EESTI STANDARD

Methods for calculating size specific dose estimates , β aph, Were a constant of the second of t (SSDE) on computed tomography



EESTI STANDARDI EESSÕNA

NATIONAL FOREWORD

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|---|--|--|--|--|
| See Eesti standard EVS-EN IEC 62985:2019 sisaldab Euroopa standardi EN IEC 62985:2019 ingliskeelset teksti. | This Estonian standard EVS-EN IEC 62985:2019 consists of the English text of the European standard EN IEC 62985:2019. | | | |
| Standard on jõustunud sellekohase teate avaldamisega EVS Teatajas | This standard has been endorsed with a notification published in the official bulletin of the Estonian Centre for Standardisation. | | | |
| Euroopa standardimisorganisatsioonid on teinud Euroopa standardi rahvuslikele liikmetele kättesaadavaks 01.11.2019. | Date of Availability of the European standard is 01.11.2019. | | | |
| Standard on kättesaadav Eesti Standardikeskusest. | The standard is available from the Estonian Centre for Standardisation. | | | |
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EUROPEAN STANDARD NORME EUROPÉENNE EUROPÄISCHE NORM

EN IEC 62985

November 2019

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English Version

Methods for calculating size specific dose estimates (SSDE) on computed tomography (IEC 62985:2019)

Méthodes de calcul de l'estimateur de dose morphologique (SSDE) en tomodensitométrie (IEC 62985:2019) To be completed (IEC 62985:2019)

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European foreword

The text of document 62B/1133/FDIS, future edition 1 of IEC 62985, prepared by SC 62B "Diagnostic imaging equipment" of IEC/TC 62 "Electrical equipment in medical practice" was submitted to the IEC-CENELEC parallel vote and approved by CENELEC as EN IEC 62985:2019.

The following dates are fixed:

- latest date by which the document has to be implemented at national (dop) 2020-07-18 level by publication of an identical national standard or by endorsement
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Annex ZA

(normative)

Normative references to international publications with their corresponding European publications

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

NOTE 1 Where an International Publication has been modified by common modifications, indicated by (mod), the relevant EN/HD applies.

NOTE 2 Up-to-date information on the latest versions of the European Standards listed in this annex is available here: www.cenelec.eu.

| Publication | Year | <u>Title</u> | <u>EN/HD</u> | Year |
|----------------|------|--|--------------|------|
| IEC/TR 60788 | 2004 | Medical electrical equipment - Glossary o defined terms | f - | - |
| IEC 60601-1 | 2005 | Medical electrical equipment - Part 1 General requirements for basic safety and essential performance | | 2006 |
| + A1 | 2012 | Q, | + A1 | 2013 |
| - | - | L. | + A12 | 2014 |
| IEC 60601-1-3 | 2008 | Medical electrical equipment - Part 1-3 General requirements for basic safety and essential performance - Collatera Standard: Radiation protection in diagnostic X-ray equipment | d I | 2008 |
| - | - | Q | + A11 | 2016 |
| IEC 60601-2-44 | 2009 | Medical electrical equipment - Part 2-44 Particular requirements for the basic safet and essential performance of X-ra equipment for computed tomography | y | 2009 |
| - | - | | + A11 | 2011 |
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INTERNATIONAL ELECTROTECHNICAL COMMISSION

METHODS FOR CALCULATING SIZE SPECIFIC DOSE ESTIMATES (SSDE) FOR COMPUTED TOMOGRAPHY

FOREWORD

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International Standard IEC 62985 has been prepared by subcommittee 62B: Diagnostic imaging equipment, of IEC technical committee 62: Electrical equipment in medical practice.

The text of this International Standard is based on the following documents:

| FDIS | Report on voting |
|---------------|------------------|
| 62B/1133/FDIS | 62B/1144/RVD |

Full information on the voting for the approval of this International Standard can be found in the report on voting indicated in the above table.

This document has been drafted in accordance with the ISO/IEC Directives, Part 2.

In this document, the following print types are used:

requirements and definitions: roman type;

- informative material appearing outside of tables, such as notes, examples and references: in smaller type. Normative text of tables is also in a smaller type;
- DEFINED CLAUSE 3, IN CLAUSE 3 OF IEC 60601-1:2005 AND TERMS IN IEC 60601-1:2005/AMD1:2012, OF THE COLLATERAL STANDARDS, OF IEC TR 60788:2004 OR AS NOTED: SMALL CAPITALS.

The committee has decided that the contents of this document will remain unchanged until the stability date indicated on the IEC website under "http://webstore.iec.ch" in the data related to the specific document. At this date, the document will be

- reconfirmed, .
- withdrawn, .
- replaced by a revised edition, or
- amended. •

NOTE The attention of the user of this document is drawn to the fact that equipment MANUFACTURERS and testing organizations may need a transitional period following publication of a new, amended or revised IEC publication in which to make products in accordance with the new requirements and to equip themselves for conducting new or revised tests. It is the recommendation of the committee that the content of this publication be adopted for implementation nationally not earlier than 3 years from the date of publication.

A bilingual version of this publication may be issued at a later date.

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INTRODUCTION

The SIZE SPECIFIC DOSE ESTIMATE (SSDE) is an estimate of the average ABSORBED DOSE to the scan volume that takes into account the ATTENUATION of the anatomy being scanned (using the WATER EQUIVALENT DIAMETER $D_{\rm W}$) and the RADIATION OUTPUT of the CT SCANNER (using CTDI_{VOL}).

SSDE is intended to provide a dose estimate for PATIENTS of all sizes. SSDE, which is given in units of mGy, is especially important for small paediatric PATIENTS since the corresponding applied level of RADIATION (CTDI_{VOL}, also given in units of mGy) does not adequately indicate the absorbed RADIATION DOSE.

SSDE is calculated using a SSDE CONVERSION FACTOR AT LONGITUDINAL POSITION Z (f) and the CTDI_{VOL} AT LONGITUDINAL POSITION Z, CTDI_{VOL}(z), where f is a function of the WATER EQUIVALENT DIAMETER AT LONGITUDINAL POSITION Z, $D_W(z)$, and the size of the CTDI PHANTOM used to report CTDI_{VOL}. f is given in normative Annex A.

This document provides a methodology (in Clause 4) for a MANUFACTURER to validate their method for calculating $D_W(z)$, which is used for the determination of f and the calculation of SSDE. This method calculates a reference WATER EQUIVALENT DIAMETER AT LONGITUDINAL POSITION Z, $D_{W,REF}(z)$, and compares it against a known PHANTOM dimension and the implemented values of WATER EQUIVALENT DIAMETER AT LONGITUDINAL POSITION Z, $D_{W,REF}(z)$, and compares are also specified.

NOTE 1 The definition of SSDE used in this document differs from that of AAPM Report No. 204 [1]¹ in that AAPM Report No. 204 estimates the average dose at the centre of the scan volume, whereas in this document, SSDE estimates the average dose across the whole scan volume.

NOTE 2 $CTDI_{VOL}$ is a dose index that allows quantitation of the RADIATION OUTPUT of CT SCANNERS in terms of one of two PMMA test objects. These test objects are 16 cm and 32 cm in diameter. SSDE is calculated by conversion of one of these PHANTOM-based dose indices to an estimate of the RADIATION dose absorbed by a PATIENT of a specific size. The magnitude of the difference between SSDE and $CTDI_{VOL}$ values increases as the difference between the PATIENT size and the size of the CTDI PHANTOM used to measure the $CTDI_{VOL}$ increases. For infants, the calculated SSDE value may be 3 times as much as the corresponding $CTDI_{VOL}$ dose index value. Conversely, the $CTDI_{VOL}$ value for large PATIENTs overestimates SSDE, which is representative of the PATIENT's actual absorbed RADIATION DOSE. For extra-large adult PATIENTs, the $CTDI_{VOL}$ dose index can overestimate the SSDE by as much as 40 % [1].

Potential uses of SSDE include the following:

- 1) evaluating PATIENT ABSORBED DOSE for quality assurance programs;
- 2) establishing diagnostic reference levels across PATIENT sizes;
- 3) displaying to the OPERATOR an estimate of PATIENT ABSORBED DOSE prior to initiation of the CT scan;
- 4) providing an estimate of ABSORBED DOSE for the DICOM RDSR;
- 5) developing DOSE NOTIFICATION VALUE and DOSE ALERT VALUES that better take into account PATIENT size;
- 6) providing an estimate of PATIENT ABSORBED DOSE for dose registries.

¹ Numbers in square brackets refer to the Bibliography.