

STN	Infúzne prístroje na zdravotnícke účely Časť 3: Hliníkové uzávery na infúzne fľaše (ISO 8536-3:2009/Adm 1: 2022) Zmena A1	STN EN ISO 8536-3/A1 70 3350
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Infusion equipment for medical use - Part 3: Aluminium caps for infusion bottles (ISO 8536-3:2009)

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. 12/22

Obsahuje: EN ISO 8536-3:2009/A1:2022, ISO 8536-3:2009/Amd 1:2022

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EUROPEAN STANDARD

EN ISO 8536-3:2009/A1

NORME EUROPÉENNE

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English Version

Infusion equipment for medical use - Part 3: Aluminium caps for infusion bottles - Amendment 1 (ISO 8536-3:2009/Amd 1:2022)

Matériel de perfusion à usage médical - Partie 3:
Capsules en aluminium pour flacons de perfusion -
Amendement 1 (ISO 8536-3:2009/Amd 1:2022)

Infusionsgeräte zur medizinischen Verwendung - Teil
3: Aluminium-Bördelkappen für Infusionsflaschen -
Änderung 1 (ISO 8536-3:2009/Amd 1:2022)

This amendment A1 modifies the European Standard EN ISO 8536-3:2009; it was approved by CEN on 19 March 2022.

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EN ISO 8536-3:2009/A1:2022 (E)

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European foreword

This document (EN ISO 8536-3:2009/A1:2022) has been prepared by Technical Committee ISO/TC 76 "Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use" in collaboration with CCMC.

This Amendment to the European Standard EN ISO 8536-3:2009 shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by March 2023, and conflicting national standards shall be withdrawn at the latest by March 2023.

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 8536-3:2009/Amd 1:2022 has been approved by CEN as EN ISO 8536-3:2009/A1:2022 without any modification.

INTERNATIONAL STANDARD

ISO 8536-3

Third edition
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AMENDMENT 1
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Infusion equipment for medical use — Part 3: Aluminium caps for infusion bottles AMENDMENT 1

Matériel de perfusion à usage médical —

Partie 3: Capsules en aluminium pour flacons de perfusion

AMENDEMENT 1



Reference number
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Foreword

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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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This document was prepared by Technical Committee ISO/TC 76, *Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use*, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/SS S02, *Transfusion equipment*, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

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