
**Elastomeric parts for parenterals and for
devices for pharmaceutical use —**

**Part 3:
Determination of released-particle count**

*Éléments en élastomère pour administration parentérale et dispositifs à
usage pharmaceutique —*

Partie 3: Dénombrement des particules libérées



PDF disclaimer

This PDF file may contain embedded typefaces. In accordance with Adobe's licensing policy, this file may be printed or viewed but shall not be edited unless the typefaces which are embedded are licensed to and installed on the computer performing the editing. In downloading this file, parties accept therein the responsibility of not infringing Adobe's licensing policy. The ISO Central Secretariat accepts no liability in this area.

Adobe is a trademark of Adobe Systems Incorporated.

Details of the software products used to create this PDF file can be found in the General Info relative to the file; the PDF-creation parameters were optimized for printing. Every care has been taken to ensure that the file is suitable for use by ISO member bodies. In the unlikely event that a problem relating to it is found, please inform the Central Secretariat at the address given below.

© ISO 2003

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized in any form or by any means, electronic or mechanical, including photocopying and microfilm, without permission in writing from either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office
Case postale 56 • CH-1211 Geneva 20
Tel. + 41 22 749 01 11
Fax + 41 22 749 09 47
E-mail copyright@iso.org
Web www.iso.org

Published in Switzerland

Contents

Page

Foreword	iv
Introduction	v
1 Scope	1
2 Normative references	1
3 Determination of visible-particle count	1
4 Determination of subvisible-particle count	3
Bibliography	6

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 liaison 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 committees 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 approval 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 8871-3 was prepared by Technical Committee ISO/TC 76, *Transfusion, infusion and injection equipment for medical and pharmaceutical use*.

Together with the other parts (see below), this part of ISO 8871 cancels and replaces ISO 8871:1990, which has been technically revised.

ISO 8871 consists of the following parts, under the general title *Elastomeric parts for parenterals and for devices for pharmaceutical use*:

- *Part 1: Extractables in aqueous autoclavates*
- *Part 2: Identification and characterization*
- *Part 3: Determination of released-particle count*
- *Part 4: Biological requirements and test methods*
- *Part 5: Functional requirements and testing*

Introduction

When elastomeric closures are used as primary packaging materials in direct contact with pharmaceutical preparations, the pharmaceutical industry requires, to an increasing extent, definite details from the rubber manufacturer about the presence of particles the closures may release into an injectable. The test methods specified in Clauses 3 and 4 make it possible to meet this request.

Elastomeric parts for parenterals and for devices for pharmaceutical use —

Part 3: Determination of released-particle count

1 Scope

Elastomeric closures may be superficially contaminated with visible and subvisible particles, and fragments can also be produced when the closure is pierced by a needle.

Such particles may be transferred to pharmaceutical preparations in contact with the elastomeric parts and affect the quality of such preparations.

This part of ISO 8871 specifies methods for the determination of the number of visible and subvisible particles, respectively, detached from elastomeric parts by rinsing.

It does not specify particle contamination limits. These will have to be agreed upon between manufacturer and user.

2 Normative references

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

ISO 3696:1997, *Water for analytical laboratory use — Specification and test methods*

3 Determination of visible-particle count

3.1 Principle

This method evaluates the potential for contamination by collecting and counting the particles detached from elastomeric parts on rinsing with water.

3.2 Classification

For the purposes of this method, particles are divided into size classes as follows, using the longest dimension as the classifying parameter:

Class I: > 25 µm but ≤ 50 µm;

Class II: > 50 µm but ≤ 100 µm;

Class III: > 100 µm.

3.3 Apparatus and materials

3.3.1 Shaking machine, moving in a horizontal circle of $12 \text{ mm} \pm 1 \text{ mm}$ diameter at 300 min^{-1} to 350 min^{-1} .

3.3.2 Membrane filters, with a maximum pore size of $0,8 \text{ }\mu\text{m}$, provided with grid lines dividing the surface into $3 \text{ mm} \times 3 \text{ mm}$ squares.

NOTE The colour of the filter may significantly affect the test results.

If no specific agreement has been made between the interested parties, the basic colour shall be medium grey and lie within the following coordinate ranges in the CIE (International Commission on Illumination) system:

L^* between 60 % and 70 %;

a^* between $-4,7 \%$ and $-3,7 \%$;

b^* between $-4,7 \%$ and $-3,7 \%$.

These specifications apply to the grid-imprinted surface of the filter, and assume this surface has a 3-mm-square grid made up of green lines.

3.3.3 Clean, wide-mouthed conical flasks, of capacity 300 ml.

3.3.4 Rinse fluid, prepared by dissolving 3 g of commercially available highly concentrated polysorbate 80 (Tween 80) in 10 l of purified water (grade 1 or grade 2 as specified in ISO 3696).

3.3.5 Equipment for supplying the rinse fluid under a suitable pressure, including a terminal filter with a maximum pore size of $1,2 \text{ }\mu\text{m}$.

3.3.6 Microscope, magnification about $\times 50$, with suitable direct illumination at an angle of 0° to 10° with the microscope stage.

3.4 Preparation for the test

3.4.1 Ensure that the environment in which the test is to be carried out is free from extraneous particles which could cause interference. This involves wearing suitable garments and gloves and using a suitable clean-air work station, for example a laminar-flow cabinet meeting the requirements of Class 8 of ISO 14644-1 (Class 100,000 of US Federal Standard 209E), as well as suitably decontaminated tools and sample-handling equipment.

3.4.2 Carry out a blank test, as follows:

- Place 50 ml of filtered rinse fluid in a conical flask.
- Shake for 20 s.
- Immediately filter the fluid through a membrane filter.
- Add another 50 ml portion of rinse fluid to the flask, shake, and filter in the same way.
- Transfer the filter to the microscope, taking care not to contaminate it.
- Count the particles on the filter.

No more than five particles of Class I and no more than one particle of Class II shall be found. No particles of Class III shall be present.

If these requirements are not met, investigate the possible causes of failure, rectify, and repeat the blank test until satisfactory results are obtained.

Carry out a blank test both before and after each series of tests. Only when satisfactory values are obtained both before and after the series of tests can the results of the tests be considered valid.

3.5 Test

- Place a number of complete elastomeric parts with a total surface area of approximately 100 cm² into a conical flask.
- Add 50 ml of filtered rinse fluid.
- Shake for 20 s.
- Immediately filter the fluid through a membrane filter.
- Add another 50 ml portion of rinse fluid to the flask, shake and filter in the same way.
- Transfer the filter to the microscope, taking care not to contaminate it.
- Count the particles on the filter.

3.6 Test report

For each test, report the following:

- a) the total surface area of the elastomeric parts tested, and the number of complete parts tested;
- b) the total particle count in each of the three particle-size classes;
- c) the particle counts in each of the three classes for the blank tests performed;
- d) the average particle count in each class per 10 cm² of surface, rounded to one place of decimals.

4 Determination of subvisible-particle count

4.1 Principle

In contact with liquid pharmaceutical preparations, elastomeric parts may release particles having dimensions of 25 µm or smaller, and hence not visible to the naked eye. Their presence can be detected by means of electrical or optical instruments.

This method evaluates the potential of elastomeric parts to release such particles by bringing the parts in contact with water and examining the contact fluid with a suitable particle counter operating on the light extinction principle.

4.2 Classification

For the purpose of this method, particles are divided into size classes as follows, using the equivalent sphere diameter as the classifying parameter:

- ≥ 2 µm but < 5 µm;
- ≥ 5 µm but < 10 µm;
- ≥ 10 µm but < 25 µm;
- ≥ 25 µm.

4.3 Apparatus and materials

- 4.3.1 Shaking machine**, moving in a horizontal circle of 12 mm ± 1 mm diameter at 300 min⁻¹ to 350 min⁻¹.
- 4.3.2 Clean, wide-mouthed conical flasks**, of capacity 300 ml.
- 4.3.3 Particle-free water**, containing not more than 100 particles larger than 2 µm per 5 ml.
- 4.3.4 Particle counter**, operating on the light extinction (light obscuration or light blockage) principle, capable of classifying particles in the rinse fluid into the categories defined in 4.2.

4.4 Preparations

4.4.1 Ensure that the environment in which the test is to be carried out is free from extraneous particles which could cause interference. This involves wearing suitable garments and gloves and using a suitable clean-air work station, for example a laminar-flow cabinet meeting the requirements of Class 8 of ISO 14644-1 (Class 100,000 of US Federal Standard 209E), as well as suitably decontaminated tools and sample-handling equipment.

4.4.2 Carry out a blank test as follows:

- Place 100 ml of particle-free water into a conical flask.
- Shake for 20 s.
- Examine the water in the particle counter within the time interval starting 15 min and ending 30 min after shaking, and record the particle count in each of the four particle-size classes.
- Repeat the entire operation twice.

In no case shall more than 100 particles larger than 2 µm per 5 ml be found.

If this requirement is not met, investigate the possible causes of failure, rectify, and repeat the blank test until satisfactory results are obtained.

Carry out a blank test both before and after each series of tests. Only when satisfactory values are obtained both before and after the series of tests can the results of the tests be considered valid.

4.5 Test

- Place a number of complete elastomeric parts with a total surface area of approximately 100 cm² into a conical flask.
- Add 100 ml of particle-free water.
- Shake for 20 s.
- Examine the water in the particle counter within the time interval starting 15 min and ending 30 min after shaking, and record the particle count in each of the four particle-size classes.
- Repeat the entire operation twice.

4.6 Test report

For each test, report the following:

- a) the total surface area of the elastomeric parts tested, and the number of complete parts tested;
- b) the three individual counts in each of the four particle-size classes;
- c) the average count in each of the four classes, calculated from the individual values, per 10 cm² of surface, rounded to the nearest whole number;
- d) the total particle count in each of the four classes;
- e) the particle counts in each of the four classes for the blank tests performed.

Bibliography

- [1] ISO 14644-1, *Cleanrooms and associated controlled environments — Part 1: Classification of air cleanliness*

© ISO 2015

11.040.20

ICS 11.040.20

Price based on 6 pages