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STN	**Očné implantáty Vnútroočné šošovky Časť 4: Označovanie a informácie (ISO 11979-4: 2026)**	**STN EN ISO 11979-4 19 5300**

Ophthalmic implants - Intraocular lenses - Part 4: Labelling and information (ISO 11979-4:2026)

Táto norma obsahuje anglickú verziu európskej normy.
This standard includes the English version of the European Standard.

Táto norma bola oznámená vo Vestníku ÚNMS SR č. 05/26

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EN ISO 11979-4

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English Version

Ophthalmic implants - Intraocular lenses - Part 4: Labelling and information (ISO 11979-4:2026)

Implants ophtalmiques - Lentilles intraoculaires - Partie 4: Étiquetage et informations (ISO 11979-4:2026)

Ophthalmische Implantate - Intraokularlinsen - Teil 4: Etikettierung und Information (ISO 11979-4:2026)

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EUROPEAN COMMITTEE FOR STANDARDIZATION
COMITÉ EUROPÉEN DE NORMALISATION
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CEN-CENELEC Management Centre: Rue de la Science 23, B-1040 Brussels

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Contents

Page

European foreword

This document (EN ISO 11979-4:2026) 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 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of an identical text or by endorsement, at the latest by September 2026, and conflicting national standards shall be withdrawn at the latest by September 2026.

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

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Endorsement notice

The text of ISO 11979-4:2026 has been approved by CEN as EN ISO 11979-4:2026 without any modification.

International Standard

ISO 11979-4

Ophthalmic implants — Intraocular lenses —

Part 4: Labelling and information

Implants ophtalmiques — Lentilles intraoculaires —
Partie 4: Étiquetage et informations

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ISO copyright office
CP 401 • Ch. de Blandonnet 8
CH-1214 Vernier, Geneva
Phone: +41 22 749 01 11
Email: copyright@iso.org
Website: www.iso.org

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Contents

Page

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.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular, the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

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For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT), see www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 172, *Optics and photonics*, Subcommittee SC 7, *Ophthalmic optics and instruments*, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/TC 170, *Ophthalmic optics,* in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

This third edition cancels and replaces the second edition (ISO 11979-4:2008), which has been technically revised. It also incorporates the Amendment ISO 11979-4:2008/Amd.1:2012.

The main changes are as follows:

— normative references have been updated and retired standards have been removed or replaced;

— Table 1 has been updated with additional information to be included in the packaging; e.g. expiration date on primary package;

— new categories and clinical requirements for SVIOLs were published in ISO 11979-7. ISO 11979-4 is updated to reflect these changes also in labelling;

— information can be provided electronically if national regulations allow.

A list of all parts in the ISO 11979 series can be found on the ISO website.

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.

Introduction

Labelling requirements for medical devices in general are given in ISO 20417[5]. However, in order to ensure correct and necessary information to the ophthalmic surgeon, additional specific information is required for intraocular lenses. Such information concerns technical and optical data as well as information about the materials used.

International Standard **ISO 11979-4:2026(en)**

Ophthalmic implants — Intraocular lenses —

Part 4:
Labelling and information

1 Scope

This document specifies the labelling requirements for intraocular lenses (IOLs) and the information to be provided within or on the packaging.

NOTE This document attempts to harmonize the recognized labelling requirements for IOLs throughout the world. However, there can be additional national requirements.

2 Normative references

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.

ISO 11979-1, *Ophthalmic implants — Intraocular lenses — Part 1: Vocabulary*

ISO 11979-7, *Ophthalmic implants — Intraocular lenses — Part 7: Clinical investigations of intraocular lenses for the correction of aphakia*