



SLOVENSKI STANDARD SIST EN ISO 80369-6:2025

01-julij-2025

Nadomešča:

SIST EN ISO 80369-6:2016

Priključki z majhnim premerom za tekočine in pline za uporabo v zdravstvu - 6. del: Priključki za nevronske uporabe (ISO 80369-6:2025)

Small bore connectors for liquids and gases in healthcare applications - Part 6:
Connectors for neural applications (ISO 80369-6:2025)

Verbindungsstücke mit kleinem Durchmesser für Flüssigkeiten und Gase in
medizinischen Anwendungen - Teil 6: Verbindungsstücke für neurale Anwendungen
(ISO 80369-6:2025)

Raccords de petite taille pour liquides et gaz utilisés dans le domaine de la santé - Partie
6: Raccords pour applications neurales (ISO 80369-6:2025)

Ta slovenski standard je istoveten z: EN ISO 80369-6:2025

ICS:

11.040.25	Injekcijske brizge, igle in katetri	Syringes, needles and catheters
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en

EUROPEAN STANDARD

EN ISO 80369-6

NORME EUROPÉENNE

EUROPÄISCHE NORM

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ICS 11.040.25

Supersedes EN ISO 80369-6:2016

English version

Small bore connectors for liquids and gases in healthcare applications - Part 6: Connectors for neural applications (ISO 80369-6:2025)

Raccords de petite taille pour liquides et gaz utilisés dans le domaine de la santé - Partie 6: Raccords pour applications neurales (ISO 80369-6:2025)

Verbindungsstücke mit kleinem Durchmesser für Flüssigkeiten und Gase in medizinischen Anwendungen - Teil 6: Verbindungsstücke für neurale Anwendungen (ISO 80369-6:2025)

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

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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 and CENELEC member into its own language and notified to the CEN-CENELEC Management Centre has the same status as the official versions.

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European foreword

This document (EN ISO 80369-6:2025) has been prepared by Technical Committee ISO/TC 210 "Quality management and corresponding general aspects for products with a health purpose including medical devices" in collaboration with Technical Committee CEN-CENELEC/ JTC 3 "Quality management and corresponding general aspects for medical devices" 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 December 2025, and conflicting national standards shall be withdrawn at the latest by December 2025.

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

This document supersedes EN ISO 80369-6:2016.

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 and CENELEC websites.

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.

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Endorsement notice

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