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**Snap-on bottles for metering pumps —  
Part 2:  
Moulded glass**

*Flacons encliquetables pour pompes doseuses —  
Partie 2: Verre moulé*

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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 liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

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](http://www.iso.org/directives)).

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. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see [www.iso.org/patents](http://www.iso.org/patents)).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT), see [www.iso.org/iso/foreword.html](http://www.iso.org/iso/foreword.html).

This document was prepared by Technical Committee ISO/TC 76, *Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use*.

A list of all parts in the ISO 24166 series can be found on the ISO website.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at [www.iso.org/members.html](http://www.iso.org/members.html).

## Introduction

This document is part of a series of International Standards for containers made from different materials used in combination with metering pumps for medicinal applications.

Glass containers are mainly used for that purpose. Plastic containers can be used as an alternative.

This document can be used for the development of standardized filling and assembling equipment.

Based on the dimensions of the containers, appropriate components, such as metering pumps and other closure systems can also be standardized. As such this document provides important inputs for developing entire packaging systems for medicinal applications.

Primary packaging materials are an integral part of medicinal products. Thus, depending on the jurisdiction, the principles of the current Good Manufacturing Practices (cGMP) can apply to the manufacturing of these components (e.g. ISO 15378).

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# Snap-on bottles for metering pumps —

## Part 2: Moulded glass

### 1 Scope

This document specifies the shape, dimensions, fill capacities and performance requirements of moulded glass bottles for metering pumps. It also specifies the material for the manufacturing of such containers as well as the secondary packaging.

This document also provides requirements for packaging of the moulded glass bottles and addresses nonsterile, ready to sterilize or sterile as three possible options.

This document is applicable to colourless or amber glass containers moulded from borosilicate or soda-lime-silica glass and intended to be used in the packaging, storage or transportation of products intended for medicinal use.

### 2 Normative references

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements 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 719, *Glass — Hydrolytic resistance of glass grains at 98 °C — Method of test and classification*

ISO 720, *Glass — Hydrolytic resistance of glass grains at 121 °C — Method of test and classification*

ISO 4802-1, *Glassware — Hydrolytic resistance of the interior surfaces of glass containers — Part 1: Determination by titration method and classification*

ISO 4802-2, *Glassware — Hydrolytic resistance of the interior surfaces of glass containers — Part 2: Determination by flame spectrometry and classification*

ISO 11607-1, *Packaging for terminally sterilized medical devices — Part 1: Requirements for materials, sterile barrier systems and packaging systems*

ISO 11607-2, *Packaging for terminally sterilized medical devices — Part 2: Validation requirements for forming, sealing and assembly processes*

### 3 Terms and definitions

For the purposes of this document, the following terms and definitions apply.

ISO and IEC maintain terminology databases for use in standardization at the following addresses:

— ISO Online browsing platform: available at <https://www.iso.org/obp>

— IEC Electropedia: available at <https://www.electropedia.org/>

#### 3.1

##### **base foil**

packaging component for protection of the pallet load from contamination

**3.2**  
**bottle**  
**vial**

container made of a suitable material, which is intended for packaging, storage and application of liquid medicinal products

EXAMPLE Tubular glass, moulded glass or plastic.

**3.3**  
**brimful capacity**

volume of water required to fill a container, placed on a flat, horizontal surface

[SOURCE: ISO 4802-1:2016, 3.3]

**3.4**  
**chemical indicator**

test system that reveals change in one or more pre-defined process variables based on a chemical change resulting from exposure to a process

[SOURCE: ISO 11139: 2018, 3.43, modified — "pre-specified" was replaced with "pre-defined", "or physical" was deleted.]

**3.5**  
**customer**

business entity that purchases bottles for metering pumps and conducts further processing or filling as appropriate

[SOURCE: ISO 21882:2019, 3.1 modified — "sterilized ready for filling vials" was replaced with "bottles for metering pumps".]

**3.6**  
**dip-tube**

conduit, part of the metering pump, which delivers the product from the bottle to the metering mechanism

**3.7**  
**foil bag**

gas permeable or non-permeable bag

Note 1 to entry: Foil bags can be used as *sterile barrier system* (3.16) or as *protective packaging* (3.13), depending on final usage.

**3.8**  
**label**

written, printed or graphic information appearing on the item itself, on the packaging of each item or on the packaging of multiple items

[SOURCE: IMDRF/GRRP WG/N52:2019, 3.17, modified – replaced "unit" and "devices" with "item".]

**3.9**  
**manufacturer**

business entity that manufactures or is otherwise responsible for the manufacturing of the bottles ready to be filled by the *customer* (3.5)

[SOURCE: ISO 21882:2019, 3.4. modified — "performs" was replaced with "manufactures", "vials" was replaced with "bottles".]

**3.10**  
**metering pump**

device actuated by the user to deliver a specific dose of liquid



**3.11****packaging system**

combination of a sterile barrier system and protective packaging

[SOURCE: ISO 11139:2018, 3.192]

**3.12****pallet**

construction for transportation and storage of goods

**3.13****protective packaging**

configuration of materials designed to prevent damage to the sterile barrier system and its contents from the time of their assembly until the point of use

[SOURCE: ISO 11139:2018, 3.219]

**3.14****sealing ring**

bottle feature on top of the flange to support tightness

**3.15****snap-on bottle**

container with press-fit for metering pumps

**3.16****sterile barrier system****SBS**

minimum package that minimizes the risk of ingress of microorganisms and allows aseptic presentation of the sterile contents at the point of use

[SOURCE: ISO 11139:2018, 3.272]

**3.17****transport cover**

outside packaging layer protecting against environmental hazards and keeping the package units in place

**4 Symbols**

$t$	bottom concavity
$d_1$	outer diameter body
$h_1$	total length
$h_2$	length cylindrical part of the body
$h_3$	inner depth
$r_1$	shoulder radius
$r_2$	bottom radius
M	moulded

## 5 Dimensions

### 5.1 General

The dimensions including tolerances are a function of manufacturing processes and materials.

To achieve the specified dimensions and tolerances, it is recommended that the offset of the mould halve and insert for forming the neck finish are horizontally max. 0,1 mm.

The dimensions and tolerances are established to achieve a neck that allows a snap-on functionality with the metering pump. This functionality can be challenged by a tightness test. An example of a snap-on process including positioning of the dip-tube as well as a typical tightness test is given in [Annex A](#) and [Annex B](#).

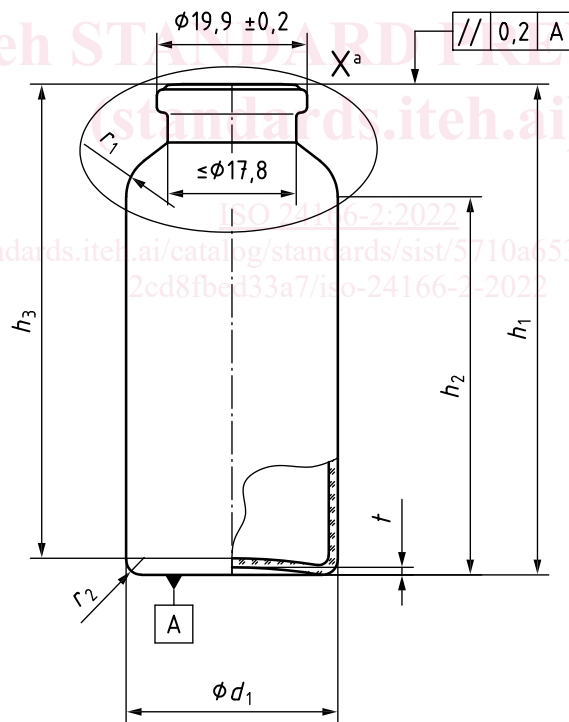
A test method for the inner depth as shown in [Table 1](#) and [Table 2](#) is described in [Annex C](#).

The bottles can be manufactured with or without a sealing ring.

### 5.2 Bottles without sealing ring

The dimensions of snap-on bottles made of moulded glass shall meet the requirements of [Figure 1](#) and [Figure 2](#), as appropriate, and [Table 1](#). The brimful capacity shall be as shown in [Table 1](#).

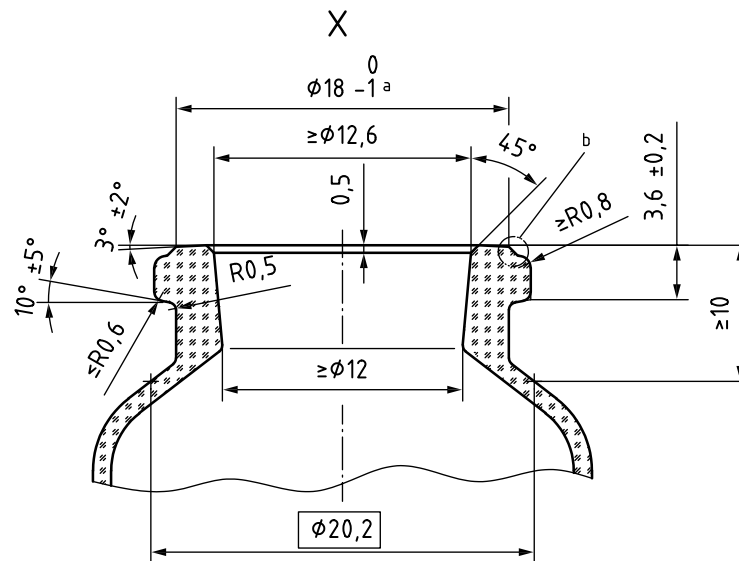
Dimensions in millimetres



<sup>a</sup> See [Figure 2](#).

**Figure 1 — Snap-on bottle without sealing ring**

Dimensions in millimetres



- a The mould split should be in between a diameter of 17 mm and 18 mm.
- b The geometry in this area of the bottle neck finish can be designed as an offset or as a chamfer (free design).

Figure 2 — Neck finish without sealing ring

Table 1 — Brimful capacity and dimensions of snap-on bottles made of moulded glass without sealing ring

Dimensions in millimetres

Size of the bottle	Brimful capacity ml	$d_1$		$h_1$		$h_2$	$h_3$		$r_1$	$r_2$	$t$
			Tol.		Tol.	min.		Tol.			
10M	15,0	25,4	$\pm 0,4$	53,3	$\pm 0,6$	35,3	49,0	$\pm 1,5$	10,0	2,0	1,0
20M	23,5	28,0	$\pm 0,6$	64,7	$\pm 0,5$	50,0	58,9		3,8	2,0	2,0
30M	35,0	31,8		75,2	$\pm 0,8$	57,0	69,4		7,8	2,5	1,5
50M	60,0	39,0	$\pm 0,5$	82,9	$\pm 0,8$	59,0	78,6	$\pm 2,0$	14,0	2,0	2,0
100M	106,0	45,5	$\pm 0,8$	102,9		76,5	97,2		14,0	3,0	2,0

### 5.3 Bottles with sealing ring

The dimensions of snap-on bottles made of moulded glass shall meet the requirements of [Figure 3](#) and [Figure 4](#), as appropriate, and [Table 2](#). The brimful capacity shall be as shown in [Table 2](#).