves should be verified by irradiation at the user's facility.

7.8 An example of a calibration verification of a calorimetric dosimetry system is given in Annex A1.

7.9 *Measurement Instrument Calibration and Performance Verification*—For the calibration of the measuring instruments, and for the verification of instrument performance between calibrations, see ISO/ASTM Guide 51261 or instrument-specific operating manuals, or both.

8. Dose measurement procedures

8.1 *Conveyor Irradiation Off-Line Measurements*—For calorimeters carried on conveyors through scanned electron beams, the calorimeter is usually disconnected from the temperature measurement system just prior to placing the calorimeter on the conveyor and reconnected for measurement as soon as practical after irradiation (7).

8.1.1 Before irradiation, the temperature of the calorimetric body is measured. It should remain stable, typically less than 0.1°C change over a period of at least 10 min.

NOTE 9—It is recommended to store calorimeters in close proximity to the irradiation area, so that the calorimeters are in approximate thermal equilibrium with the conditions of the irradiation area prior to use.

8.1.2 The measurement wires are disconnected and the calorimeter is placed on the conveyor for transport through the irradiation zone.

8.1.3 The calorimeter is transported through the irradiation zone on the conveyor system.

8.1.4 The time of irradiation, and the irradiation parameters (for example, electron energy, electron current, scanned beam width, and conveyor speed) should be recorded.

8.1.5 After passage of the calorimeter through the irradiation zone, the wires for measurement of temperature are reconnected as soon as practical and the temperature is measured.

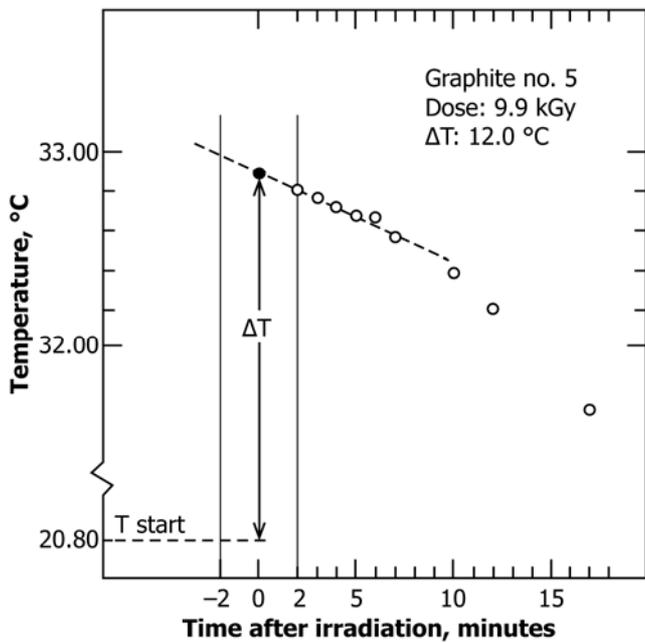
8.1.6 The temperature can be recorded as a function of time after irradiation for a period long enough to establish the thermal decay characteristics of the calorimeter. Typically a time of 10-20 min is required.

8.1.7 The temperature as a function of time before and after irradiation can be plotted.

8.1.8 The curves before and after irradiation can be extrapolated to the midpoint of the irradiation time. The two values of temperature obtained from the extrapolations are used as the temperature before irradiation (T_1) and after irradiation (T_2), respectively, that reflects the temperature rise solely due to absorbed dose. An example of data obtained by this measurement technique is shown in Fig. 3.

8.1.9 Based on the temperature difference, $T_2 - T_1$, the average absorbed dose in the calorimetric body can be determined from the established calibration curve (see 7.4.2).

8.1.10 For well-established, reproducible irradiation conditions the extrapolation procedure in 8.1.6 – 8.1.8 is not needed.



NOTE 1— ΔT is the temperature rise found by extrapolation and used for dose calculation (see 8.1.9). Measurement wires were disconnected during irradiation.

FIG. 3 Example of measurements of temperature of a graphite calorimeter before and after irradiation (7)

One measurement of temperature before and one after irradiation might suffice, and the temperature difference at the time of irradiation is found by use of a correction factor derived during the establishment of the irradiation procedures (4, 5, 7, 8). The times of the measurements relative to time of irradiation should be specified.

8.1.11 Software can be developed by the user for the calculation of dose as measured with the calorimetric dosimetry systems. Suppliers of calorimetric dosimetry systems usually supply such software. An example is Caldose from Risø High Dose Reference Laboratory (Denmark).

8.2 *On-line Measurements*—It is possible to measure the temperature of the calorimetric body during irradiation as the calorimeter is transported through the irradiation zone on the conveyor with measurement wires attached. Four-wire measurement technique (see 6.7.1) is preferred in order to reduce measurement uncertainty.

8.3 *Stationary Irradiation*—The calorimeters described in this practice might also be used in a stationary configuration instead of being transported on a conveyor system through the electron beam. In this arrangement the electron beam can be made almost uniform (see 8.3.2) over the area of the calorimetric body either by the use of metallic scattering foils or by raster scanning. The irradiation period is controlled by turning the electron beam on and off.

8.3.1 The measurement of the temperature of the calorimeters in a stationary configuration might be done continuously before, during, and after irradiation rather than measuring only before and after irradiation as described in 8.1.

8.3.2 The calorimeter should be located on the beam axis at an appropriate distance from the accelerator beam exit window

such that the beam profile is uniform to within $\pm 2\%$ across the diameter of the calorimeter body. The beam profile should be measured and if it varies more than $\pm 2\%$ across the calorimetric body, corrections for the non-uniformity might have to be carried out. If the temperature measurement system is located outside of the irradiation area for on-line measurement, and long connecting wires are used, then the use of four-wire measurement technique is essential.

8.3.3 The temperature of the calorimetric body is measured as a function of time to ensure that the initial drift rate is less than the desired limits (see 8.1.1).

8.3.4 While the calorimetric body temperature is recorded continuously, the electron beam is turned on for the desired irradiation time. After the beam is turned off, the temperature is measured for enough time to establish the thermal decay characteristics.

8.3.5 The time versus temperature curves are extrapolated before and after irradiation to the midpoint of the irradiation time in the same manner as in 8.1.7. The average absorbed dose in the calorimetric body is calculated in the same manner as in 8.1.8 or 8.1.9. An example of data obtained by this measurement technique is shown in Fig. 4 (2, 5).

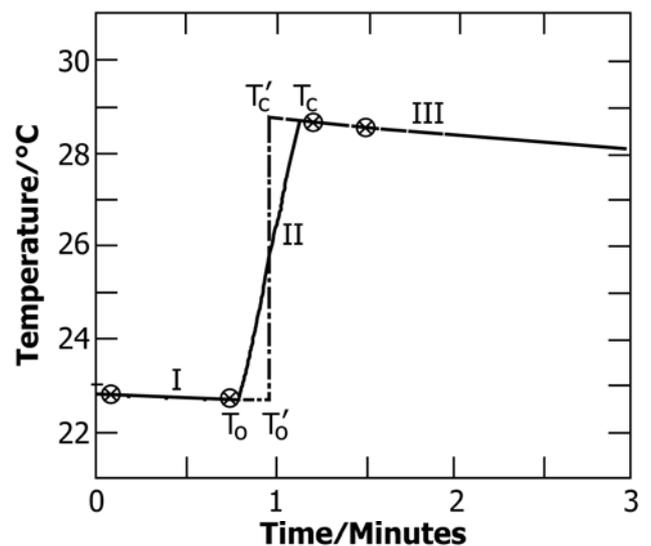
9. Calibration of other dosimetry systems

9.1 The absorbed dose measured by the calorimetric dosimetry system is the average dose in the calorimetric body.

9.2 The calibrated calorimetric dosimetry systems can be used for calibration of other dosimetry systems irradiated at the same irradiation facility.

NOTE 10—In this method of calibration the calorimeter is used as an internal reference standard dosimeter. This extra link in the traceability chain should be considered when establishing the uncertainty budget. The same considerations as given in Section 7 apply here.

9.3 The dosimeters to be calibrated must be irradiated so as to receive the same average absorbed dose as the calorimeter



NOTE 1—The temperature curves were extrapolated from T_o and T_c to the midpoint of the irradiation time at T_o' and T_c' , respectively. $\Delta T = T_c' - T_o'$ is used for dose calculation.

FIG. 4 Example of on-line measurements of a graphite calorimeter (5)

when placed at specific depths in absorbers that simulate the calorimetric body (see Fig. 2). The dosimeters for calibration shall be small enough to fit inside the thickness of the absorber (1, 5).

9.4 By successive (or simultaneous) irradiation of the absorbers and the calorimeters under identical conditions, the dosimeters being calibrated are irradiated to the doses measured by the calorimeters.

9.5 A relation between the average dose in the calorimetric body and the dose delivered to the dosimeter being calibrated can be established, if the dose distribution throughout the calorimetric body is known and the dosimeters being calibrated are small. This depth dose distribution in the absorber can be measured by irradiating a dummy calorimetric absorber containing a calibrated film dosimeter placed at an angle (preferably more than 45°) to the beam direction, and analyzing the dose distribution. If the thickness of dosimeters being calibrated is not small compared with the calorimeter absorber thickness, then the average dose delivered to the dosimeter must be related to the average dose delivered to the absorber by measurement of the dose distribution in both.

9.6 The mean temperature of the dosimeters during irradiation for calibration should be determined (5). This is necessary if temperature corrections of the response are to be applied.

10. Documentation

10.1 For the calibration and application of the calorimetric dosimetry systems, record or reference the temperature sensor calibration data and the calibration data for the calorimetric dosimetry systems. Record the total accumulated dose of the calorimeter. See 5.3.

10.2 Record or reference the irradiation date, temperature and time data, analysis instrumentation, dose interpretation of calorimetry data, electron beam characteristics, and accelerator operating parameters.

11. Measurement uncertainty

11.1 All dose measurements need to be accompanied by an estimate of uncertainty. Appropriate procedures are recommended in ISO/ASTM Guide 51707 (see also GUM).

11.2 All components of uncertainty should be included in the estimate, including those arising from calibration, dosimeter reproducibility, instrument reproducibility, and the effect of influence quantities. A full quantitative analysis of components of uncertainty is referred to as an uncertainty budget, and is then often presented in the form of a table. Typically, the uncertainty budget will identify all significant components of uncertainty, together with their methods of estimation, statistical distributions and magnitudes.

11.3 The estimate of the expanded uncertainty achievable with measurements made using a routine dosimetry system such as calorimeters is typically of the order of 4 % ($k = 2$), which corresponds approximately to a 95 % level of confidence for normally distributed data. Table 1 (9) gives examples of uncertainty components for a specific calorimetric system. The values in this table are not meant to be representative of other calorimetric systems or types and are only for illustrative purposes.

12. Keywords

12.1 absorbed dose; calibration; calorimeter; calorimetry; dosimeter; dosimetry; electron accelerator; electron beam; graphite; ionizing radiation; irradiation; polystyrene; radiation; radiation processing; water; ICS 17.240

TABLE 1 Measurement uncertainties of routine polystyrene calorimetric dosimetry systems from Risø high dose reference laboratory (in percent, at $k = 2$) (9)

NOTE 1—At doses higher than 10 kGy, (2) and (3) are reduced to 0.2 %.

	Sources of Uncertainty	Type A	Type B
1	Calibration		3.2
2	Temperature measurement of calorimeter (at 3 kGy)	1.0	
3	Temperature extrapolation of calorimeter (at 3 kGy)	1.0	
4	Change of temperature sensitivity of specific heat of polystyrene		0.5
5	Heating effects	0.5	
	Quadratic sum	1.5	3.2
	Combined	3.6 %	

ANNEXES

Informative

A1. EXAMPLE OF A CALIBRATION VERIFICATION OF CALORIMETRIC DOSIMETRY SYSTEMS

A1.1 Irradiation of calorimeters and transfer-standard dosimeters at an electron accelerator for verification of calibration.

A1.2 *Calorimeters*—Risø HDRL 10 MeV calorimeters (see Fig. 1)