
**Aseptic processing of health care
products —**

**Part 1:
General requirements**

Traitement aseptique des produits de santé —

Partie 1: Exigences générales



Reference number
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Contents

Page

Foreword.....	v
Introduction	vi
1 Scope	1
2 Normative references	1
3 Terms and definitions.....	2
4 Quality system elements.....	7
4.1 General.....	7
4.2 Assignment of responsibilities	7
4.3 Calibration	7
5 Aseptic process definition	8
5.1 General.....	8
5.2 Risk management	8
6 Manufacturing environment	10
6.1 General.....	10
6.2 Manufacturing environment design	11
6.3 Layout	12
6.4 Material and personnel flow	14
6.5 HVAC system	15
6.6 Cleanroom qualification	17
6.7 Utility services and ancillary equipment	17
6.8 Environmental and personnel monitoring programmes	17
7 Equipment	21
7.1 Qualification	21
7.2 Maintenance of equipment	23
8 Personnel.....	23
8.1 General.....	23
8.2 Training for APA qualification	24
8.3 Gowning procedures	25
8.4 General employee health	26
9 Manufacture of the product	27
9.1 Attainment and maintenance of sterility	27
9.2 Duration of the manufacturing process	27
9.3 Aseptic manufacturing procedures	28
9.4 Cleaning and disinfection of facilities	28
9.5 Cleaning, disinfection and sterilization of equipment	30
10 Process simulation	31
10.1 General.....	31
10.2 Media selection and growth support	32
10.3 Simulation procedures	32
10.4 Incubation and inspection of media filled units	33
10.5 Initial performance qualification	33
10.6 Periodic performance requalification	34
10.7 Repeat of initial performance qualification	35
10.8 Documentation of process simulations	35
10.9 Disposition of filled product	36
11 Test for sterility	37

11.1	General	37
11.2	Investigation of positive units from tests for sterility	37
Annex A (informative)	Example of a flow chart	38
Annex B (informative)	Typical elements of an aseptic process definition	39
Annex C (informative)	Examples of specific risks	40
Annex D (informative)	Comparison of classification of cleanrooms	41
Annex E (informative)	Specification for water used in the process	42
Annex F (informative)	Aseptic processing area	44
Bibliography	45

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 13408-1 was prepared by Technical Committee ISO/TC 198, *Sterilization of health care products*.

This second edition cancels and replaces the first edition (ISO 13408-1:1998), which has been technically revised. Any normative and informative clauses on subjects which have meanwhile been addressed in Part 2 to Part 6 of ISO 13408 have been removed from this part.

ISO 13408 consists of the following parts, under the general title *Aseptic processing of health care products*:

- *Part 1: General requirements*
- *Part 2: Filtration*
- *Part 3: Lyophilization*
- *Part 4: Clean-in-place technologies*
- *Part 5: Sterilization in place*
- *Part 6: Isolator systems*

Introduction

Health care products that are labelled “sterile” are prepared using appropriate and validated methods under stringent control as part of a quality management system. For pharmaceuticals and medical devices there might be various requirements including compliance with ISO standards, GMP regulations and pharmacopoeial requirements.

Wherever possible, healthcare products intended to be sterile should be sterilized in their final sealed container (terminal sterilization). ISO/TC 198 has prepared standards for terminal sterilization of health care products by irradiation (series ISO 11137), by moist heat (ISO 17665-1), by dry heat (ISO 20857, in preparation) and by ethylene oxide (ISO 11135-1).

When a health care product is intended to be sterile and cannot be terminally sterilized, aseptic processing provides an alternative. Presterilization of product, product parts and/or components and all equipment coming into direct contact with the aseptically-processed product is required. Aseptic processing intends to maintain the sterility of the pre-sterilized components and products during assembling. The resulting product is required to be sterile in its final container. Aseptic processing can also be used to prevent contamination of biological product or biological systems (e.g. tissues, vaccines).

While terminal sterilization involves the control of a well-defined process of known lethality delivered to the product and a sterility assurance level (SAL) can be extrapolated from sterilization data, this is not applicable to aseptic processing.

Examples of applications in which aseptic processing are used include:

- aseptic handling and filling of solutions, suspensions, semisolids and powders;
- aseptic handling, transfer and packaging of solid products including solid medical devices;
- aseptic handling, transfer and packaging of combination products;
- aseptic handling of tissues or biological production systems.

Sterilization procedures which render components and/or parts sterile as a prerequisite for further aseptic processing can be treated as separate procedures. They have to be evaluated and validated separately and it is important that their risk of failure is minimal. The aseptic process definition encompasses all production steps following the sterilization of product and components until the final container or package is sealed. To keep the aseptic process definition clear and workable, this part of ISO 13408 is focused on the risks to the maintenance of sterility.

It is important to control all possible sources of contamination in order to maintain the sterility of each and every component. To achieve this, a risk-based aseptic process definition is established encompassing each product and applied in a comprehensive way considering product, package design, environment and manufacturing process designs. The product is processed in a controlled environment where microbial and particulate levels are maintained at defined minimal levels and where human intervention is minimized. Validated systems, adequately trained personnel, controlled environments and well-documented systematic processes are applied to assure a sterile finished product.

The aseptic process is divided into unit operations (e.g. sterilization of product or components including sterile filtration, assembly of components, handling and storage of sterilized product) and it is necessary that potential sources of contamination from materials, components, product, personnel, facility, equipment and utilities such as water systems be considered and minimized. Only if all risks of contamination have been recognised, wherever possible minimized, eliminated or controlled and finally have been evaluated as

acceptable, can the controls on the aseptic process be considered to be acceptable. Appropriate validation of the specified elements of the aseptic process is needed, of which process simulation studies are an essential.

This revision of ISO 13408-1:1998 is intended to adopt this International Standard to the actual state of technology in the field.