



SLOVENSKI STANDARD

SIST EN ISO/IEEE 11073-10103:2025

01-december-2025

Nadomešča:

SIST EN ISO 11073-10103:2014

Zdravstvena informatika - Interoperabilnost naprav - 10103. del: Nomenklatura, pripomoček za vsaditev, srčni (ISO/IEEE 11073-10103:2025)

Health informatics - Device interoperability - Part 10103: Nomenclature, implantable device, cardiac (ISO/IEEE 11073-10103:2025)

Medizinische Informatik - Kommunikation patientennaher medizinischer Geräte - Teil 10103: Nomenklatur - Implantierbare kardiologische Geräte (ISO/IEEE 11073-10103:2025)

Informatique de santé - Interopérabilité des dispositifs - Partie 10103: Nomenclature, dispositif implantable, cardiaque (ISO/IEEE 11073-10103:2025)

Ta slovenski standard je istoveten z: **EN ISO/IEEE 11073-10103:2025**

ICS:

11.040.40	Implantanti za kirurgijo, protetiko in ortetiko	Implants for surgery, prosthetics and orthotics
35.240.80	Uporabniške rešitve IT v zdravstveni tehniki	IT applications in health care technology

SIST EN ISO/IEEE 11073-10103:2025 en,fr,de

EUROPEAN STANDARD
NORME EUROPÉENNE
EUROPÄISCHE NORM

**EN ISO/IEEE 11073-
10103**

October 2025

ICS 11.040.40; 35.240.80

Supersedes EN ISO 11073-10103:2013

English Version

**Health informatics - Device interoperability - Part 10103:
Nomenclature, implantable device, cardiac (ISO/IEEE
11073-10103:2025)**

Informatique de santé - Interopérabilité des dispositifs
- Partie 10103: Nomenclature, dispositif implantable,
cardiaque (ISO/IEEE 11073-10103:2025)

Medizinische Informatik - Kommunikation
patientennaher medizinischer Geräte - Teil 10103:
Nomenklatur - Implantierbare kardiologische Geräte
(ISO/IEEE 11073-10103:2025)

This European Standard was approved by CEN on 26 September 2025.

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.



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

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

EN ISO/IEEE 11073-10103:2025 (E)

Contents	Page
European foreword.....	3

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

[SIST EN ISO/IEEE 11073-10103:2025](https://standards.iteh.ai/catalog/standards/sist/fd498e15-936d-4b4a-b2ad-9748efa207b9/sist-en-iso-ieee-11073-10103-2025)

<https://standards.iteh.ai/catalog/standards/sist/fd498e15-936d-4b4a-b2ad-9748efa207b9/sist-en-iso-ieee-11073-10103-2025>

European foreword

This document (EN ISO/IEEE 11073-10103:2025) has been prepared by Technical Committee ISO/TC 215 "Health informatics" in collaboration with Technical Committee CEN/TC 251 "Health informatics" the secretariat of which is held by NEN.

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 April 2026, and conflicting national standards shall be withdrawn at the latest by April 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.

This document supersedes EN ISO 11073-10103:2013.

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

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, 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.

(<https://standards.iteh.ai>)
Endorsement notice

The text of ISO/IEEE 11073-10103 has been approved by CEN as EN ISO/IEEE 11073-10103:2025 without any modification.

<https://standards.iteh.ai/catalog/standards/sist/fd498e15-936d-4b4a-b2ad-9748efa207b9/sist-en-iso-ieee-11073-10103-2025>



**International
Standard**

**ISO/IEEE
11073-10103**

**Health informatics — Device
interoperability —**

Part 10103:

Nomenclature, implantable Standards

device, cardiac

Informatique de santé — Interopérabilité des dispositifs —

Partie 10103: Nomenclature, dispositif implantable, cardiaque

**Second edition
2025-10**

SIST EN ISO/IEEE 11073-10103:2025

<https://standards.iteh.ai/catalog/standards/sist/fd498e15-936d-4b4a-b2ad-9748efa207b9/sist-en-iso-ieee-11073-10103-2025>