

Irish Standard
I.S. EN ISO 11979-9:2006&A1:2014

Ophthalmic implants - Intraocular lenses -Part 9: Multifocal intraocular lenses (ISO 11979-9:2006)

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I.S. EN ISO 11979-9:2006&A1:2014

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English Version

Ophthalmic implants - Intraocular lenses - Part 9: Multifocal intraocular lenses (ISO 11979-9:2006/Amd 1:2014)

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This amendment A1 modifies the European Standard EN ISO 11979-9:2006; it was approved by CEN on 18 July 2014.

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EN ISO 11979-9:2006/A1:2014 (E)

Contents	Pag
Foreword	

EN ISO 11979-9:2006/A1:2014 (E)

Foreword

This document (EN ISO 11979-9:2006/A1:2014) has been prepared by Technical Committee ISO/TC 172 "Optics and photonics" in collaboration with Technical Committee CEN/TC 170 "Ophthalmic optics" the secretariat of which is held by DIN.

This Amendment to the European Standard EN ISO 11979-9:2006 shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by February 2015, and conflicting national standards shall be withdrawn at the latest by February 2015.

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Endorsement notice

The text of ISO 11979-9:2006/Amd, 1:2014 has been approved by CEN as EN ISO 11979-9:2006/A1:2014 without any modification.

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EN ISO 11979-9

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EN ISO 11979-9:2006 (E)

Foreword

This document (EN ISO 11979-9:2006) has been prepared by Technical Committee ISO/TC 172 "Optics and optical instruments" in collaboration with 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of an identical text or by endorsement, at the latest by March 2007, and conflicting national standards shall be withdrawn at the latest by March 2007.

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INTERNATIONAL STANDARD

ISO 11979-9

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Ophthalmic implants — Intraocular lenses —

Part 9:

Multifocal intraocular lenses

Implants ophtalmiques — Lentilles intraoculaires —
Partie 9: Lentilles intraoculaires multifocales



ISO 11979-9:2006(E)

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Contents Page Forewordiv Scope 1 2 3 4 Physical requirements 2 4.1 4.2 5 Optical requirements ______2 5.1 5.2 5.3 5.4 6 Clinical investigation ______3 6.1 6.2 Additional requirements for the clinical investigation plan4 7 Information supplied by the manufacturer4

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.

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ISO 11979-9 was prepared by Technical Committee ISO/TC 172, *Optics and photonics*, Subcommittee SC 7, *Ophthalmic optics and instruments*.

ISO 11979 consists of the following parts, under the general title Ophthalmic implants — Intraocular lenses:

- Part 1: Vocabulary
- Part 2: Optical properties and test methods
- Part 3: Mechanical properties and test methods
- Part 4: Labelling and information
- Part 5: Biocompatibility
- Part 6: Shelf-life and transport stability
- Part 7: Clinical investigations
- Part 8: Fundamental requirements
- Part 9: Multifocal intraocular lenses
- Part 10: Phakic intraocular lenses

Ophthalmic implants — Intraocular lenses —

Part 9:

Multifocal intraocular lenses

1 Scope

This part of ISO 11979 is applicable to any intraocular lens whose optic provides two or more rotationally symmetric powers and whose primary indication is the correction of aphakia with the added benefit of useful vision at more than one distance (e.g. far and near).

NOTE The term "near vision" as used in this part of ISO 11979 includes useful vision at a distance of claimed benefit; e.g. near and/or intermediate distances.

2 Normative references

The following referenced documents are indispensable for the application 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.

- ISO 11979-1, Ophthalmic implants Intraocular lenses Part 1: Vocabulary
- ISO 11979-2, Ophthalmic implants Intraocular lenses Part 2: Optical properties and test methods
- ISO 11979-3, Ophthalmic implants Intraocular lenses Part 3: Mechanical properties and test methods
- ISO 11979-4, Ophthalmic implants Intraocular lenses Part 4: Labelling and information
- ISO 11979-7, Ophthalmic implants Intraocular lenses Part 7: Clinical investigations
- ISO 14155-1, Clinical investigation of medical devices for human subjects Part 1: General requirements
- ISO 14155-2, Clinical investigation of medical devices for human subjects Part 2: Clinical investigation plans
- ISO 14971, Medical devices Application of risk management to medical devices

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 11979-1, ISO 14155-1 and ISO 14155-2 apply.



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