INTERNATIONAL STANDARD

ISO 8536-4

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Infusion equipment for medical use — Part 4: Infusion sets for single use, gravity feed

Matériel de perfusion à usage médical —

Partie 4: Appareils de perfusion non réutilisables, à alimentation par gravité

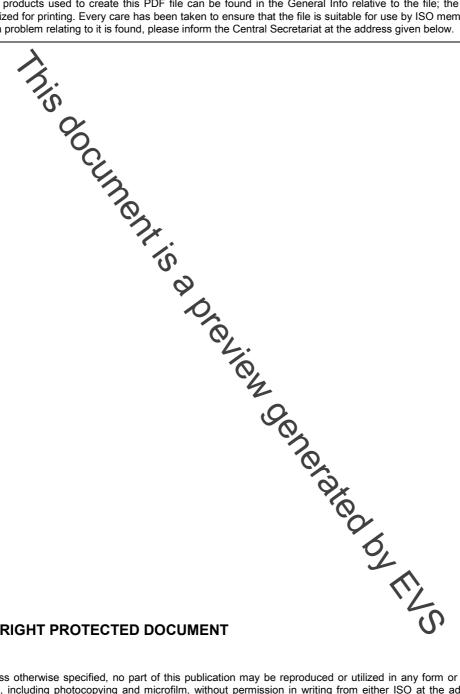


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Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in Maison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical confirmtees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires applying by at least 75 % of the member bodies casting a vote.

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

ISO 8536-4 was prepared by Technical Computere ISO/TC 76, Transfusion, infusion and injection equipment for medical and pharmaceutical use.

This fifth edition cancels and replaces the fourth edition (ISO 8536-4:2007), of which it constitutes a minor revision. In detail, 7.1 was more clarified in alignment with B.2, and A.2.2 was changed in order to go back with the leakage test pressure to 20 kPa and to restrict the leakage test for (40 ± 1) °C.

ISO 8536 consists of the following parts, under the general ute Infusion equipment for medical use:

- Part 1: Infusion glass bottles
- Part 2: Closures for infusion bottles
- Part 3: Aluminium caps for infusion bottles
- Part 4: Infusion sets for single use, gravity feed
- Part 5: Burette infusion sets for single use, gravity feed
- Part 6: Freeze drying closures for infusion bottles
- Generated by the Part 7: Caps made of aluminium-plastics combinations for infusion bottles
- Part 8: Infusion equipment for use with pressure infusion apparatus
- Part 9: Fluid lines for use with pressure infusion equipment
- Part 10: Accessories for fluid lines for use with pressure infusion equipment
- Part 11: Infusion filters for use with pressure infusion equipment
- Part 12: Check valves

Infusion equipment for medical use —

Part 4:

Infusion sets for single use, gravity feed

1 Scope

This part of ISO 8536 specifies requirements for single use, gravity feed infusion sets for medical use in order to ensure their compatibility with containers for infusion solutions and intravenous equipment.

Secondary aims of this part of 150,8536 are to provide guidance on specifications relating to the quality and performance of materials used in massion sets and to present designations for infusion set components.

In some countries, the national pharmacopoeia or other national regulations are legally binding and take precedence over this part of ISO 8536.

2 Normative references

The following referenced documents are indispensable for the application of this document. For dated references, only the edition cited applies. For untated references, the latest edition of the referenced document (including any amendments) applies.

ISO 594-1, Conical fittings with a 6 % (Luer) taper syringes, needles and certain other medical equipment — Part 1: General requirements

ISO 594-2, Conical fittings with 6 % (Luer) taper for syringes, heales and certain other medical equipment — Part 2: Lock fittings

ISO 3696, Water for analytical laboratory use — Specification and test methods

ISO 7864, Sterile hypodermic needles for single use

ISO 14644-1, Cleanrooms and associated controlled environments — Part 4. Classification of air cleanliness 1)

ISO 15223-1, Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied — Part 1: General requirements²⁾

3 General requirements

3.1 The nomenclature to be used for components of infusion sets and of a separate air-inlet device is given in Figures 1, 2 and 3. These figures illustrate examples of the configuration of infusion sets and air-inlet devices; other configurations may be used provided they lead to the same results. Infusion sets as illustrated in Figure 2 should only be used for collapsible plastic containers. Infusion sets as illustrated in Figure 2 used

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¹⁾ Under preparation. (Revision of ISO 14644-1:1999)

²⁾ To be published. (Revision of ISO 15223-1:2007)