



SLOVENSKI STANDARD

SIST EN 556-1:2024

01-september-2024

Nadomešča:

SIST EN 556-1:2002/AC:2006

Sterilizacija medicinskih pripomočkov - Zahteve za medicinske pripomočke, ki morajo biti označeni s "STERILNO" - 1. del: Zahteve za končno sterilizirane medicinske pripomočke

Sterilization of medical devices - Requirements for medical devices to be designated "STERILE" - Part 1: Requirements for terminally sterilized medical devices

Sterilisation von Medizinprodukten - Anforderungen an Medizinprodukte, die als "STERIL" gekennzeichnet werden - Teil 1: Anforderungen an Medizinprodukte, die in der Endpackung sterilisiert wurden

Stérilisation des dispositifs médicaux - Exigences relatives aux dispositifs médicaux en vue d'obtenir l'étiquetage « STÉRILE » - Partie 1 : Exigences relatives aux dispositifs médicaux stérilisés de façon terminale

Ta slovenski standard je istoveten z: EN 556-1:2024

ICS:

11.080.01 Sterilizacija in dezinfekcija na splošno Sterilization and disinfection in general

SIST EN 556-1:2024

en,fr,de

EUROPEAN STANDARD

EN 556-1

NORME EUROPÉENNE

EUROPÄISCHE NORM

July 2024

ICS 11.080.01

Supersedes EN 556-1:2001

English Version

Sterilization of medical devices - Requirements for medical devices to be designated "STERILE" - Part 1: Requirements for terminally sterilized medical devices

Stérilisation des dispositifs médicaux - Exigences relatives aux dispositifs médicaux en vue d'obtenir l'étiquetage " STÉRILE " - Partie 1 : Exigences relatives aux dispositifs médicaux stérilisés de façon terminale

Sterilisation von Medizinprodukten - Anforderungen an Medizinprodukte, die als "STERIL" gekennzeichnet werden - Teil 1: Anforderungen an Medizinprodukte, die in der Endpackung sterilisiert wurden

This European Standard was approved by CEN on 19 May 2024.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard 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 CEN member.

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

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

<https://standards.iteh.ai/catalog/standards/sist/c13cad91-5d6c-48f2-9aba-f329ce97a2f7/sist-en-556-1-2024>



EUROPEAN COMMITTEE FOR STANDARDIZATION
COMITÉ EUROPÉEN DE NORMALISATION
EUROPÄISCHES KOMITEE FÜR NORMUNG

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

Contents	Page
European foreword	3
Introduction	4
1 Scope	5
2 Normative references	5
3 Terms and definitions	5
4 Requirements	7
Annex ZA (informative) Relationship between this European Standard and the General Safety and Performance Requirements of Regulation (EU) 2017/745 aimed to be covered	8
Annex ZB (informative) Relationship between this European Standard and the General Safety and Performance Requirements of Regulation (EU) 2017/746 aimed to be covered	11
Bibliography	13

iTeh Standards
(<https://standards.iteh.ai>)
Document Preview

[SIST EN 556-1:2024](https://standards.iteh.ai/catalog/standards/sist/c13cad91-5d6c-48f2-9aba-f329ce97a2f7/sist-en-556-1-2024)

<https://standards.iteh.ai/catalog/standards/sist/c13cad91-5d6c-48f2-9aba-f329ce97a2f7/sist-en-556-1-2024>

European foreword

This document (EN 556-1:2024) has been prepared by Technical Committee CEN/TC 204, Sterilization of medical devices, the secretariat of which is held by BSI.

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 January 2025, and conflicting national standards shall be withdrawn at the latest by January 2025.

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.

This document supersedes EN 556-1:2001 and EN 556-1:2001/AC:2006.

This document has been prepared under a Standardization Request given to CEN by the European Commission and the European Free Trade Association, and supports essential requirements of EU Directive(s) / Regulation(s). For relationship with EU Directive(s) / Regulation(s), see informative Annexes ZA and ZB, which is an integral part of this document.

EN 556, *Sterilization of medical devices — Requirements for medical devices to be designated “STERILE”*, is currently composed with the following parts:

- *Part 1: Requirements for terminally sterilized medical devices* [this document];
- *Part 2: Requirements for aseptically processed medical devices*.

EN 556-1:2024 includes the following significant technical changes with respect to EN 556-1:2001 and EN 556-1:2001/AC:2006:

- definitions have been aligned with EN ISO 11139;
- the normative reference has been updated to the latest edition;
- informative Annex ZA has been replaced with informative Annexes ZA and ZB giving the relationship with the European Regulations for medical devices and *in vitro* diagnostic medical devices respectively;
- the Bibliography has been updated.

Any feedback and questions on this document should be directed to the users' national standards body. A complete listing of these bodies can be found on the CEN website.

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

EN 556-1:2024 (E)

Introduction

A sterile product item is one which is free of viable microorganisms. European standards for *medical devices* require, when it is necessary to supply a *sterile* product item, that adventitious microbiological contamination of a medical device from all sources is minimized by all practical means. Even so, product items produced under standard manufacturing conditions in accordance with their requirements for quality management systems for medical devices (see EN ISO 13485:2016 and EN ISO 13485:2016/A11:2021) can, prior to sterilization, have microorganisms on them, albeit in low numbers. Such product items are non-sterile. The purpose of sterilization processing is to inactivate the microbiological contaminants and thereby transform the non-sterile items into sterile ones.

The inactivation of a pure culture of microorganisms by physical and/or chemical agents used to sterilize *medical devices* often approximates to an exponential relationship; inevitably this means that, regardless of the extent of treatment applied, there is always a finite probability that a microorganism will survive. For a given treatment, the probability of survival is determined by the number and resistance of microorganisms and by the environment in which the organisms exist during treatment. It follows that the sterility of any one item subjected to sterilization processing cannot be guaranteed and the sterility of the processed items has to be defined in terms of the probability of the existence of a surviving microorganism on/in an item. The standards for quality management systems recognize that there are processes used which cannot be fully verified by subsequent inspection and testing of product. Sterilization is an example of such a process. Sterilization processes are validated before use, the performance of the process monitored routinely and the equipment maintained.

It is important to be aware that exposure to a properly validated and accurately controlled sterilization process is not the only factor associated with the provision of assurance that the product item is *sterile* and, in this respect, suitable for its intended use. Attention has also to be given to a number of factors including the microbiological status (*bioburden*) of incoming raw materials and/or components, their subsequent storage and to the control of the environment in which the product is manufactured, assembled and packaged.

[SIST EN 556-1:2024](https://standards.iteh.ai/catalog/standards/sist/c13cad91-5d6c-48f2-9aba-f329ce97a2f7/sist-en-556-1-2024)

<https://standards.iteh.ai/catalog/standards/sist/c13cad91-5d6c-48f2-9aba-f329ce97a2f7/sist-en-556-1-2024>