



SLOVENSKI STANDARD
SIST EN ISO 8362-1:2010/A1:2016
01-marec-2016

Vsebniki za parenteralne farmacevtske oblike in dodatna oprema - 1. del: Viale iz cevnega stekla (ISO 8362-1:2009/Amd 1:2015)

Injection containers and accessories - Part 1: Injection vials made of glass tubing (ISO 8362-1:2009/Amd 1:2015)

Injektionsbehälter und Zubehör - Teil 1: Injektionsflaschen aus Röhrglas (ISO 8362-1:2009/Amd 1:2015)

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Réipients et accessoires pour produits injectables - Partie 1: Flacons en verre étiré (ISO 8362-1:2009/Amd 1:2015)

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Ta slovenski standard je istoveten z: EN ISO 8362-1:2009/A1:2015

ICS:

11.040.20	Transfuzijska, infuzijska in injekcijska oprema	Transfusion, infusion and injection equipment
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SIST EN ISO 8362-1:2010/A1:2016 **en**

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EUROPEAN STANDARD

EN ISO 8362-1:2009/A1

NORME EUROPÉENNE

EUROPÄISCHE NORM

December 2015

ICS 11.040.20

English Version

Injection containers and accessories - Part 1: Injection vials made of glass tubing (ISO 8362-1:2009/Amd 1:2015)

Réipients et accessoires pour produits injectables -
Partie 1: Flacons en verre étiré (ISO 8362-1:2009/Amd
1:2015)

Injektionsbehältnisse und Zubehör - Teil 1:
Injektionsflaschen aus Röhrenglas (ISO 8362-
1:2009/Amd 1:2015)

This amendment A1 modifies the European Standard EN ISO 8362-1:2009; it was approved by CEN on 3 October 2015.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for inclusion of this amendment into the relevant 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 amendment 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, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and United Kingdom.



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

CEN-CENELEC Management Centre: Avenue Marnix 17, B-1000 Brussels

Contents	Page
European foreword.....	3

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[SIST EN ISO 8362-1:2010/A1:2016](https://standards.iteh.ai/catalog/standards/sist/6486fa16-b69a-453f-966f-43f0f757e491/sist-en-iso-8362-1-2010-a1-2016)
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European foreword

This document (EN ISO 8362-1:2009/A1:2015) has been prepared by Technical Committee ISO/TC 76 "Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use".

This Amendment to the European Standard EN ISO 8362-1:2009 shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by June 2016, and conflicting national standards shall be withdrawn at the latest by June 2016.

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

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Endorsement notice

The text of ISO 8362-1:2009/Amd 1:2015 has been approved by CEN as EN ISO 8362-1:2009/A1:2015 without any modification.

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<https://standards.iteh.ai/catalog/standards/sist/6486fa16-b69a-453f-966f-43f0f757e491/sist-en-iso-8362-1-2010-a1-2016>

INTERNATIONAL
STANDARD

ISO
8362-1

Third edition
2009-12-15

AMENDMENT 1
2015-12-15

**Injection containers and
accessories —**

Part 1:
Injection vials made of glass tubing

AMENDMENT 1

iTeh STANDARD PREVIEW

*Réipients et accessoires pour produits injectables —
Partie 1: Flacons en verre étiré*

AMENDEMENT 1

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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. www.iso.org/directives

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For an explanation on the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the WTO principles in the Technical Barriers to Trade (TBT), see the following URL: [Foreword - Supplementary information](#)

The committee responsible for this document is ISO/TC 76, *Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use*.

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