EESTI STANDARD

LASERTOODETE OHUTUS. OSA 1: SEADMETE KLASSIFIKATSIOON JA NÕUDED

Safety of laser products - Part 1: Equipment classification and requirements (IEC 60825-1:2014)



EESTI STANDARDI EESSÕNA

NATIONAL FOREWORD

See Eesti standard EVS-EN 60825-1:2014 +A11:2021 sisaldab Euroopa standardi EN 60825- 1:2014 ja selle muudatuste A11:2021 ja paranduse AC:2017 ja muudatuse A11 paranduse AC:2022 ingliskeelset teksti.	This Estonian standard EVS-EN 60825-1:2014 +A11:2021 consists of the English text of the European standard EN 60825-1:2014 and its amendment A11:2021 and its corrigendum AC:2017 and AC:2022.
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 and Accreditation.
Euroopa standardimisorganisatsioonid on teinud Euroopa standardi rahvuslikele liikmetele kättesaadavaks 08.08.2014, muudatus A11 19.02.2021.	Date of Availability of the European standard is 08.08.2014, for A11 19.02.2021.
Muudatusega A11 lisatud või muudetud teksti algus ja lõpp on tekstis tähistatud sümbolitega A_{11} A_{11} .	The start and finish of text introduced or altered by amendment A11 is indicated in the text by tags A_{11} A_{11} .
Parandusega AC:2017 lisatud või muudetud teksti algus ja lõpp on tekstis tähistatud sümbolitega \boxed{AC} $\langle AC \end{bmatrix}$.	The start and finish of text introduced or altered by corrigendum AC:2017 is indicated in the text by tags \overrightarrow{AC} $\langle \overrightarrow{AC} \rangle$.
Parandusega AC:2022 lisatud või muudetud teksti algus ja lõpp on tekstis tähistatud sümbolitega $\boxed{AC_2}$ $\langle \overrightarrow{AC_2}$.	The start and finish of text introduced or altered by corrigendum AC:2022 is indicated in the text by tags AC_2 AC_2
Standard on kättesaadav Eesti Standardimis-ja Akrediteerimiskeskusest.	The standard is available from the Estonian Centre for Standardisation and Accreditation.

Tagasisidet standardi sisu kohta on võimalik edastada, kasutades EVS-i veebilehel asuvat tagasiside vormi või saates e-kirja meiliaadressile <u>standardiosakond@evs.ee</u>.

ICS 13.110; 31.260

Standardite reprodutseerimise ja levitamise õigus kuulub Eesti Standardimis- ja Akrediteerimiskeskusele

Andmete paljundamine, taastekitamine, kopeerimine, salvestamine elektroonsesse süsteemi või edastamine ükskõik millises vormis või millisel teel ilma Eesti Standardimis-ja Akrediteerimiskeskuse kirjaliku loata on keelatud.

Kui Teil on küsimusi standardite autorikaitse kohta, võtke palun ühendust Eesti Standardimis-ja Akrediteerimiskeskusega: Koduleht <u>www.evs.ee</u>; telefon 605 5050; e-post <u>info@evs.ee</u>

The right to reproduce and distribute standards belongs to the Estonian Centre for Standardisation and Accreditation

No part of this publication may be reproduced or utilized in any form or by any means, electronic or mechanical, including photocopying, without a written permission from the Estonian Centre for Standardisation and Accreditation.

If you have any questions about copyright, please contact Estonian Centre for Standardisation and Accreditation: Homepage <u>www.evs.ee</u>; phone +372 605 5050; e-mail <u>info@evs.ee</u>

EUROPEAN STANDARD NORME EUROPÉENNE EUROPÄISCHE NORM

EN 60825-1 + A11

August 2014, February 2021

ICS 13.110; 31.260

Supersedes EN 60825-1:2007

English Version

Safety of laser products - Part 1: Equipment classification and requirements (IEC 60825-1:2014)

Sécurité des appareils à laser - Partie 1: Classification des matériels et exigences (CEI 60825-1:2014) Sicherheit von Lasereinrichtungen - Teil 1: Klassifizierung von Anlagen und Anforderungen (IEC 60825-1:2014)

This European Standard was approved by CENELEC on 2014-06-19. Amendment A11 was approved by CENELEC on 2021-01-18. CENELEC members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard and its amendment the status of a national standard without any alteration.

Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the CEN-CENELEC Management Centre or to any CENELEC member.

This European Standard and its amendment A11 exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CENELEC member into its own language and notified to the CEN-CENELEC Management Centre has the same status as the official versions.

CENELEC members are the national electrotechnical committees of Austria, Belgium, Bulgaria, Croatia, Cyprus, the Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, the Netherlands, Norway, Poland, Portugal, Republic of North Macedonia, Romania, Serbia, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and the United Kingdom.



European Committee for Electrotechnical Standardization Comité Européen de Normalisation Electrotechnique Europäisches Komitee für Elektrotechnische Normung

CEN-CENELEC Management Centre: Rue de la Science 23, B-1040 Brussels

© 2021 CENELEC All rights of exploitation in any form and by any means reserved worldwide for CENELEC Members.

- 2 -

Foreword

The text of document 76/502/FDIS, future edition 3 of IEC 60825-1, prepared by IEC/TC 76 "Optical radiation safety and laser equipment" was submitted to the IEC-CENELEC parallel vote and approved by CENELEC as EN 60825-1:2014.

The following dates are fixed:

- latest date by which the document has to be implemented (dop) 2015-03-19 at national level by publication of an identical national standard or by endorsement
- latest date by which the national standards conflicting with (dow) AC 2019-06-19 (AC the document have to be withdrawn

This document supersedes EN 60825-1:2007.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. CENELEC [and/or CEN] shall not be held responsible for identifying any or all such patent rights.

Endorsement notice

The text of the International Standard IEC 60825-1:2014 was approved by CENELEC as a European Standard without any modification.

IEC 60027-1	NOTE	Harmonised in EN 60027-1.
IEC 60065	NOTE	Harmonised as EN 60065.
IEC 60079 (Series)	NOTE	Harmonised as EN 60079 (Series).
IEC 60204-1	NOTE	Harmonised as EN 60204-1.
IEC 60601-2-22	NOTE	Harmonised as EN 60601-2-22.
IEC 60825-2	NOTE	Harmonised as EN 60825-2.
IEC 60825-4	NOTE	Harmonised as EN 60825-4.
IEC 60825-12	NOTE	Harmonised as EN 60825-12.
IEC 60950 (Series)	NOTE	Harmonised as EN 60950 (Series).
IEC 61010-1	NOTE	Harmonised as EN 61010-1.
IEC 61508 (Series)	NOTE	Harmonised as EN 61508 (Series).
IEC 62115	NOTE	Harmonised as EN 62115.
IEC 62368-1	NOTE	Harmonised as EN 62368-1.
IEC/ISO 11553 (Series)	NOTE	Harmonised as EN ISO 11553 (Series).
ISO 11146-1	NOTE	Harmonised as EN ISO 11146-1.
ISO 12100	NOTE	Harmonised as EN ISO 12100.
ISO 13694	NOTE	Harmonised as EN ISO 13694.
ISO 13849 (Series)	NOTE	Harmonised as EN ISO 13849 (Series).
ISO 15004-2:2007	NOTE	Harmonised as EN ISO 15004-2:2007.
ISO 80000-1	NOTE	Harmonised as EN ISO 80000-1.

An Amendment A11 European foreword

This document (EN 60825-1:2014/A11:2021) has been prepared by CLC/TC 76 "Optical radiation safety and laser equipment".

The following dates are fixed:

have to be withdrawn

•	latest date by which this document has to be implemented at national level by publication of an identical national standard or by endorsement	(dop)	2022-01-18
•	latest date by which the national standards conflicting with this document	(dow)	2024-01-18

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. CENELEC shall not be held responsible for identifying any or all such patent rights.

This document has been prepared under a mandate given to CENELEC by the European Commission and the European Free Trade Association, and supports essential requirements of EU Directive(s).

For the relationship with EU Directive(s) see informative Annex ZZ, which is an integral part of this document.

This document is expected to be read in conjunction with EN 506891 'Safety of laser products - Particular Requirements for Consumer Laser Products', when available. (An

with L

¹ Under preparation. Stage at the time of publication: prEN 50689:2019.

CONTENTS

FOF	REWORD)		7
1	Scope a	ind object		9
2	Normati	ve referenc	ces	11
3	Terms a	and definition	ons	11
4	Classific	cation princ	iples	25
	4.1	General	·	25
	4.2	Classificat	tion responsibilities	25
	4.3		tion rules	
	4.4	Laser proc	ducts designed to function as conventional lamps	31
5	Determi	nation of th	e accessible emission level and product classification	31
	5.1	Tests		31
	5.2	Measurem	nent of laser radiation	33
	5.3	Determina	ation of the class of the laser product	33
	5.4	Measurem	nent geometry	42
		5.4.1	General	
		5.4.2	Default (simplified) evaluation	
		5.4.3	Evaluation condition for extended sources	
6	-	•	ications	
	6.1		emarks and modifications	
	6.2		housing	
		6.2.1	General	
		6.2.2	Service	
		6.2.3	Removable laser system	
	6.3		anels and safety interlocks	
	6.4		iterlock connector	
	6.5		set	
	6.6		ol	
	6.7		ation emission warning o or attenuator	
	6.8	•	o or attenuator	
	6.9 6.10			
	6.10 6.11		ptics safeguard	
	6.12		I for Class 1C products	
	6.13		access	
	6.14		ental conditions	
	6.15		against other hazards	
	0.10	6.15.1	Non-optical hazards	
		6.15.2	Collateral radiation	
	6.16	Power lim	iting circuit	
7				
	7.1	•		
	7.2		nd Class 1M	
	7.3			
	7.4		nd Class 2M	
	7.5			
	7.6			

	7.7	Class 4		57
	7.8	Aperture la	abel	58
	7.9	Radiation	output and standards information	58
	7.10	Labels for	access panels	59
	2.	7.10.1	Labels for panels	59
	10	7.10.2	Labels for safety interlocked panels	60
	7.11	Warning for	or invisible laser radiation	60
	7.12	Warning for	or visible laser radiation	60
	7.13		or potential hazard to the skin or anterior parts of the eye	
8	Other in	formational	l requirements	61
	8.1	Informatio	n for the user	61
	8.2	Purchasin	g and servicing information	62
9	Addition	al requirem	nents for specific laser products	63
	9.1	Other part	s of the standard series IEC 60825	63
	9.2	Medical la	ser products	63
	9.3	Laser proc	cessing machines	63
	9.4	Electric to	ys	63
	9.5	Consumer	electronic products	63
Ann	ex A (info	ormative) N	Maximum permissible exposure values	65
	A.1	General re	emarks	65
	A.2	Limiting a	pertures	70
	A.3	Repetitive	ly pulsed or modulated lasers	71
	A.4	Measurem	ent conditions	
		A.4.1	General	72
		A.4.2	Limiting aperture	72
		A.4.3	Angle of acceptance	72
	A.5		source lasers	
Ann	ex B (inf	ormative) E	Examples of calculations	74
	B.1	Symbols u	used in the examples of this annex	74
	B.2		ion of a laser product – Introduction	
	B.3	Examples		79
Ann	ex C (inf	ormative) [Description of the classes and potentially associated hazards	84
	C.1	General		84
	C.2	Descriptio	n of classes	84
		C.2.1	Class 1	84
		C.2.2	Class 1M	84
		C.2.3	Class 1C	85
		C.2.4	Class 2	85
		C.2.5	Class 2M	85
		C.2.6	Class 3R	86
		C.2.7	Class 3B	
		C.2.8	Class 4	
		C.2.9	Note on nomenclature	87
	C.3	Limitations	s of the classification scheme	
	C.4	Reference	S	89
Ann	ex D (inf	ormative) I	Biophysical considerations	90
	D.1	Anatomy o	of the eye	90
	D.2	The effect	s of laser radiation on biological tissue	91

	D.2.1	General	91	
	D.2.2	Hazards to the eye	93	
2	D.2.3	Skin hazards	97	
D.3		irradiance averaging		
D.4		documents		
Annex E (inf	ormative) N	MPEs and AELs expressed as radiance	99	
E.1		d		
E.2		/alues		
E.3				
•		ummary tables		
		Overview of associated parts of IEC 60825	106	
		lormative references to international publications with their pean publications	108	
Annex ZE	3 (informativ	(e) Information for the Interpretation of 4.3, 4.4 and 6.3.2 (A11)	109	
ZB.1	General re	marks	109	
ZB.2	Subclause	4.3 Classification rules (IEC 60825-1:2014/ISH1:2017)	109	
	ZB.2.1	General remarks	109	
	ZB.2.2	Subclause 4.3 c) (Radiation from extended sources)	109	
	ZB.2.3	Subclause 4.3 d) (Non-uniform, non-circular or multiple apparent sources)	109	
	ZB.2.4	Subclause 4.3 f) 3); determination of α	111	
	ZB.2.5	Subclause 4.3 f) 3); groups of pulses with group duration longer than T_1	112	
	ZB.2.6	Subclause 4.3 f); simplifications	114	
ZB.3		4.4 conventional lamp replacement 5-1:2014/ISH2:2017)	117	
ZB.4	Subclause	6.3.2 - safety interlocks	118	
		e) Relationship between this European standard and the Directive 2014/35/EU [2014 OJ L96] aimed to be covered Ant	120	
Bibliography.		<u> </u>	121	
0				
Figure 1 – Me	easurement	set-up to limit angle of acceptance by imaging the apparent		
		the field stop	45	
		set-up to limit angle of acceptance by placing a circular ving as field stop) close to the apparent source	46	
Figure 3 – W	arning labe	I – Hazard symbol	53	
		abel		
		bel for Class 1		
		bel for Class 1M		
		bel for Class 1C		
0		bel for Class 2		
		bel for Class 2M		
-		abel for Class 3R		
-		abel for Class 3B		
-		abel for Class 4		
Figure 13 – A	igure 13 – Alternative label for laser aperture58			

Figure B.1 – Flowchart guide for the classification of laser products from supplied output parameters
Figure B.2 – Flowchart guide for the classification of Class 1M and Class 2M laser products
Figure B.3 – AEL for Class 1 ultra-violet laser products for selected emission durations from 10 ⁻⁹ s to 10 ³ s
Figure B.4 – AEL for Class 1 ultra-violet laser products for emission durations from 10 ⁻ ⁹ s to 10 ³ s at selected wavelengths78
Figure B.5 – AEL for Class 1 visible and selected infra-red laser products (case $C_6 = 1$)
Figure D.1 – Anatomy of the eye90
Figure D.2 – Diagram of laser-induced damage in biological systems
Figure E.1 – Radiance as a function of wavelength
Figure ZB.1 — Retinal image of a source pattern for the example of 20 emitters. Two possible groupings are defined by the respective angle of acceptance γ_X and γ_Y 110
Figure ZB.2 — Example of three groups of pulses (each group duration is longer than T_i) where each group is considered as one "effective" pulse and C_5 is applied to the AEL that applies to the group duration, where C_5 is determined with the number of
pulse groups within the evaluation duration (in the example of the figure $N = 3$)
Figure ZB.3 — Example of a train of pulses consisting of pulses with a duration of 3 μ s and 7 μ s
Figure ZB.4 — Flow diagram to illustrate the interpretation of subclause 6.3.2
Table 1 – Additivity of effects on eye and skin of radiation of different spectral regions27
Table 2 – Times below which pulse groups are summed
Table 3 – Accessible emission limits for Class 1 and Class 1M laser products and $C_6 = 1$
Table 4 – Accessible emission limits for Class 1 and Class 1M laser products in the wavelength range from 400 nm to 1 400 nm (retinal hazard region): extended sources37
Table 5 – Accessible emission limits for Class 2 and Class 2M laser products
Table 6 – Accessible emission limits for Class 3R laser products and $C_6 = 1$
Table 7 – Accessible emission limits for Class 3R laser products in the wavelengthrange from 400 nm to 1 400 nm (retinal hazard region): extended sources40
Table 8 – Accessible emission limits for Class 3B laser products 41
Table 9 – Correction factors and breakpoints for use in AEL and MPE evaluations41
Table 10 – Measurement aperture diameters and measurement distances for the default (simplified) evaluation
Table 11 – Reference points for Condition 344
Table 12 – Limiting angle of acceptance γ_{ph} 46
Table 13 – Requirements for safety interlocking
Table A.1 – Maximum permissible exposure (MPE) for $C_6 = 1$ at the cornea expressed
as irradiance or radiant exposure
Table A.2 – Maximum permissible exposure (MPE) at the cornea for extended sourcesin the wavelength range from 400 nm to 1 400 nm (retinal hazard region) expressed asirradiance or radiant exposure ^d

EVS-EN 60825-1:2014+A11:2021

Table A.3 – Maximum permissible exposure (MPE) of Table A.1 ($C_6 = 1$) for the	
wavelength range from 400 nm to 1 400 nm expressed as power or energy ^{a, b}	68
Table A.4 – Maximum permissible exposure (MPE) of Table A.2 (extended sources)	
for the wavelength range from 400 nm to 1 400 nm expressed as power or energy a, b,	
c, d, e, f, g	69
Table A.5 – Maximum permissible exposure (MPE) of the skin to laser radiation	70
Table A.6 – Aperture diameters for measuring laser irradiance and radiant exposure	71
Table D.1 – Summary of pathological effects associated with excessive exposure	
to light	94
Table D.2 – Explanation of measurement apertures applied to the eye MPEs	97
Table E.1 – Maximum radiance of a diffused source for Class 1	100
Table F.1 – Summary of the physical quantities used in this Part 1	103
Table F.2 – Summary of manufacturer's requirements (1 of 2)	104
Table G.1 – Overview of additional data in associated parts of IEC 60825	107
Table ZZ.1 — Correspondence between this European standard and Annex I of	
Directive 2014/35/EU [2014 OJ L96]	120

- 6 -

INTERNATIONAL ELECTROTECHNICAL COMMISSION

SAFETY OF LASER PRODUCTS -

Part 1: Equipment classification and requirements

FOREWORD

- 1) The International Electrotechnical Commission (IEC) is a worldwide organization for standardization comprising all national electrotechnical committees (IEC National Committees). The object of IEC is to promote international co-operation on all questions concerning standardization in the electrical and electronic fields. To this end and in addition to other activities, IEC publishes International Standards, Technical Specifications, Technical Reports, Publicly Available Specifications (PAS) and Guides (hereafter referred to as "IEC Publication(s)"). Their preparation is entrusted to technical committees; any IEC National Committee interested in the subject dealt with may participate in this preparatory work. International, governmental and non-governmental organizations liaising with the IEC also participate in this preparation. IEC collaborates closely with the International Organization for Standardization (ISO) in accordance with conditions determined by agreement between the two organizations.
- 2) The formal decisions or agreements of IEC on technical matters express, as nearly as possible, an international consensus of opinion on the relevant subjects since each technical committee has representation from all interested IEC National Committees.
- 3) IEC Publications have the form of recommendations for international use and are accepted by IEC National Committees in that sense. While all reasonable efforts are made to ensure that the technical content of IEC Publications is accurate, IEC cannot be held responsible for the way in which they are used or for any misinterpretation by any end user.
- 4) In order to promote international uniformity, IEC National Committees undertake to apply IEC Publications transparently to the maximum extent possible in their national and regional publications. Any divergence between any IEC Publication and the corresponding national or regional publication shall be clearly indicated in the latter.
- 5) IEC itself does not provide any attestation of conformity. Independent certification bodies provide conformity assessment services and, in some areas, access to IEC marks of conformity. IEC is not responsible for any services carried out by independent certification bodies.
- 6) All users should ensure that they have the latest edition of this publication.
- 7) No liability shall attach to IEC or its directors, employees, servants or agents including individual experts and members of its technical committees and IEC National Committees for any personal injury, property damage or other damage of any nature whatsoever, whether direct or indirect, or for costs (including legal fees) and expenses arising out of the publication, use of, or reliance upon, this IEC Publication or any other IEC Publications.
- 8) Attention is drawn to the Normative references cited in this publication. Use of the referenced publications is indispensable for the correct application of this publication.
- 9) Attention is drawn to the possibility that some of the elements of this IEC Publication may be the subject of patent rights. IEC shall not be held responsible for identifying any or all such patent rights.

International Standard IEC 60825-1 has been prepared by IEC technical committee 76: Optical radiation safety and laser equipment.

This third edition of IEC 60825-1 cancels and replaces the second edition published in 2007. It constitutes a technical revision.

This edition includes the following significant technical changes with respect to the previous edition:

- a new class, Class 1C, was introduced;
- the measurement condition 2 ("eye loupe" condition) was removed;
- classification of the emission of laser products below a certain radiance level that are intended to be used as replacement for conventional light sources can, as an option, be based on the IEC 62471 series;

the accessible emission limits (AELs) for Class 1, 1M, 2, 2M and 3R of pulsed sources, particularly of pulsed extended sources, were updated to reflect the latest revision of the ICNIRP guidelines on exposure limits (accepted for publication in Health Physics 105 (3): 271 – 295; 2013, see also www.icnirp.org).

This part of IEC 60825 has the status of a Group Safety Publication, in accordance with IEC Guide 104²), for aspects of laser radiation pertaining to human safety.

The text of this standard is based on the following documents:

FDIS	Report on voting	
76/502/FDIS	76/506/RVD	

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

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

The list of all parts of the IEC 60825 series, published under the title *Safety of laser products*, can be found on the IEC website.

This part of IEC 60825 is also referred to as "Part 1" in this publication.

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

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

The contents of the corrigendum of the interpretation sheets 1 (December 2017) and 2 (December 2017) have been included in this copy.

IMPORTANT – The 'colour inside' logo on the cover page of this publication indicates that it contains colours which are considered to be useful for the correct understanding of its contents. Users should therefore print this document using a colour printer.

IEC Guide 104:2010, The preparation of safety publications and the use of basic safety publications and group safety publications

It gives guidance to IEC technical committees and to writers of specifications concerning the manner in which safety publications should be drafted.

This guide does not constitute a normative reference and reference to it is given for information only.

SAFETY OF LASER PRODUCTS -

Part 1: Equipment classification and requirements

1 Scope and object

IEC 60825-1 is applicable to safety of laser products emitting laser radiation in the wavelength range 180 nm to 1 mm.

Although lasers exist which emit at wavelengths less than 180 nm (within the vacuum ultraviolet), these are not included in the scope of the standard since the laser beam normally has to be enclosed in an evacuated enclosure, and, therefore, the potential optical radiation hazards are inherently minimal.

A laser product may consist of a single laser with or without a separate power supply or may incorporate one or more lasers in a complex optical, electrical, or mechanical system. Typically, laser products are used for demonstration of physical and optical phenomena, materials processing, data reading and storage, transmission and display of information, etc. Such systems have found use in industry, business, entertainment, research, education, medicine and consumer products.

Laser products that are sold to other manufacturers for use as components of any system for subsequent sale are not subject to IEC 60825-1, since the final product will itself be subject to this standard. Laser products that are sold by or for manufacturers of end products for use as repair parts for the end products are also not subject to IEC 60825-1. However, if the laser system within the laser product is operable when removed from the end product, the requirements of this Part 1 apply to the removable laser system.

NOTE 1 Operable equipment does not require a tool to prepare for operation.

Any laser product is exempt from all further requirements of this Part 1 if classification by the manufacturer of that product according to Clauses 4 and 5 shows that the emission level does not exceed the AEL (accessible emission limit) of Class 1 under all conditions of operation, maintenance, service and failure. Such a laser product may be referred to as an exempt laser product.

NOTE 2 The above exemption is to ensure that inherently safe laser products are exempt from Clauses 6,7,8 and 9.

In addition to the adverse effects potentially resulting from exposure to laser radiation, some laser equipment may also have other associated hazards, such as electricity, chemicals and high or low temperatures. Laser radiation may cause temporary visual impairment, such as dazzle and glare. Such effects depend on the task and ambient lighting level and are beyond the scope of this Part 1. The classification and other requirements of this standard are intended to address only the laser radiation hazards to the eyes and skin. Other hazards are not included within its scope.

EVS-EN 60825-1:2014+A11:2021 - 10 -

And the MPE values of the skin), if applicable, as well as specific requirements for the specific requirements of the skin is not necessarily limited to the skin performance and testing of the skin), if applicable as the exposure of the skin is not necessarily limited to the skin the specific requirements for the safe system and testing of the safe system and testing of the safe system and testing of the safe product share the text as the exposure of the skin is not necessarily limited to the model that prevents hazardous emission towards the eye. Depending on the type of the product, laser products such as for example medical lasers, machines or toys can be required to conform to the applicable performance and testing required to safety standards.

NOTE 3 See 3.92 for "general laser product".

Where a laser system forms a part of equipment which is subject to another IEC product safety standard, e.g. for medical equipment (IEC 60601-2-22), IT equipment (IEC 60950 series), audio and video equipment (IEC 60065), audio-video and IT equipment (IEC 62368-1), electrical equipment for measurement, control, and laboratory use (IEC 61010-1), equipment for use in hazardous atmospheres (IEC 60079), or electric toys (IEC 62115), this Part 1 will apply in accordance with the provisions of IEC Guide 104² for hazards resulting from laser radiation. (And

For ophthalmic instruments, to ensure patient safety, ISO 15004-2 should be consulted and the principles of the limits provided there should be applied for laser radiation (see also Annex C and D).

In previous editions, light-emitting diodes (LEDs) were included in the scope of IEC 60825-1, and they may be still included in other parts of the IEC 60825 series. However, with the development of lamp safety standards, optical radiation safety of LEDs in general can be more appropriately addressed by lamp safety standards. The removal of LEDs from the scope of this Part 1 does not preclude other standards from including LEDs whenever they refer to lasers. IEC 62471 may be applied to determine the risk group of an LED or product incorporating one or more LEDs. Some other (vertical) standards may require the application of the measurement, classification, engineering specifications and labelling requirements of this standard (IEC 60825-1) to LED products.

Laser products with accessible radiance below the criteria specified in 4.4, designed to function as conventional light sources, and which satisfy the requirements specified in 4.4 may alternatively be evaluated under the IEC 62471 series of standards, "Photobiological safety of lamps and lamp systems". Such a product remains within the scope of this part of IEC 60825, except that the above-described optical radiation emission need not be considered for classification.

The MPE (maximum permissible exposure) values provided in Annex A were developed for laser radiation and do not apply to collateral radiation. However, if a concern exists that accessible collateral radiation might be hazardous, the laser MPE values may be applied to conservatively evaluate this potential hazard, or the exposure limit values in IEC 62471 should be consulted.

The MPE values in Annex A are not applicable to intentional human exposure to laser radiation for the purpose of medical or cosmetic/aesthetic treatment.

NOTE 4 Informative Annexes A to G have been included for purposes of general guidance and to illustrate many typical cases. However, the annexes are not regarded as definitive or exhaustive.

The objectives of this part of IEC 60825 are the following:

- to introduce a system of classification of lasers and laser products emitting radiation in the wavelength range 180 nm to 1 mm according to their degree of optical radiation hazard in order to aid hazard evaluation and to aid the determination of user control measures;
- to establish requirements for the manufacturer to supply information so that proper precautions can be adopted;
- to ensure, through labels and instructions, adequate warning to individuals of hazards associated with accessible radiation from laser products;
- to reduce the possibility of injury by minimizing unnecessary accessible radiation and to give improved control of the laser radiation hazards through protective features.

2 Normative references

The following documents, in whole or in part, are normatively referenced in this document and are indispensable for its application. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

IEC 60050 (all parts), International Electrotechnical Vocabulary (available at http://www.electropedia.org)

IEC 62471 (all parts), Photobiological safety of lamps and lamp systems

3 Terms and definitions

For the purposes of this document, the definitions given in IEC 60050-845 as well as the following apply.

NOTE For convenience here, the definitions have been arranged in English alphabetical order. Departures from IEC 60050-845 are intentional and are indicated. In such cases, reference is made, between brackets, to the definition of Part 845 of IEC 60050, with the mention "modified".

3.1

access panel

part of the protective housing which provides access to laser radiation when removed or displaced

3.2

accessible emission

level of radiation determined at a position and with aperture stops (when the AEL is given in units of Watts or Joules) or limiting apertures (when the AEL is given in units of $W \cdot m^{-2}$ or $J \cdot m^{-2}$) as described in Clause 5

Note 1 to entry: The accessible emission is determined where human access is considered, as specified in Definition 3.40. The accessible emission (determined during operation) is compared with the accessible emission limit (Entry 3.3) in order to determine the class of the laser product. In the body of the standard, whenever the term "emission level" is used, it is to be understood as accessible emission.

Note 2 to entry: When the beam is larger than the aperture stop, the accessible emission when given in units of watts or joules is less than the total emitted power or energy of the laser product. When the beam is smaller than the limiting aperture, the accessible emission when given in units of $W \cdot m^{-2}$ or $J \cdot m^{-2}$, i.e. as irradiance or radiant exposure averaged over the limiting aperture, is smaller than the actual irradiance or radiant exposure of the beam. See also aperture stop (3.9) and limiting aperture (3.55).

3.3 accessible emission limit AEL

maximum accessible emission permitted within a particular class