

Ophthalmic optics — Spectacle lenses — Fundamental requirements for uncut finished lenses (ISO 14889:2003)

ICS 11.040.70

National foreword

This British Standard is the UK implementation of EN ISO 14889:2009. It is identical to ISO 14889:2003. It supersedes BS EN ISO 14889:2003 which is withdrawn.

The UK participation in its preparation was entrusted to Technical Committee CH/172/3, Spectacles.

A list of organizations represented on this committee can be obtained on request to its secretary.

This publication does not purport to include all the necessary provisions of a contract. Users are responsible for its correct application.

Compliance with a British Standard cannot confer immunity from legal obligations.

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Amendments/corrigenda issued since publication

Date	Comments

English Version

**Ophthalmic optics - Spectacle lenses - Fundamental
requirements for uncut finished lenses (ISO 14889:2003)**

Optique ophtalmique - Verres de lunettes - Exigences
fondamentales relatives aux verres finis non détournés (ISO
14889:2003)

Augenoptik - Brillengläser - Grundlegende Anforderungen
an rohkantige fertige Brillengläser (ISO 14889:2003)

This European Standard was approved by CEN on 7 March 2009.

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Foreword

The text of ISO 14889:2003 has been prepared by Technical Committee ISO/TC 172 “Optics and optical instruments” of the International Organization for Standardization (ISO) and has been taken over as EN ISO 14889:2009 by Technical Committee CEN/TC 170 “Ophthalmic optics” the secretariat of which is held by DIN.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by September 2009, and conflicting national standards shall be withdrawn at the latest by March 2010.

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

This document supersedes EN ISO 14889:2003.

This document has been prepared under a mandate given to CEN by the European Commission and the European Free Trade Association, and supports essential requirements of EU Directive 93/42/EEC, as amended by Directive 2007/47/EC.

For relationship with EU Directive 93/42/EEC as amended by Directive 2007/47/EC, see informative Annex ZA, which is an integral part of this document.

According to the CEN/CENELEC Internal Regulations, the national standards organizations of the following countries are bound to implement this European Standard: Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland and the United Kingdom.

Endorsement notice

The text of ISO 14889:2003 has been approved by CEN as a EN ISO 14889:2009 without any modification.

Annex ZA (informative)

Relationship between this European Standard and the Essential Requirements of EU Directive 93/42/EEC

This European Standard has been prepared under a mandate given to CEN by the European Commission and the European Free Trade Association to provide a means of conforming to Essential Requirements of the New Approach Directive 93/42/EEC on medical devices.

Once this standard is cited in the Official Journal of the European Communities under that Directive and has been implemented as a national standard in at least one Member State, compliance with the clauses of this standard given in Table ZA.1 confers, within the limits of the scope of this standard, a presumption of conformity with the corresponding Essential Requirements of that Directive and associated EFTA regulations.

Table ZA.1 – Correspondence between this European Standard and Directive 93/42/EEC

Clause(s)/sub-clause(s) of this EN	Essential Requirements (ERs) of Directive 93/42/EEC	Qualifying remarks/Notes
4, 5	I.1, I.3	
4.2, 4.3.1	I.2, I.6, I.6 a)	New ER I.6 a) is implicitly covered by subclause 4.3.1 of EN ISO 14889. Manufacturers will have to take new ER I.6.a) into consideration when preparing their declaration of conformity.
4.3.1, 4.3.2, 4.4, 5.2, 5.3	II.7.1	
4.4, 5.3	II.9.2	
4.3.2, 5.2	II.9.3	
6	II.13.1, II.13.3	The relevant Essential Requirement of II.13.3 a) is only partly addressed in subclause 6.1 e) of EN ISO 14889.